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**PILLAI, S. K. K. ET AL.: 'Triple-negative breast cancer is associated with EGFR, CK5/6 and c-KIT expression in Malaysian women' BMC CLINICAL PATHOLOGY vol. 12, no. 18, 26 September 2012, pages 12 - 18, XP021139347**  
**SETON-ROGERS, S. ET AL.: 'On the origins of tumour subtypes' NATURE REVIEWS CANCER vol. 7, October 2007, XP055189503**

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# DESCRIPTION

## BACKGROUND

**[0001]** World wide, greater than 1 million women are diagnosed with breast cancer each year. Breast cancer is a very heterogeneous disease made up of dozens of different types that are distinguished using a histological classification system. A large subtype and a majority of cases are histologically identified as luminal A or luminal B which can be grossly characterized as exhibiting estrogen receptor (ER) expression with low grade or higher grade histology, respectively (Santana-Davila and Perez, 2010). Immunohistochemical methods are used to measure the expression of progesterone receptor (PgR) which, when coupled with ER-positive status allows the classification of a tumor as being hormone responsive. Furthermore, the over-expression or amplification of human epidermal growth factor receptor 2 (HER2) can be monitored either with immunohistochemistry or fluorescence *in situ* hybridization (FISH). Generally, the expression of these three markers in breast tumors is associated with a better clinical outcome because there are several treatment options available for patients that target these proteins (de Ruijter et al., 2011), including tamoxifen, Arimidex™ (anastrozole), Aromasin™ (exemestane), Femara™ (letrozole), Faslodex™ (fulvestrant), Herceptin™ (trastuzumab) or Tykerb™ (lapatinib).

**[0002]** Another histological subtype of breast cancer consists of the basal-like cancers which are associated with, among others, a higher histological grade, increase mitotic index and high Ki67 expression (Santana-Davila and Perez, 2010). The vast majority of basal-like cancers are comprised of triple-negative breast cancer (TNBC) cases, which make up a between 15-20% of all diagnosed breast cancer cases (Ismail-Khan and Bui, 2010). TNBC is defined by the lack of protein expression of ER, PgR and the absence of HER2 protein over-expression. The relationship between basal-like cancer and TNBC is not easily delineated since not all TNBC are basal-like and not all basal-like cancers are TNBC, but approximately 75% of cases in these categories share characteristics of both. TNBC is associated with poor prognosis consisting of low five-year survival rates and high recurrence.

**[0003]** Patients with TNBC develop their disease earlier in life compared with other breast cancer subtypes and are often diagnoses at the pre-menopausal stage (Carey et al., 2006). Triple-negative breast cancer shows an increased propensity of recurrence after treatment and seem to be more aggressive than other breast carcinoma subtypes (Nofech-Mozes et al., 2009), similar to those of the basal-like breast cancer subtype. Consequently, the overall five-year survival of TNBC patients is significantly lower than those diagnosed with other subtypes of breast cancer. There is currently no acceptable specific molecular marker for TNBC. Despite this lack, these tumors do respond to chemotherapy (Krieger et al., 2009). Patients have shown better response to cytotoxic agents in the adjuvant setting as well as in the neoadjuvant setting when administered agents such as 5-fluorouracil, doxorubicin and cyclophosphamide (Rouzier et al. 2005). Other agents that have shown some efficacy include platinum based compounds

such as cisplatin and anti-tubulin compounds such as taxanes (Santana-Davila and Perez, 2010).

**[0004]** As mentioned above, there are no specific targets for TNBC but this has not impeded the trial of target agents such as the inhibition of Poly [ADP-ribose] polymerase 1 (PARP1). PARP1 is an enzyme that participates in the repair of DNA single-strand breaks by associating with corrupted DNA strands and mediating the recruitment of enzymes needed to repair single-strand breaks (de Ruijter et al., 2011). Thus the strategy has been to inhibit PARP1 activity as a means of allowing cancer cells to accumulate more DNA single-strand breaks, which ultimately leads to genetic instability, mitotic arrest and apoptosis. Promising clinical results were achieved in patients that showed mutations in *BRCA1* and/or *BRCA2*, important mediators of genetic maintenance and homologous recombination required for proper cell division. Indeed, patients with *BRCA1* mutations, which are presumably deficient in these genetic stability pathways, showed greater response to PARP1 inhibitors compared with those who were wild type for *BRCA1* (Fong et al., 2009). It is clear that targeting PARP1 in TNBC patients who are carriers of *BRCA* mutation represents a promising strategy. The combination of ER/PgR/HER2 status with that of the genetic profile of the *BRCA1/2* genes might offer the best characterization for deciding the proper treatment options for TNBC patients.

**[0005]** Other strategies also examined the use of EGFR inhibitors, either as monoclonal antibodies or small molecule inhibitors or anti-angiogenic compounds to target VEGF. Several clinical trials have evaluated the efficacy of these compounds but none of them have shown significant response when administered alone. However, mild efficacy was observed in patients treated with these inhibitors in combination with other cytotoxic agents (Santana-Davila and Perez, 2010).

**[0006]** Notwithstanding the recent advances in the understanding and the treatment for breast cancer, the use of chemotherapy is invariably associated with severe adverse reactions, which limit their use. Consequently, the need for more specific strategies such as combining antigen tissue specificity with the selectivity of monoclonal antibodies should permit a significant reduction in off-target-associated side effects. There are no TNBC specific antigens that are currently under investigation as therapeutic targets for monoclonal antibodies. Thus, TNBC patients have little options because of the inability to target a specific marker of protein that is expressed in these tumors. There are urgent needs to identify new proteins expressed in TNBC for applications as new diagnostic markers and novel targeted therapies.

**[0007]** Kidney associated antigen 1 (KAAG1), the protein sequence which is identified herein as SEQ ID NO.:2, was originally cloned from a cDNA library derived from a histocompatibility leukocyte antigen-B7 renal carcinoma cell line as an antigenic peptide presented to cytotoxic T lymphocytes (Van den Eynde et al., 1999; Genebank accession no. Q9UBP8, the cDNA sequence is represented by nucleotides 738-992 of SEQ ID NO.:1). The locus containing *KAAG1* was found to encode two genes transcribed in both directions on opposite strands. The sense strand was found to encode a transcript that encodes a protein termed DCDC2. Expression studies by these authors found that the *KAAG1* antisense transcript was tumor

specific and exhibited very little expression in normal tissues whereas the *DCDC2* sense transcript was ubiquitously expressed (Van den Eynde et al., 1999). The expression of the *KAAG1* transcript in cancer, and in particular ovarian cancer, renal cancer, lung cancer, colon cancer, breast cancer and melanoma was disclosed in international application No. PCT/CA2007/001134 published on December 27, 2007 under No. WO 2007/147265. Van den Eynde et al., also observed RNA expression in renal carcinomas, colorectal carcinomas, melanomas, sarcomas, leukemias, brain tumors, thyroid tumors, mammary carcinomas, prostatic carcinomas, oesophageal carcinomas, bladder tumor, lung carcinomas and head and neck tumors. Recently, strong genetic evidence obtained through linkage disequilibrium studies found that the *VMP/DCDC2/KAAG1* locus was associated with dyslexia (Schumacher et al., 2006; Cope et al., 2005). One of these reports pointed to the *DCDC2* marker as the culprit in dyslexic patients since the function of this protein in cortical neuron migration was in accordance with symptoms of these patients who often display abnormal neuronal migration and maturation (Schumacher et al., 2006).

**[0008]** The Applicant has obtained a panel of antibodies and antigen binding fragment that bind to the *KAAG1* protein. These antibodies or antigen binding fragments were shown to target three regions of the protein; amino acids 1 to 35, amino acids 36 to 60 amino acids 61 to 84. The Applicant found that antibodies targeting a region between amino acids 30 to 84 were the most advantageous for therapeutic purposes as they recognized *KAAG1* located at the surface of tumor cells. The Applicant has shown that some of these antibodies and antigen binding fragments can mediate antibody-dependent cell cytotoxicity and/or are internalized by tumor cells, which makes them good candidates to deliver a payload to tumor cells. The Applicant has also generated chimeric and humanized antibodies based on selected antibody candidates and has shown that these antibodies can inhibit tumor cell formation and invasion (see PCT/CA2009/001586 published on June 3, 2010 under No. WO2010/060186 and PCT/CA2010/001785 published on May 12, 2011 under No. WO2011/054112). US 2011/0223107 also describes monoclonal antibodies that specifically bind to *KAAG1*. Finally, the Applicant found that these antibodies could be used for the treatment and diagnosis of ovarian cancer, skin cancer, renal cancer, colorectal cancer, sarcoma, leukemia, brain tumor, thyroid tumor, breast cancer, prostate cancer, oesophageal tumor, bladder tumor, lung tumor and head and neck tumor and metastatic form of these cancers.

**[0009]** The Applicant has now come to the unexpected discovery that breast cancer cells lacking ER protein expression, PgR protein expression and/or showing absence of HER2 protein over-expression (*i.e.*, triple-negative breast cancer cells, basal-like) can be efficiently targeted with an antibody or antigen binding fragment that specifically binds to *KAAG1*. Anti-*KAAG1* antibodies may thus be used for the, detection and therapeutic treatment of breast cancer cells that are negative for at least one of these markers.

#### BRIEF DESCRIPTION OF THE DRAWINGS

**[0010]**

**Figure 1a** is an amino acid sequence alignment of the 3A4 variable domains of the murine and humanized light chains. The light chain has two humanized variants (Lh1 an Lh2). The CDRs are shown in bold and indicated by CDRL1, CDRL2 and CDRL3. Back mutations in the human framework regions that are murine amino acids are underlined in the humanized sequences.

**Figure 1b** is an amino acid sequence alignment of the 3A4 variable domains of the murine and humanized heavy chains. The heavy chain has four humanized variants (Hh1 to Hh4). The CDRs are shown in bold and indicated by CDRH1, CDRH2 and CDRH3. Back mutations in the human framework regions that are murine amino acids are underlined in the humanized sequences.

**Figure 2a** is an alignment of murine 3A4 light chain variable region (SEQ ID NO.:4) with a light chain variable region variant (SEQ ID NO.:189) using the ClustalW2 program (Larkin M.A., et al., (2007) ClustalW and ClustalX version 2. Bioinformatics 2007 23(21): 2947-2948) where an "\*" (asterisk) indicates positions which have a single, fully conserved residue, wherein ":" (colon) indicates conservation between groups of strongly similar properties - scoring > 0.5 in the Gonnet PAM 250 matrix and where "." (period) indicates conservation between groups of weakly similar properties - scoring <= 0.5 in the Gonnet PAM 250 matrix.

**Figure 2b** is an alignment of murine 3A4 heavy chain variable region (SEQ ID NO.:2) with a light chain variable region variant (SEQ ID NO.:194) using the ClustalW2 program (Larkin M.A., et al., (2007) ClustalW and ClustalX version 2. Bioinformatics 2007 23(21): 2947-2948) where an "\*" (asterisk) indicates positions which have a single, fully conserved residue, wherein ":" (colon) indicates conservation between groups of strongly similar properties - scoring > 0.5 in the Gonnet PAM 250 matrix and where "." (period) indicates conservation between groups of weakly similar properties - scoring <= 0.5 in the Gonnet PAM 250 matrix.

**Figure 3a** represents plasmid map of pKCR5-3A4-HC-Variant 1. The heavy chains of the humanized 3A4 variants were cloned in the same manner into the *Hind*III site of pK-CR5. Consequently the resulting plasmids are identical to pKCR5-3A4-HC variant 1 except for the sequence of the heavy chain immunoglobulin variable domain.

**Figure 3b** represents plasmid map of pMPG-CR5-3A4-LC-Variant 1. The light chains of the humanized variants 1 and 2 of 3A4 antibody were cloned in the same manner into the *Bam*HI site of pMPG-CR5. Consequently, the resulting plasmid is identical to pMPG-CR5-3A4-LC-Variant 1, except for the sequence of the light chain immunoglobulin variable domain.

**Figure 4** represents an analysis of antibody production after transient transfection in CHO cells. Supernatant (13 days post-transfection) of CHOcTA cells transfected with the different combinations of light and heavy chains of humanized 3A4 antibody were analyzed by western blot. Quantification of antibody produced in the supernatants was determined after scanning the bands of the western blot against dilution of a known standard (human purified IgG antibody). Mr molecular weight marker (kDa).

**Figure 5** is a graph of a Superdex G75 gel filtration of recombinant KAAG1 sample. KAAG1 was injected over the gel filtration and separated at 0.4 ml/min. The largest peak between

fractions 15 - 19.

**Figure 6** is a Table listing the rate and affinity constants for the murine and humanized variants of the 3A4 antibody.

**Figure 7a** is an histogram illustrating the association rates ( $K_a$ ) of the humanized antibodies.

**Figure 7b** is an histogram illustrating the dissociation rates ( $K_d$ ) of the humanized antibodies.

**Figure 7c** is an histogram illustrating the affinity constants ( $K_D$ ) of the humanized antibodies.

**Figure 8a** illustrates humanized 3A4 variants binding to KAAG1 in an ELISA. This figure shows the comparative binding of 3A4 humanized antibody variants and the murine 3A4. Concentration-dependent binding profiles of the humanized heavy chains (Hh1, Hh2, Hh3 and Hh4) assembled with the Lh1 light chain variant.

**Figure 8b** illustrates humanized 3A4 variants binding to KAAG1 in an ELISA. This figure shows the comparative binding of 3A4 humanized antibody variants and the murine 3A4. Concentration-dependent binding profiles of the humanized heavy chains (Hh1, Hh2, Hh3 and Hh4) assembled with the Lh2 light chain variant.

**Figure 9** illustrates humanized 3A4 variants binding to KAAG1 on the surface of cancer cells. This illustration shows the comparative binding activity of the humanized and the murine 3A4 antibodies on the unpermeabilized SKOV-3 ovarian cancer cells.

**Figure 10** shows a scan of a tissue microarray containing 139 biopsy samples obtained from breast cancer patients. The samples were blotted with the 3A4 anti-KAAG1 antibody and showed that the vast majority of the breast tumors expressed very high level of KAAG1 antigen. The confirmed TNBC samples are marked with an asterisk.

**Figure 11** shows the results of flow cytometry performed using MDA-MB-231, MDA-MB-436, MDA-MB-468, BT-20, BT-549, T47D, MCF-7 and 293-6E cell lines incubated with the 3A4 anti-KAAG1 antibody (blue bars of the histogram) compared with a control IgG (red bars). This is a representative results from an experiment that was performed in triplicate. The TNBC cell lines are marked with an asterisk.

**Figure 12** represents the detection of the KAAG1 antigen on the surface of MDA-MB-231 cells by flow cytometry with the 3A4 anti-KAAG1 antibody. The fluorescence signal decreases with time when the cells were incubated at 37°C, which suggests that the KAAG1/antibody complex was internalized during the incubation when the cells were incubated with 3A4.

**Figure 13** represents the detection of the KAAG1 antigen on the surface of MDA-MB-436 cells by flow cytometry with the 3A4 anti-KAAG1 antibody. The fluorescence signal decreases with time when the cells were incubated at 37°C, which suggests that the KAAG1/antibody complex was internalized during the incubation when the cells were incubated with 3A4.

**Figure 14** represents the detection of the KAAG1 antigen on the surface of BT-20 cells by flow cytometry with the 3A4 anti-KAAG1 antibody. The fluorescence signal decreases with time

when the cells were incubated at 37°C, which suggests that the KAAG1/antibody complex was internalized during the incubation when the cells were incubated with 3A4.

**Figure 15** represents the detection of the KAAG1 antigen on the surface of T47D cells by flow cytometry with the 3A4 anti-KAAG1 antibody. The fluorescence signal decreases with time when the cells were incubated at 37°C, which suggests that the KAAG1/antibody complex was internalized during the incubation when the cells were incubated with 3A4.

**Figure 16** represents immunofluorescence data performed on live MDA-MB-231 cells with the 3A4 anti-KAAG1 antibody and the anti-LAMP1 antibody. The immunofluorescence signal associated with the anti-KAAG1 antibody is shown in the left panel, the immunofluorescence signal associated LAMP1 is shown in the middle panel and the merging of both images is shown in the right panel. These data illustrates the co-localization of KAAG1 and LAMP1 near the peri-nuclear area.

**Figure 17** represents immunofluorescence data performed on live MDA-MB-231 cells with the 3A4 anti-KAAG1 antibody and the anti-LAMP1 antibody. The immunofluorescence signal associated with the anti-KAAG1 antibody is shown in the left panel, the immunofluorescence signal associated LAMP1 is shown in the middle panel and the merging of both images is shown in the right panel. These data illustrates the localization of KAAG1 with LAMP1 a marker of late endosomes/lysosomes.

## SUMMARY OF THE INVENTION

**[0011]** The present invention provides an antibody or antigen binding fragment thereof which is capable of specific binding to Kidney associated antigen 1 (KAAG1) and is conjugated with a therapeutic moiety for use in the treatment of triple negative breast cancer, wherein the antibody or antigen binding fragment thereof comprises a CDRH1 as set forth in SEQ ID NO.:49, a CDRH2 as set forth in SEQ ID NO.:50 or in SEQ ID NO.:212, a CDRH3 as set forth in SEQ ID NO.:51, a CDRL1 as set forth in SEQ ID NO.: 52, a CDRL2 as set forth in SEQ ID NO.:53 and a CDRL3 as set forth in SEQ ID NO.: 54.

**[0012]** Preferred embodiments of the invention are set forth in the claims.

**[0013]** Disclosed herein is a method of treating or detecting cancer or cancer cells (*in vitro* or *in vivo*) in an individual in need.

**[0014]** In accordance with the present invention, treatment is carried out with an antibody capable of binding to KAAG1 or an antigen binding fragment thereof that is conjugated with a therapeutic moiety.

**[0015]** The individual in need may comprise, for example, an individual having or suspected of

having cancer. Such individual may have a cancer or cancer cells originating from a breast carcinoma.

**[0016]** The cancer or cancer cells may more particularly originate from a breast carcinoma characterized as being triple-negative or basal-like.

**[0017]** Therefore, the individuals who may benefit from methods of treatment or detection described herein may include those suffering from breast carcinoma.

**[0018]** The breast carcinoma may comprise tumors cells showing a decrease or a lost in the expression of the estrogen receptor.

**[0019]** The breast carcinoma may comprise tumor cells showing a decrease or a lost in the expression of the progesterone receptor.

**[0020]** The breast carcinoma may comprise tumor cells showing a decrease or a lost in the expression of Her2.

**[0021]** The breast carcinoma may comprise tumor cells showing a decrease or a lost in Her2 overexpression.

**[0022]** More particularly, the breast carcinoma may comprise tumor cells showing either 1) a decrease or a loss in expression of the estrogen receptor and the progesterone receptor, 2) a decrease or a loss in expression of the estrogen receptor and a decrease or a loss of Her2 overexpression, 3) a decrease or a loss in expression of the progesterone receptor and a decrease or a loss of Her2 overexpression or 4) a decrease or a loss in expression of the estrogen receptor, a decrease or a loss in expression of the progesterone receptor and a decrease or a loss of Her2 overexpression.

**[0023]** Even more particularly, the breast carcinoma may comprise tumor cells showing either 1) a loss in expression of the estrogen receptor and the progesterone receptor, 2) a loss in expression of the estrogen receptor and a loss of Her2 expression, 3) a loss in expression of the progesterone receptor and a loss of Her2 expression or 4) a loss in expression of the estrogen receptor, a loss in expression of the progesterone receptor and a loss of Her2 expression.

**[0024]** In accordance with the present invention, the individual may carry breast cancer cells that are characterized as being triple-negative or may have a tumor categorized as being a triple-negative breast cancer.

**[0025]** In accordance with the present invention, the individual may carry breast cancer cells that are characterized as basal-like, or may have a tumor categorized as being a basal-like breast cancer.

**[0026]** Other individuals who would benefit from treatment with an anti-KAAG1 include those having carcinoma comprising tumors cells exhibiting an epithelial-to-mesenchymal transition (EMT) phenotype.

**[0027]** Commonly used molecular markers of EMT include, for example, a reduced expression of E-cadherin, cytokeratin and  $\beta$ -catenin (in the membrane) and/or an increased expression of Snail, Slug, Twist, ZEB1, ZEB2, N-cadherin, vimentin,  $\alpha$ -smooth muscle actin, matrix metalloproteinases etc. (see for example, Kalluri and Weinberg, *The Journal of Clinical Investigation*, 119(6), p1420-1428; 2009; Fassina et al., *Modern Pathology*, 25; p86-99; 2012; Lee et al., *JCB*; 172; p973-981; 2006). An EMT phenotype may also be distinguished by an increased capacity for migration, invasion or by resistance to anoikis/apoptosis. Cells that are undergoing epithelial-to-mesenchymal transition may thus be detected by a reduction of epithelial markers and apparition of mesenchymal markers or EMT phenotypes.

**[0028]** In accordance with the present invention, the therapeutic use may thus comprise, for example, administering an antibody or antigen binding fragment which is capable of specific binding to KAAG1 to an individual in need. The individual in need is preferentially selected on the basis of their tumor lacking ER expression, PgR expression and/or by the absence of HER2 protein over-expression. Clinical testing for these markers is usually performed using histopathologic methods (immunohistochemistry, FISH, etc.) and/or by gene expression studies (see for example Dent et al, 2007, Bernstein and Lacey, 2011). The individual in need may thus be an individual who has received a diagnosis of triple-negative breast cancer or basal-like breast cancer. The individual in need may be an individual which is unresponsive to hormonal therapy and/or to trastuzumab therapy (or other anti-Her2 antibodies). Alternatively, the individual in need may be an individual carrying tumor cells that have the ability of undergoing epithelial-to-mesenchymal transition or that have acquired a mesenchymal phenotype.

**[0029]** Disclosed herein is a method of treating triple-negative breast cancer or basal-like breast cancer by administering an inhibitor of KAAG1 activity or expression to an individual in need.

**[0030]** In accordance with the present disclosure, the KAAG1 inhibitor may thus comprise an antibody described herein or an antigen binding fragment thereof.

**[0031]** Also in accordance with the present disclosure, the KAAG1 inhibitor may comprise a nucleotide sequence complementary to SEQ ID NO.:1 or to a fragment thereof. More particularly, the KAAG1 inhibitor may comprise a nucleotide sequence complementary to nucleotides 738 to 992 (inclusively) of SEQ ID NO.:1 or to a fragment thereof. For example, the inhibitor may include at least 10 consecutive nucleotides (at least 15, at least 20) which are complementary to SEQ ID NO.:1 or to nucleotides 738 to 992 (inclusively) of SEQ ID NO.:1. More particular type of KAAG1 inhibitor includes a siRNA which inhibit expression of SEQ ID NO.:1.

**[0032]** Suitable antibodies or antigen binding fragments include those that are capable of binding to KAAG1 at the surface of tumor cells. Such antibodies or antigen binding fragments thereof may preferentially bind an epitope included within amino acids 30 to 84 of KAAG1 inclusively.

**[0033]** Alternatively such antibodies or antigen binding fragments thereof may bind an epitope located within amino acids 36 to 60 (inclusively) or within amino acids 61 to 84 (inclusively) of KAAG1.

**[0034]** The epitope may particularly be located or comprised within amino acids 50 to 70, 50 to 65, 51 to 65, 52 to 65, 53 to 65, 54 to 65, 54 to 64, 54 to 63, 54 to 62, 54 to 61, 54 to 60, 50 to 62; 50 to 61, or 50 to 60 (inclusively or exclusively).

**[0035]** In accordance with an embodiment of the invention, the antibody or antigen binding fragment may bind an epitope comprised within amino acids 50 to 70 of KAAG1.

**[0036]** In a further embodiment of the invention, the antibody or antigen binding fragment may bind an epitope comprised within amino acids 50 to 62 of KAAG1.

**[0037]** In yet a further embodiment, the antibody or antigen binding fragment may bind an epitope comprised within amino acids 54 to 65 of KAAG1.

**[0038]** Suitable antibodies for therapeutic treatment include for example, those which mediate antibody-dependent cell cytotoxicity.

**[0039]** Other even more suitable antibodies for therapeutic treatment include those that are conjugated with a therapeutic moiety.

**[0040]** In accordance with the present invention, the antibody may be, for example, a monoclonal antibody, a chimeric antibody, a humanized antibody a human antibody or an antigen binding fragment thereof.

## **DETAILED DESCRIPTION OF THE INVENTION**

### **Therapeutic use**

**[0041]** As indicated herein, the present invention encompass administering an antibody or antigen binding fragment to an individual having a breast cancer characterized as being "triple negative breast cancer". The breast cancer may be "basal-like breast cancer".

**[0042]** Classification of breast cancer subtypes as being "triple negative breast cancer" or

"basal-like breast cancer" is known in the art (see for example, Foulkes et al., *N. Engl. J. Med.*, 2010; 363:1938-1948) and includes, for example, the following definitions:

"Basal-like breast cancer", may include for example, a subtype of breast cancer comprising a heterogenous group of tumors characterized by the absence of or low levels of expression of estrogen receptors, very low prevalence of Her2 overexpression and expression of genes usually found in the basal or myoepithelial cells of the human breast. Such expression may be determined by microarray analysis.

"Triple-negative breast cancer", may include for example, a tumor characterized by lack of estrogen receptor (ER), progesterone receptor (PR) and Her2 expression. Some investigators accept tumors as being negative for expression of ER or PR only if less than 1% of the cells are positive for ER or PR expression; others consider tumors to be negative for ER or PR expression when up to 10% of cells are positive for expression. Different definitions of HER2-negativity have been used. The two most frequently adopted include tumors with immunohistochemical scores of 0/1 + or 2+ that are lacking HER2 gene amplification after *in situ* hybridization. Such expression may be especially determined by immunohistochemical staining.

**[0043]** Disclosed herein is a method of treatment that includes administering a KAAG1 inhibitor to an individual in need. Such KAAG1 inhibitor includes, for example, an antibody or antigen binding fragment thereof which specifically binds to KAAG1.

**[0044]** It is likely that the most potent antibodies or antigen binding fragments may be those having a high affinity for KAAG1. It is also likely that the most potent antibodies or antigen binding fragments may be those that are internalized within a cells compartment such as, for example, a lysosome or an endosome.

**[0045]** As such, the present invention especially encompasses antibodies or antigen binding fragments having a high affinity for KAAG1.

**[0046]** Suitable antibodies or antigen binding fragments include those that are capable of binding to KAAG1 at the surface of tumor cells with a high affinity. Such high affinity antibodies or antigen binding fragments thereof may preferentially bind an epitope included within amino acids 30 to 84 of KAAG1 inclusively.

**[0047]** Alternatively such high affinity antibodies or antigen binding fragments thereof may bind an epitope located within amino acids 36 to 60 (inclusively) or within amino acids 61 to 84 (inclusively) of KAAG1.

**[0048]** The high affinity antibodies or antigen binding fragments may bind, for example, an epitope may particularly be located or comprised within amino acids 50 to 70, 50 to 65, 51 to 65, 52 to 65, 53 to 65, 54 to 65, 54 to 64, 54 to 63, 54 to 62, 54 to 61, 54 to 60, 50 to 62; 50 to

61, or 50 to 60 (inclusively or exclusively).

**[0049]** In accordance with an embodiment of the invention, the high affinity antibody or antigen binding fragment may bind an epitope comprised within amino acids 50 to 70 of KAAG1.

**[0050]** In a further embodiment of the invention, the high affinity antibody or antigen binding fragment may bind an epitope comprised within amino acids 50 to 62 of KAAG1.

**[0051]** In yet a further embodiment, the high affinity antibody or antigen binding fragment may bind an epitope comprised within amino acids 54 to 65 of KAAG1.

**[0052]** Preferred antibodies including high affinity antibodies are those than may be internalized in a cell or cell compartment (e.g., lysosomes or endosomes). The ability of antibodies to be internalized may be determined by method known in the art such as for example and without limitation, by immunofluorescence studies similar to those performed herein.

**[0053]** The present invention relates to antibodies having CDRs identical to those of the 3A4. As such, antibodies having a light chain variable region and/or heavy chain variable region consensus sequences set forth in any of SEQ ID NOs.:186 to 188 and 191 to 193 and specific sequences set forth in SEQ ID No.:46, 48, 189, 190, or 194 to 198 are encompassed by the present invention. Among those, antibodies having a light chain variable region and/or heavy chain variable region consensus sequences set forth in any of SEQ ID NO.: 188 and 196 or specific sequences set forth in SEQ ID NO.:46, 48, 189, 190, or 194 to 198 are particularly contemplated.

**[0054]** In accordance with the present invention antibodies or antigen binding fragments thereof are conjugated with a therapeutic moiety.

**[0055]** The antibodies or antigen binding fragments thereof, may have a human constant region. Preferably the antibodies or antigen binding fragments thereof may have a human IgG1 constant region. Alternatively, the antibodies or antigen binding fragments thereof may have an IgG2 constant region.

**[0056]** The antibody for use of the present invention may also include administering a KAAG1 antibody conjugated with a therapeutic moiety or antigen binding fragment conjugated with a therapeutic moiety in combination with an anticancer agent such as for example, a small molecule drug, an antibody or antigen binding fragment binding to a target other than KAAG1, a chemotherapeutic or a cytotoxic agent. Example of anticancer agent that could be administered with the KAAG1 antibody may include for example, doxorubicin, taxanes, anti-angiogenic agents, platinum salts, PARP inhibitors.

**[0057]** Other methods of treatment disclosed herein include administering other types of KAAG1 inhibitors such as antisense-based therapeutics (siRNA, antisenses, ribozymes, etc.).

**Antibodies and antigen binding fragments that binds to KAAG1**

**[0058]** The term "antibody or antigen binding fragment" or similar terms such as "antibodies and antigen binding fragments" encompasses, for example "variant antibody or antigen binding fragment" such as, for example, "humanized antibody or antigen binding fragment".

**[0059]** The term "antibody" refers to intact antibody, monoclonal or polyclonal antibodies. The term "antibody" also encompasses multispecific antibodies such as bispecific antibodies. Human antibodies are usually made of two light chains and two heavy chains each comprising variable regions and constant regions. The light chain variable region comprises 3 CDRs, identified herein as CDRL1, CDRL2 and CDRL3 flanked by framework regions. The heavy chain variable region comprises 3 CDRs, identified herein as CDRH1, CDRH2 and CDRH3 flanked by framework regions.

**[0060]** The term "antigen-binding fragment", as used herein, refers to one or more fragments of an antibody that retain the ability to bind to an antigen (e.g., KAAG1, secreted form of KAAG1 or variants thereof). It has been shown that the antigen-binding function of an antibody can be performed by fragments of an intact antibody. Examples of binding fragments encompassed within the term "antigen-binding fragment" of an antibody include (i) a Fab fragment, a monovalent fragment consisting of the  $V_L$ ,  $V_H$ ,  $C_L$  and  $C_H1$  domains; (ii) a  $F(ab')_2$  fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the  $V_H$  and  $C_H1$  domains; (iv) a Fv fragment consisting of the  $V_L$  and  $V_H$  domains of a single arm of an antibody, (v) a dAb fragment (Ward et al., (1989) *Nature* 341:544-546), which consists of a  $V_H$  domain; and (vi) an isolated complementarity determining region (CDR), e.g.,  $V_H$  CDR3. Furthermore, although the two domains of the Fv fragment,  $V_L$  and  $V_H$ , 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 polypeptide chain in which the  $V_L$  and  $V_H$  regions pair to form monovalent molecules (known as single chain Fv (scFv); see e.g., Bird et al. (1988) *Science* 242:423-426; and Huston et al. (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antigen-binding fragment" of an antibody. Furthermore, the antigen-binding fragments include binding-domain immunoglobulin fusion proteins comprising (i) a binding domain polypeptide (such as a heavy chain variable region, a light chain variable region, or a heavy chain variable region fused to a light chain variable region via a linker peptide) that is fused to an immunoglobulin hinge region polypeptide, (ii) an immunoglobulin heavy chain CH2 constant region fused to the hinge region, and (iii) an immunoglobulin heavy chain CH3 constant region fused to the CH2 constant region. The hinge region may be modified by replacing one or more cysteine residues with serine residues so as to prevent dimerization. Such binding-domain immunoglobulin fusion proteins are further disclosed in US 2003/0118592 and US 2003/0133939. These antibody fragments are obtained using conventional techniques known to those with skill in the art, and the fragments are

screened for utility in the same manner as are intact antibodies.

**[0061]** A typical antigen binding site is comprised of the variable regions formed by the pairing of a light chain immunoglobulin and a heavy chain immunoglobulin. The structure of the antibody variable regions is very consistent and exhibits very similar structures. These variable regions are typically comprised of relatively homologous framework regions (FR) interspersed with three hypervariable regions termed Complementarity Determining Regions (CDRs). The overall binding activity of the antigen binding fragment is often dictated by the sequence of the CDRs. The FRs often play a role in the proper positioning and alignment in three dimensions of the CDRs for optimal antigen binding.

**[0062]** As used herein the term "high affinity" refers to an affinity of 10nM or less. The term "high affinity" especially includes antibodies having an affinity of 5nM or less. The term "high affinity" even more particularly includes antibodies having an affinity of 1 nM or less, or 0.1nM or less.

**[0063]** Antibodies and/or antigen binding fragments of the present invention may originate, for example, from a mouse, a rat or any other mammal or from other sources such as through recombinant DNA technologies.

**[0064]** An-KAAG1 antibodies were initially isolated from Fab libraries for their specificity towards the antigen of interest. Exemplary methods on how to convert Fab into full immunoglobulins are provided herein.

**[0065]** The variable regions described herein may be fused with constant regions of a desired species thereby allowing recognition of the antibody by effector cells of the desired species. The constant region may originate, for example, from an IgG1, IgG2, IgG3, or IgG4 subtype. Cloning or synthesizing a constant region in frame with a variable region is well within the scope of a person of skill in the art and may be performed, for example, by recombinant DNA technology.

**[0066]** In certain embodiments of the present invention, antibodies that bind to KAAG1 may be of the IgG1, IgG2, IgG3, or IgG4 subtype. More specific embodiments of the invention relates to an antibody of the IgG1 subtype or especially human IgG1 subtype. Other specific embodiments of the invention relates to an antibody of the IgG2 subtype or especially of the human IgG2 subtype.

**[0067]** The antibody may be a humanized antibody of the IgG1 subtype subtype or especially human IgG1 subtype. Alternatively, the antibody may be a humanized antibody of the IgG2 subtype or especially of the human IgG2 subtype.

**[0068]** The antibody may be, for example, biologically active in mediating antibody-dependent cellular cytotoxicity (ADCC), complement-mediated cytotoxicity (CMC), or associated with immune complexes. The typical ADCC involves activation of natural killer (NK) cells and is

reliant on the recognition of antibody-coated cells by Fc receptors on the surface of the NK cells. The Fc receptors recognize the Fc domain of antibodies such as is present on IgG1, which bind to the surface of a target cell, in particular a cancerous cell that expresses an antigen, such as KAAG1. Once bound to the Fc receptor of IgG the NK cell releases cytokines and cytotoxic granules that enter the target cell and promote cell death by triggering apoptosis.

**[0069]** Described herein is a collection of antibodies that bind to KAAG1 or to a KAAG1 variant. In certain embodiments, the antibodies may be selected from the group consisting of polyclonal antibodies, monoclonal antibodies such as chimeric or humanized antibodies, antibody fragments such as antigen binding fragments, single chain antibodies, domain antibodies, and polypeptides with an antigen binding region.

**[0070]** In an aspect of the invention, the isolated antibody or antigen binding fragment of the present invention may be capable of inducing killing (elimination, destruction, lysis) of KAAG1-expressing tumor cells or KAAG1 variant-expressing tumor cells (e.g., in an ADCC-dependent manner).

**[0071]** In a further aspect of the invention, the isolated antibody or antigen binding fragment of the present invention may especially be characterized by its capacity of reducing spreading of tumor cells expressing KAAG1 or a KAAG1 variant.

**[0072]** In an additional aspect of the invention, the isolated antibody or antigen binding fragment of the present invention may be characterized by its capacity of decreasing or impairing formation of tumors expressing KAAG1 or a KAAG1 variant.

**[0073]** In an exemplary embodiment of the invention, the isolated antibody or antigen binding fragment may comprise amino acids of a constant region, which may originate, for example, from a human antibody.

**[0074]** In another exemplary embodiment of the invention, the isolated antibody or antigen binding fragment may comprise framework amino acids of a human antibody.

**[0075]** The Applicant has generated specific antibodies and antigen binding fragments that may be useful for the purposes described herein.

**[0076]** The following is a list of antibodies that were generated and shown to bind in a specific manner to KAAG1; 3D3, 3A4, 3C4, 3G10, 3A2, 3F6, 3E8, 3E10, 3A9, 3B1, 3G5, 3B2, 3B8, 3G8, 3F7, 3E9, 3G12, 3C3, 3E12, 4A2, 3F10, 3F4, 3B11, 3D1, 3C2, 3E6 and 3H3. Sequences of the antibody light chain or heavy chain, variable regions or complementary determining regions (CDRs) are available in international application No. PCT/CA2009/001586 published on June 3, 2010 under No. WO2010/060186A8, in international application No. PCT/CA2010/001795 published on May 12, 2011 under No. WO2011/054112A1 or in international application No. PCT/CA2012/000296 published on Oct. 4, 2012 under No. WO2012/129668A1.

**[0077]** In most instances, the sequence of the CDRs has been provided separately or is shown in bold herein.

**[0078]** Amongst, these antibodies, the 3D3, 3A4, 3G10 and 3C4 were selected for *in vitro* and/or *in vivo* biological testing. The 3A4 antibody appeared to have the best characteristics. Based on our experiments, the 3A4 antibody when conjugated with a therapeutic moiety (e.g. a cytotoxic agent) is more effective in killing cancer cells than its non-conjugated version.

**[0079]** In an exemplary disclosure, the antibody or antigen binding fragment may comprise any individual CDR or a combination of CDR1, CDR2 and/or CDR3 of the light chain variable region. The CDR3 may more particularly be selected. Combination may include for example, CDRL1 and CDRL3; CDRL1 and CDRL2; CDRL2 and CDRL3 and; CDRL1, CDRL2 and CDRL3.

**[0080]** In another exemplary disclosure, the antibody or antigen binding fragment may comprise any individual CDR or a combination of CDR1, CDR2 and/or CDR3 of the heavy chain variable region. The CDR3 may more particularly be selected. Combination may include for example, CDRH1 and CDRH3; CDRH1 and CDRH2; CDRH2 and CDRH3 and; CDRH1, CDRH2 and CDRH3.

**[0081]** In accordance with the present disclosure, the antibody or antigen binding fragment may comprise at least two CDRs of a CDRL1, a CDRL2 or a CDRL3.

**[0082]** Also in accordance with the present disclosure, the antibody or antigen binding fragment may comprise one CDRL1, one CDRL2 and one CDRL3.

**[0083]** Further in accordance with the present disclosure, the antibody or antigen binding fragment may comprise:

1. a. At least two CDRs of a CDRL1, CDRL2 or CDRL3 and;
2. b. At least two CDRs of a CDRH1, one CDRH2 or one CDRH3.

**[0084]** The antibody or antigen binding fragment may more preferably comprise one CDRL1, one CDRL2 and one CDRL3.

**[0085]** The antibody or antigen binding fragment may also more preferably comprise one CDRH1, one CDRH2 and one CDRH3.

**[0086]** When only one of the light chain variable region or the heavy chain variable region is available, an antibody or antigen-binding fragment may be reconstituted by screening a library of complementary variable regions using methods known in the art (Portolano et al. The Journal of Immunology (1993) 150:880-887, Clarkson et al., Nature (1991) 352:624-628).

**[0087]** Exemplary antibodies or antigen binding fragments are those having the CDRs of the light chain and/or heavy chains of the 3D3, 3A4, 3C4, 3G10, 3A2, 3F6, 3E8, 3E10, 3A9, 3B1, 3G5, 3B2, 3B8, 3G8, 3F7, 3E9, 3G12, 3C3, 3E12, 4A2, 3F10, 3F4, 3B11, 3D1, 3C2, 3E6 or 3H3 antibodies. More particular examples include antibodies or antigen binding fragments having the CDRs of the light chain and/or heavy chains of the 3D3, 3A4, 3C4 or 3G10 antibodies. In accordance with the present invention, antibodies or antigen binding fragments have the CDRs of the light chain and heavy chain of the 3A4 antibody. The invention thus encompassed any monoclonal, chimeric, human, or humanized antibody comprising the CDRs of the 3A4 antibody.

**[0088]** Antibodies or antigen binding fragments that may be used in accordance with the present invention are those having CDRs of the 3A4 antibody and comprise, for example, a CDRH1 as set forth in SEQ ID NO.:49, a CDRH2 as set forth in SEQ ID NO.:50 or in SEQ ID NO.:212, a CDRH3 as set forth in SEQ ID NO.:51, a CDRL1 as set forth in SEQ ID NO.: 52, a CDRL2 as set forth in SEQ ID NO.:53 and a CDRL3 as set forth in SEQ ID NO.: 54.

**[0089]** The present invention therefore encompass, antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise sequences selected from the group consisting of:

- d. the 3CDRs of a light chain variable region defined in SEQ ID NO.:48 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:46;
- aa. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:194,
- bb. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:195,
- cc. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:196,
- dd. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:197,
- ee. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:194,
- ff. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:195,
- gg. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:196,
- hh. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:197.

