

**(12) STANDARD PATENT**  
**(19) AUSTRALIAN PATENT OFFICE**

**(11) Application No. AU 2017294551 B2**

(54) Title  
**Soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptides and uses thereof**

(51) International Patent Classification(s)  
**C07K 14/71** (2006.01)      **A61K 38/17** (2006.01)

(21) Application No: **2017294551**      (22) Date of Filing: **2017.07.07**

(87) WIPO No: **WO18/007597**

(30) Priority Data

(31) Number      (32) Date      (33) Country  
**62/359,607**      **2016.07.07**      **US**  
**62/467,478**      **2017.03.06**      **US**

(43) Publication Date: **2018.01.11**  
(44) Accepted Journal Date: **2022.03.03**

(71) Applicant(s)  
**Pfizer Inc.;Université Côte d'Azur;INSERM (Institut National de la Santé et de la Recherche Médicale)**

(72) Inventor(s)  
**Gouze, Elvire;Garcia, Stéphanie**

(74) Agent / Attorney  
**Griffith Hack, Level 10 161 Collins St, MELBOURNE, VIC, 3000, AU**

(56) Related Art  
**WO 2014/111467 A1**  
**WO 2010/017198 A2**  
**WO 2011/084711 A2**

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property

Organization

International Bureau



(10) International Publication Number

WO 2018/007597 A1

(43) International Publication Date

11 January 2018 (11.01.2018)

WO 2018/007597 A1

(51) International Patent Classification:

C07K 14/71 (2006.01) A61K 38/17 (2006.01)

Published:

- with international search report (Art. 21(3))
- with sequence listing part of description (Rule 5.2(a))

(21) International Application Number:

PCT/EP2017/067119

(22) International Filing Date:

07 July 2017 (07.07.2017)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/359,607 07 July 2016 (07.07.2016) US  
62/467,478 06 March 2017 (06.03.2017) US

(71) **Applicants:** THERACHON [FR/FR]; 2000 route des lucioles, Les Algorithmes - Aristote A, 06410 Biot (FR). INSERM (INSTITUT NATIONAL DE LA SANTÉ ET DE LA RECHERCHE MÉDICALE) [FR/FR]; 101 rue de Tolbiac, 75013 Paris (FR). UNIVERSITÉ NICE SOPHIA ANTIPOLIS [FR/FR]; 28 avenue de Valrose, 06103 Nice (FR).



(72) **Inventors:** GOUZE, Elvire; 420 chemin des darboussières, 06620 Vallauris (FR). GARCIA, Stéphanie; 51 Corniche Fleurie Immeuble Apogée B2, 06200 Nice (FR).

(74) **Agent:** ZWICKER, Jörk; ZSP Patentanwälte PartG mbB, Hansastrasse 32, 80686 München (DE).

(81) **Designated States** (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) **Designated States** (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

WO 2018/007597 A1

(54) **Title:** SOLUBLE FIBROBLAST GROWTH FACTOR RECEPTOR 3 (SFGFR3) POLYPEPTIDES AND USES THEREOF

(57) **Abstract:** The invention features soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptides. The invention also features methods of using sFGFR3 polypeptides to treat skeletal growth retardation disorders, such as achondroplasia.

**SOLUBLE FIBROBLAST GROWTH FACTOR RECEPTOR 3 (sFGFR3) POLYPEPTIDES  
AND USES THEREOF**

**FIELD OF THE INVENTION**

5 The invention features soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptides and compositions thereof. The invention also features methods to treat skeletal growth retardation disorders, such as achondroplasia.

