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(54) Title: COMPOSITIONS COMPRISING ALKALINE PHOSPHATASE AND/OR NATRIURETIC PEPTIDE AND METHODS OF USE THEREOF

(57) Abstract: The present invention provides methods, compositions, and kits for the treatment of neurocutaneous syndromes, such as neurofibromatosis type I disorders associated with overactivation of FGFR3, such as achondroplasia; bone or cartilage disorders; or vascular smooth muscle disorders; or for the elongation of bone. Polypeptides having an alkaline phosphatase peptide fused to an Fc domain of an immunoglobulin or a natriuretic peptide fused to an Fc domain of an immunoglobulin can be administered to subjects, e.g., subcutaneously, to treat a neurocutaneous syndromes. The invention also features nucleic acid molecules encoding such polypeptides and the use of the nucleic acid molecules for treating neurocutaneous syndromes.

**COMPOSITIONS COMPRISING ALKALINE PHOSPHATASE AND/OR NATRIURETIC PEPTIDE AND METHODS OF USE THEREOF**

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**CROSS-REFERENCE TO RELATED APPLICATIONS**

This application claims benefit of priority to U.S. Provisional Application No. 61/549,047, filed October 19, 2011, and U.S. Provisional Application No. 61/649,717, filed May 21, 2012, each of which is hereby incorporated by reference.

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**REFERENCE TO A SEQUENCE LISTING**

A Sequence Listing is provided in this patent document as a .txt file entitled, "50694\_039WO3\_ST25\_Seq\_Listing.txt," created May 21, 2012 (file size 716 kB). The content of this file is hereby incorporated by reference.

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**BACKGROUND OF THE INVENTION**

In general, this invention relates to the treatment of various diseases using alkaline phosphatase and/or natriuretic peptide.

Numerous diseases and conditions involve abnormal skeletal function, structure, or growth of bone or cartilage. For some diseases, the etiology of these skeletal manifestations is known, such as in 20 hypophosphatasia (HPP) and achondroplasia (ACH), but treatment options are limited. In other diseases, the etiology is unknown. For example, neurofibromatosis type I (NF1 or Von Recklinghausen disease) is an autosomal dominant genetic disorder having an incidence of approximately 1 in 3,500 live births. NF1 encodes neurofibromin, a member of the GTPase Activating Protein (GAP) family known to suppress the Ras kinase. Neurofibromin is a specific suppressor of p21-RAS, and mutations in the NF1 gene cause 25 unsuppressed activation of RAS that lead to abnormal cell growth and differentiation. Accordingly, the clinical features of NF1 include various oncogenic transformations, such as neurocutaneous neurofibromas and optic pathway tumors, and other non-cancer manifestations, such as cognitive defects and skeletal abnormalities. Some of the NF1 skeletal manifestations have high morbidity (e.g., dystrophic scoliosis, long bone bowing, and pseudarthrosis) and unsatisfactory treatment options, which 30 has been complicated by the fact that the etiology of these manifestations is unclear.

Many skeletal diseases arise from loss of function of one or more proteins. For example, hypophosphatasia (HPP) is a rare, heritable disease caused by one or more loss-of-function mutations in the gene *ALPL*, which encodes tissue-nonspecific alkaline phosphatase (TNALP; a.k.a. liver/bone/kidney type ALP). Alkaline phosphatase deficiency in osteoblasts and chondrocytes impairs skeletal 35 mineralization, leading to symptoms of varying severity, from rickets or osteomalacia to almost complete absence of bone mineralization *in utero*. However, enzyme replacement therapy with unmodified alkaline phosphatase (e.g., infusion of native alkaline phosphatase) has been largely unsuccessful.

In another example, achondroplasia (ACH) is the most common form of short limb dwarfism in human beings, affecting more than 250,000 individuals worldwide, and is caused by mutations in the gene encoding fibroblast growth factor receptor 3 (FGFR3), which cause gain of FGFR3 function. The severity of the clinical phenotype is related to the capacity of the mutation to overactivate FGFR3 5 signaling pathways in chondrocytes, such as the MAP-kinase pathway. This pathway can be inhibited by activating the natriuretic peptide receptor B (NPR-B), which produces the second messenger cGMP, and cGMP, in turn, inhibits the MAP-kinase pathway inside the cell. In the cellular environment, the immature and mature forms of C-type natriuretic peptide (CNP), such as CNP53 and CNP22, bind to 10 NPR-B and induce cGMP production in a dose-dependent and similar fashion. Thus, use of CNP or a CNP analog that could activate the NPR-B signaling pathway for the treatment of skeletal dysplasia has been considered. However, a major drawback of the therapeutic use of CNP is its extremely short half-life.

There is thus a need in the art to develop therapeutic molecules and methods for treating diseases having skeletal manifestations, such as neurofibromatosis. In addition, more therapeutic molecules are 15 needed that have an appreciable half-life and/or other favorable pharmacokinetic and therapeutic properties, and these molecules can be used to treat a variety of disorders that would benefit from their underlying mode of action, such as neurofibromatosis, hypophosphatasia, and achondroplasia.

## SUMMARY OF THE INVENTION

20 It has surprisingly been discovered that neurofibromatosis, a neurocutaneous syndrome resulting in tumors in the nervous system, results in bone manifestations that arise from accumulation of inorganic pyrophosphate (PP<sub>i</sub>). As alkaline phosphatase-Fc fusion proteins (either with or without a bone-targeting moiety) can reduce PP<sub>i</sub> accumulation, the present invention provides a polypeptide including a soluble alkaline phosphatase (sALP) domain (i.e., an sALP polypeptide), as well as compositions and uses 25 thereof. In addition, bone manifestations in neurofibromatosis may also arise from overactivation of the MAP-kinase pathway, and a natriuretic peptide (NP) or NP analog could be used to inhibit this pathway. Accordingly, the present invention provides a polypeptide including an NP (e.g., a CNP) domain (i.e., an NP polypeptide) and compositions and uses thereof.

Furthermore, the present invention includes a combination of an sALP polypeptide and an NP 30 polypeptide, where these polypeptides can be administered separately or together. This combination would be particularly useful in diseases that would benefit from increased ALP levels (e.g., disorders associated with increased levels of PP<sub>i</sub>, such as neurofibromatosis, or disorders associated with ALP deficiency, such as hypophosphatasia) and/or diseases that would benefit from inactivation of a signaling pathway involving FGFR3 (e.g., disorders associated with overactivation of the MAP-kinase pathway, 35 such as neurofibromatosis or achondroplasia, or disorders associated with overactivation of FGFR3, such as achondroplasia or craniosynostosis or cancer) and/or diseases that would benefit from increased NP levels (e.g., disorders associated with CNP deficiency, such as skeletal dysplasia, or vascular smooth

muscle disorders). The polypeptides of the invention can also be provided in kits, either separately or together.

In a first aspect, the invention features a method of treating a neurocutaneous syndrome in a subject, the method including administering to the subject a therapeutically effective amount of a pharmaceutical composition including: (a) a polypeptide including the structure A-sALP-B; and (b) a pharmaceutically acceptable excipient, where sALP is the extracellular domain of an alkaline phosphatase, A is absent or is an amino acid sequence of at least one amino acid, and B is absent or is an amino acid sequence of at least one amino acid. In some embodiments, the syndrome in the subject is thereby treated.

10 In some embodiments, the amino acid sequence of the sALP includes or consists of amino acid residues 23-508 of SEQ ID NO: 1215, amino acid residues 18-498 of SEQ ID NO: 1216, amino acid residues 23-508 of SEQ ID NO: 1218, or amino acid residues 18-498 of SEQ ID NO: 1219, or the amino acid sequence of the sALP includes or consists of amino acid residues 23-512 of SEQ ID NO: 1215, amino acid residues 18-502 of SEQ ID NO: 1216, amino acid residues 23-512 of SEQ ID NO: 1218, or 15 amino acid residues 18-502 of SEQ ID NO: 1219. In some embodiments, the amino acid sequence of the sALP includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NO: 1205. In some embodiments, the amino acid sequence of the sALP includes or consists of the amino acid sequence of SEQ ID NO: 1205.

20 In some embodiments, the amino acid sequence of the sALP includes or consists of amino acid residues 1-497, 1-498, 1-499, 1-500, 1-501, 1-502, 1-503, 1-504, 1-505, 1-506, 1-507, 1-508, 1-509, 1-510, 1-511, 1-512, 23-497, 23-498, 23-499, 23-500, 23-501, 23-502, 23-503, 23-504, 23-505, 23-506, 23-507, 23-508, 23-509, 23-510, 23-511, or 23-512 of SEQ ID NO: 1218, where X is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP, e.g., not an amino acid corresponding to a pathogenic mutation provided in Table 1. In some embodiments, the 25 amino acid sequence of the sALP includes or consists of amino acid residues 18-497, 18-498, 18-499, 18-500, 18-501, 18-502, 18-503, 18-504, 18-505, 18-506, 18-507, 18-508, 18-509, 18-510, 18-511, or 18-512 of SEQ ID NO: 1219, where X is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP, e.g., not an amino acid corresponding to a pathogenic mutation provided in Table 1.

30 In some embodiments, the amino acid sequence of the sALP includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NOs: 1218 or 1219, where X is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP, e.g., not an amino acid corresponding to a pathogenic mutation provided in Table 1. In some embodiments, the amino acid sequence of the sALP includes or consists of an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to amino acid residues 23-508 of SEQ ID NO: 1215, amino acid residues 18-498 of SEQ ID NO: 1216, amino acid residues 23-508 of SEQ ID NO: 1218, or amino acid residues 18-498 of SEQ ID NO: 1219, or the amino acid sequence of the sALP

consists of an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to amino acid residues 23-512 of SEQ ID NO: 1215, amino acid residues 18-502 of SEQ ID NO: 1216, amino acid residues 23-512 of SEQ ID NO: 1218, or amino acid residues 18-502 of SEQ ID NO: 1219, where X in SEQ ID NO: 1218 or 1219 is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP, e.g., not an amino acid corresponding to a pathogenic mutation provided in Table 1.

5 In some embodiments, A and/or B are absent.

In some embodiments, A and/or B includes a fragment crystallizable region (Fc). In some embodiments, the Fc includes a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region, or the Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, and IgG-4, e.g., IgG-1. In some embodiments, the amino acid sequence of the Fc includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NO: 401. In some embodiments, the amino acid sequence of the Fc includes or consists of the amino acid sequence of SEQ ID NO: 401.

15 In some embodiments, A and/or B includes I<sub>n</sub>, where I represents an aspartic acid or a glutamic acid and n=10 to 16, e.g., n is 10, 11, 12, 13, 14, 15, or 16. In some embodiments, the I<sub>n</sub> is E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>10</sub> or D<sub>10</sub>.

In some embodiments, A and/or B includes a bone-targeting moiety, e.g., including 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid. In some 20 embodiments, the bone-targeting moiety includes or consists of E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>. In some embodiments, the polypeptide does not include a polyaspartic acid or polyglutamic acid region longer than three consecutive aspartic acid or glutamic acid residues, or the polypeptide does not include a polyaspartic acid or polyglutamic acid region longer than two consecutive aspartic acid or glutamic acid residues.

25 In some embodiments, the polypeptide does not include a bone-targeting moiety.

In some embodiments, the polypeptide includes the structure C-sALP-D-Fc-E or the structure C-Fc-D-sALP-E, where C is absent or is an amino acid sequence of at least one amino acid, D is absent or is an amino acid sequence of at least one amino acid, and E is absent or is an amino acid sequence of at least one amino acid.

30 In some embodiments, C and/or E are absent. In some embodiments, D is two amino acid residues, e.g., leucine-lysine or aspartic acid-isoleucine. In some embodiments, D is any linker described herein, e.g., the amino acid sequence of any one of SEQ ID NOs: 301-391.

In some embodiments, the polypeptide includes the structure C-sALP-D-Fc-G-I<sub>n</sub>-H or the structure C-Fc-D-sALP-G-I<sub>n</sub>-H, where C is absent or is an amino acid sequence of at least one amino acid, D is absent or is an amino acid sequence of at least one amino acid, G is absent or is an amino acid sequence of at least one amino acid, H is absent or is an amino acid sequence of at least one amino acid, I represents an aspartic acid or a glutamic acid, and n=10 to 16, e.g., n is 10, 11, 12, 13, 14, 15, or 16. In

some embodiments, the  $I_n$  is  $E_{10}$ ,  $E_{11}$ ,  $E_{12}$ ,  $E_{13}$ ,  $E_{14}$ ,  $E_{15}$ ,  $E_{16}$ ,  $D_{10}$ ,  $D_{11}$ ,  $D_{12}$ ,  $D_{13}$ ,  $D_{14}$ ,  $D_{15}$ , or  $D_{16}$ , e.g.,  $E_{10}$  or  $D_{10}$ .

In some embodiments, C and/or H are absent. In some embodiments, G is two amino acid residues, e.g., leucine-lysine or aspartic acid-isoleucine, e.g., aspartic acid-isoleucine. In some 5 embodiments, I is an aspartic acid or a glutamic acid and  $n=10$  to 16, e.g.,  $n$  is 10, 11, 12, 13, 14, 15, or 16, e.g., I is aspartic acid and  $n=10$ . In some embodiments, D is two amino acid residues, e.g., leucine-lysine or aspartic acid-isoleucine, e.g., leucine-lysine. In some embodiments, D or G is any linker described herein, e.g., the amino acid sequence of any one of SEQ ID NOs: 301-391.

In some embodiments, the amino acid sequence of the polypeptide includes an amino acid 10 sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity of any one of SEQ ID NOs: 1201, 1204, 1220, or 1221, e.g., SEQ ID NO: 1204. In some embodiments, the amino acid sequence of the polypeptide includes or consists of the amino acid sequence of any one of SEQ ID NOs: 1201, 1204, 1220, or 1221, e.g., SEQ ID NO: 1204. In some embodiments, the amino acid sequence of the polypeptide consists of the amino acid sequence of SEQ ID NO: 1204.

15 In a second aspect, the invention features a method of treating a neurocutaneous syndrome in a subject, the method including administering to the subject a therapeutically effective amount of a pharmaceutical composition including: (a) a polypeptide including the structure V-NP-W; and (b) a pharmaceutically acceptable excipient, where NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), V is absent or is an amino acid sequence of at least one amino acid, and W 20 is absent or is an amino acid sequence of at least one amino acid. In some embodiments, the syndrome in the subject is thereby treated.

In some embodiments, the NP includes the structure: [N-terminal extension]-[short segment]-[ring domain]-[C-terminal extension], where the ring domain includes the amino acid sequence of SEQ 25 ID NO: 6, amino acid residues 11-27 of SEQ ID NO: 30, or SEQ ID NO: 95, and each of the N-terminal extension, short segment, and C-terminal extension is, independently, absent or is an amino acid sequence of at least one amino acid. In some embodiments, the ring domain includes amino acid residues 6-22 of SEQ ID NO: 126. In some embodiments, the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Val, Ala, Ser, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe, Leu, Ile, Thr, Val, Ala, or Ser, e.g., Phe or 30 Leu, e.g., Phe, e.g., Leu. In some embodiments, the ring domain includes the amino acid sequence of SEQ ID NO: 12. In some embodiments, the short segment and the ring domain together include the amino acid sequence of any one of SEQ ID NOs: 4, 13-30, 119-122, 126, or 156-161, e.g., SEQ ID NOs: 4 or 13-30. In some embodiments, the amino acid sequence of the short segment includes or consists of amino acid residues 1-5, 2-5, 3-5, 4-5, or 5, e.g., consists of amino acid residues 1-5, of SEQ ID NO: 4; 35 amino acid residues 1-10 of SEQ ID NO: 17; amino acid residues 1-5 of SEQ ID NO: 19; amino acid residues 1-3 of SEQ ID NO: 20; amino acid residues 1-5 of SEQ ID NO: 21; or amino acid residues 1-6 of SEQ ID NO: 29. In some embodiments, the amino acid sequence of the short segment and the ring

domain together includes or consists of the amino acid sequence of SEQ ID NO: 4. In some embodiments, the amino acid sequence of the short segment and the ring domain together includes or consists of the amino acid sequence of any one of SEQ ID NOs: 119-122, 126, or 156-161 (e.g., where X in SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe, Leu, Ile, Thr, Val, Ala, or Ser, e.g., Phe or Leu, e.g., Phe, e.g., Leu).

5 In some embodiments, the amino acid sequence of the N-terminal extension includes amino acid residues 1-31 or 17-31 of SEQ ID NO: 11. In some embodiments, the amino acid sequence of the N-terminal extension includes amino acid residues 17-31 of SEQ ID NO: 11. In some embodiments, the amino acid sequence of the N-terminal extension includes KGANKK (SEQ ID NO: 314) or KGANQK (SEQ ID NO: 315). In some embodiments, the N-terminal extension, short segment, and ring domain together include the amino acid sequence of SEQ ID NO: 11. In some embodiments, the C-terminal extension includes the amino acid sequence of SEQ ID NOs: 117 or 118 or includes amino acid residues 23-37 selected from any one of SEQ ID NOs: 101-116. In some embodiments, the amino acid sequence 10 of the NP consists of SEQ ID NOs: 4 or 11, or the amino acid sequence of any one of SEQ ID NOs: 31-94, or a fragment thereof including at least a ring domain, or the amino acid sequence of any one of SEQ ID NOs: 13-29, 100-116, 119-125, 127-233, or 1001-1155.

15 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Phe.

20 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Leu.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Ile.

25 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Thr.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Glu.

30 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Arg.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Tyr.

35 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Cys.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Pro.

40 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Asp.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Gly.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Ala.

5 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Ser.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Val.

10 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Trp.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Asn.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Gln.

15 In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be His.

In any of the aspects described herein, the amino acid sequence of the NP may include amino acids 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 may be Lys.

In some embodiments, V and/or W are absent.

20 In some embodiments, V and/or W includes a fragment crystallizable region (Fc). In some embodiments, the Fc includes a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region, or the Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, and IgG-4, e.g., IgG-1. In some embodiments, the amino acid sequence of the Fc includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NO: 401. In some 25 embodiments, the amino acid sequence of the Fc includes or consists of the amino acid sequence of SEQ ID NO: 401.

In some embodiments, V and/or W includes a glycine-rich region.

In some embodiments, the amino acid sequence of V or W consists of one or more glycines and one or more serines. In some embodiments, the amino acid sequence of V or W includes 30 [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, where each of m, n, and p is, independently, between 0 and 20. In some embodiments, m is between 1 and 6; n is between 1 and 10; and p is between 0 and 4, e.g., m is 4 and n is 1-6. In some embodiments, combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of V or W includes the amino acid sequence of any one of SEQ ID NOs: 301-391.

35 In some embodiments, V and/or W does not include a bone-targeting moiety.

In some embodiments, V and/or W includes a bone-targeting moiety, e.g., including 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid. In some

embodiments, the bone-targeting moiety includes or consists of E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

In some embodiments, V and/or W includes a cathepsin (e.g., cathepsin K) cleavage sequence.

In some embodiments, the cathepsin cleavage sequence is HGPQG (SEQ ID NO: 374) or HKLRG (SEQ 5 ID NO: 375).

In some embodiments, the polypeptide includes the structure V-NP-W, where NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), each of V and W is, independently, absent or is an amino acid sequence of at least one amino acid, and the NP includes the amino acid sequence of any one of SEQ ID NOs: 17-29, 31-40, 42-94, 101-116, 119-122, 128-161, or 163-233, or V 10 or W includes the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

In some embodiments, the polypeptide includes the structure V-NP or NP-W, where NP is the natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and each of V and W includes, independently, the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

15 In some embodiments, the polypeptide includes the structure X-Fc-Y-NP-Z or the structure X-NP-Y-Fc-Z, where NP is the natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B) (e.g., any NP described herein), and each of X, Y, and Z is, independently, absent or is an amino acid sequence of at least one amino acid. In some embodiments, Y includes a glycine-rich region. In some 20 embodiments, the amino acid sequence of Y consists of one or more glycines and one or more serines. In some embodiments, the amino acid sequence of Y includes [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, where each of m, n, and p is, independently, between 0 and 20. In some embodiments, m is between 1 and 6; n is between 1 and 10; and p is between 0 and 4, e.g., m is 4 and n is 1-6. In some embodiments, 25 combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of Y includes the amino acid sequence of any one of SEQ ID NOs: 301-389. In some embodiments, the amino acid sequence of Y consists of [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, wherein combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of Y consists of the amino acid sequence of any one of SEQ ID NOs: 301-389. In some embodiments, X is absent, Z is absent, or X and Z are both absent. In some embodiments, X, Y, or Z includes a bone-targeting moiety, e.g., including 30 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid. In some embodiments, the bone-targeting moiety includes or consists of E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

In some embodiments, X, Y, or Z includes a cathepsin (e.g., cathepsin K) cleavage sequence. In some embodiments, the cathepsin cleavage sequence includes or consists of HGPQG (SEQ ID NO: 374) or HKLRG (SEQ ID NO: 375).

35 In some embodiments, the amino acid of the polypeptide includes or consists of the amino acid sequence of any one of SEQ ID NOs: 501-608, e.g., SEQ ID NOs: 502, 504, 506, 512, 514, 516, 530, 560, 562, 564, 572, 574, 576, 584, 586, 588, 596, 598, 600, or 608, e.g., SEQ ID NOs: 504, 512, 530,

554, 572, or 578, e.g., SEQ ID NO: 512. In some embodiments, the amino acid of the polypeptide includes a bone-targeting moiety, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

10 In some embodiments, the amino acid sequence of the polypeptide includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to any one of SEQ ID NOS: 504, 512, 530, 554, 572, or 578, e.g., SEQ ID NOS: 504, 512, 530, or 572, e.g., SEQ ID NO: 512. In some embodiments, the amino acid sequence of the polypeptide includes or consists of the amino acid sequence of any one of SEQ ID NOS: 504, 512, 530, 554, 572, or 578, e.g., SEQ ID NOS: 504, 512, 530, or 572, e.g., SEQ ID NO: 512. In some embodiments, the amino acid sequence of the polypeptide 15 consists of the amino acid sequence of SEQ ID NOS: 504, 512, 530, or 572, e.g., SEQ ID NO: 512.

20 In some embodiments, the polypeptide includes the structure X-Fc-Y-NP-Z or the structure X-NP-Y-Fc-Z, wherein NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and wherein either: (i) NP includes amino acids 6-22 of SEQ ID NO: 126, wherein the amino acid at position 17 is not Met; and each of X, Y, and Z is, independently, absent or is an amino acid sequence of at least one amino acid; or (ii) each of X and Z is, independently, absent or is an amino acid sequence of 15 at least one amino acid; and the amino acid sequence of Y includes [(Gly)<sub>4</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>4</sub>]<sub>n</sub>, wherein n is between 1 and 10 and p is between 0 and 4 or wherein combinations of m, n, and p are selected from a single row of Table 2, or wherein the amino acid sequence of Y includes the amino acid sequence of any one of SEQ ID NOS: 304-313, 322-333, or 337-391.

25 In some embodiments, the polypeptide includes the structure X-Fc-Y-NP-Z.

30 In some embodiments, (i) NP includes amino acids 6-22 of SEQ ID NO: 126, wherein the amino acid at position 17 is not Met; and each of X, Y, and Z is, independently, absent or is an amino acid sequence of at least one amino acid. In some embodiments, the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Val, Ala, or Ser. In some embodiments, the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu. In some embodiments, the NP includes the structure: [N-terminal extension]-[short segment]-[ring domain]-[C-terminal extension], wherein said ring domain comprises amino acids 6-22 of SEQ ID NO: 126, wherein the amino acid at position 17 is not Met, and each of said N-terminal extension, short segment, and C-terminal extension is, independently, absent or is an amino acid sequence of at least one amino acid. In some embodiments, the amino acid sequence of said NP includes or consists of the amino acid sequence of any one of SEQ ID NOS: 119-125 or 156-220, wherein position 17 relative to SEQ ID NO: 126 is not Met, or the amino acid sequence of any one of SEQ ID NOS: 221-233.

35 In some embodiments, (ii) each of X and Z is, independently, absent or is an amino acid sequence of at least one amino acid; and the amino acid sequence of Y comprises [(Gly)<sub>4</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>4</sub>]<sub>n</sub>, wherein n is between 1 and 10 and p is between 0 and 4, or wherein the amino acid sequence of Y comprises the amino acid sequence of any one of SEQ ID NOS: 304-313, 322-333, or 337-

391. In some embodiments, the NP includes the structure: [N-terminal extension]-[short segment]-[ring domain]-[C-terminal extension], wherein the ring domain includes the amino acid sequence of SEQ ID NO: 6, amino acids 11-27 of SEQ ID NO: 30, or SEQ ID NO: 95, and each of the N-terminal extension, short segment, and C-terminal extension is, independently, absent or is an amino acid sequence of at least 5 one amino acid. In some embodiments, the ring domain includes amino acids 6-22 of SEQ ID NO: 126. In some embodiments, the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Val, Ala, or Ser. In some embodiments, the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu. In some embodiments, the ring domain 10 includes the amino acid sequence of SEQ ID NO: 12. In some embodiments, the short segment and the ring domain together include the amino acid sequence of any one of SEQ ID NOs: 4 or 13-30. In some embodiments, the amino acid sequence of the short segment and the ring domain together consists of the amino acid sequence of SEQ ID NO: 4. In some embodiments, the amino acid sequence of the short segment and the ring domain together consists of the amino acid sequence of any one of SEQ ID NOs: 15 119-122, 126, or 156-161 (e.g., where X in SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu). In some embodiments, the N-terminal extension, short segment, and ring domain together include the amino acid sequence of SEQ ID NO: 11. In some 20 embodiments, the amino acid sequence of the NP consists of SEQ ID NO: 4. In some embodiments, the amino acid sequence of the NP consists of SEQ ID NO: 11. In some embodiments, the amino acid sequence of the NP consists of the amino acid sequence of any one of SEQ ID NOs: 31-94, or a fragment thereof including at least a ring domain. In some embodiments, the amino acid sequence of the NP includes or consists of the amino acid sequence of any one of SEQ ID NOs: 13-29, 100-116, 119-125, 127-233, or 1001-1155.

25 In some embodiments, the amino acid sequence of the short segment consists of amino acids 1-5 of SEQ ID NO: 4. In some embodiments, the amino acid sequence of the short segment consists of amino acids 1-5, 2-5, 3-5, 4-5, or 5 of SEQ ID NO: 4, amino acids 1-10 of SEQ ID NO: 17, amino acids 1-5 of SEQ ID NO: 19, amino acids 1-3 of SEQ ID NO: 20, amino acids 1-5 of SEQ ID NO: 21, or amino acids 1-6 of SEQ ID NO: 29. In some embodiments, the amino acid sequence of the N-terminal 30 extension includes amino acids 1-31 of SEQ ID NO: 11. In some embodiments, the amino acid sequence of the N-terminal extension includes amino acids 17-31 of SEQ ID NO: 11. In some embodiments, the amino acid sequence of the N-terminal extension includes KGANKK (SEQ ID NO: 314) or KGANQK (SEQ ID NO: 315). In some embodiments, the C-terminal extension includes the amino acid sequence of SEQ ID NO: 118, SEQ ID NO: 117, or amino acids 23-37 selected from any one of SEQ ID NOs: 101- 35 116.

In some embodiments, the NP is selective for NPR-B over NPR-A, wherein the  $EC_{50(NPR-A)}/EC_{50(NPR-B)}$  ratio for the NP, as determined in an *in vivo* pharmacokinetic assay, is at least 30.

In some embodiments, the Fc includes a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region. In some embodiments, the Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, IgG-3 and IgG-4. In some embodiments, the Fc includes the amino acid sequence of SEQ ID NO: 401. In some embodiments, the immunoglobulin is IgG-1. In some embodiments, the amino acid sequence of the Fc includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NO: 401, or includes or consists of the amino acid sequence of SEQ ID NO: 401.

In some embodiments, Y includes a glycine-rich region, or the amino acid sequence of Y consists of one or more glycines and one or more serines. For example, the amino acid sequence of Y may include [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, wherein each of m, n, and p is, independently, between 0 and 20. In some embodiments, m is 0-20 (e.g., m is 3-6, 3-7, 3-8, 3-9, 3-10, 3-11, 3-12, 3-14, 3-15, 3-16, 3-17, 3-18, 3-19, 3-20, 4-6, 4-7, 4-8, 4-9, 4-10, 4-11, 4-12, 4-14, 4-15, 4-16, 4-17, 4-18, 4-19, 4-20, 5-6, 5-7, 5-8, 5-9, 5-10, 5-11, 5-12, 5-14, 5-15, 5-16, 5-17, 5-18, 5-19, 5-20, 6-7, 6-8, 6-9, 6-10, 6-11, 6-12, 6-14, 6-15, 6-16, 6-17, 6-18, 6-19, 6-20, 7-8, 7-9, 7-10, 7-11, 7-12, 7-14, 7-15, 7-16, 7-17, 7-18, 7-19, 7-20, 8-9, 8-10, 8-11, 8-12, 8-14, 8-15, 8-16, 8-17, 8-18, 8-19, 8-20, 9-10, 9-11, 9-12, 9-14, 9-15, 9-16, 9-17, 9-18, 9-19, 9-20, 10-11, 10-12, 10-14, 10-15, 10-16, 10-17, 10-18, 10-19, or 10-20). In some embodiments, m is 4 and n is 1-6. In some embodiments, combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of Y includes the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391. In some embodiments, the amino acid sequence of Y consists of [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, wherein combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of Y consists of the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

In some embodiments, X is absent, Z is absent, or X and Z are both absent.

In some embodiments, X, Y, or Z includes a bone-targeting moiety, e.g., including 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid. In some embodiments, the bone-targeting moiety includes or consists of E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

In some embodiments, X, Y, or Z includes a cathepsin (e.g., cathepsin K) cleavage sequence. In some embodiments, the cathepsin cleavage sequence includes or consists of HGPQG (SEQ ID NO: 374) or HKLRG (SEQ ID NO: 375).

In some embodiments, the polypeptide includes or consists of the amino acid sequence of any one of SEQ ID NOs: 501-608, e.g., SEQ ID NOs: 502, 504, 506, 512, 514, 516, 530, 560, 562, 564, 572, 574, 576, 584, 586, 588, 596, 598, 600, or 608. In some embodiments, the polypeptide includes a bone-targeting moiety, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 504.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 512.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 530.

5 In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 554.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 572.

10 In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 578.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 560.

In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 566.

15 In some embodiments, the polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 538 (e.g., where X in SEQ ID NO: 538 can be any amino acid, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu).

In some embodiments, the polypeptide includes the structure V-NP.

20 In some embodiments, any of the NPs or polypeptides described herein may be used in conjunction with the method (e.g., NPs or polypeptides described in any of the aspects described herein).

In some embodiments, the amino acid sequence of V or W includes  $[(\text{Gly})_m(\text{Ser})]_n(\text{Gly})_p$  or  $(\text{Gly})_p[(\text{Ser})(\text{Gly})_m]_n$ , wherein each of m, n, and p is, independently, between 0 and 20. In some embodiments, m is 4 and n is 1-6. In some embodiments, V is absent, W is absent, or V and W are both absent. In some embodiments, V and/or W includes a bone-targeting moiety, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>. In some embodiments, V and/or W includes a cathepsin (e.g., cathepsin K) cleavage sequence, e.g., HGPQG (SEQ ID NO: 374) or HKLRG (SEQ ID NO: 375).

30 In some embodiments of any of the aspects described herein, the polypeptide of the invention, e.g., the sALP polypeptide and/or the NP polypeptide, is in dimeric form. In some embodiments, the polypeptide is glycosylated or pegylated.

In some embodiments of any of the aspects described herein, the pharmaceutical composition is administered in a dosage between about 0.2 mg/kg to about 20 mg/kg of the polypeptide of the invention, e.g., the sALP polypeptide and/or the NP polypeptide, e.g., the sALP polypeptide. In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 0.2 mg/kg to about 20 mg/kg once, twice, three times, or four times daily. The dosage may be between, e.g., about 0.5 mg/kg to about 10 mg/kg, e.g., about 1 mg/kg to about 5 mg/kg, once, twice, three times, or four times

daily. In some embodiments, the dosage is about 1 mg/kg, about 2 mg/kg, or about 3 mg/kg once, twice, or three times daily. In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 0.2 mg/kg to about 20 mg/kg once, twice, three times, or four times weekly. The dosage may be between, e.g., about 0.5 mg/kg to about 10 mg/kg, c.g., about 1 mg/kg to 5 mg/kg, once, twice, three times, or four times weekly. In some embodiments, the dosage is about 1 mg/kg, about 2 mg/kg, or about 3 mg/kg once, twice, or three times weekly.

In some embodiments of any of the aspects described herein, the pharmaceutical composition is administered in a dosage between about 0.5 mg/kg to about 500 mg/kg of the polypeptide of the invention, e.g., the sALP polypeptide and/or the NP polypeptide, e.g., the NP polypeptide. In some 10 embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 0.5 mg/kg to about 500 mg/kg once, twice, three times, or four times daily. The dosage may be between, e.g., about 5 mg/kg to about 200 mg/kg, e.g., about 10 mg/kg to about 100 mg/kg, once, twice, three times, or four times daily. In some embodiments, the dosage is about 10 mg/kg or about 100 mg/kg twice daily. In some embodiments, the pharmaceutical composition is administered to the subject in a dosage 15 between about 0.5 mg/kg to about 500 mg/kg once, twice, three times, or four times weekly. The dosage may be between, e.g., about 5 mg/kg to about 200 mg/kg, e.g., about 10 mg/kg to about 100 mg/kg, e.g., about 20 mg/kg to about 40 mg/kg, once, twice, three times, or four times weekly. In some embodiments, the dosage is about 10 mg/kg, about 30 mg/kg, or about 100 mg/kg, once, twice, or three times weekly.

In some embodiments of any of the aspects described herein, the pharmaceutical composition is 20 administered in a dosage between about 10  $\mu$ g/kg to about 1,000  $\mu$ g/kg of the polypeptide of the invention, e.g., the sALP polypeptide and/or the NP polypeptide, e.g., the NP polypeptide. In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 10  $\mu$ g/kg to about 1,000  $\mu$ g/kg once, twice, three times, or four times weekly. The dosage may be between, e.g., about 20  $\mu$ g/kg to about 800  $\mu$ g/kg, e.g., about 30  $\mu$ g/kg to about 600  $\mu$ g/kg, e.g., about 50 25  $\mu$ g/kg to about 500  $\mu$ g/kg, e.g., about 100  $\mu$ g/kg to about 400  $\mu$ g/kg, e.g., about 200  $\mu$ g/kg to about 300  $\mu$ g/kg, once, twice, three times, or four times weekly. In some embodiments, the dosage is about 30  $\mu$ g/kg, about 100  $\mu$ g/kg, about 300  $\mu$ g/kg, or about 500  $\mu$ g/kg, once, twice, or three times weekly.

In some embodiments of any of the aspects described herein, the pharmaceutical composition is administered subcutaneously. In some embodiments, the pharmaceutical composition is administered one 30 time, two times, or three times per week.

In a third aspect, the invention features a composition including a first polypeptide and a second polypeptide, where a) the first polypeptide includes the structure A-sALP-B, where i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent or is an amino acid sequence of at least one amino acid, and iii) B is absent or is an amino acid sequence of at least one amino acid; and b) the 35 second polypeptide includes the structure V-NP-W, where i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii) V is absent or is an amino acid sequence of at least one amino acid, and iii) W is absent or is an amino acid sequence of at least one amino acid.

In some embodiments, the first polypeptide is any sALP polypeptide described herein, e.g., as described for the first aspect. In some embodiments, the amino acid sequence of the first polypeptide includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity of any one of SEQ ID NOS: 1201, 1204, 1220, or 1221, e.g., SEQ ID NO: 1204. In some embodiments, the 5 amino acid sequence of the first polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 1204.

In some embodiments, the second polypeptide is any NP polypeptide described herein, e.g., as described for the second aspect. In some embodiments, the amino acid sequence of the second polypeptide includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence 10 identity to any one of SEQ ID NOS: 504, 512, 530, 554, 572, or 578, e.g., SEQ ID NO: 512. In some embodiments, the amino acid sequence of the second polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 512.

In some embodiments, the amino acid sequence of the first polypeptide includes or consists of the amino acid sequence of SEQ ID NOS: 1204 or 1221, and the amino acid sequence of the second 15 polypeptide includes or consists of the amino acid sequence of SEQ ID NOS: 504, 512, 530, or 572.

In some embodiments, the first polypeptide and/or the second polypeptide are in dimeric form. In some embodiments, the first polypeptide and/or the second polypeptide are glycosylated or pegylated.

In some embodiments, the composition is a pharmaceutical composition including a pharmaceutically acceptable excipient, e.g., saline. In some embodiments, the composition is lyophilized.

20 In some embodiments, the first polypeptide is present in a dosage between about 0.2 mg/kg to about 20 mg/kg and the second polypeptide is present in a dosage between about 0.5 mg/kg to about 500 mg/kg. In some embodiments, the first polypeptide is present in a dosage between about 0.2 mg/kg to about 20 mg/kg, e.g., about 0.5 mg/kg to about 10 mg/kg, e.g., about 1 mg/kg to about 5 mg/kg, for once daily, twice daily, three times daily, four times daily, once weekly, twice weekly, three times weekly, or 25 four times weekly administration, e.g., about 1 mg/kg, about 2 mg/kg, or about 3 mg/kg for once daily, twice daily, four times daily, once weekly, twice weekly, or three times weekly administration; and the second polypeptide is present in a dosage between about 0.5 mg/kg to about 500 mg/kg, e.g., about 5 mg/kg to about 200 mg/kg, e.g., about 10 mg/kg to about 100 mg/kg, e.g., about 20 mg/kg to about 40 mg/kg, for once daily, twice daily, three times daily, four times daily, once weekly, twice weekly, three 30 times weekly, or four times weekly administration, e.g., about 10 mg/kg, about 30 mg/kg, or about 100 mg/kg, for once daily, twice daily, three times daily, once weekly, twice weekly, or three times weekly administration.

In some embodiments, the amino acid sequence of the first polypeptide includes the amino acid sequence of SEQ ID NO: 1204 and the amino acid sequence of the second polypeptide includes the 35 amino acid sequence of SEQ ID NO: 512.

In a fourth aspect, the invention features a method of treating a disease or a condition in a subject, the method including administering to the subject a therapeutically effective amount of a first polypeptide

and a second polypeptide, where a) the first polypeptide includes the structure A-sALP-B, where i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent or is an amino acid sequence of at least one amino acid, and iii) B is absent or is an amino acid sequence of at least one amino acid; and b) the second polypeptide includes the structure V-NP-W, where i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii) V is absent or is an amino acid sequence of at least one amino acid, and iii) W is absent or is an amino acid sequence of at least one amino acid; and the disease or the condition is selected from the group consisting of a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, a bone or cartilage disorder, a vascular smooth muscle disorder, and a condition for elongation of bone. In some embodiments, the disease or the condition in the subject 10 is thereby treated.

In some embodiments, the first polypeptide is any polypeptide described herein including sALP, e.g., any sALP polypeptide described herein, e.g., as described for the first or third aspect. In some embodiments, the amino acid sequence of the first polypeptide includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity of any one of SEQ ID NOs: 1201, 1204, 1220, or 15 1221, e.g., SEQ ID NO: 1204. In some embodiments, the amino acid sequence of the first polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 1204.

In some embodiments, the second polypeptide is any polypeptide described herein including NP, e.g., any NP polypeptide described herein, e.g., as described for the second or third aspect. In some embodiments, the amino acid sequence of the second polypeptide includes an amino acid sequence 20 having at least 80%, 85%, 90%, 95%, or 99% sequence identity to any one of SEQ ID NOs: 504, 512, 530, 554, 572, or 578, e.g., SEQ ID NO: 512. In some embodiments, the amino acid sequence of the second polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 512.

In some embodiments, the first polypeptide and the second polypeptide are administered within ten days, five days, or twenty-four hours of each other, e.g., within ten, nine, eight, seven, six, five, four, 25 three, or two days of each other or within twenty-four, twelve, eleven, ten, nine, eight, seven, six, five, four, three, two, or one hour(s) of each other.

In some embodiments, the first polypeptide and the second polypeptide are administered simultaneously. In some embodiments, the first polypeptide and the second polypeptide are formulated together in a composition or each separately in a composition. In some embodiments, the composition is 30 a pharmaceutical composition comprising a pharmaceutically acceptable excipient, e.g., saline. In some embodiments, the composition is lyophilized. In some embodiments, the composition is any composition, e.g., a pharmaceutical composition, described herein, e.g., as described for the first, second, or third aspect.

In a fifth aspect, the invention features a kit including: a) a first polypeptide including the 35 structure A-sALP-B, where i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent or is an amino acid sequence of at least one amino acid, and iii) B is absent or is an amino acid sequence of at least one amino acid; and b) instructions for administering the first polypeptide to a patient

diagnosed with or at risk of developing a neurocutaneous syndrome. In some embodiments, the first polypeptide is any polypeptide described herein including sALP, e.g., any sALP polypeptide described herein, e.g., as described in any of the above aspects.

In some embodiments, the kit further includes (c) a second polypeptide including the structure V-  
5 NP-W, where i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii)  
V is absent or is an amino acid sequence of at least one amino acid, and iii) W is absent or is an amino  
acid sequence of at least one amino acid. In some embodiments, the second polypeptide is any  
polypeptide described herein including NP, e.g., any NP polypeptide described herein, e.g., as described  
in any of the above aspects.

10 In a sixth aspect, the invention features a kit including: a) a polypeptide including the structure V-  
NP-W, where i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii)  
V is absent or is an amino acid sequence of at least one amino acid, and iii) W is absent or is an amino  
acid sequence of at least one amino acid; and b) instructions for administering the polypeptide to a patient  
diagnosed with or at risk of developing a neurocutaneous syndrome. In some embodiments, the  
15 polypeptide is any polypeptide described herein including NP, e.g., any NP polypeptide described herein,  
e.g., as described in any of the above aspects.

In a seventh aspect, the invention features a kit including: a) a first polypeptide including the  
structure A-sALP-B, where i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent  
or is an amino acid sequence of at least one amino acid, and iii) B is absent or is an amino acid sequence  
20 of at least one amino acid; and b) a second polypeptide including the structure V-NP-W, where i) NP is a  
natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii) V is absent or is an  
amino acid sequence of at least one amino acid, and iii) W is absent or is an amino acid sequence of at  
least one amino acid. In some embodiments, the first polypeptide is any polypeptide described herein  
including sALP, e.g., any sALP polypeptide described herein, e.g., as described in any of the above  
25 aspects; and the second polypeptide is any polypeptide described herein including NP, e.g., any NP  
polypeptide described herein, e.g., as described in any of the above aspects. In some embodiments, the  
first polypeptide and the second polypeptide are formulated together. In some embodiments, the first  
polypeptide and the second polypeptide are formulated separately and in individual dosage amount.

In any of the aspects described herein, the amino acid sequence of the sALP includes an amino  
30 acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to any one of SEQ ID  
NOs: 1202, 1205, 1218, or 1219.

In any of the aspects described herein, the amino acid sequence of the sALP includes or consists  
of an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to amino acid  
residues 23-508 of SEQ ID NO: 1215, amino acid residues 18-498 of SEQ ID NO: 1216, amino acid  
35 residues 23-508 of SEQ ID NO: 1218, amino acid residues 18-498 of SEQ ID NO: 1219, amino acid  
residues 23-512 of SEQ ID NO: 1215, amino acid residues 18-502 of SEQ ID NO: 1216, amino acid  
residues 23-512 of SEQ ID NO: 1218, or amino acid residues 18-502 of SEQ ID NO: 1219, where X in

SEQ ID NO: 1218 or 1219 is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP, e.g., not an amino acid corresponding to a pathogenic mutation provided in Table 1.

In any of the aspects described herein, the amino acid sequence of the NP includes or consists of 5 amino acid residues 6-22 of SEQ ID NO: 126, and the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Val, Ala, Ser, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe, Leu, Ile, Thr, Val, Ala, or Ser, e.g., e.g., Phe, Leu, Arg, Tyr, e.g., Phe or Leu, e.g., Phe, e.g., Leu.

In any of the aspects described herein, the amino acid sequence of the NP comprises or consists 10 of the amino acid sequence of any one of SEQ ID NOs: 13-29, 100-116, 119-125, 127-233, or 1001-1155.

In any of the aspects described herein, the NP is selective for NPR-B over NPR-A, where the EC<sub>50(NPR-A)</sub>/EC<sub>50(NPR-B)</sub> ratio for the NP, as determined in an *in vivo* pharmacokinetic assay, is at least 30, e.g., at least 35, 40, 35, 50, 55, or 60.

15 In any of the aspects described herein, the Fc includes a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region, or where the Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, and IgG-4, e.g., IgG-1. In some embodiments, the amino acid sequence of the Fc includes an amino acid sequence having at least 80%, 85%, 90%, 95%, or 99% sequence identity to SEQ ID NO: 401. In some embodiments, the amino acid sequence of the Fc includes or consists of SEQ ID 20 NO: 401.

In any of the aspects described herein, the amino acids sequence of D or Y includes a glycine-rich region, or the amino acid sequence of D or Y consists of one or more glycines and one or more serines. For example, the amino acid sequence of D or Y may include or consist of [(Gly)<sub>m</sub>(Ser)]<sub>n</sub>(Gly)<sub>p</sub> or (Gly)<sub>p</sub>[(Ser)(Gly)<sub>m</sub>]<sub>n</sub>, where each of m, n, and p is, independently, between 0 and 20. In some 25 embodiments, m is between 1 and 6; n is between 1 and 10; and p is between 0 and 4, e.g., m is 4 and n is 1-6. In some embodiments, combinations of m, n, and p are selected from a single row of Table 2, or the amino acid sequence of D or Y includes or consists of the amino acid sequence of any one of SEQ ID NOs: 301-391.

In any of the aspects described herein, the amino acids sequence of D, G, or Y is optionally two 30 amino acid residues (e.g., leucine-lysine or aspartic acid-isoleucine). In some embodiments of an sALP polypeptide, C and E may both be absent, and D may be absent or may be an amino acid sequence of at least one amino acid. For example, the polypeptide may consist of the structure sALP-D-Fc or the structure Fc-D-sALP, where D may be any linker described herein or may optionally consist of two amino acid residues, e.g., leucine-lysine. For example, the polypeptide may consist of the structure 35 sALP-D-Fc. Optionally, the amino acid sequence of sALP is the amino acid sequence of SEQ ID NO: 1205, the amino acid sequence of D is leucine-lysine, and/or the amino acid sequence of Fc is the amino acid sequence of SEQ ID NO: 401. In other embodiments of an sALP polypeptide, C may be absent, and

D and E may both be absent or may be an amino acid sequence of at least one amino acid. For example, the polypeptide may include the structure sALP-D-Fc-E or the structure Fc-D-sALP-E. In other embodiments of an sALP polypeptide, C may be absent, and D, G, and H may all be absent or may be an amino acid sequence of at least one amino acid. For example, the polypeptide may include the structure 5 sALP-D-Fc-G-I<sub>n</sub>-H. In some embodiments of an NP polypeptide, X and Z may both be absent, and Y may be absent or may be an amino acid sequence of at least one amino acid. For example, the polypeptide may consist of the structure NP-Y-Fc or the structure Fc-Y-NP, where Y may be any linker described herein or may optionally consist of two amino acid residues, e.g., leucine-lysine. For example, the polypeptide may consist of the structure NP-Y-Fc.

10 In any of the aspects described herein, one or more of A, B, C, D, E, G, H, V, W, X, Y, or Z are absent. In some embodiments, A is absent, B is absent, or A and B are both absent. In some embodiments, C is absent, E is absent, or C and E are both absent. In some embodiments, C is absent, H is absent, or C and H are both absent. In some embodiments, X is absent, Z is absent, or X and Z are both absent.

15 In any of the aspects described herein, the polypeptide of the invention, e.g., the sALP polypeptide and/or the NP polypeptide, may include a bone-targeting moiety, e.g., including 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>. In some embodiments of any of the above aspects, one or more of A, B, C, D, E, V, W, X, Y, or Z includes a bone-targeting moiety, e.g., including 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, or 16 consecutive acidic residues, e.g., aspartic acid or glutamic acid, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>, e.g., E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

20 In any of the aspects described herein, the polypeptide of the invention optionally does not include a bone-targeting moiety (e.g., a polyaspartic acid or polyglutamic acid region longer than two consecutive aspartic acid or glutamic acid residues).

25 In any of the aspects described herein, the polypeptide of the invention may include a cathepsin (e.g., cathepsin K) cleavage sequence, e.g., HGPQG (SEQ ID NO: 374) or HKLRG (SEQ ID NO: 375). In some embodiments of any of the above aspects, A, B, C, D, E, G, H, V, W, X, Y, or Z includes a cathepsin (e.g., cathepsin K) cleavage sequence.

30 In some embodiments of any of the above aspects, the polypeptide of the invention may include a polypeptide having reduced (e.g., by about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, or about 100%) degradation (e.g., by neutral endopeptidase (NEP), insulin degrading enzyme (IDE), or any other enzyme that cleaves a 35 natriuretic peptide *in vivo*), as compared to a control (e.g., CNP22, CNP53, or any polypeptide described herein, such as a peptide described in International Application Pub. No. WO2010/135541 or U.S. Application Pub. No. 2010-0331256).

In some embodiments of any of the above aspects, the polypeptide of the invention may have increased (e.g., by about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, about 100%, or more) efficacy and/or reduced (e.g., by about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, or about 100%) dose-dependent side effects (e.g., decreased adverse hemodynamic effects, such as decreased lowering of blood pressure), as compared to a control (e.g., any polypeptide described herein, such as a peptide described in International Application Pub. No. 10 WO2010/135541 or U.S. Application Pub. No. 2010-0331256).

In any of the aspects described herein, the polypeptide of the invention (e.g., an sALP polypeptide or an NP polypeptide) is glycosylated or pegylated. In some embodiments, the pharmaceutical composition includes a dimer of one or more of the polypeptides of the invention. In some embodiments, the pharmaceutically acceptable excipient includes saline. In some embodiments, the pharmaceutical composition is lyophilized. In some embodiments, the pharmaceutical composition is administered subcutaneously, intravenously, orally, nasally, intramuscularly, sublingually, intrathecally, or intradermally, e.g., subcutaneously.

In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 0.5 mg/kg to about 500 mg/kg once, twice, three times, or four times daily. The dosage may be between, e.g., about 5 mg/kg to about 200 mg/kg, e.g., about 10 mg/kg to about 100 mg/kg, once, twice, three times, or four times daily. In some embodiments, the dosage is about 10 mg/kg or about 100 mg/kg twice daily.

In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 0.5 mg/kg to about 500 mg/kg once or twice weekly. The dosage may be between, e.g., about 5 mg/kg to about 200 mg/kg, e.g., about 10 mg/kg to about 100 mg/kg, e.g., about 20 mg/kg to about 40 mg/kg, once or twice weekly. In some embodiments, the dosage is about 10 mg/kg, about 30 mg/kg, or about 100 mg/kg, once or twice weekly.

In some embodiments, the pharmaceutical composition is administered to the subject in a dosage between about 10  $\mu$ g/kg to about 1,000  $\mu$ g/kg once or twice weekly. The dosage may be between, e.g., about 20  $\mu$ g/kg to about 800  $\mu$ g/kg, e.g., about 30  $\mu$ g/kg to about 600  $\mu$ g/kg, e.g., about 50  $\mu$ g/kg to about 500  $\mu$ g/kg, e.g., about 100  $\mu$ g/kg to about 400  $\mu$ g/kg, e.g., about 200  $\mu$ g/kg to about 300  $\mu$ g/kg, once or twice weekly. In some embodiments, the dosage is about 30  $\mu$ g/kg, about 100  $\mu$ g/kg, about 300  $\mu$ g/kg, or about 500  $\mu$ g/kg, once or twice weekly.

In any of the aspects described herein, the pharmaceutical composition is administered to the subject in a dosage between about 0.1 mg/kg to about 500 mg/kg of one or more of the polypeptides of the invention in a dosage regimen of once, twice, three times, or four times daily. The dosage may be between, e.g., about 1 mg/kg to about 200 mg/kg, e.g., about 2 mg/kg to about 100 mg/kg or about 10

mg/kg to about 100 mg/kg of one or more of the polypeptides of the invention in a dosage regimen of once, twice, three times, or four times daily. In some embodiments, the dosage is about 10 mg/kg or about 100 mg/kg of one or more of the polypeptides of the invention in a dosage regimen of twice daily. For example, the methods of the invention may optionally include administering a pharmaceutical composition including the polypeptides of the invention (e.g., an sALP polypeptide and/or an NP polypeptide) to the subject in a dosage of about 0.5 mg/kg/day to about 10 mg/kg/day (e.g., about 2 mg/kg/day to about 3 mg/kg/day).

5 In any of the aspects described herein, the pharmaceutical composition is administered to the subject between one and fourteen times a week, or is administered at least once daily for at least one month. In some embodiments, the pharmaceutical composition is administered to the subject once weekly for at least one month. In some embodiments, the pharmaceutical composition is administered one time, two times, three times, or four times a week, e.g., three times a week.

10 Any of the pharmaceutical compositions of the invention may optionally be formulated for treating a disease or condition (e.g., any described herein) in a subject.

15 Any of the pharmaceutical composition of the invention featuring an isolated nucleic acid may optionally include a recombinant expression vector (e.g., a lentiviral vector) including the isolated nucleic acid. In some embodiments, the pharmaceutical composition includes about 0.1 mg to about 10 mg of the isolated nucleic acid.

20 In any embodiment described herein, the polypeptide may or may not be isolated.

25 In any embodiment described herein, the subject may be human.

In any of the aspects described herein, the combination of two or more polypeptides of the invention (e.g., any sALP polypeptide and/or any NP polypeptide described herein) or two or more compositions of the invention provides a synergistic effect. In particular embodiments, the synergistic effect is a therapeutic effect that is observed for the combination of two or more polypeptides of the invention, wherein one or more of the polypeptides of the invention is present at a dose that is normally non-therapeutic; or a therapeutic effect that results in an unexpected decrease in one or more adverse events (e.g., hemodynamic effects, such as a decrease in blood pressure, such as systolic arterial blood pressure, diastolic arterial blood pressure, or mean arterial blood pressure, that results in adverse hypotensive effects); or a therapeutic effect that results in reduced dose-dependent side effects, as compared to the level of dose-dependent side effects observed for a single polypeptide of the invention at a therapeutic dose.

30 In some embodiments of any of the methods described herein, the disease is the neurocutaneous syndrome (e.g., neurofibromatosis (e.g., classic von Recklinghausen type (type I), either with gastrointestinal stromal tumors (i.e., as in intestinal neurofibromatosis (type 3B)) or without such tumors; an acoustic neuroma type (type II); a mixed type that combines the features of types I and II with predominant features, such as bilateral acoustic neuromas, posterior fossa and upper cervical meningiomas, and spinal/paraspinal neurofibromas (type III, Riccardi type or type 3A); an atypical type

that is distinguished from by the lack of iris Lisch nodules that are characteristic of type I (type VI); segmental neurofibromatosis, which is a variant of type I having lesions affecting a specific area of the body, such as a single segment of the body or an area that crosses the midline (type V); a type having only the symptoms of café au lait spots without other manifestations of neurofibromatosis (type VI);

5 familial spinal neurofibromatosis, which is caused by mutation in the neurofibromin gene NF1 and considered a distinguishable variant of type I; other variants of type I, such as neurofibromatosis-pheochromocytoma-duodenal carcinoid syndrome; neurofibromatosis with manifestations of Noonan syndrome, such as short stature, ptosis, midface hypoplasia, webbed neck, learning disabilities, and muscle weakness; and schwannomatosis, where any of these disorders can include or exclude one or

10 more bone manifestations); tuberous sclerosis; Sturge-Weber disease; ataxia telangiectasia; von Hippel-Lindau disease; incontinentia pigmenti; epidermal nevus syndromes, such as linear sebaceous nevus of Jadassohn; nevoid basal cell carcinoma syndrome; hypomelanosis of Ito; neurocutaneous melanosis; Klippel-Trenaunay syndrome; and Waardenburg syndrome, including types I, II, III, and IV). In some embodiments, the neurocutaneous syndrome has one or more bone manifestations. In some

15 embodiments, the neurocutaneous syndrome is neurofibromatosis type I. In some embodiments of any of the methods described herein, the disease is any syndrome (e.g., a neurocutaneous syndrome) with overactivated RAS and/or ERK signaling (e.g., Noonan syndrome, Costello syndrome, Noonan syndrome with multiple lentigines/LEOPARD syndrome, neurofibromatosis type 1, hereditary gingival fibromatosis type 1, NF1-Noonan syndrome, capillary malformation-AV malformation syndrome, Legius syndrome,

20 Noonan syndrome-like disorder with loose anagen hair, Noonan syndrome-like disorder with juvenile myelomonocytic leukemia (JMML), cardio-facio-cutaneous syndrome, or autoimmune lymphoproliferative syndrome, where any of these disorders can include or exclude one or more bone manifestations). In some embodiments of any of the methods described herein, the disorder associated with overactivation of FGFR3 is a bone or cartilage disorder, e.g., a skeletal dysplasia, such as any

25 described herein, e.g., achondroplasia or craniosynostosis. In some embodiments of any of the methods described herein, the disorder is a bone or cartilage disorder, e.g., a skeletal dysplasia, such as any described herein. In some embodiments, the bone or cartilage disorder is a skeletal dysplasia, e.g., achondroplasia, homozygous achondroplasia, heterozygous achondroplasia, achondrogenesis, acrodysostosis, acromesomelic dysplasia, atelosteogenesis, camptomelic dysplasia, chondrodysplasia

30 punctata, rhizomelic type of chondrodysplasia punctata, cleidocranial dysostosis, congenital short femur, craniosynostosis (e.g., Muenke syndrome, Crouzon syndrome, Apert syndrome, Jackson-Weiss syndrome, Pfeiffer syndrome, or Crouzonodermoskeletal syndrome), dactyly, brachydactyly, camptodactyly, polydactyly, syndactyly, diastrophic dysplasia, dwarfism, dyssegmental dysplasia, enchondromatosis, fibrochondrogenesis, fibrous dysplasia, hereditary multiple exostoses,

35 hypochondroplasia, hypophosphatasia, hypophosphatemic rickets, Jaffe-Lichtenstein syndrome, Kniest dysplasia, Kniest syndrome, Langer-type mesomelic dysplasia, Marfan syndrome, McCune-Albright syndrome, micromelia, metaphyseal dysplasia, Jansen-type metaphyseal dysplasia, metatrophic dysplasia,

Morquio syndrome, Nievergelt-type mesomelic dysplasia, neurofibromatosis (e.g., type 1, e.g., with bone manifestations or without bone manifestations; type 2; schwannomatosis; or any described herein), osteoarthritis, osteochondrodysplasia, osteogenesis imperfecta, perinatal lethal type of osteogenesis imperfecta, osteopetrosis, osteopoikilosis, peripheral dysostosis, Reinhardt syndrome, Roberts syndrome, 5 Robinow syndrome, short-rib polydactyly syndromes, short stature, spondyloepiphyseal dysplasia congenita, spondyloepimetaphyseal dysplasia, or thanatophoric dysplasia. In some embodiments, the bone or cartilage disorder is optionally hypophosphatasia (e.g., infantile HPP, childhood HPP, perinatal HPP, adult HPP, or odontohypophosphatasia). In some embodiments, the pharmaceutical composition is administered in an amount that is therapeutically effective to treat an achondroplasia phenotype selected 10 from the group consisting of growth retardation, skull deformities, and orthodontic defects. In some embodiments, the pharmaceutical composition is administered in an amount that is therapeutically effective to treat an achondroplasia phenotype selected from the group consisting of cervical cord compression, spinal stenosis, hydrocephalus, hearing loss due to chronic otitis, cardiovascular disease, neurological disease, and obesity. In some embodiments of any of the methods described herein, the 15 disorder associated with overactivation of FGFR3 is cancer, e.g., multiple myeloma, myeloproliferative syndrome, leukemia, plasma cell leukemia, lymphoma, glioblastoma, prostate cancer, bladder cancer, or mammary cancer. In some embodiments of any of the methods described herein, the vascular smooth muscle disorder is hypertension, restenosis, arteriosclerosis, acute decompensated heart failure, congestive heart failure, cardiac edema, nephredema, hepatic edema, acute renal insufficiency, or chronic 20 renal insufficiency. In some embodiments of any of the methods described herein, the condition for elongation of bone is insufficient or impaired bone growth arising from fractures, renal failure or insufficiency, poor diet, vitamin deficiency, hormone deficiency, or any skeletal dysplasia described herein.

In any method described herein, the neurocutaneous syndrome, disorder associated with 25 overactivation of FGFR3, bone or cartilage disorder, vascular smooth muscle disorder, or condition for elongation of bone in the subject is thereby treated.

In some embodiments of any of the methods described herein, one or more polypeptides of the invention or one or more compositions of the invention is optionally administered in an amount that is therapeutically effective to treat a HPP phenotype selected from the group consisting of HPP-related 30 seizure, premature loss of deciduous teeth, incomplete bone mineralization, elevated blood and/or urine levels of inorganic pyrophosphate (PP<sub>i</sub>), elevated blood and/or urine levels of phosphoethanolamine (PEA), elevated blood and/or urine levels of pyridoxal 5'-phosphate (PLP), inadequate weight gain, rickets, bone pain, calcium pyrophosphate dihydrate crystal deposition, aplasia, hypoplasia, and dysplasia of the dental cementum. In some embodiments, the incomplete bone mineralization is incomplete 35 femoral bone mineralization, incomplete tibial bone mineralization, incomplete metatarsal bone mineralization, or incomplete rib bone mineralization.

*Definitions*

As used herein, the term “about” means  $\pm 10\%$  of the recited value.

By “area under the curve” or “AUC” in the context of an *in vivo* pharmacokinetic assay is meant the area under the serum concentration vs. time curve after administration in an animal.

5 By “bone or cartilage disorder” is meant any disorder, disease, or other abnormality that affects the function, structure, or growth of bone or cartilage.

By “bone-targeting moiety” is meant an amino acid sequence of between 6 and 20 amino acid residues in length having a sufficient affinity to the bone matrix such that the bone-targeting moiety, taken alone, has an *in vivo* binding affinity to the bone matrix that is at least  $10^{-6}$  M or better (e.g.,  $10^{-7}$  M, 10 $^{-8}$  M, 10 $^{-9}$  M, or better).