**[0090]** Disclosed herein are antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise sequences selected from the group consisting of:

- a. the 3CDRs of a light chain variable region defined in SEQ ID NO.:16 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:18;
- b. the 3CDRs of a light chain variable region defined in SEQ ID NO.:20 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:22;
- c. the 3CDRs of a light chain variable region defined in SEQ ID NO.:24 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:26;
- d. the 3CDRs of a light chain variable region defined in SEQ ID NO.:48 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:46;
- e. the 3CDRs of a light chain variable region defined in SEQ ID NO.:103 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:126;
- f. the 3CDRs of a light chain variable region defined in SEQ ID NO.:104 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:127;
- g. the 3CDRs of a light chain variable region defined in SEQ ID NO.:105 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:128;
- h. the 3CDRs of a light chain variable region defined in SEQ ID NO.:106 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:145;
- i. the 3CDRs of a light chain variable region defined in SEQ ID NO.:107 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:129;
- j. the 3CDRs of a light chain variable region defined in SEQ ID NO.:108 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:130;
- k. the 3CDRs of a light chain variable region defined in SEQ ID NO.:109 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:141;
- l. the 3CDRs of a light chain variable region defined in SEQ ID NO.:110 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:131;
- m. the 3CDRs of a light chain variable region defined in SEQ ID NO.:111 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:134;
- n. the 3CDRs of a light chain variable region defined in SEQ ID NO.:112 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:135;
- o. the 3CDRs of a light chain variable region defined in SEQ ID NO.:113 and/or the 3CDRs of a

heavy chain variable region defined in SEQ ID NO.:136,

p. the 3CDRs of a light chain variable region defined in SEQ ID NO.:114 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:133,

q. the 3CDRs of a light chain variable region defined in SEQ ID NO.:115 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:140,

r. the 3CDRs of a light chain variable region defined in SEQ ID NO.:116 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:137,

s. the 3CDRs of a light chain variable region defined in SEQ ID NO.:117 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:144,

t. the 3CDRs of a light chain variable region defined in SEQ ID NO.:118 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:139,

u. the 3CDRs of a light chain variable region defined in SEQ ID NO.:119 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:132,

v. the 3CDRs of a light chain variable region defined in SEQ ID NO.:120 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:142,

w. the 3CDRs of a light chain variable region defined in SEQ ID NO.:121 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:138,

x. the 3CDRs of a light chain variable region defined in SEQ ID NO.:122 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:146,

y. the 3CDRs of a light chain variable region defined in SEQ ID NO.:123 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:153,

z. the 3CDRs of a light chain variable region defined in SEQ ID NO.:124 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:143,

aa. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:194,

bb. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:195,

cc. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:196,

dd. the 3CDRs of a light chain variable region defined in SEQ ID NO.:189 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:197,

ee. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:194,

ff. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:195,

gg. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:196, or

hh. the 3CDRs of a light chain variable region defined in SEQ ID NO.:190 and/or the 3CDRs of a heavy chain variable region defined in SEQ ID NO.:197.

**[0091]** Other exemplary antibodies or antigen binding fragments have the light chain and/or heavy chains of the 3D3, 3A4, 3C4, 3G10, 3A2, 3F6, 3E8, 3E10, 3A9, 3B1, 3G5, 3B2, 3B8, 3G8, 3F7, 3E9, 3G12, 3C3, 3E12, 4A2, 3F10, 3F4, 3B11, 3D1, 3C2, 3E6 or 3H3 antibodies. More particular examples include antibodies or antigen binding fragments having the light chain and/or heavy chains of the 3D3, 3A4, 3C4 or 3G10 antibodies. Even more particular embodiments of the invention include antibodies or antigen binding fragments having the light chain and/or heavy chains of the 3A4 antibody (humanized and non-humanized).

**[0092]** The present invention therefore encompass, antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise sequences selected from the group consisting of:

d. the light chain variable region defined in SEQ ID NO.:48 and/or the heavy chain variable region defined in SEQ ID NO.:46,

aa. the light chain variable region defined in SEQ ID NO.:189 and/or the heavy chain variable region defined in SEQ ID NO.:194,

bb. the light chain variable region defined in SEQ ID NO.:189 and/or the heavy chain variable region defined in SEQ ID NO.:195,

cc. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:194,

dd. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:195,

ee. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:196, or

ff. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:197.

**[0093]** Disclosed herein are antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise sequences selected from the group

consisting of:

- a. the light chain variable region defined in SEQ ID NO.:16 (encoded by SEQ ID NO.:15) and/or the heavy chain variable region defined in SEQ ID NO.:18 (encoded by SEQ ID NO.:17),
- b. the light chain variable region defined in SEQ ID NO.:20 (encoded by SEQ ID NO.:19) and/or the heavy chain variable region defined in SEQ ID NO.:22 (encoded by SEQ ID NO.:21);
- c. the light chain variable region defined in SEQ ID NO.:24 (encoded by SEQ ID NO.:23) and/or the heavy chain variable region defined in SEQ ID NO.:26 (encoded by SEQ ID NO.:25);
- d. the light chain variable region defined in SEQ ID NO.:48 and/or the heavy chain variable region defined in SEQ ID NO.:46,
- e. the light chain variable region defined in SEQ ID NO.:103 and/or the heavy chain variable region defined in SEQ ID NO.:126,
- f. the light chain variable region defined in SEQ ID NO.:104 and/or the heavy chain variable region defined in SEQ ID NO.:127,
- g. the light chain variable region defined in SEQ ID NO.:105 and/or the heavy chain variable region defined in SEQ ID NO.:128,
- h. the light chain variable region defined in SEQ ID NO.:106 and/or the heavy chain variable region defined in SEQ ID NO.:145,
- i. the light chain variable region defined in SEQ ID NO.:107 and/or the heavy chain variable region defined in SEQ ID NO.:129,
- j. the light chain variable region defined in SEQ ID NO.:108 and/or the heavy chain variable region defined in SEQ ID NO.:130,
- k. the light chain variable region defined in SEQ ID NO.:109 and/or the heavy chain variable region defined in SEQ ID NO.:141,
- l. the light chain variable region defined in SEQ ID NO.:110 and/or the heavy chain variable region defined in SEQ ID NO.:131,
- m. the light chain variable region defined in SEQ ID NO.:111 and/or the heavy chain variable region defined in SEQ ID NO.:134,
- n. the light chain variable region defined in SEQ ID NO.:112 and/or the heavy chain variable region defined in SEQ ID NO.:135,
- o. the light chain variable region defined in SEQ ID NO.:113 and/or the heavy chain variable region defined in SEQ ID NO.:140,
- p. the light chain variable region defined in SEQ ID NO.:114 and/or the heavy chain variable region defined in SEQ ID NO.:133,

- q. the light chain variable region defined in SEQ ID NO.:115 and/or the heavy chain variable region defined in SEQ ID NO.:140,
- r. the light chain variable region defined in SEQ ID NO.:116 and/or the heavy chain variable region defined in SEQ ID NO.:137,
- s. the light chain variable region defined in SEQ ID NO.:117 and/or the heavy chain variable region defined in SEQ ID NO.:144,
- t. the light chain variable region defined in SEQ ID NO.:118 and/or the heavy chain variable region defined in SEQ ID NO.:139,
- u. the light chain variable region defined in SEQ ID NO.:119 and/or the heavy chain variable region defined in SEQ ID NO.:132,
- v. the light chain variable region defined in SEQ ID NO.:120 and/or the heavy chain variable region defined in SEQ ID NO.:142,
- w. the light chain variable region defined in SEQ ID NO.:121 and/or the heavy chain variable region defined in SEQ ID NO.:138,
- x. the light chain variable region defined in SEQ ID NO.:122 and/or the heavy chain variable region defined in SEQ ID NO.:146,
- y. the light chain variable region defined in SEQ ID NO.:123 and/or the heavy chain variable region defined in SEQ ID NO.:147;
- z. the light chain variable region defined in SEQ ID NO.:124 and/or the heavy chain variable region defined in SEQ ID NO.:144;
- aa. the light chain variable region defined in SEQ ID NO.:189 and/or the heavy chain variable region defined in SEQ ID NO.:194,
- bb. the light chain variable region defined in SEQ ID NO.:189 and/or the heavy chain variable region defined in SEQ ID NO.:195,
- cc. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:194,
- dd. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:195,
- ee. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:196, or
- ff. the light chain variable region defined in SEQ ID NO.:190 and/or the heavy chain variable region defined in SEQ ID NO.:197.

**[0094]** The framework region of the heavy and/or light chains described herein may be derived from one or more of the framework regions illustrated in the antibodies described herein. The antibody or antigen binding fragments may thus comprise one or more of the CDRs described herein (e.g., selected from the specific CDRs or consensus CDRs of SEQ ID NO.:72 to 88 or CDR variants of SEQ ID NO.:89-102) and framework regions originating from those described herein. In SEQ ID Nos. 103-154, the expected CDRs are shown in bold, while the framework regions are not.

**[0095]** Table 1 refers to the complete sequences of light and heavy chain of some of the anti-KAAG1 antibodies which were selected for biological testing.

Table 1.

Antibody designation	Chain type	Nucleotide sequence (SEQ ID NO.:)	Amino acid sequence (SEQ ID NO.:)
3D3	Light (L)	3	4
3D3	Heavy (H)	5	6
3G10	Light	7	8
3G10	Heavy	9	10
3C4	Light	11	12
3C4	Heavy	13	14
Humanized 3D3	Light		166
Humanized 3D3	Heavy		167
Humanized 3C4	Light		170
Humanized 3C4	Heavy		171
Humanized 3A4	Light (Lh1)		199
Humanized 3A4	Light (Lh2)		200
Humanized 3A4	Heavy (Hh1)		202
Humanized 3A4	Heavy (Hh2)		203
Humanized 3A4	Heavy (Hh3)		204
Humanized 3A4	Heavy (Hh4)		205

**[0096]** Epitope mapping studies revealed that the 3D3 antibody interacts with a KAAG1 epitope spanned by amino acids 36 - 60, inclusively. The 3G10 and 3A4 antibodies interact with a KAAG1 epitope spanned by amino acids 61 - 84, inclusively and the 3C4 antibody interacts

with a KAAG1 epitope spanned by amino acids 1 - 35. Although, the 3G10 and 3A4 binds a similar region, the 3G10 antibody does not bind to KAAG1 as efficiently as the 3A4 antibody.

**[0097]** It is to be understood herein, that the light chain variable region of the specific combination provided above may be changed for any other light chain variable region. Similarly, the heavy chain variable region of the specific combination provided above may be changed for any other heavy chain variable region.

**[0098]** Sequences of light and heavy chain variable regions of selected antibodies that bind to KAAG1 are disclosed in Table 2.

Table 2

Ab. designation	Variable region type	Nucleotide (SEQ ID NO.:)	Amino acid (SEQ ID NO.:)
3D3	Light (VL)	15	16
3D3	Heavy (VH)	17	18
3G10	Light	19	20
3G10	Heavy	21	22
3C4	Light	23	24
3C4	Heavy	25	26
3A2	Light		103
3A2	Heavy		126
3E10	Light		106
3E10	Heavy		145
3G12	Light		121
3G12	Heavy		138
3A4	Light	47	48
3A4	Heavy	45	46
Humanized 3D3	Light		168
Humanized 3D3	Heavy		169
Humanized 3C4	Light		172
Humanized 3C4	Heavy		173
Humanized 3A4	Light (Lvh1)		189
Humanized 3A4	Light (Lvh2)		190
Humanized	Heavy (Hvh1)		194

Ab. designation	Variable region type	Nucleotide (SEQ ID NO.:)	Amino acid (SEQ ID NO.:)
3A4			
Humanized 3A4	Heavy (Hvh2)		195
Humanized 3A4	Heavy (Hvh3)		197
Humanized 3A4	Heavy (Hvh4)		198

**[0099]** SEQ ID NOs. 103-154 correspond to the light chain and heavy chain variable regions of other antibodies which were shown to bind KAAG1.

**[0100]** CDR sequence of the light and heavy chain variable regions of selected antibodies that bind to KAAG1 are disclosed in Table 3.

Table 3

Ab. designation	Chain type	CDR	SEQ ID NO.:	a.a. sequence
3D3	Light (L)	CDR L1	27	KSSQSLLNSNFQKNFLA
3D3	Light	CDR L2	28	FASTRES
3D3	Light	CDR L3	29	QQHYSTPLT
3D3	Heavy (H)	CDR H1	30	GYIFTDYEIH
3D3	Heavy	CDR H2	31	VIDPETGNTA
3D3	Heavy	CDR H3	32	MGYSDY
3G10	Light	CDR L1	33	RSSQSLLHSNGNTYLE
3G10	Light	CDR L2	34	KVSNRFS
3G10	Light	CDR L3	35	FQGSHVPLT
3G10	Heavy	CDR H1	36	GYTFTDNYMN
3G10	Heavy	CDR H2	37	DINPYYGT
3G10	Heavy	CDR	38	ARDDWFDY

Ab. designation	Chain type	CDR	SEQ ID NO.:	a.a. sequence
		H3		
3C4	Light	CDR L1	39	KASQDIHNFLN
3C4	Light	CDR L2	40	RANRLVD
3C4	Light	CDR L3	41	LQYDEIPLT
3C4	Heavy	CDR H1	42	GFSITSGYGWH
3C4	Heavy	CDR H2	43	YINYDGHND
3C4	Heavy	CDR H3	44	ASSYDGLFAY
3A2	Light	CDR L1	148	KSSQSLLHSDGKTYLN
3A2	Light	CDR L2	149	LVSKLDS
3A2	Light	CDR L3	150	WQGTHFPRT
3A2	Heavy	CDR H1	151	GYTFTD YNMH
3A2	Heavy	CDR H2	152	YINPYNDVTE
3A2	Heavy	CDR H3	153	AWFGL RQ
3E10	Light	CDR L1	154	RSSKSLLHSNGN TYLY
3E10	Light	CDR L2	155	RMSNLAS
3E10	Light	CDR L3	156	MQHLEYPYT
3E10	Heavy	CDR H1	157	GDTFTD YYMN
3E10	Heavy	CDR H2	158	DINPNYGGIT
3E10	Heavy	CDR H3	159	QAYYRNS DY
3G12	Light	CDR L1	160	KASQDVGTAVA
3G12	Light	CDR	161	WTSTRHT

Ab. designation	Chain type	CDR	SEQ ID NO.:	a.a. sequence
		L2		
3G12	Light	CDR L3	162	QQHYSIPLT
3G12	Heavy	CDR H1	163	GYIFTDYEIH
3G12	Heavy	CDR H2	164	VIDPETGNTA
3G12	Heavy	CDR H3	165	MGYSDY
3A4	Light	CDR L1	52	RSSQSLLHSNGNTYLE
3A4	Light	CDR L2	53	TVSNRFS
3A4	Light	CDR L3	54	FQGSHVPLT
3A4	Heavy	CDR H1	49	GYTFTDDYMS
3A4	Heavy	CDR H2	50 or 212	DINPYNGDTNYNQKFKG or DINPYNGDTN
3A4	Heavy	CDR H3	51	DPGAMDY

**Variant antibody and antigen binding fragments**

**[0101]** The present invention also encompasses variants of the antibodies or antigen binding fragments described herein. Variant antibodies or antigen binding fragments included are those having a variation in the amino acid sequence. For example, variant antibodies or antigen binding fragments included in the present disclosure are those having at least one variant CDR (two, three, four, five or six variant CDRs, etc. or even twelve variant CDRs). Variant antibodies or antigen binding fragments include those having a variant light chain variable region, a variant heavy chain variable region, a variant light chain and/or a variant heavy chain. Variant antibodies or antigen binding fragments included in the present invention are those having, for example, similar or improved binding affinity in comparison with the original antibody or antigen binding fragment.

**[0102]** As used herein the term "variant" applies to any of the sequence described herein and includes for example, a variant CDR (either CDRL1, CDRL2, CDRL3, CDRH1, CDRH2 and/or CDRH3), a variant light chain variable region, a variant heavy chain variable region, a variant

light chain, a variant heavy chain, a variant antibody, a variant antigen binding fragment and a KAAG1 variant.

**[0103]** The sites of greatest interest for substitutional mutagenesis include the hypervariable regions (CDRs), but modifications in the framework region or even in the constant region are also contemplated. Exemplary embodiments of CDR variants are provided in SEQ ID NOs.: 72-102.

**[0104]** Conservative substitutions may be made by exchanging an amino acid (of a CDR, variable chain, antibody, etc.) from one of the groups listed below (group 1 to 6) for another amino acid of the same group.

**[0105]** Other exemplary embodiments of conservative substitutions are shown in Table 1A under the heading of "preferred substitutions". If such substitutions result in a undesired property, then more substantial changes, denominated "exemplary substitutions" in Table 1A, or as further described below in reference to amino acid classes, may be introduced and the products screened.

**[0106]** It is known in the art that variants may be generated by substitutional mutagenesis and retain the biological activity of the polypeptides of the present invention. These variants have at least one amino acid residue in the amino acid sequence removed and a different residue inserted in its place. For example, one site of interest for substitutional mutagenesis may include a site in which particular residues obtained from various species are identical. Examples of substitutions identified as "conservative substitutions" are shown in Table 1A. If such substitutions result in a change not desired, then other type of substitutions, denominated "exemplary substitutions" in Table 1A, or as further described herein in reference to amino acid classes, are introduced and the products screened.

**[0107]** Substantial modifications in function or immunological identity 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:

(group 1) hydrophobic: norleucine, methionine (Met), Alanine (Ala), Valine (Val), Leucine (Leu), Isoleucine (Ile)

(group 2) neutral hydrophilic: Cysteine (Cys), Serine (Ser), Threonine (Thr)

(group 3) acidic: Aspartic acid (Asp), Glutamic acid (Glu)

(group 4) basic: Asparagine (Asn), Glutamine (Gln), Histidine (His), Lysine (Lys), Arginine (Arg)

(group 5) residues that influence chain orientation: Glycine (Gly), Proline (Pro); and

(group 6) aromatic: Tryptophan (Trp), Tyrosine (Tyr), Phenylalanine (Phe)

**[0108]** Non-conservative substitutions will entail exchanging a member of one of these classes for another.

Table 1A. Amino acid substitution

Original residue	Exemplary substitution	Conservative substitution
Ala (A)	Val, Leu, Ile	Val
Arg (R)	Lys, Gln, Asn	Lys
Asn (N)	Gln, His, Lys, Arg, Asp	Gln
Asp (D)	Glu, Asn	Glu
Cys (C)	Ser, Ala	Ser
Gln (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, norleucine	Leu
Leu (L)	Norleucine, Ile, Val, Met, Ala, Phe	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, Norleucine	Leu

**[0109]** Variation in the amino acid sequence of the variant antibody or antigen binding fragment may include an amino acid addition, deletion, insertion, substitution etc., one or more modification in the backbone or side-chain of one or more amino acid, or an addition of a group or another molecule to one or more amino acids (side-chains or backbone).

**[0110]** Variant antibody or antigen binding fragment may have substantial sequence similarity and/or sequence identity in its amino acid sequence in comparison with that the original antibody or antigen binding fragment amino acid sequence. The degree of similarity between two sequences is based upon the percentage of identities (identical amino acids) and of

conservative substitution.

**[0111]** Generally, the degree of similarity and identity between variable chains has been determined herein using the Blast2 sequence program (Tatiana A. Tatusova, Thomas L. Madden (1999), "Blast 2 sequences - a new tool for comparing protein and nucleotide sequences", FEMS Microbiol Lett. 174:247-250) using default settings, i.e., blastp program, BLOSUM62 matrix (open gap 11 and extension gap penalty 1; gapx dropoff 50, expect 10.0, word size 3) and activated filters.

**[0112]** Percent identity will therefore be indicative of amino acids which are identical in comparison with the original peptide and which may occupy the same or similar position. Percent similarity will be indicative of amino acids that are identical and those that are replaced with conservative amino acid substitution in comparison with the original peptide at the same or similar position.

**[0113]** Variants of the present invention therefore comprise those which may have at least 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% sequence identity with an original sequence or a portion of an original sequence.

**[0114]** Exemplary embodiments of variants are those having at least 81% sequence identity to a sequence described herein and 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0115]** Other exemplary embodiments of variants are those having at least 82% sequence identity to a sequence described herein and 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0116]** Further exemplary embodiments of variants are those having at least 85% sequence identity to a sequence described herein and 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0117]** Other exemplary embodiments of variants are those having at least 90% sequence identity to a sequence described herein and 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0118]** Additional exemplary embodiments of variants are those having at least 95% sequence identity to a sequence described herein and 95%, 96%, 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0119]** Yet additional exemplary embodiments of variants are those having at least 97% sequence identity to a sequence described herein and 97%, 98%, 99% or 100% sequence similarity with an original sequence or a portion of an original sequence.

**[0120]** For a purpose of concision the applicant provides herein a Table 1B illustrating exemplary embodiments of individual variants encompassed by the present invention and comprising the specified % sequence identity and % sequence similarity. Each "X" is to be construed as defining a given variant.

Table 1B		Percent (%) sequence identity																				
Percent t (%)	Percent s (%)	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
		80	X																			
		81	X	X																		

Table 1B		Percent (%) sequence identity																			
Percent t (%)	Percent s (%)	82	X	X	X																
		83	X	X	X	X															
84	X	X	X	X	X																
85	X	X	X	X	X	X															
86	X	X	X	X	X	X	X														
87	X	X	X	X	X	X	X	X	X												
88	X	X	X	X	X	X	X	X	X	X											
89	X	X	X	X	X	X	X	X	X	X	X										
90	X	X	X	X	X	X	X	X	X	X	X	X									
91	X	X	X	X	X	X	X	X	X	X	X	X	X								
92	X	X	X	X	X	X	X	X	X	X	X	X	X	X							
93	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X						
94	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X					
95	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X				
96	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X		
97	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X		
98	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	
99	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
100	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

**[0121]** The present disclosure encompasses CDRs, light chain variable regions, heavy chain variable regions, light chains, heavy chains, antibodies and/or antigen binding fragments which comprise at least 70% identity or at least 80% identity with the sequence described herein.

**[0122]** The present invention therefore encompass, antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise:

d. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:48 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:46;

**[0123]** Disclosed herein are antibodies and antigen binding fragment which are capable of specific binding to KAAG1 and which may comprise sequences selected from the group consisting of:

1. a. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:16 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:18,
2. b. a light chain variable region having at least 70% sequence identity with SEQ ID

- NO.:20 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:22;
- 3. c. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:24 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:26;
  - 4. d. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:48 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:46;
  - 5. e. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:103 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:126,
  - 6. f. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:104 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:127,
  - 7. g. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:105 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:128,
  - 8. h. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:106 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:145,
  - 9. i. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:107 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:128,
  - 10. j. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:108 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:130,
  - 11. k. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:109 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:141,
  - 12. l. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:110 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:131,
  - 13. m. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:111 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:134,
  - 14. n. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:112 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:135,
  - 15. o. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:113 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:136,
  - 16. p. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:114 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:133,

17. q. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:115 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:140,
18. r. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:116 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:137,
19. s. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:117 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:144,
20. t. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:118 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:139,
21. u. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:119 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:132,
22. v. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:120 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:142,
23. w. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:121 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:138,
24. x. the light chain variable region having at least 70% sequence identity with SEQ ID NO.:122 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:146,
25. y. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:123 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:147, or;
26. z. a light chain variable region having at least 70% sequence identity with SEQ ID NO.:124 and a heavy chain variable region having at least 70% sequence identity with SEQ ID NO.:143.

**[0124]** In accordance with the present invention, the variant antibodies or antigen binding fragments comprise CDRs that are identical to those of the corresponding light chain and/or heavy chain variable region. In other examples the variant antibodies or antigen binding fragments may comprise variant CDR(s).

**[0125]** Therefore, examples of a variant antibody or antigen binding fragment are those comprising a light chain variable region comprising a sequence which is at least 70%, 75%, 80% identical to SEQ ID NOS.:16, 20, 24, 103, 106 or 121. The CDRs of such variant may be identical to those of the corresponding non-variant (wild type sequence) antibody or antigen binding fragment or may vary by 1-3 amino acids.

**[0126]** Another exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:16 and having for example from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:16. A SEQ ID NO.:16 variant is provided in SEQ ID NO.:168.

**[0127]** An exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:20 and having for example from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:20.

**[0128]** An exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:24 and having for example from 1 to 21 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:24. A SEQ ID NO.:24 variant is provided in SEQ ID NO.:172.

**[0129]** An exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:103 and having for example from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:103.

**[0130]** An exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:106 and having for example from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:106.

**[0131]** An exemplary variant antibody light chain variable region encompasses a light chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:121 and having for example from 1 to 21 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:121.

**[0132]** In some instances, the variant antibody light chain variable region may comprise amino acid deletions or additions (in combination or not with amino acid substitutions). Often 1, 2, 3, 4 or 5 amino acid deletions or additions may be tolerated.

**[0133]** Other examples of a variant antibody or antigen binding fragment of the present invention are those comprising a heavy chain variable region comprising a sequence which is

at least 70%, 75%, 80% identical to 18, 22, 26, 126, 138 or 145. The CDRs of such variant may be identical to those of the corresponding non-variant (wild type sequence) antibody or antigen binding fragment or may vary by 1-3 amino acids.

**[0134]** An example of a variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:18 and having, for example, from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:18. A SEQ ID NO.:18 variant is provided in SEQ ID NO.:169.

**[0135]** An exemplary variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:22 and having, for example, from 1 to 23 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:22.

**[0136]** An exemplary variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:26 and having, for example, from 1 to 23 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:26. A SEQ ID NO.:26 variant is provided in SEQ ID NO.:173.

**[0137]** An exemplary variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:126 and having, for example, from 1 to 23 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:126.

**[0138]** An exemplary variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:145 and having, for example, from 1 to 23 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:145.

**[0139]** An exemplary variant antibody heavy chain variable region encompasses a heavy chain variable region having CDR amino acid sequences that are 100% identical to the CDR amino acid sequence of SEQ ID NO.:138 and having, for example, from 1 to 22 amino acid modifications (e.g., conservative or non-conservative amino acid substitutions) in its framework region in comparison with the framework region of SEQ ID NO.:138.

**[0140]** In some instances, the variant antibody heavy chain variable region may comprise amino acid deletions or additions (in combination or not with amino acid substitutions). Often 1,

2, 3, 4 or 5 amino acid deletions or additions may be tolerated.

**Variant CDRS**

**[0141]** Also disclosed herein are polypeptides, antibodies or antigen binding fragments comprising variable chains having at least one conservative amino acid substitution in at least one of the CDRs described herein (in comparison with the original CDR).

**[0142]** The present disclosure also encompasses polypeptides, antibodies or antigen binding fragments comprising variable chains having at least one conservative amino acid substitution in at least two of the CDRs (in comparison with the original CDRs).

**[0143]** The present disclosure also encompasses polypeptides, antibodies or antigen binding fragments comprising variable chains having at least one conservative amino acid substitution in the 3 CDRs (in comparison with the original CDRs).

**[0144]** The present disclosure also encompasses polypeptides, antibodies or antigen binding fragments comprising variable chains having at least two conservative amino acid substitutions in at least one of the CDRs (in comparison with the original CDRs).

**[0145]** The present disclosure also encompasses polypeptides, antibodies or antigen binding fragments comprising variable chains having at least two conservative amino acid substitutions in at least two of the CDRs (in comparison with the original CDRs).

**[0146]** The present disclosure also encompasses polypeptides, antibodies or antigen binding fragments comprising variable chains having at least two conservative amino acid substitutions in the 3 CDRs (in comparison with the original CDRs).

**[0147]** Comparison of the amino acid sequences of the light chain variable regions or the heavy chain variable regions of antibodies showing the greatest characteristics allowed us to derive consensus sequences within the CDRs and within the variable regions. The consensus for CDRs are provided in SEQ ID Nos: 72 to 88.

**[0148]** The present disclosure therefore provides in an example, an isolated antibody or antigen binding fragment comprising a light chain variable region having;

1. a. a CDRL1 sequence selected from the group consisting of SEQ ID and SEQ ID NO.:73;
2. b. a CDRL2 sequence selected from the group consisting of SEQ ID NO.:74, SEQ ID NO.: 75 and SEQ ID NO.:76, or;
3. c. a CDRL3 sequence selected from the group consisting of SEQ ID NO.:77, SEQ ID NO.:78 and SEQ ID NO.:79.

**[0149]** The present disclosure therefore provides in an example, an isolated antibody or antigen binding fragment comprising a heavy chain variable region having;

1. a. a CDRH1 sequence comprising SEQ ID NO.:80;
2. b. a CDRH2 sequence selected from the group consisting of SEQ ID NO.:81, SEQ ID NO.:82, SEQ ID NO.:83, SEQ ID NO.:84 and SEQ ID NO.:85, or;
3. c. a CDRH3 sequence selected from the group consisting of SEQ ID NO.:86, SEQ ID NO.:87 and SEQ ID NO.:88.

**[0150]** In accordance with the present disclosure, the antibody may comprise a CDR1 sequence comprising or consisting of formula:

$X_{1a}SSX_{2a}SLLX_{3a}X_{4a}X_{5a}X_{6a}X_{7a}X_{8a}X_{9a}X_{10a}LX_{11a}$  (SEQ ID NO.:72)

wherein  $X_{1a}$  may be a basic amino acid;

wherein  $X_{2a}$  may be a basic amino acid;

wherein  $X_{3a}$  may be H, Y or N;

wherein  $X_{4a}$  may be S, T, N or R;

wherein  $X_{5a}$  may be absent, S or N;

wherein  $X_{6a}$  may be D, F or N;

wherein  $X_{7a}$  may be G or Q;

wherein  $X_{8a}$  may be K, L or N;

wherein  $X_{9a}$  may be T or N;

wherein  $X_{10a}$  may be an aromatic amino acid, and;

wherein  $X_{11a}$  may be A, N, E or Y.

**[0151]** In an exemplary disclosure  $X_{1a}$  may be K or R.

**[0152]** In a further disclosure  $X_{2a}$  may be Q or K.

**[0153]** In yet a further disclosure  $X_{3a}$  may be N or H.

**[0154]** In an additional disclosure  $X_{10a}$  may be Y or F.

**[0155]** More specific disclosures include CDRL1 of SEQ ID where:  $X_{1a}$  is K;  $X_{2a}$  is Q;  $X_{3a}$  is N;  $X_{3a}$  is H;  $X_{4a}$  is S;  $X_{4a}$  is T;  $X_{5a}$  is S;  $X_{5a}$  is absent;  $X_{6a}$  is N;  $X_{7a}$  is Q;  $X_{7a}$  is G;  $X_{8a}$  is K;  $X_{9a}$  is N;  $X_{9a}$  is T;  $X_{10a}$  is Y; or  $X_{11a}$  is A.

**[0156]** In accordance with the present disclosure, the antibody may comprise a CDRL1 sequence comprising or consisting of formula:

KASQDX $X_{1b}$ X $X_{2b}$ X $X_{3b}$ X $X_{4b}$ X $X_{5b}$ X $X_{6b}$ (SEQ ID NO.:73)

wherein  $X_{1b}$  may be an hydrophobic amino acid;

wherein  $X_{2b}$  may be G or H;

wherein  $X_{3b}$  may be T, N or R;

wherein  $X_{4b}$  may be F, Y or A;

wherein  $X_{5b}$  may be an hydrophobic amino acid, and;

wherein  $X_{6b}$  may be N or A.

**[0157]** In an exemplary disclosure  $X_{1b}$  may be V or I.

**[0158]** In another exemplary disclosure  $X_{5b}$  may be V or L.

**[0159]** More specific disclosures include CDRL1 of SEQ ID NO.:73 where  $X_{1b}$  is I;  $X_{2b}$  is H;  $X_{3b}$  is T;  $X_{3b}$  is N;  $X_{4b}$  is Y;  $X_{4b}$  is F;  $X_{5b}$  is L or  $X_{6b}$  is N.

**[0160]** Other exemplary disclosures of CDRL1 are provided in SEQ ID NOs. 89 and 90.

**[0161]** In accordance with the present disclosure, the antibody may comprise a CDRL2 sequence comprising or consisting of formula:

FX $X_{1c}$ STX $X_{2c}$ X $X_{3c}$ S (SEQ ID NO.:74)

Wherein  $X_{1c}$  is A or G;

Wherein  $X_{2c}$  is R or T, and;

Wherein  $X_{3c}$  is E, K or A.

**[0162]** In an exemplary disclosure  $X_{1c}$  may be A and  $X_{2c}$  may be T.

**[0163]** In another exemplary disclosure  $X_{1c}$  may be A and  $X_{2c}$  may be R.

**[0164]** Other specific disclosures include CDRL2 of SEQ ID NO.:74 where  $X_{1c}$  is A;  $X_{2c}$  is R or  $X_{3c}$  is E.

**[0165]** In accordance with the present disclosure, the antibody may comprise a CDRL2 sequence comprising or consisting of formula:

$X_{1d}VX_{2d}X_{3d}X_{4d}S$  (SEQ ID NO.:75)

Wherein  $X_{1d}$  may be L or K;

Wherein  $X_{2d}$  may be a basic amino acid;

Wherein  $X_{3d}$  may be L or R and;

Wherein  $X_{4d}$  may be D or F.

**[0166]** In an exemplary disclosure  $X_{2d}$  may be K or N.

**[0167]** Other specific disclosures include CDRL2 of SEQ ID NO.:75 where  $X_{1d}$  is L;  $X_{2d}$  is K;  $X_{3d}$  is L or  $X_{4d}$  is D.

**[0168]** In accordance with the present disclosure, the antibody may comprise a CDRL2 sequence comprising or consisting of formula:

$X_{1e}ANRLVX_{2e}$  (SEQ ID NO.:76)

Wherein  $X_{1e}$  may be a basic amino acid, and;

Wherein  $X_{2e}$  may be D or A.

**[0169]** In an exemplary disclosure  $X_{1e}$  may be R or H.

**[0170]** Other specific disclosures include CDRL2 of SEQ ID NO.:76 where  $X_{1e}$  is R or  $X_{2e}$  is D.

**[0171]** Other exemplary disclosures of CDRL2 are provided in SEQ ID NOs.: 91-93.

**[0172]** In accordance with the present disclosure, the antibody may comprise a CDRL3 sequence comprising or consisting of formula:

$X_{1f}QX_{2f}X_{3f}X_{4f}X_{5f}PLT$  (SEQ ID NO.:77)

Wherein  $X_{1f}$  may be Q or L;

Wherein  $X_{2f}$  may be an aromatic amino acid;

Wherein  $X_{3f}$  may be D, F or Y;

Wherein  $X_{4f}$  may be E, A, N or S, and;

Wherein  $X_{5f}$  may be I, F or T.

**[0173]** In an exemplary disclosure of the invention  $X_{2f}$  may be Y or H.

**[0174]** In another exemplary disclosure of the invention  $X_{3f}$  may be Y or D.

**[0175]** In yet another exemplary disclosure of the invention  $X_{5f}$  may be I or T.

**[0176]** Other specific disclosures include CDRL3 of SEQ ID NO.:77 where  $X_{1f}$  is Q;  $X_{2f}$  is H;  $X_{3f}$  is D;  $X_{3f}$  is Y;  $X_{4f}$  is S;  $X_{4f}$  is E;  $X_{4f}$  is A;  $X_{5f}$  is T, or  $X_{5f}$  is I.

**[0177]** In accordance with the present disclosure, the antibody may comprise a CDRL3 sequence comprising or consisting of formula:

$QQHX_{1g}X_{2g}X_{3g}PLT$  (SEQ ID NO.:78)

Wherein  $X_{1g}$  may be an aromatic amino acid;

Wherein  $X_{2g}$  may be N or S, and;

Wherein  $X_{3g}$  may be I or T.

**[0178]** In an exemplary disclosures  $X_{1g}$  may be F or Y

**[0179]** Other specific disclosures include CDRL3 of SEQ ID NO.:78 where  $X_{2g}$  is S or  $X_{3g}$  is T.

**[0180]** In accordance with the present disclosure, the antibody may comprise a CDRL3 sequence comprising or consisting of formula:

$X_{1h}QGX_{2h}HX_{3h}PX_{4h}T$  (SEQ ID NO.:79)

Wherein  $X_{1h}$  may be an aromatic amino acid;

Wherein  $X_{2h}$  may be a neutral hydrophilic amino acid;

Wherein  $X_{3h}$  may be F or V, and;

Wherein  $X_{4h}$  may be R or L.

**[0181]** In an exemplary disclosure  $X_{1h}$  may be W or F.

**[0182]** In another exemplary disclosure  $X_{2h}$  may be S or T.

**[0183]** Other specific disclosures include CDRL3 of SEQ ID NO.:79 where  $X_{1h}$  is W;  $X_{2h}$  is T;  $X_{3h}$  is F, or  $X_{4h}$  is R.

**[0184]** Other exemplary disclosures of CDRL3 are provided in SEQ ID NOs. 94 and 95.

**[0185]** In accordance with the present disclosure, the antibody may comprise a CDRH1 sequence comprising or consisting of formula:

$GYX_{1i}FX_{2i}X_{3i}YX_{4i}X_{5i}H$  (SEQ ID NO.:80)

Wherein  $X_{1i}$  may be T, I or K;

Wherein  $X_{2i}$  may be a neutral hydrophilic amino acid;

Wherein  $X_{3i}$  may be an acidic amino acid;

Wherein  $X_{4i}$  may be E, N or D, and;

Wherein  $X_{5i}$  may be hydrophobic amino acid.

**[0186]** In an exemplary disclosure  $X_{2i}$  may be T or S.

**[0187]** In another exemplary disclosure  $X_{3i}$  may be D or E.

**[0188]** In yet another exemplary disclosure  $X_{4i}$  may be N or E.

**[0189]** In a further exemplary disclosure  $X_{5i}$  may be M, I or V.

**[0190]** Other specific disclosures include CDRH1 of SEQ ID NO.:80 where  $X_{2j}$  is T;  $X_{3j}$  is D;  $X_{4j}$  is E;  $X_{5j}$  is I or  $X_{5j}$  is M.

**[0191]** Other exemplary disclosures of CDRH1 are provided in SEQ ID NOs.: 96 and 97.

**[0192]** In accordance with the present disclosure, the antibody may comprise a CDRH2 sequence comprising or consisting of formula:

$X_{1j}X_{2j}DPX_{3j}TGX_{4j}TX_{5j}$  (SEQ ID NO.:81)

Wherein  $X_{1j}$  may be V or G

Wherein  $X_{2j}$  may be a hydrophobic amino acid;

Wherein  $X_{3j}$  may be A, G or E;

Wherein  $X_{4j}$  may be R, G, D, A, S, N or V, and;

Wherein  $X_{5j}$  may be a hydrophobic amino acid.

**[0193]** In an exemplary disclosure  $X_{2j}$  may be I or L.

**[0194]** In another exemplary disclosure  $X_{5j}$  may be A or V.

**[0195]** Other specific disclosures include CDRH2 of SEQ ID NO.:81 where  $X_{1j}$  is V;  $X_{2j}$  is I;  $X_{3j}$  is E;  $X_{4j}$  is D or  $X_{5j}$  is A.

**[0196]** In accordance with the present disclosure, the antibody may comprise a CDRH2 sequence comprising or consisting of formula:

$VX_{1k}DPX_{2k}TGX_{3k}TA$  (SEQ ID NO.:82)

Wherein  $X_{1k}$  may be an hydrophobic amino acid;

Wherein  $X_{2k}$  may be A, E or G;

Wherein  $X_{3k}$  may be R, G, A, S, N V or D.

**[0197]** In an exemplary disclosure  $X_{1k}$  may be L or I.

**[0198]** Other specific disclosures include CDRH2 of SEQ ID NO.:82 where  $X_{1k}$  is I;  $X_{2k}$  is E, or

$X_{3k}$  is D.

**[0199]** In accordance with the present disclosure, the antibody may comprise a CDRH2 sequence comprising or consisting of formula:

$Y|X_{1l}|X_{2l}|X_{3l}|GX_{4l}|X_{5l}|X_{6l}$  (SEQ ID NO.:83)

Wherein  $X_{1l}$  may be S or N;

Wherein  $X_{2l}$  may be an aromatic amino acid

Wherein  $X_{3l}$  may be D, E or N;

Wherein  $X_{4l}$  may be a D or H;

Wherein  $X_{5l}$  may be Y, S or N;

Wherein  $X_{6l}$  may be D, E or N.

**[0200]** In an exemplary disclosure  $X_{3l}$  may be D or N.

**[0201]** In another exemplary disclosure  $X_{6l}$  may be D or N.

**[0202]** Other specific disclosures include CDRH2 of SEQ ID NO.:83 where  $X_{2l}$  is F or Y,  $X_{3l}$  is N,  $X_{4l}$  is D or  $X_{6l}$  is N.

**[0203]** In accordance with the present disclosure, the antibody may comprise a CDRH2 sequence comprising or consisting of formula:

$X_{1m}|NPYNX_{2m}|VTE$  (SEQ ID NO.:84)

wherein  $X_{1m}$  may be N or Y, and;

wherein  $X_{2m}$  may be E, D or N.

**[0204]** In an exemplary disclosure  $X_{2m}$  may be D or N.

**[0205]** Other specific disclosures include CDRH2 of SEQ ID NO.:84 where  $X_{1m}$  is N or  $X_{2m}$  is D.

**[0206]** In accordance with the present disclosure, the antibody may comprise a CDRH2 sequence comprising or consisting of formula:

DINPX<sub>1n</sub>YGX<sub>2n</sub>X<sub>3n</sub>T (SEQ ID NO.:85)

Wherein X<sub>1n</sub> may be N or Y;

Wherein X<sub>2n</sub> may be G or T and;

wherein X<sub>3n</sub> may be I or T.

**[0207]** Other exemplary disclosures of CDRH2 are provided in SEQ ID NOs. 98 and 99.

**[0208]** In accordance with the present disclosure, the antibody may comprise a CDRH3 sequence comprising or consisting of formula:

MX<sub>1o</sub>X<sub>2o</sub>X<sub>3o</sub>DY (SEQ ID NO.:86)

Wherein X<sub>1o</sub> may be G or S;

Wherein X<sub>2o</sub> may be Y or H, and;

wherein X<sub>3o</sub> may be A or S.

**[0209]** Other specific disclosures include CDRH3 of SEQ ID NO.:86 where X<sub>1o</sub> is G; X<sub>2o</sub> is Y or X<sub>3o</sub> is S.

**[0210]** In accordance with the present disclosure, the antibody may comprise a CDRH3 sequence comprising or consisting of formula:

IX<sub>1p</sub>YAX<sub>2p</sub>DY (SEQ ID NO.:87)

Wherein X<sub>1p</sub> may be G or S and;

Wherein X<sub>2p</sub> may be absent or M.