**BACKGROUND OF THE INVENTION**

10 Fibroblast growth factor receptor 3 (FGFR3) is a member of the fibroblast growth factor (FGFR) family, in which there is high amino acid sequence conservation between family members. Members of the FGFR family are differentiated by both ligand binding affinities and tissue distribution. A full-length FGFR polypeptide contains an extracellular domain (ECD), a hydrophobic transmembrane domain, and a cytoplasmic tyrosine kinase domain. The ECD of  
15 FGFR polypeptides interacts with fibroblast growth factors (FGFs) to mediate downstream signaling, which ultimately influences cellular differentiation. In particular, activation of the FGFR3 protein plays a role in bone development by inhibiting chondrocyte proliferation at the growth plate and limiting bone elongation.

Gain-of-function point mutations in FGFR3 are known to cause several types of human  
20 skeletal growth retardation disorders, such as achondroplasia, thanatophoric dysplasia type I (TDI), thanatophoric dysplasia type II (TDII), severe achondroplasia with developmental delay and acanthosis nigricans (SADDAN), hypochondroplasia, and craniosynostosis syndromes (e.g., Muenke syndrome, Crouzon syndrome, and Crouzonodermoskeletal syndrome). Loss-of-function point mutations in FGFR3 are also known to cause skeletal growth retardation  
25 disorders, such as camptodactyly, tall stature, and hearing loss syndrome (CATSHL). Achondroplasia is the most common form of short-limb dwarfism and is characterized by disproportionate shortness of limbs and relative macrocephaly. Approximately 97% of achondroplasia is caused by a single point mutation in the gene encoding FGFR3, in which a glycine residue is substituted with an arginine residue at position 380 of the FGFR3 amino acid  
30 sequence. Upon ligand binding, the mutation decreases the elimination of the receptor/ligand complex resulting in prolonged intracellular signaling. This prolonged FGFR3 signaling inhibits the proliferation and differentiation of the cartilage growth plate, consequently impairing endochondral bone growth.

35 There exists a need for improved therapeutics that target dysfunctional FGFR3 for treating skeletal growth retardation disorders, such as achondroplasia.

It is to be understood that if any prior art publication is referred to herein, such reference does not constitute an admission that the publication forms a part of the common general knowledge in the art in Australia or any other country.

## SUMMARY OF THE INVENTION

A first aspect provides a soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide comprising an amino acid sequence of SEQ ID NO: 33.

A second aspect provides a soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide encodable by nucleotides 52-1098 of SEQ ID NO: 21.

A third aspect provides a soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide encoded by nucleotides 52-1098 of SEQ ID NO: 21.

A fourth aspect provides a nucleic acid encoding the sFGFR3 polypeptide of any one of the first to third aspects.

A fifth aspect provides a vector comprising the nucleic acid of the fourth aspect.

A sixth aspect provides a host cell comprising the nucleic acid of the fourth aspect or the vector of the fifth aspect.

A seventh aspect provides a method of producing the sFGFR3 polypeptide of any one of the first to third aspects, the method comprising:

- (i) culturing the host cell of the sixth aspect in a culture medium under conditions suitable for expression of the sFGFR3 polypeptide; and
- (ii) recovering the sFGFR3 polypeptide from the culture medium.

An eighth aspect provides a sFGFR3 polypeptide when produced by the method of the seventh aspect.

A ninth aspect provides a composition comprising:

the sFGFR3 polypeptide of any one of the first to third or eighth aspects; and  
a pharmaceutically acceptable excipient.

A tenth aspect provides a method comprising administering to a subject in need thereof the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect.

Also disclosed is use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect in a therapy in a subject in need thereof.

Also disclosed is use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects in the manufacture of a medicament for a therapy in a subject in need thereof.

An eleventh aspect provides a method for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof, the method comprising administering to the subject the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect.

A twelfth aspect provides use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof.

A thirteenth aspect provides use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects in the manufacture of a medicament for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof.

A fourteenth aspect provides a method for treating achondroplasia in a human subject in need thereof, the method comprising administering to the human subject the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect.

A fifteenth aspect provides use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects or the composition of the ninth aspect for treating achondroplasia in a human subject in need thereof.