10 By “cathepsin cleavage sequence” is meant an amino acid sequence having a site that can be cleaved by cathepsin with a  $k_{cat}/K_M$  rate constant of at least  $10^3$  M $^{-1}$ s $^{-1}$  (e.g.,  $10^4$  M $^{-1}$ s $^{-1}$ ,  $10^5$  M $^{-1}$ s $^{-1}$ ,  $10^6$  M $^{-1}$ s $^{-1}$ ,  $10^7$  M $^{-1}$ s $^{-1}$ , or  $10^8$  M $^{-1}$ s $^{-1}$ ) at 30°C or higher (e.g., 37°C). In particular embodiments, the cathepsin cleavage sequence is specific for cathepsin K. Exemplary cathepsin cleavage sequences are P2-P1-P1’, where cleavage by the enzyme would occur at the P1-P1’ peptide bond; P2 is preferentially composed of Pro, Leu, Ile, but could also be Val, Norleucine, Met, or Ala; P1 is preferentially Arg, Lys, Gln, but could also be Met, Norleucine, Leu, Ile, or Thr; and P1’ can be any amino acid but is preferentially Gly.

15 Additional cathepsin cleavage sequences are provided in Choe et al., J. Biol. Chem. 281(18):12824-832, 2006, which is incorporated herein by reference.

20 By “CNP22” is meant human CNP22 (SEQ ID NO: 4), unless a different meaning is expressly indicated.

By “CNP53” is meant human CNP53 (SEQ ID NO: 11), unless a different meaning is expressly indicated.

25 By “condition for elongation of bone” is meant any disorder, disease, or other abnormality that would benefit from lengthening of one or more segments of bone. After administration of any polypeptide described herein, the lengthening of one or more segments of bone can be increased by more than about 1%, about 2%, about 5%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, or about 30 100%, or more.

By “disorder associated with overactivation of FGFR3” is meant any disorder, disease, or other abnormality that is caused by, or is associated with, overactivation of FGFR3, e.g., stemming from a gain-of-function FGFR3 mutation.

By “efficacy” is meant the  $E_{max}$  value of a compound in a dose-response assay.

35 By “Fc” is meant a fragment crystallizable region of an immunoglobulin, e.g., IgG-1, IgG-2, IgG-3, IgG-3 or IgG-4, including the C<sub>H2</sub> and C<sub>H3</sub> domains of the immunoglobulin heavy chain. Fc may also include any portion of the hinge region joining the Fab and Fc regions. The Fc can be of any mammal,

including human, and may be post-translationally modified (e.g., by glycosylation). In a non-limiting example, Fc can be the fragment crystallizable region of human IgG-1 having the amino acid sequence of SEQ ID NO: 401.

By "fragment" is meant a portion of a polypeptide or nucleic acid molecule that contains, 5 preferably, at least 10%, 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or more of the entire length of the reference nucleic acid molecule or polypeptide. A fragment may contain, e.g., 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, 200, 210, 220, 230, 240, 250, 260, 270, 280, 290, 300, 310, 320, 330, 340, 350, 360, 370, 380, 390, 400, 500, 600, 700, 800, 900, 1,000, 1100, 1200, 1300, 1400, 1500, 1600, 1700, 10 1800, 1900, 2000, 2100, or more nucleotides, up to the entire length of the nucleic acid molecule, or 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, 200, 210, 220, 230, 240, 250, 260, 270, 280, 290, 300, 400, 500, 600, 700, or more amino acid residues, up to the entire length of the polypeptide. Exemplary sALP fragments have amino acid residues 18-498, 18-499, 18-500, 18-501, 18-502, 18-503, 18-504, 18-505, 18-506, 18-507, 18-508, 18-509, 18- 15 510, 18-511, or 18-512 of a consensus sequence for ALP (e.g., SEQ ID NOs: 1215, 1216, 1218, or 1219), and may include additional N-terminal and/or C-terminal portions. Exemplary NP fragments have at least a consensus ring domain, e.g., of SEQ ID NOs: 6, 30, or 95, and may include additional N-terminal and/or C-terminal portions.

By "homolog" is meant a polypeptide or nucleic acid molecule exhibiting at least 50% identity to 20 a reference amino acid sequence or nucleic acid sequence. Such a sequence is generally at least, e.g., 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% identical at the amino acid level or nucleic acid to a reference sequence. In general, for polypeptides, the length of comparison sequences can be at least five amino acid residues, e.g., 10, 20, 30, 40, 50, 60, 70, 80, 90, 100, 125, 150, 175, 200, 250, 300, 400, 500, 600, 700, or more amino acid residues, up to the entire length of the polypeptide. For 25 nucleic acids, the length of comparison sequences can generally be at least 10, 20, 30, 40, 50, 60, 70, 80, 90, 100, 125, 150, 175, 200, 250, 300, 400, 500, 600, 700, 800, 900, 1000, 1100, 1200, 1300, 1400, 1500, 1600, 1700, 1800, 1900, 2000, 2100, or more nucleotides, up to the entire length of the nucleic acid molecule. It is understood that for the purposes of determining sequence identity when comparing a DNA sequence to an RNA sequence, a thymine nucleotide is equivalent to a uracil nucleotide.

30 As used herein, when a polypeptide or nucleic acid sequence is referred to as having "at least X% sequence identity" to a reference sequence, it is meant that at least X percent of the amino acid residues or nucleotides in the polypeptide or nucleic acid are identical to those of the reference sequence when the sequences are optimally aligned. An optimal alignment of sequences can be determined in various ways that are within the skill in the art, for instance, the Smith Waterman alignment algorithm (Smith et al., J. 35 Mol. Biol. 147:195-7, 1981) and BLAST (Basic Local Alignment Search Tool; Altschul et al., J. Mol. Biol. 215: 403-10, 1990). These and other alignment algorithms are accessible using publicly available computer software such as "Best Fit" (Smith and Waterman, Advances in Applied Mathematics, 482-489,

1981) as incorporated into GeneMatcher PlusTM (Schwarz and Dayhof, *Atlas of Protein Sequence and Structure*, Dayhoff, M.O., Ed pp 353-358, 1979), BLAST, BLAST-2, BLAST-P, BLAST-N, BLAST-X, WU-BLAST-2, ALIGN, ALIGN-2, CLUSTAL, or Megalign (DNASTAR). In addition, those skilled in the art can determine appropriate parameters for measuring alignment, including any algorithms needed 5 to achieve optimal alignment over the length of the sequences being compared.

By "hybridize" is meant to pair to form a double-stranded molecule between complementary polynucleotides, or portions thereof, under various conditions of stringency. (See, e.g., Wahl, G. M. and S. L. Berger (1987) *Methods Enzymol.* 152:399; Kimmel, A. R. (1987) *Methods Enzymol.* 152:507.) For example, high stringency salt concentration will ordinarily be less than about 750 mM NaCl and 75 mM 10 trisodium citrate, less than about 500 mM NaCl and 50 mM trisodium citrate, or less than about 250 mM NaCl and 25 mM trisodium citrate. Low stringency hybridization can be obtained in the absence of organic solvent, e.g., formamide, while high stringency hybridization can be obtained in the presence of at least about 35% formamide or at least about 50% formamide. High stringency temperature conditions will ordinarily include temperatures of at least about 30°C, 37°C, or 42°C. Varying additional 15 parameters, such as hybridization time, the concentration of detergent, e.g., sodium dodecyl sulfate (SDS), and the inclusion or exclusion of carrier DNA, are well known to those skilled in the art. Various levels of stringency are accomplished by combining these various conditions as needed. In one embodiment, hybridization will occur at 30°C in 750 mM NaCl, 75 mM trisodium citrate, and 1% SDS. In an alternative embodiment, hybridization will occur at 37°C in 500 mM NaCl, 50 mM trisodium 20 citrate, 1% SDS, 35% formamide, and 100 µg/ml denatured salmon sperm DNA (ssDNA). In a further alternative embodiment, hybridization will occur at 42°C in 250 mM NaCl, 25 mM trisodium citrate, 1% SDS, 50% formamide, and 200 µg/ml ssDNA. Useful variations on these conditions will be readily apparent to those skilled in the art.

For most applications, washing steps that follow hybridization will also vary in stringency. Wash 25 stringency conditions can be defined by salt concentration and by temperature. As above, wash stringency can be increased by decreasing salt concentration or by increasing temperature. For example, high stringency salt concentrations for the wash steps may be, e.g., less than about 30 mM NaCl and 3 mM trisodium citrate, or less than about 15 mM NaCl and 1.5 mM trisodium citrate. High stringency temperature conditions for the wash steps will ordinarily include a temperature of, e.g., at least about 25°C, 42°C, or 68°C. In one embodiment, wash steps will occur at 25°C in 30 mM NaCl, 3 mM 30 trisodium citrate, and 0.1% SDS. In an alternative embodiment, wash steps will occur at 42°C in 15 mM NaCl, 1.5 mM trisodium citrate, and 0.1% SDS. In a further alternative embodiment, wash steps will occur at 68°C in 15 mM NaCl, 1.5 mM trisodium citrate, and 0.1% SDS. Additional variations on these conditions will be readily apparent to those skilled in the art. Hybridization techniques are well known to those skilled in the art and are described, for example, in Benton and Davis (*Science* 196:180, 1977); Grunstein and Hogness (*Proc. Natl. Acad. Sci., USA* 72:3961, 1975); Ausubel et al. (*Current Protocols in Molecular Biology*, Wiley Interscience, New York, 2001); Berger and Kimmel (*Guide to Molecular*

Cloning Techniques, 1987, Academic Press, New York); and Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, New York.

By “isolated” or “purified” is meant separated from other naturally accompanying components. Typically, a compound (e.g., polypeptide, nucleic acid, or small molecule), factor, cell, or other component is considered isolated when it is at least, e.g., 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, or even 99%, by weight, free from proteins, antibodies, naturally-occurring organic molecules, and other components with which it is naturally associated. In some instances, the component is at least 75%, 90%, or even 99%, by weight, pure. An isolated component may be obtained by chemical synthesis, separation of the factor from natural sources, or production of the component in a recombinant host cell that does not naturally produce the component. Proteins and small molecules may be purified by one skilled in the art using standard techniques such as those described by Ausubel *et al.* (Current Protocols in Molecular Biology, John Wiley & Sons, New York, 2000). The component is preferably at least, e.g., 2, 5, or 10 times as pure as the starting material, as measured using, e.g., polyacrylamide gel electrophoresis, column chromatography, optical density, HPLC analysis, or Western analysis (Ausubel *et al.*, *supra*). Exemplary methods of purification are column chromatography, immunoprecipitation, and magnetic bead immunoaffinity purification.

By “natriuretic peptide that is an agonist of natriuretic peptide receptor B” (abbreviated “NP”) is meant a natriuretic peptide as described herein, e.g., human CNP22 (SEQ ID NO: 4), or variant thereof, that is capable of agonizing NPR-B, e.g., human NPR-B, with at least 0.000001, 0.000005, 0.00001, 0.00005, 0.0001, 0.0005, 0.001, 0.005, 0.01, 0.05, 0.1, 0.5, 0.9, or 1 times the potency, and at least 1%, 2%, 5%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 95%, or even 100% times the efficacy of CNP22 as measured in a standard NPR-B activation assay, e.g., a membrane assay or whole cell assay, as described herein. Variant NPs may include one or more substitutions, additions or deletions relative to CNP22 and have the ability to agonize NPR-B. An NP as described herein may include any other sequence or moiety, attached covalently or non-covalently, provided that the NP has the ability to agonize NPR-B.

By “neurocutaneous syndrome” is meant a neurological disorder with one or more cutaneous manifestations, such as lesions on the skin and/or the eye. Such syndromes can optionally be accompanied by benign or malignant tumors in multiple sites of the body.

By “NP polypeptide” is meant any sequence including an NP sequence, as defined herein. Exemplary NP polypeptides include those having the structure V-NP-W, wherein each of V and W is absent or is an amino acid sequence of at least one amino acid (e.g., any NP fusion polypeptide described herein).

By “nucleic acid molecule” is meant a molecule, e.g., RNA or DNA, having a sequence of two or more covalently bonded, naturally occurring or modified nucleotides. The nucleic acid molecule may be, e.g., single or double stranded, and may include modified or unmodified nucleotides, or mixtures or combinations thereof. Various salts, mixed salts, and free acid forms are also included.

The terms “peptide,” “polypeptide,” and “protein” are used interchangeably and refer to any chain of two or more natural or unnatural amino acid residues, regardless of post-translational modification (e.g., glycosylation or phosphorylation), constituting all or part of a naturally-occurring or non-naturally occurring polypeptide or peptide, as is described herein.

5 As used herein, a natural amino acid is a natural  $\alpha$ -amino acid having the L-configuration, such as those normally occurring in natural polypeptides. Unnatural amino acid refers to an amino acid that normally does not occur in polypeptides, e.g., an epimer of a natural  $\alpha$ -amino acid having the L configuration, that is to say an amino acid having the unnatural D-configuration; or a (D,L)-isomeric mixture thereof; or a homolog of such an amino acid, for example, a  $\beta$ -amino acid, an  $\alpha,\alpha$ -disubstituted 10 amino acid, or an  $\alpha$ -amino acid wherein the amino acid side chain has been shortened by one or two methylene groups or lengthened to up to 10 carbon atoms, such as an  $\alpha$ -amino alcanoic acid with 5 up to and including 10 carbon atoms in a linear chain, an unsubstituted or substituted aromatic ( $\alpha$ -aryl or  $\alpha$ -aryl lower alkyl), for example, a substituted phenylalanine or phenylglycine.

15 By “pharmaceutically acceptable carrier” or “pharmaceutically acceptable excipient” is meant a carrier or excipient that is physiologically acceptable to the treated patient while retaining the therapeutic properties of the compound with which it is administered. One exemplary pharmaceutically acceptable carrier substance is physiological saline. Other physiologically acceptable carriers and their formulations are known to those skilled in the art and described, for example, in Remington’s Pharmaceutical Sciences, (20th edition), ed. A. Gennaro, 2000, Lippincott, Williams & Wilkins, Philadelphia, PA.

20 By “pharmaceutical composition” is meant a composition containing a polypeptide or nucleic acid molecule as described herein formulated with a pharmaceutically acceptable excipient, and manufactured or sold with the approval of a governmental regulatory agency as part of a therapeutic regimen for the treatment or prevention of a disease or event in a subject. Pharmaceutical compositions can be formulated, for example, for subcutaneous administration, intravenous administration (e.g., as a 25 sterile solution free of particulate emboli and in a solvent system suitable for intravenous use), for oral administration (e.g., a tablet, capsule, caplet, gelcap, or syrup), or any other formulation described herein, e.g., in unit dosage form.

30 By “potency” is meant the reciprocal of the  $EC_{50}$  value of a compound in a dose-response assay. When comparing potency between a compound and a control or between an assay and a control assay, decreased potency indicates an increased  $EC_{50}$  value, and increased potency indicates a decreased  $EC_{50}$  value, as compared to the  $EC_{50}$  value for the control or the control assay.

35 By “reduced degradation” is meant having a lower percentage of degraded peptide after exposure to an enzyme for at least 5, 10, 15, 20, 25, 30, 60, 120, 180, or 240 minutes, or higher, or any range between any two of these values, as compared to a percentage of degraded control, such as CNP22, CNP53, or any polypeptide described herein, such as a peptide described in International Application Pub. No. WO2010/135541 or U.S. Application Pub. No. 2010-0331256. The percentage of degraded peptide can be lower by about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about

50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, or about 100%, where the percentage of degraded peptide can be determined by measuring the percentage of degraded peptide directly or indirectly by measuring the percentage of remaining peptide after exposure to an enzyme (e.g., neutral 5 endopeptidase, insulin degrading enzyme, and any other enzyme that cleaves a natriuretic peptide *in vivo*) and subtracting this percentage of remaining peptide from 100%. Percentage of degraded peptide or remaining peptide can be measured by any useful method, such as liquid chromatography (e.g., high performance liquid chromatography (HPLC)), mass spectrometry (MS), or combined analytic techniques (e.g., LC-MS).

10 By “reduced dose-dependent side effect” is meant a decrease in one or more adverse effects as a function of a dosage of a compound, as compared to a control (e.g., any polypeptide described herein, such as a peptide described in International Application Pub. No. WO2010/135541 or U.S. Application Pub. No. 2010-0331256). The decrease in one or more adverse effects can be by about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 15 70%, about 75%, about 80%, about 85%, about 90%, about 95%, about 96%, about 97%, about 98%, about 99%, or about 100%, as determined by any useful assay for detecting the adverse effect.

Exemplary adverse effects include hemodynamic effects, such as a decrease in blood pressure, such as systolic arterial blood pressure, diastolic arterial blood pressure, or mean arterial blood pressure, that results in adverse hypotensive effects, and assays to detect such hemodynamic effects include a 20 sphygmomanometer or an implanted pressure transducer.

The terms “sALP,” “soluble alkaline phosphatase,” and “extracellular domain of an alkaline phosphatase” are used interchangeably and mean a soluble, non-membrane-bound alkaline phosphatase or a domain, biologically active fragment, or biologically active variant thereof. sALPs include, for example, an alkaline phosphatase lacking a C-terminal GPI signal sequence, e.g., a polypeptide including 25 or consisting of the amino acid residues 18-502 of human TNALP (SEQ ID NO: 1208). sALPs further include, for example, soluble, non-membrane-bound forms of mammalian orthologs of human TNALP (e.g., polypeptides including or consisting of amino acid residues 16-502 or 18-502 of SEQ ID NO: 1206, amino acid residues 18-502 of SEQ ID NO: 1207, amino acid residues 18-502 of SEQ ID NO: 1209, amino acid residues 18-502 of SEQ ID NO: 1210, or amino acid residues 1-480 of SEQ ID NO: 1211), 30 soluble, non-membrane-bound forms of human IALP, GALP, and PLALP (e.g., polypeptides including or consisting of amino acid residues 20-503 of SEQ ID NO: 1212, amino acid residues 20-503 of SEQ ID NO: 1213, or amino acid residues 23-506 of SEQ ID NO: 1214), and additional variants and analogs thereof which retain alkaline phosphatase activity, e.g., the ability to hydrolyze PP<sub>i</sub>.

35 By “sALP polypeptide” is meant any sequence including an sALP sequence, as defined herein. Exemplary sALP polypeptides include those having the structure A-sALP-B, wherein each of A and B is absent or is an amino acid sequence of at least one amino acid (e.g., any sALP fusion polypeptide described herein).

By “selective for NPR-B over NPR-A” is meant having an  $EC_{50(NPR-A)}/EC_{50(NPR-B)}$  ratio that is at least 1.25, 1.5, 2, 2.5, 3, 3.5, 4, 4.5, 5, 7.5, 10, 12.5, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 110, 120, 125, 130, 140, 150, 160, 170, 180, 190, 200, 225, 250, 275, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1,000, 1,100, 1,200, 1,250, 1,300, 1,400, 1,500, 1,750, 2,000, 2,500, 3,000, 4,000, 5,000, 10,000, or higher, or any range between any two of these values, in an *in vivo* or *in vitro* dose-response assay, e.g., measuring cGMP production, as described herein.

Alternatively, or in addition, the term “selective for NPR-B over NPR-A” means having an  $AUC_{(NPR-B)}/AUC_{(NPR-A)}$  ratio that is at least 1.1, 1.2, 1.25, 1.3, 1.4, 1.5, 1.6, 1.7, 1.75, 1.8, 1.9, 2, 2.25, 2.5, 2.75, 3, 3.5, 4, 4.5, 5, 7.5, 10, 12.5, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 110, 120, 125, 130, 140, 150, 160, 170, 180, 190, 200, 225, 250, 275, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1,000, 1,100, 1,200, 1,250, 1,300, 1,400, 1,500, 1,750, 2,000, 2,500, 3,000, 4,000, 5,000, 10,000, or higher, or any range between any two of these values, as described herein.

By “signal peptide” or “signal sequence” is meant an amino acid sequence that directs a polypeptide to the cellular membrane such that the polypeptide is secreted. Alternatively, the signal sequence may direct the polypeptide to an intracellular compartment or organelle, such as the Golgi apparatus. A signal sequence may be identified by homology, or biological activity, to a peptide sequence with the known function of targeting a polypeptide to a particular region of the cell. One of ordinary skill in the art can identify a signal sequence by using readily available software (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, WI 53705, BLAST, or PILEUP/PRETTYBOX programs). A signal sequence can be one that is, for example, substantially identical to amino acid residues 1-25 of SEQ ID NO: 501 or to amino acid residues 1-17 of SEQ ID NO: 1201.

By “skeletal dysplasia” is meant a bone or cartilage disorder characterized by short stature or dwarfism.

By “subject” is meant a mammal, including, but not limited to, a human or non-human mammal, such as a bovine, equine, canine, ovine, or feline.

By a “synergistic” effect is meant a therapeutic effect observed following administration of two or more agents that is greater than the sum of the therapeutic effects observed following the administration of each single agent. In one example of synergy, a therapeutic effect is observed for the combination of two or more agents, wherein one or more of the agents is present at a dose that is normally non-therapeutic. In another example of synergy, the combination of two or more agents results in an unexpected decrease in one or more adverse events (i.e., a level or number of adverse events that is less than the sum of adverse events observed following administration of the single agents). In another example, the combination of two or more agents at a therapeutic dose results in reduced dose-dependent side effects, as compared to the level of dose-dependent side effects observed for a single agent at a therapeutic dose.

By "therapeutically effective amount" is meant an amount of a polypeptide or nucleic acid molecule described herein that is sufficient to substantially treat, prevent, delay, suppress, or arrest any symptom of a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, a bone or cartilage disorder (e.g., achondroplasia), or a vascular smooth muscle disorder, or that is sufficient to substantially elongate bone. A therapeutically effective amount of a composition described herein may depend on the severity of the disorder being treated and the condition, weight, and general state of the subject and can be determined by an ordinarily-skilled artisan with consideration of such factors. A therapeutically effective amount of a composition described herein can be administered to a subject in a single dose or in multiple doses administered over a period of time.

By "treating," "treat," or "treatment" is meant the medical management of a patient with the intent to cure, ameliorate, stabilize, reduce the likelihood of, or prevent a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, a bone or cartilage disorder (e.g., achondroplasia), or a vascular smooth muscle disorder, or management of a healthy subject with the intent to elongate bone, e.g., by administering a pharmaceutical composition. This term includes active treatment, that is, treatment directed specifically toward the improvement or associated with the cure of a disease, pathological condition, disorder, or event, and also includes causal treatment, that is, treatment directed toward removal of the cause of the associated disease, pathological condition, disorder, or event. In addition, this term includes palliative treatment, that is, treatment designed for the relief of symptoms rather than the curing of the disease, pathological condition, disorder, or event; symptomatic treatment, that is, treatment directed toward constitutional symptoms of the associated disease, pathological condition, disorder, or event; preventative treatment, that is, treatment directed to minimizing or partially or completely inhibiting the development of the associated disease, pathological condition, disorder, or event, e.g., in a patient who is not yet ill, but who is susceptible to, or otherwise at risk of, a particular disease, pathological condition, disorder, or event; and supportive treatment, that is, treatment employed to supplement another specific therapy directed toward the improvement of the associated disease, pathological condition, disorder, or event.

By "vascular smooth muscle disorder" is meant any disorder, disease, or other abnormality that affects the function, structure, or growth of vascular smooth muscle.

By "vector" is meant a DNA molecule, usually derived from a plasmid or bacteriophage, into which fragments of DNA may be inserted or cloned. A recombinant vector will contain one or more unique restriction sites, and may be capable of autonomous replication in a defined host or vehicle organism such that the cloned sequence is reproducible. A vector contains a promoter operably linked to a gene or coding region such that, upon transfection into a recipient cell, an RNA is expressed.

Other features and advantages of the invention will be apparent from the detailed description and from the claims.

## BRIEF DESCRIPTION OF THE DRAWINGS

In figures showing a multiple sequence alignment, “\*” represents identity; “:” represents a conserved substitution; and “.” represents a semi-conserved substitution.

Fig. 1A is a graph showing levels of pyrophosphate (PP<sub>i</sub>) in osteoblasts from wild-type mice 5 (left) and NF1<sub>col2</sub><sup>-/-</sup> mice (right).

Fig. 1B is a graph showing levels of progressive ankylosis gene (*Ank*) mRNA expression in osteoblasts from wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice (labeled “KO”). Osteoblasts were treated with vehicle, a MEK1/MEK2 kinase inhibitor U0126 (1,4-diamino-2,3-dicyano-1,4-bis(2-aminophenylthio) butadiene), or bone morphogenetic protein 10 2 (BMP2).

Fig. 2 is a schematic showing a non-limiting, hypothetical working model for defective bone matrix mineralization in NF1<sub>col2</sub><sup>-/-</sup> mice.

Fig. 3A shows the effect of recombinant human bone morphogenetic protein 2 (rhBMP2) in osteoblasts from wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. Cell plates are shown with 15 histological staining for alkaline phosphatase (ALP) alone or in combination with alizarin red S (ALP/Alizarin) and for cell number with crystal violet.

Fig. 3B shows the effect of an sALP fusion polypeptide (sTNALP-FcD<sub>10</sub>, SEQ ID NO: 1204) in bone marrow stromal cells from wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. Cells were then treated for 8 days with increasing doses of sTNALP-FcD<sub>10</sub>. Cell plates are shown with histological 20 staining for alizarin red S to determine the presence of calcific deposition (top), and the relative intensity of staining with alizarin red S was quantified (bottom).

Fig. 3C shows the effect of an sALP fusion polypeptide (sTNALP-FcD<sub>10</sub>, SEQ ID NO: 1204) in bone marrow stromal cells (BMSCs) from wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. BMSCs were plated in culture and differentiation was induced for 14 days using vitamin C and beta-glycerophosphate. Cells were then treated with 0.5 µg/mL of sTNALP-FcD<sub>10</sub> and stained with alizarin 25 red to assess level of mineralization, with picrosirius red to demonstrate presence of extracellular matrix, or with crystal violet to stain cells.

Fig. 3D shows the effect of an sALP fusion polypeptide (sTNALP-FcD<sub>10</sub>, SEQ ID NO: 1204) in bone marrow stromal cells (BMSCs) from floxed NF1 gene mice. BMSCs were plated in culture, and 30 differentiation was induced for 14 days using vitamin C and beta-glycerophosphate. Cells were treated for 8 days with an adenovirus coding for GFP (“Ad-GFP,” as control) or an adenovirus coding for CRE recombinase (“Ad-Cre”) either with vehicle or with 0.5 µg/mL of sTNALP-FcD<sub>10</sub>. Cells were then stained with alizarin red to assess level of mineralization (first and second columns) or with picrosirius red to demonstrate presence of extracellular matrix (third and fourth columns).

Fig. 3E shows the *in vivo* effect of an sALP fusion polypeptide (sTNALP-FcD<sub>10</sub>, SEQ ID NO: 1204) in wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. NF1<sub>col2</sub><sup>-/-</sup> mice were treated from day 1 to day 18 with 8.2 mg/kg of sTNALP-FcD<sub>10</sub> and compared to untreated or WT mice. Mice were sacrificed 35

at day 19, and vertebrae were analysed with micro-CT imaging to measure bone volume over total volume (BV/TV). Treatment of mice with sTNALP-FcD<sub>10</sub> increased bone mineral density deficit in NF1<sub>col2</sub><sup>-/-</sup> mice compared to vehicle (\* p < 0.5).

5 Fig. 4A shows the expression of the *NPR-B* gene in NF1<sub>col2</sub><sup>-/-</sup> mice. Bone marrow stromal cells were cultured for 3 weeks under osteogenic conditions (with ascorbate acetate) and lysed. RT-PCR was performed using *NPR-B* and housekeeping gene *GAPDH* primers.

10 Fig. 4B shows the effect of an NP fusion polypeptide (NC2-KGANKK, SEQ ID NO: 512) in chondrocytes from wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. Western blot analysis provides levels of ERK and phosphorylated ERK (p-ERK) for chondrocytes from wild-type mice or from NF1<sub>col2</sub><sup>-/-</sup> mice. Primary chondrocytes extracted from ribs of newborn NF1<sub>col2</sub><sup>-/-</sup> mice or WT mice were cultured and then treated for 30 minutes with increasing concentrations of NC2-KGANKK. Cells were then lysed, and western blotting was performed on lysates using anti-ERK or anti-phospho-ERK specific antibodies.

15 Fig. 4C shows the *in vivo* effect of an NP fusion polypeptide (NC2B, SEQ ID NO: 504) on naso-anal length in wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. NF1<sub>col2</sub><sup>-/-</sup> mice were treated from day 1 to day 18 with 100 or 300 mg/kg of NC2B and compared to vehicle-treated or WT mice. Mice were measured for naso-anal length at day 19.

20 Fig. 4D shows the *in vivo* effect of an NP fusion polypeptide (NC2B, SEQ ID NO: 504) on tibia length in wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. NF1<sub>col2</sub><sup>-/-</sup> mice were treated from day 1 to day 18 with 100 or 300 mg/kg of NC2B and compared to vehicle-treated or WT mice. Mice were measured for tibia length at day 19.

Fig. 4E shows the *in vivo* effect of an NP fusion polypeptide (NC2B, SEQ ID NO: 504) on growth plates in wild-type mice (labeled “WT”) and NF1<sub>col2</sub><sup>-/-</sup> mice. NF1<sub>col2</sub><sup>-/-</sup> mice were treated from day 1 to day 18 with 300 mg/kg of NC2B and compared to vehicle-treated or WT mice. At day 19, tibias were used to analyze proximal growth plates in histology and to measure the size of chondrocyte zones.

25 Fig. 5A is a schematic diagram of the structure of human tissue nonspecific alkaline phosphatase (hsTNALP) as described herein, which includes a polypeptide having a TNALP ectodomain, an N-terminal signaling sequence, and a GPI signal sequence (top); a hsTNALP-FcD<sub>10</sub> having a TNALP ectodomain, an N-terminal signaling sequence, an IgG<sub>1</sub>-Fc sequence, and a bone-targeting D<sub>10</sub> moiety (middle); and a hsTNALP-FcD<sub>10</sub> without a signal sequence having a TNALP ectodomain, an IgG<sub>1</sub>-Fc sequence, and a bone-targeting D<sub>10</sub> moiety (bottom).

30 Fig. 5B shows the amino acid sequence of hsTNALP-FcD<sub>10</sub> (SEQ ID NO: 1201), including the N-terminal signal sequence (the first 17 amino acid residues, underlined and italicized; position 2, a valine, differs from the wild-type residue in that position, isoleucine); and the amino acid sequence of secreted hsTNALP-FcD<sub>10</sub> (SEQ ID NO: 1204), which lacks the N-terminal signal sequence. Amino acid residues of the hsTNALP portion of the polypeptides, which correspond to amino acid residues 18-502 of hsTNALP (SEQ ID NO: 1205), are italicized. The signal sequence in SEQ ID NO: 1201 is italicized and underlined. The Fc portions of the polypeptides (SEQ ID NO: 401) are underlined. A dipeptide leucine-

lysine (LK) linker between the hsTNALP and Fc portions and a dipeptide aspartic acid-isoleucine (DI) linker between the hsTNALP and bone-targeting moiety are shown in bold.

Fig. 5C shows the amino acid sequence of sALP fusion proteins lacking a bone-targeting moiety: hsTNALP-Fc (SEQ ID NO: 1220) and secreted hsTNALP-FcD<sub>10</sub> (SEQ ID NO: 1221), which lacks the N-terminal signal sequence. Amino acid residues of the hsTNALP portion of the polypeptides, which correspond to amino acid residues 18-502 of hsTNALP (SEQ ID NO: 1205), are italicized. The signal sequence in SEQ ID NO: 1201 is italicized and underlined. The Fc portions of the polypeptides (SEQ ID NO: 401) are underlined. A dipeptide leucine-lysine (LK) linker between the hsTNALP and Fc portions are shown in bold.

Fig. 5D shows a nucleic acid sequence (SEQ ID NO: 1217) encoding the hsTNALP-Fc polypeptide depicted in Fig. 5B.

Fig. 6 shows amino acid sequences for human soluble tissue nonspecific alkaline phosphatase (hsTNALP), including the sequence of hsTNALP having the N-terminal signal sequence (amino acid residues 1-502) (SEQ ID NO: 1202) and the sequence of secreted hsTNALP lacking the signal sequence (amino acid residues 18-502) (SEQ ID NO: 1205). The signal sequence is underlined.

Fig. 7 is a listing of the amino acid sequence of an exemplary Fc from human IgG-1 (SEQ ID NO: 401).

Fig. 8 shows a CLUSTAL™ W (1.82) multiple sequence alignment of mammalian tissue nonspecific alkaline phosphatase (TNALP) orthologs. Mammalian TNALP orthologs include cow TNALP (“P09487|PPBT\_BOVIN”; Accession No. P09487; SEQ ID NO: 1206); cat TNALP (“Q29486|PPBT\_FELCA”; Accession No. Q29486; SEQ ID NO: 1207), human TNALP (“P05186|PPBT\_HUMAN”; Accession No. P05186; SEQ ID NO: 1208), mouse TNALP (“P09242|PPBT\_MOUSE”; Accession No. P09242; SEQ ID NO: 1209), rat TNALP (“P08289|PPBT\_RAT”; Accession No. P08289; SEQ ID NO: 1210), and a partial sequence of dog TNALP (“Q9N0V0|Q9N0V0\_CANFA”; Accession No. Q9N0V0; SEQ ID NO: 1211). A consensus sequence is derived from this alignment (“Consensus”; SEQ ID NO: 1216), where X denotes degenerate positions and can be any amino acid.

Fig. 9 shows a CLUSTAL™ (2.0.5) multiple sequence alignment of mammalian tissue nonspecific alkaline phosphatase (TNALP) orthologs and human alkaline phosphatase (ALP) isozymes. Mammalian TNALP orthologs include those shown in Fig. 8, including rat TNALP (“TNALPrn,” SEQ ID NO: 1210), mouse TNALP (“TNALPmm,” SEQ ID NO: 1209), human TNALP (“TNALPhs,” SEQ ID NO: 1208), a partial sequence of dog TNALP (“TNALPcf,” SEQ ID NO: 1211), cat TNALP (“TNALPfc,” SEQ ID NO: 1207), and cow TNALP (“TNALPbt,” SEQ ID NO: 1206). Human ALP isozymes include a human gastrointestinal ALP (“GALPhs”; Accession No. P10696; SEQ ID NO: 1213), a human placental ALP (“PLALPhs”; Accession No. 05187; SEQ ID NO: 1214), and a human intestinal ALP (“IALPhs”; Accession No. P09923; SEQ ID NO: 1212). A consensus sequence is derived from this

alignment (“Consensus”; SEQ ID NO: 1215), where X denotes degenerate positions and can be any amino acid.

Fig. 10 is a consensus sequence (SEQ ID NO: 1218) for mammalian tissue nonspecific alkaline phosphatase (TNALP) orthologs and human alkaline phosphatase (ALP) isozymes excluding pathogenic mutations, where X can be any amino acid but not an amino acid corresponding to one or more pathogenic mutations provided in Table 1.

Fig. 11 is a consensus sequence (SEQ ID NO: 1219) for mammalian tissue nonspecific alkaline phosphatase (TNALP) orthologs excluding pathogenic mutations, where X can be any amino acid but not an amino acid corresponding to one or more pathogenic mutations provided in Table 1.

Fig. 12 is a multiple sequence alignment of human ANP (SEQ ID NO: 1), human urodilatin (SEQ ID NO: 2), human BNP (SEQ ID NO: 3), human CNP22 (SEQ ID NO: 4), and DNP (SEQ ID NO: 5). The 17-amino acid ring domain for each natriuretic peptide is shown in bold and enclosed in a box. A consensus sequence (SEQ ID NO: 6) is shown below, wherein each X represents any amino acid, or optionally represents any amino acid at the corresponding position in one of SEQ ID NOs: 1-5.

Fig. 13 is an alignment of human CNP53 (SEQ ID NO: 11), human CNP22, and human CNP (ring domain only) (SEQ ID NO: 12).

Fig. 14 is a multiple sequence alignment of various CNP22 homologs. The 17-amino acid ring domain for each NP is shown in bold and enclosed in a box. A consensus sequence (SEQ ID NO: 30) is shown below, wherein each X within the ring domain represents any amino acid, or optionally represents any amino acid at the corresponding position in one of SEQ ID NOs: 4 and 13-29. Each X outside the ring domain represents any amino acid or may be absent, or optionally represents any amino acid at the corresponding position in one of SEQ ID NOs: 4 and 13-29.

Figs. 15A-15G are a multiple sequence alignment of various CNP homologs, in some cases including the N-terminal pre- and pro-sequences. The 17-amino acid ring domain for each NP is shown in bold and enclosed in a box. A consensus sequence (SEQ ID NO: 95) is shown below, wherein each X represents any amino acid, or optionally represents any amino acid at the corresponding position in one of SEQ ID NOs: 31-94.

Fig. 16 is a schematic diagram of the structure of a natriuretic peptide as described herein, which includes an optional N-terminal extension, an optional short segment, a required ring domain, and an optional C-terminal extension.

Figs. 17A-17E are schematic diagrams of exemplary Fc-NP or NP-Fc constructs. Fig. 17A depicts an Fc-NP dimer. Fig. 17B depicts an NP-Fc dimer. Fig. 17C depicts an Fc:Fc-NP monomer-dimer hybrid. Fig. 17D depicts an NP-Fc:Fc monomer-dimer hybrid. Fig. 17E depicts an NP-Fc:Fc-NP hybrid dimer.

Fig. 18A is a listing of the amino acid sequence of the immature NC2 Streptag (“NC2st”) fusion protein (SEQ ID NO: 501), together with a table providing a summary of protein regions. The N-terminal signal sequence, which is cleaved during translation, is underlined. Various linker sequences are shown

in italics. The Fc domain is shown in bold. The CNP domain is shown in gray highlighting. Fig. 18B is a listing of the amino acid sequence of the NC2st fusion protein (SEQ ID NO: 502) without the signal sequence. Fig. 18C is a listing of the nucleic acid sequence (SEQ ID NO: 801) encoding the NC2st fusion protein.

5 Fig. 19A is a listing of the NC2B amino acid sequence, both with the signal sequence (SEQ ID NO: 503) and without the signal sequence (SEQ ID NO: 504), and the D10-NC2 amino acid sequence having a D<sub>10</sub> tag, both with the signal sequence (SEQ ID NO: 607) and without the signal sequence (SEQ ID NO: 608).

Fig. 19B is a listing of a nucleic acid sequence (SEQ ID NO: 802) encoding NC2B.

10 Fig. 20A is a listing of amino acid sequences for NC2B-22, NC2B-28, and NC2B-34, both with the signal sequence (SEQ ID NOs: 505, 507, and 509, respectively) and without the signal sequence (SEQ ID NOs: 506, 508, and 510, respectively). Signal sequences are underlined. The Fc domain is shown in bold. Linker sequences are shown in italics. The CNP domain is shown in gray highlighting. Fig. 20B is a listing of a nucleic acid sequence (SEQ ID NO: 803) encoding NC2B-22. Fig. 20C is a 15 listing of a nucleic acid sequence (SEQ ID NO: 804) encoding NC2B-28. Fig. 20D is a listing of a nucleic acid sequence (SEQ ID NO: 805) encoding NC2B-34.

Fig. 21 is a listing of amino acid sequences for NC2-KGANKK and NC2-KGANQK, both with the signal sequence (SEQ ID NOs: 511 and 513, respectively) and without the signal sequence (SEQ ID NOs: 512 and 514, respectively). Signal sequences are underlined. The Fc domain is shown in bold.

20 Linker sequences are shown in italics. The CNP domain is shown in gray highlighting.

Fig. 22 is a listing of amino acid sequences for NC2-CNP53mut2, both with the signal sequence (SEQ ID NO: 515) and without the signal sequence (SEQ ID NOs: 516). Signal sequence is underlined. The Fc domain is shown in bold. Linker sequences are shown in italics. The CNP domain is shown in gray highlighting.

25 Fig. 23 is a listing of amino acid sequences for Fc-CNP53-A (also referred to as Fc-CNP53wt) and Fc-CNP53-AAA (also referred to as Fc-CNP53mut), both with the signal sequence (SEQ ID NOs: 517 and 519, respectively) and without the signal sequence (SEQ ID NOs: 518 and 520, respectively). Signal sequences are underlined. The Fc domain is shown in bold. Linker sequences are shown in italics. The CNP domain is shown in gray highlighting.

30 Fig. 24 is a multiple sequence alignment of various NPs and homologs, including CDNP. The boxed region is the most conserved region of the DNP tail among NPRA-binding peptides. The sequences of numerous CDNP variants are shown in the bottom half of the figure, and a consensus sequence (SEQ ID NO: 118) for the DNP C-terminal tail is also shown. Each X in the consensus sequence represents any amino acid, or optionally represents any amino acid at the corresponding 35 position in one of SEQ ID NOs: 100-116.

Fig. 25A is a listing of amino acid sequences for CNP-16AAlinker-Fc-His10 (NC1) (SEQ ID NO: 521), CNP-6AAlinker-Fc-His10 (NC3) (SEQ ID NO: 522), CNP-6AAlinker-Fc (SEQ ID NO: 523),

CDNP-Fc (SEQ ID NO: 524), CDNP-A17saa-Fc (SEQ ID NO: 525), and CDNP-A17sra-Fc (SEQ ID NO: 526). The CNP domain is shown in gray highlighting. Linker sequences are shown in italics. The Fc domain is shown in bold. Fig. 25B is a listing of the nucleic acid sequence (SEQ ID NO: 806) of NC1.

5 Fig. 26 is a listing of various point mutants (SEQ ID NOs: 119-125) each having a mutation at position 17 of CNP22, together with a consensus sequence (SEQ ID NO: 126). X represents any amino acid, or optionally represents any amino acid at the corresponding position in one of SEQ ID NOs: 119-125.

10 Fig. 27 is a listing of amino acid sequences for several CNP variants (SEQ ID NOs: 4 and 127-150). The 17-amino acid ring domain for each variant is shown in bold. The linker region is shown in italics.

15 Figs. 28A-28E are a listing of amino acid sequences for additional CNP variants (SEQ ID NOs: 1001-1155).

20 Fig. 29 is a listing of amino acid sequences for CNP22 (SEQ ID NO: 4), CNP-L17 (SEQ ID NO: 120), CNP-F17 (SEQ ID NO: 119), CNP-T17 (SEQ ID NO: 122), D6-14AAlinker-CNP [C3] (SEQ ID NO: 147), CNP-14AAlinker-D6 [C4] (SEQ ID NO: 148), CNP-Nterm2 [C5] (SEQ ID NO: 150), CDNP-S3A4A5R6 [C13] (SEQ ID NO: 115), CDNP29-S3A4A5R6 [C14] (SEQ ID NO: 151), C1(E6) [BC1] (SEQ ID NO: 129), C2(E6) [BC2] (SEQ ID NO: 130), C3 (E6) [BC3] (SEQ ID NO: 131), C4(E6) [BC4] (SEQ ID NO: 132), C5(E6) [BC5] (SEQ ID NO: 133), C6(E6) [BC6] (SEQ ID NO: 134), C7(E6) [BC7] (SEQ ID NO: 135), C8(E6) [BC8] (SEQ ID NO: 136), C9(E6) [BC9] (SEQ ID NO: 137), C10(E6) [BC10] (SEQ ID NO: 138), C11(E6) [BC11] (SEQ ID NO: 139), PGCNP37(E6) (SEQ ID NO: 128), KA1 (SEQ ID NO: 152), KA1(E6) (SEQ ID NO: 153), KB1 (SEQ ID NO: 154), and KB1(E6) (SEQ ID NO: 155). The 17-amino acid ring domain for each variant is shown in bold. The linker sequences are shown in italics. The cathepsin cleavage sequences are shown in underline.

25 Fig. 30 is a listing of amino acid sequences for CNP variants having a point mutation at position 17 relative to CNP22 (SEQ ID NOs: 126, 119-122, and 156-172). For SEQ ID NOs: 126 and 162, X can be any amino acid, including but not limited to F, L, I, T, E, R, Y, C, P, or D. The 17-amino acid ring domain for each variant is shown in bold. The linker sequences are shown in italics.

30 Figs. 31A-31B are listings of amino acid sequences for additional CNP variants having a point mutation at position 17 relative to CNP22 (SEQ ID NOs: 173-220). X can be any amino acid, including but not limited to F, L, I, T, E, R, Y, C, P, or D. The 17-amino acid ring domain for each variant is shown in bold. The linker sequences are shown in italics. The cathepsin cleavage sequences are shown in underline.

35 Fig. 32 is a listing of amino acid sequences for CNP variants having a point mutation at position 17 relative to CNP22, where the methionine at position 17 has been substituted with a leucine (SEQ ID NOs: 221-233). The 17-amino acid ring domain for each variant is shown in bold. The linker sequences are shown in italics. The cathepsin cleavage sequences are shown in underline.

Figs. 33A-33E are listings of amino acid sequences for constructs having a point mutation at position 17 relative to CNP22 (SEQ ID NOS: 527-552). X can be any amino acid, including but not limited to F, L, I, T, E, R, Y, C, P, or D. Signal sequences are underlined. The Fc domain is shown in bold. Linker sequences are shown in italics. The CNP domain is shown in gray highlighting.

5 Figs. 34A-34J are listings of amino acid sequences for NC2 variants (SEQ ID NOS: 511-516 and 553-606) with or without the signal sequence and either with or without a D<sub>10</sub> bone-targeting moiety at the N-terminal. Signal sequences are underlined. The Fc domain is shown in bold. Linker sequences are shown in italics. The CNP domain is shown in gray highlighting.

## 10 DETAILED DESCRIPTION OF THE INVENTION

The present invention features soluble alkaline phosphatase (sALP) polypeptides, e.g., fused to an Fc domain of an immunoglobulin, nucleic acid encoding such, and their uses to treat any disease or condition described herein (e.g., neurocutaneous syndromes (e.g., neurofibromatosis, e.g., type 1), disorders associated with overactivation of FGFR3, bone or cartilage disorders (e.g., hypophosphatasia or 15 achondroplasia), vascular smooth muscle disorders, as well as to elongate bone). The present invention also features natriuretic (NP) polypeptides, e.g., fused to an Fc domain of an immunoglobulin, nucleic acid encoding such, and their uses to treat any disease or condition described herein (e.g., neurocutaneous syndromes (e.g., neurofibromatosis, e.g., type 1), disorders associated with overactivation of FGFR3, bone or cartilage disorders (e.g., hypophosphatasia or achondroplasia), vascular smooth muscle disorders, 20 as well as to elongate bone). The present invention also features a combination of such sALP polypeptides with such NP polypeptides, as described herein, and uses of this combination to treat any disease or condition described herein (e.g., neurocutaneous syndromes, disorders associated with overactivation of FGFR3, bone or cartilage disorders (e.g., hypophosphatasia or achondroplasia), vascular smooth muscle disorders, as well as to elongate bone). Additional details of the invention are provided 25 below.

### Alkaline phosphatase

Alkaline phosphatases encompass a group of enzymes that share the property of being able to cleave phosphate in a variety of contexts (e.g., hydrolysis of pyrophosphate, PP<sub>i</sub>). There are four known 30 mammalian alkaline phosphatase (ALP) isozymes: tissue nonspecific alkaline phosphatase (TNALP; described further below), placental alkaline phosphatase (PLALP) (e.g., Accession Nos. P05187, NP\_112603, and NP\_001623), germ cell alkaline phosphatase (GALP) (e.g., Accession No. P10696), and intestinal alkaline phosphatase (IALP) (e.g., Accession Nos. P09923 and NP\_001622). These isozymes possess very similar three dimensional structures. Each of their catalytic sites contains four metal binding 35 domains that bind to metal ions necessary for enzymatic activity, including two zinc ions and one magnesium ion. These enzymes catalyze the hydrolysis of monoesters of phosphoric acid and also catalyze a transphosphorylation reaction in the presence of high concentrations of phosphate acceptors. It

has been shown that PLALP is physiologically active toward phosphoethanolamine (PEA), inorganic pyrophosphate (PP<sub>i</sub>), and pyridoxal 5'-phosphate (PLP), all three being known natural substrate for TNALP (Whyte, 1995). An alignment between these isozymes is shown in Fig. 9. Additional alkaline phosphatases are described, e.g., in WO 2008/138131 and in U.S. Publication No. 2006/0014687, which

5 are hereby incorporated by reference.

Tissue nonspecific phosphatases are a family of proteins, encoded by a single gene, that differ from each other by post-translational modification. TNALPs are present predominantly in the liver, kidneys, and bone, but can occur throughout the body. Known TNALPs in mammals include, e.g., human TNALP (Accession Nos. NP\_000469, AAI10910, AAH90861, AAH66116, AAH21289, and 10 AAI26166); rhesus TNALP (Accession No. XP\_01109717); rat TNALP (Accession No. NP\_037191); dog TNALP (Accession No. AAF64516); pig TNALP (Accession No. AAN64273), mouse (Accession No. NP\_031457), cow TNALP (Accession Nos. NP\_789828, NP\_776412, AAM 8209, and AAC33858), and cat TNALP (Accession No. NP\_001036028), in addition to other examples provided herein.

## 15 Soluble alkaline phosphatase

The soluble alkaline phosphatases (sALP) of the invention include, for example, soluble (e.g., extracellular or non membrane-bound) forms of any of the alkaline phosphatases described herein. The soluble alkaline phosphatase of the invention can be, for example, a soluble form of human TNALP. A schematic representation of the domains of human TNALP (hTNALP) is shown in Fig. 5A (top).

20 TNALP is a membrane-bound protein anchored through a glycolipid bound to its C-terminal (Swiss-Prot, P05186). This glycolipid anchor (GPI) is added post translationally after the removal of a hydrophobic C-terminal end, which serves both as a temporary membrane anchor and as a signal for the addition of the 25 GPI. This GPI anchor is buried in the cell membrane, and the remaining portions of the protein are extracellular. TNALP, including hTNALP, can be engineered to replace the first amino acid of the hydrophobic C-terminal sequence (an alanine) with a stop codon. The engineered hTNALP so formed 30 contains all amino acid residues of the native anchored form of TNALP but lacks the GPI membrane anchor. An hTNALP which is soluble is herein referred to as "hsTNALP." One skilled in the art will appreciate that the position of the GPI membrane anchor will vary in different alkaline phosphatases and may include, for example, the last 10, 12, 14, 16, 18, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 32, 34, 36, 38, 40, 45, 50, or more amino acid residues on the C-terminus of the polypeptide. For example, the GPI 35 membrane anchor of the hTNALP (SEQ ID NO: 1208) is amino acid residues 503-524. The amino acid sequence of this hsTNALP (with one variation at position 2 of the signal sequence), as fused to Fc, is shown in Fig. 5B. The sequence of a nucleic acid encoding this hsTNALP-Fc fusion polypeptide is shown in Fig. 5D (SEQ ID NO: 1217).

35 In addition to the C-terminal GPI anchor, TNALP also has an N-terminal signal peptide sequence. The N-terminal signal peptide is initially present on the protein when it is synthesized, but is cleaved after translocation into the ER. Thus, the N-terminal signal peptide is absent from the secreted

form of TNALP. The sALPs of the invention include both secreted (i.e., lacking the N-terminal signal) and non-secreted (i.e., having the N-terminal signal) forms thereof. One skilled in the art will appreciate that the position of the N-terminal signal peptide will vary in different alkaline phosphatases and may include, for example, the first 5, 8, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 27, 30, or 5 more amino acid residues on the N-terminus of the polypeptide. For example, the N-terminal signal peptide of the hTNALP of SEQ ID NO: 1208 is its first 17 amino acid residues, as shown in Fig. 6. Thus, a secreted, soluble form of this hTNALP is amino acid residues 18-502 of SEQ ID NO: 1208 (SEQ ID NO: 1205), as shown in Fig. 6. The amino acid sequence of this secreted hsTNALP, as fused to Fc, is shown in Fig. 5B. The sALPs of the invention include both secreted and non-secreted forms thereof. 10 One of skill in the art can predict the position of a signal sequence cleavage site, e.g., by an appropriate computer algorithm such as that described in Bendtsen et al. (*J. Mol. Biol.* 340(4):783-795, 2004) and available on the Web at <http://www.cbs.dtu.dk/services/SignalP/>.

The sALPs of the invention also include, for example, polypeptide sequences satisfying a consensus sequence derived from the ALP extracellular domain of human ALP isozymes and of 15 mammalian TNALP orthologs (human, mouse, rat, cow, cat, and dog) (SEQ ID NO: 1215, as shown in Fig. 9), or a consensus derived from the ALP extracellular domain of just mammalian TNALP orthologs (human, mouse, rat, cow, cat, and dog) (SEQ ID NO: 1216, as shown in Fig. 8). In some embodiments, the sALP includes amino acid residues 18-498, 18-499, 18-500, 18-501, 18-502, 18-503, 18-504, 18-505, 18-506, 18-507, 18-508, 18-509, 18-510, 18-511, or 18-512 of SEQ ID NO: 1216. In some 20 embodiments, the sALP includes amino acid residues 23-498, 23-499, 23-500, 23-501, 23-502, 23-503, 23-504, 23-505, 23-506, 23-507, 23-508, 23-509, 23-510, 23-511, or 23-512 of SEQ ID NO: 1215.

The sALPs of the invention also include those which satisfy similar consensus sequences derived from various combinations of these TNALP orthologs or human ALP isozymes. Such consensus sequences are given, for example, in WO 2008/138131, herein incorporated by reference.

25 Furthermore, it has been shown that recombinant hsTNALP retaining original amino acid residues 1 to 501 (18 to 501 when secreted) (Oda et al., *J. Biochem.* 126: 694-699, 1999), amino acid residues 1 to 502 (18 to 502 when secreted) (WO 2008/138131), amino acid residues 1 to 504 (18 to 504 when secreted) (U.S. Pat. No. 6,905,689, which is herein incorporated by reference), and amino acid residues 1 to 505 (18-505 when secreted) (U.S. Pat. Pub. No. 2007/0081984, which is herein incorporated 30 by reference), are enzymatically active. This indicates that certain amino acid residues can be truncated from the C-terminal end of the soluble hsTNALP polypeptide without affecting its enzymatic activity. This also indicates that certain amino acid residues of the GPI membrane anchor, when present, do not significantly affect the solubility of the polypeptide. Hence, the sALPs of the invention also include those where, e.g., up to five (e.g., one, two, three, four, or five) amino acid residues are truncated on its 35 C-terminal end, and those where, e.g., up to five (e.g., one, two, three, four, or five) amino acid residues of the GPI membrane anchor are present. For example, non-secreted sALPs of the invention include those containing amino acid residues 1-497, 1-498, 1-499, 1-500, 1-501, 1-502, 1-503, 1-504, 1-505, 1-

506, or 1-507 of SEQ ID NO: 1208, as well as variants thereof where the amino acid at position 2 is a valine, and secreted sALPs of the invention include those containing amino acid residues 18-497, 18-498, 18-499, 18-500, 18-501, 18-502, 18-503, 18-504, 18-505, 18-506, or 18-507 of SEQ ID NO: 1208.

One skilled in the art will appreciate that many mutations in the amino acid sequence of an enzyme will not significantly disrupt the catalytic function of the enzyme. In some cases, certain mutation may even benefit the catalytic function of the enzyme in the context of therapy for any disorder or condition described herein (e.g., a neurocutaneous syndrome or a bone or cartilage disorder).

Therefore, the sALPs of the invention include not only the wild-type sequence of the alkaline phosphatases described above, but also include any polypeptide having at least 50% (e.g., 55%, 60%,

65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to these alkaline phosphatases. It is known, however, that specific mutations in TNALP are known to cause HPP. Such pathogenic mutations are preferably absent in the sALPs of the invention. Thus, the sALPs of the invention include those having an amino acid sequence including a consensus sequence for multiple mammalian TNALP orthologs and human ALP isozymes

that lack pathogenic mutations (SEQ ID NO: 1218, as shown in Fig. 10) or including a consensus sequence for multiple mammalian TNALP orthologs that lack pathogenic mutations (SEQ ID NO: 1219, as shown in Fig. 11). Exemplary sALPs of the invention include those containing amino acid residues 1-

497, 1-498, 1-499, 1-500, 1-501, 1-502, 1-503, 1-504, 1-505, 1-506, 1-507, 1-508, 1-509, 1-510, 1-511, 1-512, 23-497, 23-498, 23-499, 23-500, 23-501, 23-502, 23-503, 23-504, 23-505, 23-506, 23-507, 23-

508, 23-509, 23-510, 23-511, or 23-512 of SEQ ID NO: 1218, and secreted sALPs of the invention include those containing amino acid residues 18-497, 18-498, 18-499, 18-500, 18-501, 18-502, 18-503, 18-504, 18-505, 18-506, 18-507, 18-508, 18-509, 18-510, 18-511, or 18-512 of SEQ ID NO: 1219. In these consensus sequences (SEQ ID NOs: 1218 and 1219), X is any amino acid but is not an amino acid corresponding to a pathogenic mutation at that position of human TNALP. Examples of such pathogenic mutations are listed below and provided in Table 1.

Table 1. Pathogenic mutations for human TNALP\*

Total number of mutations 188							
Exon	Base change	Amino acid change		Reference	Clinical form in patient	Genotype of patient	% WT
		Non-standardized nomenclature	Standardized nomenclature				ref.
1	c.195C>T			Taillandier et al. 2000	perinatal	C.-195C>T/C184Y	
2	c.17T>A	L-12X	p.L6X	Taillandier et al. 2000	childhood	L-12X?	
2	c.50C>T	S-1F	p.S17F	Mornet et al. 1998	infantile	S-1F/G58S	19.0
3	c.83A>G	Y11C	p.Y28C	Taillandier et al. 2001	infantile	Y11C/R119H	7.2
3	c.98C>T	A16V	p.A33V	Henthorn et al. 1992	childhood	A16V/Y4.19H	-
3	c.110T>C	L20P	p.L37P	Versailles lab oct. 2003	perinatal	L20P/L20P	+
3	c.119C>T	A23V	p.A40V	Mornet et al. 1998	perinatal	A23V/G456S	2.3
3	c.132C>T	Q27X	p.Q44X	Mornet E, unpublished	perinatal	Q27X/c.662in	1
3	c.151G>T	A34S	p.A51S	Mumm et al. 2002	infantile	sG	
3	c.152G>T	A34V	p.A51V	Taillandier et al. 2001	infantile	A34S/T117H	+
4	c.184A>T	M45L	p.M62L	Taillandier et al. 1999	infantile	M45L/c.1172d	+
4	c.184A>G	M45V	p.M62V	Spentchian et al. 2003	infantile	elC	27.4
4	c.186G>C	M45I	p.M62I	Taillandier et al. 2005	childhood	M45V/E174K	0
4	c.187G>C	G46R	p.G63R	Spentchian et al. 2003	infantile	G46RG46R	+
4	c.188G>T	G46V	p.G63V	Lia-Baldini et al. 2001	infantile	G46V/N	0.8
4	c.203C>T	T51M	p.T68M	Orimo et al. 2002	childhood	T51M/A160T	5.2
4	c.211C>T	R54C	p.R71C	Henthorn et al. 1992	infantile	R54C/D277A	0
4	c.211C>A	R54S	p.R71S	Orimo et al. 2002	childhood	R54S?	2.9

\* In the column labeled "Non-standardized nomenclature" under "Amino acid change," the position of the mutation is provided with respect to mature human TNALP lacking the N-terminal signal sequence. In the column labeled "Standardized nomenclature" under "Amino acid change," the position is provided with respect to full length human TNALP having the 17-amino acid N-terminal signal sequence (SEQ ID NO: 1208).