**[0211]** Other specific disclosures include CDRH3 of SEQ ID NO.:87 where X<sub>1p</sub> is S or X<sub>2p</sub> is M.

**[0212]** In accordance with the present disclosure, the antibody may comprise a CDRH3 sequence comprising or consisting of formula:

AX<sub>1q</sub>X<sub>2q</sub>GLRX<sub>3q</sub> (SEQ ID NO.:88)

Wherein  $X_{1q}$  may be R or W;

Wherein  $X_{2q}$  may be an aromatic amino acid and;

wherein  $X_{3q}$  may be a basic amino acid.

**[0213]** In an exemplary disclosure  $X_{2q}$  may be W or F.

**[0214]** In another disclosure  $X_{3q}$  may be Q or N.

**[0215]** Other specific disclosures of the invention include CDRH3 of SEQ ID NO.:88 where  $X_{1q}$  is R;  $X_{2q}$  is W or  $X_{3q}$  is N.

**[0216]** Variant antibodies or antigen binding fragments encompassed by the present invention include those that may comprise an insertion, a deletion or an amino acid substitution (conservative or non-conservative). These variants may have at least one amino acid residue in its amino acid sequence removed and a different residue inserted in its place.

**Humanized antibodies**

**[0217]** Exemplary embodiments of variant antibodies and antigen binding fragments of the present invention are a group of antibodies and antigen binding fragments capable of binding to KAAG1 and characterized herein as being humanized.

**[0218]** The humanized antibodies and antigen binding fragments of the present disclosure includes more particularly, humanized 3D3, 3A4 or 3C4 antibodies and antigen binding fragments. Humanized 3A4 antibodies and antigen binding fragments may be used in accordance with the present invention. The humanized 3D3, 3A4 or 3C4 antibodies have at least one amino acid difference in a framework region in comparison with the monoclonal 3D3, 3A4 or 3C4 antibody.

**[0219]** Humanized 3A4 antibodies having CDRs identical to those of the monoclonal 3A4 antibody (VL: SEQ ID NO.:48, VH: SEQ ID NO.:46) were generated and tested. These humanized antibodies comprise up to 11 amino acid substitutions (from one to eleven) in the variable light chain framework region and up to 23 amino acid substitutions (from one to twenty-three) in the variable heavy chain framework region in comparison with the monoclonal 3A4 antibody. The applicant has shown that these humanized 3A4 antibodies bind to KAAG1 as efficiently as the monoclonal 3A4 antibody.

**[0220]** Exemplary embodiments of variant antibody or antigen binding fragments include those having a light chain variable region as set forth in SEQ ID NO.:186:

SEQ ID NO.:186

DXVMTQTPLSLVXXGXXASISCRSSQSLHSNGNTYLEWYLQKPGQSPXLLIHTVSNR

FSGVPDRFSGSGSGTDFTLKISRVEAEDXGVYYCFQGSHVPLTFGXGTLEXK,

wherein at least one of the amino acids identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48. The amino acid substitution may be, for example, an amino acid found at a corresponding position of a natural human antibody or a human antibody consensus. The amino acid substitution may be, for example conservative.

**[0221]** Another exemplary embodiment of a variant antibody or antigen binding fragment include those having a light chain variable region as set forth in SEQ ID NO.:187:

SEQ ID NO.:187

DX<sub>e1</sub>VMTQTPLSLX<sub>e2</sub>VX<sub>e3</sub>X<sub>e4</sub>GX<sub>e5</sub>X<sub>e6</sub>ASISCRSSQSLHSNGNTYLEWYLQKPGQSPX<sub>e7</sub>L

LIHTVSNRFSGVVPDRFSGSGSGTDFTLKISRVEAEDX<sub>e8</sub>GVYYCFQGSHVPLTFGX<sub>e9</sub>GT

X<sub>e10</sub>LEX<sub>e11</sub>K,

Wherein X<sub>e1</sub> may be a hydrophobic amino acid;

Wherein X<sub>e2</sub> may be A or P;

Wherein X<sub>e3</sub> may be neutral hydrophilic amino acid;

Wherein X<sub>e4</sub> may be L or P;

Wherein X<sub>e5</sub> may be an acidic amino acid;

Wherein X<sub>e6</sub> may be Q or P;

Wherein X<sub>e7</sub> may be a basic amino acid;

Wherein X<sub>e8</sub> may be a hydrophobic amino acid;

Wherein X<sub>e9</sub> may be A or Q;

Wherein X<sub>e10</sub> may be a basic amino acid; or

Wherein X<sub>e11</sub> may be a hydrophobic amino acid,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48.

**[0222]** An additional exemplary embodiment of a variant antibody or antigen binding fragment include those having a light chain variable region as set forth in SEQ ID NO.:188:

SEQ ID NO.:188

DX<sub>E1</sub>VMTQTPLSLX<sub>E2</sub>VX<sub>E3</sub>X<sub>E4</sub>GX<sub>E5</sub>X<sub>E6</sub>ASISCRSSQSLLHSNGNTYLEWYLQKPGQSPX<sub>E7</sub>  
 LLIHTVSNRFSGPDRFSGSGSGTDFTLKISRVEAEDX<sub>E8</sub>GVYYCFQGSHVPLTFGX<sub>E9</sub>G  
 TX<sub>E10</sub>LEX<sub>E11</sub>K

Wherein X<sub>E1</sub> may be V or I

Wherein X<sub>E2</sub> may be A or P

Wherein X<sub>E3</sub> may be S or T

Wherein X<sub>E4</sub> may be L or P

Wherein X<sub>E5</sub> may be D or E

Wherein X<sub>E6</sub> may be Q or P

Wherein X<sub>E7</sub> may be K or Q

Wherein X<sub>E8</sub> may be L or V

Wherein X<sub>E9</sub> may be A or Q

Wherein X<sub>E10</sub> may be R or K or

Wherein X<sub>E11</sub> may be L or I,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48.

**[0223]** In accordance with an embodiment, the light chain variable domain variant may have a sequence as set forth in SEQ ID NO.:189 or 190:

SEQ ID NO.:189

DIVMTQTPLSLPVTPGEPASISCRSSQSLLHSNGNTYLEWYLQKPGQSPQLLIYTVSNR  
 FSGVPDRFSGSGSGTDFTLKISRVEAEDVGVYYCFQGSHVPLTFGQGTKLEIK.

SEQ ID NO.:190

DVVMTQTPLSLPVTPGEPASISCRSSQSLLHSNGNTYLEWYLQKPGQSPKLLIYTVSNR  
 FSGVPDRFSGSGSGTDFTLKISRVEAEDVGVYYCFQGSHVPLTFGQGTKLEIK.

**[0224]** Exemplary embodiments of variant antibody or antigen binding fragments include those having a heavy chain variable region as set forth in SEQ ID NO.:191.

SEQ ID NO.:191

QXQI VQSGXFXXKPGASV/KXSCKASGYTFTDDYMSWV/XQXXGXXI FWXGDINPYNG

DTNYNQKFKGXXXXTDXSXSTAYMLXSLXSEDXAVYYCARDPGAMDYWGQGTXVT

VSS,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:46. The amino acid substitution may be, for example, an amino acid found at a corresponding position of a natural human antibody or a human antibody consensus. The amino acid substitution may be, for example conservative.

**[0225]** Another exemplary embodiment of a variant antibody or antigen binding fragment include those having a heavy chain variable region as set forth in SEQ ID NO.:192:

SEQ ID NO.:192

QX<sub>f1</sub>QLVQSGX<sub>f2</sub>EX<sub>f3</sub>X<sub>bf4</sub>KPGASVKX<sub>f5</sub>SCKASGYTFTDDYMSWVX<sub>f6</sub>QX<sub>f7</sub>X<sub>f8</sub>GX<sub>f9</sub>X<sub>f10</sub>LEW  
X<sub>f11</sub>GDINPYNGDTNYNQKFKGX<sub>f12</sub>X<sub>f13</sub>X<sub>bf14</sub>X<sub>f15</sub>TX<sub>f16</sub>DX<sub>f17</sub>SX<sub>f18</sub>STAYMX<sub>f19</sub>LX<sub>f20</sub>SLX<sub>f21</sub>SED  
X<sub>f22</sub>AVYYCARDPGAMDYWGQGTX<sub>f23</sub>VTVSS,

Wherein X<sub>f1</sub> may be a hydrophobic amino acid;

Wherein X<sub>bf2</sub> may be P or A;

Wherein X<sub>f3</sub> may be a hydrophobic amino acid;

Wherein X<sub>f4</sub> may be V or K;

Wherein X<sub>f5</sub> may be a hydrophobic amino acid;

Wherein X<sub>f6</sub> may be a basic amino acid;

Wherein X<sub>f7</sub> may be S or A;

Wherein X<sub>f8</sub> may be H or P;

Wherein X<sub>f9</sub> may be a basic amino acid;

Wherein X<sub>f10</sub> may be S or G;

Wherein X<sub>f11</sub> may be a hydrophobic amino acid;

Wherein X<sub>f12</sub> may be a basic amino acid;

Wherein X<sub>f13</sub> may be a hydrophobic amino acid;

Wherein X<sub>f14</sub> may be I or T;

Wherein X<sub>f15</sub> may be a hydrophobic amino acid;

Wherein X<sub>f16</sub> may be a hydrophobic amino acid;

Wherein  $X_{f17}$  may be K or T;

Wherein  $X_{f18}$  may be a neutral hydrophilic amino acid;

Wherein  $X_{f19}$  may be Q or E;

Wherein  $X_{f20}$  may be N or S;

Wherein  $X_{f21}$  may be T or R;

Wherein  $X_{f22}$  may be a neutral hydrophilic amino acid; or

Wherein  $X_{f23}$  may be S or L,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:46.

**[0226]** An additional exemplary embodiment of a variant antibody or antigen binding fragment include those having a heavy chain variable region as set forth in SEQ ID NO.:193:

SEQ ID NO.:193

$QX_{F1}QLVQSGX_{F2}EX_{F3}X_{F4}KPGASVKX_{F5}SCKASGYTFTDDYMSWVX_{F6}QX_{F7}X_{F8}GX_{F9}X_{F10}L$   
 $EWX_{F11}GDINPYNGDTNYNQKFKGX_{F12}X_{F13}X_{F14}X_{F15}TX_{F16}DX_{F17}SX_{F18}STAYMX_{F19}LX_{F20}SL$   
 $X_{F21}SEDX_{F22}AVYYCARDPGAMDYWGQQGTX_{F23}VTVSS$

Wherein  $X_{F1}$  may be I or V;

Wherein  $X_{F2}$  may be P or A;

Wherein  $X_{F3}$  may be M or V;

Wherein  $X_{F4}$  may be V or K;

Wherein  $X_{F5}$  may be M or V;

Wherein  $X_{F6}$  may be K or R;

Wherein  $X_{F7}$  may be S or A;

Wherein  $X_{F8}$  may be H or P;

Wherein  $X_{F9}$  may be K or Q;

Wherein  $X_{F10}$  may be S or G;

Wherein  $X_{F11}$  may be I or M;

Wherein  $X_{F12}$  may be K or R;

Wherein  $X_{F13}$  may be A or V;

Wherein  $X_{F14}$  may be I or T;

Wherein  $X_{F15}$  may be L or I;

Wherein  $X_{F16}$  may be V or A;

Wherein  $X_{F17}$  may be K or T;

Wherein  $X_{F18}$  may be S or T;

Wherein  $X_{F19}$  may be Q or E;

Wherein  $X_{F20}$  may be N or S;

Wherein  $X_{F21}$  may be T or R;

Wherein  $X_{F22}$  may be S or T; or

Wherein  $X_{F23}$  is S or L,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:46.

**[0227]** In accordance with an embodiment, the heavy chain variable domain variant may have a sequence as set forth in any one of SEQ ID NO.194 to 197:

SEQ ID NO.:194

QVQLVQSGAEVKKPGASVKVSCKASGYTFTDDYMSWVRQAPGQGLEWMGDINPYNG  
DTNYNQKFKGRVTITADTSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQQTLTV  
SS.

SEQ ID NO.:195

QIQLVQSGAEVKKPGASVKVSCKASGYTFTDDYMSWVRQAPGQGLEWMGDINPYNG  
DTNYNQKFKGRVTITADKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQQTLVT  
VSS.

SEQ ID NO.:196

QIQLVQSGAEVKKPGASVKVSCKASGYTFTDDYMSWVRQAPGQGLEWIGDINPYNGD  
TNYNQKFKGRATLTVDKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQQTLTV  
SS.

SEQ ID NO.:197

QIQLVQSGAEVKPGASVKVSKASGYFTDDYMSWVKQAPGQGLEWIGDINPYNGDT  
NYNQKFKGKATLTVDKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVS  
S.

**[0228]** In accordance with the present disclosure, the humanized 3D3 antibody may have a light chain variable region of formula:

DIVMTQSPXSLAVSXGXXXTNCKSSQSLLNSNFQKNFLAWYQQKPGQXPKLLIYFAS  
TRESXPDRFXGSGSGTDFTLTISSXQAEDXAXYXCQQHYSTPLTFGXGKLEXK (SEQ  
ID NO.:174);

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:16. The amino acid substitution may be, for example conservative.

**[0229]** In accordance with a more specific disclosure, the humanized 3D3 antibody may have a light chain variable region of formula:

DIVMTQSPX<sub>A1</sub>SLAVSX<sub>A2</sub>GX<sub>A3</sub>X<sub>A4</sub>X<sub>A5</sub>TX<sub>A6</sub>NCKSSQSLLNSNFQKNFLAWYQQK  
PQX<sub>A7</sub>PKLLIYFASTRESSX<sub>A8</sub>PDRFX<sub>A9</sub>GSGSGTDFTLTISSX<sub>A10</sub>QAEDX<sub>A11</sub>AX<sub>A12</sub>YX<sub>A13</sub>CQ  
QHYSTPLTFGX<sub>A14</sub>GKLEX<sub>A15</sub>K (SEQ ID NO.:175);

Wherein X<sub>A1</sub> may be, for example, D or S;

Wherein X<sub>A2</sub> may be, for example, a hydrophobic amino acid or more particularly L or I;

Wherein X<sub>A3</sub> may be, for example, E or Q;

Wherein X<sub>A4</sub> may be, for example, a basic amino acid or more particularly R or K;

Wherein X<sub>A5</sub> may be, for example, a hydrophobic amino acid or more particularly A or V;

Wherein X<sub>A6</sub> may be, for example, a hydrophobic amino acid or more particularly I or M;

Wherein X<sub>A7</sub> may be, for example, P or S;

Wherein X<sub>A8</sub> may be, for example, a hydrophobic amino acid or more particularly V or I;

Wherein X<sub>A9</sub> may be, for example, S or I;

Wherein X<sub>A10</sub> may be, for example, a hydrophobic amino acid or more particularly L or V;

Wherein X<sub>A11</sub> may be, for example, a hydrophobic amino acid or more particularly V or L;

Wherein X<sub>A12</sub> may be, for example, V or D;

Wherein X<sub>A13</sub> may be, for example, an aromatic amino acid or more particularly Y or F;

Wherein X<sub>A14</sub> may be, for example, Q or A and;

Wherein X<sub>A15</sub> may be, for example, a hydrophobic amino acid or more particularly I or L.

**[0230]** In accordance with an even more specific disclosure, the humanized 3D3 antibody may have a light chain variable region of formula:

DIVMTQSPX<sub>a1</sub>SLAVSX<sub>a2</sub>GX<sub>a3</sub>X<sub>a4</sub>X<sub>a5</sub>TX<sub>a6</sub>NCKSSQSLLNSNFQKNFLAWYQQKP  
GQX<sub>a7</sub>PKLLIYFASTRESSX<sub>a8</sub>PDRFX<sub>a9</sub>GSGSGTDFTLTISSSX<sub>a10</sub>QAEDX<sub>a11</sub>AX<sub>a12</sub>YX<sub>a13</sub>CQQ  
HYSTPLTGFGX<sub>a14</sub>GTKLEX<sub>a15</sub>K (SEQ ID NO.:176);

Wherein X<sub>a1</sub> may be, for example, D or S;

Wherein X<sub>a2</sub> may be, for example, L or I;

Wherein X<sub>a3</sub> may be, for example, E or Q;

Wherein X<sub>a4</sub> may be, for example, R or K;

Wherein X<sub>a5</sub> may be, for example, A or V;

Wherein X<sub>a6</sub> may be, for example, I or M;

Wherein X<sub>a7</sub> may be, for example, P or S;

Wherein X<sub>a8</sub> may be, for example, V or I;

Wherein X<sub>a9</sub> may be, for example, S or I;

Wherein X<sub>a10</sub> may be, for example, L or V;

Wherein X<sub>a11</sub> may be, for example, V or L;

Wherein X<sub>a12</sub> may be, for example, V or D;

Wherein X<sub>a13</sub> may be, for example, Y or F;

Wherein X<sub>a14</sub> may be, for example, Q or A and;

Wherein X<sub>a15</sub> is for example, I or L.

**[0231]** In accordance with the present disclosure, the humanized 3D3 antibody may have a

heavy chain variable region of formula:

EVQLXQSXAEXXXPGASVXSCKAS**GYIFTDYEIH**WVXQXPXXGLEWXG**VIDPE**

**TGNTAFNQKFKGXXXTADXSXSTAYMELSSLTSEDXAVYYC**MGYSDY**WGQGTXTV** SS (SEQ ID NO.:177); wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:18. The amino acid substitution may be, for example conservative.

**[0232]** In accordance with a more specific disclosure, the humanized 3D3 antibody may have a heavy chain variable region of formula:

EVQLX<sub>B1</sub>QSX<sub>B2</sub>AEX<sub>B3</sub>X<sub>B4</sub>X<sub>B5</sub>PGASVX<sub>B6</sub>X<sub>B7</sub>SCKAS**GYIFTDYEIH**WVX<sub>B8</sub>QX<sub>B9</sub>PX<sub>B10</sub>X<sub>B11</sub>GLEWX<sub>B12</sub>**VIDPETGNTAFNQKFKGX**<sub>B13</sub>X<sub>B14</sub>TX<sub>B15</sub>TADX<sub>B16</sub>SX<sub>B17</sub>STAYMELSSLTS**EDX**<sub>B18</sub>AVYYC**MGYSDY**WGQGTX<sub>B19</sub>X<sub>B20</sub>TVSS (SEQ ID NO.:178),

Wherein X<sub>B1</sub> may be, for example, V or Q;

Wherein X<sub>B2</sub> may be, for example, G or V;

Wherein X<sub>B3</sub> may be, for example, a hydrophobic amino acid or more particularly V or L;

Wherein X<sub>B4</sub> may be, for example, K or V;

Wherein X<sub>B5</sub> may be, for example, a basic amino acid or more particularly K or R;

Wherein X<sub>B6</sub> may be, for example, K or T;

Wherein X<sub>B7</sub> may be, for example, a hydrophobic amino acid or more particularly V or L;

Wherein X<sub>B8</sub> may be, for example, a basic amino acid or more particularly R or K;

Wherein X<sub>B9</sub> may be, for example, A or T;

Wherein X<sub>B10</sub> may be, for example, G or V;

Wherein X<sub>B11</sub> may be, for example, Q or H;

Wherein X<sub>B12</sub> may be, for example, a hydrophobic amino acid or more particularly M or I;

Wherein X<sub>B13</sub> may be, for example, a basic amino acid or more particularly R or K;

Wherein X<sub>B14</sub> may be, for example, a hydrophobic amino acid or more particularly V or A;

Wherein X<sub>B15</sub> may be, for example, a hydrophobic amino acid or more particularly I or L;

Wherein X<sub>B16</sub> may be, for example, T or I;

Wherein X<sub>B17</sub> may be, for example, a neutral hydrophilic amino acid or more particularly T or

S;

Wherein  $X_{B18}$  may be, for example, a neutral hydrophilic amino acid or more particularly T or S;

Wherein  $X_{B19}$  may be, for example, L or T and;

Wherein  $X_{B20}$  may be, for example, a hydrophobic amino acid or more particularly V or L.

**[0233]** In accordance with a more specific disclosure, the humanized 3D3 antibody may have a heavy chain variable region of formula:

EVQLX<sub>b1</sub>QSX<sub>b2</sub>AEX<sub>b3</sub>X<sub>b4</sub>X<sub>b5</sub>PGASVX<sub>b6</sub>X<sub>b7</sub>SCKASGYIFTDYEIHVVX<sub>b8</sub>QX<sub>b9</sub>PX<sub>b10</sub>  
X<sub>b11</sub>GLEWX<sub>b12</sub>**GVIDPETGNTAFNQKFKGX<sub>b13</sub>X<sub>b14</sub>TX<sub>b15</sub>TADX<sub>b16</sub>SX<sub>b17</sub>STAYMELSSLTSE**  
DX<sub>b18</sub>AVYYC**MGYSDYWGQGTX<sub>b19</sub>X<sub>b20</sub>TVSS** (SEQ ID NO.:179);

Wherein  $X_{b1}$  may be, for example, V or Q;

Wherein  $X_{b2}$  may be, for example, G or V;

Wherein  $X_{b3}$  may be, for example, V or L;

Wherein  $X_{b4}$  may be, for example, K or V;

Wherein  $X_{b5}$  may be, for example, K or R;

Wherein  $X_{b6}$  may be, for example, K or T;

Wherein  $X_{b7}$  may be, for example, V or L;

Wherein  $X_{b8}$  may be, for example, R or K;

Wherein  $X_{b9}$  may be, for example, A or T;

Wherein  $X_{b10}$  may be, for example, G or V;

Wherein  $X_{b11}$  may be, for example, Q or H;

Wherein  $X_{b12}$  may be, for example, M or I;

Wherein  $X_{b13}$  may be, for example, R or K;

Wherein  $X_{b14}$  may be, for example, V or A;

Wherein  $X_{b15}$  may be, for example, I or L;

Wherein  $X_{b16}$  may be, for example, T or I;

Wherein  $X_{b17}$  may be, for example, T or S;

Wherein  $X_{b18}$  may be, for example, T or S;

Wherein  $X_{b19}$  may be, for example, L or T;

Wherein  $X_{b20}$  may be, for example, V or L.

**[0234]** In accordance with the present disclosure, the humanized 3C4 antibody may have a light chain variable region of formula:

**DIVMXQSPSSXXASGXRVТИTCKASQDIHNFLNWFQQKPGKXPKTLIFRANRL  
VDGVPSRSGSGSGDXYLTISLXXEDXXXSCLQYDEIPLTFGXGKLEXX** (SEQ ID NO.:180); wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:24. The amino acid substitution may be, for example conservative.

**[0235]** In accordance with a more specific disclosure, the humanized 3C4 antibody may have a light chain variable region of formula:

**DIVMX<sub>C1</sub>QSPSSX<sub>C2</sub>X<sub>C3</sub>ASX<sub>C4</sub>GX<sub>C5</sub>RVTITCKASQDIHNFLNWFQQKPGKX<sub>C6</sub>PKT  
LIFRANRLVDGVPSRSGSGSGX<sub>C7</sub>DYX<sub>C8</sub>LTISLX<sub>C9</sub>X<sub>C10</sub>EDX<sub>C11</sub>X<sub>C12</sub>X<sub>C13</sub>YSCLQYDEIP  
LTFGX<sub>C14</sub>GTKLEX<sub>C15</sub>X<sub>C16</sub>** (SEQ ID NO.:181);

Wherein  $X_{C1}$  may be, for example, a neutral hydrophilic amino acid or more particularly T or S;

Wherein  $X_{C2}$  may be, for example, a hydrophobic amino acid or more particularly L or M;

Wherein  $X_{C3}$  may be, for example, S or Y;

Wherein  $X_{C4}$  may be, for example, a hydrophobic amino acid or more particularly V or L;

Wherein  $X_{C5}$  may be, for example, an acidic amino acid or more particularly D or E;

Wherein  $X_{C6}$  may be, for example, A or S;

Wherein  $X_{C7}$  may be, for example, T or Q;

Wherein  $X_{C8}$  may be, for example, a neutral hydrophilic amino acid or more particularly T or S;

Wherein  $X_{C9}$  may be, for example, Q or E;

Wherein  $X_{C10}$  may be, for example, P or F;

Wherein  $X_{C11}$  may be, for example, F or L;

Wherein  $X_{C12}$  may be, for example, A or G;

Wherein  $X_{C13}$  may be, for example, T or I;

Wherein  $X_{C14}$  may be, for example, Q or A;

Wherein  $X_{C15}$  may be, for example, a hydrophobic amino acid or more particularly I or L, and; wherein  $X_{C16}$  may be, for example, a basic amino acid or more particularly K or R.

**[0236]** In accordance with a more specific disclosure, the humanized 3C4 antibody may have a light chain variable region of formula:

DIVMX<sub>c1</sub>QSPSSX<sub>c2</sub>X<sub>c3</sub>ASX<sub>c4</sub>GX<sub>c5</sub>RVТИTCK**A**SQDIHNFLNW**F**QQKPGKX<sub>c6</sub>PKTLI  
**F**RANRLVDGVPSRFSGSGGX<sub>c7</sub>DYX<sub>c8</sub>LTSSLX<sub>c9</sub>X<sub>c10</sub>EDX<sub>c11</sub>X<sub>c12</sub>X<sub>c13</sub>YSCLQY**D**E**I**PLTF  
GX<sub>c14</sub>GTKLEX<sub>c15</sub>X<sub>c16</sub> (SEQ ID NO.:182);

Wherein  $X_{c1}$  may be, for example, T or S;

Wherein  $X_{c2}$  may be, for example, L or M;

Wherein  $X_{c3}$  may be, for example, S or Y;

Wherein  $X_{c4}$  may be, for example, V or L;

Wherein  $X_{c5}$  may be, for example, D or E;

Wherein  $X_{c6}$  may be, for example, A or S;

Wherein  $X_{c7}$  may be, for example, T or Q;

Wherein  $X_{c8}$  may be, for example, T or S;

Wherein  $X_{c9}$  may be, for example, Q or E;

Wherein  $X_{c10}$  may be, for example, P or F;

Wherein  $X_{c11}$  may be, for example, F or L;

Wherein  $X_{c12}$  may be, for example, A or G;

Wherein  $X_{c13}$  may be, for example, T or I;

Wherein  $X_{c14}$  may be, for example, Q or A;

Wherein  $X_{c15}$  may be, for example, I or L and;

wherein  $X_{c16}$  may be, for example, K or R.

**[0237]** In accordance with the present disclosure, the humanized 3C4 antibody may have a heavy chain variable region of formula:

EVQLQESGPX<sub>D1</sub>LVKPSQX<sub>D2</sub>LSLTCTVX<sub>D3</sub>**GFSITSGYGWHWIRQX<sub>D4</sub>PGX<sub>D5</sub>X<sub>D6</sub>L**  
**YDGHNDYNPSLKSRX<sub>D7</sub>X<sub>D8</sub>X<sub>D9</sub>IX<sub>D10</sub>QDTSKNQFX<sub>D11</sub>LX<sub>D12</sub>LX<sub>D13</sub>SVTX<sub>D14</sub>X<sub>D15</sub>D**  
(SEQ ID NO.:183); wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:26. The amino acid substitution may be, for example conservative.

**[0238]** In accordance with a more specific disclosure, the humanized 3C4 antibody may have a heavy chain variable region of formula:

EVQLQESGPX<sub>D1</sub>LVKPSQX<sub>D2</sub>LSLTCTVX<sub>D3</sub>**GFSITSGYGWHWIRQX<sub>D4</sub>PGX<sub>D5</sub>X<sub>D6</sub>L**  
**EWX<sub>D7</sub>GYINYDGHNDYNPSLKSRX<sub>D8</sub>X<sub>D9</sub>IX<sub>D10</sub>QDTSKNQFX<sub>D11</sub>LX<sub>D12</sub>LX<sub>D13</sub>SVTX<sub>D14</sub>X<sub>D15</sub>D**  
**TAX<sub>D16</sub>YYCASSYDGLFAYWGQGTLVTVSX<sub>D17</sub>** (SEQ ID NO.:184);

Wherein X<sub>D1</sub> may be, for example, G or D;

Wherein X<sub>D2</sub> may be, for example, a neutral hydrophilic amino acid or more particularly T or S;

Wherein X<sub>D3</sub> may be, for example, a neutral hydrophilic amino acid or more particularly S or T;

Wherein X<sub>D4</sub> may be, for example, H or F;

Wherein X<sub>D5</sub> may be, for example, K or N;

Wherein X<sub>D6</sub> may be, for example, G or K;

Wherein X<sub>D7</sub> may be, for example, a hydrophobic amino acid or more particularly I or M;

Wherein X<sub>D8</sub> may be, for example, a hydrophobic amino acid or more particularly V or I;

Wherein X<sub>D9</sub> may be, for example, a neutral hydrophilic amino acid or more particularly T or S;

Wherein X<sub>D10</sub> may be, for example, a neutral hydrophilic amino acid or more particularly S or T;

Wherein X<sub>D11</sub> may be, for example, a neutral hydrophilic amino acid or more particularly S or F;

Wherein X<sub>D12</sub> may be, for example, a basic amino acid or more particularly K or Q;

Wherein X<sub>D13</sub> may be, for example, S or N;

Wherein X<sub>D14</sub> may be, for example, A or T;

Wherein  $X_{D15}$  may be, for example, A or E;

Wherein  $X_{D16}$  may be, for example, V or T and;

Wherein  $X_{D17}$  may be any amino acid, A or absent.

**[0239]** In accordance with a more specific disclosure, the humanized 3C4 antibody may have a heavy chain variable region of formula:

EVQLQESGPX<sub>d1</sub>LVKPSQX<sub>d2</sub>LSLTCTVX<sub>d3</sub>**GFSITSGYGYWHWIRQX<sub>d4</sub>PGX<sub>d5</sub>X<sub>d6</sub>LE**  
WX<sub>d7</sub>**GYINYGHNDYNPSLKSRX<sub>d8</sub>X<sub>d9</sub>IX<sub>d10</sub>QDTSKNQFX<sub>d11</sub>LX<sub>d12</sub>LX<sub>d13</sub>SVTX<sub>d14</sub>X<sub>d15</sub>DTAX**  
d<sub>16</sub>YYC**ASSYDGLFAYWQQGTLVTVSX<sub>d17</sub>** (SEQ ID NO.:185);

Wherein  $X_{d1}$  may be, for example, G or D;

Wherein  $X_{d2}$  may be, for example, T or S;

Wherein  $X_{d3}$  may be, for example, S or T;

Wherein  $X_{d4}$  may be, for example, H or F;

Wherein  $X_{d5}$  may be, for example, K or N;

Wherein  $X_{d6}$  may be, for example, G or K;

Wherein  $X_{d7}$  may be, for example, I or M;

Wherein  $X_{d8}$  may be, for example, V or I;

Wherein  $X_{d9}$  may be, for example, T or S;

Wherein  $X_{d10}$  may be, for example, S or T;

Wherein  $X_{d11}$  may be, for example, S or F;

Wherein  $X_{d12}$  may be, for example, K or Q;

Wherein  $X_{d13}$  may be, for example, S or N;

Wherein  $X_{d14}$  may be, for example, A or T;

Wherein  $X_{d15}$  may be, for example, A or E;

Wherein  $X_{d16}$  may be, for example, V or T and;

Wherein  $X_{d17}$ , A or absent.

**[0240]** Accordingly, disclosed herein is an antibody or antigen binding fragment thereof capable of specific binding to Kidney associated antigen 1 (KAAG1) which may have a light chain variable region at least 70% identical to SEQ ID NO.:16 and/or a heavy chain variable region at least 70% identical to SEQ ID NO.:18. The antibody or antigen binding fragment thereof may also comprise at least one amino acid substitution in comparison with SEQ ID NO.:16 or SEQ ID NO.:18.

**[0241]** Also disclosed herein is an antibody or antigen binding fragment thereof which may have a light chain variable region at least 70% identical to SEQ ID NO.:24 and/or a heavy chain variable region at least 70% identical to SEQ ID NO.:26. The antibody or antigen binding fragment thereof may also comprise at least one amino acid substitution in comparison with SEQ ID NO.:24 or SEQ ID NO.:26.

**[0242]** The present invention also provides in another embodiment, an antibody or antigen binding fragment thereof for use in accordance with the invention which may have a light chain variable region at least 70% identical to SEQ ID NO.:48 and/or a heavy chain variable region at least 70% identical to SEQ ID NO.:46. The antibody or antigen binding fragment thereof may also comprise at least one amino acid substitution in comparison with SEQ ID NO.:48 or SEQ ID NO.:46.

**[0243]** In accordance with an embodiment of the invention, the amino acid substitution is outside of a complementarity determining region (CDR). An antibody or antigen binding fragment having such an amino acid sequence encompasses, for example, a humanized antibody or antigen binding fragment.

**[0244]** As used herein the term "from one to twenty-five" includes every individual values and ranges such as for example, 1, 2, 3, and up to 25; 1 to 25; 1 to 24, 1 to 23, 1 to 22, 1 to 21, 1 to 20, 1 to 19; 1 to 18; 1 to 17; 1 to 16; 1 to 15 and so on; 2 to 25, 2 to 24, 2 to 23, 2 to 22, 2 to 21, 2 to 20; 2 to 19; 2 to 18; 2 to 17 and so on; 3 to 25, 3 to 24, 3 to 23, 3 to 22, 3 to 21, 3 to 20; 3 to 19; 3 to 18 and so on; 4 to 25, 4 to 24, 4 to 23, 4 to 22, 4 to 21, 4 to 20; 4 to 19; 4 to 18; 4 to 17; 4 to 16 and so on; 5 to 25, 5 to 24, 5 to 23, 5 to 22, 5 to 21, 5 to 20; 5 to 19; 5 to 18; 5 to 17 and so on, etc.

**[0245]** As used herein the term "from one to twenty-three" includes every individual values and ranges such as for example, 1, 2, 3, and up to 23; 1 to 23, 1 to 22, 1 to 21, 1 to 20, 1 to 19; 1 to 18; 1 to 17; 1 to 16; 1 to 15 and so on; 2 to 23, 2 to 22, 2 to 21, 2 to 20; 2 to 19; 2 to 18; 2 to 17 and so on; 3 to 23, 3 to 22, 3 to 21, 3 to 20; 3 to 19; 3 to 18 and so on; 4 to 23, 4 to 22, 4 to 21, 4 to 20; 4 to 19; 4 to 18; 4 to 17; 4 to 16 and so on; 5 to 25, 5 to 24, 5 to 23, 5 to 22, 5 to 21, 5 to 20; 5 to 19; 5 to 18; 5 to 17 and so on, etc.

**[0246]** As used herein the term "from one to twenty" includes every individual values and ranges such as for example, 1, 2, 3, and up to 20; 1 to 20; 1 to 19; 1 to 18; 1 to 17; 1 to 16; 1

to 15 and so on; 2 to 20; 2 to 19; 2 to 18; 2 to 17 and so on; 3 to 20; 3 to 19; 3 to 18 and so on; 4 to 20; 4 to 19; 4 to 18; 4 to 17; 4 to 16 and so on; 5 to 20; 5 to 19; 5 to 18; 5 to 17 and so on, etc.

**[0247]** Likewise, the term "from one to fifteen" includes every individual values and ranges such as for example, 1, 2, 3, and up to 15; 1 to 15; 1 to 14; 1 to 13; 1 to 12; 1 to 11; 1 to 10 and so on; 2 to 15; 2 to 14; 2 to 13; 2 to 12 and so on; 3 to 15; 3 to 14; 3 to 13 and so on; 4 to 15; 4 to 14; 4 to 13; 4 to 12; 4 to 11 and so on; 5 to 15; 5 to 14; 5 to 13; 5 to 12 and so on, etc.

**[0248]** Likewise, the term "from one to eleven" includes every individual values and ranges such as for example, 1, 2, 3, and up to 11; 1 to 11; 1 to 10, 1 to 9, 1 to 8, 1 to 7, and so on; 2 to 11; 2 to 10; 2 to 9; 2 to 8 and so on; 3 to 11; 3 to 10; 3 to 9 and so on; 4 to 11; 4 to 10; 4 to 9; 4 to 8; 4 to 7 and so on; 5 to 11; 5 to 10; 5 to 9; 5 to 8 and so on, etc.

**[0249]** In a more specific disclosure, the number of amino acid substitutions that may be accommodated in a humanized light chain variable region derived from SEQ ID NO.:16 may be for example, from 1 to 15 amino acid substitutions.

**[0250]** In yet a more specific disclosure, the number of amino acid substitutions that may be accommodated in a humanized heavy chain variable region derived from SEQ ID NO.:18 may be for example, from 1 to 20 amino acid substitutions. In some instances, when considering a humanized version of SEQ ID NO.:18, it may be useful to have at least three amino acid substitutions.

**[0251]** In a further more specific disclosure, the number of amino acid substitutions that may be accommodated in a humanized light chain variable region derived from SEQ ID NO.:24 may be for example, from 1 to 16 amino acid substitutions.

**[0252]** In yet a further more specific disclosure, the number of amino acid substitutions that may be accommodated in a humanized heavy chain variable region of SEQ ID NO.:26 may be for example, from 1 to 17 amino acid substitutions.

**[0253]** In a further more specific embodiment of the invention, the number of amino acid substitutions that may be accommodated in a humanized light chain variable region derived from SEQ ID NO.:48 may be for example, from 1 to 11 amino acid substitutions.

**[0254]** In yet a further more specific embodiment of the invention, the number of amino acid substitutions that may be accommodated in a humanized heavy chain variable region of SEQ ID NO.:46 may be for example, from 1 to 23 amino acid substitutions.

**[0255]** In accordance with an embodiment of the invention, the one to twenty amino acid substitutions may be for example, in the light chain variable region.

**[0256]** In accordance with an embodiment of the invention, the one to twenty amino acid

substitutions may be for example, in the heavy chain variable region.

**[0257]** A humanized antibody or antigen binding fragment may therefore have a light chain variable region having up to twenty amino acid substitutions in comparison with SEQ ID NO.:16 or SEQ ID NO.:24 and may have a heavy chain variable region having up to twenty amino acid substitutions in comparison with SEQ ID NO.:18 or SEQ ID NO.:26. A humanized antibody or antigen binding fragment may therefore have a light chain variable region having up to twenty-five amino acid substitutions in comparison with SEQ ID NO.:48 and may have a heavy chain variable region having up to twenty-five amino acid substitutions in comparison with SEQ ID NO.:46.

**[0258]** It is to be understood herein that when the humanized antibody or antigen binding fragment has two light chain variable regions and two heavy chain variable regions, each one of the light chain variable regions may independently have up to twenty-five, twenty-four, twenty-three, twenty-two, twenty-one, twenty, nineteen, eighteen, seventeen, sixteen, fifteen, fourteen, thirteen, twelve, eleven, ten, nine, eight, seven, six, five, four, three, two, one amino acid substitutions and each one of the heavy chain variable regions may have up to twenty-five, twenty-four, twenty-three, twenty-two, twenty-one, twenty, nineteen, eighteen, seventeen, sixteen, fifteen, fourteen, thirteen, twelve, eleven, ten, nine, eight, seven, six, five, four, three, two, one amino acid substitutions.

**[0259]** As discussed herein the amino acid substitutions may be conservative or non-conservative. In an exemplary embodiment the amino acid substitutions may be conservative.

**[0260]** It is to be understood herein that the humanized antibody or antigen binding fragment disclosed herein may also have a light chain variable region and/or heavy chain variable region showing a deletion in comparison with SEQ ID NO.:16, SEQ ID NO.:18, SEQ ID NO.:189, SEQ ID NO.:190, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196, SEQ ID NO.:197, SEQ ID NO.:24 and/or SEQ ID NO.:26. It is to be understood herein that the humanized antibody or antigen binding fragment for use in accordance with the invention may also have a light chain variable region and/or heavy chain variable region showing a deletion in comparison with SEQ ID NO.:189, SEQ ID NO.:190, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 and/or SEQ ID NO.:197. Such deletion may be found, for example, at an amino- or carboxy-terminus of the light chain variable region and/or heavy chain variable region.

**[0261]** Another exemplary embodiment of the humanized antibody or antigen binding fragment for use in accordance with the present invention includes for example, an antibody or antigen binding fragment having a light chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:186, SEQ ID NO.:187, SEQ ID NO.:188, SEQ ID NO.:189 or SEQ ID NO.:190.

**[0262]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:186" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, or at least 112 consecutive amino acids". The term "at least 90

consecutive amino acids of SEQ ID NO.:186" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:186 and especially those sequences which include the 3 CDRs of SEQ ID NO.:186, such as, for example a sequence comprising amino acids 6 to 108, 5 to 109, 13 to 103, 14 to 111 of SEQ ID NO.:186 and so on.

**[0263]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:187" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, or at least 112 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:187" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:187 and especially those sequences which include the 3 CDRs of SEQ ID NO.:187, such as, for example a sequence comprising amino acids 7 to 109, 12 to 104, 22 to 113, 18 to 112 of SEQ ID NO.:187 and so on.

**[0264]** The terms "at least 90 consecutive amino acids of SEQ ID NO.:188", "at least 90 consecutive amino acids of SEQ ID NO.:189" or "at least 90 consecutive amino acids of SEQ ID NO.:190" has a similar meaning.

**[0265]** In accordance with the present invention, the antibody or antigen binding fragment of the present invention may have, for example, a light chain variable region as set forth in SEQ ID NO.:189 or 190.

**[0266]** The humanized antibody or antigen binding fragment for use in accordance with the invention includes (or further includes) for example, a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NOs.:191, 192, 193, 194, 195, 196 or 197.

**[0267]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:191" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115 or at least 116 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:191" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:191 and especially those sequences which include the 3 CDRs of SEQ ID NO.:191, such as, for example a sequence comprising amino acids 1 to 106, 2 to 112, 11 to 113, 7 to 102 of SEQ ID NO.:191 and so on.

**[0268]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:192" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115 or at least 116 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:192" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:192 and especially those sequences which include the 3 CDRs of SEQ ID NO.:192, for example a sequence comprising amino acids 6 to 109, 8 to 113, 1 to 102, 2 to 105 of SEQ ID NO.:192 and so on.

**[0269]** The terms "at least 90 consecutive amino acids of SEQ ID NO.: 193", "at least 90

consecutive amino acids of SEQ ID NO.:194", "at least 90 consecutive amino acids of SEQ ID NO.:195", "at least 90 consecutive amino acids of SEQ ID NO.:196" or "at least 90 consecutive amino acids of SEQ ID NO.:197" has a similar meaning.