A sixteenth aspect provides use of the sFGFR3 polypeptide of any one of the first to third or eighth aspects in the manufacture of a medicament for treating achondroplasia in a human subject in need thereof.

Also disclosed is a soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide comprising an amino acid sequence at least 90% identical to SEQ ID NO: 2.

The disclosure features soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptides and uses thereof, including the use of the sFGFR3 polypeptides for the treatment of skeletal growth retardation disorders (e.g., achondroplasia) in a patient (e.g., a human, particularly an infant, a child, or an adolescent). In particular, the sFGFR3 polypeptides of the disclosure feature a deletion of, e.g., amino acids 289 to 400 of the amino acid sequence of the wildtype FGFR3 polypeptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5 or 32), to provide the following sFGFR3 polypeptides: sFGFR3\_Del4 including an amino acid substitution of a cysteine residue with a serine residue at position 253 (sFGFR3\_Del4-C253S; SEQ ID NO: 2) and sFGFR3\_Del4 including an extended Ig-like C2-type domain 3 (sFGFR3\_Del4-D3; SEQ ID NO: 33) and variants thereof, such as a sFGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 4. Additionally, the sFGFR3 polypeptides of the disclosure may include a signal peptide, such as a sFGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34.

Disclosed herein is a soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide including a polypeptide sequence having at least 90% amino acid sequence identity (e.g., at least 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more (e.g., 100%) sequence identity) to amino acid residues 23 to 357 of SEQ ID NO: 32. In particular, the polypeptide lacks a signal peptide (e.g., amino acids 1-22 of SEQ ID NO: 32) and a transmembrane domain of FGFR3 (e.g., amino acids of 367-399 of SEQ ID NO: 32) and (i) is less than 500 amino acids in length (e.g., less than 475, 450, 425, 400, 375, or 350 amino acids in length); (ii) includes 200

consecutive amino acids or fewer (e.g., 175, 150, 125, 100, 75, 50, 40, 30, 20, 15, or fewer consecutive amino acids) of an intracellular domain of FGFR3; and/or (iii) lacks a tyrosine kinase domain of FGFR3. The sFGFR3 polypeptide can also include an intracellular domain of FGFR3, such as amino acid residues 423 to 435 of SEQ ID NO: 32 or an amino acid sequence having at least 90%, 92%, 95%, 97%, or 99% sequence identity to amino acid residues 423 to 435 of SEQ ID NO: 32. In particular, the polypeptide includes an amino acid sequence having at least 92%, 95%, 97%, 99%, or 100% sequence identity to SEQ ID NO: 33 (e.g., the polypeptide includes or consists of SEQ ID NO: 33). The sFGFR3 polypeptides can also include a signal peptide (e.g., the signal peptide can have the amino acid sequence of SEQ ID NO: 6 or 35 or an amino acid sequence having at least 92%, 95%, 97%, or 99% sequence identity to SEQ ID NO: 6 or 35). For example, the sFGFR3 polypeptide may have an amino acid sequence with at least 92%, 95%, 97%, 99%, or 100% sequence identity to SEQ ID NO: 34 (e.g., the sFGFR3 polypeptide includes or consists of SEQ ID NO: 34). The sFGFR3 polypeptide may also have a heterologous signal peptide (e.g., the polypeptide includes a heterologous signal peptide having the amino acid sequence of SEQ ID NO: 35).