4	c.212G>C	R54P	p.R71P	Henthorn et al. 1992	perinatal	R54P/Q190P	
4	c.212G>A	R54H	p.R71H	Taillandier et al. 2001	perinatal	A23V/R54H	+
4	c.219T>C	I55T	p.I72T	Versailles lab oct. 2004	odonto	I55T/N	+
4	c.223G>A	G58S	p.G75S	Mornet et al. 1998	infantile	S-1F/G58S	-
4	c.227A>G	Q59R	p.Q76R	Mornet et al. 2001	infantile	Q59R/T117N	+
IVS4	c.298-2A>G			Taillandier et al. 2000	perinatal	c.298-2A>G/c.997+3 A>C	This mutation affects splicing and not coding sequence
5	c.299C>T	T83M	p.T100M	Mornet et al. 2001	infantile	T83M/E174K	+
5	c.303_311del	N85_N87del	p.N102_N104del	Versailles lab Jul 2007	perinatal	C.303_311del/ G474R	na Deletion
5	c.323C>T	P91L	p.P108L	Herasse et al. 2003	odonto	P91L/N	0.4
5	c.331G>A	A94T	p.A111T	Goseki-Sone et al. 1998	odonto	A94T/?	unp. +
5	c.334G>A	G95S	p.G112S	Witters et al. 2004	infantile	G95S/R374C	-
5	c.340G>A	A97T	p.A114T	Mumm et al. 2001	infantile	A97T/D277A	+
5	c.341C>G	A97G	p.A114G	Draquet et al. 2004	perinatal	A97G+c.348- 349insACCGT C/G309R	+
5	c.348_349insA CCGTC			Draquet et al. 2004	perinatal	A97G+c.348- 349insACCGT C/G309R	na Two missense mutations and insertion
5	c.346G>T	A99S	p.A116S	Versailles lab Jul 2007	adult	A98S/N400S	+
5	c.346G>A	A99T	p.A116T	Hu et al. 2000	adult	A99T/N	0.8 3
5	c.358G>A	G103R	p.G120R	Mornet et al. 1998	perinatal	G103R/648+1 G>A	+
5	c.368C>A	A106D	p.A123D	Spentchian et al. 2006	perinatal	A106DS249_- H250del	-

5	c.382G>A	V111M	p.V128M	Mumm et al. 2002	perinatal	V111M/R206W	
5	c.385G>A	G112R	p.G129R	Mornet et al. 1998	perinatal	G112R/G474R	+
5	c.388_391delG	TAA		Spentchian et al. 2003	perinatal	E294K/388_391delGTA	na
5	c.389delT			Spentchian et al. 2003	perinatal	c.389delT/c.389delT	Frameshift mutation
5	c.392delG			Mumm et al. 2002	perinat/infant	c.392delG/A331T	na
5	c.394G>A	A115T	p.A132T	Versailles lab Jul 2006	adult	A115T/E174K	
5	c.395C>T	A115V	p.A132V	Watanabe et al. 2001	adult	A115V/?	16.9
5	c.400_401AC>CA	T117H	p.T134H	Mumm et al. 2002	perinatal	T117H/F310del	-
5	c.401C>A	T117N	p.T134N	Taillandier et al. 2000	perinatal	T117N/T117N	20.5
5	c.406C>T	R119C	p.R136C	Versailles lab oct. 2003	odont/o	R119C/R119H	-
5	c.407G>A	R119H	p.R136H	Taillandier et al. 1999	infantile	R119H/G145V	33.4
5	c.442A>G	T131A	p.T148A	Michigan et al. 2005	perinatal	T131A?	-
5	c.443C>T	T131I	p.T148I	Spentchian et al. 2003	infantile	T131I/G145S	-
6	c.480delT			Versailles lab. Jan. 2008	perinatal	c.480delT/R206W	na deletion
6	c.484G>A	G145S	p.G162S	Spentchian et al. 2003	infantile	T131I/G145S	+
6	c.485G>T	G145V	p.G162V	Taillandier et al. 1999	infantile	R119H/G145V	1.3
6	c.500C>T	T150M	p.T167M	Versailles lab oct. 2003	infantile	T150M/E174K0	+
6	c.508A>G	N153D	p.N170D	Mornet et al. 1998	perinatal	N153D/N153D	0
6	c.511C>T	H154Y	p.H171Y	Taillandier et al. 1999	infantile	H154Y/E174K2.1	1

6	c.512A>G	H154R	p.H171R	Mornet E, unpublished	adult	H154R/E174K		
6	c.526G>A	A159T	p.A176T	Taillandier et al. 2000	childhood	A159T/R229S	45.4	5
6	c.529G>A	A160T	p.A177T	Goseki-Sone et al. 1998	adult	A160T/F310L	83.8	4
6	c.535G>A	A162T	p.A179T	Weiss et al. 1988	perinatal	A162T/A162T	18	6
6	c.542C>T	S164L	p.S181L	Lia-Baldini et al. 2001	infantile	S164L/del(ex1-2)	1.3	3
6	c.544delG			Taillandier et al. 1999	perinatal	G232V/S544delG		na
6	c.550C>T	R167W	p.R184W	Mornet et al. 1998	perinatal	R167WW253X	0.6	3
6	c.567C>A	D172E	p.D189E	Spentchian et al. 2003	perinatal	D172E/D172E		-
6	c.568_570delA AC	N173del	p.N190del	Michigani et al. 2005	perinatal	c.1559del1/T 173del		Deletion of 1 a.a.
6	c.571G>A	E174K	p.E191K	Henthorn et al. 1992	infantile	E174K/D361V	88.0	1
6	c.572A>G	E174G	p.E191G	Goseki-Sone et al. 1998	odontia	E174G/c.1559 delT		-
6	c.575T>C	M175T	p.M192T	Versailles lab Jul 2007	infantile	M175T/E294K		-
6	c.577C>G	P176A	p.P193A	Mumm et al. 2002	adult	A97T/P176A		+
6	c.602G>A	C184Y	p.C201Y	Taillandier et al. 1999	perinatal	c.- 195C>T/C184Y		-
6	c.609C>G	D186E	p.D203E	Versailles lab oct. 2004	perinatal	D186E/D186E		-
6	c.620A>C	Q190P	p.Q207P	Henthorn et al. 1992	perinatal	R54P/Q190P		+
6	c.631A>G	N194D	p.N211D	Taillandier et al. 2001	infantile	A99T/N194D		+
6	c.634A>T	I195F	p.I212F	Souka et al. 2002	perinatal	I195F/E337D		-
IVS6	c.648+1G>T			Brun-Heath et al. 2005	perinatal	C.648+1G>T/ D277A		Affects splicing
IVS6	c.648+1G>A			Mornet et al. 1998	perinatal	G103R/c.648+ 1G>A		Affects splicing

IVS6	c.649-1_3delinsAA			Versailles lab Jul 2006	perinatal	c.649-1_3delinsAA/c c.649-1_3delinsAA			Frameshift mutation
7	c.653T>C	1201T	p.I218T	Utsch et al., 2005, contact	perinatal	I201T/R374C	3.7	unp. -	
7	659G>T	G203V	p.G220V	Tailandier et al. 2001	odonto	E174K/G203V		+	
7	659G>C	G203A	p.G220A	Spentchian et al. 2003	perinatal	G203A/G203A		+	
7	662insG			Mornet E, unpublished	perinatal	Q27X/662insG		na	Frameshift mutation
7	c.662delG			Spentchian et al. 2003	perinatal	R255L/c.662delG		na	Frameshift mutation
7	c.662G>T	G204V	p.G221V	Versailles lab oct. 2004	perinatal	G204V/M338T		+	
7	c.667C>T	R206W	p.R223W	Mornet et al. 1998	perinatal	R206W/T	2.8	3	Frameshift mutation
7	c.668G>A	R206Q	p.R223Q	Mumm et al. 2002	perinatal	R206Q/deletio		-	
7	c.670A>G	K207E	p.K224E	Mochizuki et al. 2000	infantile	K207E/G409C	43	15	+
7	c.677T>C	M209T	p.M226T	Baumgartner-Sigl et al. 2007	infantile	M209TT354I		-	
7	c.704A>G	E218G	p.E235G	Tailandier et al. 2001	adult	E218G/A382S	3.6	7	+
7	c.738G>T	R229S	p.R246S	Tailandier et al. 2000	childhood	A159T/R229S	4.4	5	-
7	c.746G>T	G232V	p.G249V	Fedde et al. 1996	perinatal	G232V/N	34.5	3	+
7	c.971A>G	K247R	p.K264R	Versailles lab Jan.2007	perinatal	K247R/D361V		-	
8	c.797_802del	S249_H250del	p.S266_H267del	Spentchian et al. 2006	perinatal	A106D/S249-H250del			Deletion of 2 a.a.
8	c.809G>A	W253X	p.W270X	Mornet et al. 1998	perinatal	R167W/W253X		na	Nonsense mutation
8	c.814C>T	R255C	p.R272C	Spentchian et al. 2006	perinatal	R255C/T117H		-	

8	c.815G>T	R255L	p.R272L	Spentchian et al. 2003	perinatal	R255L/c.662d elG	-
8	c.815G>A	R255H	p.R272H	Brun-Heath et al. 2005	infantile	R255H/R255 H	16 -
8	c.824T>C	L258P	p.L275P	Orimo et al. 2002	childhood	L258P/A160T	3.3 4 -
8	c.853_854insG ATC	Y268X	p.Y285X	Michigani et al. 2005	perinatal	C1559delT/Y2 68X	na Nonsense mutation
IVS8	c.862+5G>A			Taillandier et al. 1999	infantile	C.862+5G>A/C .862+5G>A	na Affects splicing
9	c.865C>T	L272F	p.L289F	Sugimoto et al. 1998	infantile	L272F?	50 8 -
9	c.871G>A	E274K	p.E291K	Mornet et al. 1998	infantile	E174K/E274K	8.3 1 -
9	c.871G>T	E274X	p.E291X	Taillandier et al. 2000	perinatal	A94T/E274X	-
9	c.874C>A	P275T	p.P292T	Brun-Heath et al. 2005	infantile	P275T/A16V	4.0 16 +
9	c.876_881delA GGGGA	G276_D277del		Spentchian et al. 2003	perinatal	G276_D277d el/c.962delG	na
9	c.880G>T	D277Y	p.D294Y	Taillandier et al. 2001	infantile	A153T/D277Y	-
9	c.881A>C	D277A	p.D294A	Henthorn et al. 1992	infantile	R54C/D277A	0 17 -
9	c.883A>G	M278V	p.M295V	Mornet et al. 2001	childhood	E174K/M278V	-
9	c.884T>C	M278T	p.M295T	Brun-Heath et al. 2005	perinatal	M278T/R206 W	8.5 16 -
9	c.885G>A	M278I	p.M295I	Michigani et al. 2005	perinatal	M278I/c.1559 delT	-
9	c.889T>G	Y280D	p.Y297D	Brun-Heath et al. 2005	childhood	R119H/Y280D	1.3 16 -
9	c.892G>A	E281K	p.E298K	Orimo et al. 1994	infantile	E281K/1559d elT	-
9	c.896T>C	L282P	p.L299P	Versailles lab oct. 2003	infantile	L282P/L282P	9.7 15 -
9	c.917A>T	D289V	p.D306V	Taillandier et al. 1999	infantile	D289V/D289V	0 12 -
9	c.919C>T	P290S	p.P307S	Versailles lab oct. 2004	infantile	P290S/M450T	+
9	c.920C>T	P290L	p.P307L	Versailles lab Jul. 2006	childhood	P290L/S164L	

9	C.928_929delT C		Brun-Heath et al. 2005	perinatal	T394A/c.928_929delTC		Frameshift mutation
9	c.931G>A	E294K	Spentchian et al. 2003	perinatal	E294K/c.388_391delCTAA	-	
9	c.962delG		Spentchian et al. 2003	perinatal	G276_D277d del/c.962delG	na	Frameshift mutation
9	c.976G>C	G309R	Litmanovitz et al. 2002	perinatal	G309R/E274K	+	
9	c.981_983delC TT	p.G326R	Orimo et al. 1997	infantile	F310del/c.155_9delT	15	Amino acid deletion
9	c.979T>G	F310C	Momet et al. 2001	perinatal	T117N/F310C	+	
9	c.979_980TT> GG	p.F327G	Taillandier et al. 2001	adult	E174K/F310G	+	
9	c.979T>C	F310L	Ozono et al. 1996	infantile	F310L/G439R	72	
9	c.982T>A	F311L	Michigani et al. 2005	perinatal non-lethal	F311L/T83M	10	
IVS9	c.997+2T>A	p.F328L	Taillandier et al. 2000	perinatal	c.997+2T>A/C 472S	15	
IVS9	c.997+2T>G		Brun-Heath et al. 2005	perinatal	C.997+2T>G/c .997+2T>G	na	Affects splicing
IVS9	c.997+3A>C		Momet et al. 1998	perinatal	C.997+3A>C/c .997+3A>C	na	Affects splicing
IVS9	c.998-1G>T		Taillandier et al. 2001	perinatal	E174K/c.988-1G>T	na	Affects splicing
10	c.1001G>A	G317D	Greenberg et al. 1993	perinatal	G317D/G317D	0	
10	c.1015G>A	G322R	Mumm et al. 2002	perinatal	G322R/A159T M	-	
10	c.1016G>A	G322E	Versailles lab oct. 2004	infantile	G322E/V111M	-	
10	c.1042G>A	A331T	Taillandier et al. 2000	infantile	E174K/A331T	33.2	
10	c.1044_1055del _332_A335del	p.I.349_A3 52del	Spentchian et al. 2006	perinatal	L332_A335del /G474R	5	Deletion of 4 a.a.

10	c.1062G>C	E337D	p.E3354D	Souka et al. 2002	perinatal	I195F/E337D	
10	c.1064A>C	M338T	p.M355T	Versailles lab oct. 2004	perinatal	G204V/M338T	
10	c.1065G>A	M338I	p.M355I	Versailles lab. Jan. 2008	infantile	M338I/R374C	
10	c.1101_1103de (CTC	S351del	p.S368del	Versailles lab oct. 2004	perinatal	c.1101_1103d eICTC/T372I	
10	c.1112C>T	T354I	p.T371I	Baumgartner-Sigl et al. 2007	infantile	M209T/T354I	Deletion of 1 a.a.
10	c.1120G>A	V357M	p.V374M	Versailles lab oct. 2004	adult	V357M/E281K	
10	c.1130C>T	A360V	p.A377V	Mornet et al. 2001	perinatal	A360V/A360V	
10	c.1133A>T	D361V	p.D378V	Henthorn et al. 1992	infantile	E174K/D361V	1.2
10	c.1142A>G	H364R	p.H381R	Tailandier et al. 2000	infantile	A23V/H364R	
10	c.1144G>A	V365I	p.V382I	Goseki-Sone et al. 1998	childhood	F310L/V365I	0 11
10	c.1166C>T	T372I	p.T389I	Versailles lab oct. 2004	perinatal	T372/S351de	
10	c.1171C>T	R374C	p.R391C	Zurutuza et al. 1999	childhood	E174K/R374C	10.3 1
10	c.1172G>A	R374H	p.R391H	Orimo et al. 2002	childhood	R374H?	3.7 4
10	c.1172delC			Tailandier et al. 1999	infantile	M45L/c.1172d elC	na
10	c.1175G>C	G375A	p.G392A	Versailles lab. Jan. 2008	perinatal	G375A/R119C	Frameshift mutation
10	c.1182T>C	I378T	p.I396T	Versailles lab Jul. 2006	perinatal	I378T/E174K	
11	c.1195G>T	A382S	p.A399S	Tailandier et al. 2001	adult	E218G/A382S	
11	c.1196C>T	A382V	p.A398V	Spentchian et al. 2006	adult	A382V/A16V	
11	c.1199C>T	P383L	p.P400L	Spentchian et al. 2006	infantile	P383L/P383L	+
11	c.1214_1215de (CA			Versailles lab Jul. 2006	adult	C.1214_1215d elCA/E174K	Frameshift mutation
11	c.1216_1219de			Brun-Heath et al. 2005	perinatal	C.1216_1219d	

	[GACA						[GACA?/		
11	c.1217A>G	D389G	p.D406G	Taillandier et al. 2000	odont.		D389G/R433	14.9	5
11	c.1228T>C	F393L	p.F410L	Versailles lab oct. 2004	infantile		H		+
11	c.1231A>G	T394A	p.T411A	Brun-Heath et al. 2005	perinatal		F393L/E174K		-
11	c.1240C>A	L397M	p.L414M	Mumm et al. 2002	perinatal		T394A/c.926_927delTC	0.3	16
11	c.1250A>G	N400S	p.N417S	Sergi et al. 2001	perinatal		L397M/D277A		-
11	c.1256delC			Taillandier et al. 2000	perinatal		N400S/c.648+1G>A	3	unp. +
11	c.1258G>A	G403S	p.G420S	Glaser et al. 2004	perinatal		c.1256delC?		na
11	c.1268T>C	V406A	p.V423A	Taillandier et al. 2001	perinatal		G403S/G403S	0.4	unp. -
11	c.1270G>A	V407M	p.V424M	Versailles lab jan. 2007	adult		A89T/V406A	15.7	2
11	c.1276G>T	G409C	p.G426C	Mochizuki et al. 2000	infantile		V407M/V407M		-
11	c.1277G>A	G409D	p.G426D	Mumm et al. 2002	childhood		K207A/G409C	18.5	15
11	c.1282C>T	R411X	p.R428X	Taillandier et al. 1999	perinatal		G409D/E174K		-
11	c.1283G>C	R411P	p.R428P	Spentchian et al. 2006	perinatal		R411X/R411X		na
11	c.1285G>A	E412K	p.E429K	Versailles lab Jul. 2006	odont.		R411P/c.997+2T>A		Nonsense mutation
11	c.1306T>C	Y419H	p.Y436H	Henthorn et al. 1992	childhood		E412K?		
12	c.1333T>C	S428P	p.S445P	Mornet et al. 1998	infantile		A16W/Y419H		na
12	c.1349G>A	R433H	p.R450H	Taillandier et al. 2000	odont.		S428P/?	2.1	1
12	c.1348C>T	R433C	p.R450C	Mornet et al. 1998	infantile		D389G/R433H		-
12	c.1354G>A	E435K	p.E452K	Spentchian et al. 2003	perinatal		R433C/R433C	4.0	1
12	c.1361A>G	H437R	p.H454R	Versailles lab oct. 2003	childhood		A94T/E435K		+
							E174K/H437R		+

12	c.1363G>A	G438S	p.G455S	Draquett et al. 2004	adult	G438S/G474R	
12	c.1364G>A	G438D	p.G455D	Versailles lab jan. 2007	perinatal	G438D/G438D	-
12	c.1366G>T	G439W	p.G456W	Versailles lab oct. 2003	childhood	G439W/?	+
12	c.1366G>A	G439R	p.G456R	Ozono et al. 1996	infantile	G439R/?	1.5 unpub.
12	c.1375G>A	V442M	p.V459M	Taillandier et al. 2000	infantile	A34VN/V442M	+
12	c.1375G>T	V442L	p.V459L	Versailles lab oct. 2004	perinatal	V442L/E435K	-
12	c.1396C>T	P449L	p.P466L	Versailles lab oct. 2003	perinatal	P449L/?	+
12	c.1400T>C	M450T	p.M467T	Versailles lab oct. 2004	infantile	M450T/P290S	-
12	c.1402G>A	A451T	p.A468T	Spentchian et al. 2003	perinatal	A451T/A451T	+
12	c.1417G>A	G456S	p.G473S	Mornet et al. 1998	perinatal	A23V/G456S	+
12	c.1426G>A	E459K	p.E476K	Taillandier et al. 1999	perinatal	A94T/E459K	+
12	c.1427A>G	E459G	p.E476G	Mornet et al. 2001	perinatal	E459G/E459G	+
12	c.1433A>T	N461I	p.N478I	Taillandier et al. 2000	childhood	N461I/N	1.1
12	c.1444_1445inSC			Brun-Heath et al. 2005	perinatal	c.1444_1445i InsC/G317D	Frameshift mutation
12	c.1456G>C	C472S	p.C489S	Taillandier et al. 2000	perinatal	C472S/c.997+2T>A	9.4 5 -
12	c.1468A>T	I473F	p.I490F	Lia-Baldini et al. 2001	adult	I473F/?	37.1 3 -
12	c.1471G>A	G474R	p.G491R	Mornet et al. 1998	perinatal	G112R/G474R	-
12	c.1471delG			Brun-Heath et al. 2005	odontic	c.1471delG/R 119H	Frameshift mutation
12	c.1559delT			Orlino et al. 1994	infantile	E281K/c.1559delT	28 18 na
<b>Large deletions</b>							
				Spentchian et al. 2006	perinatal	Homozygote	
				Spentchian et al. 2006	infantile	Compound heterozygote with S164L	

In some embodiments, the sALP polypeptides of the invention do not include any of the mutations provided in Table 1. In particular, the sALP polypeptides of the invention, using the numbering of the consensus sequence of SEQ ID NO: 1218 (Fig. 10), the amino acid at position 22 is not a phenylalanine residue; the amino acid at position 33 (position 11 in the sequence without signal peptide) is not a cysteine residue; the amino acid at position 38 (position 16 in the sequence without signal peptide) is not a valine residue; the amino acid at position 42 (position 20 in the sequence without signal peptide) is not a proline residue; the amino acid at position 45 (position 23 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 56 (position 34 in the sequence without signal peptide) is not a serine or a valine residue; the amino acid residue at position 67 (position 5 45 in the sequence without signal peptide) is not a leucine, an isoleucine or a valine residue; the amino acid residue at position 68 (position 46 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 73 (position 51 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 76 (position 54 in the sequence without signal peptide) is not a cysteine, a serine, a proline or a histidine residue; the amino acid residue at position 77 (position 10 55 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 80 (position 58 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 81 (position 59 in the sequence without signal peptide) is not an asparagine residue; the amino acid residue at position 105 (position 83 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 113 (position 89 in the sequence without signal peptide) is not 15 a leucine residue; the amino acid residue at position 116 (position 94 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 117 (position 95 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 119 (position 97 in the sequence without signal peptide) is not a glycine residue; the amino acid residue at position 121 (position 20 99 in the sequence without signal peptide) is not a serine or a threonine residue; the amino acid residue at position 125 (position 103 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 128 (position 106 in the sequence without signal peptide) is not a aspartic acid residue; the amino acid residue at position 133 (position 111 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 134 (position 112 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 137 (position 115 in the sequence without signal peptide) is not a threonine or a valine residue; the amino acid residue at position 30 139 (position 117 in the sequence without signal peptide) is not a histidine or an asparagine residue; the amino acid residue at position 141 (position 119 in the sequence without signal peptide) is not a histidine residue; the amino acid residue at position 153 (position 131 in the sequence without signal peptide) is not an alanine or an isoleucine residue; the amino acid residue at position 167 (position 145 in the sequence without signal peptide) is not a serine or a valine residue; the amino acid residue at position 172 (position 150 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 175 (position 153 in the sequence without signal peptide) is not an aspartic acid residue; the

amino acid residue at position 176 (position 154 in the sequence without signal peptide) is not a tyrosine or an arginine residue; the amino acid residue at position 181 (position 159 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 182 (position 160 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 184 (position 162 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 186 (position 164 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 189 (position 167 in the sequence without signal peptide) is not a tryptophan residue; the amino acid residue at position 194 (position 172 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 196 (position 174 in the sequence without signal peptide) is not 5 a lysine or a glycine residue; the amino acid residue at position 197 (position 175 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 198 (position 176 in the sequence without signal peptide) is not an alanine residue; the amino acid residue at position 206 (position 184 in the sequence without signal peptide) is not a tyrosine residue; the amino acid residue at position 208 (position 186 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 207 (position 190 in the sequence without signal peptide) is not a proline residue; 10 the amino acid residue at position 216 (position 194 in the sequence without signal peptide) is not an aspartic acid residue; the amino acid residue at position 217 (position 195 in the sequence without signal peptide) is not a phenylalanine residue; the amino acid residue at position 223 (position 201 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 225 15 (position 203 in the sequence without signal peptide) is not a valine or an alanine residue; the amino acid residue at position 226 (position 204 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 228 (position 206 in the sequence without signal peptide) is not a tryptophan or a glutamine residue; the amino acid residue at position 229 (position 207 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 231 (position 209 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 240 20 (position 218 in the sequence without signal peptide) is not a glycine residue; the amino acid residue at position 251 (position 229 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 254 (position 232 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 269 (position 247 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 277 (position 255 in the sequence without signal peptide) is not 25 a cysteine, a leucine or a histidine residue; the amino acid residue at position 280 (position 258 in the sequence without signal peptide) is not a proline residue; the amino acid residue at position 295 (position 273 in the sequence without signal peptide) is not a phenylalanine residue; the amino acid residue at position 297 (position 275 in the sequence without signal peptide) is not a lysine residue; the amino acid residue at position 298 (position 276 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 300 (position 278 in the sequence without signal peptide) is not a tyrosine or an alanine residue; the amino acid residue at position 301 (position 279 in the sequence 30

without signal peptide) is not a valine, a threonine or an isoleucine residue; the amino acid residue at position 303 (position 281 in the sequence without signal peptide) is not an aspirate residue; the amino acid residue at position 304 (position 282 in the sequence without signal peptide) is not a lysine residue; the amino acid residue at position 305 (position 283 in the sequence without signal peptide) is not a proline residue; the amino acid residue at position 312 (position 290 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 313 (position 291 in the sequence without signal peptide) is not a serine or a leucine residue; the amino acid residue at position 317 (position 295 in the sequence without signal peptide) is not a lysine residue; the amino acid residue at position 332 (position 310 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 333 (position 311 in the sequence without signal peptide) is not a cysteine, a glycine or a leucine residue; the amino acid residue at position 334 (position 312 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 340 (position 318 in the sequence without signal peptide) is not an aspartic acid residue; the amino acid residue at position 345 (position 323 in the sequence without signal peptide) is not an arginine or a glutamate residue; the amino acid residue at position 354 (position 332 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 360 (position 338 in the sequence without signal peptide) is not an aspartic acid residue; the amino acid residue at position 361 (position 339 in the sequence without signal peptide) is not a threonine or an isoleucine residue; the amino acid residue at position 377 (position 355 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 380 (position 358 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 383 (position 361 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 384 (position 362 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 387 (position 365 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 388 (position 366 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 395 (position 373 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 397 (position 375 in the sequence without signal peptide) is not a cysteine or a histidine residue; the amino acid residue at position 398 (position 376 in the sequence without signal peptide) is not an alanine residue; the amino acid residue at position 401 (position 379 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 405 (position 383 in the sequence without signal peptide) is not a serine or a valine residue; the amino acid residue at position 406 (position 384 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 412 (position 390 in the sequence without signal peptide) is not a glycine residue; the amino acid residue at position 416 (position 394 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 417 (position 395 in the sequence without signal peptide) is not an alanine residue; the amino acid residue at position 420 (position 398 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 423 (position 401 in the sequence without signal peptide) is not a serine residue.

residue; the amino acid residue at position 426 (position 404 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 429 (position 407 in the sequence without signal peptide) is not an alanine residue; the amino acid residue at position 430 (position 408 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 432 (position 410 in the sequence without signal peptide) is not a cysteine or an aspartic acid residue; amino acid residue at position 434 (position 412 in the sequence without signal peptide) is not a proline residue; amino acid residue at position 435 (position 413 in the sequence without signal peptide) is not a lysine residue; amino acid residue at position 442 (position 420 in the sequence without signal peptide) is not a histidine residue; amino acid residue at position 451 (position 429 in the sequence without signal peptide) is not a proline residue; amino acid residue at position 456 (position 434 in the sequence without signal peptide) is not a histidine or a cysteine residue; amino acid residue at position 458 (position 436 in the sequence without signal peptide) is not a lysine residue; amino acid residue at position 460 (position 438 in the sequence without signal peptide) is not an arginine residue; amino acid residue at position 461 (position 439 in the sequence without signal peptide) is not a serine or an aspartic acid residue; amino acid residue at position 462 (position 440 in the sequence without signal peptide) is not a tryptophan or an arginine residue; amino acid residue at position 465 (position 443 in the sequence without signal peptide) is not a methionine or a leucine residue; amino acid residue at position 472 (position 450 in the sequence without signal peptide) is not a leucine residue; amino acid residue at position 473 (position 451 in the sequence without signal peptide) is not a threonine residue; amino acid residue at position 474 (position 452 in the sequence without signal peptide) is not a threonine residue; amino acid residue at position 479 (position 457 in the sequence without signal peptide) is not a serine residue; amino acid residue at position 482 (position 460 in the sequence without signal peptide) is not a lysine or a glycine residue; amino acid residue at position 484 (position 462 in the sequence without signal peptide) is not a leucine residue; amino acid residue at position 495 (position 473 in the sequence without signal peptide) is not a serine residue; amino acid residue at position 496 (position 474 in the sequence without signal peptide) is not a phenylalanine residue; and amino acid residue at position 497 (position 475 in the sequence without signal peptide) is not an arginine residue.

Also more specifically, when a sTNALP is used in the bone targeted sALPs of the present invention, using the numbering of the human TNALP sequence, the amino acid at position 17 is not a phenylalanine residue; the amino acid at position 28 (position 11 in the sequence without signal peptide) is not a cysteine residue; the amino acid at position 33 (position 16 in the sequence without signal peptide) is not a valine residue; the amino acid at position 37 (position 20 in the sequence without signal peptide) is not a proline residue; the amino acid at position 40 (position 23 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 51 (position 34 in the sequence without signal peptide) is not a serine or a valine residue; the amino acid residue at position 62 (position 45 in the sequence without signal peptide) is not a leucine, an isoleucine or a valine residue; the amino acid residue at position 63 (position 46 in the sequence without signal peptide) is not a valine residue; the amino acid

residue at position 68 (position 51 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 71 (position 54 in the sequence without signal peptide) is not a cysteine, a serine, a proline or a histidine residue; the amino acid residue at position 72 (position 55 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 75 (position 58 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 76 (position 59 in the sequence without signal peptide) is not an asparagine residue; the amino acid residue at position 100 (position 83 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 108 (position 89 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 111 (position 94 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 112 (position 95 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 114 (position 97 in the sequence without signal peptide) is not a glycine residue; the amino acid residue at position 116 (position 99 in the sequence without signal peptide) is not a serine or a threonine residue; the amino acid residue at position 120 (position 103 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 123 (position 106 in the sequence without signal peptide) is not a aspartic acid residue; the amino acid residue at position 128 (position 111 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 129 (position 112 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 132 (position 115 in the sequence without signal peptide) is not a threonine or a valine residue; the amino acid residue at position 134 (position 117 in the sequence without signal peptide) is not a histidine or an asparagine residue; the amino acid residue at position 136 (position 119 in the sequence without signal peptide) is not a histidine residue; the amino acid residue at position 148 (position 131 in the sequence without signal peptide) is not an alanine or an isoleucine residue; the amino acid residue at position 162 (position 145 in the sequence without signal peptide) is not a serine or a valine residue; the amino acid residue at position 167 (position 150 in the sequence without signal peptide) is not a methionine residue; the amino acid residue at position 170 (position 153 in the sequence without signal peptide) is not an aspartic acid residue; the amino acid residue at position 171 (position 154 in the sequence without signal peptide) is not a tyrosine or an arginine residue; the amino acid residue at position 176 (position 159 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 177 (position 160 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 179 (position 162 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 181 (position 164 in the sequence without signal peptide) is not a leucine residue; the amino acid residue at position 184 (position 167 in the sequence without signal peptide) is not a tryptophan residue; the amino acid residue at position 189 (position 172 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 191 (position 174 in the sequence without signal peptide) is not a lysine or a glycine residue; the amino acid residue at position 192 (position 175 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 193 (position 176 in the sequence without signal peptide) is not a threonine residue.

the sequence without signal peptide) is not an alanine residue; the amino acid residue at position 201 (position 184 in the sequence without signal peptide) is not a tyrosine residue; the amino acid residue at position 203 (position 186 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 207 (position 190 in the sequence without signal peptide) is not a proline residue;

5 the amino acid residue at position 211 (position 194 in the sequence without signal peptide) is not a aspartic acid residue; the amino acid residue at position 212 (position 195 in the sequence without signal peptide) is not a phenylalanine residue; the amino acid residue at position 218 (position 201 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 220 (position 203 in the sequence without signal peptide) is not a valine or an alanine residue; the amino acid

10 residue at position 221 (position 204 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 223 (position 206 in the sequence without signal peptide) is not a tryptophan or a glutamine residue; the amino acid residue at position 224 (position 207 in the sequence without signal peptide) is not a glutamate residue; the amino acid residue at position 226 (position 209 in the sequence without signal peptide) is not a threonine residue; the amino acid residue at position 235

15 (position 218 in the sequence without signal peptide) is not a glycine residue; the amino acid residue at position 246 (position 229 in the sequence without signal peptide) is not a serine residue; the amino acid residue at position 249 (position 232 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 264 (position 247 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 272 (position 255 in the sequence without signal peptide) is

20 not a cysteine, a leucine or a histidine residue; the amino acid residue at position 275 (position 258 in the sequence without signal peptide) is not a proline residue; the amino acid residue at position 289 (position 272 in the sequence without signal peptide) is not a phenylalanine residue; the amino acid residue at position 291 (position 274 in the sequence without signal peptide) is not a lysine residue; the amino acid residue at position 292 (position 275 in the sequence without signal peptide) is not a threonine residue;

25 the amino acid residue at position 294 (position 277 in the sequence without signal peptide) is not a tyrosine or an alanine residue; the amino acid residue at position 295 (position 278 in the sequence without signal peptide) is not a valine, a threonine or an isoleucine residue; the amino acid residue at position 297 (position 280 in the sequence without signal peptide) is not an aspirate residue; the amino acid residue at position 298 (position 281 in the sequence without signal peptide) is not a lysine residue;

30 the amino acid residue at position 299 (position 282 in the sequence without signal peptide) is not a proline residue; the amino acid residue at position 306 (position 289 in the sequence without signal peptide) is not a valine residue; the amino acid residue at position 307 (position 290 in the sequence without signal peptide) is not a serine or a leucine residue; the amino acid residue at position 311 (position 294 in the sequence without signal peptide) is not a lysine residue; the amino acid residue at

35 position 326 (position 309 in the sequence without signal peptide) is not an arginine residue; the amino acid residue at position 327 (position 310 in the sequence without signal peptide) is not a cysteine, a glycine or a leucine residuc; the amino acid residue at position 328 (position 311 in the sequence without



without signal peptide) is not a lysine residue; amino acid residue at position 454 (position 437 in the sequence without signal peptide) is not an arginine residue; amino acid residue at position 455 (position 438 in the sequence without signal peptide) is not a serine or an aspartic acid residue; amino acid residue at position 456 (position 439 in the sequence without signal peptide) is not a tryptophan or an arginine 5 residue; amino acid residue at position 459 (position 442 in the sequence without signal peptide) is not a methionine or a leucine residue; amino acid residue at position 466 (position 449 in the sequence without signal peptide) is not a leucine residue; amino acid residue at position 467 (position 450 in the sequence without signal peptide) is not a threonine residue; amino acid residue at position 468 (position 451 in the sequence without signal peptide) is not a threonine residue; amino acid residue at position 473 (position 10 456 in the sequence without signal peptide) is not a serine residue; amino acid residue at position 476 (position 459 in the sequence without signal peptide) is not a lysine or a glycine residue; amino acid residue at position 478 (position 461 in the sequence without signal peptide) is not a leucine residue; amino acid residue at position 489 (position 472 in the sequence without signal peptide) is not a serine residue; amino acid residue at position 490 (position 473 in the sequence without signal peptide) is not a 15 phenylalanine residue; and amino acid residue at position 491 (position 474 in the sequence without signal peptide) is not an arginine residue. In other specific embodiments, one or more Xs are defined as being any of the amino acid residues found at that position in the sequences of the alignment or a residue that constitutes a conserved or semi-conserved substitution of any of these amino acid residues. In other specific embodiments, X's are defined as being any of the amino acid residues found at that position in 20 the sequences of the alignment. For instance, the amino acid residue at position 51 (position 34 in the sequence without signal peptide) is an alanine or a valine residue; the amino acid residue at position 177 (position 160 in the sequence without signal peptide) is an alanine or a serine residue; the amino acid residue at position 212 (position 195 in the sequence without signal peptide) is an isoleucine or a valine residue; the amino acid residue at position 291 (position 274 in the sequence without signal peptide) is a 25 glutamic acid or an aspartic acid residue; and the amino acid residue at position 374 (position 357 in the sequence without signal peptide) is a valine or an isoleucine residue.

An sALP may optionally be glycosylated at any appropriate one or more amino acid residues.

In addition, an sALP may have at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 30 more) sequence identity to any of the sALPs described herein.

An sALP may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more additions, deletions, or substitutions relative to any of the sALPs described herein.

#### NPs

35 Any natriuretic peptide or variant thereof that is an agonist of natriuretic peptide receptor B (“NPR-B”), e.g., human NPR-B, may be used in any of the methods and compositions described herein.

Natriuretic peptides as described herein are peptides that are capable of agonizing NPR-B. Natriuretic peptides include atrial natriuretic peptide (ANP), brain natriuretic peptide (BNP), and C-type natriuretic peptide (CNP). These peptides bind to three types of receptors that signal intracellularly to modulate physiological functions. ANP and BNP bind preferentially to natriuretic peptide receptor A (NPR-A) (also known as guanylyl cyclase A (GC-A)), and CNP binds preferentially to natriuretic peptide receptor B (NPR-B) (also known as guanylyl cyclase B (GC-B)). All three peptides have similar affinity for natriuretic peptide receptor C (NPR-C), which has both signaling and peptide clearance functions. 5 Clearance of natriuretic peptides also occurs through the action of membrane-bound neutral endopeptidase (NEP). Peptide binding to NPR-A or NPR-B activates the intracellular guanylyl cyclase domain of these receptors, which produces the second messenger cGMP. cGMP activates or inhibits 10 multiple signaling pathways inside the cell.

15 Natriuretic peptides, including CNP, which primarily agonizes NPR-B, and ANP and BNP, which primarily agonize NPR-A, have important roles in multiple biological processes. Multiple sequence alignments of various NP family members and consensus sequences are shown in Figs. 12-14 and 15A-15G.

A key downstream effect of CNP22 and CNP53, and variants thereof as described herein, in agonizing NPR-B is endochondral ossification. Thus, the NPs described herein are useful, e.g., for treating a wide array of disorders associated with overactivation of FGFR3 and vascular smooth muscle disorders.

20 NPs include the schematic structure shown in Fig. 16, wherein the ring domain is required and each of the N-terminal extension, short segment, and C-terminal extension is optional. The ring domain is 17 amino acid residues long, with cysteine residues at each terminus of the ring domain (positions 1 and 17) that form a disulfide bond. In some embodiments, the ring domain has an amino acid sequence that falls within one of the consensus sequences shown in Figs. 12, 14, or 15A-15G (SEQ ID NO: 6, 25 amino acid residues 11-27 of SEQ ID NO: 30, or SEQ ID NO: 95, respectively). Any of the ring domains shown in Figs. 12-14 and 15A-15G may be used in an NP as described herein.

30 The short segment is a segment immediately N-terminal to the ring domain that is between 0 and 10 amino acid residues (e.g., 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 amino acid residues) in length. Exemplary short segments are shown immediately N-terminal to the boxed region in Fig. 12 or Fig. 14, e.g., residues 1-5 of SEQ ID NO: 4, or the 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 amino acid residues immediately N-terminal to the conserved ring domain in any of the species shown in Figs. 14 and 15A-15G. In some embodiments, the short segment consists of the 5-amino acid portion immediately N-terminal to the conserved ring domain in any of the species shown in Figs. 12, 14, or 15A-15G. In some embodiments, the short segment confers increased selectivity for NPR-B relative to NPR-A.

35 The N-terminal extension is a region immediately N-terminal to the short segment (if the short segment is present) or the ring domain (if the short segment is not present) and may be of any length, e.g., 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30,

31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, or even more amino acid residues. This region is absent in CNP22 but is present in CNP53 (residues 1-31 of SEQ ID NO: 11). Exemplary N-terminal extensions are the 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 5 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 110, 120, 130, 140, 150, 160, 170, 10 180, 190, 200, 225, 250, or more residues immediately N-terminal to the short segment, e.g., of 5 amino acid residues (if short segment is present), or immediately N-terminal to the ring domain (if short segment is not present), of any of the species shown in Figs. 15A-15G. In some embodiments, the N-terminal extension provides increased selectivity for NPR-B relative to NPR-A.

The C-terminal extension is a region immediately C-terminal to the ring domain and may be of any length, e.g., 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 55, 60, 65, 70, 15 75, 80, 85, 90, 95, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, or even more amino acid residues. This region is absent in CNP22 and CNP53 but is present in the hybrid peptide CDNP (SEQ ID NO: 100, Fig. 24). Additional variants of CDNP include those that have one or more mutations that provide reduced NEP degradation, such as those provided in Fig. 24: CDNP-N1 (SEQ ID NO: 101), CDNP-G1 (SEQ ID NO: 102), CDNP-H1 (SEQ ID NO: 103), and CDNP-K1 (SEQ ID NO: 104).

20 Exemplary C-terminal extensions are shown immediately C-terminal to the boxed region in Fig. 12, e.g., amino acid residues 24-28 of SEQ ID NO: 1, amino acid residues 28-32 of SEQ ID NO: 2, amino acid residues 27-32 of SEQ ID NO: 3, or amino acid residues 24-38 of SEQ ID NO: 5. In some embodiments, the C-terminal tail includes, or consists of, the DNP C-terminal tail (SEQ ID NO: 117), or a variant thereof having one or more addition, deletion, or substitution mutations (e.g., SEQ ID NO: 118). 25 For example, a C-terminal tail of an NP may include any of the DNP C-terminal tail mutations shown in Fig. 24. In particular, residues 1, 3, 4, 5, 6, and/or 7 of the DNP C-terminal tail (SEQ ID NO: 117) may be mutated, e.g., as in any of the mutations shown in Fig. 24. In some embodiments, the C-terminal extension confers increased selectivity for NPR-B relative to NPR-A.

An NP may optionally be glycosylated at any appropriate one or more amino acid residues. 30 In addition, an NP may have at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to any of the NPs described herein, or to one or more of the ring domain, the short segment, the C-terminal extension, or the N-terminal extension.

An NP may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more additions, deletions, or substitutions relative 35 to any of the NPs described herein, or to one or more of the ring domain, the short segment, the C-terminal extension, or the N-terminal extension.

In one example, the NP can have one or more mutations that are less sensitive to oxidation without substantially reducing potency or efficacy. For example, residue 17 of CNP22 is one of the less well-conserved positions in CNP22, with naturally-occurring homologs having (without limitation) Phe, Leu, Ile, Thr, Val, or Ser at this position (see, e.g., Fig. 14). Exemplary CNP22 variants include CNP-5 F17 (SEQ ID NO: 119); CNP-L17 (SEQ ID NO: 120); CNP-I17 (SEQ ID NO: 121); CNP-T17 (SEQ ID NO: 122); CNP-V17 (SEQ ID NO: 123); CNP-A17 (SEQ ID NO: 124); CNP-S17 (SEQ ID NO: 125); CNP-E17 (SEQ ID NO: 156); CNP-R17 (SEQ ID NO: 157); and CNP-Y17 (SEQ ID NO: 158), where the consensus sequence is shown in SEQ ID NO: 126 (where X can be any amino acid, including, without limitation, Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Val, Ala, or Ser) (Fig. 26). Additional 10 exemplary CNP22 variants include those having a point mutation at position 17, as shown in Fig. 30 (SEQ ID NOs: 126, 119-122, and 156-172), where X in SEQ ID NOs: 126 or 162 can be any amino acid, e.g., F, L, I, T, E, R, Y, C, P, or D.

In another example, the NP can have one or more mutations that provide increased resistance to one or more enzymes that cleave CNP *in vivo* (e.g., neutral endopeptidase (NEP) and/or insulin degrading 15 enzyme (IDE)). Exemplary molecules are shown in Fig. 27 (SEQ ID NOs: 127-150), Figs. 28A-28E (SEQ ID NOs: 1001-1155), and Fig. 29 (SEQ ID NOs: 4, 115, 119, 120, 122, 128-139, 147, 148, and 150-155).

An NP as described herein may include any other sequence or moiety, attached covalently or non-covalently, provided that the NP has the ability to agonize NPR-B.

20 In some embodiments, an NP as described herein may be no more than 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 70, 80, 90, 100, 110, or 120 amino acid residues in length. Furthermore, in some embodiments, an NP as described herein may be no more than 1.8, 2.0, 2.2, 2.4, 2.6, 2.8, 3.0, 3.2, 3.4, 3.6, 3.8, 4.0, 4.2, 4.4, 4.6, 4.8, 5.0, 5.2, 5.4, 5.6, 5.8, 6.0, 6.2, 6.4, 6.6, 6.8, 7.0, 7.2, 25 7.4, 7.6, 7.8, 8.0, 8.2, 8.4, 8.6, 8.8, 9.0, 9.2, 9.4, 9.6, 9.8, or 10.0 kilodaltons (kDa) in molecular weight.

NPs that are suitable for use in the compositions and methods described herein include those described, e.g., in U.S. Patent Nos. 5,352,770; 5,434,133; 6,020,168; 6,034,231; 6,407,211; 6,743,425; 6,818,619; 7,276,481; 7,384,917; and 7,754,852; U.S. Application Pub. Nos. 2007-0197434; 2008-0181903; 2008-0312142; 2009-0170756; 2010-0055150; and 2010-0297021; International Application 30 Pub. Nos. WO 94/20534; WO 02/047871; WO 2004/047871; WO 2005/098490; WO 2008/154226; and WO 2009/067639; European Application Pub. Nos. EP 0497368 and EP 0466174; Furuya et al., Biochem. Biophys. Res. Comm. 183: 964-969 (1992); Takano et al., Zool. Sci., 11: 451-454 (1994); Plater et al., Toxicol., 36(6): 847-857 (1998); and Inoue et al., Proc. Nat. Acad. Sci., 100(17): 10079-10084 (2003), each of which is hereby incorporated by reference in its entirety, including all formulas, 35 structures, and sequences for natriuretic peptides and variants thereof. In alternative embodiments, the NPs referenced in the present paragraph are excluded from the compositions and methods described herein.

In some embodiments, any of the NPs described or incorporated by reference herein may be used in the compositions and methods described herein without fusion to an Fc domain or to a linker, or alternatively may be fused to any of the linkers described herein but not to an Fc domain. Such NPs may be used to treat a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, e.g., 5 achondroplasia, a bone or cartilage disorder, a vascular smooth muscle disorder, or a condition for elongation of bone, as described herein.

In other embodiments, any of the NPs described or incorporated by reference herein may include a point mutation at position 17 relative to CNP22. Wild-type CNP22 has a methionine at position 17 relative to CNP22, which can be oxidized *in vivo* and/or which can provide a peptide that is degradable 10 by a protease. As described herein, point mutations at position 17 relative to CNP22 could provide polypeptides having decreased degradation, while maintaining potency. Exemplary amino acid residues at position 17 relative to CNP22 are Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, Asp, Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys, e.g., Phe, Leu, Ile, Thr, Glu, Arg, Tyr, Cys, Pro, or Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu. For example, the amino acid at position 17 relative to CNP22 could be Phe, Leu, Ile, Thr, 15 Glu, Arg, Tyr, Cys, Pro, and Asp, e.g., Phe or Leu, e.g., Phe, e.g., Leu. In another example, the amino acid at position 17 relative to CNP22 could be Phe, Leu, Ile, Thr, Val, Ala, or Ser. Alternatively, exemplary amino acid residues at position 17 relative to CNP22 are Gly, Ala, Ser, Val, Trp, Asn, Gln, His, or Lys.

Furthermore, included in the compositions and methods described herein are nucleic acid 20 molecules encoding any of the NPs and fusion polypeptides described herein, as well as nucleic acid molecules that hybridize under high stringency conditions to at least a portion, e.g., to 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or even 100%, of a nucleic acid molecule that encodes any of the NPs or fusion polypeptides described herein.

## 25 **Fragment crystallizable region (Fc) fragments**

The fusion polypeptides of the invention may include an N-terminal or C-terminal domain such as Fc, a fragment crystallizable region of an immunoglobulin. For example, an sALP polypeptide and/or an NP polypeptide of the invention can be a fusion polypeptide including an Fc. An immunoglobulin molecule has a structure that is well known in the art. It includes two light chains (~23 kD each) and two 30 heavy chains (~50-70 kD each) joined by inter-chain disulfide bonds. Immunoglobulins are readily cleaved proteolytically (e.g., by papain cleavage) into Fab (containing the light chain and the VH and CH1 domains of the heavy chain) and Fc (containing the C<sub>H2</sub> and C<sub>H3</sub> domains of the heavy chain, along with adjoining sequences). Cleavage typically occurs in a flexible hinge region joining the Fab and Fc regions. For example, papain cleaves the hinge region immediately before the disulfide bonds joining the 35 two heavy chains.

Useful Fc fragments as described herein include the Fc fragment of any immunoglobulin molecule, including IgG, IgM, IgA, IgD, or IgE, and their various subclasses (e.g., IgG-1, IgG-2, IgG-3,

IgG-4, IgA-1, IgA-2), taken from any mammal (e.g., human). The Fc fragments of the invention may include, for example, the C<sub>H2</sub> and C<sub>H3</sub> domains of the heavy chain, as well as any portion of the hinge region. Furthermore, the Fc region may optionally be glycosylated at any appropriate one or more amino acid residues, e.g., various amino acid residues known to those skilled in the art. In some embodiments, 5 the Fc fragment is of human IgG-1. In particular embodiments, the Fc fragment of the fusion polypeptide has the amino acid sequence of SEQ ID NO: 401, or has at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to SEQ ID NO: 401 (Fig. 7).

In some embodiments, engineered, e.g., non-naturally occurring, Fc regions may be utilized in 10 the compositions and methods of the invention, e.g., as described in International Application Pub. No. WO2005/007809, which is hereby incorporated by reference.

An Fc fragment as described herein may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 50, or more additions, deletions, or substitutions relative to any of the Fc fragments described herein.

15

### Linkers

The fusion proteins described herein may include a peptide linker region between the Fc fragment and the sALP or between the Fc fragment and the NP. In addition, a peptide linker region may be included between the Fc fragment and the optional bone-targeting moiety. The linker region may be 20 of any sequence and length that allows the sALP or the NP to remain biologically active, e.g., not sterically hindered. Exemplary linker lengths are between 1 and 200 amino acid residues, e.g., 1-5, 6-10, 11-15, 16-20, 21-25, 26-30, 31-35, 36-40, 41-45, 46-50, 51-55, 56-60, 61-65, 66-70, 71-75, 76-80, 81-85, 86-90, 91-95, 96-100, 101-110, 111-120, 121-130, 131-140, 141-150, 151-160, 161-170, 171-180, 181-190, or 191-200 amino acid residues. Additional exemplary linker lengths are 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, or 200 amino acid residues. Additional exemplary linker lengths are 14-18, 20-24, 26-30, 32-36, 38-42, and 44-48 amino acid 30 residues.

In some embodiments, linkers include or consist of flexible portions, e.g., regions without significant fixed secondary or tertiary structure. Exemplary flexible linkers are glycine-rich linkers, e.g., containing at least 50%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or even 100% glycine residues. Linkers may also contain, e.g., serine residues. In some cases, the amino acid sequence of linkers 35 consists only of glycine and serine residues.

In some cases, the amino acid sequence of the linker sequence includes or consists of a sequence according to the formula  $[(\text{Gly})_m(\text{Ser})]_n(\text{Gly})_p$ , where each of m, n, and p is, independently, 0, 1, 2, 3, 4, 5,

6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20. In some embodiments, m = 1, 2, 3, 4, 5, or 6; n = 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10; and p = 0, 1, 2, 3, or 4. Alternatively, the linker sequence includes or consists of a sequence according to the formula  $(\text{Gly})_p[(\text{Ser})(\text{Gly})_m]_n$ , where each of m, n, and p is, independently, 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20. In some embodiments, m = 1, 2, 3, 4, 5, or 6; n = 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10; and p = 0, 1, 2, 3, or 4.

5 Exemplary combinations of m, n, and p values for either of the preceding two formulae are listed in Table 2.

A

Table 2.

m	n	p
N/A	0	1
N/A	0	2
N/A	0	3
N/A	0	4
N/A	0	5
N/A	0	6
N/A	0	7
N/A	0	8
N/A	0	9
N/A	0	10
1	1	0
1	2	0
1	3	0
1	4	0
1	5	0
1	6	0
1	7	0
1	8	0
1	9	0
1	10	0
2	1	0
2	2	0
2	3	0
2	4	0
2	5	0
2	6	0
2	7	0
2	8	0
2	9	0
2	10	0
3	1	0
3	2	0
3	3	0
3	4	0
3	5	0
3	6	0
3	7	0
3	8	0
3	9	0
3	10	0
4	1	0
4	2	0
4	3	0
4	4	0
4	5	0
4	6	0
4	7	0
4	8	0
4	9	0
4	10	0
5	1	0
5	2	0
5	3	0
5	4	0
5	5	0
5	6	0
5	7	0
5	8	0
5	9	0
5	10	0
6	1	0
6	2	0
6	3	0
6	4	0
6	5	0
6	6	0
6	7	0
6	8	0
6	9	0
6	10	0
7	1	0
7	2	0
7	3	0
7	4	0
7	5	0
7	6	0
7	7	0
7	8	0
7	9	0
7	10	0
8	1	0
8	2	0
8	3	0
8	4	0
8	5	0
8	6	0
8	7	0
8	8	0
8	9	0
8	10	0
9	1	0
9	2	0
9	3	0
9	4	0
9	5	0
9	6	0
9	7	0
9	8	0
9	9	0
9	10	0
10	1	0
10	2	0
10	3	0
10	4	0
10	5	0
10	6	0
10	7	0
10	8	0
10	9	0
10	10	0
11	1	0
11	2	0
11	3	0
11	4	0
11	5	0
11	6	0
11	7	0
11	8	0
11	9	0
11	10	0
12	1	0
12	2	0
12	3	0
12	4	0
12	5	0
12	6	0
12	7	0
12	8	0
12	9	0
12	10	0
13	1	0
13	2	0
13	3	0
13	4	0
13	5	0
13	6	0
13	7	0
13	8	0
13	9	0
13	10	0
14	1	0
14	2	0
14	3	0
14	4	0
14	5	0
14	6	0
14	7	0
14	8	0
14	9	0
14	10	0
15	1	0
15	2	0
15	3	0
15	4	0
15	5	0
15	6	0
15	7	0
15	8	0
15	9	0
15	10	0
16	1	0
16	2	0
16	3	0
16	4	0
16	5	0
16	6	0
16	7	0
16	8	0
16	9	0
16	10	0
17	1	0
17	2	0
17	3	0
17	4	0
17	5	0
17	6	0
17	7	0
17	8	0
17	9	0
17	10	0
18	1	0
18	2	0
18	3	0
18	4	0
18	5	0
18	6	0
18	7	0
18	8	0
18	9	0
18	10	0
19	1	0
19	2	0
19	3	0
19	4	0
19	5	0
19	6	0
19	7	0
19	8	0
19	9	0
19	10	0
20	1	0
20	2	0
20	3	0
20	4	0
20	5	0
20	6	0
20	7	0
20	8	0
20	9	0
20	10	0
21	1	0
21	2	0
21	3	0
21	4	0
21	5	0
21	6	0
21	7	0
21	8	0
21	9	0
21	10	0
22	1	0
22	2	0
22	3	0
22	4	0
22	5	0
22	6	0
22	7	0
22	8	0
22	9	0
22	10	0
23	1	0
23	2	0
23	3	0
23	4	0
23	5	0
23	6	0
23	7	0
23	8	0
23	9	0
23	10	0
24	1	0
24	2	0
24	3	0
24	4	0
24	5	0
24	6	0
24	7	0
24	8	0
24	9	0
24	10	0
25	1	0
25	2	0
25	3	0
25	4	0
25	5	0
25	6	0
25	7	0
25	8	0
25	9	0
25	10	0
26	1	0
26	2	0
26	3	0
26	4	0
26	5	0
26	6	0
26	7	0
26	8	0
26	9	0
26	10	0
27	1	0
27	2	0
27	3	0
27	4	0
27	5	0
27	6	0
27	7	0
27	8	0
27	9	0
27	10	0
28	1	0
28	2	0
28	3	0
28	4	0
28	5	0
28	6	0
28	7	0
28	8	0
28	9	0
28	10	0
29	1	0
29	2	0
29	3	0
29	4	0
29	5	0
29	6	0
29	7	0
29	8	0
29	9	0
29	10	0
30	1	0
30	2	0
30	3	0
30	4	0
30	5	0
30	6	0
30	7	0
30	8	0
30	9	0
30	10	0
31	1	0
31	2	0
31	3	0
31	4	0
31	5	0
31	6	0
31	7	0
31	8	0
31	9	0
31	10	0
32	1	0
32	2	0
32	3	0
32	4	0
32	5	0
32	6	0
32	7	0
32	8	0
32	9	0
32	10	0
33	1	0
33	2	0
33	3	0
33	4	0
33	5	0
33	6	0
33	7	0
33	8	0
33	9	0
33	10	0
34	1	0
34	2	0
34	3	0
34	4	0
34	5	0
34	6	0
34	7	0
34	8	0
34	9	0
34	10	0
35	1	0
35	2	0
35	3	0
35	4	0
35	5	0
35	6	0
35	7	0
35	8	0
35	9	0
35	10	0
36	1	0
36	2	0
36	3	0
36	4	0
36	5	0
36	6	0
36	7	0
36	8	0
36	9	0
36	10	0
37	1	0
37	2	0
37	3	0
37	4	0
37	5	0
37	6	0
37	7	0
37	8	0
37	9	0
37	10	0
38	1	0
38	2	0
38	3	0
38	4	0
38	5	0
38	6	0
38	7	0
38	8	0
38	9	0
38	10	0
39	1	0
39	2	0
39	3	0
39	4	0
39	5	0
39	6	0
39	7	0
39	8	0
39	9	0
39	10	0
40	1	0
40	2	0
40	3	0
40	4	0
40	5	0
40	6	0
40	7	0
40	8	0
40	9	0
40	10	0
41	1	0
41	2	0
41	3	0
41	4	0
41	5	0
41	6	0
41	7	0
41	8	0
41	9	0
41	10	0
42	1	0
42	2	0
42	3	0
42	4	0
42	5	0
42	6	0
42	7	0
42	8	0
42	9	0
42	10	0
43	1	0
43	2	0
43	3	0
43	4	0
43	5	0
43	6	0
43	7	0
43	8	0
43	9	0
43	10	0
44	1	0
44	2	0
44	3	0
44	4	0
44	5	0
44	6	0
44	7	0
44	8	0
44	9	0
44	10	0
45	1	0
45	2	0
45	3	0
45	4	0
45	5	0
45	6	0
45	7	0
45	8	0
45	9	0
45	10	0
46	1	0
46	2	0
46	3	0
46	4	0
46	5	0
46	6	0
46	7	0
46	8	0
46	9	0
46	10	0
47	1	0

In some embodiments, the amino acid sequence of the linker (e.g., between the Fc and the sALP, or between the Fc and the NP, or between the Fc and the optional bone-targeting moiety) includes or consists of a sequence in Table 3.

5

**Table 3.**

Linker sequence	SEQ ID NO.
G	301
GG	302
GGG	303
GGGG	304
GGGS	305
GGGS	306
GGGGSGGGGSGGGG	307
GGGGSGGGGSGGGGSG	308
GGGGSGGGGSGGGGSGGGGSGGG	309
GGGGSGGGGSGGGGSGGGGSGGGGGSGGG	310
GGGGSGGGGSGGGGSGGGGSGGGGGSGGGGGSGGG	311
GGGGSGGGGSGGGGSGGGGSGGGGSGGGGSGGGGS	312
GGGGSGGGGSGGGGSGGGGSGGGGSGGGGSGGGGS	313
KGANKK	314
KGANQK	315
KGANKQ	316
KGANQQ	317
QGANKK	318
QGANQK	319
QGANQK	320
QGANQQ	321
GGGGSGGGGSKGANKK	322
GGGGSGGGGSKGANQK	323
GGGGSGGGGSKGANQ	324
GGGGSGGGGSKGANQQ	325
GGGGSGGGGSQGANKK	326
GGGGSGGGGSQGANQK	327
GGGGSGGGGSQGANQ	328
GGGGSGGGGSQGANQQ	329
GGGDLQVDTQSQAAWAQLLQEHPNAQQYKGANKK	330
GGGGSGGGGSGGGGSGGGGSGGGGSGGGKGANKK	331
GGGGSGGGGSGGGGSGGGGSGGGGSGGGKGANQK	332
GGGGSGGGGSGGGGSGGGGSGGGGSGGGQGANQQ	333
QEHPNARKYKGANKK	334
QEHPNARKYKGANKK	335
PQEHPNARKYKGANKK	336
SGGGGSGGGGSGGGG	337
ASTSPANPQPAASSP	338
PSSAAPQPAPSTSA	339
SGGGGSGGGKGANKK	340
SGGGGSGGGQGANQQ	341
SGGGGSGGGKGANKQ	342
SGGGGSGGGKGANQK	343
SGGGGSGGGQGANKK	344
SGGGGSGGGKGANQQ	345
SGGGGSGGGQGANQK	346
SGGGGSGGGQGANQK	347
ASTSPANPQPAASSG	348

GSSAAPQPNAPSTSA	349
GSSAAPPNAPSTSAGLSKG	350
ASTSPANPRPAASSG	351
HGPQGQEHPNARKYKGANKK	352
HKLRGQEHPNARKYKGANKK	353
GHGPQGQEHPNARKYKGANKK	354
GHKLRGQEHPNARKYKGANKK	355
GGHGPQGQEHPNARKYKGANKK	356
GGHKLRGQEHPNARKYKGANKK	357
GGGHGPQGQEHPNARKYKGANKK	358
GGGHKLRGQEHPNARKYKGANKK	359
GGGGHGPQGQEHPNARKYKGANKK	360
GGGGHKLRGQEHPNARKYKGANKK	361
GGGGGHGPQGQEHPNARKYKGANKK	362
GGGGGHKLRGQEHPNARKYKGANKK	363
HGPQGSGGGSGGGKGANKK	364
HKLRGSGGGSGGGKGANKK	365
GGHGPQGSGGGSGGGKGANKK	366
GGGHKLRGSGGGSGGGKGANKK	367
SGGGGQEHPNARKYKGANKK	368
GGSGGGGQEHPNARKYKGANKK	369
SGGGGSGGGSGGGKGANKK	370
SGGGGSGGGSGGGSGGGKGANKK	371
GGGSGGGGSGGGSGGGKGANKK	372
GGGSGGGGSGGGSGGGSGGGKGANKK	373
HGPQG	374
HKLRG	375
GHGPQG	376
GGHGPQG	377
GGGHGPQG	378
GGGGHGPQG	379
GGGGGHGPQG	380
GHKLRG	381
GGHKLRG	382
GGGHKLRG	383
GGGGHKLRG	384
GGGGGHKLRG	385
GGQEIIIPNARKYKGANKK	386
GGGQEHPNARKYKGANKK	387
GGGGQEHPNARKYKGANKK	388
GGGGGQEHPNARKYKGANKK	389
LK	390
DI	391

In some embodiments, the linker may include or consist of a  $[(\text{Gly})_m(\text{Ser})]_n(\text{Gly})_p$  or  $(\text{Gly})_p[(\text{Ser})(\text{Gly})_m]_n$  linker as described above, followed by one of SEQ ID NOS: 314-321, e.g., one of SEQ ID NOS: 314, 315, or 321.

5 In other embodiments of polypeptides including an sALP, the linker may include or consist of all or a fragment of an sALP. For example, the 17-amino acid portion of human TNALP that is an N-terminal signal sequence, or homologs or variants or fragments thereof (e.g., residues 1-17 of SEQ ID NOS: 1202 or 1208), may be used as a linker. Homologs of this 17-amino acid region may be identified, e.g., by consulting a sequence alignment such as Fig. 8 (residues 1-17 of SEQ ID NO:

1216, where X can be any amino acid) or in Fig. 11 (residues 1-17 of SEQ ID NO: 1219, where X can be any amino acid but not a pathogenic mutation provided in Table 1). In another example, the C-terminal GPI anchor portion of human TNALP, or homologs or variants or fragments thereof (e.g., residues 503-524 of SEQ ID NO: 1208) may be used as a linker. Homologs of this C-terminal region 5 may be identified, e.g., by consulting a sequence alignment such as Fig. 8 (residues 503-524 of SEQ ID NO: 1216, where X can be any amino acid) or in Fig. 11 (residues 503-524 of SEQ ID NO: 1219, where X can be any amino acid but not a pathogenic mutation provided in Table 1).

In other embodiments of polypeptides including an NP, the linker may include or consist of all or a fragment of an NP. For example, the 31-amino acid portion of human CNP53 that is N-terminal to CNP22, or homologs or variants thereof (e.g., residues 4-34 of SEQ ID NO: 320), may be used as a linker. Homologs of this 31-amino acid region may be identified, e.g., by consulting a sequence alignment such as Figs. 15A-15G and identifying the regions corresponding to the N-terminal 31 amino acid residues of human CNP53. Other suitable linkers may also be identified, e.g., by choosing any portion of an NP, optionally excluding a ring domain, as shown in Figs. 15A-15G, or 10 in any other NP or region of an NP not shown in Figs. 15A-15G. For example, the C-terminal extension of DNP (SEQ ID NO: 117), or fragments or variants thereof, may be used as a linker. 15

A linker may optionally be glycosylated at any appropriate one or more amino acid residues.

In addition, a linker may have at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 20 99%, or more) sequence identity to any of the linkers described herein. In addition, a linker may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more additions, deletions, or substitutions relative to any of the linkers described herein.

A linker as described herein may include any other sequence or moiety, attached covalently or non-covalently.