**[0270]** In accordance with the present invention, the antibody or antigen binding fragment of the present invention may have, for example, a heavy chain variable region as set forth in SEQ ID NO.:194, 195, 196 or 197.

**[0271]** In accordance with the present invention the antibody or antigen binding fragment may comprise, for example,

1. a) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:186 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:191, SEQ ID NO.:192, SEQ ID NO.:193, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 or SEQ ID NO.:197;
2. b) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:187 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:191, SEQ ID NO.:192, SEQ ID NO.:193, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 or SEQ ID NO.:197;
3. c) a light chain variable region which may comprise amino acids at least 90 consecutive amino acids of SEQ ID NO.:188 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:191, SEQ ID NO.:192, SEQ ID NO.:193, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 or SEQ ID NO.:197;
4. d) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:189 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:191, SEQ ID NO.:192, SEQ ID NO.:193, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 or SEQ ID NO.:197 or
5. e) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:190 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:191, SEQ ID NO.:192, SEQ ID NO.:193, SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 or SEQ ID NO.:197.

**[0272]** In accordance with a more specific embodiment of the invention, the light chain variable region may comprise at least 90 consecutive amino acids of SEQ ID NO.:189 or 190 and the heavy chain variable region may comprise at least 90 consecutive amino acids of SEQ ID NO.:194, 195, 196 or 197.

**[0273]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:189 and the heavy chain variable region may be as set forth in SEQ ID NO.:194.

**[0274]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:189 and the heavy chain variable region

may be as set forth in SEQ ID NO.:195.

**[0275]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:189 and the heavy chain variable region may be as set forth in SEQ ID NO.:196.

**[0276]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:189 and the heavy chain variable region may be as set forth in SEQ ID NO.:197.

**[0277]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:190 and the heavy chain variable region may be as set forth in SEQ ID NO.:194.

**[0278]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:190 and the heavy chain variable region may be as set forth in SEQ ID NO.:195.

**[0279]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:190 and the heavy chain variable region may be as set forth in SEQ ID NO.:196.

**[0280]** In accordance with an even more specific embodiment of the invention, the light chain variable region may be as set forth in SEQ ID NO.:190 and the heavy chain variable region may be as set forth in SEQ ID NO.:197.

**[0281]** Another example of the humanized antibody or antigen binding fragment of the present disclosure includes for example, an antibody or antigen binding fragment having a light chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:174, SEQ ID NO.:175, SEQ ID NO.:176 or SEQ ID NO.:168.

**[0282]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:174" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112 or at least 113 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:174" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:174 and especially those sequences which include the 3 CDRs of SEQ ID NO.:174, such as, for example a sequence comprising amino acids 6 to 108, 5 to 109, 13 to 103, 14 to 111 of SEQ ID NO.:174 and so on.

**[0283]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:175" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112 or at least 113 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:175" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:175 and especially those sequences

which include the 3 CDRs of SEQ ID NO.:175, such as, for example a sequence comprising amino acids 7 to 109, 12 to 104, 22 to 113, 18 to 112 of SEQ ID NO.:175 and so on.

**[0284]** The terms "at least 90 consecutive amino acids of SEQ ID NO.:176" or "at least 90 consecutive amino acids of SEQ ID NO.:168" has a similar meaning.

**[0285]** In accordance with the present disclosure, the antibody or antigen binding fragment of the present invention may have, for example, a light chain variable region as set forth in SEQ ID NO.:168.

**[0286]** The humanized antibody or antigen binding fragment of the present disclosure includes (or further includes) for example, a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NOs.:177, 178, 179 or 169.

**[0287]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:177" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112 or at least 113 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:177" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:177 and especially those sequences which include the 3 CDRs of SEQ ID NO.:177, such as, for example a sequence comprising amino acids 1 to 106, 2 to 112, 11 to 113, 7 to 102 of SEQ ID NO.:177 and so on.

**[0288]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:178" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112 or at least 113 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:178" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:178 and especially those sequences which include the 3 CDRs of SEQ ID NO.:178, for example a sequence comprising amino acids 6 to 109, 8 to 113, 1 to 102, 2 to 105 of SEQ ID NO.:178 and so on.

**[0289]** The terms "at least 90 consecutive amino acids of SEQ ID NO.:179" or "at least 90 consecutive amino acids of SEQ ID NO.:169" has a similar meaning.

**[0290]** In accordance with the present disclosure, the antibody or antigen binding fragment of the present invention may have, for example, a heavy chain variable region as set forth in SEQ ID NO.:169.

**[0291]** In accordance with the present disclosure the antibody or antigen binding fragment may comprise, for example,

f) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:174 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:177, SEQ ID NO.:178, SEQ ID NO.:179 or SEQ ID NO.:169;

g) a light chain variable region which may comprise at least 90 consecutive amino acids of

SEQ ID NO.:175 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:177, SEQ ID NO.:178, SEQ ID NO.:179 or SEQ ID NO.:169;

h) a light chain variable region which may comprise amino acids at least 90 consecutive amino acids of SEQ ID NO.:176 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:177, SEQ ID NO.:178, SEQ ID NO.:179 or SEQ ID NO.:169 or;

i) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:168 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:177, SEQ ID NO.:178, SEQ ID NO.:179 or SEQ ID NO.:169.

**[0292]** In accordance with a more specific disclosure, the light chain variable region may comprise at least 90 consecutive amino acids of SEQ ID NO.:168 and the heavy chain variable region may comprise at least 90 consecutive amino acids of SEQ ID NO.:169.

**[0293]** In accordance with an even more specific disclosure, the light chain variable region may be as set forth in SEQ ID NO.:168 and the heavy chain variable region may be as set forth in SEQ ID NO.:169.

**[0294]** Other examples of the humanized antibodies or antigen binding fragments of the present disclosure are those which may comprise a light chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID Nos. 180, 181, 182 or 172.

**[0295]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:180" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106 or at least 107, consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:180" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:180 and especially those sequences which include the 3 CDRs of SEQ ID NO.:180, for example a sequence comprising amino acids 6 to 102, 11 to 106, 1 to 106, 3 to 95, 5 to 95 of SEQ ID NO.:180 and so on.

**[0296]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:181" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106 or at least 107, consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:181" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:181 and especially those sequences which include the 3 CDRs of SEQ ID NO.:181, for example a sequence comprising amino acids 9 to 106, 10 to 101, 1 to 98, 3 to 99, 7 to 107 of SEQ ID NO.:181 and so on.

**[0297]** The terms "at least 90 consecutive amino acids of SEQ ID NO.:182" or "at least 90 consecutive amino acids of SEQ ID NO.:172" has a similar meaning.

**[0298]** In accordance with the present disclosure, the antibody or antigen binding fragment of the present invention may have, for example, a light chain variable region as set forth in SEQ ID NO.:172.

**[0299]** The humanized antibody or antigen binding fragment of the present disclosure includes (or further includes) for example, a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NOs.:183, 184, 185 or 173.

**[0300]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:183" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115 or at least 116 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:183" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:183 and especially those sequences which include the 3 CDRs of SEQ ID NO.:183, such as, for example a sequence comprising amino acids 6 to 111, 1 to 106, 2 to 104, 5 to 106, 10 to 107 of SEQ ID NO.:183 and so on.

**[0301]** As used herein the term "at least 90 consecutive amino acids of SEQ ID NO.:185" also includes the terms "at least 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115 or at least 116 consecutive amino acids". The term "at least 90 consecutive amino acids of SEQ ID NO.:185" encompasses any possible sequence of at least 90 consecutive amino acids found in SEQ ID NO.:185 and especially those sequences which include the 3 CDRs of SEQ ID NO.:185, such as, for example a sequence comprising amino acids 3 to 107, 1 to 115, 1 to 110, 22 to 116, 20 to 115 of SEQ ID NO.:185 and so on.

**[0302]** The terms "at least 90 consecutive amino acids of SEQ ID NO.:184" or "at least 90 consecutive amino acids of SEQ ID NO.:173" has a similar meaning.

**[0303]** In accordance with the present disclosure, the antibody or antigen binding fragment of the present invention may have, for example, a heavy chain variable region as set forth in SEQ ID NO.:173.

**[0304]** In accordance with the present disclosure the antibody or antigen binding fragment may comprise, for example,

1. a) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:180 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:183, SEQ ID NO.:184, SEQ ID NO.:185 or SEQ ID NO.:173;
2. b) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:181 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:183, SEQ ID NO.:184, SEQ ID NO.:185 or SEQ ID NO.:173;

3. c) a light chain variable region which may comprise amino acids at least 90 consecutive amino acids of SEQ ID NO.:182 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:183, SEQ ID NO.:184, SEQ ID NO.:185 or SEQ ID NO.:173 or;
4. d) a light chain variable region which may comprise at least 90 consecutive amino acids of SEQ ID NO.:172 and a heavy chain variable region which may comprise at least 90 consecutive amino acids of any of SEQ ID NO.:183, SEQ ID NO.:184, SEQ ID NO.:185 or SEQ ID NO.:173.

**[0305]** In accordance with a more specific disclosure, the light chain variable region may have at least 90 consecutive amino acids of SEQ ID NO.:172 and the heavy chain variable region may have at least 90 consecutive amino acids of SEQ ID NO.:173.

**[0306]** In accordance with an even more specific disclosure, the light chain variable region may be as set forth in SEQ ID NO.:172 and the heavy chain variable region may be as set forth in SEQ ID NO.:173.

**[0307]** The antibody or antigen binding fragment of the present invention may have a light chain variable region and/or heavy chain variable region in accordance with the invention as described above and may further comprise amino acids of a constant region, such as, for example, amino acids of a constant region of a human antibody.

**[0308]** In an exemplary embodiment, the antibody or antigen binding fragment of the present invention may comprise, for example, a human IgG1 constant region.

**[0309]** In accordance with another exemplary embodiment of the invention, the antigen binding fragment may be, for example, a scFv, a Fab, a Fab' or a (Fab')<sub>2</sub>.

**Production of the antibodies in cells**

**[0310]** The anti-KAAG1 antibodies that are disclosed herein can be made by a variety of methods familiar to those skilled in the art, such as hybridoma methodology or by recombinant DNA methods.

**[0311]** The anti-KAAG1 antibodies may be produced by the conventional hybridoma technology, where a mouse is immunized with an antigen, spleen cells isolated and fused with myeloma cells lacking HGPRT expression and hybrid cells selected by hypoxanthine, aminopterin and thymine (HAT) containing media.

**[0312]** The anti-KAAG1 antibodies may be produced by recombinant DNA methods.

**[0313]** In order to express the anti-KAAG1 antibodies, nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein or any other may be inserted into an expression vector, i.e., a vector that contains the elements for transcriptional and translational control of the inserted coding sequence in a particular host. These elements may include regulatory sequences, such as enhancers, constitutive and inducible promoters, and 5' and 3' un-translated regions. Methods that are well known to those skilled in the art may be used to construct such expression vectors. These methods include *in vitro* recombinant DNA techniques, synthetic techniques, and *in vivo* genetic recombination.

**[0314]** A variety of expression vector/host cell systems known to those of skill in the art may be utilized to express a polypeptide or RNA derived from nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein. These include, but are not limited to, microorganisms such as bacteria transformed with recombinant bacteriophage, plasmid, or cosmid DNA expression vectors; yeast transformed with yeast expression vectors; insect cell systems infected with baculovirus vectors; plant cell systems transformed with viral or bacterial expression vectors; or animal cell systems. For long-term production of recombinant proteins in mammalian systems, stable expression in cell lines may be effected. For example, nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein may be transformed into cell lines using expression vectors that may contain viral origins of replication and/or endogenous expression elements and a selectable or visible marker gene on the same or on a separate vector. The invention is not to be limited by the vector or host cell employed. In certain examples, the nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein may each be ligated into a separate expression vector and each chain expressed separately. In another example, both the light and heavy chains able to encode any one of a light and heavy immunoglobulin chains described herein may be ligated into a single expression vector and expressed simultaneously.

**[0315]** Alternatively, RNA and/or polypeptide may be expressed from a vector comprising nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein using an *in vitro* transcription system or a coupled *in vitro* transcription/translation system respectively.

**[0316]** In general, host cells that contain nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein and/or that express a polypeptide encoded by the nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein, or a portion thereof, may be identified by a variety of procedures known to those of skill in the art. These procedures include, but are not limited to, DNA/DNA or DNA/RNA hybridizations, PCR amplification, and protein bioassay or immunoassay techniques that include membrane, solution, or chip based technologies for the detection and/or quantification of nucleic acid or amino acid sequences. Immunological methods for detecting and measuring the expression of polypeptides using either specific polyclonal or monoclonal antibodies are known in the art. Examples of such techniques include enzyme-linked immunosorbent assays (ELISAs), radioimmunoassays (RIAs), and fluorescence

activated cell sorting (FACS). Those of skill in the art may readily adapt these methodologies to the present invention.

**[0317]** Host cells comprising nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein may thus be cultured under conditions for the transcription of the corresponding RNA (mRNA, siRNA, shRNA etc.) and/or the expression of the polypeptide from cell culture. The polypeptide produced by a cell may be secreted or may be retained intracellularly depending on the sequence and/or the vector used. In an exemplary disclosure, expression vectors containing nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein may be designed to contain signal sequences that direct secretion of the polypeptide through a prokaryotic or eukaryotic cell membrane.

**[0318]** Due to the inherent degeneracy of the genetic code, other DNA sequences that encode the same, substantially the same or a functionally equivalent amino acid sequence may be produced and used, for example, to express a polypeptide encoded by nucleotide sequences able to encode any one of a light and heavy immunoglobulin chains described herein. The nucleotide sequences disclosed herein may be engineered using methods generally known in the art in order to alter the nucleotide sequences for a variety of purposes including, but not limited to, modification of the cloning, processing, and/or expression of the gene product. DNA shuffling by random fragmentation and PCR reassembly of gene fragments and synthetic oligonucleotides may be used to engineer the nucleotide sequences. For example, oligonucleotide-mediated site-directed mutagenesis may be used to introduce mutations that create new restriction sites, alter glycosylation patterns, change codon preference, produce splice variants, and so forth.

**[0319]** In addition, a host cell strain may be chosen for its ability to modulate expression of the inserted sequences or to process the expressed polypeptide in the desired fashion. Such modifications of the polypeptide include, but are not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation, and acylation. In an exemplary embodiment, anti-KAAG1 antibodies that contain particular glycosylation structures or patterns may be desired. Post-translational processing, which cleaves a "prepro" form of the polypeptide, may also be used to specify protein targeting, folding, and/or activity. Different host cells that have specific cellular machinery and characteristic mechanisms for post-translational activities (e.g., CHO, HeLa, MDCK, HEK293, and W138) are available commercially and from the American Type Culture Collection (ATCC) and may be chosen to ensure the correct modification and processing of the expressed polypeptide.

**[0320]** Those of skill in the art will readily appreciate that natural, modified, or recombinant nucleic acid sequences may be ligated to a heterologous sequence resulting in translation of a fusion polypeptide containing heterologous polypeptide moieties in any of the aforementioned host systems. Such heterologous polypeptide moieties may facilitate purification of fusion polypeptides using commercially available affinity matrices. Such moieties include, but are not limited to, glutathione S-transferase (GST), maltose binding protein, thioredoxin, calmodulin

binding peptide, 6-His (His), FLAG, c-myc, hemagglutinin (HA), and antibody epitopes such as monoclonal antibody epitopes.

**[0321]** Also disclosed herein is a polynucleotide which may comprise a nucleotide sequence encoding a fusion protein. The fusion protein may comprise a fusion partner (e.g., HA, Fc, etc.) fused to the polypeptide (e.g., complete light chain, complete heavy chain, variable regions, CDRs etc.) described herein.

**[0322]** Those of skill in the art will also readily recognize that the nucleic acid and polypeptide sequences may be synthesized, in whole or in part, using chemical or enzymatic methods well known in the art. For example, peptide synthesis may be performed using various solid-phase techniques and machines such as the ABI 431A Peptide synthesizer (PE Biosystems) may be used to automate synthesis. If desired, the amino acid sequence may be altered during synthesis and/or combined with sequences from other proteins to produce a variant protein.

### **Antibody conjugates**

**[0323]** The antibody or antigen binding fragment disclosed herein may be conjugated with a detectable moiety (i.e., for detection or diagnostic purposes) or with a therapeutic moiety (for therapeutic purposes). In accordance with the present invention the antibody or antigen binding fragment is conjugated with a therapeutic moiety (for therapeutic purposes).

**[0324]** A "detectable moiety" is a moiety detectable by spectroscopic, photochemical, biochemical, immunochemical, chemical and/or other physical means. A detectable moiety may be coupled either directly and/or indirectly (for example via a linkage, such as, without limitation, a DOTA or NHS linkage) to antibodies and antigen binding fragments thereof of the present invention using methods well known in the art. A wide variety of detectable moieties may be used, with the choice depending on the sensitivity required, ease of conjugation, stability requirements and available instrumentation. A suitable detectable moiety include, but is not limited to, a fluorescent label, a radioactive label (for example, without limitation, <sup>125</sup>I, <sup>111</sup>In, <sup>99</sup>Tc, <sup>131</sup>I and including positron emitting isotopes for PET scanner etc), a nuclear magnetic resonance active label, a luminescent label, a chemiluminescent label, a chromophore label, an enzyme label (for example and without limitation horseradish peroxidase, alkaline phosphatase, etc.), quantum dots and/or a nanoparticle. Detectable moiety may cause and/or produce a detectable signal thereby allowing for a signal from the detectable moiety to be detected.

**[0325]** In another exemplary embodiment of the invention, the antibody or antigen binding fragment thereof may be coupled (modified) with a therapeutic moiety (e.g., drug, cytotoxic moiety).

**[0326]** In an exemplary embodiment, the anti-KAAG1 antibodies and antigen binding

fragments may comprise an inhibitor, a chemotherapeutic or cytotoxic agent. For example, the antibody and antigen binding fragments may be conjugated to the chemotherapeutic or cytotoxic agent. Such chemotherapeutic or cytotoxic agents include, but are not limited to, Yttrium-90, Scandium-47, Rhenium-186, Iodine-131, Iodine-125, and many others recognized by those skilled in the art (e.g., lutetium (e.g., Lu<sup>177</sup>), bismuth (e.g., Bi<sup>213</sup>), copper (e.g., Cu<sup>67</sup>)). In other instances, the chemotherapeutic or cytotoxic agent may comprise, without limitation, 5-fluorouracil, adriamycin, irinotecan, platinum-based compounds such as cisplatin and anti-tubulin or anti-mitotic compounds such as, taxanes, doxorubicin and cyclophosphamide, pseudomonas endotoxin, ricin and other toxins. Suitable antibody drug conjugates are selected amongst those having an IC<sub>50</sub> in the range of 0.001nM to 150nM, 0.001nM to 100nM, 0.001nM to 50nM, 0.001nM to 20nM or 0.001nM to 10nM (inclusively). The cytotoxic drug used for conjugation is thus selected on the basis of these criteria.

**[0327]** Alternatively, in order to carry out the therapeutic uses of the present invention and as known in the art, the antibody or antigen binding fragment of the present invention (conjugated) may be used in combination with a second molecule (e.g., a secondary antibody, etc.) which is able to specifically bind to the antibody or antigen binding fragment of the present invention and which may carry a desirable detectable, diagnostic or therapeutic moiety.

**Pharmaceutical compositions of the antibodies and their use**

**[0328]** Pharmaceutical compositions of the anti-KAAG1 antibodies or antigen binding fragments (conjugated or not) are also disclosed herein. The pharmaceutical composition may comprise an anti-KAAG1 antibody or an antigen binding fragment and may also contain a pharmaceutically acceptable carrier.

**[0329]** Other disclosures relate to a composition which may comprise the antibody or antigen binding fragment described herein and a carrier.

**[0330]** The present disclosure also relates to a pharmaceutical composition which may comprise the antibody or antigen binding fragment described herein and a pharmaceutically acceptable carrier.

**[0331]** In addition to the active ingredients, a pharmaceutical composition may contain pharmaceutically acceptable carriers comprising water, PBS, salt solutions, gelatins, oils, alcohols, and other excipients and auxiliaries that facilitate processing of the active compounds into preparations that may be used pharmaceutically. In other instances, such preparations may be sterilized.

**[0332]** As used herein, "pharmaceutical composition" means therapeutically effective amounts of the agent together with pharmaceutically acceptable diluents, preservatives, solubilizers, emulsifiers, adjuvant and/or carriers. A "therapeutically effective amount" as used herein refers

to that amount which provides a therapeutic effect for a given condition and administration regimen. Such compositions are liquids or lyophilized or otherwise dried formulations and include diluents of various buffer content (e.g., Tris-HCl., acetate, phosphate), pH and ionic strength, additives such as albumin or gelatin to prevent absorption to surfaces, detergents (e.g., Tween 20, Tween 80, Pluronic F68, bile acid salts). Solubilizing agents (e.g., glycerol, polyethylene glycerol), anti-oxidants (e.g., ascorbic acid, sodium metabisulfite), preservatives (e.g., thimerosal, benzyl alcohol, parabens), bulking substances or tonicity modifiers (e.g., lactose, mannitol), covalent attachment of polymers such as polyethylene glycol to the protein, complexation with metal ions, or incorporation of the material into or onto particulate preparations of polymeric compounds such as polylactic acid, polyglycolic acid, hydrogels, etc, or onto liposomes, microemulsions, micelles, unilamellar or multilamellar vesicles, erythrocyte ghosts, or spheroplasts. Such compositions will influence the physical state, solubility, stability, rate of in vivo release, and rate of in vivo clearance. Controlled or sustained release compositions include formulation in lipophilic depots (e.g., fatty acids, waxes, oils). Also disclosed are particulate compositions coated with polymers (e.g., poloxamers or poloxamines). Other disclosed compositions incorporate particulate forms protective coatings, protease inhibitors or permeation enhancers for various routes of administration, including parenteral,

pulmonary, nasal, oral, vaginal, rectal routes. In one example the pharmaceutical composition is administered parenterally, paracancerally, transmucosally, transdermally, intramuscularly, intravenously, intradermally, subcutaneously, intraperitonealy, intraventricularly, intracranially and intratumorally.

**[0333]** Further, as used herein "pharmaceutically acceptable carrier" or "pharmaceutical carrier" are known in the art and include, but are not limited to, 0.01-0.1 M or 0.05 M phosphate buffer or 0.8 % saline. Additionally, such pharmaceutically acceptable carriers may be 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, antioxidants, collating agents, inert gases and the like.

**[0334]** For any compound, the therapeutically effective dose may be estimated initially either in cell culture assays or in animal models such as mice, rats, rabbits, dogs, or pigs. An animal model may also be used to determine the concentration range and route of administration. Such information may then be used to determine useful doses and routes for administration in humans. These techniques are well known to one skilled in the art and a therapeutically effective dose refers to that amount of active ingredient that ameliorates the symptoms or condition. Therapeutic efficacy and toxicity may be determined by standard pharmaceutical procedures in cell cultures or with experimental animals, such as by calculating and contrasting

the ED<sub>50</sub> (the dose therapeutically effective in 50% of the population) and LD<sub>50</sub> (the dose lethal to 50% of the population) statistics. Any of the therapeutic compositions described above may be applied to any subject in need of such therapy, including, but not limited to, mammals such as dogs, cats, cows, horses, rabbits, monkeys, and humans.

**[0335]** The pharmaceutical compositions utilized in this invention may be administered by any number of routes including, but not limited to, oral, intravenous, intramuscular, intra-arterial, intramedullary, intrathecal, intraventricular, transdermal, subcutaneous, intraperitoneal, intranasal, enteral, topical, sublingual, or rectal means.

**Methods of use**

**[0336]** The term "treatment" for purposes of this disclosure refers to both therapeutic treatment and prophylactic or preventative measures, wherein the object is to prevent or slow down (lessen) the targeted pathologic condition or disorder. Those in need of treatment include those already having the disorder as well as those prone to have the disorder or those in whom the disorder is to be prevented.

**[0337]** The present disclosure provides a method of treating an individual having or suspected of having breast cancer with an antibody or antigen binding fragment which is capable of specific binding to KAAG1.

**[0338]** In accordance with the present invention, the individual may have a breast cancer that is negative for the estrogen receptor expression, the progesterone receptor expression and/or Her2 expression (or overexpression).

**[0339]** Also in accordance with the present invention, the individual may have a breast cancer that has low expression for at least one of estrogen receptor, progesterone receptor and/or Her2.

**[0340]** For example, the tumor may be negative for (or have low expression of) both estrogen receptor expression and progesterone receptor expression.

**[0341]** In accordance with the present invention, the individual may have a breast cancer that is characterized as being triple-negative. The breast cancer may be basal-like.

**[0342]** Also disclosed herein is the use of the isolated antibody or antigen binding fragment described herein in the treatment or diagnosis of breast cancer characterized by a lack of estrogen receptor expression, progesterone receptor expression and/or Her2 overexpression or by low expression of at least one of these three markers.

**[0343]** In accordance with the present invention, the therapeutic use may comprise, for example, administering an antibody or antigen binding fragment which is capable of specific

binding to KAAG1 in accordance with the invention to an individual in need. The individual in need is preferentially selected on the basis of a lack of ER expression, PgR expression and/or by the absence of HER2 protein over-expression. Clinical testing for these markers is usually performed using histopathologic methods (immunohistochemistry, FISH, etc.) and/or by gene expression studies (see for example Dent et al, 2007, Bernstein and Lacey, 2011). The individual in need may thus be an individual who has received a diagnosis of triple-negative breast cancer. The cancer may be basal-like breast cancer.

**[0344]** The present invention thus particularly relates to the therapeutic treatment of individual having triple-negative breast cancer with an anti-KAAG1 antibody. The cancer may be basal-like cancer.

**[0345]** Suitable antibodies or antigen binding fragments include those that are capable of specific binding to KAAG1 at the surface of tumor cells. Such antibodies may preferentially bind an epitope included within amino acids 30 to 84 of KAAG1 inclusively (e.g., within amino acids 36 to 60 (inclusively) or within amino acids 61 to 84 (inclusively) of KAAG1).

**[0346]** Suitable antibodies may be those which mediate antibody-dependent cell cytotoxicity and those that are conjugated with a therapeutic moiety.

**[0347]** In accordance with the present invention, the antibody may be, for example, a monoclonal antibody, a chimeric antibody or a humanized antibody or an antigen binding fragment thereof.

**[0348]** The therapeutic use of the present invention may include administering the antibody or antigen binding fragment in combination with an inhibitor, a chemotherapeutic or a cytotoxic agent.

**[0349]** Other methods of treatment disclosed herein include administering other types of KAAG1 inhibitors such as antisense-based therapeutics (siRNA, antisenses, ribozymes, etc.).

**[0350]** Disclosed herein is a method of treating triple-negative breast cancer or basal-like breast cancer by administering an inhibitor of KAAG1 activity or expression to an individual in need.

**[0351]** The inhibitor may comprise a nucleotide sequence complementary to SEQ ID NO.:1 or to a fragment thereof. More particularly, the inhibitor may comprise a nucleotide sequence complementary to nucleotides 738 to 992 (inclusively) of SEQ ID NO.:1 or to a fragment thereof. For example, the inhibitor may include at least 10 consecutive nucleotides (at least 15, at least 20) which are complementary to SEQ ID NO.:1 or to nucleotides 738 to 992 (inclusively) of SEQ ID NO.:1.

**[0352]** In certain instances, the anti-KAAG1 antibodies and fragments may interact with cancer cells that express KAAG1 and induce an immunological reaction by mediating ADCC. In other

instances, the anti-KAAG1 antibodies and fragments may block the interaction of KAAG1 with its protein partners.

**[0353]** In certain instances, the anti-KAAG1 antibodies and antigen binding fragments thereof may be administered concurrently with other treatments given for the same condition (inhibitors, chemotherapeutics or cytotoxic agents). As such, the antibodies may be administered with a PARP1 inhibitor, a EGFR inhibitor, anti-mitotics (eg., taxanes), platinum-based agents (eg., cisplatin), DNA damaging agents (eg. Doxorubicin) and other anti-cancer therapies that are known to those skilled in the art. In other instances, the anti-KAAG1 antibodies and antigen binding fragments thereof may be administered with other therapeutic antibodies. These include, but are not limited to, antibodies that target EGFR, CD-20, and Her2.

**[0354]** Disclosed herein is a method for inhibiting the growth of KAAG1-expressing cell that are estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-), the method may comprise contacting the cell with an effective amount of the antibody or antigen binding fragment described herein.

**[0355]** Also disclosed herein is a method of treating cancer or inhibiting the growth of a KAAG1 expressing cells that are estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-), in a mammal, the method may comprise administering the antibody or antigen binding fragment described herein to a mammal in need.

**[0356]** The present disclosure provides method of treatment, diagnostic methods and method of detection using the antibody or antigen binding fragment disclosed herein and the use of these antibodies or antigen binding fragment in the manufacture of a pharmaceutical composition or drug for such purposes.

**[0357]** Method of treatment disclosed herein includes administering an antibody or antigen binding fragment described herein to a mammal in need, and especially to a patient having or susceptible of having a cancer characterized as being estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-),

**[0358]** Also disclosed herein are methods for reducing tumor spread, tumor invasion, tumor formation or for inducing tumor lysis, which may comprise administering an isolated antibody or antigen binding fragment to a mammal in need.

**[0359]** Disclosed herein is the use of the isolated antibody or antigen binding fragment described herein in the (manufacture of a pharmaceutical composition for) treatment of cancer, reduction of tumor spread, tumor invasion, tumor formation or for inducing tumor lysis of KAAG1-expressing tumor cells that are estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-).

**[0360]** The antibody or antigen binding fragment may more particularly be applicable for

malignant tumor including, for example, a malignant tumor having the ability to metastasize and/or tumor cells characterized by anchorage-independent growth. The antibody or antigen binding fragment disclosed herein may also be used in the diagnosis of cancer. The diagnosis of cancer may be performed *in vivo* by administering the antibody or antigen binding fragment disclosed herein to a mammal having or suspected of having a cancer. The diagnosis may also be performed *ex vivo* by contacting a sample obtained from the mammal with the antibody or antigen binding fragment and determining the presence or absence of cells (tumor cells) expressing KAAG1 or a KAAG1 variant.

**[0361]** Also disclosed is a method of detecting cancer or detecting a KAAG1 expressing cells that are estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-), in a mammal, the method may comprise administering the antibody or antigen binding fragment described herein to a mammal in need.

**[0362]** Also disclosed is a method for detecting a cell expressing KAAG1 or a KAAG1 variant, the method may comprise contacting the cell with an antibody or antigen binding fragment described herein and detecting a complex formed by the antibody and the KAAG1-or KAAG1 variant-expressing cell. Exemplary embodiments of antibodies or antigen binding fragments used in detection methods are those which are capable of binding to the extracellular region of KAAG1.

**[0363]** Other exemplary embodiments of antibodies or antigen binding fragments used in detection methods are those which bind to KAAG1 or KAAG1 variant expressed at the surface of tumor cells that are estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-).

**[0364]** Also disclosed is a method for detecting KAAG1 (SEQ ID NO.:2), a KAAG1 variant having at least 80% sequence identity with SEQ ID NO.:2 or a secreted form of circulating form of KAAG1 or KAAG1 variant, the method may comprise contacting a cell expressing KAAG1 or the KAAG1 variant or a sample (biopsy, serum, plasma, urine etc.) comprising or suspected of comprising KAAG1 or the KAAG1 variant with the antibody or antigen binding fragments described herein and measuring binding. The sample may originate from a mammal (e.g., a human) which may have cancer (e.g., breast cancer that is characterized as being estrogen receptor-negative (ER-), progesterone receptor negative (PgR-) and/or that lacks Her2 overexpression (Her2-), such as basal-like breast cancer or triple-negative breast cancer) or may be suspected of having cancer. The sample may be a tissue sample obtained from the mammal or a cell culture supernatant.

**[0365]** The sample may be a serum sample, a plasma sample, a blood sample or ascitic fluid obtained from the mammal. The antibody or antigen binding fragment described herein may advantageously detect a secreted or circulating form (circulating in blood) of KAAG1.

**[0366]** The method may comprise quantifying the complex formed by the antibody or antigen binding fragment bound to KAAG1 or to the KAAG1 variant.

**[0367]** The binding of an antibody to an antigen will cause an increase in the expected molecular weight of the antigen. A physical change therefore occurs upon specific binding of the antibody or antigen binding fragment and the antigen.

**[0368]** Such changes may be detected using, for example, electrophoresis followed by Western blot and coloration of the gel or blot, mass spectrometry, HPLC coupled with a computer or else. Apparatus capable of computing a shift in molecular weight are known in the art and include for example, Phosphorimager™.

**[0369]** When the antibody comprises for example a detectable label, the antigen-antibody complex may be detected by the fluorescence emitted by the label, radiation emission of the label, enzymatic activity of a label provided with its substrate or else.

**[0370]** Detection and/or measurement of binding between an antibody or antigen binding fragment and an antigen may be performed by various methods known in the art. Binding between an antibody or antigen binding fragment and an antigen may be monitored with an apparatus capable of detecting the signal emitted by the detectable label (radiation emission, fluorescence, color change etc.). Such apparatus provides data which indicates that binding has occurred and may also provide indication as to the amount of antibody bound to the antigen. The apparatus (usually coupled with a computer) may also be capable of calculating the difference between a background signal (e.g., signal obtained in the absence of antigen-antibody binding) or background noise and the signal obtained upon specific antibody-antigen binding. Such apparatuses may thus provide the user with indications and conclusions as to whether the antigen has been detected or not.

**[0371]** Also disclosed are kits which may include one or more container containing one or more antibodies or antigen binding fragments described herein.

#### **Nucleic acids, vectors and cells**

**[0372]** Antibodies are usually made in cells allowing expression of the light chain and heavy chain expressed from a vector(s) comprising a nucleic acid sequence encoding the light chain and/or heavy chain.

**[0373]** Disclosed herein are nucleic acids capable of encoding any of the CDRs, light chain variable regions, heavy chain variable regions, light chains, heavy chains described herein.

**[0374]** Disclosed herein is a nucleic acid encoding a light chain variable region and/or a heavy chain variable region of an antibody which is capable of specific binding to KAAG1.

**[0375]** Exemplary nucleic acids include a nucleic acid selected from the group consisting of a nucleic acid having at least 70% sequence identity (i.e., at least 75%, at least 80% sequence

identity) with any one of SEQ ID NOs.:3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 45 and 47, fragments (e.g., of at least 10, at least 15, at least 20 consecutive nucleotides) and complement thereof.

**[0376]** In accordance with the present disclosure, the nucleic acid may especially encode a light chain variable region and/or heavy chain variable region of an antibody which may be capable of inducing killing (elimination, destruction, lysis) of KAAG1- or KAAG1 variant-expressing tumor cells.

**[0377]** In accordance with the present disclosure, the nucleic acid may especially encode a light chain variable region and/or heavy chain variable region of an antibody which may be capable of reducing spreading of KAAG1- or KAAG1 variant-expressing tumor cells.

**[0378]** In accordance with the present disclosure, the nucleic acid may particularly encode a light chain variable region and/or heavy chain variable region of an antibody which may be capable of decreasing or impairing formation of KAAG1- or KAAG1 variant-expressing tumors.

**[0379]** Exemplary nucleic acids include nucleic acids encoding a light chain variable region comprising:

1. a. a CDRL1 sequence selected from the group consisting of SEQ ID NO.:72 and SEQ ID NO.:73;
2. b. a CDRL2 sequence selected from the group consisting of SEQ ID NO.:74, SEQ ID NO.: 75 and SEQ ID NO.:76, or;
3. c. a CDRL3 sequence selected from the group consisting of SEQ ID NO.:77, SEQ ID NO.:78 and SEQ ID NO.:79.

**[0380]** In accordance with the present disclosure, the nucleic acid may encode a light chain variable region which may comprise at least two CDRs of a CDRL1, a CDRL2 or a CDRL3.

**[0381]** Also in accordance with the present disclosure, the nucleic acid may encode a light chain variable region which may comprise one CDRL1, one CDRL2 and one CDRL3.

**[0382]** The present disclosure also relates to a nucleic acid encoding a heavy chain variable region comprising:

1. a. a CDRH1 sequence comprising SEQ ID NO.:80;
2. b. a CDRH2 sequence selected from the group consisting of SEQ ID NO.:81, SEQ ID NO.:82, SEQ ID NO.:83, SEQ ID NO.:84 and SEQ ID NO.:85, or;
3. c. a CDRH3 sequence selected from the group consisting of SEQ ID NO.:86, SEQ ID NO.:87 and SEQ ID NO.:88.

**[0383]** In accordance with the present disclosure, the nucleic acid may encode a heavy chain variable region which may comprise at least two CDRs of a CDRH1, a CDRH2 or a CDRH3.

**[0384]** In accordance with the present disclosure, the nucleic acid may encode a heavy chain variable region which may comprise one CDRH1, one CDRH2 and one CDRH3.

**[0385]** Also encompassed by the present disclosure are nucleic acids encoding antibody variants having at least one conservative amino acid substitution.

**[0386]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least one conservative amino acid substitution.

**[0387]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least one conservative amino acid substitution in at least two of the CDRs.

**[0388]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least one conservative amino acid substitution in the 3 CDRs.

**[0389]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least two conservative amino acid substitutions in at least one of the CDRs.

**[0390]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least two conservative amino acid substitutions in at least two of the CDRs.

**[0391]** In accordance with the present disclosure, the nucleic acid may encode a CDR comprising at least two conservative amino acid substitutions in the 3 CDRs.

**[0392]** Disclosed herein is a nucleic acid encoding a light chain variable region having at least 70%, 75%, 80% sequence identity with a sequence selected from the group consisting of SEQ ID NO.:16, SEQ ID NO.:20, SEQ ID NO.:24, SEQ ID NO.:103, SEQ ID NO.:104, SEQ ID NO.:105, SEQ ID NO.:106, SEQ ID NO.:107, SEQ ID NO.:108, SEQ ID NO.:109, SEQ ID NO.:110, SEQ ID NO.:111, SEQ ID NO.:112, SEQ ID NO.:113, SEQ ID NO.:114, SEQ ID NO.:115, SEQ ID NO.:116, SEQ ID NO.:117, SEQ ID NO.:118, SEQ ID NO.:119, SEQ ID NO.:120, SEQ ID NO.:121, SEQ ID NO.:122, SEQ ID NO.:123, SEQ ID NO.:124 and SEQ ID NO.:125.

**[0393]** Also disclosed herein is a nucleic acid encoding a heavy chain variable region having at least 70%, 75%, 80% sequence identity to a sequence selected from the group consisting of SEQ ID NO.:18, SEQ ID NO.:22, SEQ ID NO.:26, SEQ ID NO.:126, SEQ ID NO.:127, SEQ ID NO.:128, SEQ ID NO.:129, SEQ ID NO.:130, SEQ ID NO.:131, SEQ ID NO.:132, SEQ ID NO.:133, SEQ ID NO.:134, SEQ ID NO.:135, SEQ ID NO.:136, SEQ ID NO.:137, SEQ ID NO.:138, SEQ ID NO.:139, SEQ ID NO.:140, SEQ ID NO.:141, SEQ ID NO.:142, SEQ ID NO.:143, SEQ ID NO.:144, SEQ ID NO.:145, SEQ ID NO.:146 and SEQ ID NO.:147.

**[0394]** Also disclosed herein is a vector comprising the nucleic acids described herein.

**[0395]** In accordance with the present disclosure, the vector may be an expression vector.

**[0396]** Vector that contains the elements for transcriptional and translational control of the inserted coding sequence in a particular host are known in the art. These elements may include regulatory sequences, such as enhancers, constitutive and inducible promoters, and 5' and 3' un-translated regions. Methods that are well known to those skilled in the art may be used to construct such expression vectors. These methods include *in vitro* recombinant DNA techniques, synthetic techniques, and *in vivo* genetic recombination.

**[0397]** Disclosed herein is an isolated cell that may comprise the nucleic acid described herein.

**[0398]** The isolated cell may comprise a nucleic acid encoding a light chain variable region and a nucleic acid encoding a heavy chain variable region either on separate vectors or on the same vector. The isolated cell may also comprise a nucleic acid encoding a light chain and a nucleic acid encoding a heavy chain either on separate vectors or on the same vector.

**[0399]** In accordance with the present disclosure, the cell may be capable of expressing, assembling and/or secreting an antibody or antigen binding fragment thereof.

**[0400]** Also disclosed herein is a cell which may comprise and/or may express the antibody described herein.

**[0401]** In accordance with the present disclosure, the cell may comprise a nucleic acid encoding a light chain variable region and a nucleic acid encoding a heavy chain variable region.

**[0402]** The cell may be capable of expressing, assembling and/or secreting an antibody or antigen binding fragment thereof.

**[0403]** The examples below are presented to further outline details of the present invention.

## EXAMPLES

### *Example 1*

**[0404]** This example discloses the methods used to convert the Fabs into full IgG1 chimeric monoclonal antibodies.

**[0405]** Aside from the possibility of conducting interaction studies between the Fab

monoclonals and the KAAG1 protein, the use of Fabs may be limited with respect to conducting meaningful *in vitro* and *in vivo* studies to validate the biological function of the antigen. Thus, it was necessary to transfer the light and heavy chain variable regions contained in the Fabs to full antibody scaffolds, to generate mouse-human chimeric IgG1s. The expression vectors for both the light and heavy immunoglobulin chains were constructed such that i) the original bacterial signal peptide sequences upstream of the Fab expression vectors were replaced by mammalian signal peptides and ii) the light and heavy chain constant regions in the mouse antibodies were replaced with human constant regions. The methods to accomplish this transfer utilized standard molecular biology techniques that are familiar to those skilled in the art.

