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(57) Abstract: The present invention provides to a antibody construct comprising a first human binding domain capable of binding to human CDH19 on the surface of a target cell and a second domain capable of binding to human CD3 on the surface of a T cell. Moreover, the invention relates to a nucleic acid sequence encoding the antibody construct, a vector comprising said nucleic acid seqquence and a host cell transformed or transfected with said vector. Furthermore, the invention relates a process for the production of the antibody construct of the invention, a medical use of said antibody construct and a kit comprising said antibody construct.

Antibody constructs for CDH19 and CD3

Related Applications

5 This application is related to a U.S. provisional application entitled "Antibodies targeting CDH19 for melanoma", filed on March 15, 2013, the same day as the present application is filed. This related application is incorporated in its entirety by reference.

Field of the Invention

The present invention relates to an antibody construct comprising a first human binding domain capable of binding to human CDH19 on the surface of a target cell and a second domain capable of binding to human CD3 on the surface of a T cell. Moreover, the invention provides a nucleic acid sequence encoding the antibody construct, a vector comprising said nucleic acid sequence and a host cell transformed or transfected with said vector.

Furthermore, the invention provides a process for the production of the antibody construct of the invention, a medical use of said antibody construct and a kit comprising said antibody construct.

Background of the Invention

Melanoma is a skin cancer that is caused by the oncogenic transformation of melanocytes, which are pigment producing skin cells. As of 2009, Melanoma had a prevalence of more than 870,000 cases in the US alone (US National Institutes of Health). Each year, over 75,000 new cases of melanoma are diagnosed in the US, and approximately 25% of patients have advanced disease at the time of diagnosis. Despite the fact that cases of primary melanoma can be cured by surgery if they are detected early enough, melanoma is the leading cause of death from skin disease in the US, responsible for about 10,000 deaths per year in the US. Once the disease has spread and became metastatic, the prognosis is poor, with a 5 year relative survival of 15%.

There are four basic types of melanomas. Three types are found in the top layers of the skin and the fourth one is invasive and has penetrated deeper into the skin and may have spread to other areas of the body.

Superficial spreading melanoma is the most common type of melanoma which accounts for about 70% of all cases. It grows along the top layer of the skin for a fairly long time before penetrating more deeply. It first appears as a flat or slightly raised discolored patch that has irregular borders and may be somewhat asymmetrical in form. The color varies, and you

may see areas of tan, brown, black, red, blue or white. This type of melanoma can occur in a previously benign mole and is found most often in young people.

Lentigo maligna is similar to the superficial spreading type, as it also remains close to the skin surface for quite a while, and usually appears as a flat or mildly elevated mottled tan, brown or dark brown discoloration. It is found most often in the elderly. When this cancer becomes invasive, it is referred to as lentigo maligna melanoma.

Acral lentiginous melanoma also spreads superficially before penetrating more deeply. It is quite different from the others, though, as it usually appears as a black or brown discoloration under the nails or on the soles of the feet or palms of the hands. This type of melanoma is sometimes found on dark-skinned people, and can often advance more quickly than superficial spreading melanoma and lentigo maligna.

Nodular melanoma is usually invasive at the time it is first diagnosed. The malignancy is recognized when it becomes a bump. It is usually black, but occasionally is blue, gray, white, brown, tan, red or skin tone. This is the most aggressive of the melanomas, and is found in 10 to 15 percent of cases.

Common treatments for metastatic melanoma include chemotherapy, targeted therapies for eligible patients (e.g. BRAF inhibitor treatment for patients with BRAF mutations) and immunotherapy. Metastatic melanoma is a tumor type where immunotherapy has been demonstrated to not only slow disease progression, but to lead to cures in late stage patients. Interleukin-2 was approved for the use in metastatic melanoma in 1998, and in 2011 an antibody targeting CTLA4, a member of a new generation of immune checkpoint inhibitors, gained approval by the FDA.

CDH19 is a type II cadherin transmembrane protein of unknown function. The human gene was cloned in 2000 based on its sequence similarity to CDH7 (Kools, P. et al. Genomics. 2000). Expressed Sequence Tags (ESTs) for CDH19 were isolated from melanocyte cDNA libraries, indicating that expression of CDH19 may be limited to cells of neural crest origin (Kools, P. et al. Genomics. 2000). In support of this notion, rat CDH19 was found to be expressed primarily in nerve ganglia and in Schwann cells during rat embryonic development (Takahashi, M. and Osumi, O. Devl Dynamics. 2005.).

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Diagnostic antibodies detecting CDH19 in Western Blot, immunohistochemitstry or flow cytometry are known in the art and commercially available. Those antibodies comprise polyand monoclonal antibodies generated in animal hosts.

5 Summary of the Invention

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The present invention provides an isolated multispecific antibody construct comprising a first human binding domain capable of binding to human CDH19 on the surface of a target cell and a second domain capable of binding to human CD3 on the surface of a T cell.

- In one embodiment the antibody construct of the invention the first binding domain comprises a VH region comprising CDR-H1, CDR-H2 and CDR-H3 and a VL region comprising CDR-L1, CDR-L2 and CDR-L3 selected from the group consisting of:
 - (a) CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927, CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 909, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927, CDR-H1 as depicted in SEQ ID NO: 909, CDR-L1 as depicted in SEQ ID NO: 927, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927, CDR-L2 as depicted in SEQ ID NO: 927, CDR-L2 as depicted in SEQ ID NO: 909, CDR-L1 as depicted in SEQ ID NO: 927, CDR-L2 as depicted in SEQ ID NO: 910, CDR-L2 as depicted in SEQ ID NO: 927, CDR-L2 as depicted in SEQ
- CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926, CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR
 - as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926, CDR-H1 as depicted in SEQ ID NO: 1126, CDR-H2 as depicted in SEQ ID NO: 1127, CDR-H3 as depicted in SEQ ID NO: 1128, CDR-L1 as depicted in SEQ ID NO: 1129, CDR-L2 as depicted in SEQ ID NO: 1130 and CDR-L3 as depicted in SEQ ID NO: 1131,

H3 as depicted in SEQ ID NO: 904, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2

35 CDR-H1 as depicted in SEQ ID NO: 1165, CDR-H2 as depicted in SEQ ID NO: 1166, CDR-H3 as depicted in SEQ ID NO: 1167, CDR-L1 as depicted in SEQ ID NO: 1168, CDR-L2 as depicted in SEQ ID NO: 1169 and CDR-L3 as depicted in SEQ ID

NO: 1170,

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CDR-H1 as depicted in SEQ ID NO: 1334, CDR-H2 as depicted in SEQ ID NO: 1335, CDR-H3 as depicted in SEQ ID NO: 1336, CDR-L1 as depicted in SEQ ID NO: 1337, CDR-L2 as depicted in SEQ ID NO: 1338 and CDR-L3 as depicted in SEQ ID NO: 1339,

CDR-H1 as depicted in SEQ ID NO: 1347, CDR-H2 as depicted in SEQ ID NO: 1348, CDR-H3 as depicted in SEQ ID NO: 1349, CDR-L1 as depicted in SEQ ID NO: 1350, CDR-L2 as depicted in SEQ ID NO: 1351 and CDR-L3 as depicted in SEQ ID NO: 1352,

10 CDR-H1 as depicted in SEQ ID NO: 1360 CDR-H2 as depicted in SEQ ID NO: 1361, CDR-H3 as depicted in SEQ ID NO: 1362, CDR-L1 as depicted in SEQ ID NO: 1363, CDR-L2 as depicted in SEQ ID NO: 1364 and CDR-L3 as depicted in SEQ ID NO: 1365,

CDR-H1 as depicted in SEQ ID NO: 1425 CDR-H2 as depicted in SEQ ID NO: 1426, CDR-H3 as depicted in SEQ ID NO: 1427, CDR-L1 as depicted in SEQ ID NO: 1428, CDR-L2 as depicted in SEQ ID NO: 1429 and CDR-L3 as depicted in SEQ ID NO: 1430.

CDR-H1 as depicted in SEQ ID NO: 1438 CDR-H2 as depicted in SEQ ID NO: 1439, CDR-H3 as depicted in SEQ ID NO: 1440, CDR-L1 as depicted in SEQ ID NO: 1441, CDR-L2 as depicted in SEQ ID NO: 1442 and CDR-L3 as depicted in SEQ ID NO: 1443, and

CDR-H1 as depicted in SEQ ID NO: 2167 CDR-H2 as depicted in SEQ ID NO: 2168, CDR-H3 as depicted in SEQ ID NO: 2169, CDR-L1 as depicted in SEQ ID NO: 2170, CDR-L2 as depicted in SEQ ID NO: 2171 and CDR-L3 as depicted in SEQ ID NO: 2172:

(b) CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 126, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294, CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 300, CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 306, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 312,

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CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336, CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294, CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 928, CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 929, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 942, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 943, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 937, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 919, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143,

CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, 5 CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 936, CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, 10 CDR-H3 as depicted in SEQ ID NO: 138, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 933. CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 917, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 934. 15 CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 930. CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, 20 CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 931, CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 932, CDR-H1 as depicted in SEQ ID NO: 1009, CDR-H2 as depicted in SEQ ID NO: 1010, 25 CDR-H3 as depicted in SEQ ID NO: 1011, CDR-L1 as depicted in SEQ ID NO: 1012, CDR-L2 as depicted in SEQ ID NO: 1013 and CDR-L3 as depicted in SEQ ID NO: 1014, CDR-H1 as depicted in SEQ ID NO: 1022, CDR-H2 as depicted in SEQ ID NO: 1023, CDR-H3 as depicted in SEQ ID NO: 1024, CDR-L1 as depicted in SEQ ID NO: 1025, 30 CDR-L2 as depicted in SEQ ID NO: 1026 and CDR-L3 as depicted in SEQ ID NO: 1027, CDR-H1 as depicted in SEQ ID NO: 1035, CDR-H2 as depicted in SEQ ID NO: 1036, CDR-H3 as depicted in SEQ ID NO: 1037, CDR-L1 as depicted in SEQ ID NO: 1038, CDR-L2 as depicted in SEQ ID NO: 1039 and CDR-L3 as depicted in SEQ ID 35 NO: 1040. CDR-H1 as depicted in SEQ ID NO: 1074, CDR-H2 as depicted in SEQ ID NO: 1075,

CDR-H3 as depicted in SEQ ID NO: 1076, CDR-L1 as depicted in SEQ ID NO: 1077,

CDR-L2 as depicted in SEQ ID NO: 1078 and CDR-L3 as depicted in SEQ ID NO: 1079,

CDR-H1 as depicted in SEQ ID NO: 1100, CDR-H2 as depicted in SEQ ID NO: 1101, CDR-H3 as depicted in SEQ ID NO: 1102, CDR-L1 as depicted in SEQ ID NO: 1103, CDR-L2 as depicted in SEQ ID NO: 1104 and CDR-L3 as depicted in SEQ ID NO: 1105,

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CDR-H1 as depicted in SEQ ID NO: 1113, CDR-H2 as depicted in SEQ ID NO: 1114, CDR-H3 as depicted in SEQ ID NO: 1115, CDR-L1 as depicted in SEQ ID NO: 1116, CDR-L2 as depicted in SEQ ID NO: 1117 and CDR-L3 as depicted in SEQ ID NO: 1118.

CDR-H1 as depicted in SEQ ID NO: 1243, CDR-H2 as depicted in SEQ ID NO: 1244, CDR-H3 as depicted in SEQ ID NO: 1245, CDR-L1 as depicted in SEQ ID NO: 1246, CDR-L2 as depicted in SEQ ID NO: 1247 and CDR-L3 as depicted in SEQ ID NO: 1248.

CDR-H1 as depicted in SEQ ID NO: 1256, CDR-H2 as depicted in SEQ ID NO: 1257, CDR-H3 as depicted in SEQ ID NO: 1258, CDR-L1 as depicted in SEQ ID NO: 1259, CDR-L2 as depicted in SEQ ID NO: 1260 and CDR-L3 as depicted in SEQ ID NO: 1261,

CDR-H1 as depicted in SEQ ID NO: 1269, CDR-H2 as depicted in SEQ ID NO: 1270, CDR-H3 as depicted in SEQ ID NO: 1271, CDR-L1 as depicted in SEQ ID NO: 1272, CDR-L2 as depicted in SEQ ID NO: 1273 and CDR-L3 as depicted in SEQ ID NO: 1274.

CDR-H1 as depicted in SEQ ID NO: 1282, CDR-H2 as depicted in SEQ ID NO: 1283, CDR-H3 as depicted in SEQ ID NO: 1284, CDR-L1 as depicted in SEQ ID NO: 1285, CDR-L2 as depicted in SEQ ID NO: 1286 and CDR-L3 as depicted in SEQ ID NO: 1287.

CDR-H1 as depicted in SEQ ID NO: 1295, CDR-H2 as depicted in SEQ ID NO: 1296, CDR-H3 as depicted in SEQ ID NO: 1297, CDR-L1 as depicted in SEQ ID NO: 1298, CDR-L2 as depicted in SEQ ID NO: 1299 and CDR-L3 as depicted in SEQ ID NO: 1300,

CDR-H1 as depicted in SEQ ID NO: 1647, CDR-H2 as depicted in SEQ ID NO: 1648, CDR-H3 as depicted in SEQ ID NO: 1649, CDR-L1 as depicted in SEQ ID NO: 1650, CDR-L2 as depicted in SEQ ID NO: 1651 and CDR-L3 as depicted in SEQ ID NO: 1652,

35 CDR-H1 as depicted in SEQ ID NO: 1660, CDR-H2 as depicted in SEQ ID NO: 1661, CDR-H3 as depicted in SEQ ID NO: 1662, CDR-L1 as depicted in SEQ ID NO: 1663, CDR-L2 as depicted in SEQ ID NO: 1664 and CDR-L3 as depicted in SEQ ID

NO: 1665,

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CDR-H1 as depicted in SEQ ID NO: 1894, CDR-H2 as depicted in SEQ ID NO: 1895, CDR-H3 as depicted in SEQ ID NO: 1896, CDR-L1 as depicted in SEQ ID NO: 1897, CDR-L2 as depicted in SEQ ID NO: 1898 and CDR-L3 as depicted in SEQ ID NO: 1899,

CDR-H1 as depicted in SEQ ID NO: 1907, CDR-H2 as depicted in SEQ ID NO: 1908, CDR-H3 as depicted in SEQ ID NO: 1909, CDR-L1 as depicted in SEQ ID NO: 1910, CDR-L2 as depicted in SEQ ID NO: 1911 and CDR-L3 as depicted in SEQ ID NO: 1912,

10 CDR-H1 as depicted in SEQ ID NO: 1933, CDR-H2 as depicted in SEQ ID NO: 1934, CDR-H3 as depicted in SEQ ID NO: 1935, CDR-L1 as depicted in SEQ ID NO: 1936, CDR-L2 as depicted in SEQ ID NO: 1937 and CDR-L3 as depicted in SEQ ID NO: 1938,

CDR-H1 as depicted in SEQ ID NO: 1946, CDR-H2 as depicted in SEQ ID NO: 1947, CDR-H3 as depicted in SEQ ID NO: 1948, CDR-L1 as depicted in SEQ ID NO: 1949, CDR-L2 as depicted in SEQ ID NO: 1950 and CDR-L3 as depicted in SEQ ID NO: 1951.

CDR-H1 as depicted in SEQ ID NO: 1959, CDR-H2 as depicted in SEQ ID NO: 1960, CDR-H3 as depicted in SEQ ID NO: 1961, CDR-L1 as depicted in SEQ ID NO: 1962, CDR-L2 as depicted in SEQ ID NO: 1963 and CDR-L3 as depicted in SEQ ID NO: 1964,

CDR-H1 as depicted in SEQ ID NO: 1972, CDR-H2 as depicted in SEQ ID NO: 1973, CDR-H3 as depicted in SEQ ID NO: 1974, CDR-L1 as depicted in SEQ ID NO: 1975, CDR-L2 as depicted in SEQ ID NO: 1976 and CDR-L3 as depicted in SEQ ID NO: 1977,

CDR-H1 as depicted in SEQ ID NO: 1985, CDR-H2 as depicted in SEQ ID NO: 1986, CDR-H3 as depicted in SEQ ID NO: 1987, CDR-L1 as depicted in SEQ ID NO: 1988, CDR-L2 as depicted in SEQ ID NO: 1989 and CDR-L3 as depicted in SEQ ID NO: 1990,

CDR-H1 as depicted in SEQ ID NO: 1998, CDR-H2 as depicted in SEQ ID NO: 1999, CDR-H3 as depicted in SEQ ID NO: 2000, CDR-L1 as depicted in SEQ ID NO: 2001, CDR-L2 as depicted in SEQ ID NO: 2002 and CDR-L3 as depicted in SEQ ID NO: 2003,

CDR-H1 as depicted in SEQ ID NO: 2011, CDR-H2 as depicted in SEQ ID NO: 2012, CDR-H3 as depicted in SEQ ID NO: 2013, CDR-L1 as depicted in SEQ ID NO: 2014, CDR-L2 as depicted in SEQ ID NO: 2015 and CDR-L3 as depicted in SEQ ID NO: 2016,

CDR-H1 as depicted in SEQ ID NO: 2024, CDR-H2 as depicted in SEQ ID NO: 2025, CDR-H3 as depicted in SEQ ID NO: 2026, CDR-L1 as depicted in SEQ ID NO: 2027, CDR-L2 as depicted in SEQ ID NO: 2028 and CDR-L3 as depicted in SEQ ID NO: 2029,

CDR-H1 as depicted in SEQ ID NO: 2037, CDR-H2 as depicted in SEQ ID NO: 2038, CDR-H3 as depicted in SEQ ID NO: 2039, CDR-L1 as depicted in SEQ ID NO: 2040, CDR-L2 as depicted in SEQ ID NO: 2041 and CDR-L3 as depicted in SEQ ID NO: 2042, and

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CDR-H1 as depicted in SEQ ID NO: 2050, CDR-H2 as depicted in SEQ ID NO: 2051, CDR-H3 as depicted in SEQ ID NO: 2052, CDR-L1 as depicted in SEQ ID NO: 2053, CDR-L2 as depicted in SEQ ID NO: 2054 and CDR-L3 as depicted in SEQ ID NO: 2055;

CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-(c) H3 as depicted in SEQ ID NO: 96, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 15 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 102, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119, 20 CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 156, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 324, 25 CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 912, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 913, CDR-L1 as depicted in SEQ ID NO: 268, 30 CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 910, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-35 H3 as depicted in SEQ ID NO: 911, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119,

CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 914, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 920, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 997, CDR-H3 as depicted in SEQ ID NO: 998, CDR-L1 as depicted in SEQ ID NO: 999, CDR-L2 as depicted in SEQ ID NO: 999, CDR-L2 as depicted in SEQ ID NO: 1000 and CDR-L3 as depicted in SEQ ID NO: 909, CDR-L2 as depicted in SEQ ID NO: 1000 and CDR-L3 as depicted in SEQ ID NO: 1001,

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CDR-H1 as depicted in SEQ ID NO: 1048, CDR-H2 as depicted in SEQ ID NO: 1049, CDR-H3 as depicted in SEQ ID NO: 1050, CDR-L1 as depicted in SEQ ID NO: 1051, CDR-L2 as depicted in SEQ ID NO: 1052 and CDR-L3 as depicted in SEQ ID NO: 1053.

CDR-H1 as depicted in SEQ ID NO: 1087, CDR-H2 as depicted in SEQ ID NO: 1088, CDR-H3 as depicted in SEQ ID NO: 1089, CDR-L1 as depicted in SEQ ID NO: 1090, CDR-L2 as depicted in SEQ ID NO: 1091 and CDR-L3 as depicted in SEQ ID NO: 1092.

CDR-H1 as depicted in SEQ ID NO: 1608, CDR-H2 as depicted in SEQ ID NO: 1609, CDR-H3 as depicted in SEQ ID NO: 1610, CDR-L1 as depicted in SEQ ID NO: 1611, CDR-L2 as depicted in SEQ ID NO: 1612 and CDR-L3 as depicted in SEQ ID NO: 1613,

CDR-H1 as depicted in SEQ ID NO: 1621, CDR-H2 as depicted in SEQ ID NO: 1622, CDR-H3 as depicted in SEQ ID NO: 1623, CDR-L1 as depicted in SEQ ID NO: 1624, CDR-L2 as depicted in SEQ ID NO: 1625 and CDR-L3 as depicted in SEQ ID NO: 1626,

CDR-H1 as depicted in SEQ ID NO: 1634, CDR-H2 as depicted in SEQ ID NO: 1635, CDR-H3 as depicted in SEQ ID NO: 1636, CDR-L1 as depicted in SEQ ID NO: 1637, CDR-L2 as depicted in SEQ ID NO: 1638 and CDR-L3 as depicted in SEQ ID NO: 1639,

CDR-H1 as depicted in SEQ ID NO: 1673, CDR-H2 as depicted in SEQ ID NO: 1674, CDR-H3 as depicted in SEQ ID NO: 1675, CDR-L1 as depicted in SEQ ID NO: 1676, CDR-L2 as depicted in SEQ ID NO: 1677 and CDR-L3 as depicted in SEQ ID NO: 1678,

CDR-H1 as depicted in SEQ ID NO: 1686, CDR-H2 as depicted in SEQ ID NO: 1687,

CDR-H3 as depicted in SEQ ID NO: 1688, CDR-L1 as depicted in SEQ ID NO: 1689, CDR-L2 as depicted in SEQ ID NO: 1690 and CDR-L3 as depicted in SEQ ID NO: 1691,

CDR-H1 as depicted in SEQ ID NO: 1699, CDR-H2 as depicted in SEQ ID NO: 1700, CDR-H3 as depicted in SEQ ID NO: 1701, CDR-L1 as depicted in SEQ ID NO: 1702, CDR-L2 as depicted in SEQ ID NO: 1703 and CDR-L3 as depicted in SEQ ID NO: 1704,

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CDR-H1 as depicted in SEQ ID NO: 1712, CDR-H2 as depicted in SEQ ID NO: 1713, CDR-H3 as depicted in SEQ ID NO: 1714, CDR-L1 as depicted in SEQ ID NO: 1715, CDR-L2 as depicted in SEQ ID NO: 1716 and CDR-L3 as depicted in SEQ ID NO: 1717.

CDR-H1 as depicted in SEQ ID NO: 1725, CDR-H2 as depicted in SEQ ID NO: 1726, CDR-H3 as depicted in SEQ ID NO: 1727, CDR-L1 as depicted in SEQ ID NO: 1728, CDR-L2 as depicted in SEQ ID NO: 1729 and CDR-L3 as depicted in SEQ ID NO: 1730,

CDR-H1 as depicted in SEQ ID NO: 1738, CDR-H2 as depicted in SEQ ID NO: 1739, CDR-H3 as depicted in SEQ ID NO: 1740, CDR-L1 as depicted in SEQ ID NO: 1741, CDR-L2 as depicted in SEQ ID NO: 1742 and CDR-L3 as depicted in SEQ ID NO: 1743.

CDR-H1 as depicted in SEQ ID NO: 1751, CDR-H2 as depicted in SEQ ID NO: 1752, CDR-H3 as depicted in SEQ ID NO: 1753, CDR-L1 as depicted in SEQ ID NO: 1754, CDR-L2 as depicted in SEQ ID NO: 1755 and CDR-L3 as depicted in SEQ ID NO: 1756,

CDR-H1 as depicted in SEQ ID NO: 1764, CDR-H2 as depicted in SEQ ID NO: 1765, CDR-H3 as depicted in SEQ ID NO: 1766, CDR-L1 as depicted in SEQ ID NO: 1767, CDR-L2 as depicted in SEQ ID NO: 1768 and CDR-L3 as depicted in SEQ ID NO: 1769, and

CDR-H1 as depicted in SEQ ID NO: 1920, CDR-H2 as depicted in SEQ ID NO: 1921, CDR-H3 as depicted in SEQ ID NO: 1922, CDR-L1 as depicted in SEQ ID NO: 1923, CDR-L2 as depicted in SEQ ID NO: 1924 and CDR-L3 as depicted in SEQ ID NO: 1925;

(d) CDR-H1 as depicted in SEQ ID NO: 4, CDR-H2 as depicted in SEQ ID NO: 5, CDR-H3 as depicted in SEQ ID NO: 6, CDR-L1 as depicted in SEQ ID NO: 172, CDR-L2 as depicted in SEQ ID NO: 173 and CDR-L3 as depicted in SEQ ID NO: 174, CDR-H1 as depicted in SEQ ID NO: 10, CDR-H2 as depicted in SEQ ID NO: 11, CDR-H3 as depicted in SEQ ID NO: 12, CDR-L1 as depicted in SEQ ID NO: 178, CDR-L2 as depicted in SEQ ID NO: 179 and CDR-L3 as depicted in SEQ ID NO: 180,

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CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 196, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 198, CDR-H1 as depicted in SEQ ID NO: 34, CDR-H2 as depicted in SEQ ID NO: 35, CDR-H3 as depicted in SEQ ID NO: 36, CDR-L1 as depicted in SEQ ID NO: 202, CDR-L2 as depicted in SEQ ID NO: 203 and CDR-L3 as depicted in SEQ ID NO: 204, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 214, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 59, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228, CDR-H1 as depicted in SEQ ID NO: 64, CDR-H2 as depicted in SEQ ID NO: 65, CDR-H3 as depicted in SEQ ID NO: 66, CDR-L1 as depicted in SEQ ID NO: 232, CDR-L2 as depicted in SEQ ID NO: 233 and CDR-L3 as depicted in SEQ ID NO: 234, CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 71, CDR-H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 328, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 902, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 903, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 925, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907, CDR-H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907,

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CDR-H3 as depicted in SEQ ID NO: 908, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 901, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 906, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228. CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330. CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 921, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 940. CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 941, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 196, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 901, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 970, CDR-H2 as depicted in SEQ ID NO: 971, CDR-H3 as depicted in SEQ ID NO: 972, CDR-L1 as depicted in SEQ ID NO: 973, CDR-L2 as depicted in SEQ ID NO: 974 and CDR-L3 as depicted in SEQ ID NO: 975, CDR-H1 as depicted in SEQ ID NO: 1061, CDR-H2 as depicted in SEQ ID NO: 1062, CDR-H3 as depicted in SEQ ID NO: 1063, CDR-L1 as depicted in SEQ ID NO: 1064,

CDR-L2 as depicted in SEQ ID NO: 1065 and CDR-L3 as depicted in SEQ ID NO: 1066,

CDR-H1 as depicted in SEQ ID NO: 1139, CDR-H2 as depicted in SEQ ID NO: 1140, CDR-H3 as depicted in SEQ ID NO: 1141, CDR-L1 as depicted in SEQ ID NO: 1142, CDR-L2 as depicted in SEQ ID NO: 1143 and CDR-L3 as depicted in SEQ ID NO: 1144,

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CDR-H1 as depicted in SEQ ID NO: 1152, CDR-H2 as depicted in SEQ ID NO: 1153, CDR-H3 as depicted in SEQ ID NO: 1154, CDR-L1 as depicted in SEQ ID NO: 1155, CDR-L2 as depicted in SEQ ID NO: 1156 and CDR-L3 as depicted in SEQ ID NO: 1157.

CDR-H1 as depicted in SEQ ID NO: 1178, CDR-H2 as depicted in SEQ ID NO: 1179, CDR-H3 as depicted in SEQ ID NO: 1180, CDR-L1 as depicted in SEQ ID NO: 1181, CDR-L2 as depicted in SEQ ID NO: 1182 and CDR-L3 as depicted in SEQ ID NO: 1183.

CDR-H1 as depicted in SEQ ID NO: 1191, CDR-H2 as depicted in SEQ ID NO: 1192, CDR-H3 as depicted in SEQ ID NO: 1193, CDR-L1 as depicted in SEQ ID NO: 1194, CDR-L2 as depicted in SEQ ID NO: 1195 and CDR-L3 as depicted in SEQ ID NO: 1196,

CDR-H1 as depicted in SEQ ID NO: 1204, CDR-H2 as depicted in SEQ ID NO: 1205, CDR-H3 as depicted in SEQ ID NO: 1206, CDR-L1 as depicted in SEQ ID NO: 1207, CDR-L2 as depicted in SEQ ID NO: 1208 and CDR-L3 as depicted in SEQ ID NO: 1209,

CDR-H1 as depicted in SEQ ID NO: 1217, CDR-H2 as depicted in SEQ ID NO: 1218, CDR-H3 as depicted in SEQ ID NO: 1219, CDR-L1 as depicted in SEQ ID NO: 1220, CDR-L2 as depicted in SEQ ID NO: 1221 and CDR-L3 as depicted in SEQ ID NO: 1222.

CDR-H1 as depicted in SEQ ID NO: 1230, CDR-H2 as depicted in SEQ ID NO: 1231, CDR-H3 as depicted in SEQ ID NO: 1232, CDR-L1 as depicted in SEQ ID NO: 1233, CDR-L2 as depicted in SEQ ID NO: 1234 and CDR-L3 as depicted in SEQ ID NO: 1235,

CDR-H1 as depicted in SEQ ID NO: 1308, CDR-H2 as depicted in SEQ ID NO: 1309, CDR-H3 as depicted in SEQ ID NO: 1310, CDR-L1 as depicted in SEQ ID NO: 1311, CDR-L2 as depicted in SEQ ID NO: 1312 and CDR-L3 as depicted in SEQ ID NO: 1313,

35 CDR-H1 as depicted in SEQ ID NO: 1321, CDR-H2 as depicted in SEQ ID NO: 1322, CDR-H3 as depicted in SEQ ID NO: 1323, CDR-L1 as depicted in SEQ ID NO: 1324, CDR-L2 as depicted in SEQ ID NO: 1325 and CDR-L3 as depicted in SEQ ID

NO: 1326,

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CDR-H1 as depicted in SEQ ID NO: 1373, CDR-H2 as depicted in SEQ ID NO: 1374, CDR-H3 as depicted in SEQ ID NO: 1375, CDR-L1 as depicted in SEQ ID NO: 1376, CDR-L2 as depicted in SEQ ID NO: 1377 and CDR-L3 as depicted in SEQ ID NO: 1378,

CDR-H1 as depicted in SEQ ID NO: 1386, CDR-H2 as depicted in SEQ ID NO: 1387, CDR-H3 as depicted in SEQ ID NO: 1388, CDR-L1 as depicted in SEQ ID NO: 1389, CDR-L2 as depicted in SEQ ID NO: 1390 and CDR-L3 as depicted in SEQ ID NO: 1391,

10 CDR-H1 as depicted in SEQ ID NO: 1399, CDR-H2 as depicted in SEQ ID NO: 1400, CDR-H3 as depicted in SEQ ID NO: 1401, CDR-L1 as depicted in SEQ ID NO: 1402, CDR-L2 as depicted in SEQ ID NO: 1403 and CDR-L3 as depicted in SEQ ID NO: 1404,

CDR-H1 as depicted in SEQ ID NO: 1412, CDR-H2 as depicted in SEQ ID NO: 1413, CDR-H3 as depicted in SEQ ID NO: 1414, CDR-L1 as depicted in SEQ ID NO: 1415, CDR-L2 as depicted in SEQ ID NO: 1416 and CDR-L3 as depicted in SEQ ID NO: 1417,

CDR-H1 as depicted in SEQ ID NO: 1777, CDR-H2 as depicted in SEQ ID NO: 1778, CDR-H3 as depicted in SEQ ID NO: 1779, CDR-L1 as depicted in SEQ ID NO: 1780, CDR-L2 as depicted in SEQ ID NO: 1781 and CDR-L3 as depicted in SEQ ID NO: 1782,

CDR-H1 as depicted in SEQ ID NO: 1790, CDR-H2 as depicted in SEQ ID NO: 1791, CDR-H3 as depicted in SEQ ID NO: 1792, CDR-L1 as depicted in SEQ ID NO: 1793, CDR-L2 as depicted in SEQ ID NO: 1794 and CDR-L3 as depicted in SEQ ID NO: 1795,

CDR-H1 as depicted in SEQ ID NO: 1803, CDR-H2 as depicted in SEQ ID NO: 1804, CDR-H3 as depicted in SEQ ID NO: 1805, CDR-L1 as depicted in SEQ ID NO: 1806, CDR-L2 as depicted in SEQ ID NO: 1807 and CDR-L3 as depicted in SEQ ID NO: 1808.

CDR-H1 as depicted in SEQ ID NO: 1816, CDR-H2 as depicted in SEQ ID NO: 1817, CDR-H3 as depicted in SEQ ID NO: 1818, CDR-L1 as depicted in SEQ ID NO: 1819, CDR-L2 as depicted in SEQ ID NO: 1820 and CDR-L3 as depicted in SEQ ID NO: 1821,

CDR-H1 as depicted in SEQ ID NO: 1829, CDR-H2 as depicted in SEQ ID NO: 1830, CDR-H3 as depicted in SEQ ID NO: 1831, CDR-L1 as depicted in SEQ ID NO: 1832, CDR-L2 as depicted in SEQ ID NO: 1833 and CDR-L3 as depicted in SEQ ID NO: 1834,

CDR-H1 as depicted in SEQ ID NO: 1842, CDR-H2 as depicted in SEQ ID NO: 1843, CDR-H3 as depicted in SEQ ID NO: 1844, CDR-L1 as depicted in SEQ ID NO: 1845, CDR-L2 as depicted in SEQ ID NO: 1846 and CDR-L3 as depicted in SEQ ID NO: 1847,

CDR-H1 as depicted in SEQ ID NO: 1855, CDR-H2 as depicted in SEQ ID NO: 1856, CDR-H3 as depicted in SEQ ID NO: 1857, CDR-L1 as depicted in SEQ ID NO: 1858, CDR-L2 as depicted in SEQ ID NO: 1859 and CDR-L3 as depicted in SEQ ID NO: 1860,

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CDR-H1 as depicted in SEQ ID NO: 1868, CDR-H2 as depicted in SEQ ID NO: 1869, CDR-H3 as depicted in SEQ ID NO: 1870, CDR-L1 as depicted in SEQ ID NO: 1871, CDR-L2 as depicted in SEQ ID NO: 1872 and CDR-L3 as depicted in SEQ ID NO: 1873,

CDR-H1 as depicted in SEQ ID NO: 1881, CDR-H2 as depicted in SEQ ID NO: 1882, CDR-H3 as depicted in SEQ ID NO: 1883, CDR-L1 as depicted in SEQ ID NO: 1884, CDR-L2 as depicted in SEQ ID NO: 1885 and CDR-L3 as depicted in SEQ ID NO: 1886.

CDR-H1 as depicted in SEQ ID NO: 2063, CDR-H2 as depicted in SEQ ID NO: 2064, CDR-H3 as depicted in SEQ ID NO: 2065, CDR-L1 as depicted in SEQ ID NO: 2066, CDR-L2 as depicted in SEQ ID NO: 2067 and CDR-L3 as depicted in SEQ ID NO: 2068.

CDR-H1 as depicted in SEQ ID NO: 2076, CDR-H2 as depicted in SEQ ID NO: 2077, CDR-H3 as depicted in SEQ ID NO: 2078, CDR-L1 as depicted in SEQ ID NO: 2079, CDR-L2 as depicted in SEQ ID NO: 2080 and CDR-L3 as depicted in SEQ ID NO: 2081,

CDR-H1 as depicted in SEQ ID NO: 2089, CDR-H2 as depicted in SEQ ID NO: 2090, CDR-H3 as depicted in SEQ ID NO: 2091, CDR-L1 as depicted in SEQ ID NO: 2092, CDR-L2 as depicted in SEQ ID NO: 2093 and CDR-L3 as depicted in SEQ ID NO: 2094,

CDR-H1 as depicted in SEQ ID NO: 2102, CDR-H2 as depicted in SEQ ID NO: 2103, CDR-H3 as depicted in SEQ ID NO: 2104, CDR-L1 as depicted in SEQ ID NO: 2105, CDR-L2 as depicted in SEQ ID NO: 2106 and CDR-L3 as depicted in SEQ ID NO: 2107,

CDR-H1 as depicted in SEQ ID NO: 2115, CDR-H2 as depicted in SEQ ID NO: 2116, CDR-H3 as depicted in SEQ ID NO: 2117, CDR-L1 as depicted in SEQ ID NO: 2118, CDR-L2 as depicted in SEQ ID NO: 2119 and CDR-L3 as depicted in SEQ ID NO: 2120,

CDR-H1 as depicted in SEQ ID NO: 2128, CDR-H2 as depicted in SEQ ID NO: 2129,

CDR-H3 as depicted in SEQ ID NO: 2130, CDR-L1 as depicted in SEQ ID NO: 2131, CDR-L2 as depicted in SEQ ID NO: 2132 and CDR-L3 as depicted in SEQ ID NO: 2133,

CDR-H1 as depicted in SEQ ID NO: 2141, CDR-H2 as depicted in SEQ ID NO: 2142, CDR-H3 as depicted in SEQ ID NO: 2143, CDR-L1 as depicted in SEQ ID NO: 2144, CDR-L2 as depicted in SEQ ID NO: 2145 and CDR-L3 as depicted in SEQ ID NO: 2146,

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CDR-H1 as depicted in SEQ ID NO: 2154, CDR-H2 as depicted in SEQ ID NO: 2155, CDR-H3 as depicted in SEQ ID NO: 2156, CDR-L1 as depicted in SEQ ID NO: 2157, CDR-L2 as depicted in SEQ ID NO: 2158 and CDR-L3 as depicted in SEQ ID NO: 2159.

CDR-H1 as depicted in SEQ ID NO: 2180, CDR-H2 as depicted in SEQ ID NO: 2181, CDR-H3 as depicted in SEQ ID NO: 2182, CDR-L1 as depicted in SEQ ID NO: 2183, CDR-L2 as depicted in SEQ ID NO: 2184 and CDR-L3 as depicted in SEQ ID NO: 2185,

CDR-H1 as depicted in SEQ ID NO: 2193, CDR-H2 as depicted in SEQ ID NO: 2194, CDR-H3 as depicted in SEQ ID NO: 2195, CDR-L1 as depicted in SEQ ID NO: 2196, CDR-L2 as depicted in SEQ ID NO: 2197 and CDR-L3 as depicted in SEQ ID NO: 2198, and

CDR-H1 as depicted in SEQ ID NO: 2206, CDR-H2 as depicted in SEQ ID NO: 2207, CDR-H3 as depicted in SEQ ID NO: 2208, CDR-L1 as depicted in SEQ ID NO: 2209, CDR-L2 as depicted in SEQ ID NO: 2210 and CDR-L3 as depicted in SEQ ID NO: 2211; and

CDR-H1 as depicted in SEQ ID NO: 76, CDR-H2 as depicted in SEQ ID NO: 77, CDR-(e) H3 as depicted in SEQ ID NO: 78, CDR-L1 as depicted in SEQ ID NO: 244, CDR-L2 25 as depicted in SEQ ID NO: 245 and CDR-L3 as depicted in SEQ ID NO: 246. CDR-H1 as depicted in SEQ ID NO: 88, CDR-H2 as depicted in SEQ ID NO: 89, CDR-H3 as depicted in SEQ ID NO: 90, CDR-L1 as depicted in SEQ ID NO: 256, CDR-L2 as depicted in SEQ ID NO: 257 and CDR-L3 as depicted in SEQ ID NO: 258, 30 CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274, CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276, CDR-H1 as depicted in SEQ ID NO: 112, CDR-H2 as depicted in SEQ ID NO: 113, CDR-H3 as depicted in SEQ ID NO: 114, CDR-L1 as depicted in SEQ ID NO: 280, 35 CDR-L2 as depicted in SEQ ID NO: 281 and CDR-L3 as depicted in SEQ ID NO: 282, CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274,

CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276, CDR-H1 as depicted in SEQ ID NO: 983, CDR-H2 as depicted in SEQ ID NO: 984, CDR-H3 as depicted in SEQ ID NO: 985, CDR-L1 as depicted in SEQ ID NO: 986, CDR-L2 as depicted in SEQ ID NO: 987 and CDR-L3 as depicted in SEQ ID NO: 988,

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CDR-H1 as depicted in SEQ ID NO: 1582, CDR-H2 as depicted in SEQ ID NO: 1583, CDR-H3 as depicted in SEQ ID NO: 1584, CDR-L1 as depicted in SEQ ID NO: 1585, CDR-L2 as depicted in SEQ ID NO: 1586 and CDR-L3 as depicted in SEQ ID NO: 1587, and

10 CDR-H1 as depicted in SEQ ID NO: 1595, CDR-H2 as depicted in SEQ ID NO: 1596, CDR-H3 as depicted in SEQ ID NO: 1597, CDR-L1 as depicted in SEQ ID NO: 1598, CDR-L2 as depicted in SEQ ID NO: 1599 and CDR-L3 as depicted in SEQ ID NO: 1600.

- In a further embodiment of the antibody construct of the invention the first binding domain comprises a VH region selected from the group consisting of VH regions
 - (a) as depicted in SEQ ID NO: 362, SEQ ID NO: 364, SEQ ID NO: 485, SEQ ID NO: 486, SEQ ID NO: 487, SEQ ID NO: 492, SEQ ID NO: 493, SEQ ID NO: 494, SEQ ID NO: 495, SEQ ID NO: 1133, SEQ ID NO: 1172, SEQ ID NO: 1341, SEQ ID NO: 1354, SEQ ID NO: 1367, SEQ ID NO: 1432, SEQ ID NO: 1445 and SEQ ID NO: 2174;
- as depicted in SEQ ID NO: 342, SEQ ID NO: 366, SEQ ID NO: 370, SEQ ID NO: 344, SEQ ID NO: 372, SEQ ID NO: 368, SEQ ID NO: 496, SEQ ID NO: 497, SEQ ID NO: 498, SEQ ID NO: 499, SEQ ID NO: 500, SEQ ID NO: 508, SEQ ID NO: 509, SEQ ID NO: 510, SEQ ID NO: 511, SEQ ID NO: 512, SEQ ID NO: 519, SEQ ID 25 NO: 520, SEQ ID NO: 521, SEQ ID NO: 522, SEQ ID NO: 523, SEQ ID NO: 524, SEQ ID NO: 525, SEQ ID NO: 526, SEQ ID NO: 527, SEQ ID NO: 528, SEQ ID NO: 529, SEQ ID NO: 530, SEQ ID NO: 531, SEQ ID NO: 532, SEQ ID NO: 533, SEQ ID NO: 534, SEQ ID NO: 535, SEQ ID NO: 536, SEQ ID NO: 537, SEQ ID NO: 538, SEQ ID NO: 1016, SEQ ID NO: 1029, SEQ ID NO: 1042, SEQ ID NO: 1081, 30 SEQ ID NO: 1107, SEQ ID NO: 1120, SEQ ID NO: 1250, SEQ ID NO: 1263, SEQ ID NO: 1276, SEQ ID NO: 1289, SEQ ID NO: 1302, SEQ ID NO: 1654, SEQ ID NO: 1667, SEQ ID NO: 1901, SEQ ID NO: 1914, SEQ ID NO: 1940, SEQ ID NO: 1953, SEQ ID NO: 1966, SEQ ID NO: 1979, SEQ ID NO: 1992, SEQ ID NO: 2005, SEQ ID NO: 2018, SEQ ID NO: 2031, SEQ ID NO: 2044, and SEQ ID 35 NO: 2057:
 - (c) as depicted in SEQ ID NO: 338, SEQ ID NO: 354, SEQ ID NO: 378, SEQ ID NO: 356, SEQ ID NO: 476, SEQ ID NO: 477, SEQ ID NO: 478, SEQ ID NO: 479, SEQ ID

NO: 480, SEQ ID NO: 481, SEQ ID NO: 482, SEQ ID NO: 483, SEQ ID NO: 484, SEQ ID NO: 501, SEQ ID NO: 502, SEQ ID NO: 503, SEQ ID NO: 504, SEQ ID NO: 505, SEQ ID NO: 506, SEQ ID NO: 517, SEQ ID NO: 518, SEQ ID NO: 1003, SEQ ID NO: 1055, SEQ ID NO: 1094, SEQ ID NO: 1615, SEQ ID NO: 1628, SEQ ID NO: 1641, SEQ ID NO: 1680, SEQ ID NO: 1693, SEQ ID NO: 1706, SEQ ID NO: 1719, SEQ ID NO: 1732, SEQ ID NO: 1745, SEQ ID NO: 1758, SEQ ID NO: 1771, and SEQ ID NO: 1927;

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- as depicted in SEQ ID NO: 352, SEQ ID NO: 360, SEQ ID NO: 388, SEQ ID NO: 386, (d) SEQ ID NO: 340, SEQ ID NO: 346, SEQ ID NO: 374, SEQ ID NO: 348, SEQ ID NO: 390, SEQ ID NO: 463, SEQ ID NO: 464, SEQ ID NO: 465, SEQ ID NO: 466, SEQ ID NO: 467, SEQ ID NO: 468, SEQ ID NO: 469, SEQ ID NO: 470, SEQ ID NO: 471, SEQ ID NO: 472, SEQ ID NO: 473, SEQ ID NO: 474, SEQ ID NO: 475, SEQ ID NO: 488, SEQ ID NO: 489, SEQ ID NO: 490, SEQ ID NO: 491, SEQ ID NO: 513, SEQ ID NO: 514, SEQ ID NO: 515, SEQ ID NO: 516, SEQ ID NO: 540, SEQ ID NO: 541, SEQ ID NO: 542, SEQ ID NO: 543, SEQ ID NO: 977, SEQ ID NO: 1068, SEQ ID NO: 1146, SEQ ID NO: 1159, SEQ ID NO: 1185, SEQ ID NO: 1198, SEQ ID NO: 1211, SEQ ID NO: 1224, SEQ ID NO: 1237, SEQ ID NO: 1315, SEQ ID NO: 1328, SEQ ID NO: 1380, SEQ ID NO: 1393, SEQ ID NO: 1406, SEQ ID NO: 1419, SEQ ID NO: 1469, SEQ ID NO: 1478, SEQ ID NO: 1485. SEQ ID NO: 1494. SEQ ID NO: 1501. SEQ ID NO: 1508. SEQ ID NO: 1519, SEQ ID NO: 1526, SEQ ID NO: 1533, SEQ ID NO: 1542, SEQ ID NO: 1549, SEQ ID NO: 1558, SEQ ID NO: 1565, SEQ ID NO: 1784, SEQ ID NO: 1797, SEQ ID NO: 1810, SEQ ID NO: 1823, SEQ ID NO: 1836, SEQ ID NO: 1849, SEQ ID NO: 1862, SEQ ID NO: 1875, SEQ ID NO: 1888, SEQ ID NO: 2070, SEQ ID NO: 2083, SEQ ID NO: 2096, SEQ ID NO: 2109, SEQ ID NO: 2122, SEQ ID NO: 2135, SEQ ID NO: 2148, SEQ ID NO: 2161, SEQ ID NO: 2187, SEQ ID NO: 2200, and SEQ ID NO: 2213; and
 - (e) as depicted in SEQ ID NO: 376, SEQ ID NO: 392, SEQ ID NO: 358, SEQ ID NO: 350, SEQ ID NO: 507, SEQ ID NO: 990, SEQ ID NO: 1589, and SEQ ID NO: 1602.

In another embodiment of the antibody construct of the invention the first binding domain comprises a VL region selected from the group consisting of VL regions

(a) as depicted in SEQ ID NO: 418, SEQ ID NO: 420, SEQ ID NO: 580, SEQ ID NO: 581, SEQ ID NO: 582, SEQ ID NO: 587, SEQ ID NO: 588, SEQ ID NO: 589, SEQ ID NO: 590, SEQ ID NO: 1135, SEQ ID NO: 1174, SEQ ID NO: 1343, SEQ ID NO: 1356, SEQ ID NO: 1369, SEQ ID NO: 1434, SEQ ID NO: 1447 and SEQ ID NO: 2176;

as depicted in SEQ ID NO: 398, SEQ ID NO: 422, SEQ ID NO: 426, SEQ ID NO: 400, (b) SEQ ID NO: 428, SEQ ID NO: 424, SEQ ID NO: 591, SEQ ID NO: 592, SEQ ID NO: 593, SEQ ID NO: 594, SEQ ID NO: 595, SEQ ID NO: 603, SEQ ID NO: 604, SEQ ID NO: 605, SEQ ID NO: 606, SEQ ID NO: 607, SEQ ID NO: 614, SEQ ID NO: 615, SEQ ID NO: 616, SEQ ID NO: 617, SEQ ID NO: 618, SEQ ID NO: 619, SEQ ID NO: 620, SEQ ID NO: 621, SEQ ID NO: 622, SEQ ID NO: 623, SEQ ID NO: 624, SEQ ID NO: 625, SEQ ID NO: 626, SEQ ID NO: 627, SEQ ID NO: 628, SEQ ID NO: 629, SEQ ID NO: 630, SEQ ID NO: 631, SEQ ID NO: 632, SEQ ID NO: 633, SEQ ID NO: 1018, SEQ ID NO: 1031, SEQ ID NO: 1044, SEQ ID NO: 1083, SEQ ID NO: 1109, SEQ ID NO: 1122, SEQ ID NO: 1252, SEQ ID NO: 1265, SEQ ID NO: 1278, SEQ ID NO: 1291, SEQ ID NO: 1304, SEQ ID NO: 1656, SEQ ID NO: 1669, SEQ ID NO: 1903, SEQ ID NO: 1916, SEQ ID NO: 1942, SEQ ID NO: 1955, SEQ ID NO: 1968, SEQ ID NO: 1981, SEQ ID NO: 1994, SEQ ID NO: 2007, SEQ ID NO: 2020, SEQ ID NO: 2033, SEQ ID NO: 2046, and SEQ ID NO: 2059;

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- (c) as depicted in SEQ ID NO: 394, SEQ ID NO: 410, SEQ ID NO: 434, SEQ ID NO: 412, SEQ ID NO: 571, SEQ ID NO: 572, SEQ ID NO: 573, SEQ ID NO: 574, SEQ ID NO: 575, SEQ ID NO: 576, SEQ ID NO: 577, SEQ ID NO: 578, SEQ ID NO: 579, SEQ ID NO: 596, SEQ ID NO: 597, SEQ ID NO: 598, SEQ ID NO: 599, SEQ ID NO: 600, SEQ ID NO: 601, SEQ ID NO: 612, SEQ ID NO: 613, SEQ ID NO: 1005, SEQ ID NO: 1057, SEQ ID NO: 1096, SEQ ID NO: 1617, SEQ ID NO: 1630, SEQ ID NO: 1643, SEQ ID NO: 1682, SEQ ID NO: 1695, SEQ ID NO: 1708, SEQ ID NO: 1721, SEQ ID NO: 1734, SEQ ID NO: 1747, SEQ ID NO: 1760, SEQ ID NO: 1773, and SEQ ID NO: 1929;
- as depicted in SEQ ID NO: 408, SEQ ID NO: 416, SEQ ID NO: 444, SEQ ID NO: 442, 25 (d) SEQ ID NO: 396, SEQ ID NO: 402, SEQ ID NO: 430, SEQ ID NO: 404, SEQ ID NO: 446, SEQ ID NO: 558, SEQ ID NO: 559, SEQ ID NO: 560, SEQ ID NO: 561, SEQ ID NO: 562, SEQ ID NO: 563, SEQ ID NO: 564, SEQ ID NO: 565, SEQ ID NO: 566, SEQ ID NO: 567, SEQ ID NO: 568, SEQ ID NO: 569, SEQ ID NO: 570, 30 SEQ ID NO: 583, SEQ ID NO: 584, SEQ ID NO: 585, SEQ ID NO: 586, SEQ ID NO: 608, SEQ ID NO: 609, SEQ ID NO: 610, SEQ ID NO: 611, SEQ ID NO: 635, SEQ ID NO: 636, SEQ ID NO: 637, SEQ ID NO: 638, SEQ ID NO: 979, SEQ ID NO: 1070, SEQ ID NO: 1148, SEQ ID NO: 1161, SEQ ID NO: 1187, SEQ ID NO: 1200, SEQ ID NO: 1213, SEQ ID NO: 1226, SEQ ID NO: 1239, SEQ ID 35 NO: 1317, SEQ ID NO: 1330, SEQ ID NO: 1382, SEQ ID NO: 1395, SEQ ID NO: 1408, SEQ ID NO: 1421, SEQ ID NO: 1471, SEQ ID NO: 1480, SEQ ID NO: 1487, SEQ ID NO: 1496, SEQ ID NO: 1503, SEQ ID NO: 1510, SEQ ID

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NO: 1521, SEQ ID NO: 1528, SEQ ID NO: 1535, SEQ ID NO: 1544, SEQ ID NO: 1551, SEQ ID NO: 1560, SEQ ID NO: 1567, SEQ ID NO: 1786, SEQ ID NO: 1799, SEQ ID NO: 1812, SEQ ID NO: 1825, SEQ ID NO: 1838, SEQ ID NO: 1851, SEQ ID NO: 1864, SEQ ID NO: 1877, SEQ ID NO: 1890, SEQ ID NO: 2072, SEQ ID NO: 2085, SEQ ID NO: 2098, SEQ ID NO: 2111, SEQ ID NO: 2124, SEQ ID NO: 2137, SEQ ID NO: 2150, SEQ ID NO: 2163, SEQ ID NO: 2189, SEQ ID NO: 2202, and SEQ ID NO: 2215; and
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(e) as depicted in SEQ ID NO: 432, SEQ ID NO: 448, SEQ ID NO: 414, SEQ ID NO: 406, SEQ ID NO: 602, SEQ ID NO: 992, SEQ ID NO: 1591, and SEQ ID NO: 1604.

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The invention further provides an embodiment of the antibody construct of the invention, wherein the first binding domain comprises a VH region and a VL region selected from the group consisting of:

- (1) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 362+418, SEQ ID 15 NOs: 364+420. SEQ ID NOs: 485+580, SEQ ID NOs: 486+581, SEQ ID NOs: 487+582, SEQ ID NOs: 492+587, SEQ ID NOs: 493+588, SEQ ID NOs: 494+589, SEQ ID NOs: 495+590, SEQ ID NOs: 1133+1135, SEQ ID NOs: 1172+1174, SEQ ID NOs: 1341+1343, SEQ ID NOs: 1354+1356, SEQ ID NOs: 1367+1369, SEQ ID NOs: 1432+1434, SEQ ID NOs: 1445+1447, and SEQ ID 20 NOs: 2174+2176;
- pairs of a VH region and a VL region as depicted in SEQ ID NOs: 342+398, SEQ ID (2) NOs: 366+422, SEQ ID NOs: 370+426, SEQ ID NOs: 344+400, SEQ ID NOs: 372+428, SEQ ID SEQ ID NOs: 368+424, NOs: 496+591, SEQ ID NOs: 497+592, SEQ ID NOs: 498+593, SEQ ID NOs: 499+594, SEQ ID 25 NOs: 500+595, SEQ ID NOs: 508+603, SEQ ID NOs: 509+604, SEQ ID SEQ ID NOs: 510+605. SEQ ID NOs: 511+606, NOs: 512+607, SEQ ID NOs: 519+614, SEQ ID SEQ ID SEQ ID NOs: 520+615, NOs: 521+616, NOs: 522+617, SEQ ID NOs: 523+618, SEQ ID NOs: 524+619, SEQ ID NOs: 525+620, SEQ ID NOs: 526+621, SEQ ID NOs: 527+622, SEQ ID 30 NOs: 528+623, SEQ ID NOs: 529+624, SEQ ID NOs: 530+625, SEQ ID NOs: 531+626, SEQ ID SEQ ID SEQ ID NOs: 532+627, NOs: 533+628, NOs: 534+629, SEQ ID NOs: 535+630, SEQ ID NOs: 536+631, SEQ ID NOs: 537+632, SEQ ID NOs: 538+633, SEQ ID NOs: 1016+1018, SEQ ID NOs: 1029+1031, SEQ ID NOs: 1042+1044, SEQ ID NOs: 1081+1083, SEQ ID 35 NOs: 1107+1109, SEQ ID NOs: 1120+1122, SEQ ID NOs: 1250+1252, SEQ ID NOs: 1263+1265, SEQ ID NOs: 1276+1278, SEQ ID NOs: 1289+1291, SEQ ID NOs: 1302+1304, SEQ ID NOs: 1654+1656, SEQ ID NOs: 1667+1669, SEQ ID

NOs: 1901+1903, SEQ ID NOs: 1914+1916, SEQ ID NOs: 1940+1942, SEQ ID NOs: 1953+1955, SEQ ID NOs: 1966+1968, SEQ ID NOs: 1979+1981, SEQ ID NOs: 1992+1994, SEQ ID NOs: 2005+2007, SEQ ID NOs: 2018+2020, SEQ ID NOs: 2031+2033, SEQ ID NOs: 2044+2046, and SEQ ID NOs: 2057+2059; 5 pairs of a VH region and a VL region as depicted in SEQ ID NOs: 338+394, SEQ ID (3) NOs: 354+410, SEQ ID NOs: 378+434, SEQ ID NOs: 356+412, SEQ ID NOs: 476+571, SEQ ID NOs: 477+572, SEQ ID NOs: 478+573, SEQ ID NOs: 479+574, SEQ ID NOs: 480+575, SEQ ID NOs: 481+576, SEQ ID NOs: 482+577, SEQ ID NOs: 483+578, SEQ ID NOs: 484+579, SEQ ID 10 NOs: 501+596, SEQ ID NOs: 502+597, SEQ ID NOs: 503+598, SEQ ID NOs: 504+599, SEQ ID NOs: 505+600, SEQ ID NOs: 506+601. SEQ ID NOs: 517+612, SEQ ID NOs: 518+613, SEQ ID NOs: 1003+1005, SEQ ID NOs: 1055+1057, SEQ ID NOs: 1094+1096, SEQ ID NOs: 1615+1617, SEQ ID NOs: 1628+1630, SEQ ID NOs: 1641+1643, SEQ ID NOs: 1680+1682, SEQ ID NOs: 1693+1695, SEQ ID NOs: 1706+1708, SEQ ID NOs: 1719+1721, SEQ ID 15 NOs: 1732+1734, SEQ ID NOs: 1745+1747, SEQ ID NOs: 1758+1760, SEQ ID NOs: 1771+1773, and SEQ ID NOs: 1927+1929; pairs of a VH region and a VL region as depicted in SEQ ID NOs: 352+408, SEQ ID (4) NOs: 360+416. SEQ ID NOs: 388+444, SEQ ID NOs: 386+442, SEQ ID 20 NOs: 340+396. SEQ ID NOs: 346+402, SEQ ID NOs: 374+430, SEQ ID NOs: 348+404, SEQ ID SEQ ID NOs: 390+446, NOs: 463+558, SEQ ID NOs: 464+559. SEQ ID NOs: 465+560, SEQ ID NOs: 466+561, SEQ ID SEQ ID NOs: 467+562, SEQ ID NOs: 468+563, NOs: 469+564, SEQ ID NOs: 470+565, SEQ ID NOs: 471+566, SEQ ID NOs: 472+567, SEQ ID 25 NOs: 473+568, SEQ ID NOs: 474+569, SEQ ID NOs: 475+570, SEQ ID SEQ ID NOs: 488+583. SEQ ID NOs: 489+584, NOs: 490+585. SEQ ID NOs: 491+586, SEQ ID SEQ ID NOs: 513+608, NOs: 514+609, SEQ ID NOs: 515+610, SEQ ID NOs: 516+611, SEQ ID NOs: 540+635, SEQ ID NOs: 541+636, SEQ ID NOs: 542+637. SEQ ID NOs: 543+638, SEQ ID 30 NOs: 977+979, SEQ ID NOs: 1068+1070, SEQ ID NOs: 1146+1148, SEQ ID NOs: 1159+1161, SEQ ID NOs: 1185+1187, SEQ ID NOs: 1198+1200, SEQ ID NOs: 1211+1213, SEQ ID NOs: 1224+1226, SEQ ID NOs: 1237+1239, SEQ ID NOs: 1315+1317, SEQ ID NOs: 1328+1330, SEQ ID NOs: 1380,+1382 SEQ ID NOs: 1393+1395, SEQ ID NOs: 1406+1408, SEQ ID NOs: 1419+1421, SEQ ID 35 NOs: 1469+1471, SEQ ID NOs: 1478+1480, SEQ ID NOs: 1485+1487, SEQ ID NOs: 1494+1496, SEQ ID NOs: 1501+1503, SEQ ID NOs: 1508+1510, SEQ ID NOs: 1519+1521, SEQ ID NOs: 1526+1528, SEQ ID NOs: 1533+1535, SEQ ID

NOs: 1542+1544, SEQ ID NOs: 1549+1551, SEQ ID NOs: 1558+1560, SEQ ID NOs: 1565+1567, SEQ ID NOs: 1784+1786, SEQ ID NOs: 1797+1799, SEQ ID NOs: 1810+1812, SEQ ID NOs: 1823+1825, SEQ ID NOs: 1836+1838, SEQ ID NOs: 1849+1851, SEQ ID NOs: 1862+1864, SEQ ID NOs: 1875+1877, SEQ ID NOs: 1888+1890, SEQ ID NOs: 2070+2072, SEQ ID NOs: 2083+2085, SEQ ID NOs: 2096+2098, SEQ ID NOs: 2109+2111, SEQ ID NOs: 2122+2124, SEQ ID NOs: 2135+2137, SEQ ID NOs: 2148+2150, SEQ ID NOs: 2161+2163, SEQ ID NOs: 2187+2189, SEQ ID NOs: 2200+2202, and SEQ ID NOs: 2213+2215; and

(5) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 376+432, SEQ ID NOs: 392+448, SEQ ID NOs: 358+414, SEQ ID NOs: 350+406, SEQ ID NOs: 507+602, SEQ ID NOs: 990+992, SEQ ID NOs: 1589+1591, and SEQ ID NOs: 1602+1604.

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In a further embodiment of the invention the antibody construct is in a format selected from the group consisting of (scFv)₂, (single domain mAb)₂, scFv-single domain mAb, diabodies and oligomers thereof.

In a preferred embodiment the first binding domain comprises an amino acid sequence selected from the group consisting of

- 20 (a) as depicted in SEQ ID NO: 117, SEQ ID NO: 1137, SEQ ID NO: 1176, SEQ ID NO: 1345, SEQ ID NO: 1358, SEQ ID NO: 1371, SEQ ID NO: 1436, SEQ ID NO: 1449 and SEQ ID NO: 2178;
- (b) as depicted in SEQ ID NO: 1020, SEQ ID NO: 1033, SEQ ID NO: 1046, SEQ ID NO: 1085, SEQ ID NO: 1111, SEQ ID NO: 1124, SEQ ID NO: 1254, SEQ ID NO: 1267, SEQ ID NO: 1280, SEQ ID NO: 1293, SEQ ID NO: 1306, SEQ ID NO: 1658, SEQ ID NO: 1671, SEQ ID NO: 1905, SEQ ID NO: 1918, SEQ ID NO: 1944, SEQ ID NO: 1957, SEQ ID NO: 1970, SEQ ID NO: 1983, SEQ ID NO: 1996, SEQ ID NO: 2009, SEQ ID NO: 2022, SEQ ID NO: 2035, SEQ ID NO: 2048, and SEQ ID NO: 2061;
- 30 (c) as depicted in SEQ ID NO: 1007, SEQ ID NO: 1059, SEQ ID NO: 1098, SEQ ID NO: 1619, SEQ ID NO: 1632, SEQ ID NO: 1645, SEQ ID NO: 1684, SEQ ID NO: 1697, SEQ ID NO: 1710, SEQ ID NO: 1723, SEQ ID NO: 1736, SEQ ID NO: 1749, SEQ ID NO: 1762, SEQ ID NO: 1775, and SEQ ID NO: 1931;
- (d) as depicted in SEQ ID NO: 981, SEQ ID NO: 1072, SEQ ID NO: 1150, SEQ ID NO: 1163, SEQ ID NO: 1189, SEQ ID NO: 1202, SEQ ID NO: 1215, SEQ ID NO: 1228, SEQ ID NO: 1241, SEQ ID NO: 1319, SEQ ID NO: 1332, SEQ ID NO: 1384, SEQ ID NO: 1397, SEQ ID NO: 1410, SEQ ID NO: 1423, SEQ ID

NO: 1473, SEQ ID NO: 1482, SEQ ID NO: 1489, SEQ ID NO: 1498, SEQ ID NO: 1505, SEQ ID NO: 1512, SEQ ID NO: 1523, SEQ ID NO: 1530, SEQ ID NO: 1537, SEQ ID NO: 1546, SEQ ID NO: 1553, SEQ ID NO: 1562, SEQ ID NO: 1569, SEQ ID NO: 1788, SEQ ID NO: 1801, SEQ ID NO: 1814, SEQ ID NO: 1827, SEQ ID NO: 1840, SEQ ID NO: 1853, SEQ ID NO: 1866, SEQ ID NO: 1879, SEQ ID NO: 1892, SEQ ID NO: 2074, SEQ ID NO: 2087, SEQ ID NO: 2100, SEQ ID NO: 2113, SEQ ID NO: 2126, SEQ ID NO: 2139, SEQ ID NO: 2152, SEQ ID NO: 2165, SEQ ID NO: 2191, SEQ ID NO: 2204, and SEQ ID NO: 2217; and

10 (e) as depicted in SEQ ID NO: 994, SEQ ID NO: 1593, and SEQ ID NO: 1606.

In another embodiment of the antibody construct of the invention the second binding domain is capable of binding to human and Callithrix jacchus, Saguinus Oedipus or Saimiri sciureus CD3 epsilon.

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In a preferred embodiment the antibody construct of the invention has an amino acid sequence selected from the group consisting of

- (a) as depicted in SEQ ID NO: 1138, SEQ ID NO: 1177, SEQ ID NO: 1346, SEQ ID NO: 1359, SEQ ID NO: 1372, SEQ ID NO: 1437, SEQ ID NO: 14501450 and SEQ ID NO: 2179;
- (b) as depicted in SEQ ID NO: 1021, SEQ ID NO: 1034, SEQ ID NO: 1047, SEQ ID NO: 1086, SEQ ID NO: 1112, SEQ ID NO: 1125, SEQ ID NO: 1255, SEQ ID NO: 1268, SEQ ID NO: 1281, SEQ ID NO: 1294, SEQ ID NO: 1307, SEQ ID NO: 1659, SEQ ID NO: 1672, SEQ ID NO: 1906, SEQ ID NO: 1919, SEQ ID NO: 1945, SEQ ID NO: 1958, SEQ ID NO: 1971, SEQ ID NO: 1984, SEQ ID NO: 1997, SEQ ID NO: 2010, SEQ ID NO: 2023, SEQ ID NO: 2036, SEQ ID NO: 2049, and SEQ ID NO: 2062;
- (c) as depicted in SEQ ID NO: 1008, SEQ ID NO: 1060, SEQ ID NO: 1099, SEQ ID NO: 1620, SEQ ID NO: 1633, SEQ ID NO: 1646, SEQ ID NO: 1685, SEQ ID NO: 1698, SEQ ID NO: 1711, SEQ ID NO: 1724, SEQ ID NO: 1737, SEQ ID NO: 1750, SEQ ID NO: 1763, SEQ ID NO: 1776, and SEQ ID NO: 1932;
- (d) as depicted in SEQ ID NO: 982, SEQ ID NO: 1073, SEQ ID NO: 1151, SEQ ID NO: 1164, SEQ ID NO: 1190, SEQ ID NO: 1203, SEQ ID NO: 1216, SEQ ID NO: 1229, SEQ ID NO: 1242, SEQ ID NO: 1320, SEQ ID NO: 1333, SEQ ID NO: 1385, SEQ ID NO: 1398, SEQ ID NO: 1411, SEQ ID NO: 1424, SEQ ID NO: 1474, SEQ ID NO: 1475, SEQ ID NO: 1476, SEQ ID NO: 1483, SEQ ID NO: 1490, SEQ ID NO: 1491, SEQ ID NO: 1492, SEQ ID NO: 1499, SEQ ID

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NO: 1506, SEQ ID NO: 1513, SEQ ID NO: 1514, SEQ ID NO: 1515, SEQ ID NO: 1516, SEQ ID NO: 1517, SEQ ID NO: 1524, SEQ ID NO: 1531, SEQ ID NO: 1538, SEQ ID NO: 1539, SEQ ID NO: 1540, SEQ ID NO: 1547, SEQ ID NO: 1554, SEQ ID NO: 1555, SEQ ID NO: 1556, SEQ ID NO: 1563, SEQ ID NO: 1570, SEQ ID NO: 1571, SEQ ID NO: 1572, SEQ ID NO: 1573, SEQ ID NO: 1574, SEQ ID NO: 1575, SEQ ID NO: 1576, SEQ ID NO: 1577, SEQ ID NO: 1578, SEQ ID NO: 1579, SEQ ID NO: 1580, SEQ ID NO: 1581, SEQ ID NO: 1789, SEQ ID NO: 1802, SEQ ID NO: 1815, SEQ ID NO: 1828, SEQ ID NO: 1841, SEQ ID NO: 1854, SEQ ID NO: 1867, SEQ ID NO: 1880, SEQ ID NO: 1893, SEQ ID NO: 2075, SEQ ID NO: 2088, SEQ ID NO: 2101, SEQ ID NO: 2114, SEQ ID NO: 2127, SEQ ID NO: 2140, SEQ ID NO: 2153, SEQ ID NO: 2166, SEQ ID NO: 2192, SEQ ID NO: 2205, and SEQ ID NO: 2218 to 2228; and as depicted in SEQ ID NO: 995, SEQ ID NO: 1594, and SEQ ID NO: 1607.
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The invention further provides a nucleic acid sequence encoding an antibody construct of the invention.

Furthermore, the invention provides a vector comprising a nucleic acid sequence of the invention. Moreover, the invention provides a host cell transformed or transfected with the nucleic acid sequence of the invention.

In a further embodiment the invention provides a process for the production of a antibody construct of the invention, said process comprising culturing a host cell of the invention under conditions allowing the expression of the antibody construct of the invention and recovering the produced antibody construct from the culture.

Moreover, the invention provides a pharmaceutical composition comprising an antibody construct of the invention or produced according to the process of the invention

30 In one embodiment the invention provides the antibody construct of the invention or produced according to the process of the invention for use in the prevention, treatment or amelioration of a melanoma disease or metastatic melanoma disease.

The invention also provides a method for the treatment or amelioration of a melanoma disease or metastatic melanoma disease, comprising the step of administering to a subject in need thereof the antibody construct of the invention or produced according to the process of the invention.

In a preferred embodiment method of use of the invention the melanoma disease or metastatic melanoma disease is selected from the group consisting of superficial spreading melanoma, lentigo maligna, lentigo maligna melanoma, acral lentiginous melanoma and nodular melanoma.

In a further embodiment, the invention provides a kit comprising an antibody construct of the invention, or produced according to the process of the invention, a vector of the invention, and/or a host cell of the invention.

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Brief description of the drawings

Figure 1:

FIG. 1 depicts cell viability data of Colo-699 cells that have been treated with fully human anti-CDH19 antibodies and a high concentration of a goat anti-human Fc monovalent Fab conjugated with DM1 (DM1-Fab) at a drug-antibody ratio (DAR) (~1.3).

Figure 2:

FIG. 2 depicts the average cell viability data from a CHL-1 assay plotted against the average cell viability data from the Colo-699 assay.

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Figure 3:

FIG. 3 shows the relative expression of CDH19 mRNA in metastatic and primary melanoma samples.

25 **Figure 4**:

FIG. 4 shows the expression of CDH19 protein in human tumor samples by IHC.

Figure 5:

FIG. 5 shows the results of the analysis of tumor cell lines by flow cytometry and IHC to identify model systems with CDH19 expression similar to human tumors based on the number of CDH19 receptors present on the cell surface.

Figure 6:

FACS analysis of CDH19/CD3 bispecific antibodies on indicated cell lines:

1) untransfected L1.2. 2) L1.2 cells stably transfected with human CDH19, 3) melanoma cell line CHL-1, 4) melanoma cell line A2058, 5) human CD3 positive human T cell line HBP-

ALL, 6) macaque T cell line 4119 LnPx. Negative controls [1) to 6)]: detection antibodies without prior CDH19/CD3 bispecific antibody.

Figure 7:

5 Cytotoxic activity of CDH19/CD3 bispecific antibodies as measured in a 48-hour FACS-based cytotoxicity assay. Effector cells: unstimulated human PBMC. Target cells: as indicated. Effector to target cell (E:T)-ratio: 10:1.

Figure 8:

Tumor growth *in vivo* inhibition of Colo699 cells by administration of CDH19 BiTE 2G6. The bispecific antibody construct inhibits growth of tumors at 0.5 mg/kg dose.

Figure 9:

Tumor growth *in vivo* inhibition of CHL-1 cells by administration of CDH19 BiTE 2G6. The bispecific antibody construct inhibits growth of tumors at 0.5 mg/kg dose.

Figure 10:

Cytotoxic activity of CDH19/CD3 bispecific antibodies as measured in a 48-hour imaging-based cytotoxicity assay. Effector cells: unstimulated human T cells. Target cells: as indicated. Effector to target cell (E:T)-ratio: 10:1.

Figure 11:

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Chromatogram IMAC capture and elution CH19 2G6 302 x I2C SA21

Typical IMAC elution profile obtained during purification of an CDH19 BiTE antibody. The red
line indicates absorption at 254 nm, the blue line indicates absorption at 280 nm. Brown line indicates conductivity. 1 – Capture. 2 – Pre-Elution 50 mM Imidazole. 3. BiTE Elution 500 mM Imidazole

Figure 12:

30 Chromatogram Protein A capture and elution CH19 2G6 302 x F12Q

Typical Protein_A elution profile obtained during purification of an CDH19 BiTE antibody. The red line indicates absorption at 254 nm, the blue line indicates absorption at 280 nm. Brown line indicates conductivity. Green line indicates the applied gradient percentage. 1 – Capture. 2 – BiTE Elution

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Figure 13:

SEC elution profile of CDH19 BiTE antibody 2G6 302 x I2C SA21

Typical SEC elution profile obtained during purification of an CDH19 BiTE antibody. Protein peaks corresponding to the monomeric and dimeric BiTE antibody isoforms are indicated. LMW = low molecular weight. The red line indicates absorption at 254 nm, the blue line indicates absorption at 280 nm. Brown line indicates conductivity. 1 – non BiTE aggregates in SEC exclusion volume. 2. BiTE dimer. 3. BiTE monomer. 4. Low molecular weight contaminants and salts

Figure 14:

Reduced SDS PAGE analytics of CDH19 BiTE Monomer CH19 2G6 302 x I2C SA21 (left) and molecular weight marker Novex Sharp Protein Standard (Life Technologies).

Figure 15:

HP-SEC chromatogram showing the elution of CDH19 BiTE CH19 2G6 302 x I2C SA21 after seven day of storage at 37°C. Pink line indicationg optical absorption at 210 nm wavelength. Brown line indicating conductivity.

1 BiTE Dimer. 2. BiTE Monomer

Figure 16:

20 HP-SEC chromatogram showing the elution of CDH19 BiTE CH19 2G6 302 x I2C SA21 after three freeze/thaw cycles. Pink line indicationg optical absorption at 210 nm wavelength. Brown line indicating conductivity. 1. BiTE Monomer

Figure 17:

25 CatlEX chromatogramm of elution of CDH19 BiTE CH19 2G6 302 x I2C SA21. Blue line indicating optical absorption at 280 nm. Red line indicating optical absorption at 254 nm.

Figure 18:

HIC elution profil of CDH19 BiTE CH19 2G6 302 x I2C SA21. Blue line indicating optical absorption at 280 nm. Red line indicating optical absorption at 254 nm. Brown line indicating conductivity.

Figure 19:

FACS analysis of CDH19/CD3 bispecific antibodies on indicated cell lines: 1) HEK293 cells stably transfected with human CDH19, 2) human CD3 positive human T cell line HBP-ALL; Negative controls [1) and 2)]: detection antibodies without prior CDH19/CD3 bispecific antibody cell culture supernatant.

Figure 20:

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Cytotoxic activity of CDH19/CD3 bispecific antibodies as measured in an 18-hour Chromium release-based cytotoxicity assay. Effector cells: stimulated human CD8+ T-cells. Target cells: HEK293 transfected with human CDH19. Effector to target cell (E:T)-ratio: 10:1.

Detailed Description of the Invention

Definitions:

It must be noted that as used herein, the singular forms "a", "an", and "the", include plural references unless the context clearly indicates otherwise. Thus, for example, reference to "a reagent" includes one or more of such different reagents and reference to "the method" includes reference to equivalent steps and methods known to those of ordinary skill in the art that could be modified or substituted for the methods described herein.

Unless otherwise indicated, the term "at least" preceding a series of elements is to be understood to refer to every element in the series. Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the present invention.

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The term "and/or" wherever used herein includes the meaning of "and", "or" and "all or any other combination of the elements connected by said term".

The term "about" or "approximately" as used herein means within ±20%, preferably within ±15%, more preferably within ±10%, and most preferably within ±5% of a given value or range.

Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integer or step. When used herein the term "comprising" can be substituted with the term "containing" or "including" or sometimes when used herein with the term "having".

When used herein "consisting of" excludes any element, step, or ingredient not specified in the claim element. When used herein, "consisting essentially of" does not exclude materials or steps that do not materially affect the basic and novel characteristics of the claim.

In each instance herein any of the terms "comprising", "consisting essentially of" and "consisting of" may be replaced with either of the other two terms.

The definition of the term "antibody" includes embodiments such as monoclonal, chimeric, single chain, humanized and human antibodies, as well as antibody fragments, like, inter alia, Fab fragments. Antibody fragments or derivatives further comprise F(ab')₂, Fv, scFv fragments or single domain antibodies such as domain antibodies or nanobodies, single variable domain antibodies or immunoglobulin single variable domain comprising merely one variable domain, which might be VHH, VH or VL, that specifically bind an antigen or epitope independently of other V regions or domains; see, for example, Harlow and Lane (1988) and (1999), loc. cit.; Kontermann and Dübel, Antibody Engineering, Springer, 2nd ed. 2010 and Little, Recombinant Antibodies for Immunotherapy, Cambridge University Press 2009. Such immunoglobulin single variable domain encompasses not only an isolated antibody single variable domain polypeptide, but also larger polypeptides that comprise one or more monomers of an antibody single variable domain polypeptide sequence.

In line with this definition all above described embodiments of the term antibody can be subsumed under the term "antibody construct". Said term also includes diabodies or Dual-Affinity Re-Targeting (DART) antibodies. Further envisaged are (bispecific) single chain diabodies, tandem diabodies (Tandab's), "minibodies" exemplified by a structure which is as follows: (VH-VL-CH3)₂, (scFv-CH3)₂ or (scFv-CH3-scFv)₂, "Fc DART" antibodies and "IgG DART" antibodies, and multibodies such as triabodies. Immunoglobulin single variable domains encompass not only an isolated antibody single variable domain polypeptide, but also larger polypeptides that comprise one or more monomers of an antibody single variable domain polypeptide sequence.

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Various procedures are known in the art and may be used for the production of such antibody constructs (antibodies and/or fragments). Thus, (antibody) derivatives can be produced by peptidomimetics. Further, techniques described for the production of single chain antibodies (see, inter alia, US Patent 4,946,778, Kontermann and Dübel (2010), loc. cit. and Little(2009), loc. cit.) can be adapted to produce single chain antibodies specific for elected polypeptide(s). Also, transgenic animals may be used to express humanized antibodies specific for polypeptides and fusion proteins of this invention. For the preparation of monoclonal antibodies, any technique, providing antibodies produced by continuous cell line cultures can be used. Examples for such techniques include the hybridoma technique (Köhler and Milstein Nature 256 (1975), 495-497), the trioma technique, the human B-cell

hybridoma technique (Kozbor, Immunology Today 4 (1983), 72) and the EBV-hybridoma technique to produce human monoclonal antibodies (Cole et al., Monoclonal Antibodies and Cancer Therapy, Alan R. Liss, Inc. (1985), 77-96). Surface plasmon resonance as employed in the BIAcore system can be used to increase the efficiency of phage antibodies which bind to an epitope of a target polypeptide, such as CD3 epsilon (Schier, Human Antibodies Hybridomas 7 (1996), 97-105; Malmborg, J. Immunol. Methods 183 (1995), 7-13). It is also envisaged in the context of this invention that the term "antibody" comprises antibody constructs, which may be expressed in a host as described herein below, e.g. antibody constructs which may be transfected and/or transduced via, inter alia, viruses or plasmid vectors.

Furthermore, the term "antibody" as employed in the invention also relates to derivatives or variants of the antibodies described herein which display the same specificity as the described antibodies.

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The terms "antigen-binding domain", "antigen-binding fragment" and "antibody binding region" when used herein refer to a part of an antibody molecule that comprises amino acids responsible for the specific binding between antibody and antigen. The part of the antigen that is specifically recognized and bound by the antibody is referred to as the "epitope" as described herein above. As mentioned above, an antigen-binding domain may typically comprise an antibody light chain variable region (VL) and an antibody heavy chain variable region (VH); however, it does not have to comprise both. Fd fragments, for example, have two VH regions and often retain some antigen-binding function of the intact antigen-binding domain. Examples of antigen-binding fragments of an antibody include (1) a Fab fragment, a monovalent fragment having the VL, VH, CL and CH1 domains; (2) a F(ab')2 fragment, a bivalent fragment having two Fab fragments linked by a disulfide bridge at the hinge region; (3) a Fd fragment having the two VH and CH1 domains; (4) a Fv fragment having the VL and VH domains of a single arm of an antibody, (5) a dAb fragment (Ward et al., (1989) Nature 341 :544-546), which has a VH domain; (6) an isolated complementarity determining region (CDR), and (7) a single chain Fv (scFv). Although the two domains of the Fv fragment, VL and VH are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the VL and VH regions pair to form monovalent molecules (known as single chain Fv (scFv); see e.g., Huston et al. (1988) Proc. Natl. Acad. Sci USA 85:5879-5883). These antibody fragments are obtained using conventional techniques known to those with skill in the art, and the fragments are evaluated for function in the same manner as are intact antibodies.

The term "monoclonal antibody" as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations and/or post- translation modifications (e.g., isomerizations, amidations) that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site. Furthermore, in contrast to conventional (polyclonal) antibody preparations which typically include different antibodies directed against different determinants (epitopes), each monoclonal antibody is directed against a single determinant on the antigen. In addition to their specificity, the monoclonal antibodies are advantageous in that they are synthesized by the hybridoma culture, uncontaminated by other immunoglobulins. The modifier "monoclonal" indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies to be used in accordance with the present invention may be made by the hybridoma method first described by Kohler et al., Nature, 256: 495 (1975), or may be made by recombinant DNA methods (see, e.g., U. S. Patent No. 4,816,567). The "monoclonal antibodies" may also be isolated from phage antibody libraries using the techniques described in Clackson et al., Nature, 352: 624-628 (1991) and Marks et al., J. Mol. Biol., 222: 581-597 (1991), for example.

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The term "human antibody" includes antibodies having variable and constant regions corresponding substantially to human germline immunoglobulin sequences known in the art, including, for example, those described by Kabat *et al.* (See Kabat *et al.* (1991) loc. cit.). The human antibodies of the invention may include amino acid residues not encoded by human germline immunoglobulin sequences (e.g., mutations introduced by random or site-specific mutagenesis *in vitro* or by somatic mutation *in vivo*), for example in the CDRs, and in particular, CDR3. The human antibody can have at least one, two, three, four, five, or more positions replaced with an amino acid residue that is not encoded by the human germline immunoglobulin sequence. It is emphasized that the definition of human antibodies as used herein also contemplates fully human antibodies, which include only non-artificially and/or genetically altered human sequences of antibodies as those can be derived by using technologies using systems such as the Xenomice.

Examples of "antibody variants" include humanized variants of non- human antibodies, "affinity matured" antibodies (see, e.g. Hawkins et al. J. Mol. Biol. 254, 889-896 (1992) and Lowman et al., Biochemistry 30, 10832- 10837 (1991)) and antibody mutants with altered

effector function (s) (see, e.g., US Patent 5, 648, 260, Kontermann and Dübel (2010), loc. cit. and Little(2009), loc. cit.).

As used herein, "in vitro generated antibody" refers to an antibody where all or part of the variable region (e.g., at least one CDR) is generated in a non-immune cell selection (e.g., an in vitro phage display, protein chip or any other method in which candidate sequences can be tested for their ability to bind to an antigen). This term thus preferably excludes sequences generated by genomic rearrangement in an immune cell.

The pairing of a VH and VL together forms a single antigen-binding site. The CH domain most proximal to VH is designated as CH1. Each L chain is linked to an H chain by one covalent disulfide bond, while the two H chains are linked to each other by one or more disulfide bonds depending on the H chain isotype. The VH and VL domains consist of four regions of relatively conserved sequences called framework regions (FR1, FR2, FR3, and FR4), which form a scaffold for three regions of hypervariable sequences (complementarity determining regions, CDRs). The CDRs contain most of the residues responsible for specific interactions of the antibody with the antigen. CDRs are referred to as CDR 1, CDR2, and CDR3. Accordingly, CDR constituents on the heavy chain are referred to as H1, H2, and H3, while CDR constituents on the light chain are referred to as L1, L2, and L3.

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The term "variable" refers to the portions of the immunoglobulin domains that exhibit variability in their sequence and that are involved in determining the specificity and binding affinity of a particular antibody (i.e., the "variable domain(s)"). Variability is not evenly distributed throughout the variable domains of antibodies; it is concentrated in sub-domains of each of the heavy and light chain variable regions. These sub-domains are called "hypervariable" regions or "complementarity determining regions" (CDRs). The more conserved (i.e., non-hypervariable) portions of the variable domains are called the "framework" regions (FRM). The variable domains of naturally occurring heavy and light chains each comprise four FRM regions, largely adopting a β-sheet configuration, connected by three hypervariable regions, which form loops connecting, and in some cases forming part of, the β-sheet structure. The hypervariable regions in each chain are held together in close proximity by the FRM and, with the hypervariable regions from the other chain, contribute to the formation of the antigen-binding site (see Kabat et al., loc. cit.). The constant domains are not directly involved in antigen binding, but exhibit various effector functions, such as, for example, antibody-dependent, cell-mediated cytotoxicity and complement activation.

The terms "CDR", and its plural "CDRs", refer to a complementarity determining region (CDR) of which three make up the binding character of a light chain variable region (CDRL1, CDRL2 and CDRL3) and three make up the binding character of a heavy chain variable region (CDRH1, CDRH2 and CDRH3). CDRs contribute to the functional activity of an antibody molecule and are separated by amino acid sequences that comprise scaffolding or framework regions. The exact definitional CDR boundaries and lengths are subject to different classification and numbering systems. CDRs may therefore be referred to by Kabat, Chothia, contact or any other boundary definitions, including the numbering system described herein. Despite differing boundaries, each of these systems has some degree of overlap in what constitutes the so called "hypervariable regions" within the variable sequences. CDR definitions according to these systems may therefore differ in length and boundary areas with respect to the adjacent framework region. See for example Kabat, Chothia, and/or MacCallum (Kabat et al., loc. cit.; Chothia et al., J. Mol. Biol, 1987, 196: 901; and MacCallum et al., J. Mol. Biol, 1996, 262: 732). However, the numbering in accordance with the so-called Kabat system is preferred. The CDR3 of the light chain and, particularly, CDR3 of the heavy chain may constitute the most important determinants in antigen binding within the light and heavy chain variable regions. In some antibody constructs, the heavy chain CDR3 appears to constitute the major area of contact between the antigen and the antibody. In vitro selection schemes in which CDR3 alone is varied can be used to vary the binding properties of an antibody or determine which residues contribute to the binding of an antigen.

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"Consisting essentially of" means that the amino acid sequence can vary by about 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, or 15% relative to the recited SEQ ID NO: sequence and still retain biological activity, as described herein.

In some embodiments, the antibody constructs of the invention are isolated proteins or substantially pure proteins. An "isolated" protein is unaccompanied by at least some of the material with which it is normally associated in its natural state, for example constituting at least about 5%, or at least about 50% by weight of the total protein in a given sample. It is understood that the isolated protein may constitute from 5 to 99.9% by weight of the total protein content depending on the circumstances. For example, the protein may be made at a significantly higher concentration through the use of an inducible promoter or high expression promoter, such that the protein is made at increased concentration levels. The definition includes the production of an antigen binding protein in a wide variety of organisms and/or host cells that are known in the art.

For amino acid sequences, sequence identity and/or similarity is determined by using standard techniques known in the art, including, but not limited to, the local sequence identity algorithm of Smith and Waterman, 1981, *Adv. Appl. Math.* 2:482, the sequence identity alignment algorithm of Needleman and Wunsch, 1970, *J. Mol. Biol.* 48:443, the search for similarity method of Pearson and Lipman, 1988, *Proc. Nat. Acad. Sci. U.S.A.* 85:2444, computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Drive, Madison, Wis.), the Best Fit sequence program described by Devereux *et al.*, 1984, *Nucl. Acid Res.* 12:387-395, preferably using the default settings, or by inspection. Preferably, percent identity is calculated by FastDB based upon the following parameters: mismatch penalty of 1; gap penalty of 1; gap size penalty of 0.33; and joining penalty of 30, "Current Methods in Sequence Comparison and Analysis," Macromolecule Sequencing and Synthesis, Selected Methods and Applications, pp 127-149 (1988), Alan R. Liss, Inc.

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An example of a useful algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using progressive, pairwise alignments. It can also plot a tree showing the clustering relationships used to create the alignment. PILEUP uses a simplification of the progressive alignment method of Feng & Doolittle, 1987, *J. Mol. Evol.* 35:351-360; the method is similar to that described by Higgins and Sharp, 1989, *CABIOS* 5:151-153. Useful PILEUP parameters including a default gap weight of 3.00, a default gap length weight of 0.10, and weighted end gaps.

Another example of a useful algorithm is the BLAST algorithm, described in: Altschul *et al.*, 1990, *J. Mol. Biol.* 215:403-410; Altschul *et al.*, 1997, *Nucleic Acids Res.* 25:3389-3402; and Karin *et al.*, 1993, *Proc. Natl. Acad. Sci. U.S.A.* 90:5873-5787. A particularly useful BLAST program is the WU-BLAST-2 program which was obtained from Altschul *et al.*, 1996, *Methods in Enzymology* 266:460-480. WU-BLAST-2 uses several search parameters, most of which are set to the default values. The adjustable parameters are set with the following values: overlap span=1, overlap fraction=0.125, word threshold (T)=II. The HSP S and HSP S2 parameters are dynamic values and are established by the program itself depending upon the composition of the particular sequence and composition of the particular database against which the sequence of interest is being searched; however, the values may be adjusted to increase sensitivity.

An additional useful algorithm is gapped BLAST as reported by Altschul *et al.*, 1993, *Nucl. Acids Res.* 25:3389-3402. Gapped BLAST uses BLOSUM-62 substitution scores; threshold T parameter set to 9; the two-hit method to trigger ungapped extensions, charges gap

lengths of k a cost of 10+k; Xu set to 16, and Xg set to 40 for database search stage and to 67 for the output stage of the algorithms. Gapped alignments are triggered by a score corresponding to about 22 bits.

- Generally, the amino acid homology, similarity, or identity between individual variant CDRs are at least 80% to the sequences depicted herein, and more typically with preferably increasing homologies or identities of at least 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, and almost 100%. In a similar manner, "percent (%) nucleic acid sequence identity" with respect to the nucleic acid sequence of the binding proteins identified herein is defined as the percentage of nucleotide residues in a candidate sequence that are identical with the nucleotide residues in the coding sequence of the antigen binding protein. A specific method utilizes the BLASTN module of WU-BLAST-2 set to the default parameters, with overlap span and overlap fraction set to 1 and 0.125, respectively.
- Generally, the nucleic acid sequence homology, similarity, or identity between the nucleotide sequences encoding individual variant CDRs and the nucleotide sequences depicted herein are at least 80%, and more typically with preferably increasing homologies or identities of at least 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%, and almost 100%.
- Thus, a "variant CDR" is one with the specified homology, similarity, or identity to the parent CDR of the invention, and shares biological function, including, but not limited to, at least 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% of the specificity and/or activity of the parent CDR.
- While the site or region for introducing an amino acid sequence variation is predetermined, the mutation *per se* need not be predetermined. For example, in order to optimize the performance of a mutation at a given site, random mutagenesis may be conducted at the target codon or region and the expressed antigen binding protein CDR variants screened for the optimal combination of desired activity. Techniques for making substitution mutations at predetermined sites in DNA having a known sequence are well known, for example, M13 primer mutagenesis and PCR mutagenesis. Screening of the mutants is done using assays of antigen binding protein activities, such as CDH19 binding.
- The term "amino acid" or "amino acid residue" typically refers to an amino acid having its art recognized definition such as an amino acid selected from the group consisting of: alanine (Ala or A); arginine (Arg or R); asparagine (Asn or N); aspartic acid (Asp or D); cysteine (Cys or C); glutamine (Gln or Q); glutamic acid (Glu or E); glycine (Gly or G); histidine (His or H);

isoleucine (He or I): leucine (Leu or L); lysine (Lys or K); methionine (Met or M); phenylalanine (Phe or F); pro line (Pro or P); serine (Ser or S); threonine (Thr or T); tryptophan (Trp or W); tyrosine (Tyr or Y); and valine (Val or V), although modified, synthetic, or rare amino acids may be used as desired. Generally, amino acids can be grouped as having a nonpolar side chain (e.g., Ala, Cys, He, Leu, Met, Phe, Pro, Val); a negatively charged side chain (e.g., Asp, Glu); a positively charged sidechain (e.g., Arg, His, Lys); or an uncharged polar side chain (e.g., Asn, Cys, Gln, Gly, His, Met, Phe, Ser, Thr, Trp, and Tyr).

The term "hypervariable region" (also known as "complementarity determining regions" or CDRs) when used herein refers to the amino acid residues of an antibody which are (usually three or four short regions of extreme sequence variability) within the V-region domain of an immunoglobulin which form the antigen-binding site and are the main determinants of antigen specificity. There are at least two methods for identifying the CDR residues: (1) An approach based on cross-species sequence variability (i. e., Kabat *et al.*, loc. cit.); and (2) An approach based on crystallographic studies of antigen-antibody complexes (Chothia, C. *et al.*, J. Mol. Biol. 196: 901-917 (1987)). However, to the extent that two residue identification techniques define regions of overlapping, but not identical regions, they can be combined to define a hybrid CDR. However, in general, the CDR residues are preferably identified in accordance with the so-called Kabat (numbering) system.

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The term "framework region" refers to the art-recognized portions of an antibody variable region that exist between the more divergent (i.e., hypervariable) CDRs. Such framework regions are typically referred to as frameworks 1 through 4 (FR1, FR2, FR3, and FR4) and provide a scaffold for the presentation of the six CDRs (three from the heavy chain and three from the light chain) in three dimensional space, to form an antigen-binding surface.

Typically, CDRs form a loop structure that can be classified as a canonical structure. The term "canonical structure" refers to the main chain conformation that is adopted by the antigen binding (CDR) loops. From comparative structural studies, it has been found that five of the six antigen binding loops have only a limited repertoire of available conformations. Each canonical structure can be characterized by the torsion angles of the polypeptide backbone. Correspondent loops between antibodies may, therefore, have very similar three dimensional structures, despite high amino acid sequence variability in most parts of the loops (Chothia and Lesk, J. Mol. Biol., 1987, 196: 901; Chothia *et al.*, Nature, 1989, 342: 877; Martin and Thornton, J. Mol. Biol, 1996, 263: 800, each of which is incorporated by reference in its entirety). Furthermore, there is a relationship between the adopted loop structure and the amino acid sequences surrounding it. The conformation of a particular

canonical class is determined by the length of the loop and the amino acid residues residing at key positions within the loop, as well as within the conserved framework (i.e., outside of the loop). Assignment to a particular canonical class can therefore be made based on the presence of these key amino acid residues. The term "canonical structure" may also include considerations as to the linear sequence of the antibody, for example, as catalogued by Kabat (Kabat et al., loc. cit.). The Kabat numbering scheme (system) is a widely adopted standard for numbering the amino acid residues of an antibody variable domain in a consistent manner and is the preferred scheme applied in the present invention as also mentioned elsewhere herein. Additional structural considerations can also be used to determine the canonical structure of an antibody. For example, those differences not fully reflected by Kabat numbering can be described by the numbering system of Chothia et al and/or revealed by other techniques, for example, crystallography and two or threedimensional computational modeling. Accordingly, a given antibody sequence may be placed into a canonical class which allows for, among other things, identifying appropriate chassis sequences (e.g., based on a desire to include a variety of canonical structures in a library). Kabat numbering of antibody amino acid sequences and structural considerations as described by Chothia et al., loc. cit. and their implications for construing canonical aspects of antibody structure, are described in the literature.

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CDR3 is typically the greatest source of molecular diversity within the antibody-binding site. H3, for example, can be as short as two amino acid residues or greater than 26 amino acids. The subunit structures and three-dimensional configurations of different classes of immunoglobulins are well known in the art. For a review of the antibody structure, see Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, eds. Harlow et al., 1988. One of skill in the art will recognize that each subunit structure, e.g., a CH, VH, CL, VL, CDR, FR structure, comprises active fragments, e.g., the portion of the VH, VL, or CDR subunit the binds to the antigen, i.e., the antigen-binding fragment, or, e.g., the portion of the CH subunit that binds to and/or activates, e.g., an Fc receptor and/or complement. The CDRs typically refer to the Kabat CDRs, as described in Sequences of Proteins of immunological Interest, US Department of Health and Human Services (1991), eds. Kabat et al. Another standard for characterizing the antigen binding site is to refer to the hypervariable loops as described by Chothia. See, e.g., Chothia, et al. (1987; J. Mol. Biol. 227:799-817); and Tomlinson et al. (1995) EMBO J. 14: 4628-4638. Still another standard is the AbM definition used by Oxford Molecular's AbM antibody modeling software. See, generally, e.g., Protein Sequence and Structure Analysis of Antibody Variable Domains. In: Antibody Engineering Lab Manual (Ed.: Duebel, S. and Kontermann, R., Springer-Verlag, Heidelberg).

Embodiments described with respect to Kabat CDRs can alternatively be implemented using similar described relationships with respect to Chothia hypervariable loops or to the AbM-defined loops.

5 The sequence of antibody genes after assembly and somatic mutation is highly varied, and these varied genes are estimated to encode 10¹⁰ different antibody molecules (Immunoglobulin Genes, 2nd ed., eds. Jonio et al., Academic Press, San Diego, CA, 1995). Accordingly, the immune system provides a repertoire of immunoglobulins. The term "repertoire" refers to at least one nucleotide sequence derived wholly or partially from at least one sequence encoding at least one immunoglobulin. The sequence(s) may be 10 generated by rearrangement in vivo of the V, D, and J segments of heavy chains, and the V and J segments of light chains. Alternatively, the sequence(s) can be generated from a cell in response to which rearrangement occurs, e.g., in vitro stimulation. Alternatively, part or all of the sequence(s) may be obtained by DNA splicing, nucleotide synthesis, mutagenesis, 15 and other methods, see, e.g., U.S. Patent 5,565,332. A repertoire may include only one sequence or may include a plurality of sequences, including ones in a genetically diverse collection.

The term "binding molecule" or "antibody construct" in the sense of the present disclosure indicates any molecule capable of (specifically) binding to, interacting with or recognizing the target molecules CDH19 and CD3. Such molecules or constructs may include proteinaceous parts and non-proteinaceous parts (e.g. chemical linkers or chemical cross-linking agents such as glutaraldehyde).

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In the event that a linker is used, this linker is preferably of a length and sequence sufficient to ensure that each of the first and second domains can, independently from one another, retain their differential binding specificities. Most preferably and as documented in the appended examples, the antibody construct of the invention is a "bispecific single chain antibody construct", more prefereably a bispecific single chain Fv (scFv). Bispecific single chain molecules are known in the art and are described in WO 99/54440, Mack, J. Immunol. (1997), 158, 3965-3970, Mack, PNAS, (1995), 92, 7021-7025, Kufer, Cancer Immunol. Immunother., (1997), 45, 193-197, Löffler, Blood, (2000), 95, 6, 2098-2103, Brühl, Immunol., (2001), 166, 2420-2426, Kipriyanov, J. Mol. Biol., (1999), 293, 41-56.

35 The said variable domains comprised in the herein described antibody constructs may be connected by additional linker sequences. The term "peptide linker" defines in accordance with the present invention an amino acid sequence by which the amino acid sequences of

the first domain and the second domain of the antibody construct of the invention are linked with each other. An essential technical feature of such peptide linker is that said peptide linker does not comprise any polymerization activity. Among the suitable peptide linkers are those described in U.S. Patents 4,751,180 and 4,935,233 or WO 88/09344. A preferred embodiment of a peptide linker is characterized by the amino acid sequence Gly-Gly-Gly-Gly-Ser, i.e. Gly₄Ser, or polymers thereof, i.e. (Gly₄Ser)x, where x is an integer 1 or greater. The characteristics of said peptide linker, which comprise the absence of the promotion of secondary structures are known in the art and described e.g. in Dall'Acqua et al. (Biochem. (1998) 37, 9266-9273), Cheadle et al. (Mol Immunol (1992) 29, 21-30) and Raag and Whitlow (FASEB (1995) 9(1), 73-80). Peptide linkers which also do not promote any secondary structures are preferred. The linkage of said domains to each other can be provided by, e.g. genetic engineering, as described in the examples. Methods for preparing fused and operatively linked bispecific single chain constructs and expressing them in mammalian cells or bacteria are well-known in the art (e.g. WO 99/54440 or Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York, 2001).

For peptide linkers, which connect the at least two binding domains in the antibody construct of the invention peptide linkers are preferred which comprise only a few number of amino acid residues, e.g. 12 amino acid residues or less. Thus, peptide linker of 12, 11, 10, 9, 8, 7, 6 or 5 amino acid residues are preferred. An envisaged peptide linker with less than 5 amino acids comprises 4, 3, 2 or one amino acid(s) wherein Gly-rich linkers are preferred. A particularly preferred "single" amino acid in context of said "peptide linker" is Gly. Accordingly, said peptide linker may consist of the single amino acid Gly.

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The term "multispecific" as used herein refers to a binding molecule which is an antibody construct and comprises at least a first and a second binding domain, wherein the first binding domain is capable of binding to one antigen or target, and the second binding domain is capable of binding to another antigen or target. Accordingly, antibody constructs according to the invention comprise at least specificities for two different antigens or targets and are at least bispecific. The "antibody construct" of the invention also comprises multispecific binding molecules such as e.g. trispecific binding molecules, the latter ones including three binding domains.

It is also envisaged that the antibody construct of the invention has, in addition to its function to bind to the target molecules CDH19 and CD3, a further function. In this format, the antibody construct is a tri-or multifunctional antibody construct by targeting plasma cells

through binding to CDH19, mediating cytotoxic T cell activity through CD3 binding and providing a further function such as a fully functional Fc constant domain mediating antibody-dependent cellular cytotoxicity through recruitment of effector cells like NK cells, a label (fluorescent etc.), a therapeutic agent such as, e.g. a toxin or radionuclide, and/or means to enhance serum half-life, etc.

The term "binding domain" characterizes in connection with the present invention a domain which is capable of specifically binding to / interacting with a given target epitope or a given target site on the target molecules CDH19 and CD3.

Binding domains can be derived from a binding domain donor such as for example an antibody. It is envisaged that a binding domain of the present invention comprises at least said part of any of the aforementioned binding domains that is required for binding to/interacting with a given target epitope or a given target site on the target molecules CDH19 and CD3.

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It is envisaged that the binding domain of the aforementioned binding domain donors is characterized by that part of these donors that is responsible for binding the respective target, i.e. when that part is removed from the binding domain donor, said donor loses its binding capability. "Loses" means a reduction of at least 50% of the binding capability when compared with the binding donor. Methods to map these binding sites are well known in the art – it is therefore within the standard knowledge of the skilled person to locate/map the binding site of a binding domain donor and, thereby, to "derive" said binding domain from the respective binding domain donors.

The term "epitope" refers to a site on an antigen to which a binding domain, such as an antibody or immunoglobulin or derivative or fragment of an antibody or of an immunoglobulin, specifically binds. An "epitope" is antigenic and thus the term epitope is sometimes also referred to herein as "antigenic structure" or "antigenic determinant". Thus, the binding domain is an "antigen-interaction-site". Said binding/interaction is also understood to define a "specific recognition". In one example, said binding domain which (specifically) binds to / interacts with a given target epitope or a given target site on the target molecules CDH19 and CD3 is an antibody or immunoglobulin, and said binding domain is a VH and/or VL region of an antibody or of an immunoglobulin.

35 "Epitopes" can be formed both by contiguous amino acids or non-contiguous amino acids juxtaposed by tertiary folding of a protein. A "linear epitope" is an epitope where an amino acid primary sequence comprises the recognized epitope. A linear epitope typically includes

at least 3 or at least 4, and more usually, at least 5 or at least 6 or at least 7, for example, about 8 to about 10 amino acids in a unique sequence.

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A "conformational epitope", in contrast to a linear epitope, is an epitope wherein the primary sequence of the amino acids comprising the epitope is not the sole defining component of the epitope recognized (e.g., an epitope wherein the primary sequence of amino acids is not necessarily recognized by the binding domain). Typically a conformational epitope comprises an increased number of amino acids relative to a linear epitope. With regard to recognition of conformational epitopes, the binding domain recognizes a three-dimensional structure of the antigen, preferably a peptide or protein or fragment thereof (in the context of the present invention, the antigen for one of the binding domains is comprised within the CDH19 protein). For example, when a protein molecule folds to form a three-dimensional structure, certain amino acids and/or the polypeptide backbone forming the conformational epitope become juxtaposed enabling the antibody to recognize the epitope. Methods of determining the conformation of epitopes include, but are not limited to, x-ray crystallography, two-dimensional nuclear magnetic resonance (2D-NMR) spectroscopy and site-directed spin labelling and electron paramagnetic resonance (EPR) spectroscopy. Moreover, the provided examples describe a further method to characterize a given binding domain by way of binning, which includes a test whether the given binding domain binds to one or more epitope cluster(s) of a given protein, in particular CDH19.

As used herein, the term "epitope cluster" denotes the entirety of epitopes lying in a defined contiguous stretch of an antigen. An epitope cluster can comprise one, two or more epitopes. The concept of epitope cluster is also used in the characterization of the features of the antibody constructs of the invention.

The terms "(capable of) binding to", "specifically recognizing", "directed to" and "reacting with" mean in accordance with this invention that a binding domain is capable of specifically interacting with one or more, preferably at least two, more preferably at least three and most preferably at least four amino acids of an epitope.

As used herein, the terms "specifically interacting", "specifically binding" or "specifically bind(s)" mean that a binding domain exhibits appreciable affinity for a particular protein or antigen and, generally, does not exhibit significant reactivity with proteins or antigens other than CDH19 or CD3. "Appreciable affinity" includes binding with an affinity of about 10⁻⁶M (KD) or stronger. Preferably, binding is considered specific when binding affinity is about 10⁻¹² to 10⁻⁸ M, 10⁻¹² to 10⁻⁹ M, 10⁻¹² to 10⁻¹⁰ M, 10⁻¹¹ to 10⁻⁸ M, preferably of about 10⁻¹¹ to 10⁻¹²

⁹ M. Whether a binding domain specifically reacts with or binds to a target can be tested readily by, *inter alia*, comparing the reaction of said binding domain with a target protein or antigen with the reaction of said binding domain with proteins or antigens other than CDH19 or CD3. Preferably, a binding domain of the invention does not essentially bind or is not capable of binding to proteins or antigens other than CDH19 or CD3 (i.e. the first binding domain is not capable of binding to proteins other than CDH19 and the second binding domain is not capable of binding to proteins other than CD3).

The term "does not essentially bind", or "is not capable of binding" means that a binding domain of the present invention does not bind another protein or antigen other than CDH19 or CD3, i.e., does not show reactivity of more than 30%, preferably not more than 20%, more preferably not more than 10%, particularly preferably not more than 9%, 8%, 7%, 6% or 5% with proteins or antigens other than CDH19 or CD3, whereby binding to CDH19 or CD3, respectively, is set to be 100%.

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Specific binding is believed to be effected by specific motifs in the amino acid sequence of the binding domain and the antigen. Thus, binding is achieved as a result of their primary, secondary and/or tertiary structure as well as the result of secondary modifications of said structures. The specific interaction of the antigen-interaction-site with its specific antigen may result in a simple binding of said site to the antigen. Moreover, the specific interaction of the antigen-interaction-site with its specific antigen may alternatively or additionally result in the initiation of a signal, e.g. due to the induction of a change of the conformation of the antigen, an oligomerization of the antigen, etc.

Proteins (including fragments thereof, preferably biologically active fragments, and peptides, 25 usually having less than 30 amino acids) comprise one or more amino acids coupled to each other via a covalent peptide bond (resulting in a chain of amino acids). The term "polypeptide" as used herein describes a group of molecules, which consist of more than 30 amino acids. Polypeptides may further form multimers such as dimers, trimers and higher 30 oligomers, i.e. consisting of more than one polypeptide molecule. Polypeptide molecules forming such dimers, trimers etc. may be identical or non-identical. The corresponding higher order structures of such multimers are, consequently, termed homo- or heterodimers, homo- or heterotrimers etc. An example for a hereteromultimer is an antibody molecule, which, in its naturally occurring form, consists of two identical light polypeptide chains and 35 two identical heavy polypeptide chains. The terms "polypeptide" and "protein" also refer to naturally modified polypeptides/proteins wherein the modification is effected e.g. by posttranslational modifications like glycosylation, acetylation, phosphorylation and the like. A

"polypeptide" when referred to herein may also be chemically modified such as pegylated. Such modifications are well known in the art.

"Isolated" when used to describe the antibody construct disclosed herein, means a antibody construct that has been identified, separated and/or recovered from a component of its production environment. Preferably, the isolated antibody construct is free of association with all other components from its production environment. Contaminant components of its production environment, such as that resulting from recombinant transfected cells, are materials that would typically interfere with diagnostic or therapeutic uses for the polypeptide, and may include enzymes, hormones, and other proteinaceous or non-proteinaceous solutes. In preferred embodiments, the antibody construct will be purified (1) to a degree sufficient to obtain at least 15 residues of N-terminal or internal amino acid sequence by use of a spinning cup sequenator, or (2) to homogeneity by SDS-PAGE under non-reducing or reducing conditions using Coomassie blue or, preferably, silver stain. Ordinarily, however, an isolated antibody will be prepared by at least one purification step.

Amino acid sequence modifications of the antibody constructs described herein are contemplated. For example, it may be desirable to improve the binding affinity and/or other biological properties of the antibody. Amino acid sequence variants of the antibody constructs are prepared by introducing appropriate nucleotide changes into the antibody constructs nucleic acid, or by peptide synthesis.

Such modifications include, for example, deletions from, and/or insertions into, and/or substitutions of, residues within the amino acid sequences of the antibody constructs. Any combination of deletion, insertion, and substitution is made to arrive at the final construct, provided that the final construct possesses the desired characteristics. The amino acid changes also may alter post-translational processes of the antibody constructs, such as changing the number or position of glycosylation sites. Preferably, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 amino acids may be substituted in a CDR, while 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or 25 amino acids may be substituted in the framework regions (FRs). The substitutions are preferably conservative substitutions as described herein. Additionally or alternatively, 1, 2, 3, 4, 5, or 6 amino acids may be inserted or deleted in each of the CDRs (of course, dependent on their length), while 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or 25 amino acids may be inserted or deleted in each of the FRs.

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A useful method for identification of certain residues or regions of the antibody constructs that are preferred locations for mutagenesis is called "alanine scanning mutagenesis" as

described by Cunningham and Wells in Science, 244: 1081-1085 (1989). Here, a residue or group of target residues within the antibody construct is/are identified (e.g. charged residues such as arg, asp, his, lys, and glu) and replaced by a neutral or negatively charged amino acid (most preferably alanine or polyalanine) to affect the interaction of the amino acids with the epitope.

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Those amino acid locations demonstrating functional sensitivity to the substitutions then are refined by introducing further or other variants at, or for, the sites of substitution. Thus, while the site for introducing an amino acid sequence variation is predetermined, the nature of the mutation *per se* needs not to be predetermined. For example, to analyze the performance of a mutation at a given site, ala scanning or random mutagenesis is conducted at a target codon or region and the expressed antibody construct variants are screened for the desired activity.

Preferably, amino acid sequence insertions include amino- and/or carboxyl-terminal fusions ranging in length from 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 residues to polypeptides containing a hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. An insertional variant of the antibody construct includes the fusion to the N-or C-terminus of the antibody to an enzyme or a fusion to a polypeptide which increases the serum half-life of the antibody.

Another type of variant is an amino acid substitution variant. These variants have preferably at least 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 amino acid residues in the antibody construct replaced by a different residue. The sites of greatest interest for substitutional mutagenesis include the CDRs of the heavy and/or light chain, in particular the hypervariable regions, but FR alterations in the heavy and/or light chain are also contemplated.

For example, if a CDR sequence encompasses 6 amino acids, it is envisaged that one, two or three of these amino acids are substituted. Similarly, if a CDR sequence encompasses 15 amino acids it is envisaged that one, two, three, four, five or six of these amino acids are substituted.

Generally, if amino acids are substituted in one or more or all of the CDRs of the heavy and/or light chain, it is preferred that the then-obtained "substituted" sequence is at least 60%, more preferably 65%, even more preferably 70%, particularly preferably 75%, more particularly preferably 80% identical to the "original" CDR sequence. This means that it is dependent of the length of the CDR to which degree it is identical to the "substituted"

sequence. For example, a CDR having 5 amino acids is preferably 80% identical to its substituted sequence in order to have at least one amino acid substituted. Accordingly, the CDRs of the antibody construct may have different degrees of identity to their substituted sequences, e.g., CDRL1 may have 80%, while CDRL3 may have 90%.

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Preferred substitutions (or replacements) are conservative substitutions. However, any substitution (including non-conservative substitution or one or more from the "exemplary substitutions" listed in Table 1, below) is envisaged as long as the antibody construct retains its capability to bind to CDH19 via the first binding domain and to CD3 epsilon via the second binding domain and/or its CDRs have an identity to the then substituted sequence (at least 60%, more preferably 65%, even more preferably 70%, particularly preferably 75%, more particularly preferably 80% identical to the "original" CDR sequence).

Conservative substitutions are shown in Table 1 under the heading of "preferred substitutions". If such substitutions result in a change in biological activity, then more substantial changes, denominated "exemplary substitutions" in Table 1, or as further described below in reference to amino acid classes, may be introduced and the products screened for a desired characteristic.

20 Table 1: Amino Acid Substitutions

Original	Exemplary Substitutions	Preferred Substitutions
Ala (A)	val, leu, ile	val
Arg (R)	lys, gln, asn	lys
Asn (N)	gln, his, asp, lys, arg	gln
Asp (D)	glu, asn	glu
Cys (C)	ser, ala	ser
Gin (Q)	asn, glu	asn
Glu (E)	asp, gln	asp
Gly (G)	ala	ala
His (H)	asn, gln, lys, arg	arg
Ile (I)	leu, val, met, ala, phe	leu
Leu (L)	norleucine, ile, val, met, ala	ile
Lys (K)	arg, gln, asn	arg
Met (M)	leu, phe, ile	leu
Phe (F)	leu, val, ile, ala, tyr	tyr
Pro (P)	ala	ala

Ser (S)	thr	thr
Thr (T)	ser	ser
Trp (W)	tyr, phe	tyr
Tyr (Y)	trp, phe, thr, ser	phe
Val (V)	ile, leu, met, phe, ala	leu

Substantial modifications in the biological properties of the antibody construct of the present invention are accomplished by selecting substitutions that differ significantly in their effect on maintaining (a) the structure of the polypeptide backbone in the area of the substitution, for example, as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain. Naturally occurring residues are divided into groups based on common side-chain properties: (1) hydrophobic: norleucine, met, ala, val, leu, ile; (2) neutral hydrophilic: cys, ser, thr; (3) acidic: asp, glu; (4) basic: asn, gin, his, lys, arg; (5) residues that influence chain orientation: gly, pro; and (6) aromatic: trp, tyr, phe.

Non-conservative substitutions will entail exchanging a member of one of these classes for another class. Any cysteine residue not involved in maintaining the proper conformation of the antibody construct may be substituted, generally with serine, to improve the oxidative stability of the molecule and prevent aberrant crosslinking. Conversely, cysteine bond(s) may be added to the antibody to improve its stability (particularly where the antibody is an antibody fragment such as an Fv fragment).

A particularly preferred type of substitutional variant involves substituting one or more hypervariable region residues of a parent antibody (e. g. a humanized or human antibody). Generally, the resulting variant(s) selected for further development will have improved biological properties relative to the parent antibody from which they are generated. A convenient way for generating such substitutional variants involves affinity maturation using phage display. Briefly, several hypervariable region sites (e. g. 6-7 sites) are mutated to generate all possible amino acid substitutions at each site. The antibody variants thus generated are displayed in a monovalent fashion from filamentous phage particles as fusions to the gene III product of M13 packaged within each particle. The phage-displayed variants are then screened for their biological activity (e. g. binding affinity) as herein disclosed. In order to identify candidate hypervariable region sites for modification, alanine scanning mutagenesis can be performed to identify hypervariable region residues contributing significantly to antigen binding. Alternatively, or additionally, it may be beneficial

to analyze a crystal structure of the antigen-antibody complex to identify contact points between the binding domain and, e.g., human CDH19. Such contact residues and neighbouring residues are candidates for substitution according to the techniques elaborated herein. Once such variants are generated, the panel of variants is subjected to screening as described herein and antibodies with superior properties in one or more relevant assays may be selected for further development.

Other modifications of the antibody construct are contemplated herein. For example, the antibody construct may be linked to one of a variety of non-proteinaceous polymers, e.g., polyethylene glycol, polypropylene glycol, polyoxyalkylenes, or copolymers of polyethylene glycol and polypropylene glycol. The antibody construct may also be entrapped in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization (for example, hydroxymethylcellulose or gelatine-microcapsules and poly (methylmethacylate) microcapsules, respectively), in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nanoparticles and nanocapsules), or in macroemulsions. Such techniques are disclosed in Remington's Pharmaceutical Sciences, 16th edition, Oslo, A., Ed., (1980).

The antibody constructs disclosed herein may also be formulated as immuno-liposomes. A "liposome" is a small vesicle composed of various types of lipids, phospholipids and/or surfactant which is useful for delivery of a drug to a mammal. The components of the liposome are commonly arranged in a bilayer formation, similar to the lipid arrangement of biological membranes. Liposomes containing the antibody are prepared by methods known in the art, such as described in Epstein et al., Proc. Natl. Acad. Sci. USA, 82: 3688 (1985); Hwang et al., Proc. Natl Acad. Sci. USA, 77: 4030 (1980); US Pat. Nos. 4,485,045 and 4,544,545; and W0 97/38731 published October 23, 1997. Liposomes with enhanced circulation time are disclosed in US Patent No. 5,013, 556. Particularly useful liposomes can be generated by the reverse phase evaporation method with a lipid composition comprising phosphatidylcholine, cholesterol and PEG-derivatized phosphatidylethanolamine (PEG-PE). Liposomes are extruded through filters of defined pore size to yield liposomes with the desired diameter. Fab' fragments of the antibody of the present invention can be conjugated to the liposomes as described in Martin et al. J. Biol. Chem. 257: 286-288 (1982) via a disulfide interchange reaction. A chemotherapeutic agent is optionally contained within the liposome. See Gabizon et al. J. National Cancer Inst. 81 (19) 1484 (1989).

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When using recombinant techniques, the antibody construct can be produced intracellularly, in the periplasmic space, or directly secreted into the medium. If the antibody construct is

produced intracellularly, as a first step, the particulate debris, either host cells or lysed fragments, are removed, for example, by centrifugation or ultrafiltration. Carter *et al.*, Bio/Technology 10: 163-167 (1992) describe a procedure for isolating antibodies which are secreted to the periplasmic space of *E. coli*.

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The antibody construct composition prepared from the cells can be purified using, for example, hydroxylapatite chromatography, gel electrophoresis, dialysis, and affinity chromatography, with affinity chromatography being the preferred purification technique.

The term "nucleic acid" is well known to the skilled person and encompasses DNA (such as cDNA) and RNA (such as mRNA). The nucleic acid can be double stranded and single stranded, linear and circular. Said nucleic acid molecule is preferably comprised in a vector which is preferably comprised in a host cell. Said host cell is, e.g. after transformation or transfection with the nucleic acid sequence of the invention, capable of expressing the antibody construct. For that purpose the nucleic acid molecule is operatively linked with control sequences.

A vector is a nucleic acid molecule used as a vehicle to transfer (foreign) genetic material into a cell. The term "vector" encompasses – but is not restricted to – plasmids, viruses, cosmids and artificial chromosomes. In general, engineered vectors comprise an origin of replication, a multicloning site and a selectable marker. The vector itself is generally a nucleotide sequence, commonly a DNA sequence, that comprises an insert (transgene) and a larger sequence that serves as the "backbone" of the vector. Modern vectors may encompass additional features besides the transgene insert and a backbone: promoter, genetic marker, antibiotic resistance, reporter gene, targeting sequence, protein purification tag. Vectors called expression vectors (expression constructs) specifically are for the expression of the transgene in the target cell, and generally have control sequences such as a promoter sequence that drives expression of the transgene. Insertion of a vector into the target cell is usually called "transformation" for bacteria, "transfection" for eukaryotic cells, although insertion of a viral vector is also called "transduction".

As used herein, the term "host cell" is intended to refer to a cell into which a nucleic acid encoding the antibody construct of the invention is introduced by way of transformation, transfection and the like. It should be understood that such terms refer not only to the particular subject cell but to the progeny or potential progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental

influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term as used herein.

As used herein, the term "expression" includes any step involved in the production of a antibody construct of the invention including, but not limited to, transcription, post-transcriptional modification, translation, post-translational modification, and secretion.

The term "control sequences" refers to DNA sequences necessary for the expression of an operably linked coding sequence in a particular host organism. The control sequences that are suitable for prokaryotes, for example, include a promoter, optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

A nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous, and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is accomplished by ligation at convenient restriction sites. If such sites do not exist, the synthetic oligonucleotide adaptors or linkers are used in accordance with conventional practice.

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The terms "host cell," "target cell" or "recipient cell" are intended to include any individual cell or cell culture that can be or has/have been recipients for vectors or the incorporation of exogenous nucleic acid molecules, polynucleotides and/or proteins. It also is intended to include progeny of a single cell, and the progeny may not necessarily be completely identical (in morphology or in genomic or total DNA complement) to the original parent cell due to natural, accidental, or deliberate mutation. The cells may be prokaryotic or eukaryotic, and include but are not limited to bacteria, yeast cells, animal cells, and mammalian cells, e.g., murine, rat, macaque or human.

35 Suitable host cells include prokaryotes and eukaryotic host cells including yeasts, fungi, insect cells and mammalian cells.

The antibody construct of the invention can be produced in bacteria. After expression, the antibody construct of the invention, preferably the antibody construct is isolated from the *E. coli* cell paste in a soluble fraction and can be purified through, e.g., affinity chromatography and/or size exclusion. Final purification can be carried out similar to the process for purifying antibody expressed e. g, in CHO cells.