Also disclosed is an sFGFR3 polypeptide including an amino acid sequence having at least 85% sequence identity (e.g., at least 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 NO: 1, in which the sFGFR3 polypeptide further includes an amino acid substitution that removes a cysteine residue at position 253 of SEQ ID NO: 1. For example, the cysteine residue at position 253 is substituted with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine. In particular, the sFGFR3 polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 2. For instance, the sFGFR3 polypeptide can be an isolated sFGFR3 polypeptide. The sFGFR3 polypeptides can also include a signal peptide (e.g., the signal peptide can have the amino acid sequence of SEQ ID NO: 6 or 35 or an amino acid sequence having at least 92%, 95%, 97%, or 99% sequence identity to SEQ ID NO: 6 or 35). For example, the sFGFR3 may have an amino acid sequence with at least 92%, 95%, 97%, 99%, or 100% sequence identity to SEQ ID NO: 18 (e.g., the sFGFR3 polypeptide includes or consists of SEQ ID NO: 18). The sFGFR3 polypeptide may also have a heterologous signal peptide (e.g., the polypeptide includes a heterologous signal peptide having the amino acid sequence of SEQ ID NO: 35).

Also disclosed is a sFGFR3 polypeptide including an amino acid sequence having at least 85% sequence identity (e.g., at least 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 NO: 1, in which the sFGFR3 polypeptide further includes a domain including an amino acid sequence having at least 85% sequence identity (e.g., at least 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity) to all or a fragment of the amino acid sequence of SEQ ID NO: 3 (e.g., at least 10, 20, 30, 40, 45, or more

consecutive amino acids of SEQ ID NO: 3), in which the domain is inserted between amino acid residues 288 and 289 of SEQ ID NO: 1. For example, the domain can include an amino acid sequence having at least 85%, 90%, 92%, 95%, 97%, or 99% sequence identity to the amino acid sequence of SEQ ID NO: 3 (e.g., the domain can include or consists of the amino acid sequence of SEQ ID NO: 3). Optionally, the sFGFR3 polypeptide includes an amino acid substitution of a cysteine residue with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine, at position 253 of SEQ ID NO: 1 and/or position 28 of SEQ ID NO: 3. In particular, the sFGFR3 polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 4. For example, the sFGFR3 polypeptide can be an isolated sFGFR3 polypeptide.

Also featured is a polynucleotide (e.g., an isolated polynucleotide) that encodes the sFGFR3 polypeptide of the disclosure including a nucleic acid sequence having at least 85%, 90%, 92%, 95%, 97%, or 99% sequence identity to the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37 (e.g., the polynucleotide includes or consists of the nucleic acid of SEQ ID NO: 20, 21, 36, or 37). The disclosure also features a vector (e.g., an isolated vector) including the polynucleotide, such as a plasmid, an artificial chromosome, a viral vector, or a phage vector. Additionally, the disclosure features a host cell (e.g., an isolated host cell) including the polynucleotide, such as a HEK 293 cell or CHO cell.

The disclosure features a composition including the sFGFR3 polypeptide of disclosure or the polynucleotide that encodes the sFGFR3 polypeptide of the disclosure. In addition, the vector or host cell that includes the polynucleotide encoding the sFGFR3 polypeptide can be formulated in a composition. The composition can further include a pharmaceutically acceptable excipient, carrier, or diluent. The composition including the sFGFR3 polypeptide, polynucleotide, or vector can be formulated for administration at a dose of about 0.002 mg/kg to about 30 mg/kg, such as about 0.001 mg/kg to about 10 mg/kg. The composition including the host cell can be formulated for administration at a dose of about  $1 \times 10^3$  cells/mL to about  $1 \times 10^{12}$  cells/mL. The composition can be formulated for daily, weekly, or monthly administration, such as seven times a week, six times a week, five times a week, four times a week, three times a week, twice a week, weekly, every two weeks, or once a month. For example, the composition including the sFGFR3 polypeptide, polynucleotide, or vector is formulated for administration at a dose of about 0.25 mg/kg to about 10 mg/kg once or twice a week. The composition can be formulated for parenteral administration (e.g., subcutaneous administration, intravenous administration, intramuscular administration, intra-arterial administration, intrathecal administration, or intraperitoneal administration), enteral administration, or topical administration. Preferably, the composition is formulated for subcutaneous administration. The disclosure also features a medicament that includes one or more of the compositions described above.