**[0406]** *Light chain expression vector-* an existing mammalian expression plasmid, called pTTVH8G (Durocher et al., 2002), designed to be used in the 293E transient transfection system was modified to accommodate the mouse light chain variable region. The resulting mouse-human chimeric light chain contained a mouse variable region followed by the human kappa constant domain. The cDNA sequence encoding the human kappa constant domain was amplified by PCR with primers OGS1773 and OGS1774 (SEQ ID NOS:55 and 56, respectively). The nucleotide sequence and the corresponding amino acid sequence for the human kappa constant region are shown in SEQ ID NOS: 57 and 58, respectively. The resulting 321 base pair PCR product was ligated into pTTVH8G immediately downstream of the signal peptide sequence of human VEGF A (NM\_003376). This cloning step also positioned unique restriction endonuclease sites that permitted the precise positioning of the cDNAs encoding the mouse light chain variable regions. The sequence of the final expression plasmid, called pTTVK1, is shown in SEQ ID NO.:59. Based on the sequences disclosed in Table 2, PCR primers specific for the light chain variable regions of antibodies 3D3, 3G10, 3C4 and 3A4 (SEQ ID NOS:15, 19, 23 and 47, respectively) were designed that incorporated, at their 5'-end, a sequence identical to the last 20 base pairs of the VEGF A signal peptide. The sequences of these primers are shown in SEQ ID NOS:60, 61, 62 and 213. The same reverse primer was used to amplify all three light chain variable regions of 3D3, 3G10 and 3C4 since the extreme 3'-ends were identical. This primer (SEQ ID NO.:63) incorporated, at its 3'-end, a sequence identical to the first 20 base pairs of the human kappa constant domain. Primer SE ID NO.:214 was used to amplify the 3A4 light chain variable region. Both the PCR fragments and the digested pTTVK1 were treated with the 3' - 5' exonuclease activity of T4 DNA polymerase resulting in complimentary ends that were joined by annealing. The annealing reactions were transformed into competent *E. coli* and the expression plasmids were verified by sequencing to ensure that the mouse light chain variable regions were properly inserted into the pTTVK1 expression vector. Those skilled in the art will readily recognize that the method used for construction of the light chain expression plasmids applies to all anti-KAAG1 antibodies contained in the original Fab library.

**[0407]** *Heavy chain expression vector -* the expression vector that produced the heavy chain immunoglobulins was designed in a similar manner to the pTTVK1 described above for production of the light chain immunoglobulins. Plasmid pYD11 (Durocher et al., 2002), which contains the human IgGK signal peptide sequence as well as the CH2 and CH3 regions of the

human Fc domain of IgG1, was modified by ligating the cDNA sequence encoding the human constant CH1 region. PCR primers OGS1769 and OGS1770 (SEQ ID NOS:64 and 65), designed to contain unique restriction endonuclease sites, were used to amplify the human IgG1 CH1 region containing the nucleotide sequence and corresponding amino acid sequence shown in SEQ ID NOS:66 and 67. Following ligation of the 309 base pair fragment of human CH1 immediately downstream of the IgGK signal peptide sequence, the modified plasmid (SEQ ID NO.:68) was designated pYD15. When a selected heavy chain variable region is ligated into this vector, the resulting plasmid encodes a full IgG1 heavy chain immunoglobulin with human constant regions. Based on the sequences disclosed in Table 2, PCR primers specific for the heavy chain variable regions of antibodies 3D3, 3G10, 3C4 and 3A4 (SEQ ID NOS:17, 21, 25 and 45, respectively) were designed that incorporated, at their 5'-end, a sequence identical to the last 20 base pairs of the IgGK signal peptide. The sequences of these primers are shown in SEQ ID NOS:69 (3D3 and 3G10 have the same 5'-end sequence), SEQ ID NO.: 70 or SEQ ID NO.:215 for 3A4. The same reverse primer was used to amplify all three heavy chain variable regions of 3D3, 3C4 and 3G10 since the extreme 3'-ends were identical. This primer (SEQ ID NO.:71) incorporated, at its 3'-end, a sequence identical to the first 20 base pairs of the human CH1 constant domain. For the 3A4 heavy chain variable region, SEQ ID NO.:216 was used. Both the PCR fragments and the digested pYD15 were treated with the 3' - 5' exonuclease activity of T4 DNA polymerase resulting in complimentary ends that were joined by annealing. The annealing reactions were transformed into competent *E. coli* and the expression plasmids were verified by sequencing to ensure that the mouse heavy chain variable regions were properly inserted into the pYD15 expression vector. Those skilled in the art will readily recognize that the method used for construction of the heavy chain expression plasmids applies to all anti-KAAG1 antibodies contained in the original Fab library.

**[0408]** *Expression of human IgG1s in 293E cells* - The expression vectors prepared above that encoded the light and heavy chain immunoglobulins were expressed in 293E cells using the transient transfection system (Durocher et al., 2002). Other methods of transient or stable expression may be used. The ratio of light to heavy chain was optimized in order to achieve the most yield of antibody in the tissue culture medium and it was found to be 9:1 (L:H). The ability of the anti-KAAG1 antibodies (monoclonal, chimeric or humanized) to bind to recombinant Fc-KAAG1 was measured by ELISA and compared with the original mouse Fabs.

**[0409]** The scheme used to convert other Fabs into a complete IgG (including the 3A4) and for expression of the antibodies is described in more details in international application No. PCT/CA2012/000296, the entire content of which is incorporated herein by reference.

## Example 2

### Humanization of the 3A4 mouse monoclonal antibody

**[0410]** International patents No. PCT/CA2009/001586, PCT/CA2010/001795 and No. PCT/CA2012/000296, described exemplary methodology used to generate the humanized light chain and heavy chain variable regions.

**[0411]** Humanization of the 3A4 antibody light chain variable region involved 11 mutations to its proposed humanized framework for 100% framework humanization. Humanization of the 3A4 antibody heavy chain variable region involved 23 mutations to its proposed humanized framework for 100% framework humanization. These 100% humanized variable region sequences are labelled Lvh1 and Hvh1, respectively (SEQ ID NOs:189 and 194). Additional humanized sequences were also designed in which several residues from the 3A4 mouse sequences were retained based on careful structural and comparative sequence analyses that indicate a high probability of altering antigen-binding affinity if mutations are to be introduced at these positions. These sequences of the variable regions are labelled Lvh2, Hvh2, Hvh3 and Hvh4 (SEQ ID NOs: 190, 195, 196 and 197).

**[0412]** The two humanized light chain variants (including the constant region) are identified herein as Lh1 (SEQ ID NO.: 199) and Lh2 (SEQ ID NO.:200). The four humanized heavy chain variants (including the constant region are identified herein as Hh1 (SEQ ID NO.:202), Hh2 (SEQ ID NO.:203), Hh3 (SEQ ID NO.:204) and Hh4 (SEQ ID NO.:205). The two humanized light chain and 4 humanized heavy chain can be assembled into 8 humanized antibodies (Lh1Hh1, Lh1Hh2, Lh1Hh3, Lh1Hh4, Lh2Hh1, Lh2Hh2, Lh2Hh3, and Lh2Hh4).

**[0413]** In the case of 3A4 light-chain humanized sequence Lvh2 (SEQ ID NO:190), framework residues Val-L2 and Lys-L45 were retained from the mouse sequence since residue L2 is semi-buried, contacts both CDR-L1 and CDR-L3, and has antigen-contacting propensity, while residue L45 approaches the heavy-chain. We note that both these murine residues may occur in human frameworks. In the case of 3A4 heavy-chain humanized sequence Hvh2 (SEQ ID NO:195), framework residues Ile-H2 and Lys-L73 were retained from the mouse sequence since residue H2 is semi-buried, contacts both CDR-H1 and CDR-H3, and has antigen-contacting propensity, while residue H73 belongs to the Vernier zone supporting CDR-H2, and both these murine residues may occur in human frameworks. In the case of 3A4 heavy-chain humanized sequence Hvh3 (SEQ ID NO:196), Ile-H2 and Lys-L73 back-mutations were retained and in addition to these, framework residues Ile-H48, Ala-H67, Leu-H69 and Val-H71 were retained from the mouse sequence since all these additional murine residues are buried residues and belong to the Vernier zone supporting CDR-H2, and also murine residue H71 may occur in human frameworks. In the case of 3A4 heavy-chain humanized sequence Hvh4 (SEQ ID NO:197), all 6 back-mutations of the Hvh3 humanized variant were included plus additional two mouse framework residues Lys-H38 and Lys-H66 since they represent semi-buried residues close to CDR-H2. The resulting amino acid sequences of the murine and humanized chains are listed in Table 1. The alignment of the murine and humanized light chain variable regions is shown in Figure 1a and the alignment of the murine and humanized heavy chain variable regions is shown in Figure 1b.

**[0414]** Figure 2a and 2b is an alignment of the murine light chain variable region with the

100% humanized light chain variable region and the murine heavy chain variable region with the 100% humanized heavy chain variable region respectively. This figure illustrates the amino acids that are preserved and those that have been chosen for substitution.

**Example 3.**

**Assembly and expression of 3A4 humanized variant antibodies**

**[0415]** The purpose of these investigations is to determine the kinetics parameters of anti-clusterin antibodies. In particular, to determine whether the humanization of the 3A4 anti-KAAG1 monoclonal antibody affects the kinetics parameters of its binding to human KAAG1. To this end, a kinetic analysis method was developed using the ProteOn XPR36 instrument from BioRad. Human KAAG1 was immobilized on a sensor chip. Full length antibodies or Fab fragments were injected and allowed to interact with the immobilized KAAG1.

***Construction of plasmid encoding the chimeric (murine) heavy and light chains of 3A4***

**[0416]** The heavy and light chains of the chimeric antibody were amplified by PCR from the original murine immunoglobulin chains using the following oligonucleotide primer pairs: heavy chain, 5'-oligo encoded by SEQ ID NO: 206 and 3'-oligo encoded by SEQ ID NO:207; light chain, 5'-oligo encoded by SEQ ID NO: 208 and 3'-oligo encoded by SEQ ID NO:209. The resulting PCR products were digested by Hind III and cloned into pK-CR5 (SEQ ID NO:210) previously digested with Hind III.

***Construction of plasmids encoding the humanized heavy chain 3A4 variants 1, 2, 3 and 4***

**[0417]** The fragments coding for the humanized heavy chain region of the antibody 3A4 (Hh1, Hh2, Hh3 and Hh4) were ordered from GenScript (Piscataway, USA). The DNA fragments including the kozak and stop codon sequences were digested with HindIII and cloned into the HindIII site of plasmid pK-CR5 previously dephosphorylated with calf intestinal phosphatase (NEB) to prevent recircularization. Figure 3a shows the map of the plasmid pK-CR5-3A4-HC-variant1. All heavy chain variants of the humanized 3A4 were constructed in a similar manner.

***Construction of plasmids encoding the humanized light chain 3A4 variants 1 and 2***

**[0418]** The fragments coding for the human light chain regions of the antibody 3A4 (Lh1 and

Lh2) were ordered from GenScript. The DNA fragments including the kozak and stop codon sequences was digested with BamHI and cloned into the BamHI site of plasmid pMPG-CR5 (SEQ ID NO:211) previously dephosphorylated with calf intestinal phosphatase (NEB) to prevent recircularization. Figure 3b shows the map of the plasmid pMPG-CR5-3A4-LC-variant1. All light chain variants of the humanized 3A4 were constructed in a similar manner.

#### ***Transient transfection study***

**[0419]** Plasmid DNA was isolated from small cultures of *E. coli* using the Mini-Prep kit (Qiagen Inc, Mississauga, ON) according to the manufacturer's recommendation. Briefly, 2 ml of LB medium containing 100 µg/ml of ampicillin were inoculated with a single colony picked after ligation and transformation. The cultures were incubated at 37°C overnight with vigorous shaking (250 RPM). The plasmid was then isolated from 1.5 ml of culture using the protocols, buffers, and columns provided by the kit. The DNA was eluted using 50 µl of sterile water. Plasmid DNA was isolated from large culture of *E. coli* using the Plasmid Plus Maxi kit (Qiagen Inc, Mississauga, ON) according to the manufacturer's recommendation. 200 mL of LB medium containing 100 µg/mL ampicillin were inoculated with a single fresh colony of *E. coli* and incubated overnight at 37°C with vigorous shaking (250 RPM). The bacteria (130 mL of culture for the heavy chain and 180 mL of culture for the light chain) were pelleted by centrifugation at 6000 x g, for 15 min, at 4°C and the plasmid was isolated using the protocols, buffers and columns provided by the kit. The pure plasmids was resuspended in sterile 50 mM Tris, pH8 and quantified by measuring the optical density at 260 nm. Before transfection the purified plasmid were sterilized by extraction with phenol/chloroform followed by ethanol precipitation. The plasmid were resuspended in sterile 50 mM Tris, pH 8 and quantified by optical density at 260 nm.

**[0420]** Before transfection, the cells (CHO-cTA) were washed with PBS and resuspended at a concentration of  $4.0 \times 10^6$  cell/ml in growth medium (CD-CHO, Invitrogen) without dextran sulfate for 3 h in suspension culture. For each plasmid combination, 45 ml of cells were transfected by adding slowly 5 ml of CDCHO medium supplemented with 10 µg/ml of each plasmid and 50 µg/ml of polyethylenimine (PEI Max; Polysciences). The final concentration was 1 µg/ml of each plasmid and 5 µg/ml of PEI. After 2 h, the cells were transferred at 30°C. The next days, 50 µg/mL of dextran sulfate and 3.75 ml of each supplement (Efficient Feed A and B Invitrogen) were added to the cells and they were incubated at 30°C for 13 days. 2.5 ml of Feed A and 2.5 ml of Feed B were added at day 4, 6, 8 and 11. On day 13, the supernatant was clarified by centrifugation and filtered through a 0.22 µM filter.

**[0421]** CHO cells (CHOcTA) were transfected with plasmids encoding the different variants of humanized heavy and light chains of the 3A4 antibody regulated by the CR5 promoter. Transfection with different combinations of light and heavy chains was performed. As control, cells were also transfected with plasmids encoding the chimeric/murine antibody.

#### ***Purification of antibody***

**[0422]** 15 ml of supernatant from the CHO cell transfections were concentrated by centrifugation using the Amicon Ultra (Ultacell-50k) cassette at 1500 rpm. The concentrated antibody (550  $\mu$ l) was purified using the Nab spin kit Protein A Plus (Thermo Scientific) according to the manufacturer's recommendations. The purified antibodies were then desalted using PBS and the concentrating Amicon Ultra (Ultracel-10K) cassette at 2500 rpm to a final volume of 250  $\mu$ l. The purified antibody was quantified by reading the OD<sub>280</sub> using the Nanodrop spectrophotometer and kept frozen at -20°C. An aliquote of the purified antibody was resuspended into an equal volume of Laemmli 2X and heated at 95°C for 5 min and chilled on ice. A standard curve was made using known amount of purified human IgG1 kappa from Human Myeloma plasma (Athens Research). The samples were separated on a polyacrylamide Novex 10% Tris-Glycine gel (Invitrogen Canada Inc., Burlington, ON) and transferred onto a Hybond-N nitrocellulose membrane (Amersham Bioscience Corp., Baie d'Urfée, QC) for 1 h at 275 mA. The membrane was blocked for 1 h in 0.15% Tween 20, 5% skimmed milk in PBS and incubated for 1 hr with an Goat anti-Human IgG (H+L) conjugated to Cy5 (Jackson, Cat# 109-176-099). The signal was revealed and quantified by scanning with the Typhoon Trio+ scanner (GE Healthcare). As shown in Figure 4, all combinations of the 3A4 humanized antibody variants were expressed in CHO cells.

#### **Example 4.**

#### **Kinetic analysis of murine and humanized 3A4 antibody**

#### ***Supplies***

**[0423]** GLM sensorchips, the Biorad ProteOn amine coupling kit (EDC, sNHS and ethanolamine), and 10mM sodium acetate buffers were purchased from Bio-Rad Laboratories (Mississauga, ON). HEPES buffer, EDTA, and NaCl were purchased from Sigma-Aldrich (Oakville, ON). Ten percent Tween 20 solution was purchased from Teknova (Hollister, CA). The goat anti-human IgG Fc fragment specific antibody was purchased from Jackson ImmunoResearch. The gel filtration column Superdex 75 10/300 GL was purchased from GE Healthcare.

#### ***Gel filtration***

**[0424]** The KAAG1 protein at a concentration of 3.114 mg/ml and a volume of 220  $\mu$ L was injected onto the Superdex G75 column. The separation was done at 0.4ml/min in HBST

running buffer (see below) without Tween 20. The volume of the fractions collected was 500  $\mu$ L. Concentration of KAAG1 in each fraction was determined by OD<sub>280</sub> using an extension coefficient of 5500 and a MW of 8969. Figure 5 represents the profile of the gel filtration of KAAG1. A small peak of potential aggregate is eluting at around 11 ml. The protein eluting at 13 ml was used as analyte for the SPR assay (fractions 15 - 19).

### **SPR biosensor assays**

**[0425]** All surface plasmon resonance assays were carried out using a BioRad ProteOn XPR36 instrument (Bio-Rad Laboratories Ltd. (Mississauga, ON) with HBST running buffer (10mM HEPES, 150 mM NaCl, 3.4 mM EDTA, and 0.05% Tween 20 pH 7.4) at a temperature of 25°C. The anti-mouse Fc capture surface was generated using a GLM sensorchip activated by a 1:5 dilution of the standard BioRad sNHS/EDC solutions injected for 300 s at 30  $\mu$ L/min in the analyte (horizontal) direction. Immediately after the activation, a 13  $\mu$ g/mL solution of anti-human IgG Fc fragment specific in 10 mM NaOAc pH 4.5 was injected in the analyte direction at a flow rate of 25  $\mu$ L/min until approximately 8000 resonance units (RUs) were immobilized. Remaining active groups were quenched by a 300 s injection of 1M ethanolamine at 30  $\mu$ L/min in the analyte direction, and this also ensures mock-activated interspots are created for blank referencing. The screening of the 3A4 variants for binding to KAAG1 occurred in two steps: an indirect capture of 3A4 variants from cell supernatant onto the anti-human IgG Fc fragment specific surface in the ligand direction(vertical) followed by a KAAG1 injection in the analyte direction. Firstly, one buffer injection for 30 s at 100  $\mu$ L/min in the ligand direction was used to stabilize the baseline. For each 3A4 capture, unpurified 3A4 variants in cell-culture media were diluted to 4 % in HBST, or approximately 1.25  $\mu$ g/mL of purified 3A4 in HBST was used. Four to five 3A4 variants along with wild-type 3A4 were simultaneously injected in individual ligand channels for 240 s at flow 25  $\mu$ L/min. This resulted in a saturating 3A4 capture of approximately 400-700 RUs onto the anti-human IgG Fc fragment specific surface. The first ligand channel was left empty to use as a blank control if required. This 3A4 capture step was immediately followed by two buffer injections in the analyte direction to stabilize the baseline, and then the gel filtration purified KAAG1 was injected. For a typical screen, five KAAG1 concentrations (8, 2.66, 0.89, 0.29, and 0.098 nM) and buffer control were simultaneously injected in individual analyte channels at 50  $\mu$ L/min for 120 s with a 600s dissociation phase, resulting in a set of binding sensorgrams with a buffer reference for each of the captured 3A4 variants. The anti-human IgG Fc fragment specific - 3A4 complexes were regenerated by a 18 s pulse of 0.85% phosphoric acid for 18 s at 100  $\mu$ L/min to prepare the anti-human IgG Fc fragment specific surface for the next injection cycle. Sensorgrams were aligned and double-referenced using the buffer blank injection and interspots, and the resulting sensorgrams were analyzed using ProteOn Manager software v3.0. The kinetic and affinity values were determined by fitting the referenced sensorgrams to the 1:1 Langmuir binding model using local  $R_{max}$ , and affinity constants ( $K_D$  M) were derived from the resulting rate constants ( $k_d$  s<sup>-1</sup>/ $k_a$  M<sup>-1</sup>s<sup>-1</sup>).

### **Determination of rate and affinity constants**

**[0426]** Figure 6 summarizes the association ( $k_a$ , 1/Ms) and dissociation ( $k_d$ , 1/s) rate constants as well as affinity ( $K_D$ , M) constants for the interaction of KAAG1 with purified murine 3A4, murine 3A4 transiently expressed as a chimeric and transiently expressed humanized variants. These constants are graphically represented in Figure 7a-c. The association rate constant is very similar for the pure parental, chimeric and humanized 3A4 variants (Figure 7a). The dissociation rate constants is similar for the transiently express chimeric as compared to the pure parental 3A4 with suggest that the transfection procedure did not alter the parameters of the interaction of KAAG1 with the antibody (Figure 7b). However, all humanized variants seem to have a slightly altered off rate, i.e. quicker dissociation rate (Figure 7b). This is reflected in the affinity constants (Figure 7c). In summary, there is a linear correlation between the binding affinity ( $\log K_D$ ) of the humanized variant and the number of back-mutations made in the parent antibody (LcHc) with a decrease in the binding affinity as the number of mutations is increasing. However, the difference in binding affinity is only 4 fold different between the worse variant (H1L1, 0.47 nM) which has no mouse residue retained and the best variant which has 10 mouse residues retained (H4L2, 0.1 nM). Finally, the binding affinity of all variants for KAAG1 was found to be sub-nanomolar and the best variant (H4L2, 0.1 nM) exhibited an affinity about 6-fold weaker than the murine (LcHc, 0.057 nM). Overall, these results indicate that humanization was successful as all of the variants displayed high affinity for KAAG1.

### **Example 5.**

#### **Binding of 3A4 humanized variants to KAAG1 in an ELISA**

**[0427]** ELISA methods were also used to compare the binding activity of the humanized 3A4 variants to the murine 3A4 antibody. Recombinant human KAAG1 was coated in 96-well plates O/N, washed and incubated for 1h at RT with increasing quantities of murine or humanized 3A4 variants. Following another round of washing steps, an anti-human antibody conjugated to HRP was added to the wells and the bound 3A4 antibody was measured calorimetrically at  $\text{Abs}_{450}$ . As shown in Figure 8a, the humanized variants (Lh1Hh1, Lh1Hh2, Lh1Hh3 and Lh1Hh4) displayed very similar binding to KAAG1 when compared to the murine 3A4 (LcHc), which has a high affinity of 0.016nM. This result indicated that all four humanized heavy chain variants were comparable to the original h3A4 heavy chain when assembled with the L1 variant of the humanized light chain. Figure 8a shows the results when the heavy chain variants were assembled with Lh2 variant of the 3A4 humanized light chain. In this instance, there was a difference in the binding of the variants. For example, Lh2hh4 was the variant with the closest profile compared to the murine 3A4. This was in agreement with the SPR data, which showed that the variant 4 of the heavy chain had the highest affinity for KAAG1. Taken

together, these binding results show that the humanized variants all interact with human KAAG1 in this assay. Although there were some subtle differences, the binding in ELISA was in concordance with the SPR results.

**Example 6.**

**Binding of 3A4 humanized variants on the surface of cancer cells**

**[0428]** Flow cytometry was used to evaluate the capacity of the humanized 3A4 variants to interact with KAAG1 expressed on the surface of cancer cells. To this end, SKOV-3 ovarian cancer cells, which we had previously showed were efficiently bound by 3A4 by flow cytometry, were incubated with the eight humanized variants and the original murine antibody. Briefly, SKOV-3 cells were detached from the plate with EDTA and incubated on ice with either 3.0 mg/ml, 0.3 mg/ml or 0.3 mg/ml of the antibodies for 1h. After three washing steps, the cells were incubated with the secondary antibody, anti-human IgG-conjugated to FITC for 1h on ice. Cell surface fluorescence was measured in a flow cytometer and the values are shown in the histogram of Figure 9. As depicted, all variants could detect KAAG1 on the surface on unpermeabilized and the strongest signals were obtained at the highest concentration of 3A4 antibodies (3 mg/ml) and decreased as the concentration of the antibody was decreased. Among the different variants, the ones with the most murine back-mutations (Figure 9, see Lh1Hh4 and Lh2Hh4) interacted with KAAG1 on the surface of cells with the highest activity. In fact, Lh1Hh4 and Lh2Hh4 appeared to be slightly improved cell surface binding to KAAG1 compared to the murine 3A4 antibody (LcHc).

**Example 7**

**[0429]** This example describes the use of anti-KAAG1 antibodies for detecting the expression of KAAG1 in TNBC.

**[0430]** As a means of determining if the KAAG1 antigen was present in TNBC samples, immunohistochemistry was conducted. Tissue microarrays were obtained that contained 139 breast tumor samples generated from patient biopsies. Paraffin-embedded epithelial breast tumor samples were placed on glass slides and fixed for 15 min at 50°C. Deparaffinization was conducted by treating 2x with xylene followed by dehydration in successive 5 min washes in 100%, 80%, and 70% ethanol. The slides were washed 3x in PBS for 5 min and treated with antigen retrieval solution (1 mM EDTA, pH 8.0) to unmask the antigen. Endogenous peroxide reactive species were removed by incubating slides with H<sub>2</sub>O<sub>2</sub> in methanol and blocking was performed by incubating the slides with serum-free blocking solution (Santa Cruz Biotech) for 5 min at room temperature. The primary antibody (anti-KAAG1 3A4) was added for 1 h at room temperature. KAAG1-reactive antigen was detected by incubating with biotin-conjugated

mouse anti-kappa followed by streptavidin-HRP tertiary antibody. Positive staining was revealed by treating the slides with DAB-hydrogen peroxide substrate for less than 5 min and subsequently counterstained with hematoxylin. The KAAG1 protein was found to be expressed at very high levels in the vast majority of breast tumor samples. A representative array containing 139 tumors is depicted in Figure 10. In particular, 15/20 biopsy samples confirmed to be TNBC (Figure 10, samples identified by an asterisk) were stained strongly for KAAG1 expression with the 3A4 antibody. Taken together, these immunohistochemical studies illustrate the utility of detecting KAAG1 in breast cancer, in particular TNBC, with the monoclonal antibodies.

#### **Example 8**

**[0431]** This example describes the use of anti-KAAG1 antibodies for detecting the expression of KAAG1 in TNBC cell lines.

**[0432]** Combined results from the bioinformatics analysis of the primary structure of the cDNA encoding KAAG1, biochemical studies, and immunohistochemical detection of the protein in epithelial cells suggested that the KAAG1 antigen was located at the cell surface. However, more direct evidence was required to demonstrate that KAAG1 is indeed expressed on the surface of TNBC cells. To conduct this analysis, breast cancer cell lines were obtained from a commercial vendor (ATCC, Manassas, VA) and used in flow cytometry experiments. RT-PCR expression analyses using KAAG1 mRNA specific primers previously showed that certain breast cancer cell lines expressed KAAG1 mRNA (see PCT/CA2007/001134). Therefore some of these cell lines were selected to determine the presence of the KAAG1 antigen at their surface. To verify this, the triple-negative MDA-MB-231, MDA-MB-436, MDA-MB-468, BT-20 and BT-549 cell lines were tested for surface expression of KAAG1 using the 3A4 anti-KAAG1 antibody. In addition, breast cancer cell lines, which are not triple-negative, namely T47D and MCF-7, were also included in the analysis. Finally, a control cell line, 293-6E, that exhibits undetectable level of KAAG1 antigen expression was included as a negative control for the flow cytometry experiment (FCM). For the purpose of FCM analysis, the cells were harvested using 5 mM EDTA, counted with a hemocytometer, and resuspended in FCM buffer (0.5% BSA, 0.01% goat serum in 1x PBS) at a cell density of  $2 \times 10^6$  cells/ml. Chimeric 3A4 anti-KAAG1 antibody or a control IgG were added to 100  $\mu$ l of cells at a final concentration of 0.5  $\mu$ g/ml and incubated on ice for 1h. The cells were washed in cold FCM buffer to remove unbound antibodies, resuspended in 100  $\mu$ l FCM buffer containing anti-human IgG conjugated to FITC secondary antibody (diluted 1:200) and incubated on ice for 45min. Following another washing step in cold FCM buffer, the cells were resuspended in 300  $\mu$ l FCM buffer and analyzed with a flow cytometer. 10  $\mu$ g/ml propidium iodide was added to each sample to allow for gating of dead cells. The results from three independent experiments are shown in Figure 11, where the mean fluorescence intensity (MFI) fold Induction represents the geometric mean value of the signal obtained when the cells were incubated with 3A4 antibody over that of the negative human IgG control, which was arbitrarily set to 1. Incubation of the antibodies with the control

293-6EHEK-293 cells resulted in fluorescence signals that were similar to the signal obtained when the cells were incubated in the absence of the primary antibody. Furthermore, there was no significant difference between the signal obtained with 3A4 compared to the control IgG. Moreover, when the control IgG was incubated with the breast cancer cell lines, the signals were very similar to those obtained with the control 293-6E cells. By contrast, detectable fluorescence signal was observed when the 3A4 antibody was incubated with all breast cancer cells lines. Although variable amount of fluorescence was observed, the highest amount of KAAG1 was detected on the surface of MDA-MB-231 and BT-20 cell lines, two TNBC cell lines (see Figure 11, TNBC cell lines are indicated with an asterisk). In fact all five TNBC cell lines were positive for KAAG1 expression under these conditions. T47 D and MCF-7 cells also expressed KAAG1. Taken together, this flow cytometry analysis shows that TNBC cell line express high level of KAAG1 on their cell surface.

#### **Example 9**

#### **Methods for use of the 3A4 anti-KAAG1 antibody as an antibody conjugate**

**[0433]** As demonstrated above, the KAAG1 antigen was detected by 3A4 on the surface of cancer cells using flow cytometry. There are several different molecular events that can occur upon binding of an antibody to its target on the surface of cells. These include i) blocking accessibility to another cell-surface antigen/receptor or a ligand, ii) formation of a relatively stable antibody-antigen complex to allow cells to be targeted via ADCC or CDC, iii) signalling events can occur as exemplified by agonistic antibodies, iv) the complex can be internalized, or v) the complex can be shed from the cell surface. To address this question we examined the behavior of the 3A4 antibody-KAAG1 complex on the surface of the cells. The ovarian cancer cell line, SKOV3, was used as a positive control in this experiment since it was successfully used in previous internalization experiments (see PCT/CA2009/001586). MDA-MB-231 TNBC cells were plated, washed, and incubated with 0.5 µg/ml chimeric 3A4 antibody as described in Example 3. After washing, complete medium was added and the cells placed at 37°C for up to 60 minutes. The cells were removed at the indicated times (see Figure 12), rapidly cooled, prepared for flow cytometry with FITC-conjugated anti-human IgG and the results were expressed as the percentage of mean fluorescence intensity remaining on the cell surface compared with the signal at time 0 minutes (see Figure 12, Surface signal (% remaining at 0 min)). As illustrated in Figure 12, the fluorescence signal decreased rapidly when 3A4 was incubated with MDA-MB-231 cells (Figure 12, black bars, indicated by MDA-231 in the figure) and seemed to achieve a maximum loss of signal by 30 - 45 minutes. The loss of signal was comparable to that observed when 3A4 was incubated with the SKOV3 cells (Figure 12, grey bars). This result indicates that the 3A4/KAAG1 complex disappeared from the cells which indicated that an internalization of the complex likely occurred. Preliminary studies to elucidate the mechanism responsible for this decrease in cell-surface fluorescence have revealed that the complex appears to be internalized. Similar results are expected with humanized 3A4

antibodies.

**[0434]** Similar results were observed in two additional TNBC cell lines, namely MDA-MB-436 (Figure 13) and BT-20 (Figure 14) confirming that the internalization of the 3A4/KAAG1 complex on the surface of multiple TNBC cell lines. By contrast, despite similar MFI levels of 3A4 binding on the surface of MDA-MB-436 and T47D (Figure 11), the loss of signal at the cell surface was not observed when 3A4 was incubated with the T47D cell line. This finding suggests the possibility that internalization of the 3A4/KAAG1 complex might occur to a higher degree in TNBC cells (Figure 15) compared with cells that are not triple-negative.

**[0435]** These findings were further confirmed by conducting immunofluorescence on live cells to see if this internalization could be microscopically observed. MDA-MB-231 cells were seeded on cover slips and once the cells were properly adhered, fresh medium was added containing the 3A4 anti-KAAG1 chimeric antibody at 10 ug/ml and incubating at 37 C for 4h. The cells were washed in PBS then fixed in 4% paraformaldehyde (in PBS) for 20 min. After washing, the cells were permeabilized with 0.1% Triton X-100 in PBS for 5 min. Blocking was performed with 1.5% dry milk in PBS for 1h. Lysosomal-associated membrane protein 1 (LAMP1, Chang et al., 2002) was detected by incubating with anti-LAMP1 (Santa Cruz, sc-18821, diluted 1:100) in 1.5 % milk in PBS for 2h. After washing in PBS, the secondary antibodies were added together in 1.5% milk and incubated for 1h. For the anti-KAAG1 chimeric antibody the secondary antibody was a Rhodamine Red-X conjugated donkey anti-human IgG (H+L) diluted 1:300. For the anti-LAMP1 antibody the secondary antibody was a DyLight488-conjugated goat anti-mouse IgG (H+L) diluted 1:300. Both secondary antibodies were from Jackson ImmunoResearch. The coverslips were washed in PBS and mounted in ProLong Gold antifade reagent with DAPI. As seen in Figure 7, after 4 hours of incubation at 37 C in the presence of MDA-MB-231 cancer cells, the 3A4 antibody was able to be detected in complexes predominantly near the perinuclear area (arrows, see red staining in the left panel in Figure 16), which is typical of endosomal-lysosomal-based internalization pathways. This observation was further confirmed when a lysosomal marker, LAMP1 was visualized and was found to be also expressed in these areas (arrows, see green staining in the middle panel in Figure 16). Importantly, the merging of the two images resulted in the appearance of yellow-orange structures indicating that the 3A4 and the anti-LAMP1 antibodies were present in the same structures (arrows, see yellow staining in the right panel in Figure 16). The co-localization of 3A4, which binds to KAAG1 on the surface of cancer cells, with LAMP1, a marker of late endosomes/lysosomes, shows that the antibody/antigen complex was internalized and that it follows a pathway that is amenable for the release of a payload that would be conjugated to the 3A4 antibody. Identical results were observed in another TNBC cell line, BT-20 (see Figure 17).

**[0436]** Taken together, these studies demonstrated that antibodies specific for KAAG1 such as 3A4 might have uses as an antibody conjugate, in particular, as an antibody-drug conjugate (ADC). Thus, the high level of TNBC specificity of KAAG1 coupled with the capacity of this target to be internalized in cells support the development of applications as an ADC.

#### ***Example 10***

**[0437]** In order to demonstrate that anti-KAAG1 antibodies can efficiently target and kill cells lacking ER protein expression, PgR protein expression and/or showing absence of HER2 protein over-expression, we generated two antibody drug conjugates (ADCs); 3A4-ADC1 and 3A4-ADC2.

**[0438]** To that effect, we used the chimeric 3A4 antibody and conjugated a cytotoxic drug via a highly stable peptide linker that is selectively cleaved by lysosomal enzymes after internalization (3A4-ADC1), or conjugated with another anti-mitotic drug via a non-cleavable linker (3A4-ADC2). The cytotoxic drug may become active once internalized in the cells.

**[0439]** The ability of the 3A4 ADCs to detect KAAG1 on the surface of TNBC cells was determined using flow cytometry using the methods described herein. Briefly, unconjugated 3A4, 3A4-ADC1, 3A4-ADC2 and a control IgG were incubated in the presence of MDA-231 TNBC cells, which are KAAG1 positive. Results indicated that the conjugation of 3A4 with either drug did not affect its binding to triple negative breast cancer cells such as MDA-231 (data not shown).

**[0440]** Having confirmed that the 3A4 ADCs could bind to KAAG1 expressed on the surface of TNBC cells, their cytotoxicity against these cells was evaluated in cell proliferation assays. MDA-231 or TOV-112D cells were cultured as described above in previous examples. The cells were seeded at 3000 cells/well in 96-well plates in 200  $\mu$ l of media per well overnight at 37°C, in 5% CO<sub>2</sub>. The next day, media was replaced with fresh media containing antibodies, at concentrations ranging from 0.122 nM to 500 nM, and incubated at 37° C for 72h. All conditions were performed in triplicate wells. The number of surviving cells was determined by performing a cellular proliferation assay, using CellTiter 96 Aqueous One Solution (Promega, Madison, WI), following manufacturer's protocol. Following the collection of the raw data, the results were expressed as the percentage survival compared to the number of cells in the wells treated with PBS, which was set to 100%. Results indicated that the unconjugated 3A4 did not affect the proliferation of MDA-231 cells at all concentrations tested. In contrast, the 3A4 ADCs tested showed significant cytotoxicity.

**[0441]** These results indicate that 3A4 antibody conjugates may be used as an alternative treatment for patients having triple negative breast cancer or basal-like breast cancer. Similar results are expected for conjugates based on humanized 3A4 antibodies.