25 In some embodiments, the linker is absent, meaning that the Fc fragment and the sALP are fused together directly or that the Fc fragment and the NP are fused together directly, with no intervening residues.

It should be noted that certain Fc-sALP or sALP-Fc fusion proteins may be viewed, according 30 to the present disclosure, either as 1) having no linker, or as 2) having a linker which corresponds to a portion of the sALP. For example, Fc fused directly to hsTNALP (1-502) may be viewed, e.g., either as having no linker, wherein the NP is hsTNALP (1-502), or as having a 17-amino acid linker, wherein the NP is hsTNALP (18-502).

Further, it should be noted that certain Fc-NP or NP-Fc fusion proteins may be viewed, according to the present disclosure, either as 1) having no linker, or as 2) having a linker which 35 corresponds to a portion of the NP. For example, Fc fused directly to CNP53 may be viewed, e.g., either as having no linker, wherein the NP is CNP53, or as having a 31-amino acid linker, wherein the NP is CNP22.

### **sALP Polypeptides**

Any of the sALPs and linkers described herein may be combined in an sALP polypeptide, e.g., an sALP polypeptide of A-sALP-B, wherein each of A and B is absent or is an amino acid sequence of at least one amino acid. When present, A and/or B can be any linker described herein (e.g., the amino acid sequence of any one of SEQ ID NOs: 301-391). In some embodiments, A is absent, B is absent, or A and B are both absent.

The sALP polypeptides of the invention can optionally include an Fc region to provide an sALP fusion polypeptide, as described herein.

The sALP polypeptide can optionally include a bone-targeting moiety (e.g., any described herein). In some embodiments, a linker, e.g., a flexible linker, may be included between the bone-targeting moiety and the sALP. For example, Fig. 5B provide polypeptides having both a bone-targeting moiety and a linker (shown in bold) between the Fc region and the bone-targeting moiety. In some embodiments, the linker is a dipeptide sequence (e.g., leucine-lysine or aspartic acid-isoleucine) or the amino acid sequence of any one of SEQ ID NOs: 301-391.

The sALP polypeptide can include any ALPs, mutations, N-terminal signal sequence, C-terminal GPI sequence, and/or linkers, or fragments thereof, described herein. For example, the italicized regions in Figs. 5B-5C and sequences provided in Figs. 6 or 8-11 may be used as for any of the sALPs disclosed herein (e.g., SEQ ID NOs: 1201, 1202, 1204-1216, 1218, and 1219).

### *sALP Fusion Polypeptides*

Any of the sALPs, linkers, and Fc regions described herein may be combined in a fusion polypeptide, e.g., a recombinant fusion polypeptide, that includes the structure C-sALP-D-Fc-E, C-Fc-D-sALP-E, C-sALP-D-Fc-G-I<sub>n</sub>-H, or C-Fc-D-sALP-G-I<sub>n</sub>-H, where each of C, D (the linker region), E, G (linker region), and H is absent or is an amino acid sequence of at least one amino acid; I represents an aspartic acid or a glutamic acid; and n=10 to 16. D and G can be absent or can optionally include any linker described herein (e.g., the amino acid sequence of any one of SEQ ID NOs: 301-391).

The polypeptides of the invention optionally include one or more additional amino acid residues 1) at the N-terminus of the polypeptide, 2) between the sALP and Fc regions of the polypeptide, and 3) at the C-terminus of the polypeptide. Thus, the invention includes, for example, polypeptides of the form C-sALP-D-Fc-E or the structure C-Fc-D-sALP-E, where C is one or more (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 17, 20, 25, 30, 35, 40, 45, 50, or more) additional amino acid residues at the N-terminus of the polypeptide, D is one or more (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 17, 20, 25, 30, 35, 40, 45, 50, or more) additional amino acid residues (i.e., a linker) between the sALP and Fc regions of the polypeptide, and E is one or more (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 17, 20, 25, 30, 35, 40, 45, 50, or more) additional amino acid residues at

the C-terminus of the polypeptide. In a particular example, D is the dipeptide leucine-lysine. Alternatively, any combination of C, D, and E may be present or absent. For example, in some embodiments, C and E are both absent, and D is absent or is an amino acid sequence of at least one amino acid. For example, the polypeptide may consist of the structure sALP-D-Fc or the structure Fc-D-sALP. In some embodiments of polypeptides consisting of the structure sALP-D-Fc or Fc-D-sALP, D may consist of two amino acid residues, e.g., leucine-lysine. For example, the polypeptide may consist of the structure sALP-D-Fc. Optionally, the amino acid sequence of sALP is the amino acid sequence of SEQ ID NO: 1205, the amino acid sequence of D is leucine-lysine, and/or the amino acid sequence of Fc is the amino acid sequence of SEQ ID NO: 401. In some embodiments, the 5 amino acid sequence of the polypeptide consists of the amino acid sequence of SEQ ID NO: 1204. In some embodiments, polypeptide includes an amino acid sequence of at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to the amino acid sequence of SEQ ID NOs: 1204 or 1221.

10 15 In some embodiments, the polypeptide includes a bone-targeting moiety, e.g., having a series of consecutive Asp or Glu residues, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>. The bone-targeting moiety, if present, may be positioned anywhere in the fusion polypeptide, e.g., at or near the N-terminal or C-terminal end, and/or in the linker region. For example, any one of C, D, and/or E may include a bone-targeting moiety. In some 20 25 embodiments, the bone-targeting moiety is at the C-terminal end. For example, the polypeptide may comprise or consist of the structure C-sALP-D-Fc-G-I<sub>n</sub>-H or the structure C-Fc-D-sALP-G-I<sub>n</sub>-H, where each of C, D (the linker region), G (linker region), and H is absent or is an amino acid sequence of at least one amino acid, I represents an aspartic acid or a glutamic acid, and n=10 to 16. In some embodiments, C and H are both absent, and D and G can be absent or can optionally include any linker described herein (e.g., the amino acid sequence of any one of SEQ ID NOs: 301-391). For example, the polypeptide may comprise or consist of the structure sALP-D-Fc-G-I<sub>n</sub> or the structure Fc-D-sALP-G-I<sub>n</sub>.

In some embodiments, polypeptide does not include a bone-targeting moiety.

30 A fusion polypeptide as described herein may have at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to any of the fusion polypeptides described herein, e.g., SEQ ID NOs: 1201, 1204, 1220, or 1221. In addition, a fusion polypeptide as described herein may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 50, or more additions, deletions, or substitutions relative to any of the fusion polypeptides described herein.

35 Furthermore, in some embodiments, a fusion polypeptide as described herein may be encoded by a nucleic acid molecule that hybridizes under high stringency conditions to at least a portion, e.g., to 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or even 100%,

of a nucleic acid molecule that encodes any of the polypeptides, e.g., fusion polypeptides, described herein.

In some embodiments, additional amino acid residues can be introduced into the polypeptide according to the cloning strategy used to produce the fusion polypeptides. In some embodiments, the 5 additional amino acid residues do not provide an additional GPI anchoring signal so as to maintain the polypeptide in a soluble form. Furthermore, in some embodiments, any such additional amino acid residues, when incorporated into the polypeptide of the invention, do not provide a cleavage site for endoproteases of the host cell. The likelihood that a designed sequence would be cleaved by the endoproteases of the host cell can be predicted as described, e.g., by Ikezawa (*Biol. Pharm. Bull.* 10 25:409-417, 2002).

In certain embodiments, the polypeptides of the invention are associated into dimers or tetramers. For example, two sALP-Fc monomers can covalently be linked through two disulfide bonds located in the hinge regions of the Fc fragments.

## 15 NP Polypeptides

Any of the NPs and linkers described herein may be combined in an NP polypeptide, e.g., an NP polypeptide of V-NP-W, wherein each of V and W is absent or is an amino acid sequence of at least one amino acid.

The NP polypeptides of the invention can optionally include an Fc region to provide an NP 20 fusion polypeptide, as described herein.

The NP polypeptide can optionally include a bone-targeting moiety. In some embodiments, a linker, e.g., a flexible linker, may be included between the bone-targeting moiety and the NP. For example, Fig. 27 provides polypeptides having a linker or both a bone-targeting moiety and a linker (SEQ ID NOS: 128-150); Figs. 31A-31B provide polypeptides having a linker or both a bone- 25 targeting moiety and a linker, where X in any one of SEQ ID NOS: 173-220 can be any amino acid, e.g., F, L, I, T, E, R, Y, C, P, or D; and Fig. 32 provides amino acid sequences for particular variants, where X in SEQ ID NOS: 186-198 is a leucine to provide the sequences in SEQ ID NOS: 221-233.

The NP polypeptide can include any NPs, N-terminal extensions, C-terminal extensions, and/or linkers described herein. For example, the italicized regions in Figs. 30, 31A-31B, and 32 may 30 be used as for any of the NPs disclosed herein (see, e.g., SEQ ID NOS: 511-516 and 553-558).

### *NP Fusion Polypeptides*

Any of the NPs, linkers, and Fc regions described herein may be combined in a fusion polypeptide, e.g., a recombinant fusion polypeptide, that includes the structure X-Fc-Y-NP-Z or the 35 structure X-NP-Y-Fc-Z, wherein each of X, Y (the linker region), and Z is absent or is an amino acid sequence of at least one amino acid.

5 Figs. 17A-17E depict several possible schematic structures of fusion polypeptides as described herein. Fc-NP or NP-Fc homodimers may be formed, e.g., due to disulfide bonds formed by Fc (Figs. 17A and 17B, respectively). Alternative, monomer-dimer hybrids are possible in which an NP-Fc or Fc-NP fusion polypeptide is joined to a free Fc domain (Figs. 17C and 17D, respectively). Furthermore, an NP-Fc monomer may be joined to an Fc-NP monomer, as shown in Fig. 17E. These configurations not intended to be exhaustive but are merely exemplary.

Exemplary fusion polypeptides having an N-terminal NP domain and a C-terminal Fc domain include those shown in Fig. 25A and include CNP-16AAlinker-Fc-His<sub>10</sub> (NC1) (SEQ ID NO: 521); CNP-6AAlinker-Fc-His<sub>10</sub> (NC3) (SEQ ID NO: 522); CNP-6AAlinker-Fc (SEQ ID NO: 523); CDNP-Fc (SEQ ID NO: 524), which has no linker between the CDNP and Fc moieties; CDNP-A17saa-Fc (SEQ ID NO: 525), which has a mutation to alanine at position 17 of the CNP22 region and mutations S3, A4, and A5 in the DNP tail region; and CDNP-A17sra-Fc (SEQ ID NO: 526), which has a mutation to alanine at position 17 of the CNP22 region and mutations S3 and A5 in the DNP tail region. Fig. 25B is a listing of the nucleic acid sequence (SEQ ID NO: 806) of NC1.

15 The NP domain in the fusion polypeptide can be any NP described herein. For example, any of the molecules shown in Figs. 30, 31A-31B, and 32, with or without the bone-targeting moiety, may be fused to an Fc domain and may optionally further include a linker region between the Fc and NP, as disclosed herein. For example, a CNP variant with M17X mutation can be fused to an Fc domain, e.g., as shown in Figs. 33A-33E, where X can be any amino acid, e.g., F, L, I, T, E, R, Y, C, P, D, G, 20 A, S, V, W, N, Q, H, or K, e.g., F, L, I, T, E, R, Y, C, P, or D, e.g., F or L. In some embodiments, the sequence is SEQ ID NO: 530, and X is any amino acid described herein, e.g., F, L, I, T, E, R, Y, C, P, D, G, A, S, V, W, N, Q, H, or K, e.g., F, L, I, T, E, R, Y, C, P, or D, e.g., F or L.

25 In the structure X-Fc-Y-NP-Z or the structure X-NP-Y-Fc-Z, X may include one or more (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 45, 50, or more) additional amino acid residues at the N-terminus of the polypeptide, and Z may independently include one or more (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 45, 50, or more) additional amino acid residues at the C-terminus of the polypeptide.

30 In some embodiments, the polypeptide includes a bone-targeting moiety, e.g., having a series of consecutive Asp or Glu residues, e.g., E<sub>6</sub>, E<sub>7</sub>, E<sub>8</sub>, E<sub>9</sub>, E<sub>10</sub>, E<sub>11</sub>, E<sub>12</sub>, E<sub>13</sub>, E<sub>14</sub>, E<sub>15</sub>, E<sub>16</sub>, D<sub>6</sub>, D<sub>7</sub>, D<sub>8</sub>, D<sub>9</sub>, D<sub>10</sub>, D<sub>11</sub>, D<sub>12</sub>, D<sub>13</sub>, D<sub>14</sub>, D<sub>15</sub>, or D<sub>16</sub>. The bone-targeting moiety, if present, may be positioned anywhere in the fusion polypeptide, e.g., at or near the N-terminal or C-terminal end, and/or in the linker region. For example, any one of X, Y, and/or Z may include a bone-targeting moiety.

35 In some instances, one or more amino acid residues are introduced into the fusion polypeptide, e.g., within X, Y, or Z, as a result of the cloning strategy used. In some embodiments, any such additional amino acid residues, when incorporated into the polypeptide of the invention, do not provide a cleavage site for endoproteases of the host cell. The likelihood that a designed sequence

would be cleaved by the endoproteases of the host cell can be predicted as described, e.g., by Ikezawa (*Biol. Pharm. Bull.* 25:409-417, 2002), hereby incorporated by reference.

In certain embodiments, the fusion polypeptides of the invention are associated into dimers, e.g., through two disulfide bonds located in the hinge regions of the Fc fragments.

5 In some embodiments, the fusion polypeptides of the invention have at least, e.g., 1, 1.5, 2, 2.5, 3, 3.5, 4, 5, 6, 7, 8, 9, 10, 12.5, 15, 20, 25, 30, 35, 40, 45, 50, 75, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 600, 700, 800, 900, 1,000, 1,500, 2,000, 3,000, 4,000, 5,000, 7,500, 10,000, 15,000, 20,000, 25,000, 30,000, 40,000, or 50,000 times the half-life of CNP22 *in vivo*.

10 Any NP fusion protein may be expressed with an N-terminal signal sequence to facilitate secretion, e.g., amino acid residues 1-25 of SEQ ID NO: 501, or any other signal sequence known in the art. Such sequences are generally cleaved co-translationally, resulting in secretion of the mature version of the protein.

15 A fusion polypeptide as described herein may have at least 50% (e.g., 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more) sequence identity to any of the fusion polypeptides described herein, e.g., SEQ ID NOs: 501-608. In addition, a fusion polypeptide as described herein may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 25, 30, 35, 40, 50, or more additions, deletions, or substitutions relative to any of the fusion polypeptides described herein. Furthermore, in some embodiments, a fusion polypeptide as described herein may be encoded by a nucleic acid molecule 20 that hybridizes under high stringency conditions to at least a portion, e.g., to 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or even 100%, of a nucleic acid molecule that encodes any of the polypeptides, e.g., fusion polypeptides, described herein.

25 Fusion proteins include those having one or more modifications that are cleaved during expression. For example, an Fc-CNP fusion protein was designed as shown in Figs. 18A and 18B. This protein, termed “NC2 Streptag” or “NC2st,” has a 25-amino acid N-terminal signal sequence that is cleaved during expression. The mature protein has the following domain structure, from N terminus to C terminus: Strep-tag II sequence (to facilitate purification) flanked by short linker sequences; TEV protease cleavage sequence, followed by a short linker; Fc domain of human IgG-1; 16-amino acid glycine-rich linker; and CNP22. This protein can be produced by chemically 30 synthesizing the coding sequence (Fig. 18C, SEQ ID NO: 801) and inserting the coding sequence into a small cloning plasmid using standard techniques known in the art.

35 Fusion proteins may be varied in several respects. For example, NC2st can be varied by eliminating the sequence that is N-terminal to the Fc domain (resulting, e.g., in NC2B, as shown in Figs. 19A-19B), adding a bone-targeting moiety (resulting, e.g., in D10-NC2, as shown in Fig. 19A), and/or altering of the length of the linker between Fc and CNP22 (e.g., NC2B-22 (also referred to as NC2-22), NC2B-28 (also referred to as NC2-28), and NC2B-34 (also referred to as NC2-34), as

shown in Figs. 20A-20D). Other exemplary NC2st variants are shown in Fig. 21 (NC2-KGANKK and NC2-KGANQK) and Fig. 22 (NC2-CNP53mut2).

NC2B may be varied in several respects, including having a point mutation, e.g., at position 17 relative to CNP22, having a bone-targeting moiety, and/or having modified or altered linker regions. Exemplary fusion protein include any sequences having a modified or altered linker region (e.g., SEQ ID NOS: 511-516, as shown in Fig. 34A), as compared to NC2B (as shown in Figs. 16A-16B); any sequences having a bone-targeting moiety, e.g., a D<sub>10</sub> moiety (e.g., SEQ ID NOS: 553-558, as shown in Fig. 34B); any N-terminal extensions, C-terminal extensions, and/or linkers for any of the NPs disclosed herein (e.g., the italicized regions in SEQ ID NOS: 511-516 and 553-558, as shown in Figs. 30, 31A-31B, and 32); any sequences having a phenylalanine (Phe, F) substitution at position 17 relative to CNP22 (e.g., SEQ ID NOS: 559-564, as shown in Fig. 34C), including those sequences having a further modification of a D<sub>10</sub> bone-targeting moiety at the N-terminus (e.g., SEQ ID NOS: 565-570, as shown in Fig. 34D); any sequences having a leucine (Leu, L) substitution at position 17 relative to CNP22 (e.g., SEQ ID NOS: 571-576, as shown in Fig. 34E), including those sequences having a further modification of a D<sub>10</sub> bone-targeting moiety at the N-terminus (e.g., SEQ ID NOS: 577-582, as shown in Fig. 34F); any sequences having an arginine (Arg, R) substitution at position 17 relative to CNP22 (e.g., SEQ ID NOS: 583-588, as shown in Fig. 34G), including those sequences having a further modification of a D<sub>10</sub> bone-targeting moiety at the N-terminus (e.g., SEQ ID NOS: 589-594, as shown in Fig. 34H); and any sequences having a tyrosine (Tyr, Y) substitution at position 17 relative to CNP22 (e.g., SEQ ID NOS: 595-600, as shown in Fig. 34I), including those sequences having a further modification of a D<sub>10</sub> bone-targeting moiety at the N-terminus (e.g., SEQ ID NOS: 601-606, as shown in Fig. 34J).

Additional constructs include fusion proteins in which an N-terminal Fc domain is fused to a variant of CNP53 with a short Gly<sub>3</sub> linker region. An alternative way to analyze these fusion polypeptides is that the linker region is Gly<sub>3</sub> followed by amino acid residues 1-31 of CNP53 (or variants thereof); viewed in this way, the linker region connects the Fc domain to CNP22 and is 34 amino acid residues in length. For Fc-CNP53-A (also referred to as “Fc-CNP53wt”) (SEQ ID NOS: 517 (with signal sequence) and 518 (without signal sequence); Fig. 23), position 48 of CNP53, corresponding to position 17 of CNP22, was mutated to alanine. Fc-CNP53-AAA (also referred to as “Fc-CNP53mut”) (SEQ ID NOS: 519 (with signal sequence) and 520 (without signal sequence); Fig. 23) has the same sequence as Fc-CNP53-A with the exception that residues 30 and 31 of CNP53, the two residues immediately before CNP22, are mutated to alanine in order to reduce the likelihood of proteolytic cleavage. In some cases, modifying the linker region of an Fc-CNP22 fusion to include the first 31 amino acid residues of CNP53 could result in constructs having even greater potency and efficacy than NC2st in *in vitro* membrane and whole cell assays.

### **Additional Polypeptide Features**

The polypeptides of the invention also include any polypeptide having one or more post-translational modifications such as glycosylation (e.g., mannosylation and other forms of glycosylation discussed herein), acetylation, amidation, blockage, formylation, gamma-5 carboxyglutamic acid hydroxylation, methylation, ubiquitination, phosphorylation, pyrrolidone carboxylic acid modification, and sulfation. Artificial modifications, e.g., pegylation, may also be made.

### **Production of Nucleic Acids and Polypeptides**

10 The nucleic acids and polypeptides of the invention can be produced by any method known in the art. Typically, a nucleic acid encoding the desired fusion protein is generated using molecular cloning methods, and is generally placed within a vector, such as a plasmid or virus. The vector is used to transform the nucleic acid into a host cell appropriate for the expression of the fusion protein. Representative methods are disclosed, for example, in Maniatis et al. (Cold Springs Harbor 15 Laboratory, 1989). Many cell types can be used as appropriate host cells, although mammalian cells are preferable because they are able to confer appropriate post-translational modifications. For example, Human Embryonic Kidney 293 (HEK293) cells have been used as a host for expressing the fusion proteins of the present invention, as described in more detail in the Examples below.

20 The polypeptides of the invention can be produced under any conditions suitable to effect expression of the polypeptide in the host cell. Such conditions include appropriate selection of a media prepared with components such as a buffer, bicarbonate and/or HEPES, ions like chloride, phosphate, calcium, sodium, potassium, magnesium, iron, carbon sources like simple sugars, amino acids, potentially lipids, nucleotides, vitamins and growth factors like insulin; regular commercially available media like alpha-MEM, DMEM, Ham's-F12, and IMDM supplemented with 2-4 mM L-25 glutamine and 5% Fetal bovine serum; regular commercially available animal protein free media like Hyclone™ SFM4CHO, Sigma CHO DHFR, Cambrex POWER™ CHO CD supplemented with 2-4 mM L-glutamine. These media are desirably prepared without thymidine, hypoxanthine and L-glycine to maintain selective pressure, allowing stable protein-product expression.

30 Additional details of the production of the polypeptides and nucleic acids of the invention are given in the Examples.

### **Therapeutic Applications**

The polypeptides and nucleic acid molecules described herein can have a wide variety of therapeutic applications, e.g., in the fields of neurocutaneous syndromes (e.g., neurofibromatosis) or 35 disorders associated with overactivation of FGFR3 (e.g., bone and cartilage disorders, e.g., achondroplasia, or cancers, e.g., multiple myeloma) or bone or cartilage disorders (e.g., that are not associated with overactivation of FGFR3) or vascular smooth muscle disorders. In addition, the

polypeptides and nucleic acid molecules described herein can be used for any condition or disorder that would benefit from elongation of bone.

#### *Neurocutaneous Syndromes*

5 The polypeptides and nucleic acid molecules described herein can be used to treat any disorder, disease, or other abnormality associated with elevated blood and/or urine levels of inorganic pyrophosphate (PP<sub>i</sub>) and/or overactivation of MAP kinase, such as neurocutaneous syndromes. In particular embodiments, the polypeptide and nucleic acid molecules are used to treat a neurocutaneous syndrome either with or without a bone manifestation. Exemplary neurocutaneous 10 syndromes include neurofibromatosis (e.g., any type described herein, such as type I or type II); Noonan syndrome-like disorder with loose anagen hair; Noonan syndrome-like disorder with juvenile myelomonocytic leukemia (JMML); tuberous sclerosis; Sturge-Weber disease; ataxia telangiectasia; von Hippel-Lindau disease; incontinentia pigmenti; epidermal nevus syndromes, such as linear sebaceous nevus of Jadassohn; nevoid basal cell carcinoma syndrome; hypomelanosis of Ito; 15 neurocutaneous melanosis; Klippel-Trenaunay syndrome; and Waardenburg syndrome, including types I, II, III, and IV.

#### *Neurofibromatosis*

20 Neurofibromatosis is an autosomal dominant neurocutaneous syndrome characterized by abnormal growth or proliferation of nerve tissue, such as to produce tumors (or neurofibromas) or 25 abnormal pigmentation (e.g., café au lait spots). In particular, neurofibromatosis can be accompanied by bone manifestations (e.g., manifestations arising from hypophosphatasia), such as short stature, scoliosis, osteomalacia, osseous fibrous dysplasia (e.g., one or more lesions at the ends of or within one or more bones, such as the femur, tibia, or fibula), pseudarthrosis (e.g., tibial pseudarthrosis), and/or skeletal dysplasia (e.g., tibial dysplasia, orbital dysplasia, and/or sphenoid wing dysplasia).

Neurofibromatosis, or any of its manifestations or phenotypes, can be treated using the 30 compositions and methods described herein. Examples of such disorders, manifestations, and phenotypes include the classic von Recklinghausen type (type I), either with gastrointestinal stromal tumors (i.e., as in intestinal neurofibromatosis (type 3B)) or without such tumors; an acoustic neuroma type (type II); a mixed type that combines the features of types I and II with predominant features, such as bilateral acoustic neuromas, posterior fossa and upper cervical meningiomas, and spinal/paraspinal neurofibromas (type III, Riccardi type or type 3A); an atypical type that is distinguished from by the lack of iris Lisch nodules that are characteristic of type I (type VI); segmental neurofibromatosis, which is a variant of type I having lesions affecting a specific area of 35 the body, such as a single segment of the body or an area that crosses the midline (type V); a type having only the symptoms of café au lait spots without other manifestations of neurofibromatosis (type VI); familial spinal neurofibromatosis, which is caused by mutation in the neurofibromin gene

NF1 and considered a distinguishable variant of type I; other variants of type I, such as neurofibromatosis-pheochromocytoma-duodenal carcinoid syndrome; neurofibromatosis with manifestations of Noonan syndrome, such as short stature, ptosis, midface hypoplasia, webbed neck, learning disabilities, and muscle weakness; and schwannomatosis, where any of these disorders can 5 include or exclude one or more bone manifestations.

***Disorders Associated with Overactivation of RAS and/or ERK***

Any disorder, disease, or other abnormality that is caused by, or is associated with, overactivation of RAS and/or ERK may be treated using the compositions and methods described 10 herein. These disorders, diseases, and other abnormalities include, without limitation, Noonan syndrome, Costello syndrome, Noonan syndrome with multiple lentigines/LEOPARD syndrome, neurofibromatosis type 1, NF1-Noonan syndrome, hereditary gingival fibromatosis type 1, capillary malformation-AV malformation syndrome, Legius syndrome, Noonan syndrome-like disorder with loose anagen hair, Noonan syndrome-like disorder with juvenile myelomonocytic leukemia (JMML), 15 cardio-facio-cutaneous syndrome, or autoimmune lymphoproliferative syndrome, where any of these disorders can include or exclude one or more bone manifestations.

***Disorders Associated with Overactivation of FGFR3***

Any disorder, disease, or other abnormality that is caused by, or is associated with, 20 overactivation of FGFR3, e.g., stemming from a gain-of-function FGFR3 mutation, may be treated using the compositions and methods described herein. These disorders, diseases, and other abnormalities include, without limitation, bone or cartilage disorders and cancers, each of which is described in more detail below.

**25 *Bone or Cartilage Disorders Associated with Overactivation of FGFR3***

Any disorder, disease, or other abnormality, e.g., skeletal dysplasia, that affects the function, structure, or growth of bone or cartilage, may be treated using the compositions and methods described herein. In particular, the disorder may be a skeletal dysplasia that is associated with overactivation of FGFR3, such as achondroplasia, including severe achondroplasia with 30 developmental delay and acanthosis; Muenke syndrome (Muenke coronal craniosynostosis); Crouzonodermoskeletal syndrome; hypochondroplasia; thanatophoric dysplasia type I; and thanatophoric dysplasia type II. The compositions and methods of the invention can also be used to treat bone or cartilage disorders not associated with overactivation of FGFR3, and these disorders are described in more detail below.

*Cancers*

Any cancer that is caused by, or is associated with, overactivation of FGFR3, may be treated using the compositions and methods described herein. These cancers include, e.g., multiple myeloma, 5 myeloproliferative syndromes, leukemia (e.g., plasma cell leukemia), lymphomas, glioblastoma, prostate cancer, bladder cancer, and mammary cancer.

*Bone or Cartilage Disorders*

The polypeptides and nucleic acid molecules described herein can be used to treat any 10 disorder, disease, phenotype, or other abnormality that affects the function, structure, or growth of bone or cartilage. These bone or cartilage disorders may be, but do not necessarily have to be, associated with overactivation of FGFR3.

Exemplary bone or cartilage disorders include skeletal dysplasia and any other disorders, 15 diseases, phenotypes, or other abnormalities related to the bone or cartilage, including achondroplasia (e.g., homozygous or heterozygous achondroplasia), achondrogenesis, acrodysostosis, acromesomelic dysplasia, atelosteogenesis, bone pain, calcium pyrophosphate dihydrate (CPPD) crystal deposition, camptomelic dysplasia, chondrodysplasia punctata (e.g., rhizomelic type of chondrodysplasia punctata), cleidocranial dysostosis, congenital short femur, craniosynostosis (e.g., Muenke syndrome, Crouzon syndrome, Apert syndrome, Jackson-Weiss syndrome, Pfeiffer syndrome, or 20 Crouzonodermoskeletal syndrome), dactyly (e.g., brachydactyly, camptodactyly, polydactyly, or syndactyly), diastrophic dysplasia, dental disorders (e.g., decrease in teeth mineralization and premature loss of deciduous teeth, such as through aplasia, hypoplasia or dysplasia of the dental cementum), dwarfism, dyssegmental dysplasia, enchondromatosis, fibrochondrogenesis, fibrous dysplasia, hereditary multiple exostoses, hypochondroplasia, hypophosphatasia (HPP) (e.g., infantile 25 HPP, childhood HPP, perinatal HPP, adult HPP, or odontohypophosphatasia), HPP-related seizure, hypophosphatemic rickets, incomplete bone mineralization, Jaffe-Lichtenstein syndrome, Kniest dysplasia, Kniest syndrome, Langer-type mesomelic dysplasia, Marfan syndrome, McCune-Albright syndrome, micromelia, metaphyseal dysplasia (e.g., Jansen-type metaphyseal dysplasia), metatrophic dysplasia, Morquio syndrome, Nievergelt-type mesomelic dysplasia, neurofibromatosis (e.g., type 1, 30 e.g., with bone manifestations or without bone manifestations; type 2; or schwannomatosis), osteoarthritis, osteochondrodysplasia, osteogenesis imperfecta (e.g., perinatal lethal type of osteogenesis imperfecta), osteomalacia, osteopetrosis, osteopoikilosis, osteoporosis, peripheral dysostosis, Reinhardt syndrome, Roberts syndrome, Robinow syndrome, short-rib polydactyly syndromes, short stature, spondyloepiphyseal dysplasia congenita, spondyloepimetaphyseal dysplasia, 35 or thanatophoric dysplasia. Bone or cartilage disorders also include those that can be diagnosed, for example, by elevated blood and/or urine levels of one or more clinical markers related to hypophosphatasia (e.g., elevated levels of inorganic pyrophosphate (PP<sub>i</sub>), phosphoethanolamine

(PEA), and/or pyridoxal 5'-phosphate (PLP)), growth retardation with a decrease of long bone length (such as femur, tibia, humerus, radius, ulna), a decrease of the mean density of total bone, or a decrease of bone mineralization in bones such as femur, tibia, ribs and metatarsi, and phalange.

Without being so limited, treatment of bone or cartilage disorders may be observed by one or more of 5 the following: an increase of long bone length, an increase of mineralization in bone and/or teeth, a correction of bowing of the legs, a reduction of bone pain, and a reduction of CPPD crystal deposition in joints.

#### *Skeletal Dysplasia*

10 Skeletal dysplasias are bone or cartilage disorders characterized by short stature or dwarfism. Skeletal dysplasias are typically congenital and may include numerous abnormalities in addition to short stature, e.g., short limbs and trunk; bowlegs; a waddling gait; skull malformations, e.g., a large head, cloverleaf skull, craniosynostosis (premature fusion of the bones in the skull), or wormian bones (abnormal thread-like connections between the bones in the skull); anomalies of the hands and feet, 15 e.g., polydactyly (extra fingers), "hitchhiker" thumbs, and abnormal fingernails and toenails; or chest anomalies, e.g., pear-shaped chest or narrow thorax. Non-skeletal abnormalities may also be present in individuals having skeletal dysplasia, e.g., anomalies of the eyes, mouth, and ears, such as congenital cataracts, myopia, cleft palate, or deafness; brain malformations, such as hydrocephaly, 20 porencephaly, hydranencephaly, or agenesis of the corpus callosum; heart defects, such as atrial septal defect, patent ductus arteriosus, or transposition of the great vessels; developmental delays; or mental retardation. Skeletal dysplasias associated with overactivation of FGFR3 include achondroplasia.

Skeletal dysplasias include achondroplasia (e.g., homozygous or heterozygous 25 achondroplasia), achondrogenesis, acrodysostosis, acromesomelic dysplasia, atelosteogenesis, camptomelic dysplasia, chondrodysplasia punctata (e.g., rhizomelic type of chondrodysplasia punctata), cleidocranial dysostosis, congenital short femur, craniosynostosis (e.g., Muenke syndrome, Crouzon syndrome, Apert syndrome, Jackson-Weiss syndrome, Pfeiffer syndrome, or 30 Crouzonodermoskeletal syndrome), dactyly (e.g., brachydactyly, camptodactyly, polydactyly, or syndactyly), diastrophic dysplasia, dwarfism, dyssegmental dysplasia, enchondromatosis, fibrochondrogenesis, fibrous dysplasia, hereditary multiple exostoses, hypochondroplasia, 35 hypophosphatasia (HPP) (e.g., infantile HPP, childhood HPP, perinatal HPP, adult HPP, or odontohypophosphatasia), hypophosphatemic rickets, Jaffe-Lichtenstein syndrome, Kniest dysplasia, Kniest syndrome, Langer-type mesomelic dysplasia, Marfan syndrome, McCune-Albright syndrome, micromelia, metaphyseal dysplasia (e.g., Jansen-type metaphyseal dysplasia), metatrophic dysplasia, Morquio syndrome, Nievergelt-type mesomelic dysplasia, neurofibromatosis (e.g., type 1, e.g., with 40 bone manifestations or without bone manifestations; type 2; or schwannomatosis), osteoarthritis, osteochondrodysplasia, osteogenesis imperfecta (e.g., perinatal lethal type of osteogenesis imperfecta), osteopetrosis, osteopoikilosis, peripheral dysostosis, Reinhardt syndrome, Roberts

syndrome, Robinow syndrome, short-rib polydactyly syndromes, short stature, spondyloepiphyseal dysplasia congenita, spondyloepimetaphyseal dysplasia, or thanatophoric dysplasia.

In particular, some forms of craniosynostosis are the result of mutations in one of the fibroblast growth factor receptors (e.g., one or more of FGFR1, FGFR2, or FGFR3) that cause the activation of the MAPK pathway. This is the case for Muenke (Muenke coronal craniosynostosis), Crouzon, Apert, Jackson-Weiss, Pfeiffer, and Crouzonodermoskeletal syndromes, for example. There is genetic and biochemical evidence in the scientific literature that agents that can prevent activation of the MAP-kinase (ERK 1/2) can prevent craniosynostosis in animal models. In particular, use of a MEK1/2 inhibitor (e.g., U0126), which prevents activation of ERK1/2 can prevent craniosynostosis in an animal model of Apert syndrome (Shukla et al., *Nat. Genet.* 39:1145, 2007). Accordingly, the compounds of the present invention, which can prevent activation of the MAP-kinase pathway, could be used to treat these forms of craniosynostosis.

#### *Achondroplasia*

Achondroplasia is an autosomal dominant skeletal dysplasia that is the most common cause of dwarfism in humans. Its incidence is approximately 1 in 20,000 live births. Skeletal manifestations include growth retardation (with an average adult height of 123-131 cm (4 feet ½ in. – 4 feet 3½ in.)), skull deformities, and orthodontic defects. Extraskeletal manifestations include cervical cord compression (with risk of death, e.g., from central apnea or seizures); spinal stenosis (e.g., leg and lower back pain); hydrocephalus (e.g., requiring cerebral shunt surgery); hearing loss due to chronic otitis; cardiovascular disease; neurological disease; higher frequency of accidents; and obesity.

Babies are often diagnosed at birth. While the homozygous form is usually lethal, individuals diagnosed with the heterozygous form have a life expectancy, on average, of 15 years less than the normal population.

Heterozygous or homozygous achondroplasia, or any of its manifestations or phenotypes, can be treated using the compositions and methods described herein. Treatment of either form may be started as early as possible in the patient's life, e.g., shortly after birth, or even *in utero*; this is particularly important for treatment of the homozygous form, which is typically much more severe and is often lethal if untreated.

*Hypophosphatasia (HPP)*

HPP is a matrix mineralization disorder that is historically classified according to age at diagnosis and includes (in order from most severe to least severe) perinatal, infantile, childhood, adult, and odontohypophosphatasia forms. The most severe form, perinatal (lethal) HPP, is manifest as an almost complete absence of bone mineralization *in utero* and can cause stillbirth. Some neonates with perinatal HPP may survive for several days, but suffer increased respiratory compromise due to hypoplastic and rachitic disease of the chest. In infantile HPP, diagnosed before 6

months-of-age, postnatal development seems normal until the onset of poor feeding, inadequate weight gain, and appearance of rickets. Infantile HPP has characteristic radiological features showing impaired skeletal mineralization, sometimes with progressive skeletal demineralization leading to rib fractures and chest deformity. Childhood HPP has highly variable clinical expression. One symptom 5 of childhood HPP is the premature loss of deciduous teeth resulting from aplasia, hypoplasia or dysplasia of dental cementum that connects the tooth root with the periodontal ligament. Another symptom of childhood HPP is rickets, which causes short stature and skeletal deformities such as bowed legs and enlarged wrists, knees and ankles as a result of flared metaphysis. Adult HPP usually presents during middle age, but is frequently preceded by a history of rickets and/or early loss of teeth 10 followed by good health during adolescence and young adult life. In adult HPP, recurrent metatarsal stress fractures are common, and calcium pyrophosphate dihydrate deposition can cause attacks of arthritis and pyrophosphate arthropathy. Finally, odontohypophosphatasia is diagnosed when the only clinical abnormality is dental disease, and radiological studies and even bone biopsies reveal no signs of rickets or osteomalacia.

15 The more severe clinical forms of HPP are usually inherited as autosomal recessive traits, with parents of such patients showing subnormal levels of serum AP activity. For the milder forms of HPP, i.e., adult HPP and odontohypophosphatasia, an autosomal dominant pattern of inheritance has also been documented.

20 In the healthy skeleton, TNALP is an ectoenzyme present on the surface of the plasma membrane of osteoblasts and chondrocytes, and on the membranes of their shed matrix vesicles (MVs), where the enzyme is particularly enriched. Deposition of hydroxyapatite during bone mineralization normally initiates within the lumen of these MVs. Electron microscopy has shown that TNALP-deficient MVs from severely affected HPP patients and *Akp2<sup>-/-</sup>* mice (a TNALP null mouse model, see below) contain hydroxyapatite crystals, but that extravesicular crystal propagation appears 25 retarded. This defect is attributed to the extracellular accumulation of PP<sub>i</sub>, a potent inhibitor of calcification, due to a deficiency of TNALP activity.

30 At physiological concentrations (0.01-0.1 mM), PP<sub>i</sub> has the ability to stimulate mineralization. This has been demonstrated in organ-cultured chick femurs and in isolated rat MVs. However, at concentrations above 1 mM, PP<sub>i</sub> inhibits calcium phosphate mineral formation by coating 35 hydroxyapatite crystals, thus preventing mineral crystal growth and proliferative self-nucleation. Thus, PP<sub>i</sub> has a dual physiological role: it functions as a promoter of mineralization at low concentrations but as an inhibitor of mineralization at higher concentrations. TNALP has been shown to hydrolyze the mineralization inhibitor PP<sub>i</sub> to facilitate mineral precipitation and growth. Recent studies using the *Akp2<sup>-/-</sup>* mice have indicated that the primary role of TNALP *in vivo* is to restrict the size of the extracellular PP<sub>i</sub> pool to allow proper skeletal mineralization.

The severity of hypophosphatasia depends on the nature of the TNALP mutation. Missense mutations at a variety of positions in TNALP, including the enzyme's active site vicinity, homodimer

interface, crown domain, amino-terminal arm, and calcium-binding site, have all been found to affect its catalytic activity. In addition, missense, nonsense, frame-shift, and splice site mutations have also been shown to lead to aberrant mutant proteins or intracellular trafficking defects that lead to subnormal activity on the cell surface. The multitude of mutations that cause HPP, and the fact that 5 compound heterozygosity is a common occurrence in HPP, explain the variable expressivity and incomplete penetrance often observed in this disease.

Progress on the human form of HPP has benefited greatly from the existence of the TNALP null mice (*Akp2*<sup>-/-</sup>), an animal model of HPP. *Akp2*<sup>-/-</sup> mice phenocopy infantile HPP remarkably well: they are born with a normally mineralized skeleton but develop radiographically apparent rickets at 10 about 6 days of age, and die between days 12-16 suffering severe skeletal hypomineralization and episodes of apnea and epileptic seizures attributable to disturbances in PLP (vitamin B<sub>6</sub>) metabolism.

Both PP<sub>i</sub> and PLP are confirmed natural substrates of TNALP, and some TNALP active site mutations have been shown to have different effects on the ability of the enzyme to metabolize PP<sub>i</sub> and PLP. Abnormalities in PLP metabolism explain the epileptic seizures observed in *Akp2*<sup>-/-</sup> mice, 15 while abnormalities in PP<sub>i</sub> metabolism explain the skeletal phenotype in this mouse model of HPP.

In any of the methods of the invention, the pharmaceutical compositions described herein are optionally administered in an amount that is therapeutically effective to treat a HPP phenotype selected from the group consisting of HPP-related seizure, premature loss of deciduous teeth, incomplete bone mineralization, elevated blood and/or urine levels of PP<sub>i</sub>, elevated blood and/or urine 20 levels of PEA, elevated blood and/or urine levels of PLP, inadequate weight gain, rickets, bone pain, calcium pyrophosphate dihydrate crystal deposition, aplasia, hypoplasia, and dysplasia of the dental cementum. In some embodiments, the incomplete bone mineralization is incomplete femoral bone mineralization, incomplete tibial bone mineralization, incomplete metatarsal bone mineralization, or incomplete rib bone mineralization.

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### ***Vascular Smooth Muscle Disorders***

The polypeptides and nucleic acid molecules described herein can be used to treat any disorder, disease, or other abnormality that affects the function, structure, or growth of vascular smooth muscle. For example, natriuretic peptides modulate salt and water homeostasis in the body 30 and in this way act as regulators of blood pressure. The peptides belonging to this family have varying amino acid sequences and are secreted through different mechanisms by various tissues in the body. For example, ANP is released by muscle cells in the upper chambers (atria) of the heart (atrial myocytes) and acts as a vasodilator; BNP is secreted by the lower chambers (ventricles) of the heart in response to cardiac stress; and CNP exerts natriuretic and natriuretic effect and regulates vessel tone, 35 inhibits migration and proliferation of vascular smooth muscle cell. Accordingly, the polypeptides and compositions of the invention can be used to treat vascular smooth muscle disorders. Exemplary vascular smooth muscle disorders are hypertension, restenosis, arteriosclerosis, acute decompensated

heart failure, congestive heart failure, cardiac edema, nephredema, hepatic edema, acute renal insufficiency, and chronic renal insufficiency.

***Conditions for Elongation of Bone***

5 Any condition, disorder, disease, or other abnormality that would benefit from elongation of bone may be treated using the compositions and methods described herein. These conditions, disorders, diseases, and other abnormalities include, without limitation, insufficient or impaired bone growth arising from fractures, renal failure or insufficiency, poor diet, vitamin deficiency, or hormone deficiency. Healthy subjects, e.g., those without any conditions, disorders, diseases, or other  
10 abnormalities related to bone or cartilage, may also be treated using the compositions and methods described herein, e.g., for cosmetic purposes.

Skeletal dysplasias are also associated with shortened segments of long bones. Exemplary skeletal dysplasias include those associated with rhizomelia (or shortening in a proximal segment of a limb, e.g., in the humerus or femur), such as achondroplasia, atelosteogenesis, congenital short femur,  
15 diastrophic dysplasia, hypochondroplasia, Jansen type of metaphyseal dysplasia, rhizomelic type of chondrodysplasia punctata, spondyloepiphyseal dysplasia congenita, and thanatophoric dysplasia; mesomelia (or shortening in a middle segment of a limb, e.g., in the radius, ulna, tibia, or fibula), such as Langer and Nievergelt types of mesomelic dysplasias, Robinow syndrome, and Reinhart  
syndrome; acromelia (or shortening in a distal segment of a limb, e.g., in the metacarpals or  
20 phalanges), such as acrodysostosis and peripheral dysostosis; acromesomelia (or shortening in the middle and distal segments of limbs, e.g., in the forearms and hands), such as acromesomelic dysplasia; micromelia (or shortening in the entire limb), such as achondrogenesis, fibrochondrogenesis, dyssegmental dysplasia, Kniest dysplasia, and Roberts syndrome; or short-trunk,  
25 such as Dyggve-Melchior-Clausen disease, Kniest syndrome, metatrophic dysplasia, Morquio syndrome, spondyloepimetaphyseal dysplasia, and spondyloepiphyseal dysplasia congenita.

***Combination Therapy***

The combinations of the polypeptides of the invention (e.g., a combination of an sALP polypeptide and an NP polypeptide, such as a combination of an sALP fusion protein and an NP fusion protein) are useful for the treatment of any disease or condition described herein. Therapy may be performed alone or in conjunction with another therapy (e.g., surgery, radiation therapy, immunotherapy, or gene therapy). Additionally, a person having a greater risk of developing a disease described herein (e.g., one who is genetically predisposed or one who previously had a neurocutaneous syndrome) may receive prophylactic treatment to inhibit or delay disease formation.  
30 The duration of the combination therapy depends on the type of disease or disorder being treated, the age and condition of the patient, the stage and type of the patient's disease, and how the patient  
35

responds to the treatment. Therapy may be given in on-and-off cycles that include rest periods so that the patient's body has a chance to recover from any as yet unforeseen side-effects.

The administration of a combination of the polypeptides of the present invention (e.g., a combination of an sALP polypeptide and an NP polypeptide) allows for the administration of lower doses of each polypeptide, providing similar efficacy and lower toxicity compared to administration of either polypeptide alone. Alternatively, such combinations result in improved efficacy in treating a disease described herein (e.g., a neurocutaneous syndrome, such as neurofibromatosis) with similar or reduced adverse events over the single agent alone, at moderate or high doses.

## 10 *Formulation*

Formulation will depend on the route of administration, as well as on other therapeutic goals. The polypeptides and nucleic acid molecules described herein can be administered by any route known in the art, e.g., subcutaneous (e.g., by subcutaneous injection), intravenously, orally, nasally, intramuscularly, sublingually, intrathecally, or intradermally. By way of example, pharmaceutical compositions of the invention can be in the form of a liquid, solution, suspension, pill, capsule, tablet, gelcap, powder, gel, ointment, cream, nebulae, mist, atomized vapor, aerosol, or phytosome.

In some embodiments of the invention, the compositions of the invention can be administered subcutaneously. Subcutaneous administration is advantageous because it is relatively non-invasive and offers desirable pharmacokinetic profiles. Suitable volumes are known to those skilled in the art, and are typically 5 mL or smaller (e.g., 4 mL, 3.5 mL, 3 mL, 2.7 mL, 2.5 mL, 2.3 mL, 2.2 mL, 2.1 mL, 2.0 mL, 1.9 mL, 1.8 mL, 1.7 mL, 1.5 mL, 1.3 mL, 1.0 mL, 0.7 mL, 0.5 mL, 0.3 mL, 0.1 mL, 0.05 mL, 0.01 mL, or smaller). Typically, the compositions of the invention can be formulated at a concentration between 1 mg/mL and 500 mg/mL (e.g., between 10 mg/mL and 300 mg/mL, 20 mg/mL and 120 mg/mL, 40 mg/mL and 200 mg/mL, 30 mg/mL and 150 mg/mL, 40 mg/mL and 100 mg/mL, 50 mg/mL and 80 mg/mL, or 60 mg/mL and 70 mg/mL) for subcutaneous administration.

For oral administration, tablets or capsules can be prepared by conventional means with pharmaceutically acceptable excipients such as binding agents, fillers, lubricants, disintegrants, or wetting agents. The tablets can be coated by methods known in the art. Liquid preparations for oral administration can take the form of, for example, solutions, syrups, or suspension, or they can be presented as a dry product for constitution with saline or other suitable liquid vehicle before use. Compositions of the invention for oral administration also can contain pharmaceutically acceptable excipients such as suspending agents, emulsifying agents, non-aqueous vehicles, preservatives, buffer salts, flavoring, coloring, and sweetening agents as appropriate. Preparations for oral administration also can be suitably formulated to give controlled release of the active ingredients.

Enteric coatings can further be used on tablets of the present invention to resist prolonged contact with the strongly acidic gastric fluid, but dissolve in the mildly acidic or neutral intestinal environment. Without being so limited, cellulose acetate phthalate, Eudragit<sup>TM</sup> and hydroxypropyl

methylcellulose phthalate (HPMCP) can be used in enteric coatings of pharmaceutical compositions of the present invention. Cellulose acetate phthalate concentrations generally used are 0.5-9.0% of the core weight. The addition of plasticizers improves the water resistance of this coating material, and formulations using such plasticizers are more effective than when cellulose acetate phthalate is used alone. Cellulose acetate phthalate is compatible with many plasticizers, including acetylated monoglyceride; butyl phthalylbutyl glycolate; dibutyl tartrate; diethyl phthalate; dimethyl phthalate; ethyl phthalylethyl glycolate; glycerin; propylene glycol; triacetin; triacetin citrate; and tripropionin. It is also used in combination with other coating agents such as ethyl cellulose, in drug controlled-release preparations.

The compounds of the invention may be administered in combination with pharmaceutically acceptable, sterile, aqueous or non-aqueous solvents, suspensions or emulsions. Examples of nonaqueous solvents are propylene glycol, polyethylene glycol, vegetable oil, fish oil, and injectable organic esters. Aqueous carriers include water, water-alcohol solutions, emulsions or suspensions, including saline and buffered medical parenteral vehicles including sodium chloride solution, Ringer's dextrose solution, dextrose plus sodium chloride solution, Ringer's solution containing lactose, or fixed oils. Intravenous vehicles may include fluid and nutrient replenishers, electrolyte replenishers, such as those based upon Ringer's dextrose, and the like.

In some embodiments, the pharmaceutical compositions of the present invention can be delivered in a controlled release system. In some embodiments, polymeric materials including polylactic acid, polyorthoesters, cross-linked amphipathic block copolymers and hydrogels, polyhydroxy butyric acid and polydihydropyrans can be used (see also Smolen and Ball, Controlled Drug Bioavailability, Drug product design and performance, 1984, John Wiley & Sons; Ranade and Hollinger, Drug Delivery Systems, pharmacology and toxicology series, 2003, 2<sup>nd</sup> edition, CRC Press). In another embodiment, a pump may be used (Saudek et al., 1989, N. Engl. J. Med. 321: 574).

The compositions of the invention could be formulated in the form of a lyophilized powder using appropriate excipient solutions (e.g., sucrose) as diluents.

Furthermore, cells can be isolated from an individual having a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, e.g., achondroplasia, a bone or cartilage disorder, or a vascular smooth muscle disorder or from an individual that would benefit from bone elongation; transformed with a nucleic acid of the invention; and reintroduced to the afflicted individual (e.g., subcutaneous or intravenous injection). Alternatively, the nucleic acid can be administered directly to the afflicted individual, for example, by injection. The nucleic acid can also be delivered through a vehicle such as a liposome, which can be designed to be targeted to a specific cell type, and engineered to be administered through different routes.

The polypeptides or compositions of the present invention may also be used in combination with at least one other active ingredient to correct, e.g., an achondroplasia phenotype, neurofibromatosis, HPP, or any other disorder or condition described herein.

For combination therapy, two or more of the polypeptides of the present invention (e.g., an sALP polypeptide and an NP polypeptide) are formulated in a variety of ways that are known in the art. For example, the first and second polypeptides may be formulated together or separately. In some embodiments, the first and second polypeptides are formulated together for the simultaneous or 5 near simultaneous administration of the polypeptides. Such co-formulated compositions can include the sALP polypeptide and the NP polypeptide formulated together in the same pill, capsule, liquid, etc.

Administration of each compound in controlled release formulations is useful where the sALP polypeptide or the NP polypeptide, has (i) a narrow therapeutic index (e.g., the difference between the 10 plasma concentration leading to harmful side effects or toxic reactions and the plasma concentration leading to a therapeutic effect is small; generally, the therapeutic index, TI, is defined as the ratio of median lethal dose ( $LD_{50}$ ) to median effective dose ( $ED_{50}$ )); (ii) a narrow absorption window in the gastro-intestinal tract; (iii) a short biological half-life, such as by degradation *in vivo* by NEP and/or IDE; or (iv) the pharmacokinetic profile of each component must be modified to maximize the 15 exposure of the neoplasm to an amount of each agent, together, that is therapeutically effective. Accordingly, a sustained release formulation may be used to avoid frequent dosing that may be required in order to sustain the plasma levels of both agents at a therapeutic level.

Many strategies can be pursued to obtain controlled release in which the rate of release outweighs the rate of metabolism of the therapeutic polypeptide. For example, controlled release can 20 be obtained by the appropriate selection of formulation parameters and ingredients (e.g., appropriate controlled release compositions and coatings). Examples include single or multiple unit tablet or capsule compositions, oil solutions, suspensions, emulsions, microcapsules, microspheres, nanoparticles, patches, and liposomes. The control release mechanism can be such that the compound of the sALP polypeptide is released first, followed by the NP polypeptide, or vice versa. The release 25 mechanism can also be controlled that the two polypeptides are released at period intervals, the release could be simultaneous or a delayed release of one, when release of a particular drug is preferred over the other.

Controlled release formulations may include a degradable or nondegradable polymer, hydrogel, organogel, or other physical construct that modifies the bioabsorption, half life or 30 biodegradation of the agent. The controlled release formulation can be a material that is painted or otherwise applied onto the afflicted site, either internally or externally. In one example, hydrogels, such as those described in U.S. Patent No. 5,626,863 can be used in controlled release formulations of compositions of the invention. These biodegradable polymers can be tailored to degrade at a desired rate and with a desired kinetics by selecting the appropriate monomers, method of preparation and 35 molecular weight. Differences in crystallinity of the monomer can alter the polymeric degradation rate. Due to the relatively hydrophobic nature of most polymers, actual mass loss can begin with the

oligomeric fragments that are small enough to be water soluble; hence, even the initial molecular weight can influence the degradation rate.

The individually or separately formulated polypeptides can be packaged together as in a kit. Non-limiting examples include kits that contain, e.g., two pills, a pill and a powder, a suppository and 5 a liquid in a vial, two topical creams, among others. The kit can include optional components that aid in the administration of the unit dose to patients, such as vials for reconstituting powder forms, syringes for injection or subcutaneous administration, customized IV delivery systems, inhalers, among others. Additionally, the unit dose kit can contain instructions for preparation and administration of the compositions. The kit may be manufactured as a single use unit dose for one 10 subject, multiple uses for a particular subject (at a constant dose or in which the individual polypeptides may vary in potency as therapy progresses); or the kit may contain multiple doses suitable for administration to multiple subjects (“bulk packaging”). The kit components may be assembled in cartons, blister packs, bottles, tubes, and the like.

15 ***Gene Therapy***

The polypeptides described herein could also be advantageously delivered through gene therapy, where an exogenous nucleic acid encoding the proteins is delivered to tissues of interest and expressed *in vivo*. Gene therapy methods are discussed, e.g., in Verme et al. (*Nature* 389:239-242, 1997), Yamamoto et al. (*Molecular Therapy* 17:S67-S68, 2009), and Yamamoto et al., (*J. Bone 20 Miner. Res.* 26:135-142, 2011), each of which is hereby incorporated by reference. Both viral and non-viral vector systems can be used. The vectors may be, for example, plasmids, artificial chromosomes (e.g., bacterial, mammalian, or yeast artificial chromosomes), virus or phage vectors provided with an origin of replication, and optionally, a promoter for the expression of the nucleic acid encoding the viral polypeptide and optionally, a regulator of the promoter. The vectors may 25 contain one or more selectable marker genes, for example, an ampicillin or kanamycin resistance gene in the case of a bacterial plasmid or a resistance gene for a fungal vector. Vectors may be used in *in vitro*, for example, for the production of DNA, RNA, or the viral polypeptide, or may be used to transfect or transform a host cell, for example, a mammalian host cell, e.g., for the production of the viral polypeptide encoded by the vector. The vectors may also be adapted to be used *in vivo*, for 30 example, in a method of vaccination or gene therapy.

Examples of suitable viral vectors include, retroviral, lentiviral, adenoviral, adeno-associated viral, herpes viral, including herpes simplex viral, alpha-viral, pox viral, such as Canarypox and vaccinia-viral based systems. Gene transfer techniques using these viruses are known in the art. Retrovirus vectors, for example, may be used to stably integrate the nucleic acids of the invention into 35 the host genome. Replication-defective adenovirus vectors by contrast remain episomal and therefore allow transient expression. Vectors capable of driving expression in insect cells (e.g., baculovirus vectors), in human cells, yeast, or in bacteria may be employed in order to produce quantities of the

viral polypeptide(s) encoded by the nucleic acids of the invention, for example, for use in subunit vaccines or in immunoassays. Useful gene therapy methods include those described in WO 06/060641, U.S. Pat. No. 7,179,903 and WO 01/36620 (each of which is hereby incorporated by reference), which use an adenovirus vector to target a nucleic acid of interest to hepatocytes as protein producing cells.

In an additional example, a replication-deficient simian adenovirus vector may be used as a live vector. These viruses contain an E1 deletion and can be grown on cell lines that are transformed with an E1 gene. Examples of these replication-deficient simian adenovirus vectors are described in U.S. Patent No. 6,083,716 and WO 03/046124 (each of which is hereby incorporated by reference).

10 These vectors can be manipulated to insert a nucleic acid of the invention, such that the encoded viral polypeptide(s) may be expressed.

Promoters and other expression regulatory signals may be selected to be compatible with the host cell for which expression is designed. For example, mammalian promoters include the metallothionein promoter, which can be induced in response to heavy metals such as cadmium, and 15 the  $\beta$ -actin promoter. Viral promoters, such as the SV40 large T antigen promoter, human cytomegalovirus (CMV) immediate early (1E) promoter, rous sarcoma virus LTR promoter, adenovirus promoter, or a HPV promoter, particularly the HPV upstream regulatory region (URR) may also be used. All these promoters, as well as additional promoters, are well-described in the art.

The nucleic acid molecules described herein may also be administered using non-viral based 20 systems. For example, these administration systems include microsphere encapsulation, poly(lactide-co-glycolide), nanoparticle, and liposome-based systems. Non-viral based systems also include techniques facilitating the delivery of “naked” polynucleotides (such as electroporation, “gene gun” delivery and various other techniques used for the introduction of polynucleotides).

The introduced polynucleotide can be stably or transiently maintained in the host cell. Stable 25 maintenance typically requires that the introduced polynucleotide either contains an origin of replication compatible with the host cell or integrates into a replicon of the host cell such as an extrachromosomal replicon (e.g., a plasmid) or a nuclear or mitochondrial chromosome.

### Dosage

30 Any amount of a polypeptide or a pharmaceutical composition of the invention can be administered to a subject. The dosages will depend on many factors, including the mode of administration and the age of the subject. Typically, the amount of the composition of the invention contained within a single dose will be an amount that is effective to treat a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, a bone or cartilage disorder, or a vascular smooth 35 muscle disorder, or to elongate bone, without inducing significant toxicity. For example, the polypeptides described herein can be administered to subjects in individual doses ranging, e.g., from 0.01 mg/kg to 500 mg/kg (e.g., from 0.05 mg/kg to 500 mg/kg, from 0.2 mg/kg to 20 mg/kg, from 5

mg/kg to 500 mg/kg, from 0.1 mg/kg to 100 mg/kg, from 10 mg/kg to 100 mg/kg, from 0.1 mg/kg to 50 mg/kg, 0.5 mg/kg to 25 mg/kg, 1.0 mg/kg to 10 mg/kg, 1.5 mg/kg to 5 mg/kg, or 2.0 mg/kg to 3.0 mg/kg) or from 1  $\mu$ g/kg to 1,000  $\mu$ g/kg (e.g., from 5  $\mu$ g/kg to 1,000  $\mu$ g/kg, from 1  $\mu$ g/kg to 750  $\mu$ g/kg, from 5  $\mu$ g/kg to 750  $\mu$ g/kg, from 10  $\mu$ g/kg to 750  $\mu$ g/kg, from 1  $\mu$ g/kg to 500  $\mu$ g/kg, from 5  $\mu$ g/kg to 500  $\mu$ g/kg, from 10  $\mu$ g/kg to 500  $\mu$ g/kg, from 1  $\mu$ g/kg to 100  $\mu$ g/kg, from 5  $\mu$ g/kg to 100  $\mu$ g/kg, from 10  $\mu$ g/kg to 100  $\mu$ g/kg, from 1  $\mu$ g/kg to 50  $\mu$ g/kg, from 5  $\mu$ g/kg to 50  $\mu$ g/kg, or from 10  $\mu$ g/kg to 50  $\mu$ g/kg). Exemplary doses include, e.g., 0.01, 0.05, 0.1, 0.5, 1, 2, 2.5, 5, 10, 20, 25, 50, 100, 125, 150, 200, 250, or 500 mg/kg; or 1, 2, 2.5, 5, 10, 20, 25, 50, 100, 125, 150, 200, 250, 500, 750, 900, or 1,000  $\mu$ g/kg. For all dosages or ranges recited herein, the term “about” may be used to modify these dosages by  $\pm 10\%$  of the recited values or range endpoints.

Doses can also be adjusted when two or more polypeptides or compositions of the inventions are being administered. Exemplary doses include an sALP polypeptide (e.g., an sALP fusion protein) present in a dosage between about 0.2 mg/kg to about 20 mg/kg and an NP polypeptide (e.g., an NP fusion protein) present in a dosage of about 0.5 mg/kg to about 500 mg/kg. In particular 15 embodiments, the dose of each individual polypeptide or composition is lower than the therapeutic dose of a single polypeptide or single composition when administered alone.

Doses can be administered, e.g., hourly, bihourly, daily, bidaily, twice a week, three times a week, four times a week, five times a week, six times a week, weekly, biweekly, monthly, bimonthly, or yearly. Alternatively, doses can be administered, e.g., twice, three times, four times, five times, six 20 times, seven times, eight times, nine times, 10 times, 11 times, or 12 times per day. In particular embodiments, the dosing regimen is once weekly. The duration of the dosing regimen can be, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, or 30 day(s), week(s), or month(s), or even for the remaining lifespan of the subject. The amount, frequency, and duration of dosage will be adapted by the clinician in accordance with conventional 25 factors such as the extent of the disease and different parameters from the subject.