In addition to prokaryotes, eukaryotic microbes such as filamentous fungi or yeast are suitable cloning or expression hosts for the antibody construct of the invention. *Saccharomyces cerevisiae*, or common baker's yeast, is the most commonly used among lower eukaryotic host microorganisms. However, a number of other genera, species, and strains are commonly available and useful herein, such as *Schizosaccharomyces pombe*, Kluyveromyces hosts such as, e.g., *K. lactis, K. fragilis* (ATCC 12424), *K. bulgaricus* (ATCC 16045), *K. wickeramii* (ATCC 24178), *K. waltii* (ATCC 56500), *K. drosophilarum* (ATCC 36906), *K. thermotolerans*, and *K. marxianus*; yarrowia (EP 402 226); *Pichia pastoris* (EP 183 070); Candida; *Trichoderma reesia* (EP 244 234); *Neurospora crassa*; Schwanniomyces such as *Schwanniomyces occidentalis*; and filamentous fungi such as, e.g., Neurospora, Penicillium, Tolypocladium, and Aspergillus hosts such as *A. nidulans* and *A. niger*.

Suitable host cells for the expression of glycosylated antibody construct of the invention, preferably antibody derived antibody constructs are derived from multicellular organisms. Examples of invertebrate cells include plant and insect cells. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts such as *Spodoptera frugiperda* (caterpillar), *Aedes aegypti* (mosquito), *Aedes albopictus* (mosquito), *Drosophila melanogaster* (fruit fly), and *Bombyx mori* have been identified. A variety of viral strains for transfection are publicly available, e. g. , the L-1 variant of *Autographa californica* NPV and the Bm-5 strain of *Bombyx mori* NPV, and such viruses may be used as the virus herein according to the present invention, particularly for transfection of *Spodoptera frugiperda* cells.

Plant cell cultures of cotton, corn, potato, soybean, petunia, tomato, Arabidopsis and tobacco can also be utilized as hosts. Cloning and expression vectors useful in the production of proteins in plant cell culture are known to those of skill in the art. See e.g. Hiatt et al., Nature (1989) 342: 76-78, Owen et al. (1992) Bio/Technology 10: 790-794, Artsaenko et al. (1995) The Plant J 8: 745-750, and Fecker et al. (1996) Plant Mol Biol 32: 979-986.

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However, interest has been greatest in vertebrate cells, and propagation of vertebrate cells in culture (tissue culture) has become a routine procedure. Examples of useful mammalian

host cell lines are monkey kidney CV1 line transformed by SV40 (COS-7, ATCC CRL 1651); human embryonic kidney line (293 or 293 cells subcloned for growth in suspension culture, Graham et al. , J. Gen Virol. 36 : 59 (1977)); baby hamster kidney cells (BHK, ATCC CCL 10); Chinese hamster ovary cells/-DHFR (CHO, Urlaub *et al.* , Proc. Natl. Acad. Sci. USA 77: 4216 (1980)); mouse sertoli cells (TM4, Mather, Biol. Reprod. 23: 243-251 (1980)); monkey kidney cells (CVI ATCC CCL 70); African green monkey kidney cells (VERO-76, ATCC CRL1587); human cervical carcinoma cells (HELA, ATCC CCL 2); canine kidney cells (MDCK, ATCC CCL 34); buffalo rat liver cells (BRL 3A, ATCC CRL 1442); human lung cells (W138, ATCC CCL 75); human liver cells (Hep G2,1413 8065); mouse mammary tumor (MMT 060562, ATCC CCL5 1); TRI cells (Mather *et al.*, Annals N. Y Acad. Sci. 383 : 44-68 (1982)); MRC 5 cells; FS4 cells; and a human hepatoma line (Hep G2).

When using recombinant techniques, the antibody construct of the invention can be produced intracellularly, in the periplasmic space, or directly secreted into the medium. If the antibody construct is produced intracellularly, as a first step, the particulate debris, either host cells or lysed fragments, are removed, for example, by centrifugation or ultrafiltration. Carter *et al.*, Bio/Technology 10: 163-167 (1992) describe a procedure for isolating antibodies which are secreted to the periplasmic space of *E. coli.* Briefly, cell paste is thawed in the presence of sodium acetate (pH 3.5), EDTA, and phenylmethylsulfonylfluoride (PMSF) over about 30 min. Cell debris can be removed by centrifugation. Where the antibody is secreted into the medium, supernatants from such expression systems are generally first concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. A protease inhibitor such as PMSF may be included in any of the foregoing steps to inhibit proteolysis and antibiotics may be included to prevent the growth of adventitious contaminants.

The antibody construct of the invention prepared from the host cells can be purified using, for example, hydroxylapatite chromatography, gel electrophoresis, dialysis, and affinity chromatography, with affinity chromatography being the preferred purification technique.

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The matrix to which the affinity ligand is attached is most often agarose, but other matrices are available. Mechanically stable matrices such as controlled pore glass or poly (styrenedivinyl) benzene allow for faster flow rates and shorter processing times than can be achieved with agarose. Where the antibody construct of the invention comprises a CH3 domain, the Bakerbond ABXMresin (J. T. Baker, Phillipsburg, NJ) is useful for purification. Other techniques for protein purification such as fractionation on an ion-exchange column, ethanol precipitation, Reverse Phase HPLC, chromatography on silica, chromatography on

heparin SEPHAROSETM chromatography on an anion or cation exchange resin (such as a polyaspartic acid column), chromato-focusing, SDS-PAGE, and ammonium sulfate precipitation are also available depending on the antibody to be recovered.

5 The term "culturing" refers to the in vitro maintenance, differentiation, growth, proliferation and/or propagation of cells under suitable conditions in a medium.

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As used herein, the term "pharmaceutical composition" relates to a composition for administration to a patient, preferably a human patient. The particular preferred pharmaceutical composition of this invention comprises the antibody construct of the invention. Preferably, the pharmaceutical composition comprises suitable formulations of carriers, stabilizers and/or excipients. In a preferred embodiment, the pharmaceutical composition comprises a composition for parenteral, transdermal, intraluminal, intraarterial, intrathecal and/or intranasal administration or by direct injection into tissue. It is in particular envisaged that said composition is administered to a patient via infusion or injection. Administration of the suitable compositions may be effected by different ways, e.g., by intravenous, intraperitoneal, subcutaneous, intramuscular, topical or intradermal administration. In particular, the present invention provides for an uninterrupted administration of the suitable composition. As a non-limiting example, uninterrupted, i.e. continuous administration may be realized by a small pump system worn by the patient for metering the influx of therapeutic agent into the body of the patient. The pharmaceutical composition comprising the antibody construct of the invention can be administered by using said pump systems. Such pump systems are generally known in the art, and commonly rely on periodic exchange of cartridges containing the therapeutic agent to be infused. When exchanging the cartridge in such a pump system, a temporary interruption of the otherwise uninterrupted flow of therapeutic agent into the body of the patient may ensue. In such a case, the phase of administration prior to cartridge replacement and the phase of administration following cartridge replacement would still be considered within the meaning of the pharmaceutical means and methods of the invention together make up one "uninterrupted administration" of such therapeutic agent.

The continuous or uninterrupted administration of these antibody constructs of the invention may be intravenous or subcutaneous by way of a fluid delivery device or small pump system including a fluid driving mechanism for driving fluid out of a reservoir and an actuating mechanism for actuating the driving mechanism. Pump systems for subcutaneous administration may include a needle or a cannula for penetrating the skin of a patient and

delivering the suitable composition into the patient's body. Said pump systems may be directly fixed or attached to the skin of the patient independently of a vein, artery or blood vessel, thereby allowing a direct contact between the pump system and the skin of the patient. The pump system can be attached to the skin of the patient for 24 hours up to several days. The pump system may be of small size with a reservoir for small volumes. As a non-limiting example, the volume of the reservoir for the suitable pharmaceutical composition to be administered can be between 0.1 and 50 ml.

The continuous administration may be transdermal by way of a patch worn on the skin and replaced at intervals. One of skill in the art is aware of patch systems for drug delivery suitable for this purpose. It is of note that transdermal administration is especially amenable to uninterrupted administration, as exchange of a first exhausted patch can advantageously be accomplished simultaneously with the placement of a new, second patch, for example on the surface of the skin immediately adjacent to the first exhausted patch and immediately prior to removal of the first exhausted patch. Issues of flow interruption or power cell failure do not arise.

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The inventive compositions may further comprise a pharmaceutically acceptable carrier. Examples of suitable pharmaceutical carriers are well known in the art and include solutions, e.g. phosphate buffered saline solutions, water, emulsions, such as oil/water emulsions, various types of wetting agents, sterile solutions, liposomes, etc. Compositions comprising such carriers can be formulated by well known conventional methods. Formulations can comprise carbohydrates, buffer solutions, amino acids and/or surfactants. Carbohydrates may be non-reducing sugars, preferably trehalose, sucrose, octasulfate, sorbitol or xylitol. In general, as used herein, "pharmaceutically acceptable carrier" means any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, compatible with pharmaceutical administration. The use of such media and agents for pharmaceutically active substances is well known in the art. Acceptable carriers, excipients, or stabilizers are nontoxic to recipients at the dosages and concentrations employed and include: additional buffering agents; preservatives; co-solvents; antioxidants, including ascorbic acid and methionine; chelating agents such as EDTA; metal complexes (e.g., Zn-protein complexes); biodegradable polymers, such as polyesters; salt-forming counter-ions, such as sodium, polyhydric sugar alcohols; amino acids, such as alanine, glycine, asparagine, 2-phenylalanine, and threonine; sugars or sugar alcohols, such as trehalose, sucrose, octasulfate, sorbitol or xylitol stachyose, mannose, sorbose, xylose, ribose, myoinisitose, galactose, lactitol, ribitol, myoinisitol, galactitol, glycerol, cyclitols (e.g.,

inositol), polyethylene glycol; sulfur containing reducing agents, such as glutathione, thioctic acid, sodium thioglycolate, thioglycerol, [alpha]-monothioglycerol, and sodium thio sulfate; low molecular weight proteins, such as human serum albumin, bovine serum albumin, gelatin, or other immunoglobulins; and hydrophilic polymers, such as polyvinylpyrrolidone. Such formulations may be used for continuous administrations which may be intravenuous or subcutaneous with and/or without pump systems. Amino acids may be charged amino acids, preferably lysine, lysine acetate, arginine, glutamate and/or histidine. Surfactants may be detergents, preferably with a molecular weight of >1.2 KD and/or a polyether, preferably with a molecular weight of >3 KD. Non-limiting examples for preferred detergents are Tween 20, Tween 40, Tween 60, Tween 80 or Tween 85. Non-limiting examples for preferred polyethers are PEG 3000, PEG 3350, PEG 4000 or PEG 5000. Buffer systems used in the present invention can have a preferred pH of 5-9 and may comprise citrate, succinate, phosphate, histidine and acetate.

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The compositions of the present invention can be administered to the subject at a suitable dose which can be determined e.g. by dose escalating studies by administration of increasing doses of the polypeptide of the invention exhibiting cross-species specificity described herein to non-chimpanzee primates, for instance macaques. As set forth above, the antibody construct of the invention exhibiting cross-species specificity described herein can be advantageously used in identical form in preclinical testing in non-chimpanzee primates and as drug in humans. These compositions can also be administered in combination with other proteinaceous and non-proteinaceous drugs. These drugs may be administered simultaneously with the composition comprising the polypeptide of the invention as defined herein or separately before or after administration of said polypeptide in timely defined intervals and doses. The dosage regimen will be determined by the attending physician and clinical factors. As is well known in the medical arts, dosages for any one patient depend upon many factors, including the patient's size, body surface area, age, the particular compound to be administered, sex, time and route of administration, general health, and other drugs being administered concurrently.

30 Preparations for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's, or fixed

oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers (such as those based on Ringer's dextrose), and the like. Preservatives and other additives may also be present such as, for example, antimicrobials, anti-oxidants, chelating agents, inert gases and the like. In addition, the composition of the present invention might comprise proteinaceous carriers, like, e.g., serum albumin or immunoglobulin, preferably of human origin. It is envisaged that the composition of the invention might comprise, in addition to the polypeptide of the invention defined herein, further biologically active agents, depending on the intended use of the composition. Such agents might be drugs acting on the gastro-intestinal system, drugs acting as cytostatica, drugs preventing hyperurikemia, drugs inhibiting immunoreactions (e.g. corticosteroids), drugs modulating the inflammatory response, drugs acting on the circulatory system and/or agents such as cytokines known in the art. It is also envisaged that the antibody construct of the present invention is applied in a co-therapy, i.e., in combination with another anti-cancer medicament.

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The biological activity of the pharmaceutical composition defined herein can be determined for instance by cytotoxicity assays, as described in the following examples, in WO 99/54440 or by Schlereth et al. (Cancer Immunol. Immunother. 20 (2005), 1-12). "Efficacy" or "in vivo efficacy" as used herein refers to the response to therapy by the pharmaceutical composition of the invention, using e.g. standardized NCI response criteria. The success or in vivo efficacy of the therapy using a pharmaceutical composition of the invention refers to the effectiveness of the composition for its intended purpose, i.e. the ability of the composition to cause its desired effect, i.e. depletion of pathologic cells, e.g. tumor cells. The in vivo efficacy may be monitored by established standard methods for the respective disease entities including, but not limited to white blood cell counts, differentials, Fluorescence Activated Cell Sorting, bone marrow aspiration. In addition, various disease specific clinical chemistry parameters and other established standard methods may be used. Furthermore, computer-aided tomography, X-ray, nuclear magnetic resonance tomography (e.g. for National Cancer Institute-criteria based response assessment [Cheson BD, Horning SJ, Coiffier B, Shipp MA, Fisher RI, Connors JM, Lister TA, Vose J, Grillo-Lopez A, Hagenbeek A, Cabanillas F, Klippensten D, Hiddemann W, Castellino R, Harris NL, Armitage JO, Carter W, Hoppe R, Canellos GP. Report of an international workshop to standardize response criteria for non-Hodgkin's lymphomas. NCI Sponsored International Working Group. J Clin Oncol. 1999 Apr;17(4):1244]), positron-emission tomography scanning, white blood cell counts, differentials, Fluorescence Activated Cell Sorting, bone marrow aspiration, lymph node biopsies/histologies, and various lymphoma specific clinical chemistry parameters (e.g. lactate dehydrogenase) and other established standard methods may be used.

Another major challenge in the development of drugs such as the pharmaceutical composition of the invention is the predictable modulation of pharmacokinetic properties. To this end, a pharmacokinetic profile of the drug candidate, i.e. a profile of the pharmacokinetic parameters that affect the ability of a particular drug to treat a given condition, can be established. Pharmacokinetic parameters of the drug influencing the ability of a drug for treating a certain disease entity include, but are not limited to: half-life, volume of distribution, hepatic first-pass metabolism and the degree of blood serum binding. The efficacy of a given drug agent can be influenced by each of the parameters mentioned above.

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"Half-life" means the time where 50% of an administered drug are eliminated through biological processes, e.g. metabolism, excretion, etc.

By "hepatic first-pass metabolism" is meant the propensity of a drug to be metabolized upon first contact with the liver, i.e. during its first pass through the liver.

"Volume of distribution" means the degree of retention of a drug throughout the various compartments of the body, like e.g. intracellular and extracellular spaces, tissues and organs, etc. and the distribution of the drug within these compartments.

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"Degree of blood serum binding" means the propensity of a drug to interact with and bind to blood serum proteins, such as albumin, leading to a reduction or loss of biological activity of the drug.

Pharmacokinetic parameters also include bioavailability, lag time (Tlag), Tmax, absorption rates, more onset and/or Cmax for a given amount of drug administered. "Bioavailability" means the amount of a drug in the blood compartment. "Lag time" means the time delay between the administration of the drug and its detection and measurability in blood or plasma.

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"Tmax" is the time after which maximal blood concentration of the drug is reached, and "Cmax" is the blood concentration maximally obtained with a given drug. The time to reach a blood or tissue concentration of the drug which is required for its biological effect is

influenced by all parameters. Pharmacokinetic parameters of bispecific single chain antibodies exhibiting cross-species specificity, which may be determined in preclinical animal testing in non-chimpanzee primates as outlined above, are also set forth e.g. in the publication by Schlereth *et al.* (Cancer Immunol. Immunother. 20 (2005), 1-12).

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The term "toxicity" as used herein refers to the toxic effects of a drug manifested in adverse events or severe adverse events. These side events might refer to a lack of tolerability of the drug in general and/or a lack of local tolerance after administration. Toxicity could also include teratogenic or carcinogenic effects caused by the drug.

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The term "safety", "in vivo safety" or "tolerability" as used herein defines the administration of a drug without inducing severe adverse events directly after administration (local tolerance) and during a longer period of application of the drug. "Safety", "in vivo safety" or "tolerability" can be evaluated e.g. at regular intervals during the treatment and follow-up period. Measurements include clinical evaluation, e.g. organ manifestations, and screening of laboratory abnormalities. Clinical evaluation may be carried out and deviations to normal findings recorded/coded according to NCI-CTC and/or MedDRA standards. Organ manifestations may include criteria such as allergy/immunology, blood/bone marrow, cardiac arrhythmia, coagulation and the like, as set forth e.g. in the Common Terminology Criteria for adverse events v3.0 (CTCAE). Laboratory parameters which may be tested include for instance hematology, clinical chemistry, coagulation profile and urine analysis and examination of other body fluids such as serum, plasma, lymphoid or spinal fluid, liquor and the like. Safety can thus be assessed e.g. by physical examination, imaging techniques (i.e. ultrasound, x-ray, CT scans, Magnetic Resonance Imaging (MRI), other measures with technical devices (i.e. electrocardiogram), vital signs, by measuring laboratory parameters and recording adverse events. For example, adverse events in non-chimpanzee primates in the uses and methods according to the invention may be examined by histopathological and/or histochemical methods.

The term "effective dose" or "effective dosage" is defined as an amount sufficient to achieve or at least partially achieve the desired effect. The term "therapeutically effective dose" is defined as an amount sufficient to cure or at least partially arrest the disease and its complications in a patient already suffering from the disease. Amounts effective for this use will depend upon the severity of the infection and the general state of the subject's own immune system. The term "patient" includes human and other mammalian subjects that receive either prophylactic or therapeutic treatment.

The term "effective and non-toxic dose" as used herein refers to a tolerable dose of an inventive antibody construct which is high enough to cause depletion of pathologic cells, tumor elimination, tumor shrinkage or stabilization of disease without or essentially without major toxic effects. Such effective and non-toxic doses may be determined e.g. by dose escalation studies described in the art and should be below the dose inducing severe adverse side events (dose limiting toxicity, DLT).

The above terms are also referred to e.g. in the Preclinical safety evaluation of biotechnology-derived pharmaceuticals S6; ICH Harmonised Tripartite Guideline; ICH Steering Committee meeting on July 16, 1997.

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The appropriate dosage, or therapeutically effective amount, of the antibody construct of the invention will depend on the condition to be treated, the severity of the condition, prior therapy, and the patient's clinical history and response to the therapeutic agent. The proper dose can be adjusted according to the judgment of the attending physician such that it can be administered to the patient one time or over a series of administrations. The pharmaceutical composition can be administered as a sole therapeutic or in combination with additional therapies such as anti-cancer therapies as needed.

The pharmaceutical compositions of this invention are particularly useful for parenteral administration, i.e., subcutaneously, intramuscularly, intravenously, intra-articular and/or intra-synovial. Parenteral administration can be by bolus injection or continuous infusion.

If the pharmaceutical composition has been lyophilized, the lyophilized material is first reconstituted in an appropriate liquid prior to administration. The lyophilized material may be reconstituted in, e.g., bacteriostatic water for injection (BWFI), physiological saline, phosphate buffered saline (PBS), or the same formulation the protein had been in prior to lyophilization.

In an internal analysis of proprietary mRNA expression data it has been surprisingly found that CDH19 expression is elevated in both primary and metastatic melanoma tumors compared to normal, untransformed tissues. Internal analysis also confirmed that expression of CDH19 in normal tissues is limited to neural crest derived peripheral nerve ganglia and nerve fibers. The differential CDH19 expression in normal and tumor tissues makes this protein attractive for cell-surface targeting therapeutics. Although CDH 19 was discussed as one marker as part of long lists of markers associated with some cancer types (see e.g.

WO2009/055937) or Parkinson's disease (see e.g. WO2005/067391) CDH19 was never discussed as a prognostic marker or a drug target in connection with melanoma tumors.

As stated above, the present invention provides an isolated multispecific antibody construct comprising a first human binding domain capable of binding to human CDH19 on the surface of a target cell and a second domain capable of binding to human CD3 on the surface of a T cell.

The "CDH19 extracellular domain" or "CDH19 ECD" refers to a form of CDH19 which is essentially free of transmembrane and cytoplasmic domains of CDH19. It will be understood by the skilled artisan that the transmembrane domain identified for the CDH19 polypeptide of the present invention is identified pursuant to criteria routinely employed in the art for identifying that type of hydrophobic domain. The exact boundaries of a transmembrane domain may vary but most likely by no more than about 5 amino acids at either end of the domain specifically mentioned herein. A preferred human CDH19 ECD is shown in SEQ ID NO: 948. In this context it is understood that the CDH19 ECD represents the part of CDH19 on the surface of a target cell.

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The T cell CD3 receptor complex is a protein complex and is composed of four distinct chains. In mammals, the complex contains a CD3 γ chain, a CD3 δ chain, and two CD3 ϵ (epsilon) chains. These chains associate with a molecule known as the T cell receptor (TCR) and the ζ chain to generate an activation signal in T lymphocytes.

The redirected lysis of target cells via the recruitment of T cells by a multispecific, at least bispecific, antibody construct involves cytolytic synapse formation and delivery of perforin and granzymes. The engaged T cells are capable of serial target cell lysis, and are not affected by immune escape mechanisms interfering with peptide antigen processing and presentation, or clonal T cell differentiation; see, for example, WO 2007/042261.

The affinity of the first binding domain for human CDH19 is preferably ≤15 nM, more preferably ≤10 nM, even more preferably ≤5 nM, even more preferably ≤1 nM, even more preferably ≤0.5 nM, even more preferably ≤0.1 nM, and most preferably ≤0.05 nM. The affinity of the first binding domain for macaque CDH19 is preferably ≤15 nM, more preferably ≤10 nM, even more preferably ≤5 nM, even more preferably ≤1 nM, even more preferably ≤0.5 nM, even more preferably ≤0.5 nM or even ≤0.01 nM. The affinity can be measured for example in a Biacore assay or in a Scatchard assay, e.g. as described in the Examples. The affinity gap for binding to macaque CDH19 versus human

CDH19 is preferably [1:10-1:5] or [5:1-10:1], more preferably [1:5-5:1], and most preferably [1:2-3:1] or even [1:1-3:1]. Other methods of determining the affinity are well-known to the skilled person.

Human antibodies, respectively human antibody constructs, avoid some of the problems associated with antibodies/antibody constructs that possess murine or rat variable and/or constant regions. The presence of such murine or rat derived proteins can lead to the rapid clearance of the antibodies/antibody constructs or can lead to the generation of an immune response against the antibody/antibody construct by a patient. In order to avoid the utilization of murine or rat derived antibodies/antibody constructs, human or fully human antibodies can be generated through the introduction of human antibody function into a rodent so that the rodent produces fully human antibodies.

The ability to clone and reconstruct megabase-sized human loci in YACs and to introduce them into the mouse germline provides a powerful approach to elucidating the functional components of very large or crudely mapped loci as well as generating useful models of human disease. Furthermore, the utilization of such technology for substitution of mouse loci with their human equivalents could provide unique insights into the expression and regulation of human gene products during development, their communication with other systems, and their involvement in disease induction and progression.

An important practical application of such a strategy is the "humanization" of the mouse humoral immune system. Introduction of human immunoglobulin (Ig) loci into mice in which the endogenous Ig genes have been inactivated offers the opportunity to study the mechanisms underlying programmed expression and assembly of antibodies as well as their role in B-cell development. Furthermore, such a strategy could provide an ideal source for production of fully human monoclonal antibodies (mAbs)—an important milestone towards fulfilling the promise of antibody therapy in human disease. Fully human antibodies/antibody constructs are expected to minimize the immunogenic and allergic responses intrinsic to mouse or mouse-derivatized mAbs and thus to increase the efficacy and safety of the administered antibodies/antibody constructs. The use of fully human antibodies/antibody constructs can be expected to provide a substantial advantage in the treatment of chronic and recurring human diseases, such as inflammation, autoimmunity, and cancer, which require repeated compound administrations.

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One approach towards this goal was to engineer mouse strains deficient in mouse antibody production with large fragments of the human Ig loci in anticipation that such mice would

produce a large repertoire of human antibodies in the absence of mouse antibodies. Large human Ig fragments would preserve the large variable gene diversity as well as the proper regulation of antibody production and expression. By exploiting the mouse machinery for antibody diversification and selection and the lack of immunological tolerance to human proteins, the reproduced human antibody repertoire in these mouse strains should yield high affinity antibodies against any antigen of interest, including human antigens. Using the hybridoma technology, antigen-specific human mAbs with the desired specificity could be readily produced and selected. This general strategy was demonstrated in connection with our generation of the first XenoMouse mouse strains, as published in 1994. (See Green et al. Nature Genetics 7:13-21 (1994)) The XenoMouse strains were engineered with yeast artificial chromosomes (YACs) containing 245 kb and 190 kb-sized germline configuration fragments of the human heavy chain locus and kappa light chain locus, respectively, which contained core variable and constant region sequences. Id. The human Ig containing YACs proved to be compatible with the mouse system for both rearrangement and expression of antibodies and were capable of substituting for the inactivated mouse Ig genes. This was demonstrated by their ability to induce B-cell development, to produce an adult-like human repertoire of fully human antibodies, and to generate antigen-specific human mAbs. These results also suggested that introduction of larger portions of the human Ig loci containing greater numbers of V genes, additional regulatory elements, and human Ig constant regions might recapitulate substantially the full repertoire that is characteristic of the human humoral response to infection and immunization. The work of Green et al. was recently extended to the introduction of greater than approximately 80% of the human antibody repertoire through introduction of megabase sized, germline configuration YAC fragments of the human heavy chain loci and kappa light chain loci, respectively. See Mendez et al. Nature Genetics 15:146-156 (1997) and U.S. patent application Ser. No. 08/759,620, filed Dec. 3, 1996, the disclosures of which are hereby incorporated by reference.

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The production of the XenoMouse mice is further discussed and delineated in U.S. patent application Ser. No. 07/466,008, filed Jan. 12, 1990, Ser. No. 07/610,515, filed Nov. 8, 1990, Ser. No. 07/919,297, filed Jul. 24, 1992, Ser. No. 07/922,649, filed Jul. 30, 1992, filed Ser. No. 08/031,801, filed Mar. 15, 1993, Ser. No. 08/112,848, filed Aug. 27, 1993, Ser. No. 08/234,145, filed Apr. 28, 1994, Ser. No. 08/376,279, filed Jan. 20, 1995, Ser. No. 08/430,938, Apr. 27, 1995, Ser. No. 08/464,584, filed Jun. 5, 1995, Ser. No. 08/464,582, filed Jun. 5, 1995, Ser. No. 08/463,191, filed Jun. 5, 1995, Ser. No. 08/462,837, filed Jun. 5, 1995, Ser. No. 08/486,853, filed Jun. 5, 1995, Ser. No. 08/486,857, filed Jun. 5, 1995, Ser. No. 08/486,859, filed Jun. 5, 1995, Ser. No. 08/462,513, filed Jun. 5, 1995, Ser. No. 08/724,752, filed Oct. 2, 1996, and Ser. No. 08/759,620, filed Dec. 3, 1996 and U.S. Pat.

Nos. 6,162,963, 6,150,584, 6,114,598, 6,075,181, and 5,939,598 and Japanese Patent Nos. 3 068 180 B2, 3 068 506 B2, and 3 068 507 B2. See also Mendez et al. Nature Genetics 15:146-156 (1997) and Green and Jakobovits J. Exp. Med. 188:483-495 (1998). See also European Patent No., EP 0 463151 B1, grant published Jun. 12, 1996, International Patent Application No., WO 94/02602, published Feb. 3, 1994, International Patent Application No., WO 96/34096, published Oct. 31, 1996, WO 98/24893, published Jun. 11, 1998, WO 00/76310, published Dec. 21, 2000, WO 03/47336. The disclosures of each of the above-cited patents, applications, and references are hereby incorporated by reference in their entirety.

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In an alternative approach, others, including GenPharm International, Inc., have utilized a "minilocus" approach. In the minilocus approach, an exogenous Ig locus is mimicked through the inclusion of pieces (individual genes) from the Ig locus. Thus, one or more V.sub.H genes, one or more D.sub.H genes, one or more J.sub.H genes, a mu constant region, and a second constant region (preferably a gamma constant region) are formed into a construct for insertion into an animal. This approach is described in U.S. Pat. No. 5,545,807 to Surani et al. and U.S. Pat. Nos. 5,545,806, 5,625,825, 5,625,126, 5,633,425, 5,661,016, 5,770,429, 5,789,650, 5,814,318, 5,877,397, 5,874,299, and 6,255,458 each to Lonberg and Kay, U.S. Pat. Nos. 5,591,669 and 6,023.010 to Krimpenfort and Berns, U.S. Pat. Nos. 5,612,205, 5,721,367, and 5,789,215 to Berns et al., and U.S. Pat. No. 5,643,763 to Choi and Dunn, and GenPharm International U.S. patent application Ser. No. 07/574,748, filed Aug. 29, 1990, Ser. No. 07/575,962, filed Aug. 31, 1990, Ser. No. 07/810,279, filed Dec. 17, 1991, Ser. No. 07/853,408, filed Mar. 18, 1992, Ser. No. 07/904,068, filed Jun. 23, 1992, Ser. No. 07/990,860, filed Dec. 16, 1992, Ser. No. 08/053,131, filed Apr. 26, 1993, Ser. No. 08/096,762, filed Jul. 22, 1993, Ser. No. 08/155,301, filed Nov. 18, 1993, Ser. No. 08/161,739, filed Dec. 3, 1993, Ser. No. 08/165,699, filed Dec. 10, 1993, Ser. No. 08/209,741, filed Mar. 9, 1994, the disclosures of which are hereby incorporated by reference. See also European Patent No. 0 546 073 B 1, International Patent Application Nos. WO 92/03918, WO 92/22645, WO 92/22647, WO 92/22670, WO 93/12227, WO 94/00569, WO 94/25585, WO 96/14436, WO 97/13852, and WO 98/24884 and U.S. Pat. No. 5,981,175, the disclosures of which are hereby incorporated by reference in their entirety. See further Taylor et al., 1992, Chen et al., 1993, Tuaillon et al., 1993, Choi et al., 1993, Lonberg et al., (1994), Taylor et al., (1994), and Tuaillon et al., (1995), Fishwild et al., (1996), the disclosures of which are hereby incorporated by reference in their entirety.

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Kirin has also demonstrated the generation of human antibodies from mice in which, through microcell fusion, large pieces of chromosomes, or entire chromosomes, have been

introduced. See European Patent Application Nos. 773 288 and 843 961, the disclosures of which are hereby incorporated by reference. Xenerex Biosciences is developing a technology for the potential generation of human antibodies. In this technology, SCID mice are reconstituted with human lymphatic cells, e.g., B and/or T cells. Mice are then immunized with an antigen and can generate an immune response against the antigen. See U.S. Pat. Nos. 5,476,996, 5,698,767, and 5,958,765.

Human anti-mouse antibody (HAMA) responses have led the industry to prepare chimeric or otherwise humanized antibodies. While chimeric antibodies have a human constant region and a murine variable region, it is expected that certain human anti-chimeric antibody (HACA) responses will be observed, particularly in chronic or multi-dose utilizations of the antibody. Thus, it would be desirable to provide fully human antibodies against EGFRvIII in order to vitiate concerns and/or effects of HAMA or HACA response.

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Cytotoxicity mediated by CDH19/CD3 bispecific antibody constructs can be measured in various ways. Effector cells can be e.g. stimulated enriched (human) CD8 positive T cells or unstimulated (human) peripheral blood mononuclear cells (PBMC). If the target cells are of macaque origin or express or are transfected with macaque CDH19, the effector cells should also be of macaque origin such as a macaque T cell line, e.g. 4119LnPx. The target cells should express (at least the extracellular domain of) CDH19, e.g. human or macaque CDH19. Target cells can be a cell line (such as CHO) which is stably or transiently transfected with CDH19, e.g. human or macaque CDH19. Alternatively, the target cells can be a CDH19 positive natural expresser cell line, such as the human myeloma cell line CHL-1 or Colo-699. Usually EC50-values are expected to be lower with target cell lines expressing higher levels of CDH19on the cell surface. The effector to target cell (E:T) ratio is usually about 10:1, but can also vary. Cytotoxic activity of CDH19/CD3 bispecific antibody constructs can be measured in an 51-chromium release assay (incubation time of about 18 hours) or in a in a FACS-based cytotoxicity assay (incubation time of about 48 hours). Modifications of the assay incubation time (cytotoxic reaction) are also possible. Other methods of measuring cytotoxicity are well-known to the skilled person and comprise MTT or MTS assays, ATP-based assays including bioluminescent assays, the sulforhodamine B (SRB) assay, WST assay, clonogenic assay and the ECIS technology.

The cytotoxic activity mediated by CDH19/CD3 bispecific antibody constructs of the present invention is preferably measured in a cell-based cytotoxicity assay. It is represented by the EC₅₀ value, which corresponds to the half maximal effective concentration (concentration of the antibody construct which induces a cytotoxic response halfway between the baseline

and maximum). Preferably, the EC₅₀ value of the CDH19/CD3 bispecific antibody constructs is \leq 20.000 pg/ml, more preferably \leq 5000 pg/ml, even more preferably \leq 1000 pg/ml, even more preferably \leq 350 pg/ml, even more preferably \leq 350 pg/ml, even more preferably \leq 320 pg/ml, even more preferably \leq 250 pg/ml, even more preferably \leq 100 pg/ml, even more preferably \leq 50 pg/ml, even more preferably \leq 50 pg/ml, and most preferably \leq 5 pg/ml.

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Any of the above given EC₅₀ values can be combined with any one of the indicated scenarios of a cell-based cytotoxicity assay. For example, when (human) CD8 positive T cells or a macaque T cell line are used as effector cells, the EC₅₀ value of the CDH19/CD3 bispecific antibody construct is preferably \leq 1000 pg/ml, more preferably \leq 500 pg/ml, even more preferably \leq 500 pg/ml, even more preferably \leq 100 pg/ml, even more preferably \leq 50 pg/ml, lf in this assay the target cells are (human or macaque) CDH19 transfected cells such as CHO cells, the EC₅₀ value of the CDH19/CD3 bispecific antibody construct is preferably \leq 150 pg/ml, more preferably \leq 100 pg/ml, even more preferably \leq 50 pg/ml.

If the target cells are a CDH19positive natural expresser cell line, then the EC₅₀ value is preferably \leq 350 pg/ml, more preferably \leq 320 pg/ml, even more preferably \leq 250 pg/ml, even more preferably \leq 100 pg/ml, even more preferably \leq 100 pg/ml, even more preferably \leq 100 pg/ml, and most preferably \leq 50 pg/ml, or lower.

When (human) PBMCs are used as effector cells, the EC₅₀ value of the CDH19/CD3 bispecific antibody construct is preferably \leq 1000 pg/ml, more preferably \leq 750 pg/ml, more preferably \leq 500 pg/ml, even more preferably \leq 350 pg/ml, even more preferably \leq 320 pg/ml, even more preferably \leq 100 pg/ml, and most preferably \leq 50 pg/ml, or lower.

The difference in cytotoxic activity between the monomeric and the dimeric isoform of individual CDH19/CD3 bispecific antibody constructs is referred to as "potency gap". This potency gap can e.g. be calculated as ratio between EC₅₀ values of the molecule's monomeric and dimeric form. Potency gaps of the CDH19/CD3 bispecific antibody constructs of the present invention are preferably \leq 5, more preferably \leq 4, even more preferably \leq 3, even more preferably \leq 2 and most preferably \leq 1.

The antibody construct of the invention is a fusion protein comprising at least two binding domains, with or without peptide linkers (spacer peptides). Among the suitable peptide linkers are those described in U.S. Patents 4,751,180 and 4,935,233 or WO 88/09344.

Another method for preparing oligomeric antibody constuct derivatives involves use of a leucine zipper. Leucine zipper domains are peptides that promote oligomerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding proteins (Landschulz *et al.*, 1988, *Science* 240:1759), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize. Examples of leucine zipper domains suitable for producing soluble oligomeric proteins are described in PCT application WO 94/10308, and the leucine zipper derived from lung surfactant protein D (SPD) described in Hoppe *et al.*, 1994, *FEBS Letters* 344:191, hereby incorporated by reference. The use of a modified leucine zipper that allows for stable trimerization of a heterologous protein fused thereto is described in Fanslow *et al.*, 1994, *Semin. Immunol.* 6:267-78. In one approach, recombinant fusion proteins comprising CDH19 antibody fragment or derivative fused to a leucine zipper peptide are expressed in suitable host cells, and the soluble oligomeric CDH19 antibody fragments or derivatives that form are recovered from the culture supernatant.

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Covalent modifications of antigen binding proteins are included within the scope of this invention, and are generally, but not always, done post-translationally. For example, several types of covalent modifications of the antigen binding protein are introduced into the molecule by reacting specific amino acid residues of the antigen binding protein with an organic derivatizing agent that is capable of reacting with selected side chains or the N- or C-terminal residues.

Cysteinyl residues most commonly are reacted with α -haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteinyl residues also are derivatized by reaction with bromotrifluoroacetone, α -bromo- β -(5-imidozoyl)propionic acid, chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloromercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

Histidyl residues are derivatized by reaction with diethylpyrocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Para-bromophenacyl bromide also is useful; the reaction is preferably performed in 0.1M sodium cacodylate at pH 6.0.

35 Lysinyl and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysinyl residues. Other suitable reagents for derivatizing alpha-amino-containing residues

include imidoesters such as methyl picolinimidate; pyridoxal phosphate; pyridoxal; chloroborohydride; trinitrobenzenesulfonic acid; O-methylisourea; 2,4-pentanedione; and transaminase-catalyzed reaction with glyoxylate.

Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pKa of the guanidine functional group. Furthermore, these reagents may react with the groups of lysine as well as the arginine epsilon-amino group.

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The specific modification of tyrosyl residues may be made, with particular interest in introducing spectral labels into tyrosyl residues by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidizole and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively. Tyrosyl residues are iodinated using ¹²⁵I or ¹³¹I to prepare labeled proteins for use in radioimmunoassay, the chloramine T method described above being suitable.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction with carbodiimides (R'—N=C=N--R'), where R and R' are optionally different alkyl groups, such as 1-cyclohexyl-3-(2-morpholinyl-4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginyl and glutaminyl residues by reaction with ammonium ions.

Derivatization with bifunctional agents is useful for crosslinking antigen binding proteins to a water-insoluble support matrix or surface for use in a variety of methods. Commonly used crosslinking agents include, e.g., 1,1-bis(diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, for example, esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), and bifunctional maleimides such as bis-N-maleimido-1,8-octane. Derivatizing agents such as methyl-3-[(p-azidophenyl)dithio]propioimidate yield photoactivatable intermediates that are capable of forming crosslinks in the presence of light. Alternatively, reactive water-insoluble matrices such as cyanogen bromide-activated carbohydrates and the reactive substrates described in U.S. Pat. Nos. 3,969,287; 3,691,016; 4,195,128; 4,247,642;

4,229,537; and 4,330,440 are employed for protein immobilization.

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Glutaminyl and asparaginyl residues are frequently deamidated to the corresponding glutamyl and aspartyl residues, respectively. Alternatively, these residues are deamidated

under mildly acidic conditions. Either form of these residues falls within the scope of this invention.

Other modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the α-amino groups of lysine, arginine, and histidine side chains (T. E. Creighton, Proteins: Structure and Molecular Properties, W. H. Freeman & Co., San Francisco, 1983, pp. 79-86), acetylation of the N-terminal amine, and amidation of any C-terminal carboxyl group.

Another type of covalent modification of the antigen binding protein included within the scope of this invention comprises altering the glycosylation pattern of the protein. As is known in the art, glycosylation patterns can depend on both the sequence of the protein (e.g., the presence or absence of particular glycosylation amino acid residues, discussed below), or the host cell or organism in which the protein is produced. Particular expression systems are discussed below.

Glycosylation of polypeptides is typically either N-linked or O-linked. N-linked refers to the attachment of the carbohydrate moiety to the side chain of an asparagine residue. The tripeptide sequences asparagine-X-serine and asparagine-X-threonine, where X is any amino acid except proline, are the recognition sequences for enzymatic attachment of the carbohydrate moiety to the asparagine side chain. Thus, the presence of either of these tripeptide sequences in a polypeptide creates a potential glycosylation site. O-linked glycosylation refers to the attachment of one of the sugars N-acetylgalactosamine, galactose, or xylose, to a hydroxyamino acid, most commonly serine or threonine, although 5-hydroxyproline or 5-hydroxylysine may also be used.

Addition of glycosylation sites to the antigen binding protein is conveniently accomplished by altering the amino acid sequence such that it contains one or more of the above-described tri-peptide sequences (for N-linked glycosylation sites). The alteration may also be made by the addition of, or substitution by, one or more serine or threonine residues to the starting sequence (for O-linked glycosylation sites). For ease, the antigen binding protein amino acid sequence is preferably altered through changes at the DNA level, particularly by mutating the DNA encoding the target polypeptide at preselected bases such that codons are generated that will translate into the desired amino acids.

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Another means of increasing the number of carbohydrate moieties on the antigen binding protein is by chemical or enzymatic coupling of glycosides to the protein. These procedures

are advantageous in that they do not require production of the protein in a host cell that has glycosylation capabilities for N- and O-linked glycosylation. Depending on the coupling mode used, the sugar(s) may be attached to (a) arginine and histidine, (b) free carboxyl groups, (c) free sulfhydryl groups such as those of cysteine, (d) free hydroxyl groups such as those of serine, threonine, or hydroxyproline, (e) aromatic residues such as those of phenylalanine, tyrosine, or tryptophan, or (f) the amide group of glutamine. These methods are described in WO 87/05330 published Sep. 11, 1987, and in Aplin and Wriston, 1981, *CRC Crit. Rev. Biochem.*, pp. 259-306.

Removal of carbohydrate moieties present on the starting antigen binding protein may be accomplished chemically or enzymatically. Chemical deglycosylation requires exposure of the protein to the compound trifluoromethanesulfonic acid, or an equivalent compound. This treatment results in the cleavage of most or all sugars except the linking sugar (N-acetylglucosamine or N-acetylgalactosamine), while leaving the polypeptide intact. Chemical deglycosylation is described by Hakimuddin *et al.*, 1987, *Arch. Biochem. Biophys.* 259:52 and by Edge *et al.*, 1981, *Anal. Biochem.* 118:131. Enzymatic cleavage of carbohydrate moieties on polypeptides can be achieved by the use of a variety of endo- and exoglycosidases as described by Thotakura *et al.*, 1987, *Meth. Enzymol.* 138:350. Glycosylation at potential glycosylation sites may be prevented by the use of the compound tunicamycin as described by Duskin *et al.*, 1982, *J. Biol. Chem.* 257:3105. Tunicamycin blocks the formation of protein-N-glycoside linkages.

Another type of covalent modification of the antigen binding protein comprises linking the antigen binding protein to various non-proteinaceous polymers, including, but not limited to, various polyols such as polyethylene glycol, polypropylene glycol or polyoxyalkylenes, in the manner set forth in U.S. Pat. Nos. 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192 or 4,179,337. In addition, as is known in the art, amino acid substitutions may be made in various positions within the antigen binding protein to facilitate the addition of polymers such as PEG.

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In some embodiments, the covalent modification of the antigen binding proteins of the invention comprises the addition of one or more labels.

The term "labelling group" means any detectable label. Examples of suitable labelling groups include, but are not limited to, the following: radioisotopes or radionuclides (e.g., ³H, ¹⁴C, ¹⁵N, ³⁵S, ⁸⁹Zr, ⁹⁰Y, ⁹⁹Tc, ¹¹¹In, ¹²⁵I, ¹³¹I), fluorescent groups (e.g., FITC, rhodamine, lanthanide phosphors), enzymatic groups (e.g., horseradish peroxidase, β-galactosidase, luciferase,

alkaline phosphatase), chemiluminescent groups, biotinyl groups, or predetermined polypeptide epitopes recognized by a secondary reporter (e.g., leucine zipper pair sequences, binding sites for secondary antibodies, metal binding domains, epitope tags). In some embodiments, the labelling group is coupled to the antigen binding protein *via* spacer arms of various lengths to reduce potential steric hindrance. Various methods for labelling proteins are known in the art and may be used in performing the present invention.

In general, labels fall into a variety of classes, depending on the assay in which they are to be detected: a) isotopic labels, which may be radioactive or heavy isotopes; b) magnetic labels (e.g., magnetic particles); c) redox active moieties; d) optical dyes; enzymatic groups (e.g. horseradish peroxidase, β-galactosidase, luciferase, alkaline phosphatase); e) biotinylated groups; and f) predetermined polypeptide epitopes recognized by a secondary reporter (e.g., leucine zipper pair sequences, binding sites for secondary antibodies, metal binding domains, epitope tags, etc.). In some embodiments, the labelling group is coupled to the antigen binding protein *via* spacer arms of various lengths to reduce potential steric hindrance. Various methods for labelling proteins are known in the art and may be used in performing the present invention.

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Specific labels include optical dyes, including, but not limited to, chromophores, phosphors and fluorophores, with the latter being specific in many instances. Fluorophores can be either "small molecule" fluores, or proteinaceous fluores.

By "fluorescent label" is meant any molecule that may be detected *via* its inherent fluorescent properties. Suitable fluorescent labels include, but are not limited to, fluorescein, rhodamine, tetramethylrhodamine, eosin, erythrosin, coumarin, methyl-coumarins, pyrene, Malacite green, stilbene, Lucifer Yellow, Cascade BlueJ, Texas Red, IAEDANS, EDANS, BODIPY FL, LC Red 640, Cy 5, Cy 5.5, LC Red 705, Oregon green, the Alexa-Fluor dyes (Alexa Fluor 350, Alexa Fluor 430, Alexa Fluor 488, Alexa Fluor 546, Alexa Fluor 568, Alexa Fluor 594, Alexa Fluor 633, Alexa Fluor 660, Alexa Fluor 680), Cascade Blue, Cascade Yellow and R-phycoerythrin (PE) (Molecular Probes, Eugene, OR), FITC, Rhodamine, and Texas Red (Pierce, Rockford, IL), Cy5, Cy5.5, Cy7 (Amersham Life Science, Pittsburgh, PA). Suitable optical dyes, including fluorophores, are described in Molecular Probes Handbook by Richard P. Haugland, hereby expressly incorporated by reference.

35 Suitable proteinaceous fluorescent labels also include, but are not limited to, green fluorescent protein, including a Renilla, Ptilosarcus, or Aequorea species of GFP (Chalfie *et al.*, 1994, *Science* 263:802-805), EGFP (Clontech Laboratories, Inc., Genbank Accession

Number U55762), blue fluorescent protein (BFP, Quantum Biotechnologies, Inc. 1801 de Maisonneuve Blvd. West, 8th Floor, Montreal, Quebec, Canada H3H 1J9; Stauber, 1998, *Biotechniques* 24:462-471; Heim *et al.*, 1996, *Curr. Biol.* 6:178-182), enhanced yellow fluorescent protein (EYFP, Clontech Laboratories, Inc.), luciferase (Ichiki *et al.*, 1993, *J. Immunol.* 150:5408-5417), β galactosidase (Nolan *et al.*, 1988, *Proc. Natl. Acad. Sci. U.S.A.* 85:2603-2607) and Renilla (WO92/15673, WO95/07463, WO98/14605, WO98/26277, WO99/49019, U.S. Patent Nos. 5292658, 5418155, 5683888, 5741668, 5777079, 5804387, 5874304, 5876995, 5925558). All of the above-cited references are expressly incorporated herein by reference.

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The antibody construct of the invention may also comprise additional domains, which e.g. are helpful in the isolation of the molecule or relate to an adapted pharmacokinetic profile of the molecule.

Domains helpful for the isolation of an antibody construct may be elected from peptide motives or secondarily introduced moieties, which can be captured in an isolation method, e.g. an isolation column. A non-limiting embodiments of such additional domains comprise peptide motives known as Myc-tag, HAT-tag, HA-tag, TAP-tag, GST-tag, chitin binding domain (CBD-tag), maltose binding protein (MBP-tag), Flag-tag, Strep-tag and variants thereof (e.g. StrepII-tag) and His-tag. All herein disclosed antibody constructs characterized by the identified CDRs are preferred to comprise a His-tag domain, which is generally known as a repeat of consecutive His residues in the amino acid sequence of a molecule, preferably of six His residues.

As described in appended example 2 a broad number of CDH19 specific binder has been characterized with respect to identified binding characteristics and those binders were grouped into five different bins, which refers to five different subgroups of CDH19 specific binding domains. Accordingly, in one embodiment the antibody construct of the invention the first binding domain comprises a VH region comprising CDR-H1, CDR-H2 and CDR-H3 and a VL region comprising CDR-L1, CDR-L2 and CDR-L3 selected from the group consisting of:

(a) CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 222, CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 252, CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-

H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927, CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 909, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927, CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926,

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CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 904, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926,

CDR-H1 as depicted in SEQ ID NO: 1126, CDR-H2 as depicted in SEQ ID NO: 1127, CDR-H3 as depicted in SEQ ID NO: 1128, CDR-L1 as depicted in SEQ ID NO: 1129, CDR-L2 as depicted in SEQ ID NO: 1130 and CDR-L3 as depicted in SEQ ID NO: 1131,

CDR-H1 as depicted in SEQ ID NO: 1165, CDR-H2 as depicted in SEQ ID NO: 1166, CDR-H3 as depicted in SEQ ID NO: 1167, CDR-L1 as depicted in SEQ ID NO: 1168, CDR-L2 as depicted in SEQ ID NO: 1169 and CDR-L3 as depicted in SEQ ID NO: 1170,

CDR-H1 as depicted in SEQ ID NO: 1334, CDR-H2 as depicted in SEQ ID NO: 1335, CDR-H3 as depicted in SEQ ID NO: 1336, CDR-L1 as depicted in SEQ ID NO: 1337, CDR-L2 as depicted in SEQ ID NO: 1338 and CDR-L3 as depicted in SEQ ID NO: 1339,

CDR-H1 as depicted in SEQ ID NO: 1347, CDR-H2 as depicted in SEQ ID NO: 1348, CDR-H3 as depicted in SEQ ID NO: 1349, CDR-L1 as depicted in SEQ ID NO: 1350, CDR-L2 as depicted in SEQ ID NO: 1351 and CDR-L3 as depicted in SEQ ID NO: 1352, and

CDR-H1 as depicted in SEQ ID NO: 1360 CDR-H2 as depicted in SEQ ID NO: 1361, CDR-H3 as depicted in SEQ ID NO: 1362, CDR-L1 as depicted in SEQ ID NO: 1363, CDR-L2 as depicted in SEQ ID NO: 1364 and CDR-L3 as depicted in SEQ ID NO: 1365,

CDR-H1 as depicted in SEQ ID NO: 1425 CDR-H2 as depicted in SEQ ID NO: 1426, CDR-H3 as depicted in SEQ ID NO: 1427, CDR-L1 as depicted in SEQ ID NO: 1428, CDR-L2 as depicted in SEQ ID NO: 1429 and CDR-L3 as depicted in SEQ ID NO: 1430.

CDR-H1 as depicted in SEQ ID NO: 1438 CDR-H2 as depicted in SEQ ID NO: 1439, CDR-H3 as depicted in SEQ ID NO: 1440, CDR-L1 as depicted in SEQ ID NO: 1441,

CDR-L2 as depicted in SEQ ID NO: 1442 and CDR-L3 as depicted in SEQ ID NO: 1443, and

CDR-H1 as depicted in SEQ ID NO: 2167 CDR-H2 as depicted in SEQ ID NO: 2168, CDR-H3 as depicted in SEQ ID NO: 2169, CDR-L1 as depicted in SEQ ID NO: 2170, CDR-L2 as depicted in SEQ ID NO: 2171 and CDR-L3 as depicted in SEQ ID NO: 2172,

which all characterize binding domains for CDH19 grouped into bin 1;

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CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, (b) CDR-H3 as depicted in SEQ ID NO: 126, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294, 10 CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 300, CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, 15 CDR-H3 as depicted in SEQ ID NO: 138, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 306, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 312, 20 CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336, 25 CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294, CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, 30 CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 928, CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 929, 35 CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336,

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CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 942, CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 943, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 937, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938, CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 919, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935, CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 936, CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 138, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 933, CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 917, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 934, CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 930, CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131,

CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 931, CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 932, CDR-H1 as depicted in SEQ ID NO: 1009, CDR-H2 as depicted in SEQ ID NO: 1010, CDR-H3 as depicted in SEQ ID NO: 1011, CDR-L1 as depicted in SEQ ID NO: 1012, CDR-L2 as depicted in SEQ ID NO: 1013 and CDR-L3 as depicted in SEQ ID NO: 1014,

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10 CDR-H1 as depicted in SEQ ID NO: 1022, CDR-H2 as depicted in SEQ ID NO: 1023, CDR-H3 as depicted in SEQ ID NO: 1024, CDR-L1 as depicted in SEQ ID NO: 1025, CDR-L2 as depicted in SEQ ID NO: 1026 and CDR-L3 as depicted in SEQ ID NO: 1027,

CDR-H1 as depicted in SEQ ID NO: 1035, CDR-H2 as depicted in SEQ ID NO: 1036, CDR-H3 as depicted in SEQ ID NO: 1037, CDR-L1 as depicted in SEQ ID NO: 1038, CDR-L2 as depicted in SEQ ID NO: 1039 and CDR-L3 as depicted in SEQ ID NO: 1040.

CDR-H1 as depicted in SEQ ID NO: 1074, CDR-H2 as depicted in SEQ ID NO: 1075, CDR-H3 as depicted in SEQ ID NO: 1076, CDR-L1 as depicted in SEQ ID NO: 1077, CDR-L2 as depicted in SEQ ID NO: 1078 and CDR-L3 as depicted in SEQ ID NO: 1079,

CDR-H1 as depicted in SEQ ID NO: 1100, CDR-H2 as depicted in SEQ ID NO: 1101, CDR-H3 as depicted in SEQ ID NO: 1102, CDR-L1 as depicted in SEQ ID NO: 1103, CDR-L2 as depicted in SEQ ID NO: 1104 and CDR-L3 as depicted in SEQ ID NO: 1105,

CDR-H1 as depicted in SEQ ID NO: 1113, CDR-H2 as depicted in SEQ ID NO: 1114, CDR-H3 as depicted in SEQ ID NO: 1115, CDR-L1 as depicted in SEQ ID NO: 1116, CDR-L2 as depicted in SEQ ID NO: 1117 and CDR-L3 as depicted in SEQ ID NO: 1118.

CDR-H1 as depicted in SEQ ID NO: 1243, CDR-H2 as depicted in SEQ ID NO: 1244, CDR-H3 as depicted in SEQ ID NO: 1245, CDR-L1 as depicted in SEQ ID NO: 1246, CDR-L2 as depicted in SEQ ID NO: 1247 and CDR-L3 as depicted in SEQ ID NO: 1248,

CDR-H1 as depicted in SEQ ID NO: 1256, CDR-H2 as depicted in SEQ ID NO: 1257, CDR-H3 as depicted in SEQ ID NO: 1258, CDR-L1 as depicted in SEQ ID NO: 1259, CDR-L2 as depicted in SEQ ID NO: 1260 and CDR-L3 as depicted in SEQ ID NO: 1261,

CDR-H1 as depicted in SEQ ID NO: 1269, CDR-H2 as depicted in SEQ ID NO: 1270, CDR-H3 as depicted in SEQ ID NO: 1271, CDR-L1 as depicted in SEQ ID NO: 1272, CDR-L2 as depicted in SEQ ID NO: 1273 and CDR-L3 as depicted in SEQ ID NO: 1274,

CDR-H1 as depicted in SEQ ID NO: 1282, CDR-H2 as depicted in SEQ ID NO: 1283, CDR-H3 as depicted in SEQ ID NO: 1284, CDR-L1 as depicted in SEQ ID NO: 1285, CDR-L2 as depicted in SEQ ID NO: 1286 and CDR-L3 as depicted in SEQ ID NO: 1287, and

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CDR-H1 as depicted in SEQ ID NO: 1295, CDR-H2 as depicted in SEQ ID NO: 1296, CDR-H3 as depicted in SEQ ID NO: 1297, CDR-L1 as depicted in SEQ ID NO: 1298, CDR-L2 as depicted in SEQ ID NO: 1299 and CDR-L3 as depicted in SEQ ID NO: 1300,

CDR-H1 as depicted in SEQ ID NO: 1647, CDR-H2 as depicted in SEQ ID NO: 1648, CDR-H3 as depicted in SEQ ID NO: 1649, CDR-L1 as depicted in SEQ ID NO: 1650, CDR-L2 as depicted in SEQ ID NO: 1651 and CDR-L3 as depicted in SEQ ID NO: 1652.

CDR-H1 as depicted in SEQ ID NO: 1660, CDR-H2 as depicted in SEQ ID NO: 1661, CDR-H3 as depicted in SEQ ID NO: 1662, CDR-L1 as depicted in SEQ ID NO: 1663, CDR-L2 as depicted in SEQ ID NO: 1664 and CDR-L3 as depicted in SEQ ID NO: 1665.

CDR-H1 as depicted in SEQ ID NO: 1894, CDR-H2 as depicted in SEQ ID NO: 1895, CDR-H3 as depicted in SEQ ID NO: 1896, CDR-L1 as depicted in SEQ ID NO: 1897, CDR-L2 as depicted in SEQ ID NO: 1898 and CDR-L3 as depicted in SEQ ID NO: 1899,

CDR-H1 as depicted in SEQ ID NO: 1907, CDR-H2 as depicted in SEQ ID NO: 1908, CDR-H3 as depicted in SEQ ID NO: 1909, CDR-L1 as depicted in SEQ ID NO: 1910, CDR-L2 as depicted in SEQ ID NO: 1911 and CDR-L3 as depicted in SEQ ID NO: 1912,

CDR-H1 as depicted in SEQ ID NO: 1933, CDR-H2 as depicted in SEQ ID NO: 1934, CDR-H3 as depicted in SEQ ID NO: 1935, CDR-L1 as depicted in SEQ ID NO: 1936, CDR-L2 as depicted in SEQ ID NO: 1937 and CDR-L3 as depicted in SEQ ID NO: 1938,

CDR-H1 as depicted in SEQ ID NO: 1946, CDR-H2 as depicted in SEQ ID NO: 1947, CDR-H3 as depicted in SEQ ID NO: 1948, CDR-L1 as depicted in SEQ ID NO: 1949, CDR-L2 as depicted in SEQ ID NO: 1950 and CDR-L3 as depicted in SEQ ID NO: 1951,

CDR-H1 as depicted in SEQ ID NO: 1959, CDR-H2 as depicted in SEQ ID NO: 1960,

CDR-H3 as depicted in SEQ ID NO: 1961, CDR-L1 as depicted in SEQ ID NO: 1962, CDR-L2 as depicted in SEQ ID NO: 1963 and CDR-L3 as depicted in SEQ ID NO: 1964,

CDR-H1 as depicted in SEQ ID NO: 1972, CDR-H2 as depicted in SEQ ID NO: 1973, CDR-H3 as depicted in SEQ ID NO: 1974, CDR-L1 as depicted in SEQ ID NO: 1975, CDR-L2 as depicted in SEQ ID NO: 1976 and CDR-L3 as depicted in SEQ ID NO: 1977,

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CDR-H1 as depicted in SEQ ID NO: 1985, CDR-H2 as depicted in SEQ ID NO: 1986, CDR-H3 as depicted in SEQ ID NO: 1987, CDR-L1 as depicted in SEQ ID NO: 1988, CDR-L2 as depicted in SEQ ID NO: 1989 and CDR-L3 as depicted in SEQ ID NO: 1990.

CDR-H1 as depicted in SEQ ID NO: 1998, CDR-H2 as depicted in SEQ ID NO: 1999, CDR-H3 as depicted in SEQ ID NO: 2000, CDR-L1 as depicted in SEQ ID NO: 2001, CDR-L2 as depicted in SEQ ID NO: 2002 and CDR-L3 as depicted in SEQ ID NO: 2003.

CDR-H1 as depicted in SEQ ID NO: 2011, CDR-H2 as depicted in SEQ ID NO: 2012, CDR-H3 as depicted in SEQ ID NO: 2013, CDR-L1 as depicted in SEQ ID NO: 2014, CDR-L2 as depicted in SEQ ID NO: 2015 and CDR-L3 as depicted in SEQ ID NO: 2016.

CDR-H1 as depicted in SEQ ID NO: 2024, CDR-H2 as depicted in SEQ ID NO: 2025, CDR-H3 as depicted in SEQ ID NO: 2026, CDR-L1 as depicted in SEQ ID NO: 2027, CDR-L2 as depicted in SEQ ID NO: 2028 and CDR-L3 as depicted in SEQ ID NO: 2029,

CDR-H1 as depicted in SEQ ID NO: 2037, CDR-H2 as depicted in SEQ ID NO: 2038, CDR-H3 as depicted in SEQ ID NO: 2039, CDR-L1 as depicted in SEQ ID NO: 2040, CDR-L2 as depicted in SEQ ID NO: 2041 and CDR-L3 as depicted in SEQ ID NO: 2042, and

CDR-H1 as depicted in SEQ ID NO: 2050, CDR-H2 as depicted in SEQ ID NO: 2051, CDR-H3 as depicted in SEQ ID NO: 2052, CDR-L1 as depicted in SEQ ID NO: 2053, CDR-L2 as depicted in SEQ ID NO: 2054 and CDR-L3 as depicted in SEQ ID NO: 2055,

which all characterize binding domains for CDH19 grouped into bin 2;

(c) CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 96, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 102, CDR-L1 as depicted in SEQ ID NO: 268,

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NO: 1053,

CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 156, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 324, CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 912, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 913, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270, CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 910, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 911, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 914, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288, CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 920, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 324, CDR-H1 as depicted in SEQ ID NO: 996, CDR-H2 as depicted in SEQ ID NO: 997, CDR-H3 as depicted in SEQ ID NO: 998, CDR-L1 as depicted in SEQ ID NO: 999, CDR-L2 as depicted in SEQ ID NO: 1000 and CDR-L3 as depicted in SEQ ID NO: 1001, CDR-H1 as depicted in SEQ ID NO: 1048, CDR-H2 as depicted in SEQ ID NO: 1049, CDR-H3 as depicted in SEQ ID NO: 1050, CDR-L1 as depicted in SEQ ID NO: 1051,

CDR-H1 as depicted in SEQ ID NO: 1087, CDR-H2 as depicted in SEQ ID NO: 1088,

CDR-L2 as depicted in SEQ ID NO: 1052 and CDR-L3 as depicted in SEQ ID

CDR-H3 as depicted in SEQ ID NO: 1089, CDR-L1 as depicted in SEQ ID NO: 1090, CDR-L2 as depicted in SEQ ID NO: 1091 and CDR-L3 as depicted in SEQ ID NO: 1092,

CDR-H1 as depicted in SEQ ID NO: 1608, CDR-H2 as depicted in SEQ ID NO: 1609, CDR-H3 as depicted in SEQ ID NO: 1610, CDR-L1 as depicted in SEQ ID NO: 1611, CDR-L2 as depicted in SEQ ID NO: 1612 and CDR-L3 as depicted in SEQ ID NO: 1613,

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CDR-H1 as depicted in SEQ ID NO: 1621, CDR-H2 as depicted in SEQ ID NO: 1622, CDR-H3 as depicted in SEQ ID NO: 1623, CDR-L1 as depicted in SEQ ID NO: 1624, CDR-L2 as depicted in SEQ ID NO: 1625 and CDR-L3 as depicted in SEQ ID NO: 1626.

CDR-H1 as depicted in SEQ ID NO: 1634, CDR-H2 as depicted in SEQ ID NO: 1635, CDR-H3 as depicted in SEQ ID NO: 1636, CDR-L1 as depicted in SEQ ID NO: 1637, CDR-L2 as depicted in SEQ ID NO: 1638 and CDR-L3 as depicted in SEQ ID NO: 1639.

CDR-H1 as depicted in SEQ ID NO: 1673, CDR-H2 as depicted in SEQ ID NO: 1674, CDR-H3 as depicted in SEQ ID NO: 1675, CDR-L1 as depicted in SEQ ID NO: 1676, CDR-L2 as depicted in SEQ ID NO: 1677 and CDR-L3 as depicted in SEQ ID NO: 1678,

CDR-H1 as depicted in SEQ ID NO: 1686, CDR-H2 as depicted in SEQ ID NO: 1687, CDR-H3 as depicted in SEQ ID NO: 1688, CDR-L1 as depicted in SEQ ID NO: 1689, CDR-L2 as depicted in SEQ ID NO: 1690 and CDR-L3 as depicted in SEQ ID NO: 1691,

CDR-H1 as depicted in SEQ ID NO: 1699, CDR-H2 as depicted in SEQ ID NO: 1700, CDR-H3 as depicted in SEQ ID NO: 1701, CDR-L1 as depicted in SEQ ID NO: 1702, CDR-L2 as depicted in SEQ ID NO: 1703 and CDR-L3 as depicted in SEQ ID NO: 1704,

CDR-H1 as depicted in SEQ ID NO: 1712, CDR-H2 as depicted in SEQ ID NO: 1713, CDR-H3 as depicted in SEQ ID NO: 1714, CDR-L1 as depicted in SEQ ID NO: 1715, CDR-L2 as depicted in SEQ ID NO: 1716 and CDR-L3 as depicted in SEQ ID NO: 1717,

CDR-H1 as depicted in SEQ ID NO: 1725, CDR-H2 as depicted in SEQ ID NO: 1726, CDR-H3 as depicted in SEQ ID NO: 1727, CDR-L1 as depicted in SEQ ID NO: 1728, CDR-L2 as depicted in SEQ ID NO: 1729 and CDR-L3 as depicted in SEQ ID NO: 1730.

CDR-H1 as depicted in SEQ ID NO: 1738, CDR-H2 as depicted in SEQ ID NO: 1739, CDR-H3 as depicted in SEQ ID NO: 1740, CDR-L1 as depicted in SEQ ID NO: 1741,

CDR-L2 as depicted in SEQ ID NO: 1742 and CDR-L3 as depicted in SEQ ID NO: 1743,

CDR-H1 as depicted in SEQ ID NO: 1751, CDR-H2 as depicted in SEQ ID NO: 1752, CDR-H3 as depicted in SEQ ID NO: 1753, CDR-L1 as depicted in SEQ ID NO: 1754, CDR-L2 as depicted in SEQ ID NO: 1755 and CDR-L3 as depicted in SEQ ID NO: 1756,

CDR-H1 as depicted in SEQ ID NO: 1764, CDR-H2 as depicted in SEQ ID NO: 1765, CDR-H3 as depicted in SEQ ID NO: 1766, CDR-L1 as depicted in SEQ ID NO: 1767, CDR-L2 as depicted in SEQ ID NO: 1768 and CDR-L3 as depicted in SEQ ID NO: 1769, and

CDR-H1 as depicted in SEQ ID NO: 1920, CDR-H2 as depicted in SEQ ID NO: 1921, CDR-H3 as depicted in SEQ ID NO: 1922, CDR-L1 as depicted in SEQ ID NO: 1923, CDR-L2 as depicted in SEQ ID NO: 1924 and CDR-L3 as depicted in SEQ ID NO: 1925.

which all characterize binding domains for CDH19 grouped into bin 3;

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(d)

CDR-H1 as depicted in SEQ ID NO: 4, CDR-H2 as depicted in SEQ ID NO: 5, CDR-H3 as depicted in SEQ ID NO: 6, CDR-L1 as depicted in SEQ ID NO: 172, CDR-L2 as depicted in SEQID NO: 173 and CDR-L3 as depicted in SEQID NO: 174, CDR-H1 as depicted in SEQ ID NO: 10, CDR-H2 as depicted in SEQ ID NO: 11, CDR-H3 as depicted in SEQ ID NO: 12, CDR-L1 as depicted in SEQ ID NO: 178, CDR-L2 as depicted in SEQ ID NO: 179 and CDR-L3 as depicted in SEQ ID NO: 180, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 196, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 198, CDR-H1 as depicted in SEQ ID NO: 34, CDR-H2 as depicted in SEQ ID NO: 35, CDR-H3 as depicted in SEQ ID NO: 36, CDR-L1 as depicted in SEQ ID NO: 202, CDR-L2 as depicted in SEQ ID NO: 203 and CDR-L3 as depicted in SEQ ID NO: 204, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 214, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 59, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228, CDR-H1 as depicted in SEQ ID NO: 64, CDR-H2 as depicted in SEQ ID NO: 65, CDR-H3 as depicted in SEQ ID NO: 66, CDR-L1 as depicted in SEQ ID NO: 232, CDR-L2 as depicted in SEQ ID NO: 233 and CDR-L3 as depicted in SEQ ID NO: 234, CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 71, CDR-

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H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 328, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216, CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 902, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216. CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 903, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216. CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 925, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216. CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907, CDR-H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907, CDR-H3 as depicted in SEQ ID NO: 908, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 901, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 906, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228, CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228, CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 921, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939,

CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 940, CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 941, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 924, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 924, CDR-H3 as depicted in SEQ ID NO: 925, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923, CDR-H1 as depicted in SEQ ID NO: 923, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 921, CDR-H3 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 923, CDR-L2 as depicted in

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15 CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330, CDR-H1 as depicted in SEQ ID NO: 970, CDR-H2 as depicted in SEQ ID NO: 971, CDR-H3 as depicted in SEQ ID NO: 972, CDR-L1 as depicted in SEQ ID NO: 973, CDR-L2 as depicted in SEQ ID NO: 974 and CDR-L3 as depicted in SEQ ID NO: 975,

CDR-H1 as depicted in SEQ ID NO: 1061, CDR-H2 as depicted in SEQ ID NO: 1062, CDR-H3 as depicted in SEQ ID NO: 1063, CDR-L1 as depicted in SEQ ID NO: 1064, CDR-L2 as depicted in SEQ ID NO: 1065 and CDR-L3 as depicted in SEQ ID NO: 1066,

CDR-H1 as depicted in SEQ ID NO: 1139, CDR-H2 as depicted in SEQ ID NO: 1140, CDR-H3 as depicted in SEQ ID NO: 1141, CDR-L1 as depicted in SEQ ID NO: 1142, CDR-L2 as depicted in SEQ ID NO: 1143 and CDR-L3 as depicted in SEQ ID NO: 1144.

CDR-H1 as depicted in SEQ ID NO: 1152, CDR-H2 as depicted in SEQ ID NO: 1153, CDR-H3 as depicted in SEQ ID NO: 1154, CDR-L1 as depicted in SEQ ID NO: 1155, CDR-L2 as depicted in SEQ ID NO: 1156 and CDR-L3 as depicted in SEQ ID NO: 1157,

CDR-H1 as depicted in SEQ ID NO: 1178, CDR-H2 as depicted in SEQ ID NO: 1179, CDR-H3 as depicted in SEQ ID NO: 1180, CDR-L1 as depicted in SEQ ID NO: 1181, CDR-L2 as depicted in SEQ ID NO: 1182 and CDR-L3 as depicted in SEQ ID NO: 1183,

CDR-H1 as depicted in SEQ ID NO: 1191, CDR-H2 as depicted in SEQ ID NO: 1192, CDR-H3 as depicted in SEQ ID NO: 1193, CDR-L1 as depicted in SEQ ID NO: 1194, CDR-L2 as depicted in SEQ ID NO: 1195 and CDR-L3 as depicted in SEQ ID NO: 1196,

CDR-H1 as depicted in SEQ ID NO: 1204, CDR-H2 as depicted in SEQ ID NO: 1205, CDR-H3 as depicted in SEQ ID NO: 1206, CDR-L1 as depicted in SEQ ID NO: 1207, CDR-L2 as depicted in SEQ ID NO: 1208 and CDR-L3 as depicted in SEQ ID NO: 1209,

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CDR-H1 as depicted in SEQ ID NO: 1217, CDR-H2 as depicted in SEQ ID NO: 1218, CDR-H3 as depicted in SEQ ID NO: 1219, CDR-L1 as depicted in SEQ ID NO: 1220, CDR-L2 as depicted in SEQ ID NO: 1221 and CDR-L3 as depicted in SEQ ID NO: 1222,

CDR-H1 as depicted in SEQ ID NO: 1230, CDR-H2 as depicted in SEQ ID NO: 1231, CDR-H3 as depicted in SEQ ID NO: 1232, CDR-L1 as depicted in SEQ ID NO: 1233, CDR-L2 as depicted in SEQ ID NO: 1234 and CDR-L3 as depicted in SEQ ID NO: 1235.

CDR-H1 as depicted in SEQ ID NO: 1308, CDR-H2 as depicted in SEQ ID NO: 1309, CDR-H3 as depicted in SEQ ID NO: 1310, CDR-L1 as depicted in SEQ ID NO: 1311, CDR-L2 as depicted in SEQ ID NO: 1312 and CDR-L3 as depicted in SEQ ID NO: 1313.

CDR-H1 as depicted in SEQ ID NO: 1321, CDR-H2 as depicted in SEQ ID NO: 1322, CDR-H3 as depicted in SEQ ID NO: 1323, CDR-L1 as depicted in SEQ ID NO: 1324, CDR-L2 as depicted in SEQ ID NO: 1325 and CDR-L3 as depicted in SEQ ID NO: 1326,

CDR-H1 as depicted in SEQ ID NO: 1373, CDR-H2 as depicted in SEQ ID NO: 1374, CDR-H3 as depicted in SEQ ID NO: 1375, CDR-L1 as depicted in SEQ ID NO: 1376, CDR-L2 as depicted in SEQ ID NO: 1377 and CDR-L3 as depicted in SEQ ID NO: 1378,

CDR-H1 as depicted in SEQ ID NO: 1386, CDR-H2 as depicted in SEQ ID NO: 1387, CDR-H3 as depicted in SEQ ID NO: 1388, CDR-L1 as depicted in SEQ ID NO: 1389, CDR-L2 as depicted in SEQ ID NO: 1390 and CDR-L3 as depicted in SEQ ID NO: 1391,

CDR-H1 as depicted in SEQ ID NO: 1399, CDR-H2 as depicted in SEQ ID NO: 1400, CDR-H3 as depicted in SEQ ID NO: 1401, CDR-L1 as depicted in SEQ ID NO: 1402, CDR-L2 as depicted in SEQ ID NO: 1403 and CDR-L3 as depicted in SEQ ID NO: 1404,

CDR-H1 as depicted in SEQ ID NO: 1412, CDR-H2 as depicted in SEQ ID NO: 1413,

CDR-H3 as depicted in SEQ ID NO: 1414, CDR-L1 as depicted in SEQ ID NO: 1415, CDR-L2 as depicted in SEQ ID NO: 1416 and CDR-L3 as depicted in SEQ ID NO: 1417,

CDR-H1 as depicted in SEQ ID NO: 1777, CDR-H2 as depicted in SEQ ID NO: 1778, CDR-H3 as depicted in SEQ ID NO: 1779, CDR-L1 as depicted in SEQ ID NO: 1780, CDR-L2 as depicted in SEQ ID NO: 1781 and CDR-L3 as depicted in SEQ ID NO: 1782,

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CDR-H1 as depicted in SEQ ID NO: 1790, CDR-H2 as depicted in SEQ ID NO: 1791, CDR-H3 as depicted in SEQ ID NO: 1792, CDR-L1 as depicted in SEQ ID NO: 1793, CDR-L2 as depicted in SEQ ID NO: 1794 and CDR-L3 as depicted in SEQ ID NO: 1795.

CDR-H1 as depicted in SEQ ID NO: 1803, CDR-H2 as depicted in SEQ ID NO: 1804, CDR-H3 as depicted in SEQ ID NO: 1805, CDR-L1 as depicted in SEQ ID NO: 1806, CDR-L2 as depicted in SEQ ID NO: 1807 and CDR-L3 as depicted in SEQ ID NO: 1808.

CDR-H1 as depicted in SEQ ID NO: 1816, CDR-H2 as depicted in SEQ ID NO: 1817, CDR-H3 as depicted in SEQ ID NO: 1818, CDR-L1 as depicted in SEQ ID NO: 1819, CDR-L2 as depicted in SEQ ID NO: 1820 and CDR-L3 as depicted in SEQ ID NO: 1821,

CDR-H1 as depicted in SEQ ID NO: 1829, CDR-H2 as depicted in SEQ ID NO: 1830, CDR-H3 as depicted in SEQ ID NO: 1831, CDR-L1 as depicted in SEQ ID NO: 1832, CDR-L2 as depicted in SEQ ID NO: 1833 and CDR-L3 as depicted in SEQ ID NO: 1834,

CDR-H1 as depicted in SEQ ID NO: 1842, CDR-H2 as depicted in SEQ ID NO: 1843, CDR-H3 as depicted in SEQ ID NO: 1844, CDR-L1 as depicted in SEQ ID NO: 1845, CDR-L2 as depicted in SEQ ID NO: 1846 and CDR-L3 as depicted in SEQ ID NO: 1847,

CDR-H1 as depicted in SEQ ID NO: 1855, CDR-H2 as depicted in SEQ ID NO: 1856, CDR-H3 as depicted in SEQ ID NO: 1857, CDR-L1 as depicted in SEQ ID NO: 1858, CDR-L2 as depicted in SEQ ID NO: 1859 and CDR-L3 as depicted in SEQ ID NO: 1860,

CDR-H1 as depicted in SEQ ID NO: 1868, CDR-H2 as depicted in SEQ ID NO: 1869, CDR-H3 as depicted in SEQ ID NO: 1870, CDR-L1 as depicted in SEQ ID NO: 1871, CDR-L2 as depicted in SEQ ID NO: 1872 and CDR-L3 as depicted in SEQ ID NO: 1873.

CDR-H1 as depicted in SEQ ID NO: 1881, CDR-H2 as depicted in SEQ ID NO: 1882, CDR-H3 as depicted in SEQ ID NO: 1883, CDR-L1 as depicted in SEQ ID NO: 1884,

CDR-L2 as depicted in SEQ ID NO: 1885 and CDR-L3 as depicted in SEQ ID NO: 1886,

CDR-H1 as depicted in SEQ ID NO: 2063, CDR-H2 as depicted in SEQ ID NO: 2064, CDR-H3 as depicted in SEQ ID NO: 2065, CDR-L1 as depicted in SEQ ID NO: 2066, CDR-L2 as depicted in SEQ ID NO: 2067 and CDR-L3 as depicted in SEQ ID NO: 2068,

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CDR-H1 as depicted in SEQ ID NO: 2076, CDR-H2 as depicted in SEQ ID NO: 2077, CDR-H3 as depicted in SEQ ID NO: 2078, CDR-L1 as depicted in SEQ ID NO: 2079, CDR-L2 as depicted in SEQ ID NO: 2080 and CDR-L3 as depicted in SEQ ID NO: 2081.

CDR-H1 as depicted in SEQ ID NO: 2089, CDR-H2 as depicted in SEQ ID NO: 2090, CDR-H3 as depicted in SEQ ID NO: 2091, CDR-L1 as depicted in SEQ ID NO: 2092, CDR-L2 as depicted in SEQ ID NO: 2093 and CDR-L3 as depicted in SEQ ID NO: 2094.

CDR-H1 as depicted in SEQ ID NO: 2102, CDR-H2 as depicted in SEQ ID NO: 2103, CDR-H3 as depicted in SEQ ID NO: 2104, CDR-L1 as depicted in SEQ ID NO: 2105, CDR-L2 as depicted in SEQ ID NO: 2106 and CDR-L3 as depicted in SEQ ID NO: 2107,

CDR-H1 as depicted in SEQ ID NO: 2115, CDR-H2 as depicted in SEQ ID NO: 2116, CDR-H3 as depicted in SEQ ID NO: 2117, CDR-L1 as depicted in SEQ ID NO: 2118, CDR-L2 as depicted in SEQ ID NO: 2119 and CDR-L3 as depicted in SEQ ID NO: 2120,

CDR-H1 as depicted in SEQ ID NO: 2128, CDR-H2 as depicted in SEQ ID NO: 2129, CDR-H3 as depicted in SEQ ID NO: 2130, CDR-L1 as depicted in SEQ ID NO: 2131, CDR-L2 as depicted in SEQ ID NO: 2132 and CDR-L3 as depicted in SEQ ID NO: 2133.

CDR-H1 as depicted in SEQ ID NO: 2141, CDR-H2 as depicted in SEQ ID NO: 2142, CDR-H3 as depicted in SEQ ID NO: 2143, CDR-L1 as depicted in SEQ ID NO: 2144, CDR-L2 as depicted in SEQ ID NO: 2145 and CDR-L3 as depicted in SEQ ID NO: 2146,

CDR-H1 as depicted in SEQ ID NO: 2154, CDR-H2 as depicted in SEQ ID NO: 2155, CDR-H3 as depicted in SEQ ID NO: 2156, CDR-L1 as depicted in SEQ ID NO: 2157, CDR-L2 as depicted in SEQ ID NO: 2158 and CDR-L3 as depicted in SEQ ID NO: 2159,

35 CDR-H1 as depicted in SEQ ID NO: 2180, CDR-H2 as depicted in SEQ ID NO: 2181, CDR-H3 as depicted in SEQ ID NO: 2182, CDR-L1 as depicted in SEQ ID NO: 2183, CDR-L2 as depicted in SEQ ID NO: 2184 and CDR-L3 as depicted in SEQ ID

NO: 2185,

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CDR-H1 as depicted in SEQ ID NO: 2193, CDR-H2 as depicted in SEQ ID NO: 2194, CDR-H3 as depicted in SEQ ID NO: 2195, CDR-L1 as depicted in SEQ ID NO: 2196, CDR-L2 as depicted in SEQ ID NO: 2197 and CDR-L3 as depicted in SEQ ID NO: 2198, and

CDR-H1 as depicted in SEQ ID NO: 2206, CDR-H2 as depicted in SEQ ID NO: 2207, CDR-H3 as depicted in SEQ ID NO: 2208, CDR-L1 as depicted in SEQ ID NO: 2209, CDR-L2 as depicted in SEQ ID NO: 2210 and CDR-L3 as depicted in SEQ ID NO: 2211

which all characterize binding domains for CDH19 grouped into bin 4; and

CDR-H1 as depicted in SEQ ID NO: 76, CDR-H2 as depicted in SEQ ID NO: 77, CDR-(e) H3 as depicted in SEQ ID NO: 78, CDR-L1 as depicted in SEQ ID NO: 244, CDR-L2 as depicted in SEQ ID NO: 245 and CDR-L3 as depicted in SEQ ID NO: 246, CDR-H1 as depicted in SEQ ID NO: 88, CDR-H2 as depicted in SEQ ID NO: 89, CDR-H3 as depicted in SEQ ID NO: 90, CDR-L1 as depicted in SEQ ID NO: 256, CDR-L2 as depicted in SEQ ID NO: 257 and CDR-L3 as depicted in SEQ ID NO: 258, CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274, CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276, CDR-H1 as depicted in SEQ ID NO: 112, CDR-H2 as depicted in SEQ ID NO: 113, CDR-H3 as depicted in SEQ ID NO: 114, CDR-L1 as depicted in SEQ ID NO: 280, CDR-L2 as depicted in SEQ ID NO: 281 and CDR-L3 as depicted in SEQ ID NO: 282, CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274, CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276, CDR-H1 as depicted in SEQ ID NO: 983, CDR-H2 as depicted in SEQ ID NO: 984, CDR-H3 as depicted in SEQ ID NO: 985, CDR-L1 as depicted in SEQ ID NO: 986, CDR-L2 as depicted in SEQ ID NO: 987 and CDR-L3 as depicted in SEQ ID NO: 988, CDR-H1 as depicted in SEQ ID NO: 1582, CDR-H2 as depicted in SEQ ID NO: 1583, CDR-H3 as depicted in SEQ ID NO: 1584, CDR-L1 as depicted in SEQ ID NO: 1585, CDR-L2 as depicted in SEQ ID NO: 1586 and CDR-L3 as depicted in SEQ ID NO: 1587, and

CDR-H1 as depicted in SEQ ID NO: 1595, CDR-H2 as depicted in SEQ ID NO: 1596, CDR-H3 as depicted in SEQ ID NO: 1597, CDR-L1 as depicted in SEQ ID NO: 1598, CDR-L2 as depicted in SEQ ID NO: 1599 and CDR-L3 as depicted in SEQ ID NO: 1600,

which all characterize binding domains for CDH19 grouped into bin 5.

In a further embodiment of the antibody construct of the invention the first binding domain comprises a VH region selected from the group consisting of VH regions

- (a) as depicted in SEQ ID NO: 362, SEQ ID NO: 364, SEQ ID NO: 485, SEQ ID NO: 486, SEQ ID NO: 487, SEQ ID NO: 492, SEQ ID NO: 493, SEQ ID NO: 494, SEQ ID NO: 495, SEQ ID NO: 1133, SEQ ID NO: 1172, SEQ ID NO: 1341, SEQ ID NO: 1354, SEQ ID NO: 1367, SEQ ID NO: 1432, SEQ ID NO: 1445 and SEQ ID NO: 2174, grouped into bin 1;
- as depicted in SEQ ID NO: 342, SEQ ID NO: 366, SEQ ID NO: 370, SEQ ID NO: 344, (b) 10 SEQ ID NO: 372, SEQ ID NO: 368, SEQ ID NO: 496, SEQ ID NO: 497, SEQ ID NO: 498. SEQ ID NO: 499. SEQ ID NO: 500. SEQ ID NO: 508. SEQ ID NO: 509. SEQ ID NO: 510, SEQ ID NO: 511, SEQ ID NO: 512, SEQ ID NO: 519, SEQ ID NO: 520, SEQ ID NO: 521, SEQ ID NO: 522, SEQ ID NO: 523, SEQ ID NO: 524, SEQ ID NO: 525, SEQ ID NO: 526, SEQ ID NO: 527, SEQ ID NO: 528, SEQ ID NO: 529, SEQ ID NO: 530, SEQ ID NO: 531, SEQ ID NO: 532, SEQ ID NO: 533, 15 SEQ ID NO: 534, SEQ ID NO: 535, SEQ ID NO: 536, SEQ ID NO: 537, SEQ ID NO: 538, SEQ ID NO: 1016, SEQ ID NO: 1029, SEQ ID NO: 1042, SEQ ID NO: 1081, SEQ ID NO: 1107, SEQ ID NO: 1120, SEQ ID NO: 1250, SEQ ID NO: 1263, SEQ ID NO: 1276, SEQ ID NO: 1289, SEQ ID NO: 1302, SEQ ID NO: 1654, SEQ ID 20 NO: 1667. SEQ ID NO: 1901. SEQ ID NO: 1914. SEQ ID NO: 1940. SEQ ID NO: 1953, SEQ ID NO: 1966, SEQ ID NO: 1979, SEQ ID NO: 1992, SEQ ID NO: 2005, SEQ ID NO: 2018, SEQ ID NO: 2031, SEQ ID NO: 2044, and SEQ ID NO: 2057,

grouped into bin 2;

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- 25 (c) as depicted in SEQ ID NO: 338, SEQ ID NO: 354, SEQ ID NO: 378, SEQ ID NO: 356, SEQ ID NO: 476, SEQ ID NO: 477, SEQ ID NO: 478, SEQ ID NO: 479, SEQ ID NO: 480, SEQ ID NO: 481, SEQ ID NO: 482, SEQ ID NO: 483, SEQ ID NO: 484, SEQ ID NO: 501, SEQ ID NO: 502, SEQ ID NO: 503, SEQ ID NO: 504, SEQ ID NO: 505, SEQ ID NO: 506, SEQ ID NO: 517, SEQ ID NO: 518, SEQ ID NO: 1003, SEQ ID NO: 1055, SEQ ID NO: 1094, SEQ ID NO: 1615, SEQ ID NO: 1628, SEQ ID NO: 1641, SEQ ID NO: 1680, SEQ ID NO: 1693, SEQ ID NO: 1706, SEQ ID NO: 1719, SEQ ID NO: 1732, SEQ ID NO: 1745, SEQ ID NO: 1758, SEQ ID NO: 1771, and SEQ ID NO: 1927, grouped into bin 3;
 - grouped into biri 5,
- 35 (d) as depicted in SEQ ID NO: 352, SEQ ID NO: 360, SEQ ID NO: 388, SEQ ID NO: 386, SEQ ID NO: 340, SEQ ID NO: 346, SEQ ID NO: 374, SEQ ID NO: 348, SEQ ID NO: 390, SEQ ID NO: 463, SEQ ID NO: 464, SEQ ID NO: 465, SEQ ID NO: 466,

SEQ ID NO: 467, SEQ ID NO: 468, SEQ ID NO: 469, SEQ ID NO: 470, SEQ ID NO: 471, SEQ ID NO: 472, SEQ ID NO: 473, SEQ ID NO: 474, SEQ ID NO: 475, SEQ ID NO: 488, SEQ ID NO: 489, SEQ ID NO: 490, SEQ ID NO: 491, SEQ ID NO: 513, SEQ ID NO: 514, SEQ ID NO: 515, SEQ ID NO: 516, SEQ ID NO: 540, 5 SEQ ID NO: 541, SEQ ID NO: 542, SEQ ID NO: 543, SEQ ID NO: 977, SEQ ID NO: 1068, SEQ ID NO: 1146, SEQ ID NO: 1159, SEQ ID NO: 1185, SEQ ID NO: 1198, SEQ ID NO: 1211, SEQ ID NO: 1224, SEQ ID NO: 1237, SEQ ID NO: 1315, SEQ ID NO: 1328, SEQ ID NO: 1380, SEQ ID NO: 1393, SEQ ID NO: 1406, SEQ ID NO: 1419, SEQ ID NO: 1469, SEQ ID NO: 1478, SEQ ID 10 NO: 1485, SEQ ID NO: 1494, SEQ ID NO: 1501, SEQ ID NO: 1508, SEQ ID NO: 1519, SEQ ID NO: 1526, SEQ ID NO: 1533, SEQ ID NO: 1542, SEQ ID NO: 1549, SEQ ID NO: 1558, SEQ ID NO: 1565, SEQ ID NO: 1784, SEQ ID NO: 1797, SEQ ID NO: 1810, SEQ ID NO: 1823, SEQ ID NO: 1836, SEQ ID NO: 1849, SEQ ID NO: 1862, SEQ ID NO: 1875, SEQ ID NO: 1888, SEQ ID 15 NO: 2070, SEQ ID NO: 2083, SEQ ID NO: 2096, SEQ ID NO: 2109, SEQ ID NO: 2122, SEQ ID NO: 2135, SEQ ID NO: 2148, SEQ ID NO: 2161, SEQ ID NO: 2187, SEQ ID NO: 2200, and SEQ ID NO: 2213, grouped into bin 4; and

(e) as depicted in SEQ ID NO: 376, SEQ ID NO: 392, SEQ ID NO: 358, SEQ ID NO: 350, SEQ ID NO: 507, SEQ ID NO: 990, SEQ ID NO: 1589, and SEQ ID NO: 1602, grouped into bin 5.

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In another embodiment of the antibody construct of the invention the first binding domain comprises a VL region selected from the group consisting of VL regions

- 25 (a) as depicted in SEQ ID NO: 418, SEQ ID NO: 420, SEQ ID NO: 580, SEQ ID NO: 581, SEQ ID NO: 582, SEQ ID NO: 587, SEQ ID NO: 588, SEQ ID NO: 589, SEQ ID NO: 590, SEQ ID NO: 1135, SEQ ID NO: 1174, SEQ ID NO: 1343, SEQ ID NO: 1356, SEQ ID NO: 1369, SEQ ID NO: 1434, SEQ ID NO: 1447 and SEQ ID NO: 2176, grouped into bin 1;
- 30 (b) as depicted in SEQ ID NO: 398, SEQ ID NO: 422, SEQ ID NO: 426, SEQ ID NO: 400, SEQ ID NO: 428, SEQ ID NO: 424, SEQ ID NO: 591, SEQ ID NO: 592, SEQ ID NO: 593, SEQ ID NO: 594, SEQ ID NO: 595, SEQ ID NO: 603, SEQ ID NO: 604, SEQ ID NO: 605, SEQ ID NO: 606, SEQ ID NO: 607, SEQ ID NO: 614, SEQ ID NO: 615, SEQ ID NO: 616, SEQ ID NO: 617, SEQ ID NO: 618, SEQ ID NO: 619, SEQ ID NO: 620, SEQ ID NO: 621, SEQ ID NO: 622, SEQ ID NO: 623, SEQ ID NO: 624, SEQ ID NO: 625, SEQ ID NO: 626, SEQ ID NO: 627, SEQ ID NO: 628, SEQ ID NO: 629, SEQ ID NO: 630, SEQ ID NO: 631, SEQ ID NO: 632, SEQ ID

NO: 633, SEQ ID NO: 1018, SEQ ID NO: 1031, SEQ ID NO: 1044, SEQ ID NO: 1083, SEQ ID NO: 1109, SEQ ID NO: 1122, SEQ ID NO: 1252, SEQ ID NO: 1265, SEQ ID NO: 1278, SEQ ID NO: 1291, SEQ ID NO: 1304, SEQ ID NO: 1656, SEQ ID NO: 1669, SEQ ID NO: 1903, SEQ ID NO: 1916, SEQ ID NO: 1942, SEQ ID NO: 1955, SEQ ID NO: 1968, SEQ ID NO: 1981, SEQ ID NO: 1994, SEQ ID NO: 2007, SEQ ID NO: 2020, SEQ ID NO: 2033, SEQ ID NO: 2046, and SEQ ID NO: 2059,

grouped into bin 2;

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(c) as depicted in SEQ ID NO: 394, SEQ ID NO: 410, SEQ ID NO: 434, SEQ ID NO: 412, SEQ ID NO: 571, SEQ ID NO: 572, SEQ ID NO: 573, SEQ ID NO: 574, SEQ ID NO: 575, SEQ ID NO: 576, SEQ ID NO: 577, SEQ ID NO: 578, SEQ ID NO: 579, SEQ ID NO: 596, SEQ ID NO: 597, SEQ ID NO: 598, SEQ ID NO: 599, SEQ ID NO: 600, SEQ ID NO: 601, SEQ ID NO: 612, SEQ ID NO: 613, SEQ ID NO: 1005, SEQ ID NO: 1057, SEQ ID NO: 1096, SEQ ID NO: 1617, SEQ ID NO: 1630, SEQ ID NO: 1643, SEQ ID NO: 1682, SEQ ID NO: 1695, SEQ ID NO: 1708, SEQ ID NO: 1721, SEQ ID NO: 1734, SEQ ID NO: 1747, SEQ ID NO: 1760, SEQ ID NO: 1773, and SEQ ID NO: 1929,

grouped into bin 3;

as depicted in SEQ ID NO: 408, SEQ ID NO: 416, SEQ ID NO: 444, SEQ ID NO: 442, (d) 20 SEQ ID NO: 396, SEQ ID NO: 402, SEQ ID NO: 430, SEQ ID NO: 404, SEQ ID NO: 446, SEQ ID NO: 558, SEQ ID NO: 559, SEQ ID NO: 560, SEQ ID NO: 561, SEQ ID NO: 562, SEQ ID NO: 563, SEQ ID NO: 564, SEQ ID NO: 565, SEQ ID NO: 566, SEQ ID NO: 567, SEQ ID NO: 568, SEQ ID NO: 569, SEQ ID NO: 570, SEQ ID NO: 583, SEQ ID NO: 584, SEQ ID NO: 585, SEQ ID NO: 586, SEQ ID 25 NO: 608, SEQ ID NO: 609, SEQ ID NO: 610, SEQ ID NO: 611, SEQ ID NO: 635, SEQ ID NO: 636, SEQ ID NO: 637, SEQ ID NO: 638, SEQ ID NO: 979, SEQ ID NO: 1070, SEQ ID NO: 1148, SEQ ID NO: 1161, SEQ ID NO: 1187, SEQ ID NO: 1200, SEQ ID NO: 1213, SEQ ID NO: 1226, SEQ ID NO: 1239, SEQ ID NO: 1317, SEQ ID NO: 1330, SEQ ID NO: 1382, SEQ ID NO: 1395, SEQ ID 30 NO: 1408, SEQ ID NO: 1421, SEQ ID NO: 1471, SEQ ID NO: 1480, SEQ ID NO: 1487, SEQ ID NO: 1496, SEQ ID NO: 1503, SEQ ID NO: 1510, SEQ ID NO: 1521, SEQ ID NO: 1528, SEQ ID NO: 1535, SEQ ID NO: 1544, SEQ ID NO: 1551, SEQ ID NO: 1560, SEQ ID NO: 1567, SEQ ID NO: 1786, SEQ ID NO: 1799, SEQ ID NO: 1812, SEQ ID NO: 1825, SEQ ID NO: 1838, SEQ ID 35 NO: 1851, SEQ ID NO: 1864, SEQ ID NO: 1877, SEQ ID NO: 1890, SEQ ID NO: 2072, SEQ ID NO: 2085, SEQ ID NO: 2098, SEQ ID NO: 2111, SEQ ID NO: 2124, SEQ ID NO: 2137, SEQ ID NO: 2150, SEQ ID NO: 2163, SEQ ID

NO: 2189, SEQ ID NO: 2202, and SEQ ID NO: 2215, grouped into bin 4; and

(e) as depicted in SEQ ID NO: 432, SEQ ID NO: 448, SEQ ID NO: 414, SEQ ID NO: 406, SEQ ID NO: 602, SEQ ID NO: 992, SEQ ID NO: 1591, and SEQ ID NO: 1604, grouped into bin 5.

The invention further provides an embodiment of the antibody construct of the invention, wherein the first binding domain comprises a VH region and a VL region selected from the group consisting of:

(1) 10 pairs of a VH region and a VL region as depicted in SEQ ID NOs: 362+418, SEQ ID NOs: 364+420, SEQ ID NOs: 485+580. SEQ ID NOs: 486+581. SEQ ID NOs: 487+582, SEQ ID NOs: 492+587, SEQ ID SEQ ID NOs: 493+588, NOs: 494+589, SEQ ID NOs: 495+590, SEQ ID NOs: 1133+1135, SEQ ID NOs: 1172+1174, SEQ ID NOs: 1341+1343, SEQ ID NOs: 1354+1356, SEQ ID NOs: 1367+1369, SEQ ID NOs: 1432+1434, SEQ ID NOs: 1445+1447, and SEQ ID 15 NOs: 2174+2176,

all pairs grouped into bin 1;

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pairs of a VH region and a VL region as depicted in SEQ ID NOs: 342+398, SEQ ID (2) NOs: 366+422, SEQ ID NOs: 370+426, SEQ ID NOs: 344+400, SEQ ID NOs: 496+591, 20 NOs: 372+428, SEQ ID NOs: 368+424, SEQ ID SEQ ID NOs: 497+592, SEQ ID SEQ ID NOs: 498+593, NOs: 499+594, SEQ ID NOs: 500+595, SEQ ID NOs: 508+603, SEQ ID NOs: 509+604, SEQ ID NOs: 510+605, SEQ ID SEQ ID NOs: 511+606, NOs: 512+607, SEQ ID NOs: 519+614, NOs: 520+615, NOs: 521+616, SEQ ID SEQ ID SEQ ID 25 NOs: 522+617, SEQ ID NOs: 523+618, SEQ ID NOs: 524+619, SEQ ID SEQ ID NOs: 525+620. SEQ ID NOs: 526+621, NOs: 527+622, SEQ ID NOs: 528+623, SEQ ID NOs: 529+624, SEQ ID NOs: 530+625, SEQ ID NOs: 531+626, NOs: 532+627, SEQ ID SEQ ID NOs: 533+628, SEQ ID NOs: 534+629, SEQ ID NOs: 535+630, SEQ ID NOs: 536+631, SEQ ID 30 NOs: 537+632, SEQ ID NOs: 538+633, SEQ ID NOs: 1016+1018, SEQ ID NOs: 1029+1031, SEQ ID NOs: 1042+1044, SEQ ID NOs: 1081+1083, SEQ ID NOs: 1107+1109, SEQ ID NOs: 1120+1122, SEQ ID NOs: 1250+1252, SEQ ID NOs: 1263+1265, SEQ ID NOs: 1276+1278, SEQ ID NOs: 1289+1291, SEQ ID NOs: 1302+1304, SEQ ID NOs: 1654+1656, SEQ ID NOs: 1667+1669, SEQ ID 35 NOs: 1901+1903, SEQ ID NOs: 1914+1916, SEQ ID NOs: 1940+1942, SEQ ID NOs: 1953+1955, SEQ ID NOs: 1966+1968, SEQ ID NOs: 1979+1981, SEQ ID NOs: 1992+1994, SEQ ID NOs: 2005+2007, SEQ ID NOs: 2018+2020, SEQ ID

NOs: 2031+2033, SEQ ID NOs: 2044+2046, and SEQ ID NOs: 2057+2059, all pairs grouped into bin 2;

- (3) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 338+394, SEQ ID NOs: 354+410, SEQ ID NOs: 378+434, SEQ ID NOs: 356+412, SEQ ID 5 NOs: 476+571, SEQ ID NOs: 477+572, SEQ ID NOs: 478+573, SEQ ID NOs: 481+576, NOs: 479+574, SEQ ID NOs: 480+575, SEQ ID SEQ ID NOs: 482+577, NOs: 483+578, NOs: 484+579, SEQ ID SEQ ID SEQ ID NOs: 501+596, SEQ ID NOs: 502+597, SEQ ID NOs: 503+598, SEQ ID NOs: 505+600, NOs: 504+599, SEQ ID SEQ ID NOs: 506+601, SEQ ID NOs: 518+613, 10 NOs: 517+612, SEQ ID SEQ ID NOs: 1003+1005, SEQ ID NOs: 1055+1057, SEQ ID NOs: 1094+1096, SEQ ID NOs: 1615+1617, SEQ ID NOs: 1628+1630, SEQ ID NOs: 1641+1643, SEQ ID NOs: 1680+1682, SEQ ID NOs: 1693+1695, SEQ ID NOs: 1706+1708, SEQ ID NOs: 1719+1721, SEQ ID NOs: 1732+1734, SEQ ID NOs: 1745+1747, SEQ ID NOs: 1758+1760, SEQ ID 15 NOs: 1771+1773, and SEQ ID NOs: 1927+1929, all pairs grouped into bin 3;
- (4) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 352+408, SEQ ID NOs: 360+416, SEQ ID NOs: 388+444, SEQ ID NOs: 386+442, SEQ ID NOs: 340+396, SEQ ID NOs: 346+402, SEQ ID NOs: 374+430, SEQ ID 20 NOs: 348+404, SEQ ID NOs: 390+446, SEQ ID NOs: 463+558, SEQ ID NOs: 464+559, SEQ ID SEQ ID NOs: 465+560, NOs: 466+561, SEQ ID NOs: 467+562, SEQ ID NOs: 468+563, SEQ ID NOs: 469+564, SEQ ID NOs: 470+565, SEQ ID SEQ ID NOs: 471+566, NOs: 472+567, SEQ ID NOs: 473+568, SEQ ID NOs: 475+570, SEQ ID NOs: 474+569, SEQ ID 25 NOs: 488+583, SEQ ID NOs: 489+584, SEQ ID NOs: 490+585, SEQ ID SEQ ID NOs: 491+586, SEQ ID NOs: 513+608. NOs: 514+609. SEQ ID NOs: 515+610, SEQ ID NOs: 516+611, SEQ ID NOs: 540+635, SEQ ID NOs: 542+637, SEQ ID NOs: 543+638, NOs: 541+636, SEQ ID SEQ ID NOs: 977+979, SEQ ID NOs: 1068+1070, SEQ ID NOs: 1146+1148, SEQ ID 30 NOs: 1159+1161, SEQ ID NOs: 1185+1187, SEQ ID NOs: 1198+1200, SEQ ID NOs: 1211+1213, SEQ ID NOs: 1224+1226, SEQ ID NOs: 1237+1239, SEQ ID NOs: 1315+1317, SEQ ID NOs: 1328+1330, SEQ ID NOs: 1380,+1382 SEQ ID NOs: 1393+1395, SEQ ID NOs: 1406+1408, SEQ ID NOs: 1419+1421, SEQ ID NOs: 1469+1471, SEQ ID NOs: 1478+1480, SEQ ID NOs: 1485+1487, SEQ ID 35 NOs: 1494+1496, SEQ ID NOs: 1501+1503, SEQ ID NOs: 1508+1510, SEQ ID NOs: 1519+1521, SEQ ID NOs: 1526+1528, SEQ ID NOs: 1533+1535, SEQ ID

NOs: 1542+1544, SEQ ID NOs: 1549+1551, SEQ ID NOs: 1558+1560, SEQ ID

NOs: 1565+1567, SEQ ID NOs: 1784+1786, SEQ ID NOs: 1797+1799, SEQ ID NOs: 1810+1812, SEQ ID NOs: 1823+1825, SEQ ID NOs: 1836+1838, SEQ ID NOs: 1849+1851, SEQ ID NOs: 1862+1864, SEQ ID NOs: 1875+1877, SEQ ID NOs: 1888+1890, SEQ ID NOs: 2070+2072, SEQ ID NOs: 2083+2085, SEQ ID NOs: 2096+2098, SEQ ID NOs: 2109+2111, SEQ ID NOs: 2122+2124, SEQ ID NOs: 2135+2137, SEQ ID NOs: 2148+2150, SEQ ID NOs: 2161+2163, SEQ ID NOs: 2187+2189, SEQ ID NOs: 2200+2202, and SEQ ID NOs: 2213+2215, all pairs grouped into bin 4; and

- (5) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 376+432, SEQ ID NOs: 392+448, SEQ ID NOs: 358+414, SEQ ID NOs: 350+406, SEQ ID NOs: 507+602, SEQ ID NOs: 990+992, SEQ ID NOs: 1589+1591, and SEQ ID NOs: 1602+1604, all pairs grouped into bin 5.
- In a further embodiment of the invention the antibody construct is in a format selected from the group consisting of (scFv)₂, (single domain mAb)₂, scFv-single domain mAb, diabodies and oligomers thereof.

In a preferred embodiment the first binding domain comprises an amino acid selected from the group consisting of

- (a) as depicted in SEQ ID NO: 117, SEQ ID NO: 1137, SEQ ID NO: 1176, SEQ ID NO: 1345, SEQ ID NO: 1358, SEQ ID NO: 1371, SEQ ID NO: 1436, SEQ ID NO: 1449 and SEQ ID NO: 2178,
 - all binders grouped into bin 1;

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- (b) as depicted in SEQ ID NO: 1020, SEQ ID NO: 1033, SEQ ID NO: 1046, SEQ ID NO: 1085, SEQ ID NO: 1111, SEQ ID NO: 1124, SEQ ID NO: 1254, SEQ ID NO: 1267, SEQ ID NO: 1280, SEQ ID NO: 1293, SEQ ID NO: 1306, SEQ ID NO: 1658, SEQ ID NO: 1671, SEQ ID NO: 1905, SEQ ID NO: 1918, SEQ ID NO: 1944, SEQ ID NO: 1957, SEQ ID NO: 1970, SEQ ID NO: 1983, SEQ ID NO: 1996, SEQ ID NO: 2009, SEQ ID NO: 2022, SEQ ID NO: 2035, SEQ ID NO: 2048, and SEQ ID NO: 2061, all binders grouped into bin 2;
- (c) as depicted in SEQ ID NO: 1007, SEQ ID NO: 1059, SEQ ID NO: 1098, SEQ ID NO: 1619, SEQ ID NO: 1632, SEQ ID NO: 1645, SEQ ID NO: 1684, SEQ ID NO: 1697, SEQ ID NO: 1710, SEQ ID NO: 1723, SEQ ID NO: 1736, SEQ ID NO: 1749, SEQ ID NO: 1762, SEQ ID NO: 1775, and SEQ ID NO: 1931, all binders grouped into bin 3;

as depicted in SEQ ID NO: 981, SEQ ID NO: 1072, SEQ ID NO: 1150, SEQ ID (d) NO: 1163, SEQ ID NO: 1189, SEQ ID NO: 1202, SEQ ID NO: 1215, SEQ ID NO: 1228, SEQ ID NO: 1241, SEQ ID NO: 1319, SEQ ID NO: 1332, SEQ ID NO: 1384, SEQ ID NO: 1397, SEQ ID NO: 1410, SEQ ID NO: 1423, SEQ ID NO: 1473, SEQ ID NO: 1482, SEQ ID NO: 1489, SEQ ID NO: 1498, SEQ ID NO: 1505, SEQ ID NO: 1512, SEQ ID NO: 1523, SEQ ID NO: 1530, SEQ ID NO: 1537, SEQ ID NO: 1546, SEQ ID NO: 1553, SEQ ID NO: 1562, SEQ ID NO: 1569, SEQ ID NO: 1788, SEQ ID NO: 1801, SEQ ID NO: 1814, SEQ ID NO: 1827, SEQ ID NO: 1840, SEQ ID NO: 1853, SEQ ID NO: 1866, SEQ ID NO: 1879, SEQ ID NO: 1892, SEQ ID NO: 2074, SEQ ID NO: 2087, SEQ ID NO: 2100. SEQ ID NO: 2113. SEQ ID NO: 2126. SEQ ID NO: 2139, SEQ ID NO: 2152, SEQ ID NO: 2165, SEQ ID NO: 2191, SEQ ID NO: 2204, and SEQ ID NO: 2217,

all binders grouped into bin 4; and

15 (e) as depicted in SEQ ID NO: 994, SEQ ID NO: 1593, and SEQ ID NO: 1606, grouped into bin 5;.

In one aspect of the invention, the second binding domain is capable of binding to to human CD3 and to macaque CD3, preferably to human CD3 epsilon and to macaque CD3 epsilon. Additionally or alternatively, the second binding domain is capable of binding to *Callithrix jacchus*, *Saguinus oedipus* and/or *Saimiri sciureus* CD3 epsilon. According to these embodiments, one or both binding domains of the antibody construct of the invention are preferably cross-species specific for members of the mammalian order of primates. Cross-species specific CD3 binding domains are, for example, described in WO 2008/119567.

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It is particularly preferred for the antibody construct of the present invention that the second binding domain capable of binding to the T cell CD3 receptor complex comprises a VL region comprising CDR-L1, CDR-L2 and CDR-L3 selected from:

- (a) CDR-L1 as depicted in SEQ ID NO: 27 of WO 2008/119567, CDR-L2 as depicted in
 30 SEQ ID NO: 28 of WO 2008/119567 and CDR-L3 as depicted in SEQ ID NO: 29 of WO 2008/119567;
 - (b) CDR-L1 as depicted in SEQ ID NO: 117 of WO 2008/119567, CDR-L2 as depicted in SEQ ID NO: 118 of WO 2008/119567 and CDR-L3 as depicted in SEQ ID NO: 119 of WO 2008/119567; and
- 35 (c) CDR-L1 as depicted in SEQ ID NO: 153 of WO 2008/119567, CDR-L2 as depicted in SEQ ID NO: 154 of WO 2008/119567 and CDR-L3 as depicted in SEQ ID NO: 155 of WO 2008/119567.

In an alternatively preferred embodiment of the antibody construct of the present invention, the second binding domain capable of binding to the T cell CD3 receptor complex comprises a VH region comprising CDR-H 1, CDR-H2 and CDR-H3 selected from:

- 5 (a) CDR-H1 as depicted in SEQ ID NO: 12 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 13 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 14 of WO 2008/119567;
 - (b) CDR-H1 as depicted in SEQ ID NO: 30 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 31 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 32 of WO 2008/119567;

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- (c) CDR-H1 as depicted in SEQ ID NO: 48 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 49 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 50 of WO 2008/119567;
- (d) CDR-H1 as depicted in SEQ ID NO: 66 of WO 2008/119567, CDR-H2 as depicted in
 SEQ ID NO: 67 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 68 of WO 2008/119567;
 - (e) CDR-H1 as depicted in SEQ ID NO: 84 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 85 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 86 of WO 2008/119567;
- 20 (f) CDR-H1 as depicted in SEQ ID NO: 102 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 103 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 104 of WO 2008/119567;
- (g) CDR-H1 as depicted in SEQ ID NO: 120 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 121 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 122 of WO 2008/119567;
 - (h) CDR-H1 as depicted in SEQ ID NO: 138 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 139 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 140 of WO 2008/119567;
- (i) CDR-H1 as depicted in SEQ ID NO: 156 of WO 2008/119567, CDR-H2 as depicted
 30 in SEQ ID NO: 157 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 158 of WO 2008/119567; and
 - (j) CDR-H1 as depicted in SEQ ID NO: 174 of WO 2008/119567, CDR-H2 as depicted in SEQ ID NO: 175 of WO 2008/119567 and CDR-H3 as depicted in SEQ ID NO: 176 of WO 2008/119567.

It is further preferred for the antibody construct of the present invention that the second binding domain capable of binding to the T cell CD3 receptor complex comprises a VL

region selected from the group consisting of a VL region as depicted in SEQ ID NO: 35, 39, 125, 129, 161 or 165 of WO 2008/119567.

It is alternatively preferred that the second binding domain capable of binding to the T cell CD3 receptor complex comprises a VH region selected from the group consisting of a VH region as depicted in SEQ ID NO: 15, 19, 33, 37, 51, 55, 69, 73, 87, 91, 105, 109, 123, 127, 141, 145, 159, 163, 177 or 181 of WO 2008/119567.

More preferably, the antibody construct of the present invention is characterized by the second binding domain capable of binding to the T cell CD3 receptor complex comprising a VL region and a VH region selected from the group consisting of:

- (a) a VL region as depicted in SEQ ID NO: 17 or 21 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 15 or 19 of WO 2008/119567;
- (b) a VL region as depicted in SEQ ID NO: 35 or 39 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 33 or 37 of WO 2008/119567;

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- (c) a VL region as depicted in SEQ ID NO: 53 or 57 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 51 or 55 of WO 2008/119567;
- (d) a VL region as depicted in SEQ ID NO: 71 or 75 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 69 or 73 of WO 2008/119567;
- 20 (e) a VL region as depicted in SEQ ID NO: 89 or 93 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 87 or 91 of WO 2008/119567;
 - (f) a VL region as depicted in SEQ ID NO: 107 or 111 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 105 or 109 of WO 2008/119567;
- (g) a VL region as depicted in SEQ ID NO: 125 or 129 of WO 2008/119567 and a VH
 region as depicted in SEQ ID NO: 123 or 127 of WO 2008/119567;
 - (h) a VL region as depicted in SEQ ID NO: 143 or 147 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 141 or 145 of WO 2008/119567;
 - (i) a VL region as depicted in SEQ ID NO: 161 or 165 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 159 or 163 of WO 2008/119567; and
- 30 (j) a VL region as depicted in SEQ ID NO: 179 or 183 of WO 2008/119567 and a VH region as depicted in SEQ ID NO: 177 or 181 of WO 2008/119567.

According to a preferred embodiment of the antibody construct of the present invention, in particular the second binding domain capable of binding to the T cell CD3 receptor complex, the pairs of VH-regions and VL-regions are in the format of a single chain antibody (scFv). The VH and VL regions are arranged in the order VH-VL or VL-VH. It is preferred that the

VH-region is positioned N-terminally to a linker sequence. The VL-region is positioned C-terminally of the linker sequence.

A preferred embodiment of the above described antibody construct of the present invention is characterized by the second binding domain capable of binding to the T cell CD3 receptor complex comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 23, 25, 41, 43, 59, 61, 77, 79, 95, 97, 113, 115, 131, 133, 149, 151, 167, 169, 185 or 187 of WO 2008/119567.

- In a preferred embodiment the antibody construct of the invention has an amino acid sequence selected from the group consisting of
 - (a) as depicted in SEQ ID NO: 1138, SEQ ID NO: 1177, SEQ ID NO: 1346, SEQ ID NO: 1359, SEQ ID NO: 1372, SEQ ID NO: 1437, SEQ ID NO: 1450 and SEQ ID NO: 2179;
- (b) as depicted in SEQ ID NO: 1021, SEQ ID NO: 1034, SEQ ID NO: 1047, SEQ ID NO: 1086, SEQ ID NO: 1112, SEQ ID NO: 1125, SEQ ID NO: 1255, SEQ ID NO: 1268, SEQ ID NO: 1281, SEQ ID NO: 1294, SEQ ID NO: 1307, SEQ ID NO: 1659, SEQ ID NO: 1672, SEQ ID NO: 1906, SEQ ID NO: 1919, SEQ ID NO: 1945, SEQ ID NO: 1958, SEQ ID NO: 1971, SEQ ID NO: 1984, SEQ ID NO: 1997, SEQ ID NO: 2010, SEQ ID NO: 2023, SEQ ID NO: 2036, SEQ ID NO: 2049, and SEQ ID NO: 2062;
 - (c) as depicted in SEQ ID NO: 1008, SEQ ID NO: 1060, SEQ ID NO: 1099, SEQ ID NO: 1620, SEQ ID NO: 1633, SEQ ID NO: 1646, SEQ ID NO: 1685, SEQ ID NO: 1698, SEQ ID NO: 1711, SEQ ID NO: 1724, SEQ ID NO: 1737, SEQ ID NO: 1750, SEQ ID NO: 1763, SEQ ID NO: 1776, and SEQ ID NO: 1932;
- (d) as depicted in SEQ ID NO: 982, SEQ ID NO: 1073, SEQ ID NO: 1151, SEQ ID 25 NO: 1164, SEQ ID NO: 1190, SEQ ID NO: 1203, SEQ ID NO: 1216, SEQ ID NO: 1229, SEQ ID NO: 1242, SEQ ID NO: 1320, SEQ ID NO: 1333, SEQ ID NO: 1385, SEQ ID NO: 1398, SEQ ID NO: 1411, SEQ ID NO: 1424, SEQ ID NO: 1474, SEQ ID NO: 1475, SEQ ID NO: 1476, SEQ ID NO: 1483, SEQ ID NO: 1490, SEQ ID NO: 1491, SEQ ID 30 NO: 1492, SEQ ID NO: 1499, SEQ ID NO: 1506, SEQ ID NO: 1513, SEQ ID NO: 1514, SEQ ID NO: 1515, SEQ ID NO: 1516, SEQ ID NO: 1517, SEQ ID NO: 1524, SEQ ID NO: 1531, SEQ ID NO: 1538, SEQ ID NO: 1539, SEQ ID NO: 1540, SEQ ID NO: 1547, SEQ ID NO: 1554, SEQ ID NO: 1555, SEQ ID NO: 1556, SEQ ID NO: 1563, SEQ ID NO: 1570, SEQ ID NO: 1571, SEQ ID NO: 1572, SEQ ID NO: 1573, SEQ ID NO: 1574, SEQ ID NO: 1575, SEQ ID NO: 1576, SEQ ID NO: 1577, SEQ ID NO: 1578, SEQ ID 35 NO: 1579, SEQ ID NO: 1580, SEQ ID NO: 1581, SEQ ID NO: 1789, SEQ ID NO: 1802, SEQ ID NO: 1815, SEQ ID NO: 1828, SEQ ID NO: 1841, SEQ ID NO: 1854, SEQ ID

NO: 1867, SEQ ID NO: 1880, SEQ ID NO: 1893, SEQ ID NO: 2075, SEQ ID NO: 2088, SEQ ID NO: 2101, SEQ ID NO: 2114, SEQ ID NO: 2127, SEQ ID NO: 2140, SEQ ID NO: 2153, SEQ ID NO: 2166, SEQ ID NO: 2192, SEQ ID NO: 2205, and SEQ ID NO: 2218 to 2228; and

5 (e) as depicted in SEQ ID NO: 995, SEQ ID NO: 1594, and SEQ ID NO: 1607.

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The invention further provides a nucleic acid sequence encoding an antibody construct of the invention.

10 Furthermore, the invention provides a vector comprising a nucleic acid sequence of the invention. Moreover, the invention provides a host cell transformed or transfected with the nucleic acid sequence of the invention.

In a further embodiment the invention provides a process for the production of a antibody construct of the invention, said process comprising culturing a host cell of the invention under conditions allowing the expression of the antibody construct of the invention and recovering the produced antibody construct from the culture.

Moreover, the invention provides a pharmaceutical composition comprising an antibody construct of the invention or produced according to the process of the invention

The formulations described herein are useful as pharmaceutical compositions in the treatment, amelioration and/or prevention of the pathological medical condition as described herein in a patient in need thereof. The term "treatment" refers to both therapeutic treatment and prophylactic or preventative measures. Treatment includes the application or administration of the formulation to the body, an isolated tissue, or cell from a patient who has a disease/disorder, a symptom of a disease/disorder, or a predisposition toward a disease/disorder, with the purpose to cure, heal, alleviate, relieve, alter, remedy, ameliorate, improve, or affect the disease, the symptom of the disease, or the predisposition toward the disease.

Those "in need of treatment" include those already with the disorder, as well as those in which the disorder is to be prevented. The term "disease" is any condition that would benefit from treatment with the protein formulation described herein. This includes chronic and acute disorders or diseases including those pathological conditions that predispose the mammal to the disease in question. Non-limiting examples of diseases/disorders to be treated herein include proliferative disease, a tumorous disease, or an immunological disorder.

In some embodiments, the invention provides a pharmaceutical composition comprising a therapeutically effective amount of one or a plurality of the antibody construct of the invention together with a pharmaceutically effective diluents, carrier, solubilizer, emulsifier, preservative, and/or adjuvant. Pharmaceutical compositions of the invention include, but are not limited to, liquid, frozen, and lyophilized compositions.

Preferably, formulation materials are nontoxic to recipients at the dosages and concentrations employed. In specific embodiments, pharmaceutical compositions comprising a therapeutically effective amount of an antibody construct of the invention.

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In certain embodiments, the pharmaceutical composition may contain formulation materials for modifying, maintaining or preserving, for example, the pH, osmolarity, viscosity, clarity, color, isotonicity, odor, sterility, stability, rate of dissolution or release, adsorption or penetration of the composition. In such embodiments, suitable formulation materials include, but are not limited to, amino acids (such as glycine, glutamine, asparagine, arginine, proline, or lysine); antimicrobials; antioxidants (such as ascorbic acid, sodium sulfite or sodium hydrogen-sulfite); buffers (such as borate, bicarbonate, Tris-HCI, citrates, phosphates or other organic acids); bulking agents (such as mannitol or glycine); chelating agents (such as ethylenediamine tetraacetic acid (EDTA)); complexing agents (such as caffeine, polyvinylpyrrolidone, beta-cyclodextrin or hydroxypropyl-beta-cyclodextrin); monosaccharides; disaccharides; and other carbohydrates (such as glucose, mannose or dextrins); proteins (such as serum albumin, gelatin or immunoglobulins); coloring, flavoring and diluting agents; emulsifying agents; hydrophilic polymers (such as polyvinylpyrrolidone); low molecular weight polypeptides; salt-forming counterions (such as sodium); preservatives (such as benzalkonium chloride, benzoic acid, salicylic acid, thimerosal, phenethyl alcohol, methylparaben, propylparaben, chlorhexidine, sorbic acid or hydrogen peroxide); solvents (such as glycerin, propylene glycol or polyethylene glycol); sugar alcohols (such as mannitol or sorbitol); suspending agents; surfactants or wetting agents (such as pluronics, PEG, sorbitan esters, polysorbates such as polysorbate 20, polysorbate, triton, tromethamine, lecithin, cholesterol, tyloxapal); stability enhancing agents (such as sucrose or sorbitol); tonicity enhancing agents (such as alkali metal halides, preferably sodium or potassium chloride, mannitol sorbitol); delivery vehicles; diluents; excipients and/or pharmaceutical adjuvants. See, REMINGTON'S PHARMACEUTICAL SCIENCES, 18" Edition, (A. R. Genrmo, ed.), 1990, Mack Publishing Company.