The disclosure also features a method of delivering an sFGFR3 polypeptide to tissue

(e.g., skeletal tissue) in a patient (e.g. a human) having a skeletal growth retardation disorder (e.g., achondroplasia) including administering to the patient an effective amount of the sFGFR3 polypeptide of the disclosure, a polynucleotide encoding the sFGFR3 polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell.

Also disclosed is a method of treating a skeletal growth retardation disorder (e.g., a FGFR3-related skeletal disease) in a patient (e.g., a human) that includes administering the polypeptide of the disclosure or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell. The FGFR3-related skeletal disease is selected from the group consisting of achondroplasia, thanatophoric dysplasia type I (TDI), thanatophoric dysplasia type II (TDII), severe achondroplasia with developmental delay and acanthosis nigricans (SADDEN), hypochondroplasia, a craniosynostosis syndrome (e.g., Muenke syndrome, Crouzon syndrome, and Crouzonodermoskeletal syndrome), and camptodactyly, tall stature, and hearing loss syndrome (CATSHL). In particular, the skeletal growth retardation disorder is achondroplasia.

The FGFR3-related skeletal disease can be caused by expression in the patient of a constitutively active FGFR3, such as an amino acid substitution of a glycine residue with an arginine residue at position 380 of SEQ ID NO: 5 or 32. In particular, the patient may be diagnosed with the skeletal growth retardation disorder (e.g., prior to treatment). For instance, the patient exhibits one or more symptoms of the skeletal growth retardation disorder selected from the group consisting of short limbs, short trunk, bowlegs, a waddling gait, skull malformations, cloverleaf skull, craniosynostosis, wormian bones, anomalies of the hands, anomalies of the feet, hitchhiker thumb, and chest anomalies, such that the patient exhibits an improvement in the one or more symptoms of the skeletal growth retardation disorder after administration of the sFGFR3 polypeptide (or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell). Additionally, the patient may have not been previously administered the sFGFR3 polypeptide. For example, the patient may be an infant, a child, an adolescent, or an adult.

For example, the polypeptide is administered to the patient at a dose of about 0.002 mg/kg to about 30 mg/kg (e.g., a dose of about 0.001 mg/kg to about 10 mg/kg). The polypeptide may be administered to the patient one or more times daily, weekly (e.g., twice a week, three times a week, four times a week, five times a week, six times a week, or seven times a week), every two weeks, monthly, or yearly. For example, the polypeptide is administered to the patient at a dose of about 0.25 mg/kg to about 30 mg/kg at least about once or twice a week or more (e.g., the polypeptide is administered to the patient at a dose of about 2.5 mg/kg or about 10 mg/kg once or twice weekly). The polypeptide can be administered to

the patient in a composition including a pharmaceutically acceptable excipient, carrier, or diluent. The polypeptide can be administered to the patient parenterally (e.g., subcutaneously, intravenously, intramuscularly, intra-arterially, intrathecally, or intraperitoneally), enterally, or topically. Preferably, the composition is administered to the patient by subcutaneous injection. Additionally, the polypeptide can bind to fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), fibroblast growth factor 18 (FGF18), fibroblast growth factor 19 (FGF19), or fibroblast growth factor 21 (FGF21). In particular, the binding can be characterized by an equilibrium dissociation constant ( $K_d$ ) of about 0.2 nM to about 20 nM, such as the binding is characterized by a  $K_d$  of about 1 nM to about 10 nM (e.g., about 1 nm, about 2 nm, about 3 nm, about 4 nm, about 5 nm, about 6 nm, about 7 nm, about 8 nm, about 9 nm, or about 10 nm). The polypeptide can exhibit greater binding affinity to FGF1, FGF2, FGF9, and FGF18 relative to the binding affinity of the polypeptide to FGF19 and FGF21.