The nucleic acids of the invention can be administered according the formulations described herein to a patient in dosages suitable for gene therapy. The amount of the nucleic acids administered will depend on a number of factors known to those skilled in the art, including: the length and nature 30 of the nucleic acid, the vector (e.g., viral or non-viral) used, the activity of the polypeptide encoded, the presence of excipients, the route and method of administration, and the general condition and fitness of the subject. Exemplary dosages and routes of administration are described, e.g., in Melman et al. (*Isr. Med. Assoc. J.* 9:143-146, 2007; describing the intrapenile injection of 0.5 mg to 7.5 mg of a human cDNA in a plasmid for treating erectile dysfunction), Powell et al. (*Circulation* 118:58-65, 2008; describing the intramuscular injection of 0.4 mg to 4.0 mg of a hepatocyte growth factor 35 plasmid to treat critical limb ischemia, Waddill et al. (*AJR Am. J. Roentgenol.* 169:63-67, 1997; describing the CT-guided intratumoral injection of 0.01 mg to 0.25 mg of plasmid DNA encoding an MHC antigen to treat melanoma), Kastrup et al. (*J. Am. Coll. Cardiol.* 45:982-988, 2005; describing

the intramyocardial injection of 0.5 mg of a VEGF plasmid to treat severe angina pectoris), and Romero et al. (*Hum. Gene Ther.* 15:1065-1076, 2004; describing the intramuscular injection of 0.2 mg to 0.6 mg of a plasmid to treat Duchenne/Becker muscular dystrophy), each of which is hereby incorporated by reference.

5 In certain embodiments, the nucleic acids of the invention can be administered to the subject at a dose in the range from, e.g., 0.01 mg to 100 mg (e.g., from 0.05 mg to 50 mg, 0.1 mg to 10 mg, 0.3 mg to 3 mg, or about 1 mg) of nucleic acid. The total volume at which the nucleic acid can be administered will depend on its concentration, and can range from, e.g., 1  $\mu$ L to 10 mL (e.g. from 10  $\mu$ L to 1 mL, 50  $\mu$ L to 500  $\mu$ L, 70  $\mu$ L to 200  $\mu$ L, 90  $\mu$ L to 150  $\mu$ L, or 100  $\mu$ L to 120  $\mu$ L).

10 The nucleic acids can be administered, e.g., hourly, bihourly, daily, bidaily, twice a week, three times a week, four times a week, five times a week, six times a week, weekly, biweekly, monthly, bimonthly, or yearly. Alternatively, the nucleic acids can be administered, e.g., twice, three times, four times, five times, six times, seven times, eight times, nine times, 10 times, 11 times, or 12 times per day. The duration of the dosing regimen can be, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 15 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, or 30 day, weeks, or months, or even for the remaining lifespan of the subject.

20 These are guidelines, since the actual dose should be carefully selected and titrated by an attending physician or nutritionist based upon clinical factors unique to each subject. The optimal periodic dose will be determined by methods known in the art and will be influenced by factors such as the age of the subject, as indicated above, and other clinically relevant factors. In addition, subjects 25 may be taking medications for other diseases or conditions. The other medications may be continued during the time that a polypeptide or nucleic acid of the invention is given to the subject, but it is advisable in such cases to begin with low doses to determine if adverse side effects are experienced.

## 25 EXAMPLES

The following examples are provided for the purpose of illustrating the invention and are not meant to limit the invention in any way.

### **Example 1. Characterization of Osteoblasts in Mice lacking NF1**

30 To determine the effect of NF1 on bone matrix mineralization, mice lacking NF1 in osteochondroprogenitor cells were developed and characterized. These mice displayed skeletal dysplasia defects similar to patients with neurofibromatosis type I, where these defects included progressive scoliosis and kyphosis, tibial bowing and abnormalities in skull and anterior chest wall formation. In particular, NF1<sub>col2</sub><sup>-/-</sup> osteoblasts secreted increased levels of (PP<sub>i</sub>) as compared to 35 osteoblasts from wild-type mice (Fig. 1A). Bone marrow adherent stromal cells (BMSCs) extracted from adult NF1<sub>col2</sub><sup>-/-</sup> mice and grown *in vitro* under osteogenic medium form less alkaline phosphatase (AP)-positive and less mineralized (alizarin red-positive) nodules compared to BMSCs extracted from

wild-type mice, which is accompanied by an increased amount of PP<sub>i</sub> released in the medium over 24 hours. Accumulation of PP<sub>i</sub> prevents proper bone matrix mineralization and likely contributes, at least in part, to defects observed in NF1<sub>col2</sub><sup>-/-</sup> mice. Thus, compounds that reduce PP<sub>i</sub> accumulation, such as sALP or an sALP analog, could be useful for treating NF1.

5 In addition, NF1<sub>col2</sub><sup>-/-</sup> osteoblasts expressed increased levels of mRNA for the progressive ankylosis gene (*Ank*) as compared to osteoblasts from wild-type mice (Fig. 1B). Treatment with a dual-specificity MEK1/MEK2 kinase inhibitor (U0126) provided decreased levels of *Ank* mRNA expression in NF1<sub>col2</sub><sup>-/-</sup> osteoblasts as compared to vehicle, and these observed levels were similar to that in wild-type osteoblasts (Fig. 1B, second and third bars). Thus, compounds that inhibit a kinase pathway, such as NP or an NP analog, could be useful for treating NF1.

10 Taken together, these data suggest that an sALP polypeptide alone, an NP polypeptide alone, or the combination of both an sALP polypeptide and an NP polypeptide could be useful for treating any neurocutaneous syndrome with bone manifestations, such as neurofibromatosis, or any other disorder described herein.

15

#### **Example 2. Combination Therapy for the Treatment of Neurofibromatosis**

Fig. 2 provides a hypothetical working model for defective bone matrix mineralization in NF1<sub>col2</sub><sup>-/-</sup> mice, which can include multiple imbalances (e.g., increase of PP<sub>i</sub> or overactivation of one or more kinases, such as Ras or ERK) that contribute to the disease. Without wishing to be limited by theory, accumulation of PP<sub>i</sub> could be minimized by using any of the compositions and methods described herein including a soluble alkaline phosphatase (sALP) or sALP analog (see, e.g., the polypeptide of SEQ ID NO: 1204). Furthermore, overactivation of one or more kinases could be controlled by using any of the compositions and methods described herein including an NP or NP analog. As described herein, the intracellular production of cGMP resulting from NPR-B activation is known to inhibit the MAP-kinase pathway. Thus, an NP or NP analog that could activate the NPR-B signaling pathway can be used for the treatment of neurocutaneous syndromes, such as neurofibromatosis. Accordingly, the combination of an sALP or sALP analog (e.g., an sALP polypeptide) with an NP or NP analog (e.g., an NP polypeptide) could be particularly useful for treating such diseases.

20

#### **Example 3. *In vitro* and *In vivo* Effects of sTNALP-FcD<sub>10</sub> on NF1<sub>col2</sub><sup>-/-</sup> Phenotype**

To assess the effect of sALP polypeptides on NF1 phenotype, cultures of osteoblasts from NF1<sub>col2</sub><sup>-/-</sup> mice were treated with either bone morphogenetic protein 2 (BMP2) or the sALP fusion polypeptide of sTNALP-FcD<sub>10</sub> (SEQ ID NO: 1204).

25

Both wild-type and NF1<sub>col2</sub><sup>-/-</sup> osteoblasts were treated with increasing concentrations of recombinant human BMP2 (rhBMP2, Fig. 3A). In NF1<sub>col2</sub><sup>-/-</sup> osteoblasts, rhBMP2 rescued the differentiation defect, as evidenced by an increased presence of alkaline phosphatase upon increasing

doses of rhBMP2 (Fig. 3A, second row). However, increased mineralization was not observed, as evidenced by the lack of calcium deposition (as indicated by the lack of alizarin red S staining) (Fig. 3A, fourth row).

5 In contrast, treatment with sTNALP-FcD<sub>10</sub> rescued the mineralization defect that is present in NF1<sub>col2</sub><sup>-/-</sup> osteoblasts (Fig. 3B). Increasing doses of sTNALP-FcD<sub>10</sub> provided increased calcific deposition in a dose-dependent manner (Fig. 3B, bottom, and Fig. 3C).

Furthermore, *in vitro* targeted deletion of the NF1 gene in BMSCs resulted in significant reduction of mineralization, which is at least partially rescued by treatment with 0.5 µg/mL of sTNALP-FcD<sub>10</sub> (Fig. 3D).

10 *In vivo* experiments were also conducted with sTNALP-FcD<sub>10</sub>. NF1<sub>col2</sub><sup>-/-</sup> mice were treated from day 1 to day 18 with 8.2 mg/kg of sTNALP-FcD<sub>10</sub>. Treatment of mice with sTNALP-FcD<sub>10</sub> increased bone mineral density deficit in NF1<sub>col2</sub><sup>-/-</sup> mice compared to vehicle (\* p < 0.5), as determined by the ratio of mineralized bone volume (BV) to total bone volume (TV) (Fig. 3E).

15 Accordingly, any sALP polypeptide described herein, either alone or in combination with any NP polypeptide described herein, could be particularly useful for treating neurofibromatosis or any neurocutaneous syndrome with bone manifestations.

**Example 4. *In vitro* and *In vivo* Effects of NC2-KGANKK on NF1<sub>col2</sub><sup>-/-</sup> Phenotype**

20 To assess the effect of NP polypeptides on NF1 phenotype, expression levels of the *NPR-B* gene were assessed in NF1<sub>col2</sub><sup>-/-</sup> mice (Fig. 4A). *NPR-B* was expressed in BMSCs at all stages of differentiation, where lack of *NF1* expression did not affect *NPR-B* expression.

25 Additional experiments were conducted in which cultures of chondrocytes from NF1<sub>col2</sub><sup>-/-</sup> mice were treated with the NP fusion polypeptide of NC2-KGANKK (SEQ ID NO: 512). Rib primary chondrocytes from P0 mice were obtained from wild-type mice and NF1<sub>col2</sub><sup>-/-</sup> mice and treated for 30 minutes with increasing concentrations of NC2-KGANKK. Without any treatment, increased levels of phosphorylated ERK (p-ERK) were observed in NF1<sub>col2</sub><sup>-/-</sup> chondrocytes, as compared to levels in wild-type chondrocytes (Fig. 4B). After treatment with NC2-KGANKK, decreased levels of p-ERK were observed in NF1<sub>col2</sub><sup>-/-</sup> chondrocytes. These results support the use of 30 NP polypeptides, such as NC2-KGANKK, to inhibit the overactivation of one or more kinases, such as ERK.

*In vivo* experiments were also conducted. NF1<sub>col2</sub><sup>-/-</sup> mice were treated with NC2B (SEQ ID NO: 504). Treatment at least partially rescued the body length defect in NF1<sub>col2</sub><sup>-/-</sup> mice (Fig. 4C), the bone growth defect in NF1<sub>col2</sub><sup>-/-</sup> mice (Fig. 4D), and the proliferative and hypertrophic chondrocyte zone defects in NF1<sub>col2</sub><sup>-/-</sup> mice (Fig. 4E).

35 Accordingly, any NP polypeptide described herein, either alone or in combination with an sALP polypeptide described herein, could be particularly useful for treating disorders associated with

overactivation of one or more kinases, such as neurofibromatosis or any neurocutaneous syndrome with bone manifestations.

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### Other Embodiments

All publications, patents, and patent applications mentioned in the above specification are hereby incorporated by reference. Various modifications and variations of the described method and system of the invention will be apparent to those skilled in the art without departing from the scope and spirit of the invention. Although the invention has been described in connection with specific

embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention that are obvious to those skilled in the art are intended to be within the scope of the invention.

5        Other embodiments are in the claims.

**CLAIMS**

What is claimed is:

1. A method of treating a neurocutaneous syndrome in a subject, said method comprising administering to said subject a therapeutically effective amount of a pharmaceutical composition comprising:
  - (a) a polypeptide comprising the structure A-sALP-B; and
  - (b) a pharmaceutically acceptable excipient,  
wherein sALP is the extracellular domain of an alkaline phosphatase, A is absent or is an amino acid sequence of at least one amino acid, and B is absent or is an amino acid sequence of at least one amino acid, thereby treating said syndrome in said subject.
2. The method of claim 1, wherein the amino acid sequence of said polypeptide comprises the amino acid sequence of SEQ ID NOS: 1204 or 1221.
3. The method of claim 1, wherein the amino acid sequence of said sALP comprises amino acid residues 23-508 of SEQ ID NO: 1215, amino acid residues 18-498 of SEQ ID NO: 1216, amino acid residues 23-508 of SEQ ID NO: 1218, or amino acid residues 18-498 of SEQ ID NO: 1219, or the amino acid sequence of said sALP consists of amino acid residues 23-512 of SEQ ID NO: 1215, amino acid residues 18-502 of SEQ ID NO: 1216, amino acid residues 23-512 of SEQ ID NO: 1218, or amino acid residues 18-502 of SEQ ID NO: 1219.
4. The method of claim 1 or 3, wherein the amino acid sequence of said sALP comprises an amino acid sequence having at least 85% sequence identity to SEQ ID NO: 1205, or at least 95% sequence identity to SEQ ID NO: 1205, or at least 99% sequence identity to SEQ ID NO: 1205.
5. The method of claim 4, wherein the amino acid sequence of said sALP comprises or consists of the amino acid sequence of SEQ ID NO: 1205.
6. The method of any one of claims 1-5, wherein A and/or B are absent.

7. The method of any one of claims 1-6, wherein A or B comprises a fragment crystallizable region (Fc).

8. The method of any one of claims 1-7, wherein said Fc comprises a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region, or wherein said Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, and IgG-4.

9. The method of claim 8, wherein the amino acid sequence of said Fc comprises an amino acid sequence having at least 85% sequence identity to SEQ ID NO: 401, or at least 95% sequence identity to SEQ ID NO: 401, or at least 99% sequence identity to SEQ ID NO: 401.

10. The method of claim 9, wherein the amino acid sequence of said Fc comprises or consists of the amino acid sequence of SEQ ID NO: 401.

11. The method of any one of claims 1-10, wherein A or B comprises I<sub>n</sub>, and wherein I represents an aspartic acid or a glutamic acid and n=10 to 16.

12. The method of any one of claims 1-10, wherein said polypeptide does not comprise a polyaspartic acid or polyglutamic acid region longer than three consecutive aspartic acid or glutamic acid residues, or wherein said polypeptide does not comprise a polyaspartic acid or polyglutamic acid region longer than two consecutive aspartic acid or glutamic acid residues.

13. The method of any one of claims 1-10, wherein said polypeptide does not comprise a bone-targeting moiety.

14. The method of any one of claims 1-10, wherein said polypeptide comprises the structure C-sALP-D-Fc-E or the structure C-Fc-D-sALP-E,

C is absent or is an amino acid sequence of at least one amino acid,

D is absent or is an amino acid sequence of at least one amino acid, and

E is absent or is an amino acid sequence of at least one amino acid.

15. The method of claim 14, wherein C and/or E are absent.

16. The method of any one of claims 1-10, wherein said polypeptide comprises the structure C-sALP-D-Fc-G-I<sub>n</sub>-H or the structure C-Fc-D-sALP-G-I<sub>n</sub>-H,

C is absent or is an amino acid sequence of at least one amino acid,

D is absent or is an amino acid sequence of at least one amino acid,

G is absent or is an amino acid sequence of at least one amino acid,

H is absent or is an amino acid sequence of at least one amino acid,

I represents an aspartic acid or a glutamic acid, and n=10 to 16.

17. The method of claim 16, wherein C and/or H are absent.

18. The method of claim 16 or 17, wherein G is two amino acid residues.

19. The method of claim 18, wherein G is aspartic acid-isoleucine.

20. The method of any one of claims 16-19, wherein I is aspartic acid and n=10.

21. The method of any one of claims 14-20, wherein D is two amino acid residues.

22. The method of claim 21, wherein D is leucine-lysine.

23. The method of any one of claims 1-22, wherein the amino acid sequence of said polypeptide consists of the amino acid sequence of SEQ ID NOs: 1201, 1204, 1220, or 1221.

24. The method of claim 23, wherein the amino acid sequence of said polypeptide consists of the amino acid sequence of SEQ ID NO: 1204.

25. A method of treating a neurocutaneous syndrome in a subject, said method comprising administering to said subject a therapeutically effective amount of a pharmaceutical composition comprising:

(a) a polypeptide comprising the structure V-NP-W; and

(b) a pharmaceutically acceptable excipient,

wherein NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),

V is absent or is an amino acid sequence of at least one amino acid, and

W is absent or is an amino acid sequence of at least one amino acid,

thereby treating said syndrome in said subject.

26. The method of claim 25, wherein the amino acid sequence of said polypeptide comprises the amino acid sequence of SEQ ID NOs: 504, 512, 530, or 572.

27. The method of claim 25, wherein said NP comprises the structure:  
[N-terminal extension]-[short segment]-[ring domain]-[C-terminal extension],  
wherein said ring domain comprises the amino acid sequence of SEQ ID NO: 6, amino acid residues 11-27 of SEQ ID NO: 30, or SEQ ID NO: 95, and each of said N-terminal extension, short segment, and C-terminal extension is, independently, absent or is an amino acid sequence of at least one amino acid.

28. The method of claim 27, wherein said ring domain comprises amino acid residues 6-22 of SEQ ID NO: 126.

29. The method of claim 28, wherein the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Val, Ala, Ser, Glu, Arg, Tyr, Cys, Pro, or Asp.

30. The method of claim 27, wherein said ring domain comprises the amino acid sequence of SEQ ID NO: 12.

31. The method of any one of claims 27-30, wherein said short segment and said ring domain together comprise the amino acid sequence of any one of SEQ ID NOs: 4, 13-30, 119-122, 126, or 156-161.

32. The method of any one of claims 27-31, wherein the amino acid sequence of said short segment consists of amino acid residues 1-5, 2-5, 3-5, 4-5, or 5 of SEQ ID NO: 4, amino acid residues 1-10 of SEQ ID NO: 17, amino acid residues 1-5 of SEQ ID NO: 19, amino acid residues 1-3 of SEQ ID NO: 20, amino acid residues 1-5 of SEQ ID NO: 21, or amino acid residues 1-6 of SEQ ID NO: 29.

33. The method of any one of claims 27-32, wherein the amino acid sequence of said N-terminal extension comprises amino acid residues 1-31 or 17-31 of SEQ ID NO: 11.

34. The method of any one of claims 27-32, wherein the amino acid sequence of said N-terminal extension comprises KGANKK (SEQ ID NO: 314) or KGANQK (SEQ ID NO: 315).

35. The method of claim 27, wherein said N-terminal extension, short segment, and ring domain together comprise the amino acid sequence of SEQ ID NO: 11.

36. The method of any one of claims 27-35, wherein said C-terminal extension comprises the amino acid sequence of SEQ ID NOs: 117 or 118 or comprises amino acid residues 23-37 selected from any one of SEQ ID NOs: 101-116.

37. The method of claim 27, wherein the amino acid sequence of said NP consists of SEQ ID NOs: 4 or 11, or the amino acid sequence of any one of SEQ ID NOs: 31-94, or a fragment thereof comprising at least a ring domain, or the amino acid sequence of any one of SEQ ID NOs: 13-29, 100-116, 119-125, 127-233, or 1001-1155.

38. The method of any one of claims 27-37, wherein V and/or W are absent.

39. The method of any one of claims 27-38, wherein V or W comprises a fragment crystallizable region (Fc).

40. The method of claim 39, wherein said Fc comprises a C<sub>H2</sub> domain, a C<sub>H3</sub> domain, and a hinge region, or wherein said Fc is a constant domain of an immunoglobulin selected from the group consisting of IgG-1, IgG-2, IgG-3, and IgG-4.

41. The method of claim 40, wherein the amino acid sequence of said Fc comprises an amino acid sequence having at least 85% sequence identity to SEQ ID NO: 401, or at least 95% sequence identity to SEQ ID NO: 401, or at least 99% sequence identity to SEQ ID NO: 401.

42. The method of claim 41, wherein the amino acid sequence of said Fc comprises or consists of the amino acid sequence of SEQ ID NO: 401.

43. The method of any one of claims 27-42, wherein V or W comprises a glycine-rich region.

44. The method of claim 43, wherein the amino acid sequence of V or W consists of one or more glycines and one or more serines.

45. The method of claim 43 or 44, wherein the amino acid sequence of V or W comprises  $[(\text{Gly})_m(\text{Ser})]_n(\text{Gly})_p$  or  $(\text{Gly})_p[(\text{Ser})(\text{Gly})_m]_n$ , and wherein each of m, n, and p is, independently, between 0 and 20.

46. The method of claim 45, wherein m is between 1 and 6; n is between 1 and 10; and p is between 0 and 4.

47. The method of claim 46, wherein m is 4 and n is 1-6.

48. The method of claim 46, wherein combinations of m, n, and p are selected from a single row of Table 2, or wherein the amino acid sequence of V or W comprises the amino acid sequence of any one of SEQ ID NOS: 301-391.

49. The method of any one of claims 27-48, wherein V or W does not comprise a bone-targeting moiety.

50. The method of any one of claims 27-48, wherein V or W comprises a bone-targeting moiety.

51. The method of claim 50, wherein said bone-targeting moiety comprises six consecutive acidic residues.

52. The method of claim 51, wherein said bone-targeting moiety comprises ten consecutive acidic residues.

53. The method of claim 51 or 52, wherein said acidic residues are aspartic acid or glutamic acid.

54. The method of claim 53, wherein said bone-targeting moiety comprises  $E_6$ ,  $E_{10}$ ,  $D_6$ , or  $D_{10}$ .

55. The method of any one of claims 27-54, wherein V or W comprises a cathepsin cleavage sequence.

56. The method of claim 55, wherein said cathepsin cleavage sequence comprises a cathepsin K cleavage sequence.

57. The method of claim 55 or 56, wherein said cathepsin cleavage sequence is HGPQG (SEQ ID NO: 374) or HKLRG (SEQ ID NO: 375).

58. The method of any one of claims 27-57, wherein said polypeptide comprises the structure V-NP-W,

NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), each of V and W is, independently, absent or is an amino acid sequence of at least one amino acid, and

said NP comprises the amino acid sequence of any one of SEQ ID NOs: 17-29, 31-40, 42-94, 101-116, 119-122, 128-161, or 163-233, or V or W comprises the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

59. The method of any one of claims 27-57, wherein said polypeptide comprises the structure V-NP or NP-W,

NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and each of V and W comprises, independently, the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

60. The method of any one of claims 27-57, wherein said polypeptide comprises the structure X-Fc-Y-NP-Z or the structure X-NP-Y-Fc-Z,

NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and each of X, Y, and Z is, independently, absent or is an amino acid sequence of at least one amino acid.

61. The method of claim 60, wherein Y comprises a glycine-rich region.

62. The method of claim 61, wherein the amino acid sequence of Y consists of one or more glycines and one or more serines.

63. The method of claim 62, wherein the amino acid sequence of Y comprises  $[(\text{Gly})_m(\text{Ser})]_n(\text{Gly})_p$  or  $(\text{Gly})_p[(\text{Ser})(\text{Gly})_m]_n$ , and wherein each of m, n, and p is, independently, between 0 and 20.

64. The method of any one of claims 60-63, wherein X is absent, Z is absent, or X and Z are both absent.

65. The method of any one of claims 60-64, wherein X, Y, or Z comprises a bone-targeting moiety.

66. The method of claim 65, wherein said bone-targeting moiety comprises six or ten consecutive acidic residues.

67. The method of claim 66, wherein said acidic residues are aspartic acid or glutamic acid.

68. The method of any one of claims 65-67, wherein said bone-targeting moiety comprises E<sub>6</sub>, E<sub>10</sub>, D<sub>6</sub>, or D<sub>10</sub>.

69. The method of any one of claims 60-68, wherein X, Y, or Z comprises a cathepsin cleavage sequence.

70. The method of claim 69, wherein said cathepsin cleavage sequence comprises a cathepsin K cleavage sequence.

71. The method of any one of claims 25-70, wherein the amino acid sequence of said polypeptide comprises the amino acid sequence of any one of SEQ ID NOS: 501-608.

72. The method of claim 71, wherein the amino acid sequence of said polypeptide comprises the amino acid sequence of any one of SEQ ID NOS: 502, 504, 506, 512, 514, 516, 530, 560, 562, 564, 572, 574, 576, 584, 586, 588, 596, 598, 600, or 608.

73. The method of claim 72, wherein the amino acid sequence of said polypeptide comprises or consists of the amino acid sequence of SEQ ID NOs: 504, 512, 530, or 572.

74. The method of any one of claims 1-73, wherein said polypeptide is in dimeric form.

75. The method of any one of claims 1-74, wherein said polypeptide is glycosylated or pegylated.

76. The method of any one of claims 1-75, wherein said pharmaceutical composition is administered in a dosage between about 0.2 mg/kg to about 20 mg/kg of said polypeptide.

77. The method of any one of claims 1-75, wherein said pharmaceutical composition is administered in a dosage between about 0.5 mg/kg to about 500 mg/kg of said polypeptide.

78. The method of any one of claims 1-75, wherein said pharmaceutical composition is administered in a dosage between about 10 µg/kg to about 1,000 µg/kg of said polypeptide.

79. The method of any one of claims 1-78, wherein said pharmaceutical composition is administered subcutaneously.

80. The method of any one of claims 1-79, wherein said pharmaceutical composition is administered one time, two times, or three times per week.

81. The method of any one of claims 1-80, wherein said subject is human.

82. The method of any one of claims 1-81, wherein said neurocutaneous syndrome is neurofibromatosis type I.

83. A composition comprising a first polypeptide and a second polypeptide, wherein

a) said first polypeptide comprises the structure A-sALP-B, wherein

i) sALP is the extracellular domain of an alkaline phosphatase,

ii) A is absent or is an amino acid sequence of at least one amino acid, and

iii) B is absent or is an amino acid sequence of at least one amino acid; and

b) said second polypeptide comprises the structure V-NP-W, wherein

i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),

- ii) V is absent or is an amino acid sequence of at least one amino acid, and
- iii) W is absent or is an amino acid sequence of at least one amino acid.

84. The composition of claim 83, wherein the amino acid sequence of said first polypeptide comprises the amino acid sequence of SEQ ID NOs: 1204 or 1221 and the amino acid sequence of said second polypeptide comprises the amino acid sequence of SEQ ID NOs: 504, 512, 530, or 572.

85. The composition of claim 83, wherein the amino acid sequence of said sALP of said first polypeptide comprises amino acid residues 23-508 of SEQ ID NO: 1215, amino acid residues 18-498 of SEQ ID NO: 1216, amino acid residues 23-508 of SEQ ID NO: 1218, or amino acid residues 18-498 of SEQ ID NO: 1219, or

the amino acid sequence of said sALP of said first polypeptide consists of amino acid residues 23-512 of SEQ ID NO: 1215, amino acid residues 18-502 of SEQ ID NO: 1216, amino acid residues 23-512 of SEQ ID NO: 1218, or amino acid residues 18-502 of SEQ ID NO: 1219, or

the amino acid sequence of said sALP of said first polypeptide comprises an amino acid sequence having at least 85% sequence identity to SEQ ID NO: 1205, or at least 95% sequence identity to SEQ ID NO: 1205, or at least 99% sequence identity to SEQ ID NO: 1205.

86. The composition of claim 83 or 85, wherein A and/or B of said first polypeptide are absent.

87. The composition of any one of claims 83-86, wherein A or B of said first polypeptide comprises a fragment crystallizable region (Fc).

88. The composition of any one of claims 83-87, wherein A or B of said first polypeptide comprises I<sub>n</sub>, and wherein I represents an aspartic acid or a glutamic acid and n=10 to 16.

89. The composition of any one of claims 83-88, wherein said first polypeptide comprises the structure C-sALP-D-Fc-G-I<sub>n</sub>-H,

C is absent or is an amino acid sequence of at least one amino acid,

D is absent or is an amino acid sequence of at least one amino acid,

G is absent or is an amino acid sequence of at least one amino acid,

H is absent or is an amino acid sequence of at least one amino acid,

I represents an aspartic acid or a glutamic acid, and n=10 to 16.

90. The composition of any one of claims 83-89, wherein the amino acid sequence of said first polypeptide comprises or consists of the amino acid sequence of SEQ ID NO: 1204.

91. The composition of any one of claims 83-90, wherein said NP of said second polypeptide comprises the structure:

[N-terminal extension]-[short segment]-[ring domain]-[C-terminal extension],  
wherein said ring domain comprises the amino acid sequence of SEQ ID NO: 6, amino acid residues 11-27 of SEQ ID NO: 30, or SEQ ID NO: 95, and each of said N-terminal extension, short segment, and C-terminal extension is, independently, absent or is an amino acid sequence of at least one amino acid.

92. The composition of claim 91, wherein said ring domain comprises amino acid residues 6-22 of SEQ ID NO: 126.

93. The composition of claim 92, wherein the amino acid at position 17 of SEQ ID NO: 126 is Phe, Leu, Ile, Thr, Val, Ala, Ser, Glu, Arg, Tyr, Cys, Pro, or Asp.

94. The composition of any one of claims 91-93, wherein the amino acid sequence of said N-terminal extension comprises amino acid residues 1-31 or 17-31 of SEQ ID NO: 11, KGANKK (SEQ ID NO: 314), or KGANQK (SEQ ID NO: 315).

95. The composition of any one of claims 91-94, wherein said C-terminal extension comprises the amino acid sequence of SEQ ID NOs: 117 or 118 or comprises amino acid residues 23-37 selected from any one of SEQ ID NOs: 101-116.

96. The composition of claim 91, wherein the amino acid sequence of said NP consists of SEQ ID NOs: 4 or 11, or the amino acid sequence of any one of SEQ ID NOs: 31-94, or a fragment thereof comprising at least a ring domain, or the amino acid sequence of any one of SEQ ID NOs: 13-29, 100-116, 119-125, 127-233, or 1001-1155.

97. The composition of any one of claims 83-96, wherein V and/or W of said second polypeptide are absent.

98. The composition of any one of claims 83-97, wherein V or W of said second polypeptide comprises a fragment crystallizable region (Fc).

99. The composition of any one of claims 83-98, wherein V or W of said second polypeptide comprises a glycine-rich region.

100. The composition of any one of claims 83-99, wherein V or W of said second polypeptide comprises a bone-targeting moiety.

101. The composition of any one of claims 83-100, wherein V or W of said second polypeptide comprises a cathepsin cleavage sequence.

102. The composition of any one of claims 83-101, wherein said second polypeptide comprises the structure V-NP-W,

NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), each of V and W is, independently, absent or is an amino acid sequence of at least one amino acid, and

said NP comprises the amino acid sequence of any one of SEQ ID NOs: 17-29, 31-40, 42-94, 101-116, 119-122, 128-161, or 163-233, or V or W comprises the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

103. The composition of any one of claims 83-102, wherein said second polypeptide comprises the structure V-NP or NP-W,

NP is said natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and each of V and W comprises, independently, the amino acid sequence of any one of SEQ ID NOs: 304-313, 322-333, or 337-391.

104. The composition of any one of claims 83-103, wherein said second polypeptide comprises the structure X-Fc-Y-NP-Z or the structure X-NP-Y-Fc-Z,

NP is said natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), and each of X, Y, and Z is, independently, absent or is an amino acid sequence of at least one amino acid.

105. The composition of claim 104, wherein Y comprises a glycine-rich region.
106. The composition of claim 104 or 105, wherein X is absent, Z is absent, or X and Z are both absent.
107. The composition of any one of claims 104-106, wherein X, Y, or Z comprises a bone-targeting moiety.
108. The composition of any one of claims 104-107, wherein X, Y, or Z comprises a cathepsin cleavage sequence.
109. The composition of any one of claims 83-108, wherein the amino acid sequence of said second polypeptide comprises the amino acid sequence of any one of SEQ ID NOs: 501-608.
110. The composition of claim 109, wherein the amino acid sequence of said second polypeptide comprises the amino acid sequence of any one of SEQ ID NOs: 502, 504, 506, 512, 514, 516, 530, 560, 562, 564, 572, 574, 576, 584, 586, 588, 596, 598, 600, or 608.
111. The composition of claim 110, wherein the amino acid sequence of said second polypeptide comprises or consists of the amino acid sequence of SEQ ID NO: 512.
112. The composition of any one of claims 83-111, wherein said first polypeptide and/or said second polypeptide are in dimeric form.
113. The composition of any one of claims 83-112, wherein said first polypeptide and/or said second polypeptide are glycosylated or pegylated.
114. The composition of any one of claims 83-113, wherein the composition is a pharmaceutical composition comprising a pharmaceutically acceptable excipient.
115. The composition of any one of claims 83-114, wherein said composition is lyophilized.

116. The composition of any one of claims 83-115, wherein said first polypeptide is present in a dosage between about 0.2 mg/kg to about 20 mg/kg and said second polypeptide is present in a dosage between about 0.5 mg/kg to about 500 mg/kg.

117. The composition of any one of claims 83-116, wherein the amino acid sequence of said first polypeptide comprises the amino acid sequence of SEQ ID NO: 1204 and wherein the amino acid sequence of said second polypeptide comprises the amino acid sequence of SEQ ID NOs: 504, 512, 530, or 572.

118. A method of treating a disease or a condition in a subject, said method comprising administering to said subject a therapeutically effective amount of a first polypeptide and a second polypeptide, wherein

a) said first polypeptide comprises the structure A-sALP-B, wherein

- i) sALP is the extracellular domain of an alkaline phosphatase,
- ii) A is absent or is an amino acid sequence of at least one amino acid, and
- iii) B is absent or is an amino acid sequence of at least one amino acid; and

b) said second polypeptide comprises the structure V-NP-W, wherein

- i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),
- ii) V is absent or is an amino acid sequence of at least one amino acid, and
- iii) W is absent or is an amino acid sequence of at least one amino acid; and

said disease or condition is selected from the group consisting of a neurocutaneous syndrome, a disorder associated with overactivation of FGFR3, a bone or cartilage disorder, a vascular smooth muscle disorder, and a condition for elongation of bone,

thereby treating said disease or said condition in said subject.

119. The method of claim 118, wherein said first polypeptide and said second polypeptide are administered within ten days, five days, or twenty-four hours of each other.

120. The method of claim 118, wherein said first polypeptide and said second polypeptide are administered simultaneously.

121. The method of any one of claims 118-120, wherein said first polypeptide and said second polypeptide are formulated together in a composition or each separately in a composition.

122. The method of claim 121, wherein said composition is a pharmaceutical composition comprising a pharmaceutically acceptable excipient.

123. The method of claim 122, wherein said composition is lyophilized.

124. The method of any one of claims 121-123, wherein said composition is the composition of any one of claims 83-117.

125. The method of any one of claims 118-124, wherein the amino acid sequence of said first polypeptide comprises the amino acid sequence of SEQ ID NO: 1204 and wherein the amino acid sequence of said second polypeptide comprises the amino acid sequence of SEQ ID NOs: 504, 512, 530, or 572.

126. The method of any one of claims 118-125, wherein said first polypeptide is present in a dosage between about 0.2 mg/kg to about 20 mg/kg and said second polypeptide is present in a dosage between about 0.5 mg/kg to about 500 mg/kg.

127. The method of any one of claims 118-125, wherein said first polypeptide is present in a dosage between about 0.2 mg/kg to about 20 mg/kg and said second polypeptide is present in a dosage between about 10  $\mu$ g/kg to about 1,000  $\mu$ g/kg.

128. The method of any one of claims 118-127, wherein said first polypeptide and said second polypeptide are administered subcutaneously.

129. The method of any one of claims 118-128, wherein said first polypeptide and said second polypeptide are administered one time, two times, or three times per week.

130. The method of any one of claims 118-129, wherein said subject is human.

131. The method of any one of claims 118-130, wherein said disease is said neurocutaneous syndrome.

132. The method of claim 131, wherein said neurocutaneous syndrome is neurofibromatosis type I.

133. A kit comprising:

- a) a first polypeptide comprising the structure A-sALP-B, wherein
  - i) sALP is the extracellular domain of an alkaline phosphatase,
  - ii) A is absent or is an amino acid sequence of at least one amino acid, and
  - iii) B is absent or is an amino acid sequence of at least one amino acid; and
- b) instructions for administering said first polypeptide to a patient diagnosed with or at risk of developing a neurocutaneous syndrome.

134. The kit of claim 133, further comprising:

- (c) a second polypeptide comprising the structure V-NP-W, wherein
  - i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),
  - ii) V is absent or is an amino acid sequence of at least one amino acid, and
  - iii) W is absent or is an amino acid sequence of at least one amino acid.

135. A kit comprising:

- a) a polypeptide comprising the structure V-NP-W, wherein
  - i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),
  - ii) V is absent or is an amino acid sequence of at least one amino acid, and
  - iii) W is absent or is an amino acid sequence of at least one amino acid; and
- b) instructions for administering said polypeptide to a patient diagnosed with or at risk of developing a neurocutaneous syndrome.

136. A kit comprising:

- a) a first polypeptide comprising the structure A-sALP-B, wherein
  - i) sALP is the extracellular domain of an alkaline phosphatase,
  - ii) A is absent or is an amino acid sequence of at least one amino acid, and
  - iii) B is absent or is an amino acid sequence of at least one amino acid; and
- b) a second polypeptide comprising the structure V-NP-W, wherein
  - i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),
  - ii) V is absent or is an amino acid sequence of at least one amino acid, and
  - iii) W is absent or is an amino acid sequence of at least one amino acid.

137. The kit of claim 136, wherein said first polypeptide and said second polypeptide are formulated together.

138. The kit of claim 136, wherein said first polypeptide and said second polypeptide are formulated separately and in individual dosage amount.

FIG. 1A

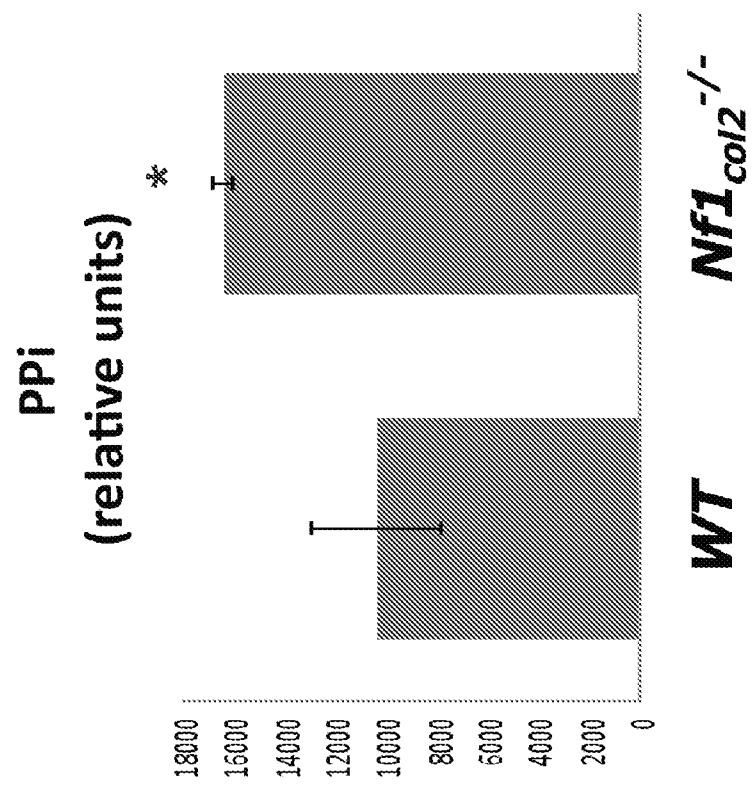
***Nf1<sub>col2</sub>*<sup>-/-</sup> Osteoblasts Secrete Increased Levels of Pyrophosphate (PP<sub>i</sub>)**

FIG. 1B

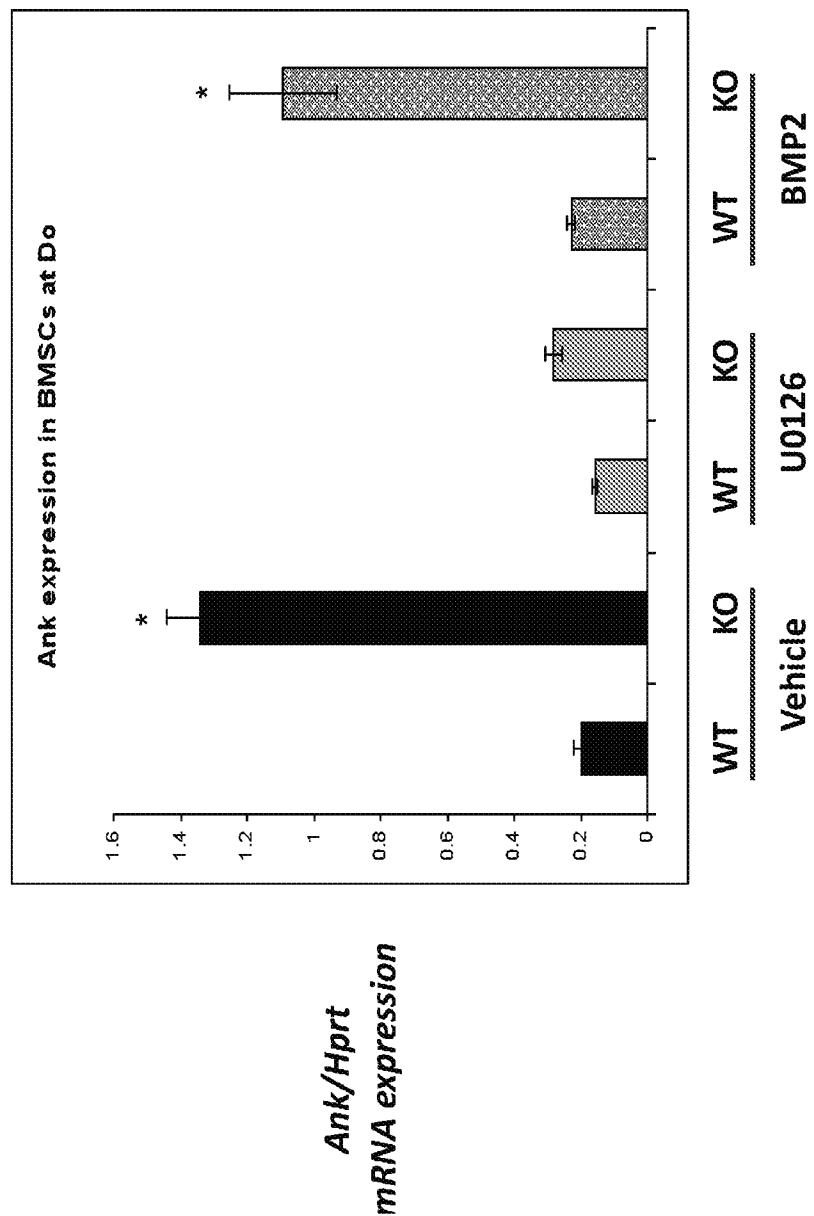
Increased *Ank* Expression in  $Nf1_{col2}^{-/-}$  Osteoblasts

FIG. 2

### Hypothetical Working Model for Defective Bone Matrix Mineralization in $\text{NF1}_{\text{col2}}^{-/-}$ Mice

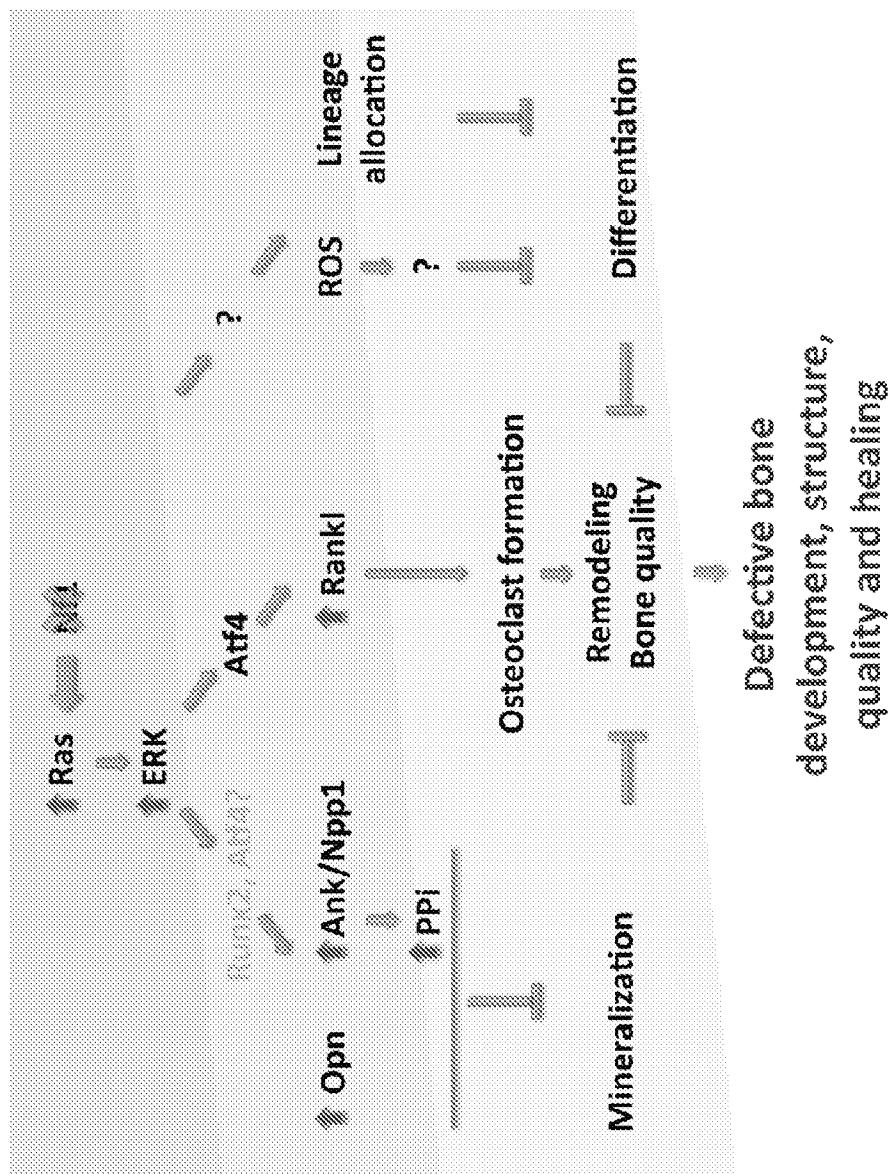


FIG. 3A

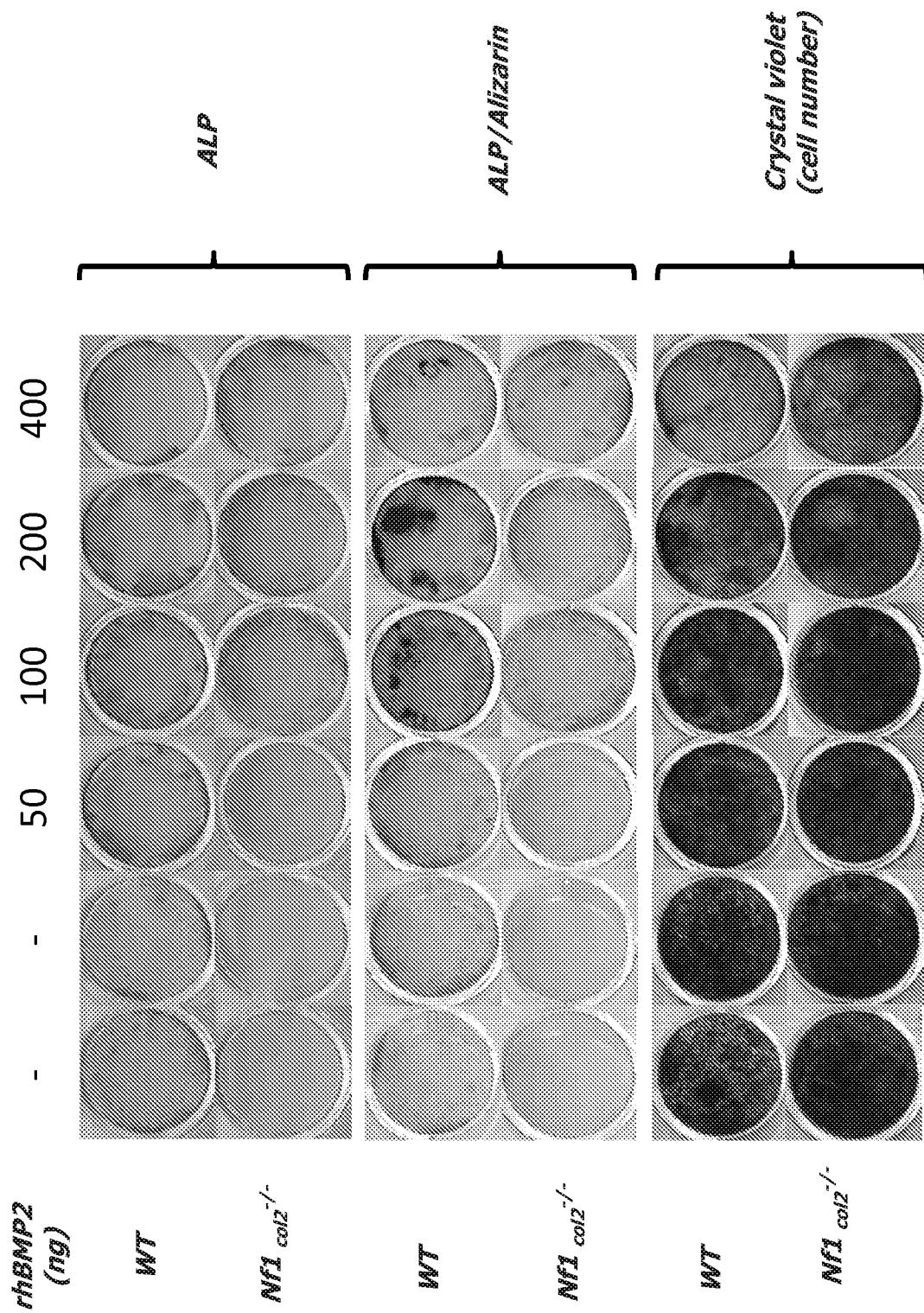
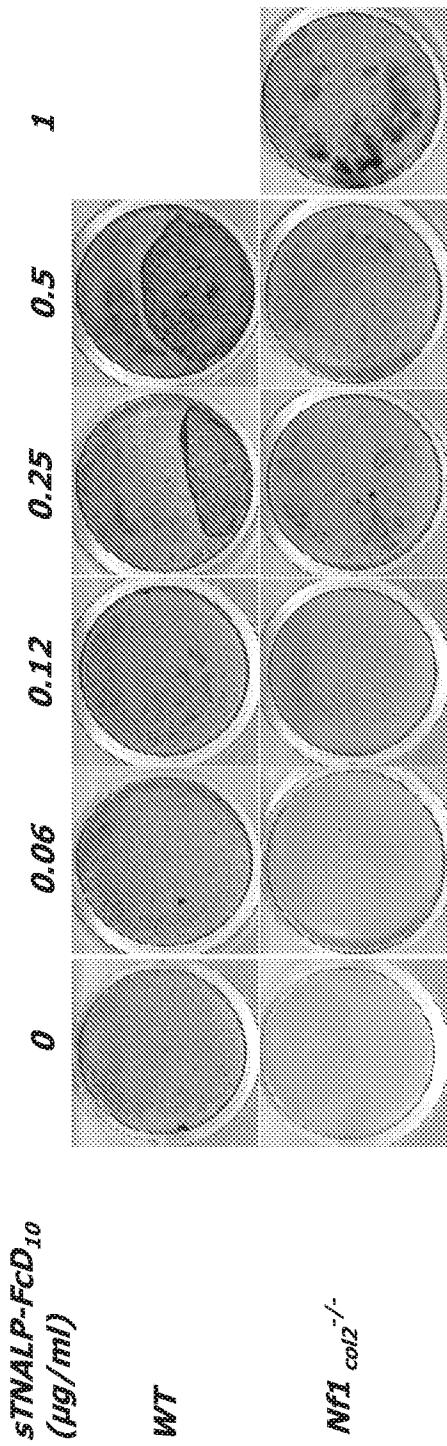
Effect of BMP2 on  $NF1_{col2}^{-/-}$  Osteoblasts

FIG. 3B

**Effect of sTNALP-FcD<sub>10</sub> on NF1<sub>col2</sub><sup>-/-</sup> Bone Marrow Stromal Cells**



**Alizarin Red S Quantification**

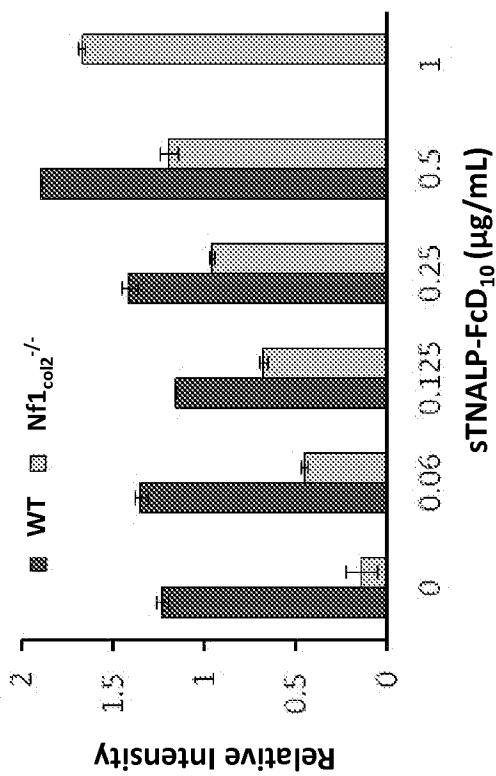


FIG. 3C

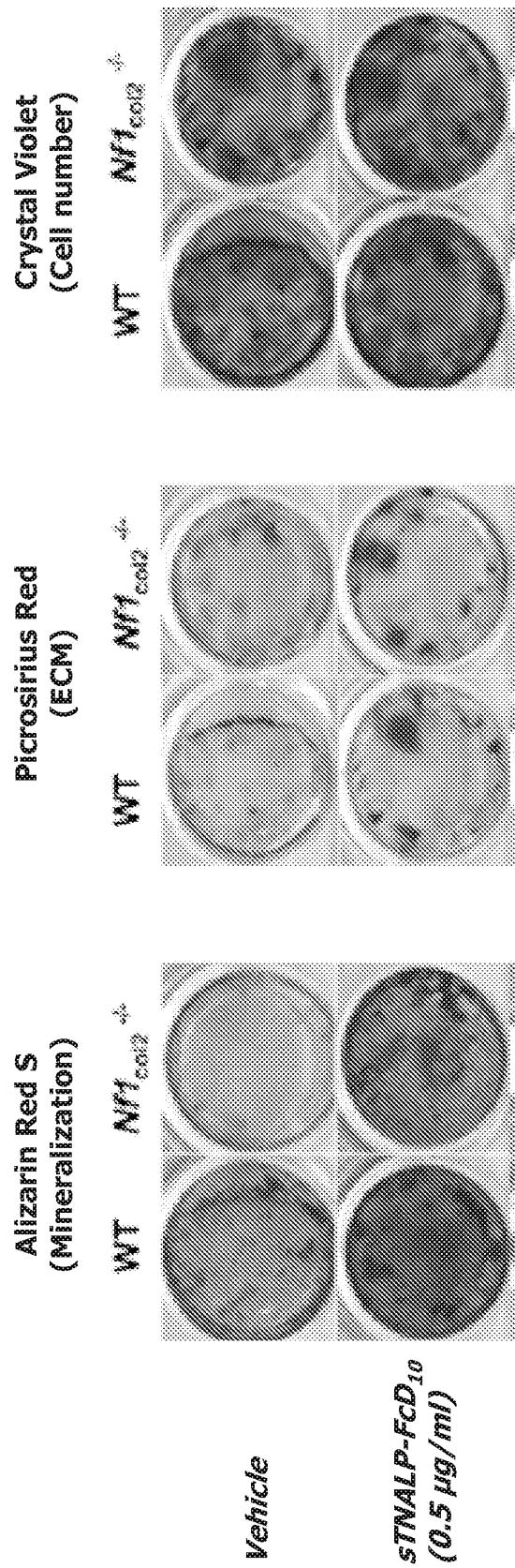
Effect of sTNALP-FcD<sub>10</sub> on NF1<sub>col2</sub><sup>-/-</sup> Bone Marrow Stromal Cells

FIG. 3D

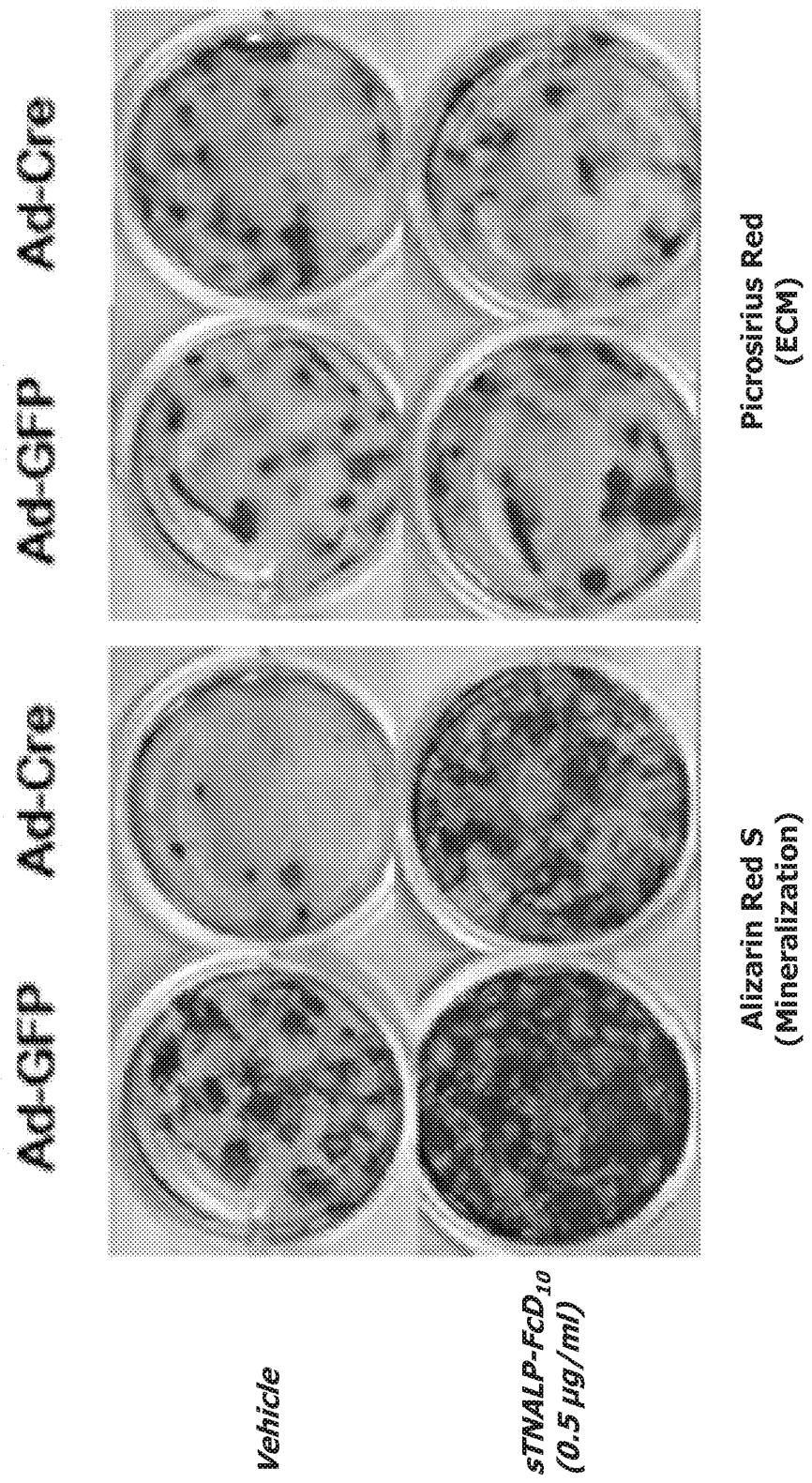
Effect of sTNALP-FcD<sub>10</sub> on Floxed NF1<sub>col2</sub><sup>-/-</sup> Bone Marrow Stromal Cells

FIG. 3E

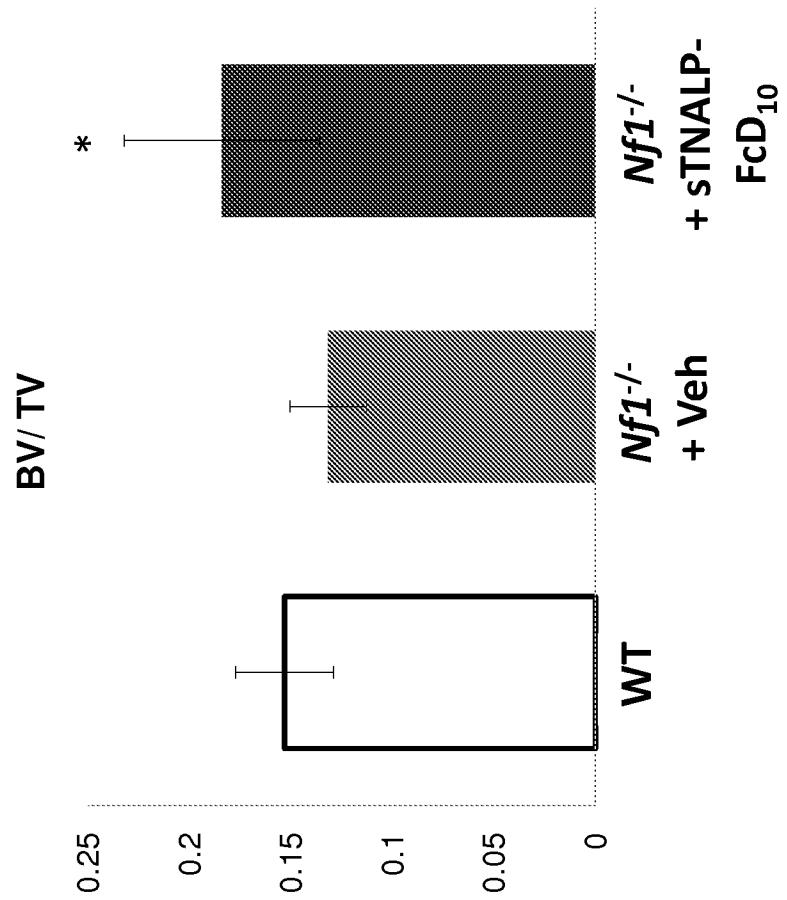
Effect of sTNALP-FcD<sub>10</sub> on Bone Volume in Nf1<sub>col2</sub><sup>-/-</sup> Mice