**[0442]** Sequences referred to in the description

SEQ ID NO.:1

GAGGGGCATCAATCACACCGAGAAGTCACAGCCCCTAACCACTGAGGTGTGGGGGGTAGGGAT  
CTGCATTCTTCATATCAACCCCACACTATAGGGCACCTAAATGGGTGGCGGTGGGGAGACCG  
ACTCACTTGAGTTCTTGAGGCTTCCTGGCCTCCAGCCACGTAATTGCCCGCTCTGGATCTG  
GTCTAGCTTCCGGATTCGGTGGCAGTCCGCGGGGTAGATGTTCTGACGGCCCCAAAGGGTG  
CCTGAACGCCGCCGGTCACCTCCTCAGGAAGACTTCGAAGCTGGACACCTTCTTCATGGATG  
ACGACGCCGCCGGCTAGAAGGGTCCCCGTTGCGGTACACAAGCACGCTCTCACGACGGG  
CTGAGACAGGTGGCTGGACCTGGCGCTGCGCTCATCTTCCCGCTGGCCGCCCTCAGCT  
CGCTGCTTCGCGTCGGGAGGCACCTCCGCTGTCCCAGCGGCCCTACCGCACCCAGGGCGCGGGAT

CGCCTCTGAAACGAACGAGAAACTGACGAATCCACAGGTGAAAGAGAACTAACGGCCGTGC  
 TAGGCCTCCACCCAGAGGGAGACACTAGGAGCTTGCAAGGACTCGGAGTAGACGCTCAAGTTTCA  
 CGTGGCGTGCACAGCAATCAGGACCCGCAGTGCACGCACCACACCAGGTTACCTGCTACGGG  
 CAGAATCAAGGTGGACAGCTCTGAGCAGGAGCCGAAACGCGGGGGCTTCAAACAGGCACGC  
 CTAGTGAGGGCAGGAGAGAGGAGACGCACACACACACACACAAATATGGTGAACACCAAT  
 TTCTTACATCATATCTGTGCTACCCCTTCCAAACAGCCTA

SEQ ID NO.:2

MDDDAAPRVEGPVAVHKHALHDGLRQVAGPGAAAHLPRWPPPQLAASRREAPPLSQRPHRTQG  
 AGSPPETNEKLTNPQVKEK

SEQ ID NO.:3

GACATTGTGATGACCCAGTCTCCATCCTCCCTGGCTGTGTCATAAGGACAGAAGGTCACTATGAA  
 CTGCAAGTCCAGTCAGAGCCTTAAATAGTAACCTTCAAAAGAACCTTTGCCCTGGTACCGC  
 AGAAACCAGGCCAGTCTCTAAACTCTGATATACTTTGCATCCACTCGGAATCTAGTATCCCT  
 GATCGCTTCATAGGCAGTGGATCTGGGACAGATTCACTCTTACCATCAGCAGTGTGCAGGCTGA  
 AGACCTGGCAGATTACTCTGTCAAGCAACATTATAGCAGTCCGTCACGTTGGTCTGGGACCA  
 AGCTGGAGCTGAAAGCTGTGGTGCACCACATGTCTTCACTTCCCGCCATCTGATGAGCAGTTG  
 AAATCTGGAACTGCCTCTGTGTGCCTGTAATAACTCTATCCCAGAGAGGCAAAGTACA  
 GTGGAAGGTGGATAACGCCCTCAATCGGTAACTCCCAGGAGAGTGTACAGAGCAGGACAGCA  
 AGGACAGCACCTACAGCCTCAGCAGCACCCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAA  
 GTCTACGCCCTGCAAGTCACCCATCAGGGCTGAGCTGCCCGTCACAAAGAGCTTCAACAGGGG  
 AGAGTGT

SEQ ID NO.:4

DIVMTQSPSSLAVSIGQKVTMNCSSQSLNSNFQKNFLAWYQQKPGQSPKLLIYFASTRESSIP  
 DRFIGSGSGTDFTLTISSVQAEDLADYFCQHQHSTPLTFGAGTKLELKAVAAPSVFIFPPSDEQL  
 KSGTASVVCLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYLSSTTLSKADYEKHK  
 VYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO.:5

GAGGTTCAGCTGCAGCAGTCTGTAGCTGAGCTGGTGAGGCCTGGGCTTCAGTGACGCTGTCCTG  
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 GCCTGGAATGGATTGGGTTATTGATCCTGAAACTGGTAATACTGCCCTCAATCAGAAGTCAAG  
 GGCAAGGCCACACTGACTGCAGACATATCCTCAGCACGCTACATGGAACTCAGCAGTTGAC  
 ATCTGAGGACTCTGCCCTATTACTGTATGGTTATTCTGATTATTGGGCAAGGCACCACTC  
 TCACAGTCTCTCAGCCTAACGAAGGGCCATCTGTCTTCCCTGGCCCCCTCCCTCAAGAGC  
 ACCTCTGGGGCACAGCGCCCTGGCTGCCTGGTCAAGGACTACTTCCCGAACCGGTGACGGT  
 GTCGTGAACTCAGGCCCTGACCAGCGCGTGCACACCTCCGGCTGTCCTACAGTCTCAG  
 GACTCTACTCCCTCAGCAGCGTGGTACCGTGCCTCAGCTGGGACCCAGACATC  
 TGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCAAATCTGTGA

ATTCACTCACACATGCCACCGTGCACCGTGAACCTGAACCTGGGGGGACCGTCAGTCTTCCCT  
 TCCCCCCTAAACCCAAGGACACCCCTCATGATCTCCGGACCCCTGAGGTACATGCGTGGTGGT  
 GACGTGAGCCACGAAGACCCCTGAGGTCAAGTCAACTGGTACGTGGACGGCGTGGAGGTGCATAA  
 TGCCAAGACAAAGCCGGGGAGGAGCAGTACACAGCACGTACCGTGTGGTCAAGGTCTCAAACAAAGCCCT  
 TCCCTGACCAAGGACTGGCTGAATGGCAAGGAGTACAAGTGAAGGTCTCAAACAAAGCCCT  
 GCCCCCATCGAGAAAACCATCTCAAAGCCAAGGGCAGCCCCGAGAACACACAGGTGTACACCC  
 GCCCCCATCCGGGATGAGCTGACCAAGAACCGGTACGCTGACCTGCTGGTCAAAGGTTCT  
 ATCCCAGCGACATGCCGTGGAGTGGAGAGCAATGGCAGCCGGAGAACAACTACAAGACCACG  
 CCTCCCGTGTGGACTCCGACGGCTCTCTCCTACAGCAAGCTACCGTGGACAAGAGCAG  
 GTGGCAGCAGGGAACGTCTCTCATGCTCCGTATGCGTGGAGGCTCTGCACAAACCACTACACGC  
 AGAAGAGCCCTCCCTGTCTCCGGAAA

SEQ ID NO.:6

EVQLQQSVAELVRPGASVTLSCSKASGYIFTDYEIHVKQTPVHGLEWIGVIDPETGNTAFNQKFK  
 GKATLTADISSSTAYMELSSLTSEDSAVYYCMGSDYWGQGTTLTVSSASTKGPSVFPLAPSSKS  
 TSGGTAALGCLVKDYFPEPVTVWSNSGALTSGVHTFPVALQSSGLYSLSSVTPSSSLGTQTYI  
 CNVNHKPSNTKVDKKVEPKSCEFTHTCPCPAPELLGGPSVFLFPPPKDLMISRTPEVTCVV  
 DVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALP  
 APIEKTISKAKGQPREPVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTT  
 PPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHTQKSLSLSPGK

SEQ ID NO.:7

GATTTGATGACCCAAACTCCACGCTCCCTGTCTGTCAGTCTGGAGATCAAGCCTCCATCTC  
 TTGTAGATCGAGTCAGAGCTTACATAGTAATGAAACACCTATTTAGAATGGTATTCAGA  
 AACCAAGGCCAGCCTCCAAAGGTCTGATCTACAAAGTTCAACCGATTTCTGGGTCCCAGAC  
 AGGTTCACTGGCAGTGGATCAGGGACAGATTCACTCAAGATCAGCGAGTGGAGGCTGAGGA  
 TCTGGGAGTTATTACTGCTTCAGGTTACATGTTCTCAGCTCGGTCTGGGACCAAGC  
 TGGAGCTGAAAGCTGGCTGCACCCTGTCTCATCTTCCGCATCTGATGAGCAGTTGAAA  
 TCTGGAACCTGCCTCTGTTGTGCTGCTGAATAACTCTATCCCAGAGAGGCCAAAGTACAGTG  
 GAAGGTGGATAACGCCCTCAACTGGGTAACCTCCAGGAGAGTGTACAGAGCAGGACAGCAAGG  
 ACAGCACCTACAGCCTCAGCAGCACCCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTC  
 TACGCCTGCGAAGTCACCCATCAGGGCTGAGCTGCCGTACAAAGAGCTTCAACAGGGAGA  
 GTGT

SEQ ID NO.:8

DVLMTQTPRSLSVSLGDQASISCRSSQSLHSNGNTYLEWYLQKPGQPKVLIYKVSNRFSGVPD  
 RFSGSGSGTDFTLKISGVEAEDLGVYYCFQGSHVPLTFGAGTKLELKAVAAPSVFIFPPSDEQLK  
 SGTAASVVCLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYLSSTLTLKADYEHKVV  
 YACEVTHQGLSSPVTKSFRGEC

SEQ ID NO.:9

GAGATCCAGCTGCAGCAGTCTGGACCTGAGTTGGTAAGCCTGGGCTTCAGTGAAGATATCCTG  
 TAAGGCTCTGGATACACCTTCACTGACAACATGAACTGGTGAAGCAGAGCCATGGAAAGA  
 GCCTTGAGTGGATTGGAGATATTAATCCTTACTATGGTACTACTACCTACAACCAGAAGTCAAG  
 GGCAAGGCCACATTGACTGTAGACAAGTCCTCCGCACAGCCTACATGGAGCTCCGGCCCTGAC  
 ATCTGAGGACTCTGCAGTCTATTACTGTGCAAGAGATGACTGGTTGATTATGGGCAAGGG  
 CTCTGGTCACTGTCTCGAGCCTCAACGAAGGGCCATCTGTTCTCCCTGGCCCTCC  
 AAGAGCACCTCTGGGGCACAGCGCCCTGGCTGCTGGTCAAGGACTACTTCCCCGAACCG  
 GACGGTGTGCTGGAACTCAGGCCCCCTGACCAAGCGGCGTGCACACCTTCCGGCTGCTACAGT  
 CCTCAGGACTCTACTCCCTCAGCAGCTGGTACCGTGCCTCCAGCAGCTGGCACCCAGACC  
 TACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAAGAAAGTTGAGCCAAATC  
 TTGTGAATTCACTCACACATGCCACCGTCCCCAGCACCTGAACCTCTGGGGACCGTCAGTCT  
 TCCTCTTCCCCAAAACCAAGGACACCCATGATCTCCGGACCCCTGAGGTACATCGTG

GTGGTGGACGTGAGCCACGAAGACCCCTGAGGTCAAGTCAACTGGTACGTGGACGGCGTGGAGGT  
 GCATAATGCAAGACAAAGCCGGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCGTCC  
 TCACCGTCTGCACCAAGGACTGGCTGAATGCAAGGAGTACAAGTCAAGGTCTCAACAAAGCC  
 CTCCCAGCCCCATCGAGAAACCATCCTCAAAGCCAAGGGCAGCCCGAGAACACAGGTGTA  
 CACCCCTGCCCATCCGGATGAGCTGACCAAGAACCCAGGTGACCTGACCTGCTGGTCAAAG  
 GCTTCTATCCCAGCGACATCGCCGTGGAGTGGAGAGCAATGGGAGCCGGAGAACAACTACAAG  
 ACCACGCCTCCGTGGACTCCGACGGCTCTCTACAGCAAGCTACCGTGGACAA  
 GAGCAGGTGGCAGCAGGGAACGTCTCTACATGCTCCGTGATGCATGAGGCTCTGCACAAACCACT  
 ACACGCAGAAGACCTCTCCGTCTCCGGAAA

SEQ ID NO.:10

EIQLQQSGPELVKPGASVKISCKASGYTFTDNYMNWVKQSHGKSLEWIGDINPYGTTYNQKFK  
 GKATLTVDKSSRTAYMELRGLTSEDAVYYCARDDWFDYWGQGLTVVSAASTKGPSVFPLAPSS  
 KSTSGGTAALGCLVKDYFPEPVTVWSNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSQLGTQT  
 YICNVNHPNSNTKVDKKVEPKSCEFTHTCPCPAPPELLGGPSVLFPPPKDLMISRTPEVTCV  
 VVDVSHEDPEVFKFNWYVGVEVHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKA  
 LPAPIEKTIKAKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYK  
 TTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVHEALHNHTQKSLSLSPGK

SEQ ID NO.:11

GACATCGTTATGTCTCAGTCTCCATCTTCCATGTATGCATCTCTAGGGAGAGAGTCACATCAC  
 TTGCAAGGCCAGTCAGGACATTCTACAACTTTAAACTGGTCCAGCAGAAACCCAGGAAATCTC  
 CAAAGACCCCTGATCTTGTGCAAACAGATTGGTAGATGGGTCCCATCAAGGTTCACTGGCAGT  
 GGATCTGGCAAGATTATTCTCTCACCATCAGCAGCCTGGAGTTGAAGATTTGGAAATTATT  
 TTGTCTACAGTATGATGAGATTCCGCTCACGTTGGTCTGGGACCAAGCTGGAGCTGAGAGCTG  
 TGGCTGCACCATCTGCTTCTACCTCCGCCATCTGATGAGCAGTTGAAATCTGAACTGCCCT  
 GTTGTGTGCTGCTGTAATAACTCTATCCCAGAGAGGCCAAAGTACAGTGGAAAGGTGGATAACGC  
 CCTCCAATCGGGTAACCTCCAGGAGAGTGTACAGAGCAGGACAGCAAGGACAGCACCTACAGCC  
 TCAGCAGCACCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCACTGCCCTGCGAAGTC  
 ACCCATCAGGGCTGAGCTGCCGTACAAAGAGCTTCAACAGGGAGAGTGT

SEQ ID NO.:12

DIVMSQSPSSMYASLGERVTITCKASQDIHNFLNWFQQKPGKSPKTLIFRANRLVDGVPSRFSGS  
 GSGQDYSLTISLLEFEDLGIYSCLQYDEIPLTFGAGTKLELRAVAAPSVFIFPPSDEQLKSGTAS  
 VVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTKADYEKHKVYACEV  
 THQGLSSPVTKSFNRGEC

SEQ ID NO.:13

GAGGTGCAGCTTCAGGAGTCAGGACCTGACCTGGTAAACCTTCTCAGTCACTTCACTCACCTG  
 CACTGTCACTGGCTTCTCCATCACCACTGGTTATGGCTGGCACTGGATCCGGCAGTTCCAGGAA  
 ACAAACTGGAGTGGATGGGCTACATAAACTACGATGGTCACAATGACTACAACCCATCTCTCAAA  
 AGTCGAATCTCTATCACTCAAGACACATCCAAGAACAGCTTCTCAGTCAGTTGAATTCTGTGAC  
 TACTGAGGACACAGCCACATATTACTGTGCAAGCAGTTACGACGGCTTATTGCTTACTGGGGCC  
 AAGGGACTCTGGTCACTGTCTCTGCAGCCTCAACGAAGGGCCCATGTCTTCCCCTGGGGCC  
 TCCTCCAAGAGCACCTCTGGGGCACAGCGGCCCTGGCTGCAGTCAGGACTACTTCCCCTGA  
 ACCGGTGACGGTGTCTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCCTGTCC  
 TACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTACCGTGCCTCCAGCAGCTGGGCACC  
 CAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGGACAAGAAAGTTGAGCC  
 CAAATTTGTGAATTCACTCACACATGCCACCGTGCAGCACCTGAACCTCTGGGGGACCGT  
 CAGTCTCCCTCTTCCCCCAAAACCCAAGGACACCCATGATCTCCCGGACCCCTGAGGTACAA  
 TGCCTGGTGGTGGACGTGAGCCACGAAGACCTGAGGTCAGTTCAACTGGTACGTGGACGGCGT  
 GGAGGTGCATAATGCCAAGACAAAGCCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCA  
 CGCTCCTACCGCTCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAGGTCTCCAAC

AAAGCCCTCCAGCCCCATCGAGAAAACCATCTCAAAGCAAAGGGCAGCCCCGAGAACACACA  
 GGTGTACACCTGCCCATCCGGATGAGCTGACCAAGAACCCAGGTCAAGCTGACCTGCCCTGG  
 TCAAAGGCTCTATCCAGCGACATGCCGTGGAGTGGAGAGCAATGGCAGCCGGAGAACACAC  
 TACAAGACCACGCCCTCCGTCTGGACTCCGACGGCTCTTCCCTACAGCAAGCTCACCGT  
 GGACAAGAGCAGGTGGCAGCAGGGAACGTCTTCTCATGCTCCGTATGAGGCTCTGCACA  
 ACCACTACACCGAGAACAGCCTCTCCCTGTCTCCGGAAA

SEQ ID NO.:14

EVQLQESGPDLVKPQSLSLTCTVTGFSITSGYGWHIRQFPGNKLEWMGYINYDGHNDYNPSLK  
 SRISITQDTSKNQFFLQLNSVTTEDATYYCASSYDGLFAYWGQGTIVTVAASTKGPSVFPLAP  
 SSKSTSGGTAALGCLVKDYFPEPVTVWSWNSGALTSGVHTFPAVLQSSGLYLSLSSVTVPSLGT  
 QTYICNVNHPNSNTKVDKKVEPKSCEFTHTCPPCPAPELGGPSVFLFPPKPKDTLMISRTPEVT  
 CVVVDVSHEDPEVKFNWYVDGVEVNAKTPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSN  
 KALPAPIEKTIASKKGQPREPVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENN  
 YKTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSPGK

SEQ ID NO.:15

GACATTGTGATGACCCAGTCTCCATCCTCCCTGGCTGTCAATAGGACAGAACGGTCACTATGAA  
 CTGCAAGTCAGTCAGAGCCTTTAAATAGTAACCTTCAAAAGAACCTTTGCCCTGGTACCAAGC  
 AGAAACCAGGCCAGTCCTAAACTCTGATATACTTGCATCCACTGGGAATCTAGTATCCCT  
 GATCGCTTCATAGGAGTGGATCTGGACAGATTCTACCTTACCATCAGCAGTGTGCAGGCTGA  
 AGACCTGGCAGATTACTCTGTCAAGAACATTAGCAGTCACGTTGGTCTGGGACCA  
 AGCTGGAGCTGAAA

SEQ ID NO.:16

DIVMTQSPSSLAVSIGQKVTMNCKSSQSLNSNFQKNFLAWYQQKPGQSPKLLIYFASTRESSIP  
 DRFIGSGSGTDFLTISVQAEDLADYFCQQHYSTPLTFGAGTKLELK

SEQ ID NO.:17

GAGGTTCAGCTGCAGCAGTCAGCTGTAGCTGAGCTGGTGGAGCCCTGGGGCTTCAGTGACGGCTGTCC  
 CAAGGCTCGGGCTACATATTACTGACTATGAGATAACACTGGGTGAAGCAGACTCCTGTGCATG  
 GCCTGGAATGGATTGGGGTATTGATCCTGAAACTGGTAATACTGCCTTCATCAGAACGGTCAAG  
 GGCAAGGCCACACTGACTGCAGACATATCCTCCAGCACAGCCTACATGGAACCTCAGCAGTTGAC  
 ATCTGAGGACTCTGCCGTCTATTACTGTATGGGTTATTCTGATTATTGGGCCAGGCACCACTC  
 TCACAGTCCTCA

SEQ ID NO.:18

EVQLQQSVAELVRPGASVTLSCASGYIFTDYEIHVKQTPVHGLEWIGVIDPETGNTAFNQKFK

GKATLTADISSSTAYMELSSLTSEDSAVYYCMGYSDYWGQGTTLVSS

SEQ ID NO.:19

GATGTTTGATGACCCAAACTCCACGCTCCCTGTCTGTCAGTCTGGAGATCAAGCCTCCATCTCTGAGATCGAGTCAGAGCCTTACATAGTAATGAAACACCTATTTAGAATGGTATTGCAGAACCAGGCCAGCCTCAAAGGTCTGATCTACAAAGTTCCAACCGATTTCTGGGTCCCAGACAGGTTCAAGTGGCAGTGGATCAGGACAGATTCAACTCAAGATCAGCGGAGTGGAGGCTGAGGATCTGGGAGTTATTACTGCTTCAGGTTACATGTTCTCACGTTCGGTCTGGGACCAAGCTGGAGCTGAAA

SEQ ID NO.:20

DVLMTQTPRSLSVSLGDQASISCRSSQSLLHSNGNTYLEWYLQKPGQPPKVLIYKVSNRFSGVPDFSGSGSGTDFTLKISGVEAEDLGVYYCFQGSHVPLTFGAGTKLELK

SEQ ID NO.:21

GAGATCCAGCTGCAGCAGTCTGGACCTGAGTTGGTGAAGCCTGGGCTTCAGTGAAGATATCCTGTAAGGCTCTGGATACACCTCACTGACAACATACATGAACGGTGAAGCAGAGCCATGGAAAGAGCCTGAGTGGATTGGAGATATTAATCCTTACATGGTACTACTACCTACAACCAGAAGTCAAGGGCAAGGCCACATTGACTGTAGACAAGTCCTCCGCACAGCCTACATGGAGCTCCGGCCTGACATCTGAGGACTCTGCAGTCTATTACTGTGCAAGAGATGACTGGTTGATTATTGGGCCAAGGGACTCTGGTCACTGTCTTGCA

SEQ ID NO.:22

EIQLQQSGPELVPGASVKISCKASGYTFTDNYMNWKQSHGKSLEWIGDINPYGTTYNQKFKGKATLTVDKSSRTAYMELRGLTSEDSAVYYCARDDWFDYWGQGTLTVSA

SEQ ID NO.:23

GACATCGTTATGTCTCAGTCTCCATCTTCCATGTATGCATCTCTAGGAGAGAGAGTCACTATCACTTGCAAGGGAGACTCAGGACATTCTACATAACTTTAACTGGTCCAGCAGAAACCAGGAAATCTCAAAGACCCGTATCTCGTCAAACAGATTGGTAGATGGGTCCCATCAAGGTTCAAGTGGCAGTGGATCTGGGCAAGATTATTCTCTACCATCAGCAGCCTGGAGTTGAAGATTGGGAATTATTC TTGTCTACAGTATGATGAGATTCCGTCACGTTGGTCTGGGACCAAGCTGGAGCTGAGA

SEQ ID NO.:24

DIVMSQSPSSMYASLGERVTITCKASQDIHNFLNWFFQQKPGKSPKTLIFRANRLVDGVPSRSGSGSGQDYSLTISLEFEDLGIYSCLQYDEIPLTFGAGTKLELR

SEQ ID NO.:25

GAGGTGCAGCTTCAGGAGTCAGGACCTGACCTGGTGAACACCTCTCAGTCACTTCACTCACCTGCACTGCACTGGCTTCATCACCAAGTGGTTATGGCTGGCACTGGATCCGGCAGTTCCAGGAAACAAACTGGAGTGGATGGCTACATAAAACTACGATGGTCAAATGACTACAACCCATCTCTCAAAGTCGAATCTCTATCACTCAAGACACATCCAAGAACCGATTCTCTGCAGTTGAATTCTGTGACTACTGAGGACACAGCACATATTACTGTGCAAGCAGTTACGACGGCTATTGCTACTGGGCCAAGGGACTCTGGTCACTGTCTTGCA

SEQ ID NO.:26

EVQLQESGPDLVKPSQSLSLTCTVTGFSITSGYGWHWIRQFPGNKLEWMGYINYDGHNDYNPSLKSRSISITQDTSKNQFFLQLNSVTTEDTATYYCASSYDGLFAYWGQGTLTVSA

SEQ ID NO.:27

KSSQSLLNSNFQKNFLA

SEQ ID NO.:28

FASTRES

SEQ ID NO.:29

QQHYSTPLT

SEQ ID NO.:30

GYIFTDYEIH

SEQ ID NO.:31

VIDPETGNTA

SEQ ID NO.:32

MGYSDY

SEQ ID NO.:33

RSSQSLLHSNGNTYLE

SEQ ID NO.:34

KVSNRFS

SEQ ID NO.:35

FQGSHVPLT

SEQ ID NO.:36

GYTFTDNYMN

SEQ ID NO.:37

DINPYYGTTT

SEQ ID NO.:38

ARDDWF DY

SEQ ID NO.:39

KASQDIHNFLN

SEQ ID NO.:40

RANRLVD

SEQ ID NO.:41

LQYDEIPLT

SEQ ID NO.:42

GFSITSGY GWH

SEQ ID NO.:43

YINYDGHND

SEQ ID NO.:44

ASSYDGLFAY

SEQ ID NO.:45 - 3A4 heavy chain variable region nucleotide sequence

CAGATCCAGTTGGTGCAATCTGGACCTGAGATGGTGAAGCCTGGGCTTCAGTGAAGATGTCCTG  
TAAGGCTCTGGATAACACATTCACTGACGACTACATGAGCTGGGTGAAACAGAGCCATGAAAGA  
GCCTTGAGTGGATTGGAGATATTAATCCTTACAACGGTGTACTAACTACAACCAGAAGTCAAG  
GGCAAGGCCATATTGACTGTAGACAAATCCTCCAGCACAGCCTACATGCAGCTAACAGCCTGAC

ATCGGAAGACTCAGCAGTCTATTACTGTGCAAGAGACCCGGGGCTATGGACTACTGGGTCAAG  
GAACCTCAGTCACCGTCTCCTCA

SEQ ID NO.:46 - 3A4 heavy chain variable region polypeptide sequence  
QIQLVQSGPEMVKGASVMSCKASGYTFDDYMSWVKQSHGKSLIEWIGDINPYNGDTNYNQKFK  
GKAILTVKSSSTAYMOLNSLTSEDAVYYCARDPGAMDYWGQGTSVTVSS

SEQ ID NO.:47 - 3A4 light chain variable region nucleotide sequence  
GATGTTGTGATGACCCAACTCCACTCTCCCTGGCTGTCAGTCTGGAGATCAAGCCTCCATCTC  
TTGCAGATCTAGTCAGAGCCTCTACATAGTAATGGAAACACCTATTTAGAATGGTACCTCAGA

AACCGGCCAGTCTCAAAGCTCCTGATCCACACAGTTCCAACCGATTTCTGGGGTCCCAGAC  
AGATTCACTGGCAGTGGATCAGGGACAGATTCACTCAAGATCAGCAGAGTGGAGGCTGAGGA  
TCTGGGAGTTTATTACTGCTTCAAGGTTCACATGTTCCGCTCACGTTCGGTGCTGGGACCAGGC  
TGGAGCTGAAA

SEQ ID NO.:48 - 3A4 light chain variable region polypeptide sequence  
DVVMTQTPLSLAVSLGDQASISCRSSQSLIHNSNGNTYLEWYLQKPGQSPKLLIHTVSNRFSGVPRFSGSGSGTDFTLKISRVEADLGVYYCFQGSHVPLTFGAGTRLEK

SEQ ID NO.:49 - 3A4 heavy chain CDR1 polypeptide sequence GYTFTDDYMS

SEQ ID NO.:50 - 3A4 heavy chain CDR2 polypeptide sequence DINPYNGDTNYNQKFKG

SEQ ID NO.:51 - 3A4 heavy chain CDR3 polypeptide sequence DPGAMDY

SEQ ID NO.:52 - 3A4 light chain CDR1 polypeptide sequence RSSQSLLHSNGNTYLE

SEQ ID NO.:53- 3A4 light chain CDR2 polypeptide sequence TVSNRFS

SEQ ID NO.:54 - 3A4 light chain CDR3 polypeptide sequence FQGSHVPLT

SEQ ID NO.:55

GTAAAGCAGGCGTGTGGCTGCAACCATCTGTCTTC

SEQ ID NO.:56

GTAAAGCGCTAGCCCTAACACTCTCCCCCTGTGAAAGC  
SEQ\_ID: NC\_0053

SEQ ID NO.: 37

GCCTGGCTGCCACATCTGCTTCTCTGGCTGAATAACTCTATCCAGAGAGGCCAAAGTACAGTGAAGGGTGGATA  
ACGCCCTCCAATCGGGTAACTCCCAGGAGAGTGTACAGAGCAGGACAGCAAGGCACGACCTAC  
AGCCTCAGCAGCACCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACGCTGCGA  
AGTCACCCATCAGGCCTGAGCTGCCGTACAAAGAGCTTCAACAGGGGAGAGTGTAG

SEQ ID NO.:58

AVAAPSVFIFPPSDEQLKSGTASVVCNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTY  
SLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO.:59

CTT GAGCCGGCGGATGGTCGAGGTGAGGTGTGGCAGGCTTGAGATCCAGCTGGGGTGAGTAC  
TCCCTCTAAAAGCGGGCATTACTCTCGCGCTAACGATTGTCAGTTCCAAAAGCAGGAGGAGTT  
GATATTACCTGGCCGATCTGGCCATACACTTGAGTGACAATGACATCCACTTGCCTTCTCT  
CCACAGGTGTCCACTCCCAGGTCCAAGTTAACGGATCTAGCGAATTGAACTTCTGCT  
GTCTTGGGTGCATTGGAGCCTTGCCTTGCTCTACCTCCACCATGCCAAGTGGTCCCAGGCTT  
GAGACGGAGCTTACAGCGCTGCGCTGCACCACATCTGTCTTCATCTCCCGCCATCTGATGAGCAG  
TTGAAATCTGGAACTGCCTCTGTTGTGCGCTGCTGAATAACTTCTATCCCAGAGAGGGCCAAGT

ACAGTGGAAAGGTGGATAACGCCCTCCAATCGGGTAACCTCCAGGAGAGTGTACAGAGCAGGACA  
 GCAAGGACAGCACCTACAGCCTCAGCAGCACCTGACGCTGAGCAAAGCAGACTACGAGAAACAC  
 AAAGTCTACGCCCTGCGAAGTCACCCATCAGGGCCTGAGCTGCCCGTCACAAAGAGCTTCAACAG  
 GGGAGAGTGTAGGGTACCGCGGGCGCTTCGAATGAGATCCCCGACCTGACCTCTGGCTAATA  
 AAGGAAATTATTTCAATTGCAATAGTGTGTGGAATTGGTGTCTCACTCGGAAGGACAT  
 ATGGGAGGGCAAATCATTGGTCAGATCCCTCGAGATCTAGCTAGCTAGAGCCCCGCCGCG  
 GAACTAAACCTGACTACGGCATCTCGCCCCCTCTCGGGGGCAGTGCATGTAATCCCTCAGT  
 TGGTTGGTACAACCTGCCAACCTGGCCCTGTTCCACATGTGACACAGGGGGGACCAAACACAAA  
 GGGGTTCTCTGACTGTAGTTGACATCCTATAAAATGGATGTGACATTTGCCAACACTGAGTGGC  
 TTTCATCCTGGAGCAGACTTGTGAGTCTGTGACTGCAACACAACTGGCTTATGTGTAAC  
 TTGGCTGAAGCTCTACACCAATGCTGGGGACATGTACCTCCCAGGGGCCAGGAAGACTACGG  
 GAGGCTACACCAACGTCACAGAGGGCCTGTGAGCTACCGATAAGCGGACCCCAAGAGGGC  
 ATTAGCAATAGTGTATAAGGCCCCCTGTTAACCTAAACGGTAGCATATGCTCCGGTA  
 GTAGTATATACTATCCAGACTAACCTAACTCAATAGCATATGTTACCCAACGGGAAGCATATGC  
 TATCGAATTAGGGTTAGTAAAAGGGCTTAAGGAACAGCGATATCTCCCACCCATGAGCTGTCA  
 CGGTTTATTACATGGGTCAGGATTCCACGGAGGGTAGTGAACCATTTAGTCACAAGGGCAGT  
 GGCTGAAGATCAAGGAGCGGGCAGTGAACCTCCTGAATCTCGCCGCTCTCATTCTCCTC  
 GTTAGCTAATAGAATAACTGCTGAGTTGTAACAGTAGTGTAGGTGAGGTGCTGAAACAA  
 GTTTTCAGGTGACGCCCGAGAATAAAATTGGACGGGGGTCTAGTGGCCATTGTGCTATGA  
 CACCAATATAACCCCTCACAAACCCCTGGGCAATAAAATACTAGTGTAGGAATGAAACATTGAA  
 TATCTTAACATAGAAATCCATGGGTGGGACAAGCCGTAAGACTGGATGTCCATCTCACAC  
 GAATTATGGCTATGGCAACACATAATCCTAGTGAATATGATACTGGGTTATTAAGATGTG  
 CCCAGGCAGGGACCAAGACAGGTGAACCATGTTGTTACACTCTATTGTAACAAGGGGAAAGAGA  
 GTGGACGCCGACAGCAGCGGACTCCACTGGTGTCTAAACACCCCGAAAATTAAACGGGCTC  
 CACGCCAATGGGGCCATAAAACAAAGACAAGTGGCCTCTTTTGAAATTGGAGGG  
 GCACGGCTCAGCCCCACACGGCCCTGCCGTTGGACTGTAAGGTTGCTAATACCTG  
 GCTGATTGTAACCCGCTAACCAACTGCGGTCAAACCAACTGCCCACAAACACTAATGGCACCC  
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 CGCTGAGGACACGGTGGCTAATGTTGCCATGGTAGCATATACTACCCAAATATCTGGATAGCA  
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 TATCCTAACTCTATCTGGTAGTATGCTATCCTAACTCTATATCTGGTAGCATAGGCTATCC  
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 CTGTATCCGGTAGCATATGCTATCCTACGATGATAAGCTGCTAAACATGAGAATTAACTCTG  
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 CATTCAAATATGTTACCGCTCATGAGACAATAACCCGATAAAATGCTCAATAATTGAAAAG  
 GAAGAGTATGAGTATCAACATTCCGTGTCGCCCTATTCCCTTTTGCGGCATTGGCTTC  
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 ACAGAAAAGCATCTACGGATGGCATGACAGTAAGAGAATTATGCACTGCTGCCATAACCATGAG  
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 TGCACAACATGGGGGATCATGTAACTCGCCTGATCGTGGGAACCGGAGCTGAATGAACCCATA  
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 GCAAAACGCCCTCTCCCCCGCGCTGGCCGATTCATTAACTGCAGCTGGCACGACAGGTTCCGAC  
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 AGATCTATACATTGAATCAATATTGGCAATTAGCCATATTAGTCATTGGTTATATAGCATAAATC  
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 GGAGGTCTATATAAGCAGAGCTGTTAGTGAACCGTCAGATCCTCACTCTTCCGATCGCTG  
 TCTGCGAGGGCCAGCTGTTGGGCTCGCGGTTGAGGACAAACTCTTCGCGGTCTTCCAGTACTCT  
 TGGATCGGAAACCCGTCGGCTCCGAACGGTACTCCGCCACCGAGGGACCTGAGCGAGTCCGCAT  
 CGACCGGATCGGAAAACCTCTCGAGAAAGCGTCAACCAGTCACAGTCGAAGGTAGGCTGAGC  
 ACCGTGGCGGGCGGAGCGGGTGGCGGTGGGGTTGTTCTGGCGGAGGTGCTGATGATGTA  
 ATTAAAGTAGGCGGT

SEQ DI NO.:60

ATGCCAAGTGGTCCCAGGCTGACATTGTGATGACCCAGTCTCC

SEQ ID NO.:61

ATGCCAAGTGGTCCCAGGCTGATGTTTGATGACCCAAACTCC

SEQ ID NO.:62

ATGCCAAGTGGTCCCAGGCTGACATCGTTATGTCTCAGTCTCC

SEQ ID NO.:63

GGGAAGATGAAGACAGATGGTGCAGCCACAGC

SEQ ID NO.:64

GTAAGCGCTAGCGCCTCAACGAAGGGCCCCTGTCTTCCCTGGCCCC

SEQ ID NO.:65

GTAAGCGAATTACAAGATTGGGCTCAACTTCTTCTG

SEQ ID NO.:66

GCCTCCACCAAGGGCCATCGGTCTTCCCCCTGGCACCCCTCCAAAGAGCACCTCTGGGGCAC  
 AGCAGCCCTGGCTGCCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGTGGAACCTCAG  
 GCGCCCTGACCAGCGCGTGCACACCTCCCGCTGTCTACAGTCTCAGGACTCTACTCCCTC  
 AGCAGCGTGGTACCGTGCCTCCAGCAGCTGGGACCCAGACCTACATCTGCAACGTGAATCA  
 CAAGCCCAGAACACCAAGGTGGACAAGAAAGTTGAGGCCAATCTTGT

SEQ ID NO.:67

ASTKGPSVPLAPSSKSTSGTAALGCLVKDYFPEPVTWSWNSALSGVHTFPALQSSGLYSL  
 SSVVTVPSSLGTQTYICNVNHPNSNTKVDKKVEPKSC

SEQ ID NO.:68

CTTGAGCCGGCGGATGGTCGAGGTGAGGTGAGGCTTGAGATCCAGCTGTTGGGTGAGTAC  
 TCCCTCTCAAAAGCGGGCATTACTCTCGCCTAAGATTGTCAGTTCCAAAACGAGGAGGATTT

GATATTCACCTGGCCCGATCTGGCCATACACTTGAGTGACAATGACATCCACTTGCCCTTCTCT  
 CCACAGGTGTCACCTCCCAGGTCCAAGTTGCCGCCACATGGAGACAGACACACTCCTGCTATG  
 GGTACTGCTGCTCTGGTTCCAGGTTCCACTGGCGGAGACGGAGCTTACGGGCCATCTGCTTT  
 CCCCTGGCCCCCTCCTCCAAGAGACACCTCTGGGGCACAGCGGCCCTGGCTGCCCTGGTCAAGGA  
 CTACTTCCCCGAACCGGTGACGGTGTGGAACTCAGGCCTGACCAGCGGTGACACCT  
 TCCCGGCTGTCCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTACCGTGCCCTCCAGC  
 AGCTTGGCACCCAGACACTACATGCAACGTGAATCACAAGCCCAGCAACACCAAGGTGGACAA  
 GAAAGTTGAGCCAAATCTTGTGAATTCACTCACACATGCCACCGTGCCAGCACCTGAACCTC  
 TGGGGGACCGTCAGTCTCCTCTCCCCAAAACCAAGGACACCCCTCATGATCTCCGGACC  
 CCTGAGGTACATGCGTGGTGGACGTGAGGCCACGAAGACCTGAGGTCAAGTTCAACTGGTA  
 CGTGGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGGAGCAGTACAACAGCACGT  
 ACCGTGTGGTCAGCCTCACCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGC  
 AAGGTCTCCAACAAAGCCCTCCAGCCCCATCGAGAAAACATCTCAAAGCCAAGGGCAGCC  
 CCGAGAACACAGGTGTACACCCCTGCCCATCCGGATGAGCTACCAAGAACAGGTGAGCC  
 TGACCTGCTGGTCAAAGGCTTCTATCCCAGCGACATGCCGTGGAGTGGAGAGCAATGGCAG  
 CCGGAGAACAACTACAAGACCACGCCCTCCGTGCTGGACTCCGACGGCTCCTCTCCTACAG  
 CAAGCTCACCGTGGACAAGAGCAGGTGGCAGCAGGGAACGTCTCTCATGCTCCGTGATGCATG  
 AGGCTCTGACAACCAACTACACGAGAACAGGCTCTCCGTCTCCGGAAATGATCCCCGAC  
 CTCGACCTCTGGCTAAAGGAAATTATTTCATTGCAATAGTGTGGAAATTGGTGTGTC  
 TCTCACTCGGAAGGACATATGGGAGGGAAATCATTGTCGAGATCCCTCGGAGATCTCTAGCT  
 AGAGCCCCGCCGCCGAGCAACTAAACCTGACTACGGCATCTGCCCTCTCGCGGGCAGT  
 GCATGTAATCCCTCAGTGGTTGGTACAACCTGCAACTGAACCTAAACGGTAGCATATGCT  
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 AGCATATGCTATCCTAATCTATATCTGGTAGCATATGCTATCCTAATCTATATCTGGTAGCATA  
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 CGTTGACGCAAATGGCGTAGGCGTGTACGGTGGGAGGTATATAAGCAGAGCTCGTTAGTG  
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 GAGGACAAACCTTCCGGCTTCCAGTACTCTGGATCGGAAACCGTCGGCTCCGAACGGT  
 ACTCCGCCACCGAGGGACCTGAGCGAGTCCGCATCGACCGGATCGGAAACCTCTCGAGAAAGGC  
 GTCTAACCAAGTCACAGTCGAAGGTAGGCTGAGCACCGTGGCGGGCGCAGCGGGTGGCGGT  
 GGTTGTTCTGGCGAGGTGCTGCTGATGATGTAATTAAAGTAGGCAGT

SEQ ID NO.:69

GGGTTCCAGGTTCCACTGGCGAGGTTCAGCTGCAGCAGTCTGT

SEQ ID NO.:70

GGGTTCCAGGTTCCACTGGCGAGGTCAGCTCAGGAGTCAGG

SEQ ID NO.:71

GGGGCCAGGGAAAGACAGATGGGCCCTCGTTGAGGC

SEQ ID NO.: 89: Exemplary embodiment of CDRL1 K-S-S-Q-S-L-L-N/H-S/T-S/N/D-N/G-Q/N/K-K/L-N-Y-L-A

SEQ ID NO.:90: Exemplary embodiment of CDRL1 K-A-S-Q-D-I-H-N/T-Y/F-L-N

SEQ ID NO.:91: Exemplary embodiment of CDRL2 F-A-S-T-R-E-S

SEQ ID NO.: 92: Exemplary embodiment of CDRL2 L-V-S-K-L-D-S

SEQ ID NO.:93: Exemplary embodiment of CDRL2 R-A-N-R-L-V-D

SEQ ID NO.:94: Exemplary embodiment of CDRL3 Q-Q-H-Y-S-T-P-L-T

SEQ ID NO.:95: Exemplary embodiment of CDRL3 W/L-Q-Y/G-D/T-A/E/H-F-P-R-T

SEQ ID NO.:96: Exemplary embodiment of CDRH1 1 G-Y-T/I-F-T-D/E-Y-E/N-M/I/V-H

SEQ ID NO.:97: Exemplary embodiment of CDRH1 G-F-T/S-I-T-S-G-Y-G-W-H

SEQ ID NO.:98: Exemplary embodiment of CDRH2 V/N/G-I/L-D-P-E/A/G-T/Y-G-X-T-A

SEQ ID NO.:99: Exemplary embodiment of CDRH2 Y-I-N/S-F/Y-N/D-G

SEQ ID NO.:100: Exemplary embodiment of CDRH3 M-G-Y-S/A-D-Y

SEQ ID NO.:101: Exemplary embodiment of CDRH3 A-S-S-Y-D-G-F-L-A-Y

SEQ ID NO.:102: Exemplary embodiment of CDRH3 3 A-R/W-W/F-G-L-R-Q/N

SEQ ID NO.103- 3A2 light chain variable region

DAVMTQIPLTLSVTIGQPASLSCKSSQSLLHSDGKTYLNWLLQRPGQSPKRLISLVS**KLD**SGV**P**D  
RFTGSGSGTDFTLKISRVEAEDLGLYYC**WQG**THFPRTFAGGTNLEIK

SEQ ID NO.104 3F6 light chain variable region

SIVMTQTPLTLSVTIGQPASITCKSSQSLLYSDGKTYLNWLLQRPGQSPKRLISLVS**KLD**SGV**P**D  
GFTGSGSGTDFTLKISRVEAEDLGVYYC**WQG**THFPRTFGGGTKLEIK

SEQ ID NO.105- 3E8 light chain variable region

DAVMTQIPLTLSVTIGQPASISCKSSQSLLHSDGKTYLNWLLQRPGQSPKRLIY**LVS****KLD**SGV**P**D  
RFTGSGSGTDFTLKISRVEAEDLGVYYC**WQG**THFPRTFGGGTKLEIK

SEQ ID NO.106- 3E10 light chain variable region

DIVMTQAAPSVPVTPGESVSISCRSSKSLLHSGNGNTYLYWFLQRPGQSPQLIY**RMSNL**ASG**V**P  
RFSGSGSGTAFTRLRISRV~~EAED~~VGVYYC**MQH**LEY**P**YTFGGGT**K**LEIK

SEQ ID NO.107- 3A9 light chain variable region

DIVMTQSPSSLAMSLGQKV**T**MSCKSSQSLLNSNNQ**LN**YLA**W**YQQKPGQSPKLLVY**FASTR**KG**V**P  
DRFIGSGSGTDFTLTITSVQAEDLADYFC**QQH**FNTPLTFGAGTKLEIK

SEQ ID NO.108- 3B1 light chain variable region

DIVMTQSPSSLAI**S**VGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLV**FFASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.109- 3G5 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLV**FFASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.110- 3B2 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.111- 3B8 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.112- 3G8 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.113- 3F7 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLIY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.114- 3E9 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTEFTLTITSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.115- 3C3 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FGSTR**ESG**V**P  
DRFIGSGSGTDFTLTISGVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.116- 3E12 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**M**N**CKSSQSLLNRSNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISSVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

SEQ ID NO.117- 4A2 light chain variable region

DIVMTQSPSSLAMSVGQKV**T**MSCKSSQSLLNSNNQ**KN**YLA**W**YQQKPGQSPKLLVY**FASTR**ESG**V**P  
DRFIGSGSGTDFTLTISGVQAEDLADYFC**QQH**YSIPLTFGAGTKLEIK

DIVMTQSPSSLTMSVGQKVTM**SCSSQSLLNTSNQ**NYLAWYQQKPGQSPKLLVY**FASTTES**GVP  
 DRFIGSGSGTYFTLT**ISSVQAEDLADYFCQQHYSTPLTFGAGTKLDLK**

SEQ ID N0.118-3F10 light chain variable region

DIVMTQSPSSLTMSVGQKVTM**SCSSQSLLNTSNQ**NYLAWYQQKPGQSPKLLVY**FASTRAS**GVP  
 DRFIGSGSGTDF**TLTISSVQAEDLADYFCQQHYSTPLTFGAGTKLELK**

SEQ ID NO.119-3F4 light chain variable region

DIVMTQSPSSLT**VTAGEKVTM**SCSSQSLLNTSNQ**NYLAWYQQKPGQSPKLLVYFASTRES**GVP  
 DRFIGSGSGTDF**TLTISSVQAEDLADYFCQQHYSTPLTFGAGTKLELK**

SEQ ID NO.120-3B11 light chain variable region

DIVMTQSPSSLAMSVGQKVTM**SCSSQSLLN**SSNQ**NYLAWYQQKPGQSPKLLVYFASTRES**GVP  
 DRFIGSGSGTDF**TLTISSVQAEDLADYFCQQHYSTPLTFGAGTKLELK**

SEQ ID NO.121-3G12 light chain variable region

DIVMTQSPKFM**STSVGDRVSITCKASQD**IVGTAVAWYQQKPGQSP**ELLIYWTSTRHTGV**PD**RFSGS**  
 GSGTDF**TLTISSVQAEDLADYFCQQHYSIPLTFGAGTKLELR**

SEQ ID NO.122-3D1 light chain variable region

DIKMTQSPSSMYASLGERV**TITCKASQD**IHTYLNWFQQKPGKSP**ETLIYRANRLV**DGVPSRFSGS  
 GSGQD**YSLT**ISSLEYED**MG**IYY**CLQYDEFPLTFGAGTKLELK**

SEQ ID NO.123-3C2 light chain variable region

DIQMTQSPSSMYASLGERV**TLTCKASQD**IHN**YLNWFQQKPGKSP**KT**L**IHRANRLVAGVPSRFSGS  
 GSGQD**YSLT**ISSLEYED**LG**IYY**CLQYDAFPLTFGAGTKLELK**