The polypeptide can have an *in vivo* half-life of between about 2 hours to about 25 hours (e.g., 2 hours, 3 hours, 4 hours, 5 hours, 6 hours, 7 hours, 8 hours, 9 hours, 10 hours, 11 hours, 12 hours, 13 hours, 14 hours, 15 hours, 16 hours, 17 hours, 18 hours, 19 hours, 20 hours, 21 hours, 22 hours, 23 hours, 24 hours, or 25 hours). Preferably, administration of the polypeptide provides one or more, or all, of the following: an increase in survival of the patient, an improvement in locomotion of the patient, an improvement in abdominal breathing in the patient, an increase in body and/or bone length of the patient, an improvement in the cranial ratio of the patient, and/or restoration of the foramen magnum shape in the patient, e.g., relative to an untreated patient (e.g., an untreated achondroplasia patient).

The disclosure also features a method of producing the sFGFR3 polypeptide of the disclosure, which includes culturing the host cell described above (e.g., a CHO cell or HEK 293 cell) in a culture medium under conditions suitable to effect expression of the sFGFR3 polypeptide and recovering the sFGFR3 polypeptide from the culture medium. In particular, the recovering includes chromatography, such as affinity chromatography (e.g., ion exchange chromatography or anti-FLAG chromatography, such as immunoprecipitation) or size exclusion chromatography.

Also disclosed is the polypeptide of the disclosure (or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell) for treating a skeletal growth retardation disorder in a patient. In particular, the sFGFR3 polypeptide can bind to fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), fibroblast growth factor 18 (FGF18), fibroblast growth factor 19 (FGF19), or fibroblast growth factor 21 (FGF21).

Also disclosed is a sFGFR3 polypeptide (or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell) including an amino

acid sequence having at least 85% sequence identity (e.g., at least 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 NO: 1 for treating a skeletal growth retardation disorder in a patient (e.g., a human), in which the sFGFR3 polypeptide further includes an amino acid substitution that removes a cysteine residue at position 253 of SEQ ID NO: 1. For example, the cysteine residue at position 253 is substituted with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine. In particular, the sFGFR3 polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 2. For example, the sFGFR3 polypeptide can be an isolated sFGFR3 polypeptide. Furthermore, the sFGFR3 polypeptide can bind to fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), fibroblast growth factor 18 (FGF18), fibroblast growth factor 19 (FGF19), or fibroblast growth factor 21 (FGF21).

Also disclosed is a sFGFR3 polypeptide (or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell) including an amino acid sequence having at least 85% sequence identity (e.g., at least 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 NO: 1 for treating a skeletal growth retardation disorder in a patient (e.g., a human), in which the sFGFR3 polypeptide further includes a domain including an amino acid sequence having at least 85% sequence identity (e.g., at least 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity) to all or a fragment of the amino acid sequence of SEQ ID NO: 3 (e.g., at least 10, 20, 30, 40, 45, or more consecutive amino acids of SEQ ID NO: 3), in which the domain is inserted between amino acid residues 288 and 289 of SEQ ID NO: 1. For example, the domain can include an amino acid sequence having at least 85%, 90%, 92%, 95%, 97%, or 99% sequence identity to the amino acid sequence of SEQ ID NO: 3 (e.g., the domain can include or consists of the amino acid sequence of SEQ ID NO: 3). Optionally, the sFGFR3 polypeptide includes an amino acid substitution of a cysteine residue with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine, at position 253 of SEQ ID NO: 1 and/or position 28 of SEQ ID NO: 3. In particular, the sFGFR3 polypeptide includes or consists of the amino acid sequence of SEQ ID NO: 4. For example, the sFGFR3 polypeptide can be an isolated sFGFR3 polypeptide. Furthermore, the sFGFR3 polypeptide can bind to fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), fibroblast growth factor 18 (FGF18), fibroblast growth factor 19 (FGF19), or fibroblast growth factor 21 (FGF21).