In certain embodiments, the optimal pharmaceutical composition will be determined by one skilled in the art depending upon, for example, the intended route of administration, delivery format and desired dosage. See, for example, REMINGTON'S PHARMACEUTICAL SCIENCES, supra. In certain embodiments, such compositions may influence the physical state, stability, rate of in vivo release and rate of in vivo clearance of the antigen binding proteins of the invention. In certain embodiments, the primary vehicle or carrier in a pharmaceutical composition may be either aqueous or non-aqueous in nature. For example, a suitable vehicle or carrier may be water for injection, physiological saline solution or artificial cerebrospinal fluid, possibly supplemented with other materials common in compositions for parenteral administration. Neutral buffered saline or saline mixed with serum albumin are further exemplary vehicles. In specific embodiments, pharmaceutical compositions comprise Tris buffer of about pH 7.0-8.5, or acetate buffer of about pH 4.0-5.5, and may further include sorbitol or a suitable substitute therefore. In certain embodiments of the invention, human antibody or antigen binding fragment thereof of the invention or the antibody construct of the invention compositions may be prepared for storage by mixing the selected composition having the desired degree of purity with optional formulation agents (REMINGTON'S PHARMACEUTICAL SCIENCES, supra) in the form of a lyophilized cake or an aqueous solution. Further, in certain embodiments, the human antibody or antigen binding fragment thereof of the invention or the antibody construct of the invention may be formulated as a lyophilizate using appropriate excipients such as sucrose.

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The pharmaceutical compositions of the invention can be selected for parenteral delivery. Alternatively, the compositions may be selected for inhalation or for delivery through the digestive tract, such as orally. Preparation of such pharmaceutically acceptable compositions is within the skill of the art. The formulation components are present preferably in concentrations that are acceptable to the site of administration. In certain embodiments, buffers are used to maintain the composition at physiological pH or at a slightly lower pH, typically within a pH range of from about 5 to about 8.

30 When parenteral administration is contemplated, the therapeutic compositions for use in this invention may be provided in the form of a pyrogen-free, parenterally acceptable aqueous solution comprising the desired human antibody or antigen binding fragment thereof of the invention or the antibody construct of the invention in a pharmaceutically acceptable vehicle. A particularly suitable vehicle for parenteral injection is sterile distilled water in which the antibody construct of the invention is formulated as a sterile, isotonic solution, properly preserved. In certain embodiments, the preparation can involve the formulation of the desired molecule with an agent, such as injectable microspheres, bio-erodible particles,

polymeric compounds (such as polylactic acid or polyglycolic acid), beads or liposomes, that may provide controlled or sustained release of the product which can be delivered via depot injection. In certain embodiments, hyaluronic acid may also be used, having the effect of promoting sustained duration in the circulation. In certain embodiments, implantable drug delivery devices may be used to introduce the desired antigen binding protein.

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Additional pharmaceutical compositions will be evident to those skilled in the art, including formulations involving h the antibody construct of the invention in sustained- or controlleddelivery formulations. Techniques for formulating a variety of other sustained- or controlleddelivery means, such as liposome carriers, bio-erodible microparticles or porous beads and depot injections, are also known to those skilled in the art. See, for example, International Patent Application No. PCT/US93/00829, which is incorporated by reference and describes controlled release of porous polymeric microparticles for delivery of pharmaceutical compositions. Sustained-release preparations may include semipermeable polymer matrices in the form of shaped articles, e.g., films, or microcapsules. Sustained release matrices may include polyesters, hydrogels, polylactides (as disclosed in U.S. Pat. No. 3,773,919 and European Patent Application Publication No. EP 058481, each of which is incorporated by reference), copolymers of L-glutamic acid and gamma ethyl-L-glutamate (Sidman et al., 1983, Biopolymers 2:547-556), poly (2-hydroxyethyl-methacrylate) (Langer et al., 1981, J. Biomed. Mater. Res. 15:167-277 and Langer, 1982, Chem. Tech. 12:98-105), ethylene vinyl acetate (Langer et al., 1981, supra) or poly-D(-)-3-hydroxybutyric acid (European Patent Application Publication No. EP 133,988). Sustained release compositions may also include liposomes that can be prepared by any of several methods known in the art. See, e.g., Eppstein et al., 1985, Proc. Natl. Acad. Sci. U.S.A. 82:3688-3692; European Patent Application Publication Nos. EP 036,676; EP 088,046 and EP 143,949, incorporated by reference.

Pharmaceutical compositions used for in vivo administration are typically provided as sterile preparations. Sterilization can be accomplished by filtration through sterile filtration membranes. When the composition is lyophilized, sterilization using this method may be conducted either prior to or following lyophilization and reconstitution. Compositions for parenteral administration can be stored in lyophilized form or in a solution. Parenteral compositions generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

Aspects of the invention includes self-buffering antibody construct of the invention formulations, which can be used as pharmaceutical compositions, as described in international patent application WO 06138181A2 (PCT/US2006/022599), which is incorporated by reference in its entirety herein.

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As discussed above, certain embodiments provide antibody construct of the invention protein compositions, particularly pharmaceutical compositions of the invention, that comprise, in addition to the antibody construct of the invention, one or more excipients such as those illustratively described in this section and elsewhere herein. Excipients can be used in the invention in this regard for a wide variety of purposes, such as adjusting physical, chemical, or biological properties of formulations, such as adjustment of viscosity, and or processes of the invention to improve effectiveness and or to stabilize such formulations and processes against degradation and spoilage due to, for instance, stresses that occur during manufacturing, shipping, storage, pre-use preparation, administration, and thereafter.

A variety of expositions are available on protein stabilization and formulation materials and methods useful in this regard, such as Arakawa et al., "Solvent interactions in pharmaceutical formulations," Pharm Res. 8(3): 285-91 (1991); Kendrick et al., "Physical stabilization of proteins in aqueous solution," in: RATIONAL DESIGN OF STABLE PROTEIN FORMULATIONS: THEORY AND PRACTICE, Carpenter and Manning, eds. Pharmaceutical Biotechnology. 13: 61-84 (2002), and Randolph et al., "Surfactant-protein interactions," Pharm Biotechnol. 13: 159-75 (2002), each of which is herein incorporated by reference in its entirety, particularly in parts pertinent to excipients and processes of the same for self-buffering protein formulations in accordance with the current invention, especially as to protein pharmaceutical products and processes for veterinary and/or human medical uses.

Salts may be used in accordance with certain embodiments of the invention to, for example, adjust the ionic strength and/or the isotonicity of a formulation and/or to improve the solubility and/or physical stability of a protein or other ingredient of a composition in accordance with the invention.

As is well known, ions can stabilize the native state of proteins by binding to charged residues on the protein's surface and by shielding charged and polar groups in the protein and reducing the strength of their electrostatic interactions, attractive, and repulsive interactions. Ions also can stabilize the denatured state of a protein by binding to, in particular, the denatured peptide linkages (--CONH) of the protein. Furthermore, ionic

interaction with charged and polar groups in a protein also can reduce intermolecular electrostatic interactions and, thereby, prevent or reduce protein aggregation and insolubility.

lonic species differ significantly in their effects on proteins. A number of categorical rankings of ions and their effects on proteins have been developed that can be used in formulating pharmaceutical compositions in accordance with the invention. One example is the Hofmeister series, which ranks ionic and polar non-ionic solutes by their effect on the conformational stability of proteins in solution. Stabilizing solutes are referred to as "kosmotropic." Destabilizing solutes are referred to as "chaotropic." Kosmotropes commonly are used at high concentrations (e.g., >1 molar ammonium sulfate) to precipitate proteins from solution ("salting-out"). Chaotropes commonly are used to denture and/or to solubilize proteins ("salting-in"). The relative effectiveness of ions to "salt-in" and "salt-out" defines their position in the Hofmeister series.

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Free amino acids can be used in the antibody construct of the invention formulations in accordance with various embodiments of the invention as bulking agents, stabilizers, and antioxidants, as well as other standard uses. Lysine, proline, serine, and alanine can be used for stabilizing proteins in a formulation. Glycine is useful in lyophilization to ensure correct cake structure and properties. Arginine may be useful to inhibit protein aggregation, in both liquid and lyophilized formulations. Methionine is useful as an antioxidant.

Polyols include sugars, e.g., mannitol, sucrose, and sorbitol and polyhydric alcohols such as, for instance, glycerol and propylene glycol, and, for purposes of discussion herein, polyethylene glycol (PEG) and related substances. Polyols are kosmotropic. They are useful stabilizing agents in both liquid and lyophilized formulations to protect proteins from physical and chemical degradation processes. Polyols also are useful for adjusting the tonicity of formulations.

Among polyols useful in select embodiments of the invention is mannitol, commonly used to ensure structural stability of the cake in lyophilized formulations. It ensures structural stability to the cake. It is generally used with a lyoprotectant, e.g., sucrose. Sorbitol and sucrose are among preferred agents for adjusting tonicity and as stabilizers to protect against freeze-thaw stresses during transport or the preparation of bulks during the manufacturing process. Reducing sugars (which contain free aldehyde or ketone groups), such as glucose and lactose, can glycate surface lysine and arginine residues. Therefore, they generally are not among preferred polyols for use in accordance with the invention. In addition, sugars that form such reactive species, such as sucrose, which is hydrolyzed to fructose and glucose

under acidic conditions, and consequently engenders glycation, also is not among preferred polyols of the invention in this regard. PEG is useful to stabilize proteins and as a cryoprotectant and can be used in the invention in this regard.

5 Embodiments of the antibody construct of the invention formulations further comprise surfactants. Protein molecules may be susceptible to adsorption on surfaces and to denaturation and consequent aggregation at air-liquid, solid-liquid, and liquid-liquid interfaces. These effects generally scale inversely with protein concentration. These deleterious interactions generally scale inversely with protein concentration and typically are exacerbated by physical agitation, such as that generated during the shipping and handling of a product.

Surfactants routinely are used to prevent, minimize, or reduce surface adsorption. Useful surfactants in the invention in this regard include polysorbate 20, polysorbate 80, other fatty acid esters of sorbitan polyethoxylates, and poloxamer 188.

Surfactants also are commonly used to control protein conformational stability. The use of surfactants in this regard is protein-specific since, any given surfactant typically will stabilize some proteins and destabilize others.

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Polysorbates are susceptible to oxidative degradation and often, as supplied, contain sufficient quantities of peroxides to cause oxidation of protein residue side-chains, especially methionine. Consequently, polysorbates should be used carefully, and when used, should be employed at their lowest effective concentration. In this regard, polysorbates exemplify the general rule that excipients should be used in their lowest effective concentrations.

Embodiments of the antibody construct of the invention formulations further comprise one or more antioxidants. To some extent deleterious oxidation of proteins can be prevented in pharmaceutical formulations by maintaining proper levels of ambient oxygen and temperature and by avoiding exposure to light. Antioxidant excipients can be used as well to prevent oxidative degradation of proteins. Among useful antioxidants in this regard are reducing agents, oxygen/free-radical scavengers, and chelating agents. Antioxidants for use in therapeutic protein formulations in accordance with the invention preferably are water-soluble and maintain their activity throughout the shelf life of a product. EDTA is a preferred antioxidant in accordance with the invention in this regard.

Antioxidants can damage proteins. For instance, reducing agents, such as glutathione in particular, can disrupt intramolecular disulfide linkages. Thus, antioxidants for use in the invention are selected to, among other things, eliminate or sufficiently reduce the possibility of themselves damaging proteins in the formulation.

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Formulations in accordance with the invention may include metal ions that are protein cofactors and that are necessary to form protein coordination complexes, such as zinc necessary to form certain insulin suspensions. Metal ions also can inhibit some processes that degrade proteins. However, metal ions also catalyze physical and chemical processes that degrade proteins.

Magnesium ions (10-120 mM) can be used to inhibit isomerization of aspartic acid to isoaspartic acid. Ca⁺² ions (up to 100 mM) can increase the stability of human deoxyribonuclease. Mg⁺², Mn⁺², and Zn⁺², however, can destabilize rhDNase. Similarly, Ca⁺² and Sr⁺² can stabilize Factor VIII, it can be destabilized by Mg⁺², Mn⁺² and Zn⁺², Cu⁺² and Fe⁺², and its aggregation can be increased by Al⁺³ ions.

Embodiments of the antibody construct of the invention formulations further comprise one or more preservatives. Preservatives are necessary when developing multi-dose parenteral formulations that involve more than one extraction from the same container. Their primary function is to inhibit microbial growth and ensure product sterility throughout the shelf-life or term of use of the drug product. Commonly used preservatives include benzyl alcohol, phenol and m-cresol. Although preservatives have a long history of use with small-molecule parenterals, the development of protein formulations that includes preservatives can be challenging. Preservatives almost always have a destabilizing effect (aggregation) on proteins, and this has become a major factor in limiting their use in multi-dose protein formulations. To date, most protein drugs have been formulated for single-use only. However, when multi-dose formulations are possible, they have the added advantage of enabling patient convenience, and increased marketability. A good example is that of human growth hormone (hGH) where the development of preserved formulations has led to commercialization of more convenient, multi-use injection pen presentations. At least four such pen devices containing preserved formulations of hGH are currently available on the market. Norditropin (liquid, Novo Nordisk), Nutropin AQ (liquid, Genentech) & Genotropin (lyophilized--dual chamber cartridge, Pharmacia & Upjohn) contain phenol while Somatrope (Eli Lilly) is formulated with m-cresol. Several aspects need to be considered during the formulation and development of preserved dosage forms. The effective preservative concentration in the drug product must be optimized. This requires testing a given

preservative in the dosage form with concentration ranges that confer anti-microbial effectiveness without compromising protein stability.

As might be expected, development of liquid formulations containing preservatives are more challenging than lyophilized formulations. Freeze-dried products can be lyophilized without the preservative and reconstituted with a preservative containing diluent at the time of use. This shortens the time for which a preservative is in contact with the protein, significantly minimizing the associated stability risks. With liquid formulations, preservative effectiveness and stability should be maintained over the entire product shelf-life (about 18 to 24 months). An important point to note is that preservative effectiveness should be demonstrated in the final formulation containing the active drug and all excipient components.

The antibody construct of the invention generally will be designed for specific routes and methods of administration, for specific administration dosages and frequencies of administration, for specific treatments of specific diseases, with ranges of bio-availability and persistence, among other things. Formulations thus may be designed in accordance with the invention for delivery by any suitable route, including but not limited to orally, aurally, opthalmically, rectally, and vaginally, and by parenteral routes, including intravenous and intraarterial injection, intramuscular injection, and subcutaneous injection.

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Once the pharmaceutical composition has been formulated, it may be stored in sterile vials as a solution, suspension, gel, emulsion, solid, crystal, or as a dehydrated or lyophilized powder. Such formulations may be stored either in a ready-to-use form or in a form (e.g., lyophilized) that is reconstituted prior to administration. The invention also provides kits for producing a single-dose administration unit. The kits of the invention may each contain both a first container having a dried protein and a second container having an aqueous formulation. In certain embodiments of this invention, kits containing single and multichambered pre-filled syringes (e.g., liquid syringes and lyosyringes) are provided. The therapeutically effective amount of an antibody construct of the invention protein-containing pharmaceutical composition to be employed will depend, for example, upon the therapeutic context and objectives. One skilled in the art will appreciate that the appropriate dosage levels for treatment will vary depending, in part, upon the molecule delivered, the indication for which the antibody construct of the invention is being used, the route of administration, and the size (body weight, body surface or organ size) and/or condition (the age and general health) of the patient. In certain embodiments, the clinician may titer the dosage and modify the route of administration to obtain the optimal therapeutic effect. A typical dosage may range from about 0.1 µg/kg to up to about 30 mg/kg or more, depending on the factors

mentioned above. In specific embodiments, the dosage may range from 1.0 μ g/kg up to about 20 mg/kg, optionally from 10 μ g/kg up to about 10 mg/kg or from 100 μ g/kg up to about 5 mg/kg.

- A therapeutic effective amount of an antibody construct of the invention preferably results in a decrease in severity of disease symptoms, in increase in frequency or duration of disease symptom-free periods or a prevention of impairment or disability due to the disease affliction. For treating CDH19-expressing tumors, a therapeutically effective amount of the antibody construct of the invention, e.g. an anti-CDH19/CD3 antibody construct, preferably inhibits cell growth or tumor growth by at least about 20%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, or at least about 90% relative to untreated patients. The ability of a compound to inhibit tumor growth may be evaluated in an animal model predictive of efficacy in human tumors.
- Pharmaceutical compositions may be administered using a medical device. Examples of medical devices for administering pharmaceutical compositions are described in U.S. Patent Nos. 4,475,196; 4,439,196; 4,447,224; 4,447, 233; 4,486,194; 4,487,603; 4,596,556; 4,790,824; 4,941,880; 5,064,413; 5,312,335; 5,312,335; 5,383,851; and 5,399,163, all incorporated by reference herein.

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- In one embodiment the invention provides the antibody construct of the invention or produced according to the process of the invention for use in the prevention, treatment or amelioration of a melanoma disease or metastatic melanoma disease.
- The invention also provides a method for the treatment or amelioration of a melanoma disease or metastatic melanoma disease, comprising the step of administering to a subject in need thereof the antibody construct of the invention or produced according to the process of the invention.
- 30 In a preferred embodiment method of use of the invention the melanoma disease or metastatic melanoma disease is selected from the group consisting of superficial spreading melanoma, lentigo maligna, lentigo maligna melanoma, acral lentiginous melanoma and nodular melanoma.
- In a further embodiment, the invention provides a kit comprising an antibody construct of the invention, or produced according to the process of the invention, a vector of the invention, and/or a host cell of the invention.

It should be understood that the inventions herein are not limited to particular methodology, protocols, or reagents, as such can vary. The discussion and examples provided herein are presented for the purpose of describing particular embodiments only and are not intended to limit the scope of the present invention, which is defined solely by the claims.

All publications and patents cited throughout the text of this specification (including all patents, patent applications, scientific publications, manufacturer's specifications, instructions, etc.), whether supra or infra, are hereby incorporated by reference in their entirety. Nothing herein is to be construed as an admission that the invention is not entitled to antedate such disclosure by virtue of prior invention. To the extent the material incorporated by reference contradicts or is inconsistent with this specification, the specification will supersede any such material.

Examples:

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The following examples are provided for the purpose of illustrating specific embodiments or features of the present invention. These examples should not be construed as to limit the scope of this invention. The examples are included for purposes of illustration, and the present invention is limited only by the claims.

20 Example 1 – Fully human monoclonal antibodies against CDH19

1.1 Immunization:

Fully human antibodies to Cadherin-19 (CDH19) were generated using XENOMOUSE® technology, transgenic mice engineered to express diverse repertoires of fully human IgGκ and IgGλ antibodies of the corresponding isotype. (United States Patent Nos. 6,114,598;

6,162,963; 6,833,268; 7,049,426; 7,064,244, which are incorporated herein by reference in their entirety; Green *et al.*, 1994, *Nature Genetics* 7:13-21; Mendez *et al.*, 1997, *Nature Genetics* 15:146-156; Green and Jakobovitis, 1998, *J. Ex. Med.* 188:483-495; Kellermann and Green, *Current Opinion in Biotechnology* 13, 593-597, 2002).

Mice were immunized with multiple forms of Cadherin-19 immunogen, including: (1) full length human and cynomologous ("cyno") monkey cadherin-19, (2) secreted Cadherin-19 ecto-domain (amino acids 1-596), and (3) a truncated membrane bound form of human cadherin-19 (amino acids 1-624). Mice were immunized over a period of 8 to 10 weeks with a range of 16-18 boosts.

Sera were collected at approximately 5 and 9 weeks after the first injection and specific titers were determined by FACs staining of recombinant Cadherin-19 receptor transiently expressed on CHO-S cells. A total of 37 animals were identified with specific immune responses, these animals were pooled into 3 groups and advanced to antibody generation.

1.2 Preparation of Monoclonal Antibodies

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Animals exhibiting suitable titers were identified, and lymphocytes were obtained from draining lymph nodes and, if necessary, pooled for each cohort. Lymphocytes were dissociated from lymphoid tissue by grinding in a suitable medium (for example, Dulbecco's Modified Eagle Medium (DMEM); obtainable from Invitrogen, Carlsbad, CA) to release the cells from the tissues, and suspended in DMEM. B cells were selected and/or expanded using standard methods, and fused with suitable fusion partner using techniques that were known in the art.

After several days of culture, the hybridoma supernatants were collected and subjected to screening assays as detailed in the examples below, including confirmation of binding to human and cynomologous monkey as well as the ability to kill cell lines in secondary antibody-drug conjugate Bioassays. Hybridoma lines that were identified to have the binding and functional properties of interest were then further selected and subjected to standard cloning and subcloning techniques. Clonal lines were expanded in vitro, and the secreted human antibodies obtained for analysis and V gene sequencing was performed.

1.3 Selection of Cadherin-19 receptor specific binding antibodies by FMAT

After 14 days of culture, hybridoma supernatants were screened for CDH19-specific monoclonal antibodies by Fluorometric Microvolume Assay Technology (FMAT) (Applied Biosystems, Foster City, CA). The supernatants were screened against adherent CHO cells transiently transfected with human Cadherin-19 and counter screened against CHO cells transiently transfected with the same expression plasmid that did not contain the Cadherin-19 gene.

After multiple screening campaigns, a panel of 1570 anti-Cadherin-19 binding hybridoma lines were identified and advanced to further characterization assays.

Example 2 – Assessment of Fully human monoclonal antibodies against CDH19

2.1 Additional Binding Characterization by Flow Cytometry (FACs)

- 30 FACS binding assays were performed to evaluate the binding of the anti-Cadherin-19 receptor specific antibodies to endogenous Cadherin-19 receptor expressed on the CHL-1 tumor cell lines. In addition, cross-reactive binding to murine and cynomologous monkey Cadherin-19 orthologues was also evaluated by FACs using recombinant forms of the various receptors transiently expressed on 293T cells.
- FACs assays were performed by incubating hybridoma supernatants with 10,000 to 25,000 cells in PBS/2%Fetal bovine serum/2mM Calcium Chloride at 4°C for one hour followed by two washes with PBS/2%Fetal bovine serum/2mM Calcium Chloride. Cells were then treated

with florochrome-labeled secondary antibodies at 4°C followed by one wash. The cells were resuspended in 50µl of PBS/2%FBS and antibody binding was analyzed using a FACSCalibur™ instrument.

5 2.2 Antibody drug conjugate screening of fully human antibodies derived from XenoMouse® hybridomas

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Cell killing through antibody drug conjugates requires the delivery of the conjugate into a cell through internalization and the catabolism of the drug-conjugate into a form that it is toxic to the cell. To identify antibodies with these properties, CDH19-positive cell lines (Colo-699 or CHL-1) were seeded at low cell densities and allowed to adhere overnight in a 384 well plate. XENOMOUSE® hybridoma samples containing fully human anti-CDH19 antibodies were then added to these cells in the presence of a high concentration of a goat anti-human Fc monovalent Fab conjugated with DM1 (DM1-Fab) at a relatively low drug-antibody ratio (DAR) (~1.3). The cells were incubated for 96 hours at 37°C and 5% CO₂ in the presence of the antibody samples and the DM1-Fab. At the end of this time, the cell viability was assessed using the CellTiter-Glo® Luminescent Cell Viability reagent (Promega) according to manufacturer's recommendations.

An example of the cell viability data with the Colo-699 cells is shown in Figure 1 and Figure 2. The antibodies capable of delivering the DM1-Fab to the cells and inhibiting the cell growth read out with a lower luminescent signal (RLU). The top antibodies of interest from this screen are observed in the lower left corner of Fig. 1 and are denoted as open circles. These antibodies were taken forward into a cell viability assay on CHL-1 cells. The average cell viability data from the CHL-1 assay is plotted against the average cell viability data from the Colo-699 assay (Fig. 2). The antibodies that had activity on both the Colo-699 and the CHL-1 cells are denoted as open circles on the left-hand side of the Figure 2.

This assay was run concurrently with the FACs antibody binding assay above (2.2), and the results from these two studies were used to select the antibodies for further characterization. In total, 1570 antibodies were run through these cell based viability assays and approximately 44 antibodies were selected on the bases of *in vitro* cell killing and/or antibody binding for sub-cloning,V gene sequencing and expressed in recombinant form for further characterization assays as described below.

These 44 antibodies were again assayed as in Example 2 and 19 antibodies were selected that contained unique sequences. Of these 19 antibodies, 18 antibodies were analyzed and their properties characterized in Table 2 below. The data in this table was generated using FACs binding on recombinant human and cynomologous CDH-19, +/- Calcium (Ca⁺²⁾ binding data on 293/CDH-19 transfectants, binding to endogenous CDH-19 on CHL-1 and Colo699

tumor cells and competition with the antibody designated as 4A9 in the table. These experiments provided the further characterizations for the grouping of these antibodies into 5 groups or bins.

5 Table 2 –Binning of Lead panel using Antibody Binding Information

Bin ID	LMR Sequence/ Ab ID	Clone ID	Bin Characteristics
1	13589	4A9	High Endogenous binding, Calcium insensitive, sequence clustered,
'	13591	4F7	moderate cyno complete 4A9 competitor
	13885	19B5	
	13880	25F8	
	13882	26D1	High Endogenous binding, Calcium
2	13881	26F12=27B3	insensitive, sequence clustered, Good cyno, partial 4A9 competitor
	13878	16H2=20D3=23E7	gire, partial in a compositor
	13879	22D1	
	13877	22G10	High Endogenous binding, moderate 293
	13874	17H8=23B6=28D10	binding, Calcium insensitive, 2 sequence
3	13883	25G10	clusters, moderate cyno, partial 4A9 competitor, 22G10 best binder in
	13875	16C1	bin.
	13590	4B10	
	13586	4F3	Low Endogenous and recombinant
4	13592	4A2	binding, Calcium sensitive, sequence diverse group, comparable cyno, No 4A9
	13884	23A10	competition
	13588	2G6	·
5	13876 16A4		Best endogenous binder, moderate recombinant binder, calcium insensitive, very weak cyno, No 4A9 competition.

Of these 18 antibodies. 8 antibodies were selected for further analysis of their epitope binding as described below. At least one representative antibody from each bin was selected for further analysis.

Example 3 – Epitope Prediction

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Epitope Prediction by 4A9 Antibody Competition and by Human/Mouse Cadherin-19 Chimeras

A 4A9 binding competition method was developed to identify antibodies that compete with 4A9 binding. In 96-well V-bottom plates (Sarstedt #82.1583.001), 50,000 transiently

transfected 293T cells were incubated with 5ug/ml of purified anti-CDH19 antibodies for 1hr at 4oC followed by one wash with PBS/2%FBS. 25µl of 5µg/ml Alexa647-labelled 4A9 was then added to each well and the plates incubated for 1 hour at 4°C. Cells were then washed two times and the amount of cell associated Alexa647-labelled 4A9 was quantitated by flow cytometry.

The experiments included negative controls consisting of PBS/2%FBS only. The average signal observed in these negative control experiments was adopted as the maximum possible signal for the assay. Antibodies were compared to this maximum signal and a percent inhibition was calculated for each well (% Inhibition = (1-(FL4 Geomean with the anti-CDH19 antibodies/Maximum FL4 Geomean signal)).

Domain binding was determined by flow cytometry as above on 293T cells transiently transfected with plasmids consisting of single or dual human CDH19 cadherin repeat domain replacements into the mouse Cadherin19 backbone cloned into the pTT5 expression vector immediately preceded by native human or murine CDH19 leader sequences and a Flag tag (SEQ ID NO: 968). The experiment included assaying the anti-CDH19 antibodies against mouse Cadherin19 to determine suitability for binning on these human/mouse chimeras.

The data from these experiments are presented in the Table below entitled as follows:

Table 3 – Calcium Sensitive Binding and Epitope Prediction Summary

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	Predicted Epitope Region						44-141								250-364			:	un- oldcapisse	assignable	Unassigned	epitope	
Mu EC1- 5	-			ı	1	•		•							•	1		+	+	+	•		+
Hu EC5	I		•		•	•	•	•						•	•	1	•	+	+	+	-	•	+
Hu EC4- 5	Ø							1						1		1		+	+	+			+
Hu EC3	L				1			1			1		+	+	+	+	+	+	+	+			+
Hu EC2- 3	ш			ı		1		1			ı		+	+	+	+	+	+	+	+			+
Hu EC2	Q							•										+	+	+	•		+
Hu EC1-	ပ	+	+	+	+	+	+	+	+	+	+	+		1		1		+	+	+	+	+	+
Hu EC1	В	+	+	+	+	+	+	+	+	+	+	+						+	+	+	+	+	+
Hu EC1- 5	∢	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
	Competes with 4A9 (13589)	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	No	N _O	N _o	No	No	S _O	No	No	No	
	Ca2+ Sensitive Binding	No	N _o	N _o	No	No	No	°N	No	No	No	No	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	No	No	
	Bin	-	_	-	2	7	7	7	7	က	က	က	4	4	4	4	4	4	4	4	5	5	
	Ab ID	13589	14056	14057	13880	14094	14096	13882	14088	13874	14045	14048	13592	14026	13590	14055	14054	13588	14304	14039	13876	14071	-FLAG
	Clone	4A9			25F8			26D1		17H8			4A2		4B10			2 G6			16A4		Rat anti-FLAG

Legend Table 3

Human and/or murine chimera constructs

A = huCDH19(44-772) (see SEQ ID NO: 944)

B = huCDH19(44-141)::muCDH19(140-770) (see SEQ ID NO: 952)

5 C = huCDH19(44-249)::muCDH19(248-770) (see SEQ ID NO: 954)

D = muCDH19(44-139)::huCDH19(142-249)::muCDH19(248-770) (see SEQ ID NO: 956)

E = muCDH19(44-139)::huCDH19(142-364)::muCDH19(363-770) (see SEQ ID NO: 958)

F = muCDH19(44-247)::huCDH19(250-364)::muCDH19(363-770) (see SEQ ID NO: 960)

G = muCDH19(44-362)::huCDH19(365-772) (see SEQ ID NO: 962)

10 H = muCDH19(44-461)::huCDH19(464-772) (see SEQ ID NO: 964)

I = muCDH19(44-770) (see SEQ ID NO: 966)

Epitope Prediction by Human/Chicken Cadherin-19 Chimeras

Domain binding was determined by flow cytometry on 293T cells transiently transfected with plasmids consisting of single human CDH19 cadherin repeat domain replacements into the chicken Cadherin19 backbone cloned into the pTT5 expression vector immediately preceded by native human or chicken CDH19 leader sequences and a Flag tag. The experiment included assaying a subset of anti-CDH19 antibodies against chicken Cadherin19 to determine suitability for binning on these human/chicken chimeras.

The following binding assay was completed in presence of 2mM CaCl2. In 96-well V-bottom plates (Costar 3897), 50,000 transiently transfected 293T cells were incubated with 5ug/ml of purified anti-CDH19 antibodies for 1hr at 4oC followed by two washes with PBS/2%FBS. 50µl of 5µg/ml Alexa647-labelled anti-human IgG secondary antibody (Jackson Immuno 109-605-098) and 2ug/ml 7AAD (Sigma A9400) was then added to each well and the plates incubated for 15 minutes at 4oC. Cells were then washed one time and the amount of cell associated Alexa647-labelled Ab was quantitated by flow cytometry. The experiments included mock transfected controls. The data from these experiments are presented in the Table below, n.d. = not determined.

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Table4 - Antibody Bin C Epitope Prediction Summary

			Hu EC1-5	Ck EC1-5	Hu EC1	Hu EC2	Hu EC3	Hu EC5	
Clone ID	Ab. ID	Bin	Α	J	К	L	M	o	Predicted Epitope Region
4A9	13589	1	+	-	+	-	-	-	
26F12	13881	2	+	-	+	-	-	-	
25F8	14096	2	+	-	+	-	-	-	44-141
26D1	13882	2	+	-	+	-	-	-	Bin A
17H8	13874	3	+	-	+	-	-	-	
16 A 4	14071	5	+	-	+	-	-	-	
4A2	13592	4	+	-	-	-	+	-	
4B10	13590	4	+	-	-	-	+	-	250-364
2G6	13588	4	+	-	-	-	+	-	Bin B
23A10	14077	4	+	-	-	-	+	-	1
Rat an	Rat anti-FLAG		+	+	+	+	+	+	control

Positive Binding (+)
Negative Binding (-)

Legend Table 4

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Human and/or chicken chimera constructs

5 A = huCDH19(44-772) (see SEQ ID NO: 944)

J = ckCDH19(44-776) (see SEQ ID NO: 1451)

K = huCDH19(44-141)::ckCDH19(142-776) (see SEQ ID NO: 1452)

L = ckCDH19(44-141)::huCDH19(142-249)::ckCDH19(250-776) (see SEQ ID NO: 1453)

M = ckCDH19(44-249)::huCDH19(250-364)::ckCDH19(365-776) (see SEQ ID NO: 1454)

10 N = ckCDH19(44-364)::huCDH19(365-463)::ckCDH19(469-776) (see SEQ ID NO: 1455)

O = ckCDH19(44-468)::huCDH19(464-772) (see SEQ ID NO: 1456)

Epitope Prediction by macaque/dog or rat/macaque Cadherin-19 Chimeras

Domain binding was determined by flow cytometry on 293T cells transiently transfected with plasmids consisting of rhesus macaque CDH19 cadherin repeat domain 1 or segments domain 1 (designated EC1a, EC1b, EC1c) replacements into the dog Cadherin19 backbone, or rat CDH19 cadherin repeat domain 2 replacement into the rhesus Cadherin19 backbone cloned into the pTT5 expression vector immediately preceded by native rhesus or canine CDH19 leader sequences and a Flag tag. The experiment included assaying a subset of anti-CDH19 antibodies against dog, rat and macaque Cadherin19 to determine suitability for binning on these macaque/dog and rat/rhesus chimeras.

The following binding assay was completed in presence of 2mM CaCl2. In 96-well V-bottom plates (Costar 3897), 50,000 transiently transfected 293T cells were incubated with 5ug/ml of purified anti-CDH19 antibodies for 1hr at 4oC followed by two washes with PBS/2%FBS. 50µl of 5µg/ml Alexa647-labelled anti-human IgG secondary antibody (Jackson Immuno 109-605-098) and 2ug/ml 7AAD (Sigma A9400) was then added to each well and the plates incubated for 15 minutes at 4oC. Cells were then washed one time and the amount of cell associated Alexa647-labelled Ab was quantitated by flow cytometry. The experiments included mock transfected controls. The data from these experiments are presented in the Table below, n.d. = not determined.

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Table 5 – Antibody BinA Epitope prediction Summary

			Rh	Ca	rh	rh	rh	ra	Ra	
			EC1-5	EC1-5	EC1	EC1a	EC1b	EC2	EC1-5	
										Predicted
Clone ID	Ab. ID	Bin	P	Q	R	S	Т	V	W	Epitope
										Region
4A9	13589	1	+	_	+	_	_	_	_	44-141
47.5	13303		•		'					Bin A.1
26F12	13881	2	+	-	+	+	+	-	-	44-141
25F8	14096	2	+	-	+	+	+	-	-	Bin A.2
26D1	13882	2	+	-	+	+	+	-	-	(44-114)
17H8	13874	3	+	_	+	+	_	_	_	44-141
17110	13074		•		'	,				Bin A.3
16A4	14071	5	+	-	+	+	-	n.d.	+	(44-65)
4A2	13592	4	+	-	n.d.	n.d.	n.d.	n.d.	+	
4B10	13590	4	+	+	n.d.	n.d.	n.d.	n.d.	+	250-364
2G6	13588	4	+	+	n.d.	n.d.	n.d.	n.d.	+	Bin B
23A10	14077	4	+	+	n.d.	n.d.	n.d.	n.d.	+	
Rat an	Rat anti-FLAG		+	+	+	+	+	+	+	

Positive Binding (+) Negative Binding (-) Not Determined (n.d.)

Legend Table 5

Rhesus macaque, dog, and/or rat chimera constructs

15 P = rhCDH19(44-772) (see SEQ ID NO: 1457)

Q = caCDH19(44-770) (see SEQ ID NO: 1458)

R = rhCDH19(44-141)::caCDH19(141-770) (see SEQ ID NO: 1459)

S = rhCDH19(44-65)::caCDH19(65-770) (see SEQ ID NO: 1460)

T = caCDH19(44-87)::rhCDH19(89-114)::caCDH19(115-770) (see SEQ ID NO: 1461)

20 U = caCDH19(44-120)::rhCDH19(122-137)::caCDH19(137-770) (see SEQ ID NO: 1462)

V = rhCDH19(44-141)::raCDH19(140-247)::rhCDH19(250-772) (see SEQ ID NO: 1463)

W = raCDH19(44-770) (see SEQ ID NO: 1464)

The data summarized in table 5 allowed for segregating the binder of Bin A 44-141 into the following subgroups:

5 Bin A.1 44-141

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Bin A.2 44-141 (44-114)

Bin A.3 44-141 (44-65)

Epitope Prediction by rat/mouse or human/mouse Cadherin-19 Chimeras

Domain binding was determined by flow cytometry on 293T cells transiently transfected with plasmids consisting of rat CDH19 cadherin repeat domain 3 substitutions (designated EC3a, EC3b) or human CDH19 cadherin repeat domain 3 substitution (designated EC3c) into the mouse Cadherin19 backbone cloned into the pTT5 expression vector immediately preceded by native mouse CDH19 leader sequence and a Flag tag. The experiment included assaying a subset of anti-CDH19 antibodies against human, rat and mouse Cadherin19 to determine suitability for binning on these rat/mouse and human/mouse chimeras.

The following binding assay was completed in presence of 2mM CaCl2. In 96-well V-bottom plates (Costar 3897), 50,000 transiently transfected 293T cells were incubated with 5ug/ml of purified anti-CDH19 antibodies for 1hr at 4oC followed by two washes with PBS/2%FBS. 50µl of 5µg/ml Alexa647-labelled anti-human IgG secondary antibody (Jackson Immuno 109-605-098) and 2ug/ml 7AAD (Sigma A9400) was then added to each well and the plates incubated for 15 minutes at 4oC. Cells were then washed one time and the amount of cell associated Alexa647-labelled Ab was quantitated by flow cytometry. The experiments included mock transfected controls. The data from these experiments are presented in the Table below, n.d. = not determined.

Table 6 – Antibody Bin B Epitope Prediction Summary

			Hu EC1-5	Mo EC1-5	Ra EC1-5	Ra EC3c	Ra EC3b	Hu EC3a	
Clone ID	Ab. ID	Bin	Α	ı	w	х	Y	z	Predicted Epitope Region
4A9	13589	1	+	-	-	n.d.	n.d.	n.d.	
26F12	13881	2	+	-	-	n.d.	n.d.	n.d.	
25F8	14096	2	+	-	-	n.d.	n.d.	n.d.	44-141
26D1	13882	2	+	-	-	n.d.	n.d.	n.d.	Bin A
17H8	13874	3	+	-	-	n.d.	n.d.	n.d.	
16A4	14071	5	+	-	+	n.d.	n.d.	n.d.	
4A2	13592	4	+	-	+	+	-	-	250-364 (324-327)
4B10	13590	4	+	-	+	+	-	-	Bin B.2
2G6	13588	4	+	+	+	+	+	+	250-364
23A10	14077	4	+	+	+	n.d.	n.d.	n.d.	Bin B.1
Rat an	Rat anti-FLAG		+	+	+	+	+	+	control

Positive Binding (+)
Negative Binding (-)
Not Determined (n.d.)

Legend Table 6

Rat/mouse or human/mouse chimera constructs

5 A = huCDH19(44-772) (see SEQ ID NO: 944)

I = muCDH19(44-770) (see SEQ ID NO: 966)

W = raCDH19(44-770) (see SEQ ID NO: 1464)

X = muCDH19(44-323)::raCDH19(324-327)::muCDH19(328-770) (see SEQ ID NO: 1465)

Y = muCDH19(44-770)::raCDH19(290,299,308) (see SEQ ID NO: 1466)

10 Z = muCDH19(44-770)::huCDH19(271) (see SEQ ID NO: 1467)

The data summarized in table 6 allowed for segregating the binder of Bin B 250-364 into the following subgroups:

Bin B.1 250-364

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Bin B.2 250-364 (324-327)) by rodent numeration as referenced in table 6, corresponding to residues (326-329) within human and macaque CDH19.

Example 4 - Hotspot/Covariant Mutants

A total of 18 antibodies were analyzed for potential hotspots and covariance violations. The designed variants (shown below) outline amino acid substitutions capable of reducing and/or avoiding isomerization, deamidation, oxidation, covariance violations, and the like. The 80 engineered variants together with the 15 parental antibodies, thus totaling 95 sequences, were taken forward to the cloning, expression, and purification processes. Site-directed

mutagenesis was performed on the engineered variants in a 96-well format. The parental antibodies and engineered variants were expressed by high throughput transient transfection in HEK 293-6E cells, purified using a modified AKTA auto-sampler and assayed for activity and biophysical characteristics. The 3 parental antibodies that had either free (unpaired) Cys or N-glycosylation site were not taken forward in this process. Those were replaced with the engineered version of the parental antibodies. The designed variants outline amino acid substitutions capable of reducing and/or avoiding isomerization, deamidation, oxidation, covariance violations, immunogenicity and the like. It will be appreciated that these variant sequences are examples of engineered antibodies within the meaning of the present application but single point and/or multiple point mutations can be combined in any combinatorial manner in order to arrive at a final desired antigen binding molecule or antibody.

Example 5 – CDH19 mRNA expression pattern

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RNA was extracted from individual patient tissues representing tumor (>70% tumor content by cell count) or normal (0% tumor content by cell count). Individual tissues were homogenized using TisssueLyzer (Qiagen, Valencia, CA) and total RNA extracted and purified by the mirVana total RNA extraction kit (Life Technologies, Foster City, CA). RNA quality and quantity checked by NanoDrop (NanoDrop, Wilmington, DE) spectrophotometer readings and Bioanalyzer RNA profiling (Agilient Technologies, Santa Clara, CA). RNA was DNAse treated with DNA-free kit (Life Technologies, Foster City, CA) and reverse transcribed according to manufacturer's specifications using random hexamers in the High Capacity cDNA Reverse Transcription Kit (Life Technologies, Foster City, CA). Quantitative Real Time Polymerase Chain Reaction (qRT-PCR) was performed on cDNA using primers to CDH19, probeset Hs00253534 m1, (Life Technologies, Foster City, CA) or the housekeeping gene human ACTB (primers CCT GGC ACC CAG CAC AA; GCC GAT CCA CAC GGA GTA CT; probe ATC AAG ATC ATT GCT CCT GAG CG). 10 µL qRT-PCR reaction components; 1.0 ng/µL cDNA, 2xUniversal PCR Master Mix (Life Technologies, Foster City, CA), gene expression assay (ACTB; 75 nM primers, 150 nM probe. EPOR; 300 nM primers, 250 nM probe) Following the qRT-PCR amplification program: (1) activation at 50°C for 2 min; (2) denaturation at 95°C for 10 min; (3) amplification 40 cycles at 95°C for 15 s and 60°C for 1 min with fluorescence capture at each step (ABI PRISM 7900HT Sequence Detection Systems, Applied Biosystems). Threshold cycle values (C_T) were determined, using Sequence Detector software version 2.3 (Applied Biosystems) and transformed to $2^{-\Delta CT}$ for relative expression of CDH19 specific transcript to ACTB. The results are shown in

Figure 3. Of 54 unique metastatic and primary melanoma samples, the majority can be seen to overexpress CDH19 mRNA relative to the expression in samples from normal tissue.

Example 6 – CDH19 protein expression

5 Expression of CDH19 protein was analyzed in human tumor samples by IHC and the results are shown in Figure 4. Samples were fixed in 10% neutral buffered formalin for 24 hours, dehydrated and paraffin embedded. 4 µm sections were cut. Sections were deparaffinized first and then heated in DIVA Decloaker solution (Biocare) for 40 minutes for antigen retrieval. Remaining IHC steps were performed at room temperature in a DAKO Autostainer. Sections were incubated for 10 minutes with Peroxidazed 1 (Biocare) to block endogenous 10 peroxidase, followed by incubation for 10 minutes with background sniper (Biocare) to reduce nonspecific background. Section were incubated for 60 minutes with CDH19 antibody (Novo Biologicals, Catalog #H00028513-B01P) at 5 µg/ml, then incubated for 30 minutes with Envision+ HRP anti-mouse polymer (DAKO), followed by DAB+ (DAKO) for 5 minutes. Sections were counterstained with hematoxylin (DAKO) approximately for 1 minute. CDH19 15 expression could be detected in 62% of tumors examined (staining intensity ≥1+ in 101 of 162 samples). 51% of the tumor samples demonstrated medium to high expression (staining intensity of 2+ to 3+ in 83 of162 samples). CDH19 showed dense and distinct membrane staining in many samples, although in some tumors heterogeneity was noted.

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Example 7 – Selection of model cell lines

Tumor cell lines were analyzed by flow cytometry and IHC to identify model systems with CDH19 expression similar to human tumors. Human anti-huCDH19 IgG4 antibody 4A2 was purified directly from hybridoma conditioned media. For flow cytometry, 2x10⁵ cells were incubated with 200 nM of the CDH19 4A2 antibody that was conjugated to PE at a 1:1 ratio. The incubation and subsequent wash steps were performed in the presence of 1.2 mM calcium. A tube of QuantiBRITE PE lyophilized beads with four levels of PE (BD, cat# 340495) was simultaneously prepared according to the manufacturer's instructions. The beads were analyzed by flow cytometry to generate a standard curve. The PE median values obtained from the melanoma lines after FACS analysis were then calibrated against the standard curve to calculate the antibodies bound per cell (ABC), which provides an estimate of the number of receptors on each cell. IHC was performed as described in Example 6 and the results are provided in Figure 5. The melanoma cell line CHL-1 expresses about 10,000 CDH19 molecules on the cell surface, while Colo699 cells express about 5,000 receptors.

Both cell lines represent tumors with medium to high expression levels based on IHC. Expression in A2058 is very low, while LOX cells do not express any detectable CDH19 protein.

Example 8

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5 Bispecific binding and interspecies cross-reactivity

For confirmation of binding to human CDH19 and to human and macaque CD3, bispecific antibodies were tested by flow cytometry using indicated cell lines. L1.2 transfected with human CDH19, the human melanoma cell lines CHL-1 and A2058 expressing native human CDH19, CD3-expressing human T cell leukemia cell line HPB-ALL (DSMZ, Braunschweig, ACC483) and the CD3-expressing macaque T cell line 4119LnPx (Knappe A, et al., Blood, 2000, 95, 3256-3261) were used as antigen positive cell lines. Moreover, untransfected L1.2 cells were used as negative control.

For flow cytometry 200,000 cells of the respective cell lines were incubated for 30 min on ice with 50 µl of purified bispecific antibody at a concentration of 5 µg/ml. The cells were washed twice in PBS/2% FCS and binding of the constructs was detected with a murine PentaHis antibody (Qiagen; diluted 1:20 in 50 µl PBS/2% FCS). After washing, bound PentaHis antibodies were detected with an Fc gamma-specific antibody (Dianova) conjugated to phycoerythrin, diluted 1:100 in PBS/2% FCS. Samples were measured by flow cytometry on a FACSCanto II instrument and analyzed by FACSDiva software (both from Becton Dickinson).

The CDH19/CD3 bispecific antibodies stained L1.2 cells transfected with human CDH19, the human CDH19-expressing melanoma cell lines CHL-1 and A2058 as well as human and macaque T cells. Moreover, there was no staining of untransfected L1.2 cells (see Figure 6).

Example 9

Cytotoxic activity

FACS-based cytotoxicity assay with unstimulated human PBMC

30 Isolation of effector cells

Human peripheral blood mononuclear cells (PBMC) were prepared by Ficoll density gradient centrifugation from enriched lymphocyte preparations (e.g. buffy coats), a side product of blood banks collecting blood for transfusions. Buffy coats were supplied by a local blood bank and PBMC were prepared on the same day of blood collection. After Ficoll density centrifugation and extensive washes with Dulbecco's PBS (Gibco), remaining erythrocytes were removed from PBMC via incubation with erythrocyte lysis buffer (155 mM NH $_4$ Cl, 10 mM KHCO $_3$, 100 μ M EDTA). Platelets were removed via the supernatant upon centrifugation

of PBMC at $100 \times g$. Remaining lymphocytes mainly encompass B and T lymphocytes, NK cells and monocytes. PBMC were kept in culture at 37° C/5% CO₂ in RPMI medium (Gibco) with 10% FCS (Gibco).

Depletion of CD14⁺ and CD56⁺ cells

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For depletion of CD14⁺ cells, human CD14 MicroBeads (Milteny Biotec, MACS, #130-050-201) were used, for depletion of NK cells human CD56 MicroBeads (MACS, #130-050-401). PBMC were counted and centrifuged for 10 min at room temperature with 300 x g. The supernatant was discarded and the cell pellet resuspended in MACS isolation buffer [80 μL/ 10^7 cells; PBS (Invitrogen, #20012-043), 0.5% (v/v) FBS (Gibco, #10270-106), 2 mM EDTA (Sigma-Aldrich, #E-6511)]. CD14 MicroBeads and CD56 MicroBeads (20 μL/ 10^7 cells) were added and incubated for 15 min at 4 - 8°C. The cells were washed with MACS isolation buffer (1 - 2 mL/ 10^7 cells). After centrifugation (see above), supernatant was discarded and cells resuspended in MACS isolation buffer (500 μL/ 10^8 cells). CD14/CD56 negative cells were then isolated using LS Columns (Miltenyi Biotec, #130-042-401). PBMC w/o CD14+/CD56+ cells were cultured in RPMI complete medium i.e. RPMI1640 (Biochrom AG, #FG1215) supplemented with 10% FBS (Biochrom AG, #S0115), 1x non-essential amino acids (Biochrom AG, #K0293), 10 mM Hepes buffer (Biochrom AG, #L1613), 1 mM sodium pyruvate (Biochrom AG, #L0473) and 100 U/mL penicillin/streptomycin (Biochrom AG, #A2213) at 37°C in an incubator until needed.

Target cell labeling

For the analysis of cell lysis in flow cytometry assays, the fluorescent membrane dye DiOC₁₈ (DiO) (Molecular Probes, #V22886) was used to label human CDH19- as target cells and distinguish them from effector cells. Briefly, cells were harvested, washed once with PBS and adjusted to 10^6 cell/mL in PBS containing 2 % (v/v) FBS and the membrane dye DiO (5 μ L/ 10^6 cells). After incubation for 3 min at 37°C, cells were washed twice in complete RPMI medium and the cell number adjusted to 1.25 x 10^5 cells/mL. The vitality of cells was determined using 0.5 % (v/v) isotonic EosinG solution (Roth, #45380).

Flow cytometry based analysis

This assay was designed to quantify the lysis of human CDH19-transfected CHO cells in the presence of serial dilutions of CDH19 bispecific antibodies.

35 Equal volumes of DiO-labeled target cells and effector cells (i.e., PBMC w/o CD14⁺ cells) were mixed, resulting in an E:T cell ratio of 10:1. 160 μL of this suspension were transferred to each well of a 96-well plate. 40 μL of serial dilutions of the CDH19 bispecific antibodies

and a negative control bispecific (an CD3-based bispecific antibody recognizing an irrelevant target antigen) or RPMI complete medium as an additional negative control were added. The bispecific antibody-mediated cytotoxic reaction proceeded for 48 hours in a 7% CO₂ humidified incubator. Then cells were transferred to a new 96-well plate and loss of target cell membrane integrity was monitored by adding propidium iodide (PI) at a final concentration of 1 μ g/mL. PI is a membrane impermeable dye that normally is excluded from viable cells, whereas dead cells take it up and become identifiable by fluorescent emission.

Samples were measured by flow cytometry on a FACSCanto II instrument and analyzed by FACSDiva software (both from Becton Dickinson).

Target cells were identified as DiO-positive cells. Pl-negative target cells were classified as living target cells. Percentage of cytotoxicity was calculated according to the following formula:

$$Cytotoxicity \, [\%] = \frac{n_{\text{ dead target cells}}}{n_{\text{target cells}}} \times 100$$

n = number of events

Using GraphPad Prism 5 software (Graph Pad Software, San Diego), the percentage of cytotoxicity was plotted against the corresponding bispecific antibody concentrations. Dose response curves were analyzed with the four parametric logistic regression models for evaluation of sigmoid dose response curves with fixed hill slope and EC50 values were calculated. The results are shown in Figure 7.

Example 10

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In vivo tumor growth inhibition experiments

5 million Colo699 or CHL-1 tumor cells were admixed with 2.5 million freshly isolated peripheral blood mononuclear cells (PBMC) and injected subcutaneously in the left flank of female athymic nude mice on Day 0. The same day, mice were treated intraperitoneally with either CDH19 BiTE 2G6 or non-specific control BiTE (MEC14) at the indicated doses. Dosing continued daily for the first 10 days post-tumor inoculation.

Tumor volumes and body weights were measured twice per week using calipers and an analytical scale, respectively.

The results of experiments with Colo699 or CHL-1 tumor cells are shown in Figures 8 and 9.

Example 11

35 Cytotoxic activity

Imaging-based cytotoxicity assay with unstimulated human T-cells

Effector cells

Purified, naïve human T cells were obtained from AllCells LLC, Alameda, USA.

Image based analysis

This assay measures the T cell mediated lysis of melanoma cells. 3000 A2058 cells (CDH19 positive) or 2500 LOX IMVI cells (CDH19 negative) are combined with naïve human T cells in a 1:10 ratio in the wells of 384 well plates. After addition of a serial dilution of CDH19 targeting BiTE molecules as well as a negative control bispecific (a CD3-based bispecific antibody recognizing an irrelevant target antigen), the cells are incubated for 48 h at 37°C. Next, the samples are treated for 2 h with 30 μ M Hoechst 33342 to stain the nuclei of all cells and 2 μ M propidium iodide (PI) to identify dead cells.

Image acquisition and analysis is performed on a ThermoFisher ArrayScan with a 10x objective. Data for two channels is collected, at 386nm (Hoechst 33342) and at 549nm (propidium iodide).

Live cells are identified as Hoechst positive, PI negative events, dead cells as Hoechst positive, PI positive.

Percentage of cytotoxicity is determined as described in example 7. Representative results are shown in figure 10.

Example 12

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20 Domain specificity and biochemical affinity determination of bi-specific binders

Purification of CDH19 sub-domains lacking post-translational modifications

A methionine initiation codon followed by nucleotide sequences encoding CDH19 subdomain protein A= huCDH19(140-367 of SEQ ID NO:944), immediately preceding a G₄S linker and poly-Histidine tag was cloned into a suitable pET vector; whereas, nucleotides sequences encoding sub-domain proteins B= huCDH19(44-367 of SEQ ID NO:944) and C= rhCDH19(44-367 of SEQ ID NO:1457) were cloned into the pET-SUMO vector (Life Technologies, Invitrogen) by methods known in the art. Each was expressed in E coli, isolated from the soluble fraction and purified to homogeneity by metal chelate affinity chromatography, followed by anion exchange, and size exclusion chromatography in HEPES buffered saline, 3mM CaCl2, pH 8. Sub-domain protein A retained its linker and C terminal polyhistidine tag, but His-SUMO tags constituent to the N termini of proteins B and C were removed by digestion with SUMO protease (Life Technologies, Invitrogen) prior to anion exchange. All proteins were determined to have their expected molecular weight by ESI LC/MS. Proteins used in binding experiments described below were randomly biotinylated by typical methods known in the art.

Purification of CDH19 sub-domains with post-translational modifications

CDH19 sub-domain proteins D= huCDH19(44-367 of SEQ ID NO:944), and E= rhCDH19(44-367 of SEQ ID NO:1457) were generated by cloning nucleotide sequences encoding respective amino acid residues 1-367 into the pSURETech235b vector (Selexis) each immediately preceded a G₄S linker and poly-Histidine tag were cloned into the pSURETech235b vector (Selexis), transfected into CHO-S cells (Life Technologies, Invitrogen), and stable pools were generated following hygromycin selection by methods known in the art. Stable pools were expanded and conditioned media was collected after 7 days culture in serum free media. CM was exchanged by UF/DF with 5 diavolumes HEPES buffered saline plus CaCl₂ using a 1 sq ft 10K PES Pellicon 2 membrane and purified to homogeneity as described above. CDH19 sub-domain proteins D and E retained constituent linker and C terminal polyhistidine tags. N terminal sequence of each protein was determined to be G44 as expected, while ESI LC/MS of purified proteins as compared with same subjected to PNGase F digestion revealed the presence of both N- and O-linked glycans. Proteins used in binding experiments described below were randomly biotinylated by methods well known in the art.

Methods for binding affinity determination by Octet

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The Octet RED384 biosensor was used to characterize kinetics and affinity of protein-protein interactions. Minimally biotinylated CDH19 domain target proteins A-E were bound to streptavidin tips in the machine while serial dilutions of analyte bi-specific binder proteins were made in 96-well or 384-well plates. Empirical target loading conditions were found from assay development to be 10-20 nM target concentration and loading for 600 seconds to give a 2nm signal. Binding experiments were performed by setting up a plate with 6-point (Tables 7-9) or 3-point (Table 10) 1:3 serial dilutions from 30nM starting concentrations of each analyte, with two reference wells per column having buffer alone. Octet Buffer: 10 mM HEPES (pH 7.5), 150 mM NaCl, +/- 1 mM CaCl₂, 0.13% Triton X-100 and 0.10 mg/ml BSA. Additional baseline and dissociation wells in the plate also contained buffer alone. The binding method was as follows: ForteBio Octet streptavidin tips were (1) soaked in buffer for 10 minutes; (2) transferred to the plate baseline wells and incubated for 5 minutes; (3) transferred to the target loading wells and incubated for 10 minutes; (4) transferred to the plate baseline wells and incubated for 5 minutes; (5) transferred to the sample wells and incubated for 5 minutes (Table 9) or 20 minutes (Tables 7, 8, 10); (6) transferred to the dissociation wells and incubated for 8.3 minutes (Table 9) or 1.5hr (Tables 7, 8, 10). Raw data was processed in the following manner: (a) reference tip curves were averaged and subtracted from sample curves; (b) the association and dissociation curves were isolated and aligned to the Y axis; (c) the association and dissociation interstep was aligned; (d) Savitzky-

Golay filtering was implemented to reduce the signal noise and (e) the resulting set of association and dissociation curves for each sample-target interaction were fit globally with a single 1:1 binding model to determine the measured values of the association (Ka) and dissociation (Kd) rate constants to calculate the equilibrium dissociation constant, KD.

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Table 7 – Domain Specificity and Biochemical Affinity of Bi-specific Binders to Isolated human CDH19 Protein Domains Lacking Post Translational Modifications

		A = huC	DH19(140-3	67) E coli	B = hu					
Clone ID	Bispecific binder ID	KD (nM)	ka (M-1s-1)	kd (s-1)	KD (pM)	ka (M-1s-1)	kd (s-1)	Predicted Epitope Region		
2G6	65254	< 0.03	3.37E+05	< 1.0E-05	< 0.04	2.31E+05	< 1.0E-05	250-364 Bin B.1		
26F12	26F12 65251 (-) (-) (-) 0.20 3.86E+05 7.56E-05 44-114 Bin A.2									
(-) negative binding, 20 min association, 1.5 hr dissociation										

Legend Table 7

Human CDH19 Protein domains lacking post translational modifications

A = E coli expressed huCDH19(140-367 of SEQ ID NO:944)

B = E coli expressed huCDH19(44-367 of SEQ ID NO:944)

The data summarized in table 7 confirmed CDH19 epitope region specificity of bi-specific binders and allowed for their relative affinity ranking.

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Table 8 – Calcium Modulated Biochemical Affinity of Bi-specific Binders to Isolated Human and Macaque CDH19 Protein Domains Lacking Post Translational Modifications

		B = hu	CDH19(44-36	7) E coli	C = rhC	') E coli					
Clone ID, Epitope Bin	Bispecific binder ID	KD (nM)	ka (M-1s-1)	kd (s-1)	KD (nM)	ka (M-1s-1)	kd (s-1)	CaCl ₂			
2G6, Bin B.1	65254	< 0.06	1.66E+05	< 1.0E-05	< 0.03	2.97E+05	< 1.0 E-05	1mM			
26F12, Bin A.2 65251 0.31 2.91E+05 9.0E-05 0.17 8.19E+05 1.36E-04 1mM											
2G6, Bin B.1	2G6, Bin B.1 65254 (-) (-) (-) (-) (-) absent										
26F12, Bin A.2	65251	2.56	1.21E+05	3.08E-04	1.16	4.68E+05	5.44E-04	absent			
(-) negative binding, 20 min association, 1.5 hr dissociation											

Legend Table 8

20 CDH19 Protein domains lacking post translational modifications

B = E coli expressed huCDH19(44-367 of SEQ ID NO:944)

C = E coli expressed rhCDH19(44-367 of SEQ ID NO:1457)

The data summarized in table 8 allowed determination of calcium sensitivity of bi-specific binders and for their relative affinity ranking. Data further suggests conformational epitopes, with Bin B.1 more dependent on CDH19/Ca2+ association than epitope Bin A.2

Table 9 – Biochemical Affinity of Bi-specific Binders to Isolated Human and Macaque CDH19 Protein Domains Lacking Post Translational Modifications

		B = huCDH19(44-367) E coli			C = rh	CDH19(44-36	7) E coli		
Clone ID	Bispecific binder ID	KD (nM)	ka (M-1s-1)	kd (s-1)	KD (nM)	ka (M-1s-1)	kd (s-1)		
2G6	65254	< 0.3	3.11E+05	< 1.0E-04	< 0.3	3.69E+05	< 1.0E-04		
2G6.001	65254.001	< 0.4	2.21E+05	< 1.0E-04	< 0.4	2.42E+05	< 1.0E-04		
2G6.003	65254.003	< 0.5	1.80E+05	< 1.0E-04	< 0.5	1.91E+05	< 1.0E-04		
2G6.007	65254.007	0.57	2.95E+05	1.69E-04	0.55	3.53E+05	1.94E-04		
4A2.002	65238.002	< 0.2	5.48E+05	< 1.0E-04	< 0.1	9.13E+05	< 1.0E-04		
4B10.002	65240.002	< 0.2	5.02E+05	< 1.0E-04	< 0.1	7.48E+05	< 1.0E-04		
4B10.003	65240.003	< 0.2	3.87E+05	< 1.0E-04	< 0.2	5.06E+05	< 1.0E-04		
4B10.005	65240.005	< 0.2	4.41E+05	< 1.0E-04	< 0.2	6.00E+05	< 1.0E-04		
19B5.1.002	65235.002	1.74	3.74E+05	6.49E-04	1.02	4.94E+05	5.02E-04		
19B5.1.003	65235.003	2.44	3.09E+05	7.54E-04	1.63	3.97E+05	6.45E-04		
23A10.001 (B1)	65237.001	< 0.4	2.55E+05	< 1.0E-04	< 0.3	3.16E+05	< 1.0E-04		
23A10.001 (B2)	65237b.001	0.57	2.95E+05	1.69E-04	0.55	3.53E+05	1.94E-04		
23A10.002	65237.002	< 0.3	2.86E+05	< 1.0E-04	< 0.3	3.61E+05	< 1.0E-04		
26D1.1.003	65250.003	0.66	3.64E+05	2.41E-04	0.50	5.20E+05	2.62E-04		
26D1.1.004	65250.004	1.08	3.39E+05	3.67E-04	0.65	4.66E+05	3.02E-04		
26D1.1.005	65250.005	2.65	3.19E+05	8.44E-04	1.42	4.42E+05	6.25E-04		
26F12.002	65251.002	0.97	3.25E+05	3.16E-04	1.70	4.33E+05	7.36E-04		
26F12.004	65251.004	1.04	2.90E+05	3.00E-04	1.85	3.46E+05	6.38E-04		
26F12.006	65251.006	3.96	4.10E+05	1.62E-03	5.39	5.95E+05	3.21E-03		
26F12.008	65251.008	3.77	4.87E+05	1.84E-03	5.14	7.45E+05	3.83E-03		
1mM CaCl ₂ , 5 min association, 8.3 min dissociation									

10 Legend Table 9

CDH19 Protein domains lacking post translational modifications

B = E coli expressed huCDH19(44-367 of SEQ ID NO:944)

C = E coli expressed rhCDH19(44-367 of SEQ ID NO:1457)

The data summarized in table 9 allowed relative affinity ranking of bi-specific binders to human and non-human primate CDH19 domains lacking glycosylation.

Table 10 – Calcium Modulated Biochemical Affinity of Bi-specific Binders to Isolated Glycosylated Human and Macaque CDH19 Protein Domains

		D = hւ	CDH19(44-36	57) CHO	E = rh0	CDH19(44-36	7) CHO	
Clone ID, Epitope Bin	Bispecific binder ID	KD (nM)	ka (M-1s-1)	kd (s-1)	KD (nM)	ka (M-1s-1)	kd (s-1)	CaCl ₂
2G6, Bin B.1	65254	< 0.041	2.44E+05	< 1.0E-05	< 0.031	3.19E+05	< 1.0E-05	1mM
2G6.003, Bin B.1	65254.003	< 0.099	1.01E+05	< 1.0E-05	< 0.09	1.10E+05	< 1.0E-05	1mM
4B10.003, Bin B.2	65240.003	0.24	2.08E+05	4.91E-05	0.29	2.70E+05	7.88E-05	1mM
19B5.1.003, Bin A.2	65235.003	1.01	4.02E+05	4.07E-04	0.27	7.12E+05	1.93E-04	1m M
23A10.002, Bin B.1	65237.002	< 0.036	2.75E+05	< 1.0E-05	< 0.035	2.82E+05	< 1.0E-05	1mM
26D1.1.005, Bin A.2	65250.005	0.97	3.13E+05	3.04E-04	0.37	4.64E+05	1.74E-04	1mM
26F12, Bin A.2	65251	0.28	5.28E+05	1.50E-04	0.22	8.72E+05	1.94E-04	1mM
26F12.006, Bin A.2	65251.006	1.24	4.92E+05	6.07E-04	1.13	6.94E+05	7.86E-04	1mM
2G6, Bin B.1	65254	(-)	(-)	(-)	(-)	(-)	(-)	absent
2G6.003, Bin B.1	65254.003	(-)	(-)	(-)	(-)	(-)	(-)	absent
4B10.003, Bin B.2	65240.003	(-)	(-)	(-)	(-)	(-)	(-)	absent
19B5.1.003, Bin A.2	65235.003	3.49	2.90E+05	1.01E-03	3.28	2.65E+05	8.68E-04	absent
23A10.002, Bin B.1	65237.002	(-)	(-)	(-)	(-)	(-)	(-)	absent
26D1.1.005, Bin A.2	65250.005	0.86	4.12E+05	3.56E-04	2.58	3.26E+05	8.41E-04	absent
26F12, Bin A.2	65251	1.91	2.66E+05	5.09E-04	1.09	5.38E+05	5.88E-04	absent
26F12.006, Bin A.2	65251.006	0.79	6.29E+05	4.95E-04	18.53	3.36E+05	6.22E-03	absent

Legend Table 10

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10 Glycosylated CDH19 Protein domains

D = CHO expressed huCDH19(44-367 of SEQ ID NO:944)

E = CHO expressed rhCDH19(44-367 of SEQ ID NO:1457)

The data summarized in table 10 allowed determination of calcium sensitivity of bi-specific binders and relative affinity ranking toward glycosylated human and non-human primate CDH19 domain proteins. As compared to data in Table 8, affinities are similar to those with domains lacking post-translational modifications. Data further suggests conformational epitopes, with epitope Bins B.1 and B.2 being more dependent on CDH19/Ca2+ association than epitope Bin A.2

Example 13

10 Bispecific binding and interspecies cross-reactivity:

For confirmation of binding to human CDH19 and to human CD3, bispecific antibodies were tested by flow cytometry using indicated cell lines. HEK293 transfected with human CDH19 (see example 14) and CD3-expressing human T cell leukemia cell line HPB-ALL (DSMZ, Braunschweig, ACC483) were used as antigen positive cell lines.

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For flow cytometry 200,000 cells of the respective cell lines were incubated for 30 min on ice with 100 µl of BiTE containing cell culture supernatant. The cells were washed twice in PBS/2% FCS and binding of the constructs was detected with a murine anti-CD3scFv antibody (3E5.A5, Amgen; diluted to 2 µg/ml PBS/2% FCS). After washing, bound anti-CD3scFv antibodies were detected with an Fc gamma-specific antibody (Dianova) conjugated to phycoerythrin, diluted 1:100 in PBS/2% FCS. Samples were measured by flow cytometry on a FACSCanto II instrument and analyzed by FACSDiva software (both from Becton Dickinson).

The CDH19/CD3 bispecific antibodies stained HEK293 cells transfected with human CDH19 as well as human and macaque T cells (see Figure 19).

Example 14

Cytotoxic activity

30 Chromium release assay with stimulated human T cells

Isolation of effector cells

A petri dish (145 mm diameter, Greiner bio-one GmbH, Kremsmünster) was coated with a commercially available anti-CD3 specific antibody (OKT3, Orthoclone) in a final concentration of 1 μ g/ml for 1 hour at 37°C. Unbound protein was removed by one washing step with PBS. $3-5 \times 10^7$ human PBMC were added to the precoated petri dish in 120 ml of RPMI 1640 with stabilized glutamine / 10% FCS / IL-2 20 U/ml (Proleukin®, Chiron) and

stimulated for 2 days. On the third day, the cells were collected and washed once with RPMI 1640. IL-2 was added to a final concentration of 20 U/ml and the cells were cultured again for one day in the same cell culture medium as above.

Depletion of CD4⁺ and CD56⁺ cells

CD8⁺ cytotoxic T lymphocytes (CTLs) were enriched by depletion of CD4⁺ T cells and CD56⁺ NK cells using Dynal-Beads according to the manufacturer's protocol.

⁵¹Cr release based analysis

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Human CDH19-transfected HEK293 target cells (production see example 14) were washed twice with PBS and labeled with 11.1 MBq ⁵¹Cr in a final volume of 50 μl supplemented RPMl for 60 minutes at 37°C. Subsequently, the labeled target cells were washed 3 times with 5 ml RPMl and then used in the cytotoxicity assay. The assay was performed in a 96-well plate in a total volume of 200 μl supplemented RPMl with an E:T ratio of 10:1. A starting concentration of 0.1 – 1 μg/ml of purified bispecific antibody and threefold dilutions thereof were used. Incubation time for the assay was 18 hours. Cytotoxicity was determined as relative values of released chromium in the supernatant relative to the difference of maximum lysis (addition of Triton-X) and spontaneous lysis (without effector cells). All measurements were carried out in quadruplicates. Measurement of chromium activity in the supernatants was performed in a Wizard 3" gamma counter (Perkin Elmer Life Sciences GmbH, Köln, Germany). Analysis of the results was carried out with Prism 6 for Windows (version 6.02, GraphPad Software Inc., San Diego, California, USA). EC50 values calculated by the analysis program from the sigmoidal dose response curves were used for comparison of cytotoxic activity (see Figure 20).

Example 15

25 Production and purification of BiTE antibodies

Standardized research scale production of CDH19 BiTE antibodies was performed in roller bottles. Harvested culture supernatant was subjected after filtration to two step BiTE antibody purification based either on immobilized metal affinity chromatography (IMAC) capture and subsequent size exclusion chromatography or Protein_A capture and subsequent size exclusion chromatography (SEC).

15.1 IMAC capture step of BiTE antibodies

Äkta® Explorer Systems (GE Healthcare) controlled by Unicorn® Software were used for chromatography. Immobilized metal affinity chromatography (IMAC) was performed using Fractogel EMD chelate® (Merck, Darmstadt) which was loaded with ZnCl2 according to the protocol provided by the manufacturer. The column was equilibrated with buffer A (20 mM sodium phosphate buffer , 0.1 M NaCl , 10 mM imidazole, pH 7.2) and the cell culture supernatant (1000 ml) applied to the column (10 ml packing volume) at a flow rate of 4

ml/min. The column was washed with buffer A to remove unbound sample. Bound protein was eluted using a two step gradient of buffer B (20 mM sodium phosphate buffer, 0.1 M NaCl, 0.5 M imidazole, pH 7.2) according to the following procedure:

Step 1: 10 % buffer B in 5 column volumes

5 Step 2: 100% buffer B in 5 column volumes

Eluted protein fractions from step 2 were pooled for further purification and concentrated to 3 ml final volume using Vivaspin (Sartorius-Stedim, Göttingen-Germany) centrifugation units with PES membran and a molecular weight cut-off of 10 kDa. All chemicals were of research grade and purchased from Merck (Darmstadt, Germany). Figure 11

10 15.2 Protein A capture of BiTE antibodies

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Äkta® Explorer Systems (GE Life Sciences) controlled by Unicorn® Software were used for chromatography. Affinity columns which containin beads with covalently bound Protein_A were used for the capture step. The column was equilibrated with equillibration buffer pH 7.4 and the cell culture supernatant applied. After washing the column with three column volumes of equillibration buffer to wash out unbound sample the bound BiTE antibodies were eluted by application of an elution buffer at pH 3.0. Eluted solution was immediately neutralized in pH by a Trishydroxymethylamine Tris solution pH 8.0 already contained in the fractionation tubes in the fraction collector.

Eluted protein fractions from step 2 were pooled for further purification and concentrated to 3 ml final volume using Vivaspin (Sartorius-Stedim, Göttingen-Germany) centrifugation units with PES membran and a molecular weight cut-off of 10 kDa. All chemicals were of research grade and purchased from Merck (Darmstadt, Germany). Figure 12

15.3 Size Exclusion Chromatography

Size exclusion chromatography was performed on a HiLoad 16/60 Superdex 200 prep grade column (GE Healthcare) equilibrated with SEC buffer (20 mM NaCl, 30 mM NaH2PO4, 100 mM L-Arginin, pH 7.0) at a flow rate of 1 ml/min. BiTE antibody monomer and dimer fractions were pooled and a 24% trehalose stock solution was added to reach a final trehalose concentration of 4%. Eluted protein samples were subjected to reducing SDS-PAGE and Anti His TAG Western Blot for analysis.

Protein pools were measured at 280 nm in polycarbonate cuvettes with 1 cm lightpath (Eppendorf, Hamburg-Germany) and protein concentration was calculated on the base of the Vector NTI sequence analysis software calculated factor for each protein.

BiTE monomer pools were adjusted to 250 μg/ml with additional BiTE formulation buffer (20 mM NaCl, 30 mM NaH2PO4, 100 mM L-Arginin, 4% Trehalose, pH 7.0). An amount of a minimum of 600 μg for each BiTE was taken and transferred for immediate protein analytics as described in example 16.

Remaining protein pools of BiTE antibody monomer and BiTE antibody dimer were aliquoted in 15 and 50 μ g protein aliquots and shock frozen in liquid nitrogen. Further storage until usage was done in a -80°C freezer until analysis of biologic activity and affinity measurements. Figure 13.

The purity of isolated BiTE antibody monomer was determined by SDS-PAGE to be >95%. As expected, purified monomeric BiTE antibody appeared as protein bands in the molecular weight range of 54-56 kDa. Figure 14

Example 16

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10 Protein Properties

The freshly prepared BiTE monomer solution generated in example 15 was applied to the following analytical methods

- High Performance Size Exclusion Chromatography (HP-SEC) of initially monomeric CDH19 BiTE antibodies after one week of incubation at 250 μg/ml and 37°C.
- BiTE monomer conversion of BiTE monomer to dimer by three freeze/thaw cycles followed by HP-SEC
 - High resolution analytical cation exchange
 - Hydrophobic interaction chromatography on a Sepharose Octyl FF matrix.
 - Concentration to 2500 μg/ml followed by over night storage and turbidity measurement
 - Aggregation temperature TA determination by heated Dynamic Light Scattering measurment

16.1 BiTE monomer conversion into dimer by incubation for 7 days

- 25 15 μg of the monomeric CDH19 BiTE antibody at a concentration of 250 μg/ml were incubated at 37°C for 7 days.
 - A high resolution SEC Column TSK Gel G3000 SWXL (Tosoh,Tokyo-Japan) was connected to an Äkta Purifier 10 FPLC (GE Lifesciences) equipped with an A905 Autosampler. Column equilibration and running buffer consisted of 100 mM KH2PO4 200 mM Na2SO4 adjusted to pH 6.6. After 7 days of incubation, the BiTE antibody solution (15 µg protein) was applied to the equilibrated column and elution was carried out at a flow rate of 0.75 ml/min at a maximum pressure of 7 MPa. The whole run was monitored at 280, 254 and 210 nm optical absorbance. Analysis was done by peak integration of the 210 nm signal recorded in the Äkta Unicorn software run evaluation sheet. Dimer content was calculated by dividing the area of the dimer peak by the total area of monomer plus dimer peak. Figure 15

16.2. BiTE monomer conversion into dimer by three freeze/thaw cycles

15 μ g of monomeric BiTE antibody at 250 μ g/ml were frozen at -80°C for 30 min followed by thawing for 30 min at room temperature. After three freeze/thaw cycles the dimer content was determined by HP-SEC as described in example 16.1. Figure 16

5 CDH19 BiTE CH19 2G6 302 x I2C SA21: 0.50 % Dimer content

16.3 High resolution analytical ion exchange chromatography

A 1 ml BioPro SP column manufactured by YMC (YMC Europe GmbH, Dinslaken-Germany) with sulphpropyl groups coupled to solid beads was connected to a Äkta Micro FPLC (GE Healthcare) device.

For column equilibration, sample dilution and washing a buffer consisting of 20 mM sodium dihydrogen phosphate and 30 mM sodium chloride adjusted with sodium hydroxide to a pH of 5.5 was used.

For elution a buffer consisting of 20 mM NaH2PO4 and 1000 mM NaCl adjusted with sodium hydroxide to a pH of 5.5 was used.

50 µg of BiTE antibody monomer were diluted with dilution buffer to 50 ml final volume.

After column equilibration 40 ml of the diluted protein solution was applied to the column followed by a wash step.

Elution was carried out by a steadily increasing gradient with elution buffer from zero to 100% over a total volume corresponding to 200 column volumes. The whole run was monitored at 280 (blue line) and 254 nm (red line) optical absorption.

Percentage of Main Peak was calculated by dividing the peak area of the main peak by the sum of peak area of all detected peaks followed by multiplication with a factor of 100. Figure 17

25 CDH19 BiTE CH19 2G6 302 x I2C SA21: 89.3 % Main Peak Percentage

16.4 Sepharose Octyl FF

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Elution of monomeric BiTE antibodies was evaluated on a hydrophobic interaction chromatography C8 Sepharose Octyl FF column (GE Healthcare) with 1 ml gel volume.

- 30 50 μg of BiTE antibody monomeric protein was filled up with buffer (10 mM Citric acid 75 mM Lysine x HCl 4% Trehalose pH 7.2) to a final volume of 300 μl. The column was connected to an Äkta Purifier 10 system (GE Healthcare). A 500 μl sample loop was connected to the system. The system and column were equilibrated with running buffer (10 mM Citric acid 75 mM Lysine x HCl 200 mM NaCl pH 7.2).
- 35 The complete sampe was injected into the sample loop and the content of the sample loop was applied to the column. After sample injection a volume of 10 ml running buffer was

applied to the column at a flow rate of 0,2 ml/min while recording the optical absorption at 254 and 280 nm together with conductivity. Figure 18

CDH19 BiTE CH19 2G6 302 x I2C SA21: Rapid and complete elution

5 16.5 Concentration of BiTE monomer to 2500 μg/ml followed by over night storage and turbidity measurement

1000 μ l of CDH19 BiTE monomer were concentrated in two Vivaspin 500 centrifugation units with 10 kDa PES membran (Sartorius-Stedim, Göttingen-Germany) to a final volume of 100 μ l. This volume as stored over night at 5°C in a cooling cabinet. Turbidity was measured three times at 340 nm optical wavelength absorption. Afterwards the mean value off he three measurement values was calculated.

OD340 Turbidity of CDH19 BiTE CH19 2G6 302 x I2C SA21: 0.034

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16.6 Aggregation temperature TA determination by heated Dynamic Light Scattering measurement

A volume of 40 μ l monomeric BiTE antibody at 250 μ g/ml was transferred into the inner core of a disposable plastic cuvette. The deeper placed outer core was filled up with generic BiTE formulation buffer. The top of the cuvette was sealed with a rubber top to avoid liquid loss by evaporation in the process of sample heating.

- The cuvette was placed in a Nanostar Dynamic Light Scattering device (Wyatt) and heated from 40°C to 70°C at a heating increment of 0.5 °C/min
 - Aggregation status was permanently monitored and recorded in the whole heating process. Evaluation was was executed with the software package supplied by the device manufacturer.
- 25 Aggregation temperature of CDH19 BiTE CH19 2G6 302 x I2C SA21: 52.4°C

16.7 PEGylation of BiTE antibodies with CysLoop

Monomeric BiTE antibody containing an c-terminal CysLoop (see for methodical details WO 2006/008096) was dialyzed against a Tris/NaCl buffer pH 7.4 and reduced by the addition of the reduction agent Tris(2-carboxyethyl)phosphine TCEP (Perbio Pierce) to create two reduced cysteins of the now opened CysLoop.

TCEP was removed by dialysis. PEG Maleimid capable of covalent binding to reduced cystein was added in molar excess and incubated for 3 hours at room temperature.

A Sepharose SP column cation exchange column (GE Healthcare) was connected to an Äkta FPLC system and equillibrated with binding buffer (low molar Phosphat/NaCl buffer of pH 5.0)

The protein solution was diluted with binding buffer adjusted to pH 5.0 to enable binding of the BiTE protein to the cation exchange column. Unbound PEG was removed in the wash step with further binding puffer pH 5.0 over 10 column volumes. Bound protein was eluted by a linear increasing percentage of elution buffer 20 mM phosphat 1 M NaCl.

5 PEGylated BiTE antibody eluted at lower molarity of the elution buffer compared to the unmodified BiTE antibody.

Sequence Table:

TABLE Ia: HEAVY CHAIN CDRs

Total Tota		100000000000000000000000000000000000000	Y CHAIN CDRS	ADD A	epp a
NA	Ab	Туре	CDR 1	CDR 2	CDR 3
NA			AGCTATGGCATGCAC		
AA	20.2	NA			
Part			SEQ ID NO: 1	SEQ ID NO: 2	SEQ ID NO: 3
SEQ 1D NO: 4 SEQ 1D NO: 5 SEQ 1D NO: 6			SYGMH	VIWYDGSNKYYADSVKG	RAGIIGTTGYYYGMDV
Seq id No: 7 Seg id No: 8 Seq id No: 9		AA	SEQ ID NO: 4	SEQ ID NO: 5	SEQ ID NO: 6
SEQ ID NO: 7 SEQ ID NO: 8 SEQ ID NO: 9	1F10	NA		ACCTACTACAACCCGTCCCTC	
AA			SEQ ID NO: 7		SEQ ID NO: 9
SEQ ID NO: 10 SEQ ID NO: 11 SEQ ID NO: 12			SGGYYWS	YIYYSGSTYYNPSLTS	DGSSGWYFQH
NA		AA	SEQ ID NO: 10	SEQ ID NO: 11	SEQ ID NO: 12
AA	2C12_LC#1	NA	AGCTATGGCATGCAC	AATAAATACTATGCAGACTCC	TACAGGCTACTACTACGGTA
AA SEQ ID NO: 16 SEQ ID NO: 17 SEQ ID NO: 18			SEQ ID NO: 13	SEQ ID NO: 14	SEQ ID NO: 15
SEQ ID NO: 16 SEQ ID NO: 17 SEQ ID NO: 18			SYGMH	VIWYDGSNKYYADSVKG	RAGIIGTTGYYYGMDV
NA		AA	SEQ ID NO: 16	SEQ ID NO: 17	SEQ ID NO: 18
SYGMH	2G6_LC#1	NA	AGCTATGGCATGCAC	AATAAATACTATGCAGACTCC	TATAGGCTACTACTACGGTA
AA			SEQ ID NO: 19	SEQ ID NO: 20	SEQ ID NO: 21
SEQ ID NO: 22 SEQ ID NO: 23 SEQ ID NO: 24		7.7	SYGMH	FIWYDGSNKYYADSVKD	RAGIIGTIGYYYGMDV
NA		AA	SEQ ID NO: 22	SEQ ID NO: 23	SEQ ID NO: 24
AA SYGMH FIWYDGSNKYYADSVKD RAGIIGTIGYYYGMDV	2G6	NA	AGCTATGGCATGCAC	AATAAATACTATGCAGACTCC	TATAGGCTACTACTACGGTA
AA			SEQ ID NO: 25	SEQ ID NO: 26	SEQ ID NO: 27
SEQ ID NO: 28 SEQ ID NO: 29 SEQ ID NO: 30			SYGMH	FIWYDGSNKYYADSVKD	RAGIIGTIGYYYGMDV
NA		AA	SEQ ID NO: 28	SEQ ID NO: 29	SEQ ID NO: 30
SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 34 SEQ ID NO: 35 SEQ ID NO: 36 2H12_LC#2 NA AGCTATGGCATGCAC SEQ ID NO: 37 SEQ ID NO: 38 SEQ ID NO: 39 SYGMH AA SEQ ID NO: 37 SEQ ID NO: 38 SEQ ID NO: 39 SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 4A2 5B4 5C5 NA AGTAGTGGTTACTACT GGAGC SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY	2H12	NA		AATAAATACTATACAGACTCC GTGAAGGGC	TACAGGCTACTACTACGGTA TGGACGTC
AA SEQ ID NO: 34 SEQ ID NO: 35 SEQ ID NO: 36 2H12_LC#2 NA AGCTATGGCATGCAC GTTATATGGTATGATGAAGT AATAAATACTATACAGACTCC GTGAAGGGC SEQ ID NO: 37 SEQ ID NO: 38 SEQ ID NO: 39 SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 4A2 5B4 5C5 NA AGTAGTGGTTACTACT GCAGTA GGAGC GTGAAGGGC SEQ ID NO: 41 SEQ ID NO: 42 CTTCCAGTAT CTTCCAGTAT CTTCCAGTAT SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY			SEQ ID NO: 31	SEQ ID NO: 32	
2H12_LC#2 AGCTATGGCATGCAC NA AGCTATGGCATGCAC SEQ ID NO: 34 AGCTATGGCATGCAC AATAAATACTATACAGACTCC GTGAAGGGC GTGAAGGGC SEQ ID NO: 37 SEQ ID NO: 38 SEQ ID NO: 39 SEQ ID NO: 39 SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 AGTAGTGGTTACTACT GGAGC SEQ ID NO: 41 SEQ ID NO: 42 AGTAGTGGTTACTACT GCAGC GCCTACTACAACCCGTCCCTC AAGAGT SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY		72 72	SYGMH	VIWYDGSNKYYTDSVKG	RAGIIGTTGYYYGMDV
AATAAATACTATACAGACTCC GTGAAGGGC SEQ ID NO: 37 SEQ ID NO: 38 SEQ ID NO: 39 SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 AGTAGTGGTTACTACT GGAGC GCCTACTACTACACTGGGAGC GCTACTACTACTCT AAGAGT SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY		m	SEQ ID NO: 34	SEQ ID NO: 35	SEQ ID NO: 36
SYGMH VIWYDGSNKYYTDSVKG RAGIIGTTGYYYGMDV SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 4A2 5B4 5C5 NA AGTAGTGGTTACTACT GGAGC GCCTACTACAACCCGTCCCTC AAGAGT SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY	2H12_LC#2	NA	AGCTATGGCATGCAC	AATAAATACTATACAGACTCC	TACAGGCTACTACTACGGTA
AA SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 4A2			SEQ ID NO: 37	SEQ ID NO: 38	SEQ ID NO: 39
AA2 5B4 5C5 NA AGTAGTGGTTACTACT GGAGC NA SEQ ID NO: 41 SEQ ID NO: 42 TACATCTATTACACTGGGAGC GCCTACTACAACCCGTCCCTC AAGAGT SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SEQ ID NO: 45 SEQ ID NO: 45 DGSSGWYFQY		7. 7.	SYGMH	VIWYDGSNKYYTDSVKG	RAGIIGTTGYYYGMDV
5B4 5C5 NA GGAGC GCCTACTACAACCCGTCCCTC AAGAGT SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY		AA	SEQ ID NO: 40	SEQ ID NO: 41	SEQ ID NO: 42
SSGYYWS YIYYTGSAYYNPSLKS DGSSGWYFQY	5B4	NA		GCCTACTACAACCCGTCCCTC	
ΔΔ			SEQ ID NO: 43	SEQ ID NO: 44	SEQ ID NO: 45
SEQ ID NO: 46 SEQ ID NO: 47 SEQ ID NO: 48		7. 7	SSGYYWS	YIYYTGSAYYNPSLKS	DGSSGWYFQY
		AA	SEQ ID NO: 46	SEQ ID NO: 47	SEQ ID NO: 48

Ab	Туре	CDR 1	CDR 2	CDR 3
4A9	NA	GGTTACTACTGGAGC	TATTTCTCTTACAGTGGGAGC ACCAACTACAACCCCTCCCTC AAGAGT	AACTGGGCCTTCCACTTTGA CTTC
		SEQ ID NO: 49	SEQ ID NO: 50	SEQ ID NO: 51
	AA	GYYWS	YFSYSGSTNYNPSLKS	NWAFHFDF
	AA	SEQ ID NO: 52	SEQ ID NO: 53	SEQ ID NO: 54
4B10 4C2	NA	AGCTATGACATGCAC	GTTATATCATATGATGGAACT AATGAATACTATGCAGACTCC GTGAAGGGC	GAACGATATTTTGACTGGTC TTTTGACTAC
		SEQ ID NO: 55	SEQ ID NO: 56	SEQ ID NO: 57
	7.7	SYDMH	VISYDGTNEYYADSVKG	ERYFDWSFDY
	AA	SEQ ID NO: 58	SEQ ID NO: 59	SEQ ID NO: 60
4D2	NA	AGTTATGACATGCAC	GTTATATCATATGATGGAACT AATGAATACTATGCAGACTCC GTGAAGGGC	GAACGATATTTTGACTGGTC TTTTGACTAC
		SEQ ID NO: 61	SEQ ID NO: 62	SEQ ID NO: 63
	7. 7.	SYDMH	VISYDGTNEYYADSVKG	ERYFDWSFDY
	AA	SEQ ID NO: 64	SEQ ID NO: 65	SEQ ID NO: 66
4D3 4F3	NA	AGCTATGACATGGAC	GTTATATGGTATGATGGAAGT AATAAAtacTATGCAGACTCC GTGAGGGGC	GAAACTGGGGAGGGCTGGTA CTTCGAtctc
		SEQ ID NO: 67	SEQ ID NO: 68	SEQ ID NO: 69
	7. 7.	SYDMD	VIWYDGSNKYYADSVRG	ETGEGWYFDL
	AA	SEQ ID NO: 70	SEQ ID NO: 71	SEQ ID NO: 72
4E10	NA	AGCTATGACATGCAC	GTTATATGGTATGATGGAAGT AATAAATACTATGCAGACTCC GTGAAGGGC	GAGTATAGGTACAGCTGGTA CTTTGACTAC
		SEQ ID NO: 73	SEQ ID NO: 74	SEQ ID NO: 75
	70.70	SYDMH	VIWYDGSNKYYADSVKG	EYRYSWYFDY
	AA	SEQ ID NO: 76	SEQ ID NO: 77	SEQ ID NO: 78
4F7	NA	AGTTACTCCTGGAGC	TATATCTATTACAGTGGGAGC ACCAACTACAACCCCTCCCTC AAGAGT	AACTGGGCCTTCCACTTTGA CTAC
		SEQ ID NO: 79	SEQ ID NO: 80	SEQ ID NO: 81
	7.7	SYSWS	YIYYSGSTNYNPSLKS	NWAFHFDY
	AA	SEQ ID NO: 82	SEQ ID NO: 83	SEQ ID NO: 84
5E3	NA	AGCTATAGCATGCAC	TCCATTAGTAGTAGTAGT TACATATACTACGCAGACTCA GTGAAGGGC	GGGGAAACTGGAACTAACTA CTACTACTACGGTATGGACG TC
		SEQ ID NO: 85	SEQ ID NO: 86	SEQ ID NO: 87
	7.7	SYSMH	SISSSSYIYYADSVKG	GETGTNYYYYGMDV
	AA	SEQ ID NO: 88	SEQ ID NO: 89	SEQ ID NO: 90
17H8 23B6 28D10	NA	AGTTACTACTGGAGC	TATATCTATTACATTGGGAGC ACCAACTACAACCCCTCCCTC AAGAGT	GATTCCCGGTATAGAAGTGG CTGGTACGATGCTTTTGATA TC
		SEQ ID NO: 91	SEQ ID NO: 92	SEQ ID NO: 93
	77.77	SYYWS	YIYYIGSTNYNPSLKS	DSRYRSGWYDAFDI
	AA	SEQ ID NO: 94	SEQ ID NO: 95	SEQ ID NO: 96
16C1	NA	GGTTACTACTGGAGC	TATATCTATTACATTGGGAGC ACCAACTACAACCCCTCCCTC AAGAGT	GATGGGAGCAGTGGCTGGTA CCGGTGGTTCGACCCC

Ab	Туре	CDR 1	CDR 2	CDR 3
		SEQ ID NO: 97	SEQ ID NO: 98	SEQ ID NO: 99
	7.7	GYYWS	YIYYIGSTNYNPSLKS	DGSSGWYRWFDP
	AA	SEQ ID NO: 100	SEQ ID NO: 101	SEQ ID NO: 102
16A4	NA	AGTTACTACTGGAGC	TATATCTATTACAGTGGGAGC ACCAATTACAACCCCTCCCTC AAGAGT	GATCAAAGGCGGATAGCAGC AGCTGGTACCCACTTCTACG GTATGGACGTC
		SEQ ID NO: 103	SEQ ID NO: 104	SEQ ID NO: 105
		SYYWS	YIYYSGSTNYNPSLKS	DQRRIAAAGTHFYGMDV
	AA	SEQ ID NO: 106	SEQ ID NO: 107	SEQ ID NO: 108
16E2 17E10 20B12	NA	AGCTATGGCATGCAC	GTGATATGGTATGATGGAAGT AATAAATACTATGCAGACTCC GTGAAGGGC	GACGGGTGGGAGCTGTCCTT TGACTAC
		SEQ ID NO: 109	SEQ ID NO: 110	SEQ ID NO: 111
	73. 73	SYGMH	VIWYDGSNKYYADSVKG	DGWELSFDY
	AA	SEQ ID NO: 112	SEQ ID NO: 113	SEQ ID NO: 114
22G10	NA	AGTTATGCCATGAAC	ACTATTAGTGGTGGTGGTGCT AACACATACTACGCAGACTCC GTGAAGGGC	GGGGGAATGGGGGGATACTA CTACGGTATGGACGTC
		SEQ ID NO: 115	SEQ ID NO: 116	SEQ ID NO: 117
	AA	SYAMN	TISGGGANTYYADSVKG	GGMGGYYYGMDV
	AA	SEQ ID NO: 118	SEQ ID NO: 119	SEQ ID NO: 120
16H2 20D3 23E7	NA	AGCTACTTTATTCAC	ATAATCAACCCTATTAGTGTT AGCACAAGCTACGCACAGAAG TTCCAGGGC	GGGGGGATACAGCTATGGTT ACATTTTGACTAC
		SEQ ID NO: 121	SEQ ID NO: 122	SEQ ID NO: 123
	AA	SYFIH	IINPISVSTSYAQKFQG	GGIQLWLHFDY
	7171	SEQ ID NO: 124	SEQ ID NO: 125	SEQ ID NO: 126
22D1	NA	AGCTACTTTATTCAC	ATAATCAACCCTATTAGTGTT AGCACAAGCTACGCACAGAAG TTCCAGGGC	GGGGGGATACAGCTATGGTT ACATTTGGACTAC
		SEQ ID NO: 127	SEQ ID NO: 128	SEQ ID NO: 129
	AA	SYFIH	IINPISVSTSYAQKFQG	GGIQLWLHLDY
	AA	SEQ ID NO: 130	SEQ ID NO: 131	SEQ ID NO: 132
25F8	NA	AGCTACTATATTCAC	ATAATCAACCCCAGTGGTGGT AGCACAAGGTACGCACAGAAG TTCCAGGGC	GGGGGAATACAGCTATGGTT ACATTttGACTAC
		SEQ ID NO: 133	SEQ ID NO: 134	SEQ ID NO: 135
	AA	SYYIH	IINPSGGSTRYAQKFQG	GGIQLWLHFDY
		SEQ ID NO: 136	SEQ ID NO: 137	SEQ ID NO: 138
26F12 27B3	NA	AACTACTATATGTCC	ATAATCAACCTAGTGGTGGT GACTCAACCTACGCACAGAAG TTCCAGGGC	GGGGGGATACAACTATGGTT ACATTTTGACTAC
		SEQ ID NO: 139	SEQ ID NO: 140	SEQ ID NO: 141
	7) 7	NYYMS	IINPSGGDSTYAQKFQG	GGIQLWLHFDY
	AA	SEQ ID NO: 142	SEQ ID NO: 143	SEQ ID NO: 144
26D1	NA	AGCTACTATATGTCC	ATAATCCACCCTAGTGGTGGT GACACAACCTACGCACAGAAG TTCCAGGGC	GGGGGGATAAAACTATGGTT ACATTTTGACTAT
		SEQ ID NO: 145	SEQ ID NO: 146	SEQ ID NO: 147
	AA	SYYMS	IIHPSGGDTTYAQKFQG	GGIKLWLHFDY

Ab	Туре	CDR 1	CDR 2	CDR 3
		SEQ ID NO: 148	SEQ ID NO: 149	SEQ ID NO: 150
25G10	NA	GGTTACTACTGGAGC	TATATCTATTACATTGGGAGC ACCAACTACAACCCCTCCCTC AAGAGT	GATGGGAGCAGTGGCTGGTA CCGGTGGTTCGACCCC
		SEQ ID NO: 151	SEQ ID NO: 152	SEQ ID NO: 153
	73. 73	GYYWS	YIYYIGSTNYNPSLKS	DGSSGWYRWFDP
	AA	SEQ ID NO: 154	SEQ ID NO: 155	SEQ ID NO: 156
23A10	NA	CGCTATGGCATACAC	GTTATATGGTATGATGGAAGT AATAAATACTATGCAGACTCC GTGAAGGGC	AGGGCCGGTATACCTGGAAC TACGGGCTACTACTATGGTA TGGACGTC
		SEQ ID NO: 157	SEQ ID NO: 158	SEQ ID NO: 159
	AA	RYGIH	VIWYDGSNKYYADSVKG	RAGIPGTTGYYYGMDV
	AA	SEQ ID NO: 160	SEQ ID NO: 161	SEQ ID NO: 162
19B5	NA	AGCTACTTTATTCAC	ATTATCAACCCTATTAGTGTT AGCACAAGCTACGCACAGAAG TTCCAGGGC	GGGGGGATACAGCTATGGTT ACATTTGGACTAC
		SEQ ID NO: 163	SEQ ID NO: 164	SEQ ID NO: 165
	73. 73	SYFIH	IINPISVSTSYAQKFQG	GGIQLWLHLDY
	AA	SEQ ID NO: 166	SEQ ID NO: 167	SEQ ID NO: 168

TABLE Ib: LIGHT CHAIN CDRs

Ab	Туре	CDR 1	CDR 2	CDR 3
1D10 2C12	NA	TCTGGAGATAGATTGG GGGAAAAATATACTTG C	CAAGATACCAAGCGGCCCTCA	CAGGCGTGGGACAGCAC TGTGGTA
		SEQ ID NO: 169	SEQ ID NO: 170	SEQ ID NO: 171
	AA	SGDRLGEKYTC	QDTKRPS	QAWDSSTVV
	AA	SEQ ID NO: 172	SEQ ID NO: 173	SEQ ID NO: 174
1F10	NA	AGGGCCAGTCGGAGTA TTAGCAGCAGCTACTT AGCC	GGTCCATCCAGCAGGGCCACT	CAGCAGTATGGTAGCTCATT CACT
		SEQ ID NO: 175	SEQ ID NO: 176	SEQ ID NO: 177
	73. 73	RASRSISSSYLA	GPSSRAT	QQYGSSFT
	AA	SEQ ID NO: 178	SEQ ID NO: 179	SEQ ID NO: 180
2C12_LC#1	NA	AGGTCTAGTCAAAGCC tcgtaTACAGTGATGG AAACAcctACTTGAAT	AAGGTTTCTAACTGGGactct	ATGCAAGGTATAGTGTGGCC GTGCAGT
		SEQ ID NO: 181	SEQ ID NO: 182	SEQ ID NO: 183
	7.7	RSSQSLVYSDGNTYLN	KVSNWDS	MQGIVWPCS
	AA	SEQ ID NO: 184	SEQ ID NO: 185	SEQ ID NO: 186
2G6_LC#1	NA	AGGTCTAGTCAAAGCC TCGTATACAGTGATGG AAACACCTACTTGAAT	CAGGTTTCTAACTGGGACTCT	ATGCAAGATACACTGTGGCC GTGCAGT
		SEQ ID NO: 187	SEQ ID NO: 188	SEQ ID NO: 189
	7) 7)	RSSQSLVYSDGNTYLN	QVSNWDS	MQDTLWPCS
	AA	SEQ ID NO: 190	SEQ ID NO: 191	SEQ ID NO: 192
2G6	NA	TCTGGAGATAGGTTGG GGGAAAAATATACTTG C	CAAGATACCAAGCGGCCCTCA	CAGGCGTGGGACAGCAC TGTGGTA

Ab	Туре	CDR 1	CDR 2	CDR 3
1 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	<i></i>	SEQ ID NO: 193	SEQ ID NO: 194	SEQ ID NO: 195
		SGDRLGEKYTC	QDTKRPS	QAWDSSTVV
	AA	SEQ ID NO: 196	SEQ ID NO: 197	SEQ ID NO: 198
2H12	NA	TCTGGAGATAGATTGG GGGAAAAATATACTTG C	CAAGATACCAAGCGGCCCTCA	CAGGCGTGGGACAGCAC TGTGGTA
		SEQ ID NO: 199	SEQ ID NO: 200	SEQ ID NO: 201
		SGDRLGEKYTC	QDTKRPS	QAWDSSTVV
	AA	SEQ ID NO: 202	SEQ ID NO: 203	SEQ ID NO: 204
2H12_LC#2	NA	AGGTCTAGTCAAAGCC TCGTATACAGTGATGG AAACACCTACTTGAAT	AAGGTTTCTAACTGGGACTCT	ATGCAAGATACACTGTGGCC GTGCAGT
		SEQ ID NO: 205	SEQ ID NO: 206	SEQ ID NO: 207
	7.7	RSSQSLVYSDGNTYLN	KVSNWDS	MQDTLWPCS
	AA	SEQ ID NO: 208	SEQ ID NO: 209	SEQ ID NO: 210
4A2 5B4 5C5	NA	AGGGCCAGTCGGAATA TTAGCAGCAGCTACtt aGCC	GGTCCATCCAGCAGGGccaCT	CAGCAGTATGGtagctCATT CACT
		SEQ ID NO: 211	SEQ ID NO: 212	SEQ ID NO: 213
	73.73	RASRNISSSYLA	GPSSRAT	QQYGSSFT
	AA	SEQ ID NO: 214	SEQ ID NO: 215	SEQ ID NO: 216
4A9	NA	ACTGGGAGCAGCTCCA ACATCGGGACAGGTTA TGCTGTACAC	GGTAACAACAATCGGCCCTCA	CAGTCCTATGACAGCagACT GAGTGGTTGGGTG
		SEQ ID NO: 217	SEQ ID NO: 218	SEQ ID NO: 219
		TGSSSNIGTGYAVH	GNNNRPS	QSYDSRLSGWV
	AA	SEQ ID NO: 220	SEQ ID NO: 221	SEQ ID NO: 222
4B10 4C2	NA	AGGGCCAGTCAGAGTG TTAGCAACACCTACTT AGCC	GGTGCATCCAGCAGGGCCACT	CAGCAGTACAGTAACTCgtg GACG
		SEQ ID NO: 223	SEQ ID NO: 224	SEQ ID NO: 225
	7.7	RASQSVSNTYLA	GASSRAT	QQYSNSWT
	AA	SEQ ID NO: 226	SEQ ID NO: 227	SEQ ID NO: 228
4D2	NA	AGGGCCAGTCAGAGTG TTAGCAACACCTACTT AGCC	GGTGCATCCAGCAGGGCCGCT	CagcagTATAGTAacTcgtg GACG
-		SEQ ID NO: 229	SEQ ID NO: 230	SEQ ID NO: 231
	AA	RASQSVSNTYLA	GASSRAA	QQYSNSWT
	AA	SEQ ID NO: 232	SEQ ID NO: 233	SEQ ID NO: 234
4D3 4F3	NA	AGGGCCAGTCAGAGTG TTAGCAGCAGCTACTT AGCC	GGTGCATCCAGCAGGGCCACT	CAGCAGTATGGTAGCTCGTG GACG
		SEQ ID NO: 235	SEQ ID NO: 236	SEQ ID NO: 237
	AA	RASQSVSSSYLA	GASSRAT	QQYGSSWT
		SEQ ID NO: 238	SEQ ID NO: 239	SEQ ID NO: 240
4E10	NA	AGGGCCAGTCAGAGTG TTGGCAGCAGCTACTT AGCC	GGTGCATCCAGCAGGGTCACT	CAGCAATATAGTAACTCGTG GACG
		SEQ ID NO: 241	SEQ ID NO: 242	SEQ ID NO: 243

Ab	Type	CDR 1	CDR 2	CDR 3
<u> </u>		RASQSVGSSYLA	GASSRVT	QQYSNSWT
	AA	SEQ ID NO: 244	SEQ ID NO: 245	SEQ ID NO: 246
4F7	NA	ACTGGGAGCAGCTCCA ATATCGGGACAGGTTA TGATGTACAC	GGTAACAGCAATCGGCCCTCA	CAGTCCTATGACAGCAGTCT GAGTGGTTGGGTG
		SEQ ID NO: 247	SEQ ID NO: 248	SEQ ID NO: 249
	AA	TGSSSNIGTGYDVH	GNSNRPS	QSYDSSLSGWV
	AA	SEQ ID NO: 250	SEQ ID NO: 251	SEQ ID NO: 252
5E3	NA	TCTGGAGATAAATTGG GGGATGAATATGCTTG C	CAAGATAGCAAGCGGCCCTCA	CAGGCGTGGGACAGCAC TGTGGTA
		SEQ ID NO: 253	SEQ ID NO: 254	SEQ ID NO: 255
	73.73	SGDKLGDEYAC	QDSKRPS	QAWDSSTVV
	AA	SEQ ID NO: 256	SEQ ID NO: 257	SEQ ID NO: 258
17H8 23B6 28D10	NA	AGGGCCAGTCAGAGTG TTGCCGGCAGCTACCT AGCC	GGTGCATCCAGCAGGGCCACT	CAGCAGTATGGTAAATCACC GATCACC
		SEQ ID NO: 259	SEQ ID NO: 260	SEQ ID NO: 261
	AA	RASQSVAGSYLA	GASSRAT	QQYGKSPIT
	AA	SEQ ID NO: 262	SEQ ID NO: 263	SEQ ID NO: 264
16C1	NA	AGGGCCAGCCAGAGTG TTAGCAGCAGCTACTT AGCC	GGTGCATCCAGCAGGGCCACT	CAGCAGTATGGTAACTCACC GCTCACT
		SEQ ID NO: 265	SEQ ID NO: 266	SEQ ID NO: 267
	7. 7.	RASQSVSSSYLA	GASSRAT	QQYGNSPLT
	AA	SEQ ID NO: 268	SEQ ID NO: 269	SEQ ID NO: 270
16A4	NA	AGGGCCAGTCAGAGTG TTAGCAGCAGTTATTT AGCC	GGTACATCCAGCAGGGCCACT	CAGCAGTACGGTAGCTCACC TTTCACT
		SEQ ID NO: 271	SEQ ID NO: 272	SEQ ID NO: 273
	AA	RASQSVSSSYLA	GTSSRAT	QQYGSSPFT
	AA	SEQ ID NO: 274	SEQ ID NO: 275	SEQ ID NO: ***276
16E2 17E10 20B12	NA	CGGGCGAGTCAGGGCA TTAGCAATTATTTAGC C	GCTGCATCCAGTTTGCAAAGT	CAACACTATTTTACTTACCC TCGGACG
		SEQ ID NO: 277	SEQ ID NO: 278	SEQ ID NO: 279
		RASQGISNYLA	AASSLQS	QHYFTYPRT
22G10	AA	SEQ ID NO: 280	SEQ ID NO: 281	SEQ ID NO: 282
22G10	NA	AGGGCCAGTCAGAGTA TTAGCAGCAACTTAGC C	GGTGCATTTACCAGGGCCACT	CAGCAGTATAATTACTGGCC GCTCACT
		SEQ ID NO: 283	SEQ ID NO: 284	SEQ ID NO: 285
	7\ 7\	RASQSISSNLA	GAFTRAT	QQYNYWPLT
	AA	SEQ ID NO: 286	SEQ ID NO: 287	SEQ ID NO: 288
16H2 20D3 23E7	NA	TCTGGAAGCAGCTCCA ACATCGGAAGTAATTT TGTAAAC	ACTAATAATCAGCGGCCCTCA	GCAACATGGGATGACAGCCT GAATGGTTGGGTG
		SEQ ID NO: 289	SEQ ID NO: 290	SEQ ID NO: 291
	7\ 7\	SGSSSNIGSNFVN	TNNQRPS	ATWDDSLNGWV
	AA	SEQ ID NO: 292	SEQ ID NO: 293	SEQ ID NO: 294

Ab	Туре	CDR 1	CDR 2	CDR 3
22D1	NA	TCTGGAAGCAGCTCCA ACATCGGAAGCAATTT TGTAAAC	ACTAATAATCAGCGGCCCTCA	GCAACATGGGATGACAGTAT GAATGGTTGGGTG
		SEQ ID NO: 295	SEQ ID NO: 296	SEQ ID NO: 297
	AA	SGSSSNIGSNFVN	TNNQRPS	ATWDDSMNGWV
	AA	SEQ ID NO: 298	SEQ ID NO: 299	SEQ ID NO: 300
25F8	NA	TCTGGAAGCAGCTCCA ACATCGGAAGGAATTT TGTAAAC	ACTAATAATCAGCGGCCCTCA	GCAGCATGGGATGACAGCCT GAATGGTTGGGTG
		SEQ ID NO: 301	SEQ ID NO: 302	SEQ ID NO: 303
	AA	SGSSSNIGRNFVN	TNNQRPS	AAWDDSLNGWV
	AA	SEQ ID NO: 304	SEQ ID NO: 305	SEQ ID NO: 306
26F12 27B3	NA	TCTGGAAGCCGCTCCA ACATCGGAAGTAATTT TGTAAAC	ACTAATTATCAGCGGCCCTCA	GCAGTATGGGATGACAGCCT GAATGGTTGGGTG
		SEQ ID NO: 307	SEQ ID NO: 308	SEQ ID NO: 309
	AA	SGSRSNIGSNFVN	TNYQRPS	AVWDDSLNGWV
	AA	SEQ ID NO: 310	SEQ ID NO: 311	SEQ ID NO: 312
26D1	NA	TCTGGAAGCCGCTCCA ACATCGGAAGTAATTT TGTAAAC	ACTAATAATCAGCGGCCCTCA	GCAGTATGGGATGACAGCCT GAATGGTTGGGTG
		SEQ ID NO: 313	SEQ ID NO: 314	SEQ ID NO: 315
	7\ 7\	SGSRSNIGSNFVN	TNNQRPS	AVWDDSLNGWV
	AA	SEQ ID NO: 316	SEQ ID NO: 317	SEQ ID NO: 318
25G10	NA	AGGGCCAGTCAGAGTG TTAGCAGCAGCTACTT AGCC	GGTGCATCCAGCAGGGCCACT	CAGCAGTATGGTAACTCACC GCTCACT
		SEQ ID NO: 319	SEQ ID NO: 320	SEQ ID NO: 321
	70.70	RASQSVSSSYLA	GASSRAT	QQYGNSPLT
	AA	SEQ ID NO: 322	SEQ ID NO: 323	SEQ ID NO: 324
23A10	NA	TCTGGAGATAGATTGG GGGAGAAATATGTTTG C	CAAGATAATAAGTGGCCCTCA	CAGGCGTGGGACAGCAC TGTGGTA
		SEQ ID NO: 325	SEQ ID NO: 326	SEQ ID NO: 327
	AA	SGDRLGEKYVC	QDNKWPS	QAWDSSTVV
	AA	SEQ ID NO: 328	SEQ ID NO: 329	SEQ ID NO: 330
19B5	NA	TCTGGAAGCAGTCCA ACATCGGAAGCAATTT TGTAAAC	ACTAATAATCAGCGGCCCTCA	GCAACATGGGATGACAGTAT GAATGGTTGGGTG
		SEQ ID NO: 331	SEQ ID NO: 332	SEQ ID NO: 333
	7\ 7\	SGSRSNIGSNFVN	TNNQRPS	ATWDDSMNGWV
	AA	SEQ ID NO: 334	SEQ ID NO: 335	SEQ ID NO: 336

Anti-CDH19 Variable Region Amino Acid Sequences and Polynucleotide Sequences

TABLE IIa: Heavy Chain Variable Region Polynucleotide and Amino acid Sequences

	,			
SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
D N O.				
337	17H8 23B6 28D10	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGCCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACGTGCACTGTCTCTGGTGGCTCCAT CAATAGTTACTTACTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATATTACATTACATTGGGAGCACCA ACTACAACAACCCTTCTACAAGAGTCGCGTCACCATATTCAGTAGAAGAACTACAACAACAACAACAACTACACTACAAAAAAAA
338	17H8	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT
	23B6 28D10			AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSS
339	4A2	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCACAGACCCTGTCCCTCACCTGCACTGCTCTCTGGTGGCTCCAT
	5B4			CAGCAGTAGTGGTTACTACTGGAGCTGGATCCGCCAGCACCCAGGGAAGGGCCTGGAGTGGATTGGGTACATCTATTACACTGGGA
	5C5			GCGCCTACTACAACCCGTCCCTCAAGAGTCGAGTTACCATATCAGTAGACACGTCTAAGAACCAGTTCTCCCTGAAGCTGAGCTCTT
				GTGACTGCCGCGGACACGGCCGTGTATTACTGTGCGAGAGATGGAAGCAGTGGCTGGTACTTCCAGTATTGGGGCCAGGGCACCCT
1 4	7.47	loioitito	6	GGICACCGICICA AMO DESCRITIVESAMI SI MAMVISCOSI SECAVAMISMI ENIDAZZOI EMICATAVECSA VANIBSI KERIMI SIMPESAMO ESI ZI SE
340	4A2 5B4	ariiiciai ariiiciai	g V	QVQLQESGFGLVKFSQTLSSLICTVSGGSLSSSGIIWSWIKQHFGRGLEWIGIIIITGSAIINFSLKSKVIISVDISKNQESLKSS VTBBDTBVVVCBRDGSGGWVFDVWGDGTIVTVSS
	5C5			
341	16H2	artificial	nt	CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGACCTGGGGCCTCAGTGAAGGTTTCCTGCAAGGTTTCTGGATACACCTT
	20D3			CACCAGCTACTTTATTCACTGGGTGCGCCAGGCCCCTGGACAAGGGCTTGAGTGGATGGGAATAATCAACCCTATTAGTGTTAGCA
	23E7			CAAGCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACAGTCTTCATGGAGCTGAGCAGCCTG
				AGATCTGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGGATACAGCTATGGTTACATTTTGACTACTGGGGCCAGGGAACCCT gentalogicalitics
242	16H2	artificial	a	
7	20D3	5	3	RSEDTAVYCARGGIQLWLHFDYWGQGTLVTVSS
343	26F12	artificial	ıt	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGCAAGGCATCTAGATACACCTT
	27B3			CACCAACTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGGCTTGAGTGGATGGGAATAATCAACCCTAGTGGTGGTGACT
				AGATCTGAGGACACGGCCGTGTATTACTGTGCGAGAGGGGGGGATACAACTATGGTTACATTTTTGACTACTGGGGGCCAGGGAACCCT
				GGTCACCGTCTCCTCA
344	26F12	artificial	aa	
	27B3			
345	4B10	artificial	Ħ	CAGGTGCAGTTGGTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCACCTT
	4C2			CAGTAGCTATGACATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
Ö. Ö.				
				AATACTATGCAGACTCCGTGAAGGGCCGATTCACCATCTCCAGAGACACTTCCAAGAACACGCTGTATTTGCAAATGAACAGCCTG AGAGCTGAGGACACGGCTGTATATTACTGTGCGAGAGAACGATATTTTGACTGGTCTTTTGACTACTGGGGCCAGGGAACCTGGT CAGTGTCTCCTCA
346	4B10 4C2	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS
347	4F3	artificial	t	CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCTCCTT CAGTAGCTATGACATGGACTGGGTCCGCCAGACTCCAGGGCGTGGAGGGGCTGGAGTGGTATGATGAATA AATACTATGCAGACTCCGTGAGGGCCCGATTCACCATCTCCAGAGACATTCCAAGAACAGCCTGTTTCTGCAAATGAACAGCCTG AGAGTCGAGGACACGGCTGTGTTACTGTGCGAGAAACTGGGGGCTGGTACTTCGATCTCTGGGGCCGTGGCACCTGGT CACTGTCTCTCC
348	4D3 4F3	artificial	aa	QVQLVESGGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS
648 143	16E2 17E10 20B12	artificial	t	CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGGTCCAGGCGTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCATCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGACTCCAGGCAAGGGGCTGGAGTGGTGGTGATGGATG
350	16E2 17E10 20B12	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFIFSSYGMHWVRQTPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDISKNTLYLQMNSL RVEDTAVYYCARDGWELSFDYWGQGTLVTVSS
351	1D10 2C12	artificial	t	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGGTCCCAGGCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGTCAGTTATATGGTATGATGGAAGTAATA AATACTATGCAGACTCCGTGAAGGGCCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTTTCTGCAAATGAATAGCCTG AGAGCTGAGGACACGGCTGTGTATTACTGCGCGAGAAGGGCCGGTATAGGAACTACAGGCTACTACGGTATGGACGTCTG GGGCCAAGGGACCACGGTCTCCTCA
352	1D10 2C12	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVSVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTTGYYYGMDVWGQGTTVTVSS
353	16C1	artificial	t	CAGGTGCAGCTGCAGGAGTCGGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACTTGTACTGTCTCTGGTGGCTCCAT CAGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATACTTTACATTGGAGCACCCA ACTACAACCCCTCCCTCAAGAGTCACCATGTCAATAGACACGTCCAAGAACCAGTTCTCCCTGACGCTGAGCTTTGACC GCTGCGGACACGGCCGTGTATTTCTGTGCGAGAGATGGGAGCAGTGGCTGGTACCGGTGGTTCGACCCTGGGGCCAGGGAACCCT GCTGCCGCACCGCTCTCTATTTCTGTGCGAGAGATGGGAGCAGTGGCTGGTACCGGTGGTTCGACCCTTGGGGGCCAGGGAACCCT GGTCCCGCTCCTCA
354	16C1	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSIDTSKNQFSLTLSSLT AADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSS
355	25G10	artificial	t	CAGGTGCAGCTGCAGGAGTCGGGCCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGCACTGTCTCTGGTGGCTCCAT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9.				
				CAGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATATCTATTACATTGGAACCACCCAACCAA
356	25G10	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSS
357	16A4	artificial	ŧ	CAGGTGCAGCAGGAGTCGGGCCCAGGACTGGCGAAGCCttcGGAGACCctgtccctcacctgCACTGTCTCTGGTGACTCCAT CACTAGTTACTACTGCAGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATATCTATTACAGTGGGAGCACCA ATTACAACCCCTCCCTCCAGAGTCACCATATCAGTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGTTCTGTGACC GCTGCGGACACGGCCGTGTATTACTGTGCGAGAGTCAAAAGGCGGATAGCAGCTGGTACCCACTTCTACGGTATGGACGTCTG GCGCCAAGGGACCACGGCCACCTCTCCTCC
358	16A4	artificial	aa	QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTTVTVSS
359	1F10	artificial	Ħ	CAGGTGCAGCTGCAGGAGTCGGGCCCCAGGACTGGTGAAGCCTTCACAGACCCTGTCCCTCACCTGCACTGTCTCTGGTGGCTCCAT
144				
360	1F10	artificial	aa	QVQLQESGPGLVKPSQTLSLTCTVSGGSISSGGYYWSWIRQHPGKGLEWIGYIYYSGSTYYNPSLTSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQHWGQGTLVTVSS
361	4A9	artificial	Ħ	CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGTCTCTGGTGGCTCCAT CAGGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGAAAGGGACTGGAGTGGTTTGCATATTTCTCTTTACAGTGGGAGCACCA ACTACAACCCCTCCCTCCAAGAGTCGAGTC
362	4A9	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSS
363	4F7	artificial	Ħ	CAGGTGCAGCTGCAGGAGTCGGGACTTGGTGAAGCCTTCGGAGACCCTGTCCTCACCTGCACTGTCTCTGGTGGCTCCAT CAGTAGTTACTCCTGGAGCTGGATCCGGCAGCCCCAGGGAAGGGACTGGAGTGGATTGGGTATTACAGTGGGAGCACCA ACTACAACCCCTCCCTCCAAGAGTCGAGTC
364	4F7	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSS
365	22D1	artificial	ıt	CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAGGGTTTCCTGCAAGGTTTCTGGATAACACCTT CACCAGCTACTTAATTCACTGGGTACGCCTAGGTTAGTGTTAGTGTTAGCA