SEQ ID NO.124-3E6 light chain variable region

DIQMTQSPSSMYASLGERV**TLTCKASQD**IHN**YLNWFQQKPGKSP**KT**L**IHRANRLVAGVPSRFSGS  
 GSGQD**YSLT**ISSLEYED**LG**IYY**CLQYDAFPLTFGAGTKLELK**

SEQ ID NO.125-3H3 light chain variable region

DIVMSQSPSSMYASLGERV**TITCKASQD**IHRFLNWFQQKPGKSP**KT**LIFHANRLV**DGVPSRFSGS**  
 GSGLD**YSLT**ISSLEYED**MG**IY**FCLQYDAFPLTFGAGTKLELK**

SEQ ID NO.126- 3A2 heavy chain variable region

HEIQLQQSGPELV**KPGASV**KMSCK**ASGYIF**TEYNIH**WVKQ**KPGQGLEWIG**YINPYNDV**TEYNEKF  
 KGRATL**TSDK**SSSTAYMDLSSLT**SD**AVY**FC**AW**GLRQ**WGQGTL**TV**ST

SEQ ID NO.127- 3F6 heavy chain variable region

HEVQLQQSGPELV**KPGASV**KMSCK**ASGYIF**TEY**NMHWV**KQ**KPGQ**GLEWIG**WIGNINPYNN**TEYNEKF  
 KGKATL**TSDK**ASSTAYMDLSSLT**SD**AVY**YCAR**W**GLRN**WG**QGTL****TV**SA

SEQ ID NO.128- 3E8 heavy chain variable region

HEVQLQQSGPELV**KPGASV**KMSCK**ASGYIF**TEY**NMHWV**KQ**KPGQ**GLEWIG**WIGNINPYNN**TEYNEKF  
 KGKATL**TSDK**SSSTAY**Y**MDLSSLT**SD**AVY**YCAR**W**GLRN**WG**QGTL****TV**SA

SEQ ID NO.129- 3A9 heavy chain variable region

HQVQLQQPG**AELVR**PGASV**TLSCK**AS**GYIF**TDY**EIH**WV**Q**RPV**H**GLEWIG**VIDP**E**TG**D**TAY**NQKF  
 KGKATL**TADK**SSSTAY**MEL**SSLT**ED**AVY**YCIG**Y**ADW**Q**GTT****TV**SS

SEQ ID NO.130-3B1 heavy chain variable region

HQVQLQQPG**AELVR**PGASV**TLSCK**AS**GYIF**TDY**EIH**WV**Q**TPV**H**GLEWIG**VIDP**E**TGG**T**AY**NQKF  
 KGKATL**TTDK**SSSTAY**MEL**SL**TS**ED**SAVY****CMG**Y**SDY**WG**QGTT****TV**SS

SEQ ID NO.131-3B2 heavy chain variable region

HEVQLQQSG**AELVR**PGASV**TLSCK**AS**GYIF**TDY**EIH**WV**Q**TPV**H**GLEWIG**VIDP**E**TG**A**TAY**NQKF  
 KGKATL**TADK**SSSTAY**MEL**SSLT**ED**AVY**YCIG**Y**ADW**Q**GTT****TV**SS

SEQ ID NO.132- 3F4 heavy chain variable region

HEVQLQQSGAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGSTAYNQKF  
KGKATLTADKASSTAYMELSSLTSEDSAVYYCMGYSDYWGQGTTLTVSS

SEQ ID NO.133- 3E9 heavy chain variable region

HEVQLQQSGAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGSTAYNQKF  
KGKATLTADKSSSTAYMELSSLTSEDSAVYYCMGYADYWGQGTTLTVSS

SEQ ID NO.134- 3B8 heavy chain variable region

HEVQLQQSGAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGDTAYNQNF  
TGKATLTADKSSSTAYMELSSLTSEDSAVYYCMGYADYWGQGTTLTVSS

SEQ ID NO.135- 3G8 heavy chain variable region

HQVQLKQSGAELVRPGASVTLSCASGYTFTDYEVHWVKQTPVHGLEWIGVIDPATGDTAYNQKF  
KGKATLTADKSSSTAYMEVSSLTSEDSAVYYCMGYSDYWGQGTTLTVSS

SEQ ID NO.136- 3F7 heavy chain variable region

HQAYLQQSGAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGDTAYNQKF  
KDKATLTADKASSTAYMELSSLTSEDSAVYYCMGYSDYWGQGTTLTVSS

SEQ ID NO.137- 3E12 heavy chain variable region

HQVQLQQSEAEVLVPGASVKSCKASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGDTAYNQKF  
KGKATLTADKSSSTAYMELSLTSEDSAVYYCMGHSDYWGQGTTLTVSS

SEQ ID NO.138- 3G12 heavy chain variable region

HEVQLQQSVAELVRPGASVTSCKASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGNTAFNQKF  
KGKATLTADISSSTAYMELSSLTSEDSAVYYCMGYSDYWGQGTTLTVSS

SEQ ID NO.139- 3F10 heavy chain variable region

HEVQLQQSVAELVRPGAPVTLSCASGYTFTDYEVHWVKQTPVHGLEWIGVIDPETGATAYNQKF  
KGKATLTADKSSSAAYMELSLTSEDSAVYYCMSYSYSDYWGQGTTLTVSS

SEQ ID NO.140- 3C3 heavy chain variable region

HEVQLQQSVAEVVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPETGVTAYNQRF  
RDKATLTTDKSSSTAYMELSSLTSEDSAVYFCMGYSDYWGQGTTLTVSS

SEQ ID NO.141- 3G5 heavy chain variable region

HQVQLQQPGAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVLDPGTGRAYNQKF  
KDKATLSADKSSSTAYMELSSLTSEDSAVYYCMSYSYSDYWGPGTTLTVSS

SEQ ID NO.142- 3B11 heavy chain variable region

HEVQLQQSVAELVRPGASVTLSCASGYTFTDYEIHVKQTPVHGLEWIGVIDPATGDTAYNQKF  
KGKATLTADKSSSAFMELSSLTSEDSAVYYCMGYSDYWGQGTTLTVSS

SEQ ID NO.143- 3E6 heavy chain variable region

HQVQLQQSGAELVRPGASVTLSCASGYTFSDYEMHWVKQTPVHGLEWIGGIDPETGDTVYNQKF  
KGKATLTADKSSSTAYMELSSLTSEDSAVYYCISYAMDYWGQGTSVTSS

SEQ ID NO.144- 4A2 heavy chain variable region

HQVQLQQSGTELVRPGASVTLSCASGYKFTDYEIHVKQTPVHGLEWIGGIDPETGGTAYNQKF  
KGKAILTADKSSTTAYMELRSLTSEDSAVYYCISYAMDYWGQGTSVTSS

SEQ ID NO.145- 3E10 heavy chain variable region

HEVQLQQSGPELVKPGASVKSCKASGDTFTDYYMNWVKQSHGKSLEWIGDINPNYGGITYNQKF  
KGKATLTVDTSSTAYMELRGLTSEDSAVYYCQAYYRNSDYWGQGTTLTVSS

SEQ ID NO.146- 3D1 heavy chain variable region

HEVQLQESGPDLVKPSQSLSLTCTVTGFSITSGYGWHWIHQFPGDKLEWMGYISFNGDYNYNPSL  
KSRISITRDTSKNQFFLQLSSVTTEDTATYYCASSYDGLFAYWGQGTLTVSA

SEQ ID NO.147- 3C2 heavy chain variable region

HDVQLQESGPDLVKPSQSLSLTCTVTGFSITSGYGWHWIHQFPGDKLEWMGYISFNGDYNYNPSL  
KSRISITRDTSKNQFFLQLNSVTSEDATYYCASSYDGLFAYWGQGPLTVSA

A

SEQ ID NO.:148

KSSQSLLHSDGKTYLN

SEQ ID NO.:149

LVSKLDS

SEQ ID NO.:150

WQGTHFPRT

SEQ ID NO.:151

GYTFTD YNMH

SEQ ID NO.:152

YINPYNDVTE

SEQ ID NO.:153

AWFGL RQ

SEQ ID NO.:154

RSSKSLLHSNGN TYLY

SEQ ID NO.:155

RMSNLAS

SEQ ID NO.:156

MQHLEYPYT

SEQ ID NO.:157

GDTFTD YYMN

SEQ ID NO.:158

DINPNYGGIT

SEQ ID NO.:159

QAYYRNS DY

SEQ ID NO.:160

KASQDVGTAVA

SEQ ID NO.:161

WTSTRHT

SEQ ID NO.:162

QQHYSIPLT

SEQ ID NO.:163

GYIFTDYEIH

SEQ ID NO.:164

VIDPETGNTA

SEQ ID NO.:165

MGYSDY

SEQ ID NO.:166

MVLQTQVFISLLLWISGAYGDIVMTQSPDSLAVSLGERATINCKSSQSLLNSNFQKNFLAWYQQK  
PGQPPKLLIYFASTRESSVPDRFSGSGSGTDFTLTISSLQAEDEVAVYYCQQHYSTPLTFQGQTKL  
EIKRTVAAPSVFIFPPSDEQLKSGTASVVCLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK  
DSTYSLSSTLTSKADYEKHKVYACEVTHQGLSSPVTKSFRGEC

SEQ ID NO.:167

MDWTWRILFLVAAATGTHAEVQLVQSGAEVKKPGASVKVSCKASGYIFTDYEIHWRQAPGQGLE  
WMGVIDPETGNTAFNQKFKGRVTITADTSTSTAYMELSSLTSEDTAVYYCMGYSYWGQGTLVTV  
SSASTKGPSVFPLAPSKSTSGGTAALGCLVKDYFPEPVTWSWNSGALTSGVHTFPAVLQSSGLY  
SLSSVVTVPSSSLGTQTYICNVNHPNTKVDKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP  
KPKDTLMISRTPEVTCVVVDVSHEDPEVFKFNWYVDGVEVHNAKTPREEQYNSTYRVVSVLTVLH  
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPS  
DIAVEWESNGQPENNYKTPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKS  
LSLSPGK

SEQ ID N:168

DIVMTQSPDSLAVSLGERATINCKSSQSLLNSNFQKNFLAWYQQKPGQPPKLLIYFASTRESSVP  
DRFSGSGSGTDFTLTISSLQAEDEVAVYYCQQHYSTPLTFQGQTKLEIK

SEQ ID NO.:169

EVQLVQSGAEVKKPGASVKVSCKASGYIFTDYEIHWRQAPGQGLEWMGVIDPETGNTAFNQKFK  
GRVTITADTSTSTAYMELSSLTSEDTAVYYCMGYSYWGQGTLVTVSS

SEQ ID NO.:170

MVLQTQVFISLLLWISGAYGDIVMTQSPSSLSAVGDRVTITCKASQDIHNFLNWFQQKPGKAPK  
TLIFRANRLVGDGVPSRFSGSGSGTDFTLTISSLQPEDFATYSCLYDEIPLTFQGQTKLEIKRTV  
AAPSVFIFPPSDEQLKSGTASVVCLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSL  
SSTLTSKADYEKHKVYACEVTHQGLSSPVTKSFRGEC

SEQ ID NO.:171

MDWTWRILFLVAAATGTHAEVQLQESGPGLVKPSQTLSLTCTVSGFSITSGYGHWRQHPGKGL  
EWIGYINYDGHNDYNPSLKSRTISQDTSKNQFSLKLSVTAAADTAVYYCASSYDGLFAYWGQGT  
LTVVSSASTKGPSVFPLAPSKSTSGGTAALGCLVKDYFPEPVTWSWNSGALTSGVHTFPAVLQ  
SGLYSLSSVVTVPSSSLGTQTYICNVNHPNTKVDKVEPKSCDKTHTCPPCPAPELLGGPSV  
LFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVFKFNWYVDGVEVHNAKTPREEQYNSTYRVV  
TVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRDELTKNQVSLTCLVK  
GFYPSDIAVEWESNGQPENNYKTPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHN  
TQKSLSLSPGK

SEQ ID NO.:172

DIVMTQSPSLSAVGDRVTITCKASQDIHNFLNWFQQKPGKAPKTLIFRANRLVDGVPSRSGS  
GSGTDYTLTISSLQPEDFATYSCLQYDEIPLTFGQGTKLEIK

SEQ ID NO.:173

EVQLQESGPGLVKPSQTLSTCTVSGFSITSGYGYWHIRQHPGKGLEWIGYINYDGHNDYNPSLK  
SRVTISQDTSKNQFSLKLSSVTAADTAVYYCASSYDGLFAYWGQGTLVTVS

SEQ ID NO.:186 (3A4 variant light chain variable region consensus 1)

DXVMTQTPLSLXVXXGXXASISCRSSQSLLHSNGNTYLEWYLQKPGQSPXLLIHTVSNRFSGVP

DRFSGSGSGTDFTLKISRVEAEDXGVYYCFQGSHVPLTFGXGTLEXK

wherein at least one of the amino acids identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48. The amino acid substitution may be, for example conservative.

SEQ ID NO.:187 (3A4 variant light chain variable region consensus 2)

DX<sub>a1</sub>VMTQTPLSLX<sub>a2</sub>VX<sub>a3</sub>X<sub>a4</sub>GX<sub>a5</sub>X<sub>a6</sub>ASISCRSSQSLLHSNGNTYLEWYLQKPGQSPX<sub>a7</sub>LLIHTV<sub>a8</sub>

NRFSGVPDRFSGSGSGTDFTLKISRVEAEDX<sub>a8</sub>GVYYCFQGSHVPLTFGX<sub>a9</sub>GTX<sub>a10</sub>LEX<sub>a11</sub>K

Wherein X<sub>a1</sub> may be a hydrophobic amino acid;

Wherein X<sub>a2</sub> may be A or P;

Wherein X<sub>a3</sub> may be neutral hydrophilic amino acid;

Wherein X<sub>a4</sub> may be L or P;

Wherein X<sub>a5</sub> may be an acidic amino acid;

Wherein X<sub>a6</sub> may be Q or P;

Wherein X<sub>a7</sub> may be a basic amino acid;

Wherein X<sub>a8</sub> may be a hydrophobic amino acid;

Wherein X<sub>a9</sub> may be A or Q;

Wherein X<sub>a10</sub> may be a basic amino acid; or

Wherein X<sub>a11</sub> may be a hydrophobic amino acid,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48.

SEQ ID NO.:188 (3A4 variant light chain variable region consensus 3)

DX<sub>A1</sub>VMTQTPLSLX<sub>A2</sub>VX<sub>A3</sub>X<sub>A4</sub>GX<sub>A5</sub>X<sub>A6</sub>ASISCRSSQSLLHSNGNTYLEWYLQKPGQSPX<sub>A7</sub>LLIHTV

SNRFSGVPDRFSGSGSGTDFTLKISRVEAEDX<sub>A8</sub>GVYYCFQGSHVPLTFGX<sub>A9</sub>GTX<sub>A10</sub>LEX<sub>A11</sub>K

Wherein X<sub>A1</sub> may be V or I

Wherein X<sub>A2</sub> may be A or P

Wherein X<sub>A3</sub> may be S or T

Wherein X<sub>A4</sub> may be L or P

Wherein X<sub>A5</sub> may be D or E

Wherein X<sub>A6</sub> may be Q or P

Wherein X<sub>A7</sub> may be K or Q

Wherein X<sub>A8</sub> may be L or V

Wherein X<sub>A9</sub> may be A or Q

Wherein X<sub>A10</sub> may be R or K or

Wherein X<sub>A11</sub> may be L or I,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:48.

SEQ ID NO.:189 (3A4 variant 1 light chain variable region: Lvh1)

DIVMTQTPLSLPVTPGEPASISCRSSQSLLHSNGNTYLEWYLQKPGQSPQLIYTVSNRFSGVPD

RFSGSGSGTDFTLKISRVEAEDVGVYYCFQGSHVPLTFGQGTKLEIK

SEQ ID NO.:190 (3A4 variant 2 light chain variable region: Lvh2)

DVVMQTPLSLPVTPGEPASISCRSSQSLLHSNGNTYLEWYLQKPGQSPKLLIYTVSNRFSGVPD

RFSGSGSGTDFTLKISRVEAEDVGVYYCFQGSHVPLTFGQGTKLEIK

SEQ ID NO.:191 (3A4 variant heavy chain variable region consensus 1)

QXQLVQSGXEXXKPGASVKXSCKASGYTFTDDYMSWVXQXXGXLEWXGDINPYNGDTNYNQ

KFKGXXXXTDXSXSTAYMLXSLXSEDXAVYYCARDPGAMDYWGQGTXTVTVSS

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:46. The amino acid substitution may be, for example conservative.

SEQ ID NO.:192 (3A4 variant heavy chain variable region consensus 2)

QX<sub>b1</sub>QLVQSGX<sub>b2</sub>EX<sub>b3</sub>X<sub>b4</sub>KPGASVKX<sub>b5</sub>SCKASGYTFTDDYMSWVX<sub>b6</sub>QX<sub>b7</sub>X<sub>b8</sub>GX<sub>b9</sub>X<sub>b10</sub>LEWX<sub>b11</sub>G

DINPYNGDTNYNQKFKGX<sub>b12</sub>X<sub>b13</sub>X<sub>b14</sub>X<sub>b15</sub>TX<sub>b16</sub>DX<sub>b17</sub>SX<sub>b18</sub>STAYMX<sub>b19</sub>LX<sub>b20</sub>SLX<sub>b21</sub>SEDX<sub>b22</sub>AVYY

CARDPGAMDYWGQGTGX<sub>b23</sub>VTVSS

Wherein  $X_{b1}$  may be a hydrophobic amino acid;

Wherein  $X_{b2}$  may be P or A;

Wherein  $X_{b3}$  may be a hydrophobic amino acid;

Wherein  $X_{b4}$  may be V or K;

Wherein  $X_{b5}$  may be a hydrophobic amino acid;

Wherein  $X_{b6}$  may be a basic amino acid;

Wherein  $X_{b7}$  may be S or A;

Wherein  $X_{b8}$  may be H or P;

Wherein  $X_{b9}$  may be a basic amino acid;

Wherein  $X_{b10}$  may be S or G;

Wherein  $X_{b11}$  may be a hydrophobic amino acid;

Wherein  $X_{b12}$  may be a basic amino acid;

Wherein  $X_{b13}$  may be a hydrophobic amino acid;

Wherein  $X_{b14}$  may be I or T;

Wherein  $X_{b15}$  may be a hydrophobic amino acid;

Wherein  $X_{b16}$  may be a hydrophobic amino acid;

Wherein  $X_{b17}$  may be K or T;

Wherein  $X_{b18}$  may be a neutral hydrophilic amino acid;

Wherein  $X_{b19}$  may be Q or E;

Wherein  $X_{b20}$  may be N or S;

Wherein  $X_{b21}$  may be T or R;

Wherein  $X_{b22}$  may be a neutral hydrophilic amino acid; or

Wherein  $X_{b23}$  may be S or L,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the

polypeptide set forth in SEQ ID NO.:46.

SEQ ID NO.:193 (3A4 variant heavy chain variable region consensus 3)

QX<sub>B1</sub>QLVQSGX<sub>B2</sub>EX<sub>B3</sub>X<sub>B4</sub>KPGASVKX<sub>B5</sub>SCKASGYTFTDDYMSWVX<sub>B6</sub>QX<sub>B7</sub>X<sub>B8</sub>GX<sub>B9</sub>X<sub>B10</sub>LEWX<sub>B11</sub>  
GDINPYNGDTNYNQKFKGX<sub>B12</sub>X<sub>B13</sub>X<sub>B14</sub>X<sub>B15</sub>TX<sub>B16</sub>DX<sub>B17</sub>SX<sub>B18</sub>STAYMX<sub>B19</sub>LX<sub>B20</sub>SLX<sub>B21</sub>SEDX<sub>B22</sub>AV  
YYCARDPGAMDYWGQQGTX<sub>B23</sub>VTVSS

Wherein X<sub>B1</sub> may be I or V;

Wherein X<sub>B2</sub> may be P or A;

Wherein X<sub>B3</sub> may be M or V;

Wherein X<sub>B4</sub> may be V or K;

Wherein X<sub>B5</sub> may be M or V;

Wherein X<sub>B6</sub> may be K or R;

Wherein X<sub>B7</sub> may be S or A;

Wherein X<sub>B8</sub> may be H or P;

Wherein X<sub>B9</sub> may be K or Q;

Wherein X<sub>B10</sub> may be S or G;

Wherein X<sub>B11</sub> may be I or M;

Wherein X<sub>B12</sub> may be K or R;

Wherein X<sub>B13</sub> may be A or V;

Wherein X<sub>B14</sub> may be I or T;

Wherein X<sub>B15</sub> may be L or I;

Wherein X<sub>B16</sub> may be V or A;

Wherein X<sub>B17</sub> may be K or T;

Wherein X<sub>B18</sub> may be S or T;

Wherein X<sub>B19</sub> may be Q or E;

Wherein X<sub>B20</sub> may be N or S;

Wherein X<sub>B21</sub> may be T or R;

Wherein X<sub>B22</sub> may be S or T; or

Wherein X<sub>B23</sub> may be S or L,

wherein at least one of the amino acid identified by X is an amino acid substitution (conservative or non-conservative) in comparison with a corresponding amino acid in the polypeptide set forth in SEQ ID NO.:46.

SEQ ID NO.:194 (3A4 variant 1 heavy chain variable region: Hvh1)

QVQLVQSGAEVKPGASVKVSCKASGYTFTDDYMSWVRQAPGQGLEWMGDINPYNGDTNYNQ  
QKFKGRVTITADTSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSS

SEQ ID NO.:195 (3A4 variant 2 heavy chain variable region: Hvh2)

QIQLVQSGAEVKPGASVKVSCKASGYTFTDDYMSWVRQAPGQGLEWIGDINPYNGDTNYNQ  
KFKGRVTITADKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSS

SEQ ID NO.:196 (3A4 variant 3 heavy chain variable region: Hvh3)

QIQLVQSGAEVKPGASVKVSCKASGYTFTDDYMSWVKQAPGQGLEWIGDINPYNGDTNYNQK  
FKGRATLTVDKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSS

SEQ ID NO.:197 (3A4 variant 4 heavy chain variable region: Hvh4)

QIQLVQSGAEVKPGASVKVSCKASGYTFTDDYMSWVKQAPGQGLEWIGDINPYNGDTNYNQK  
FKGKATLTVDKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSS

SEQ ID NO: 198 3A4 murine light (kappa) chain

DVVTMTQTPLSLAVSLGDQASISCRSSQSLLHSNGNTYLEWYLQKPGQSPKLLIHTVSNRFSGVP  
DRFSGSGSGTDFTLKISRVEAEDLGVYYCFQGSHVPLTFGAGTRLELKRTVAAPSVFIFPPSDEQ  
LKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTTLSKADYEK  
HKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO:199 3A4 humanized light (kappa) chain variant 1; Lh1

DIVMTQTPLSLPVTPGEPAISCRSSQSLLHSNGNTYLEWYLQKPGQSPKLLIYTVSNRFSGVPD  
RFSGSGSGTDFTLKISRVEAEDGVVYYCFQGSHVPLTFGQGTKLEIKRTVAAPSVFIFPPSDEQL  
KSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTTLSKADYEK  
HKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO:200 3A4 humanized light (kappa) chain variant 2; Lh2

DVVTMTQTPLSLPVTPGEPAISCRSSQSLLHSNGNTYLEWYLQKPGQSPKLLIYTVSNRFSGVPD  
RFSGSGSGTDFTLKISRVEAEDGVVYYCFQGSHVPLTFGQGTKLEIKRTVAAPSVFIFPPSDEQL  
KSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTTLSKADYEK  
HKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO:201 3A4 murine heavy (Igg1) chain

QIQLVQSGPEMVKPGASVKMSCKASGYTFTDDYMSWVKQSHGKSLEWIGDINPYNGDTNYNQ  
KFKGKAILTVDKSSSTAYMQLNSLTSEDSAVYYCARDPGAMDYWGQGTSVTSSASTKGPSVF  
PLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPALQSSGLYSLSSVTVPS  
SSLGTQTYICNVNHPNSNTKVDKKVEPKSCDKHTCPCPAPELLGGPSVFLFPPPKDTLMISR  
TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKE  
YKCKVSNKALPAPIEKTIASKAKGQPREPVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWES  
NGQPENNYKTPPVLDSDGSFFLYSKLTVDKSRWQQGVFCSCVMHEALHNHYTQKSLSLSPG  
K

SEQ ID NO:202 3A4 humanized heavy (Igg1) chain variant 1; Hh1  
 QVQLVQSGAEVKKPGASVKVSKCASGYTFTDDYMSWRQAPGQGLEWMGDINPYNGDTNYN  
 QKFKGRVTITADTSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSSASTKGPSVF  
 PLAPSSKSTSGGTAALGCLVKDYFPEPVTVWSNSGALTSGVHTFPAVLQSSGLYSLSSVTVPS  
 SSLGTQTYICNVNHPKNSNTKVDKKVEPKSCDKTHCPCPAPELLGGPSVFLFPPKPKDTLMISR  
 TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTPREEQYNSTYRVVSVLTVLHQDWLNGKE  
 YKCKVSNKALPAPIEKTIKAKGQPREPQVTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWES  
 NGQPENNYKTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG  
 K

SEQ ID NO:203 3A4 humanized heavy (Igg1) chain variant 2; Hh2  
 QIQLVQSGAEVKKPGASVKVSKCASGYTFTDDYMSWRQAPGQGLEWMGDINPYNGDTNYNQ  
 KFKGRVTITADKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSSASTKGPSVFP  
 LAPSSKSTSGGTAALGCLVKDYFPEPVTVWSNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSS  
 SLGTQTYICNVNHPKNSNTKVDKKVEPKSCDKTHCPCPAPELLGGPSVFLFPPKPKDTLMISR  
 PEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTPREEQYNSTYRVVSVLTVLHQDWLNGKEY  
 KCKVSNKALPAPIEKTIKAKGQPREPQVTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESN  
 GQPENNYKTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

SEQ ID NO:204 3A4 humanized heavy (Igg1) chain variant 3; Hh3  
 QIQLVQSGAEVKKPGASVKVSKCASGYTFTDDYMSWRQAPGQGLEWMGDINPYNGDTNYNQK  
 FKGRATLTVDKSTSTAYMELSSLRSEDTAVYYCARDPGAMDYWGQGTLTVSSASTKGPSVFP  
 LAPSSKSTSGGTAALGCLVKDYFPEPVTVWSNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSS  
 SLGTQTYICNVNHPKNSNTKVDKKVEPKSCDKTHCPCPAPELLGGPSVFLFPPKPKDTLMISR  
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SEQ IDNO.:215 - OGS1879

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EQ ID NO.:216 - OGS1810

GGGGCCAGGGAAAGACAGATGGGCCCTCGTTGAGGC

## REFERENCES

### [0443]

Santana-Davila R. and Perez E.A. (2010) "Treatment options for patients with triple-negative breast cancer" *J Hematol Oncol*. 27:42.

de Ruijter T.C., Veeck J., et al. (2011) "Characteristics of triple-negative breast cancer." *J Cancer Res Clin Oncol*. 137:183.

Ismail-Khan R. and Bui M.M. (2010) "A review of Triple-negative breast cancer" *Cancer Control* 17:173.

Carey L.A., Perou C.M. et al. (2006) "Race, breast cancer subtypes, and survival in the Carolina Breast Cancer Study." *JAMA* 295:2492.

Krieg M., Seynaeve C. et al. (2009) "Sensitivity to first-line chemotherapy for metastatic breast cancer in BRCA1 and BRCA2 mutation carriers." *J Clin Oncol* 27:3764.

Rouzier R., Perou C.M. et al. (2005) "Breast cancer molecular subtypes respond differently to preoperative chemotherapy" *Clin Cancer Res* 11:5678.

Fong P.C., Boss D.S. et al. (2009) "Inhibition of poly(ADP-ribose) polymerase in tumors from BRCA mutation carriers." *N Engl J Med* 361:123.

Dent R., Trudeau M et al. (2007) "Triple-Negative Breast Cancer: Clinical Feature and Patterns of Recurrence" *Clin. Cancer Res*. 13: 4429.

Bernstein L and J.V. Lacey Jr. (2011) "Receptors, Associations, and Risk Factor Differences by Breast Cancer Subtypes: Positive or Negative?" *J Natl Cancer Inst* 103(6): 451-453 (Advanced publication February 23, 2011).

Nofech-Mozes S. et al., (2009) "Patterns of recurrence in the basal and non-basal subtypes of triple-negative breast cancers" *Cancer Res. Treat*. 118: 131-137.

## REFERENCES CITED IN THE DESCRIPTION

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#### Patent documents cited in the description

- CA2007001134W [0007] [0432]
- WO2007147265A [0007]
- CA2009001586W [0008] [0076] [0410] [0433]
- WO2010060186A [0008]
- CA2010001785W [0008]
- WO2011054112A [0008]
- US20110223107A [0008]
- US20030118592A [0060]
- US20030133939A [0060]
- WO2010060186A8A [0076]
- CA2010001795W [0076] [0410]
- WO2011054112A1 [0076]
- CA2012000296W [0076] [0409] [0410]
- WO2012129668A1 [0076]

#### Non-patent literature cited in the description

- **LARKIN M.A. et al.** ClustalW and ClustalX version 2 Bioinformatics, 2007, vol. 23, 212947-2948 [0010] [0010]
- **KALLURIWEINBERG** The Journal of Clinical Investigation, 2009, vol. 119, 61420-1428 [0027]
- **FASSINA et al.** Modern Pathology, 2012, vol. 25, 86-99 [0027]
- **LEE et al.** JCB, 2006, vol. 172, 973-981 [0027]
- **FOULKES et al.** N. Engl. J. Med., 2010, vol. 363, 1938-1948 [0042]
- **WARD et al.** Nature, 1989, vol. 341, 544-546 [0060]
- **BIRD et al.** Science, 1988, vol. 242, 423-426 [0060]
- **HUSTON et al.** Proc. Natl. Acad. Sci. USA, 1988, vol. 85, 5879-5883 [0060]
- **PORTOLANO et al.** The Journal of Immunology, 1993, vol. 150, 880-887 [0086]
- **CLARKSON et al.** Nature, 1991, vol. 352, 624-628 [0086]
- **TATIANA ATATUSOVATHOMAS LMADDEN** Blast 2 sequences - a new tool for

comparing protein and nucleotide sequencesFEMS Microbiol Lett, 1999, vol. 174, 247-250 [0111]

- **SANTANA-DAVILA R.PEREZ E.A.**Treatment options for patients with triple-negative breast cancerJ Hematol Oncol., 2010, vol. 27, 42- [0443]
- **DE RUIJTER T.C.VEECK J. et al.**Characteristics of triple-negative breast cancerJ Cancer Res Clin Oncol., 2011, vol. 137, 183- [0443]
- **ISMAIL-KHAN RBUI M.MA** review of Triple-negative breast cancerCancer Control, 2010, vol. 17, 173- [0443]
- **CAREY L.A.PEROU C.M. et al.**Race, breast cancer subtypes, and survival in the Carolina Breast Cancer StudyJAMA, 2006, vol. 295, 2492- [0443]
- **KRIEG M.SEYNAEVE C. et al.**Sensitivity to first-line chemotherapy for metastatic breast cancer in BRCA1 and BRCA2 mutation carriersJ Clin Oncol, 2009, vol. 27, 3764- [0443]
- **ROUZIER R.PEROU C.M. et al.**Breast cancer molecular subtypes respond differently to preoperative chemotherapyClin Cancer Res, 2005, vol. 11, 5678- [0443]
- **FONG P.C.BOSS D.S. et al.**Inhibition of poly(ADP-ribose) polymerase in tumors from BRCA mutation carriersN Engl J Med, 2009, vol. 361, 123- [0443]
- **DENT R.TRUDEAU M et al.**Triple-Negative Breast Cancer: Clinical Feature and Patterns of RecurrenceClin. Cancer Res., 2007, vol. 13, 4429- [0443]
- **BERNSTEIN LJ.V. LACEY JR.**Receptors, Associations, and Risk Factor Differences by Breast Cancer Subtypes: Positive or Negative?J Natl Cancer Inst, 2011, vol. 103, 6451-453 [0443]
- **NOFECH-MOZES S. et al.**Patterns of recurrence in the basal and non-basal subtypes of triple-negative breast cancersCancer Res. Treat., 2009, vol. 118, 131-137 [0443]

**Patentkrav**

1. Antistof eller antigenbindende fragment deraf, som kan binde specifikt til nyre-associeret antigen 1 (KAAG1) og er konjugeret med en terapeutisk enhed til anvendelse ved behandling af triple-negativ brystkræft, hvor antistoffet eller det antigenbindende fragment deraf omfatter en CDRH1 som anført i SEQ ID NO.:49, en CDRH2 som anført i SEQ ID NO.:50 eller i SEQ ID NO.:212, en CDRH3 som anført i SEQ ID NO.:51, en CDRL1 som anført i SEQ ID NO.: 52, en CDRL2 som anført i SEQ ID NO.:53 og en CDRL3 som anført i SEQ ID NO.: 54.
2. Antistof eller antigenbindende fragment deraf til anvendelse ifølge krav 1, hvor antistoffet eller det antigenbindende fragment deraf binder til overfladen af kræftceller.
3. Antistof eller antigenbindende fragment deraf til anvendelse ifølge krav 1 eller 2, hvor antistoffet eller det antigenbindende fragment deraf binder en epitop mellem aminosyrerne 30 til 84 af KAAG1.
4. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 3, hvor antistoffet eller det antigenbindende fragment deraf er et monoklonalt antistof, et kimært antistof, et human antistof eller et humaniseret antistof eller et antigenbindende fragment deraf.
5. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 4, hvor antistoffet eller det antigenbindende fragment deraf anvendes i kombination med et kemoterapeutisk eller et cytotoxisk middel.
6. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5, hvor antistoffet eller det antigenbindende fragment deraf omfatter en letkæde-variabel region som anført i SEQ ID NO.:48 og en tungkæde-variabel region som anført i SEQ ID NO.:46.
7. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5, hvor antistoffet eller det antigenbindende fragment

deraf omfatter en letkæde-variabel region som anført i SEQ ID NO.:186, hvor mindst en af aminosyrerne identificeret med X er en aminosyresubstitution sammenlignet med en tilsvarende aminosyre i polypeptidet anført i SEQ ID NO.:48 og en tungkæde-variabel region som anført i SEQ ID NO.:191, hvor mindst en af aminosyrerne identificeret med X er en aminosyresubstitution sammenlignet med en tilsvarende aminosyre i polypeptidet anført i SEQ ID NO.:46.

8. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5 eller 7, hvor den lette kædes variable region er som anført i SEQ ID NO.:187, og hvor den tunge kædes variable region er som anført i SEQ ID NO.:192.

9. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5, 7 eller 8, hvor den lette kædes variable region er som anført i SEQ ID NO.:188, og hvor den tunge kædes variable region er som anført i SEQ ID NO.:193.

10. Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5 eller 7 til 9, hvor antistoffet eller det antigenbindende fragment deraf omfatter:

a. en letkæde-variabel region som anført i SEQ ID NO.: 189 eller SEQ ID NO.:190 og en tungkæde-variabel region som anført i SEQ ID NO.:194, SEQ ID NO.:195, SEQ ID NO.:196 eller SEQ ID NO.:197;

25 b. en letkæde-variabel region som anført i SEQ ID NO.:189 og en tungkæde-variabel region som anført i SEQ ID NO.:194;

c. en letkæde-variabel region som anført i SEQ ID NO.:189 og en tungkæde-variabel region som anført i SEQ ID NO.:195;

d. en letkæde-variabel region som anført i SEQ ID NO.:189 og en tungkæde-variabel region som anført i SEQ ID NO.:196;

30 e. en letkæde-variabel region som anført i SEQ ID NO.:189 og en tungkæde-variabel region som anført i SEQ ID NO.:197;

f. en letkæde-variabel region som anført i SEQ ID NO.:190 og en tungkæde-variabel region som anført i SEQ ID NO.:194;

35 g. en letkæde-variabel region som anført i SEQ ID NO.:190 og en tungkæde-variabel region som anført i SEQ ID NO.:195; eller

h. en let kæde som anført i SEQ ID NO.: 199 eller SEQ ID NO.:200 og en tung kæde som anført i SEQ ID NO.:202, SEQ ID NO.:203, SEQ ID NO.:204 eller SEQ ID NO.:205.

5 **11.** Antistof eller antigenbindende fragment til anvendelse ifølge et hvilket som helst af kravene 1 til 5 eller 8 til 10, hvor antistoffet eller det antigenbindende fragment deraf omfatter;

a. en let kæde som anført i SEQ ID NO.:199 og en tung kæde som anført i SEQ ID NO.:202;

10 b. en let kæde som anført i SEQ ID NO.:199 og en tung kæde som anført i SEQ ID NO.:203;

c. en let kæde som anført i SEQ ID NO.:199 og en tung kæde som anført i SEQ ID NO.:204;

15 d. en let kæde som anført i SEQ ID NO.:199 og en tung kæde som anført i SEQ ID NO.:205;

e. en let kæde som anført i SEQ ID NO.:200 og en tung kæde som anført i SEQ ID NO.:202;

f. en let kæde som anført i SEQ ID NO.:200 og en tung kæde som anført i SEQ ID NO.:203;

20 g. en let kæde som anført i SEQ ID NO.:200 og en tung kæde som anført i SEQ ID NO.:204; eller

h. en let kæde som anført i SEQ ID NO.:200 og en tung kæde som anført i SEQ ID NO.:205.

25 **12.** Antistof eller antigenbindende fragment deraf til anvendelse ifølge krav 1, hvor den terapeutiske enhed er et cytotoxisisk middel.

**13.** Antistof eller antigenbindende fragment deraf til anvendelse ifølge krav 1, hvor den terapeutiske enhed er en antimitotisk forbindelse.

30 **14.** Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5 eller 8 til 10, hvor antistoffet eller det antigenbindende fragment deraf har en affinitet på 1 nM eller mindre eller 0,1 nM eller mindre for KAAG1.

- 15.** Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 5, 8 til 10, 12 til 14, hvor antistoffet eller det antigenbindende fragment deraf er internaliseret i en celle.
- 5      **16.** Antistof eller antigenbindende fragment deraf til anvendelse ifølge et hvilket som helst af kravene 1 til 15, hvor antistoffet eller det antigenbindende fragment deraf anvendes sammen med et antikræftmiddel.