FIG. 4A

## Expression of *NPR-B* Gene in NF1<sub>col2</sub><sup>-/-</sup> Mice

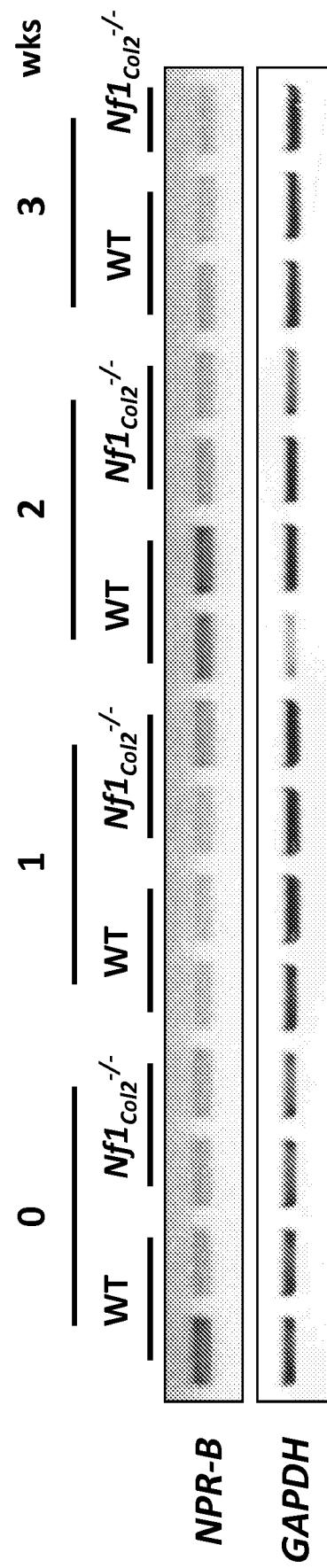


FIG. 4B

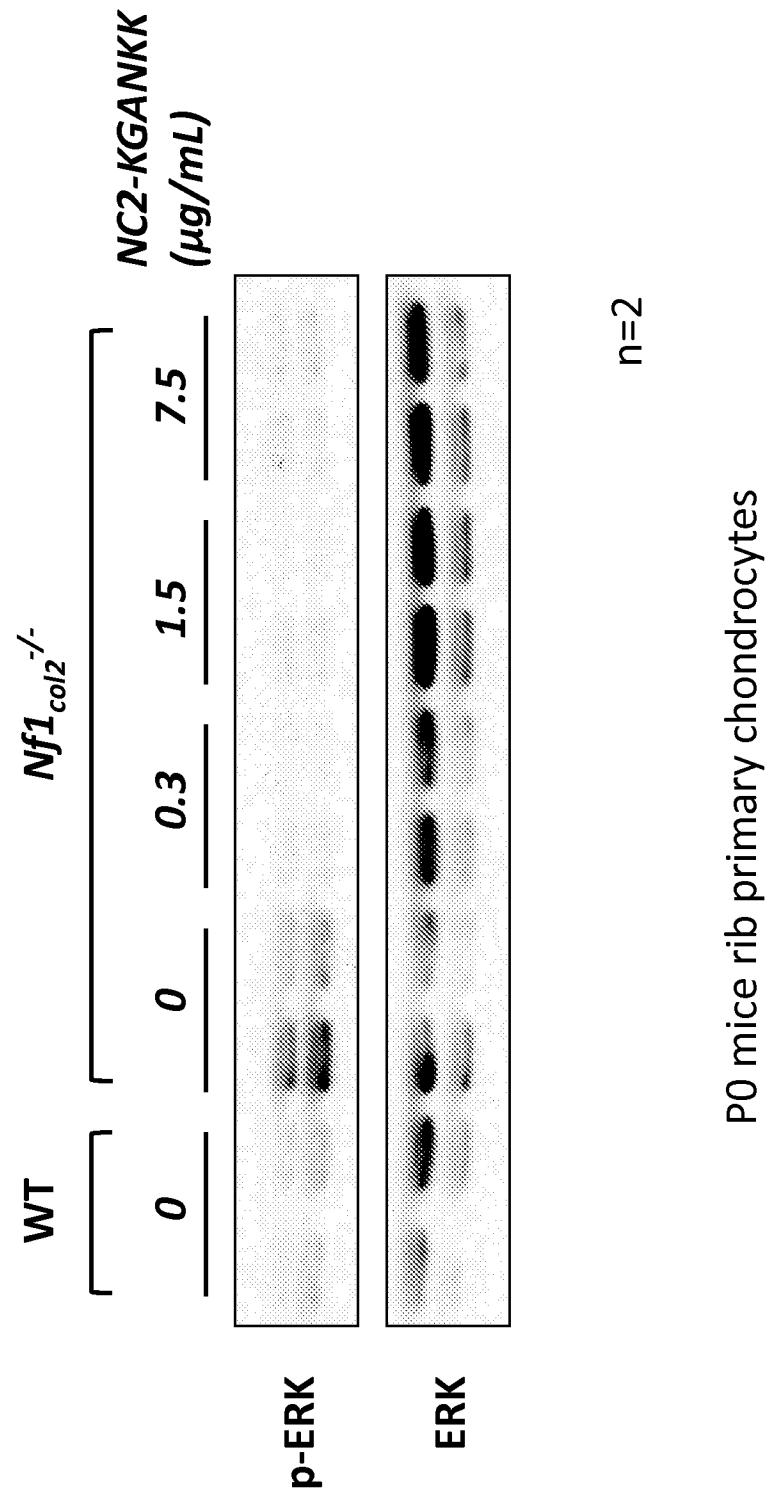
Effect of NC2-KGANKK on NF1<sub>col2</sub><sup>-/-</sup> Chondrocytes

FIG. 4C

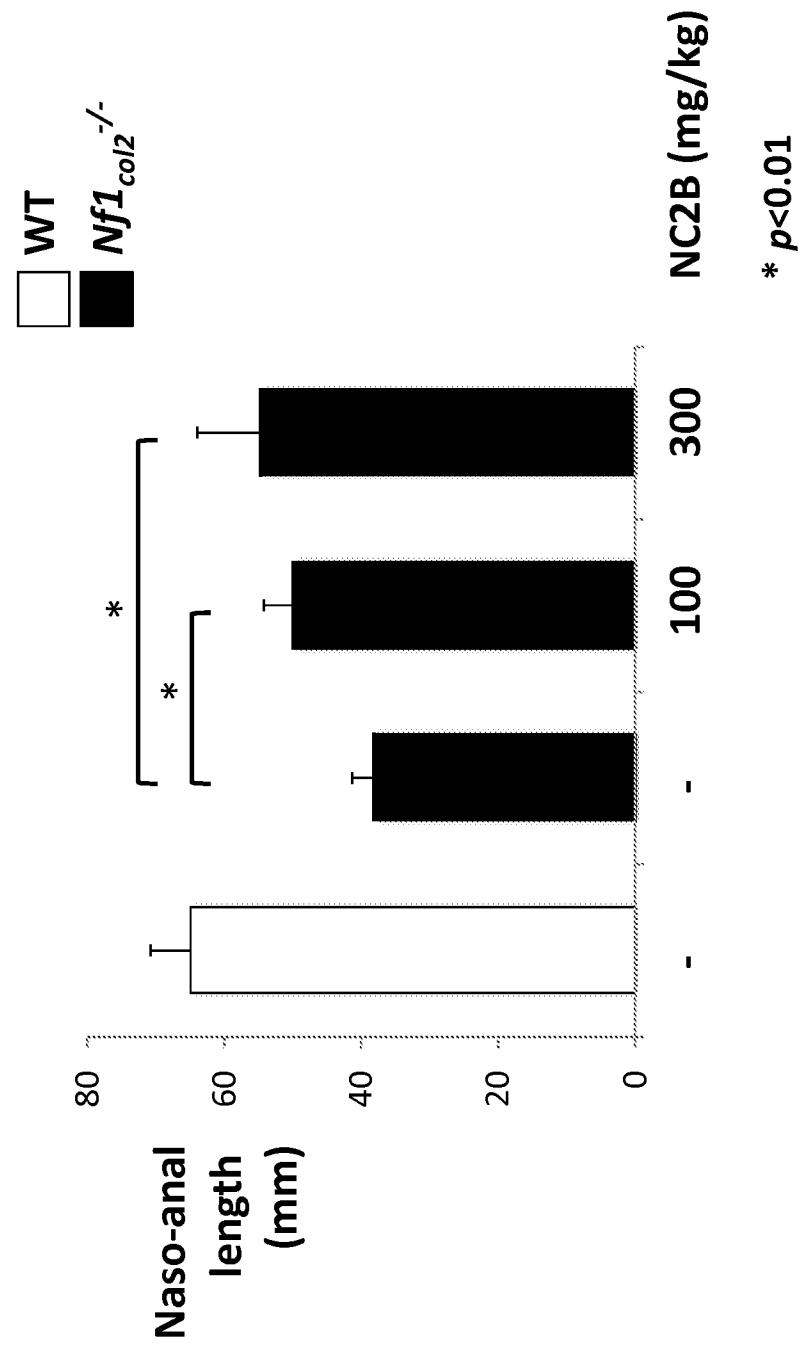
Effect of NC2B on Naso-anal Length in NF1<sub>col2</sub><sup>-/-</sup> Mice

FIG. 4D

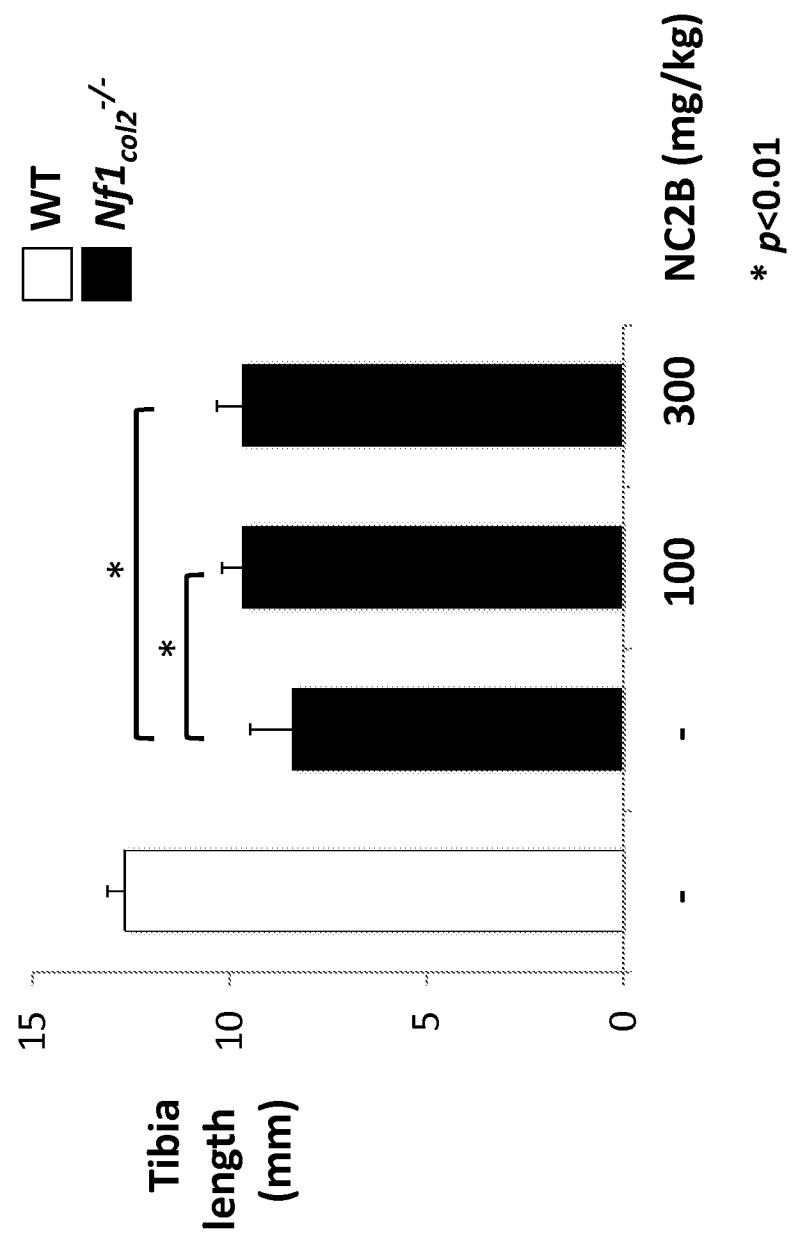
Effect of NC2B on Tibia Length in  $Nf1_{col2}^{-/-}$  Mice

FIG. 4E

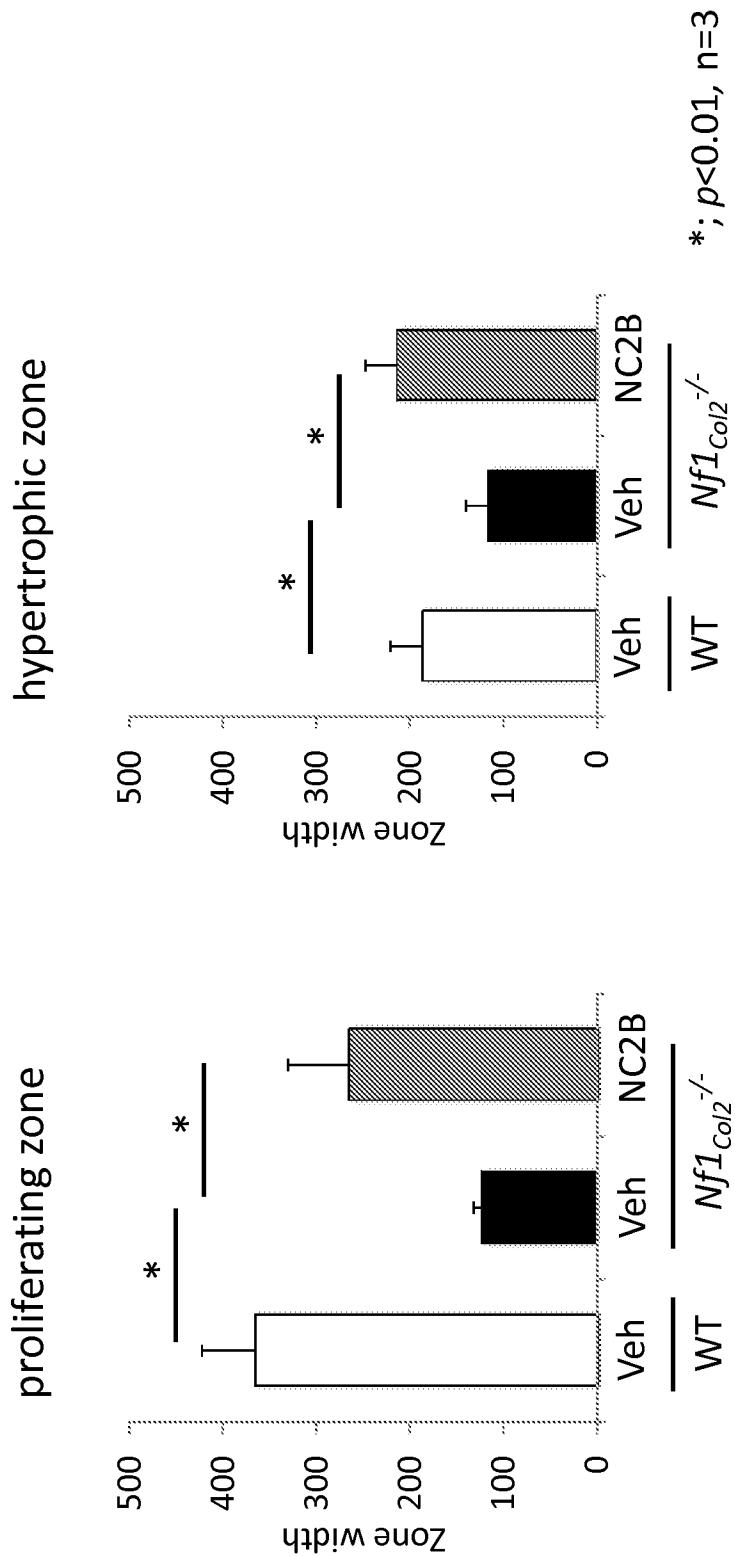
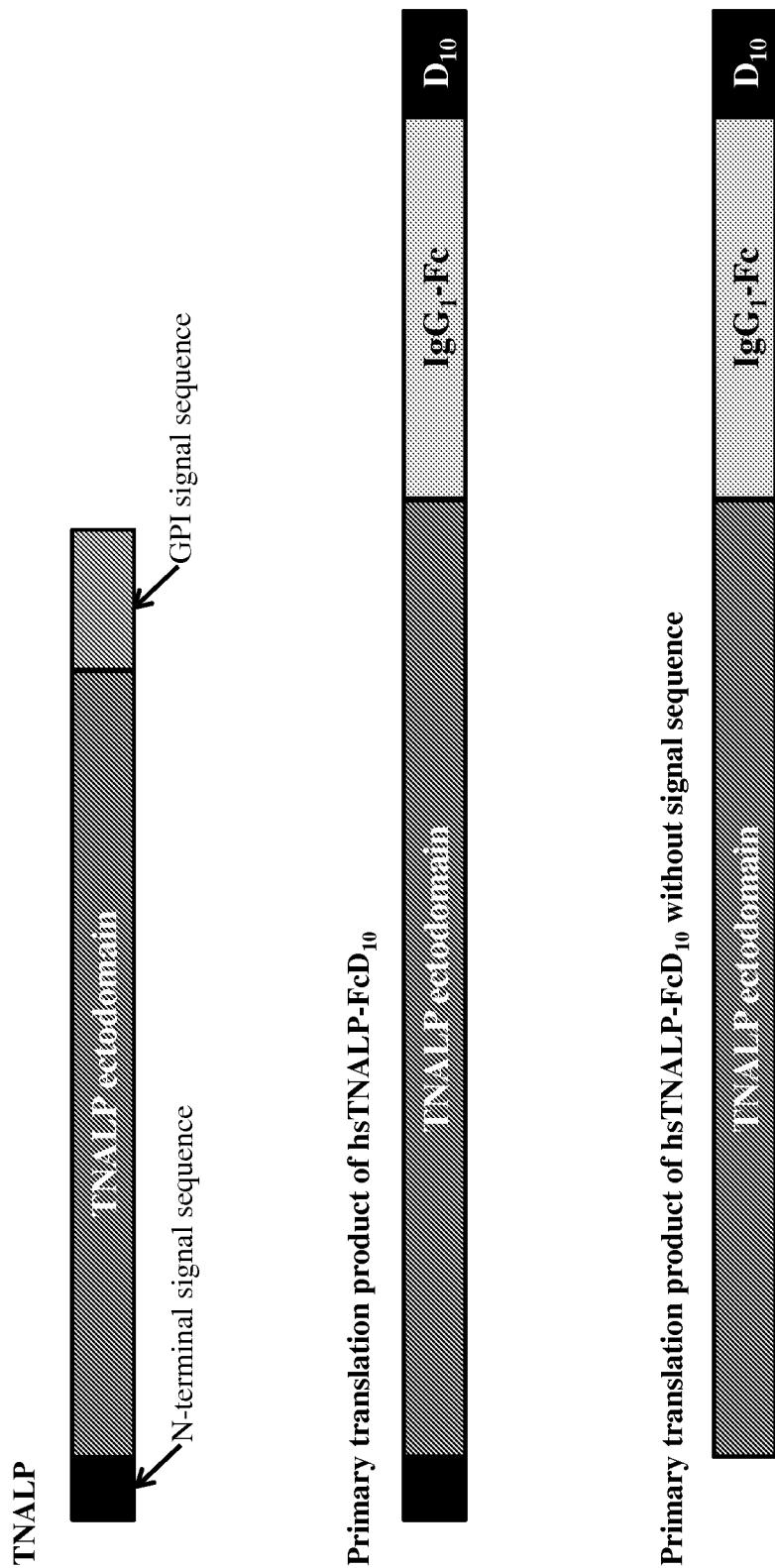
**Effect of NC2B on Growth Plates in  $Nf1_{Col2}^{-/-}$  Mice**

FIG. 5A

**Schematic Structure of TNALP and hsTNALP-FcD<sub>10</sub>  
(With and Without Signal Sequence)**



## FIG. 5B

## hsTNALP-FcD<sub>10</sub> Protein Sequence (With and Without Signal Sequence)

STNALP-FcD<sub>10</sub>  
(w/ sig. seq.)

MVSPFELVLAIGTCLNSSLVPEKEKDPKYWRDQAETILKYALELQQLNNTNVAKNVIMFLGDMGVSTVTAARILRGQLHHNPGEETRL  
RILKGQLHHNPGEETRLMDKFPFVALSKTYNTNAQVFDSAGTATAYLCGVKANEGTGVSATERSRRCN  
TTQGNEVTSILRWAKDAGKSVGIVTTTRVNHATPSAAYAHSADRDWYSDNEMPPEALSQGCKDIAYQLMH  
NIRDIDVIMGGGRKYMYPKRNKTDVEYEESDEKARGTRLDGLDLVDTWKSFKPRYKHSHETWNRTELLTLDP  
HNVDYLLLGLFEPGDMQYELNRNNVTDPSLSEMVVVAIQIILRKNPKGFFLLVEGGRIDHGHEGKAKQALH  
EAEMDRAIGQAGSLTSSEDTLTVVTADHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTALLYNGPG  
YKVVGGERENVSMVDYAHNNYQAQSAVPLRHETHGGEDDAVFTSKGPMAHLLHGVHEQNYVPHVMAYAACI  
GANLGHCAPASS**LK**DKTTHTCPPCPPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFN  
WYVDGVEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREP  
QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKLTVDKSR  
WQQGNVVESCSVMHEALHNHYTQKSILSSPGK**D**IDDDDDDDDD (SEQ ID NO: 1201)

STNALP-FcD<sub>10</sub>  
(w/o sig. seq.)

LVPEKEKDPKYWRDQAETILKYALELQQLNNTNVAKNVIMFLGDMGVSTVTAARILRGQLHHNPGEETRL  
EMDKFPFVALSKTYNTNAQVFDSAGTATAYLCGVKANEGTGVSATERSRCNTTQGNEVTSILRWAKDA  
GKSVGIVTTTRVNHATPSAAYAHSADRDWYSDNEMPPEALSQGCKDIAYQLMHNTDIDVIMGGGRKYM  
PKNKTDVEYEESDEKARGTRLDGLDLVDTWKSFKPRYKHSHETWNRTELLLDPHNVDYLLGFEPGDMQY  
ELNRRNNVTDPSLSEMVVVAIQIILRKNPKGFFLLVEGGRIDHGHEGKAKQALHEAVEMDRAIGQAGSLTS  
SEDTLTVVTADHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTALLYNGPGYKVVGGERENVSMVDY  
HNNYQAQSAVPLRHETHGGEDDAVFTSKGPMAHLLHGVHEQNYVPHVMAYAACI**GANLGHCAPASS****LK**DKT  
HTCPPCPPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGEVHNAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQV  
SLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVPHVMHEALH  
NHYTQKSILSSPGK**D**IDDDDDDDDD (SEQ ID NO: 1204)

## FIG. 5C

## hsTNALP-Fc Protein Sequence (With and Without Signal Sequence)

STNALP-Fc  
(w/ sig. seq.)

MVSPFELVLAIGTCLTNSLVP**E**KDPKYWRDQAETLKYALELQQLNNTNVAKNVIMFLGDMGVSTVTAARILKGQLHHNPGEETRL  
RILKGQLHHNPGEETRLMDKFPFVALSKTYNTNAQVPDSAGTATAYLGGVKANEGETVGVSAATERSRCN  
TTQGNEVTSILRWAKDAGKSVGIVTTTRVNHATPSAAyahSADRDWYSDNEMPPEALSQGCKDIAYQLMH  
NIRDIDVIMGGGRKYMYPKRNKTDVEYESDEKARGTRLDGLDLVDTWKSEKPRYKHSHEIWNRTELLTLDP  
HNVDYLIGLFFEPGDMQYELNRRNNVTDPSSLSEMVVVAIQILRKNPKGFFLVEGGRIDHGHHEGKAKQALH  
EAVEMDRAIGQAGSLTSSSEDTLTVVTAHDHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTAILYGNPG  
YKVVGGERENVSMDYAHNNYQAQSAPVLRHETHGGEDDVAVFSKGPMMAHLLHGVAHEQNYVPHVMAYAACI  
GANLGHCAPASS**LK**DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFN  
WYVDGVEVHNNAKTKPREEQYNSTYRVVSVLTLHQDWLNGKEYKCKVSNKALPAPIEKTIISKAKGQPREP  
QVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKLTVDKSR  
WQQGNVESCSCVVMHEALHNHHTQKSLSLSPGK (SEQ ID NO: 1220)

STNALP-Fc  
(w/o sig. seq.)

LVPEKEKDPKYWRDQAETLKYALELQQLNNTNVAKNVIMFLGDMGVSTVTAARILKGQLHHNPGEETRL  
EMDKFPFVALSKTYNTNAQVPDSAGTATAYLGGVKANEGETVGVSAATERSRCNTIQTGNEVTSILRWAKDA  
GKSVGIVTTTRVNHATPSAAyahSADRDWYSDNEMPPEALSQGCKDIAYQLMHNIRDIDVIMGGGRKYM  
PKNKTIDVEYESDEKARGTRLDGLDLVDTWKSEKPRYKHSHEIWNRTELLTLDPHNVDYLLFEPGDMQY  
ELNRRNNVTDPSSLSEMVVVAIQILRKNPKGFFLVEGGRIDHGHHEGKAKQALHEAVEMDRAIGQAGSLTS  
SEDTLTVVTAHDHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTAILYGNPGYKVVGGERENVSMDYAH  
HNNYQAQSAPVLRHETHGGEDDVAVFSKGPMMAHLLHGVAHEQNYVPHVMAYACTIGANLGHCAPASS**LK**DKT  
HTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREF  
EQYNSTYRVVSVLTLHQDWLNGKEYKCKVSNKALPAPIEKTIISKAKGQPREPQVYTLPPSREEMTKNQV  
SLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALH  
NHYTQKSLSLSPGK (SEQ ID NO: 1221)

FIG. 5D

## hsTNALP-FcD<sub>10</sub> D<sub>NA</sub> Sequence

## FIG. 6

**Human Soluble Tissue Nonspecific Alkaline Phosphatase  
(hsTNALP) Protein Sequence**

hsTNALP (1-502)	<u>MVSPFELVIALIGTCLTNSLVEPEKEKDPKYWRDQAQETLKYALELQKLNTNVAKNVI</u> <u>LKQQLHHNPGEETRLEMDKFPFVALSKTYNTNAQVPDFSAGTATAYLCGVKANE</u> <u>GTGVSAATERSRCNTTQGNEVTSILRWA</u> <u>KDAGKSVGIVTTIRVNHATPSAAAYAHSADRDIYQLMHNIRDID</u> <u>NEVTSILRWA</u> <u>KDAGKSVGIVTTIRVNHATPSAAAYAHSADRDIYQLMHNIRDID</u> <u>VIMGGGRKYMVKNKT</u> <u>DVEYESDEKARGTRLGDLVDTWKSFKPRYKHSF</u> <u>IWNRTTELLTLDPHNVDYLLG</u> <u>LFEPGDMQYELNRRNNVTDP</u> <u>SLSEMVVVAIQILRKNPKGFFLVEGGRIDHGHHEGKAKQALHEAVEMDRAIG</u> <u>QAGSLTSSEDTITVVTADHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTAILY</u> <u>GNGPGYKVVGGERENVSMVHDYAHNNYQAQS</u> <u>AVPLMVDYAHNNYQAQS</u> <u>AVPLRHETHGGEDVAVFSKGPMMAHLLHG</u> <u>VHEQNYVPHVMAACIGANLGH</u> <u>CAPASS (SEQ ID NO: 1202)</u>	hsTNALP (18-502)	<u>LYPEKEKDPKYWRDQAQETLKYALELQKLNTNVAKNVI</u> <u>MVSPFELVIALIGTCLTNSLVEPEKEKDPKYWRDQAQETLKYALELQKLNTNVAKNVI</u> <u>IMFLGDGMGVSTVTAARI</u> <u>LKGQLHHNPGEETRLEMDKFPFVALSKTYNTNAQVPDFSAGTATAYLCGVKANE</u> <u>GTGVSAATERSRCNTTQGNEVTSILRWA</u> <u>KDAGKSV</u> <u>DKFPFVALSKTYNTNAQVPDFSAGTATAYLCGVKANE</u> <u>GTGVSAATERSRCNTTQGNEVTSILRWA</u> <u>KDAGKSV</u> <u>GIVTTIRVNHATPSAAAYAHSADRDIYQLMHNIRDIDVIMGGGRKYMVKNKT</u> <u>DYESDEKARGTRLGDLVDTWKSFKPRYKHSF</u> <u>IWNRTTELLTLDPHNVDYLLG</u> <u>FEPGDMQYELNRRNNVT</u> <u>DPSLSEMVVVAIQILRKNPKGFFLVEGGRIDHGHHEGKAKQALHEAVEMDRAIG</u> <u>QAGSLTSSEDTITVVTADHSHVFTFGGYTPRGNSIFGLAPMLSDTDKKPFTAILY</u> <u>GNGPGYKVVGGERENVSMVHDYAHNNYQAQS</u> <u>AVPLMVDYAHNNYQAQS</u> <u>AVPLRHETHGGEDVAVFSKGPMMAHLLHG</u> <u>VHEQNYVPHVMAACIGANLGH</u> <u>CAPASS (SEQ ID NO: 1205)</u>
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**FIG. 7****Exemplary Fc Sequence (IgG-1)**

DKTHTCPPCPAPELLGGPSVFLFPPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEV  
HNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQV  
YTLPPSREEMTKNQVSLTCIUVKGFYPSDIAVEWESENQOPENNYKTTPPVLDSDGSFFLYSKLTIV  
DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 401)

FIG. 8

## Multiple Sequence Alignment of Tissue Nonspecific Alkaline Phosphatase (TNALP)

CLUSTAL W (1.82) multiple sequence alignment: Tissue Nonspecific Alkaline Phosphatase

<p>1</p> <pre> P09487  PPBT_BOVIN Q29486  PPBT_FELCA P05186  PPBT_HUMAN P09242  PPBT_MOUSE P08289  PPBT_RAT Q9NOVO  Q9NOVO_CANFA Consensus </pre>	<p>61</p> <pre> GMGVSTVTAAG RILKGQLHHN PGEETKLEM KEPYVVALSKT YNTNAQVPDS AGTATAYLCG GMGVSTVTAAG RILKGQLHHN PGEETRLEM KEPYVVALSKT YNTNAQVPDS AGTATAYLCG </pre>	<p>121</p> <pre> VKANEGTVGV SAATQRQCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSASYAH VKANEGTVGV SAATORTCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH VKANEGTVGV SAATERSRN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH VKANEGTVGV SAATERTRCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH VKANEGTVGV SAATORTHCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH VKANEGTVGV SAATQRTHCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH VKANEGTVGV SAATRXRCN TTQGNEVTSI LRWAKDAGKS VGIYTTTRVN EATPSAAYAH </pre>
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FIG. 8 (cont'd)

P09487   PPBT_BOVIN	SADRDWYSDN	EMPEALSQG	CKDIAYQLMY	NIKDIEVIMG	GERKYMFPKN	RTDVEYELDE
Q29486   PPBT_FELCA	SADRDWYSDN	EMPEALSQG	CKDIAYQLMH	NVRDIEVIMG	GERKYMFPKN	RTDVEYEMDE
P05186   PPBT_HUMAN	SADRDWYSDN	EMPEALSQG	CKDIAYQLMH	NIRDIVIMG	GERKYMFPKN	KIDVEYESDE
P09242   PPBT_MOUSE	SADRDWYSDN	EMPEALSQG	CKDIAYQLMH	NTKDIDVIMG	GERKYMFPKN	RTDVEYELDE
P08289   PPBT_RAT	SADRDWYSDN	EMPEALSQG	CKDIAYQLMH	NIKDIDVIMG	GERKYMFPKN	RTDVEYELDE
Q9NOVO   Q9NOVO_CANFA	SADRDWYSDN	EMPEALSQG	CKDIAYQLMH	NVKDIEVIMG	GERKYMFPKN	RTDVEYEMDE
Consensus	SADRDWYSDN	EMPEALSQG	CKDIAYQLMX	NXXDIXVIMG	GGRKYMXPKN	XTDVEYEXDE
P09487   PPBT_BOVIN	KARGTRLDGL	NLIDIWKSFK	PKHKHSHYVW	NRTDLLALDP	HSVYDYLGLF	EPGDMQYELN
Q29486   PPBT_FELCA	KARGTRLDGL	NLVDIWKSFK	PRHKHSHYVW	NRTELLTLDP	YGVYDYLGLF	EPGDMQYELN
P05186   PPBT_HUMAN	KARGTRLDGL	DLVDIWKSFK	PRYKHSHEFW	NRTELLFLDP	HNVYDYLGLF	EPGDMQYELN
P09242   PPBT_MOUSE	KARGTRLDGL	DLISIWKSFK	PRHKHSHYVW	NRTELLALDP	SRYVYDYLGLF	EPGDMQYELN
P08289   PPBT_RAT	KARGTRLDGL	DLISIWKSFK	PRHKHSHYVW	NRTELLALDP	SRYVYDYLGLF	EPGDMQYELN
Q9NOVO   Q9NOVO_CANFA	KSTGARLDGL	NLIDIWKNFK	PRHKHSHYVW	NRTELLALDP	YIVYDYLGLF	DPGDMQYELN
Consensus	XXXGXRLDGL	XLIXXXWXXXFK	XXXKHSHXXXW	NRTXLXXLDP	XXVDYLGLF	XPGDMQYELN
P09487   PPBT_BOVIN	KNNATDPSLS	EMVEMAIRIL	NKNPKGEFL	HEGKAKQALH	EAEMDQAIG	
Q29486   PPBT_FELCA	RNSTTDPSSL	EMVEIAIKIL	SKNPKGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDQAIG
P05186   PPBT_HUMAN	RNNVTDPSLS	EMVVAIAQIL	RKNPKGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDRAIG
P09242   PPBT_MOUSE	RNNLTDPSLS	EMVEVALRIL	TKNLKSFLL	VEGGRIDHGH	HEGKAKQALH	EAEMDQAIG
P08289   PPBT_RAT	RNNLTDPSLS	EMVEVALRIL	TKNPKGEFL	VEGGRIDHGH	HEGKAKOALH	EAEMDEAIG
Q9NOVO   Q9NOVO_CANFA	RNNVTDPSLS	EMVEIAIKIL	SKKPRGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDRAIG
Consensus	***	***	***	***	***	***
P09487   PPBT_BOVIN	RNNATDPSLS	EMVEMAIRIL	NKNPKGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDQAIG
Q29486   PPBT_FELCA	RNSTTDPSSL	EMVEIAIKIL	SKNPKGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDQAIG
P05186   PPBT_HUMAN	RNNVTDPSLS	EMVVAIAQIL	RKNPKGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDRAIG
P09242   PPBT_MOUSE	RNNLTDPSLS	EMVEVALRIL	TKNLKSFLL	VEGGRIDHGH	HEGKAKQALH	EAEMDQAIG
P08289   PPBT_RAT	RNNLTDPSLS	EMVEVALRIL	TKNPKGEFL	VEGGRIDHGH	HEGKAKOALH	EAEMDEAIG
Q9NOVO   Q9NOVO_CANFA	RNNVTDPSLS	EMVEIAIKIL	SKKPRGEFL	VEGGRIDHGH	HEGKAKQALH	EAEMDRAIG
Consensus	***	***	***	***	***	***

FIG. 8 (cont'd)

FIG. 9

## Multiple Sequence Alignment of Alkaline Phosphatase (TNALP, GALP, PLALP, and IALP)

CLUSTAL 2.0.5 multiple sequence alignment

TNALP <sub>Prn</sub>	MILP-----F	LVLAITGCLT	NSFVPEKEKD	PSYWQRQQAQE	TLKNALKLQK	LNTNVAKNII	55
TNALP <sub>Mm</sub>	MISP-----F	LVLAITGCLT	NSFVPEKEKD	PSYWQRQQAQE	TLKNALKLQK	LNTNVAKNVI	55
TNALP <sub>Phs</sub>	MISP-----F	LVLAITGCLT	NSLVPPEKEKD	PKYWRDQAQE	TLKYALELQK	LNTNVAKNVI	55
TNALP <sub>Cf</sub>	-----	-----	-----	-----	-----	-----	33
TNALP <sub>FC</sub>	MISP-----F	LVLAITGCLT	NSLVPPEKEKD	PKYWRDQAQQ	TLKYALRLQK	LNTNVAKNVI	55
TNALP <sub>Bt</sub>	MISP-----F	LLLAIGTCFA	SSLVPPEKEKD	PKYWRDQAQQ	TLKNALRLQK	LNTNVAKNVI	55
GALP <sub>hs</sub>	MQGPWV---L	LLLGLRLQLS	LGIIPVEEEN	PDFWNRQAAE	ALGAAKKLQP	AQT-AAKNL I	56
PLALP <sub>hs</sub>	MLGPCMLLLL	LLLGLRLQLS	LGIIPVEEEN	PDFWNREAAE	ALGAARKLQP	AQT-AAKNL I	59
IALP <sub>hs</sub>	MQGPWV---L	LLLGLRLQLS	LGVIPIAEEEN	PAFWNRQAAE	ALDAAKKLQP	IQK-VAKNL I	56
Consensus	XXXXXX	XXXXXX	XXXXXX	XXXXXX	XXXXXX	XXXXXX	
TNALP <sub>Prn</sub>	MFLGDGMGV S	TVTAARILKG	QLHHNTGEET	RLEMDFKFPFV	ALSKTYNTNA	QVPDSAGTAT	115
TNALP <sub>Mm</sub>	MFLGDGMGV S	TVTAARILKG	QLHHNTGEET	RLEMDFKFPFV	ALSKTYNTNA	QVPDSAGTAT	115
TNALP <sub>Phs</sub>	MFLGDGMGV S	TVTAARILKG	QLHHNPGEET	RLEMDFKFPFV	ALSKTYNTNA	QVPDSAGTAT	115
TNALP <sub>Cf</sub>	MFLGDGMGV S	TVTAIRILKG	QLHHNPGEET	RLEMDFKFPYV	ALSKTYNTNA	QVPDSAGTAT	93
TNALP <sub>FC</sub>	MFLGDGMGV S	TVTAARILKG	QLHHNPGEET	RLEMDFKFPYV	ALSKTYNTNA	QVPDSAGTAT	115
TNALP <sub>Bt</sub>	MFLGDGMGV S	TVTAARILKG	QLHHSPGEET	KLEMDFKFPYV	ALSKTYNTNA	QVPDSAGTAT	115
GALP <sub>hs</sub>	IFLGDGMGV S	TVTAARILKG	QKKDKLGPET	FLAMDRFPYV	ALSKTYSVDK	HVPDSGATAT	116
PLALP <sub>hs</sub>	IFLGDGMGV S	TVTAARILKG	QKKDKLGPET	PLAMDRFPYV	ALSKTYNVDK	HVPDSGATAT	119
IALP <sub>hs</sub>	LFLGDGLGV P	TVTATRILKG	QKNGKLGPEI	PLAMDRFPYL	ALSKTYNVDR	QVPDSAATAT	116
Consensus	:* * * * :* * *	* * * * :* * *	* : * * *	* * * :* * :	* * * * * :* :	* * * * * :* :	
XFLGDGXGV X	TVTAXRILKG	QXXXXGXEX	XLMXDFPXX	ALSKTYXXX	XVPDSXXAT		

FIG. 9 (cont'd)

## FIG. 9 (cont'd)

TNALPrn	MQYELNRNNL	TDPSSLSEMVE	VALRILTKNP	KGFFLLVEGG	RIDHGHHEGK	AKQALHEAVE	354
TNALPmm	MQYELNRNNV	TDPSSLSEMVV	VAIQILRKNP	KGFFLLVEGG	RIDHGHHEGK	AKQALHEAVE	354
TNALPhs	MQYELNRNNV	TDPSSLSEMVE	IAIKILSKKP	RGFFLLVEGG	RIDHGHHEGK	AKQALHEAVE	332
TNALPcf	MQYELNRNNST	TDPSSLSEMVE	IAIKILSKNP	KGFFLLVEGG	RIDHGHHEGK	AKQALHEAVE	354
TNALPbt	MQYELNRNNNA	TDPSSLSEMVE	MAIRILNKNP	KGFFLLVEGG	RIDHGHHEGK	AKQALHEAVE	354
GALPhs	MKYEIHRDST	LDPSSLMEMTE	AALLLSSRNP	RGFFFLFVEGG	RIDHGHHESR	AYRALTETIM	352
PLALPhs	MKYEIHRDST	LDPSSLMEMTE	AALRLLSRNP	RGFFFLFVEGG	RIDHGHHESR	AYRALTETIM	355
IALPhs	TKYEIHRDPT	LDPSSLMEMTE	AALRLLSRNP	RGFFFLFVEGG	RIDHGHHEGV	AYQALTEAVM	352
Consensus	: * : * : *	* : * : * : *	* : * : * : * : * : *	* : * : * : * : * : *	* : * : * : * : * : *	* : * : * : * : * : *	
	XXYEXXXRXXX	XDPSSLXEMXX	XAXXXLXXXX	XGFXLXVEGG	RIDHGHHEXX	AXXALXEXXX	
TNALPrn	MDEAIGKAGT	MTSQKDILTV	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMVD	TDKKPFTAIL	414
TNALPmm	MDQAIKGAGA	MTSQKDILTV	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMVD	TDKKPFTAIL	414
TNALPhs	MDRAIGQAGS	LTSSEDTLTV	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMLSD	TDKKPFTAIL	414
TNALPcf	MDRAIGKAGV	MTSLEDTLTV	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMVD	TDKKPFTAIL	392
TNALPbt	MDQAIQGRAGA	MTSVEDDTLTI	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMVD	TDKKPFTSIL	414
GALPhs	MDQAIQGQAGA	MTSVEDDTLTV	VTADHSHHVFT	FGGYTPRGNS	IFGLAPMVD	TDKKPFTAIL	414
PLALPhs	FDDAIERAGQ	LTSEEDTLSL	VTADHSHHVFS	FGGYPLRGSS	IFGLAPGKAR	-DRKAYTVLL	411
IALPhs	FDDAIERAGQ	LTSEEDTLSL	VTADHSHHVFS	FGGYPLRGSS	IFGLAPGKAR	-DRKAYTVLL	414
Consensus	: * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	
	XDXAIXXAGX	XTSXXDTLXX	VTADHSHHVFX	FGGYXXRGXS	IFGLAPXXXX	XDXKXXTXXL	
TNALPrn	YGNPGPYKVV	DGERENVSMV	DYAHNNYQAO	SAPLRETH	GGEDVAVFAK	GPMALILLHGV	474
TNALPmm	YGNPGPYKVV	DGERENVSMV	DYAHNNYQAO	SAPLRETH	GGEDVAVFAK	GPMALILLHGV	474
TNALPhs	YGNPGPYKVV	GGERENVSMV	DYAHNNYQAO	SAPLRETH	GGEDVAVFSK	GPMALILLHGV	474
TNALPcf	YGNPGPYKVV	GGERENVSMV	DYAHNNYQAO	SAPLRETH	GGEDVAVFAK	GPMALILLHGV	452
TNALPbt	YGNPGPYKVV	GGERENVSMV	DYAHNNYQAO	SAPLRETH	GGEDVAVFAK	GPMALILLHGV	474
GALPhs	YGNPGPYKLV	DGARPDVTE	ESGSPEYRQQ	SAPLDEETH	GGEDVAVFAK	GPMALILLHGV	474
PLALPhs	YGNPGPYKLV	DGARPDVTE	ESGSPEYRQQ	SAPLDEETH	AGEDVAVFAAR	GPOAHLVHGV	471
IALPhs	YGNPGPYVFN	SGVRPDVNES	ESGSPDYQQQ	AAVPLSETH	GGEDVAVFAAR	GPOAHLVHGV	474
Consensus	: * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	: * : * : * : * : *	
	YGNPGPYXXX	XGXRXVXXX	XXXXXXXXXQ	XAVPLXXETH	XGEDVAVFX	GPXAHLXHGV	

## FIG. 9 (cont'd)

TNALPrn	HEQNYIPHVM	AYASCIG---	-ANLDHCAWA	SSASSSPSPGA	LLPLPLALFFPL	RTLF-----	524
TNALPmm	HEQNYIPHVM	AYASCIG---	-ANLDHCAWA	GGSAPSPPGA	LLPLPLAVLSL	PTLF-----	524
TNALPhs	HEQNYVPHVM	AYAACIG---	-ANLGHCAPA	SSAGSLLAAGP	LLLALALYPL	SVLF-----	524
TNALPcf	HEQNYIPHVM	AYAACIG---	-ANQDHCASA	SSAGGSPGP	LLLLALLPV	GILF-----	502
TNALPfc	HEQNYIPHVM	AYAACIG---	-ANLDHCAWA	SSAGGSPGP	LLLLALPSL	GILF-----	524
TNALPbt	QEQNYIPHVM	AYAACIG---	-ANRDHCAWA	SSSGSPSPGP	LLLLALLPL	GSLF-----	524
GALPhs	QEQTFIAHVM	AFAACLEPYT	ACDLAPPAGT	TDAAHPGPSV	VPAALLPLAG	TLILLGTATA	531
PLALPhs	QEQTFIAHVM	AFAACLEPYT	ACDLAPPAGT	TDAAHPGRSV	VPAALLPLAG	TLILLETATA	534
IALPhs	QEQSFIAHVM	AFAACLEPYT	ACDLAPPACT	TDAAHP---	VAAISLPLAG	TLILLGASAA	527
Consensus	XEQXXXXHVM	AXAXCXXXXX	XXXXXXAXX	XXXXXX	XXXXXX	XXXXXX	XXXXXX
	-	(SEQ ID NO: 1210)					
	-	(SEQ ID NO: 1209)					
	-	(SEQ ID NO: 1208)					
	-	(SEQ ID NO: 1211)					
	-	(SEQ ID NO: 1207)					
	-	(SEQ ID NO: 1206)					
	P 532	(SEQ ID NO: 1213)					
	P 535	(SEQ ID NO: 1214)					
	P 528	(SEQ ID NO: 1212)					
	X	(SEQ ID NO: 1215)					

FIG. 10

## ALP-TNALP Consensus Sequences Excluding Pathogenic Mutations

XXXXXXXXXX	XXXXXXXXXX	XXXXXXXXXX	PXXXXXXX	XXXXXXXAAXX	XLXXXAXXLQX	XXXXXXAKNXTI	60
XFIIGDGMGVX	TVTAXRILKG	QXXXXXXGXFX	XIXMDXF	FPXX	ALSKTYXXXX	XVPDSSXXTAT	120
AYLCGVKXNX	XTXGXSAAAX	XXXCNNTXGN	EVT	SXXXXXAK	XXGKSVGXVT	TTRVXHAXXPX	180
XXYAHXXXRX	WYSDDXXXPXX	AXXXGCXDIA	XQLXXNXXDI	I	XVIXGGGRKY	MXXXXXXXDXE	240
YXXDXXXXGX	RLDGXXLXXX	WXXXXXXX	XXXXWNRTXL	XXXXXXXVX	XLXGLFXPGD	300	
MXYEXXXRXXX	XDPSSLXEMXX	XAXXXLXXXX	XGFXXLXVEGG	RIDHGHHEXXX	AXXVALXEXXXX	360	
XDXAIXXAGX	XTSXXXDTLXX	VTADHSHVFX	FGGYXXRGXS	IFGLAPXXXX	DXKXXXTXXL	420	
YGNGPGYXXX	XGXRXXXVXX	XXXXXXQ	XAVPLXXXETH	XGEDVAVFXX	GPyAHLXHGV	480	
XEQXXXXHVM	AXAXCXXXXX	XXXXXXAAXX	XXXXXXLXXXXX	XXXXXXLXXXXX	XXLXXXXXXX	540	
X							541

(SEQ ID NO: 1218)

**TNALP Consensus Sequences Excluding Pathogenic Mutations**

**FIG. 11**

MIXPFLXLAI	GTCXXXSXVP	EKEXDPXYWR	XQAQXTLKKXA	LXLQXLNTNV	AKNXIMFLGD	60
GMGVSTVTAX	RIIKGQLHHX	XGEETIXLEMD	KFPXVALSKT	YNTNAQVPDS	AGTATAYLCG	120
VKANE GTVGV	SAATXRXXCN	TTQGNEVTSI	LRWAKDXGKS	VGIVTTRVN	HATPSAXYAH	180
SADRDWYSDN	EMPPEALSQG	CKDIAYQLMX	NXXDIXVIMG	GGRKYMXPKN	XTDVEYEXDE	240
KXXGXRLDGL	XIXXXWXXFK	PXXXKHSXXXW	NRTXLIIXLDP	XXVDYLLGLF	XPGDMQYELN	300
RNXXXTDPSSL	EMVXXXAXLL	XKXXXGFFLL	VEGGRIDHGH	HEGKAKQALH	EAVEMDXAIG	360
XAGXXTSXXD	TLTXVTAHDHS	HVFITFGGYTP	RGNSIIFGLAP	MKSDDTDKKPF	TXILYGNNGPG	420
YKVVVXGEREN	VSMVDYAHNN	YQAQSAVPLR	HETHGEDVA	VFXKGPMMAHL	LHGVVXEQNYX	480
PHVMAYAXCI	GANXXXHCAXA	XSXXXXXXGX	LXLXLAXXXX	XXLXF		524

(SEQ ID NO: 1219)

FIG. 12

## Multiple Sequence Alignment of Natriuretic Peptides

Human ANP	-----SLRRLSSCFGGMDRIGAQSGLGCNSFRY-----	(SEQ ID NO: 1)
Human urodilatin	-----TAPRSLRRLSSCFGGMDRIGAQSGLGCNSFRY-----	(SEQ ID NO: 2)
Human BNP	-SPKMVQGSGCGCFGRKMDRISSSGLGCKVLRRH-----	(SEQ ID NO: 3)
Human CNP22	-----GLSKGCGFGLKLDRIGSMSGLGC-----	(SEQ ID NO: 4)
DNP	-----EVKYDPPCFGHKIDRINHVSNLGGCPSSLRDPRNAPSSTSA-----	(SEQ ID NO: 5)
CONSENSUS	***** : * * * . * . * * .	
SEQUENCE	<b>CFGXXXDRIXXSXSLGC</b>	(SEQ ID NO: 6)

FIG. 13

**Sequences of Human CNP53, CNP22, and CNP (ring only)**

Human CNP53	DLRVDTKSRAAWARLLQEHPNARKYKGANKKGLSKG	<b>CFG</b> LIKLDRIGSMSGLGC	(SEQ ID NO: 11)
Human CNP22	-----	<b>CFG</b> LIKLDRIGSMSGLGC	(SEQ ID NO: 4)
Human CNP (ring only)	-----	<b>CFG</b> LIKLDRIGSMSGLGC	(SEQ ID NO: 12)

## Multiple Sequence Alignment of C-Type Natriuretic Peptides from Various Species

FIG. 14

Human	-----GLSKG <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 4)
Bovine	-----GLSKG <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 13)
Sheep	-----GLSKG <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 14)
Mouse	-----GLSKG <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 15)
Pig	-----GLSKG <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 16)
<i>Micrurus fulvius fulvius</i>	GLAKEALGDC <u>CGFG</u> KLDRIGTSSGLGC	(SEQ ID NO: 17)
Taenioptygia guttata	-----GLSRS <u>SCFG</u> VKLDRIGTSSGLGC	(SEQ ID NO: 18)
Chicken	-----SRG <u>CGFG</u> VKLDRIGATSGLGC	(SEQ ID NO: 19)
<i>Rana catesbeiana</i>	-----GWNRC <u>CGFG</u> KLDRIGSISGLGC	(SEQ ID NO: 20)
Fel	-----GWNRC <u>CGFG</u> KLDRIGSISGLGC	(SEQ ID NO: 21)
Trout	-----GWNRC <u>CGFG</u> KLDRIGMSGLGC	(SEQ ID NO: 22)
<i>Ornithorhynchus anatinus</i>	-----GLSKG <u>CGFG</u> KLDRIGSTSGLGC	(SEQ ID NO: 23)
<i>Trimeresurus flavoviridis</i>	-----KGCFGH <u>CGFG</u> KLDRIGSTSGLGC	(SEQ ID NO: 24)
<i>Polypterus endlicheri</i>	-----SK <u>CGFG</u> KLDRIGSISGLGC	(SEQ ID NO: 25)
<i>Xenopus laevis</i>	-----LSKG <u>CGFG</u> KLDRIGVVSGLGC	(SEQ ID NO: 26)
<i>Oryzias latipes</i>	-----GCFG <u>CGFMK</u> DRIGSISGLGC	(SEQ ID NO: 27)
<i>Tetraodon nigroviridis</i>	-----GCFG <u>CGMK</u> DRIGSISGLGC	(SEQ ID NO: 28)
<i>Pseudechis australis</i>	-----SKIGDG <u>CGFG</u> PLDHIGSVSGLGC	(SEQ ID NO: 29)
	***** : * : * *	*****
CONSENSUS SEQUENCE	XXXXXXXXXXXX <u>CGFGXXDXIGXXSGLGC</u>	(SEQ ID NO: 30)

## FIG. 15A

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp  Q8AYR5  ANFC2_ORYLA	---MAVSSSS-----LILTVFILSVAEVTTRP-SSDRDE-----	30
sp  Q805D5  ANFC2_TAKRU	---MAASSSF-----VPLVLLFLAIPVEPRP-SMTRDE-----	30
sp  Q76KW6  ANFC_ACITR	---MSISSSSSSSSSSSCLLISLMLLAASCQGRPDQHNRH-----	40
sp  Q61839  ANFC_MOUSE	---MHLSQLI-----ACALLALLSLRPSEAKPGTP-----	28
tr  Q544K5  Q544K5_MOUSE	---MHLSQLI-----ACALLALLSLRPSEAKPGTP-----	28
tr  Q8VHG9  Q8VHG9_NOTAL	---MHLSQLI-----ACALLALLSLRPSEAKPGTP-----	28
sp  P55207  ANFC_RAT	---MHLSQLI-----ACALLALLSLRPSEAKPGTP-----	28
sp  P55206  ANFC_BOVIN	---MHLSQLL-----ACALLALLSLRPSEAKPGAP-----	28
sp  P56283  ANFC_SHEEP	---MHLSQLL-----ACALLLSSLLSLRPSEAKPGAP-----	28
sp  P18104  ANFC_PIG	---MHLSQLL-----ACALLLTLLSLRPSEAKPGAP-----	28
sp  P23582  ANFC_HUMAN	---MHLSQLL-----ACALLLTLLSLRPSEAKPGAP-----	28
sp  P84715  ANF39_ORNAN	---MHLSHLL-----AWALLTLLSLR-AEAKPPSPQ-----	28
tr  Q9QZ96  Q9QZ96_CAVPO	-----	
sp  Q80017  ANFC4_ORYLA	---MNLSYLV-----ACGLIVTFLSDK-MDAQPLTQA-----	29
sp  Q805D3  ANFC4_TAKRU	---MNLSYLV-----ACGLMITLLSVR-MGAKPLSQA-----	29
tr  C1BX15  C1BX15_ESOLU	---MNISYLV-----ACGLMITLLSVR-SGAKPLTAAQ-----	29
tr  D2KXA5  D2KXA5_ANGJA	---MNVSQLM-----VCGLIMALFSFS-TEAKSLIPA-----	29
tr  Q1XGY7  Q1XGY7_9ACTI	---MNISHLV-----ACGLLIVALLTVT-MEAKPLTQS-----	29
sp  P40756  ANFD_RANCA	---MHFCHIV-----GWGLVLAVLYLR-TEAKPVAQAH-----	29
sp  P0C7P5  BNP_TRIFL	---MFVSRLA-----ASGLLLLALLALSLDGKPVHQSKPGR-----	33
sp  P0C7P6  BNP_TRIGA	---MFVSRLA-----ASGLLLLALLALSLDGKPVQE-KPGR-----	32
sp  Q6LEMS5  BNP1_BOTJA	---MVLSRLA-----ASGLLLLALLALSLVDGKPVQQWAQS-----	32
sp  Q9PW56  BNP2_BOTJA	---MVLSRLA-----ASGLLLLALLALSLVDGKPVQQWAQGG-----	33
sp  P68515  BNP_BOTIN	---MVLSRLA-----ASGLLLLALLALSLVDGKPVQQWAQGG-----	33
sp  Q90Y12  BNP_CRODU	---MFVSRLA-----ASGLLLLALLALAVSLDGKPLQQWS-----	30
sp  Q2PE51  BNP_CRODO	---MFVSRLA-----ASGLLLLALLALAVSLDGKPLQQWS-----	30
sp  B0VXV8  BNP_SISCA	---MFVSRLA-----ASGLLLLALLALAVSLDGKPVQQWS-----	30
sp  Q27J49  BNP_LACMU	---MFVSRLA-----ASGLLLLALLALAVSLDGKPVQQWSH-----	31
sp  P01021  BNP_AGKHA	---MFVSRLA-----ASGLLLLALMALSLDGKPVQQWSQGRPPGPI	39
sp  Q09GK2  VNP_PHIOL	---MVASRLA-----AGGLLLLALLALALDGKAPP-QP-----	30
tr  D1MZV3  D1MZV3_RHATT	---MFASRLA-----ALGLLLLALV--LDGKAPPQP-----	28
tr  Q7T1M4  Q7T1M4_BOTJR	---HEKPSRSG-----AKSAVGAKLAASSDSADECSSGRK-----	34
tr  Q402A2  Q402A2_PETMA	-----	
tr  Q402A3  Q402A3_LAMJA	---MKLQLLMMV-----VVVGWSWTFLG--VGAKPLTSYELYD-----	32
tr  Q402A1  Q402A1_9PETR	---MRRQVLVMV-----VMVVVVVMMSGKSVTAKPVASYELLD-----	35
sp  P21805  ANFC_CHICK	-----	
tr  A9CDT6  A9CDT6_CHICK	---MKLQLFC-----PGFFLLLIVSQKQAMAKPIS-----	26
sp  P20968  ANFC_RANCA	---MSYKRGTC-----LGFIMLLMVSHHHTKGKPLS-----	28
sp  P55208  ANFC_TRISC	---MSGQTSFY-----CGLLVLILIQAQ--ARPRS-----	25
sp  P23259  ANFC_SCYCA	-----RPRS-----	4
sp  P41319  ANFC_SQUAC	---MSGHTSFY-----CGLLLPLLQVQ--ARPRA-----	25
tr  Q2MH72  Q2MH72_9CHON	---MSGNTNFY-----CGLVLLLLQVQ--GRPRS-----	25
tr  Q2MH73  Q2MH73_9CHON	---MSGNTNFY-----CGLVLLLLQVQ--GRPRS-----	25
tr  Q2MH71  Q2MH71_9CHON	---MSGNTNFY-----CGLVLLLLQVQ--GRPRS-----	25
tr  Q2MH74  Q2MH74_DASAK	---MSGNTNFY-----CGLVLLLLQVQ--GRPRS-----	25
tr  Q2PF87  Q2PF87_CALMI	---MNAHVSFP-----CGLMLLLLQVQVQ-ARPRTG-----	28
sp  Q80018  ANFC3_ORYLA	---MSLRAF-----MLCVCLLQLQSVG--ARPAS-----	23
tr  Q4ADV1  Q4ADV1_ORYLA	---MSLRAF-----MLCVCLLQLQSVG--ARPAS-----	23
sp  Q805D4  ANFC3_TAKRU	---MSLNLPGY-----ALFFILLVASSG--AKPAP-----	25
tr  C0H7B0  C0H7B0_SALSA	---MKMISNIQFF-----CLTALVLLNLNVG--ANPMS-----	27
tr  C1BWD1  C1BWD1_ESOLU	---MISNIQFC-----CLSVLVLNLNVG--AKPVS-----	25
tr  D2KXA3  D2KXA3_ANGJA	---MISNITIY-----CISSLLFLNLNVG--GKPV-----	25
tr  B3DJJ2  B3DJJ2_DANRE	---MIANISVF-----CVSSLLNLNLNVG--AKPVS-----	25
tr  Q1XGY8  Q1XGY8_9ACTI	---MVSRLTVY-----CALFIIVLSQVS--AKPVS-----	25
sp  Q8AXR2  ANFC2_ONCMY	---ML--YPA-----LLCAALLLIAPLGHTEGRTLYPSPD-----	30
sp  Q8AXR3  ANFC1_ONCMY	---ML--YPA-----LLCAALLLIAPLGHTEGRTLHPSPD-----	30
tr  C1BKS8  C1BKS8_OSMMO	---ML--CPV-----LLCATLVLPPLESEGRALHPSPE-----	30
tr  Q805E7  Q805E7_OREMO	---ML--CPV-----LLCAALLLLTPLEITEARALHPSPD-----	30
tr  C3KH23  C3KH23_ANOFI	---ML--CPV-----LLCATLLLLTPFEVTEARALHPPPD-----	30
sp  Q8AYR6  ANFC1_ORYLA	---ML--CPV-----LLCAALLLLTPVEITDARALQQPSD-----	30
sp  Q805D6  ANFC1_TAKRU	---ML--CPA-----LLCAALLLLTPVEITDARALQQPSD-----	30
sp  P18145  ANFC_ANGJA	---MM--CKA-----LVFAVILLAVPLERADSRALRTEPV-----	30
tr  Q1XGY9  Q1XGY9_9ACTI	---MMGCSAPLLTGHRLCLFLMASSLSPIHSRAFRSPP-----	38
tr  A9CDT5  A9CDT5_CHICK	---MLGLPA-----WPCSLFLLLVLLSASVQAMSSSGQR-----	31