The use of the disclosure also features the administration of a polynucleotide, vector, host cell, or composition of the disclosure. The sFGFR3 polypeptide of the disclosure can be encoded by a polynucleotide including a nucleic acid sequence having at least 85%, 90%, 92%,

95%, 97%, or 99% sequence identity to the nucleic acid sequence of SEQ ID NO: 20 or 36 (e.g., the polynucleotide includes or consists of the nucleic acid of SEQ ID NO: 20 or 36). The sFGFR3 polypeptide of the disclosure can be encoded by a polynucleotide including a nucleic acid sequence having at least 85%, 90%, 92%, 95%, 97%, or 99% sequence identity to the nucleic acid sequence of SEQ ID NO: 21 or 37 (e.g., the polynucleotide includes or consists of the nucleic acid of SEQ ID NO: 21 or 37).

The skeletal growth retardation disorder of the disclosure can be any FGFR3-related skeletal disease, such as achondroplasia, TDI, TDII, severe achondroplasia with developmental delay and acanthosis nigricans (SADDEN), hypochondroplasia, a craniosynostosis syndrome (e.g., Muenke syndrome, Crouzon syndrome, and Crouzonodermoskeletal syndrome), or CATSHL. In particular, the skeletal growth retardation disorder is achondroplasia. The FGFR3-related skeletal disease can be caused by expression in the patient of a constitutively active FGFR3, e.g., in which the constitutively active FGFR3 includes an amino acid substitution of a glycine residue with an arginine residue at position 380 of SEQ ID NO: 5.

The patient (e.g., a human) of the disclosure can be one that has been diagnosed with the skeletal growth retardation disorder (e.g., prior to treatment). The patient can exhibit one or more symptoms of the skeletal growth retardation disorder (e.g., achondroplasia) selected from the group consisting of short limbs, short trunk, bowlegs, a waddling gait, skull malformations, cloverleaf skull, craniosynostosis, wormian bones, anomalies of the hands, anomalies of the feet, hitchhiker thumb, and chest anomalies. As a result of the methods, the patient can exhibit an improvement in the one or more symptoms of the skeletal growth retardation disorder after administration of the sFGFR3 polypeptide. Moreover, administration of the sFGFR3 polypeptide can increase survival of the patient and/or restore the shape of the foramen magnum of the patient. The patient can be an infant, a child, an adolescent, or an adult. Additionally, the patient can be one that has not been previously administered the sFGFR3 polypeptide (or a polynucleotide encoding the polypeptide, a vector containing the polynucleotide, a host cell containing the polynucleotide or vector, or a composition containing the polypeptide, polynucleotide, vector, or host cell).

The sFGFR3 polypeptide, polynucleotide, or vector of the disclosure can be 30 administered to the patient (e.g., a human) at a dose of about 0.002 mg/kg to about 30 mg/kg, such as about 0.001 mg/kg to about 10 mg/kg. The composition including the host cell of the disclosure can be administered to the patient (e.g., a human) at a dose of about  $1 \times 10^3$  cells/mL to about  $1 \times 10^{12}$  cells/mL. For example, the sFGFR3 polypeptide, polynucleotide, vector, or host cell is administered to the patient one or more times daily, weekly, monthly, or 35 yearly (e.g., the sFGFR3 polypeptide is administered to the patient seven times a week, six

times a week, five times a week, four times a week, three times a week, twice a week, weekly, every two weeks, or once a month). In particular, the sFGFR3 polypeptide is administered to the patient at a dose of about 0.25 mg/kg to about 10 mg/kg once or twice a week. The sFGFR3 polypeptide can be administered to the patient in a composition including a pharmaceutically acceptable excipient, carrier, or diluent. For example, the composition is administered to the patient parenterally (e.g., subcutaneously, intravenously, intramuscularly, intra-arterially, intrathecally, or intraperitoneally), enterally, or topically. In particular, the composition is administered to the patient by subcutaneous injection.