C	DESIGNATION	SOLIDO	TVDE	CEOI IENCE
Z Z Z	DESIGNATION	SOORCE		
NO.				
				CAAGCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACAGTCTTCATGGAGCTGAGCCTG AGATCTGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGGATACAGCTATGGTTACATTTGGACTACTGGGGCCAGGGAACCCT GGTCACCGTCTCCTCA
366	22D1	artificial	aa	QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
367	1985	artificial	ŧ	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGCAAGGTTTCTGGATACACCTT CACCAGCTACTTTATTCACTGGGTGCGCCCAGGCCCCTGGACAGGCCTTGAATGGATGG
				CAAGCIACGCACAGAAGITCCAGGGCGAGGGCGATACACGICCACGAGCACAGICITCATGGAGCIGAGCCIG AGAICTGAGGACACGGCCGTGTATTACTGTGCGCGGGGGGGGATACAGCTATGGTTACATTTGGACTACTGGGGGGAACCCT GGTCACCGTCTCCTCA
368	19B5	artificial	aa	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
369	25F8	artificial	Ħ	CAGGIGCAGCIGCAGCICGGGCTGAGGIGAAGAAGCCIGGGGCCICAGIGAAGGIITICCIGCAAGGCAICIGGAIACACCIT
				CACCAGCTACTATATTCACTGGGTGCGCCCAGGCCCCTGGACAAGACTTGAGTGGATGGGAATAATCAACCCCAGTGGTGGTAGCA
				CAAGGTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACAGCACACTCTTCATGGAGCTGAGCAGCCtG
14				AGAICIGAGGACACGCCGIGIAITACIGCGCGAGGGGGGAAIACAGCIAIGGITACAITCGGACIACIGGGCGGCCAGGGAACCCI GGTCACCGTCTCCTCA
370	25F8	artificial	aa	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVFMELSSL
				RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
371	26D1	artificial	r T	CAGGIGCAGIIGGIGCAGICIGGGGCIGAGGIGAAGAAGCCIGGGGCCICAGIGAAGGIIICCIGIAAGGCAICIAGAIACACCII
				CACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGGCTTGAGTGGATGGGAATAATCCACCCTAGTGGTGGTGACA
				CAACCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCGGGGACACGTCCACGAGCACAGTCTACATGGAGCTGAGCAGCCTG
				AGAICTGAGGACACGGCCGTGTATTACTGTGCGAGAGGGGGGATAAAACTATGGTTACATTTTGACTATTGGGGCCAGGGAACCCT GGTCACCGTCTCCTCA
372	26D1	artificial	aa	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL
373	4D2	artificial	Ħ	
				CAGTAGTTATGACATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGGCTGGAGTGGGTGG
				AGAGCTGAGGACACGGCTGTATATTACTGTGCGAGAGCAACGATATTTTGACTGGTCTTTTGACTACTGGGGCCCAGGGAACCCTGGT
				CAGTGTCTCCTCA
374	4D2	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPCKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL Bardamanivacardedstringerdvægogtingss
r L	717	10:0:3:40	-	
375	4E10	artificial	ĭ	CAGGIIGCAGCIIGGIIGGAGIICIIGGGGGGGGGGGGG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>o</u> 8				
5				AGAGCCGAGGACACGGCTGTGTACTACTGTGCGAGAGAGTATAGGTACAGCTGGTACTTTGACTACTGGGGGCCCAGGGAACCCTGGT
376	4E10	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSTNTLHLQMNSP RAEDTAVYYCAREYRYSWYFDYWGQGTLVTVSS
377	22G10	artificial	t	GAGGTGCAACTGTTGGAGTCTGGGGAGGCTTGGTACAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCACCTT TAGCAGTTATGCCATGAACTGGGTCCGCCAGGCTCCCAGGGAAGGGGCTGGAGTGGGTCTCAACTATTAGTGGTGGTGGTGCTAAACA CATACTACGCAGACTCCGTGAAGGGCCGGTTCACCATCTCCAGTGACAATTCCAAGAGCACGCTGTATTGCAAATGAACAGCTG AGAGCCGCGGACACGCCCGTATATCACTGTGCGAAAGGGGGGATTGGGGGGGATACTACGGTATGGACGTTGGGGGCCAAGGGGAC
378	22G10	artificial	aa	CACGGICACCGICICCICA EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISSDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGOGTTVTVSS
379	2C12_LC#1	artificial	Ħ	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGGGTCCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGGATTCACCTT
				CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGTCAGTTATATGTGTGGTATGGAAGTAATA AATACTATGCAGACTCCGTGAAGGGCCGATTCACAATCTCCAAGAACAGAACAGAACAGTGTATCTGCAAATGAATAGCCTG
				AGAGCTGAGGACACGGCTGTGTATTACTGCGCGAGAAGGGCCCGGTATAATAGGAACTACAGGCTACTACTACGGTATGGACGTCTG GGGCCAAGGGACCACGGTCACCGTCTCCTCA
08 146	2C12_LC#1	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVSVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTTGYYYGMDVWGQGTTVTVSS
381	2H12_LC#2	artificial	nt	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGGGGTCCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT
				CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
				AATACTATACAGACTCCGTGAAGGGCCGATTCACCATCTCCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAATAGCCTG
				AGAGCIGAGGACACGCIGIGIAITACIGIGGAGAGGGCCGGIAIAAIAGGAACIACAGGCIACIACIACGGIAIGGACGICIG GGGCCAAGGGACCACGGICACCGICICCCICA
382	2H12_LC#2	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAVIWYDGSNKYYTDSVKGRFTISRDNSKNTLYLQMNSL
				RAEDTAVYYCARRAGIIGTTGYYYGMDVWGQGTTVTVSS
383	2G6_LC#1	artificial	t	
				CAGTAGCTATGGCATGCACTGGGTCCGCCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
				AATACTATGCAGACTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACAGCTGTATCTGCAAATGAAAAGCCTG
				AGAGCIGAGGGACCACGGTCACGTAIIACIGIGAGAAGGGCCGGIAIAAIAGGAACIAIAGGCIACIACIACGACGGIAIGGACGICIG GGGCCAAGGGACCACGGTCACCGTCTCCTCA
384	2G6_LC#1	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL
				RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
385	2H12	artificial	nt	CAGGIGCAGCIGGIGGAGICIGGGGGAGGCGIGGICCAGCCIGGGAGGICCCIGAGACICICCIGIGCAGCGICIGGAITCACCII
				CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
				AATACTATACAGACTCCGTGAAGGGCCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAATAGCCTG
				AGAGCTGAGGACACGGCTGTGTATTACTGTGCGAGAAGGGCCGGTATAATAGGAACTACAGGCTACTACTACGGTATGGACGTCTG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
_				
NO.				
				GGGCCAAGGGACCACGGTCACCGTCTCCTCA
386	2H12	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAVIWYDGSNKYYTDSVKGRFT1SRDNSKNTLYLQMNSL
				RAEDTAVYYCARRAGIIGTTGYYYGMDVWGQGTTVTVSS
387	2G6	artificial	nt	CAGGIGCAGITGGIGGAGICTGGGGGAGGCGIGGICCAGCCIGGGAGGICCCTGAGACICTCCTGIGCAGCGICTGGAITCACCIT
				CAGTAGCTATGGCATGCACTGGGGTCCGCCAGGCTCCAGGCAAGGGGCCTGGAGTGGGTGG
				AATACTATGCAGACTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACACCGCTGTATCTGCAAATGAAAAGCCTG
				AGAGCIGAGGACACGGCIGIGIATIACIGIGCGAGAAGGGCCGGIATAAIAGGAACIATAGGCIACIACIACGGIAIGGACGICIG
				GGGCCAAGGGACCACGGTCACCGTCTCCA
388	2G6	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL
				RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
389	23A10	artificial	nt	CAGGTGCAGCTGGTGGAGTCTGGGGGGAGGCGTGGTCCAGCCTGGGGAGGTCCCTGAGACTCTCTGTGCAGCGTCTGGATTCACCTT
				CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
				AATACTATGCAGACTCCGTGAAGGGCCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCTAATGAACAGCCTG
				AGAGCCGAGGACTCGGCTGTGTATTACTGTGCGAGAAGGGCCGGTATACCTGGAACTACGGGCTACTACTATGGTATGGACGTCTG
				GGGCCAAGGGACCACGGTCACCGTCTCCA
06E 1	23A10	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL
47				RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
391	5E3	artificial	nt	GAGGTGCAGTTGGTGGAGTCTGGGGGAGGCCTGGTCAAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCACCTT
				CAGTAGCTATAGCATGCACTGGGTCCGCCAGGCTCCAGGGAAGGGGCTGGAGTGGGTCCATCCA
				TATACTACGCAGACTCAGTGAAGGGCCGATTCACCATCTCCAGAGACAACGCCAAGAACTCACTGTATCTGCAAATGAACAGCCTG
				AGAGCCGAGGACACGGCTGTGTATTACTGTGCGAGAGGGGAAACTGGAACTAACT
				AGGGACCACGGTCACCGTCTCCTCA
392	5E3	artificial	aa	EVQLVESGGGLVKPGGSLRLSCAASGFTFSSYSMHWVRQAPGKGLEWVSSISSSSSYIYYADSVKGRFTISRDNAKNSLYLQMNSL
				RAEDTAVYYCARGETGTNYYYYGMDVWGQGTTVTVSS

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IAB	' LE IIB: Light (Chain Var	iable R	I ABLE IIB : Light Chain Variable Region Polynucleotide and Amino acid Sequences
SEQ	SEQ DESIGNATION SOURCE TYPE SEQUENCE	SOURCE	TYPE	SEQUENCE
₽				
NO.				
393	17H8	artificial	пt	GACATIGIAITGACGCAGtetCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCTCTCCTGCAGGGCCAGTCAGAGTGT
	23B6			TGCCGGCAGCTACCTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCATCTCTGGTGCATCCAGCAGGCCACTG
	28D10			GCATCCCAGACAGGTTCAGTGGCAGTGGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCTTTTTGCAGTG
				IAITACTGTCAGCAGTATGGTAAATCACCGATCACCTTCGGCCAAGGGACACGACTGGAGATGAAAGGA
394	394 17H8	artificial	aa	DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV
	23B6			YYCOOYGKSPITFGOGTRLEMKG

SEO	DESIGNATION	SOURCE	TYPE	SEOUENCE
Q Q				
	28D10			
395	4A2	artificial	nt	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGGCCCACCCTCTTGCAGGGCCAGTCGGAATAT TAGCAGCAGCTACTTAGCTTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGGCTCCTCATCTATGTCCATCCA
	5C5			GCATCCCAGACAGATTCAGTGGCAGTGGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCCTGAAGATTTTACAGTG TATTTACTAGCTAGTTAGCTTAGC
				TO CONTROL TO THE PROPERTY OF
396	4A2	artificial	aa	EIVLIQSPGTLSLSEGERATLSCRASRNISSSYLAWYQQRPGDAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFTV VYCOOVASSFFFGDGTRVDIRR
	5C5			1
397	16H2	artificial	Ħ	CAGICIGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGGGGTCACCATCTCTTGTTCTGGAAGCAGCTCCAACAT
	20D3			CGGAAGTAATTTTGTAAACTGGTACAACACCAACTCCCAGGAACGGCCCCCCAAAGTCCTCATCTATAATAATCAGCGGCCCTCAG
	23E7			GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGTCTGAT TATTACTGTGCAACATGGGATGACAGCCTGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCGTCCTAGGT
398	16H2	artificial	aa	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD
	20D3			YYCATWDDSLNGWVFGGGTKLTVLG
	23E7			
66E 14	26F12	artificial	nt	CAGTCTGTGCTGACTCAGCCTCAGCGTCTGGGACCCCGGGCAGAAGGTCACCATCTCTTGTTCTGGAAGCCGCTCCAACAT
	27B3			CGGAAGTAATTTTGTAAACTGGTACCAGCAGCTCCCAGGAACGGCCCCCCAAACTCCTCATCTATTATTATCAGCGGCCCTCAG
				GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGAT mammacmcoacaamacaancaancaaancaaaagactgagagacaaaaagacaacaagaaagaaaaaaaa
400	26F12	artificial	aa	OSVLTOSPSASGTPGOKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLOSEDEAD VYCAVWDDSINGWVFGGGTKITVI.G
	7/02			
401	4B10	artificial	r	GAAATTGTATTGACGCAGTCTCCCAGGCACCCTGTCTTTGTCTCCCAGGGGAAAGAGCCCACCCTCCTGCAGGGCCAGTCAGAGTGT
	4C2			TAGCAACACCTACTTAGCCTGGTACCATCAGAGACCTGGCCTCCCAGGCTCCTCATCTATGGTGCATCCAGCAGGCCACTG
				TATTACTGTCAGCAGTACAGTAACTCGtgGGACGTTCGCCAAGGGACAAGGGAAATCAaacGA
402	4B10	artificial	aa	EIVLIQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFALTISSLEPEDFAV
	4C2			YYCQQYSNSWIFGQGTKVEIKR
403	4D3	artificial	nt	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGAAAGAGCCACCCTCCTGCAGGGCAGTGTGT
	4F3			TAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCTTATGGTGCATCCAGCAGGGCCACTG
				GCAICCCAGACAGGITCAGIGGCAGIGGGICIGGGACAGACIICACICICACCAICAGCAGAACIGGAACCIGAGGAITIIGCAGIG TAITACIGICAGCAGIAIGGIAGCICGIGGACGIICGGCCAAGGGACCAAGGIGGAAAICAAACGA
404	4D3	artificial	aa	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPCQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV
	4F3			YYCQQYGSSWTFGQGTKVEIKR
405	16E2	artificial	nt	GACATCCAGATGACCCAGTCTCCATCCTCACTGTCTGCATCTGTAGGAGACAGAGTCACCATCACTTGTCGGGCGAGTCAGGGAT TAGCAATTATTTAGCCTGGTTACAGCAGAAACCAGGAAAGCCCCTAAAGTCCTTGATCTTATGCAGTCAAGGTCAAGGGG
	T/EIU))))));;;;;;));;;));;;));;;));;;;));;;;;

C	DICITAINOIDE	1701102	TVDL	JONAI I CAS
g ⊡ 8	DESIGNATION	SOURCE		SECONICE
	20B12			TCCCATCAAAGTTCAGCGGCAGTGGATCTGGGACAGATTTCACTCTCACCATCAGCAGCCTGCAGCCTGAAGATTTTGCAACTTAT TACTGCCAACACTATTTTACTTACCCTCGGACGTTCGGCCAAGGACCAAGGTGGAAATCAAACGA
406	16E2	artificial	aa	DIQMTQSPSSLSASVGDRVTITCRASQGISNYLAWLQQKPGKAPKSLIYAASSLQSGVPSKFSGSGSGTDFTLTISSLQPEDFATY
	17E10 20B12			YCQHYFTYPRTFGQGTKVEIKR
407	1D10	artificial	nt	
	2C12			
				CTGAGCGATTCTCTGGCTCCACCTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTAC TGTCAGGCGTGGGACAGCAGCACCTGTGGTATTCGGCGGAGGGACCAAGCTGACCGTCCTAGGT
408	1D10 2C12	artificial	aa	SYALTQPPSVSVSPGQTASLICSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSTSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVLG
409	16C1	artificial	nt	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCTGCAGGGCCAGCCA
				TAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCATCTTTGGTGCATCCAGGGCCACTG
				GCATCCCAGACAGGTTCAGTGGCAGTGGGTCTGGGACAGACTTCACTCTCACCATCAGCGGACTTGGAAGATTTTGCAGTG
410	16C1	artificial	aa	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTDFTLTISGLEPEDFAV
49				YHCQQYGNSPLTFGGGTKVEIKR
411	25G10	artificial	Ħ	GAAATTGTGTTGACGCAGTCTCCCAGGCACCCTGTCTTTGTCTCCCAGGGGAAAGAGCCCACCCTCTCCTGCAGGGCCAGTCAGAGTGT
				TAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCATCTTTGGTGCATCCAGCGGCCACTG
				TATCACTGTCAGCAGTATGGTAACTCACCGCTCACTTTCGGCGGAGGGACCAAGGTGGAGATCAAACGA
412	25G10	artificial	aa	
				YHCQQYGNSPLTFGGGTKVEIKR
413	16A4	artificial	Ħ	
				TAGCAGCAGTTATTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCATCTATGGTACATCCAGCAGGGCCACTG
				GCAICCCAGACAGGITCAGIGGCAGIGGGICIGGGACAGACIICACICICCACCAICAGCAGACIGGAGCCIGAAGAITIIGCAGIG
717	7607	Licititic	5	FIXT TO CONCINCT SOCREDULIS SOCRES SO
† †	101	<u>a</u>	0	
415	1F10	artificial	п	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCCACCCTCTCCTGCAGGGCCAGTCGGAGTAT
416	1F10	artificial	aa	EIVLTQSPGTLSLSPGERATLSCRASRSISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV
				II/VXIGOSI II GEGIN/DIN/

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> №				
417	4A9	artificial	nt	CAGTCTGTGCTGACGCAGCCGCCCTCAGTGTCTGGGGCCCCAGGACAGAGGGTCACCATCTCCTGCACTGGGAGCTCCAACAT CGGGACAGGTTATGCTGTACACTGGTACCAGCTTTCCAGGAACAGCCCCCAAACTCCTCATCTATGGTAACAACAATCGGCCCT CAGGGGTTCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCCTGGCCATCATCGGGTTCAGGATGAGGATGAGGCT GATTATTACTGCCAGTCCTATGACAGCAGCTGAGGTGGTTGGGTGTTCGGCGGAGGGACGAACCTGAGGTCTAGGT
418	4A9	artificial	aa	QSVLIQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQFPGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSRLSGWVFGGGTKLIVLG
419	4F7	artificial	nt	CAGTCTGTGCTGACGCAGCCGCCCTCAGTGTCTGGGGCCCCAGGGCAGAGGGTCACCATCTCCTGCACTGGGAGCAGCTCCAATAT CGGGACAGGTTATGATGTACACTGGTATCAGCAGCTCCCAGGAACAGCCCCCAAACTCCCTCATCCTTGGTAACAGCAATCGGCCCT CAGGGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCACTGGGCTCCAGGCTGAGGATGAGGCT GATTATTACTGCCAGTCCTATGACAGCAGTCTGAGGTTGGGTGTTCGGCGGAGGGACGAGGTTGACTAGGT
420	4F7	artificial	aa	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSSLSGWVFGGGTRLTVLG
421	22D1	artificial	nt	CAGTCTGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACTCTTGTTCTGGAAGCTCCAACAT CGGAAGCAATTTTGTAAACTGGTACAAGCAGCTCCCAGGAACGGCCCCCCAAAGTCCTCTTATACTAATAATCAGCGGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCCATCAGTGGGCTCCAGTCTGAGGATGAGTCTGAT TATTACTGTGCAACATGGATGAATGAATGGTTGGGTGTTCGCGGAGGGACCAAGCTGACCTAGGT
455	22D1	artificial	aa	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCATWDDSMNGWVFGGGTKLTVLG
423	1985	artificial	t	CAGTCTGCGCTGACTCAGCCACCCTCAACGACTGGGACCCCCGGGCAGAGGGTCACCATCTTGTTCTGGAAGGTCCAACAT CGGAAGCAATTTTGTAAACTGGTACAAGCAGCTCCCAGGAACGGCCCCCAAGAGTCCTCATCTATACTAATAATCAGCGGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCATCAGTGGGCTCCAGTCTGAGATGATGATCAGT TATTACTGCGCAACATGGGATGACAGTATGAATGGTTGGGTGTTCGGCGAGGGACCAAACTGACCTAGGT
424	1985	artificial	aa	QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCATWDDSMNGWVFGGGTKLTVLG
425	25F8	artificial	nt	CAGTCTGCGCTGactCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTCTTGTTCTGGAAGCTCCAACAT CGGAAGGAATTTTGTAAACTGGTATAAGCAGCTCCCAGGAACGGCCCCCCAAAGTCCTCATTTATACTAATAATCAGCGGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCATCAGTGGGCTCCAGTCTGAGATGATGATCTGAT TATTACTGTGCAGCATGGGATGACAGCTGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCTAGGT
426	25F8	artificial	aa	QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCAAWDDSLNGWVFGGGTKLTVLG
427	26D1	artificial	nt	CACTCTGTGCTGACTCAGTCACCCTCAGCGTCTGGGACCCCCGGACAGAGGGTCACCATCTTGTTCTGGAAGCCGCTCCAACAT CGGAAGTAATTTTGTAAACTGGTACCAGCAGCTCCCAGGAACGGCCCCCAAACTCCTCTTATACTAATAATCAGCGGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGAT TATTACTGTGCAGTATGGGATGACAACCTGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTCAAGGT
428	26D1	artificial	aa	HSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDDSLNGWVFGGGTKLTVLG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
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429	4D2	artificial	nt	GAAATTGTATTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCTGCAGGGCCAGTCAGAGTGT TAGCAACACCTATTAGCCTGGTACCATCAGAGACCTGGCCAGGCTCCCAGGCTCCTTATGGTGCATCCAGCGGCCGCTG GCATCCCAGACAGGTTCAGTGGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTTGAAGATTTTGCAGTG TATTACTGTCAGCAGTATAGTAACTCGTGGACGTTCGGCCAAGGGACCAAGGTGAAATCAAACGA
430	4D2	artificial	aa	EIVLIQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRAAGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYSNSWTFGQGTKVEIKR
431	4E10	artificial	nt	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCTGCAGGGCCAGTCAGAGTGT TGGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCTCCCAGGCTCCTCTATGGTGCATCCAGGGTCACTG GCATCCCAGACAGGTTCAGTGGCAGTGGGTCTGGGACAGATTTCACTCTCACCCATCAGCAGACTGAAGATTTTGCAGTG TATTACTGTCAGCAATATAGTAACTCGTGAACGTTCGGCCAAGGGACCAAGGTGGAAATCAAAACGA
432	4E10	artificial	aa	EIVLTQSPGTLSLSPGERATLSCRASQSVGSSYLAWYQQKPGQAPRLLIYGASSRVTGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYSNSWTFGQGTKVEIKR
433	22G10	artificial	nt	GAAATAGTGACGCAGTCTCCAGTCACCCTGTCTCTGTCTCTAGGGGAAAGAGCCCACCCTCTCTGCAGGGCCAGTCAGAGTAT TAGCAACATTAGCCTGGTTCCAGCAGAAACCTGGCCAGGCTCCCAGACTCCTCATCTATGGTGCATTTACCAGGGCCACTGGTA TCCCAGCCAGGGTCAGTGGCAGTGGGTCTGGGACAGAGTTCACTCTCACCATCAGCAGCCTGCAGTTTTGCAGTTTAT TACTGTCAGCAGTATAATTACTGGCCGCTCACTTTCGGCGAGGGACCAAGGTGAAGATTTTGCAGTTTAT
51 51	22G10	artificial	aa	EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVY YCQQYNYWPLTFGGGTKVEIKR
435	2C12_LC#1	artificial	nt	GATGTIGTGATGACTCAGtCTccActctcctgcCCGTCACCCTTGGACAGCCGGcctCCAtctcctgCAGGtCTAGTCAAGcct cgtaTACAGTGATGGAAACAcctACTTGAATTGGTTTCAGCAGGGCCAGGCCAATCTCCCAAGGcgcctaATTTATAAGGTTTCTA ACTGGGactctGGGGtCCCAGACAGATTCAGCGGCAGTGGGTCAGGCACTGATTTCACactGAAAAtCAGCAGGGTGGaggctgaG GATGTTGGGGTTTATTactgCATGCAAGGTATAGTGTGGCCGTGCAGTTTTGGCCAGGGGACCAAGCTGGAGATCAAACGA
436	2C12_LC#1	artificial	aa	DVVMTQSPLSLPVTLGQPASISCRSSQSLVYSDGNTYLNWFQQRPGQSPRRLIYKVSNWDSGVPDRFSGSGSGTDFTLKISRVEAE DVGVYYCMQGIVWPCSFGQGTKLEIKR
437	2H12_LC#2	artificial	nt	GATGTIGTGATGACTCAGTCTCCACTCTCCCTGCCCGTCACCCTTGGACAGCCGGCCTCCATCTCTGCAGGTCTAGTCAAAGCCT CGTATACAGTGATGGAAACACCTACTTGAATTGGTTTCAGCAGGGCCAGGCCAATCTCCAAGGGCCTTAATTATAAGGTTTCTA ACTGGGACTCTGGGGTCCCAGACAGAATCAGCGGCAGTGGGGTCAGGCACTTTCACACTGAAAATCAGCAGGTGGAGGCTGAG GATGTTGGGGTTTATTACTGCAAGATACACTGTGGCCGTGCAGTTTTGGCCAGGGGACCAAGCTGGAGATCAAACGA
438	2H12_LC#2	artificial	aa	DVVMTQSPLSLPVTLGQPASISCRSSQSLVYSDGNTYLNWFQQRPGQSPRRLIYKVSNWDSGVPDRISGSGSGTDFTLKISRVEAE DVGVYYCMQDTLWPCSFGQGTKLEIKR
439	2G6_LC#1	artificial	nt	GATGTIGTGATGACTCAGLCCCACTCTCCCTGCCCGTCACCCLtggacaGCCGGCCTccaTCTCCTGCAGGTCTAGTCAAGCCT CGTATACAGTGAAACACCTACTTGAATTGGTTTCAGCAGAGGCCAGGCCAATCTCCACGGCGCCCTAATTTATCAGGTTTCTA ACTGGGACTCTGGGGTCCCAGACAGATTCAGCGGCGCAGTGGGTCAGGCACTGATTTCACACTGAAAATCAGCAGGTGGAGGCTGAG GATGTTGGGATTTATTACTGCAAGATACAAGATACACTGTGGCCGTGCAGTTTTTGGCCAGGGGACCAAGCTGAGAAACCAA
440	2G6_LC#1	artificial	aa	DVVMTQSPLSLPVTLGQPASISCRSSQSLVYSDGNTYLNWFQQRPGQSPRRLIYQVSNWDSGVPDRFSGSGSGTDFTLKISRVEAE DVGIYYCMQDTLWPCSFGQGTKLEIKR

SFO	DESIGNATION	SOURCE	TVPF	SFOILENCE
∑ 0 N O O O			I : :	
441	2H12	artificial	nt	17) 🗀 1
				CTGAGCGATTCTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGCCTATGGATGAGGCTGACTATTAC TGTCAGGCGTGGGACAGCAGCACTGTGGTATTCGGCGGAGGGACCAAGCTGACCGTCCtAGGT
442	2H12	artificial	aa	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQPMDEADYY CQAWDSSTVVFGGGTKLTVLG
443	2G6	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
				TGTCAGGCGTGGGACAGCACCACTGTGGTATTCGGCGGAGGGACCAAGCTGACCGTCCTAGGT
444	2G6	artificial	aa	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
				CQAWDSSTVVFGGGTKLTVLG
445	23A10	artificial	nt	TCCTATGAGCTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
				GAAATATGTTTGCTGGTATCAGCAGAAGCCAGGCCAGTCCCCTATACTGGTCATCTATCAAGATAATAAGTGGCCCTCAGGGATCC
				CIGAGCGATICICICGGCTCCAACTCTGGGAACACACCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGATTATTAC
1				TGTCAGGCGTGGGACAGCACTGTGGTATTCGGCGGGGACCAAGCTGACCGTCCTAGGT
5 446	23A10	artificial	aa	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVCWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
).				CQAWDSSTVVFGGGTKLTVLG
447	5E3	artificial	nt	TCCTATGAGCTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
				TGAATATGCTTGCTGGTATCAGCAGAAGCCAGGCCAGTCCCCTGTGCTGGTCATCTATCAAGATAGCAAGCGGCCCTCAGGGATCC
				CTGAGCGATTCTCTGGCTCCCAACTCTGGGAACACAGCCACTCTGACCATCAGCGGGGGCCCAGGCTATGGGATGAGGCTGACTATTAC
				TGTCAGGCGTGGGACAGCACTGTGGTATTCGGCGGAGGACCAAGCTGACCGTCCTAGGT
448	5E3	artificial	aa	SYELTQPPSVSVSPGQTASITCSGDKLGDEYACWYQQKPGQSPVLVIYQDSKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
				CQAWDSSTVVFGGGTKLTVLG

TABLE IIc: Heavy Chain Variable Region Polynucleotide and Amino acid Sequences 13586 HC [hu anti-<huCDH19> 4F3 VH]

QVQLVESGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS

5 SEQ ID NO: 449

13589 HC [hu anti-<huCDH19> 4A9 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSS

10 SEQ ID NO: 450

13590 HC [hu anti-<huCDH19>4B10 VH]

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS

15 SEQ ID NO: 451

13874 HC [hu anti-<huCDH19> 17H8.2 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSS

20 SEQ ID NO: 452

13875 HC [hu anti-<huCDH19> 16C1.1 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLTLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSS

25 SEQ ID NO: 453

13876 HC [hu anti-<huCDH19> 16A4.1 VH]

QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISV 30 DTSKNQFSLKLSSVTAADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTTVTVSS SEQ ID NO: 454

13877 HC [hu anti-<huCDH19> 22G10.1 VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR 55 FTISSDNSKSTLYLQMNSLRAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 455

13878 HC [hu anti-<huCDH19> 20D3.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV 40 TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 456

13879 HC [hu anti-<huCDH19> 22D1.1 VH]

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV
45 TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
SEQ ID NO: 457

13880 HC [hu anti-<huCDH19> 25F8.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 458

13881 HC [hu anti-<huCDH19> 26F12.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG 55 RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 459

13882 HC [hu anti-<huCDH19> 26D1.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEO ID NO: 460

13883 HC [hu anti-<huCDH19> 25G10.1 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSS

5 SEQ ID NO: 461

13885 HC [hu anti-<huCDH19> 19B5.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS

10 SEQ ID NO: 462

14022 HC [hu anti-<huCDH19> 4A2 VH]

QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS

15 SEO ID NO: 463

14024 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS

20 SEQ ID NO: 464

14025 HC [hu anti-<huCDH19> 4A2 VH]

QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS

25 SEQ ID NO: 465

14026 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS

30 SEQ ID NO: 466

14027 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P,D111E) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSSVTAADTAVYYCAREGSSGWYFQYWGQGTLVTVSS

35 SEQ ID NO: 467

14028 HC [hu anti-<huCDH19>4A2 (1-472)(Q17E,H47P,D111E,W134Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCAREGSSGYYFQYWGQGTLVTVSS

40 SEQ ID NO: 468

14029 HC [hu anti-<huCDH19> 4A2 VH]

 $QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV\\TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS$

45 SEQ ID NO: 469

14030 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS

50 SEQ ID NO: 470

14031 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS

55 SEQ ID NO: 471

14032 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,R141Q) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGQGTLVTVSS

60 SEQ ID NO: 472

14033 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,D61E,D72E,R141Q) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGQGTLVTVSS SEQ ID NO: 473

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14034 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,D61E,D72E,W134Y,R141Q) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGYYFDLWGQGTLVTVSS SEO ID NO: 474

10

14039 HC [hu anti-<huCDH19>2G6 (1-477)(R17G,D61E,D72E,K94N) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKD RFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS SEO ID NO: 475

15

14040 HC [hu anti-<huCDH19> 16C1.1 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLTLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSS SEQ ID NO: 476

20

14041 HC [hu anti-<huCDH19> 16C1.1 (1-469)(T92K) VH]

 $QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS\\ IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSS$

25 SEQ ID NO: 477

14042 HC [hu anti-<huCDH19>16C1.1 (1-469)(T92K,D109E) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCAREGSSGWYRWFDPWGQGTLVTVSS

30 SEQ ID NO: 478

14043 HC [hu anti-<huCDH19>16C1.1 (1-469)(T92K,W132Y,W135Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGYYRYFDPWGQGTLVTVSS

35 SEQ ID NO: 479

14044 HC [hu anti-<huCDH19> 16C1.1 (1-469)(T92K) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSS

40 SEQ ID NO: 480

14045 HC [hu anti-<huCDH19> 17H8.2 VH]

 $QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV\\DTSKNQFSLKLSSVTAADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSS$

45 SEQ ID NO: 481

14046 HC [hu anti-<huCDH19> 17H8.2 (1-471)(D109E) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSS

50 SEQ ID NO: 482

14047 HC [hu anti-<huCDH19>17H8.2 (1-471)(D109E,W132Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGYYDAFDIWGQGTMVTVSS

55 SEQ ID NO: 483

14048 HC [hu anti-<huCDH19> 17H8.2 (1-471)(D109E) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSS

60 SEQ ID NO: 484

14049 HC [hu anti-<huCDH19>4F7 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDYWGQGTLVTVSS SEO ID NO: 485

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14050 HC [hu anti-<huCDH19> 4F7 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDYWGQGTLVTVSS SEO ID NO: 486

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14051 HC [hu anti-<huCDH19> 4F7 (1-468)(W113Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNYAFHFDYWGQGTLVTVSS SEO ID NO: 487

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14052 HC [hu anti-<huCDH19> 4B10 (1-471)(R17G,D61E,D72E,W134Y) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDYSFDYWGQGTLVSVSS SEQ ID NO: 488

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14053 HC [hu anti-<huCDH19> 4B10 VH]

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS SEO ID NO: 489

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14054 HC [hu anti-<huCDH19>4B10 (1-471)(R17G) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKG RFTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS SEQ ID NO: 490

30

14055 HC [hu anti-<huCDH19>4B10 (1-471)(R17G,D61E,D72E) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS SEQ ID NO: 491

35

14056 HC [hu anti-<huCDH19> 4A9 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSS SEQ ID NO: 492

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14057 HC [hu anti-<huCDH19> 4A9 (1-468)(F55I,A56G) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSS SEQ ID NO: 493

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14058 HC [hu anti-<huCDH19>4A9 (1-468)(F55I,A56G) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSS SEQ ID NO: 494

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14059 HC [hu anti-<huCDH19>4A9 (1-468)(F55I,A56G,W113Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNYAFHFDFWGQGTLVTVSS SEQ ID NO: 495

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14060 HC [hu anti-<huCDH19> 20D3.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 496

60

14061 HC [hu anti-<huCDH19> 20D3.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEO ID NO: 497

5 14062 HC [hu anti-<huCDH19> 20D3.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEO ID NO: 498

10 14063 HC [hu anti-<huCDH19> 20D3.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEO ID NO: 499

15 <u>14064 HC [hu anti-<huCDH19> 20D3.1 (1-469)(W133Y) VH]</u>

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEQ ID NO: 500

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14065 HC [hu anti-<huCDH19> 22G10.1 (1-470)(S82R,A99E) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISRDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS SEO ID NO: 501

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14066 HC [hu anti-<huCDH19>22G10.1 (1-470)(A99E,H105Y) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 502

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14067 HC [hu anti-<huCDH19> 22G10.1 (1-470)(A99E) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 503

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14068 HC [hu anti-<huCDH19> 22G10.1 (1-470)(A99E) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 504

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14069 HC [hu anti-<huCDH19> 22G10.1 (1-470)(D72E,A99E) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRF TISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 505

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14070 HC [hu anti-<huCDH19> 22G10.1 (1-470)(H105Y) VH]

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAADTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSS SEQ ID NO: 506

50

14071 HC [hu anti-<huCDH19> 16A4.1 (1-474)(T144L) VH]

QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSS SEQ ID NO: 507

55

14072 HC [hu anti-<huCDH19>19B5.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEQ ID NO: 508

60

14073 HC [hu anti-<huCDH19> 19B5.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSS SEO ID NO: 509

5 14074 HC [hu anti-<huCDH19>19B5.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEQ ID NO: 510

10 14075 HC [hu anti-<huCDH19> 19B5.1 VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEQ ID NO: 511

15 14076 HC [hu anti-<huCDH19> 19B5.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSS SEQ ID NO: 512

20 <u>14077 HC [hu anti-<huCDH19> 23A10.3 (1-474)(L92Q) VH]</u>

QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGR FTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS SEQ ID NO: 513

25 <u>14078 HC [hu anti-<huCDH19> 23A10.3 (1-474)(R17G,L92Q) VH]</u>

QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKG RFTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS SEQ ID NO: 514

30 <u>14079 HC [hu anti-<huCDH19> 23A10.3 (1-474)(R17G,D61E,D72E,L92Q) VH]</u>

QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGR FTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS SEQ ID NO: 515

35 <u>14080 HC [hu anti-<huCDH19> 23A10.3 VH]</u>

QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGR FTISRDNSKNTLYLLMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS SEQ ID NO: 516

40 14081 HC [hu anti-<huCDH19> 25G10.1 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSS SEQ ID NO: 517

45 14082 HC [hu anti-<huCDH19>25G10.1 (1-469)(D109E,W132Y,W135Y) VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCAREGSSGYYRYFDPWGQGTLVTVSS SEQ ID NO: 518

50 <u>14083 HC [hu anti-<huCDH19> 26D1.1 VH]</u>

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEQ ID NO: 519

55 14084 HC [hu anti-<huCDH19> 26D1.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEQ ID NO: 520

60 14085 HC [hu anti-<huCDH19> 26D1.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEO ID NO: 521

5 14086 HC [hu anti-<huCDH19> 26D1.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEO ID NO: 522

10 14087 HC [hu anti-<huCDH19> 26D1.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLYLHFDYWGQGTLVTVSS SEQ ID NO: 523

15 14088 HC [hu anti-<huCDH19> 26D1.1 (1-469)(R27G,G82R) VH]

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS SEQ ID NO: 524

20 14089 HC [hu anti-<huCDH19> 26F12.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 525

25 14090 HC [hu anti-<huCDH19> 26F12.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 526

30 <u>14091 HC [hu anti-<huCDH19> 26F12.1 (1-469)(W133Y) VH]</u>

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEQ ID NO: 527

35 <u>14092 HC [hu anti-<huCDH19> 26F12.1 (1-469)(W133Y) VH]</u>

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEQ ID NO: 528

40 14093 HC [hu anti-<huCDH19> 25F8.1 VH]

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 529

45 <u>14094 HC [hu anti-<huCDH19> 25F8.1 VH]</u>

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 530

50 <u>14095 HC [hu anti-<huCDH19> 25F8.1 (1-469)(F90Y) VH]</u>

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 531

55 14096 HC [hu anti-<huCDH19> 25F8.1 (1-469)(F90Y) VH]

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS SEQ ID NO: 532

60 14097 HC [hu anti-<huCDH19>25F8.1 (1-469)(F90Y,W133Y) VH]

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSS SEO ID NO: 533

5 14098 HC [hu anti-<huCDH19> 22D1.1 VH]

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEO ID NO: 534

10 14099 HC [hu anti-<huCDH19> 22D1.1 VH]

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEQ ID NO: 535

15 14100 HC [hu anti-<huCDH19> 22D1.1 (1-469)(W133Y) VH]

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSS SEQ ID NO: 536

20 <u>14101 HC [hu anti-<huCDH19> 22D1.1 (1-469)(W133Y) VH]</u>

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSS SEQ ID NO: 537

25 <u>14102 HC [hu anti-<huCDH19> 22D1.1 (1-469)(F90Y) VH]</u>

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS SEQ ID NO: 538

30 13591 HC [hu anti-<huCDH19> 4F7 VH]

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDYWGQGTLVTVSS SEQ ID NO: 539

35 14301 HC [hu anti-<huCDH19> 2G6 VH]

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS SEQ ID NO: 540

40 14302 HC [hu anti-<huCDH19> 2G6 (1-477)(R17G,K94N) VH]

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS SEQ ID NO: 541

45 14303 HC [hu anti-<huCDH19> 2G6 (1-477)(D61E,D72E) VH]

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS SEQ ID NO: 542

50 <u>14304 HC [hu anti-<huCDH19> 2G6 (1-477)(R17G) VH]</u>

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS SEQ ID NO: 543

55 TABLE IId: Light Chain Variable Region Amino acid Sequences

13586 LC [hu anti-<huCDH19> 4F3 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEQ ID NO: 544

13589 LC [hu anti-<huCDH19> 4A9 VL]

60

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQFPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLG SEQ ID NO: 545

5 13590 LC [hu anti-<huCDH19> 4B10 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FALTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKR SEQ ID NO: 546

10 13874 LC [hu anti-<huCDH19> 17H8.2 VL]

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKG SEQ ID NO: 547

15 13875 LC [hu anti-<huCDH19> 16C1.1 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 548

20 <u>13876 LC [hu anti-<huCDH19>16A4.1 VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSPFTFGGGTKVEIKR SEQ ID NO: 549

25 <u>13877 LC [hu anti-<huCDH19> 22G10.1 VL]</u>

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 552

30 13878 LC [hu anti-<huCDH19> 20D3.1 VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 554

35 13879 LC [hu anti-<huCDH19> 22D1.1 VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 555

40 <u>13880 LC [hu anti-<huCDH19> 25F8.1 VL]</u>

45

50

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QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 556

13881 LC [hu anti-<huCDH19> 26F12.1 VL]

QSVLTQSPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 557

13882 LC [hu anti-<huCDH19> 26D1.1 VL]

HSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 555

13883 LC [hu anti-<huCDH19> 25G10.1 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 556

13885 LC [hu anti-<huCDH19> 19B5.1 VL]

QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 557

5 14022 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q) VL]

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFTVYYCQQYGSSFTFGPGTKVDIKR SEQ ID NO: 558

10 14024 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKR SEQ ID NO: 559

15 14025 LC [hu anti-<huCDH19>4A2 (1-236)(N30Q,T102A) VL]

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKR SEQ ID NO: 560

20 <u>14026 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKR SEQ ID NO: 561

25 <u>14027 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKR SEQ ID NO: 562

30 <u>14028 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKR SEQ ID NO: 563

35 <u>14029 LC [hu anti-<huCDH19> 4A2 (1-236)(R29Q,N30S) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFTVYYCQQYGSSFTFGPGTKVDIKR SEQ ID NO: 564

40 <u>14030 LC [hu anti-<huCDH19> 4F3 VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEQ ID NO: 565

45 14031 LC [hu anti-<huCDH19> 4F3 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEQ ID NO: 566

50 <u>14032 LC [hu anti-<huCDH19> 4F3 VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEQ ID NO: 567

55 14033 LC [hu anti-<huCDH19> 4F3 VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEQ ID NO: 568

60 <u>14034 LC [hu anti-<huCDH19> 4F3 VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKR SEO ID NO: 569

5 14039 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLG SEQ ID NO: 570

10 14040 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 571

15 14041 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 572

20 <u>14042 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 573

25 <u>14043 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 574

30 <u>14044 LC [hu anti-<huCDH19> 16C1.1 (1-235)(G95R,H105Y,G141Q) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGNSPLTFGQGTKVEIKR SEQ ID NO: 575

35 <u>14045 LC [hu anti-<huCDH19> 17H8.2 (1-235)(G149R) VL]</u>

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKR SEQ ID NO: 576

40 <u>14046 LC [hu anti-<huCDH19> 17H8.2 (1-235)(G149R) VL]</u>

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKR SEQ ID NO: 577

45 <u>14047 LC [hu anti-<huCDH19> 17H8.2 (1-235)(G149R) VL]</u>

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKR SEQ ID NO: 578

50 <u>14048 LC [hu anti-<huCDH19> 17H8.2 (1-235)(S57Y,G149R) VL]</u>

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKR SEQ ID NO: 579

55 14049 LC [hu anti-<huCDH19>4F7 (1-239)(H57Y) VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLG SEQ ID NO: 580

60 14050 LC [hu anti-<huCDH19>4F7 (1-239)(H57Y,D110E) VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESSLSGWVFGGGTRLTVLG SEQ ID NO: 581

5 14051 LC [hu anti-<huCDH19>4F7 (1-239)(D110E) VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESSLSGWVFGGGTRLTVLG SEQ ID NO: 582

10 14052 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKR SEQ ID NO: 583

15 14053 LC [hu anti-<huCDH19>4B10 (1-236)(H45Q,A90T) VL]

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKR SEQ ID NO: 584

20 <u>14054 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKR SEQ ID NO: 585

25 <u>14055 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKR SEQ ID NO: 586

30 <u>14056 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L) VL]</u>

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLG SEQ ID NO: 587

35 14057 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L) VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLG SEQ ID NO: 588

40 14058 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L,D110E) VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESRLSGWVFGGGTKLTVLG SEQ ID NO: 589

45 <u>14059 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L,D110E) VL]</u>

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESRLSGWVFGGGTKLTVLG SEQ ID NO: 590

50 <u>14060 LC [hu anti-<huCDH19> 20D3.1 (1-235)(S102A) VL]</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 591

55 14061 LC [hu anti-<huCDH19> 20D3.1 (1-235)(K45Q,S102A) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 592

60 14062 LC [hu anti-<huCDH19> 20D3.1 (1-235)(K45Q,S102A) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 593

5 14063 LC [hu anti-<huCDH19>20D3.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDESLQGWVFGGGTKLTVLG SEQ ID NO: 594

10 14064 LC [hu anti-<huCDH19> 20D3.1 (1-235)(W109Y) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATYDDSLNGWVFGGGTKLTVLG SEQ ID NO: 595

15 14065 LC [hu anti-<huCDH19> 22G10.1 VL]

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 596

20 14066 LC [hu anti-<huCDH19> 22G10.1 VL]

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 597

25 <u>14067 LC [hu anti-<huCDH19> 22G10.1 (1-234)(Q97E,S98P) VL]</u>

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 598

30 <u>14068 LC [hu anti-<huCDH19> 22G10.1 (1-234)(V78F,Q97E,S98P) VL]</u>

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEF TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 599

35 <u>14069 LC [hu anti-<huCDH19> 22G10.1 (1-234)(V78F,Q97E,S98P) VL]</u>

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEF TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 600

40 14070 LC [hu anti-<huCDH19> 22G10.1 VL]

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKR SEQ ID NO: 601

45 <u>14071 LC [hu anti-<huCDH19> 16A4.1 (1-235)(G141Q) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSPFTFGQGTKVEIKR SEQ ID NO: 602

50 <u>14072 LC [hu anti-<huCDH19> 19B5.1 (1-235)(K45Q,S102A) VL]</u>

QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 603

55 14073 LC [hu anti-<huCDH19> 19B5.1 (1-235)(K45Q,S102A) VL]

QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 604

60 <u>14074 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A) VL]</u>

QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLG SEO ID NO: $605\,$

5 14075 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A,D111E,N135Q) VL]

QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLG SEQ ID NO: 606

10 14076 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A,W109Y,D111E,N135Q) VL]

QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLG SEQ ID NO: 607

15 <u>14077 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLG SEQ ID NO: 608

20 <u>14078 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLG SEQ ID NO: 609

25 <u>14079 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S,D110E) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLG SEQ ID NO: 610

30 14080 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42Y) VL]

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVYWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLG SEQ ID NO: 611

35 <u>14081 LC [hu anti-<huCDH19> 25G10.1 (1-235)(H105Y) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 612

40 <u>14082 LC [hu anti-<huCDH19> 25G10.1 (1-235)(H105Y) VL]</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKR SEQ ID NO: 613

45 14083 LC [hu anti-<huCDH19> 26D1.1 (1-235)(S7P) VL]

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 $HSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS\\ ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG\\ SEQ ID NO: 614$

14084 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P) VL]

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 615

14085 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P,W109Y) VL]

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDDSLNGWVFGGGTKLTVLG SEQ ID NO: 616

14086 LC [hu anti-<huCDH19>26D1.1 (1-235)(H1Q,S7P,W109Y,D111E,N135Q) VL]

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLG SEQ ID NO: 617

5 14087 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P,W109Y,D111E,N135Q) VL]

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLG SEQ ID NO: $618\,$

10 14088 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P) VL]

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 619

15 14089 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P) VL]

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 620

20 <u>14090 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,D111E) VL]</u>

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLG SEQ ID NO: 621

25 <u>14091 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,D111E) VL]</u>

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLG SEQ ID NO: 622

30 <u>14092 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,W109Y,D111E,N135Q) VL]</u>

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLG SEQ ID NO: 623

35 14093 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 624

40 <u>14094 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A) VL]</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 625

45 14095 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDDSLNGWVFGGGTKLTVLG SEQ ID NO: 626

50 <u>14096 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A,D111E) VL]</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDESLNGWVFGGGTKLTVLG SEQ ID NO: 627

55 14097 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDESLQGWVFGGGTKLTVLG SEQ ID NO: 628

60 <u>14098 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A) VL]</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 629

5 14099 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLG SEQ ID NO: 630

10 14100 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,W109Y,D111E,N135Q) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLG SEQ ID NO: 631

15 14101 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,W109Y) VL]

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATYDDSMNGWVFGGGTKLTVLG SEQ ID NO: 632

20 <u>14102 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A) VL]</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLG SEQ ID NO: 633

25 13591 LC [hu anti-<huCDH19> 4F7 VL]

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLG SEQ ID NO: 634

30 <u>14301 LC [hu anti-<huCDH19> 2G6 (1-234)(D110E) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLG SEQ ID NO: 635

35 14302 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLG SEQ ID NO: 636

40 <u>14303 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLG SEQ ID NO: 637

45 <u>14304 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLG SEQ ID NO: 638

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Anti-CDH19 Variable and Constant Region Polynucleotide and Amino Acid Sequences

TABLE IIIa: Heavy Chain Variable and Contant Region Polynucleotide and Amino acid Sequences

2G6

CAGGTGCAGTTGGTGGAGTCTGGGGGGGGGGGGGGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTTCAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGG

60 CTGGAGTGGCATTTATATGGTATGGAAGTAATAAATACTATGCAGACTCCGTGAAGGAC CGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAAAAGCCTGAGAGCT

GAGGACACGGCTGTGTATTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACTAC
GGTATGGACGTCTGGGGCCAAGGGACCACGGTCACCGTCTCTAGTGCCTCCACCAAGGGCCCATCG
GTCTTCCCCCTGGCACCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTC
AAGGACTACTTCCCCGAACCGGTGACGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCAC
ACCTTCCCGGCTGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCA
GCAGCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCACCAAGACCCAAGGTGGAC
CTGGGGGGACCGTCAGTCTTCCTCTTCCCCCAAAACCCAAGGACACCTCATGATCTCCCGGACC
CCTGAGGTCACATGCGTGGTGGACGTGGACCATACAACCCAAGGTCACAACTCACACGTGCCTCCAACGCACGTACCAAGTTCAACTGGTAC
CTGGACGCCTGCAGGTGCATAAATGCCAAGAACCAAGACCAAGGACCACTACAAACACCAACGT

- 10 GTGGACGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGT ACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCA AGGTCTCCAACAAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCAGGTCAGCCT GACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGC
- 15 CGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCA AGCTCACCGTGGACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAG GCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 639
- 20 QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE
- 25 KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 640

<u>4A2</u>

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- 35 GGCCAGGGCACCCTGGTCACCGTCTCTAGTGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCA CCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCC GAACCGGTGACGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGGGTGCACACCTTCCCGGCTGTC CTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACC CAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCC
- 40 CAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGTC AGTCTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATGC GTGGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGA GGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAACAGCACGTACCGTGTGGTCAGCG TCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAA
- 50 ACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 641
 - QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKST
- 55 SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 642

4A9

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CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGC ACTGTCTCTGGTGGCTCCATCAGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGAAAGGGA GTCACCTTATCAGTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGCTCTGTGACCGCTGCG GACACGGCCGTGTATTACTGTGCGAGGAACTGGGCCTTCCACTTTGACTTCTGGGGCCAGGGAACC GCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGG GACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCCAGACCTACATCT GCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCCAAATCTTGTGAC AAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGTCAGTCTTCCTCTTCCCCCAAAACCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGAC $\tt GTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGCGTGGAGGTGCATAATGC$ CAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCC TGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAGCC CCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGTGTACACCCTGCC

- CCCATCCGGGAGGAGTGACCAAGAACCAGGTCAGCCTGACCTGCTGAAAGGCTTCTATCC CAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTC CCGTGCTGGACTCCGACGGCTCCTTCTTCTCTATAGCAAGCTCACCGTGGACAAGAGCAGGTGGC AGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGCAGAAGA
- 25 QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSN TKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPR EPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVD

KSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 644

4B10

- 35 CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGT
 GCAGCCTCTGGATTCACCTTCAGTAGCTATGACATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGG
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 CGATTCACCATCTCCAGAGACACTTCCAAGAACACGCTGTATTTTGCAAATGAACAGCCTGAGAGCT
 GAGGACACGGCTGTATATTACTGTGCGAGAGAACAGATATTTTGACTGGTCTTTTTGACTACTGGGGC
- 40 CAGGGAACCCTGGTCAGCGTCTCTAGTGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCC TCCTCCAAGAGCACCTCTGGGGCACAGCGGCCCTGGCTGCCTGGTCAAGGACTACTTCCCCGAA CCGGTGACGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGGGTGCACACCTTCCCGGCTGTCCTA CAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCATTGAGCCCAG ACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCCAA
- 45 ATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGGACCGTCAGT CTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTG GTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGAGGT GCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAACAGCACGTACCGTGTGGTCAGCGTCC TCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCC
- 55 ACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 645
- QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV

KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 646

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4F3 CAGGTGCAGCTGGTGGAGTCTGGGGGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGT GCAGCGTCTGGATTCTCCTTCAGTAGCTATGACATGGACTGGGTCCGCCAGACTCCAGGCAAGGGG ${\tt CGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTTTCTGCAAATGAACAGCCTGAGAGTC}$ GAGGACACGGCTGTGTATTACTGTGCGAGAGAAACTGGGGAGGGCTGGTACTTCGATCTCTGGGGC CGTGGCACCCTGGTCACCGTCTCTAGTGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCT CCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAAC CGGTGACGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTCCTAC AGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCCAGA CCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCCAAA TCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGTCAGTC TTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTG GCATAATGCCAAGACAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCGTCC TCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCC CTCCCAGCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGTGTA GCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTACAAG

25 ACCACGCCTCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGACAAG AGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTAC ACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 647

30 QVQLVESGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 648

<u>4F7</u>

- 40 CAGGTGCAGCTGCAGGACTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGC ACTGTCTCTGGTGGCTCCATCAGTAGTTACTCCTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGA GTCACCATATCATTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGCTCTGTGACCGCTGCG GACACGGCCGTGTATTACTGTGCGAGGAACTGGGCCTTCCACTTTGACTACTGGGGCCAGGGAACC 45 CTGGTCACCGTCTCTAGTGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCTCCTCCAAGA GCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGG TGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTCCTACAGTCCTCAG GACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCCAGACCTACATCT GCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCCAAATCTTGTGAC 50 AAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGTCAGTCTTCCTCTTCCCCCAAAACCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGAC GTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGCGTGGAGGTGCATAATGC CAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCC TGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAGCC
- 60 GCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 649

PCT/EP2014/051550 WO 2014/114800

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNOFSLKLSSVTAADTAVYYCARNWAFHFDYWGOGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLOSSGLYSLSSVVTVPSSSLGTOTYICNVNHKPSNT KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEOYNSTYRVVSVLTVLHODWLNGKEYKCKVSNKALPAPIEKTISKAKGOPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 650

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CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGCGAAGCCTTCGGAGACCCTGTCCCTCACCTGC ACTGTCTCTGGTGACTCCATCACTAGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGA 15 GTCACCATATCAGTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGTTCTGTGACCGCTGCG GACACGGCCGTGTATTACTGTGCGAGAGATCAAAGGCGGATAGCAGCAGCTGGTACCCACTTCTAC GGTATGGACGTCTGGGGCCAAGGGACCACGGTCACTGTCTCCTCAGCTTCCACCAAGGGCCCATCC GTCTTCCCCTGGCGCCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTC AAGGACTACTTCCCCGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCA20 ${\tt CACCTTCCCGGCTGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCC}$ AGCAGCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGA ${\tt CCCTGAGGTCACATGCGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTA}$ 25 CGTGGACGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGT AGGTCTCCAACAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCATCCCGGGAGGAGATGACCAAGAACCAGGTCAGCCT GACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGC

30 ${\tt CGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCA}$ AGCTCACCGTGGACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAG GCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 651

35 QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTTVTVSSASTKGPSVFPLAPS SKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICN VNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDP EVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK 40

AKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLY SKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 652

- 45 ACTGTCTCTGGTGGCTCCATCAGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGA GTCACCATGTCAATAGACACGTCCAAGAACCAGTTCTCCCTGACGCTGAGCTCTTTGACCGCTGCG GACACGCCGTGTATTTCTGTGCGAGAGATGGGAGCAGTGGCTGGTACCGGTGGTTCGACCCCTGG 50 GGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGCG ${\tt CCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCC}$ GAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTC ${\tt CTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACC}$ CAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCC 55 CAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGGACCGTC
- AGTCTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATGC GTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGA GGTGCATAATGCCAAGACAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCG
- 60 GCCTCCCAGCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCGAGAACCACAGGT

AGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTACA AGACCACGCCTCCCGTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGAGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACT ACACGCAGAAGAGCCTCTCCCTGTCTCCCGGGTAAATGA

5 SEQ ID NO: 653

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLTLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 654

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17H8 CAGGTGCAGCTGCAGGACTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACGTGC ACTGTCTCTGGTGGCTCCATCAATAGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGA 20 GTCACCATATCAGTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGCTCTGTGACCGCTGCG GACACGGCCCTGTATTACTGTGCGAGAGATTCCCGGTATAGAAGTGGCTGGTACGATGCTTTTGAT ATCTGGGGCCAAGGGACAATGGTCACCGTCTTCTAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGCGCCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTAC TTCCCGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCG25 GCTGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGG GCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTT GAGCCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGA CCGTCAGTCTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCA ${\tt CATGCGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGC}$

35 CTACAAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCTCTATAGCAAGCTCACCGTG GACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAA CCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 655

40 QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK

45 GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 656

19B5

- 50 CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGC
 AAGGTTTCTGGATACACCTTCACCAGCTACTTTATTCACTGGGTGCGCCAGGCCCCTGGACAAGGG
 CTTGAATGGATGGGAATTATCAACCCTATTAGTGTTAGCACAAGCTACGCACAGAAGTTCCAGGGC
 AGAGTCACCATGACCAGGGACACGTCCCACGAGCACAGTCTTCATGGAGCTGAGCAGCCTGAGATC
 TGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGGGATACAGCTATGGTTACATTTGGACTACTG

 55 GGCCAGGGAACCCTGGTCACCGTCTCCCCAGAGGGCCCATCCGTCTTCCCCCTGGC
- GCCTCCTCAAGAGCACCTCTCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGC
 GCCCTCCTCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGCTGAAGGACTACTTCCC
 CGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT
 CCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCAC
 CCAGACCTACATCTGCAACACGTGAATCACAAGCCCAGCAACACCAAGGTTGGACAAGAAAGTTGAGC
- 60 CCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGT CAGTCTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATG

PCT/EP2014/051550 WO 2014/114800

 ${\tt CGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGCGTGG}$ AGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGC GTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAA AGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGG AAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTAC AAGACCACGCCTCCCGTGCTGGACTCCGACGCCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGAC AAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCAC TACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA

10 SEQ ID NO: 657

> OVOLVOSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVROAPGOGLEWMGIINPISVSTSYAOKFOGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLOSSGLYSLSSVVTVPSSSLGTOTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHODWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 658

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CAGGTGCAGCTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGC AAGGTTTCTGGATACACCTTCACCAGCTACTTTATTCACTGGGTGCGCCAGGCCCCTGGACAAGGG ${\tt CTTGAGTGGATGGGAATAATCAACCCTATTAGTGTTAGCACAAGCTACGCACAGAAGTTCCAGGGC}$ 25 AGAGTCACCATGACCAGGGACACGTCCACGAGCACAGTCTTCATGGAGCTGAGCAGCCTGAGATC TGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGGATACAGCTATGGTTACATTTTGACTACTG GGGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGC GCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCC CGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT30 ${\tt CCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGC}$ ${\tt CCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGT}$ CAGTCTTCCTCTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATG CGTGGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGG 35 AGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGC GTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAA AGCCCTCCCAGCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGG AAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTAC

40 AAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGAC AAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCAC TACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA **SEQ ID NO: 659**

45 QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK 50 GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 660

55 CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAGGGTTTCCTGC AAGGTTTCTGGATACACCTTCACCAGCTACTTTATTCACTGGGTACGCCAGGCCCCTGGACAAGGG $\tt CTTGAGTGGATGGGAATAATCAACCCTATTAGTGTTAGCACAAGCTACGCACAGAAGTTCCAGGGC$ TGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGGATACAGCTATGGTTACATTTGGACTACTG 60 GGGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGC GCCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCC

CGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT
CCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCAC
CCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGC
CCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGT
CAGTCTTCCTCTTCCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACATG
CGTGGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGG
AGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAAGTGCAAGGTCTCCAACAA
AGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGGCAAAGGGCAGCCCCGAGAACCACAGG
TGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCAGGTCAGCCTGACCTGCTCA
AAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTAC
AAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGAC
AAGACCACGCCTCCCGTGGACTCTCCCATGCTCCGTGATGCACAACCAC
TACACGCAGAAGAGCCTCTCCCTGTCTCCCGGGTAAATGA

15 SEQ ID NO: 661

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QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 662

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22G10 GAGGTGCAACTGTTGGAGTCTGGGGGGAGGCTTGGTACAGCCTGGGGGGTCCCTGAGACTCTCCTGT GCAGCCTCTGGATTCACCTTTAGCAGTTATGCCATGAACTGGGTCCGCCAGGCTCCAGGGAAGGGG $\tt CTGGAGTGGGTCTCAACTATTAGTGGTGGTGGTGCTAACACATACTACGCAGACTCCGTGAAGGGC$ 30 GCGGACACGCCGTATATCACTGTGCGAAAGGGGGAATGGGGGGATACTACTACGGTATGGACGT $\tt CTGGGGCCAAGGGACCACGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCT$ GGCGCCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTT CCCGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGC 35 TGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGC ACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGA GCCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACC GTCAGTCTTCCCCCCAAAACCCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCACA TGCGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGCGT40 GGAGGTGCATAATGCCAAGACAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCA GCGTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACA AAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAG GTGTACACCCTGCCCCATCCCGGGAGGAGATGACCAAGAACCAGGTCAGCCTGACCTGGCTC AAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTA

45 CAAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCTCTCTATAGCAAGCTCACCGTGGAC AAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCAC TACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 663

- 50 EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SK AKGOPREPOVYTI PPSREEMTKNOVSI TCI VKGEVPSDIAVEWESNGOPENNVKTTPPVI DSDGSFE
- 55 SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 664

23A10

60 CAGGTGCAGCTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGT GCAGCGTCTGGATTCACCTTCAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAAGGGG

CGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCTAATGAACAGCCTGAGAGCC GAGGACTCGGCTGTTATTACTGTGCGAGAAGGGCCGGTATACCTGGAACTACGGGCTACTACTAT GGTATGGACGTCTGGGGCCAAGGGACCACGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCC GTCTTCCCCCTGGCGCCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTC AAGGACTACTTCCCCGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCA ${\tt CACCTTCCCGGCTGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCC}$ AGCAGCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGA CCCTGAGGTCACATGCGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTA CGTGGACGCGTGGAGGTGCATAATGCCAAGACAAGCCGCGGGAGGAGCAGTACAACAGCACGTAGGTCTCCAACAAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCAGGTCAGCCT GACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGC ${\tt CGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCA}$

20 **SEQ ID NO: 665**

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OVOLVESGGGVVOPGRSLRLSCAASGFTFSRYGIHWVROAPGKGLEWVAVIWYDGSNKYYADSVKGR FTISRDNSKNTLYLLMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGOGTTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLOSSGLYSLSSVVTVPSSSLGTOT

AGCTCACCGTGGACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAG

GCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA

25 YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 666

CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGC AAGGCATCTGGATACACCTTCACCAGCTACTATATTCACTGGGTGCGCCAGGCCCCTGGACAAGGA ${\tt CTTGAGTGGATGGGAATAATCAACCCCAGTGGTGGTAGCACAAGGTACGCACAGAAGTTCCAGGG}$ 35 CAGAGTCACCATGACCAGGGACACGTCCACGAGCACAGTCTTCATGGAGCTGAGCAGCCTGAGAT $\tt CTGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGAATACAGCTATGGTTACATTTTGACTACT$ GGGGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGG CGCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCC 40 CCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAG CCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCG TCAGTCTTCCTCTTCCCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACCCCTGAGGTCACAT GCGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTG

- 45 GAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAG CGTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAA AGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGG AAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTAC
- AAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGAC 50 AAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCAC TACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA **SEQ ID NO: 667**
- 55 QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK ${\tt STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN}$ HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK 60 GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

PCT/EP2014/051550 WO 2014/114800

SEQ ID NO: 668

25G10

CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGC 5 ACTGTCTCTGGTGGCTCCATCAGTGGTTACTACTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGA GTCACCATGTCAGTAGACACGTCCAAGAACCAGTTCTCCCTGAAGCTGAGCTCTGTGACCGCTGCG GACACGGCCGTGTATTACTGTGCGAGAGATGGGAGCAGTGGCTGGTACCGGTGGTTCGACCCCTGG GGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGCG CCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCC

- 10 GAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTC ${\tt CTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACC}$ CAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCC CAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGGACCGTC
- 15 AGTCTTCCTCTTCCCCCAAAACCCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCACATGC GTGGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGA GGTGCATAATGCCAAGACAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCG TCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAA GCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGT
- 20 AGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTACA AGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA
- 25 **SEQ ID NO: 669**

OVOLOESGPGLVKPSETLSLTCTVSGGSISGYYWSWIROPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG OPREPOVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG SEQ ID NO: 670

30

35

26D1 CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGT AAGGCATCTAGATACACCTTCACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGG ${\tt CTTGAGTGGATGGGAATAATCCACCCTAGTGGTGGTGACACAACCTACGCACAGAAGTTCCAGGGC}$ 40 AGAGTCACCATGACCGGGGACACGTCCACGAGCACAGTCTACATGGAGCTGAGCAGCCTGAGATC TGAGGACACGGCCGTGTATTACTGTGCGAGAGGGGGGATAAAACTATGGTTACATTTTGACTATTG GGGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGC GCCTCCTCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCC CGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT45 CCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCAC CCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGC AGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGC GTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAA

- 50 AGCCCTCCCAGCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGG AAGGCTTCTATCCCAGCGACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTAC
- 55 AAGACCACGCCTCCCGTGCTGGACTCCGACGGCTCCTTCTTCCTCTATAGCAAGCTCACCGTGGAC AAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCAC TACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGA SEQ ID NO: 671
- 60 QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS

KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 672

26F12

5

- CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGC

 10 AAGGCATCTAGATACACCTTCACCAACTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGG
 CTTGAGTGGATGGGATAATCAACCCTAGTGGTGGTGACTCAACCTACGCACAGAAGTTCCAGGGC
 AGACTCACCATGACCGGGACACGTCCACGAGCACAGTCTACATGGAGCTGAGCAGCCTGAGATC
 TGAGGACACGGCCGTGTATTACTGTGCGAGAGGGGGGATACAACTATGGTTACATTTTGACTACTG
 GGGCCAGGGAACCCTGGTCACCGTCTCCTCAGCTTCCACCAAGGGCCCATCCGTCTTCCCCCTGGC
- 15 GCCCTCCCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCC CGAACCGGTGACGGTGTCGTGGAACTCAGGGGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT CCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCAC CCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGC CCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCAGCACCTGAACTCCTGGGGGGACCGT
- 20 CAGTCTTCCTCTTCCCCCAAAACCCAAGGACACCTCATGATCTCCCGGACCCCTGAGGTCACATG CGTGGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGG AGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGC GTCCTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAA AGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGG
- 30 SEQ ID NO: 673

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV

35 NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 674

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TABLE IIIb: Light Chain Variable and Contant Region Polynucleotide and Amino acid Sequences

<u>2G6</u>

- 50 GGCCACCCACTGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTCCAAGCCAACAAGGCCACACT AGTGTGTCTGATCAGTGACTTCTACCCGGGAGCTGTGACAGTGGCCTGGAAGGCAGATGGCAGCCC CGTCAAGGCGGGAGTGGAGACCACCAAACCCTCCAAACAGGCAACAACAAGTACGCGGCCAGCA GCTACCTGAGCCTGACGCCCGAGCAGTGGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCAT GAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA
- 55 SEQ ID NO: 675

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVCLISDFY PGAVTVAWKADGSPVKAGVETTKPSKOSNNKYAASSYLSLTPEOWKSHRSYSCOVTHEGSTVEKTVA

60 PTECS SEQ ID NO: 676

4A2

GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCT
GCAGGGCCAGTCGGAATATTAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCT

CCCAGGCTCCTCATCTATGGTCCATCCAGCAGGGCCACTGGCATCCCAGACAGGTTCAGTGGCAGT
GGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCCTGAAGATTTTACAGTGTATTAC
TGTCAGCAGTATGGTAGCTCATTCACTTTCGGCCCTGGGACCAAAGTGGATATCAAACGTACGGTG
GCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGAACTGCCTCTGTTG
TGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCCCTCC

- 10 AATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCAAGGACAGCACCTACAGCCTCAGC AGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCACCCA TCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA SEO ID NO: 677
- 15 EIVLTQSPGTLSLSPGERATLSCRASRNISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFTVYYCQQYGSSFTFGPGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC SEQ ID NO: 678

440

20

CAGTCTGTGCTGACGCAGCCGCCCTCAGTGTCTGGGGCCCCAGGACAGAGGGTCACCATCTCCTGCACTGGGAGCAGCTCCAACATCGGGACAGGTTATGCTGTACACTGGTACCAGCAGTTTCCAGGAACAGCCCCCAAACTCCTCATCTATGGTAACAACAATCGGCCCTCAGGGGTTCCTGACCGATTCTCTGGCT

- 25 CCAAGTCTGGCACCTCAGCCTCCCTGGCCATCACTGGGCTCAGGCTGAGGATGAGGCTGATTATT ACTGCCAGTCCTATGACAGCAGACTGAGTGGTTGGGTGTTCGGCGGAGGGACCAAGCCGACCGTCC TAGGTCAGCCCAAGCCCAACCCCACTGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTCCAAGCCAACCAGGCCACACTAGTGTCTCTGATCAGTGACTTCTACCCGGGAGCTGTGACAGTGGCCTGGAAGGCAGATGGCAGCCCCGTCAAGGCGGAGTGGAGACCACCAAACCCTCCAAACAGAGCAACAACAAG
- 30 TACGCGGCCAGCAGCTACCTGAGCCTGACGCCCGAGCAGTGGAAGTCCCACAGAAGCTACAGCTG CCAGGTCACGCATGAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA SEQ ID NO: 679
- QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQFPGTAPKLLIYGNNNRPSGVPDRFSGSKSG
 TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC
 LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
 EKTVAPTECS
 SEQ ID NO: 680
- 40 4B10

- 50 TCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA SEQ ID NO: 681
 - $EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD\\FALTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP$
- 55 REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC SEQ ID NO: 682

4F3

60 GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCT GCAGGGCCAGTCAGAGTGTTAGCAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCT

CCCAGGCTCCTCATCTATGGTGCATCCAGCAGGGCCACTGGCATCCCAGACAGGTTCAGTGGCAGT
GGGTCTGGGACAGACTTCACTCTCACCATCAGCAGGACTGGAACCTGAGGATTTTGCAGTGTATTAC
TGTCAGCAGTATGGTAGCTCGTGGACGTTCGGCCAAGGGACCAAGGTGGAAATCAAACGTACGGT
GGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGAACTGCCTCTGTT
GTGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCCCTC
CAATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCACCTACAGCCTCAG
CAGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCACCC
ATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA
SEQ ID NO: 683

10

5

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

15 SEQ ID NO: 684

4F7

ZAGTCTGTGCCGACGCAGCCGCCCTCAGTGTCTGGGGCCCCAGGGCAGAGGGTCACCATCTCCTGC
ACTGGGAGCAGCTCCAATATCGGGACAGGTTATGATGTACACTGGTATCAGCAGCTTCCAGGAACA

GCCCCCAAACTCCTCATCCATGGTAACAGCAATCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGC
TCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCACTGGGCTCCAGGCTGAGGATGAGGCTGATTAT
TACTGCCAGTCCTATGACAGCAGTCTGAGTGGTTTCGGCGGAGGGACCAGGTTGACCGTC
CTAGGTCAGCCCAAGGCCAACCCCACTGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTCCAAGCC
AACAAGGCCACACTAGTGTCTCTGATCAGTGACTTCTACCCGGGAGCTGTACAGTGGCCTGGAAG

GCAGATGGCAGCCCCGTCAAGGCGGGAGTGGAGCCCCAAACCCTCCAAACAGAGCAACAACAA

GCAGATGGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCAAACCCTCCAAACAGAGCAACAACAA GTACGCGGCCAGCAGCTACCTGAGCCTGACGCCCGAGCAGTGGAAGTCCCACAGAAGCTACAGCT GCCAGGTCACGCATGAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA SEQ ID NO: 685

30 QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS SEQ ID NO: 686

35

16A4

GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCT GCAGGGCCAGTCAGAGTGTTAGCAGCAGTTATTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCTC CCAGGCTCCTCATCTATGGTACATCCAGCAGGGCCACTGGCATCCCAGACAGGTTCAGTGGCAGTG

- 45 GCAGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCACC CATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA SEQ ID NO: 687
- EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSPFTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC SEQ ID NO: 688

55 16C1

60

TTGTGTGCCTGCAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCCC TCCAATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCAGCAGCACCTACAGCCTC AGCAGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCAC CCATCAGGGCCTGAGCTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA

5 SEQ ID NO: 689

 $EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD\\FTLTISGLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP\\REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF$

NRGEC SEQ ID NO: 690

17H8

GACATTGTATTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCT

GCAGGGCCAGTCAGAGTGTTGCCGGCAGCTACCTAGCCTGGTACCAGCAGAAACCTGGCCAGGCT
CCCAGGCTCCTCATCTCTGGTGCATCCAGCAGGGCCACTGGCATCCCAGACAGGTTCAGTGGCAGT
GGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCCTGAAGATTTTGCAGTGTATTAC
TGTCAGCAGTATGGTAAATCACCGATCACCTTCGGCCAAGGGACACGACTGGAGATGAAAGGAAC
TGTGGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGTACCGCCTCT

20 GTTGTGTGCCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCC CTCCAATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCAAGGACAGCACCTACAGCCT CAGCAGCACCCTGACGCTGAGCAAAGCAGACACACAAAGTCTACGCCTGCGAAGTCA CCCATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA SEQ ID NO: 691

25

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKGTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC

30 SEQ ID NO: 692

<u>19B5</u>

CAGTCTGCGCTGACTCAGCCACCCTCAACGACTGGGACCCCCGGGCAGAGGGTCACCATCTCTTGTTCTGGAAGCAGGTCCAACATCTGGAAGCAACTTTTGTAAACTGGTACAAGCAGCTCCCAGGAACGGC

- 35 CCCCAAAGTCCTCATCTATACTAATAATCAGCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGCTCC AAGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGTCTGATTATTACT GCGCAACATGGGATGACAGTATGAATGGTTGGGTGTTCCGGCGGAGGGACCAAACTGACCGTCCTA GGTCAGCCCAAGGCTGCCCCTCGGTCACTCTGTTCCCACCCTCCTCTGAGGAGCTTCAAGCCAAC AAGGCCACACTGGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGCCTGGAAGGCA
- 45 QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEQ ID NO: 694

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20D3

CAGTCTGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTCTTGT TCTGGAAGCAGCTCCAACATCGGAAGCAATTTTGTAAACTGGTACAAGCAGCTCCCAGGAACGGCCCCCAAAGTCCTCATCTATACTAATAATCAGCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGCTCCA

- 60 GCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGTGGAAGTCCCACAGAAGCTACAGCTGCCA GGTCACGCATGAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA

SEQ ID NO: 695

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI 5 SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEQ ID NO: 696

22D1

- 10 CAGTCTGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTCTTGT
 TCTGGAAGCAGCTCCAACATCGGAAGCAATTTTGTAAACTGGTACAAGCAGCTCCCAGGAACGGCC
 CCCAAAGTCCTCATCTATACTAATAATCAGCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGCTCCA
 AGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGTCTGATTATTACTG
 TGCAACATGGGATGACAGTATGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCGTCCTAG
- 20 SEQ ID NO: 697

 $QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS\\ ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI\\ SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK$

25 TVAPTECS SEQ ID NO: 698

22G10

- 35 TGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCCCTCC AATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCACGCAGCACCCTACAGCCTCAGC AGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCACCCA TCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA SEQ ID NO: 699
- EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC
- 45 SEQ ID NO: 700

<u>23A10</u>

- 55 GTCAAGGCGGAGTGGAGACCACCACCACCACAACAAAGCAACAACAAGTACGCGGCCAGCAG CTATCTGAGCCTGACGCCTGAGCAGTGGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGA AGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA SEQ ID NO: 701
- 60 SYELTQPPSVSVSPGQTASITCSGDRLGEKYVCWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF

YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS SEO ID NO: $702\,$

5 **25F8**

CAGTCTGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTCTTGT TCTGGAAGCAGCTCCAACATCGGAAGGAATTTTGTAAACTGGTATAAGCAGCTCCCAGGAACGGCC CCCAAAGTCCTCATTTATACTAATAATCAGCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGCTCCA AGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGTCTGATTATTACTG

- 15 GGTCACGCATGAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA SEQ ID NO: 703

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC

20 LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS SEQ ID NO: 704

25G10

- 25 GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCT GCAGGGCCAGTCAGAGTGTTAGCAGCAGCTACTTAGCCTGGTACCAGCAGAAACCTGGCCAGGCT CCCAGGCTCCTCATCTTTGGTGCATCCAGCAGGGCCACTGGCATCCCAGACAGGTTCAGTGGCAGT GGGTCTGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCCTGAAGATTTTGCAGTGTATCAC TGTCAGCAGTATGGTAACTCACCGCTCACTTTCGGCGGAGGGACCAAGGTGGAGATCAAACGAACT
- 30 GTGGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGTACCGCCTCTG
 TTGTGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGTGGATAACGCCC
 TCCAATCGGGTAACTCCCAGGAGAGTGTCACAGAGCAGGACAGCAAGGACAGCACCTACAGCCTC
 AGCAGCACCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCCTGCGAAGTCAC
 CCATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTGTTGA
- 35 SEQ ID NO: 705

 $EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD\\FTLTISRLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP\\REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF$

40 NRGEC SEQ ID NO: 706

26D1

- CACTCTGTGCTGACTCAGTCACCCTCAGCGTCTGGGACCCCCGGACAGAGGGTCACCATCTCTTGTT

 45 CTGGAAGCCGCTCCAACATCGGAAGTAATTTTGTAAACTGGTACCAGCAGCTCCCAGGAACGCCC
 CCAAACTCCTCATCTATACTAATAATCAGCGGCCCTCAGGGGTCCCTGACCGATTCTCTGGCTCCAA
 GTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGATTATTACTGT
 GCAGTATGGGATGACAGCCTGAATGGTTGGGTGTTCCGCCGGAGGGACCAAGCTGACCGTCCTAGG
 TCAGCCCAAGGCTGCCCCCTCGGTCACTCTGTTCCCACCCTCCTCTGAGGAGCTTCAAGCCAACAA
- - HSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS
- 60 SEQ ID NO: 708

26F12

10 CGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGTGGAAGTCCCACAGAAGCTACAGCTGCCAG GTCACGCATGAAGGGAGCACCGTGGAGAAGACAGTGGCCCCTACAGAATGTTCATGA SEQ ID NO: 709

QSVLTQSPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS

ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI
SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK
TVAPTECS
SEQ ID NO: 710

20 TABLE IIIc: Heavy Chain Variable and Contant Region Polynucleotide and Amino acid Sequences

13586 HC [hu anti-<huCDH19> 4F3 VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK

30 LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 711

13589 HC [hu anti-<huCDH19>4A9 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLS

VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGT

AALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSN

TKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW

YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPR

EPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVD

40 KSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 712

13590 HC [hu anti-<huCDH19>4B10 VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGR
45 FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSASTKGPSVFPLAPSSKS
TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK

50 LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 713

13874 HC [hu anti-<huCDH19> 17H8.2 VH]::huIgG1z

- QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV

 DTSKNQFSLKLSSVTAADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKS
 TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
 KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
 GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
- 60 LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 714

13875 HC [hu anti-<huCDH19> 16C1.1 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNOFSLTLSSLTAADTAVYFCARDGSSGWYRWFDPWGOGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLOSSGLYSLSSVVTVPSSSLGTOTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPOVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

10 SEQ ID NO: 715

13876 HC [hu anti-<huCDH19> 16A4.1 VH]::huIgG1z

OVOLOESGPGLAKPSETLSLTCTVSGDSITSYYWSWIROPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTTVTVSSASTKGPSVFPLAPS 15 SKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLOSSGLYSLSSVVTVPSSSLGTOTYICN VNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDP EVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK AKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLY SKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 **SEO ID NO: 716**

13877 HC [hu anti-<huCDH19> 22G10.1 VH]::huIgG1z

EVOLLESGGGLVOPGGSLRLSCAASGFTFSSYAMNWVROAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLOMNSLRAADTAVYHCAKGGMGGYYYGMDVWGOGTTVTVSSASTKGPSVFPLAP 25 SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGOPREPOVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFF

LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

30 SEQ ID NO: 717

13878 HC [hu anti-<huCDH19> 20D3.1 VH]::huIgG1z

OVOLVOSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVROAPGOGLEWMGIINPISVSTSYAOKFOGRV 35 TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK

40 LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 718

13879 HC [hu anti-<huCDH19> 22D1.1 VH]::huIgG1z

- QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV 45 TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS ${\tt TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN}$ HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
- 50 LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 719

13880 HC [hu anti-<huCDH19> 25F8.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR 55 VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK ${\tt STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN}$ HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK

LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK 60

SEQ ID NO: 720

13881 HC [hu anti-<huCDH19> 26F12.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

10 SEQ ID NO: 721

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13882 HC [hu anti-<huCDH19> 26D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPRFFQYNSTYRVVSVI TVI HODWI NGKFYKCKVSNKAI PAPIFKTISKA

NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEQ ID NO: 722

13883 HC [hu anti-<huCDH19> 25G10.1 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNOFSLKLSSVTAADTAVYYCARDGSSGWYRWFDPWGOGTLVTVSSASTKGPSVFPLAPSSKST

- 25 SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 723

13885 HC [hu anti-<huCDH19> 19B5.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS

- 35 TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 724

14022 HC [hu anti-<huCDH19> 4A2 VH]::huIgG1z

 $QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV\\TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKST$

- 45 SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 725

14024 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKSTS

- 55 GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 726

14025 HC [hu anti-<huCDH19>4A2 VH]::huIgG1z

QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 727

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14026 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 728

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14027 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P,D111E) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCAREGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

SEQ ID NO: 729

SEQ ID NO: 730

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14028 HC [hu anti-<huCDH19> 4A2 (1-472)(Q17E,H47P,D111E,W134Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVT ISVDTSKNQFSLKLSSVTAADTAVYYCAREGSSGYYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

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14029 HC [hu anti-<huCDH19> 4A2 VH]::huIgG1z

QVQLQESGPGLVKPSQTLSLTCTVSGGSISSSGYYWSWIRQHPGKGLEWIGYIYYTGSAYYNPSLKSRV TISVDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 731

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14030 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G) VH]::huIgG1z

RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 732

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRG

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14031 HC [hu anti-<huCDH19>4F3 (1-471)(R17G,T47A) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRG RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQOGNVFSCSVMHEALHNHYTQKSLSLSPGK

10 SEQ ID NO: 733

14032 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,R141Q) VH]::huIgG1z

RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSASTKGPSVFPLAPSSK
STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

OVOLVESGGGVVOPGGSLRLSCAASGFSFSSYDMDWVROAPGKGLEWVAVIWYDGSNKYYADSVRG

20 SEO ID NO: 734

14033 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,D61E,D72E,R141Q) VH]::hulgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRGRFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSASTKGPSVFPLAPSSK

- 25 STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 735

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14034 HC [hu anti-<huCDH19> 4F3 (1-471)(R17G,T47A,D61E,D72E,W134Y,R141Q) VH]::huIgG1z QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRG

RFTISRDNSKNTLFLQMNSLRVEDTAVYYCARETGEGYYFDLWGQGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLEPPKPKDTLMISRTPEVTCVVVDVSHEDPEV

- HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 736

14039 HC [hu anti-<huCDH19> 2G6 (1-477)(R17G,D61E,D72E,K94N) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKD RFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 45 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 737

14040 HC [hu anti-<huCDH19> 16C1.1 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLTLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS

- 55 GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 738

14041 HC [hu anti-<huCDH19> 16C1.1 (1-469)(T92K) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 739

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14042 HC [hu anti-<huCDH19>16C1.1 (1-469)(T92K,D109E) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCAREGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 740

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14043 HC [hu anti-<huCDH19>16C1.1 (1-469)(T92K,W132Y,W135Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGYYRYFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTSG GTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKP SNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFN

25 SNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFN WYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQP REPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 741

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14044 HC [hu anti-<huCDH19> 16C1.1 (1-469)(T92K) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS IDTSKNQFSLKLSSLTAADTAVYFCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

40 SEQ ID NO: 742

14045 HC [hu anti-<huCDH19> 17H8.2 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKS

45 TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

50 SEQ ID NO: 743

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14046 HC [hu anti-<huCDH19> 17H8.2 (1-471)(D109E) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG

 $QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL\\TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK$

60 SEQ ID NO: 744

14047 HC [hu anti-<huCDH19> 17H8.2 (1-471)(D109E,W132Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGYYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 745

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14048 HC [hu anti-<huCDH19>17H8.2 (1-471)(D109E) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSSASTKGPSVFPLAPSSKST SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 746

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14049 HC [hu anti-<huCDH19>4F7 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNT

25 KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 747

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14050 HC [hu anti-<huCDH19>4F7 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNT

35 KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 748

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14051 HC [hu anti-<huCDH19> 4F7 (1-468)(W113Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNQFSLKLSSVTAADTAVYYCARNYAFHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNT KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVELEPPKPKDTI MISRTPEVTCVVVDVSHEDPEVKENWV

45 KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 749

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14052 HC [hu anti-<huCDH19> 4B10 (1-471)(R17G,D61E,D72E,W134Y) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDYSFDYWGQGTLVSVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN

55 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 750

14053 HC [hu anti-<huCDH19>4B10 VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWOOGNVFSCSVMHEALHNHYTOKSLSLSPGK

10 SEQ ID NO: 751

14054 HC [hu anti-<huCDH19> 4B10 (1-471)(R17G) VH]::hulgG1z OVOLVESGGGVVOPGGSLRLSCAASGFTFSSYDMHWVROAPGKGLEWVAVISYDGTNEYYADSVKG

RFTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSASTKGPSVFPLAPSSK
STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEO ID NO: 752

14055 HC [hu anti-<huCDH19> 4B10 (1-471)(R17G,D61E,D72E) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGR FTISRDTSKNTLYLQMNSLRAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSASTKGPSVFPLAPSSKS

- 25 TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 753

14056 HC [hu anti-<huCDH19>4A9 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGT

- 35 AALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSN TKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPR EPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVD KSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 754

14057 HC [hu anti-<huCDH19>4A9 (1-468)(F55I,A56G) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGT

- 45 AALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSN TKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPR EPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVD KSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 755

14058 HC [hu anti-<huCDH19>4A9 (1-468)(F55I,A56G) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNWAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGT

- 55 AALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSN TKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPR EPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVD KSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 756

14059 HC [hu anti-<huCDH19>4A9 (1-468)(F55I,A56G,W113Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLS VDTSKNQFSLKLSSVTAADTAVYYCARNYAFHFDFWGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNT KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 757

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14060 HC [hu anti-<huCDH19> 20D3.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV
TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS
TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO: 758

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14061 HC [hu anti-<huCDH19> 20D3.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN

25 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 759

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14062 HC [hu anti-<huCDH19> 20D3.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKDSNTKVDKKVYEDKSCDKTHTCDDCDADELLGGDSVELEDDKDKDTI MISDTDEVTCVVVDVSHEDDEN

35 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 760

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14063 HC [hu anti-<huCDH19>20D3.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

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SEQ ID NO: 761

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14064 HC [hu anti-<huCDH19> 20D3.1 (1-469)(W133Y) VH]::huIgG1z

 $QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV\\TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSKS\\TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN$

55 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 762

14065 HC [hu anti-<huCDH19> 22G10.1 (1-470)(S82R,A99E) VH]::huIgG1z

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISRDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWOOGNVFSCSVMHEALHNHYTOKSLSLSPGK

10 SEQ ID NO: 763

14066 HC [hu anti-<huCDH19>22G10.1 (1-470)(A99E,H105Y) VH]::huIgG1z

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEO ID NO: 764

14067 HC [hu anti-<huCDH19> 22G10.1 (1-470)(A99E) VH]::huIgG1z

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP

- 25 SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 765

14068 HC [hu anti-<huCDH19>22G10.1 (1-470)(A99E) VH]::huIgG1z

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR FTISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP

- 35 SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 766

14069 HC [hu anti-<huCDH19>22G10.1 (1-470)(D72E,A99E) VH]::huIgG1z

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISSDNSKSTLYLQMNSLRAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAPS

- 45 SKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICN VNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDP EVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK AKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLY SKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 767

14070 HC [hu anti-<huCDH19> 22G10.1 (1-470)(H105Y) VH]::huIgG1z

 $EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGR\\FTISSDNSKSTLYLQMNSLRAADTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSSASTKGPSVFPLAP$

- 55 SSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHE DPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFF LYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 768

14071 HC [hu anti-<huCDH19>16A4.1 (1-474)(T144L) VH]::huIgG1z

QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISV DTSKNQFSLKLSSVTAADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSSASTKGPSVFPLAPS SKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICN VNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDP EVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK AKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLY SKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 769

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14072 HC [hu anti-<huCDH19>19B5.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 770

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14073 HC [hu anti-<huCDH19> 19B5.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GOPREPOVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSK

LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 771

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14074 HC [hu anti-<huCDH19>19B5.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN

35 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 772

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14075 HC [hu anti-<huCDH19>19B5.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKYDKKVEPKSCDKTHTCPPCPAPEILGGPSVELEPPKPKDTI MISPTPEVTCVVVDVSHEDPEV

45 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 773

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14076 HC [hu anti-<huCDH19> 19B5.1 (1-469)(W133Y) VH]::huIgG1z

 $QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV\\TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS\\TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN$

55 HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 774

14077 HC [hu anti-<huCDH19> 23A10.3 (1-474)(L92Q) VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGR FTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQOGNVFSCSVMHEALHNHYTQKSLSLSPGK

10 SEQ ID NO: 775

14078 HC [hu anti-<huCDH19> 23A10.3 (1-474)(R17G,L92Q) VH]::huIgG1z OVOLVESGGGVVOPGGSLRLSCAASGFTFSRYGIHWVROAPGKGLEWVAVIWYDGSNKYYADSVKG

RFTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSASTKGPSVF
PLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQ
TYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDV
SHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE
KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG
SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEQ ID NO: 776

14079 HC [hu anti-<huCDH19> 23A10.3 (1-474)(R17G,D61E,D72E,L92Q) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGR FTISRDNSKNTLYLQMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 25 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 777

14080 HC [hu anti-<huCDH19> 23A10.3 VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGR FTISRDNSKNTLYLLMNSLRAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 35 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 778

14081 HC [hu anti-<huCDH19> 25G10.1 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSSASTKGPSVFPLAPSSKST

- 45 SGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVK FNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG QPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 779

14082 HC [hu anti-<huCDH19>25G10.1 (1-469)(D109E,W132Y,W135Y) VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMS VDTSKNQFSLKLSSVTAADTAVYYCAREGSSGYYRYFDPWGQGTLVTVSSASTKGPSVFPLAPSSKSTS

- 55 GGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHK PSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT VDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 780

14083 HC [hu anti-<huCDH19> 26D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEO ID NO: 781

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14084 HC [hu anti-<huCDH19> 26D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 782

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15

14085 HC [hu anti-<huCDH19> 26D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

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SEQ ID NO: 783

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14086 HC [hu anti-<huCDH19> 26D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 784

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14087 HC [hu anti-<huCDH19>26D1.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR VTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIKLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 785

50

14088 HC [hu anti-<huCDH19> 26D1.1 (1-469)(R27G,G82R) VH]::huIgG1z

VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 786

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGR

60

14089 HC [hu anti-<huCDH19> 26F12.1 VH]::hulgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQOGNVFSCSVMHEALHNHYTOKSLSLSPGK

10 SEQ ID NO: 787

14090 HC [hu anti-<huCDH19> 26F12.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEQ ID NO: 788

14091 HC [hu anti-<huCDH19> 26F12.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQG RLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS

- 25 KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 789

14092 HC [hu anti-<huCDH19> 26F12.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS

- 35 KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 790

14093 HC [hu anti-<huCDH19> 25F8.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV

- HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 791

14094 HC [hu anti-<huCDH19> 25F8.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK

- 55 STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 792

14095 HC [hu anti-<huCDH19> 25F8.1 (1-469)(F90Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 793

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14096 HC [hu anti-<huCDH19> 25F8.1 (1-469)(F90Y) VH]::hulgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSASTKGPSVFPLAPSS KSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPE VKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKA KGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYS KLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 794

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14097 HC [hu anti-<huCDH19>25F8.1 (1-469)(F90Y,W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGR VTMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLYLHFDYWGQGTLVTVSSASTKGPSVFPLAPSSK STSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 795

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14098 HC [hu anti-<huCDH19> 22D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV
TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS
TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO: 796

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14099 HC [hu anti-<huCDH19> 22D1.1 VH]::huIgG1z

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 797

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV

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14100 HC [hu anti-<huCDH19> 22D1.1 (1-469)(W133Y) VH]::huIgG1z

TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 798

14101 HC [hu anti-<huCDH19> 22D1.1 (1-469)(W133Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV
TMTRDTSTSTVFMELSSLRSEDTAVYYCARGGIQLYLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS
TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQOGNVFSCSVMHEALHNHYTQKSLSLSPGK

10 SEQ ID NO: 799

14102 HC [hu anti-<huCDH19> 22D1.1 (1-469)(F90Y) VH]::huIgG1z

QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRV
TMTRDTSTSTVYMELSSLRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSASTKGPSVFPLAPSSKS
TSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVN
HKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAK
GQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSK
LTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

20 SEO ID NO: 800

13591 HC [hu anti-<huCDH19> 4F7 VH]::huIgG1z

QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISL DTSKNOFSLKLSSVTAADTAVYYCARNWAFHFDYWGOGTLVTVSSASTKGPSVFPLAPSSKSTSGGTA

- 25 ALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNT KVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY VDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKS RWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 30 SEQ ID NO: 801

14301 HC [hu anti-<huCDH19>2G6 VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 35 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 40 SEQ ID NO: 802

14302 HC [hu anti-<huCDH19> 2G6 (1-477)(R17G,K94N) VH]::huIgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 45 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 50 SEQ ID NO: 803

14303 HC [hu anti-<huCDH19> 2G6 (1-477)(D61E,D72E) VH]::huIgG1z

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP

- 55 LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
- 60 SEQ ID NO: 804

14304 HC [hu anti-<huCDH19> 2G6 (1-477)(R17G) VH]::hulgG1z

QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKD RFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDG SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK SEQ ID NO: 805

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TABLE IIId: Light Chain Variable and Contant Region Polynucleotide and Amino acid Sequences

13586 LC [hu anti-<huCDH19> 4F3 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD

FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP
REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF
NRGEC

SEQ ID NO: 806

20 13589 LC [hu anti-<huCDH19> 4A9 VL]::huLLC-C1

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQFPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

25 SEQ ID NO: 807

13590 LC [hu anti-<huCDH19>4B10 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FALTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP

30 REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 808

13874 LC [hu anti-<huCDH19> 17H8.2 VL]::huKLC

35 DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKGTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC

SEQ ID NO: 809

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13875 LC [hu anti-<huCDH19> 16C1.1 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEO ID NO: 810

13876 LC [hu anti-<huCDH19> 16A4.1 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTD

50 FTLTISRLEPEDFAVYYCQQYGSSPFTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP
REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF
NRGEC

SEQ ID NO: 811

55 <u>13877 LC [hu anti-<huCDH19> 22G10.1 VL]::huKLC</u>

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

60 SEQ ID NO: 812

13878 LC [hu anti-<huCDH19> 20D3.1 VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 813

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13879 LC [hu anti-<huCDH19> 22D1.1 VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS

ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI
SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK
TVAPTECS
SEQ ID NO: 814

15 13880 LC [hu anti-<huCDH19> 25F8.1 VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

20 SEQ ID NO: 815

13881 LC [hu anti-<huCDH19> 26F12.1 VL]::huLLC-C2

QSVLTQSPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI

25 SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 816

13882 LC [hu anti-<huCDH19> 26D1.1 VL]::huLLC-C2

30 HSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEO ID NO: 817

13883 LC [hu anti-<huCDH19> 25G10.1 VL]::huKLC

 $\label{thm:construction} EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD\\ FTLTISRLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP\\ REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF$

40 NRGEC

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SEQ ID NO: 818

13885 LC [hu anti-<huCDH19> 19B5.1 VL]::huLLC-C2

QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS

45 ASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI
SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK
TVAPTECS
SEQ ID NO: 819

50 14022 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFTVYYCQQYGSSFTFGPGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

55 SEQ ID NO: 820

14024 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]::huKLC

 $\label{thm:constraint} \hline EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF\\ TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE\\ \hline$

60 AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

SEQ ID NO: 821

14025 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

SEQ ID NO: 822

10 14026 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

15 SEQ ID NO: 823

14027 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE

20 AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

SEQ ID NO: 824

14028 LC [hu anti-<huCDH19> 4A2 (1-236)(N30Q,T102A,P141Q) VL]::huKLC

25 EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC

SEQ ID NO: 825

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14029 LC [hu anti-<huCDH19> 4A2 (1-236)(R29Q,N30S) VL]::huKLC

SEQ ID NO: 826

14030 LC [hu anti-<huCDH19> 4F3 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 827

45 <u>14031 LC [hu anti-<huCDH19> 4F3 VL]::huKLC</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

50 SEQ ID NO: 828

14032 LC [hu anti-<huCDH19> 4F3 VL]::huKLC

 $EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD\\FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP$

55 REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 829

14033 LC [hu anti-<huCDH19>4F3 VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVOWKVDNALOSGNSOESVTEODSKDSTYSLSSTLTLSKADYEKHKVYACEVTHOGLSSPVTKSF

SEQ ID NO: 830

14034 LC [hu anti-<huCDH19>4F3 VL]::huKLC

10 EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 831

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14039 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]::huLLC-C1

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVCLISDFY PGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVA

20 PTECS

SEQ ID NO: 832

14040 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 833

30 <u>14041 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]::huKLC</u>

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

35 SEQ ID NO: 834

14042 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP

40 REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 835

14043 LC [hu anti-<huCDH19> 16C1.1 (1-235)(H105Y) VL]::huKLC

45 EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISGLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 836

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14044 LC [hu anti-<huCDH19> 16C1.1 (1-235)(G95R,H105Y,G141Q) VL]::huKLC

 $EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD\\FTLTISRLEPEDFAVYYCQQYGNSPLTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP\\REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF$

55 NRGEC

SEQ ID NO: 837

14045 LC [hu anti-<huCDH19>17H8.2 (1-235)(G149R) VL]::huKLC

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF

 ${\tt YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC}$

SEQ ID NO: 838

5 14046 LC [hu anti-<huCDH19> 17H8.2 (1-235)(G149R) VL]::huKLC

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC

10 SEQ ID NO: 839

14047 LC [hu anti-<huCDH19> 17H8.2 (1-235)(G149R) VL]::huKLC

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF

15 YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC

SEQ ID NO: 840

14048 LC [hu anti-<huCDH19> 17H8.2 (1-235)(S57Y,G149R) VL]::huKLC

DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGT DFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC

SEQ ID NO: 841

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14049 LC [hu anti-<huCDH19> 4F7 (1-239)(H57Y) VL]::huLLC-C2

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

SEQ ID NO: 842

14050 LC [hu anti-<huCDH19> 4F7 (1-239)(H57Y,D110E) VL]::huLLC-C2

35 QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSG
TSASLAITGLQAEDEADYYCQSYESSLSGWVFGGGTRLTVLGQPKANPTVTLFPPSSEELQANKATLVC
LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTECS
SEQ ID NO: 843

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14051 LC [hu anti-<huCDH19> 4F7 (1-239)(D110E) VL]::huLLC-C2

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESSLSGWVFGGGTRLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV

45 EKTVAPTECS SEQ ID NO: 844

14052 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD

50 FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPR
EAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN
RGEC
SEQ ID NO: 845

55 14053 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPR EAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC

60 SEQ ID NO: 846

14054 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPR EAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC

SEQ ID NO: 847

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14055 LC [hu anti-<huCDH19> 4B10 (1-236)(H45Q,A90T) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTD

10 FTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPR
EAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN
RGEC

SEQ ID NO: 848

15 <u>14056 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L) VL]::huLLC-C1</u>

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

20 SEQ ID NO: 849

14057 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L) VL]::huLLC-C1

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC

25 LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

SEQ ID NO: 850

14058 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L,D110E) VL]::huLLC-C1

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QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

35 SEQ ID NO: 851

14059 LC [hu anti-<huCDH19> 4A9 (1-239)(F47L,D110E) VL]::huLLC-C1

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYESRLSGWVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVC

40 LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

SEQ ID NO: 852

14060 LC [hu anti-<huCDH19> 20D3.1 (1-235)(S102A) VL]::huLLC-C2

45 QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEO ID NO: 853

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14061 LC [hu anti-<huCDH19> 20D3.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK

55 TVAPTECS

SEQ ID NO: 854

14062 LC [hu anti-<huCDH19> 20D3.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI

SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 855

5 14063 LC [hu anti-<huCDH19> 20D3.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDESLQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

10 SEQ ID NO: 856

14064 LC [hu anti-<huCDH19> 20D3.1 (1-235)(W109Y) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDESDYYCATYDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI

15 SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 857

14065 LC [hu anti-<huCDH19> 22G10.1 VL]::huKLC

20 EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 858

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14066 LC [hu anti-<huCDH19> 22G10.1 VL]::huKLC

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 859

14067 LC [hu anti-<huCDH19> 22G10.1 (1-234)(Q97E,S98P) VL]::huKLC

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEF
TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP
REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF
NRGEC

SEQ ID NO: 860

40 14068 LC [hu anti-<huCDH19> 22G10.1 (1-234)(V78F,Q97E,S98P) VL]::huKLC

EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEF TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

45 SEQ ID NO: 861

14069 LC [hu anti-<huCDH19> 22G10.1 (1-234)(V78F,Q97E,S98P) VL]::huKLC

 $\label{thm:constraint} EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEF\\ TLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP$

50 REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 862

14070 LC [hu anti-<huCDH19> 22G10.1 VL]::huKLC

55 EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGSGTEF TLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC SEQ ID NO: 863

14071 LC [hu anti-<huCDH19> 16A4.1 (1-235)(G141Q) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGSSPFTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF

SEQ ID NO: 864

14072 LC [hu anti-<huCDH19> 19B5.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

10 QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCL ISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVE KTVAPTECS SEO ID NO: 865

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14073 LC [hu anti-<huCDH19> 19B5.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCL ISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVE

20 KTVAPTECS

SEQ ID NO: 866

14074 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A) VL]::huLLC-C2

QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT
SASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC
LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTECS
SEQ ID NO: 867

- 30 14075 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A,D111E,N135Q) VL]::huLLC-C2 QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS
- 35 SEQ ID NO: 868

14076 LC [hu anti-<huCDH19> 19B5.1 (1-235)(T11V,K45Q,S102A,W109Y,D111E,N135Q) VL]::huLLC-C2

QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT

40 SASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC
LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTECS
SEO ID NO: 869

45 <u>14077 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]::huLLC-C2</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS

50 SEQ ID NO: 870

14078 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]::huLLC-C2

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF

55 YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS

SEQ ID NO: 871

14079 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S,D110E) VL]::huLLC-C2

60 SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA TLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF

 $YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV\\ APTECS$

SEQ ID NO: 872

5 14080 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42Y) VL]::huLLC-C2

 $SYELTQPPSVSVSPGQTASITCSGDRLGEKYVYWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA\\TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF\\YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV\\APTECS$

10 SEQ ID NO: 873

14081 LC [hu anti-<huCDH19> 25G10.1 (1-235)(H105Y) VL]::huKLC

EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

SEQ ID NO: 874

14082 LC [hu anti-<huCDH19> 25G10.1 (1-235)(H105Y) VL]::huKLC

20 FIVI

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EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTD FTLTISRLEPEDFAVYYCQQYGNSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYP REAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF NRGEC

25 SEQ ID NO: 875

14083 LC [hu anti-<huCDH19> 26D1.1 (1-235)(S7P) VL]::huLLC-C2

HSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 876

14084 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P) VL]::huLLC-C2

35 QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEQ ID NO: 877

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14085 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P,W109Y) VL]::huLLC-C2

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK

45 TVAPTECS SEQ ID NO: 878

14086 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P,W109Y,D111E,N135Q) VL]::huLLC-C2

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS

ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI
SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK
TVAPTECS
SEQ ID NO: 879

- 55 <u>14087 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P,W109Y,D111E,N135Q) VL]::huLLC-C2</u> QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS
- 60 SEQ ID NO: 880

14088 LC [hu anti-<huCDH19> 26D1.1 (1-235)(H1Q,S7P) VL]::huLLC-C2

QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 881

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14089 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P) VL]::huLLC-C2

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS

ASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI
SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK
TVAPTECS
SEQ ID NO: 882

15 <u>14090 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,D111E) VL]::huLLC-C2</u>

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

20 SEQ ID NO: 883

14091 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,D111E) VL]::huLLC-C2

QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI

25 SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS

SEQ ID NO: 884

14092 LC [hu anti-<huCDH19> 26F12.1 (1-235)(S7P,W109Y,D111E,N135Q) VL]::huLLC-C2

30 QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEQ ID NO: 885

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14093 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV

40 EKTVAPTECS

SEQ ID NO: 886

14094 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT
45 SASLAISGLQSEDEADYYCAAWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC
LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTECS
SEQ ID NO: 887

50 <u>14095 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A) VL]::huLLC-C2</u>

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDDSLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

55 SEQ ID NO: 888

14096 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A,D111E) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT SASLAISGLQSEDEADYYCAAWDESLNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC

60 LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

SEO ID NO: 889

14097 LC [hu anti-<huCDH19> 25F8.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGT
SASLAISGLQSEDEADYYCAAWDESLQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVC
LISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTECS
SEO ID NO: 890

10 14098 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCL ISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVE KTVAPTECS

15 SEQ ID NO: 891

14099 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,D111E,N135Q) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCL

20 ISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVE KTVAPTECS SEQ ID NO: 892

14100 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,W109Y,D111E,N135Q) VL]::huLLC-C2

25 QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS SEQ ID NO: 893

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14101 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A,W109Y) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS ASLAISGLQSEDEADYYCATYDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLI SDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK

35 TVAPTECS

SEQ ID NO: 894

14102 LC [hu anti-<huCDH19> 22D1.1 (1-235)(K45Q,S102A) VL]::huLLC-C2

QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTS

40 ASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCL
ISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVE
KTVAPTECS
SEO ID NO: 895

45 <u>13591 LC [hu anti-<huCDH19> 4F7 VL]::huLLC-C1</u>

QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSG TSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLGQPKANPTVTLFPPSSEELQANKATLVC LISDFYPGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV EKTVAPTECS

50 SEQ ID NO: 896

14301 LC [hu anti-<huCDH19> 2G6 (1-234)(D110E) VL]::huLLC-C1

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVCLISDFY

55 PGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVA PTECS

SEQ ID NO: 897

14302 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]::huLLC-C1

60 SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVCLISDFY

PGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVA PTECS SEQ ID NO: $898\,$

5 <u>14303 LC [hu anti-<huCDH19> 2G6 (1-234)(C42S,D110E) VL]::huLLC-C1</u>

SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTAT LTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLGQPKANPTVTLFPPSSEELQANKATLVCLISDFY PGAVTVAWKADGSPVKAGVETTKPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVA PTECS

10 SEQ ID NO: 899

14304 LC [hu anti-<huCDH19> 23A10.3 (1-231)(C42S) VL]::huLLC-C2

SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTA
TLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQANKATLVCLISDF
YPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV
APTECS
SEQ ID NO: 900

20

TABLE IVa: HEAVY CHAIN CDRs

	: HEAV	Y CHAIN CDRs		
Ab	Туре	CDR 1	CDR 2	CDR 3
14039	7.7	SYGMH	FIWYEGSNKYYAESVKD	RAGIIGTIGYYYGMDV
14303	AA	SEQ ID NO: 28	SEQ ID NO: 901	SEQ ID NO: 30
14027	7. 7.	SSGYYWS	YIYYTGSAYYNPSLKS	EGSSGWYFQY
	AA	SEQ ID NO: 46	SEQ ID NO: 47	SEQ ID NO: 902
14028	2.2	SSGYYWS	YIYYTGSAYYNPSLKS	EGSSGYYFQY
	AA	SEQ ID NO: 46	SEQ ID NO: 47	SEQ ID NO: 903
14059	7.7	GYYWS	YFSYSGSTNYNPSLKS	NYAFHFDF
	AA	SEQ ID NO: 52	SEQ ID NO: 53	SEQ ID NO: 904
14052	7.7	SYDMH	VISYEGTNEYYAESVKG	ERYFDYSFDY
	AA	SEQ ID NO: 58	SEQ ID NO: 905	SEQ ID NO: 906
14055		SYDMH	VISYEGTNEYYAESVKG	ERYFDWSFDY
	AA	SEQ ID NO: 58	SEQ ID NO: 905	SEQ ID NO: 60
14033		SYDMD	VIWYEGSNKYYAESVRG	ETGEGWYFDL
	AA	SEQ ID NO: 70	SEQ ID NO: 907	SEQ ID NO: 72
14034		SYDMD	VIWYEGSNKYYAESVRG	ETGEGYYFDL
	AA	SEQ ID NO: 70	SEQ ID NO: 907	SEQ ID NO: 908
14051		SYSWS	YIYYSGSTNYNPSLKS	NYAFHFDY
	AA	SEQ ID NO: 82	SEQ ID NO: 83	SEQ ID NO: 909
14046		SYYWS	YIYYIGSTNYNPSLKS	ESRYRSGWYDAFDI
14048	AA	SEQ ID NO: 94	SEQ ID NO: 95	SEQ ID NO: 910
14047	7.7	SYYWS	YIYYIGSTNYNPSLKS	ESRYRSGYYDAFDI
	AA	SEQ ID NO: 94	SEQ ID NO: 95	SEQ ID NO: 911
14042		GYYWS	YIYYIGSTNYNPSLKS	EGSSGWYRWFDP
	AA	SEQ ID NO: 100	SEQ ID NO: 101	SEQ ID NO: 912
14043	7.7	GYYWS	YIYYIGSTNYNPSLKS	DGSSGYYRYFDP
	AA	SEQ ID NO: 100	SEQ ID NO: 101	SEQ ID NO: 913
14069	7.7	SYAMN	TISGGGANTYYAESVKG	GGMGGYYYGMDV
	AA	SEQ ID NO: 118	SEQ ID NO: 914	SEQ ID NO: 120
	r	1	<u> </u>	I

Ab	Туре	CDR 1	CDR 2	CDR 3
14062	73.73	SYFIH	IINPISVSTSYAQKFQG	GGIQLYLHFDY
14063 14064	AA	SEQ ID NO: 124	SEQ ID NO: 125	SEQ ID NO: 915
14100	73.73	SYFIH	IINPISVSTSYAQKFQG	GGIQLYLHLDY
14101	AA	SEQ ID NO: 130	SEQ ID NO: 131	SEQ ID NO: 916
14097	7. 7.	SYYIH	IINPSGGSTRYAQKFQG	GGIQLYLHFDY
	AA	SEQ ID NO: 136	SEQ ID NO: 137	SEQ ID NO: 917
14091	7.7	NYYMS	IINPSGGDSTYAQKFQG	GGIQLYLHFDY
14092	AA	SEQ ID NO: 142	SEQ ID NO: 143	SEQ ID NO: 918
14087	73.73	SYYMS	IIHPSGGDTTYAQKFQG	GGIKLYLHFDY
	AA	SEQ ID NO: 148	SEQ ID NO: 149	SEQ ID NO: 919
14082	73.73	GYYWS	YIYYIGSTNYNPSLKS	EGSSGYYRYFDP
	AA	SEQ ID NO: 154	SEQ ID NO: 155	SEQ ID NO: 920
44070	73. 75	RYGIH	VIWYEGSNKYYAESVKG	RAGIPGTTGYYYGMDV
14079	AA	SEQ ID NO: 160	SEQ ID NO: 921	SEQ ID NO: 162
	7. 7.	SYFIH	IINPISVSTSYAQKFQG	GGIQLYLHLDY
14073	AA	SEQ ID NO: 1	SEQ ID NO: 2	SEQ ID NO: 3
14076	7. 7.	SYGMH	VIWYDGSNKYYADSVKG	RAGIIGTTGYYYGMDV
	AA	SEQ ID NO: 4	SEQ ID NO: 5	SEQ ID NO: 6

TABLE IVb: LIGHT CHAIN CDRs

Ab	Туре	CDR 1	CDR 2	CDR 3
14039	7.7	SGDRLGEKYTS	QDTKRPS	QAWESSTVV
14302 14303	AA	SEQ ID NO: 922	SEQ ID NO: 197	SEQ ID NO: 923
14301		SGDRLGEKYTC	QDTKRPS	QAWESSTVV
	AA	SEQ ID NO: 196	SEQ ID NO: 197	SEQ ID NO: 923
14022		RASRQISSSYLA	GPSSRAT	QQYGSSFT
14024 14025 14026 14027 14028	AA	SEQ ID NO: 924	SEQ ID NO: 215	SEQ ID NO: 216
14029		RASQSISSSYLA	GPSSRAT	QQYGSSFT
	AA	SEQ ID NO: 925	SEQ ID NO: 215	SEQ ID NO: 216
14058	T	TGSSSNIGTGYAVH	GNNNRPS	QSYESRLSGWV
14059	AA	SEQ ID NO: 220	SEQ ID NO: 221	SEQ ID NO: 926
14050	T	TGSSSNIGTGYDVH	GNSNRPS	QSYESSLSGWV
14051	AA	SEQ ID NO: 250	SEQ ID NO: 251	SEQ ID NO: 927
14063		SGSSSNIGSNFVN	TNNQRPS	ATWDESLQGWV
	AA	SEQ ID NO: 292	SEQ ID NO: 293	SEQ ID NO: 928
14064		SGSSSNIGSNFVN	TNNQRPS	ATYDDSLNGWV
	AA	SEQ ID NO: 292	SEQ ID NO: 293	SEQ ID NO: 929
14099		SGSSSNIGSNFVN	TNNQRPS	ATWDESMQGWV
	AA	SEQ ID NO: 298	SEQ ID NO: 299	SEQ ID NO: 930
14100	7.7	SGSSSNIGSNFVN	TNNQRPS	ATYDESMQGWV
	AA	SEQ ID NO: 298	SEQ ID NO: 299	SEQ ID NO: 931
14101	AA	SGSSSNIGSNFVN	TNNQRPS	ATYDDSMNGWV

Ab	Туре	CDR 1	CDR 2	CDR 3
		SEQ ID NO: 298	SEQ ID NO: 299	SEQ ID NO: 932
14096	7. 7.	SGSSSNIGRNFVN	TNNQRPS	AAWDESLNGWV
	AA	SEQ ID NO: 304	SEQ ID NO: 305	SEQ ID NO: 933
14097	7.7	SGSSSNIGRNFVN	TNNQRPS	AAWDESLQGWV
	AA	SEQ ID NO: 304	SEQ ID NO: 305	SEQ ID NO: 934
14090	7.7	SGSRSNIGSNFVN	TNYQRPS	AVWDESLNGWV
14091	AA	SEQ ID NO: 310	SEQ ID NO: 311	SEQ ID NO: 935
14092	7.7	SGSRSNIGSNFVN	TNYQRPS	AVYDESLQGWV
	AA	SEQ ID NO: 310	SEQ ID NO: 311	SEQ ID NO: 936
14085	7.7	SGSRSNIGSNFVN	TNNQRPS	AVYDDSLNGWV
	AA	SEQ ID NO: 316	SEQ ID NO: 317	SEQ ID NO: 937
14086		SGSRSNIGSNFVN	TNNQRPS	AVYDESLQGWV
14087	AA	SEQ ID NO: 316	SEQ ID NO: 317	SEQ ID NO: 938
14077		SGDRLGEKYVS	QDNKWPS	QAWDSSTVV
14078 14304	AA	SEQ ID NO: 939	SEQ ID NO: 329	SEQ ID NO: 330
14079	7.7	SGDRLGEKYVS	QDNKWPS	QAWESSTVV
	AA	SEQ ID NO: 939	SEQ ID NO: 329	SEQ ID NO: 940
14080	7.7	SGDRLGEKYVY	QDNKWPS	QAWDSSTVV
	AA	SEQ ID NO: 941	SEQ ID NO: 329	SEQ ID NO: 330
14075	7.7	SGSRSNIGSNFVN	TNNQRPS	ATWDESMQGWV
	AA	SEQ ID NO: 334	SEQ ID NO: 335	SEQ ID NO: 942
14076	7.7	SGSRSNIGSNFVN	TNNQRPS	ATYDESMQGWV
	AA	SEQ ID NO: 334	SEQ ID NO: 335	SEQ ID NO: 943

human and cynomologous monkey cadherin-19 sequences

			SEQUENCE
ruman Cadherin-19	Human	e e	MNCYLLLRFMLGIPLLWPCLGATENSQTKKVKQPVRSHLRVKRGWVWNQFFVPEEMNTTSHHIGQLRSDLDNGNNSFQYKLLGAGAGAGSTFIIDERTGDIYAIQKLDREERSLYILRAQVIDIATGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTASDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKESLAYRLIVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTFDIITNHETQEGIVILKKKVDFEHQNHYGIRAKVKNHHVPEQLMKYHTEASTTFIKIQVEDVDEPPLFLLPYYVFETPQGSFVGVVSATDPDNRKSPIRYSITRSKVFNINDNGTITTSNSLDREISAWYNLSITATEKYNIEQISSIPLYVQVLNINDHAPEFSQYYETYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNNSSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLTTTHVVCDCGDSGSTQTCQYQGLVLSMGFKTEVIIAILICIMIIFGFIFLTLGLKQRRKQILFPEKSEDFRENIEQISSLOTYAFEGSLGSLAGSLSSLSAVSDODESYDYLNELGPRFKRLACMFGSAVOSNNGPDSAIFRKFILEKLEEANTDPCAPPFDSLOTYAFEGTGSLAGSLSSLESAVSDODESYDYLNELGPRFKRLACMFGSAVOSNN
Cadherin-19	Human	t t	atgaactgttatttactgctgcttttatgttgggaattcctctctatggcttgttttggaacaagaaaactctcaaaacaaa gaaagtcaagcaagcaagcagcagctgtttggaagtgaagtgaacaatgggaacaattttttgtaccagtagaggaatgaat
	ממנופו וו-דמ	במנו בו וו-דה	מַנוֹפּוּ - בּרַ