FIG. 15B

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp | Q8AYR5 | ANFC2\_ORYLA  
sp | Q805D5 | ANFC2\_TAKRU  
sp | Q76KW6 | ANFC\_ACITR  
sp | Q61839 | ANFC\_MOUSE  
tr | Q544K5 | Q544K5\_MOUSE  
tr | Q8VHG9 | Q8VHG9\_NOTAL  
sp | P55207 | ANFC\_RAT  
sp | P55206 | ANFC\_BOVIN  
sp | P56283 | ANFC\_SHEEP  
sp | P18104 | ANFC\_PIG  
sp | P23582 | ANFC\_HUMAN  
sp | P84715 | ANF39\_ORNAN  
tr | Q9QZ96 | Q9QZ96\_CAVPO  
sp | Q80017 | ANFC4\_ORYLA  
sp | Q805D3 | ANFC4\_TAKRU  
tr | C1BXI5 | C1BXI5\_ESOLU  
tr | D2KXA5 | D2KXA5\_ANGJA  
tr | Q1XGY7 | Q1XGY7\_9ACTI  
sp | P40756 | ANFD\_RANCA  
sp | P0C7P5 | BNP\_TRIFL  
sp | P0C7P6 | BNP\_TRIGA  
sp | Q6LEM5 | BNP1\_BOTJA  
sp | Q9PW56 | BNP2\_BOTJA  
sp | P68515 | BNP\_BOTIN  
sp | Q90Y12 | BNP\_CRODU  
sp | Q2PE51 | BNP\_CRODO  
sp | B0VXV8 | BNP\_SISCA  
sp | Q27J49 | BNP\_LACMU  
sp | P01021 | BNP\_AGKHA  
sp | Q09GK2 | VNP\_PHIOL  
tr | D1MZV3 | D1MZV3\_RHATT  
tr | Q7T1M4 | Q7T1M4\_BOTJR  
tr | Q402A2 | Q402A2\_PETMA  
tr | Q402A3 | Q402A3\_LAMJA  
tr | Q402A1 | Q402A1\_9PETR  
sp | P21805 | ANFC\_CHICK  
tr | A9CDT6 | A9CDT6\_CHICK  
sp | P20968 | ANFC\_RANCA  
sp | P55208 | ANFC\_TRISC  
sp | P23259 | ANFC\_SCYCA  
sp | P41319 | ANFC\_SQUAC  
tr | Q2MH72 | Q2MH72\_9CHON  
tr | Q2MH73 | Q2MH73\_9CHON  
tr | Q2MH71 | Q2MH71\_9CHON  
tr | Q2MH74 | Q2MH74\_DASAK  
tr | Q2PF87 | Q2PF87\_CALMI  
sp | Q80018 | ANFC3\_ORYLA  
tr | Q4ADV1 | Q4ADV1\_ORYLA  
sp | Q805D4 | ANFC3\_TAKRU  
tr | C0H7B0 | C0H7B0\_SALSA  
tr | C1BWD1 | C1BWD1\_ESOLU  
tr | D2KXA3 | D2KXA3\_ANGJA  
tr | B3DJJ2 | B3DJJ2\_DANRE  
tr | Q1XGY8 | Q1XGY8\_9ACTI  
sp | Q8AXR2 | ANFC2\_ONCMY  
sp | Q8AXR3 | ANFC1\_ONCMY  
tr | C1BKS8 | C1BKS8\_OSMMO  
tr | Q805E7 | Q805E7\_OREMO  
tr | C3KH23 | C3KH23\_ANOFI  
sp | Q8AYR6 | ANFC1\_ORYLA  
sp | Q805D6 | ANFC1\_TAKRU  
sp | P18145 | ANFC\_ANGJA  
tr | Q1XGY9 | Q1XGY9\_9ACTI  
tr | A9CDT5 | A9CDT5\_CHICK

## FIG. 15C

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp Q8AYR5 ANFC2_ORYLA	
sp Q805D5 ANFC2_TAKRU	
sp Q76KW6 ANFC_ACITR	
sp Q61839 ANFC_MOUSE	
tr Q544K5 Q544K5_MOUSE	
tr Q8VHG9 Q8VHG9_NOTAL	
sp P55207 ANFC_RAT	
sp P55206 ANFC_BOVIN	
sp P56283 ANFC_SHEEP	
sp P18104 ANFC_PIG	
sp P23582 ANFC_HUMAN	
sp P84715 ANF39_ORNAN	
tr Q9QZ96 Q9QZ96_CAVPO	
sp Q80017 ANFC4_ORYLA	
sp Q805D3 ANFC4_TAKRU	
tr C1BX15 C1BX15_ESOLU	
tr D2KXA5 D2KXA5_ANGJA	
tr Q1XGY7 Q1XGY7_9ACTI	
sp P40756 ANFD_RANCA	
sp P0C7P5 BNP_TRIFL	
sp P0C7P6 BNP_TRIGA	
sp Q6LEMS BNP1_BOTJA	
sp Q9PW56 BNP2_BOTJA	
sp P68515 BNP_BOTIN	
sp Q90Y12 BNP_CRODU	
sp Q2PE51 BNP_CRODO	
sp B0VXV8 BNP_SISCA	
sp Q27J49 BNP_LACMU	
sp P01021 BNP_AGKHA	
sp Q09GK2 VNP_PHIOL	
tr D1MZV3 D1MZV3_RHATT	
tr Q7T1M4 Q7T1M4_BOTJR	
tr Q402A2 Q402A2_PETMA	
tr Q402A3 Q402A3_LAMJA	
tr Q402A1 Q402A1_9PETR	
sp P21805 ANFC_CHICK	
tr A9CDT6 A9CDT6_CHICK	
sp P20968 ANFC_RANCA	
sp P55208 ANFC_TRISC	
sp P23259 ANFC_SCYCA	
sp P41319 ANFC_SQUAC	
tr Q2MH72 Q2MH72_9CHON	
tr Q2MH73 Q2MH73_9CHON	
tr Q2MH71 Q2MH71_9CHON	
tr Q2MH74 Q2MH74_DASAK	
tr Q2PF87 Q2PF87_CALMI	
sp Q80018 ANFC3_ORYLA	
tr Q4ADV1 Q4ADV1_ORYLA	
sp Q805D4 ANFC3_TAKRU	
tr C0H7B0 C0H7B0_SALSA	
tr C1BWD1 C1BWD1_ESOLU	
tr D2KXA3 D2KXA3_ANGJA	
tr B3DJJ2 B3DJJ2_DANRE	
tr Q1XGY8 Q1XGY8_9ACTI	
sp Q8AXR2 ANFC2_ONCMY	
sp Q8AXR3 ANFC1_ONCMY	
tr C1BKS8 C1BKS8_OSMMO	
tr Q805E7 Q805E7_OREMO	
tr C3KH23 C3KH23_ANOFI	
sp Q8AYR6 ANFC1_ORYLA	
sp Q805D6 ANFC1_TAKRU	
sp P18145 ANFC_ANGJA	
tr Q1XGY9 Q1XGY9_9ACTI	
tr A9CDT5 A9CDT5_CHICK	
-----PLSAQQWMPEGRPPHP-----	68
-----HHIPPLSVQKFPPGWKPTHHIPPLEVQQWSQGGP-----	89
WPHPQIPPLTVQQWAQGRAPGPP-IPPLTVQQWAQGRAPHPIPPAPLQ	117
WPHPQIPPLTVQQWAQGRAPGPP-IPPLTVQQWAQGRAPHPIPPAPLQ	121
WPHPQIPPLTVQQWAQGRAPGPP-IPPLEVQQWAQGRAPHPIPPAPLQ	121
-----HLE-IPPLVVQWNK-----	47
-----HLE-IPPLVVQWNK-----	47
-----PKVPLVVQQWSQN-----	64
-----WPHPQ-IPPLVVQWNK-----	64
-----GHHIPPLVVQQWSQKKWPPGHH-IPPLVVQWNK-----	102
-----PPRKIPPLVVQQWSQG-LPPRPK-IPPLVVQWNK-----	121
-----PPIPPLTVQQWAQAR-PPHPP-IPPAAPLQKWPVQK-----	73