# DRAWINGS

**Figure 1a**

3A4-VL

<b>murine</b>	DDVMTQTPSLAVSLCGDQASISCRSOSLILHSIGNTYLEWVLOQPGQSPNLLIHTVSNRFSQV2PDRFSGSGCTDFTLKISRVEADLGLVYCPQGSVPLAFGACTRLEIK	11/80 (86.3%)
<b>Humanized1</b>	DDVMTQTPSLAVSLCGDQASISCRSOSLILHSIGNTYLEWVLOQPGQSPNLLIHTVSNRFSQV2PDRFSGSGCTDFTLKISRVEADLGVYCPQGSVPLAFGACTRLEIK	0/80 (100%)
<b>Humanized2</b>	DDVMTQTPSLAVSLCGDQASISCRSOSLILHSIGNTYLEWVLOQPGQSPNLLIHTVSNRFSQV2PDRFSGSGCTDFTLKISRVEADLGVYCPQGSVPLAFGACTRLEIK	2/80 (97.5%)

CDR-L1

CDR-L2

CDR-L3

**Figure 1b**

3A4-VH

<b>mouse</b>	QIQLVQSGPENVKPGASIVKMSCKASGYTTDDWMSWYQSHGKSLWEIGDINPYGDTNQFKGKALITVQSSSTAYMQLNSLTSEDAVYCCARDPGMDYNGQGTSVTVSS	21/82 (74.4%)
<b>Humanized1</b>	QIQLVQSGAEVVKPGASIVKMSCKASGYTTDDWMSWYQAFQGQLEWGDINPYGDTNQFKGRTITADKSTSTAYMELSSRSEDAVYCCARDPGMDYNGQGTLVTVSS	0/82 (100%)
<b>Humanized2</b>	QIQLVQSGAEVVKPGASIVKMSCKASGYTTDDWMSWYQAFQGQLEWGDINPYGDTNQFKGRTITADKSTSTAYMELSSRSEDAVYCCARDPGMDYNGQGTLVTVSS	2/82 (91.5%)
<b>Humanized3</b>	QIQLVQSGAEVVKPGASIVKMSCKASGYTTDDWMSWYQAFQGQLEWGDINPYGDTNQFKGRTITADKSTSTAYMELSSRSEDAVYCCARDPGMDYNGQGTLVTVSS	6/82 (92.7%)
<b>Humanized4</b>	QIQLVQSGAEVVKPGASIVKMSCKASGYTTDDWMSWYQAFQGQLEWGDINPYGDTNQFKGRTITADKSTSTAYMELSSRSEDAVYCCARDPGMDYNGQGTLVTVSS	8/82 (90.2%)

CDR-H1

CDR-H2

CDR-H3

Figure 2a

## Variable light chain alignment

Mouse VL SEQ ID NO. 189	DVMTQTPPLSLAVSLIGDQASISCRSSQSLHNSGNTYLEWLYLQKPGQSPKLLIHTVSNRF DVMTQTPPLSLPVTPCGEPASISCRSSQSLHNSGNTYLEWLYLQKPGQSPQLLIHTVSNRF
Mouse VL SEQ ID NO. 189	SGVPDRFSGSGSGTDFELKISRVEADLGYYYCFQGSSHVPLTFAGTRLEIK SGVPDRFSGSGSGTDFELKISRVEADVGYYCFQGSSHVPLTFGQGTTKLEIK

Figure 2b

## Variable heavy chain alignment

Figure 3a

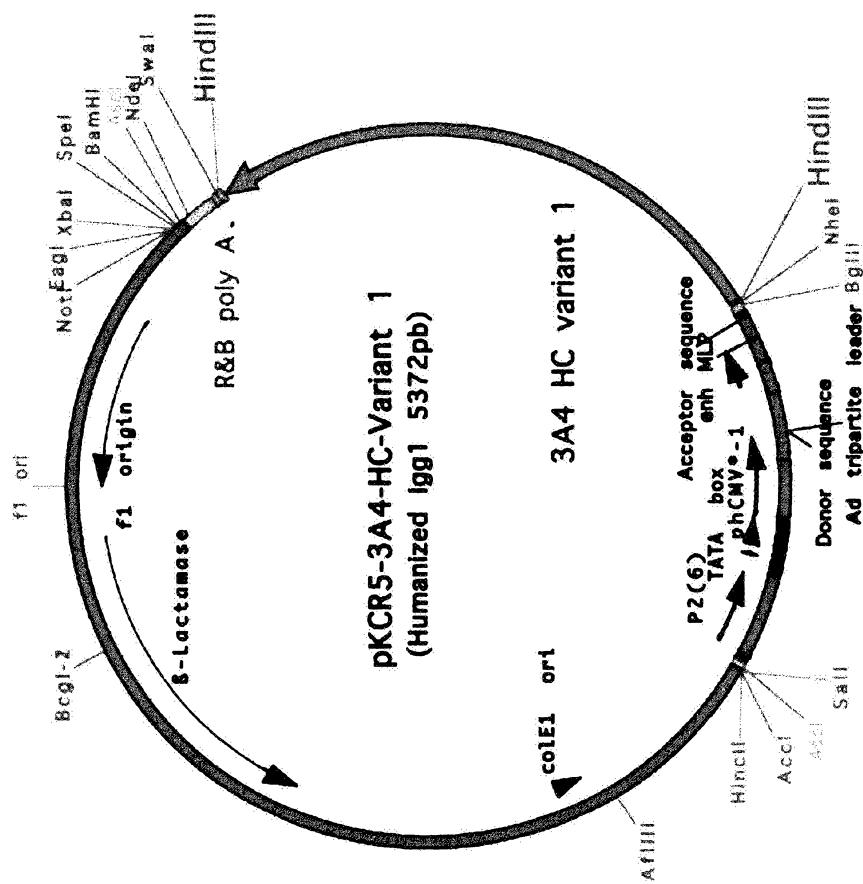
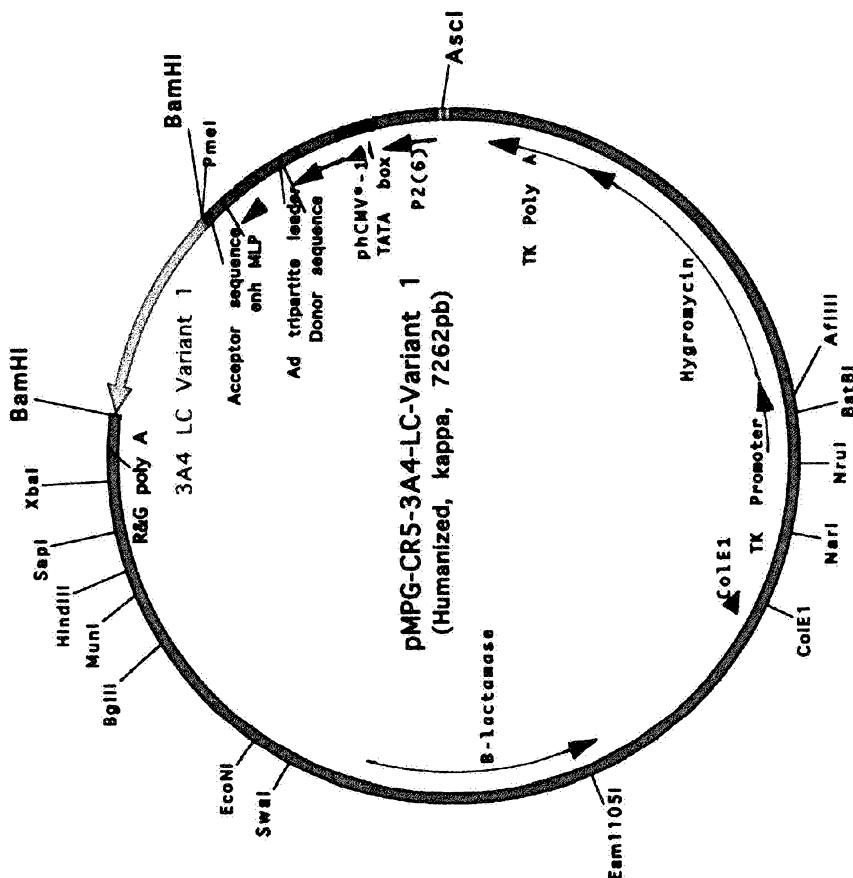


Figure 3b



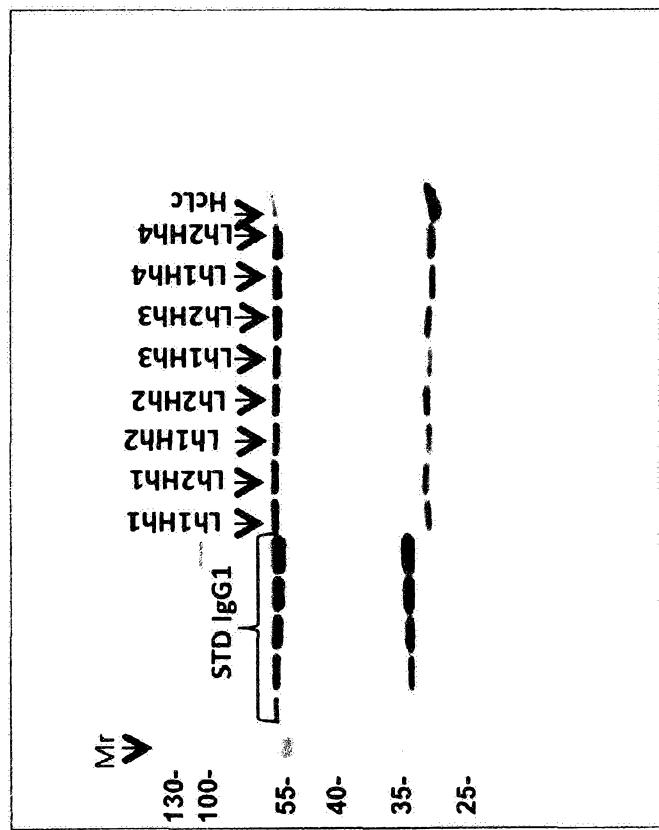


Figure 4

Figure 5

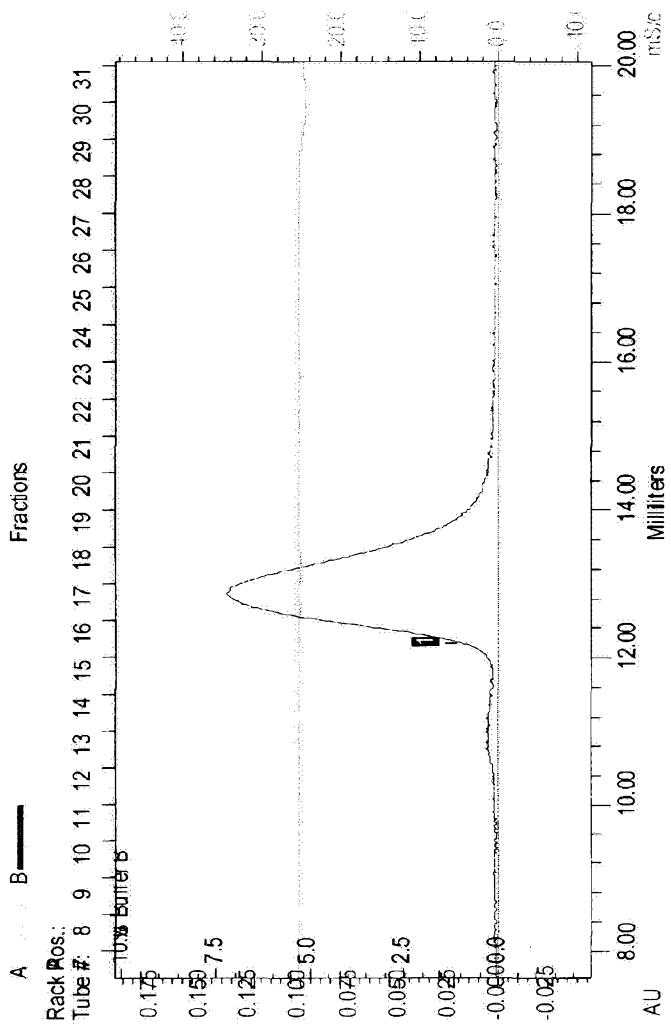


Figure 6

Antibody	$k_a$ (1/Ms)	$k_d$ (1/s)	$K_D$ (nM)	Fold diff.
LcHc	$7.72 \times 10^6$	$1.21 \times 10^{-4}$	0.016	-
Lh1Hh1	$6.93 \times 10^6$	$3.28 \times 10^{-3}$	0.474	29.6
Lh2Hh1	$6.97 \times 10^6$	$2.37 \times 10^{-3}$	0.341	21.3
Lh1Hh2	$5.65 \times 10^6$	$1.19 \times 10^{-3}$	0.211	13.2
Lh2Hh2	$7.40 \times 10^6$	$1.81 \times 10^{-3}$	0.245	15.3
Lh1Hh3	$6.46 \times 10^6$	$9.60 \times 10^{-4}$	0.149	9.3
Lh2Hh3	$4.46 \times 10^6$	$1.02 \times 10^{-3}$	0.228	14.3
Lh1Hh4	$5.14 \times 10^6$	$7.64 \times 10^{-4}$	0.149	9.3
Lh2Hh4	$4.57 \times 10^6$	$4.70 \times 10^{-4}$	0.103	6.4

Figure 7a

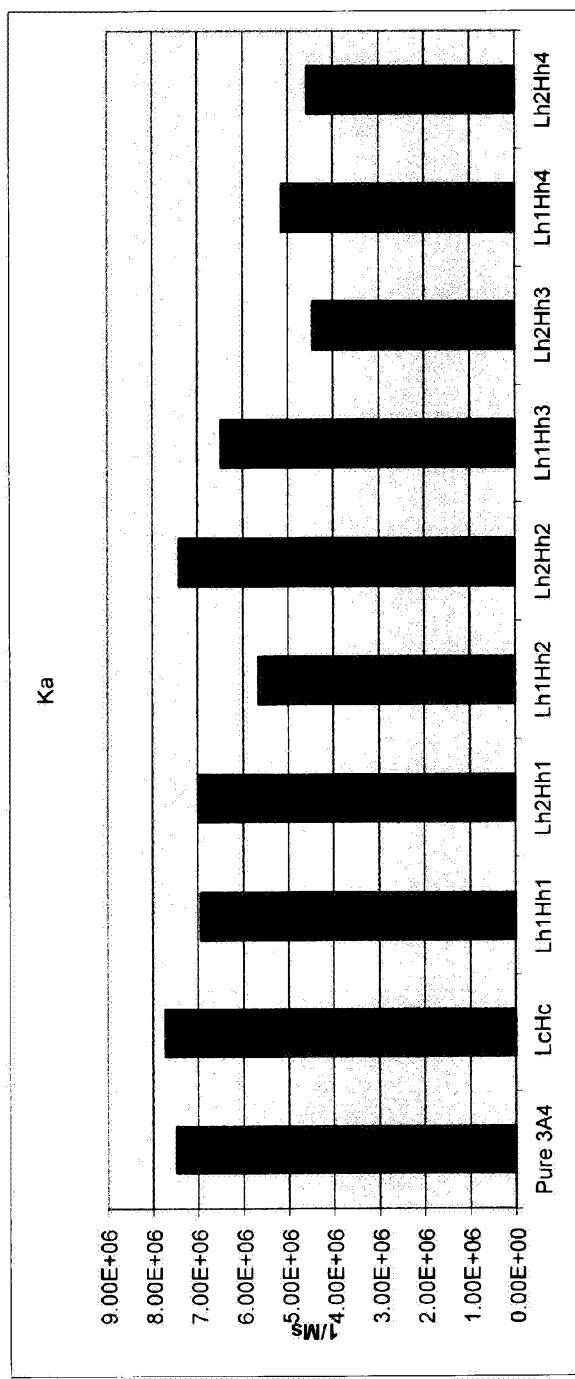


Figure 7b

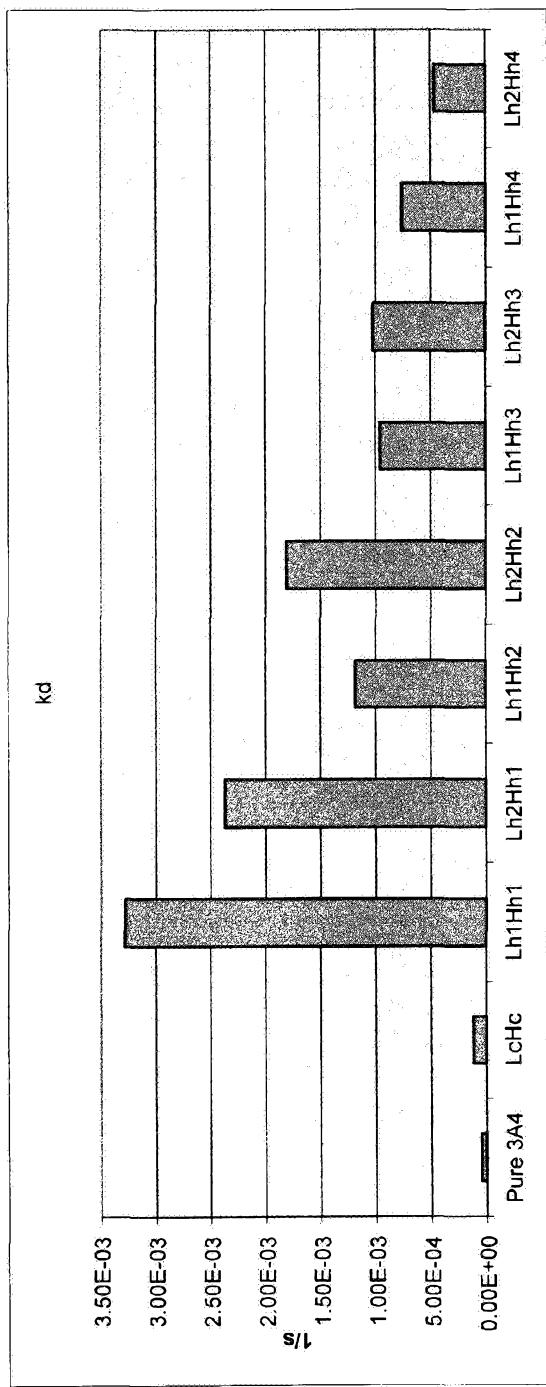


Figure 7c

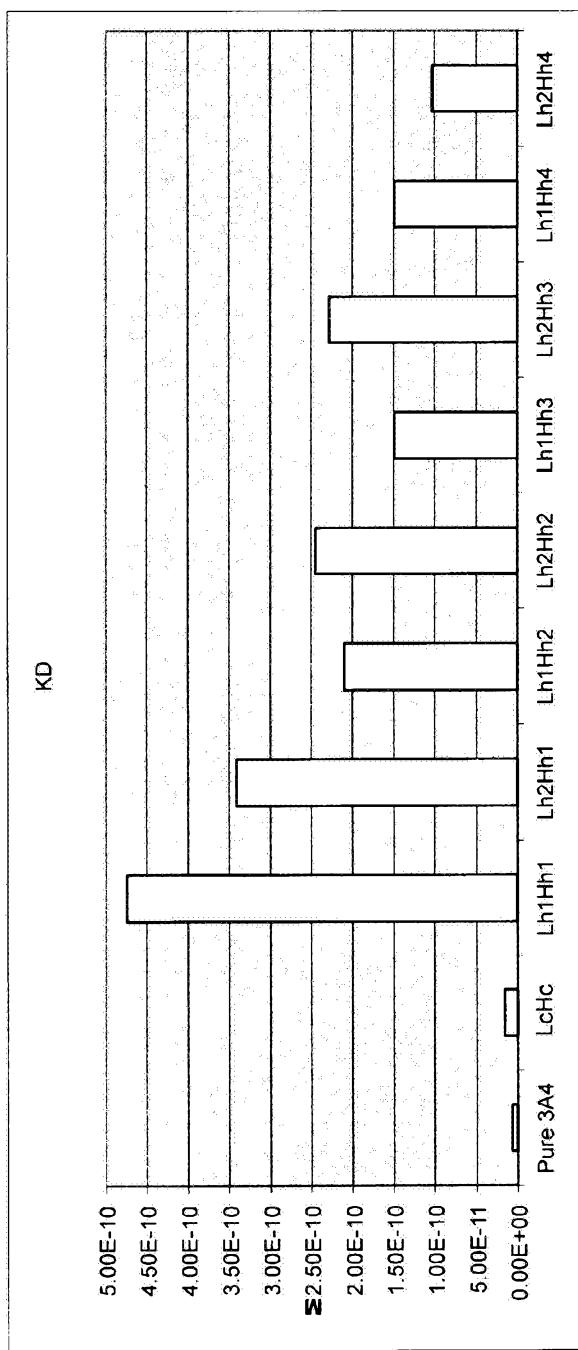


Figure 8a

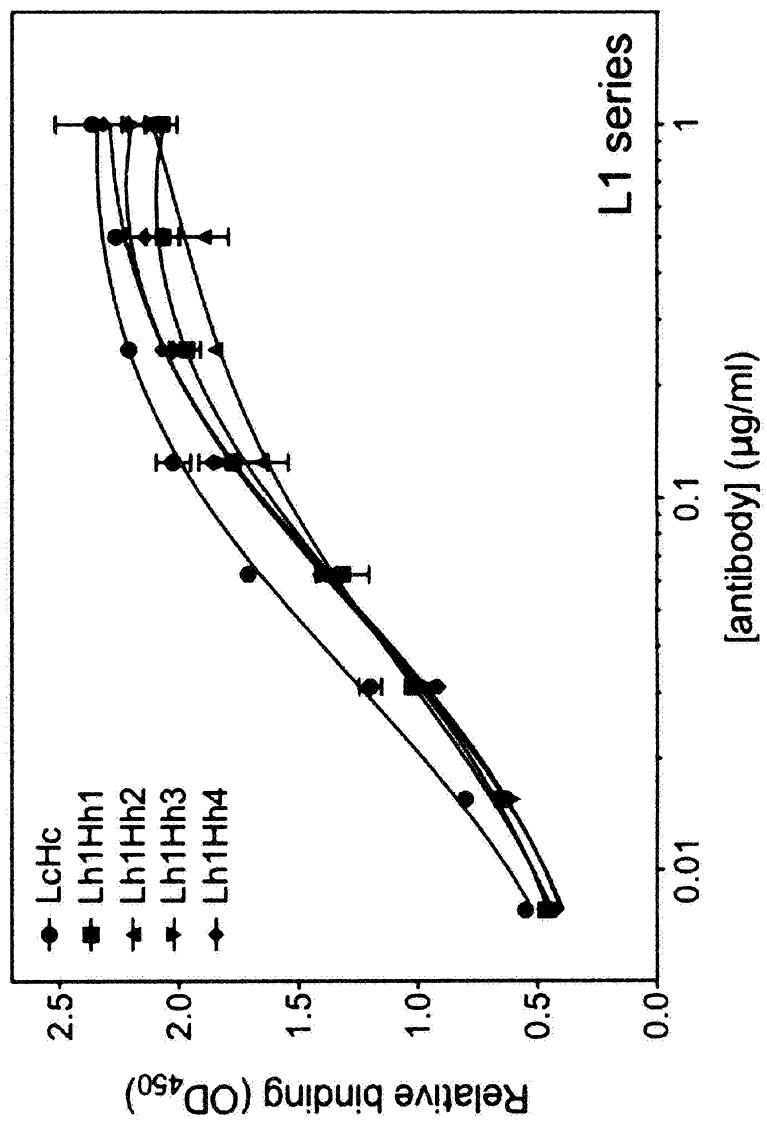
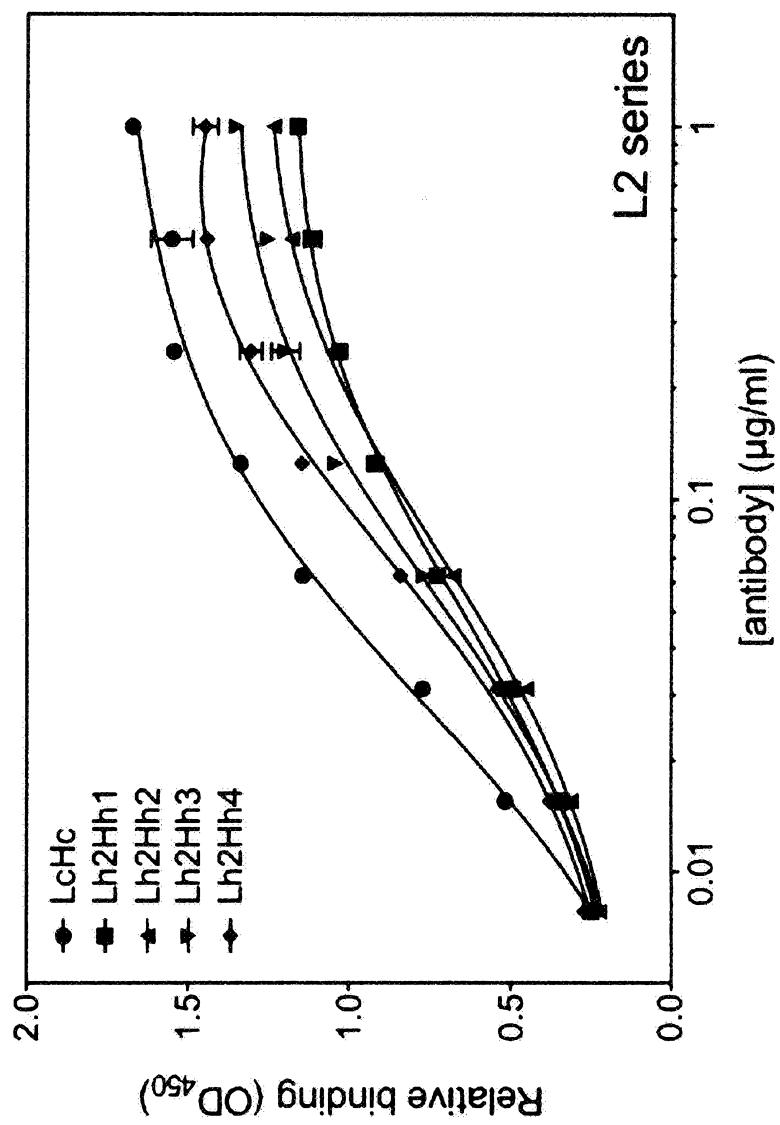


Figure 8b



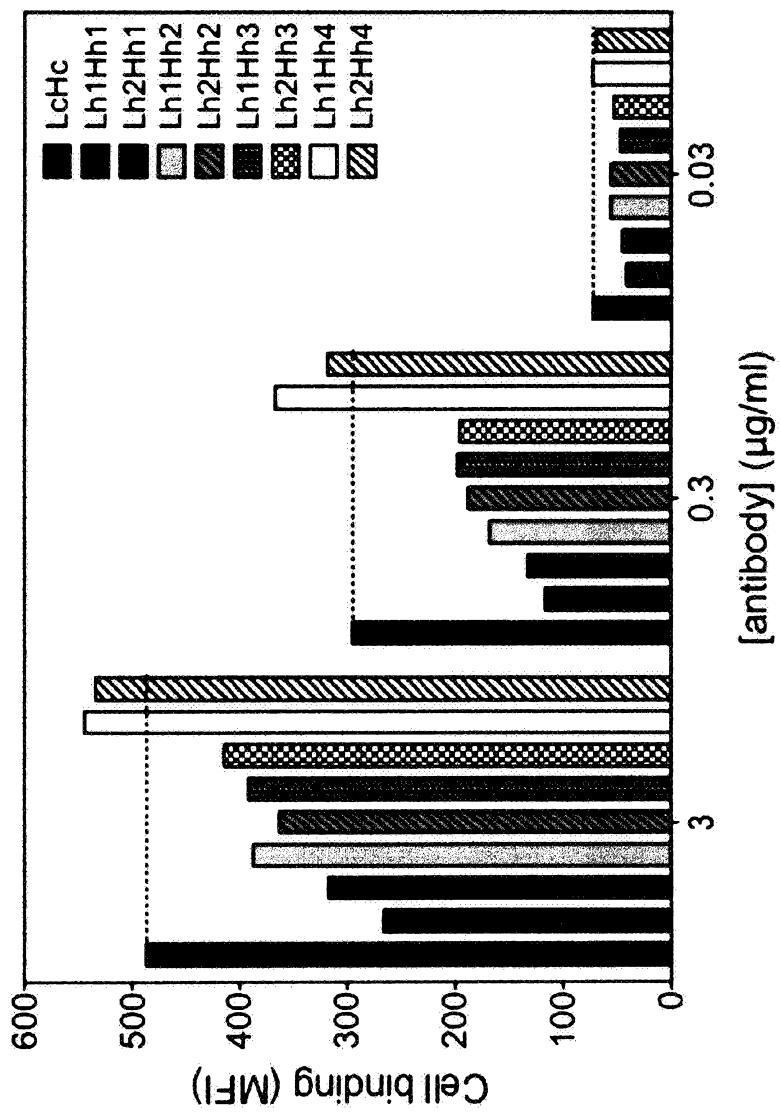


Figure 9

Figure 10

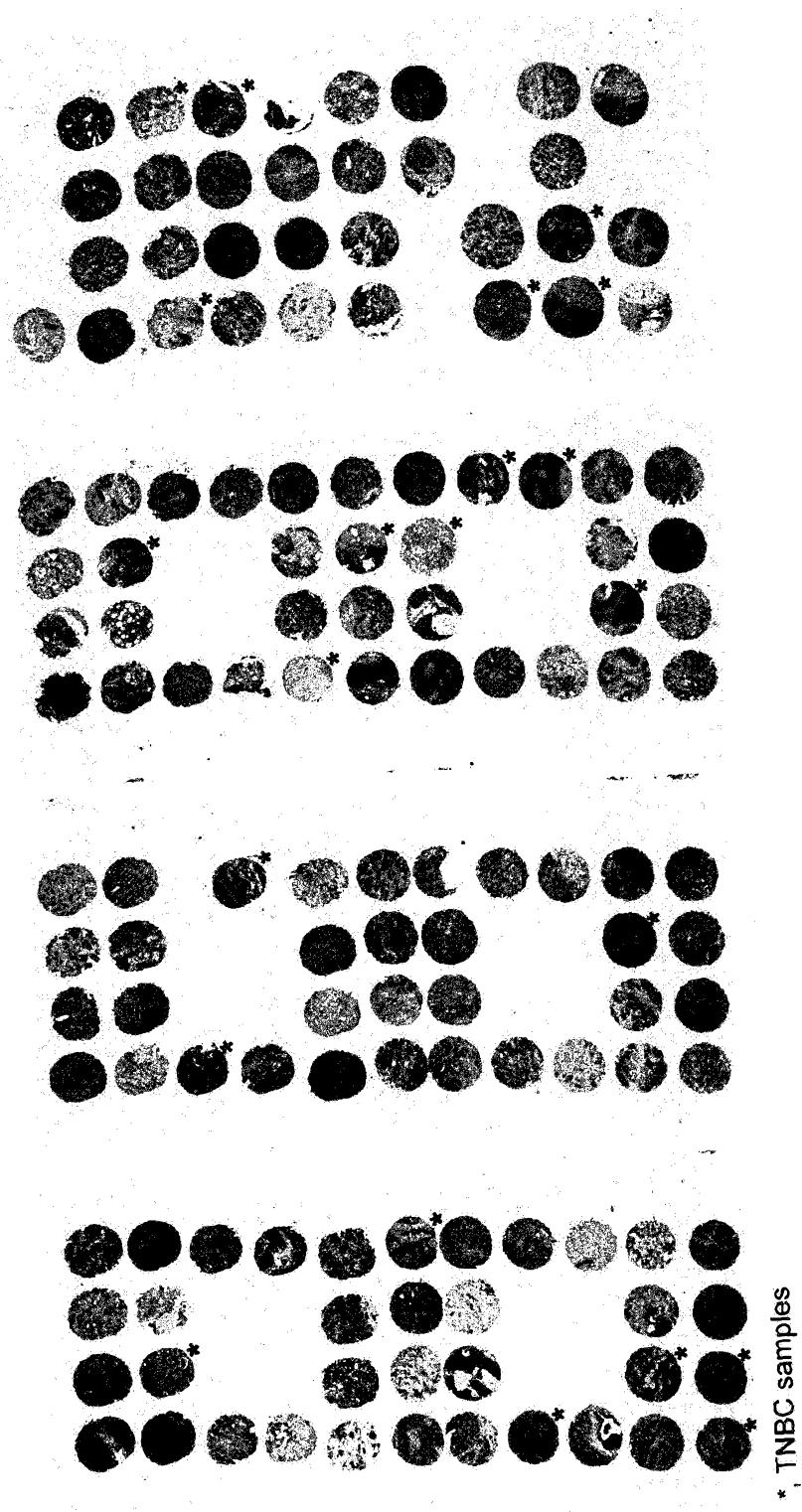


Figure 11

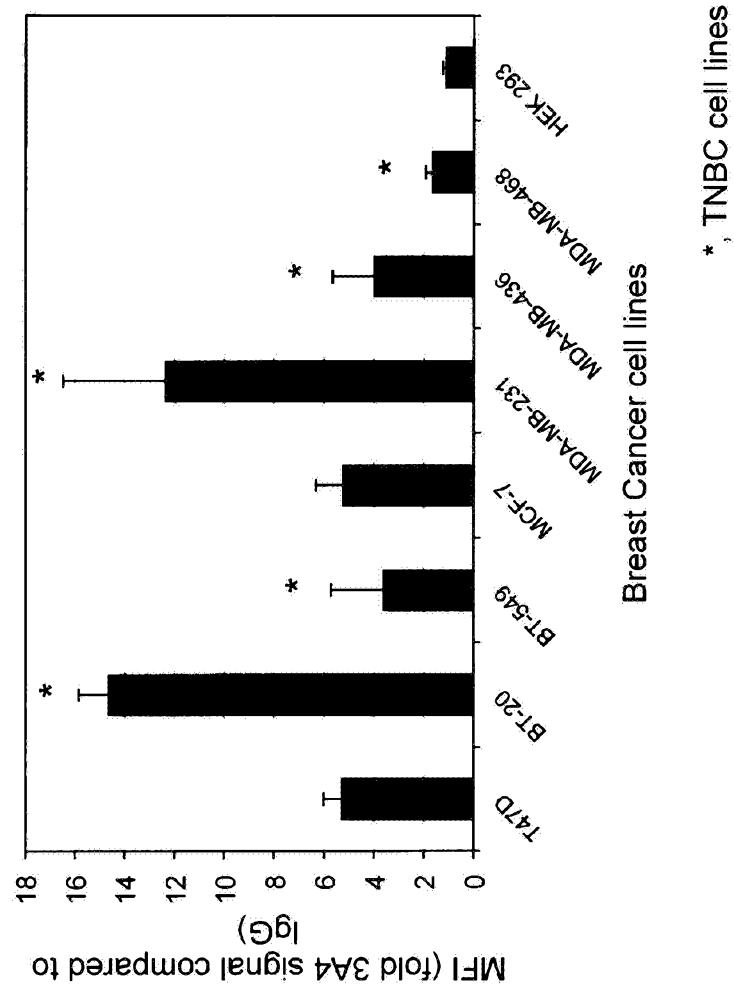


Figure 12

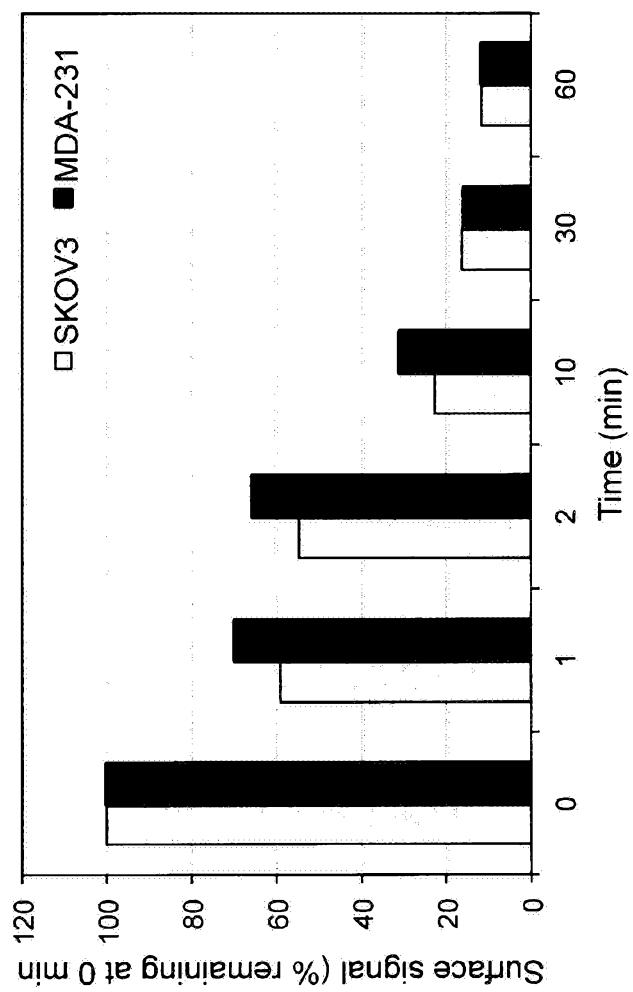


Figure 13

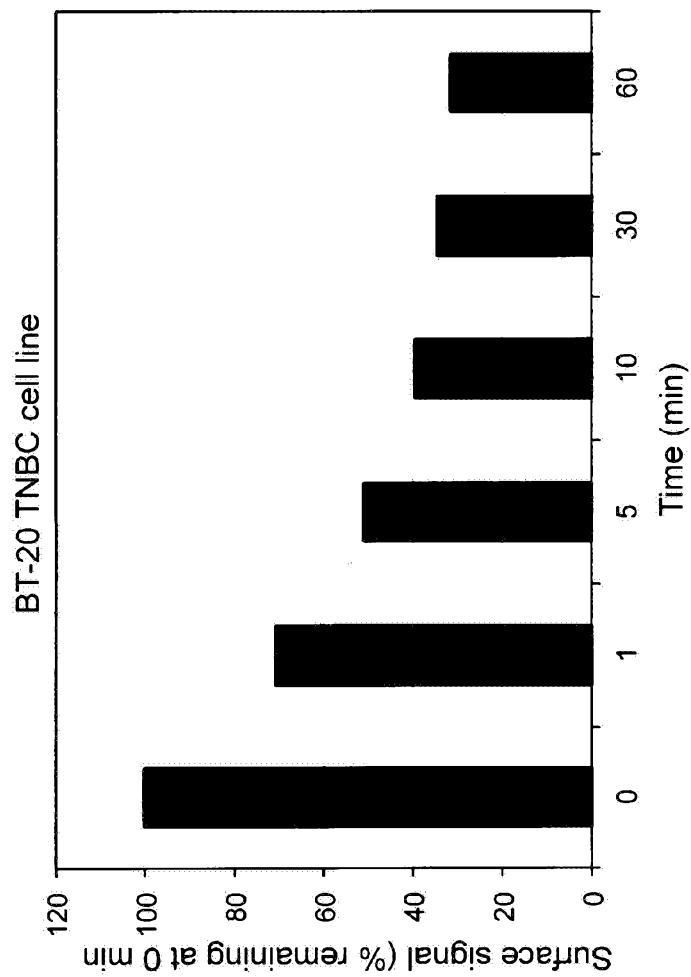


Figure 14

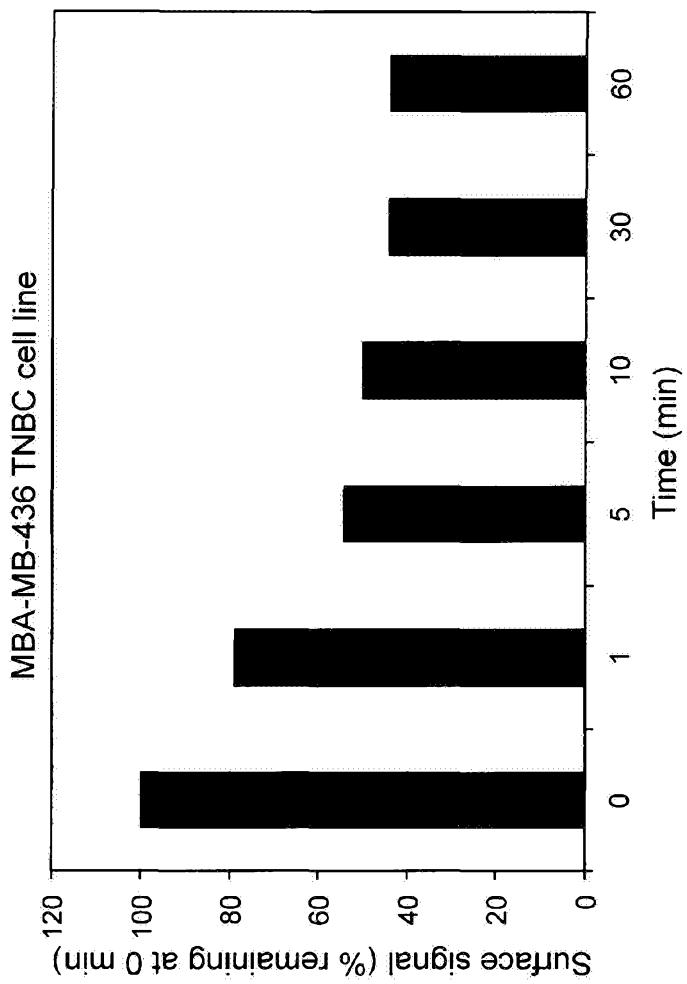


Figure 15

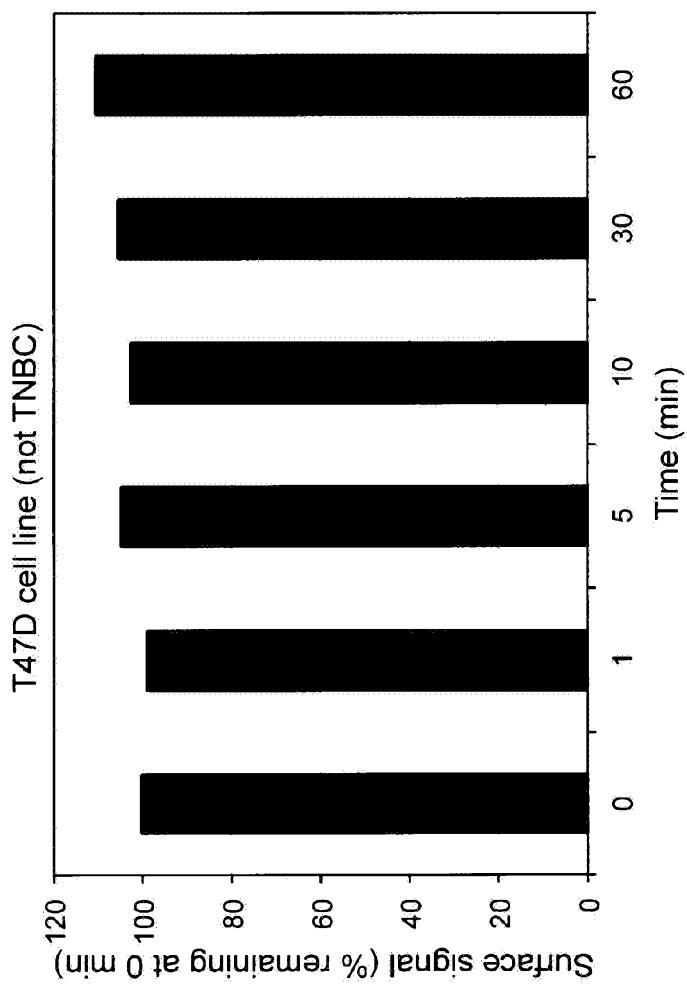


Figure 16

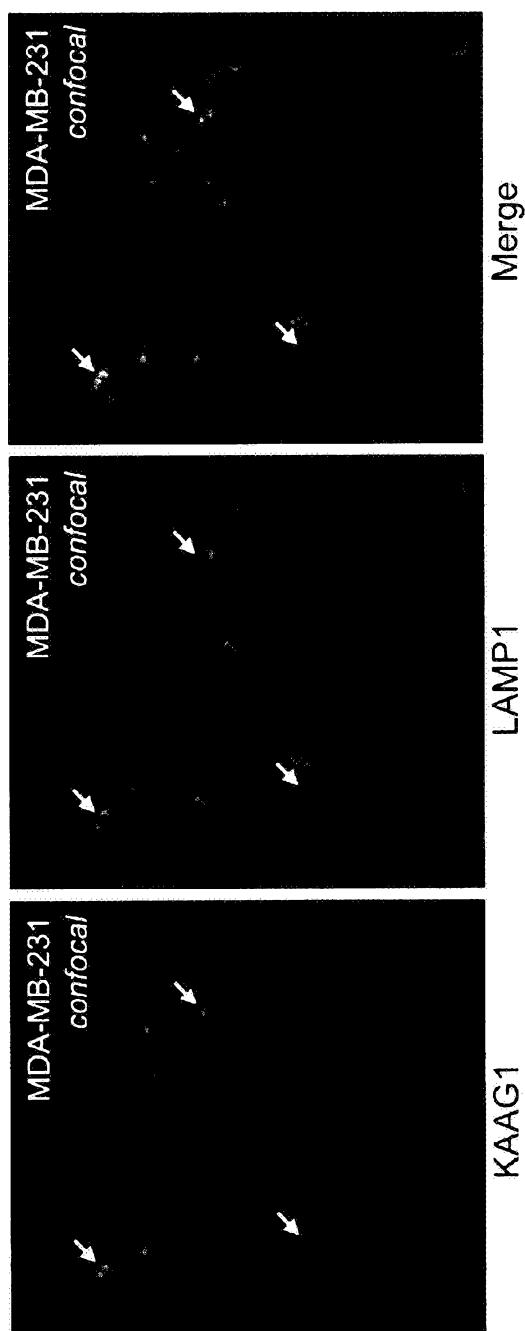




Figure 17