SFO	DESIGNATION	SOURCE	TYPF	SEOLIFINGE
N N S			I : :	
				ggccccgacagtgccatattcaggaaattcattctggaaaagctcgaagaagctaatactgatccgtgtgcccctccttttgattccctccc
946	Cyno	Macaca	aa	MNCYLLLPFMLGIPLLWPCLGATENSQTKKVQQPVGSHLRVKRGWVWNQFFVPEEMNTTSHHVGRLRSDLDNGNNSFQYKLLGAGA
	Cadherin-19	fascicular		GSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDITTGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTA
		<u>.s</u>		SDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKE
				SLYRLTVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTFDIITNHETQEGIVILKKKVNFEHQNHYGIRAKVKNHHVDEQ
				LMKYHTEASTTFIKIQVEDVDEPPLFLLPYYIFEIFETPQGSFVGVVSATDPDNRKSPIRYSITRSKVFNIDDNGTITTNSLDR
				EISAWYNLSITATEKYNIEQISSIPVYVQVLNINDHAPEFSQYYESYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNS
				SSFTIIDNQDNTAVILTNRTGFNLQEEPIFYISILIADNGIPSLTSTNTLTIHVCDCDDSGSTQTCQYQELMLSMGFKTEVIIAIL
				ICIMVIFGFIFLTLGLKQRRKQILFPEKSEDFRENIFRYDDEGGGEEDTEAFDVAALRSSTIMRERKTRKTTSAEIRSLYRQSLQV
				GPDSAIFRKFILEKLEEADTDPCAPPFDSLQTYAFEGTGSLAGSLSSLESAVSDQDESYDYLNELGPRFKRLACMFGSAVQSNN
947	Cyno	Macaca	nt	AIGAAITGITAITTACIGCIGCCTITITAIGIIGGGAAIICCICCCTAIGGCCTIGICTIGGAGCAACAGAAAACICTCAAACAAA
	Cadherin-19	fascicular		GAAAGICCAGCAGCAGTAGGAICTCAICTGAGAGTGAAGCGTGGCTGGGTGTGGAACCAAITTTTTGTACCAGAGGAAATA
		<u>.v</u>		CGACTAGTCATCACGTTGGCCGGCTAAGATCTGATTTAGACAATGGAAACAATTCTTTCCAGTACAAGCTTTTGGGAGCTGGAGCT
)		GGAAGTACTTTTATCATTGATGAAGAACAGGTGACATATATAT
21				AAGAGCCCAGGTAATAGACATCACTACTGGAAGGGCTGTGGAACCTGAGTCTGAGTTTGTCATCAAAGTTTCGGATATCAATGACA
				AIGAACCAAAAIICCIAGAIGAACCIIAIGAGGCCAIIGIACCAGAGAIGICICCCAGAAGGAACAIIAGICAICCAGGIGACAGCA
				AGTGATGCTGATGACCCTTCAAGTGGTAATAATGCTCGTCTCCTCTACAGCTTATTACAAGGCCAGCCA
				AACAACAGGAGTCATAAGAATATCTTCTAAAATGGATAGAGAACTGCAAGATGAGTATTGGGTAATCATTCAAGCCAAGGACATGA
				TIGGICAGCCAGGAGCGIIGICIGGAACAACGAGIGIAITAATIAAACITICAGAIGITAATGACAATAAGCCIAITATITAAAGAA
				AGTITATACCGCCTGACGGTCTCTGCATCTGCACCCACTGGGACTTCTATAGGAACAATCATGGCATATGATAATGACATAGGAGA
				GAATGCAGAAATGGATTACAGCATTGAAGAGGATGATTCACAGACATTTGACATTATTACTAATCATGAAACTCAAGAAGGAATAG
				TTATATTAAAAAAAAAAGAAAGTGAATTTTGAGCACCACCACTATGGTATTAGAGCAAAAAGTTAAAAAACCATCATGTTGATGAGCAG
				CICAIGAAAIACCACACIGAAGCIICCACCACTIICAIIAAGAICCAGGIGGAAGAIGIIGAIGAGAGCCICCICIIIIICCICCIICCI
				GTATTACATATTTGAAATTTTTGAAGAAACCCCACAAGGATCATTTGTAGGCGTGGTGTCTGCCACAGACCCAGACCAATAGGAAAT
				CICCIAICAGGIAIICIAIIACIAGGAGCAAAGIGIICAAIAICGAIGAIAAIGGIACAAICACIACAACIAACI
				GAAATCAGTGCTTGGTACAACCTAAGTATTACAGCCACAGAAAAATACAATATAGAGCAGATCTCTTCGATCCCAGTGTATGTGCA
				AGTICITAATATCAATGATCATGCTCCTGAGTICTCTCAATACTATGAGAGTTATGTTTGTGAAAAATGCAGGCTCTGGTCAGGTAA
				TICAGACTATCAGTGCAGTGGATAGAGATGAATCCATAGAAGAGCACCATTTTTACTTTAATCTATCT
				TCAAGITITIACAATCATAGACAATCAAGATAACACAGCTGTCATTTTGACTAATAGAACTGGTTTTAACCTTCAAGAAGAGCCCAT
				CITCTACATCTCCATCTTAATTGCCGACAATGGAATCCCGTCACTTACAAGTACAAACACCCTTACCATCCAT
				ATGACAGTGGGAGCACACACACCTGCCAGTACCAGGAGCTTATGCTTTCCATGGGATTCAAGACAGAAGTCATCATTGCTATTCTC
				ATTIGCATTATGGTAATATTTGGGTTTATTTTTTGACTTTGGGTTTAAAACAACGGAGAAAACAGATTCTATTTCCTGAGAAAAG
				TGAAGATTTCAGAGAGAATATATTCCGATATGATGACGAAGGGGGGTGGAGAAGAAGAAGATACAGAGGCCTTTGACGTAGCAGCGCTGA
				GGAGTAGCACCATAATGCGGGAACGCAAGACTCGGAAAACCACCAGCGCTGAGATCAGGAGCCTATACAGGCAGTCTTTGCAAGTT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>o</u> 8				
				GGCCCCGACAGTGCCATATTCAGGAAGTTCATCCTGGAAAAGCTCGAAGAAGCTGATACTGATCCGTGTGCCCCTCCTTTTGATTC CCTCCAGACCTACGCTTTTGAGGGAACAGGGTCATTAGCTGGATCCCTGAGCTCCTTAGAATCAGCTGTCTCTGATCAGGATGAAA GCTATGATTACCTTAACGAGTTGGGACCTCGCTTTAAAAGATTAGCATGCAT
948	secreted	Human	aa	MNCYLLLRFMLGIPLLWPCLGATENSQTKKVKQPVRSHLRVKRGWVWNQFFVPEEMNTTSHHIGQLRSDLDNGNNSFQYKLLGAGA
	Cadherin-19			GSTFIIDERTGDIYAIQKLDREERSLYILRAQVIDIATGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTA
	ecto-domain			SDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKE etvot mysboa degestatma vondicena bendystedden bendystenden i mydbendelytii syyyydbendniyych dayyynduidda
	(amino acids			SLIKLIVSESAFIGISIGILMAIDNDIGENAEMDISIEEDDSQIFDIINNEIQEGIVILKAKVUFEHQNRIGIKAKANAN HETARAKVATTINNEILIMAIGIKAKONN IMKYHTRASTTFIKIOVEDVDRPPIRIIPYVYFRYFRYFRIPOGSFVGVVSATDPDNRKSPITRSKVFNINDNGTITTSNSI,DR
	1-596)			EISAWYNLSITATEKYNIEQISSIPLYVQVLNINDHAPEFSQYYETYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNN
				SSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLTSTNTLTIHVCDCGDSGSTQTCQYQELVLSMGFKTE
949	secreted	Human	nt	atgaactgttatttactgctgcgttttatgttgggaattcctctcctatggccttgtcttggagcaacagaaaactctcaaacaaa
	Cadherin-19			gaaagtcaagcagccagtgcgatctcatttgagagtgaagcgtggctgggtgtggaaccaattttttgtaccagaggaaatgaata
	ecto-domain			cgactagtcatcacatcggccagctaagatctgatttagacaatggaaacaattctttccagtacaagcttttggggagctggagct
	(amino acids			ggaagtacttttatcattgatgaagaacaggtgacatatatgccatacagaagcttgatagagaggagcgatcctttacttt
	1-596)			aagagcccaggtaatagacatcgctactggaagggctgtggaacctgagtctgagtctgagtttgtcatcaaagtttcggatatcaatgaca
2	1			atgaaccaaaattcctagatgaaccttatgaggccattgtaccagagatgtctccagaaggaacattagttatccaggtgacagca
216				agtgatgctgacgatccctcaagtggtaataatgctcgtctcctctacagcttacttcaaggccagcca
<u>.</u>				aacaacaggagtcataagaatatcttctaaaatggatagagaactgcaagatgagtagttgggtaatcattcaagccaaggacatga
				ttggtcagccaggagcgttgtctggaacaacaagtgtattaatta
				agtttataccgcttgactgtctctgaatctgcacccactgggacttctataggaacaatcatggcatatgataatgacataggaga
				gaatgcagaaatggattacagcattgaagaggatgattcgcaaacatttgacattattactaatcatgaaactcaagaaggaatag
				ttatattaaaaaagaaagtggattttgagcaccagaaaccactacggtattagagcaaaagttaaaaaaccatcatgttcctgagcag
				ctcatgaagtaccacactgaggcttccaccactttcattaagatccaggtggaagatgttgatgagcctcctcttttcctccttcc
				atattatgtatttgaagtttttgaagaaaccccacaggggatcatttgtaggcgtggtgtctgccacagacccagacaataggaaat
				ctcctatcaggtattctattactaggagcaaagtgttcaatatcaatgataatggtacaatcactacaagtaactcactggatcgt
				gaaatcagtgcttggtacaacctaagtattacagccacagaaaaatacaatatagaacagatctcttcgatcccactgtatgtgca
				agttettaacateaatgateatgeteetgagtteteteteaataetatgagaettatgagaetttatgtgaaaatgeaggetetggteaggtaa
				ttcagactatcagtgcagtggatagagatgaatccatagaagagcaccatttttactttaatctatct
				tcaagttttacaatcatagataatcaagataacacagctgtcattttgactaatagaactggttttaaccttcaagaagaacctgt
				cttctacatctccatcttaattgccgacaatggaatcccgtcacttacaagtacaaacaccttaccatccat
				gtgacagtggggagcacacagacctgccagtaccaggagcttgtgctttccatgggattcaagacagaa
920	truncated	Human	aa	MNCYLLLRFMLGIPLLWPCLGATENSQTKKVKQPVRSHIRVKRGWVWNQFFVPEEMNTTSHHIGQLRSDLDNGNNSFQYKLLGAGA
	membrane			GOLT TIDERTIGOLIAND TREEKOLI LIKAQVI DIATORAKE PEDETA INVODINUNE PRI EDEFIELAT VERMANEGETUA. INVITATORAKE PRI EDEFIELAT VERMANEGETUA INVITATORAKE PRI EDEFIELAT VERMANEGETUA INVITATORAKE PRI EDEFIELAT VERMANEGETUA INVITATORAKE PRI EDEFIELAT VERMANEGETUA PRI EDEFIELAT VERMANEGETUA INVITATORAKE PRI EDEFIELAT VERMANEGETUA INVITATORAKE PRI EDEFIELAT VERMANEGETUA PRI EDEFIELATUA INVITATORAKE PRI EDEFIELATUA PRI EDEFI
	bound form of			SDADDPSSGNNAKLLISLLQGQPIFFSVEFTTGVIKISSKMDKELQDEIWVIIQAKDMIGQFGALSGTTSVLIKLSDVNDNKFIFKE siybi mysesadmomsiomimaydnndicenaemmysieeddesomedtimneemoeciiiiikkmydeeunnhycidarknhhydeo
	human			SELVELVSESSKI ISTSIGIIKKAIDNEIGENSEKERISISIEEDESKII ELIINNEIKEGIVIENKAVE ENKANTIIVITTINNIINVEK LMKYHTEASTTIIKIOVEDVDEPPLFILLPYYVEEVFEETPOGSFYGVVSATDPDNRKSPIRYSITRSKVFNINDNGTITTSNSLDR
	cadherin-19			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9.				
	(amino acids 1-624)			EISAWYNLSITATEKYNIEQISSIPLYVQVLNINDHAPEFSQYYETYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNN SSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLTSTNTLTIHVCDCGDSGSTQTCQYQELVLSMGFKTEVIIAIL ICIMIIFGFIFLTLGLKQRRKQ
217	truncated membrane bound form of human cadherin-19 (amino acids 1-624)	Humanl	ţ	atgaactgttatttactgctgcqttttatgttgggaattcctctctatggccttgtcttggagcaacagaaaactctcaaacaaa
952	C137897 huCDH19 (44-141) muCDH19 (140-770)	artificial	aa	GWVWNQFFVPEEMNTTSHHIGQLRSDLDNGNNSFQYKLLGAGAGSTFIIDERTGDIYAIQKLDREERSLYILRAQVIDIATGRAVE PESEFVIKVSDINDNEPRELDEPYEAIVPEMSPEGTFVIKVTANDADDPSTGYHARILYNLERGQPYFSVEPTTGVIRISSKMDRE LQDTYCVIIQAKDMLGQPGALSGTTTVSIKLSDINDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSK IFDIIIDNDTQEGIVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPVFLLPYYILEIPEGKPYGT IVGTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQF YETYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDTNNSSFMLTDNQDNTAVILSNRTGFNLKEEPVFYMIILIADNGIPS LTSTNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVWVIFGFFFLILALKQRRKKETLFPEKTEDFRENIFCYDDEG GGEEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAG SLSSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
953	C137897 huCDH19	artificial	nt	

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
218	(44-141) muCDH19 (140-770)			ccatacagaagcttgatagaagaggagccatctacatcttaagagcccaggttaatagacatcgctactggaaaggctgtggaaccttgattcatcaaggttgatcatcaatgacaattaaccattgatcctcagaagttgatcatcaaggttgaaccaattgatgccctgaatcctgaattcctagattgatcctccagaaggtgaacatttgtaccaatgagccaatgaagccaatgaaccaattagaacctttaaaactgataccttaaacaggatgtacacaacaacaggattacttctcaagatggaacattcttaaagatggatcatcaacaacaggattcaatacattaaagttgaacatattaaggccaatgacaacaacaggatcaattcaacaacaacaggatcaattaaagaaag
954	C137896 huCDH19 (44-249) muCDH19 (248-770)	artificial	99	GWVWNQFFYPEEMNTTSHHIGQLRSDLDNGNNSFQYKLLGAGAGSTFIIDERTGDIYAIQKLDREERSLYILRAQVIDIATGRAVE PESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTASDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRE LQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSK IFDIIIDNDTQEGIVVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPVFLLPYYILEIPEGKPYGT IVGTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQF YETYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDTNNSSFMLTDNQDNTAVILSNRTGFNLKEEPVFYMIILIADNGIPS LTSTNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVMVIFGFFFIILALKQRRKETLFPEKTEDFRENIFCYDDEG GGEEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAG SLSSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
955	C137896 huCDH19 (44-249)	artificial	nt	ggctgggtgtggaaccaattttttgtaccagaggaaatgaatacgactagtcatcacatcggccagctaagatctgatttagacaa tggaaaacaattctttccagtacaagcttttgggagctggaagtactggaagtacttttatcattgatgaaagaacaggtgacatatatgcatacatccagtacatatatgccatacagaaagcgatccctctacatcttaagagcccaggtaatagaaggcgatcgtgtggaa

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
219	(248-770)			cctgaytctgaytttgtcatcaaagtttcggatatcaatgacaatgaaccaaaattcctagatgaaccttatgagccattgtaccagagagtttgtcatcaaagtgataccaacaagtgatgatcataaagtgatatcatcaaagtgataccattactacaatgagaaacttcacaagtgatactacacaacaagtgatataaatctctcaaaatggataataacttctcacaaatggataataaata
956	C137913 muCDH19 (44-139) huCDH19 (142-249) muCDH19 (248-770)	artificial	e e	AWVWRPETVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE SEFVIRVLDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTASDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQ DEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSKIF DIIIDNDTQGGIVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPVFLLPYYILEIPEGKFYGTIV GTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAMYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQFYE TYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDTNNSSFMLTDNQDNTAVILSNRTGFNLKEEPVFYMIILIADNGIPSLT STNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVMVIFGFFFLILALKQRRKETLFPEKTEDFRENIFCYDDEGGG EEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAGSL SSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
957	C137913 muCDH19 (44-139) huCDH19	artificial	nt	goctgggtgtggagaccatttgttgttgttctagaaagaaatggatgatatatacaatgtgttggaaagctaagatctgacttagacaatggaaactggggattggcatac aaacaactctttccagtacaagctactggggattggcgctggaagctttagcattaatgaaagaacaggtgaaatatgtgccatac agaagcttgatagagaggaaaaatccctctacattctgagagcccaggtaatagacaccactattgggaaggctgtggaaactgaa tccgagtttgtcatcagagttttggatatcaatgacaatgaaccaaaattcctagatgaaccttatgaggccattgtaccagagat

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
N S				
220	(142-249) muCDH19 (248-770)			gtctccagaaggaacattagttatccaggtgacagcaagtgatgctgacagtgatcctcaagtggtaataatgctcctctcacaggtcctccagaaggactctcaaggagctctctcaaagtggttatcagtcaagaactgaagaacacaagaagaagaagaagaagaagaagaagaaga
958	C137847 muCDH19 (44-139) huCDH19 (142-364) muCDH19 (363-770)	artificial	вв	AWVWRPFVVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE SEFVIRVLDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTASDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQ DEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVNDNKPIFKESLYRLTVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTF DIITNHETQEGIVILKKKVDFEHQNHYGIRAKVKNHHVPEQLMKYHTEASTTFIKIQVEDVDEPPVFLLPYYILEIPEGKPYGTIV GTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQFYE TYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDTNNSSFMLTDNQDNTAVILSNRTGFNLKEEPVFYMIILIADNGIPSLT STNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVMVIFGFFFLILALKQRRKETLFPEKTEDFRENIFCYDDEGGG EEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAGSL SSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
959	C137847 muCDH19 (44-139) huCDH19 (142-364)	artificial	nt	gcctgggtgtggagaccatttgttgttctagaagaaatggatgatatacaatgtgtttggaaagctaagatctgacttagacaatgg aaacaactgtttccagtacaagctactggggattggcgctggaagctttagcattaatgaaaagaacaggtgaaatatgtgccatac agaagcttgatagagagaaaaatccctcacattctgagaagcccaggtaatagacaccactattgggaaaggctgtggaaactgaa tccgagtttgtcatcagaggacactattgggaaactgaaactgaa tccgagtttgtcatcagagttttggatatcaatgacaatgaaccaaaattcctagatgaaccttatgaggccattgtaccagagat tccgagtttgtcccagaaggaacattagaaccaaagtgatgacagatgaccatacaagtgatccctcaaagtggtaataccagaagat

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
221	muCDH19 (363-770)			gettacttcaaggccatcatttttctgttgaaccaacaacaaggagtcataagaatatcttctaaaatggatagaactgcaa gatgagtattcaaggcaattctcaagaagacatttttcaattcaagacaagaaggacattggtcaggaacagaactgtcattcaagacattcaagacaattaagacattcaagacaattagacaacatttaagacaatcatgaaattgacaacattgaagacattactaacaacattggaaacaacatttactaatacaagacaattgaagacaagaagaagaagaagaagacaacaagaaaggaattacaagaaaggattcaagaacatttactaaacaacattactaaacaacattactaaacaac
096	C137911 muCDH19 (44-247) huCDH19 (250-364) muCDH19 (363-770)	artificial	e e	AWVWRPFVVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE SEFVIRVLDINDNEPRFLDEPYEAIVPEMSPEGTFVIKVTANDADDPSTGYHARILYNLERGQPYFSVEPTTGVIRISSKMDRELQ DTYCVIIQAKDMLGQPGALSGTTTVSIKLSDINDNKPIFKESLYRLTVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTF DIITNHETQEGIVILKKKVDFEHQNHYGIRAKVKNHHVPFQLMKYHTEASTTFIKIQVEDVDEPPVFLLPYYILEIPEGKPYGTIV GTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQFYE TYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDTNNSSFMLTDNQDNTAVILSNRTGFNLKEEPVFYMIILIADNGIPSLT STNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVMVIFGFFFLILALKQRRKETLFPEKTEDFRENIFCYDDEGGG EEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAGSL SSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
961	C137911 muCDH19 (44-247) huCDH19 (250-364) muCDH19	artificial	nt	goctgggtgtggagaccatttgttgttgttctagaaatggatgatatacaatgtgttggaaagctaagatctgacttagacaatgg aaacaatgg aaacaactgtttccagtacaaggtgattgtggggattggggatgggaagctttagcattaatgaaagaacaggtgaaatatgtgccatac agaaacatctttccagtacaaggtacttggggaaggctgggaaggccaggttggatagagacaccactattgggaaaggctgtggaaactgaa tccgagtttggtcatcagagttttggatatcaatgacaatgaacccagattcctagatgaaaccatatgaggccattgtacctgagat tccgagtttgtcatcaaggttaccaatgacaatgaccaatgaccaatgaccaatgacccagattcctagatgaccatatgaggccattgtacctgagat gtctccagaagggaacattgtcatcaaggtgacaaccaatgaccaatgaccaatgaccaatgaccaatgaccaatgaccaatgaccaatgaccaatgaccaacaagaatgaccaacaacaacaacaacaacaacaacaacaacaacaac

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
222	(363-770)			gatacatactgtgtaattattcaagccaaggacatgctggtcagccttgacttgtctggaacaacaaccgtatcaattaagct gtcagtattaatgacatataagct gtcagtattaaagaaatggattatacagcttgactgtctctgaatctgcaaccatgggacttctaattagaacaattttaagaacaattttaaaaaaggaattattacatactattaaaaaggaattagattacagaattgagattcaagacacattgggacttctaaaattagaaacttaaaaaagaaag
962	C137917 muCDH19 (44-362) huCDH19 (365-772)	artificial	ee	AWVWRPFVVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE SEFVIRVLDINDNEPRFLDEPYEAIVPEMSPEGTFVIKVTANDADDPSTGYHARILYNLERGQPYFSVEPTTGVIRISSKMDRELQ DTYCVIIQAKDMLGQPGALSGTTTVSIKLSDINDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSKIF DIIIDNDTQEGIVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPLFLLPYYVFEVFEETPQGSFV GVVSATDPDNRKSPIRYSITRSKVFNINDNGTITTSNSLDREISAWYNLSITATEKYNIEQISSIPLYVQVLNINDHAPEFSQYYE TYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNNSSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLT STNTLTIHVCDCGDSGSTQTCQYQELVLSMGFKTEVIIAILLICIMIIFGFIFLTIGLKQRRKQILFPEKSEDFRENIFQYDDEGGG EEDTEAFDIAELRSSTIMRERKTRKTTSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSL SSLESAVSDQDESYDYLNELGPRFKRLACMFGSAVQSNN
963	C137917 muCDH19 (44-362) huCDH19 (365-772)	artificial	nt	gcctgggtgtggagaccatttgttgttctagaagaaatggatgatatacaatgtgttggaaagctaagatctgacttagacaatgg aaacaactgt aaacaactctttccagtacaagctactggggattggcgctggaagctttagcattaatgaaagaacaggtgaaatatgtgccatac agaagctttgctacaatgtgcattctgagagctttagcatcagatgacaccactattgggaaggctgtggaaactgaactgaactgaattggtaatagaagctgtggaaactgaactgaactgaacccaggtttgtcatcagaggttttggaataccattgaaccaatgacaatgaacccaatgacaccaatgacaaccaatgacaaccaatgacaaccaatgacaaccaatgacaacaaagagagatcatcaagagataagaagatgacaacaacaacaagagaccaacaacaacaacaacaacaa

SEO	DESIGNATION	SOURCE	TYPE	SEOUENCE
Q. No				
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				tttttactttaatctatctgtagaagacactaacaattcaagttttacaatcatagataatcaagataacacagctgtcattttga
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				catgggattcaagacagaagtcatcattgctattctcatttgcattatgatcatatttgggtttatttttttt
				aacaacggagaaaacagattctatttcctgagaaaagtgaagatttcagagaaatatattccaatatgatgatgaagggggtgga
				gaagaagatacagaggcctttgatatagcagagctgaggagtagtaccataatgcgggaacgcaagactcggaaaaccacaagcgc
				tgagatcaggagcctatacaggcagtctttgcaagttggccccgacagtgccatattcaggaaattcattc
22				aagetaataetgateegtgtgeeeeteettttgatteeeteeagaeetaegettttgaagggaaeagggteattagetggateeetg
				agctccttagaatcagcagtctctgatcaggatgaaagctatgattaccttaatgagttgggacctcgctttaaaagattagcatg
				catgittggittctgcagtgcagtcaaataattag
964	C137915	artificial	aa	AWWWRPFVVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE
	muCDH19			SEFVIRVLDINDNEPRFLDEPYEAIVPEMSPEGTFVIKVTANDADDPSTGYHARILYNLERGQPYFSVEPTTGVIRISSKMDRELQ
	(44-461)			DTYCVIIQAKDMLGQPGALSGTTTVSIKLSDINDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSKIF
	huCDH19			DIIIDNDTQEGIVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPVFLLPYYILEIPEGKPYGTIV
	(464-772)			GTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDHAPEFSQYYE
	/= · · · · · · · ·			TYVCENAGSGQVIQTISAVDRDESIEEHHFYFNLSVEDTNNSSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLT
				STNTLTIHVCDCGDSGSTQTCQYQELVLSMGFKTEVIIAILICIMIIFGFIFLTLGLKQRRKQILFPEKSEDFRENIFQYDDEGGG
				EEDTEAFDIAELRSSTIMRERKTRKTTSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSL
				SSLESAVSDQDESYDYLNELGPRFKRLACMFGSAVQSNN
965	C137915	artificial	nt	gcctgggtgtgggagaccatttgttgttctagaagaaatggatgatatacaatgtgttggaaagctaagatctgacttagacaatgg
	muCDH19			aaacaactettteceagtacaaagetaectggggattggegettggaagetttagcattaatgaaaagaacaggtgaaatatgtgeceatae
	(44-461)			agaagcttgatagagaagaaaaatccctctacattctgagagcccaggtaatagacaccactattgggaaggctgtggaaactgaa
	huCDH19			tccgagtttgtcatcagagttttggatatcaatgacaatgaacccagattcctagatgaaccatatgaggccattgtacctgagat
	(464-772)			gtctccagaaggaacatttgtcatcaaggtgacagccaatgacgcagatgatcatcttcaactggctatcatgctcgcatcctataca
				gatacatactgtgtaattattcaagccaaggacatgctcggtcagcctggagccttgtctggaacaacaacacgtatcaattaagct
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SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
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				aginicadadacaccicideccaiccaicgicing iganicaging and and and and and an arrangaccan accompany citig and in a sa Catadaatt Caanacaanaant Cataattact attot attot catt tacattat attat catatt tacatt tact the text the cantata and
				Cargagaaricaagacagaagicaricicaarigaaricaaria
				aacaacggagaaaacagattoctatttoctgagaaaagtgaagatttocagagagaatatattocaatatgatgaagggggggggg
				gaagaagatacagaggcctttgatatagcagagctgaggagtagtaccataatgcgggaacgcaagactcggaaaaccaaagcgc
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1				catgtttggttctgcagtcaaataattag
996	C71144	artificial	aa	AWVWRPFVVLEEMDDIQCVGKLRSDLDNGNNSFQYKLLGIGAGSFSINERTGEICAIQKLDREEKSLYILRAQVIDTTIGKAVETE
	muCDH19			SEFVIRVLDINDNEPRFLDEPYEAIVPEMSPEGTFVIKVTANDADDPSTGYHARILYNLERGQPYFSVEPTTGVIRISSKMDRELQ
	(44-770)			DTYCVIIQAKDMLGQPGALSGTTTVSIKLSDINDNKPIFKESFYRFTISESAPIGTSIGKIMAYDDDIGENAEMEYSIEDDDSKIF
	(0), ++			DIIIDNDTQEGIVILKKKVDFEQQSYYGIRAKVKNCHVDEELAPAHVNASTTYIKVQVEDEDEPPVFLLPYYILEIPEGKPYGTIV
				GTVSATDPDRRQSPMRYYLTGSKMFDINDNGTIITTNMLDREVSAWYNLTVTATETYNVQQISSAHVYVQVFNINDNAPEFSQFYE
				TYVCENAESGEIVQIISAIDRDESIEDHHFYFNHSLEDINNSSFMLTDNQDNTAVILSNRTGFNLKEEFVFYMIILIADNGIPSLT
				STNTLTIQVCDCGDSRNTETCANKGLLFIMGFRTEAIIAIMICVMVIFGFFFLILALKQRRKETLFPEKTEDFRENIFCYDDEGGG
				EEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAGSL
				SSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAVQPNN
296	C71144	artificial	nt	gcctgggtgtggagaccatttgttgttctagaagaaatggatgatatacaatgtgttggaaagctaagatctgacttagacaatgg
	muCDH19			aaacaactctttccagtacaagctactggggattggcgctggaagctttagcattaatgaaagaacaggtgaaatatgtgccatac
	(44-770)			agaagottgatagagaagaaaaatcoototacattotgagagoocaggtaatagacaccactattgggaaaggotgtgggaaactgaa
				tccgagtttgtcatcagagttttggatatcaatgacaatgaacccagattcctagatgaaccatatgaggccattgtacctgagat
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SEQ	DESIGNATION SOURCE TYPE	SOURCE	TYPE	SEQUENCE
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NO.				
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				agcacaaacactctcactatccaagtctgtgactgtggagacagtagaaacacagaaaacttgtgctaacaagggacttctctttat
				catgggattcagaacagaggcaataattgccatcatgatatgtgttatggtaatatttgggttttttggtttttt
				aacagcgaaggaaaggagactctatttccagagagaagactgaagactttagggagaatatattttgctatgatgatgaaggcggcggg
				gaagaagactcggaagcctttgacatcgtagagctgagacaaagtacagtaatgagagaaagaa
				ggagatcaggagcttgtacaggcagtccctgcaggtgggcccagacagtgccatatttcgaaaatttatcctagagagcttgaag
				aagccaacacagacccatgtgctccccctttgattcactacagacgtttgcctatgagggaacagggtcatcagctggctctctg
				agotcottggcatccagagacactgatcaggaggatgacttcgactaccttaatgacctgggacctcgttttaaaagattagcaag
22				catgtttggctctgcagtacaacccaacaattag
896 _L	Flag Tag	artificial	aa	DYKDDDDK
696	Flag Tag	artificial	nt	gactacaaagacgatgacgacaag

Bispecific binding molecules

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SEQ	SEQ DESIGNATION SOURCE TYPE SEQUENCE	SOURCE	J AVT	SEQUENCE
<u></u>				
NO.				
970	CDR-H1 of	artificial	ΑA	SYGMH
	CDH19 2G6			
971	CDR-H2 of	artificial AA	γγ	FIWYDGSNKYYADSVKD
	CDH19 2G6			
972	CDR-H3 of	artificial AA	٧V	RAGIIGTIGYYYGMDV
	CDH19 2G6			
973	CDR-L1 of	artificial	ΑA	SGDRLGEKYTC
	CDH19 2G6			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
8				
974	CDR-L2 of CDH19 2G6	artificial	AA	QDTKRPS
975	CDR-L3 of CDH19 2G6	artificial	AA	QAWDSSIVV
926	VH of CDH19	artificial	N	
	2G6			CTCCAGCTACGGCATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCTTCATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAAGTCCTGT CGGGCCGAGGACACCGGTGTACTACTGTGCCAGAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACGGCATGGACGTGTG GGGCCAGGGCACCCGTGTCTAGC
977	VH of CDH19 2G6	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
978	VL of CDH19	artificial	F	TACGAGCTGACCCCAGCCCCCTCCGTGTCCGTGTCTCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGAGAA
	720			
6 <u>7</u> 6 226	VL of CDH19 2G6	artificial	ΑΑ	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
980	VH-VL of	artificial	۲	CAGGIGCAGCIGGIGGAAICCGGCGGAGGCGIGGIGCAGCCIGGCCGGTCCCIGAGACIGICTIGCGCCGCCTCCGGCTTCACCIT
	CDH19 2G6			CICCAGCIACGGCATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCTTCATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAAGTCCCTG
				CGGGCCGAGGACACCGCCGTGTACTACTGTGCCCAGAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACTACTACGACGAGGACGTGTG GGGCCAGGGCACCACCGTGACCGTGTCTAGCGGAGGAGGGAG
				TGACCCCAGCCCCCTCCGTGTCCGTGTCTCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGAGAGTACACC
				TGTIGGIATCAGCAGCGGCCTGGCCAGTCCCCCCTGCTGGTCATCTACCAGGACACCAAGCGGCCCTCCGGCATCCCTGAGCGGTT
				CTCCGGCTCCAACTCCGGCAACACCGCCACCCTGACCATCTCCGGCACCCAGGCCATGGACGAGGCCGACTACTACTGCCAGGCCT GGGACTCCTCCACCGTGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTG
981	VH-VL of	artificial	ΑA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL
	CDH19 2G6			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGSGSSYELTQPPSVSVSVSPGQTASITCSGDRLGEKYT CWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVL
985	CDH19 2G6 x	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL
	12C			
				CWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNIATLIISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGIVOPGGSIKISGAA SGFTFNKYAMNWYROA PGKGIFWYAR IRSKYNNYA TYYADSYKDRFTISRDDSKNTA YLOMNNIKTED
				TAVYYCYRHGUYETT Y DYWAYWGOGTLYTY SSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
				QQKFGQAFKGLIGGIKFLAFGIFAKFSGSLLGGKAALTLSGVQFEDEAETICVLWISNKWVFGGGIKLIVLHHHHHH

SFO	DESIGNATION	SOURCE	TYPE	SFOUFINCE
Ω S O			 	
983	CDR-H1 of CDH19 16E2.1	artificial	AA	SYGMH
984	CDR-H2 of CDH19 16E2.1	artificial	AA	VIWYDGSNKYYADSVKG
985	CDR-H3 of CDH19 16E2.1	artificial	AA	DGWELSFDY
986	CDR-L1 of CDH19 16E2.1	artificial	AA	RASQGISNYLA
286	CDR-L2 of	artificial	AA	AASSLQS
000	CDH19 16E2.1		:	m ##***********************************
9886 88	CDR-L3 of CDH19 16E2.1	artificial	AA	QHYFTYPEKT.
686	VH of CDH19	artificial	N	CAGGIGCAGCIGGIGGAATCCGGCGGAGGCGIGGIGCAGCCIGGCCGGTCCCTGAGACIGICITGCGCCGCCTCCGGCTTCATCTT
227	16E2.1			CTCCAGCTACGGCATGCACTGGGTCCGACAGACCCCCGGCAAGGGCCTGGAATGGGTGGCCGTGATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACATCTCCCAAGAACACCCTGTACCTGCAGATGAACTCCCTG CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGGGACGGCTGGGAGCTGTCCTTCGATTACTGGGGCCAGGGGCACCCTGGTCAC
				CGTGTCTAGC
066	VH of CDH19 16E2.1	artificial	₹	QVQLVESGGGVVQPGRSLRLSCAASGFIFSSYGMHWVRQTPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDISKNTLYLQMNSL RVEDTAVYYCARDGWELSFDYWGQGTLVTVSS
991	VL of CDH19	artificial	N	GACATCCAGATGACCCAGTCCCCCTCCAGCCTGTCCGCCTCCGTGGGCGACAGAGTGACCATCACCTGTCGGGCCTCCCAGGGCAT
	16E2.1			CAGCAACTACCTGGCCTGGCTGCAGCAGAAGCCCGGCAAGGCCCCCAAGTCCCTGATCTACGCCGCCAGCTCCCTGCAGTCCGGCG TGCCCTCCAAGTTCTCCGGCTCTGGCTCCGGCACCGACTTCACCCTGACCATCTCCAGCCTGCAGCCCGAGGACTTCGCCACCTAC
				TACTGCCAGCACTTCACCTACCCCCGGACCTTCGGACAGGCACCAAGGTGGAAATCAAG
992	VL of CDH19	artificial	ΑΑ	DIQMTQSPSSLSASVGDRVTITCRASQGISNYLAWLQQKPGKAPKSLIYAASSLQSGVPSKFSGSGSGTDFTLTISSLQPEDFATY VCOHYFTYPRTFGOGTKVRIK
	1052.1		!	
666	VH-VL of CDH19 16E2.1	artificial	Z	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGCAGCCTGGCCGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCATCTT CTCCAGCTACGGCATGCACTGGGTCCGACAGACCCCCGGCAAGGGCCTGGAATGGGGTGGCCGTGATTTGGTACGACGGCTCAACA
				AGINCIANGE CONTROLLE CONTR
				CGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGGGGGGCTCCGACATCCAGATGACCCAGTCCCCCTCCAGCC
				TGTCCGCCTCCGTGGGCGACAGAGTGACCATCACCTGTCGGGCCTCCCAGGGCATCAGCAACTACCTGGCCTGGCTGCAGAAGAACACCCACACAACTACCTGGCTTGCAGCAGAAGAACAAAAAAAA
				COCGACTICACCCTGACCATCTCCAGCCTGCAGCCCGAGGACTICGCACCTACTACTGCCAGCACTTCACCTACCTACCTGCCGGGA
				CCTTCGGACAGGCCACCAAGGTGGAAATCAAG

SFO	DESIGNATION	SOURCE	TYPE	SEOUENCE
N N N				
994	VH-VL of CDH19 16E2.1	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFIFSSYGMHWVRQTPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDISKNTLYLQMNSL RVEDTAVYYCARDGWELSFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGJIQMTQSPSSLSASVGDRVTITCRASQGISNYLAWLQQK PGKAPKSLIYAASSLQSGVPSKFSGSGSGTDFTLTISSLQPEDFATYYCQHYFTYPRTFGQGTKVEIK
995	CDH19 16E2.1 x I2C	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFIFSSYGMHWVRQTPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDISKNTLYLQMNSL RVEDTAVYYCARDGWELSFDYWGQGTLVTVSSGGGGSGGGSGGGSDIQMTQSPSSLSASVGDRVTITCRASQGISNYLAWLQQK PGKAPKSLIYAASSLQSGVPSKFSGSGSGTDFTLTISSLQPEDFATYYCQHYFTYPRTFGQGTKVEIKSGGGGSEVQLVESGGGLV QPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYYC VRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPGQ APRGLIGGTKFLAAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
966	CDR-H1 of CDH19 17H8.2	artificial	ЧΑ	SXXWS
266	CDR-H2 of CDH19 17H8.2	artificial	AA	YIYYIGSTNYNPSLKS
866	CDR-H3 of CDH19 17H8.2	artificial	AA	DSRYRSGWYDAFDI
66 228	CDR-L1 of CDH19 17H8.2	artificial	AA	RASQSVAGSYLA
1000	CDR-L2 of CDH19 17H8.2	artificial	ΑΑ	GASSRAT
1001	CDR-L3 of CDH19 17H8.2	artificial	AA	QQYGKSPIT
1002	VH of CDH19 17H8.2	artificial	Z	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTAGAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCAT CAACTCCTACTACTGGTCCTGGATCCGGCAGCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATACTACATCGGCTCCACCA ACTACAACCCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACCATCCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCGTGACC GCCGCTGACACCCCCTGTACTACTGCGCCAGAGACTCCCGGTACAGAACCGGGTGCTCGACGGGCCAGGG CACCATGGTCACCGTGTCCTCT
1003	VH of CDH19 17H8.2	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSS
1004	VL of CDH19 17H8.2	artificial	TN	GATATCGTGCTGACCCAGTCCCCCGGCACCCTGTCTCTGAGCCCTGGCGAGAGAGCCCTGTCCTGCCAGAGCCTCTCAGTCCGTGGGCGCGCGC
1005	VL of CDH19 17H8.2	artificial	AA	DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGKSPITFGQGTRLEMK

SFO	DESIGNATION	SOURCE	TVPF	SFOLIENCE
g ⊖ 8			! :	
1006	6 VH-VL of CDH19 17H8.2	artificial	LN .	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCCAT CAACTCCTACTACTGGTCCTGGATCCGGCAGCCCCTGGCAAGGGCCTGGAATGGATGG
1007	7 VH-VL of CDH19 17H8.2	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSINSYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSGGGGGGGGGGGGGGGTULTQSPGTLSLSPGERATLSCRASQSVAGSYLA WYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMK
229		artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSINSYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYCARDSRYRSGWYDAFDIWGQGTWVTVSSGGGGSGGGGGGGDIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLA WYQQKPGQAPRLLISGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1009	9 CDR-H1 of CDH19 19B5.1	artificial	AA	SYFIH
1010	O CDR-H2 of CDH19 19B5.1	artificial	AA	IINPISVSTSYAQKFQG
1011	1 CDR-H3 of CDH19 19B5.1	artificial	AA	GGIQLWLHLDY
1012	2 CDR-L1 of CDH19 19B5.1	artificial	AA	SGSRSNIGSNFVN
1013	3 CDR-L2 of CDH19 19B5.1	artificial	AA	TNNQRPS
1014	4 CDR-L3 of CDH19 19B5.1	artificial	AA	ATWDDSMNGWV
1015	5 VH of CDH19 19B5.1	artificial	Z	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCCGCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTACTTCATCCACTGGGTCCGAAGGCCCCAGGGCCTGGGAATGGATGG

SEQ ID	DESIGNATION	SOURCE	ТУРЕ	SEQUENCE
1016	VH of CDH19 19B5.1	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
1017	VL of CDH19 19B5.1	artificial	F	CAGTCTGCCCTGACCCAGCCTCCCACCACCACCGGCACACCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT CGGCTCCAACTTCGTGAACTGGTACAAGCAGCTGCCCGGCACCGCCCCCCAAGGTGCTGATCTACACCAACAACCAGCGGCCCTCCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCTGGCCATCTCCGGCCTGCAGGACGAGGACGAGTCCGAC TACTACTGTGCCCACCTGGGACGACGTCCATGAACGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTG
1018	VL of CDH19 19B5.1	artificial	AA	QSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCATWDDSMNGWVFGGGTKLTVL
230	VH-VL of CDH19 19B5.1	artificial	L V	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTAACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCAGGGCCTGGAATGGATGG
1020	VH-VL of CDH19 19B5.1	artificial	ΑA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWY KQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVL
1021	СDH19 19В5.1 x I2C	artificial		QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYF1HWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSALTQPPSTTGTPGQRVTISCSGSRSNIGSNFVNWY KQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1022	CDR-H1 of CDH19 20D3.1	artificial	Ą	SYFIH
1023	CDR-H2 of CDH19 20D3.1	artificial	AA	IINPISVSTSYAQKFQG
1024	CDR-H3 of CDH19 20D3.1	artificial	AA	GGIQLWLHFDY
1025	CDR-L1 of CDH19 20D3.1	artificial	ΑA	SGSSSNIGSNFVN
1026	CDR-L2 of	artificial	ΑĄ	TNNQRPS

SFO.	DESIGNATION	SOURCE	TVPF	SEOLIENCE
2 0) ; ;)	! :	
NO.				
	CDH19 20D3.1			
1027	CDR-L3 of CDH19 20D3.1	artificial	AA	ATWDDSLNGWV
1028	VH of CDH19 20D3.1	artificial	LN L	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCAGGGCCTGGAATGGATGG
1029	VH of CDH19 20D3.1	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGIQLWLHFDYWGQGTLVTVSS
1030	VL of CDH19 20D3.1	artificial	FN .	CAGTCTGCCCTGACCCAGCCTCCTTCTGCCACCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCTCCTCCAACAT CGGCTCCAACTTCGTGAACTGGTACAAGCAGCTGCCCGGCACCGCCCCCAAGGTGCTGATCTACACCAACAACCAGCGGCCCTCCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCTGGCCATCTCCGGCCTGCAGTCCGAGGACGAGTCCGAC TACTACTGTGCCCACCTGGGACGACGACTCCTGAACGGCTGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTG
1031	VL of CDH19 20D3.1	artificial	ЧΑ	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCATWDDSLNGWVFGGGTKLTVL
1033	VH-VL of CDH19 20D3.1 CDH19 20D3.1 CDH19 20D3.1 x I2C	artificial artificial	AA AA	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAACCTGGCGCCTCGTGAAGGTGTCCTCAAGCTGTTCCAGGCTTACTAGCGCTTACTAGCGCTACTACTCCAGGTGTTCTGGGGCTACTACTCCAGGCCTACTACTCCAGGTGTTCTAGTGGATGGTGTTCTAGTGGATGTGTTCTAGTGGATGTGTGCTCTACTCGGTGTTCTAGTGGATGTGTTCTAGTGGATGTGTTCTAGTGGATGTGTTCTAGTGTGTGT
1035	CDR-H1 of	artificial	AA	SYFIH

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9				
	CDH19 22D1.1			
1036	CDR-H2 of	artificial	ΑA	IINPISVSTSYAQKFQG
1037	CDR-H3 of	artificial	۵۵	GGTOT.WT.HT.DY
	CDH19 22D1.1	5	;	
1038	CDR-L1 of	artificial	AA	SGSSSNIGSNFVN
	CDH19 22D1.1			
1039	CDR-L2 of CDH19 22D1.1	artificial	AA	TINIQRPS
1040	CDR-L3 of	artificial	Ą	ATWDDSMNGWV
	CDH19 22D1.1			
1041	VH of CDH19 22D1.1	artificial	LN	CAGGIGCAGCIGGIGCAGICIGGCGCCGAAGIGAAGAAACCIGGCGCCTCCGIGCGGGTGICCIGCAAGGIGICCGGCTACCIA CACCAGCIACIICAICCACIGGGICCGACAGGCCCCAGGGCCAGGGCCIGGAAIGGAIGG
23				CCTCCTACGCCCAGAAATTCCAGGGCAGAGTGACCATGACCCGGGACACCTCCACCTCCACCTGTGTGTATGGAACTGTCCTCCCTG CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCCAGCTGTGGCTGCACCTGGACTATTGGGGGCCAGGGCACCT GGTCACCGTGTCCTCT
2 1045	VH of CDH19 22D1.1	artificial	AA	
1043	VL of CDH19	artificial	ΙN	
	22D1.1			CGGCTCCCAACTTCGTGAACTGGTACAAGCAGCTGCCCGGCACCGCCCCCAAGGTGCTGATCTACACCAACAACCAGCGCCTCCG GCGTGCCCGAACTCTCTGGCTCCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGGACGAGGACGAGTCCGACA macana company control of the company of the
				IACIACIGIGCACCIGGGACGACIGCIGGGIGITCGGCGGGGGCACCAAGCIGACCGIGCIG
1044	VL of CDH19 22D1.1	artificial	ΑA	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCATWDDSMNGWVFGGGTKLTVL
1045	VH-VL of	artificial	F	
	CDH19 22D1.1			
				GGECACCGEGICCICIGGGGGGGGAGGAICIGCGGGGGGGGGAGGCGGGGGGGG
				CCGCTACCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCTCCTAACATCGGCTCCAACTTCGTGAACTGGTAC
				CAAGICIGGCACCICCGCCICCIGGCCAICICCGGCCIGCAGICCGAGGACGAGICCGACIACIACIGIGCCACCIGGGACGACI CCAIGAACGGCIGGGIGIICGGCGGAGGCACCAAGCIGACCGIGCIG
1046	VH-VL of	artificial	AA	QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWY

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
	CDH19 22D1.1			KQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVL
1047	CDH19 22D1.1 x12C	artificial		QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSIRSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWY RQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCATWDDSMNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1048	CDR-H1 of CDH19 22G10.1	artificial	AA	SYAMIN
1049	CDR-H2 of CDH19 22G10.1	artificial	АА	TISGGGANTYYADSVKG
23	CDR-H3 of CDH19 22G10.1	artificial	AA	GGMGGYYYGMDV
3	CDR-L1 of CDH19 22G10.1	artificial	AA	RASQSISSNLA
1052	CDR-L2 of CDH19 22G10.1	artificial	AA	GAFTRAT
1053	CDR-L3 of CDH19 22G10.1	artificial	АА	QQYNYWPLT
1054	VH of CDH19 22G10.1	artificial	٧	GAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTTCACCTTCACCTTCACCTTCACCTTCACCTTCACCTTCACCTTCACCTGCACACACA
1055	VH of CDH19 22G10.1	artificial	AA	EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISSDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS
1056	VL of CDH19 22G10.1	artificial	N	GAGATCGTGATGACCCAGTCCCCCGTGACCCTGTCCCTGAGCCTGGGCGAGAGACCACCCTGTCTTGCCGGGCCTCCTCCTTCCCAGCCATCCTTCCCAGCCAG

SEQ NO.	Q DESIGNATION O.	SOURCE	TYPE	SEQUENCE
				TCCCTGCCAGAGTGTCTGGCTCCGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGCAGTCCAGGACTTTGCCGTGTAC TACTGCCAGCAGTACAACTACTGGCCCCTGACCTTCGGAGGCGGCACCAAGGTGGAAATCAAG
10	1057 VL of CDH19 22G10.1	artificial	AA	EIVMTQSPVTLSLSGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVY YCQQYNYWPLTFGGGTKVEIK
10	1058 VH-VL of CDH19 22G10.1	artificial	Ľ	GAGGTGCAGCTGCTGGAGTCCGGCGGAGGACTGGTGCTGGCGGCTCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGCCATGAACTGGGTCCGACGGCCCTGGCAAGGGCCTCGGAATGGGTGTCCACCATCAGCGGCGGAGGCGCCAACA CCTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCTCCGACACTCCACCTGTACCTGCAGATGAACTCCTG AGAGCCGCCGACACCGCGTGTACCATGTGCTAAGGGCGGCATGGGGGCTACTACTACGGCATGGATGTGTGGGGCCACG CACCGTGACCGTGTCTAGCGGAGGATCTGGCGGGCTTCTTGCCGGGCTACTACTACGGCATGGTGTGGGGCCAC CCGTGACCCTGTCCTGGCGGAGGCCGAGGATCTTGCCGGGCCTCCCAGTCCTCCAGCTGCTGGTTCC CCGTGACCCTGACCCTGGCCGAGGAGCCCCCTGTTTTCCCGGGCCTCCCAGTCCTGCCTG
	1059 VH-VL of CDH19 22G10.1	artificial	AA	EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISSDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYGMDVWGQGTTVTVSSGGGGGGGGGGGGGGGGTVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIK
34	1060 CDH19 22G10.1 x I2C	artificial		EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGCGANTYYADSVKGRFTISSDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHH
1061	61 CDR-H1 of CDH19 23A10.3	artificial	ΑА	RYGIH
10	1062 CDR-H2 of CDH19 23A10.3	artificial	ΑĄ	VIMYDGSNKYYADSVKG
10	1063 CDR-H3 of CDH19 23A10.3	artificial	ΑА	RAGIPGTTGYYYGMDV
10	1064 CDR-L1 of CDH19 23A10.3	artificial	АА	SGDRLGEKYVC

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
1065	CDR-L2 of CDH19 23A10.3	artificial	¥4	QDNKWPS
1066	CDR-L3 of CDH19 23A10.3	artificial	AA	QAWDSSTVV
1067	VH of CDH19 23A10.3	artificial	LN .	CAGGIGCAGCIGGIGGAAICCGGCGGAGGCGIGGIGCAGCCIGGCCGGICCCIGAGACIGICITGCGCCGCCTCCGGCTITCACCTT CICCAGAIACGGCAICCACIGGGICCGACAGGCCCCTGGCAAGGGCCTGGAAIGGGIGGCGGTGAIITGGIACGACGGCTCCAACA AGIACIACGCCGACTCCGTGAAGGGCCGGTICACCAICTCCCGGGACAACTCCAAGAACACCCTGTACCTGCTGATGAACTCCCTG CGGGCCGAGGACTCCGCCGTGTACTACTGIGCCAGAAGGGCCGGCATCCCGGCACCACGGCTACTACGGCATGGATGTGTG GGGCCAGGGCACCACGTGTACTACTACTACTACTGTGCCAGAAGGCCGGCACCCGGCACCACCGGCTACTACTACGGCATGGTGTG
1068	VH of CDH19 23A10.3	artificial	AA	QVQLVESGGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
235	VL of CDH19 23A10.3	artificial	TN	TACGAGCTGACCCAGCCCCCCCCCCGTGTCCGTGTCTCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCAGAAATACGTGTTCTGGCGACCGGCTGGGCGAGAAAAACCCGGCTGGGCCATCTGGTTCTACCAGGACAACAAGTGGCCCTCCGGCATCCTGAAAAAAAA
1070	VL of CDH19 23A10.3	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVCWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
1071	VH-VL of CDH19 23A10.3	artificial	L	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCCGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGATACGGCATCCGCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCTGTTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCACTGGAAGGGCCCTTGGTCCTGCTGATCACACACA
1072	VH-VL of CDH19 23A10.3	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV CWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVL
1073	CDH19 23A10.3 x I2C	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGSGSYELTQPPSVSVSPGQTASITCSGDRLGEKYV CWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED

		10000		
SEQ NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGTVTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1074	CDR-H1 of CDH19 25F8.1	artificial	AA	SYYIH
1075	CDR-H2 of CDH19 25F8.1	artificial	ΑA	IINPSGGSTRYAQKFQG
1076	CDR-H3 of CDH19 25F8.1	artificial	AA	GGIQIWLHFDY
1077	CDR-L1 of CDH19 25F8.1	artificial	AA	SGSSSNIGRNFVN
1078	CDR-L2 of CDH19 25F8.1	artificial	AA	TNNQRPS
1079	CDR-L3 of CDH19 25F8.1	artificial	ΑA	AAWDDSLNGWV
236	VH of CDH19 25F8.1	artificial	T	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGCTACACCTTT CACCAGCTACTACATCCACTGGGTCCGACAGGCCCCAGGGCCTGGGAATGGAATGGGCATCATCAACCCCTCCGGCGGCTCCA CCAGATACGCCCCAGAAATTCCAGGGCAGAGTGACCATGACCCGGGACACCTCCACCTCCACCGTGTTCATGGAACTGTCCTCCTC CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCTTCGACTACTGGGGCCCAGGGCACCCT GGTCACCGTGTTCAAC
1081	VH of CDH19 25F8.1	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
1082	VL of CDH19 25F8.1	artificial	Z	CAGTCTGCCCTGACCCAGCCTCCTTCTGCCACCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCTCCTCCAACAT CGGCCGGAACTTCGTGAACTGGTACAAGCAGCTGCCCGGCACCCCCCAAGGTGCTGATCTACACCAACAACAACCTCCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCCTGGCCATCTCCGGGCCTTCTCGAGGCCGAC TACTACTGTGCCGCTGGGACGACCCTCGAACGGCTGGTGCTGCGGAGGCACCTGCAACACCTGCAC
1083	VL of CDH19 25F8.1	artificial	AA	QSALIQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYKQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESD YYCAAWDDSLNGWVFGGGTKLTVL
1084	VH-VL of CDH19 25F8.1	artificial	L	CAGGTGCAGCTGGTGCAGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGCTACACCTT CACCAGCTACTACATCCACTGGGTCCGACAGGCCCCAGGCCTGGAATGGATGG

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	DESIGNALION	SOUNCE] - -	SECUENCE
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				CAAGTCTGGCACCTCCGCTCCCTGGCCATCTCCGGCCTGCAGTCCGAGGACGAGTCCGACTACTACTGTGCCGCCTGGGACGACTACTGAGACGACTACTGTGCCGCCTGGGACGACTACTGAACGGCTGTTGCTGCCTGGGACGACTACTGAACGGCTGTTGCTGCTGCTGGGACGACTACTGAACGGCTGTTGCTGAACGACTACTGAACGTGTTGAACGGCTGTTGAACGGACGACTGAACTGAACGAAC
1085	VH-VL of CDH19 25F8.1	artificial	Ą	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWY KQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVL
1086	CDH19 25F8.1 x I2C	artificial		QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQLTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWY KQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDESDYYCAAWDDSLNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTCVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1087	CDR-H1 of	artificial	AA	GYYWS
	CDH19 25G10.1			
1088	CDR-H2 of	artificial	¥	YIYYIGSTNYNPSLKS
237	CDH19 25G10.1			
1089	CDR-H3 of	artificial	AA	DGSSGWYRWFDP
	CDH19 25G10.1			
1090	CDR-L1 of	artificial	Ą	RASQSVSSSYLA
	CDH19 25G10.1			
1091	CDR-L2 of	artificial	Ą	GASSRAT
	CDH19 25G10.1			
1092	CDR-L3 of	artificial	ΑA	QQYGNSPLT
	CDH19 25G10.1			
1093	VH of CDH19	artificial	IN	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCCAT
	25G10.1			CTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACATCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATGTCCGTGGACACCTCCAAGAAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC GCCGCTGAACCCCGTGTACTGCTGCCCAGAGATGGCTCCTCCGGCTGGTATCGTTGGTTCGACCCTTGGGGCCCAGGGCACCCT
				GGTCACCGTGTCTAGC

SEO	DESIGNATION	SOURCE	TYPE	SEQUENCE
0 N 0.				
1094	VH of CDH19 25G10.1	artificial	ΑΑ	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSS
1095	VL of CDH19	artificial	M	GAGATCGTGCTCGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGCCCACCCTGTCCTGCAGAGCCTCCCAGTCCGT
	25G10.1			GICCICCICCIACCTGGCTTGGTATCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTTCGGCGCCTCTTCCAGAGCCACCG
				TACCACTGCCAGCAGTACGGCAACAGCCCCCTGACCTTCGGCGGGGGCGCCCAAGGTGGAAATCAAG
1096	VL of CDH19	artificial	ΑA	
	25G10.1			YHCQQYGNSPLTFGGGTKVEIK
1097	VH-VL of	artificial	NT	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCAAT CTCCGGCTACTACTGCTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATGG
	25G10.1			ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATGTCCGTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC
				GCCGCTGACACCGCCGTGTACTACTGCGCCAGAGATGGCTCCTCCGGCTGGTATCGTTGGTTCGACCCTTGGGGCCCAGGGCACCCTT GCTCACCGTGTTTACCGCAAAAAAAAAA
23				TGGCTCCGGCACCGACTTCACCCTGACCATCTCCCGGCTGGAACCCGAGGACTTCGCTGTGTACCACTGCCAGCAGTACGGCAACA
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TOSO	CDH19	al cili	{	EVERENCE CHAIR STILLS TO SOCCESSION TO SECOND STANDARD STANDARD SECOND STANDARD SECOND
	25G10.1			VVnrgvarreligasoralgirdni ogogogidi ilionefedravincvvigignofeliggginveln
1099	CDH19 25G10.1 x I2C	artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSSGGGGGGGGGGGGEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWY
				QQKPGQAPKLLIFGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYHCQQYGNSPLTFGGGTKVEIKSGGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV
				YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1100	CDR-H1 of	artificial	ΑΑ	SYYMS
1101	CDR-H2 of	artificial	AA	IIHPSGGDTTYAOKFOG
 	CDH19 26D1.1			
1102	CDR-H3 of CDH19 26D1.1	artificial	ΑĄ	GGIKLWLHFDY
1103	CDR-L1 of	artificial	₩	SGSR,SNIGSNFVN
	CDH19 26D1.1			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
1104	CDR-L2 of CDH19 26D1.1	artificial	ΑΑ	TNNQRPS
1105	CDR-L3 of CDH19 26D1.1	artificial	AA	AVWDDSLNGWV
1106	VH of CDH19	artificial	۲	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTT
	26D1.1			CACCAGCTACTACATGTCCTGGGTCCGACAGGCCCCAGGCCAGGGCCTGGAATGGATGG
				GGTCACCGTGTCTAGC
1107	VH of CDH19 26D1.1	artificial	ΑA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS
1108	VL of CDH19	artificial	N	CATTCCGTGCTGACCCAGTCTCCTTCCGCCTCCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCGGTCCAACAT
	26D1.1			CGGCTCCCAACTTCGTGAACTGGTATCAGCAGCTGCCCCGGCACCGCCCCCAAGCTGCTGATCTACAACAACAACAGCGGCCCTCCG GCGTGCCCGAACTTCTCTGGCTCCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCCGGCCTGCAGTCTGAGGACGAGGCCGAC
				TACTACTGTGCCGTGTGGGACGACTCCCTGAACGGCTGGGTGTTTCGGCGGAGGCACCAAGCTGACCGTGCTG
239	VL of CDH19 26D1.1	artificial	¥	HSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDDSLNGWVFGGGTKLTVL
1110	VH-VL of	artificial	F	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTT
	CDH19 26D1.1			CACCAGCIACIACAIGICCIGGGICCGACAGGCCCCAGGGCCAGGGCCIGGAAIGGAIGG
				CCACCTACGCCCAGAAATTCCAGGGCAGAGTGACCATGACCGGCGACACACCTCCACCTCCACCGTGTATATGGAACTGTCCTCCCTG
				GGTCACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGCTTCTGGCGGCGGCGGAGGCTCCCATTCCGTGTCTGACCCAGTCTCCTT
				CCGCCTCCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACATCGGCTCCAACTTCGTGAACTGGTAT
				CAGCAGCIGCCCGGCACCGCCCCCAAGCIGCIGAICIACACCAACAACCAGCGGCCCICCGGCGIGCCCGACCGGIICICICIGGCIC
				CAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGACTACTACTGTGCCGTGTGGGACGACTACT CCCTGAACGGCTGGGTGTTTCGGCGAAGGAACAAACTGAACGTGTG
1111	VH-VL of	artificial	Ą	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL
	CDH19 26D1.1			RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGSGGGGSHSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWY
				QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVL
1112	CDH19 26D1.1	artificial		QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL
	x I2C			RSEDTAVYYCARGGIKLMLHFDYWGQGTLVTVSSGGGGSGGGSGGGSHSVLTQSPSASGTPGQRVTISCSGSRSNIGSNFVNWY
				QQLFGIAFKLLIIINQKFSGVFDRFSGSKSGISASLAISGLQSEDEADIICAVWDDSLNGWVFGGGIKLIVLSGGGGSEVQLVES GGGIVODGGSIKISCAAAGFFFNKVAMNWVROAPGKGIEWVARIFSKYNNVATVYADSVKDRFTISRDDSKNTAYIOMNNIKTFF
				AVYYCVRHGNEGNSYI SYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
				QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
1113	CDR-H1 of CDH19 26F12.1	artificial	AA	NYYMS
1114	CDR-H2 of CDH19 26F12.1	artificial	AA	IINPSGGDSTYAQKFQG
1115	CDR-H3 of CDH19 26F12.1	artificial	AA	GGIQLWLHFDY
1116	CDR-L1 of CDH19 26F12.1	artificial	AA	SGSRSNIGSNFVN
7111	CDR-L2 of CDH19 26F12.1	artificial	AA	TNYQRPS
81111	CDR-L3 of CDH19 26F12.1	artificial	AA	AVWDDSLNGWV
1119	VH of CDH19 26F12.1	artificial	N.	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACTT CACCAACTACTACTACTTCTGGGTCCGACAGGCCCAGGCCAGGGCCTGGAATGGATGG
1120	VH of CDH19 26F12.1	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
1121	VL of CDH19 26F12.1	artificial	L	CAGTCTGTGCTGACCCAGTCCCCTTCCGCCTCTGGCACCCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGCGGCCCTCCG GCGTGCCCGACCTGCTGGCTCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGAC TACTACTGTGCCGTGTGGGACGACTCCCTGAACGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTG
1122	VL of CDH19 26F12.1	artificial	AA	QSVLTQSPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDDSLNGWVFGGGTKLTVL
1123	VH-VL of CDH19 26F12.1	artificial	F	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACCTTT CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGGCCAGGGCCTGGAATGGATGG

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				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCCAGCTGTGGCTGCACTTCGACTACTGGGGGCCAGGGCACCCT GGTCACCGTGTCTAGCGGAGGGGGGGAGGATCTGGTGGCGGGGGGGG
				CCGCCTCTGGCACCCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACATCGGCTCCAACTTCGTGAACTGGTAT CAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGCGGCCCTCCGGCGTGCCCGACCGGTTCTCTGGGCTC
				CAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGACTACTACTGTGCCGTGTGGGACGACTTCTCTGAGGACGACTTCTGAACGACGACGACTTTCGAACGTGTGGGACGACGACTTTCGAACGACGACGACGACGACTTGAACGAAC
1124	VH-VL of	artificial	AA	
	CDH19 26F12.1			RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGSQSVLTQSPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVL
1125	CDH19	artificial		QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL
	26F12.1 x I2C			
				QQLFGIAFALLIINIQKESGVEDKESGSGSESGISASLAISGLQSEDEADIICAVWDDSLNGWVFGGGIALLIVLSGGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT
				AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2 1126	CDR-H1 of	artificial	AA	GYYWS
	CDH19 4A9			
1127	CDR-H2 of	artificial	ΑA	YFSYSGSTNYNPSLKS
	CDH19 4A9			
1128	CDR-H3 of CDH19 4A9	artificial	ΑA	NWAFHFDF
1129	CDR-L1 of	artificial	AA	TGSSSNIGTGYAVH
	CDH19 4A9			
1130	CDR-L2 of	artificial	Ą	GNNNRPS
	CDH19 4A9			
1131	CDR-L3 of CDH19 4A9	artificial	¥	QSYDSRLSGWV
1132	VH of CDH19	artificial	IN	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCAT CTCCGGCTACTTACTTAGTTCTTGGATCCGGCAACCCCTCTGGAAAGGGCCTTGGAATGGTTTCCCTTACTTTCTTACTTCCGGCTCTCACCA
	Q			CAACCC
				TAGC
1133	VH of CDH19 4A9	artificial	¥	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSS

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
N 0				
1134	VL of CDH19 4A9	artificial	LN L	CAGTCTGTGCTGACCCAGCCTCCTCTGTGTCTGGCGCCCCTGGCCAGAGAGTGACCATCTCCTGCACCGGCTCCTCCAGCAACATCTCCGGCTCCTCCTCCAGCAACATCTCGGCACCGGCTACTTCCCGGCCCCCTGCCCCCAACATCTACGGCAACAACAACAACAACAACAACAACAACAACAACAA
1135	VL of CDH19 4A9	artificial	ΑA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQFPGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSRLSGWVFGGGTKLTVL
1136	VH-VL of	artificial	M	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACGTGTCCGGGGGCGCTCCAT
	CDH19 4A9			CTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGTTCGCCTACTTCTCCTACTCCGGCTCCACCA
				ACIGETGACACCEGEGIACIACTACTACTGCCCCGGAACTGGCCCTTCCACTTCGATTTCTGGGCCCAGGCACCTGGTCACCGTGTC
				TAGCGGAGGCGGAGGATCTGGTGGTGGTTCTGGCGGCGCGCAGGCTCCCAGTCTGTGCTGACCCAGCCTCCCTC
				CCCCTGGCCAGAGAGTGACCATCTCCTGCACCGGCTCCTCCAGCAACATCGGCACCGGCTACGCCGTGCACTGGTATCAGCAGTTC
				CCCGGCACCGCCCCCAAGCTGCTGATCTACGGCAACAACAACGGCCTCCGGCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGG
				CACCTCCGCCTCCCTGGCTATCACCGGCCTGCAGGCTGAGGACGAGGCCGACTACTACTGCCAGTCCTACGACTCCCGGCTGTCCG
				GCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTG
242	VH-VL of CDH19 4A9	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGSGGGGSGGGSGGGSQSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQF PGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVL
1138	CDH19 4A9 x	artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT
	12C			AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
				PGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLSGGGGSEVQLVESGGG
				LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY
				YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTGGSTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1139	CDR-H1 of	artificial	ΑA	SYDMH
1140	╀	artificial	AA	VISYDGTNEYYADSVKG
1141	CDR-H3 of	artificial	¥	ERYFDWSFDY
	CDH19 4B10			
1142		artificial	¥	RASQSVSNTYLA
1143	CDR-L2 of CDH19 4B10	artificial	ΑA	GASSRAT
1144	CDR-L3 of	artificial	ΑA	QQYSNSWT

SFO	DESIGNATION	SOURCE	TYPF	SEOLIFINGE
∑ 0 9			 	
	CDH19 4B10			
1145	VH of CDH19 4B10	artificial	Z Z	CAGGTGCAGCTGGTGGAATCCGGCGAGGCGTGGTGGTGGCCGGCC
1146	VH of CDH19 4B10	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS
1147	VL of CDH19 4B10	artificial	TN	GAGATCGIGCTGACCCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGA
1148	VL of CDH19 4B10	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYHQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFALTISSLEPEDFAV YYCQQYSNSWTFGQGTKVEIK
1149	VH-VL of	artificial	N	CAGGIGCAGCIGGIGGAAICCGGCGGGGGGGGGGGCGTGCCTGGCCGGTCCTGAGACTGICTIGCGCCGCCTCCGGCTTCACCTT
243	CDH19 4B10		:	CTCCAGCTACGACATGCACTGGGTCCGACAGGCCCCTGGGAATGGGTGGCTGATCTCCTACGACGGCACCAACG AGTACTACGACGTGAAGGGCCCGACAGGCCCATCTCCGGGACCCTCCAAGGAACCTCCTGCAGATGAACTCCTGG CGGGCCGAGGACACCGCCGTGTACTGCGCCAGAGCGGTACTTCGACTGGTCCTTCGACTGTACTGGGGCCCAGGGCACCTGGT GTCCGTGTCTAGCGGAGGAGCTGTGGTGGTGGTTCTTGGCGGCGGGGGGGCTCCTGAGATCGTGCTGGCTG
1150	VH-VL of CDH19 4B10	artificial	ΑΑ	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGGGGETVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYH QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFALTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIK
1151	CDH19 4B10 x 12C	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYDMHWVRQAPGRGLEWVAVISYDGTNEYYADSVRGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGSGTVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYH QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFALTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1152	CDR-H1 of CDH19 4F3	artificial	ΑA	SYDMD
1153	CDR-H2 of	artificial	AA	VIWYDGSNKYYADSVRG

SEQ LI	DESIGNATION	SOURCE	IYPE	SEQUENCE
NO.				
	CDH19 4F3			
1154	CDR-H3 of CDH19 4F3	artificial	AA	ETGEGWYFDL
1155	CDR-L1 of CDH19 4F3	artificial	ΑA	RASQSVSSSYLA
1156	CDR-L2 of CDH19 4F3	artificial	ΑΑ	GASSRAT
1157	CDR-L3 of CDH19 4F3	artificial	AA	QQYGSSWT
1158	VH of CDH19	artificial	N	CAGGIGCAGCIGGIGGAAICCGGCGGAGGCGIGGIGCAGCCIGGCCGGICCCTGAGACIGICTIGIGCCGCCICCGGCTICAGCII
	4F3			CTCCTCCTACGACATGGACTGGGGTCCGACAGACCCCCGGCAAGGGCCTGGAATGGGTGGCCGTGATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTG
				CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCAGAGGCACCTGGT CACCGTGTCTAGC
1159	VH of CDH19	artificial	AA	
244	4F3			
1160	VL of CDH19	artificial	TN	GAGAICGIGCIGACCCAGICCCCIGGCACCCIGICCCIGAGCCCIGGCGAGAGAGA
	n F			
1161	VL of CDH19 4F3	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSWTFGQGTKVEIK
1162	VH-VL of	artificial	IN	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGCCTGGCCTGGCCGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT
	CDH19 4F3			CICCICCIACGACAIGGACIGGGCICGGGCACACACACCCCCGGGACAACGGCCIGGAAIGGGIGGCCGIGAIIIGGIACGACGGCICAACA AGTACTACGCCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTG
				CACCGIGICTAGCGGAGGGGGAGGAICTGGTGGGGGGGTGGTTCTGGCGGGGGGGGCTCCGAGATCGTGCTGACCCAGTCCCTGGCA CCCTGICCCTGAGCCCTGGCGAGAGAGCCACCCTGTCCTGCAGAGCCTCCCAGTCCGTGTCCTCCTACCTGGCCTGGTATCAG
				CAGAAGCCCGGCCAGGCCCCTCGGCTGATCTACGGCGCCTCTTCCAGAGCCACGGCATCCCTGACCGGTTCTCCGGCTCTGG
				CICCGGCACLICACCLIGACCAICAGCCGGCIGGAACCCGAGGACIICGCIGIGIACIAIIGCCAGCAGIACGGCICCICCI GGACCTTCGGCCAGGGCAACAAATCAAG
1163	VH-VL of	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVFDTAVYYCARFTGRGWYFDIMGRGTIVTVSSGGGGGSGGGGSGGGSFIVITOSPGTI.SISPGFRATI.SCRASOSVSSSYIAWYO
				QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIK
1164	CDH19 4F3 x	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL

SEO	DESIGNATION	SOURCE	TYPE	SEOUENCE
0 N 0.				
	12C			RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGTVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGGTDFTLTISRLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1165	CDR-H1 of CDH19 4F7	artificial	AA	SYSWS
1166	CDR-H2 of CDH19 4F7	artificial	AA	YIYYSGSTNYNPSLKS
1167	CDR-H3 of CDH19 4F7	artificial	AA	NWAFHFDY
1168	CDR-L1 of CDH19 4F7	artificial	AA	TGSSSNIGTGYDVH
1169	CDR-L2 of CDH19 4F7	artificial	AA	GNSNRPS
0211 245	CDR-L3 of CDH19 4F7	artificial	ΑA	QSYDSSLSGWV
1171	VH of CDH19 4F7	artificial	F	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCTGCTGCGGCGGCGCGCTCAT CTCCTCCTACTCTTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTTCCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCCTGGACACCTCCAAGAACCAGTTCTCCCTGAAGACCTGTCCTCCGTGACC GCCGCTGACACCCCAGCCTGTACTACTGCGCCCGGAACTGGGCCTTCCAAGAACCAGGGCCAGGGCACCTGGTCACCGTGTC TAGC
1172	VH of CDH19 4F7	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSS
1173	VL of CDH19 4F7	artificial	TN	CAGTCTGTGCTGACCCAGCCTCCCTCTGTGTCTGGCGCCCCTGGCCAGCGCGTGACCATTTCCTGCACCGGCTCCTCCAGCAACAT CGGCACCGGCTACGACGTGCACTGGTATCAGCAGCTGCCGGCCCCCCCC
1174	VL of CDH19 4F7	artificial	AA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSSLSGWVFGGGTRLTVL
1175	VH-VL of CDH19 4F7	artificial	TN	CAGGTGCAGCTGCAGGAATCCGGCCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCCACCGTGTCCGGCGCGCGC

SEQ O	DESIGNATION	SOURCE	TYPE	SEQUENCE
Ö				TAGCGGAGGGAGGATCTGGTGGCGGTGGTTCTGGCGGGGGGCTCCCAGTCTGTGCTGACCCCAGCCTCCCTC
1176	VH-VL of CDH19 4F7	artificial	AA	
1177	CDH19 4F7 x 12C	artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSSGGGSGGGGSGGGGSQSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQL PGTAPKLLIHGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLSGGGGSEVQLVESGGG LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1178	CDR-H1 of CDH19 14039	artificial	ΑA	SYGMH
6211 246	CDR-H2 of CDH19 14039	artificial	AA	FIWYEGSNKYYAESVKD
1180	CDR-H3 of CDH19 14039	artificial	AA	RAGIIGYIYGMDV
1181	CDR-L1 of CDH19 14039	artificial	ΑΑ	SGDRIGEKYTS
1182	CDR-L2 of CDH19 14039	artificial	AA	QDTKRPS
1183	CDR-L3 of CDH19 14039	artificial	AA	QAWESSTVV
1184	VH of CDH19 14039	artificial	Z	CAGGTGCAGTTGGTGGAGTGTGGGGGGAGGCGTGGTCCTGGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGAGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTTGGAGGGAG
1185	VH of CDH19 14039	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1186	VL of CDH19	artificial	NT	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	14039			CTGAGCGATICTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTAC TGTCAGGCGTGGGAGAGCAGCACTGTGGTATTCGGCGGGGACCAAGCTGACCGTCCTA
1187	VL of CDH19 14039	artificial	ΑA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
1188	VH-VL of	artificial	N	CAGGTGCAGTTGGTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCTCTGTGCAGCGTCTGGGATTCACCTT
	CDH19 14039			CAGTAGCTATGCCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
				AGAGCTGAGGACACGCCTGTGTATTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACGGTATGGACGTCTG
				GGGCCAAGGGACCACGGTCACCGTCTCTAGTGGTGGTGGCGGAGGATCTGGCGGAGGTGGAAGCGGAGGCGGCGGCGGTCTTCCTATGAAC TGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
				AGCTGGTATCAGCAGAGGCCAGGCCCAGTCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCCTGAGCGATT
				CTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTACTGTCAGGCGT GGGAGAGCAGCACTGTGGTATTCGGCGGAGGGACCAAGCTGACCGTCCTA
1189	VH-VL of	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL
	CDH19 14039			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
24	0007	3		DWINGTON CONTROL FOR FORE FORENCE OF THE STATE OF THE STA
1190	CDH19 14039	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGRGLEWVAFTWYEGSNRYYAESVKDRFTISRDNSKNTLYLQMNSL babbaaaascabbacticmicvvvcmbaaaccabcabcabcabcabcabcabaaaaaaaaaa
	x 17C			RAEDIAVIICARRAGIIGIIGIIGIIIGMDVWGQGIIVIVOOGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1191	CDR-H1 of	artificial	ΑA	SYGMH
	CDH19 14304			
1192	CDR-H2 of	artificial	ΑA	FIWYDGSNKYYADSVKD
1193	CDR-H3 of	artificial	Ą	RAGIIGYYYGMDV
	CDH19 14304			
1194	CDR-L1 of	artificial	AA	SGDRLGEKYVS
	CDH19 14304			
1195	CDR-L2 of	artificial	₩	QDNKWPS
,	CD1129 14504			A STATE OF THE STA
1196	CDR-L3 of CDH19 14304	artificial	AA	QAWDSSTVV
1197	VH of CDH19	artificial	N	CAGGTGCAGTTGGTGGAGTCTGGGGGGAGGCGTGGTCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> №				
	14304			CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGGGCTGGAGTGGGTGG
1198	VH of CDH19 14304	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1199	VL of CDH19 14304	artificial	Z	TCCTATGAGCTGACTCAGCCCACCCTCAGTGTCCCCCAGGACAGACA
1200	VL of CDH19 14304	artificial	ΑA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
248	VH-VL of CDH19 14304	artificial	F	CAGGTGCAGTTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCTCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCAGGCAGG
1202	VH-VL of CDH19 14304	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVL
1203	CDH19 14304 x I2C	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
1204	CDR-H1 of CDH19 14301	artificial	AA	SYGMH
1205	CDR-H2 of CDH19 14301	artificial	ΑA	FIWYDGSNKYYADSVKD
1206	CDR-H3 of	artificial	AA	RAGIIGTIGYYYGMDV

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
© 8				
	CDH19 14301			
1207	CDR-L1 of CDH19 14301	artificial	AA	SGDRIGERYTC
1208	CDR-L2 of CDH19 14301	artificial	ΑA	QDTKRPS
1209	CDR-L3 of CDH19 14301	artificial	ΑΑ	QAWESSTVV
1210	VH of CDH19 14301	artificial	F	CAGGTGCAGTTGGTGGAGTCTGGGGGGGGGTCCCAGGCGTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGAGTGGTGGTGGTATGATGAATA AATACTATGCAGACTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAAAAAAAA
1211	VH of CDH19 14301	artificial	AA A	QVQLVESGGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
249	VL of CDH19 14301	artificial	LN .	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
1213	VL of CDH19 14301	artificial	ΑΑ	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTCWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
1214	VH-VL of CDH19 14301	artificial	LN .	CAGGTGCAGTTGGTGGAGTCTGGGGGGGGCGTGGTCCAGGCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGCTGGC
1215	VH-VL of CDH19 14301	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT CWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVL
1216	CDH19 14301 x I2C	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT CWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGSEVQLVE

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
o N S				
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVYTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1217	CDR-H1 of CDH19 14079	artificial	AA	RYGIH
1218	CDR-H2 of CDH19 14079	artificial	AA	VIWYEGSNKYYAESVKG
1219	CDR-H3 of CDH19 14079	artificial	AA	RAGIPGTTGYYYGMDV
1220	CDR-L1 of CDH19 14079	artificial	AA	SGDRLGEKYVS
1221	CDR-L2 of CDH19 14079	artificial	AA	QDNKWPS
1222	CDR-L3 of CDH19 14079	artificial	AA	QAWESSTVV
1223	VH of CDH19	artificial	Ν	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGGGGTCCTGGGGGGGTCCCTGAGACTCTCTGTGCAGCGTCTGGAATTCACCTT
	14079			CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTTGGAGTGGCTGGC
1224	VH of CDH19 14079	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
1225	VL of CDH19 14079	artificial	LN .	TCCTATGAGCTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
1226	VL of CDH19 14079	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
1227	VH-VL of CDH19 14079	artificial	Z	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGCGTGGTCCAGGCGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTTGCTATGGCATACACTGGGTCCGCCAGGCTAGGGGGGGG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
				AGCTGGTATCAGCAGAAGCCAGGCCAGTCCCCTATACTGGTCATCTATCAAGATAATAAGTGGCCCTCAGGGATCCCTGAGCGATT CTCTGGCTCCAACTCTGGGAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTACTGTCAGGCGT GGGAGAGCAGCAGCATGTGGGCGGGGGGGACCAAGCTGACCGTCCTA
1228	VH-VL of CDH19 14079	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVL
1229	CDH19 14079 × I2C	artificial		
1230	CDR-H1 of CDH19 14077	artificial	AA	RYGIH
1231	CDR-H2 of CDH19 14077	artificial	AA	VIWYDGSNKYYADSVKG
1232	CDR-H3 of CDH19 14077	artificial	AA	RAGIPGTTGYYYGMDV
1233	CDR-L1 of CDH19 14077	artificial	AA	SGDRIGEKYVS
1234	CDR-L2 of CDH19 14077	artificial	AA	QDNKWPS
1235	CDR-L3 of CDH19 14077	artificial	AA	QAWDSSTVV
1236	VH of CDH19 14077	artificial	L	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGGCTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAGG
1237	VH of CDH19 14077	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
1238	VL of CDH19 14077	artificial	L	TCCTATGAGCTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
1239	VL of CDH19 14077	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
1240	VH-VL of CDH19 14077	artificial	L Z	CAGGTGCAGCTGGTGGAGGTCTGGGGGGGGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTTT CAGTCGCTATGGCATACACTGGGTCCGCCAGGCAGGCCAGGGGCTGGAGTGGTATATAGTATGGTATGATGAAGTAATA AATACTATGCAGACTCCGTGAAGGGCCCATCTCCAGGCAAGACAATTCCAAGAACACGCTGTATTGCTAATGGAATGAACAGCCTG AGAGCCGAGGACTCCGTGTATTACTGTGCGAGAAGGGCCGGTATACCTGGAACTACGGGCTACTATTGGTATTGGACGTTG GGGCCAAGGGACTCGGCTGTTACTGTGCGAGAAGGGCCGGTATACCTGGAACTACGGGCTACTATTGGTATTGACTCTGCCGAGGACTCTGCCGAGGCCGAGGCCGCGAGCCTATTGTTGTT AGACTCAGCCAAGGGACCCACGTCCCCTCAGGACACACAC
1241	VH-VL of CDH19 14077	artificial	АА	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVL
252	CDH19 14077 x I2C	artificial		QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1243	CDR-H1 of CDH19 14096	artificial	AA	HIXXS
1244	CDR-H2 of CDH19 14096	artificial	AA	IINPSGGSTRYAQKFQG
1245	CDR-H3 of CDH19 14096	artificial	АА	GGIQLWLHFDY
1246	CDR-L1 of CDH19 14096	artificial	AA	SGSSSNIGRNFVN
1247	CDR-L2 of CDH19 14096	artificial	AA	TNNQRPS
1248	CDR-L3 of CDH19 14096	artificial	АА	AAWDESLNGWV
1249	VH of CDH19 14096	artificial	LN .	CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGCAAGGCATCTGGATACACCTT CACCAGCTACTATTTCACTGGGTGCGCCAGGCCCTGGACAAGGACTTGAGTGGAATAATCAACCCCAGTGGTGGTAGCA CAAGGTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACATGAAGTTGAGCTGAGCAGCCTG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
© .				
				AGATCTGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGAATACAGCTATGGTTTTGACTACTGGGGCCCAGGGAACCCT GGTCACCGTCTCCTCA
1250	VH of CDH19 14096	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
1251	VL of CDH19 14096	artificial	Ä	CAGTOTGCGCTGACTCAGCCACCCTCAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTTGTTCTGGAAGCAGCTCCAACAT CGGAAGGAATTTTGTAAACTGGTATCAGCAGCTCCCAGGAACGGCCCCCAAAGTCCTCATTTATATATA
1252	VL of CDH19	artificial	AA	QSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAAWDESLNGWVFGGGTKLTVL
1253	VH-VL of	artificial	F	CAGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCCTCAGTGAAGGTTTCCTGCAAGGCATCTGGATACACCTT CACCAGCTAATTCACTGTGGGTGAAGGTGAAGGTGAAGGCCTTGAAGGTTTCCTGCAAGGCTTCTGGATACACCTT
	CDH19 14096			CANCERCIACIAIAIICACIGGGGGGGGGGGGGGGGGGGGGGGGG
253				CAGCGACTGGGACCCCCGGGCAGAGGGTCACCATCTTGTTCTGGAAGCAGCTCCAACATCGGAAGGAA
1254	VH-VL of CDH19 14096	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAAWDESLNGWVFGGGTKLTVL
1255	CDH19 14096 x I2C	artificial		QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYIHWVRQAPGQGLEWMGIINPSGGSTRYAQKFQGRVTMTRDTSTSTVYMELSSI RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGGSQALTQPPSATGTPGQRVTISCSGSSSNIGRNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAAWDESLNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSIKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLMYSNRWVFGGGTKLTVLHHHHHH
1256	CDR-H1 of CDH19 14088	artificial	AA	SYYMS
1257	CDR-H2 of CDH19 14088	artificial	AA	IIHPSGGDTTYAQKFQG
1258	CDR-H3 of CDH19 14088	artificial	AA	GGIKLWLHFDY
1259	CDR-L1 of	artificial	AA	SGSRSNIGSNFVN

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	CDH19 14088			
1260	CDR-L2 of CDH19 14088	artificial	AA	TNNQRPS
1261	CDR-L3 of CDH19 14088	artificial	ΑA	AVWDDSLNGWV
1262	VH of CDH19 14088	artificial	TN	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGCCTCAGTGAAGGTTTCCTGTAAGGCATCTGGATACACCTTT CACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGGCTTGAGGGATTTCCTGTAAGGCATCTGGACAA CAACCTACGCACAGAATTATGTCCTGGGTGCACAGGCCCTTGACGGGCTTGAGGGATGGAATAATCCACCTTAGTGGTGACA CAACCTACGCACAGAAGTTCCAGGGCCAGAGTCACCAGGGACACGTCCACGAGCACAGTCTACAGGAGCTGAGCCTG AGATCTGAGGACACGCCGTGTATTACTGTGCGAGAGGGGGATAAAACTATGGTTACATTTGACTATTGGGGCCAGGGAACCCT GGTCACCGTCTCCTCA
1263	VH of CDH19 14088	artificial	ΑΑ	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS
1264	VL of CDH19 14088	artificial	F	CAGTCTGTGCTGACTCAGCCACCCTCAGCGTCTGGGACCCCCGGACAGGGTCACCATCTTTTTTTT
1265	VL of CDH19 14088	artificial	AA	QSVLIQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDDSLNGWVFGGGTKLTVL
1266	VH-VL of CDH19 14088	artificial	F	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGGCCTCAGTGAAGGTTTCCTGTAAGGCATCTGGATACACCTTT CACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAGGGCTTGAGTGGAATAATCCACCCTAGTGGTGACA CAACCTACGCACAGAAGTTCCAGGGCCAGAGTCACCATGGAGCACAGTCTACATGGAGCTGAGCGGCCTG AGATCTGAGGACACGCCGTGTATTACTGTGCGAGAGGGGGGATAAAACTATGGTTACATTTTGACTATTGGGGCCAGGGGAACCCT GGTCACCGTCTCCTCAGGTGGCGGAGGAGGGGGGGGGAGGCGGCGGCGGCGGGAGCCGT CAGCGTCTCGTCCTCAGGTGGCGGAGGAGTCTTTTTTTTT
1267	VH-VL of CDH19 14088	artificial	ΑA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVL
1268	CDH19 14088 × I2C	artificial		QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDDSLNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGCYVTVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9				
				QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1269	CDR-H1 of CDH19 14085	artificial	AA	SYYMS
1270	CDR-H2 of	artificial	AA	IIHPSGGDTTYAQKFQG
	CDH19 14085			
1271	CDR-H3 of	artificial	¥	GGIKLWIHFDY
1	CDH19 14085			THE TANK THE TOPON
12/2	CDK-L1 of CDH19 14085	artificial	¥	OG OK ON LGONF VN
1273	CDR-L2 of	artificial	AA	TNNQRPS
	CDH19 14085			
1274	CDR-L3 of CDH19 14085	artificial	Ą	AVYDDSLNGWV
1275	VH of CDH19	artificial	N	CAGGTGCAGTTGGTGCAGTCTGGGGGCTGAGGTGAAGAAGAAGCCTGGGGGCCTCAGTGAAGGTTTCCTGTAAGGCATCTAGATACACCTT
	14085			CACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGGCTTGAGTGGATGGGAATAATCCACCCTAGTGGTGGTGACA
:55				CAACCIACGCACAGAAGTICCAGGGCAGAGTCACCAIGAGCATCACCGGGGACACGTCCACGAGCACACATCIACAIGGAGCTGAGCAGCCTG
				AGATOTGAGGACACGCCG1G1A11AC1G1GCGAGGGGGGGATAAAAC1A1GG11ACA1111GAC1A1GGGCCAGGGAACCC1 GGTCACCGTCTCCTCA
1276	VH of CDH19	artificial	ΑA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL PSFPTAVVVCAPCCIKIM:HFDVWGCCTIVIVOSS
	14000			
1277	VL of CDH19 14085	artificial	Z	CAGTCTGTGCTGACTCAGCCACCCTCAGCGTCTGGGACCCCCGGACAGGGTCACCATCTCTTGTTCTGGAAGCCGCTCCAACAT CGGAAGTAATTTTGTAAACTGGTACCAGCAGCTCCCAGGAACGGCCCCCAAACTCCTCTTTTTTTAATAATCAGCGGCCCTCAG
				GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCCAGGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGAT TATTACTGTGCAGTATACGATGACAGCCTGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCTCTA
1278	VL of CDH19 14085	artificial	AA	QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDDSLNGWVFGGGTKLTVL
1279	VH-VL of	artificial	F	CAGGTGCAGTTGGTGCAGTCTGGGGGCTGAGGTGAAGAAGAAGCCTGGGGGCCTCAGTGAAGGTTTCCTGTAAGGCATCTAGATACACCTT
	CDH19 14085			CACCAGCTACTATATGTCCTGGGTGCGACAGGCCCCTGGACAAGGGCTTGAGTGGATGGGAATAATCCACCCTAGTGGTGGTGACA
				CAACCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCGGGGACACGTCCACGACACACAGTCTACATGGAGCTGAGCAGCTGACCACGCTGA
				AGAICIGAGGACACGCCGIGIAIIACIGIGCGAGGGGGGGAIAAAACIAIGGIIACAIIIIGGGGCAAAGGAAGG
				CAGCGTCTGGGACCCCCGGACAGAGGGTCACCATCTCTTGTTCTGGAAGCCGCTCCAACATCGGAAGTAATTTTGTAAACTGGTAC
				CAGCAGCICCCAGGAACGGCCCCCAAACICCICAICIAIACIAAIAAICAGCGGCCCICAGGGGGICCCIGAACGAITCICIGGCIC Caacittiggcalciticagaacgcaattaataattagaattaaiaaagaattaataataaataa
				[

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>.</u> 8				
				GCCTGAATGGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCGTCCTA
1280	VH-VL of CDH19 14085	artificial	АА	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
1281	CDH19 14085 x l2C	artificial		QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEMMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAVYYCARGGIKLMIHFDYWGQGTLVTVSSGGGSGGGSGGGSGGSGSCVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY GGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNPGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGGGSGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKITVLHHHHHH
1282	CDR-H1 of CDH19 14074	artificial	AA	SYFIH
1283	CDR-H2 of CDH19 14074	artificial	AA A	IINPISVSTSYAQKFQG
1284	CDR-H3 of CDH19 14074	artificial	AA	GGIQLWLHLDY
90 1285	CDR-L1 of CDH19 14074	artificial	AA	SGSRSNIGSNFVN
1286	CDR-L2 of CDH19 14074	artificial	AA	TNNQRPS
1287	CDR-L3 of CDH19 14074	artificial	ΑA	ATWDDSMNGWV
1288	VH of CDH19 14074	artificial	L	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGACCTGGGGGCCTCAGTGAAGGTTTCCTGCAAGGTTTCTGGATACACCTT CACCAGCTACTTTATTCACTGGGTGCCCAGGCCCCTGGACAAGGCCTTGAATGGATGG
1289	VH of CDH19 14074	artificial	¥¥	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
1290	VL of CDH19 14074	artificial	TN	CAGTCTGCGCTGACTCAGCCACCCTCAGTGACTGGGACCCCCGGGCAGAGGGTCACCATCTTGTTCTGGAAGCAGGTCCAACAT CGGAAGCAATTTTGTAAAACTGGTACCAGCAGCTCCCAGGAACGGCCCCCCAAAGTCCTCATCTATACTAATAATCAGCGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGAT TATTACTGCCCAACATGGGATGAATGAATGGTTGGGTGTTCGGCGGAGGGACCAAACTGACCGTCCTA
1291	VL of CDH19	artificial	AA	QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATWDDSMNGWVFGGGTKLTVL

CH	DESIGNATION	SOLIBOE	TVDE	CEOI IENCE
7 4		2000		
<u>2</u> 8				
	14074			
1292	VH-VL of	artificial	N	CAGGTGCAGTTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCCTGGGGGCCTCAGTGAAGGTTTCCTGCAAGGTTTCTGGATACACCTT
	CDH19 14074			CACCAGCTACTTTATTCACTGGGTGCGCCCAGGCCCCTGGACAAGGGCTTGAATGGATGG
				CAAGCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACAGTCTTCATGGAGCTGAGCAGCCTG
				AGATCTGAGGACACGCCGTGTATTACTGTGCGCGAGGGGGATACAGCTATGGTTACATTTGGACTACTGGGGCCAGGGAACCCT Ggataaccataacaacaagaagaagaagaacaagaagaagaagaagaaga
				CAGCAGCICCCAGGAACGGCCCCCAAAGICCTCATATAATAATCAGCGGCCCTCAGGGGGICCCTGACCAAITCTCTGGCTC
				CAAGICIGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGATTATTACTGCGCAACATGGGATGACA
1203	VH-VI of	artificial	۸۸	GIAIGAAIGGIIGGGIGIICGGCGGAGGGACCAAACIGACCGICCIA OVOIVOSGAFVKKPGASVKVSCKVSGYTFTSYFTHWVROAPGOGIEWMGIINPISVAOKFOGRVTMTRDTSTSTVFMFISST.
1455	CDH19 14074		{	RSEDTAVYCARGGIQLWLHLDYWGQGTLVTVSSGGGGGGGGGGGGGQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY
				QQLPGTAPKVL1YTNNQRPSGVPDRFSGSKSGTSASLA1SGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVL
1294	CDH19 14074	artificial		QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL
	x I2C			RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
				QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLSGGGGSEVQLVES
57				GGGLVQPGGSLKLSCAASGFTFINKYAMNWVKQAFGKGLEWVAKLKSKYNNYATYYADSVKDKFTLSKDDSKNTAYLQMINNLKTEDT
				AVIICVRHGNFGNSIISIWAIWGQGTLVIVSSGGGGGGGGGGGGGGGGGTVTQLFSLIVSFGGTVILTGGSSTGAVISGNIFNWVQ QRPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1295	CDR-H1 of	artificial	ΑA	SYFIH
	CDH19 14075			
1296	CDR-H2 of	artificial	AA	IINPISVSTSYAQKFQG
	CDH19 14075			
1297	CDR-H3 of	artificial	AA	ССІОТИТНІ Т
,	CDH19 140/5		:	ATT AMERICAN CONTRACTOR OF THE PROPERTY OF THE
1798	CDK-L1 of	artificial	¥	OGDKONI GONF VN
1299	CDR-12 of	artificial	۵۵	TINNORPS
14.0	CDH19 14075	5	{	
1300	CDR-L3 of	artificial	AA	ATWDESMQGWV
	CDH19 14075			
1301	VH of CDH19	artificial	nt	
	14075			
				CAAGCIACGCACACAGAAGITCCAGGGCAGAGICACCAIGACCAGGGACACGICCACGAGGCACTATTGAAGITCTICAIGGAGGTGGAAAGCTGAGCAGCCIG AGAICTTGAGGACACGGCCGTGTATTACTGAGGGGAGAGAGAGA

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u> </u>				
				GGTCACCGTCTCCTCA
1302	VH of CDH19 14075	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
1303	VL of CDH19 14075	artificial	nt	CAGTOTGCGCTGACTCAGCCACCCTCAGTGACTGGGACCCCCGGGCAGGGGTCACCATCTTGTTCTGGAAGCAGGTCCAACAT CGGAAGCAATTTTGTAAACTGGTACCAGCAGCTCCCAGGAACGGCCCCCAAAGTCCTCATCTATACTAATAATCAGCGCCCTCAG GGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCAGTGGGCTCCAGTCTGAGGATGAGGCTGAA TATTACTGCGCAACATGGGATGAGGATATGCAGGCTTGGGTGTTCGGCGAGGGACCAAACTGACCGTCCTA
1304	VL of CDH19 14075	artificial	ΑA	QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATWDESMQGWVFGGGTKLTVL
1305	VH-VL of CDH19 14075	artificial	nt	CAGGIGCAGITGGIGCAGICTGGGGCCTGAAGGTGAAGCCTGGGGCCCTCAGTGAAGGITTCCTGCAAGGITTCTGGATACACCTT CACCAGCTACTTTATTCACTGGGTGCGCCCAGGCCCTGGACAAGGGCTTGAATGGATGG
				CAAGCTACGCACAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACGTCCACGAGCACACATTTCATGGAGCTGAGCAGCCTG AGATCTGAGGACACGGCCGTGTATTACTGTGCGCGAGGGGGATACAGCTATGGTTACATTTGGACTACTGGGGCCAGGGAACCCT GGTCACCGTCTCCTCAGGTGGCGGAGGATCTGGCGGAGGGGAAGCGGAGGGGGGGG
258				CAGCAGCTCCCAGGAACGGCCCCCCAAGTCCTCTTTATTATTATCAGCGGCCCTCAGGGGTCCCTGACCATTCTTTTTTTT
1306	VH-VL of CDH19 14075	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVL
1307	CDH19 14075 x I2C	artificial		QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSGGGSGSALTQPPSVTGTPGQRVTFGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1308	CDR-H1 of CDH19 14024	artificial	ΑA	SSGYY
1309	CDR-H2 of CDH19 14024	artificial	AA	YIYYTGSAYYNPSLKS
1310	CDR-H3 of CDH19 14024	artificial	AA	DGSSGWYFQY
1311	CDR-L1 of CDH19 14024	artificial	AA	RASRQISSSYLA

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
1312	CDR-L2 of CDH19 14024	artificial	АА	GPSSRAT
1313	CDR-L3 of CDH19 14024	artificial	AA	QQYGSSFT
1314	VH of CDH19	artificial	nt	CAGGIGCAGCIGCAGGAGICGGGCCCCAGGACIGGIGAAGCCIICAGAGACCCIGICCCICACCIGCACIGICICIGGIGGCICCAI
	14024			CAGCAGTAGTGGTTACTACTGGAGCTGGATCCGCCAGCCCCCAGGGAAGGGCCTGGAGTGGGTTGGGTACATCTATTACACTGGGAAGCGCTACTAGTAGAGCTGGGAAGCCCTACTACAACAACCCGTCCCTGAAGAGTCGAGTTACCATATCAGTAGAACGTCTTAGAGAACCTGAAGCTGAGCTCTGAGAAGCTGAGCTCTGTACCAGTACTACCAGTATTACTGTGCAAGCTGAAGCAGTGCTGGTACTTCCAGTATTGGGGCCAGGGCACCCTGAACCCCTGAACCCCTTACCTCTACTTCCAGTATTGGGGCCAGGGCACCCTACTCCACTACTTCCAGTATTGGGGCCAGGGCCACCCTACTCCACTCTAGTAACTACTAAAAAAAA
1315	VH of CDH19 14024	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS
1316	VL of CDH19	artificial	nt	GAAATTGTGTTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCTGCAGGGCCAGTCGGCAGAT
	14024			TAGCAGCAGCTACTTAGCCTGSTACCAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCATCTATGGTCCATCAGCAGGGCACTG GCATCCCAGACAGGTTCAGTGGCAGTGGGTCTGGGACAGACTTCACTCTCACCTCAGCAGACTGGAAGACTGAAGATTTTGCAGTG TATTACTGTCAGCAGTATGGTAGCTCATTCACTTTCGGCCAGGGAACAAAGTGGATATCAAA
259	VL of CDH19 14024	artificial	Ą	EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSFTFGQGTKVDIK
1318	VH-VL of	artificial	nt	CAGGIGCAGCIGCAGGAGICGGGCCCAGGACTGGIGAAGCCIICAGAGACCCIGICCCTCACCIGCACTGICICTGGIGGCTCCAT
	CDH19			CAGCAGTAGTGGTTACTACTGGAGCTGGATCCGCCAGCCCCCAGGGAAGGGCCTGGAGTGGGTTGGGTACATCTATTACACTGGGA GCGCCTACTACAACCCGTCCCTCAAGAGTCGAGTTACCATATCAGTAGACACGTCTAAGAACCAGTTCTCCCTGAAGCTGAGCGTCT
				GTGACTGCCGCGCACACGCCCTGTATTACTGTGCGAGATGGAAGCAGTGCTGGTACTTCCAGTATTGGGGCCCAGGGCACCTT
				GGTCAUGGTUTUTAGTGGTGGTGGAGGATUTGGGGGGGGGG
				CAGCAGAAACCTGGCCAGGCTCCCAGGCTCCTCTATGGTCCATCCA
				TGGGTCTGGGGACAGACTTCACTCTCACCATCAGCAGACTGGAGCCTGAAGATTTTGCAGTGTATTACTGTCAGCAGTATGGTAGCT CATTCACTTTCGGCCAGGGGACCAAAGTGGATATCAAA
1319	VH-VL of	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS
	CDH19 14024			VIAADIAVYYCARDGSSGWY£QYWGQGTLVTVSSGGGGSGGGGGGSEIVLTQSPGTLSLSBYGERATLSCRASRQISSSYLAWY QQRPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIK
1320	CDH19 14024	artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS
	× 12C			VTAADIAVYYCARDGSSGWYFQYWGQGTLVTVSSGGGGSGGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWY Ookbeoapriitygpssratgipdrfsgsgsggturtitasissbedyvoooygssftfgogtkydiksgggsfydiveggg
				LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY
				YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
N .				
1321	CDR-H1 of CDH19 14054	artificial	AA	SYDMH
1322	CDR-H2 of CDH19 14054	artificial	¥	VISYDGTNEYYADSVKG
1323	CDR-H3 of CDH19 14054	artificial	Ą	ERYFDWSFDY
1324	CDR-L1 of CDH19 14054	artificial	ΑΑ	RASQSVSNTYLA
1325	CDR-L2 of CDH19 14054	artificial	ΑA	GASSRAT
1326	CDR-L3 of CDH19 14054	artificial	¥	QQYSNSWT
260	VH of CDH19 14054	artificial	nt	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCACCTT CAGTAGCTATGACATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGGTGGG
1328	VH of CDH19 14054	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS
1329	VL of CDH19 14054	artificial	Ħ	GAAATTGTATTGACGCAGTCTCCAGGCACCCTGTCTTTGTCTCCAGGGGAAAGAGCCACCCTCTCCTGCAGGGCCAGTCAGAGTGT TAGCAACACCTTACTTAGCCTGGTACCAGCAGACCTGGCCTCCCAGGCTCCTCATCTATGGTGCATCCAGGGCCACTG GCATCCCAGACAGATTCAGTGGCAGTGGGTCTGGGACAGACTTCACTCTCACCATCAGCAGTCTGGAGCCTGAAGATTTTGCAGTG TATTACTGTCAGCAGTACAGTA
1330	VL of CDH19 14054	artificial	¥	EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYSNSWTFGQGTKVEIK
1331	VH-VL of CDH19 14054	artificial	nt	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCACCTT CAGGTGCAGCTGGTGGAGTCCGCCAGGCTCCAGGCAGGGGTGGGT

SFO	DESIGNATION	SOURCE	TYPE	SFOUENCE
Ω NO.				
1332	VH-VL of CDH19 14054	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIK
1333	CDH19 14054 x I2C	artificial		QVQLVESGGGVVQPGGSLRLSCAASGETFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALITLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1334	CDR-H1 of CDH19 14056	artificial	AA	GYYWS
1335	CDR-H2 of CDH19 14056	artificial	AA	YFSYSGSTNYNPSLKS
1336	CDR-H3 of CDH19 14056	artificial	ΑA	NWAEHEDF
1337	CDR-L1 of CDH19 14056	artificial	AA	TGSSSNIGTGYAVH
1338	CDR-L2 of CDH19 14056	artificial	Ą	GNNNRPS
1339	CDR-L3 of CDH19 14056	artificial	ΑĄ	QSYDSRLSGWV
1340	VH of CDH19 14056	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCTCACCTGCACTGTCTCTGGTGGCTCAT CAGTGGTTACTACTGCAGGATCCGGCAGCCCCCAGGAAAGGGACTGGAGTGGTTTGCATATTTCTCTTACAGTGGAGCACCA ACTACAACCCCTCCTCAAGAGTCGAGTC
1341	VH of CDH19 14056	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSS
1342	VL of CDH19 14056	artificial	nt	CAGTCTGTGCTGACGCCGCCCTCAGTGTCTGGGGCCCCAGGACAGAGGGTCACCATCTCCTGCACTGGGAGCTCCAACAT CGGGACAGGTTATGCTGTACACTGGTACCAGCAGCTTCCAGGAACAGCCCCCAAACTCCTCATCTATGGTAACAACAATCGGCCCT CAGGGGTTCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCACTGGGCTGAGGATGAGGCT GATTATTACTGCCAGTCCTATGACAGCAGATGAGTGGTTGGGTTTCGGCGGAGGGACCAAGCTGACCGTCCTA
1343	VL of CDH19 14056	artificial	AA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSRLSGWVFGGGTKLTVL

₩ 0	SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
ΙŹ	NO.				
 ii	1344	VH-VL of CDH19 14056	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGTCTCTGGTGGTCGTCGTCCTCTACAGTGGTGGTCCTTACAGTGGTGGTCCTTACAGTGGTGGTGGTTACTACTAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT
∺	1345	VH-VL of CDH19 14056	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
262	1346	CDH19 14056 x I2C	artificial		QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWFAYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGSGGGGSGGGGSQSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQL PGTAPKLLIYGNNNRPPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVLSGGGGSEVQLVESGGG LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
H	1347	CDR-H1 of CDH19 14057	artificial	AA	GYYWS
Ħ	1348	CDR-H2 of CDH19 14057	artificial	AA	YFSYSGSTNYNPSLKS
Ι Π	1349	CDR-H3 of CDH19 14057	artificial	AA	NWAFHFDF
H	1350	CDR-L1 of CDH19 14057	artificial	AA	TGSSSNIGTGYAVH
∺	1351	CDR-L2 of CDH19 14057	artificial	AA	GNNNRPS
H	1352	CDR-L3 of CDH19 14057	artificial	AA	QSYDSRLSGWV
Ħ	1353	VH of CDH19 14057	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGCCTG

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO. 1354	VH of CDH19 14057	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSS
1355	VL of CDH19 14057	artificial	nt	CAGTOTGTGOTGACGCCGCCCTCAGTGTCTGGGGCCCCCAGGACAGGGTCACCATCTCCTGCACTGGGAGCAGCTCCAACAT CGGGACAGGTTATGCTGTACACTGGTACCAGCAGCTTCCAGGAACAGCCCCCAAACTCCTCATGGTAACAACAATCGGCCCT CAGGGGTTCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCTGGCCCATCATGGGCTCAGGCTGAGGATGAGGCT GATTATTATTATTACTGCCAGTCCTATGACAGACTGAGTTGGGTGTTCGGCGGAGGGACCAAGCTGACCTGA
1356	VL of CDH19 14057	artificial	ЧΑ	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQLPGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSRLSGWVFGGGTKLTVL
263	VH-VL of CDH19 14057	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGCACCTGTCTCTGGTGGCTCCAT CAGTGGTTACTACTACAGAGCTGGATCGGCAGCCCCCAGGAAAGGGACTGGAGTGGATTGGATATTTCTCTTACAGTGGGAGCACCA ACTACAACCCCTCCTCAAGAGTCGAGTC
1358	VH-VL of CDH19 14057	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQL PGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSRLSGWVFGGGTKLTVL
1359	CDH19 14057 x I2C	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYFSYSGSTNYNPSLKSRVTLSVDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDFWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYAVHWYQQL PGTAPKLLIYGNNNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSRLSGWYFGGGTKLTVLSGGGGSEVQLVESGGG LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1360	CDR-H1 of CDH19 14049	artificial	ΑΑ	SYSWS
1361	CDR-H2 of CDH19 14049	artificial	AA	YIYYSGSTNYNPSLKS
1362	CDR-H3 of CDH19 14049	artificial	AA	NWAFHFDY
1363	CDR-L1 of CDH19 14049	artificial	AA	TGSSSNIGTGYDVH
1364	CDR-L2 of	artificial	ΑA	GNSNRPS

CEO	DESIGNATION	COLIBCE	TVDE	CECI IENCE
7 5		30015		פרעייי
<u>.</u> 8				
	CDH19 14049			
1365	CDR-L3 of CDH19 14049	artificial	AA	QSYDSSLSGWV
1366	VH of CDH19 14049	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGCACCTGTCTCTGGTGGCTCCAT CAGTAGTTACTCCTGGAGCTGGAGCCCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATATTACAGTGGAGCACCA ACTACAACCCCTCCACCAAGAGTCGAGTC
1367	VH of CDH19 14049	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSS
1368	VL of CDH19 14049	artificial	Ħ	CAGTCTGTGCTGACGCCGCCCCCCCAGTGTCTGGGGCCCCCAGGGCCACCACCATCTCCTGCACTGGGAGCAGCTCCAATAT CGGGACAGGTTATGATGTACACTGGTATCAGCAGCTTCCAGGAACACCCCCAAACTCCTCTTATGGTAACAGCAATCGGCCCT CAGGGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCATCACTGGGCTCCAGGCTGAGGATGAGGCT GATTATTACTGCCAGTCCTATGACAGCAGTCTGAGTTGGGTTGGGTTCGGCGAGGGACCAGGTTGACTCAGCTCTA
1369	VL of CDH19 14049	artificial	AA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSSLSGWVFGGGTRLTVL
1371	VH-VL of CDH19 14049 CDH19 14049 x I2C	artificial artificial	A A A	CAGGRACAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
1373	CDR-H1 of	artificial	AA	SYGMH

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
S 6				
	CDH19 14302			
1374	CDR-H2 of CDH19 14302	artificial	AA	FIWYDGSNKYYADSVKD
1375	CDR-H3 of CDH19 14302	artificial	AA	RAGIIGTIGYYYGMDV
1376	CDR-L1 of	artificial	AA	SGDRLGEKYTS
	CDH19 14302			
1377	CDR-L2 of CDH19 14302	artificial	ΑA	QDTKRPS
1378	CDR-L3 of CDH19 14302	artificial	AA	QAWESSTVV
1379	VH of CDH19	artificial	ţ	<u>であららずららならですららならなででもららららならないらずららずららなららららららららららららないできならないでででです。</u>
	14302	5	<u>:</u>	CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCCAGGGGGCTGGGGTGGGT
6.				GGGCCAAGGGACCACGGTCACCGTCTCTAGT
1380	VH of CDH19 14302	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1381	VL of CDH19	artificial	пţ	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
	14302			AAAATATACTAGCTGGTATCAGCAGAGGCCAGGCCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC CTGAGCGATTCTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGATTAC TGTCAGGCGTGGGAGGAGCAGCAGTAGTGGTAATTCGGCGGAGGAAGGGAACTTGACCGTCCTA
1382	VL of CDH19	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
	14302			CQAWESSTVVFGGGTKLTVL
1383	VH-VL of CDH19 14302	artificial	nt	CAGGTGCAGTTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCCAG
				GGGAGAGCAGCACTGTGGTATTCGGCGGAGGACCAAGCTGACCGTCCTA
1384	VH-VL of	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT

CEO	DESIGNATION	SOLIBCE	TVDE	CEOLIENCE
N ⊡ N			<u>-</u>	
	CDH19 14302			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVL
1385	CDH19 14302 x I2C	artificial	e e	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV OOKPGOAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHH
1386	CDR-H1 of CDH19 14303	artificial	AA	SYGMH
1387	CDR-H2 of CDH19 14303	artificial	ΑA	FIWYEGSNKYYAESVKD
1388	CDR-H3 of CDH19 14303	artificial	ΑA	RAGIIGTIGYYYGMDV
1389	CDR-L1 of CDH19 14303	artificial	ΑA	SGDRIGEKYTS
0621 266	CDR-L2 of CDH19 14303	artificial	AA	QDTKRPS
1391	CDR-L3 of CDH19 14303	artificial	ΑA	QAWESSIVV
1392	VH of CDH19 14303	artificial	t	CAGGTGCAGTTGGTGGAGGGGGGGGGGGCTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGTGGTATTATATATGGTATGAAGGAAG
1393	VH of CDH19 14303	artificial	ΑA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1394	VL of CDH19 14303	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
1395	VL of CDH19 14303	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
1396	VH-VL of CDH19 14303	artificial	nt	CAGGTGCAGTTGGTGGAGTGTGGGGGGGGGGGTGCTCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTTTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
				AATACTATGCAGAGTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAAAAGCCTG AGAGCTGAGGACACGGCTGTGTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACTACGTATGGACGTCTG GGGCCTACGACGCTGTGTGTGTGCGAGAGGGCCGGTTAATAGGAACTATAGGCTACTACTACGGTTGAACGTCTG GGGCCAAGGGACCACGGTCTCTTAGTGGTGGCGGAGGTCTGGCGGAGGCGGAGGCGGAGGCGGATCTTCCTATGAAC TGACTCAGCCACCCTCAGTGTCCCCAGGACAGACAGACAG
1397	VH-VL of CDH19 14303	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGSGYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVL
1398	x I2C	artificial	g	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
6621 267	CDR-H1 of CDH19 14078	artificial	AA	RYGIH
1400	CDR-H2 of CDH19 14078	artificial	AA	VIWYDGSNKYYADSVKG
1401	CDR-H3 of CDH19 14078	artificial	AA	RAGIPGTTGYYYGMDV
1402	CDR-L1 of CDH19 14078	artificial	AA	SGDRIGEKYVS
1403	CDR-L2 of CDH19 14078	artificial	AA	QDNKWPS
1404	CDR-L3 of CDH19 14078	artificial	AA	QAWDSSTVV
1405	VH of CDH19 14078	artificial	t	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGGCGTGGTCCAGCCTGGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
1406	VH of CDH19 14078	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS

VL of CDH19 artificial nt 14078 VL of CDH19 artificial AA 14078 VH-VL of artificial nt CDH19 14078 CDH19 14078 CDH19 14078 CDH19 14080 CDR-H1 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080	S	SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
1407 VL of CDH19 artificial nt 14078 vL of CDH19 artificial AA 1409 vH-VL of artificial nt CDH19 14078 artificial AA 1410 vH-VL of artificial AA 1411 CDH19 14078 artificial AA 1412 CDR-H1 of artificial AA 1413 CDR-H2 of artificial AA 1414 CDR-H3 of artificial AA 1415 CDR-H3 of artificial AA 1415 CDR-L1 of artificial AA 1416 CDR-L2 of artificial AA 1416 CDR-L2 of artificial AA 1416 CDR-L3 of artificial AA CDH19 14080 artificial AA CDH19 14080 artificial AA	= Z	o <u>o</u>				
1408 VL of CDH19 artificial AA 14078 artificial nt CDH19 14078 artificial AA CDH19 14078 artificial AA 1411 CDH19 14078 artificial AA 1412 CDR-H1 of artificial AA 1413 CDR-H2 of artificial AA 1414 CDR-H3 of artificial AA 1415 CDR-H3 of artificial AA 1416 CDR-L1 of artificial AA 1416 CDR-L2 of artificial AA 1416 CDR-L2 of artificial AA 1417 CDR-L3 of artificial AA 1416 CDR-L2 of artificial AA CDH19 14080 artificial AA CDH19 14080 artificial AA	—		VL of CDH19 14078	artificial	t	TCCTATGAGCTGACTCAGCCCACCTCAGTGTCCCCAGGACAGACA
1409 VH-VL of CDH19 14078 artificial nt 1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial aa 1412 CDR-H1 of CDH19 14080 artificial AA 1413 CDR-H2 of CDH19 14080 artificial AA 1414 CDR-H3 of CDH19 14080 artificial AA 1415 CDR-L1 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1416 CDR-L3 of CDH19 14080 artificial AA 1417 CDR-L3 of CDH19 14080 artificial AA	Π.	408	VL of CDH19 14078	artificial	АА	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
CDH19 14078 AA		409	VH-VL of	artificial	ıt	CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGCTGGGGGGGTCCCTGAGACTCTCTGTGCAGCGTCTGGATTCACCTT
1410 VH-VL of artificial AA 1411 CDH19 14078 artificial aa			CDH19 14078			CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAAGGGGCTGGAGTGGGTGG
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial aa artificial AA 1412 CDR-H1 of CDH19 14080 artificial AA 1413 CDR-H2 of CDH19 14080 artificial AA 1414 CDR-H3 of CDH19 14080 artificial AA 1415 CDR-L1 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1417 CDR-L3 of CDH19 14080 artificial AA						AATACTATGCAGACTCCGTGTATTACTGTGCGAGAAGGCCCGGTATACCTGGAACACACGCTGTATTTGCAATATGAACAGCCTG
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial AA 1412 CDR-H1 of CDH19 14080 artificial AA 1413 CDR-H2 of CDH19 14080 artificial AA 1414 CDR-H3 of CDH19 14080 artificial AA 1415 CDR-L1 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1417 CDR-L3 of CDH19 14080 artificial AA 1417 CDR-L3 of CDR-L3 of CDR-L3 of CDH19 14080						GGGCCAAGGGACCACGGTCACCGTCTCCTCAGGTGGCGGAGGATCTGGCGGAGGTGGAAGCGGAGGCGGCGGAGGCGGATCTTCCTATGAGC
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial aa artificial AA 1412 CDR-H1 of artificial AA CDH19 14080 artificial AA 1414 CDR-H2 of CDH19 14080 artificial AA 1415 CDR-H3 of artificial AA CDH19 14080 artificial AA						TGACTCAGCCACCCTCAGTGTCCCGTGTCCCCAGGACAGACA
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial aa 1412 CDR-H1 of artificial AA CDH19 14080 artificial AA CDH19 14080 artificial AA CDH19 14080 artificial AA 1415 CDR-H3 of artificial AA CDH19 14080 artificial AA						AGCTGGTATCAGCAGAAGCCAGGCCAGTCCCCTATACTGGTCATCTATCAAGATAATAAGTGGCCCTCAGGGATCCCTGAGCGATT
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial AA 1412 CDR-H1 of CDH19 14080 artificial AA 1413 CDR-H2 of CDH19 14080 artificial AA 1414 CDR-H3 of CDH19 14080 artificial AA 1415 CDR-L1 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1417 CDR-L3 of CDH19 14080 artificial AA 1417 CDR-L3 of CDH19 14080 artificial AA						CICTGGCICCAACTCTGGGAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTACTGTCAGGCGT
1410 VH-VL of CDH19 14078 artificial AA 1411 CDH19 14078 artificial AA 1412 CDR-H1 of CDH19 14080 artificial AA 1413 CDR-H2 of CDR-H3 of CDH19 14080 artificial AA 1414 CDR-H3 of CDR-H3 of CDH19 14080 artificial AA 1415 CDR-L1 of CDR-L3 of CDH19 14080 artificial AA 1416 CDR-L2 of CDH19 14080 artificial AA 1416 CDR-L3 of CDR-L3 of CDH19 14080 artificial AA 1417 CDR-L3 of CDR-						GGGACAGCAGCACTGTGGTATTCGGCGGGGGGGCCAAGCTGACCGTCCTA
CDH19 14078 artificial aa x I2C CDR-H1 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L3 of artificial AA CDH19 14080		410	VH-VL of CDH19 14078	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYOOKPGOSPIIJYTYODNKWPSGIPPRFSGSNSGNYTATTTTSGTOAMDRADYYCOAWDSSTYJYFGGGTKITVI
CDR-H1 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L3 of artificial AA CDH19 14080		711	CDH19 1/078		;	OVOTEX TO THE SECOND OF THE SE
CDR-H1 of artificial AA CDH19 14080 CDR-H2 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L3 of artificial AA CDH19 14080	1		x 12C	5	3	RAEDSAVYYCARRAGIPGTIGYYYGMDVWGOGTTVTVSSGGGGSGGGGGGGSSYELTOPPSVSVSPGOTASITCSGDRLGEKYV
CDR-H1 of artificial AA CDH19 14080 CDR-H2 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L3 of artificial AA CDH19 14080)			SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE
CDR-H1 of artificial AA CDH19 14080 CDR-H2 of artificial AA CDH19 14080 CDR-H3 of artificial AA CDH19 14080 CDR-L1 of artificial AA CDH19 14080 CDR-L2 of artificial AA CDH19 14080 CDR-L3 of artificial AA CDH19 14080						SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
CDR-H1 of artificial AA RYGIH CDH19 14080 artificial AA VIWYDGSNK CDR-H2 of artificial AA RAGIPGTTG CDR-H3 of artificial AA RAGIPGTTG CDR-L1 of artificial AA SGDRLGEKY CDR-L2 of artificial AA QDNKWPS CDR-I3 of artificial AA QDNKWPS						TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
CDH19 14080 CDR-H2 of CDR-H2 of CDH19 14080 artificial AA VIWYDGSNK CDH19 14080 CDR-H3 of CDR-L1 of CDR-L1 of CDH19 14080 artificial AA SGDRLGEKY CDH19 14080 CDR-L2 of CDH19 14080 artificial AA QDNKWPS CDH19 14080 artificial AA QDNKWPS CDR-L3 of ABDISSTVV AA OAWDSSTVVV	1	412	CDR-H1 of	artificial	AA	RYGIH
CDR-H2 of artificial AA VIWYDGSNK CDH19 14080 Artificial AA RAGIPGTTG CDR-H3 of artificial AA SGDRLGEKY CDR-L1 of artificial AA SGDRLGEKY CDH19 14080 artificial AA QDNKWPS CDH23 of artificial AA QDNKWPS CDR-L3 of artificial AA OAWDSSTVV			CDH19 14080			
CDH19 14080 CDR-H3 of artificial AA RAGIPGITG CDH19 14080 CDR-L1 of artificial AA SGDRLGEKY CDH19 14080 CDR-L2 of artificial AA QDNKWPS CDH19 14080 artificial AA QDNKWPS CDR-L3 of artificial AA OAWDSSTVV	1	413	CDR-H2 of	artificial	¥	
CDR-H3 of artificial AA RAGIPGTTG CDH19 14080 artificial AA SGDRLGEKY CDH19 14080 artificial AA QDNKWPS CDH19 14080 artificial AA QDNKWPS CDH3 3 of artificial AA OAWDSSTVV			CDH19 14080			
CDH19 14080 AA SGDRLGEKY CDR-L1 of CDR19 14080 artificial AA QDNKWPS CDH19 14080 artificial AA OAWDSSTVV	1	414	CDR-H3 of	artificial	ΑĄ	RAGIPGTIGYYGMDV
CDR-L1 of artificial AA SGDRLGEKY CDH19 14080 artificial AA QDNKWPS CDH19 14080 artificial AA ODNKWPS CDR-I3 of artificial AA OAWDSSTVV			CDH19 14080			
CDR-L2 of artificial AA CDR-13 of artificial AA	1	415	CDR-L1 of	artificial	AA	SGDRIGEKYVY
CDR-L2 of artificial AA CDH19 14080 CDR-I3 of artificial AA			CDH19 14080			
CDR-13 of artificial AA		416	CDR-L2 of CDH19 14080	artificial	Ą	QDNKWPS
ai miciai De la Civica	1	1417	CDR-L3 of	artificial	AA	QAWDSSTVV

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> №				
	CDH19 14080			
1418	VH of CDH19 14080	artificial	nt	CAGGTGCAGCTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTCGCTATGGCATACCACTGGGTCCGCCAGGCTCCAGGCCTGGAGTGGGTGG
1419	VH of CDH19 14080	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
1420	VL of CDH19 14080	artificial	t	TCCTATGAGCTGACTCAGCCCACCCTCAGTGTCCCCAGGACAGACA
1421	VL of CDH19 14080	artificial	Ą	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVYWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVL
1422	VH-VL of	artificial	ıt	
269	CDH19 14080			CAGTCGCTATGGCATACACTGGGTCCGCCAGGCTCCAGGCAGG
1423	VH-VL of CDH19 14080	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV YWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVL
1424	CDH19 14080 x I2C	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLLMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV YWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
1425	CDR-H1 of CDH19 13591	artificial	Ą	SYSWS
1426	CDR-H2 of	artificial	AA	YIYYSGSTNYNPSLKS

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9				
	CDH19 13591			
1427	CDR-H3 of CDH19 13591	artificial	AA	NWAFHFDY
1428	CDR-L1 of CDH19 13591	artificial	AA	TGSSSNIGTGYDVH
1429	CDR-L2 of	artificial	ΑA	GNSNRPS
1430	CDR-L3 of CDH19 13591	artificial	AA	QSYDSSLSGWV
1431	VH of CDH19	artificial	nt	
	13591			CAGTAGTTACTCCTGGAGCTGGATCCGGCAGCCCCCAGGGAAGGGACTGGAGTGGATTGGGTATATCTATTACAGTGGAGCACCAAACCAACTACAAGCACCCCAAACCAACC
				TAGT
1432	VH of CDH19 13591	artificial	₹	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSS
1433	VL of CDH19 13591	artificial	nt	CAGTCTGTGCTGACGCCGCCCCCCAGTGTCTGGGGCCCCCAGGGCAGAGGGTCACCATCTCCTGCACTGGGAGCAGCTCCAATAT CGGGACAGGTTATGATGATGTACACTGGTATCAGCAGCTTCCAGGAACAGCCCCCAAACTCCCTCATCCTTGGTAACAGCAATCGGCCCT CAGGGGTCCCTGACCGATTCTCTGGCTCCAAGTCTGGCACCTCAGCCTCCCTGGCCTATGGGCTCAGGCTGAGGATGAGGCT GATTATTACTGCCAGTCCTATGACAGCCAGTCTGAGTGTTGGGTGTTCGGCGGAGGGGACCAGGTTGACCGTCCTA
1434	VL of CDH19 13591	artificial	ΑA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIHGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYDSSLSGWVFGGGTRLTVL
1435	VH-VL of CDH19 1591	artificial	Ħ	
				ACTACAACCCCTCCAAGAGTCGAGTCACCATATCATTAGACACGTCCAAGAACCAGTTCTCCTGAAGCTGTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCTGTGACCGACC
1436	VH-VL of CDH19 13591	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
1437	CDH19 13591	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT

SEQ 10 0. 0. 14438 1440 1440 1440 1440 1440 1440 1440 144	x I2C CDR-H1 of CDR-H2 of CDH19 14299 CDR-H3 of CDH19 14299 CDR-L1 of CDH19 14299 CDR-L2 of CDH19 14299 CDR-L3 of CDH19 14299 CDR-L3 of CDH19 14299 CDR-L3 of CDH19 14299	artificial artificial artificial artificial artificial artificial artificial artificial artificial	TYPE AA	SEQUENCE SEQUENCE AADTAVYYCARNWAPHFDYWGQGTLVTTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGG
1444	VH of CDH19 14299 VH of CDH19 14299	artificial	Nt AA	CAGGIGCAGGAGAGACCCAGGACTGGIGGAGGGAGGGACTGIGTCCCTGACCTGCACTGCA
1446	VL of CDH19 14299 VL of CDH19	artificial artificial	nt AA	
1448	14299 VH-VL of CDH19 14299	artificial	nt	CAGGTGCAGCTGCAGGAGTCGGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTCACCTGTCTCTGTCTCTGGTGGCTCCAT CAGGTGCAGCTGCAGGAGCTCGGAGCTCGGACCCCAGGGACTTCGGAGACCCTGTCCTCTGTTATCTATTACAGTGGGAGCACCCA ACTACAACCCCTCCTCAAGAGTCGAGTC

SEQ ID NO.	DESIGNATION	SOURCE	ТҮРЕ	SEQUENCE
				TAGTGGTGGCGGAGGATCTGGCGGAGGTGGAAGCGGAGGCGCGCGGATCTCAGTCTGTGCTGACGCCGCCCCTCAGTGTCTGGGGGCCCCCTCAGTGTCTGGGGGCCCCCCCAGGGGGGGG
1449	VH-VL of CDH19 14299	artificial	Ą	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
1450	CDH19 14299 x I2C	artificial	aa	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPCKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQL PGTAPYLLIHGNSNRPPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYDSSLSGWVFGGGTRLTVLSGGGGSEVQLVESGGG LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
272	ckCDH19(1- 43)::FLAG::ckC DH19(44-776)	artificial	ве	MNCSTFLSLVLALVQLQLCSPTTQIFSAQKTDQSYTTIRRVKRDYKDDDDKGWVWEPLFVTEEETSTMPMYVGQLKSDLDKEDGSL QYILTGEGADSIFFINEHGKIYVRQKLDREKKSFYILRAQVINRKTRHPIEPDSEFIIKVRDINDHEPQFLDGPYVATVPEMSPEG TSVTQVTATDGDDPSYGNNARLLYSLIQGQPYFSVEPKTGVIRMTSQMDRETKDQYLVVIQAKDMVGQAGAFSATATVTINLSDVN DNPPKFQQRLYYLNVSEEAPVGTTVGRLLAEDSDIGENAAMNYFIEEDSSDVFGIITDRETQEGIIILKKRVDYESKRKHSVRVA VNRYIDDRFLKEGPFEDITIVQISVVDADEPPVFTLESYVMEIAEGVVSGSLVGTVSARDLDNDDSSVRYSIVQGLHLKRLFSINE HNGTIITTEPLDREKASWHNITVTATETRNPEKISEANVYIQVLDVNDHAPEFSKYYETFVCENAVPGQLIQNISAVDBDSAENH RFYFSLAQATNSSHFTVKDNQDNTAGIFTAGSGFSRKEQFYFTLPILILDNGSPPLTSTNTLTVTVCDCDTEVNTLYCRYGAFLYS IGLSTEALVAVLACLLILLVFFLAIIGIRQQRKKTLFSEKVEEFRENIVRYDDEGGGEEDTEAFDISALRTRAVLRTHKPRKKITT EIHSLYRQSLQVGPDSAIFRQFISEKLEEANTDPSVPPYDSLQTYAFEGTGSLAGSLSSLGSNTSDVDQNYEYLVGWGPPFKQLAG MYTSQRSTRD
1452	huCDH19(1- 43)::FLAG::hu(44- 141)::ckCDH1 9(142-776)	artificial	ee	MNCYLLRFMLGIPLLWPCLGATENSQTKKVKQPVRSHLRVKRDYKDDDDKGWVWNQFFVPEEMNTTSHHIGQLRSDLDNGNNSFQ YKLLGAGAGAGSTFIIDERTGDIYAIQKLDREERSLYILRAQVIDIATGRAVEPESEFVIKVSDINDHEPQFLDGPYVATVPEMSPEG TSVTQVTATDGDDPSYGNNARLLYSLIQGQPYFSVEPKTGVIRMTSQMDRETKDQYLVVIQAKDWVGQAGAFSATATVTINLSDVN DNPPKFQQRLYYLNVSEEAPVGTTVGRLLAEDSDIGENAAMNYFIEEDSSDVFGIITDRETQEGIIILKRRVDYESKRHKHSVRVA VNRYIDDRFLKEGPFEDITIVQISVVDADEPPVFTLESYVMEIAEGVVSGSLVGTVSARDLDNDDSSVRYSIVQGLHLKRLFSINE HNGTIITTEPLDREKASWHNITVTATETRNPEKISEANVYIQVLDVNDHAPEFSKYYETFVCENAVPGQLIQNISAVDBDSAENH RFYFSLAQATNSSHFTVKDNQDNTAGIFTAGSGFSRKEQFYFFLPILILDNGSPPLTSTNTLTVTVCDCDTEVNTLYCRYGAFLYS IGLSTEALVAVLACLLILLVFFLAIIGIRQQRKKTLFSEKVEEFRENIVRYDDEGGGEEDTEAFDISALRTRAVLRTHKPRKKITT EIHSLYRQSLQVGPDSAIFRQFISEKLEEANTDPSVPPYDSLQTYAFEGTGSLAGSLSSLGSNTSDVDQNYEYLVGWGPPFKQLAG MYTSQRSTRD
1453	ckCDH19(1- 43)::FLAG::ckC	artificial	aa	MNCSTFLSLVLALVQLQLCSPTTQIFSAQKTDQSYTTIRRVKRDYKDDDDKGWVWEPLFVTEEETSTMPMYVGQLKSDLDKEDGSL QYILTGEGADSIFFINEHGKIYVRQKLDREKKSFYILRAQVINRKTRHPIEPDSEFIIKVRDINDNEPKFLDEPYEAIVPEMSPEG

DESIGNATION DH19(44- 141)::huCDH1 9(142- 249)::ckCDH1 9(250-776) ckCDH19(1- 43)::FLAG::ckC DH19(44- 249)::huCDH1 9(250- 364)::ckCDH19(1- 43)::FLAG::ckC DH19(44- 364)::ckCDH1 9(365- 463)::ckCDH1	artificial artificial artificial	аа аа	### SEQUENCE ### SEQUENCE ### SEQUENCE ### TILVIQVTASDADDPSSGNNARLIYSILQGQPYESVEPTTGVIRISSKNORELQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVN ### DIPPKEYQRLYTLANVSEEAPVGTTVGRLAAEDSDLGENAAMNYFIEBDSSDVGIIYDARDDNDDSSVRXSIVOGENARVRA ***WIXTIDDREFLEGPPEDITIVORSULAEDSDLGENAAMNYFIEBDSSDVGIIYDARDDNDDSSVRXSIVOGENARRAVRA ***WIXTIDDREFLEGPPEDITIVORSULAEDSONGEPPYTLAEGVVSGSLOGTVSARDDNDDSSVRXSIVOGDDENARLASINE ### RRYTELADALANACLILILAVPTATETRANPERTUSEKVEBPREDITYIONGSPPTTYTYVCDCDTGVNTTYTYCRDABALSINE ### RRYTELADALANACLILILAVPTATIONGPREDEXTYTORYBODDRGWWEDELTVTPEETSTMPHYVCDCDTGVNTTYCRGAFLYS ### TILGARTELANALACLILILAVPTATIONGPREDEXTYREWRENDENARDSDDRGWWEDELTWYTDEETSTMPHYVCDCDTGVNTTYCRGAFLYS ### TILGARTELANALACLILILAVPTATIONGPREDEXTYREWRENDENARDSDDRGWWEDELTWYTDEETSTMPHYVCDCDTGVNTTYCRGAFLYS ### TILGARTELANALACLILILAVPTATIONGPREDEXTYREWRENDENARDSDDRGWWEDELTWYTDEETSTMPHYVCDCDTGVNTTYCRGAFLYS ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTATT ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTHT ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTHT ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTHT ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTHT ### TILGARTELANALACLILILAVPTATIONGPREDEXTREATHYREWRENGTHT ### TILGARTELANALACLILILATYRERGETSTATHYREWRENGTHT ### TILGARTELANALACLILILATYREWRENGTHT ### TILGARTELANALACLILILATYREWRENGTHT ### TILGARTELANALACLILILATYREWRENGTHT ### TILGARTELANALACLILILATYREWRENGTHT ### TILGARTELANALACLILILATYREAGTHT ### TILGARTELANALACLILILATYREWRENGTHT ### TILGARTELANALA
DH19(44- 468)::huCDH1 9(464-772)			TSVTQVTATDGDDPSYGNNARLLYSLIQGQPYFSVEPKTGVIRMTSQMDRETKDQYLVVIQAKDMVGQAGAFSATATVTINLSDVN DNPPKFQQRLYYLNVSEEAPVGTTVGRLLAEDSDIGENAAMNYFIEEDSSDVFGIITDRETQEGIILLKKRVDYESKRKHSVRVKA VNRYIDDRFLKEGPFEDITIVQISVVDADEPPVFTLESYVMEIAEGVVSGSLVGTVSARDLDNDDSSVRYSIVQGLHLKRLFSINE HNGTIITTEPLDREKASWHNITVTATETRNPEKISEANVYIQVLDVNDHAPEFSQYYETYVCENAGSGQVIQTISAVDRDESIEEH HFYFNLSVEDTNNSSFTIIDNQDNTAVILTNRTGFNLQEEPVFYISILIADNGIPSLTSTNTLTHVCDCGDSGSTQTCQYQELVL SMGFKTEVIIALLICIMIIFGFIFLTLGLKQRRKQILFPEKSEDFRENIFQYDDEGGGEEDTEAFDIAELRSSTIMRERKTRKTTS
	DESIGNATION DH19(44- 141)::huCDH1 9(142- 249)::ckCDH1 9(250-776) ckCDH19(1- 43)::FLAG::ckC DH19(44- 249)::huCDH1 9(365-776) 364)::ckCDH19(1- 43)::FLAG::ckC DH19(44- 463)::ckCDH1 9(365- 463)::ckCDH1 9(469-776)	HATION H4- WCDH1 Y76) H4- WCDH1 Y76) H4- WCDH1 Y76) Y6::ckC H4- WCDH1 Y76) H4- WCDH1 Y76)	HATION SOURCE H4- UCDH1 776) H6:CC H4- UCDH1 776) AG::CKC H4- UCDH1 776) AG::CKC H4- UCDH1 AG::CKC H4- UCDH1 776) AG::CKC H4- UCDH1 776)

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
Q. 0.				
				AEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSLSSLESAVSDQDESYDYLNELGPRFKRLA CMFGSAVQSNN
1457	rhCDH19(1- 43)::FLAG::rhC DH19(44-772)	artificial	e e	MNCYLLLPFMLGIPLLWPCLGATENSQTKKVQQPVGSHLRVKRDYKDDDDKGWVWNQFFVPEEMNTTSHHVGRLRSDLDNGNNSFQ YKLLGAGAGSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDITTGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEG TLVIQVTASDADDPSSGNNARLLYSLLQGQPYFSVEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGQPGALSGTTSVLIKLSDVN DNKPIFKESLYRLTVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTFDIITNHETQEGIVILKKKVNFEHQNHYGIRAKV KNHHVDEQLMKYHTEASTTFIKIQVEDVDEPPLFLLPYYIFEIFETPQGSFVGVVSATDPDNRKSPIRYSITRSKVFNIDDNGTI TTTNSLDREISAWYNLSITATEKYNIEQISSIPVYVQVLNINDHAPEFSQYYESYVCENAGSGQVIQTISAVDRDESIEEHHFYFN LSVEDTNSSSFTIIDNQDNTAVILTNRTGFNLQEEPIFYISILIADNGIPSLTSTNTLTIHVCDCDDSGSTQTCQYQELMLSMGFK TEVIIAILICIMVIFGFIFLTLGLKQRRKQILFPEKSEDFRENIFRYDDEGGGEEDTEAFDVAALRSSTIMRERKTRKTTSAEIRS LYRQSLQVGPDSAIFRKFILEKLEEADTDPCAPPFDSLQTYAFEGTGSLAGSLSSLESAVSDQDESYDYLNELGPRFFKRLACMFGS AVQSNN
274	caCDH19(1- 42)::FLAG::caC DH19(43-770)	artificial	a a	QFFVPEEMNKTDYHIGQLRSDLDNGNNSFQYKLLGAGAGSIFVIDERTGDIYAIQKLDREERSLYTLRAQVIDSTTGRAVEPESEF VIRVSDINDNEPKFLDEPYEAIVPEMSPEGTLVIQVTATDADDPASGNNARLLYSLLQGQPYFSIEPTTGVIRISSKMDRELQDEY WVIIQAKDMIGLPGALSGTTSVLIKLSDVNDNKPIFKERLYRLTVSESAPTGTSIGRIMAYDNDIGENAEMDYSIEDDSQTFDIIT NNETQEGIVILKKKVDFEHQNHYLIRANVKNRHVAEHLMEYHVEASTTFVRVQVEDEDEPPVFLLPYYLFEILEESPHGSFVGMVS ATDPDQRKSPIRYSITRSKVFSIDDNGTIITTNPLDREISAMYNLSITATEKYNVQQISAVPVYVQVLNINDHAPEFSEFYDSYVC ENAGSGQVIQTISAVDRDESVEDHHFYFNLSVEDTKNSSFIIIDNEDNTAVILTNRTGFSLQEEPVFISVIIADNGIPSLTSTNT LTIHICDCDDYGSTQTCRDKDLLLSMGFRTEVILAILISIMIIFGFIFLLGLKQRRKPTLFPEKGEDFRENIFRYDDEGGGEEDT EAFDIVQLRSSTIMRERKTRKTAAAEIRSLYRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSLSSLG SAVSDQDENYDYLNELGPRFKRLACMFGSAMQSNN
1459	rhCDH19(1- 43):FLAG::rhC DH19(44- 141)::caCDH1 9(141-770)	artificial	ee	MNCYLLLPFMLGIPLLWPCLGATENSQTKKVQQPVGSHLRVKRDYKDDDKGWVWNQFFVPEEMNTTSHHVGRLRSDLDNGNNSFQ YKLLGAGAGSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDITTGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEG YKLLGAGAGSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDITTGRAVEPESEFVIKVSDINDNEPKFLDEPYEAIVPEMSPEG TLVIQVTATDADDPASGNNARLLYSLLQGQPYFSIEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGLPGALSGTTSVLIKLSDVN DNKPIFKERLYKLTVSESAPTGTSIGENMAYDNDIGENAEMDYSIEDDSQTFDIITNNETQEGIVILKKKVDFEHQNHYLIRANVK NRHVAEHLMEYHVEASTTFVRVQVEDEDEPPVFLLPYYLFEILEESPHGSFVGMVSATDPDQRKSPIRYSITRKKVDFEHQNHYLITRANVK TTNPLDREISAWYNLSITATEKYNVQQISAVPVYVQVLNINDHAPEFSEYYDSYVCENAGSGQVIQTISAVDRDESVEDHHFYFNL SVEDTKNSSFIIIDNEDNTAVILTNRTGFSLQEEPVFYISVLIADNGIPSLTSTHTHICDCDDYGSTQTCRDKDLLLSMGFRT EVILAILISIMIIFGFIFLILGLKQRRKPTLFPEKGEDFRENIFRYDDEGGGEEDTEAFDIVQLRSSTIMRERKTRKTAAAEIRSL YRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFFGTGSLAGSSLGSAVSDQDENYDYLNELGPRFKRLACMFGSA MQSNN
1460	rhCDH19(1- 43)::FLAG::rhC DH19(44- 65)::caCDH19(artificial	вв	MNCYLLLPFMLGIPLLWPCLGATENSQTKKVQQPVGSHLRVKRDYKDDDDKGWVWNQFFVPEEMNTTSHHVGRLRSDLDNGNNSFQ YKLLGAGAGSIFVIDERTGDIYAIQKLDREERSLYTLRAQVIDSTTGRAVEPESEFVIRVSDINDNEPKFLDEPYEAIVPEMSPEG TLVIQVTATDADDPASGNNARLLYSLLQGQPYFSIEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGLPGALSGTTSVLIKLSDVN DNKPIFKERLYRLTVSESAPTGTSIGRIMAYDNDIGENAEMDYSIEDDSQTFDIITNNETQEGIVILKKKVDFEHQNHYLIRANVK NRHVAEHLMEYHVEASTTFVRVQVEDEDEPPVFLLPYYLFEILEESPHGSFVGMVSATDPDQRKSPIRYSITRSKVFSIDDNGTII

SEQ ID NO.	DESIGNATION	SOURCE	ТҮРЕ	SEQUENCE
				TTNPLDREISAWYNLSITATEKYNVQQISAVPVYVQVLNINDHAPEFSEYYDSYVCENAGSGQVIQTISAVDRDESVEDHHFYFNL SVEDTKNSSFIIIDNEDNTAVILTNRTGFSLQEEPVFYISVLIADNGIPSLTSTNTLTIHICDCDDYGSTQTCRDKDLLLSMGFRT EVILAILISIMIIFGFIFLILGLKQRRKPTLFPEKGEDFRENIFRYDDEGGGEEDTEAFDIVQLRSSTIMRERKTRKTAAAEIRSL YRQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSLSSLGSAVSDQDENYDYLNELGPRFKRLACMFGSA MQSNN
1461	caCDH19(1- 43)::FLAG::caC DH19(44- 87)::rhCDH19(89- 114)::caCDH1 9(115-770)	artificial	ее	MNYCFLLPLMLGIPLIWPCFTASESSKTEVKHQAGSHLRVKRDYKDDDKGWMWNQFFVPEEMNKTDYHIGQLRSDLDNGNNSFQY KLLGAGAGSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDSTTGRAVEPESEFVIRVSDINDNEPKFLDEPYEAIVPEMSPEGT LVIQVTATDADDPASGNNARLLYSLLQGQPYFSIEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGLPGALSGTTSVLIKLSDVND NKPIFKERLYRLTVSESAPTGTSIGRIMAYDNDIGENAEMDYSIEDDSQTFDIITNNETQEGIVILKKKVDFEHQNHYLIRANVKN RHVAEHLMEYHVEASTTFVRVQVEDEDEPPVFLLPYYLFEILEESPHGSFVGMVSATDPDQRKSPIRYSITRSKVFSIDDNGTIIT TNPLDREISAWYNLSITATEKYNVQQISAVPVYVQVLNINDHAPEFSEYYDSYVCENAGSGQVIQTISAVDRDSVVEDHHFYFNLS VEDTKNSSFIIIDNEDNTAVILTNRTGFSLQEEPVFYISVLIADNGIPSLTSTNTLTHICDCDDYGSTQTCRDKDLLLSMGFRTE VILAILISIMIIFGFIFLIIGLKQRRKPTLFPEKGEDFRENIFRYDDEGGGEEDTEAFDIVQLRSSTIMRERKTRKTAAAEIRSLY RQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSLSSLGSAVSDQDENYDYLNELGPRFKRLACMFGSAM QSNN
275	caCDH19(1- 43)::FLAG::caC DH19(44- 120)::rhCDH1 9(122- 137)::caCDH1 9(137-770)	artificial	е е	MNYCFLLPLMLGIPLIWPCFTASESSKTEVKHQAGSHLRVKRDYKDDDKGWMWNQFFVPEEMNKTDYHIGQLRSDLDNGNNSFOY KLLGAGAGSIFVIDERTGDIYALQKLDREERSLYTLRAQVIDITTGRAVEPESEFVIKVSDINDNEPKFLDEPYEALVPEMSPEGT LVIQVTATDADDPASGNNARLLYSLLQGQPYFSIEPTTGVIRISSKMDRELQDEYWVIIQAKDMIGLPGALSGTTSVLIKLSDVND NKPIFKERLYRLTVSESAPTGTSIGRIMAYDDIGENAEMDYSIEDDSQTFDIITNNETQEGIVILKKKVDFEHQNHYLIRANVKN RHVAEHLMEYHVEASTTFVRVQVEDEDEPPVFLLPYYLFEILEESPHGSFVGMVSATDPDQRKSPIRYSITRSKVFSIDDNGTIIT TNPLDREISAWYNLSITATEKYNVQQISAVPVYVQVLNINDHAPEFSEYYDSYVCENAGSGQVIQTISAVDRDESVEDHHFYFNLS VEDTKNSSFIIIDNEDNTAVILTNRTGFSLQEEPVFYISVLIADNGIPSLTSTNTLTHICDCDDYGSTQTCRDKDLLLSMGFRTE VILAILISIMIIFGFIFLIIGLKQRRKPTLFPEKGEDFRENIFRYDDEGGGEEDTEAFDIVQLRSSTIMRERKTRRTAAAEIRSLY RQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTYAFEGTGSLAGSLSSLGSAVSDQDENYDYLNELGPRFKRLACMFGSAM QSNN
1463	rhcDH19(1- 43)::FLAG::rhC DH19(44- 141)::raCDH19 (140- 247)::rhCDH1 9(250-772)	artificial	вв	MNCYLLPFMLGIPLLWPCLGATENSQTKKVQQPVGSHLRVKRDYKDDDDKGWVWNQFFVPEEMNTTSHHVGRLRSDLDNGNNSFQ YKLLGAGAGSTFIIDERTGDIYAIEKLDREERSLYILRAQVIDITTGRAVEPESEFVIKVSDINDNEPRFLDEPYEAIVPEMSPEG TFVIKVTANDADDPTSGYHARILYNLEQGQPYFSVEPTTGVIRISSKMDRELQDTYCVIIQAKDMLGQPGALSGTTTISIKLSDIN DNKPIFKESLYRLTVSESAPTGTSIGTIMAYDNDIGENAEMDYSIEEDDSQTFDIITNHETQEGIVILKKKVNFEHQNHYGIRAKV KNHHYDEQLMKYHTEASTTFIKIQVEDVDEPPLFLLPYYIFEIFETPQGSFVGVVSATDPDNRKSPIRYSITRSKVFNIDDNGTI TTTNSLDREISAWYNLSITATEKYNIEQISSIPVYVQVLNINDHAPEFSQYYESYVCENAGSGQVIQTISAVDRDESIEEHHFYFN LSVEDTNSSSFTIIDNQDNTAVILTNRTGFNLQEEPIFYISILIADNGIPSLTSTNTLTIHVCDCDDSGSTQTCQYQELMLSMGFK TEVIIAILICIMVIFGFIFLTLGLKQRRKQILFPEKSEDFRENIFRYDDEGGGEEDTEAFDVAALRSSTIMRERKTRTASAEIRS LYRQSLQVGPDSAIFRKFILEKLEEADTDPCAPPFDSLQTYAFEGTGSLAGSSLESAVSDQDESYDYLNELGPRFKRLACMFGS AVQSNN
1464	raCDH19(1-	artificial	aa	MNHYFLKYWILMVPLIWPCLKVAETLKIEKAQRAVPSLGRAKRDYKDDDDKGWVWKQFVVPEEMDTIQHVGRLRSDLDNGNNSFQY

1416 CH19(44- 233)::aCDH19(44- 232)::aCDH19(44- 233)::aCDH19(44- 232)::aCDH19(44- 24)::ACDH19(14- 24):ACDH19(14- 24)::ACDH19(14- 24):ACDH19(14- 24):ACDH19(14- 24):ACDH19(14- 24):ACDH19(14- 24):ACDH19(14- 24):ACDH19(14- 24):ACDH1
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SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9				
				AIIAIMICVMVIFGFFFLILALKQRRKETLFPEKTEDFRENIFCYDDEGGGEEDSEAFDIVELRQSTVMRERKPQRSKSAEIRSLY RQSLQVGPDSAIFRKFILEKLEEANTDPCAPPFDSLQTFAYEGTGSSAGSLSSLASRDTDQEDDFDYLNDLGPRFKRLASMFGSAV QPNN
1468	VH of CDH19	artificial	nt	CAGGTGCAGTTGGTGGAGTCTGGGGGAGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCCTGTGCAGCGTCTGGATTCACCTT
	14302 CC x I2C			CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGTGTCTGGAGTGGGGTGGCATTTATATATGGTATGAAGTAATA AATACTATGCAGACTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAATAGCCTG
				AGAGCTGAGGACACGGCTGTGTTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACTACGATATGGACGTCTG
				GGGCCAAGGGACCACGGTCACCGTCTTAGT
1469	VH of CDH19 14302 CC x I2C	artificial	¥	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1470	VL of CDH19	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
	14302 CC x 12C			AAAATATACTAGCTGGTATCAGCAGGGCCAGGCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC
				CTGAGCGATTCTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGATTAC TGTCAGGCGTGGGAGAGAACTAGTAGTATTTGGCTGCAGCAACTGAAGCTGACCATCATAA
1471	VI of CDH19	artificial	AA	SYELTOPPSYSVSPGOTASITCSGDRIGEKYTSWYOORPGOSPILVIYODTKRPSGIPFRFSGSNSGNTATLTISGTOAMDEADYY
	14302 CC x 12C	5		CQAWESSTVVFGCGTKLTVL
1472	VH-VL of	artificial	nt	CAGGTGCAGTTGGTGGAGTCTGGGGGGAGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCCTGTGCAGCGTCTGGGATTCACCTT
	CDH19 14302			CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGTGTCTGGAGTGGGTGG
	CC x 12C			AATACTATGCAGACTCCGTGAAGGACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAATAGCCTG
				AGAGCTGAGGACACGGCTGTGTTACTGTGCGAGAAGGGCCCGGTATAATAGGAACTATAGGCTACTACTACTACGGTATGGACGTCTG
				GGGCCAAGGGACCACGGTCACCGTCTCTAGTGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGGGGGGCTCCTCCTATGAAC
				TGACTCAGCCACCCTCAGTGTCCCGTGTCCCCAGGACAGACA
				AGCTGGTATCAGCAGAGGCCCAGGCCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCCCTGAGCGATT
				CTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTACTGTCAGGCGT GGGAGAGCAGCAGCACTGTGGTATTCGGCTGCGGGACCAAGCTGACCGTCCTA
1473	VH-VL of	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPCKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CDH19 14302			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
	CC x 12C			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVL
1474	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x 12C			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
				SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRIGNYGNSYLSYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV Ookpaoaprattagatkttapattapaktssasttagaattatsgvoptdeatyyvovtwysnrwvtgagatkttvthhhhhh
1475	CDH19 14302	artificial	ממ	OVOLVESGGGVVOPGGSTRTSCAASGFTFSSYGMHWVROAPGRGTFWYDGSNKYYADSVKDRFTTSRDNSKNTTYTOMNST.
7 17	CDIIIO 14004	alulua	۵۵	X.X.I. 100000. 1XI 00000000000000000000000000000000000

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	x F12q0			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKGRFTISRDDSKNTAYLQMNSLKTED TAVYYCVRHGNFGNSYVSWWAYWGQGTLVTVSSGGGGSGGGGSGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVL
1476	CD x F12q0	artificial	ee	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTIYLQMNSL RAEDTAVYYCARRAGIIGTIGYYGMDVWGQGTTVTVSSGGGGSGGGGGGSGYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKGRFTISRDDSKNTAYLQMNSLKTED TAVYYCVRHGNFGNSYVSWWAYWGQGTLVTVSSGGGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVL
1477	VH of CDH19	artificial	nt	CGGCTGATCGAGGACATCTGCCTGCCCAGATGGGGCCTGCCT
	21-14302 x I2C			GGTCCAGCCIGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTTCAGTAGCTATGGCATGCACTGGGTCCGCCAGG CTCCAGGCAAGGGGGGTCCCTGAGACTTTATATATGGTATGATGGAAGTAATAAATA
273	VH of CDH19	artificial	Ą	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRF
8	21-14302 x 12C			TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1479	VL of CDH19	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
	21-14302 x 12C			AAAATATACTAGCTGGTATCAGCAGAGGCCAGGCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC CTGAGCGATTCTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTAC TGTCAGGCGTGGGAGAGAGACACTGTGGTATTCGGCGGAGGGACCAAGCTGACCGTCCTA
1480	VL of CDH19 21-14302 x 12C	artificial	Ą	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
1481	VH-VL of	artificial	nt	CGGCTGATCGAGGACATCTGCCTGCCCAGATGGGGCTGCCTGTGGGACGACCAGCTGCAGGTTGGTGGAGGTTTGGGGGGGG
	14302 × I2C			CCCAGGCAGGGGCTGGAGGGGCATTTATATGGAGGAGGTGAATAATAATACTGTGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGA

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
				ACCAAGCTGACCGTCCTA
1482	VH-VL of CDH19 21- 14302 x I2C	artificial	AA	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVSVSP GQTASIICSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGG TKLTVL
1483	CDH19 21- 14302 x I 2 C	artificial	o o	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGGSGGGGGGGGGSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGG TKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISR DDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTL TCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKL TVLHHHHHH
1484	VH of CDH19	artificial	t	CGGCTGATCGAGGACATCTGCCCTGCCCAGATGGGGCTGCCTGTGGGAGGACGACCAGGTGCAGTTGGTGGAGTCTGGGGGAGGCGT
27	21-14302 CC x 12C		!	
1485	VH of CDH19 21-14302 CC x 12C	artificial	AA	RLIEDICLPRWGCIWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1486	VL of CDH19	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
	21-14302 CC x 12C			AAAATATACTAGCTGGTATCAGCAGGGCCAGGCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC CTGAGCGGTTCTTTTAC CTGAGCGATTCTTGGTATGATGAGCTGACTATTAC TGTCAGGCGTGCTGTGGTATTGGCTGACTATTAC TGTCAGGCGTGGCAGCTGTGGTATTCGGCTGCGGGACCCAGGCTCCTA
1487	VL of CDH19	artificial	ΑΑ	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
	21-14302 CC x 12C			CQAWESSTVVFGCGTKLTVL
1488	VH-VL of	artificial	Ħ	CGGCTGATCGAGGACATCTGCCCAGATGGGGCTGCCTGTGGGGAGGACGACCAGGTGCAGTTGGTGGAGTCTGGGGGGAGGCGT
	14302 CC x 12C			
	333			ACCATCTCCAGAGACAATTCCCAAGAACACGCTGTATCTGCAAATGAATAGCCTGAGAGCTGAGGACACGGCTGTGTATTACTGTGC
				GAGGCGGAGGATICTGGTGGCGGTGGTTCTGGCGGCGGGGGGGCTCCTCTATGAACTGAACTCAGCCACCTCAGTGTGTCCCAA CCacaacaacaacaacaacaacaacaacaacaacaaaaaa

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
o Š				
				TGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTACTGTCAGGCGTGGGAGAGCAGCACTGTGGTATTCGGCTGCGGG
1489		artificial	ΑA	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSYELTQPPSVSVSP
	14302 CC x I2C			GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCG TKLTVL
1490	_	artificial	aa	
	14302 CC x I2C			TISRDNSKNTLYLÖMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGGGGGGGGGGGGSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTRRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCG
				TKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISR
				CCSNICATED CONTROLL OF THE CON
1/0/1	CDH19 1/13/02	ortificial	o o	IVLHHHHHH OVOTVESGGGVVODGGSTRTSCAASGFTFSSVGMHWVROAPGKGTFWVDGSNKYVADSVKDRFTTSPDNSKNTTVTOMNST.
1		5	3	RAEDTAVYYCARRAGIIGTIGYYYGMDVWGOGTTVTVSSGGGGSGGGGGGGSSYELTOPPSVSVSPGOTASITCSGDRLGEKYT
	1)			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE
280				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
<u> </u>				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				OOKFGOAPRGLIGGIKFLAPGIPARFSGSLLGGKAALILSGVOPEDEAEYYCVLWYSNRWVFGGGIKLIVLRLIEDICLPRWGCLW EDDHHHHHH
1492	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
				RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAFGKGLEWVAKLRSKYNNYATYYADSVKDKFTLSKDDSKNTAYLQMNNLKTED TAXYVYNYBHONFCNSVISVMAVMOCTIXTYSSGGGGSGGGSGGGSGGGSGTTXTTTTTTTTTTTTTT
				OOKPGOAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVVPEDFAFFYCVLWYSNRWVFGGTKLTVLRLIEDICLPRWGCLW
				БОДНИНН
1493	_	artificial	nt	CAACGITICIGIACCGGICACITCGGIGGICIGIACCCGIGIAAIGGIGGIGGIGGIGGIICGCAGGIGCAGGIIGGIGGAGICIGG
	14302 x I2C x			GGGAGGCGTGGTCCAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTTCAGTAGCTATGCCATGCACTGGG
	FcBY			TUCGCUAGGUTUCAGGCAAGGGGCTGGAGTGGGTGGCATTTTATATGGTATGGAAGTAATAATAATAATACTATGGAAGT Oxoocxiiiiiicxaaxiiiiiiiiiiiiiiiiiiiiiiii
				THACTGTGCGAGAGGGCCGGTATAATAGGAACTATAGGCTACTACTACGGTATGGACGTCTGGGGCCAAGGGACCACGGTCACGG
				TCTCTAGT
1494		artificial	AA	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVK
	14302 × I2C ×			DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
	FCBY			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
8				
1495	VL of CDH19 14302 x I2C x FcBY	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
1496	VL of CDH19 14302 x I2C x FcBY	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
7641	VH-VL of CDH19 14302 x I2C x FcBY	artificial	ıt	CAACGTTTCTGTACCGGTCACTTCGGTGGTCTGTACCCGTGTAATGGTGGTGGTGGTGGTGCTGCAGGTGCAGTTGGTGGTGGTGGTGGGGGGTCTGGGGGGGG
1498	VH-VL of CDH19 14302 x I2C x FcBY	artificial	AA	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVL
1499	CDH19 14302 x I2C x FcBY	artificial	аа	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADASVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPCNGHHHHHH
1500	VH of CDH19 14302 CC x I2C x FcBY	artificial	t t	CAACGTTTCTGTACCGGTCACTTCGGTGGTCTGTACCCGTGTAATGGTGGTGGTGGTTCGCAGGTGCAGGTTGTGGTGGAGTCTGGGGGGGG
1501	VH of CDH19 14302 CC x I2C	artificial	Ą	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS

SFO	DESIGNATION	SOURCE	TYPE	SFOUENCE
O				
	x FcBY			
1502	VL of CDH19 14302 CC x I2C x FcBY	artificial	nt	TCCTATGAACTGACTCAGCCCACCCTCAGTGTCCCCTGGTCCCCAGGACAGCCAGC
1503	VL of CDH19 14302 CC x I2C x FcBY	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
282	VH-VL of CDH19 14302 CC x I2C x FCBY	artificial	nt	CAACGTTTCTGTACCGGTCACTTCGGTGGTCCTGTACCCGTGTAATGGTGGTGGTGGTGGTTCCAGGGTGCAGTTGGTGGAGTCTGGGGGGGG
1505	VH-VL of CDH19 14302 CC x I2C x FcBY	artificial	АА	QRFCTGHFGGLYPCNGGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVL
1506	CDH19 14302 CC x I 2 C x FcBY	artificial	aa	QRFCTGHFGGLYPCNGGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPCNGHHHHHH
1507	VH of CDH19 14303 CC x I2C	artificial	nt	CAGGTGCAGTTGGTGGAGTCTGGGGGGGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCTCT
1508	VH of CDH19	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
ID NO.				
	14303 CC x 12C			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1509	VL of CDH19 14303 CC x I2C	artificial	nt	TCCTATGAACTGACTCAGCCCCCCCCCGGTGTCCCCAGGACAGACA
1510	VL of CDH19 14303 CC x I2C	artificial	ΑA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
1511	VH-VL of CDH19 14303 CC x I2C	artificial	nt	CAGGTGCAGTTGGTGGAGGGGGAGGCGTGCTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCATTGGTATGGCATGCACTGGGTCCCCAGGCTCTCAGGTGAGTGGTGTGTATTATATATA
8 1512 %	VH-VL of CDH19 14303 CC x I2C	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVL
1513	CC x I2C	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1514	CDH19 14303 x F12q0	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKGRFTISRDDSKNTAYLQMNSLKTED TAVYYCVRHGNFGNSYVSWWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFTAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVL
1515	CDH19 14303 CC x F12q0	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSSVKGRFTISRDDSKNTAYLQMNSLKTED

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
8 8				
				TAVYYCVRHGNFGNSYVSWWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGTVTTTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVL
1516	CDH19 14303	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL
	x I2C-21			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
				SWIQQRFGQSFLLVIIQDIRRFSGIPERFSGSNSGNIATLIISGIQAMDEADIICQAWESSIVVFGGGIRLIVLSGGGGSEVQLVE SGGGLVOPGGSLKLSCAASGFTFNKYAMNWVROAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLOMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGUTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLRLIEDICLPRWGCLW EDDHHHHHH
1517	CDH19 14303	artificial	aa	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMKSL
	CC x I2C-21			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
				SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGIIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLRLIEDICLPRWGCLW EDDHHHHHH
28 28	VH of CDH19	artificial	nt	CAACGITICIGIACCGGICACTICGGIGGICIGIACCCGIGIAAIGGIGGIGGIGGIGGIICGCAGGIGCAGIIGGIGGAGICIGG
	14303 x I2C x			GGGAGGCGTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCTGTGCAGCGTCTGGATTCACCTTCAGTAGCTATGGCATGCACTGGG
	FcBV			TCCGCCAGGCTCCAGGCCAAGGGGCTGGAGTGGCTTGCATTTATATATGGTATGAGGGAAGTAATAAATA
)))			GACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAAAAGCCTGAGAGCTGAGGACACGGCTGTGTA
				TTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACTACGTATGGACGTCTGGGGCCAAGGGACCACGGTCACGGTCACCG
				TCTCTAGT
1519	VH of CDH19	artificial	Ą	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVK
	14303 x I2C x FcBY			DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1520	VL of CDH19	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTCCGTGTCCCCAGGACAGACA
	14303 x I2C x			AAAATATACTAGCTGGTATCAGCAGAGGCCAGGCCAGTCCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC
	FcBY			CTGAGCGATTCTCTGGCTCCAACTCTGGTAACACAGCCACTCTGACCATCAGCGGGACCCAGGCTATGGATGAGGCTGACTATTAC TGTCAGGCGTGGGAGAGCAGCACTGTGGGTATTCGGCGGAGGGACCAAGCTGACCGTCCTA
1521	VL of CDH19	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
	14303 x I2C x			CQAWESSTVVFGGGTKLTVL
	FcBY			
1522	VH-VL of	artificial	nt	CAACGITICIGIACCGGICACTICGGIGGICIGIACCCGIGIAAIGGIGGIGGIGGIGGIICGCAGGIGCAGGIIGGAGIIGGAGICIGG
	CDH19 14303 v I2C v EcBV			GGGAGGCGTGGTCCAGGCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTTTGGATTCACCTTCAGTAGCTATGGCATGCACTGGG TCCGCCAGGCTCCAGGCGAAGGGGCTGGAGTGGGTGTATATATA
	X 12 C A 1 CD 1			

SEQ NO	DESIGNATION	SOURCE	TYPE	SEQUENCE
5				GACCGATTCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCTGCAAATGAAAAGCCTGAGAGCTGAGGACACGGCTGTGTA TTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACGGTATGGACGTCTGGGGCCCAAGGGACCACGGTCACCG TCTCTAGTGGTGGCGGAGGATCTGGCGGAGGTGGAAGCGGAGGCGCGGATCTTCCTATGAACTGACTCAGCCACCCTCAGTGTCC GTGTCCCCAGGACAGACAGCCAGCATCACCTGCTCTGGAAGAGTTGGGGGAAAAATATACTAGCTGGTATCAGCAGAGGCCAGG CCAGTCCCCAGGACAGACATCTATCAAGATACCAAGCGGCCCTCAGGGAACCTGAGCGAATTCTCTGGCTCCAACTCTGGTAACA CAGCCCACTCTGACCATCAGCGGGCCCAGGCTATGGATGAGCTTATTACTGTCAGGCGAGGAGCACTCTGGTATTC GGCGGAGGGACCAAGCTCCAGCCTCTATCAAGATGAGCTTATTACTGTCAGGCGTGGGAGGCCACTCTGTGTTATTC GGCGGAGGGACCAAGCTCCAACCTCTA
1523	VH-VL of CDH19 14303 x I2C x FcBY	artificial	Ą	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVL
1524	CDH19 14303 x I2C x FcBY	artificial	о о	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVLSGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPHHHH
1525	VH of CDH19 14303 CC x I2C x FcBY	artificial	nt	CAACGTTTCTGTACCGGTCACTTCGGTGGTCCCGTGTAATGGTGGTGGTGGTGGTTCGCAGGTGCAGTTGGTGGAGTCTGGGGGGGG
1526	VH of CDH19 14303 CC x I2C x FcBY	artificial	AA	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1527	VL of CDH19 14303 CC x I2C x FcBY	artificial	nt	TCCTATGAACTGACTCAGCCCCCCCTCAGTGTCCCCAGGACAGACA
1528	VL of CDH19 14303 CC x I2C x FcBY	artificial	AA	SYELTQPPSVSVSPGGTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
1529	VH-VL of	artificial	nt	CAACGTTTCTGTACCGGTCACTTCGGTGGTCTGTACCCGTGTAATGGTGGTGGTGGTGGTTCGCAGGTGCAGGTTGGTGGTGGAGTCTGG

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
	CDH19 14303 CC x I2C x FcBY			GGGAGCCTGGTCCAGCCTGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTTCAGTAGCTATGGCATGCACTGGGGGTCCGGGGGAGGAGGTCCCTGGGGAAGTAATAATAATACTATGCCATGCACTGGGGGGCCCCAGGCCCAGGCCTGCAGGGCTCCTGAGGAGTCCGTGAAGGGCTCCAGGCTCCTGAGGCTCCTGAGGACTCCTGAGGCTCCTGAGGACTCCTGAGGACTCCTGAGGACACGGCTGTGTATTACTGTGCGAGAAGGCTGAGGACACGGCTGTGTATTAATAGGAACTATAGGAACTACTACTACGGTATGGACGTCTGGGGCCCAAGGGACCACGGTCACCGTTATAGTGGAGGCCGAGGACCTACTATAGGAGGCTACTACTACGGAGGCCTACTATGGGGGAGCCTCTATGGAGCCCAAGGGACCACCTCAGTGTCCCAGGACCCACCTCTAGTGGAGGCCAAGGCCAGGACCTCTGGTACTACTAGCTTGCTGGTATCAGCAGCAGAGCCAGGGCCAGGGCCAGGGCCAGGCCCTTTGCTGGTCATCTACAGCAGCAGCCAGGGCCTATGGTTACTGGTCATCTGGCAGAGCCAGGCCAGGCCAGGCCCAGGCCCTTTGCTGGTAACAACTTACTAGCTTGTATCAGCAGGCCAGGCCAGGCCAGGCCCTTTGCTGACATTACTGGCAGCAGCACTCTGGTAACAACATTACTGGCACCAACCTGAGCAACTTTCCAGCCGGACCAAGCCAGCACTGTGATATCGCTGCGCACCTCTGACCAACCTGCGCACCTCTGACCAACCTGCGCACCTCTGACCAACCTGCGCACCTCTGCCAACCTGTCCTGCTAATTCCTGCCACCTCTGACCAACCTGTCCTAATTACTTGCTCCTGGCAGCCACCTGTGCTAATTCGCTCCGCACCTCTGCTAATTCCTGCCACCACCTCTGCTAATTCCTGCCCAACCTGTCCTAATTCCTGCCCAACCTGTCCTAATTCCTGCCCAACCTGTGCTAATTCCTGCCCAACCTGTCCTAATTCCTGCCTCCAACCTGTGCTAATTCCTGCCCAACCTGTCCTAATTACTTCTGCCTCCAACCTGTCTGT
1530	VH-VL of CDH19 14303 CC x I2C x FcBY	artificial	AA	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVS VSPGQIASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVL
286	CDH19 14303 CC x I2C x FcBY	artificial	a a	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMKSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSGGCSQTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPCNGHHHHHH
1532	VH of CDH19 14039 CC x I2C	artificial	t	CAGGTGCAGTTGGTGGAGGTCTGGGGGAGGCGTGGTCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCCAGGCAAGTGTCTGGAGTGGGTGG
1533	VH of CDH19 14039 CC x I2C	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
1534	VL of CDH19 14039 CC x I2C	artificial	Ħ	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCCAGGACAGACA
1535	VL of CDH19 14039 CC x I2C	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
1536	VH-VL of CDH19 14039	artificial	nt	CAGGTGCAGTTGGTGGAGGTCTGGGGGGGGGGTGCTCCAGCCTGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGGATTCACCTT CAGTAGCTATGGCATGCACTGGGTCCGCCAGGCTCCAGGCAAGTGTCTGGAGTGGGTGG

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	CC × 12 C			AATACTATGCAGAGTCCGTGAAGGACCGATTCACCATCTCCAGAGACTAATTCCAAGAACACGCTGTATCTGCAAATGAATAGCCTG AGAGCTGAGGACCACGGTGTGTTTACTGTGCGAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACTACGGTGTCTG AGAGCTGAGGGACCACGGTCTCTTAGTGCGAGAAAGGGCCGGTATTAGGTTCTGCCGCGGAGGCTCTTG TGACTCAGCCACGCTCTCTTAGTGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGAGGCTCCTTTGAAC TGACTCAGCCACCCTCAGTCCCCTGTCCCCAGGACACACAC
1537	VH-VL of CDH19 14039 CC x I2C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVL
1538	CC x 12C	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
87	CDH19 14039 ×F12q0	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKGRFTISRDDSKNTAYLQMNSLKTED TAVYYCVRHGNFGNSYVSWWAYWQQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVL
1540	CC x F12q0	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLRLSCAASGFTFNSYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKGRFTISRDDSKNTAYLQMNSLKTED TAVYYCVRHGNFGNSYVSWWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
1541	VH of CDH19 21-14039 x 12C	artificial	nt	CGGCTGATCGAGGACATCTGCCTGCCCCAGATGGGGCTGCCTGTGGGAGGACGACGAGGTGCAGTTGGTGGAGTCTGGGGGGGG
1542	VH of CDH19 21-14039 x	artificial	AA	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS

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	ISC			
1543	3 VL of CDH19 21-14039 x 12C	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCCCAGGACAGACA
1544	4 VL of CDH19 21-14039 x 12C	artificial	AA	SYELTQPPSVSYSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVL
288 288	5 VH-VL of CDH19 21- 14039 x I2C	artificial	ŧ	CGGCTGAJCGAGGACATCTGCCCAGAJGGGGCTGCCTGTGGGAGGACCAGGTGCAGTTGGTGGAGTTGGTGGAGTCTGGGAGGCGT GGTCCAGCCTGAGGACTCTGCCTGTGCAGCGTTTGGATTCACCTTCAGTAGCTAJGGCTATGGCATGCACTGGGTCCGCCAGG CTCCAGCCAAGGGGCTGGAGTGGGTGCATTTATATGGTATGAGTATAATAATAATAATACTAJGCATGCAGTCCGTGAAGGACCGATTC ACCATCTCCAGAGACAATTCCAAGAACAGCGTGTATCTGCAAATGAATAAGCCTGAGGACACGGTCCGTGTGTTACTGTGC GAGAAGGGCCGGTATAATAGGAACTATAGGCTACTACGGTATGGAGGCTTGGGGGCCAAGGGACCACGGTCACCTTTAGTG GTGGCGGAGGCCGGTATAATAGGAACTTACTACGGCGAAATATACTTGGGGCCCAAGGGACCACGGTCCCCTTAGTG GTGGCGGAGGCCCACCCTCTGCAGCGGCGGCGGCGGATCTTCCTATGAACTTCAGCTGGTCACTGTCCCCC TTTGCTGGTCATCTATCAAGATACCAAGCGGCCTCAGGGATCCTGAGCGATTCTCTGGTATCAGGAACACTCCCC TTTGCTGGTCATCTATCAAGATACCAAGCGGCCTCAGGGATCCTGAGCGAGC
1546	5 VH-VL of CDH19 21- 14039 x I2C	artificial	AA	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGG TKLTVL
1547	7 CDH19 21- 14039 x I 2 C	artificial		RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGG TKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISR DDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYMAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGTVTL TCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKL TVLHHHHHH
1548	3 VH of CDH19 21-14039 CC x 12C	artificial	nt	CGGCTGAICGAGGACATCTGCCCTGCCCAGAIGGGGCTGCCTGTGGGAGGACCACCAGGTGCAGTTGGTGGAGTCTGGGGAGGCGT GGTCCAGCCTGGGGGGGTCCCTGAGACTCTCCTGTGCAGCGTTCACCTTCAGTAGCTATGGCATGCACTGGGTCCGCCCAGG CTCCAGGCAAGTGTCTGGAGTGGGTGGCATTTATATGGTATGAGGGAAGTAATAAATA
1549	9 VH of CDH19	artificial	AA	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
0 NO.				
	21-14039 CC x 12C			
1550	VL of CDH19 21-14039 CC x 12C	artificial	nt	TCCTATGAACTGACTCAGCCACCCTCAGTGTCCCCTGGACCAGACAGCCAGC
1551	VL of CDH19 21-14039 CC x 12C	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
289	VH-VL of CDH19 21- 14039 CC x I 2C	artificial	nt	CGGCTGATCGAGGACATCTGCCCAGATGGGGCTGCCTGTGGGAGGACCAGCTGGTGGAGTTGGTGGAGGTCTGGGGGAGGCGT GGTCCAGCCTGAGGACTCTCCCAGATGCAGCCTTCTGGATTCACCTTCAGTAGCTATGGCATGCACTGGGGCCCAGG CTCCAGGCCAAGTGGGGGGGCATTTATATGCTATGC
1553	VH-VL of CDH19 21- 14039 CC x I2C	artificial	ΑА	RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCG TKLTVL
1554	CDH19 21- 14039 CC x I2C	artificial		RLIEDICLPRWGCLWEDDQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVKDRF TISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSP GQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCG TKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISR DDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYMAYWCQGTLVTVSSGGGGGGGGGGGGGGGGGGGGTVVTGFFTTTT TCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKL TVLHHHHHH
1555	CDH19 14039 x I2C-21	artificial		QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG

SEQ ID NO.	CDH19 14039 x 12C x FCBY CDH19 14039 v 13C x FCBV	SOURCE	ТУРЕ	SEQUENCE DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVL QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYTOMNSTRAEDTAVYYCARRAGITGTTGYYYGMDVWGOGTTVYTVSSGGGGSGGGGSSYFLTDPPSVS
1564	X I Z C X FCB Y VH of CDH 19 14039 C C V 17 C	artificial	nt	USPGQTASITCSGDRIGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGTTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPCNGHHHHHHH CAACGTTTCTGTACCGGTCCAGTGGTGCTCTGTAATGGTGGTGGTGGTGGTTCGCAGGTTGGTGGAGTCTGGGAGGTCTGGGAGGTCTGGGAGGTCTGGGAGGTCTGGGAGGTCTGGGAGGTCTGGATTCAGTAGGCATGGAGTGGAGGGGGGGG
29	x FcBY	artificial	AA	GOGGEGOCGIGGICCAGGCGGGGGGGGGGGGGGGGGGGGGGGGGG
	x FcBY VL of CDH19	artificial	± }	DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS TCCTATGAACTGACTCAGCCACCCTCAGTGTCCGTGTCCCAGGACAGACA
8	14039 CC x 12C x FcBY		<u> </u>	AAAATATACTAGCTGGTATCAGCAGAGGCCAGGCCAGTCCCTTTGCTGGTCATCTATCAAGATACCAAGCGGCCCTCAGGGATCC CTGAGCGATTCTCTGGTAACACAGGCCAGCCACTCTGACCATCAGCGGGGCTATGGATGAGGCTATTAC TGTCAGGCGTGGGAGAGCAGCAGCTGTGGTATTCGGCTGCGGGACCAAGCTGACCGTCCTA
1567	VL of CDH19 14039 CC x I2C x FcBY	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGCGTKLTVL
1568	VH-VL of CDH19 14039 CC x I2C x FCBY	artificial	ļ t	CAACGITICIGIACOGGICACTICGGIGGICTACCCGTGIAATGGIGGIGGIGGITGGITGGTGCAGGIGCAGGITGGAGTCIGG GGAAGGCGITTCIGIACOGGICACTICGGIGGICTGTACCCTGIAATGGIGGITCGCAGGTGCAGGAGTCTGG GGGAGGCGIGGICCAGGCAAGGACTCTCCTGTGCAGGAGTAAIAAATAAATAGGAATGCAGGACTGGAG TCCGCCAGGCTCCCAGGACAATTCCAAGAACAGTTTAIATATGGAATGAATGAATAAATAAATAAATAAGGACACGGTTGTA TTACTGTGCGAAGAGGCCGGTATAATAGGAACTATAGGCTACTACTACGGACCTTGGGGCCCAAGGGACCACGGTCACCG TCTCTAGTGGAGGCCGGAGGATCTGGTGGCGGTGGTTTTGCGGCGGAGGCTCCTTTGGGGGCCAAGGGACCACGGTCACGG GTGTCCCCAGGACAGCCGAGCAGCAGCGGTGGTTCTGGCGGCGGGGGGCCCTCTGGGGGACCAGGGCCAGG GTGTCCCCAGGCCGACCAGCCAGCCAGCTCTGGAGATAGGTTGGGGGAAATATACTTGGGCGAGGCTCCAGGTGTACCAGGGCCAGG CCAGTCCCCAGGCCAGCCAGCCAACCCAAGCGGCCTCAGGGAAAATATACTTGGGCGAGACCCAACTGGTAACA CCAGTCCCCTTTGCTGGTCATCAAGATACCAAGCGGCCTCAGGGAAACATTACTTGCTGGAGACCAGCAGCTGTGGTAATTC CAGCCCACTCTGACCATCAGGGCTATGGATGAGGCTATTACTTCAGGCGAGACCAGCACTGTGGTAATTC CAGCCCACTCTGACCATCAGCGGCTATTGAGGCTATTACTTCAGGCGAGGCAGCCACTGTGGTAATTC CAGCCCACTCTGACCACCAGCGGACTATGAGCTATTACTTCACGGAGAGCAGCCACTGTGGTAATTC CAGCCCACTCTGACCACCAGCCACTATGAGCTATTACTTCACGGAGAGCCAGCC

SEQ O	DESIGNATION	SOURCE	TYPE	SEQUENCE
2				GGCTGCGGGACCAAGCTGACCGTCCTA
1569	VH-VL of CDH19 14039 CC x I2C x FcBY	artificial	AA	QRFCTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLIVL
1570	CDH19 14039 CC x I 2C x FcBY	artificial		QRECTGHFGGLYPCNGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYEGSNKYYAESVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSGTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQQRFCTGHFGGLHPCNGHHHHHH
292	CDH19 14302 x l2C-156	artificial	a a	TALLY MESSESSES CONTROLLED TO THE CONTROLLED TO THE CONTROLLED THE CONTROLLED TO THE CONTROLLED TO THE CONTROLLED THE CONTROLLED TO THE CONTROLLED THE CONTROLLED TO THE CONTROLLED TO THE CONTROLLED THE CONTROLLED TO THE CONTROLLED THE CONTROLL
1572	CDH19 14302 x I2C-LFcBY	artificial	aa	QRFVTGHFGGLYPANGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVLSGGGGSEVQLVESGGGIVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGSQRFCTGHFGGLHPHHHH
1573	CDH19 14302 x I2C-LFGBY- 156	artificial	o o	QRFVIGHFGGLYPANGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GGGTKLTVLSGGGGSEVQLVESGGGIVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGS QRFCTGHFGGLHPCNGGGGGSSGBDWDFDVFGGGTPVGGHHHHHHH
1574	CDH19 14302 x I2C-Cys-Loop	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE

SEQ ID NO.	DESIGNATION	SOURCE	ТҮРЕ	SEQUENCE
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
1575	CDH19 14302 × I2C-HALB	artificial	ва	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGTVLTUCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTVLTVLPGGGGSDAHKSEVAH RFKDLGEENFKALVLIAFAQYLQQCPFEDHVKLVNEVTFFAKTCVADESAENCDKSLHTLFFGDKLCTVATLRETYGEMADCCAKQE PERNECFLQHKDDNPNLPRLVRPEVDVWCTAFHDNEETFLKKYLYEIARRHPYFYAPELLFFAKRYKAAFTTCCQAADKAACLLPR LDELRDEGKASSAKQRLKCASLQKFGERAFKAWAVARLSQRFPKAEFASKLVTDLTKVHTECCHGDLLECADDRADLAKYICEN QDSISSKLKECCEKPLLEKSHCIAEVENDEMPADLPSLAADFVESKDVCKNYAEAKDVFLGMFLYEYARRHPDYSVVLLRLAKTY ETTLEKCCAAADPHECYAKVFDEFKPLVEEPQNLIKQNCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRRRLAKT HPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKCCTESLVNRRPCFSALEVDETYVPKEFNAETFFHADICTLSEKERQIKK QTALVELVKHKPRATKEQLKAVMDDFAAFVERCCKADDKETCFAEEGKKLVAASQAALGLDYHHHHHH
9251 293	CDH19 14302 x I2C-GS- D3HSA	artificial	ва	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGTVTTTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGTKLTVLPGGGGSEFPQNLIKQ NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC CTESLVNRRPCFSALEVDETYVPKEFNAEFTFHADICTLSEKERQIKKQTALVELVKHKPRATKEQLKAVMDDFAAFVEKCCKAD DKETCFAEEGKKLVAASQAALGLHHHHHH
1577	CDH19 14302 x I2C-3GS- D3HSA	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGGGTLVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
1578	CDH19 14302 x I2C-GS-	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGGGGGGGGGGSYELTQPPSVSVSPGQTASITCSGDRLGEKYT

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
	D3HSA-156			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGSEEPQNLIKQ NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC CTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFAAFVEKCCKAD DKETCFAEEGKKLVAASQAALGLGGGGSGGGSRDWDFDVFGGGTPVGGHHHHHH
1579	CDH19 14302 x I2C-3GS- D3HSA-156	artificial	e e	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG
0851 294	CDH19 14302 x I2C-6S- D3HSA-21	artificial	ee	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGTGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGSGGESPQNLIKQ NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC CTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPRATKEQLKAVMDDFAAFVEKCCKAD DKETCFAEEGKKLVAASQAALGLGGGSSGLIEDICLPRWGCLWEDDHHHHHH
1581	CDH19 14302 x I2C-3GS- D3HSA-21	artificial	ت ت	QVQIVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGSGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGSGGGGGGGGGGGGGSGGGSGGGSPRLIKQNCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEK TPVSDRVTKCCTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHRPKATKEQLKAVMDDFA AFVEKCCKADDKETCFAEEGKKLVAASQAALGLGGGSGGGSGGSRLIEDICLPRWGCLWEDDHHHHHH
1582	CDR-H1 of CDH19	artificial	AA	SYYWS

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
	65231.002			
1583	CDR-H2 of CDH19 65231.002	artificial	AA	YIYYSGSTNYNPSLKS
1584	CDR-H3 of CDH19 65231.002	artificial	AA	DQRRIAAAGTHFYGMDV
1585	CDR-L1 of CDH19 65231.002	artificial	AA	RASQSVSSSYLA
1586	CDR-L2 of CDH19 65231.002	artificial	AA	GISSRAT
295	CDR-L3 of CDH19 65231.002	artificial	AA	QQYGSSPFT
1588	VH of CDH19 65231.002	artificial	TN	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTAGGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGACTCCAT CACCTCCTACTACTAGTCCTGGATCCGGCAGCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACTCCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGAACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCGTGACC GCCGCTGACACCCCAGGCCGTGTACTGCGCCCAGGAACCAGCTGCCGCGCACCCACGTGTCTCCGTGAACCTGTCTCTACGGCATGTGTGTG
1589	VH of CDH19 65231.002	artificial	ΑA	QVQLQESGPGLAKPSETLSLTCTVSGDSITSYVWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSS
1590	VL of CDH19 65231.002	artificial	L	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCCCTGTCCTGCAGAGCCTCCCAGTCCGT GTCCTCCTCCTACCTGGCCTGG
1591	VL of CDH19 65231.002	artificial	ΑA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSPFTFGGGTKVEIKS
1592	VH-VL of CDH19 65231.002	artificial	TN	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTAGGCCCTCCGAGACACTGTCCCTGACCTGCCCGGTGTCCGGCGACTCCAT CACCTCCTACTACTGGTCCTGGATCCGGCAGCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACTCCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACACCTCCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCGTGACC GCCGCTGACCCCAGCGCTGTACTGCGCCAGGGACCAGCGGAATCGCCGGCACCCGCGCACCCTTCTACGGCATGGATGTGTG GGGCCAGGGCACCCTCGTGACCGTGTCTAGCGGAGGCGGAGGCTCTGGTGGCTGCCGCGGGGGCTCCCGAGATCGTGC

SEQ NO	DESIGNATION	SOURCE	TYPE	SEQUENCE
				TGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGCCACCCTGTCCTGCAGAGCCTCCCAGTCCGTGTCCTCCTCCTCCTCCTCCTCCTCCTCCTCCTCC
1593	VH-VL of CDH19 65231.002	artificial	АА	QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSSGGGGSGGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPFTFGGGTKVEIKS
1594	CDH19 65231.002 x I2C	artificial	AA	QVQLQESGPGLAKPSETLSLTCTVSGDSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSSGGGGSGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSTDFTLTISRLEPEDFAVYYCQQYGSSPFTFGGGTKVEIKSGGGSEVQL VESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKT EDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPN WVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSCVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1595	CDR-H1 of CDH19 65231.003	artificial	АА	SYYWS
96 1596	CDR-H2 of CDH19 65231.003	artificial	АА	YIYYSGSTNYNPSLKS
1597	CDR-H3 of CDH19 65231.003	artificial	АА	DQRRIAAAGTHFYGMDV
1598	CDR-L1 of CDH19 65231.003	artificial	АА	RASQSVSSSYLA
1599	CDR-L2 of CDH19 65231.003	artificial	АА	GISSRAT
1600	CDR-L3 of CDH19 65231.003	artificial	АА	QQYGSSPFT
1601	VH of CDH19 65231.003	artificial	H	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGCCTCCGAAGCCCTGTGTCCCTGACCTGACCTGCACCTGTGCACCGTGTCCGGCGCGCTCCAT CACCTCCTACTACTACTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTCCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCGTGACC

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				GCCGCTGACACCCGCCGTGTACTACTGCGCCAGGGACCAGCGGAATCGCCGCTGCCGGCACCCACTTCTACGGCATGTGTGTG
1602	VH of CDH19 65231.003	artificial	Ą	QVQLQESGPGLAKPSETLSLTCTVSGGSITSYWWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSS
1603	VL of CDH19 65231.003	artificial	LN	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCTGCAGAGCCTCCCCGTCCGT
1604	VL of CDH19 65231.003	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSPFTFGQGTKVEIKS
1605	VH-VL of CDH19	artificial	TN.	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCTGGCCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGCGCGC
297	65231.003			ACTACAACCCCAGCTGAAGTCCAGAGTGACCATCTCCGTGGACAACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC GCCGCTGACACCCCCGCGCTGTACTACTGCGCCAGGGACCAGCGCTGCCGGCCCCCAGTTCTACGGATGTGTG GGCCCAGGGCACCCTCGTGACCGTGTCTAGCGAGGCCGGAGGATCTTGGTGGTGGTTCTGGCGGCGCGGCGGGGGCTCCGAGATCGTGC TGACCCAGGCCACCCTCGTCCCTGAGCCGAGGCGGAGGCTCTGCTGCTGCTGCTGCTCCTCC TACCTGGCCTGG
1606	VH-VL of CDH19 65231.003	artificial	AA	QVQLQESGPGLAKPSETLSLTCTVSGGSITSYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTIVTVSSGGGGSGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPFTFGQGTKVEIKS
1607	CDH19 65231.003 x 12C	artificial	AA	QVQLQESGPGLAKPSETLSLTCTVSGGSITSYYWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTAVYYCARDQRRIAAAGTHFYGMDVWGQGTLVTVSSGGGGSGGGGGGGGGSGGGGSTLTTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGTSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPFTFGQGTKVEIKSGGGSEVQL VESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKT EDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGCGUVTVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPN WVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHH
1608	CDR-H1 of CDH19 65234.001	artificial	АА	SYYWS
1609	CDR-H2 of CDH19 65234.001	artificial	АА	YIYYIGSTNYNPSLKS
1610	CDR-H3 of	artificial	AA	DSRYRSGWYDAFDI

TYPE SEQUENCE		AA RASQSVAGSYLA	AA GASSRAT	AA QQYGKSPIT	NT CAGGTGCAGCTGCAGGAATCCGGCCCTGGCTGAGCCCTCCGAGACACTGTCCCTGACCTGTGTCCGGCGGCTCCAT CAACTCCTACTACTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACTACTACTACTACTCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACCCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC GCCGCTGAACCCCTGTACTGCGCCAGAGACTCCCGGTACAGATCCGGGTGGTACGACGCTTCGAACATCTGGGGCCAGGG CACCATGGTCACCGTGTCCTCT	AA QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSS	NT GATATCGTGACCCAGTCCCCCGGCACCCTGTCTGAGCCCTGGCGAGAGAGCCCTCTGTCTG	AA DIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV	NT CAGGTGCAGCTGCAGGATCCGGCCCTGGTCAAGCCCTCCGAGACACTGTCCTGACCTGTACTGCTGTCCGGCGGCTCCAT CAACTCCTACTACTACTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACATCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACACCTCCAGAGACCTGCTGCTCCTCCGTGACC GCCGCTGACCCCTGTACTACTGCGCCAGAGACTCCCGGTACAGAACCGGGTACGACGCTTCCTCCTGGGGCCTGCTCCTGGGGCCAGGG CACCATGGTCACCCGTGTCCTGGCGAGACTCCGGTACAGACCGGGTGGTACGACGCTTCGACATCTGGGGGCCAGGG CACCATGGTCACCTGTCTCTGGCGAGAGCCTCTGGCGGAGCTGCAGGCCGCTCTCAGTCCGTGGCCGGCTGCCTGGCCCAGG CCCCGGCACCCTGTCTCTGGCGAGAGCCTCTGGCGGAGCCTCTTCTAGAGCCCGTGTACTACTGCCGGCTTCTCTGGCCTGACCCGGTTCTCTGGCCGGCTTCTCTAGAGCCCGGCTTCTCTAGAGCCCGGCTTCTCTGGCCTGACCCGGTTCTCTCTAGAGCCCCTGTACTTTGCCAGCCGGTTCTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGACCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTAGAGCCCCTTCTCTCTAGAGCCCTTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGAGCCCCTTCTCTCTAGACCCCTTCTCTCTAGAGCCCTTCTCTCTAGAGCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTAGACCCCTTCTCTCTC	AA QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSGGGGGGGGGGGGDIVLTQSPGTLSLSCRATLSCRASQSVAGSYLA
SOURCE		artificial	artificial	artificial	artificial	artificial	artificial	artificial	artificial	artificial
DESIGNATION	CDH19 65234.001	CDR-L1 of CDH19 65234.001	CDR-L2 of CDH19 65234.001	CDR-L3 of CDH19 65234.001	VH of CDH19 65234.001	VH of CDH19 65234.001	VL of CDH19 65234.001	VL of CDH19 65234.001	VH-VL of CDH19 65234.001	VH-VL of
SEQ ID NO.		1611	1612	1613	1614	²² 1615	1616	1617	1618	1619

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
⊇ 8				
	CDH19 65234.001			WYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKS
1620	CDH19 65234.001 x 12C	artificial	Ą	QVQLQESGPGLVKPSETLSLTCTVSGGSINSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARDSRYRSGWYDAFDIWGQGTMVTVSSGGGGSGGGGGGGGGDIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLA WYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALITLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1621	CDR-H1 of CDH19 65234.004	artificial	AA	SYYWS
1622	CDR-H2 of CDH19 65234.004	artificial	АА	YIYYIGSTNYNPSLKS
299	CDR-H3 of CDH19 65234.004	artificial	AA	ESRYRSGWYDAFDI
1624	CDR-L1 of CDH19 65234.004	artificial	AA	RASQSVAGSYLA
1625	CDR-L2 of CDH19 65234.004	artificial	AA	GASSRAT
1626	CDR-L3 of CDH19 65234.004	artificial	AA	QQYGKSPIT
1627	VH of CDH19 65234.004	artificial	L	CAGGTGCAGCTGCAGGAATCCGGCCCTGGCCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCCAT CAGCTCCTACTACTACTGGTCCTGGATCCGGCAGCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACTACATCGGCTCCACA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCGTGACC GCCGCTGACACCCCTGTACTACTGCGCCAGAGAGTCCCGGTACAGAACCGGGTGCTCCTGGGGCCAGGG CACCATGGTCACCGTGTCCTCT
1628	VH of CDH19 65234.004	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSS
1629	VL of CDH19	artificial	N	GATATCGTGCTGACCCAGTCCCCCGGCACCCTGTCTCTGAGCCCTGGCGAGAGAGCCCTGTCCTGCAGAGAGAG

ATION SOURCE TYPE SEQUENCE DH19 artificial AA DIVLTQSPGT TACTATTGCC DO4 artificial AA DIVLTQSPGT TYCQQYGKSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYGRSP TYCQQYBCGC TYCCTGGC TYCCTTGC TYCCTTTGC TYCCTTGC TYCCTTG					
65234.004 A GGCCGCCTCGA GCA24.004 artificial AA DIVLTQSPGT GCA16.004 A YYCQXYGKSF GCA34.004 artificial NT CAGGTGCAGG GCA34.004 artificial NT CAGGTGCAGG GCA34.004 artificial AA ACTACAACCC GCA34.004 artificial AA GCGCGGCAC GCA34.004 artificial AA GVQLQESGPG GCA35.005 artificial AA SYYCVRHGN GCA35.005 artificial AA SYYCVRHGN GCA35.005 artificial AA GGGLUMLHLL GCDH19 G5235.005	SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
1630 VL of CDH19 artificial AA DIVLTQSPGT 1631 VH-VL of artificial NT CAGGTGCAGC CDH19 ACTACAACCC GCGCTGACACCC 65234.004 ACTACAACCC GCGCTGACACCC 65234.004 ACTACAACCC GCCCTGGCACACCC 1632 VH-VL of artificial AA QVQLQESGPG 65234.004 artificial AA QVQLQESGPG 1633 CDH19 AADTALYYCAGC GGGTCTGGC 65234.004 x artificial AA QVQLQESGPG 12C GGGLVQPGGS AADTALYYCAGC AADTALYYCAGC 12C CDH19 AADTALYYCAGC AADTALYCAGC 1634 CDR-H1 of artificial AA SYF1H 1635 CDR-H2 of artificial AA SYF1H 1635 CDR-H3 of artificial AA SYF1H 1636 CDR-H3 of artificial AA SYF1H 1637 CDH19 GG1QLWIHILD GG1QLWIHILD <tr< th=""><th></th><th>65234,004</th><th></th><th></th><th>GGCCGGCTCCTACCTGGCTTGGTATCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCCCTCTTCTAGAGCCACCGGCGCCGCCTCTTCTAGAGCCACCGGCCGCTCTCTAGAGCCACCGGCTCCTGACCCTGACCGTGGACCTTCGCCGTGTTCTAGACCATCGCCGGCTGGAAACCGAGGAACCCGGCTGCTTCGCCGTGTATTGCCAGCAGAAATGAAGTACCAGCAAAATGAAGTACC</th></tr<>		65234,004			GGCCGGCTCCTACCTGGCTTGGTATCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCCCTCTTCTAGAGCCACCGGCGCCGCCTCTTCTAGAGCCACCGGCCGCTCTCTAGAGCCACCGGCTCCTGACCCTGACCGTGGACCTTCGCCGTGTTCTAGACCATCGCCGGCTGGAAACCGAGGAACCCGGCTGCTTCGCCGTGTATTGCCAGCAGAAATGAAGTACCAGCAAAATGAAGTACC
1631 VH-VL of CDH19 artificial NT CAGGTGCAGC CACCTCTAC CACCTCTAC CACCTCTACCTCTACCTCTACCTCTACCTCTACTCTACTCACCTCTACTCACTCACTCTACTCACTACT	1630	VL of CDH19 65234.004	artificial	Ą	
March Marc	1631	VH-VL of CDH19	artificial	TN	
1632 VH-VL of artificial AA TGGTATCAGC CGCTCTGGC GCAAGTCCCC GCAAGTCCCCGCACAC AADTALYYCAAA GCGCAAGTCCCC GCAAGTCCCCGCACAC AADTALYYCAAA AADTALYYCAAAC GCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAGTCCCCGCAAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCCGCAACTCCCGCAACTCCCCGCAACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCGCACACTCCCCCGCACACTCCCCGCACACTCCCCCGCACACTCCCCCGCACACTCCCCCGCACACTCCCCCGCACACTCCCCCGCACACTCCCCCGCACACTCCCCCCGCACACTCCCCCCGCACACTCCCCCCCC		65234.004			
1632 VH-VL of artificial AA CDH19 65234.004 1633 CDH19 artificial AA 1634 CDR-H1 of CDH19 artificial AA 1635 CDR-H2 of CDH19 artificial AA 1635 CDR-H3 of CDR-H3 of CDH19 artificial AA 1636 CDR-H3 of CDR-H3 of CDH19 artificial AA 1637 CDR-L1 of CDR-L1 of CDR-L1 of CDH19 artificial AA					CCCCCGGCACCCTGTCTCTGAGCCCTGGCGAGAGAGACCACCCTGTCTGCAGAGCCTCTCTCAGTCCGTGGCCGGCTCCTACCTGGCT TGGTATCAGCAGAAGCCCGGCCCCGGCCCCTCTGCTGCTGCTGCTGCGCGCCCTCTTCT
1632 VH-VL of CDH19 artificial AA CDH19 artificial AA 1633 CDH19 artificial AA 12C artificial AA 1634 CDR-H1 of CDH19 artificial AA 65235.005 artificial AA 1635 CDR-H2 of CDR-H3 of CDH19 artificial AA 65235.005 artificial AA 1636 CDR-H3 of CDR-H3 of CDH19 artificial AA 1637 CDR-L1 of CDR-L1 of CDH19 artificial AA					GGCAGTCCCCCATCGGCCAGGGAACCCGGCTGGAATGAAGTCC
1633 CDH19 artificial AA 65234.004 x CDR-H1 of artificial CDH19 CDH1		VH-VL of CDH19 65234,004	artificial	ЧΑ	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNQFSLKLSSVT AADTALYYCARESRYRSGWYDAFDIWGQGTMVTVSSGGGGGGGGGGGGGGDIVLTQSPGTLSLSPGERATLSCRASQSVAGSYLA WYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKS
CDR-H1 of artificial AA CDH19 65235.005 artificial AA CDH19 65235.005 CDR-H3 of CDH19 65235.005 CDR-H3 of CDH19 65235.005 CDR-H1 of artificial AA CDH19 65235.005 CDR-H1 of artificial AA CDH19		CDH19 65234 004 x	artificial	AA	OVOLOESGPGLVKPSETLSLTCTVSGGSISSYYWSWIROPPGKGLEWIGYIYYIGSTNYNPSLKSRVTISVDTSKNOFSLKLSSVT AADTALYYCARESRYRSGWYDAFDIWGOGTMVTVSSGGGGSGGGGSGDIVLTOSPGTLSLSPGERATLSCRASOSVAGSYLA
CDR-H1 of CDR-H1 of CDH19 artificial AA AA SYFIH CDH19 AA IINPISVSTS CDH19 AA IINPISVSTS CDR-H3 of CDR-H3 of CDH19 AA GGIQLWLHID CDH19 AA GGIQLWLHID CDR-L1 of artificial AA AA GGIQLWLHID CDR-L1 of artificial AA SGSRSNIGSN		12C			WYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGKSPITFGQGTRLEMKSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ OKPGOAPRGIIGGTKFIAPGRPARFSGSIIGGKAAITTSGVOPEDFAFYYCVIMYSNRWYFGGGTKITVIHHHHHH
CDH19 G5235.005 Antificial AA IINPISVSTS CDH19 65235.005 Artificial AA GGIQLWLHID CDR-H3 of CDH19 GGIQLWLHID CDH19 CDR-L1 of artificial AA GGIQLWLHID CDR-L1 of artificial AA SGSRSNIGSN CDH19 CDR-L1 of artificial AA SGSRSNIGSN	1634	CDR-H1 of	artificial	AA	
CDR-H2 of CDR-H2 of CDH19 artificial AA A IINPISVSTS CDH19 65235.005 CDR-H3 of CDH19 661QLWLHLD CDH19 CDR-L1 of artificial AA AA GGIQLWLHLD CDR-L1 of artificial AA SGSRSNIGSN CDH19		CDH19 65235.005			
CDH19 65235.005 artificial AA GGIQLWLHLL CDH19 65235.005 CDR-L1 of artificial AA SGSRSNIGSN CDH19 CDH1	1635	CDR-H2 of	artificial	ΑA	IINPISVSTSYAQKFQG
CDR-H3 of artificial AA GGIQLWLHILD CDH19 65235.005 CDR-L1 of artificial AA SGSRSNIGSN CDH19 CDH19 AA SGSRSNIGSN		CDH19 65235.005			
CDH19 65235.005 AA SGSRSNIGSN CDH19 CDH19	1636	CDR-H3 of	artificial	AA	GGIQLWIHIDY
CDR-L1 of artificial AA SGSRSNIGSN CDH19		CDH19 65235.005			
	1637	CDR-L1 of CDH19	artificial	AA	SGSRSNIGSNFVN

SEQ D	DESIGNATION	SOURCE	TYPE	SEQUENCE
2				
	65235.005			
1638	CDR-L2 of CDH19 65235.005	artificial	AA	TNNQRPS
1639	CDR-L3 of CDH19 65235.005	artificial	AA	ATYDESMQGWV
1640	VH of CDH19 65235.005	artificial	L L	CAGGTGCAGCTGGTGCAGTCTGGCGCGCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTACTTCATCCACTGGGTCCGACGGCCCCAGGCCTGGGAATGGATGG
1641	VH of CDH19 65235.005	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
1642	VL of CDH19	artificial	۲	CAGTCTGCCCTGACCCAGCCTCCCTCCGTCACCGGCACACCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
301	65235.005			CGGCTCCAACTTCGTGAACTGGTACCAGCAGCTGCCCGGCACCGCCCCCAAGGTGCTGATCTACACCAACAACAGGGGCCCTCCGGCCTCCGGCTCCGACCAGCGGCCCTCCGACCACCACCACGAGGCCGACCAACAACAACAACAACA
1643	VL of CDH19 65235.005	artificial	AA	QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATYDESMQGWVFGGGTKLTVLS
1644	VH-VL of CDH19 65234.005	artificial	L L	CAGGTGCAGCTGGTGCAGTCTGGCGCGCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCCAGGGCCTGGAATGGATGG
				GGTCACCGTGTCTCTGGTGGCGGAGGCTCTGGCGGAGGTGGAAGCGGAGGCGGCGGGGTCCCAGTCTGCCCTGACCCAGCCTCCCT CCGTCACCGGCACCGCCCAGCGCGTGACCATCTCCTGCTCCGGCTCCAACATCGGCTCCAACTTCGTGAACTGGTAC CAGCAGCTGCCCGGCACCGCCCCCCAAGGTGCTGATCTACACCAACAACAGCGGGCCCTCCGGCGTGCCCGACCGGTTCTTCTGGCTC CAAGTCTGGCACCTCCCTGGCCATCTCCGGCCTGCAGTCCAAGGACGAGGCCGACTACTACTACGACGAGT CCATGCAGGGCTGGCTGGCGGAGGCCCAAGCTGCCGACCTACGACGAGT
1645	VH-VL of	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL
	CDH19 65234.005			RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGGGGGGSQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLS
1646	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIOLWLHIDYWGOGTIAATVSSGGGGSGGGGGGGGGSOSALTOPPSVTGTPGORVTISCSGSRSNIGSNFYAWY
	03234.003 A			QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLSGGGGSEVQLVES

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
	12C			GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1647	CDR-H1 of	artificial	AA	SYFIH
	CDH19			
	65235.002			
1648	CDR-H2 of	artificial	¥	IINPISVSTSYAQKFQG
	CDH19 65235 002	_		
1649	CDR-H3 of	artificial	AA	GGIQLWLHLDY
	CDH19			
	65235.002			
1650	CDR-L1 of	artificial	AA	SGSRSNIGSNFVN
	CDH19			
	65235.002			
1651	CDR-L2 of	artificial	Ą	TNNQRPS
	CDH19			
	65235.002			
1652	CDR-L3 of	artificial	Ą	ATWDDSMNGWV
	CDH19 65235.002			
1653	VH of CDH19	artificial	N	CAGGIGCAGCIGGIGCAGICIGGCGCCGAAGIGAAGAAACCIGGCGCCICCGIGAAGGIGICCIGCAAGGIGICCGGCIACACCII
	65235.002			CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCCAGGCCAGGGCCTGGAATGGATGG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCACCTGGACTATTGGGGCCAGGGCACCCT GGTCACCGTGTCCTCT
1654	VH of CDH19 65235.002	artificial	АА	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
1655	VL of CDH19	artificial	NT	CAGTCTGCCCTGACCCAGCCTCCCTCCGTCACCGGCACACCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65235.002			CGGCTCCAACTTCGTGAACTGGTACCAGCAGCTGCCCGGCACCGCCCCCAAGGTGCTGATCTACACCAACAACCAGCGCCCTCCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACTCCGCCTCCCTGGCCATCTCCGGCCTGCAGGTCGAGGACGAGGCCGAC TACTACTGTGCCACCTGGGACGACGACTCCATGAACGGCTGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1656	VL of CDH19	artificial	AA	OSALTOPPSVTGTPGORVTISCSGSRSNIGSNFVNWYOOLPGTAPKVLIYTNNORPSGVPDRFSGSKSGTSASLAISGLOSEDEAD
)))	65235.002	5		YYCATWDDSMNGWVFGGGTKLTVLS

SEQ NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
1657	VH-VL of CDH19 65235.002	artificial	۲ ا	CAGGTGCAGCTGGTGCAGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTGCTGCTGGTCCGACGGCCCAGGCCCAGGCCTCGGAATGGATGG
1658	VH-VL of CDH19 65235.002	artificial	АА	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDDSMNGWVFGGGTKLTVLS
303	CDH19 65235.002 x 12C	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGGSQALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDDSWNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ
1660	CDR-H1 of CDH19 65235.003	artificial	АА	SYFIH
1661	CDR-H2 of CDH19 65235.003	artificial	АА	IINPISVSTSYAQKFQG
1662	CDR-H3 of CDH19 65235.003	artificial	AA	GGIQLWLHLDY
1663	CDR-L1 of CDH19 65235.003	artificial	АА	SGSRSNIGSNFVN
1664	CDR-L2 of CDH19 65235.003	artificial	AA	TNNQRPS
1665	CDR-L3 of	artificial	AA	ATWDESMQGWV

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
_ 0 0 0 0				
	CDH19 65235.003			
1666	VH of CDH19	artificial	N	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGGCTACACCTT
	65235.003			CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCCAGGCCAGGGCCTGGAATGGATGG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCACCTGGACTATTGGGGCCAGGGCACCTTGGACTATTGGGGCCAGGGCACCTT
1667	VH of CDH19 65235 003	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL RSEDTAVYYCARGGIQLWLHLDYWGOGTLVTVSS
1668	VL of CDH19	artificial	N	CAGTCTGCCCTGACCCAGCCTCCCTCCGTCACCGGCACACCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65235.003			CGGCTCCAACTTCGTGAACTGGTACCAGCAGCTGCCCGGCACCGCCCCCAAGGTGCTGATCTACACCAACAACCAGCGGCCCTCCG
				GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCGCTGGCCATCTCCGGCCTGCAGTCCGAGGACGAGGCCGAC TACTACTGTGCCACTGCAGTCCGAGGACGAGGCCGAC
1669	VL of CDH19 65235 003	artificial	ΑΑ	QSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATWDESMOGWYFGGGTKLTVLS
1670	VH VI of	Lioiai+rc	ΤΙΛ	一番のできた。
Q 404	VII-VE OI	alcia	2	くされていることできない。このでは、このできないできないできないできない。このできないできないできないできないできない。このできないできないできない。このできないできないできない。このできないできないできない。このできないできないできない。このできないできない。このできないできないできない。このできないできない。このできないできない。このできないできない。このできないである。このできないできない。このできないできない。このできないでは、このでは、このでは、このでは、このでは、このでは、このでは、このでは、この
	65235 003			CCTCCTACGCCCAGAAATTCCAGGGCAGAGTGACCATGACCCGGGACACCTCCACCTCCACCGTGTTCATGGAACTGTCCTCCTCCTC
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCCAGCTGTGGCTGCACCTGGACTATTGGGGCCCAGGGCACCTT
				GGTCAUCGTGTCTCTGGTGGCGGAGGATCTGGCGGAGGTGGAAGCGGAGGCGGCGGCGGCGGATCTCTGCCCTGACCCAGGCTTCTCT CCCTTAACCCCCCAACTTCTCCCCTTAACCTCTCTTTTCTTC
				CCGICACCGGCACACCIGGCCAGCGCGIGACCAICICCIGCICCGGCICCGGICCAACAICGCCICCAACAICGGCICCAACAIGGGAACIGGTAC
				CAGLAGLIGULUGGLALUGULULUAAGGIGUIGAIGIALIAALLAALAALLAGLGGGLULULGGLGIGULUGALUGAITUILIGGLIG CAAGICIGGCACCICCGCCICCCIGGCCAICICCGGCCIGCAGICCGAGGACGAGGCCGACIACIACIGIGCCACCIGGGACGGAGGI
				CCATGCAGGCTGGTGTTCGGCGGAGCACCAAGCTGACCGTGCTGTCC
1671	Jo JV-HV	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL
	CDH19 65235 003			RSEDTAVYYCARGGIQLMLHLDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLS
1672	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVFMELSSL
	65235.003 x			RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGALTQPPSVTGTPGQRVTISCSGSRSNIGSNFVNWY
	12C			QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDESMQGWVFGGGTKLTVLSGGGGSEVQLVES GGGIMOPGGSIKISCAASGFTFNKYAMNWYBOAPGKGIEMMABIRSKYNNYATVYAPSVKDRFTISPDDSKNTAYIOMNNIKTFDT
				AVYYCVRHGNEGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ OKPGOAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVOPEDEAEYYCVLWYSNRWYFGGGTKLTVLHHHHHH
1673	CDR-H1 of	artificial	AA	SYAMN
	CDH19			

SEQ ID NO.	DESIGNATION	SOURCE	ТҮРЕ	SEQUENCE
	65236.001			
1674	CDR-H2 of CDH19 65236.001	artificial	AA	TISGGGANTYYADSVKG
1675	CDR-H3 of CDH19 65236.001	artificial	AA	GGMGGYYYGMDV
1676		artificial	AA	RASQSISSNLA
1677	CDR-L2 of CDH19 65236.001	artificial	AA	GAFTRAT
305	CDR-L3 of CDH19 65236.001	artificial	AA	QQYNYWPLT
1679	VH of CDH19 65236.001	artificial	LN L	CAGGTGCAGCTGCAGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGCCATGAACTGGGTCCGACAGGCCCTGGCAATGGGTGTCCACCCATCAGCGGCGGCGGCGCCAACA CCTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGCGACAACTCCAAGTCCACCTGTACTGCAGATGAACTCCCTG AGAGCCGCCGACACCGCCGTGTACCACTGTGCTAAGGGCGGCGATGGGCGGCGCTACTACTACGGCATGATGTGTGGGGCCACGGGCCACCGTGTACTACGGCCCAGGGCCACCGTGTACTACGGCCCAGGGCCACCGTGTACTACGGCCACGGGCCACCGTGTACTACGGCCACGGGCCACCGTGTACTACGGCCACGGGCCACCGTGTACTACGCCACGGGCCACCGTGTACTACCTAC
1680	VH of CDH19 65236.001	artificial	ΑA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS
1681	VL of CDH19 65236.001	artificial	L	GAGATCGTGATGACCCAGTCCCCCGTGACCCTGACCCTGAGCCTGGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTCCAT CTCCAGCAACCTGGCTTCCAGCAGAAGCCCGGCCAGGCCCTTCGGCTGTTACCGGGCCTTTTACCCGGGCCACCGGCA TCCCTGCCAGAGTGTCTGGCTCCGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGCAGTCCAGGGACTTTGCCGTGTAC TACTGCCAGCAGTACAACTACTGGCCCCTGACCTTCGGAGGCGGCACCAAGGTGGAAATCAAGTTC
1682	VL of CDH19 65236.001	artificial	ΑA	EIVMTQSPVTLSLSGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVY YCQQYNYWPLTFGGGTKVEIKS
1683	VH-VL of CDH19 65236.001	artificial	TN	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGCCATGAACTGGGTCCGACAGGCCCTGGCAAGGGCCTGGAATGGGTGTCCACCATCAGCGGCGGAGGGCCCAACA CCTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGCGACAACTCCAAGTTCCACCTGTACTGCAGATGAACTCCCTG AGAGCCGCCGACACCCGCGTGTACCACTGTGCTAAGGGCCGGCGACAACTCCAAGTACTACGGCATGGATGTGTGGGGCCACCCTC CACCGTGACCGACACCGCGGAGGATCTGGCCGTGGTGCTGTTCTGGCGGCGCGCGC

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
				COGTGACCCTGTCCCTGAGCCTGGGGGAGAGGCCACCCTGTCTTGCCGGGCCTCCCAGTCCTCCAGCAACCTGGCCTGGTTC CAGCAGAAGCCCGGCCCAGGCCCTCGGCTGATCTACGGCGCCTTTACCCGGGCCACCGGCATCCTGCCAGAGTGTCTGGCTC CGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCTGCAGTCCGAGGACTTTGCCGTGTACTACTGCCAGCAGTACAACTACT GGCTCCGGCACCTTCGGAGGCGCGCACCAAGGTGGAAATCAAGTCC
1684	VH-VL of CDH19 65236.001	artificial	АА	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1685	CDH19 65236.001 x I2C	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAADTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSEIVMTQSPVTL.SLSLGERATL.SCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGQQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1686	CDR-H1 of CDH19 65236.007	artificial	АА	SYAMN
1687	CDR-H2 of CDH19 65236.007	artificial	AA	TISGGGANTYYAESVKG
1688	CDR-H3 of CDH19 65236.007	artificial	АА	GGMGGYYYGMDV
1689	CDR-L1 of CDH19 65236.007	artificial	АА	RASQSISSNLA
1690	CDR-L2 of CDH19 65236.007	artificial	АА	GAFTRAT
1691	CDR-L3 of CDH19 65236.007	artificial	АА	QOYNYWPLT
1692	VH of CDH19 65236.007	artificial	TN	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCCGCC

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NO.				
				AGAGCCGAGGACACCGCCGTGTACCACTGTGCTAAGGGCGGCATGGGCGGCTACTACTACGGCATGGATGTGTGGGGCCCAGGGCAC CCTCGTGACCGTGTCTAGC
1693	VH of CDH19 65236.007	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISSDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTLVTVSS
1694	VL of CDH19	artificial	LN	GAGATCGTGATGACCCAGTCCCCCGTGACCCTGTCCCTGAGCCTGGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTACAT
	65236,007			
1695	VL of CDH19 65236.007	artificial	Ą	EIVMTQSPVTLSLSGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVY YCQQYNYWPLTFGGGTKVEIKS
1696	VH-VL of	artificial	_N	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCTCCGGCTTCACCTT
	CDH19			
	65236.007			CCTACTACGCCGAGTCCGTGAAGGGCCGGTTCACCATCTCCTCCGACAACTCCAAGTCCACCTGTACCTGCAGATGAACTCCCTG
				AGAGCCGAGGACCGTGTGTACCACTGTGCTAAGGGCGGGC
30				CCGTGACCCTGTCCCTGAGCCTGGGCGAGAGAGACCACCCTGTCTTGCCGGGCCTCCCAGTCCATCTCCAGCAACCTGGCCTGGTTC
7				
				GGCCCCTGACCTTCGGAGGCGCACCAAGGTGGAATCAAGTCC
1697	VH-VL of	artificial	Ą	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVST1SGGGANTYYAESVKGRFT1SSDNSKSTLYLQMNSL RAEDTAVYHCAKGAMGGYYYGMDVWGOGT1VTVSSGGGGSGGGSGETVMTOSPVT1.S1.S1.S1.GFRAT1.SCRASOS1SSN1.AWF
	65236.007			QQKPGQAPKLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1698	CDH19	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPCKGLEWVST1SGGGANTYYAESVKGRFT1SSDNSKSTLYLQMNSL
	65236.007 x			RAEDTAVYHCAKGGMGGYYGMDVWGQGTLVTVSSGGGGSGGGGGGGGGELVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF Ookbgoadriiyagaftratgidarbsgggggggfffttjtissifpavyycooynywditegggtryyfiksgggggfynniyfsgg
	126			GENVOPGGSLKISCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV
				YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1699	CDR-H1 of CDH19	artificial	ΑA	SYAMN
	65236.009			
1700	CDR-H2 of	artificial	AA	TISGGGANTYYADSVKG
	CDH19 65236.009			
1701	CDR-H3 of	artificial	ΑA	GGMGGYYYGMDV

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
2	CDH19 65236.009			QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1711	CDH19 65236.009 x 12C	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1712	CDR-H1 of CDH19 65236.010	artificial	AA	SYAMN
1713	CDR-H2 of CDH19 65236.010	artificial	AA	TISGGGANTYYADSVKG
309	CDR-H3 of CDH19 65236.010	artificial	АА	GGMGGYYYGMDV
1715	CDR-L1 of CDH19 65236.010	artificial	АА	RASQSISSNLA
1716	CDR-L2 of CDH19 65236.010	artificial	AA	GAFTRAT
1717	CDR-L3 of CDH19 65236.010	artificial	AA	QQYNYWPLT
1718	VH of CDH19 65236.010	artificial	Z	CAGGIGCAGCIGCIGGAAICCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCTGAGACTGTCTTGCGCCGGCTCCGGCTTCACCTT CICCAGCTACGCCATGAACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGTCCACCATCAGGGCGGGGGGGG
1719	VH of CDH19 65236.010	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS
1720	VL of CDH19	artificial	N	GAGATCGTGATGACCCAGTCCCCCGTGACCCTGTCCCTGAGCCTGGGCGAGAGAGCCACCTGTCTTGCCGGGCCTCCCAGTCCAT

CEC	DESIGNATION	SOLIBCE	TVDE	CEOI IENCE
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	65236.010			CTCCAGCAACCTGGCCTGGTTCCAGCAGAAGCCCGGCCCCTCGGCTGCTGATCTACGGCGCGCCTTTACCCGGGCCACCGGCAATCTGCCTGC
1721	VL of CDH19 65236.010	artificial	AA	EIVMTQSPVTLSLSGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLEPEDFAVY YCQQYNYWPLTFGGGTKVEIKS
1722	VH-VL of	artificial	F	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGATGGCTACGATGGCTACGATGGCTTCACCATGAACTGGCGGGGGGGG
	65236.010			CCTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGCGACAACTCCAAGTCCACCCTGTACCTGCAGATGAACTCCCTG AGAGCCGAGGACACCGCGTGTACCACTGTGCTAAGGGCGGCATGGCGGCTACTACTACGACGCATGGATGTGTGGGGCCAGGGCAC CACCGTGACCGTGTCTAGCGGAGGCGGAGGATCTGGCGGTGGTGGTGTTCTGGCGGAGGCGGCGCTCCGAGATCGTGATGACCAGTCCC
				CGGCTCCGGCACCTCGCGGCGCGCATCAGCTCCTGGAGCCCGAGGACTTTGCCGTGTACTACTGCAGCAGTACAACTACT GGCCCCTGACCTTCGGAGGCGGCACCAAGGTGGAAATCAAGTCC
1723	VH-VL of	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL
310	CDH19 65236.010			RAEDTAVIHCAKGGMGGIIIGMDVWGQGITTVTVSSGGGGSGGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQRPGQAPRLIIYGAFTRATGIPARVSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1724	CDH19	artificial	AA	OVOLLESGGGLVOPGGSLRLSCAASGFTFSSYAMNWVROAPGRGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLOMNSL RAEDTAVYHCAKGAMGGYYYGMDVWGOGTTVTVSSGGGGGSGGGGGGGGGGGTVMTOSPVTLSISLGFRATLSCRASOSISSNIAWF
	12C			QQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV
				YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1725	CDR-H1 of	artificial	AA	SYAMIN
	65236.011			
1726	CDR-H2 of	artificial	AA A	TISGGGANTYYADSVKG
	CDH19 65236.011			
1727	CDR-H3 of	artificial	¥₩	GGMGGYYYGMDV
	CDH19 65236.011			
1728	CDR-L1 of CDH19	artificial	AA	RASQSISSNLA

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9				
	65236.011			
1729	CDR-L2 of CDH19 65236.011	artificial	АА	GAFTRAT
1730	CDR-L3 of	artificial	AA	OOYNYWPLT
<u> </u>	CDH19 65236.011			
1731	VH of CDH19	artificial	IN	
	65236.011			CTCCAGCTACGCCATGAACTGGGTCCGACAGGCCCCTGGCAAGGGCCTTGGAATGGGTGTCCACCATCAGCGGCGGGGGGGCGAACACA CTCCAACTACTACTACTACTACTACTACTACTACTACTAC
1732	VH of CDH19	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL
	65236.011			RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS
1733	VL of CDH19	artificial	N	GAGATCGTGATCCCCCGGGCCTCCCCTGTCCCTGAGCCTGGGCGAGAGAGCCTGTCTTGCCGGGCCTCTCCCAGTCCAT
31	65236.011			CICCAGCAACCIGGCCIGGTICCAGCAGAAGCCCGGCCAGGCCCCICGGCIGCIGAICIACGGCGCCCTITACCCGGGCCACCGGCA
1				TCCCTGCCAGATTCTCTGGCTCCGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGGAGCCCGAGGACTTTGCCGTGTAC
100				TITLY TO CALL OTHER THE TOTAL OF THE TANK THE TA
1/34	VL of CDH19	artificial	ΑA	ELVMTQSPVTLSLSLGERATLSCRASQSLSSNLAWFQQRPGQAPRLLLYGAFTRATGLPARFSGSGSGTEFTLTLSSLEPEDFAVY YCOONNWPTTFGGGTKVFTKS
	0.05200		!	
1735	VH-VL of	artificial	Z	CAGGTGCAGCTGCTGGAGGATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACTTT CTCCAGCTGCCATCAACAACAGGGTCCGAACAGGCCTCTTGGCAAAAGGGCTTTGGAAATGGGTTCCACCAAATGAGGGGGGGG
	CDHI9 65236 011			
	11000000			AGAGCCGAGGACACCGCCGTGTACCACTGTGCTAAGGGCGGCATGGGCGGCTACTACTACGGCCATGGATGTGTGTG
				CACCGIGACCGIGICIAGCGGAGGCGGAGGAICIGGCGGIGGIGGIGGIICIGGCGGAGGCGGCICCGAGAICGIGAIGACCCAGICCC
				CCGTGACCCTGTCCCTGAGCCTGGGCGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTCCATCTCCAGCAACCTGGCCTGGTTC
				CGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGGAGCCCGAGGACTTTGCCGTGTACTACTGCCAGCAGTACAACTACT
				GGCCCCTGACCTTCGGAGGCGCCACCAAGGTGGAAATCAAGTCC
1736	VH-VL of	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL
	CDH19			RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGGGTTVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF
	65236.011			QQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1737	CDH19	artificial	ΑA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL
	65236.011 x			RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF OOK PGDA PRIJ TYGA FTRATGI PAR FSGSGGGGGGGGGGTTT TISSIR PEDFAVYYCOOYNYWPI TFGGGTKVFTKSGGGSFVOIVFSGG
				XXII (XXIII) (XXIII) (XXIII) (XXIII) (XXIII) (XXIII) (XXIIII) (XXIIIII) (XXIIIII) (XXIIIII) (XXIIIII) (XXIIIII) (XXIIIII) (XXIIIIII) (XXIIIIII) (XXIIIIII) (XXIIIIII) (XXIIIIIII) (XXIIIIIIIII) (XXIIIIIIIIII

SEQ NO.	DESIGNATION	SOURCE	IYPE	SEQUENCE
	12C			GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHH
1738	CDR-H1 of	artificial	AA	SYAMN
	CDH19			
1739	CDR-H2 of	artificial	AA	TISGGGANTYYAESVKG
i	CDH19			
	65236.012			
1740	CDR-H3 of	artificial	ΑA	GGMGGYYGMDV
	CDH19 65236 012			
1771	CDB 11 of	10:01:1	<	DAGOGICANITA
1/41	CDR-L1 0	arundal	¥	NASCO LOSNILA
	CDH19 65236.012			
31	CDR-L2 of	artificial	AA	GAFTRAT
	CDH19			
	65236.012			
1743	CDR-L3 of	artificial	AA	QQYNYWPLT
	CDH19			
1777	VH of CDH10	Licitit	Ę	
1/44	65236.012	al cilicia	Z	
				AGAGCCGAGGACACCGCCGTGTACCACTGTGCTAAGGGCGGCATGGGCGGCTACTACTACGGCATGGATGTGTGTG
1745	VH of CDH19 65236.012	artificial	AA	QVQILESGGGLVQPGGSIRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSS
1746	VL of CDH19	artificial	N	GAGATCGTGATCCCCAGTCCCCCGTGACCCTGTCCCTGAGCCTGGGCGAGAGACCCACCC
	65236.012			CTCCAGCAACCTGGCCTGGTTCCAGCAGAAGCCCGGCCCCTCGGCTGCTGATCTACGGCGCGCCTTTACCCGGGCCACCGGCAATTCTCTGGCTCTCTGGCTCCGGGCGCCCTGAGGCCCGGGGCCACCGGGCAATTCTCTGGCGTCTCTGGCGTCTTCGCCGAGGCCCGAGGACTTTGCCGTGTAC
				TACTGCCAGTACAACTACTGGCCCCTTCGGAGGCGGCCACCAAGGTCAAGTC
1747	VL of CDH19 65236.012	artificial	ΑA	EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVY YCQQYNYWPLTFGGGTKVEIKS

SEQ ID NO.	DESIGNATION	SOURCE	ТУРЕ	SEQUENCE
1748	VH-VL of CDH19 65236.012	artificial	L	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTTGGTGCAGCCTGGCGGCTCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTTTCCCGCTGCACTTCAGCTGCTTCAGCGGCGGAGGCCTTCACCTTTCCCAGCTTCAGCGGCGAGCGCCAACACACAC
1749	VH-VL of CDH19 65236.012	artificial	АА	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARFSGSGGGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKS
313	CDH19 65236.012 x 12C	artificial	AA	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPCKGLEWVST1SGGGANTYYAESVKGRFT1SRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLIIYGAFTRATGIPARFSGSGSTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGSEVQLVESGG GLVQPGGSIKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1751	CDR-H1 of CDH19 65236.013	artificial	АА	SYAMN
1752	CDR-H2 of CDH19 65236.013	artificial	АА	TISGGGANTYYADSVKG
1753	CDR-H3 of CDH19 65236.013	artificial	AA	GGMGGYYYGMDV
1754	CDR-L1 of CDH19 65236.013	artificial	АА	RASQSISSNLA
1755	CDR-L2 of CDH19 65236.013	artificial	АА	GAFTRAT
1756	CDR-L3 of	artificial	AA	QQYNYWPLT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
_ 0 N				
	CDH19 65236.013			
1757	VH of CDH19	artificial	۲	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65236.013			CTCCAGCTACGCCATGAACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGTCCACCATCAGCGGGGGGGG
				AGAGCCGCCGACACCGCCGTGTACTACTGTGCTAAGGGCGGCATGGGCGGCTACTACTACGGCATGGATGTGTGGGGCCAGGGCAC CACCGTGACCGTGTCTAGC
1758	VH of CDH19	artificial	ΑΑ	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL RAADTAVYYCAKGGMGGYYYGMDVWGOGTTVTVSS
1750	VI of CDH19	artificial	F	できる。
17.39	65236.013	alılıcıal	<u>-</u>	CTCCAGCAACCTGGCTCCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCGCCCTTTACCCGGGCCACCGCA
				TCCCTGCCAGAGTGTCTGGCTCCGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGCAGTCCGAGGACTTTGCCGTGTAC TACTGCCAGCAGTACAACTACTGGCCCCTGACCTTCGGAGGGGGGCACCAAGGTGGAAATCAAGTCC
1760	VL of CDH19	artificial	ΑA	EIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWFQQKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVY
	65236.013			YCQQYNYWPLTFGGGTKVEIKS
1761	Jo JV-HV	artificial	NT	CAGGTGCAGCTGCTGGAATCCGGCGGAGGACTGGTGCAGCCTGGCGGCTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	CDH19			CTCCAGCTACGCCATGAACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGTCCACCATCAGCGGCGGAGGCGCCAACA
	65236.013			. CUTACITACECUGACITCGIVEAAGGGCCGGITICACCAICITCCGCGACAACITCCAAGITCCACCTGIATACTGCAGATGAACITCCCTG
				AGAGCCCGCCGACACCGCCGTGTACTACTACTGTGCGCGCGC
				CCGTGACCCTGTCCTGAGCCTGGGCGAGAGAGACCCTGTCTTGCCGGGCCTCCCAGTCCATCTCCAGGAACCTGGCCTGGTTC
				CAGCAGAAGCCCGGCCCAGGCCCCTCGGCTGCTGATCTACGGCGCCTTTACCCGGGCCCACCGGCATCCCTGCCAGAGTGTCTGGCTC
				CGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCCTGCAGTCCGAGGACTTTGCCGTGTACTACTGCCAGCAGTACAACTACT
				GGCCCCTGACCTTCGGAGGCGGCACCAAGGTGGAAATCAAGTCC
1762	VH-VL of	artificial	¥	QVQLLESGGGIVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYADSVKGRFTISRDNSKSTLYLQMNSL
	CDH19 65236 013			RAADTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGETVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF OOKPGOAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLOSEDFAVYYCOOYNYWPLTFGGGTKVEIKS
1763	CDH19	artificial	۷۷	OVOLI ESCECTIVOPERSTRISCA A SCETT-SSYAMNIMVROA PERCIENIVSTISCECANTYYA DSVIKERFTISRDNSKSTIVIOMNSI.
3	65236.013 x	5	{	RAADTAVYYCAKGGMGGYYYGMDVWGQGTTVTVSSGGGGSGGGSGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF
	12C			QOKPGQAPRLLIYGAFTRATGIPARVSGSGSGTEFTLTISSLQSEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGGSEVQLVESGG
				GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISKDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK
				PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1764	CDR-H1 of CDH19	artificial	AA	SYAMN
	1			

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
				CCGTGACCCTGTCCCTGAGCCTGGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTCTCCAGCAACCTGGCCTGGTTC CAGCAGAAGCCCGGCCCAGGCCCTCGGCTGCTGATCTACGGCGCCTTTACCCGGGCCACCGGCATCCCTGCCAGATTCTCTGGCTC CGGCTCCGGCACCGAGTTCACCCTGACCATCAGCTCCTGGAGCCCTTTGCCGTGTACTACTGCCAGCAGTACAACTACT GGCCCTGACCTTCGGAGGCGCACCAAGGTGGAAATCAAGTCC
1775	VH-VL of CDH19 65236.014	artificial	АА	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTLVTVSSGGGGSGGGGGGGGGGSEIVMTQSPVTLSLSLGERATLSCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKS
1776	CDH19 65236.014 x 12C	artificial	AA A	QVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAPGKGLEWVSTISGGGANTYYAESVKGRFTISRDNSKSTLYLQMNSL RAEDTAVYHCAKGGMGGYYYGMDVWGQGTLVTVSSGGGGSGGGGGGGGGGGGTIVMTQSPVTL.SL.SLGERATL.SCRASQSISSNLAWF QQKPGQAPRLLIYGAFTRATGIPARFSGSGSGTEFTLTISSLEPEDFAVYYCQQYNYWPLTFGGGTKVEIKSGGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSCVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
37	CDR-H1 of CDH19 65237.001	artificial	АА	RYGIH
16	CDR-H2 of CDH19 65237.001	artificial	АА	VIWYDGSNKYYADSVKG
1779	CDR-H3 of CDH19 65237.001	artificial	AA	RAGIPGTTGYYYGMDV
1780	CDR-L1 of CDH19 65237.001	artificial	АА	SGDRLGEKYVS
1781	CDR-L2 of CDH19 65237.001	artificial	АА	QDNKWPS
1782	CDR-L3 of CDH19 65237.001	artificial	AA	QAWDSSTVV
1783	VH of CDH19 65237.001	artificial	LN L	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGCAGCCTGGCCGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGATACGGCATCCACTGGGCGACAGGCCCTGGCAAGGGCCTGGAATGGGTGGCTGATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAACTCCCTG

CEO	DESIGNATION	COLIDA	TVDE	CECHIENCE
2 O S		3000		אַרעטנועטר
2				CGGGCCGAGGACTCCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCCCCGGCACCACCGGCTACTACTACGGCATGGATGTGTG GGGCCAGGGCACCACGTGACCGTGTCTAGC
1784	VH of CDH19 65237.001	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
1785	VL of CDH19 65237.001	artificial	F.	TCTTACGAGCTGACCCCGCCCCCCCTCCGTGTCCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGA GAAATACGTGAGCTGGTATCAGCAGAAGCCCGGCCAGTCCCCCATCCTGGTCATCTACCAGGACAACAAGTGGCCCTCCGGCATCC CTGAGCGGTTCTCCGGCTCCAACTCCGGCAACACCGCCACCCTGACCATCTCCGGCACCAGGCCATGGACGAGGCCGACTACTAC TGCCAGGCCTGGGACTCCTCCACCGTGGTGTTCGGCGAGGCACCAAGCTGACCGTGCTGTCC
1786	VL of CDH19 65237.001	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVLS
1787	VH-VL of CDH19	artificial	F	CAGGTGCAGCTGGTGGAATCCGGCGGGGGGGGGGTGCTGGCCGGTCCTGGAGACTGTCTTGCGCCGCCTCGGCTTCACCTT CTCCAGATACGGCATCCACTGGGTCCGACAGGCCCTTGGCAAGGGCCTGGAATGGGTGGCCGTGATTTGGTACGACGGCTCCAACA
317	65237.001			AGTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACAACTCCAAGAACCCCTGTACCTGCAGATGAACTCCCTG CGGGCCGAGGACTCCGCCGTGTACTGTGCCAGAAGGGCCGGCATCCCCGGCCACCCGGCTACTACTACTGCGATGGTGTG GGGCCAGGGCACCACCGTGTCTACTGTGCCAGAGGGCCGAGGCTCTTGGTGGCGGCGACCCGCCTCTTACGAGC TGACCCAGGCCCCCCTCCGTGTCTCTAGCGGAGGCGAGGAGCTCTTACGAGC TGACCCAGCCCCCCTCCGTGTCTCCTGGCCAGACCGCCTCCATCTGGCGACCCGCCTGGGCGAGAATACGTG AGCTGGTATCAGCAGAAGCCCGGCCAGTCCCCATCCTGGTCATCTACCAGGACAACACCGGCTTCCTGGCCGATCCCTGAGCGGTT CTCCGGCTCCAACTCCGGCAACCCCACCCTGACCATCTCCGGCCATGGACGAGGCCGACTACTGCCAGGCCT GGGACTCCTCCAACTCCGGCAACCCAGGCACCCAAGCTGCTGCTGTCC GGGACTCCTCCAACTCCGGCAACCCAAGCTGACCTACTGCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCCAGGCCTACTCCACCCAGGCCTACTCCCAGGCCCACCTACTCCCAGGCCCACCCA
1788	VH-VL of CDH19 65237.001	artificial	АА	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLS
1789	CDH19 65237.001 x 12C	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGGGSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGCQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1790	CDR-H1 of CDH19 65237b.001	artificial	АА	RYGIH
1791	CDR-H2 of CDH19 65237b.001	artificial	ΑΑ	VIWYEGSNKYYAESVKG
1792	CDR-H3 of	artificial	AA	RAGIPGTTGYYYGMDV

IRCE TYPE SEQUENCE		icial AA SGDRLGEKYVS	icial AA QDNKWPS	:	AA	icial NT CAGGTGCAGCTGGTAGCGGGAGGCGTGGTGCTGCCGGGTCCCTGAGACTGTCTTGCGCCTCCGGCTTCACCTT CTCCAGATACGGCATCCACTGGGCACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGTTTTGGTACGAGGGCTCCAACA AGTACTACGCCGAGTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAACTCCCTG CGGGCCGAGGACTCCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCCCCGGCACCACGGCTACTACTACGGCATGGATGTGTG GGGCCAGGGCACCACGTGTACTACTACTACTAGC	icial AA QVQLVESGGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS	icial NT TCTTACGAGCTGACCCCCCCCCCCCCCCCCGGCCGGCCGG	icial AA SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWESSTVVFGGGTKLTVLS	Icadetecadetegraphecegecegecegecegecegecegecegecegecegece	icial AA QVQLVESGGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGRFTISRDNSKNTLYLQMNSL RAFDSALVYYCAPPAGT DGTTGVVVCANDYMGOGTTTVVVSGGGGGGGGGGGGGGGSSVF1.TOPDSVSVSDGOTASTTGGDDF1.GFKVVV
SOURCE		artificial	artificial	-	artificia	artificial	artificial	artificia	artificia	artificia	artificial
DESIGNATION	CDH19 65237b.001	3 CDR-L1 of CDH19 65237b.001		65237b	CDR-L3 of CDH19 65237b.001	65237b.001	7 VH of CDH19 65237b.001	65237b.001	9 VL of CDH19 65237b.001	CDH19 65237b.001	1 VH-VL of
SEQ ID NO.		1793	1794		1795	31	1797	1798	1799	1800	1801

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
© N				
	CDH19 65237b.001			SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLS
1802	CDH19 65237b 001 x	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYEGSNKYYAESVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGOGTTVTVSSGGGGSGGGGSGGGGSSYELTOPPSVSVSPGOTASITCSGDRLGEKYV
	120			SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QOKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1803	CDR-H1 of	artificial	AA	RYGIH
	CDH19 65237.002			
1804	CDR-H2 of	artificial	ΑA	VIWYDGSNKYYADSVKG
	CDH19 65237.002			
1805	CDR-H3 of	artificial	AA	RAGIPGTTGYYYGMDV
31	CDH19			
a	65237.002			
1806	CDR-L1 of	artificial	ΑA	SGDRLGEKYVS
	CDH19 65237.002			
1807	CDR-12 of	artificial	AA	ODNKWPS
	CDH19			
	65237.002			
1808	CDR-L3 of	artificial	AA	QAWDSSTVV
	CDH19 65237.002			
1809	VH of CDH19	artificial	M	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGGTGCTGGCGGCGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65237.002			AGIACIACGCCGACICCGTGAGGCCGGTTCACCATCICCGGGACAACTCCAGGAACACCCTGIACCAGATGAACTCCCTG
				CGGGCCGAGGACTCCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCCCCGGCACCACCGGCTACTACTACGGCATGGATGTGG GGGCCAGGGCACCACCGTGACCGTGTCTAGC
1810	VH of CDH19 65237.002	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSS
1811	VL of CDH19	artificial	N	TCTTACGAGCTGACCCAGCCCCCCTCCGTGTCCCGTGTCTCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGCGAA
1)	5	:	

SFO	DESIGNATION	SOURCE	TYPF	SFOLIENCE
∑ 0 N 0.		; ; ; ; ; ;	-	
	65237.002			GAAATACGTGAGCTGGTATCAGCAGAAGCCCGGCCAGTCCCCCATCCTGGTCATCTACCAGGACAAGTGGCCCTCCGGCATCC CTGAGCGGTTCTCCGGCTCCAACTCCGGCAACACCGCCACCTGACCATCTCCGGCACCAGGCCATGGACGAGGCCGACTACTAC TGCCAGGCCTGGGACTCCTCCACCGTGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1812	VL of CDH19 65237.002	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYVSWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVLS
1813	VH-VL of	artificial	F	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65237.002			
				TGACCCCAGCCCTCCGTGTCCCTGTCCTCGCCAGACCACACCCTCCATCCCTGTCTCTCCCCCCGCTGGCCGACCGCGCGCG
				AGCIGGIAICAGCAGAAGCCCGGCCAGICCCCAICCIGGICAICIAGGAGAGAAGAGIGGCCGICCGGGCAIGCTGGAGCGGTI CTCCGGCTCCAACTCCGGCAACACCGCCACCCTGACCATCTCCGGCACCCAGGCCCATGGACGAGGCCGACTACTACTGCCAGGCCT GGGACTCCTCCACCGTGGTTCGGCGGAGGCACCAAGCTGACCGTGTCC
1814	VH-VL of	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL
320	CDH19 65237.002			RAEDSAVYYCARRAGIPGTTGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVSVSPGQTASITCSGDRLGEKYV SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLS
1815	CDH19	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSRYGIHWVRQAPGKGLEWVAVIWYDGSNKYYADSVKGRFTISRDNSKNTLYLQMNSL RAFDSAVYYCARRAGIPGTTGXYYCMDYMGOGTTTYTTYS SGGGGGSGGGGGSSYFITOPPSVSVSPFOTTASITCSGDRIGERXV
	63237.002 x 12C			SWYQQKPGQSPILVIYQDNKWPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE
	!			SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1816	CDR-H1 of	artificial	AA A	SSGYYWS
	CDH19 65238.002			
1817	CDR-H2 of	artificial	AA	YIYYTGSAYYNPSLKS
	CDH19 65238.002			
1818	CDR-H3 of	artificial	AA	DGSSGWYFQY
	CDH19 65238.002			
1819	CDR-L1 of CDH19	artificial	AA	RASRQISSSYLA

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	65238.002			
1820	CDR-L2 of CDH19 65238.002	artificial	AA	GPSSRAT
1821	CDR-L3 of CDH19 65238.002	artificial	AA	QQYGSSFT
1822	VH of CDH19 65238.002	artificial	Z	CAGGTGCAGCTGCAGGAATCCGGCCCTGGTCAAGCCCTCCGAGACCCTGTCCCTGACCTGCACCGTGTCCGGCGGCGCTCAT CTCCTCCTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCGGCAAGGGCCTGGAATGGATCGGCTACATCTACAACCGGCT CCGCCTACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCC GTGACCGCCGCTGACACCGCCGTGTACTGCGCCCAGAGATGGCTCCAGGGCTGGTACTTCCAGTACTGGGGCCAGGGCACCCT GGTCACCGCCGTGTACTACAGCCCCAGAGATGGCTCCAGGGCTGGTACTTCCAGTACTGGGGCCAGGGCACCCT GGTCACCGTCTTACA
1823	VH of CDH19 65238.002	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS
321	VL of CDH19 65238.002	artificial	TN	GAGATCGTGCTGACCCCAGTCCCCCGGCACCCTGTCTCTGAGCCCTGGCGAGAGAGA
1825	VL of CDH19 65238.002	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSFTFGQGTKVDIKS
1826	VH-VL of CDH19 65238.002	artificial	F	
1827	VH-VL of CDH19 65238.002	artificial	ΑΑ	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWY QQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKS
1828	CDH19 65238.002 x	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSGGGGSGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWY QQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSFTFGQGTKVDIKSGGGGSEVQLVESGGG

SEQ D	DESIGNATION	SOURCE	TYPE	SEQUENCE
5	12C			LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQIVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1829	CDR-H1 of CDH19 65238.004	artificial	АА	SSGYYWS
1830	CDR-H2 of CDH19 65238.004	artificial	AA	YIYYTGSAYYNPSLKS
1831	CDR-H3 of CDH19 65238.004	artificial	AA	DGSSGWYFQY
1832	CDR-L1 of CDH19 65238.004	artificial	AA	RASRQISSSYLA
322 322	CDR-L2 of CDH19 65238.004	artificial	AA	GPSSRAT
1834	CDR-L3 of CDH19 65238.004	artificial	AA	QQYGSSFT
1835	VH of CDH19 65238.004	artificial	Z	CAGGTGCAGCTGCAGGAATCCGGCCTGGCCTGGTCAAGCCCTCCGAGACCCTGTCCTGACCTGCACCGGTGTCCGGCGGCGTCCAT CTCCTCCTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCGGCAAGGGCCTGGAATGGATCGGCTACATCTACAACCGGCT CCGCCTACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCGTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCTCC GTGACCGCCGCTGACACCCCCTGTACTACTGCGCCAGAGATGGCTCCCAGGGCCAGGGCCCCT GGTACCGCCGCTGTACTACTGCGCCCAGAGATGGCTCCCAGGCTGCTCCAGTACTGGGGCCAGGGCCCCCT GGTACCGCCGTGTACTACTACTGCCCAGAGATGGCTCCAGGCGCTGGTACTTCCAGTACTGGGGCCAGGGCCCCCT GGTCACCGTCTTACTACTACTACTACTACTACTACTACTACTACTACT
1836	VH of CDH19 65238.004	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSS
1837	VL of CDH19 65238.004	artificial	Z	GAGATCGTGCTGACCCAGTCCCCCGGCACCCTGTCTCTGAGCCCTGGCGAGAGAGCCCTCTCTGCCGGGGCCTCCCGGCAGAT CTCCTCCAGCTACCTGGCTTGGTATCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGAGCCTAGCTCCAGAGCCACG GCATCCCTGACCGGTTCTCCGGCTCTGGCTCCGGCACCTCACCCTGACCCTGACCCGGCTGGAACCCGAGGACTTCGCCGTG TACTATTGCCAGCAGTACGGCTCCTTCACCTTCACCTGGCACCAAGGTGAACATCAAGTCC
1838	VL of CDH19 65238.004	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWYQQKPGQAPRLLIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGSSFTFGPGTKVDIKS

SEQ ID NO.	DESIGNATION	SOURCE	ТУРЕ	SEQUENCE
1839	VH-VL of CDH19 65238.004	artificial	L	CAGGTGCAGCTGCAGGAATCCGGCCCTGGTCAAGCCCTCCGAGACCCTGTCCCTGACCTGCACCGTGTCCGGCGCTCCAT CTCCTCCTCCTCCGCTACTACTGGTCCTGGATCCGGCAGCCCCCGGCAGGGCCTGGAATGGATCGGCTACATCTACTACTACACCGGCT CCGCCTACTACACACCCCAGCCTGAAGTCCAGAGCCCCCTGGAACCGCTCCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCC GTGACCGCCGCTGACCACCGCCGTGTACTACTGCGCAGAGATGGCTCCAGGGGCTGGTACTTCCAGTACTGGGGCCAGGCCCC GGTCACCGTGTCTAGCGGAGGCGGAGGTGCTGGTGGTGGTTCTGGCGGCGGGGCTGGTTCTCCAGTTCCTGGGGCCAGGCCCCC GCACCCTGTCTCTGAGCCGAGGAGGAGCCACCCTGTCCTGCCGGGCCTCCCGGGCCAGATCTCCTGGCGCTTGGTTAT CAGCACCGTGTCTTCTGAGCCCTGGCTGCTGATCTTACGGCCCTTACTGCCGGCAGCCTCCTGGCTTCTCCGGCTCCTTGGCTTCTTGCCAGCCTTGGCTTCTCCGGCTCCTTATTGCCAGCCTTGGCTTCTCCGGCTCCTTCCCTCCTTCCGGCTCCTTCCTCC
1840	VH-VL of CDH19 65238.004	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSGGGGSGGGGGGGGGGSTIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWY QQKPGQAPRLLIYGPSSRATGIPDRFSGSGGGGGGTLTIISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKS
353	CDH19 65238.004 x 12C	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSGYYWSWIRQPPGKGLEWIGYIYYTGSAYYNPSLKSRVTISVDTSKNQFSLKLSS VTAADTAVYYCARDGSSGWYFQYWGQGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASRQISSSYLAWY QQKPGQAPRLIIYGPSSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSFTFGPGTKVDIKSGGGGSEVQLVESGGG LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGCTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1842	CDR-H1 of CDH19 65240.002	artificial	АА	SYDMH
1843	CDR-H2 of CDH19 65240.002	artificial	AA	VISYDGTNEYYADSVKG
1844	CDR-H3 of CDH19 65240.002	artificial	АА	ERYFDWSFDY
1845	CDR-L1 of CDH19 65240.002	artificial	AA	RASQSVSNTYLA
1846	CDR-L2 of CDH19 65240.002	artificial	АА	GASSRAT
1847	CDR-L3 of	artificial	AA	QQYSNSWT