## FIG. 15D

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

```

sp |Q8AYR5 |ANFC2_ORYLA          -----EQVLKSLFGPHLTS----- IL 47
sp |Q805D5 |ANFC2_TAKRU          -----AQVLRALFGARLSSI----- IS 47
sp |Q76KW6 |ANFC_ACITR          -----KSQLAGLGAEVAA----- LE 57
sp |Q61839 |ANFC_MOUSE          -----PKVPRTPPGEELADS----- QAAG 47
tr |Q544K5 |Q544K5_MOUSE          -----PKVPRTPPGEELADS----- QAAG 47
tr |Q8VHG9 |Q8VHG9_NOTAL          -----PKVPRTPPGEELADS----- QAAG 47
sp |P55207 |ANFC_RAT          -----PKVPRTPPGEELADS----- QAAG 47
sp |P55206 |ANFC_BOVIN          -----PKVPRTPPGEELAEP----- QAAG 47
sp |P56283 |ANFC_SHEEP          -----PKVPRTPSCEEVAEP----- QAAG 47
sp |P18104 |ANFC_PIG          -----PKVPRTPGEEVAEP----- QAAG 47
sp |P23582 |ANFC_HUMAN          -----PKVPRTPPAEELAEP----- QAAG 47
sp |P84715 |ANF39_ORNAN          -----PKVPRSP-GDEASEA----- VAAN 46
tr |Q9QZ96 |Q9QZ96_CAVPO          -----
sp |Q80017 |ANFC4_ORYLA          -----QKSLRSLLGEEELAEP----- LESG 48
sp |Q805D3 |ANFC4_TAKRU          -----QKSFRSLLGEEELAEP----- LESE 48
tr |C1BX15 |C1BX15_ESOLU          -----QKSLRNLGEELSEF----- LASG 48
tr |D2KXA5 |D2KXA5_ANGJA          -----EKSLRNLGEELSEY----- LASG 48
tr |Q1XGY7 |Q1XGY7_9ACTI          -----QKSLRNLGEEL SAY----- LTSD 48
sp |P40756 |ANFD_RANCA          -----QKSLRALLGEELA EY----- LVSG 48
sp |P0C7P5 |BNP_TRIFL          -----RSEVPPVVVQHESPAGTTAFREELSPG----- PEAASG 103
sp |P0C7P6 |BNP_TRIGA          -----RSEL---VQHESPAGTTAFREELSLG----- PEAASG 120
KWAPLQWAPLLQPHESPASGTTALREELSLG----- PEAASG 155
KWAPVQKWAPLLQPHESPASGTTALREELSLG----- PEAASG 159
KWAPVQKWAPLLQPHESPASGTTALREELSLG----- PEAASG 159
-----SPTQLQARESPAGTTALREELSLG----- PEAALD 78
-----SPTQLQARESPAGTTALREELSLG----- PEAALD 78
-----SPTQLQRESPAGTTALREELSLG----- PDAALD 95
---PPI -SPPLLKPHESPAGTTALREELSLG----- PEAALD 136
---PPVSPPLLQPHESPAGTTALREELSLG----- PEAASG 156
-----LRKAPAGGTTAWRRELTEQ----- PEGASR 55
-----LRKAPAGGTTALQRQLTEQQQQQQ----- QAEQSSG 59
-WAPVQKWAPLLQPHESPAGTTALREELSLG----- PEAASG 110
-----SGSEPWE GG-FLPRVSSSSSSSS----- EVEPLAE 30
-----DAGSEPWE GG-FLPRISSSSSS----- AERLAD 60
-----DTNSEPWE GGSLLPSLPSQGED----- SHPLS- 63
-----SLQSLSLM LDEE----- L 39
-----SLQNLSLRLL EDN----- F 41
-----DDSLQTL SRLLEDE----- Y 40
-----DDSLQTL SRLLEDE----- Y 19
-----DDSLQVLSRLL ED----- Y 40
-----DDSLQALTRLLEDE----- Y 40
-----DDSLQALTRLLEDE----- Y 40
-----DDSLQALTRLLEDE----- Y 40
-----VDSLQILSRLLEDE----- Y 43
-----ELQNLERLL----- 32
-----ELQNLERLL----- 32
-----DLQILEPPLSSLEEQQEEMQEEVQE KVQEQQEEVQ----- 59
-----NLQSLKQLLEEE----- SH 41
-----NLQSLKLFLEE----- SN 38
-----SLQSLKELLEEE----- SN 39
-----SLQSLKQLLDEE----- VN 39
-----SLQSFQAQLLEDE----- SN 39
-----AIQFVEQFLDR----- YN 43
-----AIQFVEQFLDR----- YN 43
-----GLQFVEQFLER----- CT 43
-----AVQFVEQFLER----- YN 43
-----AVQFMEQFLER----- YN 43
-----AVQFVEQFLDR----- YN 43
-----AAQFMEQFLES----- YN 43
-----AIQFVEQFLEH----- YN 43
-----LQFLSTLLEKE----- YG 51
-----LQVLLS QLLPS----- DS 44

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## FIG. 15E

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp  Q8AYR5  ANFC2_ORYLA	APPTSNDs--TEGSSGSPEPP-----	TPSEAPVLIH-- 76
sp  Q805D5  ANFC2_TAKRU	TPVNTDDI--AELLPRPGPPR-----	SFGASPGALRGLTR-- 81
sp  Q76KW6  ANFC_ACITR	DAGAADGSSGEEAALSQRAPPS-----	IRALHPRSGRLGLRDD 95
sp  Q61839  ANFC_MOUSE	GNQKKGDKTPGGGANLKGDRS-----	RLLRDLR 76
tr  Q544K5  Q544K5_MOUSE	GNQKKGDKTPGGGANLKGDRS-----	RLLRDLR 76
tr  Q8VHG9  Q8VHG9_NOTAL	GNQKKGDKTPGGGANLKGDRS-----	RLLRDLR 97
sp  P55207  ANFC_RAT	GNQKKGDKTPGGGANLKGDRS-----	RLLRDLR 76
sp  P55206  ANFC_BOVIN	GGQKKGDKTPGGGANLKDDRS-----	RLLRDLR 76
sp  P56283  ANFC_SHEEP	GGQKKGDKTPGGGANLKDDRS-----	RLLRDLR 76
sp  P18104  ANFC_PIG	GGQKKGDKTPGGGANLKGDRS-----	RLLRDLR 76
sp  P23582  ANFC_HUMAN	GGQKKGDKAPGGGANLKGDRS-----	RLLRDLR 76
sp  P84715  ANF39_ORNAN	GGGKKGDKEP-----KGDRP-----	RLLREL 68
tr  Q9QZ96  Q9QZ96_CAVPO	-----	
sp  Q80017  ANFC4_ORYLA	ENENRLDDVRSRM-----	RLLRDLR 68
sp  Q805D3  ANFC4_TAKRU	EKERRLDAVRSRL-----	RLLRDLR 68
tr  C1BX15  C1BX15_ESOLU	ERERRLDTVRSRV-----	RLLRDLR 68
tr  D2KXA5  D2KXA5_ANGJA	ERERNLESARS-----	RLLRDLR 66
tr  Q1XGY7  Q1XGY7_9ACTI	EQESGSENLRSA-----	RLLRDLR 68
sp  P40756  ANFD_RANCA	ERGERSIDPKTRA-----	RLLRDIR 68
sp  P0C7P5  BNP_TRIFL	-----PAAPHRLPKSKG-----ASATS-AASRPMRDLR 130	
sp  P0C7P6  BNP_TRIGA	-----PAAPQRLPKRKG-----ASATS-AASRSMRDLR 147	
sp  Q6LEMS5  BNP1_BOTJA	VPSAGAEVGRSGSK-APAAPHRLSKSKG-----AAAT-----RPMRDLR 193	
sp  Q9PW56  BNP2_BOTJA	VPSAGAEVGRSGSK-APAAPHRLSKSKG-----AAATS-AASRPMRDLR 201	
sp  P68515  BNP_BOTIN	VPSAGAEVGRSGSK-APAAPHRLSKSKG-----AAATS-AASRPMRDLR 201	
sp  Q90Y12  BNP_CRODU	TPPAGPDGGPRGSKAAAAAPQRLSKSKG-----ASATS-AASR---DLR 118	
sp  Q2PE51  BNP_CRODO	TPPAGPDGGPRGSKAAAAAPQRLSKSKG-----ASATS-AASR---DLR 118	
sp  B0VXV8  BNP_SISCA	TPPAGPDVGPRGSK-AAAAPQRLSKSKG-----ASATS-TASRPMRDLR 137	
sp  Q27J49  BNP_LACMU	TPPAGPDVGPRGSK-APAAPHRLPKSKG-----ASATS-AASRPMRDLR 178	
sp  P01021  BNP_AGKHA	PAAAGADGGRSGSK-APAALHRLSKSKG-----ASATSASASRPMRDLR 199	
sp  Q09GK2  VNP_PHIOL	PAAGGGGGGRSGSKAANAAPTAPEKSKGG-----AAAAAAAAALMRDLR 100	
tr  D1MZV3  D1MZV3_RHATT	PAAGGGGG--RSGSKTANAAPTAPEKSKGG-----AAASAASRLLRDLR 99	
tr  Q7T1M4  Q7T1M4_BOTJR	VPSAGAEGRAQRLEGARCTPSAVEEQRGG-----GDLGGVAADAGLAPRR 155	
tr  Q402A2  Q402A2_PETMA	VTATGGVSGGVSGVLLGGTPWVGPGRHQ-----RPSRGLA 65	
tr  Q402A3  Q402A3_LAMJA	ATVTR-HGGGGGGGGDDGTSWELPQGPONG-----PPRSSRGLA 97	
tr  Q402A1  Q402A1_9PETR	-----AEGG-----PWERGSGPQ-----RSSRGIG 83	
sp  P21805  ANFC_CHICK	-----	
tr  A9CDT6  A9CDT6_CHICK	QHPLVSEERDREQDGSIPVGAF-----DQEDAEFQWTRNTRD----- 76	
sp  P20968  ANFC_RANCA	ERSFGSDEADQQL-----VPTDSL-----DQLDPELQWNKNRLE----- 75	
sp  P55208  ANFC_TRISC	GHYLPSPDELNNEAQEMSPAASLPEFNADQSDLELPWDRESRE----- 82	
sp  P23259  ANFC_SCYCA	GHYLPSPDELNNEAEEMSPAASLPELNADQSDLELPWERESRE----- 61	
sp  P41319  ANFC_SQUAC	GHFN-SEELNNEAQEIISPAASLPELNDTQSDLELPWDRESRE----- 81	
tr  Q2MH72  Q2MH72_9CHON	GQYFTTEDLNNEAPEIPPAASLPELNDTQSDLELPWDRESRE----- 82	
tr  Q2MH73  Q2MH73_9CHON	GQYFTTEDLNNEAPEIPPAASLPELNDTQSDLELPWDRESRE----- 82	
tr  Q2MH71  Q2MH71_9CHON	GQYFTSEDLNNEAPEIPPAASLPELNDTQSDLELPWDRESRE----- 82	
tr  Q2MH74  Q2MH74_DASAK	GQYFTSEDLNNEAPEIPPAASLPELNDTQSDLELPWDRESRE----- 82	
tr  Q2PF87  Q2PF87_CALMI	GQYFTSEDLNNEAPEIPPAASLPELNDTQSDLELPWDRESRE----- 82	
sp  Q80018  ANFC3_ORYLA	GPYLSSEDSDMEEAEASRAGTLRDLNLDQADM DLLWDRDARD----- 85	
tr  Q4ADV1  Q4ADV1_ORYLA	---QDQLSSTEHPEE---DRLD-----RTREEPQLGG----- 58	
sp  Q805D4  ANFC3_TAKRU	---QDQLSSTEHPEE---DRLD-----RTREEPQLGG----- 58	
tr  C0H7B0  C0H7B0_SALSA	EKVQEQQEEVQEQQEEVQEQQE-----EQQEVEQERGRCTGDVLLRAQLD 104	
tr  C1BWD1  C1BWD1_ESOLU	VPYYASDEMVGDKDLNTEN-----IAEEVPPWDSEDAR----- 75	
tr  D2KXA3  D2KXA3_ANGJA	VPYYGSEEVDDKDLNTEKEA-----FTGEVVQPWLTEDR----- 74	
tr  B3DJJ2  B3DJJ2_DANRE	APYLDSEEAEVQGKEMNAENAA-----FTKASLHSWDPNSR----- 75	
tr  Q1XGY8  Q1XGY8_9ACTI	KPFEESQESVMEQKDATAEKS-----LDERAEQLWESDAR----- 75	
sp  Q8AXR2  ANFC2_ONCMY	HPyVSDDDTRGGLDVS A E I A T-----DDSEADIPWNHNFRLD----- 77	
sp  Q8AXR3  ANFC1_ONCMY	DLT-----LDDLENLVLSSQPEEPSS-----AFTSGVKIAEYPKW 77	
tr  C1BKS8  C1BKS8_OSMMO	DLT-----LDDLENLVLSSQPEEPSS-----AFTSGVKVAEYPKW 77	
tr  Q805E7  Q805E7_OREMO	DLLN-----LDDLENAGSNQPEEPS-----DYANGVKVAEYPKW 77	
tr  C3KH23  C3KH23_ANOFI	DLLT-----LDDLENMLNSQAAEQS-----TLSSGSKAVEYPKW 77	
sp  Q8AYR6  ANFC1_ORYLA	DLLT-----LDDLENLLNSQPEEQS-----TFSSGVKAAEYPKW 77	
sp  Q805D6  ANFC1_TAKRU	DLLT-----LDDLENMLNSHPEDQS-----NLSS-SKADEYPKW 76	
sp  P18145  ANFC_ANGJA	DLLN-----IDDLENQTGDQLESPQ-----PLSSGLKVAEYPKW 77	
tr  Q1XGY9  Q1XGY9_9ACTI	NLQSGPVNIHNVSSQYEDPQ-----PWADVSSVSKEQIW 86	
tr  A9CDT5  A9CDT5_CHICK	ESMPAEEDMKEGSSESPQLLSP-----PLPLLPSRAR--- 76	

## FIG. 15F

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp |Q8AYR5 |ANFC2\_ORYLA  
 sp |Q805D5 |ANFC2\_TAKRU  
 sp |Q76KW6 |ANFC\_ACITR  
 sp |Q61839 |ANFC\_MOUSE  
 tr |Q544K5 |Q544K5\_MOUSE  
 tr |Q8VHG9 |Q8VHG9\_NOTAL  
 sp |P55207 |ANFC\_RAT  
 sp |P55206 |ANFC\_BOVIN  
 sp |P56283 |ANFC\_SHEEP  
 sp |P18104 |ANFC\_PIG  
 sp |P23582 |ANFC\_HUMAN  
 sp |P84715 |ANF39\_ORNAN  
 tr |Q9QZ96 |Q9QZ96\_CAVPO  
 sp |Q80017 |ANFC4\_ORYLA  
 sp |Q805D3 |ANFC4\_TAKRU  
 tr |C1BX15 |C1BX15\_ESOLU  
 tr |D2KXA5 |D2KXA5\_ANGJA  
 tr |Q1XGY7 |Q1XGY7\_9ACTI  
 sp |P40756 |ANFD\_RANCA  
 sp |P0C7P5 |BNP\_TRIFL  
 sp |P0C7P6 |BNP\_TRIGA  
 sp |Q6LEMS5 |BNP1\_BOTJA  
 sp |Q9PW56 |BNP2\_BOTJA  
 sp |P68515 |BNP\_BOTIN  
 sp |Q90Y12 |BNP\_CRODU  
 sp |Q2PE51 |BNP\_CRODO  
 sp |B0VXV8 |BNP\_SISCA  
 sp |Q27J49 |BNP\_LACMU  
 sp |P01021 |BNP\_AGKHA  
 sp |Q09GK2 |VNP\_PHIOL  
 tr |D1MZV3 |D1MZV3\_RHATT  
 tr |Q7T1M4 |Q7T1M4\_BOTJR  
 tr |Q402A2 |Q402A2\_PETMA  
 tr |Q402A3 |Q402A3\_LAMJA  
 tr |Q402A1 |Q402A1\_9PETR  
 sp |P21805 |ANFC\_CHICK  
 tr |A9CDT6 |A9CDT6\_CHICK  
 sp |P20968 |ANFC\_RANCA  
 sp |P55208 |ANFC\_TRISC  
 sp |P23259 |ANFC\_SCYCA  
 sp |P41319 |ANFC\_SQUAC  
 tr |Q2MH72 |Q2MH72\_9CHON  
 tr |Q2MH73 |Q2MH73\_9CHON  
 tr |Q2MH71 |Q2MH71\_9CHON  
 tr |Q2MH74 |Q2MH74\_DASAK  
 tr |Q2PF87 |Q2PF87\_CALMI  
 sp |Q80018 |ANFC3\_ORYLA  
 tr |Q4ADV1 |Q4ADV1\_ORYLA  
 sp |Q805D4 |ANFC3\_TAKRU  
 tr |C0H7B0 |C0H7B0\_SALSA  
 tr |C1BWD1 |C1BWD1\_ESOLU  
 tr |D2KXA3 |D2KXA3\_ANGJA  
 tr |B3DJJ2 |B3DJJ2\_DANRE  
 tr |Q1XGY8 |Q1XGY8\_9ACTI  
 sp |Q8AXR2 |ANFC2\_ONCMY  
 sp |Q8AXR3 |ANFC1\_ONCMY  
 tr |C1BKS8 |C1BKS8\_OSMMO  
 tr |Q805E7 |Q805E7\_OREMO  
 tr |C3KH23 |C3KH23\_ANOFI  
 sp |Q8AYR6 |ANFC1\_ORYLA  
 sp |Q805D6 |ANFC1\_TAKRU  
 sp |P18145 |ANFC\_ANGJA  
 tr |Q1XGY9 |Q1XGY9\_9ACTI  
 tr |A9CDT5 |A9CDT5\_CHICK

-----GDRGTASQILRSFLRQRE-----KTRRWG--RKPMVAG- 107  
 -----GSEG-GSRFLLDLQQQS-----KTTRRG--RSSMVGG- 111  
 LEAEPPAENKPRRLLKDFMSSR-----KMFGRG--TKKMQQG- 131  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWARLLHEHPN-----ARKNKG--GNKKGLS- 128  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWTRLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWTRLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 VDTKSR---AAWARLLHEHPN-----ARKYKG--GNKKGLS- 107  
 LDTRSRSGSRGVWTRLLHDHPN-----PRKYKP--ANKKGLS- 102  
 -----N-----ARKYKG--GNKKGLS- 14  
 VDTRAR---GMWARRLNDQPA-----SRRHKS--GSKKGGST 100  
 MDTRAR---GVWARRLNDQPV-----PRRHKT--GIKKGGS- 99  
 MDTRAK---GMWARRLNDQPN-----ARRHKQ--NSKKGTV- 99  
 LNTRAR---GMWSRIMNDQPA-----SRKQKT--GVKKGAST 98  
 LDIRAK---AAWARLLNDHPN-----PRKSKG--INKKGLS- 99  
 ADTRSR---AAWTRLLNEHPN-----SRKIKG--INKKGTs- 99  
 TDGKQERQKWG--RMVQPDHHAAPGGGGGGGG-ARRMKG--LAKKAMG- 174  
 ADGKQARQKWG--RMVQPDHHAAPGGGGGGGG-ARRLKG--LAKKAVG- 191  
 PDGKQARQNWG--RMAHHDDHAAAGGGGGGGGG-ARRLKG--LAKKGAA- 237  
 PDGKQARQNWG--RMVHDDHAAVGGGGGGGGGARRLKG--LAKKGAA- 246  
 PDGKQARQNWG--RMVHDDHAAVGGGGGGGGGARRLKG--LAKKGAA- 246  
 TDGKQARQNWG--RLVSPDHSAAGGGGGGGGG-ARRLKG--LAKKRAG- 162  
 TDGKQARQNWG--RLVSPDHSAAGGGGGGGGG-ARRLKG--LAKKRAG- 162  
 TDGKQARQNWG--RLMLNPDHHSAPGGGGGGGGGARRLKG--LAKKRAG- 182  
 TDGKQARQNWG--RMMNPDDHVGCCCCGGGGGGGARRLKG--LVKKGVA- 220  
 TDGKQARQNWA--RMVNPDDHVGCCCCGGGGGGGARRLKG--LVKKGVA- 244  
 PDSKQARAAWG--RMVHPEHHAGGGGGGGGGGGGASRRLKG--VAKKGLG- 145  
 PDGKQSRRAAWG--RMVHPEHHAGGGGGGGGGGG-ARRLKG--LPKKGLG- 142  
 QAGAAELGPDPGAPRPPRPPRSRRAAAAAAERGARRLKG--LAKKGAA- 202  
 EGGSQVSG--GVWQRLFNDFVSN-----QRRFRG--RTKKGKG- 99  
 EGGSQVSG--GVWQRLFNDFVSN-----QRRFRG--RTKKGKG- 131  
 EGGSQVSG--EVWQRLFNDFVSN-----QRRFRG--RTKKGKG- 116  
 -----GLS- 3  
 QPASTSTADSDVQRILSDLGL-----PQRYQN--RSKKGLS- 111  
 QGDSPHVNEMTLQQLNDPVGT-----SRRYRQ--RNKKGYS- 110  
 IGGRPFQEAFLARLLKDLSSN-----PLRFRG--RSKKGPS- 117  
 IGGRPFQEAFLARLLKDLSSN-----PLRFRG--RSKKGPS- 96  
 IGGRSFRQEALLARLQLDLSNN-----PLRFRG--RSKKGPS- 116  
 IASR-----PILARILKDLSSN-----PLRFRG--RSKKGPS- 112  
 IASR-----PILARILKDLSSN-----PLRFRG--RSKKGPS- 112  
 IASR-----PILARILKDLSSN-----PLRFRG--RSKKGPS- 112  
 IASR-----PILARILKDLNNI-----PLRFRG--RSKKGPS- 112  
 IGGRSFQHGDLLLRLKDLTIS-----PLRFRG--RSKKGPS- 120  
 SSSREAADESALTRLFADLLRT-----SKRSWG--RYKKGGM- 93  
 SSSREAADESALTRLFADLLRT-----SKRSWG--RYKKGGM- 93  
 SSTWALQKDDVLMRLFKDLLRT-----SKRSRS--RYKKGGL- 139  
 -NSALTGKEDVIARLLNDIMTT-----PKRSWS--RFKKGGL- 109  
 --SALTGKENAVARLLSDIMTT-----PKRSWG--RFKKGGM- 107  
 -DAALSSNENALVRLNLDILSS-----SKRSWS--RFKKGGL- 109  
 -NSALAGKYGMFERLLGDLLST-----SKRSWS--RFKKGDL- 109  
 -QHRQAAHSSRMLKLLKDLILTS-----SGRSWD--REKKSGL- 111  
 ADIP-AQGDSTWLRLKGTLAN-----QKRAVT--DRSRRGWN- 112  
 ADIP-AQGDSTWLRLKGTLAN-----QKRAVM--DRSRRGWN- 112  
 ADLPAAQEDSAWLRLKAALAN-----QKRAEP--DRSRRAWN- 113  
 ADAQTQPE--TPWLRLKGALAN-----QKRAEP--DRSRRGWN- 112  
 ADAQTQAE--TPWLRLKGAVAN-----QKRAEP--DRSRRGWN- 112  
 ADLQTQPE--TPWFRLKGALTN-----QKRAEP--DRSRRGWN- 112  
 AEAD-----TPWLRLRGALAN-----QKRAEP--DRSRRGWN- 107  
 VDVPSQND-NTWFRLLRGALAN-----RKRALP--DRAKRGWN- 112  
 GDEPPANE-NALYLLRRAAAN-----RTWISA--DRVKKAWS- 121  
 -----AAHPLLWRKALAS-----RKRALSGDWAWKAVP- 104

## FIG. 15G

## Multiple Alignment of C-Type Natriuretic Peptides from Various Species

sp |Q8AYR5 |ANFC2\_ORYLA  
 sp |Q805D5 |ANFC2\_TAKRU  
 sp |Q76KW6 |ANFC\_ACITR  
 sp |Q61839 |ANFC\_MOUSE  
 tr |Q544K5 |Q544K5\_MOUSE  
 tr |Q8VHG9 |Q8VHG9\_NOTAL  
 sp |P55207 |ANFC\_RAT  
 sp |P55206 |ANFC\_BOVIN  
 sp |P56283 |ANFC\_SHEEP  
 sp |P18104 |ANFC\_PIG  
 sp |P23582 |ANFC\_HUMAN  
 sp |P84715 |ANF39\_ORNAN  
 tr |Q9QZ96 |Q9QZ96\_CAVPO  
 sp |Q80017 |ANFC4\_ORYLA  
 sp |Q805D3 |ANFC4\_TAKRU  
 tr |C1BX15 |C1BX15\_ESOLU  
 tr |D2KXA5 |D2KXA5\_ANGJA  
 tr |Q1XGY7 |Q1XGY7\_9ACTI  
 sp |P40756 |ANFD\_RANCA  
 sp |P0C7P5 |BNP\_TRIFL  
 sp |P0C7P6 |BNP\_TRIGA  
 sp |Q6LEMS5 |BNP1\_BOTJA  
 sp |Q9PW56 |BNP2\_BOTJA  
 sp |P68515 |BNP\_BOTIN  
 sp |Q90Y12 |BNP\_CRODU  
 sp |Q2PE51 |BNP\_CRODO  
 sp |B0VXV8 |BNP\_SISCA  
 sp |Q27J49 |BNP\_LACMU  
 sp |P01021 |BNP\_AGKHA  
 sp |Q09GK2 |VNP\_PHIOL  
 tr |D1MZV3 |D1MZV3\_RHATT  
 tr |Q7T1M4 |Q7T1M4\_BOTJR  
 tr |Q402A2 |Q402A2\_PETMA  
 tr |Q402A3 |Q402A3\_LAMJA  
 tr |Q402A1 |Q402A1\_9PETR  
 sp |P21805 |ANFC\_CHICK  
 tr |A9CDT6 |A9CDT6\_CHICK  
 sp |P20968 |ANFC\_RANCA  
 sp |P55208 |ANFC\_TRISC  
 sp |P23259 |ANFC\_SCYCA  
 sp |P41319 |ANFC\_SQUAC  
 tr |Q2MH72 |Q2MH72\_9CHON  
 tr |Q2MH73 |Q2MH73\_9CHON  
 tr |Q2MH71 |Q2MH71\_9CHON  
 tr |Q2MH74 |Q2MH74\_DASAK  
 tr |Q2PF87 |Q2PF87\_CALMI  
 sp |Q80018 |ANFC3\_ORYLA  
 tr |Q4ADV1 |Q4ADV1\_ORYLA  
 sp |Q805D4 |ANFC3\_TAKRU  
 tr |C0H7B0 |C0H7B0\_SALSA  
 tr |C1BWD1 |C1BWD1\_ESOLU  
 tr |D2KXA3 |D2KXA3\_ANGJA  
 tr |B3DJJ2 |B3DJJ2\_DANRE  
 tr |Q1XGY8 |Q1XGY8\_9ACTI  
 sp |Q8AXR2 |ANFC2\_ONCMY  
 sp |Q8AXR3 |ANFC1\_ONCMY  
 tr |C1BKS8 |C1BKS8\_OSMMO  
 tr |Q805E7 |Q805E7\_OREMO  
 tr |C3KH23 |C3KH23\_ANOFI  
 sp |Q8AYR6 |ANFC1\_ORYLA  
 sp |Q805D6 |ANFC1\_TAKRU  
 sp |P18145 |ANFC\_ANGJA  
 tr |Q1XGY9 |Q1XGY9\_9ACTI  
 tr |A9CDT5 |A9CDT5\_CHICK

---GC~~CFGMKMDRIGSISGLGC~~ 126 (SEQ ID NO: 31)  
 ---RG~~CFGMKIDRIGSISGLGC~~ 130 (SEQ ID NO: 32)  
 ---RG~~CFGMKLDRIGSMSGLGC~~ 150 (SEQ ID NO: 33)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 34)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 35)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 147 (SEQ ID NO: 36)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 37)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 38)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 39)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 40)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 41)  
 ---KC~~CFGKLDRIGSTSGLGC~~ 121 (SEQ ID NO: 42)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 33 (SEQ ID NO: 43)  
 ---SRSC~~CFGHKMDRIGTISGMGC~~ 121 (SEQ ID NO: 44)  
 ---SRSC~~CFGHKMDRIGTISGMGC~~ 120 (SEQ ID NO: 45)  
 ---PRSG~~CFGQKLDRIGTLSGMGC~~ 121 (SEQ ID NO: 46)  
 PARGC~~CFGHKLDRISTLSGMGC~~ 120 (SEQ ID NO: 47)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 118 (SEQ ID NO: 48)  
 ---KC~~CFGKLDRIGAMSGLGC~~ 118 (SEQ ID NO: 49)  
 ---KC~~CFGHKLDRIGSTSGLGC~~ 193 (SEQ ID NO: 50)  
 ---KC~~CFGGLPDRIGSMSGMGC~~ 210 (SEQ ID NO: 51)  
 ---KC~~CFGKLDRIGTMSGLGC~~ 256 (SEQ ID NO: 52)  
 ---KC~~CFGGLKVRIGTMSGLGC~~ 265 (SEQ ID NO: 53)  
 ---KC~~CFGKLDRIGTMSGLGC~~ 265 (SEQ ID NO: 54)  
 ---NC~~CFGKLDRIGSMSGLGC~~ 181 (SEQ ID NO: 55)  
 ---NC~~CFGKLDRIGSMSGLGC~~ 181 (SEQ ID NO: 56)  
 ---SC~~CFGKLDRIGSMSGLGC~~ 201 (SEQ ID NO: 57)  
 ---DC~~CFGKLDRIGSMSGLGC~~ 239 (SEQ ID NO: 58)  
 ---KC~~CFGKLDRIGTMSGLGC~~ 263 (SEQ ID NO: 59)  
 ---KC~~CFGKLDRIGSMSGLGC~~ 164 (SEQ ID NO: 60)  
 ---SC~~CFGKLDRIGSMSGLGC~~ 161 (SEQ ID NO: 61)  
 ---KC~~CFGKLDRIGTMSGLGC~~ 221 (SEQ ID NO: 62)  
 ---CFGV~~KLDRIGSMSGLGC~~ 116 (SEQ ID NO: 63)  
 ---CFGV~~KLDRIGSMSGLGC~~ 148 (SEQ ID NO: 64)  
 ---CFGV~~KLDRIGSMSGLGC~~ 133 (SEQ ID NO: 65)  
 ---RSC~~CFGVKLDRIGSMSGLGC~~ 22 (SEQ ID NO: 66)  
 ---RSC~~CFGVKLDRIGSMSGLGC~~ 130 (SEQ ID NO: 67)  
 ---RGC~~CFGVKLDRIGAFSGLGC~~ 129 (SEQ ID NO: 68)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 136 (SEQ ID NO: 69)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 115 (SEQ ID NO: 70)  
 ---RSC~~CFGKLDRIGAMSGLGC~~ 135 (SEQ ID NO: 71)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 131 (SEQ ID NO: 72)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 131 (SEQ ID NO: 73)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 131 (SEQ ID NO: 74)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 131 (SEQ ID NO: 75)  
 ---RGC~~CFGVKLDRIGAMSGLGC~~ 139 (SEQ ID NO: 76)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 112 (SEQ ID NO: 77)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 112 (SEQ ID NO: 78)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 158 (SEQ ID NO: 79)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 128 (SEQ ID NO: 80)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 126 (SEQ ID NO: 81)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 128 (SEQ ID NO: 82)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 128 (SEQ ID NO: 83)  
 ---RSC~~CFGVRLERIGSFSGLGC~~ 130 (SEQ ID NO: 84)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 131 (SEQ ID NO: 85)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 131 (SEQ ID NO: 86)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 132 (SEQ ID NO: 87)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 131 (SEQ ID NO: 88)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 131 (SEQ ID NO: 89)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 131 (SEQ ID NO: 90)  
 ---RGC~~CFGKLDRIGSMSGLGC~~ 126 (SEQ ID NO: 91)  
 ---RGC~~CFGKLDRIGSLSGLGC~~ 131 (SEQ ID NO: 92)  
 ---KC~~CFGKLDRIGSISGLGC~~ 140 (SEQ ID NO: 93)  
 ---RGC~~CFGKLMDRIGAFSGLGC~~ 123 (SEQ ID NO: 94)

\*\*\* : \*\*\* : \*\*\* : \*\*\*  
 CFGXXXRIXXXSGXGC (SEQ ID NO: 95)

CONSENSUS SEQUENCE

FIG. 16

**Schematic Structure of Natriuretic Peptide**

## Schematic Structures of Exemplary Fc-NP or NP-Fc Fusion Proteins

FIG. 17A

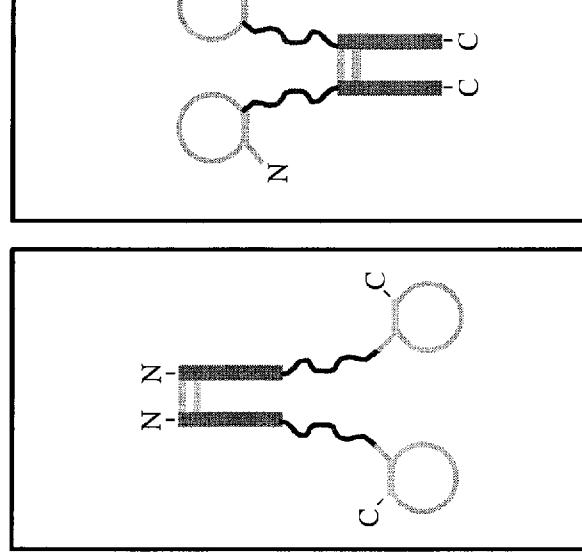


FIG. 17B

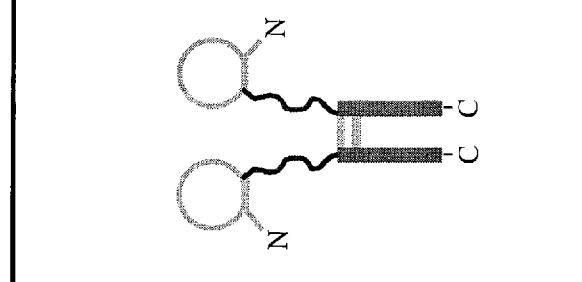


FIG. 17C

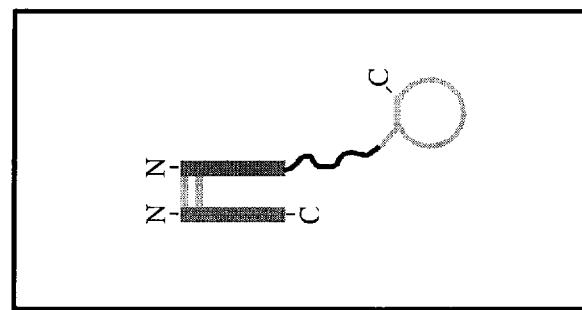


FIG. 17D

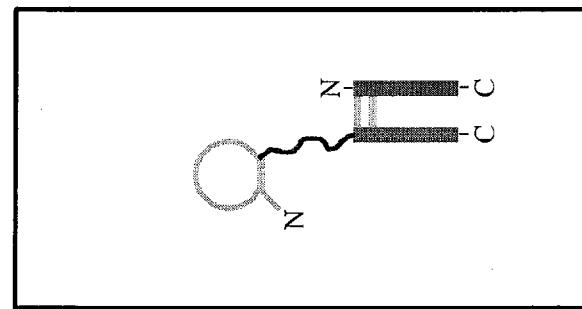
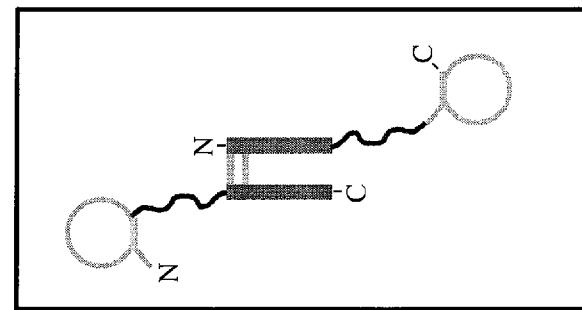


FIG. 17E



## FIG. 18A

## NC2st Protein Sequence (With Signal Sequence)

**Sequence:**

<u>10</u>	<u>20</u>	<u>30</u>	<u>40</u>	<u>50</u>
<u>MGVHECPAWL</u>	<u>WLLLSSLLSLW</u>	<u>PGAYAAASWSH</u>	<u>PQFEQSQQGG</u>	<u>GENLYFQGGGD</u>
<u>70</u>	<u>80</u>	<u>90</u>	<u>100</u>	<u>110</u>
<b>PEI<sub>2</sub>GGGPSVF</b>	<b>LFPPKPKD<sub>2</sub>TL</b>	<b>MISRTPEVTC</b>	<b>VVVDVSHEDP</b>	<b>EVKETNNWYVDG</b>
<u>130</u>	<u>140</u>	<u>150</u>	<u>160</u>	<u>170</u>
<b>REEQYNSTYR</b>	<b>VVS<sub>2</sub>VLT<sub>2</sub>VHLQ</b>	<b>DWLNGKEYKC</b>	<b>KVS<sub>2</sub>NKALPAP</b>	<b>IERTISKARG</b>
<u>190</u>	<u>200</u>	<u>210</u>	<u>220</u>	<u>230</u>
<b>PPSREEMTRN</b>	<b>QVS<sub>2</sub>L<sub>2</sub>T<sub>2</sub>LYRG</b>	<b>FYPSDIAE<sub>2</sub>W</b>	<b>ESNGQOPENNY</b>	<b>KTTTPPVLDSD</b>
<u>250</u>	<u>260</u>	<u>270</u>	<u>280</u>	<u>290</u>
<b>VDKSRWQ<sub>2</sub>GN</b>	<b>VFSCSVMHEA</b>	<b>LHNHYTQ<sub>2</sub>KSL</b>	<b>SLS<sub>2</sub>PGK<sub>2</sub>GGG</b>	<b>S<sub>2</sub>GGGS<sub>2</sub>GGGG</b>
<u>310</u>				<u>300</u>
<b>LKLD<sub>2</sub>RICSMS</b>	<b>GI<sub>2</sub>CC</b>			

Number of amino acids: 314  
Molecular weight of monomer: 34053.5

(SED ID NO: 501)

1-25 = Signal peptide  
26-27 = Linker  
28-35 = Strep-tag II  
36-41 = Linker  
42-47 = Tobacco etch virus protease cleavage sequence  
48-49 = Linker  
50-276 = Fc domain of human IgG1  
277-292 = Glycine-rich linker  
293-314 = C-type natriuretic peptide

FIG. 18B

## NC2st Protein Sequence (Without Signal Sequence)

FIG. 18C

## NC2<sup>st</sup> DNA Sequence

ATGGGCGTGCACGAGTGTCCCTGGCTGGCTGCTGAGCCCTGCTGGCCTGGCTCTGGCTGGCCCTACGCCCTCTGGAGC  
CACCCCCCAGTTGAGCAGCTGGGGGGAGGGAGGAGAACCTGTACTTCAAGGGGGAGAACCCACACCTGCCCCCTGTGATGATCAGCCGGACCCGAA  
GGTGAACCTGGGTGGACGGTGTCCCACGAGGACCCCTGAAGTTCAATTGGTACGTTGAGGGTGGACGGGGTGGCACAAAGCC  
AAGACCAAGCCCCGGAGGAACAGTACACAGCACCTACCGGGTGTCCGTGCTGACCCGTGCTGCACCCAGGACTGGCTGAACGGC  
AAAGAGTACAAGTGCAGGTCTCCAAACAAGGCCCTGCCTGCCCCCATCGAGAAACCATCAGCAAGGCCAAGGGCCAGCCCCAGAGAA  
CCCCCAGGGTACACCCCTGGGGAGGAATGACCAAGAACAGGTGTCCCTGACCTGGCTGGCTGGACAGCGGACGGCCCT  
AGCGATAATCGCCGTGGAGTGGAGCAACGGCAACGGCCAGCCTGAGAACAAACTACAAGAACCCCCCTGTCAGCGGACGGCCCT  
TCTCTGTACTTCCAAACTGACCCGTGGACAAGGGGGTGGCAGCAGGGCAACGTTGTCAGCTGCACCGGGTGTGATGCCAGGGGG  
CACAAACCAACTACACCCAGAAGTCCCTGAGCCTGTCCTGGCAAGGGGGAGGGAAAGTGGAGGGAGGAAGGGGGAGGGAGGG  
AGCGGGACTGAGCAAGGGCTGCTCGGCCTGAAGCTGGACGGATGGCAGCATGAGCGGGCTGGCTGA (SEQ ID NO:  
801)

## FIG. 19A

## NC2B Protein Sequence

NC2B (w/ sig. seq.)

NC2B (w/o sig. seq.)

D10-NC2 (w/ sig. seq.)

MGVHECPAWLWLLSLLSLWPGAYADDDDDDDDKTHTCPPCAPELLGGPSVFLFPPKPKDTLMISRTP  
VTCVVVDVSHEDEWKENWYDGEVHNAAKTRPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISKAKGOPREPQVYTLPSSREEMTKNOVSLTICLVKGFYPSDIAVEWESNGOPENNYKTTPVVLDSDGSE  
FLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGGSGGGGGSGGGGGGGGGGGGGGGGG  
GSMSCIGC (SEQ ID NO: 607)

**FIG. 19B**  
**NC2B DNA Sequ**

## FIG. 20A

## NC2B-22, NC2B-28, and NC2B-34 Sequences

NC2B-22  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 505)

NC2B-28  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 506)

NC2B-28  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 507)

NC2B-34  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 508)

NC2B-34  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 509)

NC2B-34  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFNVWYDGEVHNAAKTKPREEQYNSTYRVS  
VSLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
SRWQQGNVFS  
CSVMHEALHNHYTQKSLSLSPGKGGGGGGGGGGGGGGGG  
GIGC (SEQ ID NO: 510)

FIG. 20B  
NC2B-22 DNA Sequence

FIG. 20C  
NC2B-28 DNA Sequence

FIG. 20D  
NC2B-34 DNA Sequence

## FIG. 21

## NC2 Variants

NC2-KGANQK  
(w/o sig. seq.)

MGVHECPAWLLLSSLWPGAYADKTHTCPPCAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEVHN~~AK~~TKPREEQYNSTYRVS~~V~~SLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPOVYTLPPSREEMTKNQVS~~L~~TCLVKGFYPSDI~~A~~VEWESENQOPENNYKTT~~P~~PD~~S~~SDGSEFLYSKLTVDK  
SRWQ~~Q~~GNVFECSVMHEALHNHYTQ~~K~~SLSPGKGGGGSGGGSKGAN~~K~~GLSKGCEGLKLDRTIGMSGI~~GC~~  
(SEQ ID NO: 511)

NC2-KGANQK  
(w/o sig. seq.)

DKTHTCPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHDEPKFNWYVDGVEVHN~~AK~~TKP  
EQYNSTYRVS~~V~~SLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPOVYTLPPSREEMTKNQVS~~L~~T  
CLVKGFYPSDI~~A~~VEWESENQOPENNYKTT~~P~~PD~~S~~SDGSEFLYSKLTVDKSRWQ~~Q~~GNVFECSVMHEALHNHYTQ~~K~~  
SI~~S~~SLSPGKGGGGSGGGSKGAN~~K~~GLSKGCEGLKLDRTIGMSGI~~GC~~ (SEQ ID NO: 512)

NC2-KGANQK  
(w/o sig. seq.)

MGVHECPAWLLLSSLWPGAYADKTHTCPPCAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEVHN~~AK~~TKPREEQYNSTYRVS~~V~~SLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPOVYTLPPSREEMTKNQVS~~L~~TCLVKGFYPSDI~~A~~VEWESENQOPENNYKTT~~P~~PD~~S~~SDGSEFLYSKLTVDK  
SRWQ~~Q~~GNVFECSVMHEALHNHYTQ~~K~~SLSPGKGGGGSGGGSKGAN~~K~~GLSKGCEGLKLDRTIGMSGI~~GC~~  
(SEQ ID NO: 513)

NC2-KGANQK  
(w/o sig. seq.)

DKTHTCPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHDEPKFNWYVDGVEVHN~~AK~~TKP  
EQYNSTYRVS~~V~~SLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPOVYTLPPSREEMTKNQVS~~L~~T  
CLVKGFYPSDI~~A~~VEWESENQOPENNYKTT~~P~~PD~~S~~SDGSEFLYSKLTVDKSRWQ~~Q~~GNVFECSVMHEALHNHYTQ~~K~~  
SI~~S~~SLSPGKGGGGSGGGSKGAN~~K~~GLSKGCEGLKLDRTIGMSGI~~GC~~ (SEQ ID NO: 514)

## FIG. 22

## NC2 Variants

NC2-CNP53mut2  
(w/o sig. seq.)

MGVHECPAMWLILLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMIISRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEVHNAAKTKPREEYQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIASKAG  
QPREPQVYTLPPSRREEMTKRNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKRTTPVLDSDGSEFLYSKLTVDK  
SRWQOGNVFSCCSVMEALHNHYTQKSLSLSPGKGGDLQVDTQSQAAWAQLLQEHPNAQQYKGANKKGLSKGCG  
FGIKLDRIGSMSGLGC (SEQ ID NO: 515)

NC2-CNP53mut2  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMIISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPRE  
EQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIASKAGQPREPQVYTLPPSREEMTKKNQVSLT  
CLVKGFYPSDIAVEWESNGQOPENNYKRTTPVLDSDGSEFLYSKLTVDKSRWQOGNTFSCSVMEALHNHYTQK  
SISLSPGKGGDLQVDTQSQAAWAQLLQEHPNAQQYKGANKKGLSKGCG (SEQ ID NO: 516)

## FIG. 23

## Fc-CNP53 Constructs

Fc-CNP53-A  
(w/o sig. seq.)

MGVHECPAW<sub>1</sub>W<sub>2</sub>W<sub>3</sub>W<sub>4</sub>W<sub>5</sub>W<sub>6</sub>W<sub>7</sub>W<sub>8</sub>W<sub>9</sub>W<sub>10</sub>W<sub>11</sub>W<sub>12</sub>W<sub>13</sub>W<sub>14</sub>W<sub>15</sub>W<sub>16</sub>W<sub>17</sub>W<sub>18</sub>W<sub>19</sub>W<sub>20</sub>W<sub>21</sub>W<sub>22</sub>W<sub>23</sub>W<sub>24</sub>W<sub>25</sub>W<sub>26</sub>W<sub>27</sub>W<sub>28</sub>W<sub>29</sub>W<sub>30</sub>W<sub>31</sub>W<sub>32</sub>W<sub>33</sub>W<sub>34</sub>W<sub>35</sub>W<sub>36</sub>W<sub>37</sub>W<sub>38</sub>W<sub>39</sub>W<sub>40</sub>W<sub>41</sub>W<sub>42</sub>W<sub>43</sub>W<sub>44</sub>W<sub>45</sub>W<sub>46</sub>W<sub>47</sub>W<sub>48</sub>W<sub>49</sub>W<sub>50</sub>W<sub>51</sub>W<sub>52</sub>W<sub>53</sub>W<sub>54</sub>W<sub>55</sub>W<sub>56</sub>W<sub>57</sub>W<sub>58</sub>W<sub>59</sub>W<sub>60</sub>W<sub>61</sub>W<sub>62</sub>W<sub>63</sub>W<sub>64</sub>W<sub>65</sub>W<sub>66</sub>W<sub>67</sub>W<sub>68</sub>W<sub>69</sub>W<sub>70</sub>W<sub>71</sub>W<sub>72</sub>W<sub>73</sub>W<sub>74</sub>W<sub>75</sub>W<sub>76</sub>W<sub>77</sub>W<sub>78</sub>W<sub>79</sub>W<sub>80</sub>W<sub>81</sub>W<sub>82</sub>W<sub>83</sub>W<sub>84</sub>W<sub>85</sub>W<sub>86</sub>W<sub>87</sub>W<sub>88</sub>W<sub>89</sub>W<sub>90</sub>W<sub>91</sub>W<sub>92</sub>W<sub>93</sub>W<sub>94</sub>W<sub>95</sub>W<sub>96</sub>W<sub>97</sub>W<sub>98</sub>W<sub>99</sub>W<sub>100</sub>W<sub>101</sub>W<sub>102</sub>W<sub>103</sub>W<sub>104</sub>W<sub>105</sub>W<sub>106</sub>W<sub>107</sub>W<sub>108</sub>W<sub>109</sub>W<sub>110</sub>W<sub>111</sub>W<sub>112</sub>W<sub>113</sub>W<sub>114</sub>W<sub>115</sub>W<sub>116</sub>W<sub>117</sub>W<sub>118</sub>W<sub>119</sub>W<sub>120</sub>W<sub>121</sub>W<sub>122</sub>W<sub>123</sub>W<sub>124</sub>W<sub>125</sub>W<sub>126</sub>W<sub>127</sub>W<sub>128</sub>W<sub>129</sub>W<sub>130</sub>W<sub>131</sub>W<sub>132</sub>W<sub>133</sub>W<sub>134</sub>W<sub>135</sub>W<sub>136</sub>W<sub>137</sub>W<sub>138</sub>W<sub>139</sub>W<sub>140</sub>W<sub>141</sub>W<sub>142</sub>W<sub>143</sub>W<sub>144</sub>W<sub>145</sub>W<sub>146</sub>W<sub>147</sub>W<sub>148</sub>W<sub>149</sub>W<sub>150</sub>W<sub>151</sub>W<sub>152</sub>W<sub>153</sub>W<sub>154</sub>W<sub>155</sub>W<sub>156</sub>W<sub>157</sub>W<sub>158</sub>W<sub>159</sub>W<sub>160</sub>W<sub>161</sub>W<sub>162</sub>W<sub>163</sub>W<sub>164</sub>W<sub>165</sub>W<sub>166</sub>W<sub>167</sub>W<sub>168</sub>W<sub>169</sub>W<sub>170</sub>W<sub>171</sub>W<sub>172</sub>W<sub>173</sub>W<sub>174</sub>W<sub>175</sub>W<sub>176</sub>W<sub>177</sub>W<sub>178</sub>W<sub>179</sub>W<sub>180</sub>W<sub>181</sub>W<sub>182</sub>W<sub>183</sub>W<sub>184</sub>W<sub>185</sub>W<sub>186</sub>W<sub>187</sub>W<sub>188</sub>W<sub>189</sub>W<sub>190</sub>W<sub>191</sub>W<sub>192</sub>W<sub>193</sub>W<sub>194</sub>W<sub>195</sub>W<sub>196</sub>W<sub>197</sub>W<sub>198</sub>W<sub>199</sub>W<sub>200</sub>W<sub>201</sub>W<sub>202</sub>W<sub>203</sub>W<sub>204</sub>W<sub>205</sub>W<sub>206</sub>W<sub>207</sub>W<sub>208</sub>W<sub>209</sub>W<sub>210</sub>W<sub>211</sub>W<sub>212</sub>W<sub>213</sub>W<sub>214</sub>W<sub>215</sub>W<sub>216</sub>W<sub>217</sub>W<sub>218</sub>W<sub>219</sub>W<sub>220</sub>W<sub>221</sub>W<sub>222</sub>W<sub>223</sub>W<sub>224</sub>W<sub>225</sub>W<sub>226</sub>W<sub>227</sub>W<sub>228</sub>W<sub>229</sub>W<sub>230</sub>W<sub>231</sub>W<sub>232</sub>W<sub>233</sub>W<sub>234</sub>W<sub>235</sub>W<sub>236</sub>W<sub>237</sub>W<sub>238</sub>W<sub>239</sub>W<sub>240</sub>W<sub>241</sub>W<sub>242</sub>W<sub>243</sub>W<sub>244</sub>W<sub>245</sub>W<sub>246</sub>W<sub>247</sub>W<sub>248</sub>W<sub>249</sub>W<sub>250</sub>W<sub>251</sub>W<sub>252</sub>W<sub>253</sub>W<sub>254</sub>W<sub>255</sub>W<sub>256</sub>W<sub>257</sub>W<sub>258</sub>W<sub>259</sub>W<sub>260</sub>W<sub>261</sub>W<sub>262</sub>W<sub>263</sub>W<sub>264</sub>W<sub>265</sub>W<sub>266</sub>W<sub>267</sub>W<sub>268</sub>W<sub>269</sub>W<sub>270</sub>W<sub>271</sub>W<sub>272</sub>W<sub>273</sub>W<sub>274</sub>W<sub>275</sub>W<sub>276</sub>W<sub>277</sub>W<sub>278</sub>W<sub>279</sub>W<sub>280</sub>W<sub>281</sub>W<sub>282</sub>W<sub>283</sub>W<sub>284</sub>W<sub>285</sub>W<sub>286</sub>W<sub>287</sub>W<sub>288</sub>W<sub>289</sub>W<sub>290</sub>W<sub>291</sub>W<sub>292</sub>W<sub>293</sub>W<sub>294</sub>W<sub>295</sub>W<sub>296</sub>W<sub>297</sub>W<sub>298</sub>W<sub>299</sub>W<sub>300</sub>W<sub>301</sub>W<sub>302</sub>W<sub>303</sub>W<sub>304</sub>W<sub>305</sub>W<sub>306</sub>W<sub>307</sub>W<sub>308</sub>W<sub>309</sub>W<sub>310</sub>W<sub>311</sub>W<sub>312</sub>W<sub>313</sub>W<sub>314</sub>W<sub>315</sub>W<sub>316</sub>W<sub>317</sub>W<sub>318</sub>W<sub>319</sub>W<sub>320</sub>W<sub>321</sub>W<sub>322</sub>W<sub>323</sub>W<sub>324</sub>W<sub>325</sub>W<sub>326</sub>W<sub>327</sub>W<sub>328</sub>W<sub>329</sub>W<sub>330</sub>W<sub>331</sub>W<sub>332</sub>W<sub>333</sub>W<sub>334</sub>W<sub>335</sub>W<sub>336</sub>W<sub>337</sub>W<sub>338</sub>W<sub>339</sub>W<sub>340</sub>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>674</sub>W<sub>675</sub>W<sub>676</sub>W<sub>677</sub>W<sub>678</sub>W<sub>679</sub>W<sub>680</sub>W<sub>681</sub>W<sub>682</sub>W<sub>683</sub>W<sub>684</sub>W<sub>685</sub>W<sub>686</sub>W<sub>687</sub>W<sub>688</sub>W<sub>689</sub>W<sub>690</sub>W<sub>691</sub>W<sub>692</sub>W<sub>693</sub>W<sub>694</sub>W<sub>695</sub>W<sub>696</sub>W<sub>697</sub>W<sub>698</sub>W<sub>699</sub>W<sub>700</sub>W<sub>701</sub>W<sub>702</sub>W<sub>703</sub>W<sub>704</sub>W<sub>705</sub>W<sub>706</sub>W<sub>707</sub>W<sub>708</sub>W<sub>709</sub>W<sub>710</sub>W<sub>711</sub>W<sub>712</sub>W<sub>713</sub>W<sub>714</sub>W<sub>715</sub>W<sub>716</sub>W<sub>717</sub>W<sub>718</sub>W<sub>719</sub>W<sub>720</sub>W<sub>721</sub>W<sub>722</sub>W<sub>723</sub>W<sub>724</sub>W<sub>725</sub>W<sub>726</sub>W<sub>727</sub>W<sub>728</sub>W<sub>729</sub>W<sub>730</sub>W<sub>731</sub>W<sub>732</sub>W<sub>733</sub>W<sub>734</sub>W<sub>735</sub>W<sub>736</sub>W<sub>737</sub>W<sub>738</sub>W<sub>739</sub>W<sub>740</sub>W<sub>741</sub>W<sub>742</sub>W<sub>743</sub>W<sub>744</sub>W<sub>745</sub>W<sub>746</sub>W<sub>747</sub>W<sub>748</sub>W<sub>749</sub>W<sub>750</sub>W<sub>751</sub>W<sub>752</sub>W<sub>753</sub>W<sub>754</sub>W<sub>755</sub>W<sub>756</sub>W<sub>757</sub>W<sub>758</sub>W<sub>759</sub>W<sub>760</sub>W<sub>761</sub>W<sub>762</sub>W<sub>763</sub>W<sub>764</sub>W<sub>765</sub>W<sub>766</sub>W<sub>767</sub>W<sub>768</sub>W<sub>769</sub>W<sub>770</sub>W<sub>771</sub>W<sub>772</sub>W<sub>773</sub>W<sub>774</sub>W<sub>775</sub>W<sub>776</sub>W<sub>777</sub>W<sub>778</sub>W<sub>779</sub>W<sub>780</sub>W<sub>781</sub>W<sub>782</sub>W<sub>783</sub>W<sub>784</sub>W<sub>785</sub>W<sub>786</sub>W<sub>787</sub>W<sub>788</sub>W<sub>789</sub>W<sub>790</sub>W<sub>791</sub>W<sub>792</sub>W<sub>793</sub>W<sub>794</sub>W<sub>795</sub>W<sub>796</sub>W<sub>797</sub>W<sub>798</sub>W<sub>799</sub>W<sub>800</sub>W<sub>801</sub>W<sub>802</sub>W<sub>803</sub>W<sub>804</sub>W<sub>805</sub>W<sub>806</sub>W<sub>807</sub>W<sub>808</sub>W<sub>809</sub>W<sub>810</sub>W<sub>811</sub>W<sub>812</sub>W<sub>813</sub>W<sub>814</sub>W<sub>815</sub>W<sub>816</sub>W<sub>817</sub>W<sub>818</sub>W<sub>819</sub>W<sub>820</sub>W<sub>821</sub>W<sub>822</sub>W<sub>823</sub>W<sub>824</sub>W<sub>825</sub>W<sub>826</sub>W<sub>827</sub>W<sub>828</sub>W<sub>829</sub>W<sub>830</sub>W<sub>831</sub>W<sub>832</sub>W<sub>833</sub>W<sub>834</sub>W<sub>835</sub>W<sub>836</sub>W<sub>837</sub>W<sub>838</sub>W<sub>839</sub>W<sub>840</sub>W<sub>841</sub>W<sub>842</sub>W<sub>843</sub>W<sub>844</sub>W<sub>845</sub>W<sub>846</sub>W<sub>847</sub>W<sub>848</sub>W<sub>849</sub>W<sub>850</sub>W<sub>851</sub>W<sub>852</sub>W<sub>853</sub>W<sub>854</sub>W<sub>855</sub>W<sub>856</sub>W<sub>857</sub>W<sub>858</sub>W<sub>859</sub>W<sub>860</sub>W<sub>861</sub>W<sub>862</sub>W<sub>863</sub>W<sub>864</sub>W<sub>865</sub>W<sub>866</sub>W<sub>867</sub>W<sub>868</sub>W<sub>869</sub>W<sub>870</sub>W<sub>871</sub>W<sub>872</sub>W<sub>873</sub>W<sub>874</sub>W<sub>875</sub>W<sub>876</sub>W<sub>877</sub>W<sub>878</sub>W<sub>879</sub>W<sub>880</sub>W<sub>881</sub>W<sub>882</sub>W<sub>883</sub>W<sub>884</sub>W<sub>885</sub>W<sub>886</sub>W<sub>887</sub>W<sub>888</sub>W<sub>889</sub>W<sub>890</sub>W<sub>891</sub>W<sub>892</sub>W<sub>893</sub>W<sub>894</sub>W<sub>895</sub>W<sub>896</sub>W<sub>897</sub>W<sub>898</sub>W<sub>899</sub>W<sub>900</sub>W<sub>901</sub>W<sub>902</sub>W<sub>903</sub>W<sub>904</sub>W<sub>905</sub>W<sub>906</sub>W<sub>907</sub>W<sub>908</sub>W<sub>909</sub>W<sub>910</sub>W<sub>911</sub>W<sub>912</sub>W<sub>913</sub>W<sub>914</sub>W<sub>915</sub>W<sub>916</sub>W<sub>917</sub>W<sub>918</sub>W<sub>919</sub>W<sub>920</sub>W<sub>921</sub>W<sub>922</sub>W<sub>923</sub>W<sub>924</sub>W<sub>925</sub>W<sub>926</sub>W<sub>927</sub>W<sub>928</sub>W<sub>929</sub>W<sub>930</sub>W<sub>931</sub>W<sub>932</sub>W<sub>933</sub>W<sub>934</sub>W<sub>935</sub>W<sub>936</sub>W<sub>937</sub>W<sub>938</sub>W<sub>939</sub>W<sub>940</sub>W<sub>941</sub>W<sub>942</sub>W<sub>943</sub>W<sub>944</sub>W<sub>945</sub>W<sub>946</sub>W<sub>947</sub>W<sub>948</sub>W<sub>949</sub>W<sub>950</sub>W<sub>951</sub>W<sub>952</sub>W<sub>953</sub>W<sub>954</sub>W<sub>955</sub>W<sub>956</sub>W<sub>957</sub>W<sub>958</sub>W<sub>959</sub>W<sub>960</sub>W<sub>961</sub>W<sub>962</sub>W<sub>963</sub>W<sub>964</sub>W<sub>965</sub>W<sub>966</sub>W<sub>967</sub>W<sub>968</sub>W<sub>969</sub>W<sub>970</sub>W<sub>971</sub>W<sub>972</sub>W<sub>973</sub>W<sub>974</sub>W<sub>975</sub>W<sub>976</sub>W<sub>977</sub>W<sub>978</sub>W<sub>979</sub>W<sub>980</sub>W<sub>981</sub>W<sub>982</sub>W<sub>983</sub>W<sub>984</sub>W<sub>985</sub>W<sub>986</sub>W<sub>987</sub>W<sub>988</sub>W<sub>989</sub>W<sub>990</sub>W<sub>991</sub>W<sub>992</sub>W<sub>993</sub>W<sub>994</sub>W<sub>995</sub>W<sub>996</sub>W<sub>997</sub>W<sub>998</sub>W<sub>999</sub>W<sub>1000</sub>W<sub>1001</sub>W<sub>1002</sub>W<sub>1003</sub>W<sub>1004</sub>W<sub>1005</sub>W<sub>1006</sub>W<sub>1007</sub>W<sub>1008</sub>W<sub>1009</sub>W<sub>1010</sub>W<sub>1011</sub>W<sub>1012</sub>W<sub>1013</sub>W<sub>1014</sub>W<sub>1015</sub>W<sub>1016</sub>W<sub>1017</sub>W<sub>1018</sub>W<sub>1019</sub>W<sub>1020</sub>W<sub>1021</sub>W<sub>1022</sub>W<sub>1023</sub>W<sub>1024</sub>W<sub>1025</sub>W<sub>1026</sub>W<sub>1027</sub>W<sub>1028</sub>W<sub>1029</sub>W<sub>1030</sub>W<sub>1031</sub>W<sub>1032</sub>W<sub>1033</sub>W<sub>1034</sub>W<sub>1035</sub>W<sub>1036</sub>W<sub>1037</sub>W<sub>1038</sub>W<sub>1039</sub>W<sub>1040</sub>W<sub>1041</sub>W<sub>1042</sub>W<sub>1043</sub>W<sub>1044</sub>W<sub>1045</sub>W<sub>1046</sub>W<sub>1047</sub>W<sub>1048</sub>W<sub>1049</sub>W<sub>1050</sub>W<sub>1051</sub>W<sub>1052</sub>W<sub>1053</sub>W<sub>1054</sub>W<sub>1055</sub>W<sub>1056</sub>W<sub>1057</sub>W<sub>1058</sub>W<sub>1059</sub>W<sub>1060</sub>W<sub>1061</sub>W<sub>1062</sub>W<sub>1063</sub>W<sub>1064</sub>W<sub>1065</sub>W<sub>1066</sub>W<sub>1067</sub>W<sub>1068</sub>W<sub>1069</sub>W<sub>1070</sub>W<sub>1071</sub>W<sub>1072</sub>W<sub>1073</sub>W<sub>1074</sub>W<sub>1075</sub>W<sub>1076</sub>W<sub>1077</sub>W<sub>1078</sub>W<sub>1079</sub>W<sub>1080</sub>W<sub>1081</sub>W<sub>1082</sub>W<sub>1083</sub>W<sub>1084</sub>W<sub>1085</sub>W<sub>1086</sub>W<sub>1087</sub>W<sub>1088</sub>W<sub>1089</sub>W<sub>1090</sub>W<sub>1091</sub>W<sub>1092</</sub>

FIG. 24

## NP Multiple Sequence Alignment and CDNP Constructs

Human ANP	SLRRSSCFGGRMDRIGAQSSGLGCNSFRY	(SEQ ID NO: 1)
Rat ANP	SLRRSSCFGGRIDRIGAQSSGLGCNSFRY	(SEQ ID NO: 96)
Urodilatin	TAPRSIIRRSSCFGGRMDRIGAQSSGLGCNSFRY	(SEQ ID NO: 2)
Human BNP	SPKMYQGSGCFGRKMDRISSSSGLGCKVILRRH	(SEQ ID NO: 3)
Rat BNP	SPKTMRDSGCFGRRLDRIGSLSGLCNVILRRY	(SEQ ID NO: 97)
Pig BNP	EVKYDPCFGHKIDRINHVSNLGCPSLRLDPRPINAPSTSAA	(SEQ ID NO: 98)
DNP	SDSKIGNGCFFGFPIDRIGSVSGLCNRIMQNPPKKFSGE	(SEQ ID NO: 99)
TNP-C	GLSKGCFFGLKLDRIGSMSGLGCPSLRLDPRPINAPSTSAA	(SEQ ID NO: 100)
CDNP	* * * : * * * * . * * * * : * * * * :	
CDNP-N1	GLSKGCFFGLKLDRIGSMSGLGCNSLRLDPRPINAPSTSAA	(SEQ ID NO: 101)
CDNP-G1	GLSKGCFFGLKLDRIGSMSGLGCSSLRLDPRPINAPSTSAA	(SEQ ID NO: 102)
CDNP-H1	GLSKGCFFGLKLDRIGSMSGLGCSSLRLDPRPINAPSTSAA	(SEQ ID NO: 103)
CDNP-K1	GLSKGCFFGLKLDRIGSMSGLGCSSLRLDPRPINAPSTSAA	(SEQ ID NO: 104)
CDNP-Z1 (Z=hydroxyproline)	GLSKGCFFGLKLDRIGSMSGLGCSSLRLDPRPINAPSTSAA	(SEQ ID NO: 105)
CDNP-S3	GLSKGCFFGLKLDRIGSMSGLGCSSLRLDPRPINAPSTSAA	(SEQ ID NO: 106)
CDNP-A4	GLSKGCFFGLKLDRIGSMSGLGCSSLADPRPINAPSTSAA	(SEQ ID NO: 107)
CDNP-A5	GLSKGCFFGLKLDRIGSMSGLGCSSLADPRPINAPSTSAA	(SEQ ID NO: 108)
CDNP-S3A4	GLSKGCFFGLKLDRIGSMSGLGCSSSADPRPINAPSTSAA	(SEQ ID NO: 109)
CDNP-A4A5	GLSKGCFFGLKLDRIGSMSGLGCSSSGLGCPSLAAAPRPINAPSTSAA	(SEQ ID NO: 110)
CDNP-S3A5	GLSKGCFFGLKLDRIGSMSGLGCSSSGLGCPSLAAAPRPINAPSTSAA	(SEQ ID NO: 111)
CDNP-(A17)S3A5	GLSKGCFFGLKLDRIGSASGLGCPSSSAAPRPINAPSTSAA	(SEQ ID NO: 112)
CDNP-S3A4A5	GLSKGCFFGLKLDRIGSMSGLGCPSSSAAPRPINAPSTSAA	(SEQ ID NO: 113)
CDNP-(A17)S3A4A5	GLSKGCFFGLKLDRIGSMSGLGCPSSSAAPRPINAPSTSAA	(SEQ ID NO: 114)
CDNP-S3A4A5R6	GLSKGCFFGLKLDRIGSMSGLGCPSSSAARRPINAPSTSAA	(SEQ ID NO: 115)
CDNP-S3A4A5S7	GLSKGCFFGLKLDRIGSMSGLGCPSSSAAPSINAPSTSAA	(SEQ ID NO: 116)
DNP TAIL	PSLRDPRPINAPSTSAA	(SEQ ID NO: 117)
CONSENSUS SEQUENCE FOR DNP C-TERMINAL TAIL	XXXXXXXPINAPSTSAA	(SEQ ID NO: 118)

FIG. 25A

## Exemplary Constructs Having N-terminal NP fused to C-terminal Fc Domain

CNP-16AA1linker-FC-His<sub>10</sub>  
(NC1)

CNP-6AA1inker-FC-His<sub>10</sub>  
(NC3)

CNP-6AAlinker-FC

GLSKGCFIGKLDRIGSMSGIGCGGGGGDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVD  
VSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK  
AKGQPREPQVTLPSSREEMTKNQVSILTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDGSFFFLYSKLT  
VDKRSRWQOGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 523)

CDNP-FC

GLSKGCFIGKLDIGSMMSGIGCPSSLRDPRNAPSTSADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLIMISRT  
PEVTCVWVVDVSHEDEVKFNWYDGVFDEVHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALP  
APIEKTISAKGOPREPQVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTTPVLDSDG  
SFFFLYSKLTVDKSRWQOGNVEFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 524)

GLSKGCFIGKLDIGSASGIGCPSSAAPRPNAPSTSADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLIMISRT  
PEVTCVWVVDVSHEDEVKFNWYDGVFDEVHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALP  
APIEKTISAKGOPREPQVYTLPPSREEMTKNOVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTTPVLDSDG  
SFFFLYSKLTVDKSRWQOGNVEFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 525)

CDNP-A17 Saa-FC

GLSKGCFGIKLDRIGSASGLCPSSAMPPRPNAPSTSADKTHTCPPCPAPELLGGPSVFLFPPKKDTLMISRT  
PEVTCVVDVSHEDPEVKFNWYVDGVEVHNATKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALP  
APIEKTISKAKGQPREQYVTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYTTPPVLDSDG  
SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 525)  
SFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 526)

## FIG. 25B NC1 DNA Sequ

FIG. 26

## Human CNP22 and Point Mutations at Position 17

CNP22	GLSKGCGFGLKLDRIGSMSGLGC	(SEQ ID NO: 4)
CNP-F17 (C15)	GLSKGCGFGLKLDRIGSF <u>S</u> GLGC	(SEQ ID NO: 119)
CNP-L17 (C16)	GLSKGCGFGLKLDRIG <u>S</u> LSGLGC	(SEQ ID NO: 120)
CNP-I17 (C17)	GLSKGCGFGLKLDRIG <u>I</u> SGLGC	(SEQ ID NO: 121)
CNP-T17 (C18)	GLSKGCGFGLKLDRIG <u>T</u> SGLGC	(SEQ ID NO: 122)
CNP-V17 (C19)	GLSKGCGFGLKLDRIG <u>V</u> SGLGC	(SEQ ID NO: 123)
CNP-A17 (C1)	GLSKGCGFGLKLDRIG <u>A</u> SGLGC	(SEQ ID NO: 124)
CNP-S17	GLSKGCGFGLKLDRIG <u>S</u> SSGLGC	(SEQ ID NO: 125)
CNP-E17	GLSKGCGFGLKLDRIG <u>E</u> SEGLGC	(SEQ ID NO: 156)
CNP-R17	GLSKGCGFGLKLDRIG <u>R</u> SGLGC	(SEQ ID NO: 157)
CNP-Y17	GLSKGCGFGLKLDRIG <u>Y</u> SGLGC	(SEQ ID NO: 158)
CONSENSUS:		
	GLSKGCGFGLKLDRIGSX <u>S</u> GLGC	(SEQ ID NO: 126)

FIG. 27 CNP variance

CNP22 (SEQ ID NO: 4)  
 CNP37 (SEQ ID NO: 127)  
 E6 PGNCNP37 (SEQ ID NO: 128)  
 C1 (E6) (SEQ ID NO: 129)  
 C2 (E6) (SEQ ID NO: 130)  
 C3 (E6) (SEQ ID NO: 131)  
 C4 (E6) (SEQ ID NO: 132)  
 C5 (E6) (SEQ ID NO: 133)  
 C6 (E6) (SEQ ID NO: 134)  
 C7 (E6) (SEQ ID NO: 135)  
 C8 (E6) (SEQ ID NO: 136)  
 C9 (E6) (SEQ ID NO: 137)  
 C10 (E6) (SEQ ID NO: 138)  
 C11 (E6) (SEQ ID NO: 139)  
 D6 CNP37 (SEQ ID NO: 140)  
 C1 (D6) (SEQ ID NO: 141)  
 C2 (D6) (SEQ ID NO: 142)  
 C3 (D6) (SEQ ID NO: 143)  
 C4 (D6) (SEQ ID NO: 144)  
 C5 (D6) (SEQ ID NO: 145)  
 C6 (D6) (SEQ ID NO: 146)  
 D6-14AA1 linker-CNP (SEQ ID NO: 147)  
 CNP-1-4AA1 linker-D6 (SEQ ID NO: 148)  
  
 GSSAAPRPNAPSTSAGLSSKGKGLSGMSGGLGC (SEQ ID NO: 149)  
 ASTSPANPRPAASSGGLSSKGKGLSGMSGGLGC (SEQ ID NO: 150)

## FIG. 28A

## Additional CNP Variants

GDLRVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1001)  
 PDLRVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1002)  
 MDLRVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1003)  
 DLRVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1004)  
 LRVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1005)  
 RVDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1006)  
 VDTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1007)  
 DTKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1008)  
 TKSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1009)  
 KSRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1010)  
 SRAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1011)  
 RAAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1012)  
 AAWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1013)  
 AWARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1014)  
 WARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1015)  
 ARLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1016)  
 RLLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1017)  
 LLQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1018)  
 LQEHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1019)  
 EHPNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1020)  
 PNARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1021)  
 NARKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1022)  
 YKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1023)  
 KGANKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1024)  
 RKYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1025)  
 KYKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1026)  
 YKGANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1027)  
 KGANKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1028)  
 GANKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1029)  
 ANKKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1030)  
 NKKKKGLSKGCFGGLKLDRIGSMSGLGC (SEQ ID NO: 1031)

**FIG. 28B**

KKGLSKGCFGGLKLDRIGSMSGLGC  
 (SEQ ID NO:1032)  
 RGLSKGCFGGLKLDRIGSMSGLGC  
 (SEQ ID NO:1033)  
 LSKGCFGGLKLDRIGSMSGLGC  
 (SEQ ID NO:1034)  
 SKGCFGGLKLDRIGSMSGLGC  
 (SEQ ID NO:1035)  
 KGCFCGLKLDRIGSMSGLGC  
 (SEQ ID NO:1036)  
 GCFGLKLDRIGSMSGLGC  
 (SEQ ID NO:1037)  
 SEQ ID NO:1038)  
 (SEQ ID NO:1039)  
 (SEQ ID NO:1040)  
 (SEQ ID NO:1041)  
 (SEQ ID NO:1042)  
 (SEQ ID NO:1043)  
 (SEQ ID NO:1044)  
 (SEQ ID NO:1045)  
 (SEQ ID NO:1046)  
 (SEQ ID NO:1047)  
 (SEQ ID NO:1048)  
 (SEQ ID NO:1049)  
 (SEQ ID NO:1050)  
 (SEQ ID NO:1051)  
 (SEQ ID NO:1052)  
 (SEQ ID NO:1053)  
 (SEQ ID NO:1054)  
 (SEQ ID NO:1055)  
 (SEQ ID NO:1056)  
 (SEQ ID NO:1057)  
 (SEQ ID NO:1058)  
 (SEQ ID NO:1059)  
 (SEQ ID NO:1060)  
 (SEQ ID NO:1061)  
 (SEQ ID NO:1062)

PQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 MQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANPKGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANQKGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANQQGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANKPKGLSKGCFGGLKLDRIGSMSGLGC  
 GQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 PGQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 MGQEHPNARKYKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 MGANRRGLSRGCFGGLKLDRIGSMSGLGC  
 GANRRGLSRGCFGGLKLDRIGSMSGLGC  
 PGANRRGLSRGCFGGLKLDRIGSMSGLGC  
 MGANRRGLSRGCFGGLKLDRIGSMSGLGC  
 GHKSEVAHRFKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 GHKSEVAHRFKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 PGHKSEVAHRFKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 MGHKSEVAHRFKGANKKGLSKGCFGGLKLDRIGSMSGLGC  
 RGLSRGCFGGLKLDRIGSMSGLGC  
 ERGLSRGCFGGLKLDRIGSMSGLGC  
 GANQQGLSKGCFGGLKLDRIGSMSGLGC  
 GANRRGLSKGCFGGLKLDRIGSMSGLGC  
 GANPRGLSKGCFGGLKLDRIGSMSGLGC  
 GANSSGLSKGCFGGLKLDRIGSMSGLGC

## FIG. 28C

(X: D-Phe)  
(X: 3-amino-2-phenylpropionic acid)  
(X: Cit)  
(X: [CH<sub>2</sub>NH] bond)

GANPRGLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1063)  
GANRRGLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1064)  
GANQQGLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1065)  
GANSSGLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1066)  
AAWARLLQEHPNAGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1067)  
AAWARLLQEHPNARGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1068)  
DLRVDTKSRRAAWARGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1069)  
GQEHPNARKYKGANPRGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1070)  
GQEHPNARKYKGANQKGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1071)  
GLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1072)  
ERGLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1073)  
RGLSKGCFGLRLLDRIGSMMSGI GLGC  
(SEQ ID NO:1074)  
GLSRGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1075)  
GLSKGCGXGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1076)  
GLSKGCGXGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1077)  
RLSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1078)  
ELSKGCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1079)  
GLSKRCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1080)  
GLSKQCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1081)  
GLSKSCFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1082)  
GLSKGCFGLKLDRISSMSGI GLGC  
(SEQ ID NO:1083)  
GLSKGCFGLKLDRINMSGI GLGC  
(SEQ ID NO:1084)  
GLSKGCFGLKLDRIRMSGI GLGC  
(SEQ ID NO:1085)  
GLSKGCFGLKLDRIIXMSGI GLGC  
(SEQ ID NO:1086)  
GLSKGCFGLKLDRIGSMSNLGC  
(SEQ ID NO:1087)  
GLSKGCFGLKLDRIGSMSRLGC  
(SEQ ID NO:1088)  
GLSKGCFGLKLDRIGSMSNLGC  
(SEQ ID NO:1089)  
GLSKGCFGLKLDRIGSMSGI GLSC  
(SEQ ID NO:1090)  
GLSKGCFGLKLDRIGSMSGI LTC  
(SEQ ID NO:1091)  
GLSKGCFGLKLDRIGSMSGI RLC  
(SEQ ID NO:1092)  
GLSKGCFXFGLKLDRIGSMSGI GLGC  
(SEQ ID NO:1093)

## FIG. 28D

(X: N-Me-Phe)  
 (X: t-Bu-Gly)  
 (X: NHCH<sub>2</sub>CH (Ph) CO)

(X<sub>1</sub>: [CH<sub>2</sub>NH] bond; X<sub>2</sub> and X<sub>3</sub>: N-Me-Leu)  
 (X<sub>1</sub> and X<sub>2</sub>: N-Me-Leu)  
 (X<sub>1</sub>, X<sub>2</sub>, and X<sub>3</sub>: N-Me-Leu)  
 (X<sub>1</sub> and X<sub>2</sub>: N-Me-Leu)

(X: Cit)  
 (X: t-Bu-Ala)

GLSKGCX<sub>1</sub>GLKLDRIGSMSGLGC (SEQ ID NO: 1094)  
 GLSKGCFX<sub>1</sub>KLKLDRIGSMSGLGC (SEQ ID NO: 1095)  
 GLSKGCX<sub>1</sub>GLKLDRIGSMSGLGC (SEQ ID NO: 1096)  
 GLSKGCFTLPPSGLKLDRIGSMSGLGC (SEQ ID NO: 1097)  
 GQPREPQVYTLPSSGLKLDRIGSMSGLGC (SEQ ID NO: 1098)  
 GQHKKDDNPNLPRGANPRLGLKLDRIGSMSGLGC (SEQ ID NO: 1099)  
 GHSHEQHPHGANQQGLSKGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1100)  
 GAHHPHEHDTHGANQQGLSKGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1101)  
 FGIPMDRIGRNPRGLSKGPKGLSKGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1102)  
 GKRTGQYKLGSKTGPGPKGLSKGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1103)  
 SPKMYQGSGCFGLKLDRIGSMSGLGCKVLRHH (SEQ ID NO: 1104)  
 GQPREPQVYTGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1105)  
 GVPQVSTSTGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1106)  
 GQPSSSSQSTGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1107)  
 GSTGQWHSESGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1108)  
 GQTHSSGTQSGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1109)  
 GSSSSSSSSGANQQGLSRGCFGLKLDRIGSMSGLGC (SEQ ID NO: 1110)  
 GLSKGCX<sub>1</sub>FGX<sub>2</sub>KLDRIGSMSGX<sub>3</sub>GC (SEQ ID NO: 1111)  
 GLSKGCFGX<sub>1</sub>KLDRIGSMSGX<sub>2</sub>GC (SEQ ID NO: 1112)  
 GLSKGCFGX<sub>1</sub>KX<sub>2</sub>DRIGSMSGX<sub>3</sub>GC (SEQ ID NO: 1113)  
 GLSKGCFGX<sub>1</sub>KX<sub>2</sub>DRIGSMSGLGC (SEQ ID NO: 1114)  
 GLSRGCFVVKLKLDRIGSMSGLGC (SEQ ID NO: 1115)  
 GLSRGCFSLKLDRIGSMSGLGC (SEQ ID NO: 1116)  
 GLSRGCFTLKLDRIGSMSGLGC (SEQ ID NO: 1117)  
 GLSRGCFGTKLKLDRIGSMSGLGC (SEQ ID NO: 1118)  
 GLSRGCFGTKLKLDRIRSMSGLGC (SEQ ID NO: 1119)  
 GLSRGCFGLKLDRIRSMSGLGC (SEQ ID NO: 1120)  
 GLSRGCFGLKLDRIXSMSGLGC (SEQ ID NO: 1121)  
 GLSRGCFGLKLDRIGSVSGLGC (SEQ ID NO: 1122)  
 GLSRGCFGLKLDRIGSMSGVGC (SEQ ID NO: 1123)  
 GLSRGCFGLKLDRIGSMSGXGC (SEQ ID NO: 1124)

## FIG. 28E

(X: pentanoic acid)  
 (X: heptanoic acid)  
 (X: pentanoic acid)  
 (X: heptanoic acid)  
 (X: Cit)

(all D-amino acids)  
 (X: N-Me-Leu)  
 (X: N-Me-Leu)  
 (X: N-Me-Leu)  
 (X: N-Me-Leu)  
 (X: 3, 4-dichloro-Phe)  
 (X: 3-Me-Phe)

ELSEGGCFGGLKLDRIGSMSGLGC (SEQ ID NO:1125)  
 XELSEGCGFGLKLDRIGSMSGLGC (SEQ ID NO:1126)  
 XELSEGCGFGLKLDRIGSMSGLGC (SEQ ID NO:1127)  
 XELSKRGCFGGLKLDRIGSMSGLGC (SEQ ID NO:1128)  
 XELSKGGCFGGLKLDRIGSMSGLGC (SEQ ID NO:1129)  
 GLSKRGCFGGLXLDRIGSMSGLGC (SEQ ID NO:1130)  
 GLSKGGCFGGLQLDRIGSMSGLGC (SEQ ID NO:1131)  
 GLSKRGCFGGLRLDRIGSMSGLGC (SEQ ID NO:1132)  
 GLSKGGCFGGLKLDRINSMSGLGC (SEQ ID NO:1133)  
 GLSKGGCFGGLKLDRITSMSGLGC (SEQ ID NO:1134)  
 GLSKGGCFGGLKLDRIGSMSGLGC (SEQ ID NO:1135)  
 GLSKGGCFGGXKLDRIGSMSGLGC (SEQ ID NO:1136)  
 GLSKGGCFGGLKXDRIGSMSGLGC (SEQ ID NO:1137)  
 GLSKGGCFGGLKLDRIGSMSGXGC (SEQ ID NO:1138)  
 GLSGGGCXGLKLDRIGSMSGLGC (SEQ ID NO:1139)  
 GLSGGGCXGLKLDRIGSMSGLGC (SEQ ID NO:1140)  
 GLSRSCFGGLKLDRIGSMSGLGC (SEQ ID NO:1141)  
 GLSRRRCFGGLKLDRIGSMSGLGC (SEQ ID NO:1142)  
 GLSGGGCFGGLKLDRIGSMSGLGC (SEQ ID NO:1143)  
 GLSGGGCFGGLSLDRIGSMSGLGC (SEQ ID NO:1144)  
 GLSRGCFGGLKLDRIGQMSGLGC (SEQ ID NO:1145)  
 GLSRGCFGGLKLDRIGNSNGLGC (SEQ ID NO:1146)  
 GLSRGCFGGLKLDRIGSMSLGLC (SEQ ID NO:1147)  
 GLSRGCFGGLKLDRIGSMSRGLC (SEQ ID NO:1148)  
 GLSRGCFGGLKLDRIGSMSGRGC (SEQ ID NO:1149)  
 GLSRGCFGGLKLDRIGSMSGLSC (SEQ ID NO:1150)  
 GLSRGCFGGLKLDRIGSMSGLTC (SEQ ID NO:1151)  
 GLSRGCFGGLKLDRIGSMSGLRC (SEQ ID NO:1152)  
 GLSKGGCXGLKLDRIGSMSGLGC (SEQ ID NO:1153)  
 GLSKGGCXGLKLDRIGSMSGLGC (SEQ ID NO:1154)  
 RSSCCFGGRISRIGACFFFFF (SEQ ID NO:1155)

FIG. 29

## CNP variants

## FIG. 30

## CNP variants having Point Mutations at Position 17

CNP-X17	GLSKGCFGKLDLDRIGSXSGLGC	(SEQ ID NO: 126)
CNP-F17	GLSKGCFGKLDLDRIGFESGLGC	(SEQ ID NO: 119)
CNP-L17	GLSKGCFGKLDLDRIGSISGLGC	(SEQ ID NO: 120)
CNP-I17	GLSKGCFGKLDLDRIGSISGLGC	(SEQ ID NO: 121)
CNP-T17	GLSKGCFGKLDLDRIGSTSGLGC	(SEQ ID NO: 122)
CNP-E17	GLSKGCFGKLDLDRIGSESGLGC	(SEQ ID NO: 156)
CNP-R17	GLSKGCFGKLDLDRIGRSGLGC	(SEQ ID NO: 157)
CNP-Y17	GLSKGCFGKLDLDRIGSYSGLGC	(SEQ ID NO: 158)
CNP-C17	GLSKGCFGKLDLDRIGSCSGLGC	(SEQ ID NO: 159)
CNP-P17	GLSKGCFGKLDLDRIGSPSGLGC	(SEQ ID NO: 160)
CNP-D17	GLSKGCFGKLDLDRIGSDSGLGC	(SEQ ID NO: 161)
CNP37-X17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSXSGLGC	(SEQ ID NO: 162)
CNP37-F17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGFESGLGC	(SEQ ID NO: 163)
CNP37-L17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSISGLGC	(SEQ ID NO: 164)
CNP37-I17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSISGLGC	(SEQ ID NO: 165)
CNP37-T17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSTSGLGC	(SEQ ID NO: 166)
CNP37-E17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSESGLGC	(SEQ ID NO: 167)
CNP37-R17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGRSGLGC	(SEQ ID NO: 168)
CNP37-Y17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSYSGLGC	(SEQ ID NO: 169)
CNP37-C17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSCSGLGC	(SEQ ID NO: 170)
CNP37-P17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSPSGLGC	(SEQ ID NO: 171)
CNP37-D17	QEHPNARYKGANKKGLSKGCFGKLDLDRIGSDSGLGC	(SEQ ID NO: 172)

## FIG. 31A

## Additional CNP variants having Point Mutations at Position 17

KA1-X17	<u>HGPQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 173)
KA2-X17	<u>SGGGQQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 174)
KA3-X17	<u>GGGHGPQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 175)
KA4-X17	<u>GGSGGGQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 176)
KA5-X17	<u>HGPQSGGGGSGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 177)
KA6-X17	<u>GGGHHGPQGSGGGGSGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 178)
KA7-X17	<u>HKLRGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 179)
KA8-X17	<u>HKLRGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 180)
KB1-X17	<u>HGGHKLRGSGGGGSGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 181)
KB2-X17	<u>HKLRGSGGGGSGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 182)
KB3-X17	<u>HGGHKLRGSGGGGSGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 183)
KB4-X17	<u>PGQEHPQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 184)
PGCNP37-X17	<u>EEEEEEHGPQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 185)
KA1 (E6)-X17	<u>EEEEEESGGGHGPQGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 186)
KA2 (E6)-X17	<u>EEEEEEGGSGGGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 187)
KA3 (E6)-X17	<u>EEEEEEGGSGGGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 188)
KA4 (E6)-X17	<u>EEEEEEGGSGGGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 189)
KA5 (E6)-X17	<u>EEEEEEGGSGGGGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 190)
KA6 (E6)-X17	<u>EEEEEEHGPQGSGGGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 191)
KA7 (E6)-X17	<u>EEEEEEGGGGHGPQGSGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 192)
KA8 (E6)-X17	<u>EEEEEEGGGHKLRGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 193)
KB1 (E6)-X17	<u>EEEEEEGGGHKLRGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 194)
KB2 (E6)-X17	<u>EEEEEEGGGHKLRGSGGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 195)
KB3 (E6)-X17	<u>EEEEEEGGGHKLRGSGGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 196)
KB4 (E6)-X17	<u>EEEEEEGGGHKLRGSGGGGGGGGGKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 197)
E6PGCNP37-X17	<u>EEEEEEPGQEHPNARYKGANKKGLSKGCFGLKLDRIGSXGLGC</u>	(SEQ ID NO: 198)

FIG. 31B

## Additional CNP variants having Point Mutations at Position 17

C1 (E6) -X17  
 C2 (E6) -X17  
 C3 (E6) -X17  
 C4 (E6) -X17  
 C5 (E6) -X17  
 C6 (E6) -X17  
 C7 (E6) -X17  
 C8 (E6) -X17  
 C9 (E6) -X17  
 C10 (E6) -X17  
 C11 (E6) -X17  
 D6CNP37-X17  
 C1 (D6) -X17  
 C2 (D6) -X17  
 C3 (D6) -X17  
 C4 (D6) -X17  
 C5 (D6) -X17  
 C6 (D6) -X17  
 D6-14AA linker-CNP-X17  
 CNP-14AA linker-D6-X17

EEEEEEEASTSPANPQPAASSPGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 199)  
 EEEEEEEASTSPANPQPAASSPGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 200)  
 EEEEEEEPSSAAPQPNAPSTSAGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 201)  
 EEEEEEESGGGSGGGKGANKKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 202)  
 EEEEEEESGGGSGGGQGANQQGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 203)  
 EEEEEEESGGGSGGGKGANQKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 204)  
 EEEEEEESGGGSGGGKGANQKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 205)  
 EEEEEEESGGGSGGGQGANKKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 206)  
 EEEEEEESGGGSGGGKGANQQGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 207)  
 EEEEEEESGGGSGGGQGANQKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 208)  
 EEEEEEESGGGSGGGQGANQKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 209)  
 DDDDDDDQEHPNARKYKGANKKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 210)  
 DDDDDDSGGGGSGGGGGSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 211)  
 DDDDDDASTSPANPQPAASSGGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 212)  
 DDDDDDDGSSAAPQPNAPSTSAGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 213)  
 DDDDDDSGGGGSGGGKGANKKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 214)  
 DDDDDDSGGGGSGGGQGANQQGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 215)  
 DDDDDDSGGGGSGGGKGANQKGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 216)  
 DDDDDDDGGGGSGGGGGSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 217)  
 GLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 218)

(SEQ ID NO: 218)  
 GSSAAPRPNAPSTSAGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 219)  
 ASTSPANPRPAASSGGLSKGCFGLKLDRIGSXSLGLGC (SEQ ID NO: 220)

FIG. 32

## CNP variants having M17L

## FIG. 33A

## NC2st-X17 and NC2B-X17 Protein Sequences

NC2st-X17  
(w/ sig. seq.)

MGVHECPAWLWLLSLLSSLWPAYAASWSHQPQEQSGGGGGENLYFQSGGGGENLYFQGGDKTHTCPPCPAPELLGGPSVFLFP  
PKPKDTLMSRTPEVTCVVVDSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVVSVLTVLHQDWLNGK  
EYKCKVSNKALPAPIERTISKARGQPREPQVYTLPPSREEMTKRNQVSLLTCLVKGFYPSPSDIAVEWESENQOPENN  
YKTTPPVLDSDGSFFFLYSKLTVDKSRWQOQGNVFSCSVMHEALHNHYTQKSLSSLSPGKGGGGSGGGSGGGSG  
GLSKGGCFGLKLDRIGSXSGLGC (SEQ ID NO: 527)

NC2st-X17  
(w/o sig. seq.)

ASWSHQPQEQSGGGGGENLYFQGGDKTHTCPPCPAPELLGGPSVFLFPKKPKDTLMSRTPEVTCVVVDSHE  
DPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ  
PREPQVYTLPPSREEMTKRNQVSLLTCLVKGFYPSPSDIAVEWESENQOPENNYKTTPVLDSDGSFFFLYSKLTVDKS  
RWQOQGNVFSCSVMHEALHNHYTQKSLSSLSPGKGGGGSGGGSGGGSGGG  
(SEQ ID NO: 528)

NC2B-X17  
(w/ sig. seq.)

MGVHECPAWLWLLSLLSSLWPAYADKTHTCPPCPAPELLGGPSVFLFPKKPKDTLMSRTPEVTCVVVDSHE  
EDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQ  
PREPQVYTLPPSREEMTKRNQVSLLTCLVKGFYPSPSDIAVEWESENQOPENNYKTTPVLDSDGSFFFLYSKLTVDK  
SRWQOQGNVFSCSVMHEALHNHYTQKSLSSLSPGKGGGGSGGGSGGG  
(SEQ ID NO: 529)

NC2B-X17  
(w/o sig. seq.)

DKTHTCPPCPAPELLGGPSVFLFPKKPKDTLMSRTPEVTCVVVDSHEDEPVKFNWYVDGVEVHNAAKTPRE  
EQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKRNQVSLLT  
CLVKGFYPSPSDIAVEWESENQOPENNYKTTPVLDSDGSFFFLYSKLTVDKSRWQOQGNVFSCSVMHEALHNHYTQK  
SLSLSLSPGKGGGGSGGGSGGGSGGG  
(SEQ ID NO: 530)

FIG. 33B

## NC2B-22-X17, NC2B-28-X17, and NC2B-34-X17 Sequences

NC2B-22-X17  
(w/ sig. seq.)

NC2B-22-X17  
(w/o sig. seq.)

NC2B-28-X17  
(w/sig, seq.)

NC2B-28-X17 (W/O sig, seq.)

NC2B-34-X17  
(w/sig. seq.)

NC2B-34-X17  
(w/o sig. seq.)

# FIG. 33C

## NC2-X17 Variants

NC2-KGANQK-X17  
(w/o sig. seq.)

NC2-CNP53mut2-X17  
(w/o sig. seq.)

NC2-CNP53mut2-X17  
(w/o sig. seq.)

NC2-CNP53mut2-X17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGGSGGGSKGANKKGLSKG  
(SEQ ID NO: 537)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGGSGGGSKGANKKGLSKG  
(SEQ ID NO: 538)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGGSGGGSKGANKKGLSKG  
(SEQ ID NO: 539)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGGSGGGSKGANKKGLSKG  
(SEQ ID NO: 540)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGLQVDTQSQAQWQ  
LLQEHPNAAQYKGANKKKGLSKG  
(SEQ ID NO: 541)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLLSPGKGGGLQVDTQSQAQWQ  
LLQEHPNAAQYKGANKKKGLSKG  
(SEQ ID NO: 542)

## FIG. 33D

## Fc-CNP53-X17 Constructs

Fc-CNP53-X17  
(w/o sig. seq.)

MGVHECPAWILLILLSLWMPGAYADKTHTCBPCPAELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCVKVSNKALPAPIEKTISKARG  
QREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPVLSDGFFFLYSKLTVDK  
SRWQQGNVFSCSVMHEALHNHYTQKSLSSPGKGGDIRVDTKSRAAWARLQEHPNARGYGANKKGLISKGC  
FGLKIDRIGSXSGIGC (SEQ ID NO: 543)

Fc-CNP53-X17  
(w/o sig. seq.)

DKTHTCPPCPAELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCVKVSNKALPAPIEKTISKARG  
CLVKGFYPSDIAVEWESNGOPENNYKTTPVLSDGFFFLYSKLTVDK  
SISLSPGKGGDIRVDTKSRAAWARLQEHPNARGYGANKKGLISKGC  
NO: 544)

Fc-CNP53-AA-X17  
(w/o sig. seq.)

MGVHECPAWILLILLSLWMPGAYADKTHTCBPCPAELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSH  
EDPEVKFNWYVDGVEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCVKVSNKALPAPIEKTISKARG  
QREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPVLSDGFFFLYSKLTVDK  
SRWQQGNVFSCSVMHEALHNHYTQKSLSSPGKGGDIRVDTKSRAAWARLQEHPNARGYGANKKGLISKGC  
FGLKIDRIGSXSGIGC (SEQ ID NO: 545)

Fc-CNP53-AA-X17  
(w/o sig. seq.)

DKTHTCPPCPAELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCVKVSNKALPAPIEKTISKARG  
CLVKGFYPSDIAVEWESNGOPENNYKTTPVLSDGFFFLYSKLTVDK  
SISLSPGKGGDIRVDTKSRAAWARLQEHPNARGYGANKKGLISKGC  
NO: 546)

FIG. 33E

## Constructs Having N-terminal NP-X17 fused to C-terminal Fc Domain

CNP-X17-16AAlinker-FC-His<sub>10</sub> (NC1)

CNP-X17-6AAlinker-FC-His<sub>10</sub> (NC3)

CNP-X17-6AAlinker-FC

CDNP-X17-F.C.

CDNP-X17-Saa-FC

CDNP-X17-Sra-FC

GLSKGCFIGKLDRIGSXSGIGCGGGGSGDKTHTCPPCPAPELLGGPSVFLPPPKDTLMISRTPEVTCVVVD  
VSHEDEPVKFNWYVDGVEVHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK  
AKGQPREPQVTLPSSREEMTKNQVSILTCLVKGFYPSDIAVEWESENQOPENNYKTPVLDSDGSFFLYSRLT  
VDKRSRWQQGNVFSCSVMEALHNHYTQKSSLSPGK (SEQ ID NO: 549)

GLSKGCGFGLKLDRIGSXSGLGCPSLRDPRPNAPSTSADKTHTCPPCKPKDTLIMISRT  
PEVTCVWDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALP  
APIEKTISKAKGOPREPQVYTLPPSREEMTKNOVSLSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDG  
SEFLYSKITVTKSPVTCOCNTYTESCSYMTHEAIHNUHYTOKSISI SPCK / SEQ ID NO: 550

GLSKKGCEGLKLDRIGSXSGLCPSSMAAPRPNAPSTSADKTHTCPPCPAPELGGPSVFLFPPKPKDITMISRT  
PEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKT~~KPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALP~~  
APIEKTISAKGQP~~REPQVTLLPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGOPENNYKTPVLDSDG~~  
SFEFLYSKLTVDKSRWOOGNVFTSCSVMEALHNHYTOKS~~SSLSPGK~~ (SEO ID NO: 551)

GLSKGCFIGIKLDRIGSXGICCPSSRAPRPNAPST SADKTHTCPPCPAPEILGGPSVFLFPPKPKDITLIMISRT  
PEVTCVWDVSHEDPEVKFNWYDGVETEVHNAKTRPREEQYNSTYRVSVLTVLHQDWLNGKEYCKVSNKALP  
APIEKTISSAKGQPREPQVTYLPPSREEMTKNQVSLSLTCLVKGFYPSDIAVEWESNGOPENNYKTTPPVLDSDG  
SFLFLYSKLTVDKSRWOOGNVFTSCSVMEALHNHYTOKSLLSPGK (SEQ ID NO : 552)

# FIG. 34A

## NC2 Variants

NC2-KGANQK  
(w/o sig. seq.)

NC2-CNP53mut2  
(w/o sig. seq.)

NC2-CNP53mut2  
(w/o sig. seq.)

NC2-CNP53mut2  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
KKLDRIGSMSGLGC  
(SEQ ID NO: 511)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLT  
CLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
KKLDRIGSMSGLGC  
(SEQ ID NO: 512)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLT  
CLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
KKLDRIGSMSGLGC  
(SEQ ID NO: 513)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLT  
CLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
KKLDRIGSMSGLGC  
(SEQ ID NO: 514)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLT  
CLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
AAWQLLQEHPNAAQYKGANKKKLDRIGSMSGLGC  
(SEQ ID NO: 515)

DKTHHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNAAKTKPRE  
EQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
ETKTSKAKGQPREPQVYTLPPSREEMTKNQVSLLT  
CLVKGFYPSDI  
AVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
AAWQLLQEHPNAAQYKGANKKKLDRIGSMSGLGC  
(SEQ ID NO: 516)

## FIG. 34B

### D10-NC2 Variants

D10-NC2-KGANQK  
(w/o sig. seq.)

D10-NC2-CNP53mut2  
(w/o sig. seq.)

D10-NC2-KGANQK  
(w/o sig. seq.)

D10-NC2-CNP53mut2  
(w/o sig. seq.)

D10-NC2-KGANQK  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWYVTDGVEVHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAP  
IEKTISRAKGQPREPQYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGSGGSKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 553)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAPIEKTISRAKGQPREPQYTLPPSRE  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGSGGSKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 554)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWYVTDGVEVHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAP  
IEKTISRAKGQPREPQYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGSGGSKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 555)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAPIEKTISRAKGQPREPQYTLPPSRE  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGSGGSKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 556)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWYVTDGVEVHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAP  
IEKTISRAKGQPREPQYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
AAQQLQEHPNAAQQYKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 557)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKTPREEQYNSTYRVSVLTVLDNGKEYKCKVSNKALPAPIEKTISRAKGQPREPQYTLPPSRE  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPVLSDGSF  
FLYSKLTVDKSRWQQGNVSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
AAQQLQEHPNAAQQYKGANKGLSKGCFGKKLDR  
MSGLGC (SEQ ID NO: 558)

# FIG. 34C

## NC2-F17 Variants

NC2-KGANQK-F17  
(w/o sig. seq.)

NC2-CNP53mut2-F17  
(w/o sig. seq.)

NC2-CNP53mut2-F17  
(w/o sig. seq.)

NC2-CNP53mut2-F17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
QFVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
GGKGLKLDRIGSFSGLGC  
(SEQ ID NO: 559)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNNAKTKPREE  
QYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
GGKGLKLDRIGSFSGLGC  
(SEQ ID NO: 560)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
GGKGLKLDRIGSFSGLGC  
(SEQ ID NO: 561)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGDI  
GGGDLQVDTQSQAQWQHNAQYKGANKKGLSKGC  
FCGLKLDRIGSFSGLGC  
(SEQ ID NO: 562)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGDI  
GGGDLQVDTQSQAQWQHNAQYKGANKKGLSKGC  
FCGLKLDRIGSFSGLGC  
(SEQ ID NO: 563)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTCVVYDVSHEDPEVFKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPI  
EKTKISKAKGQPREPQVYTLPPSREEMTKNQVS  
LTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSEFLYSKLTVDK  
SRWQQGNVFSCSVMSHEALHNHYTQKSLSLSPGKGGGDI  
GGGDLQVDTQSQAQWQHNAQYKGANKKGLSKGC  
(SEQ ID NO: 564)

## FIG. 34D

### D10-NC2-F17 Variants

D10-NC2-KGANQK-F17  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWEDPEVKFNWYVTDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
FLYSKLTVDKSRWQQGNVFSCSVMVH  
GSESGLGCC (SEQ ID NO: 565)

D10-NC2-KGANQK-F17  
(w/o sig. seq.)

DDDDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPEPNNYKTPPVLDSDGFE (SEQ ID NO: 566)

D10-NC2-KGANQK-F17  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWEDPEVKFNWYVTDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
FLYSKLTVDKSRWQQGNVFSCSVMVH  
GSESGLGCC (SEQ ID NO: 567)

D10-NC2-KGANQK-F17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPEPNNYKTPPVLDSDGFE (SEQ ID NO: 568)

D10-NC2-KGANQK-F17  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
TCVYVVDWEDPEVKFNWYVTDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
FLYSKLTVDKSRWQQGNVFSCSVMVH  
GSESGLGCC (SEQ ID NO: 569)

D10-NC2-CNP53mut2-F17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPPKPKDTLMSRTPEVTKFNWYVVDGVE  
VHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTISKLTVDKSRWQGNVFSCSVMVH  
EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPEPNNYKTPPVLDSDGFE (SEQ ID NO: 570)

## FIG. 34E

### NC2-L17 Variants

NC2-KGANQK-L17  
(w/o sig. seq.)

NC2-CNP53mut2-L17  
(w/o sig. seq.)

NC2-CNP53mut2-L17  
(w/o sig. seq.)

NC2-CNP53mut2-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANQKGLSKGCGFLKLDRIGSLSGLGC  
(SEQ ID NO: 571)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDKSRWQQGNVFTSCSVMHEALHNHYTQK  
SISSLSPGKGGGSGGGSKGANQKGLSKGCGFLKLDRIGSLSGLGC (SEQ ID NO: 572)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANQKGLSKGCGFLKLDRIGSLSGLGC  
(SEQ ID NO: 573)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDKSRWQQGNVFTSCSVMHEALHNHYTQK  
SISSLSPGKGGGSGGGSKGANQKGLSKGCGFLKLDRIGSLSGLGC (SEQ ID NO: 574)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGDLQVDTQSQAAWAQLQEHPNAAQYKGANKKGLSKGC  
FGFLKLDRIGSLSGLGC (SEQ ID NO: 575)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SDPEVKFNWYVDGVEHNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SISSLSPGKGGDLQVDTQSQAAWAQLQEHPNAAQYKGANKKGLSKGC (SEQ ID NO: 576)

# FIG. 34F

## D10-NC2-L17 Variants

D10-NC2-KGANQK-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRTI  
GSLSGLGC (SEQ ID NO: 577)

D10-NC2-KGANQK-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRTI  
GSLSGLGC (SEQ ID NO: 578)

D10-NC2-KGANQK-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRTI  
GSLSGLGC (SEQ ID NO: 579)

D10-NC2-KGANQK-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKKGLSKGCFGKLKDRTGSLGLGC (SEQ ID NO: 580)

D10-NC2-CNP53mut2-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKKGLSKGCFGKLKDRTGSLGLGC (SEQ ID NO: 581)

D10-NC2-CNP53mut2-L17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGFFGLKLDRTGSLGLGC (SEQ ID NO: 582)

# FIG. 34G

## NC2-R17 Variants

NC2-KGANQK-R17  
(w/o sig. seq.)

NC2-CNP53mut2-R17  
(w/o sig. seq.)

NC2-CNP53mut2-R17  
(w/o sig. seq.)

NC2-CNP53mut2-R17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGGSGGGSKGANKKGLSKGCGFLKLDRIGSRSGLGC  
(SEQ ID NO: 583)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDKSRWQQGNVFTSCSVMHEALHNHYTQK  
SISSSPGKGGGGSGGGSKGANKKGLSKGCGFLKLDRIGSRSGLGC (SEQ ID NO: 584)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGGSGGGSKGANQKGLSKGCGFLKLDRIGSRSGLGC  
(SEQ ID NO: 585)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDKSRWQQGNVFTSCSVMHEALHNHYTQK  
SISSSPGKGGGGSGGGSKGANQKGLSKGCGFLKLDRIGSRSGLGC (SEQ ID NO: 586)

MGVHECPAHLWLLSLSLWMPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGLQVDTQSQAAWAQLQEHPNAAQYKGANKKGLSKGC  
FGFLKLDRIGSRSGLGC (SEQ ID NO: 587)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNAAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDIAVEWESNGQOPENNYKTTPPVLDSDGSFFFLYSKLTVDKSRWQQGNVFTSCSVMHEALHNHYTQK  
SISSSPGKGGGLQVDTQSQAAWAQLQEHPNAAQYKGANKKGLSKGC (SEQ ID NO: 588)

# FIG. 34H

## D10-NC2-R17 Variants

D10-NC2-KGANQK-R17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRI  
GSRSGLGC (SEQ ID NO: 589)

D10-NC2-KGANQK-R17  
(w/o sig. seq.)

DDDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRI  
GSRSGLGC (SEQ ID NO: 590)

D10-NC2-KGANQK-R17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWMPGAYADDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGANKKGLSKGCFGKLKDRI  
GSRSGLGC (SEQ ID NO: 591)

D10-NC2-KGANQK-R17  
(w/o sig. seq.)

DDDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKGLSKGCFGKLKDRI GSRSGLGC (SEQ ID NO: 592)

D10-NC2-CNP53mut2-R17  
(w/o sig. seq.)

DDDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKGLSKGCFGKLKDRI GSRSGLGC (SEQ ID NO: 593)

D10-NC2-CNP53mut2-R17  
(w/o sig. seq.)

DDDDDDDDDDKTHTCPPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKISRAKGQPREPOVYTLPPSREEMTKNOVSLLTCLVKGFYP SDIAVENE SNGQOPENNYKTTPPVLDSDGSF  
FLYSKLTVDKSRWQOGNVE SCSVMHEALHNHYTQKSLSLSPGKGFFGLKLDRI GSRSGLGC  
C (SEQ ID NO: 594)

# FIG. 34I

## NC2-Y17 Variants

NC2-KGANQK-Y17  
(w/o sig. seq.)

NC2-CNP53mut2-Y17  
(w/o sig. seq.)

NC2-CNP53mut2-Y17  
(w/o sig. seq.)

NC2-CNP53mut2-Y17  
(w/o sig. seq.)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
(SEQ ID NO: 595)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
(SEQ ID NO: 596)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
(SEQ ID NO: 597)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGSGGGSKGAN  
(SEQ ID NO: 598)

MGVHECPAHLWLLSLSLWPGAYADKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
CLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
(SEQ ID NO: 599)

DKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMI  
SRTPEVTKFENWYDGEVHNNAKTKPREEQYNSTYRVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKG  
QPREPQVYTLPPSREEMTKNQVSLLTCLVKGFYPSDI  
AVEWESENQOPENNYKTTPPVLDSDGSFFFLYSKLTVDK  
SRWQQGNVFTSCSVMHEALHNHYTQKSLSLSPGKGGGDLQVDTQSQA  
(SEQ ID NO: 600)

# FIG. 34J

## D10-NC2-Y17 Variants

D10-NC2-KGANQK-Y17  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAQGQPREPQVTILPPSREEMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGSGGGSKGANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 601)

D10-NC2-KGANQK-Y17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIASKAKGQPREPQVTLPSSRE  
EMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGSGGGSKGANKKGLSKGCEGKLKDRTI  
EALHNHYTQKSISLSPGKGGGSGGGSKGANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 602)

D10-NC2-KGANQK-Y17  
(w/o sig. seq.)

MGVHECPAWLWLLSLSLWMPGAYADDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAQGQPREPQVTILPPSREEMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGSGGGSKGANKKGLSKGCEGKLKDRTI  
EALHNHYTQKSISLSPGKGGGSGGGSKGANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 603)

D10-NC2-KGANQK-Y17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIASKAKGQPREPQVTLPSSRE  
EMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGDQLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 604)

D10-NC2-CNP53mut2-Y17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
TCVYVVDVSHEDPEVKFNWYVDGVEVHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAP  
IETKTISKAQGQPREPQVTILPPSREEMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGDQLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 605)

D10-NC2-CNP53mut2-Y17  
(w/o sig. seq.)

DDDDDDDDDKTHTCPCPAPELLGGPSVFTLFPKKPKDTLMSRTPE  
VHNAAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIASKAKGQPREPQVTLPSSRE  
EMTKNOVSILTCLVKGFYP SDIAVENE SNGQOPENNYKTTTPVVLDSDGSE  
FLYSKLTVDKSRWQOGNVE SC SVMHEALHNHYTQKSISLSPGKGGGDQLQVDTQSQAAWAQLQEHPNAAQQYKG  
ANKKGLSKGCEGKLKDRTI  
GSYSGLGC (SEQ ID NO: 606)

**INTERNATIONAL SEARCH REPORT**

International application No.

PCT/US 12/39004

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(8) - A61K 39/00; A61K 38/00; A61P 13/06; C12N 9/16 (2012.01)  
USPC - 424/134.1; 514/12.1

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
USPC: 424/134.1; 514/12.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
USPC: 435/196; 424/134.1; 514/12.1  
(keyword limited; terms below)

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

PubWEST (PGPB,USPT,USOC,EPAB,JPAB); Google; PubMed

Search terms: alkaline phosphatase, ALP, sALP, TNALP, neurocutaneous, neurofibromatosis, extracellular, bone, skeletal, mineralization, natriuretic peptide, NPR-B, SEQ ID NOs:1204, 1216, 1205, 572

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- A	US 2010/0184680 A1 (BEVEC) 22 July 2010 (22.07.2010) para [0007], [0116], [0141]-[0142], [0201]	25 --- 26
Y	US 2010/0297119 A1 (CRINE et al.) 25 November 2010 (25.11.2010) para [0040], [0064], [0089], [0117], [0163], [0176]; SEQ ID NOs:4-6	1-5
Y	RAMACHANDRAN et al. Treatment of an anabolic bone deficiency in neurofibromatosis with bone morphogenetic proteins and its potential application for congenital pseudoarthrosis of the tibia. J. Bone Joint Surg. Br. (2009), Vol. 91-B, No. Supp. I, abstract 137; entire abstract	1-5
A	US 7,803,769 B2 (SULLIVAN et al.) 28 September 2010 (28.09.2010) SEQ ID NO:83	26

Further documents are listed in the continuation of Box C.

\* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

24 October 2012 (24.10.2012)

Date of mailing of the international search report

02 NOV 2012

Name and mailing address of the ISA/US

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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 12/39004

Continuation of Box III - observations where unity of invention is lacking:

Group II+: claims 25-31 and 37, drawn to a method of treating a neurocutaneous syndrome in a subject, said method comprising administering to said subject a therapeutically effective amount of a pharmaceutical composition comprising:  
 (a) a polypeptide comprising the structure V-NP-W; and  
 (b) a pharmaceutically acceptable excipient,  
 wherein NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), V is absent or is an amino acid sequence of at least one amino acid, and W is absent or is an amino acid sequence of at least one amino acid, thereby treating said syndrome in said subject. The first invention is restricted to SEQ ID NO: 504 (Claims 25 and 26). Should an additional fee(s) be paid, Applicant is invited to elect an additional sequence(s) to be searched. The exact claims searched will depend on Applicant's election.

Group III: claims 83-86 and 118-123, drawn to a composition comprising a first polypeptide and a second polypeptide, wherein  
 a) said first polypeptide comprises the structure A-sALP-B, wherein  
 i) sALP is the extracellular domain of an alkaline phosphatase,  
 ii) A is absent or is an amino acid sequence of at least one amino acid, and  
 iii) B is absent or is an amino acid sequence of at least one amino acid; and  
 b) said second polypeptide comprises the structure V-NP-W, wherein  
 i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B),  
 ii) V is absent or is an amino acid sequence of at least one amino acid, and  
 iii) W is absent or is an amino acid sequence of at least one amino acid.

The inventions listed as Groups I through III do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The inventions of Groups I-II+ do not include the inventive concept of a composition comprising a first polypeptide A-sALP-B and a second polypeptide V-NP-W, as required by Group III.

The inventions of Group I do not include the inventive concept of a polypeptide comprising the structure V-NP-W, as required by Group II+.

The inventions of Group II+ do not include the inventive concept of a polypeptide comprising the structure A-sALP-B, as required by Group I.

The inventions of Groups I and II+ share the technical feature of a method of treating a neurocutaneous syndrome in a subject, said method comprising administering to said subject a therapeutically effective amount of a pharmaceutical composition comprising a therapeutic polypeptide. However, this shared technical feature does not represent a contribution over prior art as being anticipated by US 2010/0184680 A1 (Bevec). Bevec discloses Claim 25, a method of treating a neurocutaneous syndrome in a subject (para [0141], neurocutaneous melanosis), said method comprising administering to said subject a therapeutically effective amount of a pharmaceutical composition (para [0201]) comprising:

(a) a polypeptide comprising the structure V-NP-W; wherein NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), V is absent and W is absent (para [0007]), and  
 (b) a pharmaceutically acceptable excipient (para [0142]), thereby treating said syndrome in said subject (para [0116]). As said composition was known in the art at the time of the invention, this cannot be considered a special technical feature that would otherwise unify the groups.

The inventions of Groups I and III share the technical feature of a polypeptide comprises the structure A-sALP-B, wherein i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent or is an amino acid sequence of at least one amino acid, and iii) B is absent or is an amino acid sequence of at least one amino acid. However, this shared technical feature does not represent a contribution over prior art as being anticipated by US 2010/0297119 A1 to Crine et al. (hereafter 'Crine'). Crine teaches a polypeptide comprises the structure A-sALP-B, wherein i) sALP is the extracellular domain of an alkaline phosphatase, ii) A is absent, and iii) B is an amino acid sequence of at least one amino acid (Fig. 1C, hsTNALP-FcD10 w/o signal peptide). As said composition was known in the art at the time of the invention, this cannot be considered a special technical feature that would otherwise unify the groups.

The inventions of Groups II+ and III share the technical feature of a polypeptide comprises the structure V-NP-W, wherein i) NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), ii) V is absent or is an amino acid sequence of at least one amino acid, and iii) W is absent or is an amino acid sequence of at least one amino acid. However, this shared technical feature does not represent a contribution over prior art as being anticipated by Bevec. Bevec discloses a polypeptide comprising the structure V-NP-W; wherein NP is a natriuretic peptide that is an agonist of natriuretic peptide receptor B (NPR-B), V is absent and W is absent (para [0007]). As said composition was known in the art at the time of the invention, this cannot be considered a special technical feature that would otherwise unify the groups.

Groups I through III therefore lack unity under PCT Rule 13 because they do not share a same or corresponding special technical feature.