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<u> </u>			-	
	CDH19 65240.002			
1848	VH of CDH19	artificial	Ł	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
?	65240.002			CTCCAGCTACGACATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCGTGATCTCCTACGACGGCCAACG
				GGGCCCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGTACTTCGACTGGTCCTTCGACTACTGGGGCCAGGGCCACCTGGGT GTCCGTGTCTAGC
1849	VH of CDH19 65240.002	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPCKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS
1850	VL of CDH19	artificial	M	GAGATCGIGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTCCGT
	65240.002			GTCCAACACCTACCTGGCCTGGTATCAGCAGCGCCCTGGCCAGGCCCCTCGGCTGGTGCTGATCTACGGCGCCTCTTCCAGAGCCACCG GCATCCTGGACCGGTTCTCCGGCTTCTGGCTTCTTGGCACCACTTTCACCTTGACCAATTTCACATTTCACAATTTCACAAGAACCACCG
				TACTATIGCCAGCAGTACTCCAACTCCTGGACCTTCGGACAGGGCACCAAGGTGGAAATCAAGTCC
1851	VL of CDH19 65240 002	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCOOYSNSWTFGOGTKVEIKS
1957	VH-VI of	er+ificial	ΤΝ	である。 では、これのようには、これのようについては、または、これのようには、これのは、これのは、これのようには、これのようには、これのようには、これのようには、これのようには、これのようには、これのようには、これのようには、これのようには、これのようには、これのよりには、これのようには、これのようには、これのよりには、これのようには、これのは、これのよりには、これのようには、これのよりには、これのは、これのよりには、これのは、これのよりには、これのよりにはいる。これのは、これのは、これのは、これのは、これのは、これのは、これのは、これのは、
7 24	CDH10	a cilicia	2	CITICA DOLINGATOR TO THE SECOND CONTROLL OF T
	65240.002			AGTACTACGCCGACTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACACCTCCAAGAACACCTGTACCTGCAGATGAACTCCCTG
				CGGCCCGAGGACACCGCCGTGTACTACTGCGCCCAGAGCGGTACTTCGACTGGTCCTTCGACTACTGGGGCCCAGGGCACCTGGT
				GICCGIGICIAGCGGAGGCGGAGGAICIGGIGGCGGIGGTICIGGCGGCGGAGGCICCGAGAICGIGCIGACCCAGICCCIGGCA
				CCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCAGTCCGTGTCCAACACCTACCT
				CAGCGCCCTGGCCCAGGCCCCTCGGCTGCTGATCTACGGCGCCTCTTCCAGAGCCACCGGCATCCCTGACCGGTTCTCCGGCTTCTGG
				CTCTGGCACCACCACCCTGACCATCTCCAGCCTGGAACCCGAGGATTTCGCTGTGTACTATTGCCAGCAGTACTCCAACTCCT GGACCTTCGGACAGGGCACCAAGGTGGAAATCAAGTCC
1853	VH-VL of	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL
	CDH19			RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGGGGGETVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ
	65240.002			QRPGQAPRLLIYGASSRATG1PDRFSGSGSGTDFTLTSSLEPEDFAVYYCQQYSNSWTFGQGTRVE1RS
1854	CDH19	artificial	ΑĄ	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYDGTNEYYADSVKGRFTISRDTSKNTLYLQMNSL bardhaityvcaabbrovedagedvedagedvecchiiigaygeccccecccertii moedchiei sisbcedahi scodagoshiva amotaamotaamo
	65240.002 X 135			INTERPRETATION TO THE PROTECTION OF SOCIOETIC STEPEDEAVY COOKSING TO SHOULD THE SECTION OF THE S
	7			
				CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLMYSNRWVFGGGTKLTVLHHHHHH
1855	CDR-H1 of	artificial	AA	SYDMH
	CDHI9			

SEQ PD CA	DESIGNATION	SOURCE	TYPE	SEQUENCE
	65240.003			
1856	CDR-H2 of	artificial	AA	VISYEGTNEYYAESVKG
	CDH19 65240.003			
1857	CDR-H3 of	artificial	AA	ERYFDWSFDY
	CDH19			
1959	CDP-11 of	or+ificiol	<	RASOSVANITYT.A
201	CDH19	5	{	
	65240.003			
1859	CDR-L2 of	artificial	ΑA	GASSRAT
	CDH19 65240.003			
1860	CDR-L3 of	artificial	AA	QYSNSWT
	CDH19			
25	65240.003			
1861	VH of CDH19	artificial	N	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65240.003			CTCCAGCTACGACATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCGTGATCTCCTACGAGGGCACCAACG AGTACTACGACGAGTCCGTGAAGGCCGGGTTCACCATCTCCGGGACACCTCCAAGAACACCTGTGTACCTGCAGATGAACTCCTG
				CGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGAGAGCGGTACTTCGACTGGTCCTTCGACTACTGGGGCCAGGGCCACCTGGT GTCCGTGTCTAGC
1862	VH of CDH19	artificial	AA	
	65240.003		!	
1863	VL of CDH19	artificial	¥	
	65240.003			GICCAACATTACCIGGETTICAGCAGCGCCCTGGCACTTCAGCCCCTCGGCTGCTGATCTACGGCGCCTTTCCAGAGCCACTGTTTCGCTGTGTTTCGCTGTGTTTCGCTGTGTGTG
1864	VL of CDH19	artificial	AA	
	65240.003			
1865	VH-VL of	artificial	۲	
	CDH19 65240 003			LICCAGCIACGACAIGCACIGGGICCGACAGGCCCCIGGCAAGGGCCIGGAAIGGGIGGCCGIGAICICCIACGAGGGCACCACCTG AGTACTACGCCCGAGTCCGTGAAGGCCCGGTTCACCATCTCCCGGGACACCTCCAAGAACACCTGTACCTGCAGATGAACTCCCTG
				CGGGCCGAAGGACACCGCCGTGTACTACTGCGCCAGAGAGCGGTACTTCGACTGGTCCTTCGACTACTAGGGCCAGGGCACCCTGGT GTCCGTGTCTAACGGAAGGAAGAATCTGGTGGCGGTGGTGGTTGGACGGGAAGGAA

SEQ ID NO.	DESIGNATION	SOURCE	TYPE	SEQUENCE
				CCCTGTCCCTGAGCCCTGGCGAGAGACCACCTGTCTTGCCGGGCCTCCCAGTCCGTGTCCAACACCTGCCTG
1866	VH-VL of CDH19 65240.003	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGGGGGGGGGGGGTIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKS
1867	CDH19 65240.003 x 12C	artificial	A A	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAAGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1868	CDR-H1 of CDH19 65240.004	artificial	АА	SYDMH
1869	CDR-H2 of CDH19 65240.004	artificial	AA	VISYEGTNEYYAESVKG
1870	CDR-H3 of CDH19 65240.004	artificial	AA	ERYFDWSFDY
1871	CDR-L1 of CDH19 65240.004	artificial	АА	RASQSVSNTYLA
1872	CDR-L2 of CDH19 65240.004	artificial	АА	GASSRAT
1873	CDR-L3 of CDH19 65240.004	artificial	AA	QQYSNSWT
1874	VH of CDH19 65240.004	artificial	Z	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGACAGGCCCTCCGGCACAGGGCCTTGGGAATGGGTGGCCGTGATCTCCTACGAGGGCCACAACG AGTACTACGACGTGATGCTGCAGGGCCCTGTACGAGGGCCCTGTACGAGGGCCCTGTACGAGGGCCCTGTACGAGGGCCCTGTACGAGGGCCCTGTACGAGATGAACTCCCTG

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				CGGGCCGAGGACACCGCCGTGTACTACTGCGCCCAGAGAGCGGTACTTCGACTGGTCCTTCGACTACTGGGGGCCAGGGCACCCTGGT GTCCGTGTCTAGC
1875	VH of CDH19 65240.004	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSS
1876	VL of CDH19	artificial	F	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGCCACCTGTCTTGCCGGGCCTCCCCGTCGTCCGT
	65240.004			GLOCAMONOLIACOLGGIALONGONGANGOLGGIOCOLGGIAGO
1877	VL of CDH19 65240.004	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYSNSWTFGQGTKVEIKS
1878	VH-VL of	artificial	N	CAGGTGCAGCTGGAGGCGTGGAGGCGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	CDH19			CTCCAGCTACGACATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCGTGATCTCCTACGAGGGCACCAACG
	65240.004			AGTACTACGCCGAGTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACACCTCCAAGAACACCTGTACCTGCAGATGAACTCCCTG CGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGAGAGGGGTACTTCGACTGGTCCTTTCGACTACGACTACTACGAGGCACCAAGAGAACTCCTGG
				GTCCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGCTTCTGGCGGCGGAGGCTCCGAGATCGTGCTGACCCAGTCCCTGGCA
32				CCCTGICCCIGAGCCCTGGCGAGAGAGCCACCCTGICITGCCGGGCCTCCCAGICCGIGICCAACACCTACCTGGCTGTATCAG
7				CTCTGGCACCTTCACCCTGACCATCTCCAGCCTGGAACCCGAGGATTTCGCTGTGTACTATTGCCAGCAGCAACTCCT
				GGACCTTCGGACAGGGCACCAAGGTGGAAATCAAGTCC
1879	VH-VL of CDH19	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVSVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ
	65240.004			QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKS
1880	CDH19	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL
	65240.004 x			RAEDTAVIICAKERIFDWSFDIWGQGTLVSVSSGGGGSGGGGGGGGETTVLTQSPGTLSLSPGERATLSCRASQSVSNTILAWIQ OKPGOAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCOOYSNSWTFGOGTKVEIKSGGGGGSEVOLVESGGG
	27			
				CVRHGNFGNSY1SYMGYGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLMYSNRWVFGGGTKLTVLHHHHHH
1881	CDR-H1 of CDH19	artificial	AA	SYDMH
	65240.005			
1882	CDR-H2 of	artificial	AA	VISYEGINEYYAESVKG
	65240.005			
1883	CDR-H3 of	artificial	AA	ERYFDWSFDY

SEQ ID NO.	DESIGNATION	SOURCE	ТУРЕ	SEQUENCE
	CDH19 65240.005			
1884	CDR-L1 of CDH19 65240 005	artificial	AA	RASQSVSNTYLA
1885	CDR-L2 of CDH19 65240 005	artificial	АА	GASSRAT
1886	CDR-L3 of CDH19 65240.005	artificial	AA	QQYSNSWT
32	VH of CDH19 65240.005	artificial	TN	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTTC CTCCAGCTACGACATGCACTGGGTCCGACAGGCCCTGGCAGGGCCTGGAATGGGTGGCTGTTTCCTACGAGGGCACCAACG AGTACTACGCCGAGTCCGTGAAGGGCCGGTTCACCATCTCCCGGGACACCTCCAAGAACACCCTGTACCTGCAGATGAACTCCCTG CGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGAGAGCGGTACTTCGACTGGTCCTTCGACTACTGGGGCCAGGGCACCCTGGT GACCGTGTCTAGC
8 1888	VH of CDH19 65240.005	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVTVSS
1889	VL of CDH19 65240.005	artificial	TN	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGAGCCCTGGCGAGAGAGCCACCCTGTCTTGCCGGGCCTCCCCAGTCCGT GTCCAACACCTACCTGGCTGTATCAGCAGAAGCCTGGCCAGGCCCTCGGCTGATCTACGGCGCCTCTTCCAGAGCCACG GCATCCCTGACCGGTTCTCCGGCTCTGGCTCTGGCACCTTCACCCTGACCATCTCCAGCCTGGAACCCGAGGATTTCGCTGTG TACTATTGCCAGCAGTACTCCAACTCCTGGACCTTCGGACAGGGGGAAATCAAGTCC
1890	VL of CDH19 65240.005	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYSNSWTFGQGTKVEIKS
1891	VH-VL of CDH19 65240.005	artificial	LN .	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTTTTCCTACGGTGGCTTCACCTTTTCCCACGGCTTCACCTTTTCCAGGCTTCACCTTTCACCTTTCACCTTTCACCTTCACCTTTCACCTTACGGTGGCCGTGATCCTTACGGGCCCCAGGGCCCAACGAGTACTACGCCGAGTTCACCTGCGAGTTCACCTGCGGTTCACCTGCGGTTCACCTGGTGATCCTTCAGTTCACTGGGGCCCTGGTTCACTTCGGTGCTTCAGTTCAGTTCATTCGGTGGTTCTTTCGGTGGTTCTTTCGGTGGTTCTTTCGGTGGT
1892	VH-VL of	artificial	AA	QVQLVESGGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVTVSSGGGGGGGGGGGGGGGTIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
₽ NO.				
	CDH19 65240.005			QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKS
1893	CDH19 65240.005 x 12C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYDMHWVRQAPGKGLEWVAVISYEGTNEYYAESVKGRFTISRDTSKNTLYLQMNSL RAEDTAVYYCARERYFDWSFDYWGQGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSNTYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYSNSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1894	CDR-H1 of CDH19 65246.004	artificial	AA	SYFIH
1895	CDR-H2 of CDH19 65246.004	artificial	AA	IINPISVSTSYAQKFQG
1896	CDR-H3 of CDH19 65246.004	artificial	AA	GGIQLWLHFDY
1897	CDR-L1 of CDH19 65246.004	artificial	AA	SGSSSNIGSNFVN
1898	CDR-L2 of CDH19 65246.004	artificial	AA	TNNQRPS
1899	CDR-L3 of CDH19 65246.004	artificial	AA	ATWDESLQGWV
1900	VH of CDH19 65246.004	artificial	Z	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCAGGGCCTGGAATGGATGG
1901	VH of CDH19 65246.004	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
1902	VL of CDH19	artificial	NT	CAGTCTGCCCTGACCCAGCCTCCTTCTGCCACCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCTCCTCCTCCAACAT

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N 0 8.	DESIGNATION	300RCE	<u> </u>	SECOLINCE
	65246.004			CGGCTCCAACTTCGTGAACTGGTACCAGCAGCTGCCCGGCACCGCCCCCAAGGGTGCTGATCTACACCAACAACAGCGGCCCTCCGGGCTCCCGGTGCCCGACCAGCGGCCCTCCGGTGCCCGACCAGCGGCCCTCCGAGGCCGACCAACAACAACAACAACAACAACAACAACA
1903	VL of CDH19 65246.004	artificial	AA	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATWDESLQGWVFGGGTKLTVLS
1904	VH-VL of CDH19	artificial	N	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGTGTCCGGCTACACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCCAGGGCCTGGAATGGATGG
	65246.004			CCTCCTACGCCCAGAAATTCCAGGGCAGAGTGACCATGACCCGGGACACCTCCACCTCCACCGTGTACATGGAACTGTCCTCCTG CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCACTTCGACTACTGGGGCCAGGGCACCT GGTCACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGGGGGGG
				CTGCCACCGGCACCCCTGGCCCAGCGCGCGTGTCTCCTGCTCCGGCTCCTCCTCCAACATCGGCTCCAACTTCGTGAACTGGTTAC CAGCAGCTGCCCGGCCCCCCCAAGGTGCTGATCTACACCAACAACAGGGGCCCTCCGGCGTGCCCGACCGGTTCTCTGGCTC CAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGGACGAGGCCGACCAACTACTGTGCCACCTGGGACGAGT
				CCCTGCAGGGCTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
330	VH-VL of CDH19 65246,004	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATWDESLQGWVFGGGTKLTVLS
1906	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL PSFDTAVVVCAPGGIOLMIHFDVWGOGTIVTVSSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
	65246.004 x 12C			COLPGIANCES EMBLICATIONS OF THE STATE OF THE
1907	CDR-H1 of	artificial	AA	KAT GENT INCLUDING THE GIFFING OGDING OF THE DEAD INCLUDED ON THE CONTROL OF THE STATE OF THE ST
	CDH19 65247.004			
1908	CDR-H2 of	artificial	AA	IINPISVSTSYAQKFQG
	65247.004			
1909	CDR-H3 of	artificial	AA	ССІЎГИТНІ Т
	65247.004			
1910	CDR-L1 of CDH19	artificial	AA	SGSSSNIGSNFVN

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	65247.004			
1911	CDR-L2 of CDH19 65247.004	artificial	AA	TNNQRPS
1912	CDR-L3 of CDH19 65247.004	artificial	AA A	ATYDESMQGWV
1913	VH of CDH19 65247.004	artificial	Z	CAGGTECAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGCGGGTGTCCTGCAAGGTGTCCGGCTACCTT CACCAGCTACTTCATCCACTGGGTCCGACAGGCCCAGGGCCTGGGAATGGATGG
1914	VH of CDH19 65247.004	artificial	ΑΑ	QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSS
331	VL of CDH19 65247.004	artificial	LN L	CAGTCTGCCCTGACCCAGCCTCCTTCCGCTACCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCTCCTCCAACAT CGGCTCCAACTTCGTGAACTGGTACCAGCAGCTGCCCGGCACCGCCCCCCAAGGTGCTGATCTACACCAACAACCAGCGGCCCTCCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTTCCTGGCCTTCTCCGGCCTGCAGGACGAGGCCGAC TACTACTGTGCCCACCTACGACGACCATGCAGGCTGGGTGTTCGGCGGAGGCACCAAGCTGCCGTGCTGTCC
1916	VL of CDH19 65247.004	artificial	AA	QSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWYQQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCATYDESMQGWVFGGGTKLTVLS
1917	VH-VL of CDH19 65247.004	artificial	٢	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAACCTGGCGCCTCCGTGCGGGTGTCCTGCAAGGTGTCCGGCTACCCTT CACCAGCTACTTCATCCACTGGTCCGACAGGCCCAGGGCCTGGAATGGATGG
1918	VH-VL of CDH19 65247.004	artificial	AA	QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGSGGGSQSALTQPPSATGTPGQRVTISCSGSSSNIGSNFVNWY QQLPGTAPKVLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCATYDESMQGWVFGGGTKLTVLS
1919	CDH19 65247.004 x	artificial	AA	QVQLVQSGAEVKKPGASVRVSCKVSGYTFTSYFIHWVRQAPGQGLEWMGIINPISVSTSYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHLDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGGG

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.	15C			GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1920	CDR-H1 of CDH19 65249.002	artificial	AA	GYYWS
1921	CDR-H2 of CDH19 65249.002	artificial	AA	YIYYIGSTNYNPSLKS
1922	CDR-H3 of CDH19 65249.002	artificial	AA	DGSSGWYRWFDP
1923	CDR-L1 of CDH19 65249.002	artificial	AA	RASQSVSSSYLA
332 332	CDR-L2 of CDH19 65249.002	artificial	AA	GASSRAT
1925	CDR-L3 of CDH19 65249.002	artificial	AA	QQYGNSPLT
1926	VH of CDH19 65249.002	artificial	L	CAGGTGCAGCTGCAGGAATCCGGCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCTGACCTGCACCGGTGTCCGGCGGCGTCCAT CTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACATCGGCTCCACCA ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATGTCCGTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC GCCGCTGACACCCCAGAGTGCCCAGAGATGGCTCCTCCGGCTGGTACGTTCGACCCTTGGGGCCAGGGCACCCT GCCGCTGACACCCTTACTACTGCGCCCAGAGATGCTCCTCCGGCTGGTATCGTTCGACCCTTGGGGCCAGGGCCACCCT GGTCACCGTGTTCGACCTTGCGCCCAGAGATGCTCCTCCGGCTGGTATCGTTCGACCCTTGGGGCCAGGGCCACCCT
1927	VH of CDH19 65249.002	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSS
1928	VL of CDH19 65249.002	artificial	Z	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGACCACCCTGTCCTGCAGAGCCTCCCAGTCCGT GTCCTCCTCCTACCTGGCTTGGTATCAGCAGAAGCCCGGCCCGGCCTGCTGCTGCTGGTTCTTCGGCCTCTTCCAGAGCCACCG GCATCCCTGACCGGTTCTCCGGCTCTGGCTCCGGCACTTCACCCTGACCATCTCCCGGCTGGAACCCGAGGACTTCGCTGTG TACTACTGCCAGCAGTACGGCAACAGCCCCTGACCTTCGGCCAAGGCACAAGGTGAAATCAAGTCC
1929	VL of CDH19 65249.002	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAV YYCQQYGNSPLTFGQGTKVEIKS

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
1930	VH-VL of CDH19 65249.002	artificial	LN .	CAGGTGCAGCTGCAGGAATCCGGCCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCTGTCCGGGGGGCGCTCCAT CTCCGGCTACTACTGGTCCTGGATCCGGCAGCCCCTGGCAATGGAATGGATCGGCTACATCTACTACATCGGCTCCACCA ACTACAACCCCCAGAGTGACTCCAGGTGACCATGTCCTTGGAATGGATCGGCTTCCCTGAACCCAGGCTGACC GCCGTGACCCGGTGTACTACTGCGCCCAGAGATGGCTCCTCCGGTGGTTCGTCGTTCGT
1931	VH-VL of CDH19 65249.002	artificial	АА	QVQLQESGPGLVKPSETLSLTCTVSGGSISGYYWSWIRQPPGKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSSGGGGSGGGGGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWY QQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGNSPLTFGQGTKVEIKS
1932	CDH19 65249.002 x 12C	artificial	AA	QVQLQESGPCLVKPSETLSLTCTVSGGSISGYWWWIRQPPCKGLEWIGYIYYIGSTNYNPSLKSRVTMSVDTSKNQFSLKLSSVT AADTAVYYCARDGSSGWYRWFDPWGQGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWY QQKPGQAPRLLIFGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGNSPLTFGQGTKVEIKSGGGGSEVQLVESGG GLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAV YYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQK PGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1933	CDR-H1 of CDH19 65250.003	artificial	AA	SYYMS
1934	CDR-H2 of CDH19 65250.003	artificial	AA	IIHPSGGDTTYAQKFQG
1935	CDR-H3 of CDH19 65250.003	artificial	AA	GGIKLWLHFDY
1936	CDR-L1 of CDH19 65250.003	artificial	AA	SGSRSNIGSNFVN
1937	CDR-L2 of CDH19 65250.003	artificial	АА	TNNQRPS
1938	CDR-L3 of	artificial	AA	AVYDDSLNGWV

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
Ω NO.				
	CDH19 65250.003			
1939	VH of CDH19	artificial	M	CAGGTGCAGCTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACTT
	65250.003			CACCAGCTACTACATGTCCTGGGTCCGACAGGCCCCAGGCCAGGGCCTGGAATGGATGG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCAAGCTGTGGCTGCACTTCGACTACTGGGGGCCAGGGCACCCT GGTCACCGTGTCTAGC
1940	VH of CDH19 65250.003	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS
1941	VL of CDH19	artificial	F	CAGTCCGTGCTGACCCAGCCTCCTTCCGCCTCCGGCACCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65250.003			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACAACCAGCGGCCCTCCG
				GCGTGCCCGACCGGTTCTCTGGCTCCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGAC TACTACTGTGCCGTGTACGACGACGACGACGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1942	VL of CDH19 65250.003	artificial	ΑĄ	QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDDSLNGWVFGGGTKLTVLS
1943	VH-VI of	artificial	Ā	CAGGTGCAGCTGGTAGCAGTCGGCGCGCGAAGTAAAAACTTGGCGCCCTCCGTGAAAGTAAACCTTGGAAAAAAAA
34	CDH19	5	Ē	CINCLECTION OF THE CONTROLL OF
	65250.003			CCACCTACGCCCAGAAATTCCAGGGCAGAGTGACCATGACCGGCGACACCTCCACCTCCACCTGTATATGGAACTGTCCTCCTC
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCTTCAAGCTGCGCTGCACTTCGACTACTGGGGCCAGGGCACCT
				GGTCACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTTGGCGCGGCGGAGGCTCCCAGTCCGTGCTGACCCAGCCTCCTT
				CCGCCTCCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACATCGGCTCCAACTTCGTGAACTGGTTAT
				CAGCAGCTGCCCGGCACCGCCCCCCAAGCTGCTGATCTACACCAAGCGGGCGCCCTCCGGCGTGCCCGACCGGGTTCTCTGGCTC
				CAAGICIGGCACCICCGCCIGCCGIGGCCAICICGGCCIGCAGICIGAGGACGAGGCGGACIACIACIGIGCGGIGIACGACGACIA CCCTGAACGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGTCC
1944	VH-VL of	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL
	CDH19			RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGSGGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY Oot detta datti tatanmoddsgydddffsgraegraegraegraegraegraegraegraegraegrae
70.4	65250.003	3.1		XXII OIMI MHI IIMXXII OOVI DIA GOOMGI DAGAADA OODXOOLAAA OODXOOLAAA IYAA IYAA IYAA OOTIMII AADOO OOTIMII AADOO
1945	CDH19 65250.003 x	artificial	{	QVQLVQSGAEVKKRGASVKVSCKASKITFTSIIMSWVKQARGGLEWMGIIHRSGGDTTIAQKKFQCKVIMTGDTSTSTVIMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY
	120			QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDDSLNGWVFGGGTKLTVLSGGGGSEVQLVES
) 			GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT
				AVIICVILIONI GNOITTOIMAINGEGIIVIVOSSGGGGGGGGGGGGGGGGGTVVILGI VIIVOIVIIVOITTOININAVE QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLMYSNRWVFGGGTKLTVLHHHHHH
1946	CDR-H1 of	artificial	ΑΑ	SYYMS
	CDUTA			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
⊇ .				
	65250.004			
1947	CDR-H2 of CDH19	artificial	ΑA	IIHPSGGDTTYAQKFQG
	65250.004			
1948	CDR-H3 of	artificial	ΑA	GGIKLWIHFDY
	CDH19 65250.004			
1949	CDR-L1 of	artificial	AA	SGSRSNIGSNFVN
	CDH19			
	65250.004			
1950	CDR-L2 of	artificial	Ą	TNNQRPS
	CDH19 65250.004			
1951	CDR-L3 of	artificial	AA	AVYDESLQGWV
33	CDH19			
35	65250.004			
1952	VH of CDH19	artificial	N	
	65250.004			CACCAGCTACTACATGTCCTGGGTCCGACAGGCCCAGGCCTGGAATGGAATGGGCATGGGCATCATCCACCCCTCTGGCGGCGACA CCACCTACGCCGCGCGCGCGCCACA CCACCTACGCAAAATTCCAGGGCAGAGAGTGACCATGACCGGCGACACCTCCACCTCCACCTGTATAGGAACTGTCCTCCTCCTG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCAAGCTGTGGCTGCACTTCGACTACTGGGGCCAGGGCACCCTGGGGCCACCCTTCGACGTGTTCGACTACTGGGGCCACCCTTCGACGTGTTCGACTACTGGGGCCACCCTTCGACGTGTTCGACTACTACTACTACTACTACTACTACTACTACTACTACTA
1953	VH of CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL
	65250.004		ļ	
1954	VL of CDH19	artificial	Z	CAGICCGIGGLIGACCCAGCCICCGGCACCCCIGGCCAGCGCIGACCAICTCCGGCTCCAGCACAI
	65250.004			CGGCTCCGACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCCCCCAGGCTGCTGATCTACACCAACAACAGCGGCCCTCCGGCCCTCCGGCCCTGCAGCAGCGGGCCCTCCGGCCCTGCCGGCCTGCAGGACGAGGCCGGACCAACAACAACAACAACAACAACA
				TACTACTGTGCCGTGTACGACGACGCTCCTGCAGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1955	VL of CDH19 65250.004	artificial	ΑA	QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDESLQGWVFGGGTKLTVLS
1956	VH-VL of	artificial	NT	
	CDH19 65250 004			CACCAGCIACIACAIGICCIGGGICCGAGGCCCCAGGCCCAGGCCIGGAAIGGGCAIGGGCAICAICCACCCITCIGGCGCGCGGCGACA CCACCIACGCCCAGAAATICCAGGGCAGAGIGACCAIGACCGGCGACACCTCCACCTCCACCGTGIATAIGGAACIGICCICCTCCTCCTC
	+00.0020			

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 8				
				CCGCCTCCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGGTCCCGGCTCCAACATCGGCTCCAACTTCGTGAACTGGTAT CAGCAGCTGCCGGCACCGCCCCCAAGCTGATCTACACCAACAACAGCGGGCCTCCGGCGTGCCCGGACCGGTTCTTGGCTC CAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGGTCTGAGGACGAGGCCGACTACTACTGTGCCGTGTACGACGAGT CCCTGCAGGGCTGGTTCGGCGAGGCCAACCAAGCTGTTCTCTACGAGGCCGACTACTACTGTGCCGTGTACGACGAGT CCCTGCAGGGCTGGTTCGGCGAGCCACCAAGCTGACCTGTCC
1957	VH-VL of CDH19 65250.004	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLS
1958	CDH19 65250.004 x 12C	artificial	AA A	QVQLVQSGAEVKKPGASVKVSCKASRYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQUSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLSGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1959	CDR-H1 of CDH19 65250.005	artificial	AA	SYYMS
1960	CDR-H2 of CDH19 65250.005	artificial	АА	IIHPSGGDTTYAQKFQG
1961	CDR-H3 of CDH19 65250.005	artificial	AA	GGIKLWLHFDY
1962	CDR-L1 of CDH19 65250.005	artificial	АА	SGSRSNIGSNFVN
1963	CDR-L2 of CDH19 65250.005	artificial	АА	TNNQRPS
1964	CDR-L3 of CDH19 65250.005	artificial	АА	AVYDESLQGWV
1965	VH of CDH19 65250.005	artificial	LN	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCCCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT CACCAGCTACTACTACATGTCCTGGGTCCGACAGGCCCCAGGGCCTGGAATGGATGG

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ਮ ਹ ਹ		30016		
Š.				H C C C C C C C C C C C C C C C C C C C
				CGGAGCGAGGACACCGCGTGTACTACTGCGCCAGAGGCGGCATCAAGCTGTGGCTGCACTTCGACTGCGGGGCCAGGGCACCTT GGTCACCGTGTCTAGC
1966	VH of CDH19 65250.005	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSS
1967	VL of CDH19	artificial	N	
	65250.005			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACAACAGCGGCCCTCCG GCGTCCCCAACTTCTTCTGAGACGACGGCCCTCCG GCGTGCCCGACCTGCAGTCTGAGACGAGGCCGAC
				TACTACTGTGCCGTGTACGACGAGTCCCTGCAGGGCTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1968	VL of CDH19	artificial	AA	QSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDESIOGWYFGGGTKI,TVI,S
1969	VH-VI of	artificial	LN	
	CDH19	8	2	CACCAGCTACTACATGTCCTGGGTCCGACAGGCCCCCAGGCCTGGAATGGATGG
	65250,005			CCACCTACGCCCCAGAAATTCCAGGGCAGAGTGACCATGACCCGCGACACCTCCACCTCCACCGTGTATATGGAACTGTCCTCCTG
				GGTCACCGIGICTAGCGGAGGCGGAGGATCIGGIGGCGGTGGTICIGGCGGCGGAGGCTCCCAGICCGIGCTGACCCAGCCTCCTI
33				CCGCCTCCGGCACCCCTGGCCAGCGCGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACATCGGCTCCAACTTCGTGAACTGGTAT
_				CAGCAGCTIGUCUGGCACUGCUCUCUCAAGCTIGATICTACACCAACAACAACAGCGGCUCTICUGGCGTGCUCAACGGGCTTICTICTIGGCTCT
				CAAGICIGGCACCICCGCICCCIGGCCAICICCGGCCIGCAGICIGAGGACGAGGCCGACIACIGIGCCGIGIACGACGAGI CCCTGCAGGGCTGGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
1070	VH-VI of	ortificial	~	OVOTAVO GGA FLVIK PGA SLVIKUSCIK A SGVITFITS YVMSMADA PGOGT FILMIGT I H PSGGDITTI YA OK FOGRATIMIP PITSTILVYMETISST.
19/0	VH-VL 0 CDH19	al cilicia	{	ZVZHVZSCAEVAARTGASVAVSCAASCIIIISSIASAVARASATGASCERMIGIIIIISSGASTIIAASAI ZGAVIIIIADISISIVIMEERSSE RSEDTAVYYCARGGIKLWIHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY
	65250.005			QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLS
1971	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYYMSWVRQAPGQGLEWMGIIHPSGGDTTYAQKFQGRVTMTRDTSTSTVYMELSSL
	65250.005 x			RSEDTAVYYCARGGIKLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGGGGGGTTQPPSASGTPGQRVTISCSGSRSNIGSNFVNWY
	12C			QQLPGTAPKLLIYTNNQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVOPGGSLKLSCAASGFTFNKYAMNWVROAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLOMNNLKTEDT
1972	CDR-H1 of	artificial	AA	NYMS
	CDH19 65251.002			
1973	CDR-H2 of	artificial	AA	IINPSGGDSTYAQKFQG
	CDH19 65251 002			
7	2001TCZC0	1 - 1 - 1 - 1	•	OTOTIVITIES V
1974	CDR-H3 of	artificial	ΑA	GGIQLWLHFDY

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>.</u> ≥ 8				
	CDH19 65251.002			
1975		artificial	ΑA	SGSRSNIGSNFVN
	CDH19 65251.002			
1976	CDR-L2 of	artificial	AA	TNYQRPS
	CDH19 65251.002			
1977		artificial	AA	AVWDESLNGWV
	65251.002			
3 8 8	VH of CDH19 65251.002	artificial	F	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTT CACCAACTACTACTACTACTAGTCCTGGGCCCAGGCCCAGGGCCTGGGAATGGAATGGGCATCATCAACCCCTCTGGCGGCGACT CCACCTACGCCCAGAAGTTCCAGGGCCGGCTGACCATGACCGGCGACACCTCCACCTCCACCGTGTATAGGAACTGTCCTCCCTG CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGCACTTCGACTTCGACTACTGGGGCCAGGGCACCT
88 1979	VH of CDH19	artificial	AA	GGICACCOIGICIAGO QVQLVQSGAEVKKFGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIOLWIHFDVWGOGTLVTVSS
,	+		!	
1980	VL of CDH19 65251.002	artificial	E	CAGTOTGIGOTGACOCCAGOCOCCOTTOCGCOTOTGGCACOCOTGGCCAAAAGTGACCATOTOCTGCTOCGGCTOCCGGTOCAACAT CGGCTOCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCCCCAAGCTGCTGATOTACCAACTACCAGGCGCCTOCG GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGGTCTGAGGACGAGGCCGAC machine control contr
1981	VL of CDH19 65251.002	artificial	\{	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDESLNGWVFGGGTKLTVLS
1982		artificial	ځ	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCCTT CACCAACTACTACTACTAGTCTGGGTCCGACAGGCCCAGGGCCTGGAATGGATGG
1983	VH-VL of	artificial	ΑΑ	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGYLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
N S				
	CDH19 65251.002			QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLS
1984	СDH19 65251.002 x 12C	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1985	CDR-H1 of CDH19 65251.003	artificial	АА	NYYMS
1986	CDR-H2 of CDH19 65251.003	artificial	AA	IINPSGGDSTYAQKFQG
1987	CDR-H3 of CDH19 65251.003	artificial	AA	GGIQLWLHFDY
1988	CDR-L1 of CDH19 65251.003	artificial	AA	SGSRSNIGSNEVN
1989	CDR-L2 of CDH19 65251.003	artificial	AA	TNYQRPS
1990	CDR-L3 of CDH19 65251.003	artificial	AA	AVWDESLQGWV
1991	VH of CDH19 65251.003	artificial	Z	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTTT CACCAACTACTACATGTCCTGGGTCCGACAGGCCCAGGCCAGGGCCTGGAATGGATGG
1992	VH of CDH19 65251.003	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
1993	VL of CDH19	artificial	NT	CAGTCTGTGCTGACCCAGCCCCCTTCCGCCTCTGGCACCCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT

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S Q Q	~ .	DESIGNATION	SOURCE	IYPE	SEQUENCE
	-	65251.003			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGGGCCCTCCGGCCTCCGGCCTCCGAGCCGGCCCTCCGAGCCGACCGA
11	1994	VL of CDH19 65251.003	artificial	ΑΑ	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDESLQGWVFGGGTKLTVLS
15	1995	VH-VL of CDH19 65251.003	artificial	٢	CAGGTGCAACTGGTGCAGGTGCGCGCGAAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTTT CACCAACTACTACATGTCCTGGGTCCGACGGCCCCAGGGCCTGGAATGGATGG
					CCCTGCAGGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
340	1996	VH-VL of CDH19 65251.003	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFYGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLQGWVFGGGTKLTVLS
	1997	CDH19 65251.003 x 12C	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLQGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
1,5	1998	CDR-H1 of CDH19 65251.004	artificial	AA	NYYMS
11	1999 (CDR-H2 of CDH19 65251.004	artificial	AA	IINPSGGDSTYAQKFQG
7	2000	CDR-H3 of CDH19 65251.004	artificial	AA	GGIQLWLHFDY
20	2001 (CDR-L1 of CDH19	artificial	ΑA	SGSRSNIGSNFVN

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	65251.004			
2002	CDR-L2 of CDH19 65251.004	artificial	AA	TNYQRPS
2003	CDR-L3 of CDH19 65251.004	artificial	AA	AVYDESLQGWV
2004	VH of CDH19 65251.004	artificial	Z	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTT CACCAACTACTACTACTACTGTCCTGGGTCCGACAGGCCCAGGGCCTGGGAATGGATGG
2005	VH of CDH19 65251.004	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
341	VL of CDH19 65251.004	artificial	TN	CAGTCTGTGCTGACCCCAGCCCCCTTCCGCCTCTGGCACCCCTGGCCAACAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCTGCCCGCCC
1				GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCGAC TACTACTGTGCCGTGTACGACGACGAGGCTGGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
2007	VL of CDH19 65251.004	artificial	AA	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDESLQGWVFGGGTKLTVLS
2008	VH-VL of CDH19 65251.004	artificial	L	CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCCGGTACACCTT CACCAACTACTACTACATGTCCTGGGTCCGACAGGCCCAGGCCAGGGCCTGGAATGGATGG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCCAGTGGGCTGCACTTCGACTACTGGGGGCCAGGGCCCCT GGTCACCGTGTCTAGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGGGCGTCCCAGTCTGTGCTGACCCAGCCCCTT CCGCCTCTGGCACCGCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACTTCGTGAAAGTGACCCTT CAGCAGCTGCCCGCCCCCCAAAGTGACCTGTTCTACACCAACTACCAGCGGCCCTCCGGCGTCCCAACTTCGTGAAACTGGCTC CAGCAGCTGCCCGCCCCCCAAGCTGCTGATCTACACCAACTACAACGCGCCCTCCGGCGTGCCCGACCGGTTCTCTGGCTC CAAGTCTGGCACCTCCGCCTCCTGGCCATCTCCGGCCTGCAGGACGAGGCCGACTACTACTGTGCCGTGTACGACGAGT CCCTGCAGGCCTGCGCGGAGGCACCAAGCTGACCGTGCTGTCC
2009	VH-VL of CDH19 65251.004	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLS
2010	CDH19 65251.004 x	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASRYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTGDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLSGGGGSEVQLVES

SEQ ID NO.	DESIGNATION	SOURCE	ТҮРЕ	SEQUENCE
	12C			GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2011	CDR-H1 of	artificial	AA	NYYMS
	CDH19 65251 005			
2012	CDR-H2 of	artificial	Ą	IINPSGGDSTYAQKFQG
	CDH19 65251 005			
2013	CDR-H3 of	artificial	AA	GGIQLWLHFDY
	CDH19			
	65251.005			
2014	CDR-L1 of	artificial	ΑA	SGSRSNIGSNFVN
	CDH19 65251 005			
- 1	CDD.1.2.00	10:0:5:4:0	4	SAGONA
STO7 42	CDK-LZ of	artificial	¥	INIÇAFO
	65251.005			
2016	CDR-L3 of	artificial	Ą	AVWDDSINGWV
	CDH19 65251.005			
2017	VH of CDH19	artificial	N	
	65251.005			CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGCCCTGGAATGGATGG
2018	VH of CDH19 65251.005	artificial	АА	QVQIVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
2019	VL of CDH19	artificial	IN	CAGTCTGTGCTGACCCAGCCCCCTTCCGCCTCTGGCACCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65251.005			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGGGCCCTCCGGCCTCCGGCTCCCGACTCTGCAGGACGGCCCTCCGGTTCTCCGGCCTTCTGAGGACGAGGCCGACTACTACTACTGCAGTCTGAGGACGACGACCGAC
2020	VL of CDH19 65251.005	artificial	ΑΑ	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDDSLNGWVFGGGTKLTVLS

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
2021	VH-VL of CDH19 65251.005	artificial	Z	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGCCTTGGAATGGATGG
2022	VH-VL of CDH19 65251.005	artificial	АА	
343	CDH19 65251.005 x 12C	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGCGGSQXVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDDSLNGWYFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALITLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2024	CDR-H1 of CDH19 65251.006	artificial	АА	NYYMS
2025	CDR-H2 of CDH19 65251.006	artificial	AA	IINPSGGDSTYAQKFQG
2026	CDR-H3 of CDH19 65251.006	artificial	АА	GGIQLWLHFDY
2027	CDR-L1 of CDH19 65251.006	artificial	АА	SGSRSNIGSNFVN
2028	CDR-L2 of CDH19 65251.006	artificial	АА	TNYQRPS
2029	CDR-L3 of	artificial	AA	AVWDESLNGWV

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
Ö.				
	CDH19 65251.006			
2030	VH of CDH19	artificial	ΙN	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGGTACACCTT
	65251.006			CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGCCAGGGCCTGGAATGGATGG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCCAGAGGCGGCATCCAGCTGTGGCTGCACTTCGACTACTGGGGCCAGGGCACCT GGTCACCGTGTCTAGC
2031	VH of CDH19 65251.006	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPCQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
2032	VL of CDH19	artificial	F	CAGTCTGTGCTGACCCCAGCCCCCTTCCGCCTCTGGCACCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCAGCATCT
	65251.006			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGCGGCCCTCCG
				GCGTGCCCGACCGGTTCTCTGGCTCCAAGTCTGGCACCTCCGCCTCCCTGGCCATCTCCGGGCCTGCAGTCTGAGGACGAGGCCGAC TACTACTGTGCCGTGTGGGACGAGTCCCTGAACGGCTGGGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
2033	VL of CDH19 65251 006	artificial	ΑA	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVWDESLNGWVFGGGTKLTVLS
700	000777000		Ę	
5607 344	VH-VL 0	alcillicia	Ē	
4	CDH19 65251 006			CACCTACTACTACTATATCTGGGTCCGGCCGGCTGACCATGGCCAGGGCTTGGAATGGGCATGGGCATCATCAACCCTCTTGGGACTGCGCGGCGACTG
	00271:000			CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCACTTCGACTACTGGGGCCCAGGGCACCCT
				GGICACCGIGICIAGCGGAGGAGGAGGAICIGGIGGCGGIGGIICIGGCGGCGGAGGCICCCAGICIGIGCIGACCCAGCCCCII
				CCGCCTCTGGCACCCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACATCGGCTCCAACTTCGTGAACTGGGTAT
				CAGCAGCIGCCCGGCACCGCCCCCAAGCIGCIGAICIACACCAACIACCAGCGGCGCCCICCGGCGIGCCCGACCGGIICICIGGCIC
				CAAGICIGGCACCICCGCCICCCTGGCCAICICCGGCCTGCAGICIGAGGACGAGGCGGACIACIACIGIGCCGIGIGGGACGAGI CCCIGAACGACIGGGTGGGGGGGAGGGAACCAAGCIGACGTGTGTCC
2035	VH-VL of	artificial	Ą	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL
	CDH19			RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY
	65251.006			QQLPGTAPKLL1YTNYQRPSGVPDRF:SGSKSGTSASLA1SGLQSEDEADYYCAVWDESLNGWVFGGGTKLTVLS
2036	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL
	65251.006 x			RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSGGGSGGGSGYLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY
	12C			QQLFGIAFKLLIIINIQRFSGVFDRFSGSKSGISASLAISGLQSEDEADIICAVWDESLNGWVFGGGIRLIVLSGGGGSEVQLVES GGGIVOPGGSIRISCAASGFTFNKYAMNWVROAPGKGIEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYIOMNNIKTFDT
				AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ
				ZNFGZAFKGLIGGIRFLAFGIFAKFJGSDLGGKAALILJGGVZFEDEAEIICVLWISNKWVFGGGIRLIVLHHHHH
2037	CDR-H1 of CDH19	artificial	¥	NYYMS

C L	i	1000	Į.	
Х С	DESIGNATION	SOURCE	IYPE	SEQUENCE
S				
	65251.007			
2038	CDR-H2 of	artificial	ΑA	IINPSGGDSTYAQKFQG
	CDH19			
	65251.007			
2039	CDR-H3 of	artificial	Ą	GGIQLWIHFDY
	CDH19			
	65251.007			
2040	CDR-L1 of	artificial	ΑĄ	SGSRSNIGSNFVN
	CDH19			
	65251.007			
2041	CDR-L2 of	artificial	AA	INYQRPS
	CDH19			
	65251.00/			
2042	CDR-L3 of	artificial	AA	AVWDESLQGWV
-2	CDH19			
45	65251.007			
2043	VH of CDH19	artificial	F	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT
	65251.007			CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGGCCTGGAATGGATGG
				CCACCTACGCCCAGAAGTTCCAGGGCCGGCTGACCATGACCCGCGACACCTCCACCTCCACCGTGTATATGGAACTGTCCTCCTG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCGCACTTCGACTACTGGGGGCCCAGGGCACCCT
7700	0 F 1 P 1 P 1 P 1 P 1 P 1 P 1 P 1 P 1 P 1		< <	OUTO INTO CALABOTA STRITE OF A
7044	65251.007	al IIIIcial	{	ZVZIV ZOGAŁYNIK GASVIV SCIASCIII IN INSWYNZAF CZGŁEWMGIINE SGGŁEJIAŻNI ZGNIMINDIEJ IN INEBESE RSEDTAVYYCARGGIQLWIHFDYWGOGTLVTVSS
2045	VL of CDH19	artificial	F	CAGTCTGTGCTCAGCCCCCTTCCGCCTCTGGCACCCTTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65251.007			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGCGGCCCTCCG
				GCGIGCCCGACCGGTICITCITGGCICCAAGICIGGCACCICCGCCICCCTGGCCAICITCGGCCGTGCAGITGIGAGGACGAGGCCGAC TACIACIACIAACIAACAAGGACGAGGCCGAC TACIAACIAACAAAGAAAAAAAAAAAAAAAAAAAAAAAAA
2046	VL of CDH19	artificial	AA	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD
	65251.007			YYCAVWDESLQGWVFGGGTKLTVLS
2047	VH-VL of	artificial	L	CAGGTGCAGCTGGTGCAGTCTGGCGCCCCGAAGTGAAGAAACCTGGCGCCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT
	CDH19			CACCAACTACTACATGICCIGGGICCGACAGGCCCCCAGGCCTGGAAIGGAAI
	00271:00/			CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGCATCCAGCTGTGGCTGCTGCTACTGGGGCCCAGGGCCACCTTCGACTACTGGGGCCCCCT
				GGTCACCGTGTCTAGCGGAGGCGGGAGGATCTGGTGGTGGTTCTGGCGGCGGAGGCTCCCAGTCTGTGCTGACCCAGCCCCTTT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
o S O S				
				CCGCCTCTGGCACCCCTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCAGCTCCAACATCGGCTCCAACTTCGTGAACTGGTAT CAGCAGCTGCCCGGCACCGCCCCCAAGCTGATCTACACCAACTACCAGCGGCCCTCCGGCGTGCCCGACCGGTTCTCTGGCTC CAAGTCTGGCACCTCCCTCGCCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGACTACTACTGTGCCGTGTGGGACGAGT CCCTGCAGGGCTGGTTCGGCGAGGCCAACTACCAAGCTGTTCC
2048	VH-VL of CDH19 65251.007	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLQGWVFGGGTKLTVLS
2049	CDH19 65251.007 x 12C	artificial	Ą	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGSQSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVWDESLQGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDT AVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQ QKPGQAPRGLIGGTKFLAAPGTPARFSGSLLGGKAALITLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2050	CDR-H1 of CDH19 65251.008	artificial	AA	NYYMS
9 2051	CDR-H2 of CDH19 65251.008	artificial	АА	IINPSGGDSTYAQKFQG
2052	CDR-H3 of CDH19 65251.008	artificial	AA	GGIQLWLHFDY
2053	CDR-L1 of CDH19 65251.008	artificial	АА	SGSRSNIGSNFVN
2054	CDR-L2 of CDH19 65251.008	artificial	АА	INYQRPS
2055	CDR-L3 of CDH19 65251.008	artificial	АА	AVYDESLQGWV
2056	VH of CDH19 65251.008	artificial	LN	CAGGTGCAGCTGGTGCAGTCTGGCGCCCGAAGTGAAGAAACCTGGCCCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT CACCAACTACTACATGTCCTGGGTCCGACAGGCCCCAGGGCCTGGAATGGATGG

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N O				
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAGGCGGCATCCAGCTGTGGCTGCACTTCGACTACTGGGGGCCAGGGCACCTTCGACGACTACTGGGGGCCAGGGCACCCT
2057	VH of CDH19 65251.008	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSS
2058	VL of CDH19	artificial	IN	CAGTCTGTGCTGACCCAGCCCCCTTCCGCCTCTGGCACCCTTGGCCAGAAAGTGACCATCTCCTGCTCCGGCTCCCGGTCCAACAT
	65251.008			CGGCTCCAACTTCGTGAACTGGTATCAGCAGCTGCCCGGCACCGCCCCCAAGCTGCTGATCTACACCAACTACCAGCGGCCTCCG GCGTGCCCGACTTCTCTGAGGACGACGCCCTCCG
				TACTACTGTGCCGTGTACGACGAGTCCCTGCAGGGCTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
2059	VL of CDH19 65251.008	artificial	AA	QSVLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWYQQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEAD YYCAVYDESLQGWVFGGGTKLTVLS
2060	VH-VL of	artificial	NT	CAGGTGCAGCTGCAGTCTGGCGCCGAAGTGAAGAAACCTGGCGCCTCCGTGAAGGTGTCCTGCAAGGCCTCCGGGTACACCTT
	CDH19			CACCAACTACTACATGTCCTGGGTCCGACAGGCCCAGGCCAGGGCCTGGAATGGATGG
	65251.008			CCACCTACGCCCAGAAGTTCCAGGGCCGGCTGACCATGACCCGCGACACCTCCACCTCCACCGTGTATATGGAACTGTCCTCCTG
				CGGAGCGAGGACACCGCCGTGTACTACTGCGCCGGCATCCAGCGCGCTGTGGCCTTCTCGACTTTCGACTACTGGGGCCCAGGCACCCT GGTGACCGAGTTTACGCGAAGGAATCTGCTGCGCGGTGCTTTTGGCGCGCCGAAGTTCTTTCGAAGTTCTTTTTTTT
				のなるとなっては、これでは、これないでは、これでは、これでは、これでは、これでは、これでは、これでは、これでは、これ
347				COGCO GCIGOCO COCCO COCCO COCCO AGO TO TO COCO COCO COCO COCO COMO COCO COC
7				CAAGTUTGGCACCTCCGCCTCCCTGGCCATCTCCGGCCTGCAGTCTGAGGACGAGGCCGACTACTACTGTGCCGTGTACGACGAGT
				C.C.T.T.G.G.G.G.T.G.G.G.T.T.T.C.G.G.G.G.
2061	VH-VL of	artificial	ΑA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL
	CDH19 65251.008			RSEDTAVIYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGSGGGGSQSVLTQPPSASGTPGQKVT1SCSGSRSN1GSNFVNWY QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLS
2062	CDH19	artificial	AA	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMSWVRQAPGQGLEWMGIINPSGGDSTYAQKFQGRLTMTRDTSTSTVYMELSSL
	65251.008 x			RSEDTAVYYCARGGIQLWLHFDYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGYLTQPPSASGTPGQKVTISCSGSRSNIGSNFVNWY
	12C			QQLPGTAPKLLIYTNYQRPSGVPDRFSGSKSGTSASLAISGLQSEDEADYYCAVYDESLQGWVFGGGTKLTVLSGGGGSEVQLVES GGGLVOPGGSLKLSCAASGFTFNKYAMNWVROAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLOMNNLKTEDT
2063	CDR-H1 of	artificial	AA	SYDMD
	65252.005			
2064	CDR-H2 of	artificial	AA	VIMYDGSNKYYADSVRG
	CDH19 65252.005			
3000	CDP U2 of	cioiii+rc	«	PHOEDIMYENT
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SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
© .				
	CDH19 65252.005			
2066	CDR-L1 of	artificial	ΑA	RASQSVSSSYLA
	CDH19 65252.005			
2067	CDR-L2 of	artificial	AA	GASSRAT
	CDH19 65252 005			
2068	CDR-L3 of	artificial	Ą	QQYGSSWT
	CDH19 65252.005			
2069	VH of CDH19	artificial	F	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTTCAGCTTCAGCTTCAGCTTCAGCTTCAGCTTCAACAACAAGACTTCAACAACAACAACAACAACAACAACAACAACAACAA
34	000.202.000			
8 2070	VH of CDH19 65252.005	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFSFSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS
2071	VL of CDH19	artificial	ΤN	
	65252.005			GTCCTCCTCCTACCTGGCCTGGTATCAGCAGAAGCCCGGCCAGGCCCTCGGCTGGTGATCTACGGCGCCTCTTCCAGAGCCACCG GCATCCCTGACCGGTTCTCCCGGCTCTGGCTCCGGCACCGACTTCACCCTGACCATCAGCTCGCTGGAACCCGAGGACTTCGCTGTG TACTATTGCCAGCAGTACGGCTCCTTGGACTTTCGGCCAGGGCACCAAGGTGGAAATCAAGTTC
2072	VI of CDH19	artificial	AA	EIVLTOSPGTLSLSPGERATLSCRASOSVSSSYLAWYOOKPGOAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV
 	65252.005	5	, i	YYCQQYGSSWIFGQGTKVEIKS
2073	VH-VL of CDH19	artificial	Ä	CAGGIGCAGCIGGIGGAAICCGGCGGAGGCGIGGIGCAGCCIGGCGGGGTCCCIGAGACIGITIGICCIGCCGCCICCGGCITCAGCIT CICCICCIACGACAIGGACIGGGICCGACAGACCCCCGGCAAGGGCCIGGAAIGGGIGGCCGIGAITIGGIACGACGGCITCAACA
	65252.005			AGTACTACGCCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCTGTTTCTGCAGATGAACTCCCTG CGGGTGGAAGATACCGCCGTGTACTACTGCGCCCAGAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCAGAGGCACCCTGGT
				CACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGGAGGCTCCGAGATCGTGCTGACCCAGTCCCTGGCA
				CCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCCTGCAGAGCCTCCCAGTCCGTGTCTTCTTCTTCTTGTTGTTCTACCTGGTATCAG
				GGACCTTCGGCCAGGGCACCAAGGTGGAAATCAAGTCC
2074	VH-VL of	artificial	Ą	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGGGGGGGGGGGGTIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ

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N 0 %	DESIGNATION	SOURCE	YPE	SEQUENCE
	CDH19 65252.005			QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2075	CDH19 65252.005 x 12C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2076	CDR-H1 of CDH19 65252.006	artificial	AA	SYDMD
2077	CDR-H2 of CDH19 65252.006	artificial	ΑΑ	VIWYDGSNKYYADSVRG
349	CDR-H3 of CDH19 65252.006	artificial	ΑΑ	ETGEGWYFDL
2079	CDR-L1 of CDH19 65252.006	artificial	AA	RASQSVSSSYLA
2080	CDR-L2 of CDH19 65252.006	artificial	АА	GASSRAT
2081	CDR-L3 of CDH19 65252.006	artificial	AA	QQYGSSWT
2082	VH of CDH19 65252.006	artificial	Z	CAGGTGCAGCTGGTGGAATCCGGCGGGGGGTGGTGCAGCCTGGCGGGTCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGACATGGACTGGGTCCGACAGGCCCCCGGCAAGGGCCTGGAATGGGTGGCTGGTTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGCGGGCAGATTCACCATCTCCCGGGACAACACTCCAAGAACACCCTGTTTCTGCAGATGAATGA
2083	VH of CDH19 65252.006	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS
2084	VL of CDH19	artificial	N	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCCCTGTCCTGCAGAGCCTCCCAGTCCGT

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	65252.006			GTCCTCCTCCTACCTGGCCTGGTATCAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCGCCTCTTCCAGAGCCACCGGCCACCGGCTCTTCCAGAGCCACCGAGGCCACCGAGCCTCTGGCTCCGGGCTTCTCGCTGTGGTATCCTTCGCTGGAACCCGAGGAACCCGAGGAACCCGAGGACTTCGCTGTGTAATTGCCAGCAGAATCAAGTTCCAGCAGGAACCCTGGACCTTCGGCCGAGGCCAACAAGTACAAGTTCCAGCAGAAATCAAGTTCCAGCAGAAATCAAGTTCCAAGAAATCAAGTTCCAAGAAAAAAAA
2085	VL of CDH19 65252.006	artificial	ΑΑ	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYGSSWTFGQGTKVEIKS
2086	VH-VL of CDH19	artificial	N	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGAATGGAATG
	65252.006			AGTACTACGCCGACTCCGTGCGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTG CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCAGAGGCACCTGGT CACCGTGTCTAGCGGAGGAGGAGGATCTGGTGGCGGTGGTTCTGGCGGGGGGGG
				CCCTGTCCCTGAGCCCTGGCGAGAGAGACCACCCTGTCCTGCAGAGCCTCCCAGTCCGTGTCCTCCTACCTGGCCTGGTATCAGCCAGGAAGCCCGGCTCCTGACCTGGCCTGGTTCTCCGGCTTCTGG
				CTCCGGCACCACCACCACCAGCACCATCAGCTCGCTGGAACCCGAGGACTTCGCTGTGTACTATTGCCAGCAGTACGCTCCTCCT GGACCTTCGGCCAGGGCACCAAGGTGGAAATCAAGTCC
2087	VH-VL of	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL
350	CDH19 65252.006			RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGGGSETVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2088	CDH19 65252 006 v	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYXCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGGGGSEIVLTOSPGTLSLSPGERATLSCRASOSVSSSYLAWYO
	12C			
				CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHH
2089	CDR-H1 of	artificial	AA	SYDMD
	CDH19 65252.007			
2090	CDR-H2 of	artificial	ΑA	VIWYDGSNKYYADSVRG
	CDH19 65252.007			
2091	CDR-H3 of	artificial	ΑA	ETGEGWYFDL
	CDH19 65252.007			
2092	CDR-L1 of CDH19	artificial	AA	RASQSVSSSYLA

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	65252.007			
2093	CDR-L2 of CDH19 65252.007	artificial	AA	GASSRAT
2094	CDR-L3 of CDH19 65252.007	artificial	AA	QQYGSSWT
2095	VH of CDH19 65252.007	artificial	Z	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGACTGGGTCCGACAGGCCCCCGGCAAGGGCCTGGAATGGGTGGCGGTGATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGCGGGCAGATTCACCATCTCCCGGGACAGAACACCCTGTTTCTGCAGATGAATGA
2096	VH of CDH19 65252.007	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSS
351	VL of CDH19 65252.007	artificial	TN	GAGATCGTGCTGACCCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCCACCCTGTCCTGCAGAGCCTCCCAGTCCGT GTCCTCCTCCTACCTGGCCTGG
2098	VL of CDH19 65252.007	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYGSSWTFGQGTKVEIKS
2099	VH-VL of CDH19 65252.007	artificial	F	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGTCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGACATGGACTGGAC
2100	VH-VL of CDH19 65252.007	artificial	ΑΑ	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2101	CDH19 65252.007 x	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL

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SEQ. NO.	DESIGNATION	SOURCE	IYPE	SEQUENCE
	12C			VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGGGGGGGGGGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2102	CDR-H1 of	artificial	AA	SYDMD
	CDH19			
	65252.008			
2103	CDR-H2 of	artificial	¥	VIWYEGSNKYYAESVRG
	CDH19 65252 008			
2104	CDR-H3 of	artificial	AA	ETGEGWYFDL
	CDH19			
	65252.008			
2105	CDR-L1 of	artificial	ΑA	RASQSVSSYLA
	CDH19			
	65252.008			
52	CDR-L2 of	artificial	Ą	GASSRAT
2	CDH19			
	65252.008			
2107	CDR-L3 of	artificial	Ą	QQYGSSWT
	CDH19 65252 008			
2108	VH of CDH19	artificial	¥	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCCGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT
	65252,008			
				AGTACTACGCCGAGTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTG
				CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCCAAGGCACCCTGGT CACCGTGTCTAGC
2109	VH of CDH19 65252.008	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSS
2110	VL of CDH19	artificial	N	GAGATCGTGCTGACCCAGTCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGACCACCCTGTCCTGCAGAGACCTCCCAGTCCGT
	65252.008			GICCICCICCTACCIGGCOTGGIAICAGCAGAAGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCGCCCTCTTCCAGAGCCACCG
				GCATCCCTGACCGGTTCTCCGGCTCTGGCTCCGGCACCGACTTCACCCTGACCATCAGCTCGGTGGAACCCGAGGACTTCGCTGTG TACTATTGCCAGCAGTACGGCTCCTCCTCCTGGACCTTCGGCCAGGGCAAGGTGGAAATCAAGTCC
2111	VL of CDH19 65252.008	artificial	AA	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYGSSWTFGQGTKVEIKS

SEQ ID NO.	DESIGNATION	SOURCE	ТУРЕ	SEQUENCE
2112	VH-VL of CDH19 65252.008	artificial	TN .	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGGCCGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGACATGGACTGGAC
2113	VH-VL of CDH19 65252.008	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
353	CDH19 65252.008 x 12C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QKPGQAPRLLIYGASSRATGIPDRFSGSGSTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2115	CDR-H1 of CDH19 65252.009	artificial	АА	SYDMD
2116	CDR-H2 of CDH19 65252.009	artificial	AA	VIWYDGSNKYYADSVRG
2117	CDR-H3 of CDH19 65252.009	artificial	AA	ETGEGWYFDL
2118	CDR-L1 of CDH19 65252.009	artificial	АА	RASQSVSSSYLA
2119	CDR-L2 of CDH19 65252.009	artificial	АА	GASSRAT
2120	CDR-L3 of	artificial	AA	QQYGSSWT

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	CDH19 65252.009			
2121	VH of CDH19	artificial	M	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT
	65252.009			CTCCTCCTACGACATGGACTGGGGTCCGACAGACCCCCGGCAAGGGCCTGGAATGGGGTGGCCGTGATTTGGTACGACGCCTCCAACAAGAACTACCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAAACACCCTGTTTCTGCAGATGAACTCCCTG
				CGGGTGGAAGATACCGCCGTGTACTACTGCGCCCAGAGAGACAGGCGGGGGGGG
2122	VH of CDH19	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPCKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS
2123	VI of CD110	lcio!jit	F	(1973年) は、日本の日本の日本の日本の日本の日本の日本の日本の日本の日本の日本の日本の日本の日
6717	65252.009	al III clai	2	GTCCTCCTCCTACCTGGCCTGGTATCAGCAGAGGCCCGGCCAGGCCCTCGGCTGCTGATCTACGGCGCCCTCTTCCAGAGCCACC
				GCATCCCTGACCGGTTCTCCGGCTCTGGCTCCGGCACCGACTTCACCCTGACCATCAGCTCGCTGGAACCCGAGGACTTCGCTGTG TACTATTGCCAGCAGTACGGCTCCTCCTGGACCTTCGGCCAGGGCACCAAGGTGGAAATCAAGTCC
2124	VI of CDH19	artificial	ΔΔ	FIVI.TOSPGTI.ST.SPGERATT.SCRASOSVSSSVI.AWYOORPGOAPRII.TYGASSRATGIPDRESGSGSGTDFTI.TISSI.EPRDEAV
+ 217	65252.009	מונובומ	{	YYCQQYGSSWTFGQGTKVEIKS
2125	VH-VL of	artificial	N	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT
- 1	CDH19			CICCICCIACGACAIGGACIGGGICCGACAGACCCCCGGCAAGGGCCTGGAAIGGGIGGCCGIGAIIIGGIACGACGGCICCAACA
	6252.009			AGTACTACGCCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCCAAGAACACCCCTGTTTCTGCAGATGAACTCCCTG
				CGGGTGGAAGATACCGCCGTGTACTACTGCGCCCAGAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCCAGAGGCACCCTGGT
				CACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGTGGTGGTTCTGGCGGCGGAGGCTCCGAGATCGTGCTGACCCAGTCCCTGGCT
				CCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGCAGAGCCTCCCAGTCCGTGTCCTCCTCCTCCTACCTGGCTATCAG
				CAGAGGCCCGGCCAGGCCCCTCGGCTGCTGATCTACGGCGCCTCTTCCAGAGCCACCGGCATCCCTGACCGGTTCTCCGGCTCTTGG
				CTCCGGCACCGACTTCACCCTGACCATCAGCTCGCTGGAACCCGAGGACTTCGCTGTGTACTATTGCCAGCAGTACGGCTCCTCCT CCACCTTTCGCCCAGCCAACAACGTCCAAAAAAAAAA
2476	50 IV IIV	leie itie	<	OSITICATE CONTROLOGIA CONTROLO
2772	VH-VL UI	alıllıcıal	{	ZVŽBVEDGGGVVVŽEGGGDINDOCAZAGGGGGGGGGGGGGGGGGGGGGGGGGGGTTULTOGGNNTLIANDOVINGNILLIANDONALILLEDINGNILLIANDO RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGGGGGGGGGGGTTVLTOSPGTLSLSPGERATLSCRASOSVSSSYLAWYO
	65252.009			QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2127	CDH19	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQTPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL
	65252.009 x			RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGGGGGGGGGGGGGTLVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ
	12C			QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGSEVQLVESGGGI
				VQFGSSLKLSCASGFTFNKIAMNWVKQAFGKGLEWVAKIKSKINNIATIIADSVKDKFTISKDDSKNIAILQMNNLKIEDIAVII CVRHGNFGNSYISYWAYWGOGTLVTVSSGGGGGGGGGGGGGSOTVVTOEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVOOKPG
				QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2128	CDR-H1 of CDH19	artificial	AA	SYDMD
	2			

SOURCE TYPE SEQUENCE		artificial AA VIWYDGSNKYYADSVRG	artificial AA ETGEGWYFDL		artificial AA RASQSVSSSYLA		TASSRAT		artificial AA QQYGSSWT		artificial NT CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGTGCCCGCCTCCGGCTTCAGCTT	CTCCTCCTACGACATGGGCTCCGACAGGCCCCCGGCAAGGGCCTGGAATGGGTGGCTGATTTGGTACGACGGCTCCAACAACAACAACAACAACAACAACAACTCCCTG AGTACTACGCCGACTCCGGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTGT CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGAGAGACAGGCGAGGGCTGGTACTTCGACCTGTGGGGCAAGAGCCCTGGT	artificial AA QVQLVESGGGVVQPGGSIRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSS	artificial NT GAGATOGTGCTGACCCAGTCCCCTGGCACCCTGAGCCCTGGGCGAGAGAGCCACCCTGTCCTGCAGAGCCTCCTCCCAGTCCGT	GICCICCICCIACCIGGCTICGGTAICAGCAGAGGCCCGGCCC	artificial AA EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV	artificial NT CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCAGCCTGGCGGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT	CICCICCIACGACAIGGACIGGGICCGACAGGCCCCCGGCAAGGGCCIGGAAIGGGIGGCCGIGAIITIGGIACGACGGCICCAACA AGIACIACGCGGACICCGIGCGGGGCAGAITCACCAICICCCGGGACAACICCAAGAACACCTGIITICIGCAGAIGAACICCCTG CGGGIGGAAGAIACCGCCGIGIACIACIGCGCCAGAGAGAG
		artifi	artifi		artifi		ort:	5	artifi					artifi		artifi	artifi	
DESIGNATION	65252.010	CDR-H2 of CDH19 65252.010	CDR-H3 of	CDH19 65252.010	CDR-L1 of	CDH19	+		CDR-L3 of	CDH19 65252.010	-	65252.010	VH of CDH19 65252,010	VL of CDH19	65252.010	VL of CDH19 65252.010		CDH19 65252.010
SEQ ID NO.		2129	2130		2131		2132		2133	355	2134		2135	2136		2137	2138	

C I	DESIGNATION	SOLIBCE	TVDE	CEOI IENCE
NO .			- -	
				CCCTGTCCCTGAGCCCTGGCGAGAGACCACCCTGTCCTGCAGAGCCTCCCAGTCCGTGTCCTCCTACCTGGCCTGGTATCAG CAGAGGCCCGGCCAGGCCCTCGGCTGCTGATCTACGGCGCCTCTTCCAGAGCCACCGGCATCCCTGACCGGTTCTCCGGCTCTGG CTCCGGCACCTTCACCCTGACCATCAGCTCGCTGGAACCCGAGGACTTCGCTGTGTACTATTGCCAGCAGTACGGCTCCTCCT GGACCTTCGGCCAGGGCACCAAGGTGGAAATCAAGTCC
2139	VH-VL of CDH19 65252.010	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2140	CDH19 65252.010 x 12C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGRGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QRPGQAPRLIYGASSRATGIPDRFSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAYWGQGTLVTVSSGGGSGGGGSGTVVTUQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2141	CDR-H1 of CDH19 65252.011	artificial	АА	SYDMD
99 2142	CDR-H2 of CDH19 65252.011	artificial	AA	VIWYDGSNKYYADSVRG
2143	CDR-H3 of CDH19 65252.011	artificial	АА	ETGEGWYFDL
2144	CDR-L1 of CDH19 65252.011	artificial	АА	RASQSVSSSYLA
2145	CDR-L2 of CDH19 65252.011	artificial	АА	GASSRAT
2146	CDR-L3 of CDH19 65252.011	artificial	АА	QQYGSSWT
2147	VH of CDH19 65252.011	artificial	TN	CAGGTGCAGCTGGTGGAATCCGGCGGGGGGGTGCTGGCGGGTCCCTGAGACTGTCTTGTGCCGCCTCCGGCTTCAGCTT CTCCTCCTACGACATGGACTGGGTCCGACAGGCCCCCGGCAAGGGCCTGGAATGGGTGGCCGTGATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGCGGGGCAGATTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTTTCTGCAGATGAACTCCCTG

CEO.	DESIGNATION	SOLIBCE	TVDE	CECI IENCE
5 0 S			- -	
				CGGGTGGAAGATACCGCCGTGTACTACTGCGCCAGAGACAGGCCGAGGGCTGGTACTTCGACCTGTGGGGCCAAGGCACCTGGGT CACCGTGTCTAGC
2148	VH of CDH19 65252.011	artificial	ЧΑ	QVQLVESGGGVVQPGGSLRLSCAASGFSFSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSS
2149	VL of CDH19 65252.011	artificial	Ä	GAGATCGTGCTGACCCAGTCCCCTGGCACCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCCTGCAGAGCCTCCCAGTCCGT GTCCTCCTCCTACCTGGCCTGG
2150	VL of CDH19 65252.011	artificial	АА	EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAV YYCQQYGSSWTFGQGTKVEIKS
2151	VH-VL of CDH19	artificial	LN L	CAGGIGCAGCIGGIGGAAICCGGCGGAGGCGIGGIGCAGCCIGGCGGGICCCIGAGACIGICTIGIGCCGCCTCCGGCTTCAGCTT CTCAGCTTCAGCTTCAGCTTCAGCTTCAGCTTCAGCTTCGAACGGCCTCCAACA
357	65252.011			AGTACTACGCCGACTCCGTGCGGGCAGATTCACCCATCTCCCGGGACAACTCCAAGAACCCTGTTTCTGCAGATGAACTCCTG AGTACTACGCCGACTCCGTGCGCCCAGAGAGACAGGCGAGGCTGGTACTTCGACCTGTGGGGCCAAGGCACCTGGT CGGGTGGTACTACCGCCGTGTACTGCGCGCGGAGGCTGGTACTTCGACCTGTGGGGCCCAAGCCCTGGTA CACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGGGAGGCTCCGAGATCGTGCTGACCCCTGGCA CCCTGTCCCTGAGCCCTGGCGAGAGAGCCACCCTGTCCTGCAGAGCCTCCTGCTGCTCCTGCTGGCAGGCCCTGGCTACTGGCCTGGCTACTGGCCTGGCTCCTGGCCTGGTATCAGGCCCCGGCTCTTCCGGCACCCGGCACCCGGCTTCTCCGGCTCTTCCGGCCTCTTCGGCACCCGGCACCTGACCGGCTCTTCGGCTCCTTCGGCACCCGGCACCTGACCGGCTCCTCGGCTCCTCGGCCCCTCGGCACCCGGCACCAGGCCCCTGACCAGAGACCCAGGAACCCGAGAACCCGAGAACCCGAGAACCCGAGAACCCGAGAACCCAGGGAACCCGAGAACCCGAGAACCCGAGAACCCGAGAACCCGAGAACCCAGGGAACCCGAGAACCCGAGAACCCGAGAACCCAGAGAACCCAGGAACCCAGAGAACCCAGAGAACCCAGAGAACCCCTGACCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAAGAACCCAACACAACA
2152	VH-VL of CDH19 65252.011	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGGGGGGGTIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2153	CDH19 65252.011 x 12C	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYDGSNKYYADSVRGRFTISRDNSKNTLFLQMNSL RVEDTAVYYCARETGEGWYFDLWGQGTLVTVSSGGGGSGGGGSGGGGSEIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQ QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWAQWGGTLVTVSSGGGGSGGGGSGGGCSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKPG QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2154	CDR-H1 of CDH19 65252.012	artificial	АА	SYDMD
2155	CDR-H2 of CDH19 65252.012	artificial	AA	VIWYEGSNKYYAESVRG
2156	CDR-H3 of	artificial	AA	ETGEGWYFDL

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
<u>0</u> 9				
	CDH19 65252.012			QRPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKS
2166	CDH19	artificial	₽¥	QVQLVESGGGGVVQPGGSLRLSCAASGFSFSSYDMDWVRQAPGKGLEWVAVIWYEGSNKYYAESVRGRFTISRDNSKNTLFLQMNSL
	65252.012 x 12C			RVEDIAVIICAREIGEGWIFDLWGVGILVIVSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGTLSGGGTLGCRASQSVSSSILAWIQ QRPGQAPRLIIYGASSRATGIPDRFSGSGSGTDFTLTISSLEPEDFAVYYCQQYGSSWTFGQGTKVEIKSGGGGSEVQLVESGGGL VQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVYY CVRHGNFGNSYISYWACGGTLVTVSSGGGGSGGGGSGGGGSOTVVTOEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVOOKPG
				QAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2167	CDR-H1 of	artificial	AA	SYSWS
	CDH19 65253.003			
2168	CDR-H2 of	artificial	AA	YIYYSGSTNYNPSLKS
	CDH19			
	65253.003			
2169	CDR-H3 of	artificial	ΑA	NWAFHFDY
359	CDH19			
	65253.003			
2170	CDR-L1 of	artificial	ΑĄ	IGSSSNIGTGYDVH
	CDH19			
	65253.003			
2171	CDR-L2 of	artificial	Ą	GNSNRPS
	CDH19			
	65253.003			
2172	CDR-L3 of	artificial	Ą	QSYESSLSGWV
	CDH19 65253.003			
2173	VH of CDH19	artificial	NT	CAGGIGCAGCIGCAGGAAICCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCCAI
	65253.003			CICCICCIACITALICIIGGILCIIGGAILCCGGCAGCCCCCIIGGCAGGGCCCIIGGAAIIGGAILCGGCIIACAICIACIIALICICGGCILCACCA ACIACAACCCCAGCCIGAAGILCCAGAGIGACCAICICCCIGGACACCACAAGAACCAGTICICCCIGAAGGCGIICICCIIGAAGCIIGICCIIGAAGCIIGICCIIGA
2174	VH of CDH19 65253.003	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT AADTAVYYCARNWAFHFDYWGQGTLVTVSS
2175	VI of CDH19	artificial	ΙN	CAGTICTICTICACICIA GICTACICTICTICTICTICTICTICTICTICTICTICTICTICT
C/T7	VL VI CUPITS	dı ullua	Ξ	[

Ĺ	H 4 4 0 10 L 4	100	i di	
S O S	DESIGNATION	SOURCE	7 Y P E	SEQUENCE
	65253.003			CGGCACCGGCTACGACGTGCTATCAGCAGCTGCCCGGCACCGCCCCCCAAGCTGATCTACGGCAACTCCAACCGGCCTTCTCCCAACCGGCCCTTCTCTACGGCAACCGGCCCTTTCTCTCTC
2176	VL of CDH19 65253.003	artificial	AA	QSVLTQPPSVSGAPGQRVTISCTGSSSNIGTGYDVHWYQQLPGTAPKLLIYGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEA DYYCQSYESSLSGWVFGGGTKLTVLS
2177	VH-VL of CDH19	artificial	TN	CAGGIGCAGCIGCAGGAATCCGGCCCTGGCCTGGTCAAGCCCTCCGAGACACTGTCCCTGACCTGCACCGTGTCCGGCGGCTCCAT CTCCTCCTACTCTTGGTCCTGGATCCGGCAGCCCCCTGGCAAGGGCCTGGAATGGATCGGCTACATCTACTACTACTCCGGCTCCACCA
	65253.003			ACTACAACCCCAGCCTGAAGTCCAGAGTGACCATCTCCCTGGACACCTCCAAGAACCAGTTCTCCCTGAAGCTGTCCTCCGTGACC GCCGCTGACACCGCCGTGTACTACTGCGCCCGGAACTGGGCCTTCCACTTCGACTACTGGGGCCCAGGGCACCTGGTCACCGTGTC TAGCGGAGGAGGAGGATCTGGTGGCGGTGGTTCTGGCGGGGGGCCCAGTCTGTGTCTGTGTCTGGCC
				CACCICCCICCITACAACGACCIACCAGACIGAGACGAGAC
2178	VH-VL of	artificial	AA	
360	CDH19 65253.003			AADTAVIICAKNWAFHFDIWGQGTLVIVSSGGGGGSGGGGGGGGGGGGGGGGGGGGGGGGGTGGTGAFGQKVIISCIGSSSNIGIGIGIDVHWIQQL PGTAPKLLIYGNSNRPSGVPDRFSGSKSGTSASLAIIGLQAEDEADYYCQSYESSLSGWVFGGGTKLIVLS
2179	CDH19	artificial	AA	QVQLQESGPGLVKPSETLSLTCTVSGGSISSYSWSWIRQPPGKGLEWIGYIYYSGSTNYNPSLKSRVTISLDTSKNQFSLKLSSVT
	65253.003 x 12C			AADIAVIICAKNWAFRIDIMGGGILVIVSSGGGGGGGGGGGGGGGGGGGSXVLIZFFSVSGAFGGFVIISCIGSSSNIGIGIDVRWIRGE PGTAPKLLIYGNSNRPSGVPDRFSGSKSGTSASLAITGLQAEDEADYYCQSYESSLSGWVFGGGTKLTVLSGGGGSEVQLVESGGG
				LVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTEDTAVY YCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWVQQKP GOAPRGITGGTKFTAPGTPARFSGSILGGKAALTLSGVOPFDFAFSVYCVLMYSNRWVFGGGTKTTWYHHHHHHH
2180	CDR-H1 of	artificial	Ą	SYGMH
	CDH19 65254.001			
2181	CDR-H2 of	artificial	ΑA	FIWYDGSNKYYADSVKD
	CDH19 65254.001			
2182	CDR-H3 of	artificial	ΑA	RAGIIGTIGYYYGMDV
	CDH19 65254.001			
2183	CDR-L1 of CDH19	artificial	AA	SGDRLGEKYTS

SEQ NO	DESIGNATION	SOURCE	TYPE	SEQUENCE
	65254.001			
2184	CDR-L2 of CDH19 65254.001	artificial	АА	QDTKRPS
2185	CDR-L3 of CDH19 65254.001	artificial	AA	QAWDSSTVV
2186	VH of CDH19 65254.001	artificial	Z	CAGGTGCAGCTGGTGGAATCCGGCGGGGCGTGGTGCAGCCTGGCCGGTCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT CTCCAGCTACGCATGCACTGGGCGACAGGCCCTGGCAAGGGCCTGGGAATGGGTGGCTCTTCATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAGAACTCCAAGAACACCCTGTACCTGCAGATGAAGTCCTG CGGGCCGAGGACACCGCCGTGTACTACTGTGCCCAGAAGGGCCGGCATCATCGGCACCTTACGGCTACTACGGCATGGACGTGTG GGGCCGAGGACACCGCCGTGTACTACTACTGTGCCAGAAGGGCCGGCATCGGCACCATCGGCTACTACGGCATGGACGTGTG GGGCCAGGGCCACCGTGTACTACTACTACTAGC
2187	VH of CDH19 65254.001	artificial	AA	QVQLVESGGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
361	VL of CDH19 65254.001	artificial	TN	TCTTACGAGCTGACCCCAGCCCCCCCTCCGTGTCCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGA GAAGTACACCAGTTGGTATCAGCAGCGGCCTGGCCAGTCCCCCTGCTGGTCATCTACCAGGACACCAAGCGGCCCTCCGGCATCC CTGAGCGGTTCTCCGGCTCCCAACTCCGGCAACACCCGCCACCCTGACCATCTCCGGCACCAGGCCATGGACGAGGCCGACTACTAC TGCCAGGCCTGGGACTCCTCCACCGTGGTGTTCGGCGGAGGCACAAGCTGACCGTGCTGTCC
2189	VL of CDH19 65254.001	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVLS
2190	VH-VL of CDH19 65254.001	artificial	LN .	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCCGGCCTGGCCTGAGACTGTCTTGCGCCCCCCCGGCTTCACCTT CTCCAGCTACGGCATGCACTGGGTCCGACAGGCCCTGGCAAGGGCCTGGAATGGGTGGCTTCATTTGGTACGACGGCTCCAACA AGTACTACGCCGACTCCGTGAAGGACCGGTTCACCTTCCCGGGACAACTCCAAGAACACCCTTGTACCTGCAGATGAAGTCCTG CGGGCCGAGGACACCGCCGTGTACTGTGCCAAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACGGCATGAAGTCCTG GGGCCAGGGCACCCCCGTGTACTTACTGTGCCAGACGGCCGCATCTCGGCGCTACTACTACGGCATGCTGTG TGACCCAGGCCACCCCTCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGGAGGCTCCTTACGAGC AGTTGGTATCAGCAGCGCCTGCTCTCTAGCGGAGCCGCACCCTCCATCTACCAGGCCCTCCGGCTTACTACCACC AGTTGGTATCAGCAGCGCCTGCCCTGC
2191	VH-VL of CDH19 65254.001	artificial	AA	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLS
2192	CDH19 65254.001 x	artificial	ΑА	QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE

SEQ	DESIGNATION	SOURCE	TYPE	SEQUENCE
Q. 0.				
	12C			SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2193	CDR-H1 of	artificial	AA	SYGMH
	CDH19 65254.003			
2194	CDR-H2 of	artificial	AA	FIWYDGSNKYYADSVKD
	CDH19 65254.003			
2195	CDR-H3 of	artificial	AA	RAGIIGTIGYYYGMDV
	CDH19			
	65254.003			
2196	CDR-L1 of	artificial	ΑA	SGDRLGEKYIS
	CDH19 65254 003			
	03234.003	1 - 1 - 1 - 1 - 1	4	Sagamao
/617 62	CDK-LZ of	artificial	¥	VDINKES
	65254.003			
2198	CDR-L3 of	artificial	AA	QAWDSSTVV
	CDH19 65254 003			
2199	VH of CDH19	artificial	M	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65254.003			CICCAGCTACGGCATGCACTGGGTCCGACAGGGCCCTGGCAAGGGCCTGGAATGGGTGGCCTTCATTTGGTACGACGGCTCCAACA
				AGTACTACGCCGACTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCTGTGTACCTGCAGATGAAGTCCCTG
				CGGGCCGAGGACACCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACTACGGCATGGACGTGTG GGGCCAGGGCACCACCGTGACCGTGTCTAGC
2200	VH of CDH19	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGTIGTIGYYYGMDVWGOGTTYTTSS
2000	0227.003	101019144	H	
7701	VL of CDH19 65254.003	artificial	Z	TUTTALGAGETIGALECCAGECEGECTEGECCAGTECCCTGCTGGTCATCTACCAGGACAGCGGCCCTCCGGCATCCGGCAGCGGCCTCCGGCATCCGGCAGCGGCCTCCGGCATCC
				CTGAGCGGTTCTCCGGCTCCCAACTCCGGCAACACCGCCACCCTGACCATCTCCGGCACCCAGGCCATGGACGAGGCCGACTACTAC TGCCAGGCCTGGGACTCCTCCACCGTGTTCGGCGGAGGCACCAAGCTGACCGTGCTGTCC
2202	VL of CDH19 65254.003	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY CQAWDSSTVVFGGGTKLTVLS

			1	
NO.	DESIGNATION	SOURCE	YPE	SEQUENCE
2203	VH-VL of CDH19 65254.003	artificial	LN .	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACTTCACTTGGTACGACGCTTCACTTCACTTGGTACGACGCTTCACTTCACTTTGGTACGACGCTTCACTTCACTTCACTTCACTTCACTTCACTACGACGCTCCAACAACACCCTCCAGCAGCTCCAACAAACA
2204	VH-VL of CDH19 65254.003	artificial	АА	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLS
363	CDH19 65254.003 x 12C	artificial	AA	QVQLVESCGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMKSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWDSSTVVFGGGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
2206	CDR-H1 of CDH19 65254.007	artificial	АА	SYGMH
2207	CDR-H2 of CDH19 65254.007	artificial	АА	FIWYEGSNKYYAESVKD
2208	CDR-H3 of CDH19 65254.007	artificial	AA	RAGIIGTIGYYYGMDV
2209	CDR-L1 of CDH19 65254.007	artificial	АА	SGDRLGEKYTS
2210	CDR-L2 of CDH19 65254.007	artificial	AA	QDTKRPS
2211	CDR-L3 of	artificial	AA	QAWESSIVV

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
	CDH19 65254.007			
2212	VH of CDH19	artificial	ΙN	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
	65254.007			CTCCAGCTACGGCATGCACTGGGTCCGACAGGCCCCTGGCAAGGGCCTGGAATGGGTGGCCTTCATTTGGTACGAGGGCTCCAACA AGTACTACGCCGAGTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAATTCCCTG
				CGGGCCGAGGACACCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACGGCATGGACGTGTG
2213	VH of CDH19	artificial	AA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL
	65254.007			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSS
2214	VL of CDH19	artificial	ΙN	TCTTACGAGCTGACCCCAGCCCCCCTCCGTGTCCTGTCTCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGA
	65254.007			GAAGTACACCAGTTGGTATCAGCAGCGGCCTGGCCAGTCCCCCTGGTGGTCATCTACCAGGGACACCAAGCGGCCCTCCGGCATCC
				CIGAGCGGIICICCGGCICCAACICCGGCAACACCGCCACCCIGACCAICICCGGCACCCAGGCCAIGGACGAGGCGGCIACIAC IGCCAGGCCIGGGAGICCICCACCGIGGIGIITCGGCGGAGGCACCAAGCIGACCGIGCIGICC
2215	VL of CDH19	artificial	AA	SYELTQPPSVSVSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYY
	65254.007			CQAWESSTVVFGGGTKLTVLS
2216	VH-VL of	artificial	NT	CAGGTGCAGCTGGTGGAATCCGGCGGAGGCGTGGTGCAGCCTGGCGGGTCCCTGAGACTGTCTTGCGCCGCCTCCGGCTTCACCTT
- 1	CDH19			CICCAGCIACGGCAIGCACIGGGICCGACAGGCCCCIGGCAAGGGCCIGGAAIGGGIGGCCIICAIIIGGIACGAGGGCICCAACA
	65254.007			AGTACTACGCCGAGTCCGTGAAGGACCGGTTCACCATCTCCCGGGACAACTCCAAGAACACCCTGTACCTGCAGATGAATTCCCTG
				CGGGCCGAGGACACCGCCGTGTACTACTGTGCCAGAAGGGCCGGCATCATCGGCACCATCGGCTACTACTACGGCATGGACGTGTG
				GGGCCAGGGCACCACCGTGACCGTGTCTAGCGGAGGCGGAGGATCTGGTGGCGGTGGTTCTGGCGGCGCGGGGGCTCCTTACGAGC
				TGACCCAGCCCCCTCCGTGTCCGTGTCTCCTGGCCAGACCGCCTCCATCACCTGTTCTGGCGACCGGCTGGGCGAGAGTACACC
				AGTIGGIAICAGCAGCGGCCIGGCCAGICCCCCCTGCIGGICAICIACCAGGACACCAAGCGGCCCICCGGCAICCCIGAGCGGTI
				CICCGGCTCCAACTCCGGCAACACCCCCCTGACCATCTCCGGCACCCATGGCCCATGGACGAGGCCGACTACTACTGCAGGCCT
				GGGAGTCCTCCACCGTGGTGGTTCGGCGGAGGCACCAAGCTGACGTGCTGTCC
2217	VH-VL of	artificial	₹	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL
	CDH19			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT smyooddoogdiiiiidddagddaciidddgcanscmmamimictscmoamdaaadyycoamgcmyygcccmyimiis
	65254.007			ONIXXING COLUMN 1.XIII.NO COLUMN 13.111.001XIII.DEAU 1.VXIVIDEAU 1
2218	CDH19	artificial	ΑA	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKGLEWVAFIWYEGSNKYYAESVKDRFTISRDNSKNTLYLQMNSL Rabdtavyyygarragiigtigyyygmdwmogettyvytssgggggssggggssgyfitoppsyyygpggtastffkyt
	03234.007 x			SWOOD POOR TO STATE THE PROPERTY OF THE PROPER
	127			SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGTVVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLHHHHHH
2219	CDH19 14302 CC x I2C-LFcBY	artificial	aa	QRFVTGHFGGLYPANGGGGGGSQVQLVESGGGVVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSYELTQPPSVS

SEQ NO	DESIGNATION	SOURCE	TYPE	SEQUENCE
				VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGGSEVQLVESGGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGGSGTVVTQEPSLTVSPGGTVTLCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTTKLTVLGGGGS QRFCTGHFGGLHPCNGHHHHHH
2220	CDH19 14302 CC x I2C- LFcBY-156	artificial	e e	QREVTGHFGGLYPANGGGGGSQVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVK DRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGGGGGGGGGGGSYELTQPPSVS VSPGQTASITCSGDRLGEKYTSWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVF GCGTKLTVLSGGGGSEVQLVESGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFT ISRDDSKNTAYLQMNNLKTEDTAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSGTVVTQEPSLTVSPGGT VTLTCGSSTGAVTSGNYPNWVQQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGG TKLTVLGGGGS QRFCTGHFGGLHPCNG GGGGS RDWDFDVFGGGTPVGGHHHHHHH
365	CDH19 14302 CC x I2C-Cys- Loop	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGSSGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLGCGGGCCHHHHHH
2222	CC x I2C-HALB	artificial	e e	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED TAVYYCKPGNFGNSTISYWAXWGQGTLVTVSSGGGGSGGGGGGGGGGTVVTQFPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTVTLTCGSSTGAVTSGNYPNWV QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTVTLTVLPGGGGSDAHKSEVAH RFKDLGEENFKALVLIAFAQYLQQCPFEDHVKLVNEVTEFAKTCVADESAENCDKSLHTLFGDKLCTVATLRETYGEMADCCAKQE PERNECFLQHKDDNPNLPRLVRPEVDVMCTAFHDNEETFLKKYLYEIARRHPYYAPPELLFFAKRYKAAFTTCCCQAADKAACLLFK LDELRDGKASSAKQRLKCASLQKFGERAFKAWAVARLSQRFPEFAKTVYAPEAKNYTGMFLYTKKVPQVSTPTLVEVSRNLGKVGSKCCK HPEAKRMPCAADPHECYAKVFDEFKPLVEEPQNLIKQNCELFEQLGEYKFQNALLLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCK HPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKCCTESLVNRRPCFSALEVDETYVPKEFNAETTFHADICTLSEKERQIKK QTALVELVKHKPRATKEQLKAVMDDFAAFVEKCCKADDKETCFAEEGKKLVAASQAALGLDYHHHHH
2223	CDH19 14302 CC x I2C-GS- D3HSA	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNIKTED TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGSGGGSGGGSQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV

SEQ ID	DESIGNATION	SOURCE	TYPE	SEQUENCE
NO.				
		_		QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSEEPQNLIKQ NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC CTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFAAFVEKCCKAD
				DKETCFAEEGKKLVAASQAALGLHHHHHH
2224	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x 12C-3GS-			RAEDTAVÝYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKÝT SWYOORPGOSPLLVIYODTKRPSGIPERFSGSNSGNTATLTISGTOAMDEADYYCOAWESSTVVFGCGTKLTVLSGGGGSEVOLVE
	USHSA			SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSGGGSGGGG
				SEEPQNLIKQNCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEK
				TPVSDRVTKCCTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFA
				AFVEKCCKADDKETCFAEEGKKIVAASQAALGLHHHHHH
2225	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x 12C-GS-			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGSGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
-	D3HSA-156			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE
36				SGGGLVQPGGSIKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
6				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSEEPQNLIKQ
				NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC
				CTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFAAFVEKCCKAD
				DKETCFAEEGKKLVAASQAALGL GGGGSGGGS RDWDFDVFGGGTPVGG HHHHHH
2226	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x I2C-3GS-			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
	D3HSA-156			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE
				SGGGLVQPGGSIKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGGGGGQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSGGGGSGGGG
				SEEPQNLIKQNCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEK
				HADICTLSEKERQIKKQI
				AFVEKCCKADDKETCFAEEGKKLVAASQAALGL GGGGSGGGS RDWDFDVFGGGTPVGG HHHHHH
2227	CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x I2C-GS-			RAEDIAVIICARRAGIIGIIGIIGIIGVAGOOTIVSSGGGGSGGGGSGGGGSSIELIQPPSVSVSPGGIASIICSGDRLGERYI smxooppcostivivodhrebbscibppppgsnsongnhamimischoamdeadvooamiessmyyeccomumivi
	D3HSA-21			OWIÇKEGÇOFELVIIÇDIRKEGGIFEKEGGGEGNOGNIAILIIGGIÇAMDEADIICÇAMEGOIVVEGCGINLIVDGGGGGGEVÇLVE SGGGTAVOPGGSIKISCAA SGFTFNKVAMNMAVROA PGKGLEMAAA TRSKYNNYATYYA DSYKDRFTISRDDSKNTA VIOMNNIKTED
				TAVYYCVRHGNEGNSYTSYWAYWGOGTTATVSGGGGGGGGGGGGGGGGGGGGGGGGGGGGGTTATTATTATT

SEQ ID NO.	DESIGNATION SOURCE	SOURCE	TYPE	SEQUENCE
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSEEPQNLIKQ NCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKC
				CTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFAAFVEKCCKAD
				DKETCFAEEGKKLVAASQAALGL GGGGSGGGS RLIEDICLPRWGCLWEDD HHHHHH
2228	2228 CDH19 14302	artificial	aa	QVQLVESGGGVVQPGGSLRLSCAASGFTFSSYGMHWVRQAPGKCLEWVAFIWYDGSNKYYADSVKDRFTISRDNSKNTLYLQMNSL
	CC x I2C-3GS-			RAEDTAVYYCARRAGIIGTIGYYYGMDVWGQGTTVTVSSGGGGSGGGGGGGGSSYELTQPPSVSVSPGQTASITCSGDRLGEKYT
	D3HSA-21			SWYQQRPGQSPLLVIYQDTKRPSGIPERFSGSNSGNTATLTISGTQAMDEADYYCQAWESSTVVFGCGTKLTVLSGGGGSEVQLVE
				SGGGLVQPGGSLKLSCAASGFTFNKYAMNWVRQAPGKGLEWVARIRSKYNNYATYYADSVKDRFTISRDDSKNTAYLQMNNLKTED
				TAVYYCVRHGNFGNSYISYWAYWGQGTLVTVSSGGGGGGGGGGGGGGQQTVVTQEPSLTVSPGGTVTLTCGSSTGAVTSGNYPNWV
				QQKPGQAPRGLIGGTKFLAPGTPARFSGSLLGGKAALTLSGVQPEDEAEYYCVLWYSNRWVFGGGTKLTVLPGGGGSGGGGSGGGG
				SEEPQNLIKQNCELFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCCKHPEAKRMPCAEDYLSVVLNQLCVLHEK
				TPVSDRVTKCCTESLVNRRPCFSALEVDETYVPKEFNAETFTFHADICTLSEKERQIKKQTALVELVKHKPKATKEQLKAVMDDFA
				AFTVEKCCKADDKETCFAEEGKKLVAASQAALGL GGGGSGGGS RLIEDICLPRWGCLWEDD HHHHHH

Claims

1. An isolated multispecific antibody construct comprising a first human binding domain capable of binding to human CDH19 on the surface of a target cell and a second domain capable of binding to human CD3 on the surface of a T cell.

- 2. The antibody construct according to claim 1, wherein the first binding domain comprises a VH region comprising CDR-H1, CDR-H2 and CDR-H3 and a VL region comprising CDR-L1, CDR-L2 and CDR-L3 selected from the group consisting of:
 - (a) CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 222,

CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 252,

CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 84, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927,

CDR-H1 as depicted in SEQ ID NO: 82, CDR-H2 as depicted in SEQ ID NO: 83, CDR-H3 as depicted in SEQ ID NO: 909, CDR-L1 as depicted in SEQ ID NO: 250, CDR-L2 as depicted in SEQ ID NO: 251 and CDR-L3 as depicted in SEQ ID NO: 927,

CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 54, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926.

CDR-H1 as depicted in SEQ ID NO: 52, CDR-H2 as depicted in SEQ ID NO: 53, CDR-H3 as depicted in SEQ ID NO: 904, CDR-L1 as depicted in SEQ ID NO: 220, CDR-L2 as depicted in SEQ ID NO: 221 and CDR-L3 as depicted in SEQ ID NO: 926,

CDR-H1 as depicted in SEQ ID NO: 1126, CDR-H2 as depicted in SEQ ID NO: 1127, CDR-H3 as depicted in SEQ ID NO: 1128, CDR-L1 as depicted in SEQ ID NO: 1129, CDR-L2 as depicted in SEQ ID NO: 1130 and CDR-L3 as depicted in SEQ ID NO: 1131,

CDR-H1 as depicted in SEQ ID NO: 1165, CDR-H2 as depicted in SEQ ID NO: 1166, CDR-H3 as depicted in SEQ ID NO: 1167, CDR-L1 as depicted in SEQ ID NO: 1168, CDR-L2 as depicted in SEQ ID NO: 1169 and CDR-L3 as depicted in SEQ ID NO: 1170,

CDR-H1 as depicted in SEQ ID NO: 1334, CDR-H2 as depicted in SEQ ID NO: 1335, CDR-H3 as depicted in SEQ ID NO: 1336, CDR-L1 as depicted in SEQ ID NO: 1337, CDR-L2 as depicted in SEQ ID NO: 1338 and CDR-L3 as depicted in SEQ ID NO: 1339,

CDR-H1 as depicted in SEQ ID NO: 1347, CDR-H2 as depicted in SEQ ID NO: 1348, CDR-H3 as depicted in SEQ ID NO: 1349, CDR-L1 as depicted in SEQ ID NO: 1350, CDR-L2 as depicted in SEQ ID NO: 1351 and CDR-L3 as depicted in SEQ ID NO: 1352,

CDR-H1 as depicted in SEQ ID NO: 1360 CDR-H2 as depicted in SEQ ID NO: 1361, CDR-H3 as depicted in SEQ ID NO: 1362, CDR-L1 as depicted in SEQ ID NO: 1363, CDR-L2 as depicted in SEQ ID NO: 1364 and CDR-L3 as depicted in SEQ ID NO: 1365,

CDR-H1 as depicted in SEQ ID NO: 1425 CDR-H2 as depicted in SEQ ID NO: 1426, CDR-H3 as depicted in SEQ ID NO: 1427, CDR-L1 as depicted in SEQ ID NO: 1428, CDR-L2 as depicted in SEQ ID NO: 1429 and CDR-L3 as depicted in SEQ ID NO: 1430,

CDR-H1 as depicted in SEQ ID NO: 1438 CDR-H2 as depicted in SEQ ID NO: 1439, CDR-H3 as depicted in SEQ ID NO: 1440, CDR-L1 as depicted in SEQ ID NO: 1441, CDR-L2 as depicted in SEQ ID NO: 1442 and CDR-L3 as depicted in SEQ ID NO: 1443, and

CDR-H1 as depicted in SEQ ID NO: 2167 CDR-H2 as depicted in SEQ ID NO: 2168, CDR-H3 as depicted in SEQ ID NO: 2169, CDR-L1 as depicted in SEQ ID NO: 2170, CDR-L2 as depicted in SEQ ID NO: 2171 and CDR-L3 as depicted in SEQ ID NO: 2172;

(b) CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 126, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294,

CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 300,

CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID

NO: 137, CDR-H3 as depicted in SEQ ID NO: 138, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 306,

CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 312,

CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318,

CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336,

CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 294,

CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 928,

CDR-H1 as depicted in SEQ ID NO: 124, CDR-H2 as depicted in SEQ ID NO: 125, CDR-H3 as depicted in SEQ ID NO: 915, CDR-L1 as depicted in SEQ ID NO: 292, CDR-L2 as depicted in SEQ ID NO: 293 and CDR-L3 as depicted in SEQ ID NO: 929,

CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 336,

CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 942,

CDR-H1 as depicted in SEQ ID NO: 166, CDR-H2 as depicted in SEQ ID NO: 167, CDR-H3 as depicted in SEQ ID NO: 168, CDR-L1 as depicted in

SEQ ID NO: 334, CDR-L2 as depicted in SEQ ID NO: 335 and CDR-L3 as depicted in SEQ ID NO: 943,

CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 318,

CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 937,

CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 150, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938,

CDR-H1 as depicted in SEQ ID NO: 148, CDR-H2 as depicted in SEQ ID NO: 149, CDR-H3 as depicted in SEQ ID NO: 919, CDR-L1 as depicted in SEQ ID NO: 316, CDR-L2 as depicted in SEQ ID NO: 317 and CDR-L3 as depicted in SEQ ID NO: 938,

CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 144, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935,

CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 935,

CDR-H1 as depicted in SEQ ID NO: 142, CDR-H2 as depicted in SEQ ID NO: 143, CDR-H3 as depicted in SEQ ID NO: 918, CDR-L1 as depicted in SEQ ID NO: 310, CDR-L2 as depicted in SEQ ID NO: 311 and CDR-L3 as depicted in SEQ ID NO: 936,

CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 138, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as depicted in SEQ ID NO: 933.

CDR-H1 as depicted in SEQ ID NO: 136, CDR-H2 as depicted in SEQ ID NO: 137, CDR-H3 as depicted in SEQ ID NO: 917, CDR-L1 as depicted in SEQ ID NO: 304, CDR-L2 as depicted in SEQ ID NO: 305 and CDR-L3 as

depicted in SEQ ID NO: 934,

CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 132, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 930,

CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 931,

CDR-H1 as depicted in SEQ ID NO: 130, CDR-H2 as depicted in SEQ ID NO: 131, CDR-H3 as depicted in SEQ ID NO: 916, CDR-L1 as depicted in SEQ ID NO: 298, CDR-L2 as depicted in SEQ ID NO: 299 and CDR-L3 as depicted in SEQ ID NO: 932,

CDR-H1 as depicted in SEQ ID NO: 1009, CDR-H2 as depicted in SEQ ID NO: 1010, CDR-H3 as depicted in SEQ ID NO: 1011, CDR-L1 as depicted in SEQ ID NO: 1012, CDR-L2 as depicted in SEQ ID NO: 1013 and CDR-L3 as depicted in SEQ ID NO: 1014,

CDR-H1 as depicted in SEQ ID NO: 1022, CDR-H2 as depicted in SEQ ID NO: 1023, CDR-H3 as depicted in SEQ ID NO: 1024, CDR-L1 as depicted in SEQ ID NO: 1025, CDR-L2 as depicted in SEQ ID NO: 1026 and CDR-L3 as depicted in SEQ ID NO: 1027,

CDR-H1 as depicted in SEQ ID NO: 1035, CDR-H2 as depicted in SEQ ID NO: 1036, CDR-H3 as depicted in SEQ ID NO: 1037, CDR-L1 as depicted in SEQ ID NO: 1038, CDR-L2 as depicted in SEQ ID NO: 1039 and CDR-L3 as depicted in SEQ ID NO: 1040,

CDR-H1 as depicted in SEQ ID NO: 1074, CDR-H2 as depicted in SEQ ID NO: 1075, CDR-H3 as depicted in SEQ ID NO: 1076, CDR-L1 as depicted in SEQ ID NO: 1077, CDR-L2 as depicted in SEQ ID NO: 1078 and CDR-L3 as depicted in SEQ ID NO: 1079,

CDR-H1 as depicted in SEQ ID NO: 1100, CDR-H2 as depicted in SEQ ID NO: 1101, CDR-H3 as depicted in SEQ ID NO: 1102, CDR-L1 as depicted in SEQ ID NO: 1103, CDR-L2 as depicted in SEQ ID NO: 1104 and CDR-L3 as depicted in SEQ ID NO: 1105,

CDR-H1 as depicted in SEQ ID NO: 1113, CDR-H2 as depicted in SEQ ID NO: 1114, CDR-H3 as depicted in SEQ ID NO: 1115, CDR-L1 as depicted in SEQ ID NO: 1116, CDR-L2 as depicted in SEQ ID NO: 1117 and CDR-L3 as depicted in SEQ ID NO: 1118,

CDR-H1 as depicted in SEQ ID NO: 1243, CDR-H2 as depicted in SEQ ID NO: 1244, CDR-H3 as depicted in SEQ ID NO: 1245, CDR-L1 as depicted in SEQ ID NO: 1246, CDR-L2 as depicted in SEQ ID NO: 1247 and CDR-L3 as depicted in SEQ ID NO: 1248,

CDR-H1 as depicted in SEQ ID NO: 1256, CDR-H2 as depicted in SEQ ID NO: 1257, CDR-H3 as depicted in SEQ ID NO: 1258, CDR-L1 as depicted in SEQ ID NO: 1259, CDR-L2 as depicted in SEQ ID NO: 1260 and CDR-L3 as depicted in SEQ ID NO: 1261,

CDR-H1 as depicted in SEQ ID NO: 1269, CDR-H2 as depicted in SEQ ID NO: 1270, CDR-H3 as depicted in SEQ ID NO: 1271, CDR-L1 as depicted in SEQ ID NO: 1272, CDR-L2 as depicted in SEQ ID NO: 1273 and CDR-L3 as depicted in SEQ ID NO: 1274,

CDR-H1 as depicted in SEQ ID NO: 1282, CDR-H2 as depicted in SEQ ID NO: 1283, CDR-H3 as depicted in SEQ ID NO: 1284, CDR-L1 as depicted in SEQ ID NO: 1285, CDR-L2 as depicted in SEQ ID NO: 1286 and CDR-L3 as depicted in SEQ ID NO: 1287,

CDR-H1 as depicted in SEQ ID NO: 1295, CDR-H2 as depicted in SEQ ID NO: 1296, CDR-H3 as depicted in SEQ ID NO: 1297, CDR-L1 as depicted in SEQ ID NO: 1298, CDR-L2 as depicted in SEQ ID NO: 1299 and CDR-L3 as depicted in SEQ ID NO: 1300,

CDR-H1 as depicted in SEQ ID NO: 1647, CDR-H2 as depicted in SEQ ID NO: 1648, CDR-H3 as depicted in SEQ ID NO: 1649, CDR-L1 as depicted in SEQ ID NO: 1650, CDR-L2 as depicted in SEQ ID NO: 1651 and CDR-L3 as depicted in SEQ ID NO: 1652,

CDR-H1 as depicted in SEQ ID NO: 1660, CDR-H2 as depicted in SEQ ID NO: 1661, CDR-H3 as depicted in SEQ ID NO: 1662, CDR-L1 as depicted in SEQ ID NO: 1663, CDR-L2 as depicted in SEQ ID NO: 1664 and CDR-L3 as depicted in SEQ ID NO: 1665,

CDR-H1 as depicted in SEQ ID NO: 1894, CDR-H2 as depicted in SEQ ID NO: 1895, CDR-H3 as depicted in SEQ ID NO: 1896, CDR-L1 as depicted in SEQ ID NO: 1897, CDR-L2 as depicted in SEQ ID NO: 1898 and CDR-L3 as depicted in SEQ ID NO: 1899,

CDR-H1 as depicted in SEQ ID NO: 1907, CDR-H2 as depicted in SEQ ID NO: 1908, CDR-H3 as depicted in SEQ ID NO: 1909, CDR-L1 as depicted in SEQ ID NO: 1910, CDR-L2 as depicted in SEQ ID NO: 1911 and CDR-L3 as depicted in SEQ ID NO: 1912,

CDR-H1 as depicted in SEQ ID NO: 1933, CDR-H2 as depicted in SEQ ID

NO: 1934, CDR-H3 as depicted in SEQ ID NO: 1935, CDR-L1 as depicted in SEQ ID NO: 1936, CDR-L2 as depicted in SEQ ID NO: 1937 and CDR-L3 as depicted in SEQ ID NO: 1938,

CDR-H1 as depicted in SEQ ID NO: 1946, CDR-H2 as depicted in SEQ ID NO: 1947, CDR-H3 as depicted in SEQ ID NO: 1948, CDR-L1 as depicted in SEQ ID NO: 1949, CDR-L2 as depicted in SEQ ID NO: 1950 and CDR-L3 as depicted in SEQ ID NO: 1951,

CDR-H1 as depicted in SEQ ID NO: 1959, CDR-H2 as depicted in SEQ ID NO: 1960, CDR-H3 as depicted in SEQ ID NO: 1961, CDR-L1 as depicted in SEQ ID NO: 1962, CDR-L2 as depicted in SEQ ID NO: 1963 and CDR-L3 as depicted in SEQ ID NO: 1964,

CDR-H1 as depicted in SEQ ID NO: 1972, CDR-H2 as depicted in SEQ ID NO: 1973, CDR-H3 as depicted in SEQ ID NO: 1974, CDR-L1 as depicted in SEQ ID NO: 1975, CDR-L2 as depicted in SEQ ID NO: 1976 and CDR-L3 as depicted in SEQ ID NO: 1977,

CDR-H1 as depicted in SEQ ID NO: 1985, CDR-H2 as depicted in SEQ ID NO: 1986, CDR-H3 as depicted in SEQ ID NO: 1987, CDR-L1 as depicted in SEQ ID NO: 1988, CDR-L2 as depicted in SEQ ID NO: 1989 and CDR-L3 as depicted in SEQ ID NO: 1990,

CDR-H1 as depicted in SEQ ID NO: 1998, CDR-H2 as depicted in SEQ ID NO: 1999, CDR-H3 as depicted in SEQ ID NO: 2000, CDR-L1 as depicted in SEQ ID NO: 2001, CDR-L2 as depicted in SEQ ID NO: 2002 and CDR-L3 as depicted in SEQ ID NO: 2003,

CDR-H1 as depicted in SEQ ID NO: 2011, CDR-H2 as depicted in SEQ ID NO: 2012, CDR-H3 as depicted in SEQ ID NO: 2013, CDR-L1 as depicted in SEQ ID NO: 2014, CDR-L2 as depicted in SEQ ID NO: 2015 and CDR-L3 as depicted in SEQ ID NO: 2016,

CDR-H1 as depicted in SEQ ID NO: 2024, CDR-H2 as depicted in SEQ ID NO: 2025, CDR-H3 as depicted in SEQ ID NO: 2026, CDR-L1 as depicted in SEQ ID NO: 2027, CDR-L2 as depicted in SEQ ID NO: 2028 and CDR-L3 as depicted in SEQ ID NO: 2029,

CDR-H1 as depicted in SEQ ID NO: 2037, CDR-H2 as depicted in SEQ ID NO: 2038, CDR-H3 as depicted in SEQ ID NO: 2039, CDR-L1 as depicted in SEQ ID NO: 2040, CDR-L2 as depicted in SEQ ID NO: 2041 and CDR-L3 as depicted in SEQ ID NO: 2042, and

CDR-H1 as depicted in SEQ ID NO: 2050, CDR-H2 as depicted in SEQ ID NO: 2051, CDR-H3 as depicted in SEQ ID NO: 2052, CDR-L1 as depicted in

SEQ ID NO: 2053, CDR-L2 as depicted in SEQ ID NO: 2054 and CDR-L3 as depicted in SEQ ID NO: 2055;

(c) CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 96, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264,

CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 102, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270,

CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288,

CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 156, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 324,

CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 912, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270,

CDR-H1 as depicted in SEQ ID NO: 100, CDR-H2 as depicted in SEQ ID NO: 101, CDR-H3 as depicted in SEQ ID NO: 913, CDR-L1 as depicted in SEQ ID NO: 268, CDR-L2 as depicted in SEQ ID NO: 269 and CDR-L3 as depicted in SEQ ID NO: 270,

CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 910, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264,

CDR-H1 as depicted in SEQ ID NO: 94, CDR-H2 as depicted in SEQ ID NO: 95, CDR-H3 as depicted in SEQ ID NO: 911, CDR-L1 as depicted in SEQ ID NO: 262, CDR-L2 as depicted in SEQ ID NO: 263 and CDR-L3 as depicted in SEQ ID NO: 264,

CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 119, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as

depicted in SEQ ID NO: 288,

CDR-H1 as depicted in SEQ ID NO: 118, CDR-H2 as depicted in SEQ ID NO: 914, CDR-H3 as depicted in SEQ ID NO: 120, CDR-L1 as depicted in SEQ ID NO: 286, CDR-L2 as depicted in SEQ ID NO: 287 and CDR-L3 as depicted in SEQ ID NO: 288,

CDR-H1 as depicted in SEQ ID NO: 154, CDR-H2 as depicted in SEQ ID NO: 155, CDR-H3 as depicted in SEQ ID NO: 920, CDR-L1 as depicted in SEQ ID NO: 322, CDR-L2 as depicted in SEQ ID NO: 323 and CDR-L3 as depicted in SEQ ID NO: 324,

CDR-H1 as depicted in SEQ ID NO: 996, CDR-H2 as depicted in SEQ ID NO: 997, CDR-H3 as depicted in SEQ ID NO: 998, CDR-L1 as depicted in SEQ ID NO: 999, CDR-L2 as depicted in SEQ ID NO: 1000 and CDR-L3 as depicted in SEQ ID NO: 1001,

CDR-H1 as depicted in SEQ ID NO: 1048, CDR-H2 as depicted in SEQ ID NO: 1049, CDR-H3 as depicted in SEQ ID NO: 1050, CDR-L1 as depicted in SEQ ID NO: 1051, CDR-L2 as depicted in SEQ ID NO: 1052 and CDR-L3 as depicted in SEQ ID NO: 1053,

CDR-H1 as depicted in SEQ ID NO: 1087, CDR-H2 as depicted in SEQ ID NO: 1088, CDR-H3 as depicted in SEQ ID NO: 1089, CDR-L1 as depicted in SEQ ID NO: 1090, CDR-L2 as depicted in SEQ ID NO: 1091 and CDR-L3 as depicted in SEQ ID NO: 1092,

CDR-H1 as depicted in SEQ ID NO: 1608, CDR-H2 as depicted in SEQ ID NO: 1609, CDR-H3 as depicted in SEQ ID NO: 1610, CDR-L1 as depicted in SEQ ID NO: 1611, CDR-L2 as depicted in SEQ ID NO: 1612 and CDR-L3 as depicted in SEQ ID NO: 1613,

CDR-H1 as depicted in SEQ ID NO: 1621, CDR-H2 as depicted in SEQ ID NO: 1622, CDR-H3 as depicted in SEQ ID NO: 1623, CDR-L1 as depicted in SEQ ID NO: 1624, CDR-L2 as depicted in SEQ ID NO: 1625 and CDR-L3 as depicted in SEQ ID NO: 1626,

CDR-H1 as depicted in SEQ ID NO: 1634, CDR-H2 as depicted in SEQ ID NO: 1635, CDR-H3 as depicted in SEQ ID NO: 1636, CDR-L1 as depicted in SEQ ID NO: 1637, CDR-L2 as depicted in SEQ ID NO: 1638 and CDR-L3 as depicted in SEQ ID NO: 1639,

CDR-H1 as depicted in SEQ ID NO: 1673, CDR-H2 as depicted in SEQ ID NO: 1674, CDR-H3 as depicted in SEQ ID NO: 1675, CDR-L1 as depicted in SEQ ID NO: 1676, CDR-L2 as depicted in SEQ ID NO: 1677 and CDR-L3 as depicted in SEQ ID NO: 1678,

CDR-H1 as depicted in SEQ ID NO: 1686, CDR-H2 as depicted in SEQ ID NO: 1687, CDR-H3 as depicted in SEQ ID NO: 1688, CDR-L1 as depicted in SEQ ID NO: 1689, CDR-L2 as depicted in SEQ ID NO: 1690 and CDR-L3 as depicted in SEQ ID NO: 1691,

CDR-H1 as depicted in SEQ ID NO: 1699, CDR-H2 as depicted in SEQ ID NO: 1700, CDR-H3 as depicted in SEQ ID NO: 1701, CDR-L1 as depicted in SEQ ID NO: 1702, CDR-L2 as depicted in SEQ ID NO: 1703 and CDR-L3 as depicted in SEQ ID NO: 1704,

CDR-H1 as depicted in SEQ ID NO: 1712, CDR-H2 as depicted in SEQ ID NO: 1713, CDR-H3 as depicted in SEQ ID NO: 1714, CDR-L1 as depicted in SEQ ID NO: 1715, CDR-L2 as depicted in SEQ ID NO: 1716 and CDR-L3 as depicted in SEQ ID NO: 1717,

CDR-H1 as depicted in SEQ ID NO: 1725, CDR-H2 as depicted in SEQ ID NO: 1726, CDR-H3 as depicted in SEQ ID NO: 1727, CDR-L1 as depicted in SEQ ID NO: 1728, CDR-L2 as depicted in SEQ ID NO: 1729 and CDR-L3 as depicted in SEQ ID NO: 1730,

CDR-H1 as depicted in SEQ ID NO: 1738, CDR-H2 as depicted in SEQ ID NO: 1739, CDR-H3 as depicted in SEQ ID NO: 1740, CDR-L1 as depicted in SEQ ID NO: 1741, CDR-L2 as depicted in SEQ ID NO: 1742 and CDR-L3 as depicted in SEQ ID NO: 1743,

CDR-H1 as depicted in SEQ ID NO: 1751, CDR-H2 as depicted in SEQ ID NO: 1752, CDR-H3 as depicted in SEQ ID NO: 1753, CDR-L1 as depicted in SEQ ID NO: 1754, CDR-L2 as depicted in SEQ ID NO: 1755 and CDR-L3 as depicted in SEQ ID NO: 1756,

CDR-H1 as depicted in SEQ ID NO: 1764, CDR-H2 as depicted in SEQ ID NO: 1765, CDR-H3 as depicted in SEQ ID NO: 1766, CDR-L1 as depicted in SEQ ID NO: 1767, CDR-L2 as depicted in SEQ ID NO: 1768 and CDR-L3 as depicted in SEQ ID NO: 1769, and

CDR-H1 as depicted in SEQ ID NO: 1920, CDR-H2 as depicted in SEQ ID NO: 1921, CDR-H3 as depicted in SEQ ID NO: 1922, CDR-L1 as depicted in SEQ ID NO: 1923, CDR-L2 as depicted in SEQ ID NO: 1924 and CDR-L3 as depicted in SEQ ID NO: 1925;

(d) CDR-H1 as depicted in SEQ ID NO: 4, CDR-H2 as depicted in SEQ ID NO: 5, CDR-H3 as depicted in SEQ ID NO: 6, CDR-L1 as depicted in SEQ ID NO: 172, CDR-L2 as depicted in SEQ ID NO: 173 and CDR-L3 as depicted in SEQ ID NO: 174,

CDR-H1 as depicted in SEQ ID NO: 10, CDR-H2 as depicted in SEQ ID NO: 11,

CDR-H3 as depicted in SEQ ID NO: 12, CDR-L1 as depicted in SEQ ID NO: 178, CDR-L2 as depicted in SEQ ID NO: 179 and CDR-L3 as depicted in SEQ ID NO: 180,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 196, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 198,

CDR-H1 as depicted in SEQ ID NO: 34, CDR-H2 as depicted in SEQ ID NO: 35, CDR-H3 as depicted in SEQ ID NO: 36, CDR-L1 as depicted in SEQ ID NO: 202, CDR-L2 as depicted in SEQ ID NO: 203 and CDR-L3 as depicted in SEQ ID NO: 204,

CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 214, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216,

CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 59, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228,

CDR-H1 as depicted in SEQ ID NO: 64, CDR-H2 as depicted in SEQ ID NO: 65, CDR-H3 as depicted in SEQ ID NO: 66, CDR-L1 as depicted in SEQ ID NO: 232, CDR-L2 as depicted in SEQ ID NO: 233 and CDR-L3 as depicted in SEQ ID NO: 234,

CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 71, CDR-H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240,

CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 328, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330,

CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216,

CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 902, CDR-L1 as depicted in SEQ ID

NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216,

CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 903, CDR-L1 as depicted in SEQ ID NO: 924, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216,

CDR-H1 as depicted in SEQ ID NO: 46, CDR-H2 as depicted in SEQ ID NO: 47, CDR-H3 as depicted in SEQ ID NO: 48, CDR-L1 as depicted in SEQ ID NO: 925, CDR-L2 as depicted in SEQ ID NO: 215 and CDR-L3 as depicted in SEQ ID NO: 216,

CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907, CDR-H3 as depicted in SEQ ID NO: 72, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240,

CDR-H1 as depicted in SEQ ID NO: 70, CDR-H2 as depicted in SEQ ID NO: 907, CDR-H3 as depicted in SEQ ID NO: 908, CDR-L1 as depicted in SEQ ID NO: 238, CDR-L2 as depicted in SEQ ID NO: 239 and CDR-L3 as depicted in SEQ ID NO: 240,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 901, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923,

CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 906, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228,

CDR-H1 as depicted in SEQ ID NO: 58, CDR-H2 as depicted in SEQ ID NO: 905, CDR-H3 as depicted in SEQ ID NO: 60, CDR-L1 as depicted in SEQ ID NO: 226, CDR-L2 as depicted in SEQ ID NO: 227 and CDR-L3 as depicted in SEQ ID NO: 228,

CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330,

CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 921, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as

depicted in SEQ ID NO: 940,

CDR-H1 as depicted in SEQ ID NO: 160, CDR-H2 as depicted in SEQ ID NO: 161, CDR-H3 as depicted in SEQ ID NO: 162, CDR-L1 as depicted in SEQ ID NO: 941, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 196, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 901, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 922, CDR-L2 as depicted in SEQ ID NO: 197 and CDR-L3 as depicted in SEQ ID NO: 923,

CDR-H1 as depicted in SEQ ID NO: 28, CDR-H2 as depicted in SEQ ID NO: 29, CDR-H3 as depicted in SEQ ID NO: 30, CDR-L1 as depicted in SEQ ID NO: 939, CDR-L2 as depicted in SEQ ID NO: 329 and CDR-L3 as depicted in SEQ ID NO: 330,

CDR-H1 as depicted in SEQ ID NO: 970, CDR-H2 as depicted in SEQ ID NO: 971, CDR-H3 as depicted in SEQ ID NO: 972, CDR-L1 as depicted in SEQ ID NO: 973, CDR-L2 as depicted in SEQ ID NO: 974 and CDR-L3 as depicted in SEQ ID NO: 975,

CDR-H1 as depicted in SEQ ID NO: 1061, CDR-H2 as depicted in SEQ ID NO: 1062, CDR-H3 as depicted in SEQ ID NO: 1063, CDR-L1 as depicted in SEQ ID NO: 1064, CDR-L2 as depicted in SEQ ID NO: 1065 and CDR-L3 as depicted in SEQ ID NO: 1066,

CDR-H1 as depicted in SEQ ID NO: 1139, CDR-H2 as depicted in SEQ ID NO: 1140, CDR-H3 as depicted in SEQ ID NO: 1141, CDR-L1 as depicted in SEQ ID NO: 1142, CDR-L2 as depicted in SEQ ID NO: 1143 and CDR-L3 as depicted in SEQ ID NO: 1144,

CDR-H1 as depicted in SEQ ID NO: 1152, CDR-H2 as depicted in SEQ ID NO: 1153, CDR-H3 as depicted in SEQ ID NO: 1154, CDR-L1 as depicted in SEQ ID NO: 1155, CDR-L2 as depicted in SEQ ID NO: 1156 and CDR-L3 as depicted in SEQ ID NO: 1157,

CDR-H1 as depicted in SEQ ID NO: 1178, CDR-H2 as depicted in SEQ ID NO: 1179, CDR-H3 as depicted in SEQ ID NO: 1180, CDR-L1 as depicted in SEQ ID NO: 1181, CDR-L2 as depicted in SEQ ID NO: 1182 and CDR-L3 as depicted in SEQ ID NO: 1183,

CDR-H1 as depicted in SEQ ID NO: 1191, CDR-H2 as depicted in SEQ ID NO: 1192, CDR-H3 as depicted in SEQ ID NO: 1193, CDR-L1 as depicted in SEQ ID NO: 1194, CDR-L2 as depicted in SEQ ID NO: 1195 and CDR-L3 as depicted in SEQ ID NO: 1196,

CDR-H1 as depicted in SEQ ID NO: 1204, CDR-H2 as depicted in SEQ ID NO: 1205, CDR-H3 as depicted in SEQ ID NO: 1206, CDR-L1 as depicted in SEQ ID NO: 1207, CDR-L2 as depicted in SEQ ID NO: 1208 and CDR-L3 as depicted in SEQ ID NO: 1209,

CDR-H1 as depicted in SEQ ID NO: 1217, CDR-H2 as depicted in SEQ ID NO: 1218, CDR-H3 as depicted in SEQ ID NO: 1219, CDR-L1 as depicted in SEQ ID NO: 1220, CDR-L2 as depicted in SEQ ID NO: 1221 and CDR-L3 as depicted in SEQ ID NO: 1222,

CDR-H1 as depicted in SEQ ID NO: 1230, CDR-H2 as depicted in SEQ ID NO: 1231, CDR-H3 as depicted in SEQ ID NO: 1232, CDR-L1 as depicted in SEQ ID NO: 1233, CDR-L2 as depicted in SEQ ID NO: 1234 and CDR-L3 as depicted in SEQ ID NO: 1235,

CDR-H1 as depicted in SEQ ID NO: 1308, CDR-H2 as depicted in SEQ ID NO: 1309, CDR-H3 as depicted in SEQ ID NO: 1310, CDR-L1 as depicted in SEQ ID NO: 1311, CDR-L2 as depicted in SEQ ID NO: 1312 and CDR-L3 as depicted in SEQ ID NO: 1313,

CDR-H1 as depicted in SEQ ID NO: 1321, CDR-H2 as depicted in SEQ ID NO: 1322, CDR-H3 as depicted in SEQ ID NO: 1323, CDR-L1 as depicted in SEQ ID NO: 1324, CDR-L2 as depicted in SEQ ID NO: 1325 and CDR-L3 as depicted in SEQ ID NO: 1326,

CDR-H1 as depicted in SEQ ID NO: 1373, CDR-H2 as depicted in SEQ ID NO: 1374, CDR-H3 as depicted in SEQ ID NO: 1375, CDR-L1 as depicted in SEQ ID NO: 1376, CDR-L2 as depicted in SEQ ID NO: 1377 and CDR-L3 as depicted in SEQ ID NO: 1378,

CDR-H1 as depicted in SEQ ID NO: 1386, CDR-H2 as depicted in SEQ ID NO: 1387, CDR-H3 as depicted in SEQ ID NO: 1388, CDR-L1 as depicted in SEQ ID NO: 1389, CDR-L2 as depicted in SEQ ID NO: 1390 and CDR-L3 as depicted in SEQ ID NO: 1391,

CDR-H1 as depicted in SEQ ID NO: 1399, CDR-H2 as depicted in SEQ ID

NO: 1400, CDR-H3 as depicted in SEQ ID NO: 1401, CDR-L1 as depicted in SEQ ID NO: 1402, CDR-L2 as depicted in SEQ ID NO: 1403 and CDR-L3 as depicted in SEQ ID NO: 1404,

CDR-H1 as depicted in SEQ ID NO: 1412, CDR-H2 as depicted in SEQ ID NO: 1413, CDR-H3 as depicted in SEQ ID NO: 1414, CDR-L1 as depicted in SEQ ID NO: 1415, CDR-L2 as depicted in SEQ ID NO: 1416 and CDR-L3 as depicted in SEQ ID NO: 1417,

CDR-H1 as depicted in SEQ ID NO: 1777, CDR-H2 as depicted in SEQ ID NO: 1778, CDR-H3 as depicted in SEQ ID NO: 1779, CDR-L1 as depicted in SEQ ID NO: 1780, CDR-L2 as depicted in SEQ ID NO: 1781 and CDR-L3 as depicted in SEQ ID NO: 1782,

CDR-H1 as depicted in SEQ ID NO: 1790, CDR-H2 as depicted in SEQ ID NO: 1791, CDR-H3 as depicted in SEQ ID NO: 1792, CDR-L1 as depicted in SEQ ID NO: 1793, CDR-L2 as depicted in SEQ ID NO: 1794 and CDR-L3 as depicted in SEQ ID NO: 1795,

CDR-H1 as depicted in SEQ ID NO: 1803, CDR-H2 as depicted in SEQ ID NO: 1804, CDR-H3 as depicted in SEQ ID NO: 1805, CDR-L1 as depicted in SEQ ID NO: 1806, CDR-L2 as depicted in SEQ ID NO: 1807 and CDR-L3 as depicted in SEQ ID NO: 1808,

CDR-H1 as depicted in SEQ ID NO: 1816, CDR-H2 as depicted in SEQ ID NO: 1817, CDR-H3 as depicted in SEQ ID NO: 1818, CDR-L1 as depicted in SEQ ID NO: 1819, CDR-L2 as depicted in SEQ ID NO: 1820 and CDR-L3 as depicted in SEQ ID NO: 1821,

CDR-H1 as depicted in SEQ ID NO: 1829, CDR-H2 as depicted in SEQ ID NO: 1830, CDR-H3 as depicted in SEQ ID NO: 1831, CDR-L1 as depicted in SEQ ID NO: 1832, CDR-L2 as depicted in SEQ ID NO: 1833 and CDR-L3 as depicted in SEQ ID NO: 1834,

CDR-H1 as depicted in SEQ ID NO: 1842, CDR-H2 as depicted in SEQ ID NO: 1843, CDR-H3 as depicted in SEQ ID NO: 1844, CDR-L1 as depicted in SEQ ID NO: 1845, CDR-L2 as depicted in SEQ ID NO: 1846 and CDR-L3 as depicted in SEQ ID NO: 1847,

CDR-H1 as depicted in SEQ ID NO: 1855, CDR-H2 as depicted in SEQ ID NO: 1856, CDR-H3 as depicted in SEQ ID NO: 1857, CDR-L1 as depicted in SEQ ID NO: 1858, CDR-L2 as depicted in SEQ ID NO: 1859 and CDR-L3 as depicted in SEQ ID NO: 1860,

CDR-H1 as depicted in SEQ ID NO: 1868, CDR-H2 as depicted in SEQ ID NO: 1869, CDR-H3 as depicted in SEQ ID NO: 1870, CDR-L1 as depicted in

SEQ ID NO: 1871, CDR-L2 as depicted in SEQ ID NO: 1872 and CDR-L3 as depicted in SEQ ID NO: 1873,

CDR-H1 as depicted in SEQ ID NO: 1881, CDR-H2 as depicted in SEQ ID NO: 1882, CDR-H3 as depicted in SEQ ID NO: 1883, CDR-L1 as depicted in SEQ ID NO: 1884, CDR-L2 as depicted in SEQ ID NO: 1885 and CDR-L3 as depicted in SEQ ID NO: 1886,

CDR-H1 as depicted in SEQ ID NO: 2063, CDR-H2 as depicted in SEQ ID NO: 2064, CDR-H3 as depicted in SEQ ID NO: 2065, CDR-L1 as depicted in SEQ ID NO: 2066, CDR-L2 as depicted in SEQ ID NO: 2067 and CDR-L3 as depicted in SEQ ID NO: 2068,

CDR-H1 as depicted in SEQ ID NO: 2076, CDR-H2 as depicted in SEQ ID NO: 2077, CDR-H3 as depicted in SEQ ID NO: 2078, CDR-L1 as depicted in SEQ ID NO: 2079, CDR-L2 as depicted in SEQ ID NO: 2080 and CDR-L3 as depicted in SEQ ID NO: 2081,

CDR-H1 as depicted in SEQ ID NO: 2089, CDR-H2 as depicted in SEQ ID NO: 2090, CDR-H3 as depicted in SEQ ID NO: 2091, CDR-L1 as depicted in SEQ ID NO: 2092, CDR-L2 as depicted in SEQ ID NO: 2093 and CDR-L3 as depicted in SEQ ID NO: 2094,

CDR-H1 as depicted in SEQ ID NO: 2102, CDR-H2 as depicted in SEQ ID NO: 2103, CDR-H3 as depicted in SEQ ID NO: 2104, CDR-L1 as depicted in SEQ ID NO: 2105, CDR-L2 as depicted in SEQ ID NO: 2106 and CDR-L3 as depicted in SEQ ID NO: 2107,

CDR-H1 as depicted in SEQ ID NO: 2115, CDR-H2 as depicted in SEQ ID NO: 2116, CDR-H3 as depicted in SEQ ID NO: 2117, CDR-L1 as depicted in SEQ ID NO: 2118, CDR-L2 as depicted in SEQ ID NO: 2119 and CDR-L3 as depicted in SEQ ID NO: 2120,

CDR-H1 as depicted in SEQ ID NO: 2128, CDR-H2 as depicted in SEQ ID NO: 2129, CDR-H3 as depicted in SEQ ID NO: 2130, CDR-L1 as depicted in SEQ ID NO: 2131, CDR-L2 as depicted in SEQ ID NO: 2132 and CDR-L3 as depicted in SEQ ID NO: 2133,

CDR-H1 as depicted in SEQ ID NO: 2141, CDR-H2 as depicted in SEQ ID NO: 2142, CDR-H3 as depicted in SEQ ID NO: 2143, CDR-L1 as depicted in SEQ ID NO: 2144, CDR-L2 as depicted in SEQ ID NO: 2145 and CDR-L3 as depicted in SEQ ID NO: 2146,

CDR-H1 as depicted in SEQ ID NO: 2154, CDR-H2 as depicted in SEQ ID NO: 2155, CDR-H3 as depicted in SEQ ID NO: 2156, CDR-L1 as depicted in SEQ ID NO: 2157, CDR-L2 as depicted in SEQ ID NO: 2158 and CDR-L3 as

depicted in SEQ ID NO: 2159,

CDR-H1 as depicted in SEQ ID NO: 2180, CDR-H2 as depicted in SEQ ID NO: 2181, CDR-H3 as depicted in SEQ ID NO: 2182, CDR-L1 as depicted in SEQ ID NO: 2183, CDR-L2 as depicted in SEQ ID NO: 2184 and CDR-L3 as depicted in SEQ ID NO: 2185,

CDR-H1 as depicted in SEQ ID NO: 2193, CDR-H2 as depicted in SEQ ID NO: 2194, CDR-H3 as depicted in SEQ ID NO: 2195, CDR-L1 as depicted in SEQ ID NO: 2196, CDR-L2 as depicted in SEQ ID NO: 2197 and CDR-L3 as depicted in SEQ ID NO: 2198, and

CDR-H1 as depicted in SEQ ID NO: 2206, CDR-H2 as depicted in SEQ ID NO: 2207, CDR-H3 as depicted in SEQ ID NO: 2208, CDR-L1 as depicted in SEQ ID NO: 2209, CDR-L2 as depicted in SEQ ID NO: 2210 and CDR-L3 as depicted in SEQ ID NO: 2211; and

(e) CDR-H1 as depicted in SEQ ID NO: 76, CDR-H2 as depicted in SEQ ID NO: 77, CDR-H3 as depicted in SEQ ID NO: 78, CDR-L1 as depicted in SEQ ID NO: 244, CDR-L2 as depicted in SEQ ID NO: 245 and CDR-L3 as depicted in SEQ ID NO: 246,

CDR-H1 as depicted in SEQ ID NO: 88, CDR-H2 as depicted in SEQ ID NO: 89, CDR-H3 as depicted in SEQ ID NO: 90, CDR-L1 as depicted in SEQ ID NO: 256, CDR-L2 as depicted in SEQ ID NO: 257 and CDR-L3 as depicted in SEQ ID NO: 258,

CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274, CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276

CDR-H1 as depicted in SEQ ID NO: 112, CDR-H2 as depicted in SEQ ID NO: 113, CDR-H3 as depicted in SEQ ID NO: 114, CDR-L1 as depicted in SEQ ID NO: 280, CDR-L2 as depicted in SEQ ID NO: 281 and CDR-L3 as depicted in SEQ ID NO: 282,

CDR-H1 as depicted in SEQ ID NO: 106, CDR-H2 as depicted in SEQ ID NO: 107, CDR-H3 as depicted in SEQ ID NO: 108, CDR-L1 as depicted in SEQ ID NO: 274, CDR-L2 as depicted in SEQ ID NO: 275 and CDR-L3 as depicted in SEQ ID NO: 276,

CDR-H1 as depicted in SEQ ID NO: 983, CDR-H2 as depicted in SEQ ID NO: 984, CDR-H3 as depicted in SEQ ID NO: 985, CDR-L1 as depicted in SEQ ID NO: 986, CDR-L2 as depicted in SEQ ID NO: 987 and CDR-L3 as depicted in SEQ ID NO: 988,

CDR-H1 as depicted in SEQ ID NO: 1582, CDR-H2 as depicted in SEQ ID NO: 1583, CDR-H3 as depicted in SEQ ID NO: 1584, CDR-L1 as depicted in SEQ ID NO: 1585, CDR-L2 as depicted in SEQ ID NO: 1586 and CDR-L3 as depicted in SEQ ID NO: 1587, and

CDR-H1 as depicted in SEQ ID NO: 1595, CDR-H2 as depicted in SEQ ID NO: 1596, CDR-H3 as depicted in SEQ ID NO: 1597, CDR-L1 as depicted in SEQ ID NO: 1598, CDR-L2 as depicted in SEQ ID NO: 1599 and CDR-L3 as depicted in SEQ ID NO: 1600.

- 3. The antibody construct according to claim 1 or 2, wherein the first binding domain comprises a VH region selected from the group consisting of VH regions
 - (a) as depicted in SEQ ID NO: 362, SEQ ID NO: 364, SEQ ID NO: 485, SEQ ID NO: 486, SEQ ID NO: 487, SEQ ID NO: 492, SEQ ID NO: 493, SEQ ID NO: 494, SEQ ID NO: 495, SEQ ID NO: 1133, SEQ ID NO: 1172, SEQ ID NO: 1341, SEQ ID NO: 1354, SEQ ID NO: 1367, SEQ ID NO: 1432, SEQ ID NO: 1445 and SEQ ID NO: 2174;
 - (b) as depicted in SEQ ID NO: 342, SEQ ID NO: 366, SEQ ID NO: 370, SEQ ID NO: 344, SEQ ID NO: 372, SEQ ID NO: 368, SEQ ID NO: 496, SEQ ID NO: 497, SEQ ID NO: 498, SEQ ID NO: 499, SEQ ID NO: 500, SEQ ID NO: 508, SEQ ID NO: 509, SEQ ID NO: 510, SEQ ID NO: 511, SEQ ID NO: 512, SEQ ID NO: 519, SEQ ID NO: 520, SEQ ID NO: 521, SEQ ID NO: 522, SEQ ID NO: 523, SEQ ID NO: 524, SEQ ID NO: 525, SEQ ID NO: 526, SEQ ID NO: 527, SEQ ID NO: 528, SEQ ID NO: 529, SEQ ID NO: 530, SEQ ID NO: 531, SEQ ID NO: 532, SEQ ID NO: 533, SEQ ID NO: 534, SEQ ID NO: 535, SEQ ID NO: 536, SEQ ID NO: 537, SEQ ID NO: 538, SEQ ID NO: 1016, SEQ ID NO: 1029, SEQ ID NO: 1042, SEQ ID NO: 1081, SEQ ID NO: 1107, SEQ ID NO: 1120, SEQ ID NO: 1250, SEQ ID NO: 1263, SEQ ID NO: 1276, SEQ ID NO: 1289, SEQ ID NO: 1302, SEQ ID NO: 1654, SEQ ID NO: 1667, SEQ ID NO: 1901, SEQ ID NO: 1914, SEQ ID NO: 1940, SEQ ID NO: 1953, SEQ ID NO: 1966, SEQ ID NO: 1979, SEQ ID NO: 1992, SEQ ID NO: 2005, SEQ ID NO: 2018, SEQ ID NO: 2031, SEQ ID NO: 2044, and SEQ ID NO: 2057;
 - (c) as depicted in SEQ ID NO: 338, SEQ ID NO: 354, SEQ ID NO: 378, SEQ ID NO: 356, SEQ ID NO: 476, SEQ ID NO: 477, SEQ ID NO: 478, SEQ ID NO: 479, SEQ ID NO: 480, SEQ ID NO: 481, SEQ ID NO: 482, SEQ ID NO: 483, SEQ ID NO: 484, SEQ ID NO: 501, SEQ ID NO: 502, SEQ ID NO: 503, SEQ ID NO: 504, SEQ ID NO: 505, SEQ ID NO: 506, SEQ ID NO: 517, SEQ ID NO: 518, SEQ ID NO: 1003, SEQ ID NO: 1055, SEQ ID NO: 1094, SEQ ID NO: 1615, SEQ ID

NO: 1628, SEQ ID NO: 1641, SEQ ID NO: 1680, SEQ ID NO: 1693, SEQ ID NO: 1706, SEQ ID NO: 1719, SEQ ID NO: 1732, SEQ ID NO: 1745, SEQ ID NO: 1758, SEQ ID NO: 1771, and SEQ ID NO: 1927;

- (d) as depicted in SEQ ID NO: 352, SEQ ID NO: 360, SEQ ID NO: 388, SEQ ID NO: 386, SEQ ID NO: 340, SEQ ID NO: 346, SEQ ID NO: 374, SEQ ID NO: 348, SEQ ID NO: 390, SEQ ID NO: 463, SEQ ID NO: 464, SEQ ID NO: 465, SEQ ID NO: 466, SEQ ID NO: 467, SEQ ID NO: 468, SEQ ID NO: 469, SEQ ID NO: 470, SEQ ID NO: 471, SEQ ID NO: 472, SEQ ID NO: 473, SEQ ID NO: 474, SEQ ID NO: 475, SEQ ID NO: 488, SEQ ID NO: 489, SEQ ID NO: 490, SEQ ID NO: 491, SEQ ID NO: 513, SEQ ID NO: 514, SEQ ID NO: 515, SEQ ID NO: 516, SEQ ID NO: 540, SEQ ID NO: 541, SEQ ID NO: 542, SEQ ID NO: 543, SEQ ID NO: 977, SEQ ID NO: 1068, SEQ ID NO: 1146, SEQ ID NO: 1159, SEQ ID NO: 1185, SEQ ID NO: 1198, SEQ ID NO: 1211, SEQ ID NO: 1224, SEQ ID NO: 1237, SEQ ID NO: 1315, SEQ ID NO: 1328, SEQ ID NO: 1380, SEQ ID NO: 1393, SEQ ID NO: 1406, SEQ ID NO: 1419, SEQ ID NO: 1469, SEQ ID NO: 1478, SEQ ID NO: 1485, SEQ ID NO: 1494, SEQ ID NO: 1501, SEQ ID NO: 1508, SEQ ID NO: 1519, SEQ ID NO: 1526, SEQ ID NO: 1533, SEQ ID NO: 1542, SEQ ID NO: 1549, SEQ ID NO: 1558, SEQ ID NO: 1565, SEQ ID NO: 1784, SEQ ID NO: 1797, SEQ ID NO: 1810, SEQ ID NO: 1823, SEQ ID NO: 1836, SEQ ID NO: 1849, SEQ ID NO: 1862, SEQ ID NO: 1875, SEQ ID NO: 1888, SEQ ID NO: 2070, SEQ ID NO: 2083, SEQ ID NO: 2096, SEQ ID NO: 2109, SEQ ID NO: 2122. SEQ ID NO: 2135. SEQ ID NO: 2148. SEQ ID NO: 2161. SEQ ID NO: 2187, SEQ ID NO: 2200, and SEQ ID NO: 2213; and
- (e) as depicted in SEQ ID NO: 376, SEQ ID NO: 392, SEQ ID NO: 358, SEQ ID NO: 350, SEQ ID NO: 507, SEQ ID NO: 990, SEQ ID NO: 1589, and SEQ ID NO: 1602.
- 4. The antibody construct according to any one of the preceding claims, wherein the first binding domain comprises a VL region selected from the group consisting of VL regions
 - (a) as depicted in SEQ ID NO: 418, SEQ ID NO: 420, SEQ ID NO: 580, SEQ ID NO: 581, SEQ ID NO: 582, SEQ ID NO: 587, SEQ ID NO: 588, SEQ ID NO: 589, SEQ ID NO: 590, SEQ ID NO: 1135, SEQ ID NO: 1174, SEQ ID NO: 1343, SEQ ID NO: 1356, SEQ ID NO: 1369, SEQ ID NO: 1434, SEQ ID NO: 1447 and SEQ ID NO: 2176;
 - (b) as depicted in SEQ ID NO: 398, SEQ ID NO: 422, SEQ ID NO: 426, SEQ ID NO: 400, SEQ ID NO: 428, SEQ ID NO: 424, SEQ ID NO: 591, SEQ ID NO: 592,

SEQ ID NO: 593, SEQ ID NO: 594, SEQ ID NO: 595, SEQ ID NO: 603, SEQ ID NO: 604, SEQ ID NO: 605, SEQ ID NO: 606, SEQ ID NO: 607, SEQ ID NO: 614, SEQ ID NO: 615, SEQ ID NO: 616, SEQ ID NO: 617, SEQ ID NO: 618, SEQ ID NO: 619, SEQ ID NO: 620, SEQ ID NO: 621, SEQ ID NO: 622, SEQ ID NO: 623, SEQ ID NO: 624, SEQ ID NO: 625, SEQ ID NO: 626, SEQ ID NO: 627, SEQ ID NO: 628, SEQ ID NO: 629, SEQ ID NO: 630, SEQ ID NO: 631, SEQ ID NO: 632, SEQ ID NO: 633, SEQ ID NO: 1018, SEQ ID NO: 1031, SEQ ID NO: 1044, SEQ ID NO: 1083, SEQ ID NO: 1109, SEQ ID NO: 1122, SEQ ID NO: 1252, SEQ ID NO: 1265, SEQ ID NO: 1278, SEQ ID NO: 1291, SEQ ID NO: 1304, SEQ ID NO: 1656, SEQ ID NO: 1669, SEQ ID NO: 1903, SEQ ID NO: 1916, SEQ ID NO: 1942, SEQ ID NO: 1955, SEQ ID NO: 1968, SEQ ID NO: 1981, SEQ ID NO: 1994, SEQ ID NO: 2007, SEQ ID NO: 2020, SEQ ID NO: 2033, SEQ ID NO: 2046, and SEQ ID NO: 2059;

- (c) as depicted in SEQ ID NO: 394, SEQ ID NO: 410, SEQ ID NO: 434, SEQ ID NO: 412, SEQ ID NO: 571, SEQ ID NO: 572, SEQ ID NO: 573, SEQ ID NO: 574, SEQ ID NO: 575, SEQ ID NO: 576, SEQ ID NO: 577, SEQ ID NO: 578, SEQ ID NO: 579, SEQ ID NO: 596, SEQ ID NO: 597, SEQ ID NO: 598, SEQ ID NO: 599, SEQ ID NO: 600, SEQ ID NO: 601, SEQ ID NO: 612, SEQ ID NO: 613, SEQ ID NO: 1005, SEQ ID NO: 1057, SEQ ID NO: 1096, SEQ ID NO: 1617, SEQ ID NO: 1630, SEQ ID NO: 1643, SEQ ID NO: 1682, SEQ ID NO: 1695, SEQ ID NO: 1708, SEQ ID NO: 1721, SEQ ID NO: 1734, SEQ ID NO: 1747, SEQ ID NO: 1760, SEQ ID NO: 1773, and SEQ ID NO: 1929;
- (d) as depicted in SEQ ID NO: 408, SEQ ID NO: 416, SEQ ID NO: 444, SEQ ID NO: 442, SEQ ID NO: 396, SEQ ID NO: 402, SEQ ID NO: 430, SEQ ID NO: 404, SEQ ID NO: 446, SEQ ID NO: 558, SEQ ID NO: 559, SEQ ID NO: 560, SEQ ID NO: 561, SEQ ID NO: 562, SEQ ID NO: 563, SEQ ID NO: 564, SEQ ID NO: 565, SEQ ID NO: 566, SEQ ID NO: 567, SEQ ID NO: 568, SEQ ID NO: 569, SEQ ID NO: 570, SEQ ID NO: 583, SEQ ID NO: 584, SEQ ID NO: 585, SEQ ID NO: 586, SEQ ID NO: 608, SEQ ID NO: 609, SEQ ID NO: 610, SEQ ID NO: 611, SEQ ID NO: 635, SEQ ID NO: 636, SEQ ID NO: 637, SEQ ID NO: 638, SEQ ID NO: 979, SEQ ID NO: 1070, SEQ ID NO: 1148, SEQ ID NO: 1161, SEQ ID NO: 1187, SEQ ID NO: 1200, SEQ ID NO: 1213, SEQ ID NO: 1226, SEQ ID NO: 1239, SEQ ID NO: 1317, SEQ ID NO: 1330, SEQ ID NO: 1382, SEQ ID NO: 1395, SEQ ID NO: 1408, SEQ ID NO: 1421, SEQ ID NO: 1471, SEQ ID NO: 1480, SEQ ID NO: 1487, SEQ ID NO: 1487, SEQ ID NO: 1496, SEQ ID NO: 1503, SEQ ID NO: 1510, SEQ ID NO: 1521, SEQ ID NO: 1528, SEQ ID NO: 1535, SEQ ID NO: 1544, SEQ ID NO: 1551, SEQ ID NO: 1560, SEQ ID NO: 1567, SEQ ID NO: 1786,

SEQ ID NO: 1799, SEQ ID NO: 1812, SEQ ID NO: 1825, SEQ ID NO: 1838, SEQ ID NO: 1851, SEQ ID NO: 1864, SEQ ID NO: 1877, SEQ ID NO: 1890, SEQ ID NO: 2072, SEQ ID NO: 2085, SEQ ID NO: 2098, SEQ ID NO: 2111, SEQ ID NO: 2124, SEQ ID NO: 2137, SEQ ID NO: 2150, SEQ ID NO: 2163, SEQ ID NO: 2189, SEQ ID NO: 2202, and SEQ ID NO: 2215; and

- (e) as depicted in SEQ ID NO: 432, SEQ ID NO: 448, SEQ ID NO: 414, SEQ ID NO: 406, SEQ ID NO: 602, SEQ ID NO: 992, SEQ ID NO: 1591, and SEQ ID NO: 1604.
- 5. The antibody construct according to any one of the preceding claims, wherein the first binding domain comprises a VH region and a VL region selected from the group consisting of:
 - (1) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 362+418, SEQ ID NOs: 364+420, SEQ ID NOs: 485+580, SEQ ID NOs: 486+581, SEQ ID NOs: 487+582, SEQ ID NOs: 492+587, SEQ ID NOs: 493+588, SEQ ID NOs: 494+589, SEQ ID NOs: 495+590, SEQ ID NOs: 1133+1135, SEQ ID NOs: 1172+1174, SEQ ID NOs: 1341+1343, SEQ ID NOs: 1354+1356, SEQ ID NOs: 1367+1369, SEQ ID NOs: 1432+1434, SEQ ID NOs: 1445+1447, and SEQ ID NOs: 2174+2176;
 - (2) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 342+398, SEQ ID NOs: 366+422, SEQ ID NOs: 370+426, SEQ ID NOs: 344+400, SEQ ID NOs: 372+428, SEQ ID NOs: 368+424, SEQ ID NOs: 496+591, SEQ ID NOs: 497+592, SEQ ID NOs: 498+593, SEQ ID NOs: 499+594, SEQ ID NOs: 500+595, SEQ ID NOs: 508+603, SEQ ID NOs: 509+604, SEQ ID NOs: 510+605, SEQ ID NOs: 511+606, SEQ ID NOs: 512+607, SEQ ID NOs: 519+614, SEQ ID NOs: 520+615, SEQ ID NOs: 521+616, SEQ ID NOs: 522+617, SEQ ID NOs: 523+618, SEQ ID NOs: 524+619, SEQ ID NOs: 525+620, SEQ ID NOs: 526+621, SEQ ID NOs: 527+622, SEQ ID NOs: 528+623, SEQ ID NOs: 529+624, SEQ ID NOs: 530+625, SEQ ID NOs: 531+626, SEQ ID NOs: 532+627, SEQ ID NOs: 533+628, SEQ ID NOs: 534+629, SEQ ID NOs: 535+630, SEQ ID NOs: 536+631, SEQ ID NOs: 537+632, SEQ ID NOs: 538+633, SEQ ID NOs: 1016+1018, SEQ ID NOs: 1029+1031, SEQ ID NOs: 1042+1044, SEQ ID NOs: 1081+1083, SEQ ID NOs: 1107+1109, SEQ ID NOs: 1120+1122, SEQ ID NOs: 1250+1252, SEQ ID NOs: 1263+1265, SEQ ID NOs: 1276+1278, SEQ ID NOs: 1289+1291, SEQ ID NOs: 1302+1304, SEQ ID NOs: 1654+1656, SEQ ID NOs: 1667+1669, SEQ ID NOs: 1901+1903, SEQ ID NOs: 1914+1916, SEQ ID NOs: 1940+1942, SEQ ID

NOs: 1953+1955, SEQ ID NOs: 1966+1968, SEQ ID NOs: 1979+1981, SEQ ID NOs: 1992+1994, SEQ ID NOs: 2005+2007, SEQ ID NOs: 2018+2020, SEQ ID NOs: 2031+2033, SEQ ID NOs: 2044+2046, and SEQ ID NOs: 2057+2059;

- (3) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 338+394, SEQ ID NOs: 354+410, SEQ ID NOs: 378+434, SEQ ID NOs: 356+412, SEQ ID NOs: 476+571, SEQ ID NOs: 477+572, SEQ ID NOs: 478+573, SEQ ID NOs: 479+574, SEQ ID NOs: 480+575, SEQ ID NOs: 481+576, SEQ ID NOs: 482+577, SEQ ID NOs: 483+578, SEQ ID NOs: 484+579, SEQ ID NOs: 501+596, SEQ ID NOs: 502+597, SEQ ID NOs: 503+598, SEQ ID NOs: 504+599, SEQ ID NOs: 505+600, SEQ ID NOs: 506+601, SEQ ID NOs: 517+612, SEQ ID NOs: 518+613, SEQ ID NOs: 1003+1005, SEQ ID NOs: 1055+1057, SEQ ID NOs: 1094+1096, SEQ ID NOs: 1615+1617, SEQ ID NOs: 1628+1630, SEQ ID NOs: 1641+1643, SEQ ID NOs: 1680+1682, SEQ ID NOs: 1732+1734, SEQ ID NOs: 1745+1747, SEQ ID NOs: 1758+1760, SEQ ID NOs: 1771+1773, and SEQ ID NOs: 1927+1929;
- (4) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 352+408, SEQ ID NOs: 360+416, SEQ ID NOs: 388+444, SEQ ID NOs: 386+442, SEQ ID NOs: 340+396, SEQ ID NOs: 346+402, SEQ ID NOs: 374+430, SEQ ID NOs: 348+404, SEQ ID NOs: 390+446, SEQ ID NOs: 463+558, SEQ ID NOs: 464+559, SEQ ID NOs: 465+560, SEQ ID NOs: 466+561, SEQ ID NOs: 467+562, SEQ ID NOs: 468+563, SEQ ID NOs: 469+564, SEQ ID NOs: 470+565, SEQ ID NOs: 471+566, SEQ ID NOs: 472+567, SEQ ID NOs: 473+568, SEQ ID NOs: 474+569, SEQ ID NOs: 475+570, SEQ ID NOs: 488+583, SEQ ID NOs: 489+584, SEQ ID NOs: 490+585, SEQ ID NOs: 491+586, SEQ ID NOs: 513+608, SEQ ID NOs: 514+609, SEQ ID NOs: 515+610, SEQ ID NOs: 516+611, SEQ ID NOs: 540+635, SEQ ID NOs: 541+636, SEQ ID NOs: 542+637, SEQ ID NOs: 543+638, SEQ ID NOs: 977+979, SEQ ID NOs: 1068+1070, SEQ ID NOs: 1146+1148, SEQ ID NOs: 1159+1161, SEQ ID NOs: 1185+1187, SEQ ID NOs: 1198+1200, SEQ ID NOs: 1211+1213, SEQ ID NOs: 1224+1226, SEQ ID NOs: 1237+1239, SEQ ID NOs: 1315+1317, SEQ ID NOs: 1328+1330, SEQ ID NOs: 1380+1382 SEQ ID NOs: 1393+1395, SEQ ID NOs: 1406+1408, SEQ ID NOs: 1419+1421, SEQ ID NOs: 1469+1471, SEQ ID NOs: 1478+1480, SEQ ID NOs: 1485+1487, SEQ ID NOs: 1494+1496, SEQ ID NOs: 1501+1503, SEQ ID NOs: 1508+1510, SEQ ID NOs: 1519+1521, SEQ ID NOs: 1526+1528, SEQ ID NOs: 1533+1535, SEQ ID NOs: 1542+1544, SEQ ID NOs: 1549+1551, SEQ ID NOs: 1558+1560, SEQ ID

NOs: 1565+1567, SEQ ID NOs: 1784+1786, SEQ ID NOs: 1797+1799, SEQ ID NOs: 1810+1812, SEQ ID NOs: 1823+1825, SEQ ID NOs: 1836+1838, SEQ ID NOs: 1849+1851, SEQ ID NOs: 1862+1864, SEQ ID NOs: 1875+1877, SEQ ID NOs: 1888+1890, SEQ ID NOs: 2070+2072, SEQ ID NOs: 2083+2085, SEQ ID NOs: 2096+2098, SEQ ID NOs: 2109+2111, SEQ ID NOs: 2122+2124, SEQ ID NOs: 2135+2137, SEQ ID NOs: 2148+2150, SEQ ID NOs: 2161+2163, SEQ ID NOs: 2187+2189, SEQ ID NOs: 2200+2202, and SEQ ID NOs: 2213+2215; and

- (5) pairs of a VH region and a VL region as depicted in SEQ ID NOs: 376+432, SEQ ID NOs: 392+448, SEQ ID NOs: 358+414, SEQ ID NOs: 350+406, SEQ ID NOs: 507+602, SEQ ID NOs: 990+992, SEQ ID NOs: 1589+1591, and SEQ ID NOs: 1602+1604.
- 6. The antibody construct according to any one of the preceding claims, wherein the antibody construct is in a format selected from the group consisting of (scFv)₂, (single domain mAb)₂, scFv-single domain mAb, diabodies and oligomers thereof.
- 7. The antibody construct according to claim 6, wherein the first binding domain comprises an amino acid sequence selected from the group consisting of
 - (a) as depicted in SEQ ID NO: 117, SEQ ID NO: 1137, SEQ ID NO: 1176, SEQ ID NO: 1345, SEQ ID NO: 1358, SEQ ID NO: 1371, SEQ ID NO: 1436, SEQ ID NO: 1449 and SEQ ID NO: 2178;
 - (b) as depicted in SEQ ID NO: 1020, SEQ ID NO: 1033, SEQ ID NO: 1046, SEQ ID NO: 1085, SEQ ID NO: 1111, SEQ ID NO: 1124, SEQ ID NO: 1254, SEQ ID NO: 1267, SEQ ID NO: 1280, SEQ ID NO: 1293, SEQ ID NO: 1306, SEQ ID NO: 1658, SEQ ID NO: 1671, SEQ ID NO: 1905, SEQ ID NO: 1918, SEQ ID NO: 1944, SEQ ID NO: 1957, SEQ ID NO: 1970, SEQ ID NO: 1983, SEQ ID NO: 1996, SEQ ID NO: 2009, SEQ ID NO: 2022, SEQ ID NO: 2035, SEQ ID NO: 2048, and SEQ ID NO: 2061;
 - (c) as depicted in SEQ ID NO: 1007, SEQ ID NO: 1059, SEQ ID NO: 1098, SEQ ID NO: 1619, SEQ ID NO: 1632, SEQ ID NO: 1645, SEQ ID NO: 1684, SEQ ID NO: 1697, SEQ ID NO: 1710, SEQ ID NO: 1723, SEQ ID NO: 1736, SEQ ID NO: 1749, SEQ ID NO: 1762, SEQ ID NO: 1775, and SEQ ID NO: 1931;
 - (d) as depicted in SEQ ID NO: 981, SEQ ID NO: 1072, SEQ ID NO: 1150, SEQ ID NO: 1163, SEQ ID NO: 1189, SEQ ID NO: 1202, SEQ ID NO: 1215, SEQ ID NO: 1228, SEQ ID NO: 1241, SEQ ID NO: 1319, SEQ ID NO: 1332, SEQ ID NO: 1384, SEQ ID NO: 1397, SEQ ID NO: 1410, SEQ ID NO: 1423, SEQ ID NO: 1473, SEQ ID NO: 1482, SEQ ID NO: 1489, SEQ ID NO: 1498, SEQ ID

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NO: 1505, SEQ ID NO: 1512, SEQ ID NO: 1523, SEQ ID NO: 1530, SEQ ID NO: 1537, SEQ ID NO: 1546, SEQ ID NO: 1553, SEQ ID NO: 1562, SEQ ID NO: 1569, SEQ ID NO: 1788, SEQ ID NO: 1801, SEQ ID NO: 1814, SEQ ID NO: 1827, SEQ ID NO: 1840, SEQ ID NO: 1853, SEQ ID NO: 1866, SEQ ID NO: 1879, SEQ ID NO: 1892, SEQ ID NO: 2074, SEQ ID NO: 2087, SEQ ID NO: 2100, SEQ ID NO: 2113, SEQ ID NO: 2126, SEQ ID NO: 2139, SEQ ID NO: 2152, SEQ ID NO: 2165, SEQ ID NO: 2191, SEQ ID NO: 2204, and SEQ ID NO: 2217; and
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- (e) as depicted in SEQ ID NO: 994, SEQ ID NO: 1593, and SEQ ID NO: 1606.
- 8. The antibody construct according to any one of the preceding claims, wherein the second binding domain is capable of binding to human and Callithrix jacchus, Saguinus Oedipus or Saimiri sciureus CD3 epsilon.
- 9. The antibody construct according claim 8, having the amino acid sequence selected from the group consisting of
 - (a) as depicted in SEQ ID NO: 1138, SEQ ID NO: 1177, SEQ ID NO: 1346, SEQ ID NO: 1359, SEQ ID NO: 1372, SEQ ID NO: 1437, SEQ ID NO: 1450 and SEQ ID NO: 2179;
 - (b) as depicted in SEQ ID NO: 1021, SEQ ID NO: 1034, SEQ ID NO: 1047, SEQ ID NO: 1086, SEQ ID NO: 1112, SEQ ID NO: 1125, SEQ ID NO: 1255, SEQ ID NO: 1268, SEQ ID NO: 1281, SEQ ID NO: 1294, SEQ ID NO: 1307, SEQ ID NO: 1659, SEQ ID NO: 1672, SEQ ID NO: 1906, SEQ ID NO: 1919, SEQ ID NO: 1945, SEQ ID NO: 1958, SEQ ID NO: 1971, SEQ ID NO: 1984, SEQ ID NO: 1997, SEQ ID NO: 2010, SEQ ID NO: 2023, SEQ ID NO: 2036, SEQ ID NO: 2049, and SEQ ID NO: 2062;
 - (c) as depicted in SEQ ID NO: 1008, SEQ ID NO: 1060, SEQ ID NO: 1099, SEQ ID NO: 1620, SEQ ID NO: 1633, SEQ ID NO: 1646, SEQ ID NO: 1685, SEQ ID NO: 1698, SEQ ID NO: 1711, SEQ ID NO: 1724, SEQ ID NO: 1737, SEQ ID NO: 1750, SEQ ID NO: 1763, SEQ ID NO: 1776, and SEQ ID NO: 1932;
 - (d) as depicted in SEQ ID NO: 982, SEQ ID NO: 1073, SEQ ID NO: 1151, SEQ ID NO: 1164, SEQ ID NO: 1190, SEQ ID NO: 1203, SEQ ID NO: 1216, SEQ ID NO: 1229, SEQ ID NO: 1242, SEQ ID NO: 1320, SEQ ID NO: 1333, SEQ ID NO: 1385, SEQ ID NO: 1398, SEQ ID NO: 1411, SEQ ID NO: 1424, SEQ ID NO: 1474, SEQ ID NO: 1475, SEQ ID NO: 1476, SEQ ID NO: 1483, SEQ ID NO: 1490, SEQ ID NO: 1491, SEQ ID NO: 1492, SEQ ID NO: 1499, SEQ ID NO: 1506, SEQ ID NO: 1513, SEQ ID NO: 1514, SEQ ID NO: 1515, SEQ ID

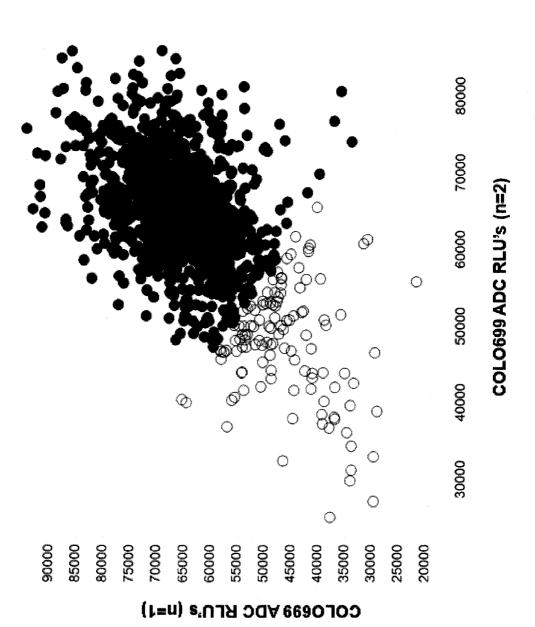
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- (e) as depicted in SEQ ID NO: 995, SEQ ID NO: 1594, and SEQ ID NO: 1607.
- 10. A nucleic acid sequence encoding an antibody construct as defined in any one of claims 1 to 9.
- 11. A vector comprising a nucleic acid sequence as defined in claim 10.
- 12. A host cell transformed or transfected with the nucleic acid sequence as defined in claim 10 or with the vector as defined in claim 11.
- 13. A process for the production of a antibody construct according to any one of claims 1 to 9, said process comprising culturing a host cell as defined in claim 12 under conditions allowing the expression of the antibody construct as defined in any one of claims 1 to 9 and recovering the produced antibody construct from the culture.
- 14. A pharmaceutical composition comprising an antibody construct according to any one of claims 1 to 9, or produced according to the process of claim 13.
- 15. The antibody construct according to any one of claims 1 to 9, or produced according to the process of claim 13 for use in the prevention, treatment or amelioration of a melanoma disease or metastatic melanoma disease.
- 16. A method for the treatment or amelioration of a melanoma disease or metastatic melanoma disease, comprising the step of administering to a subject in need thereof

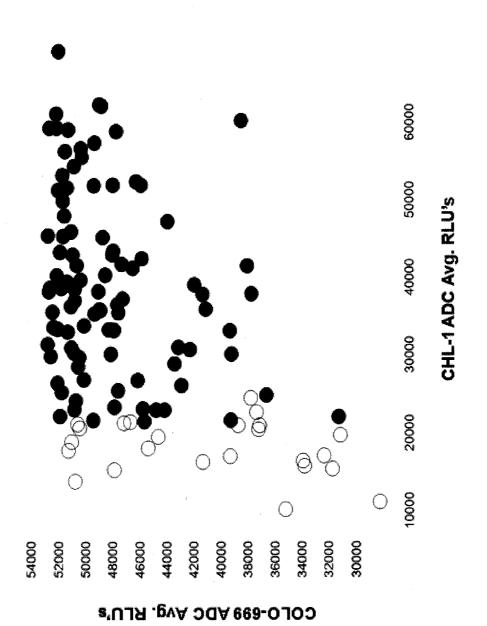
the antibody construct according to any one of claims 1 to 9, or produced according to the process of claim 13.

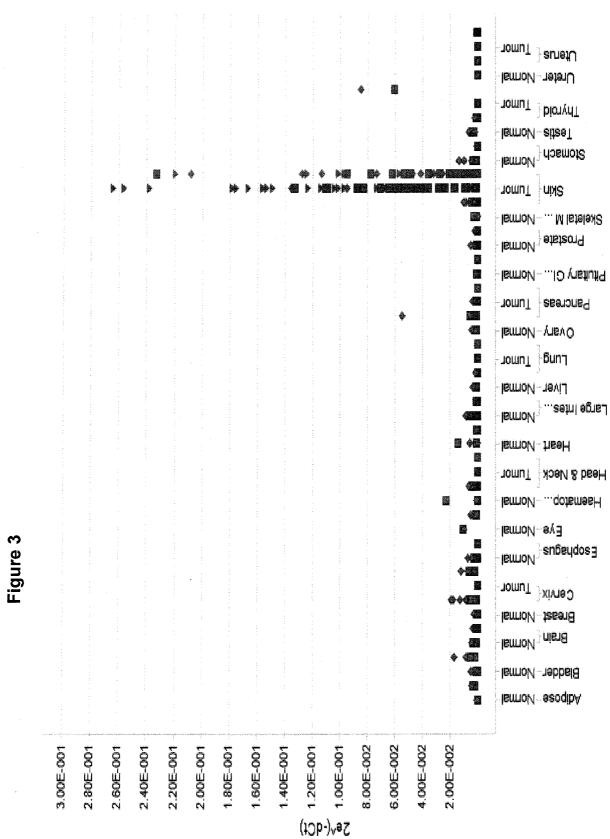
17. The method according to claim 16, wherein the melanoma disease or metastatic melanoma disease is selected from the group consisting of superficial spreading melanoma, lentigo maligna, lentigo maligna melanoma, acral lentiginous melanoma and nodular melanoma.

18. A kit comprising an antibody construct according to any one of claims 1 to 9, or produced according to the process of claim 13, a vector as defined in claim 11, and/or a host cell as defined in claim 12.



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Tissue of Origin, Normal, tumor, metastasis

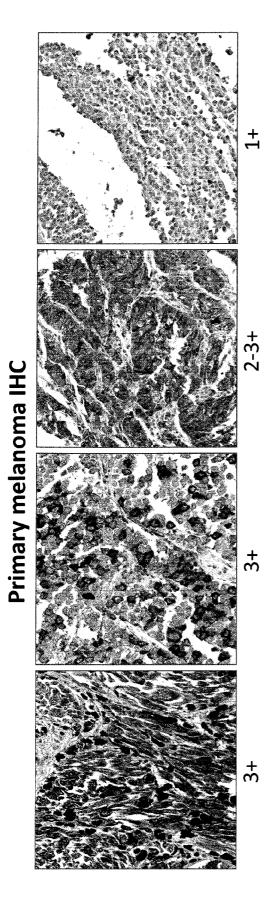


Figure 4

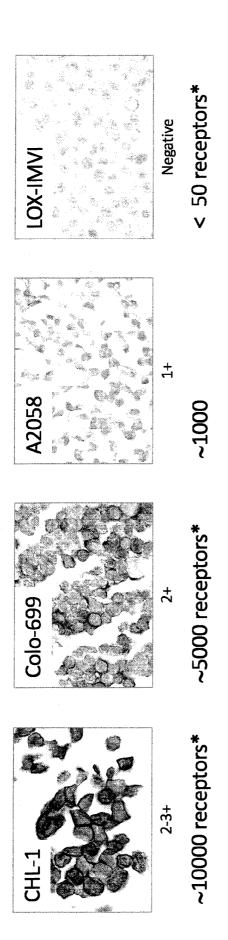
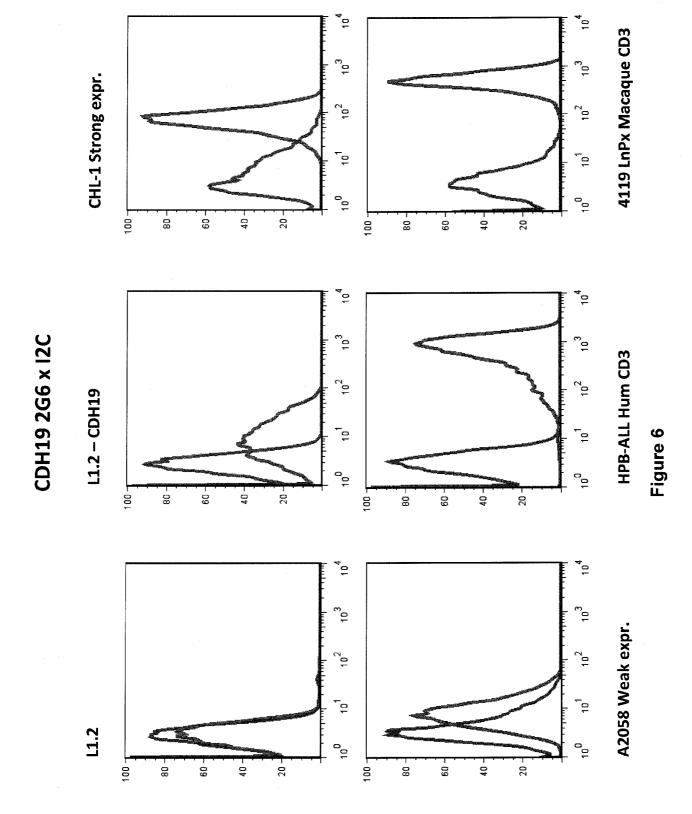


Figure (



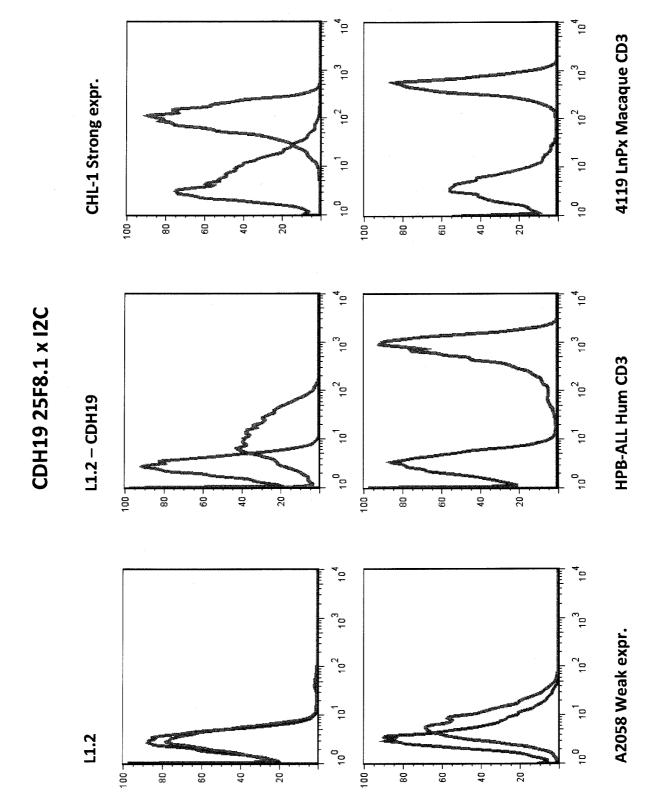


Figure 6 (continued)

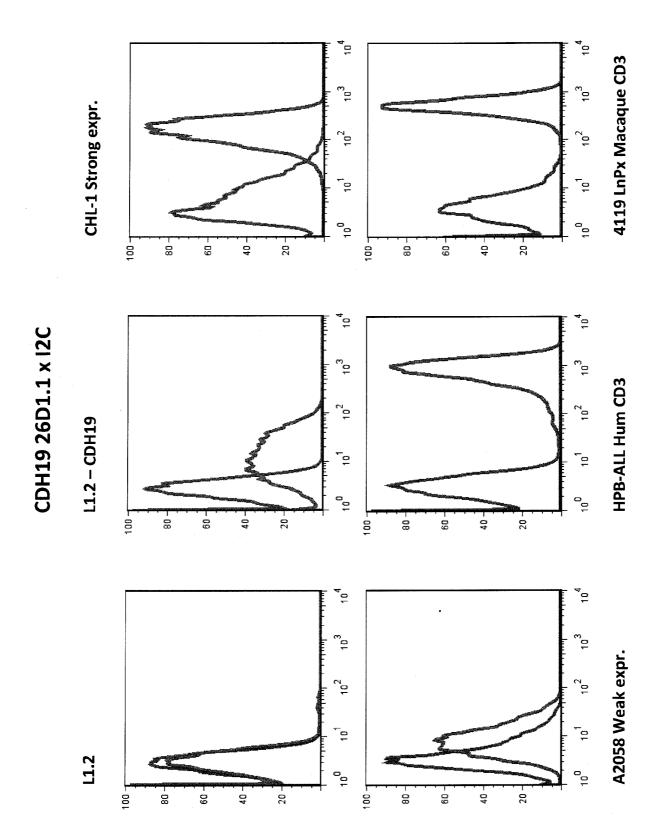


Figure 6 (continued)

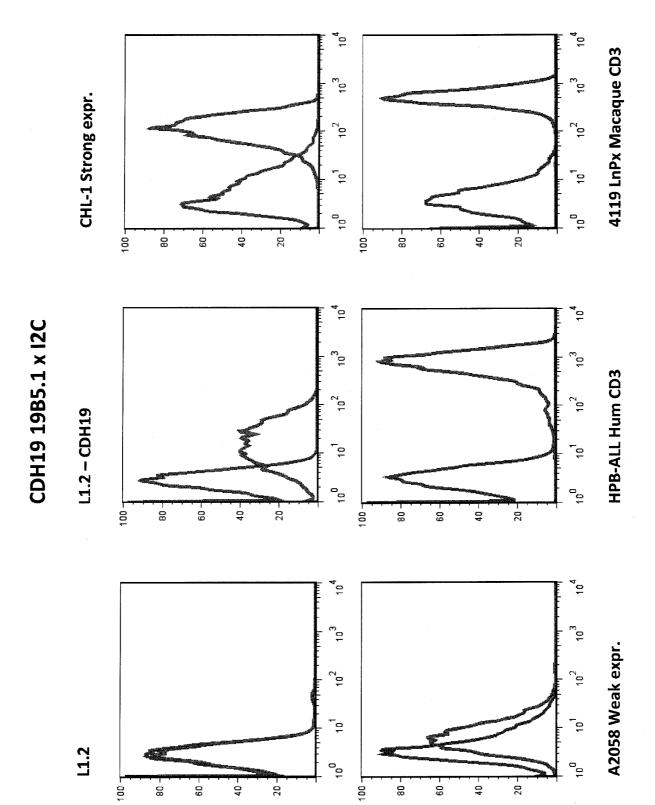


Figure 6 (continued)

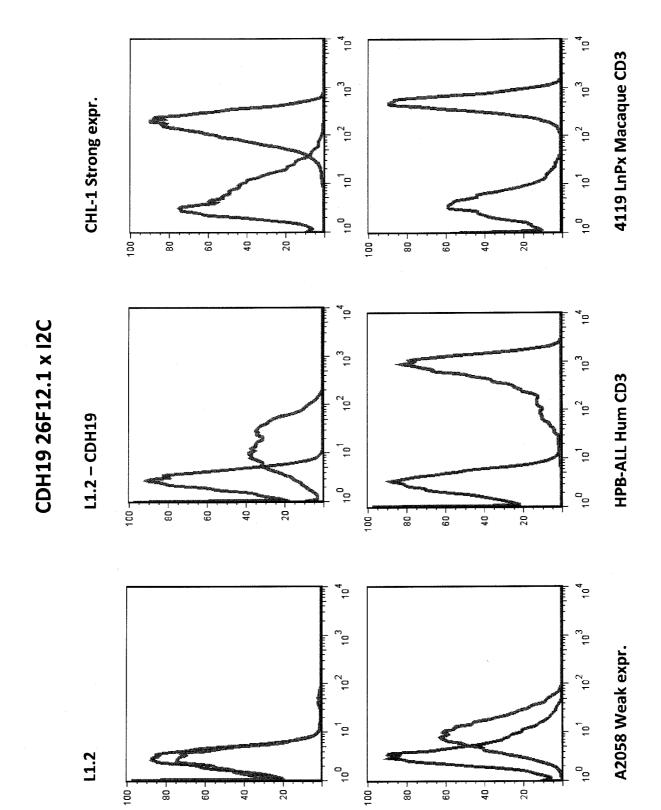
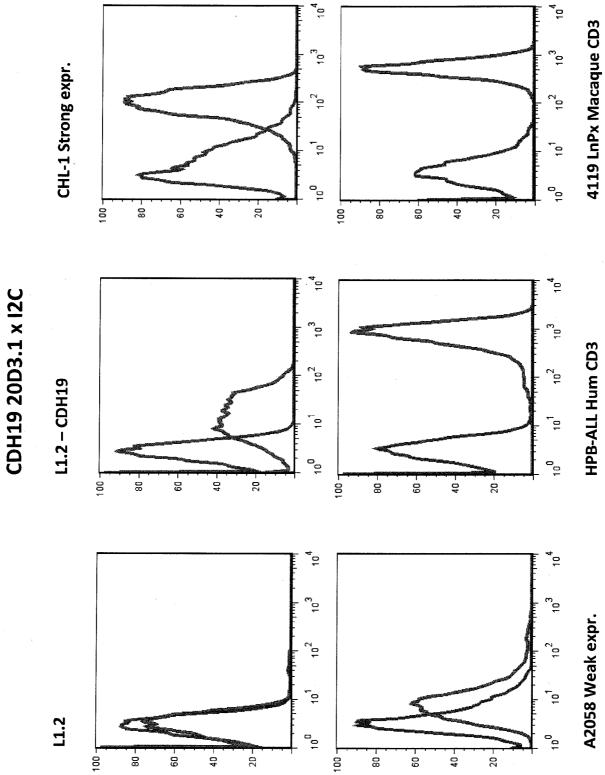


Figure 6 (continued)





- 08

- 09

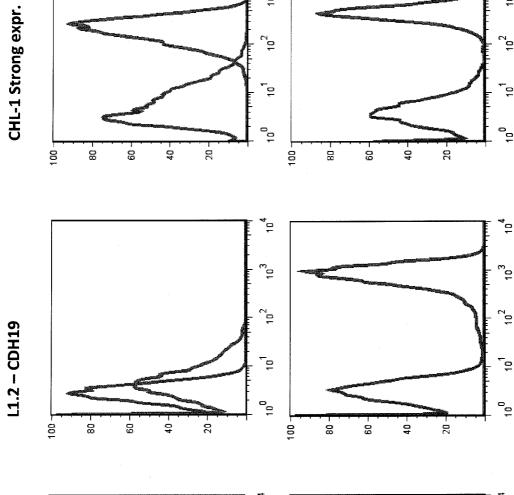
Figure 6 (continued)

4119 LnPx Macaque CD3

40



11.2



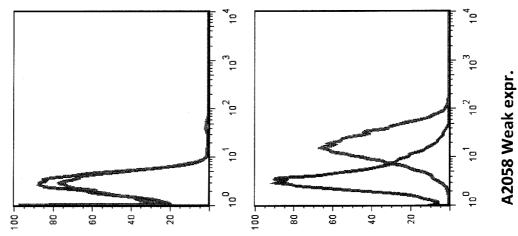


Figure 6 (continued)

HPB-ALL Hum CD3

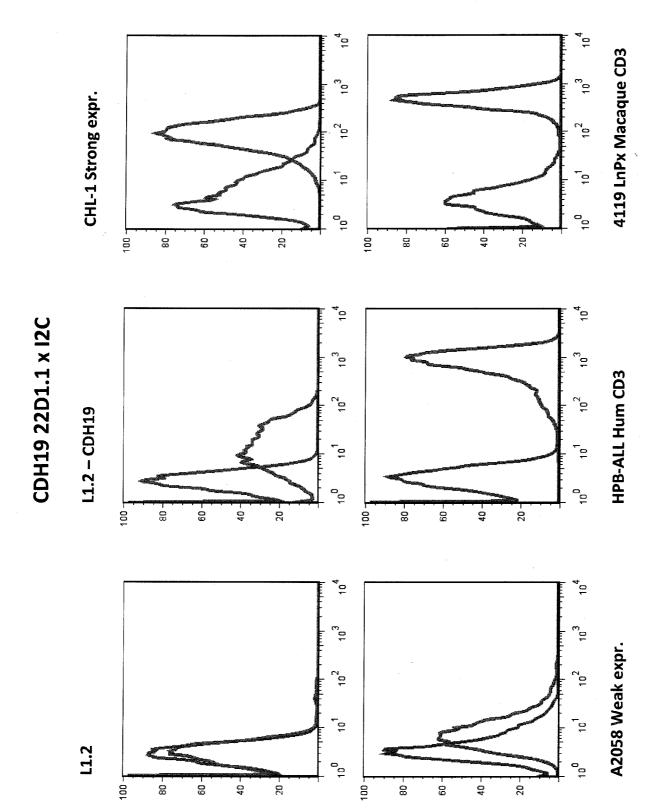


Figure 6 (continued)

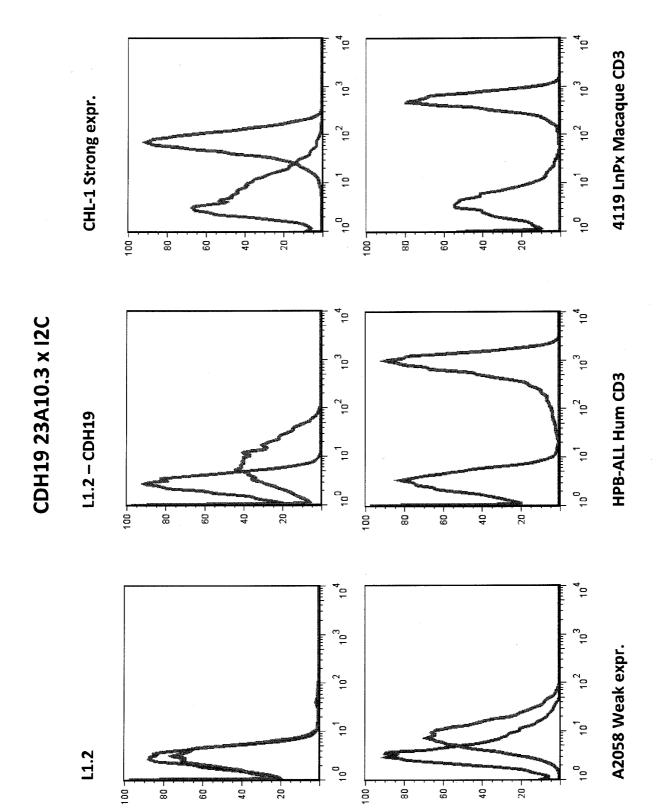


Figure 6 (continued)

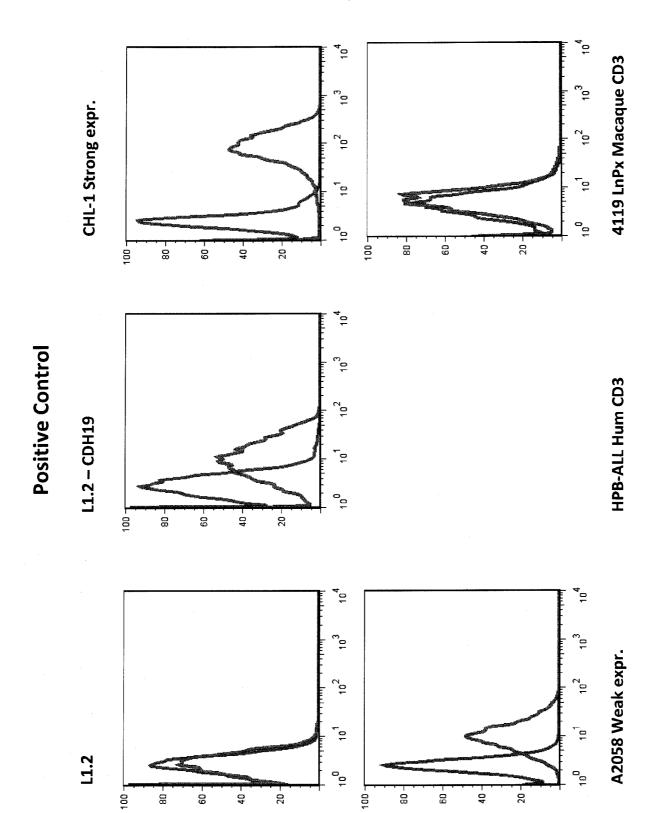


Figure 6 (continued)

6.5 346 23A10.3 x 12C 10-6 10-4 10-2 100 102 BiTE Conc. [pg/ml] 270 14379 22D1.1 x 12C CH19 23A10.3xI2C CH19 26F12.1x/2C CH19 26D1.1xI2C CH19 19B5.1xI2C CH19 20D3.1xI2C CH19 22D1.1xl2C CH19 25F8.1xI2C 304 16168 CH19 2G6x12C CH19 4F7x12C x 12C **4F7** 447 23796 20D3.1 x 12C 127 6748 **189** -08 20-8 \$ 26F12.1 x 12C (%) siskī 268 14236 19B5.1 x 12C 65.6 3487 106 104 102 100 102 104 26D1.1 x 12C BiTE Conc. [pg/ml] 131 6973 25F8.1 x 12C CH19 23A10.3xI2C CH19 26F12, 1x12C L1.2 CDH19 CH19 19B5.1xI2C CH19 22D1.1xI2C CH19 26D1.1xI2C CH19 20D3.1xI2C CH19 25F8, 1xt2C CH19 2G6x12C CH19 4F7x12C 261 x 12C 266 [bg/m]] pMolar 300 8 8

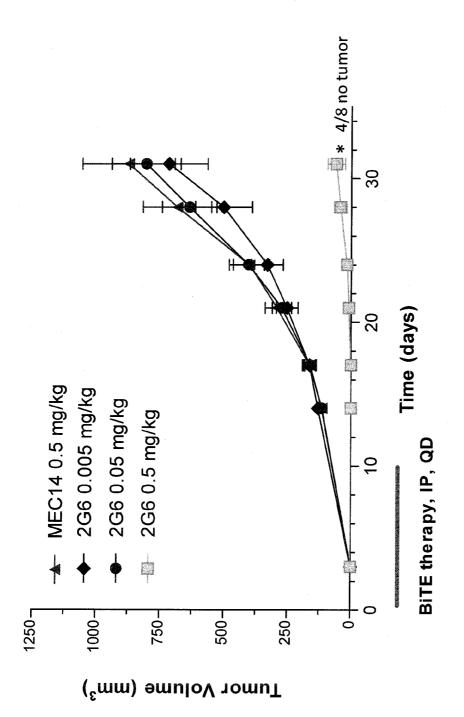
Figure 7

10-6 10-4 10-2 100 102 104 BITE Conc. [pg/ml] CH19 23A10.3 x I2C CH19 22D1.1 x I2C Neg. control BITE CH19 2G6 x I2C CH19 4F7 x 12C CHL-1 (CDH19 high expresser) 1001 8 -09 80 40-Lysis (%) BITE Conc. [pg/ml] 10-6 10-4 10-2 100 10-2 CH19 26F12.1 x 12C CH19 26D1.1 x 12C CH19 19B5.1 x I2C CH19 20D3.1 x I2C CH19 25F8.1 x 12C Neg. control BiTE 1001 40-8 5 -08 9 Lysis (%)

EC ₅₀	2G6 x I2C	25F8.1 x I2C	25F8.1 26D1.1 19B5.1 26F12.1 20D3.1 4F7 x12C x12C x12C x12C x12C	19B5.1 x I2C	26F12.1 x I2C	20D3.1 x I2C	4F7 x 12C	22D1.1 x I2C	22D1.1 23A10.3 x 2C x 2C
[lm/gd]	0.4	28	27	6.9	21	20	1842	28	3.6
fMolar	7.5	526	508	130	395	376	34630	526	67.7

Figure 7 (continued)





@ Tumors and human PBMC inoculated at a 2:1 ratio on Day 0.

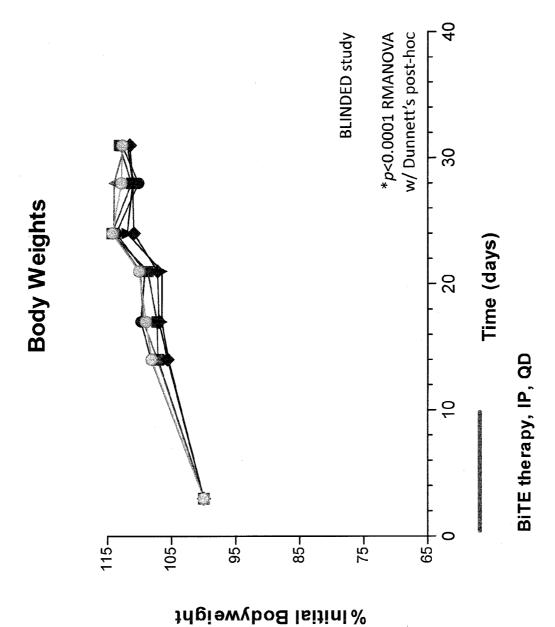


Figure 8 (continued)



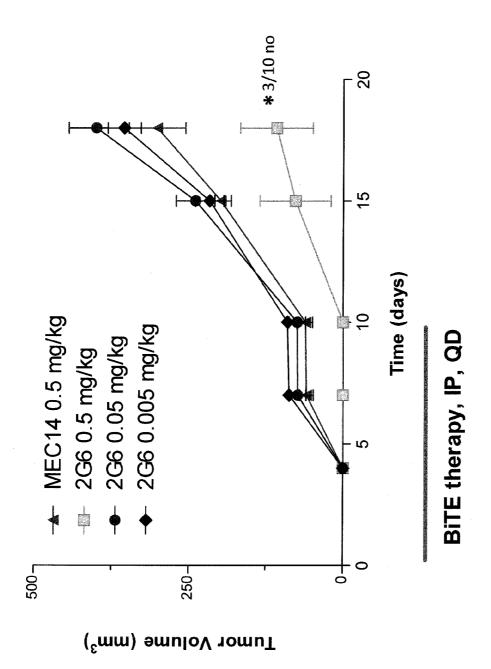
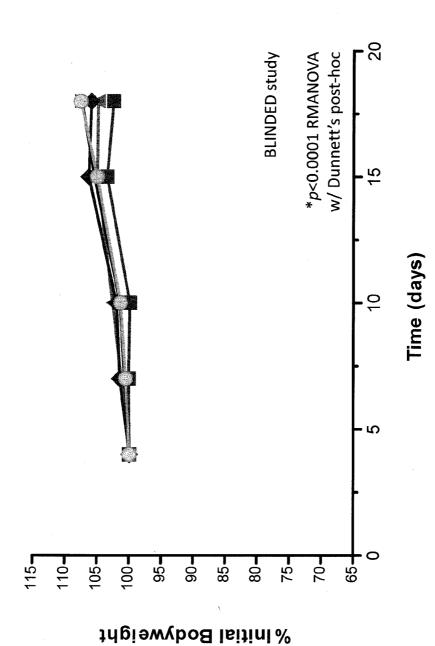


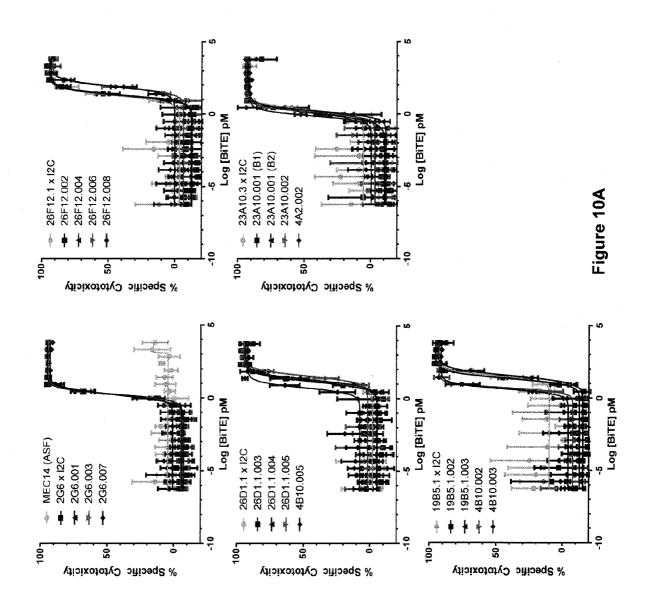
Figure 9



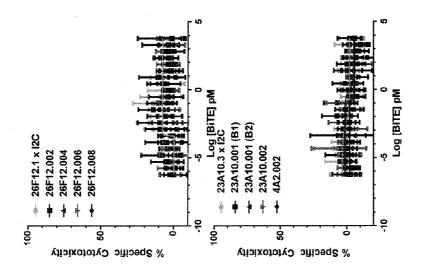


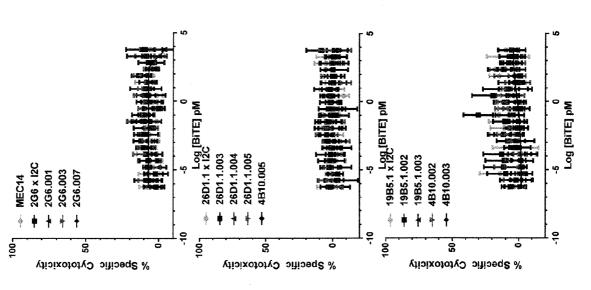
BiTE therapy, IP, QD Figure 9 (continued)

EC50[pM]	1.729	1.576	1.721	1.783	18.48	19.4	20.74	72.22	83.43	33.59	22.37	34.62	4.83	4.088	5.583	19.19	15.53	19.77	36.42	2.086	1.078	1.623	1.351	0.7547
СDH19 ВіТЕ	2G6 x 12C	2G6.001	2G6.003	2G6.007	26F12.1 x I2C	26F12.002	26F12.004	26F12.006	26F12.008	1985.1 x 12C	1985.1.002	1985.1.003	4810.002	4B10.003	4810.005	26D1.1 x I2C	26D1.1.003	26D1.1.004	26D1.1.005	23A10.3 x I2C	23A10.001 (B1)	23A10.001 (B2)	23A10.002	4A2.002



112C	CDH19 BiTE	ECSO[BM]
2C no specific 3 no specific 7 no specific 92 no specific 94 no specific 96 no specific 96 no specific 03 no specific 93 no specific 93 no specific 94 no specific 95 no specific 96 no specific 97 no specific 98 no specific 99 no specific 12C no specific		
1 no specific 2 no specific 12 no specific 22 no specific 24 no specific 26 no specific 27 no specific 28 no specific 29 no specific 20 no specific 31 no specific 32 no specific 34 no specific 35 no specific 36 no specific 37 no specific 38 no specific 39 no specific 481 no specific 482 no specific 682 no specific 683 no specific 684 no specific 685 no specific 686 no specific 687 no specific 687 no specific	2G6 x I2C	no specific activity
3 no specific 7 no specific 12C no specific 04 no specific 06 no specific 06 no specific 07 no specific 03 no specific 03 no specific 04 no specific 05 no specific 06 no specific 07 no specific 08 no specific 09 no specific 04 no specific 05 no specific 06 no specific 07 no specific 08 no specific 09 no specific 005 no specific 02 no specific 02 no specific 02 no specific	266.001	no specific activity
7 no specific 12C no specific 04 no specific 04 no specific 06 no specific 08 no specific 02 no specific 03 no specific 04 no specific 04 no specific 05 no specific 04 no specific 05 no specific 065 no specific 07 no specific (B1) no specific 02 no specific 04 no specific 05 no specific 06 no specific 07 no specific 08 no specific 09 no specific	2G6.003	no specific activity
12C no specific 04 no specific 04 no specific 06 no specific 08 no specific 12C no specific 03 no specific 03 no specific 12C no specific 03 no specific 04 no specific 05 no specific 04 no specific 05 no specific (B1) no specific 02 no specific	266.007	no specific activity
02 no specific 04 no specific 06 no specific 08 no specific 03 no specific 03 no specific 03 no specific 04 no specific 04 no specific 05 no specific 04 no specific 05 no specific 05 no specific 05 no specific 05 no specific 02 no specific	26F12.1 x I2C	no specific activity
04 no specific 06 no specific 08 no specific 02 no specific 03 no specific 03 no specific 12C no specific 03 no specific 04 no specific 05 no specific 05 no specific (B1) no specific (B2) no specific 02 no specific 03 no specific 04 no specific	26F12.002	no specific activity
06 no specific 08 no specific 12C no specific 03 no specific 03 no specific 12C no specific 03 no specific 12C no specific 04 no specific 05 no specific (B1) no specific (B2) no specific 02 no specific 02 no specific 03 no specific 04 no specific 05 no specific 06 no specific	26F12.004	no specific activity
08 no specific 12C no specific 03 no specific 12 no specific 12 no specific 12C no specific 03 no specific 04 no specific 05 no specific 12C no specific 05 no specific (B1) no specific 02 no specific 2 no specific 2 no specific	26F12.006	no specific activity
I2C no specific 02 no specific 03 no specific 12 no specific 15 no specific 12C no specific 04 no specific 05 no specific (B1) no specific (B2) no specific 02 no specific 2 no specific	26F12.008	no specific activity
02 no specific 03 no specific 12 no specific 13 no specific 14 no specific 15 no specific 16 no specific 17 no specific 18 no specific 18 no specific 10 no specific		no specific activity
03 no specific 12 no specific 13 no specific 12C no specific 03 no specific 04 no specific 05 no specific (B1) no specific 02 no specific 2 no specific	1985.1.002	no specific activity
12 no specific 13 no specific 15 no specific 12C no specific 03 no specific 04 no specific 05 no specific (B1) no specific 02 no specific 2 no specific	1985.1.003	no specific activity
B3 no specific L2C no specific 12C no specific 03 no specific 04 no specific 12C no specific (B1) no specific (B2) no specific 02 no specific 2 no specific	4810.002	no specific activity
12C no specific 03 no specific 04 no specific 05 no specific (12C no specific (12C) no specific	4B10.003	no specific activity
12C no specific 03 no specific 04 no specific 05 no specific (12C no specific (B1) no specific 02 no specific 2 no specific	4810.005	no specific activity
03 no specific 04 no specific 05 no specific 12C no specific (B1) no specific 02 no specific 2 no specific		no specific activity
04 no specific 05 no specific (12C no specific (B1) no specific 02 no specific 2 no specific	26D1.1.003	no specific activity
(12C no specific (12C no specific (12) no specific (12) no specific (13) no specific (14) no specific	26D1.1.004	no specific activity
(B1) no specific (B2) no specific 02 no specific 2	26D1.1.005	no specific activity
(B1) no specific (B2) no specific 02 no specific 2 no specific		no specific activity
(B2) no specific 02 no specific 2 no specific	1	no specific activity
no specific no specific		no specific activity
no specific	23A10.002	no specific activity
	4A2.002	no specific activity





igure 10B

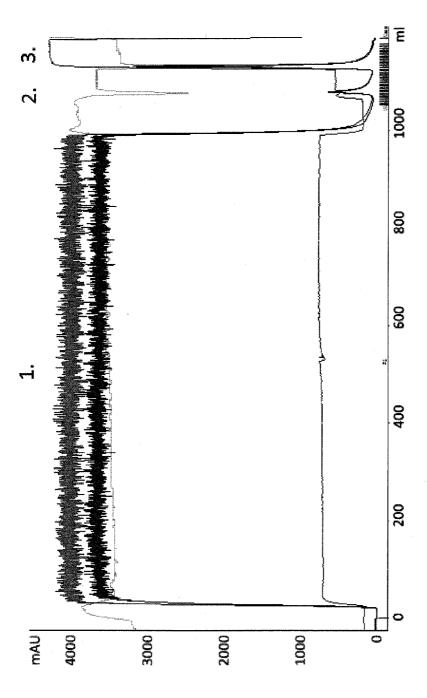
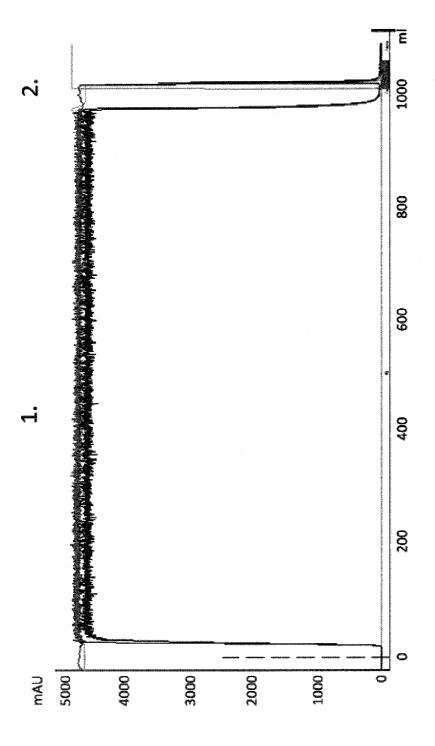
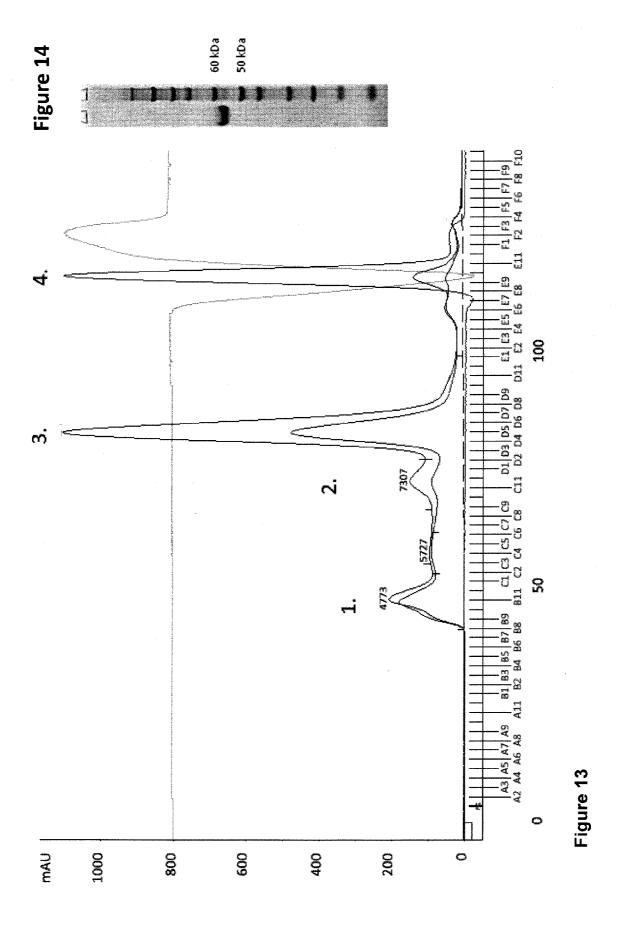


Figure 11







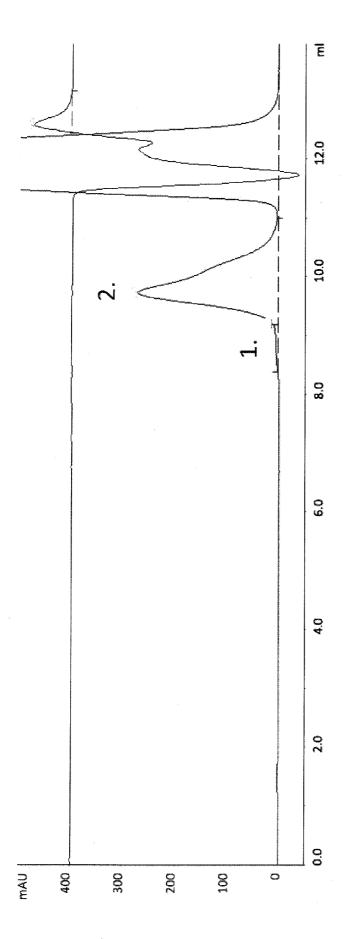


Figure 1

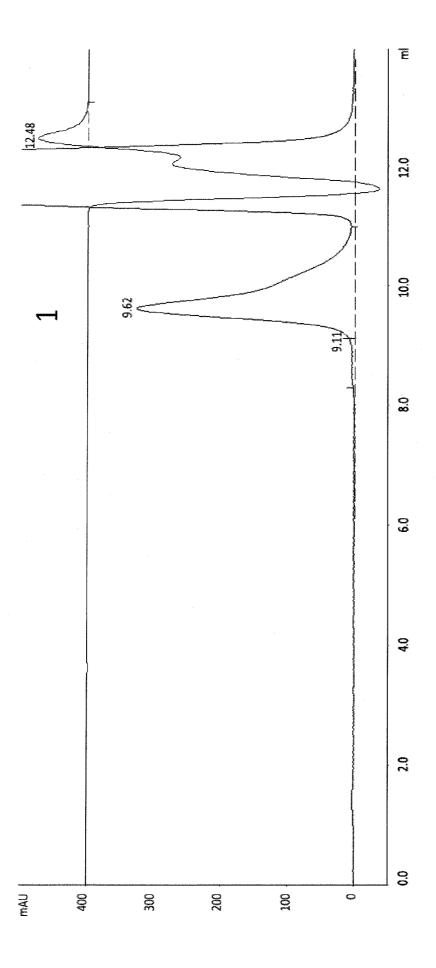
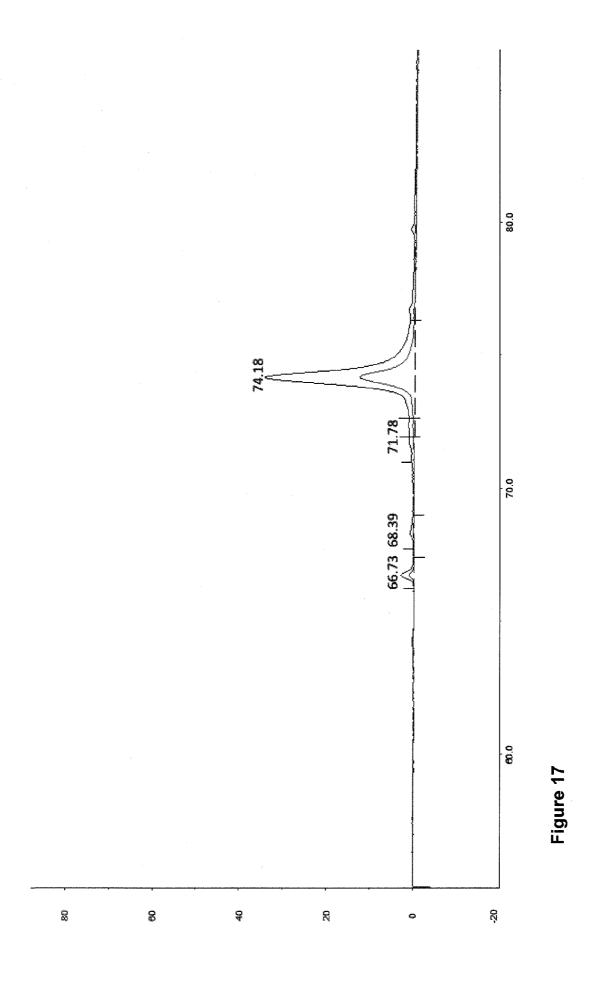


Figure 16



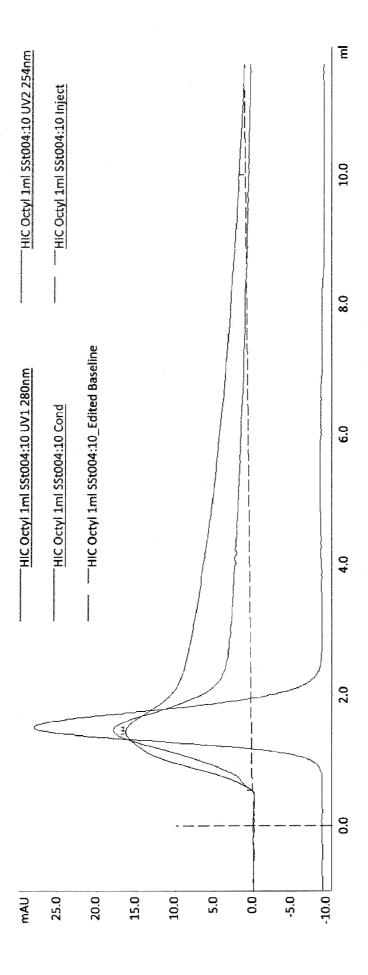
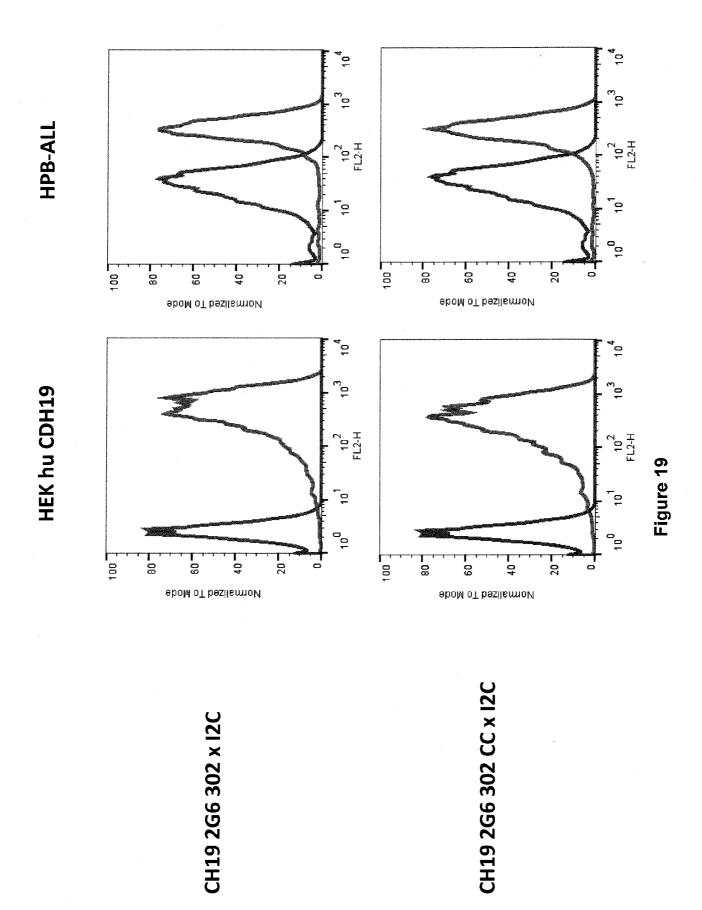


Figure 18



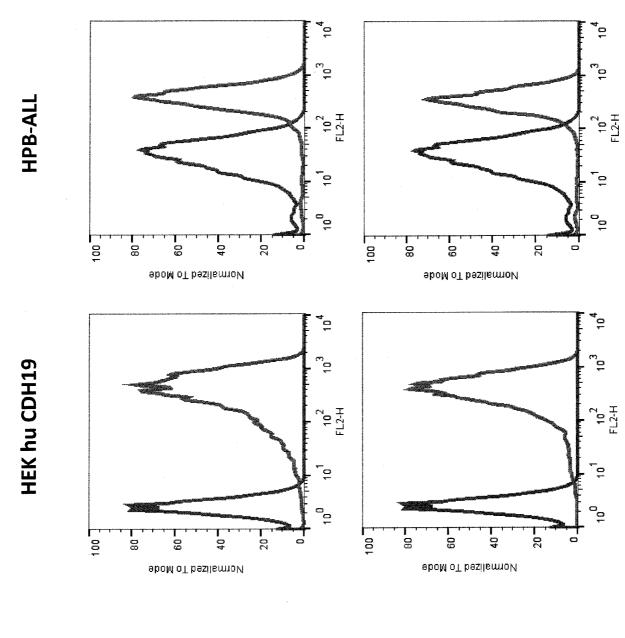


Figure 19 (continued)

CH19 2G6 303 x I2C

CH19 2G6 303 CC x I2C

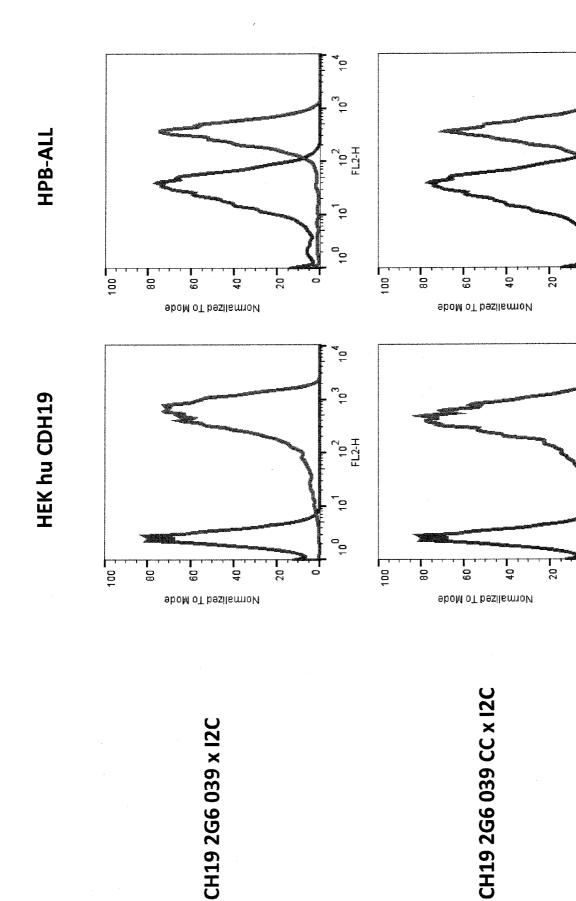


Figure 19 (continued)

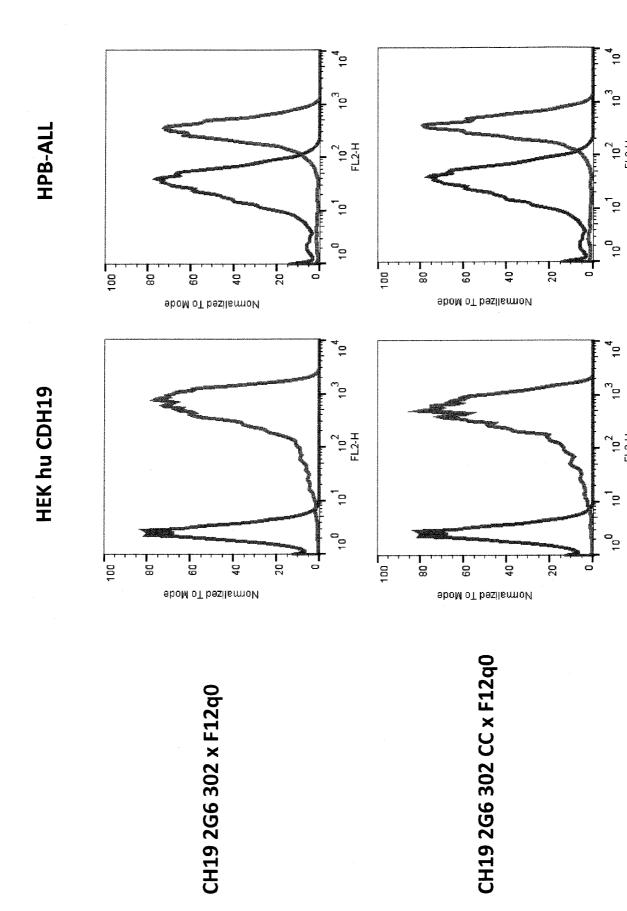


Figure 19 (continued)

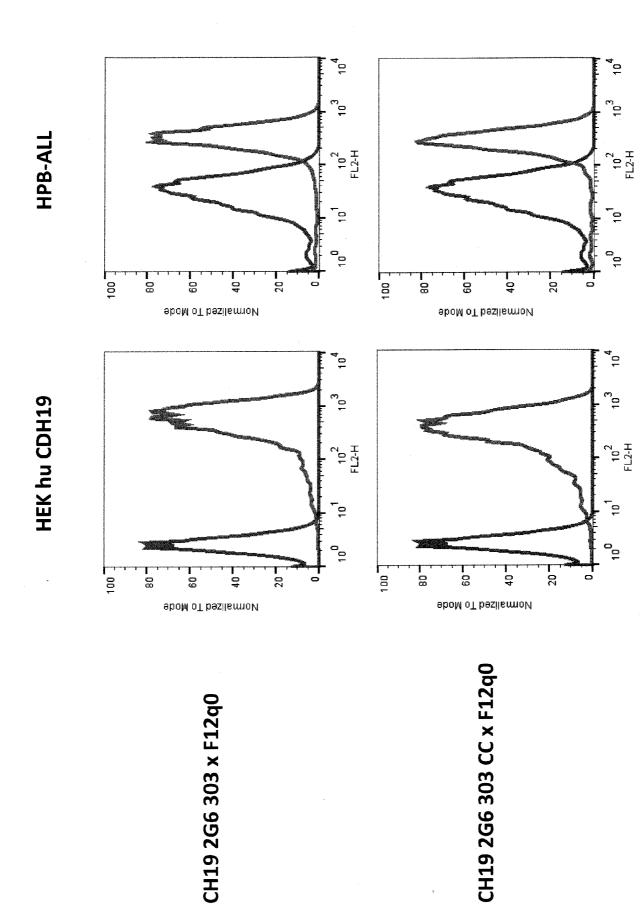


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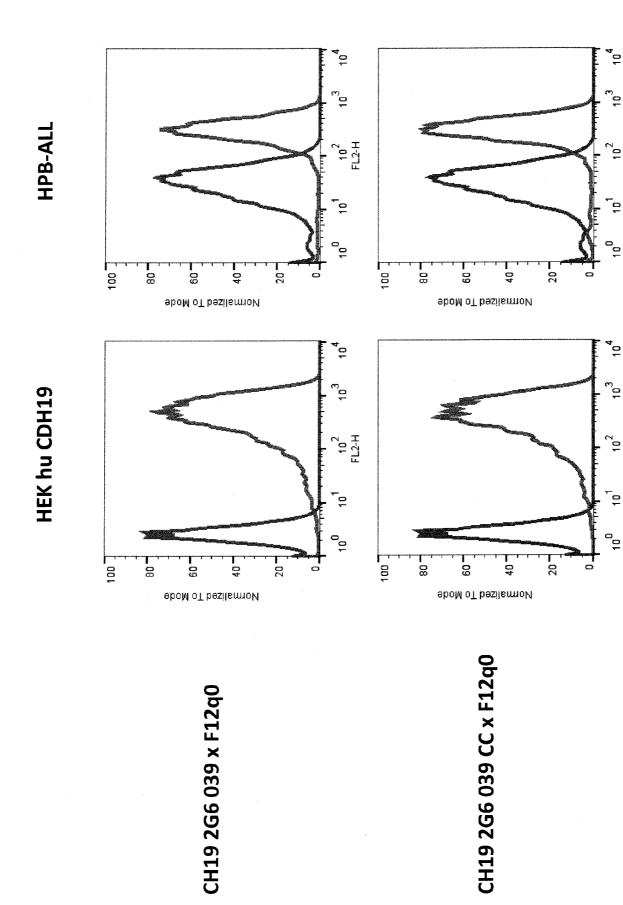


Figure 19 (continued)

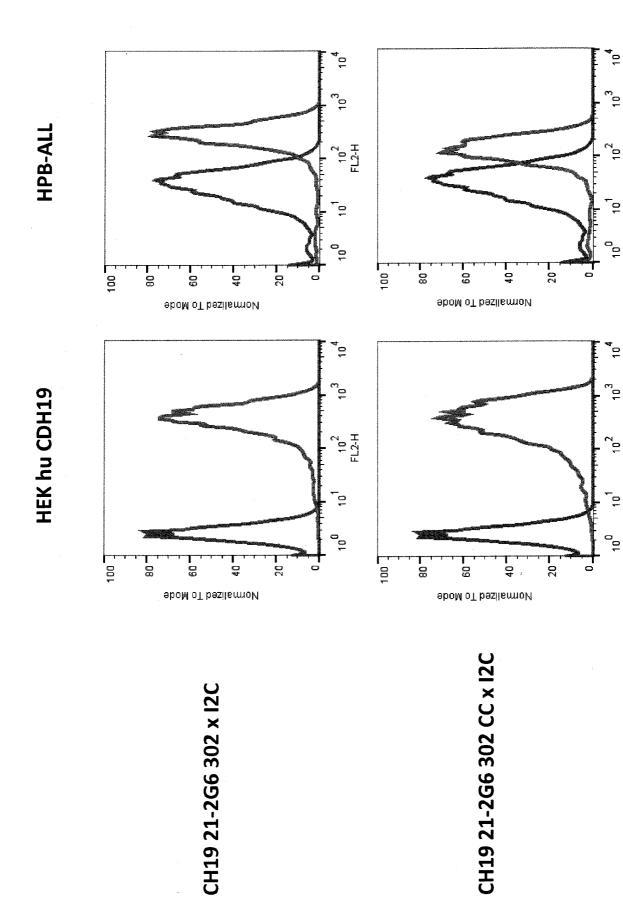


Figure 19 (continued)

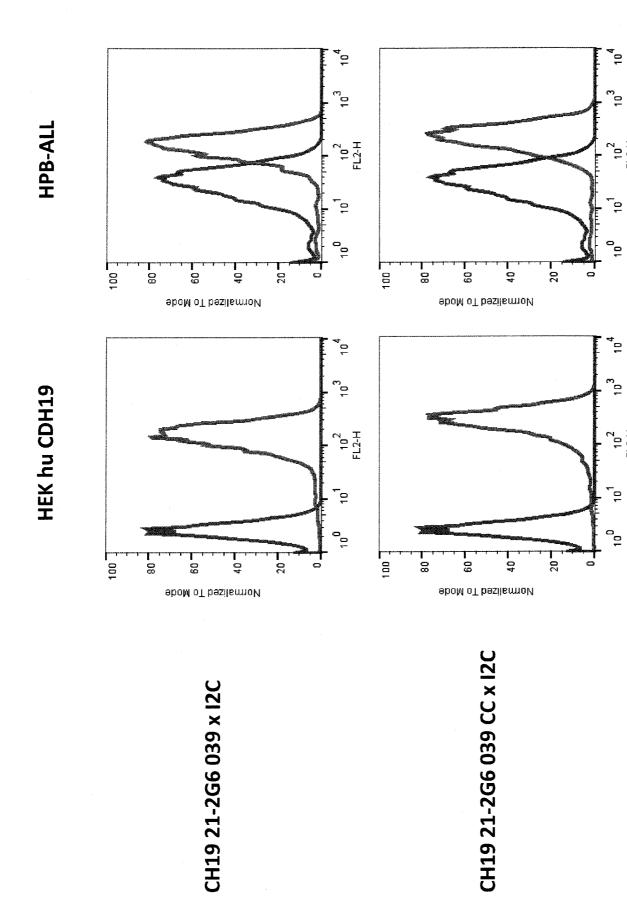


Figure 19 (continued)

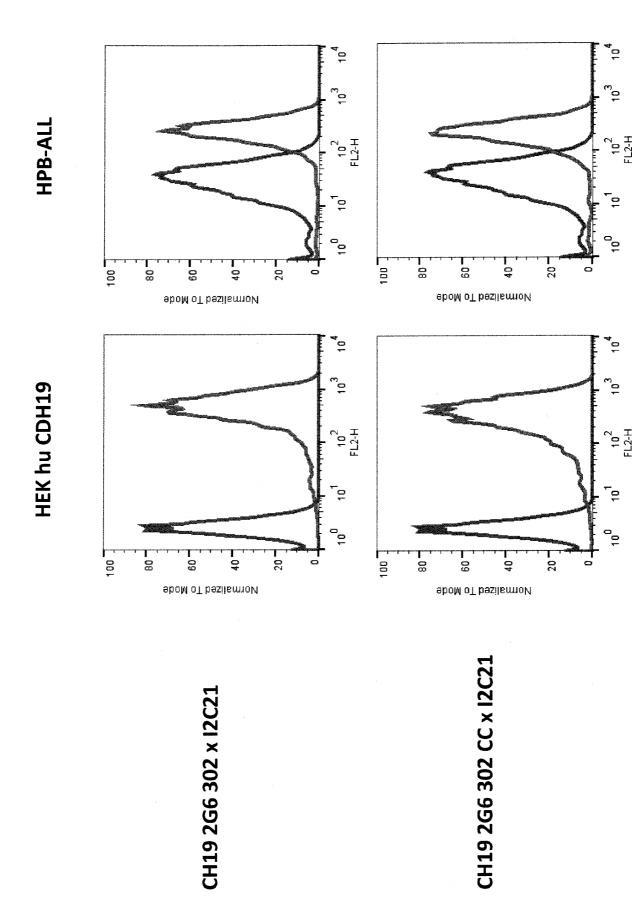


Figure 19 (continued)

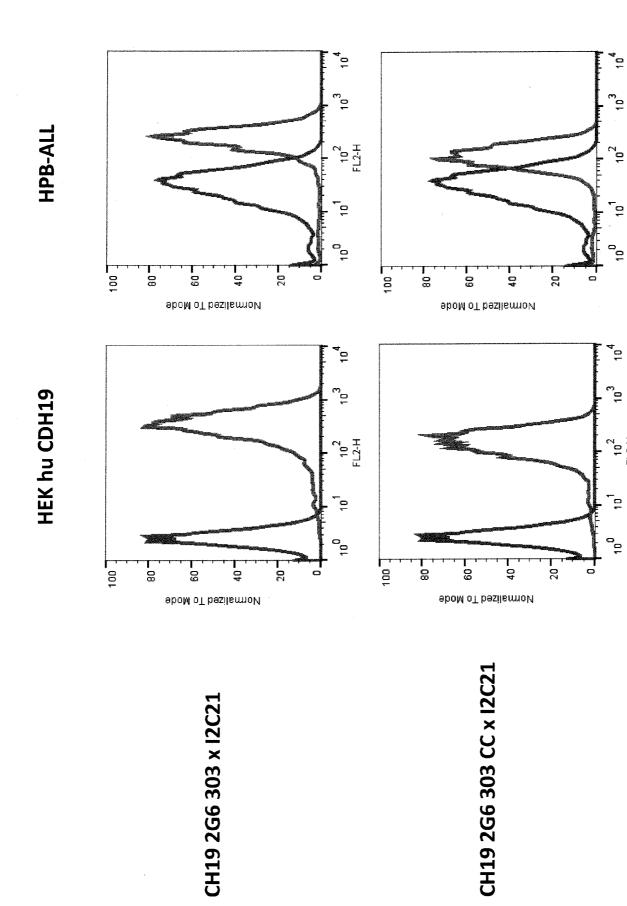


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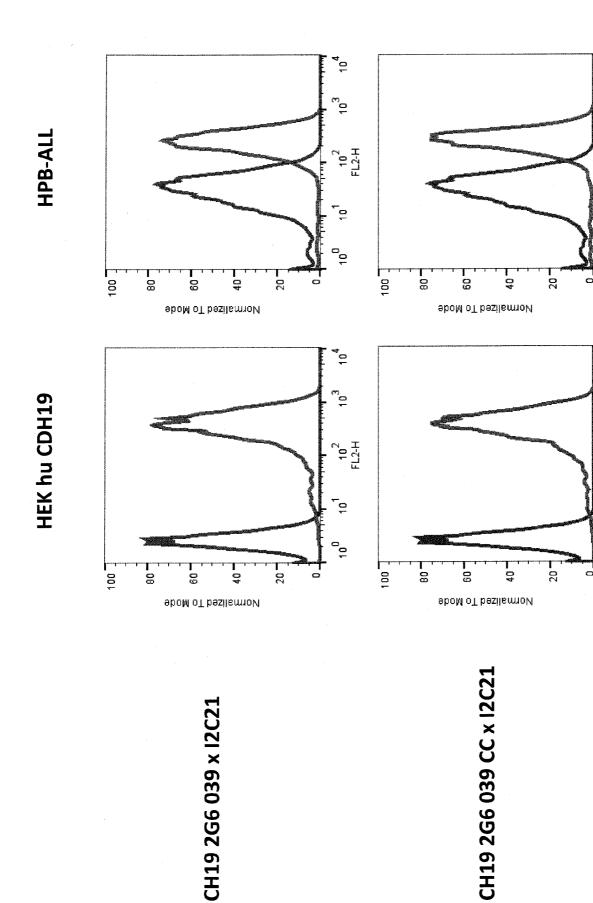


Figure 19 (continued)

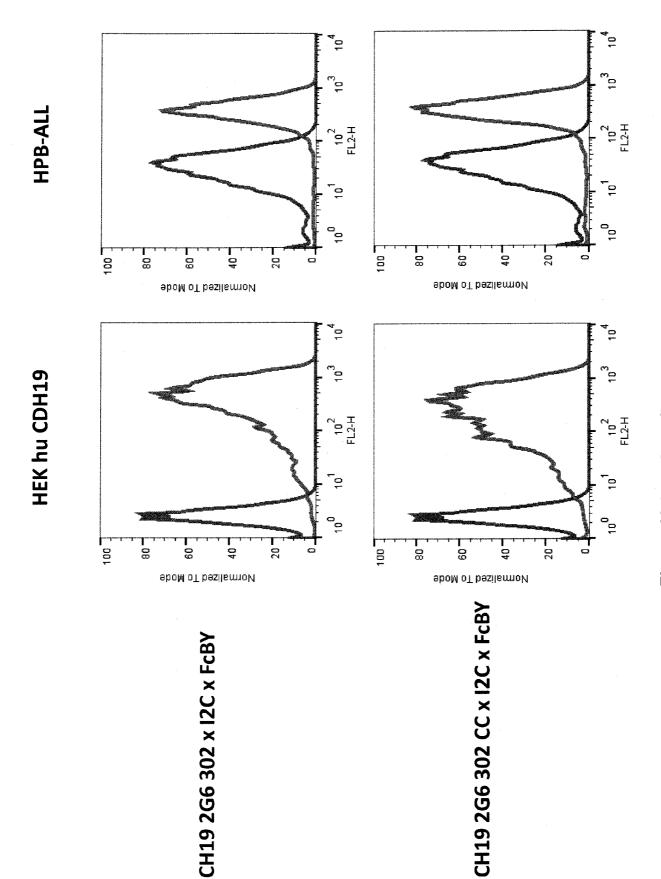


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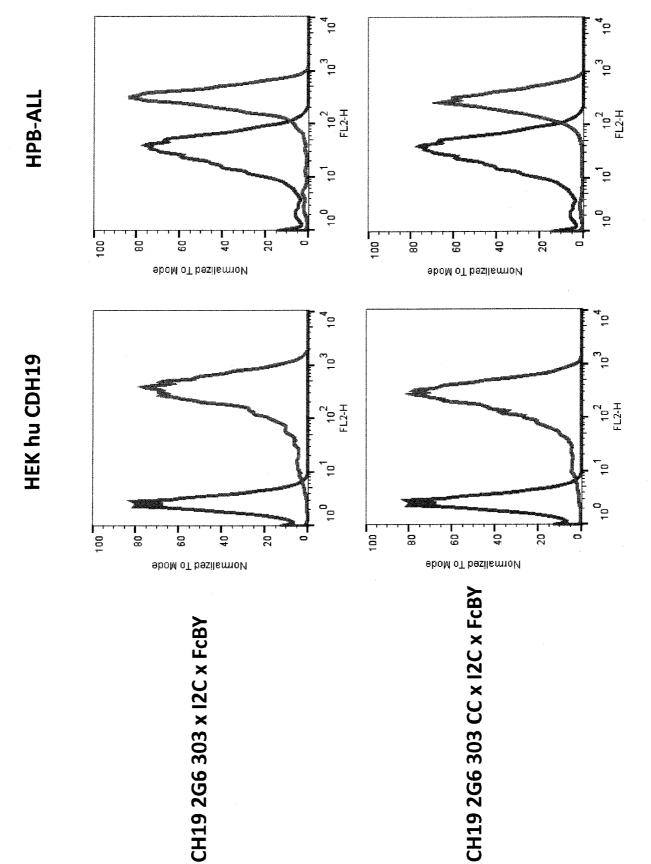


Figure 19 (continued)

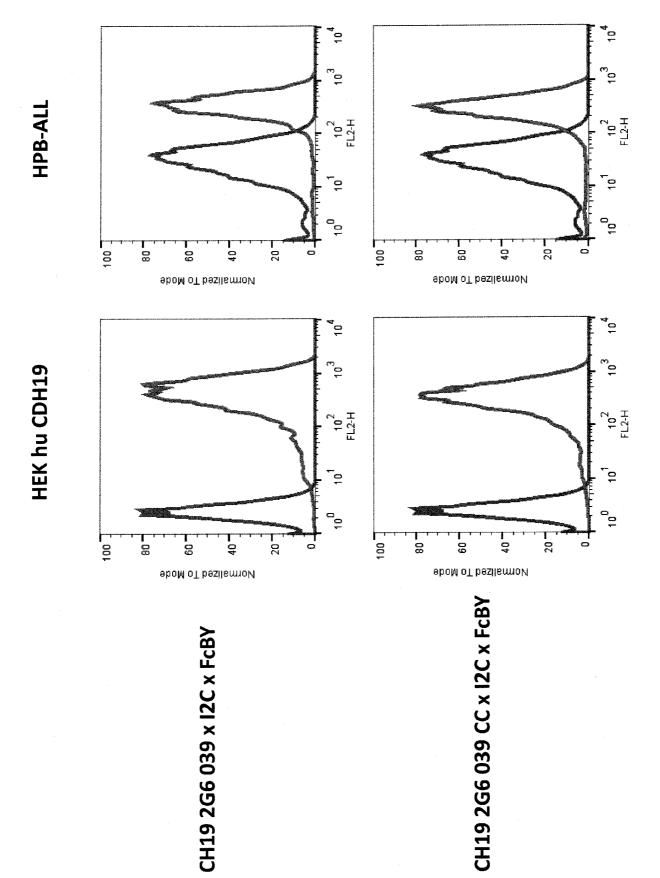


Figure 19 (continued)



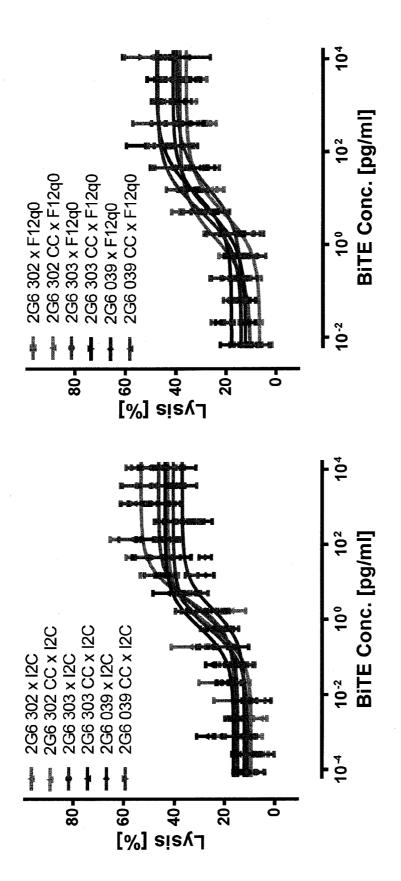


Figure 20



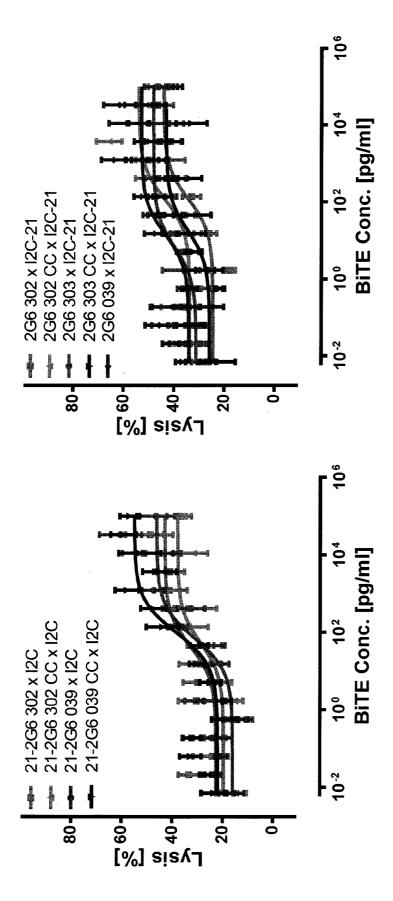


Figure 20 (continued)

International application No.

INTERNATIONAL SEARCH REPORT PCT/EP2014/051550 Box No. I Nucleotide and/or amino acid sequence(s) (Continuation of item 1.c of the first sheet) With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, the international search was carried out on the basis of: (means) on paper in electronic form (time) in the international application as filed together with the international application in electronic form subsequently to this Authority for the purpose of search 2. In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed Х or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished. 3. Additional comments:

International application No
PCT/EP2014/051550

A. CLASSIFICATION OF SUBJECT MATTER
INV. C07K16/28 C07K16/30 A61K47/48 A61K39/395 A61P35/00
C07K16/46

ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) $C07\,K-A61\,K$

OOTK MOIK

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, BIOSIS, EMBASE, WPI Data

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 2006/071441 A2 (CURAGEN CORP [US]; ABGENIX INC [US]; XIAO FENG [US]; JIA XIAO-CHI [US]) 6 July 2006 (2006-07-06) whole document, especially Examples 13, 19, 20, 28; Figures 1, 4, 5, 20	1-18

Further documents are listed in the continuation of Box C.	X See patent family annex.	
* Special categories of cited documents : "A" document defining the general state of the art which is not considered	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	
to be of particular relevance "E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art	
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"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed	"&" document member of the same patent family	
Date of the actual completion of the international search	Date of mailing of the international search report	
14 May 2014	22/05/2014	
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2	Authorized officer	
NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Luyten, Kattie	

1

International application No
PCT/EP2014/051550

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT					
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.			
Y	KISCHEL ROMAN ET AL: "Characterization of novel CD33-and MCSP-specific BiTE antibodies for the treatment of acute myeloid leukemia and melanoma, respectively, that are fully human in sequence", AMERICAN ASSOCIATION FOR CANCER RESEARCH. PROCEEDINGS OF THE ANNUAL MEETING, AMERICAN ASSOCIATION FOR CANCER RESEARCH, US, vol. 49, 12 April 2008 (2008-04-12), pages 567-568, XP009175475, ISSN: 0197-016X the whole document	1-18			
Y	WO 2008/119567 A2 (MICROMET AG [DE]; EBERT EVELYN [DE]; MEIER PETRA [DE]; SRISKANDARAJAH) 9 October 2008 (2008-10-09) cited in the application whole document, especially Table 2; Examples 16-18	1-18			
Y	BERTUCCI FRANÇOIS ET AL: "Gene expression profiling of human melanoma cell lines with distinct metastatic potential identifies new progression markers", ANTICANCER RESEARCH - INTERNATIONAL JOURNAL OF CANCER RESEARCH AND TREATMENT, INTERNATIONAL INSTITUTE OF ANTICANCER RESEARCH, GR, vol. 27, no. 5A, 1 September 2007 (2007-09-01), pages 3441-3449, XP009154071, ISSN: 0250-7005 whole document, especially the Abstract; Table I; page 3446, left-hand column, lines 44-46	1-18			
A	Anonymous: "Anti-CDH19 Product Datasheet", December 2012 (2012-12), XP055117756, Retrieved from the Internet: URL:https://atlasantibodies.com/print_data sheet/R74953 [retrieved on 2014-05-13] the whole document	1-18			

1

International application No
PCT/EP2014/051550

C(Continua	tion). DOCUMENTS CONSIDERED TO BE RELEVANT	
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Α	Anonymous: "CDH19 monoclonal antibody (M01), clone 1G4",	1-18
	January 2008 (2008-01-01), page 1, XP055117753, Retrieved from the Internet: URL:http://www.abnova.com/protocol_pdf/DS_H00028513-M01.pdf [retrieved on 2014-05-13]	
А	J. NIU ET AL: "Monocyte Chemotactic Protein (MCP)-1 Promotes Angiogenesis via a Novel Transcription Factor, MCP-1-induced Protein (MCPIP)", JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 283, no. 21, 23 May 2008 (2008-05-23), pages 14542-14551, XP055116978, ISSN: 0021-9258, DOI: 10.1074/jbc.M802139200 page 14545, left-hand column, line 15	1-18

1

Information on patent family members

International application No
PCT/EP2014/051550

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2006071441 A2	06-07-2006	AT 476994 T AU 2005322410 A1 CA 2589374 A1 DK 1827492 T3 EP 1827492 A2 EP 2305716 A2 JP 2008521411 A JP 2012120544 A JP 2014003986 A US 2013022597 A1 WO 2006071441 A2	15-08-2010 06-07-2006 06-07-2006 22-11-2010 05-09-2007 06-04-2011 26-06-2008 28-06-2012 16-01-2014 24-01-2013 06-07-2006
WO 2008119567 A2	09-10-2008	CN 101687915 A CN 103694350 A EP 2155783 A2 EP 2520590 A2 ES 2432792 T3 NZ 580755 A RU 2009136912 A SG 182234 A1 SG 195609 A1 WO 2008119567 A2	31-03-2010 02-04-2014 24-02-2010 07-11-2012 05-12-2013 25-05-2012 10-05-2011 30-07-2012 30-12-2013 09-10-2008