The disclosure features a method of manufacturing the sFGFR3 polypeptide of the disclosure by deleting the signal peptide, the transmembrane domain, and a portion of the intracellular domain from a FGFR3 polypeptide (e.g., to manufacture a polypeptide having the amino acid sequence of SEQ ID NO: 33). In particular, the intracellular domain consists of amino acid residues 436 to 806 of SEQ ID NO: 32. The disclosure also features a method of manufacturing the sFGFR3 polypeptide of the disclosure by introducing an amino acid substitution that removes a cysteine residue at position 253 of SEQ ID NO: 1 (e.g., to manufacture a polypeptide having the amino acid sequence of SEQ ID NO: 2). For example, the cysteine residue at position 253 is substituted with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine.

The disclosure also features a kit including the sFGFR3 polypeptide of the disclosure (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33), the polynucleotide of the disclosure (e.g., a polynucleotide having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37), the vector of the disclosure (e.g., a plasmid, an artificial chromosome, a viral vector, or a phage vector), or the host cell of the disclosure (e.g., a HEK 293 cell or a CHO cell), in which the kit optionally includes instructions for using the kit.

## Definitions

As used herein, “a” and “an” means “at least one” or “one or more” unless otherwise indicated. In addition, the singular forms “a,” “an,” and “the” include plural referents unless the context clearly dictates otherwise.

As used herein, “about” refers to an amount that is  $\pm$  10% of the recited value and is preferably  $\pm$  5% of the recited value, or more preferably  $\pm$  2% of the recited value. For instance, the term “about” can be used to modify all dosages or ranges recited herein by  $\pm$  10% of the recited values or range endpoints.

In the claims which follow and in the description of the invention, except where the context requires otherwise due to express language or necessary implication, the word “comprise” or variations such as “comprises” or “comprising” is used in an inclusive sense, i.e. to specify the presence of the stated features but not to preclude the presence or addition of further features in various embodiments of the invention.

The term “domain” refers to a conserved region of the amino acid sequence of a polypeptide (e.g. a FGFR3 polypeptide) having an identifiable structure and/or function within the polypeptide. A domain can vary in length from, e.g., about 20 amino acids to about 600 amino acids. Exemplary domains include the immunoglobulin domains of FGFR3 (e.g., Ig-like C2-type domain 1, Ig-like C2-type domain 2, and Ig-like C2-type domain 3).

The term “dosage” refers to a determined quantity of an active agent (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33) calculated to produce a desired therapeutic effect (e.g., treatment of a skeletal growth retardation disorder, such as achondroplasia) when the active agent is administered to a patient (e.g., a patient having a skeletal growth retardation disorder, such as achondroplasia). A dosage may be defined in terms of a defined amount of the active agent or a defined amount coupled with a particular frequency of administration. A dosage form can include an sFGFR3 polypeptide or fragment thereof in association with any suitable pharmaceutical excipient, carrier, or diluent.

The terms “effective amount,” “amount effective to,” and “therapeutically effective amount” refer to an amount of an sFGFR3 polypeptide, a vector encoding a sFGFR3, and/or an sFGFR3 composition that is sufficient to produce a desired result, for example, the treatment of a skeletal growth retardation disorder (e.g., achondroplasia).

The terms “extracellular domain” and “ECD” refer to the portion of a FGFR3 polypeptide that extends beyond the transmembrane domain into the extracellular space. The ECD mediates binding of a FGFR3 to one or more fibroblast growth factors (FGFs). For instance, an ECD includes the Ig-like C2-type domains 1-3 of a FGFR3 polypeptide. In particular, the ECD 5 includes the Ig-like C2-type domain 1 of a wildtype (wt) FGFR3 polypeptide (e.g., amino acids 36-88 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 5 (a mature FGFR3 protein without a signal sequence) or amino acids 57-110 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 32 (a precursor FGFR3 protein with the signal sequence)), the Ig-like C2-type domain 2 of a wildtype (wt) FGFR3 polypeptide (e.g., amino acids 139-234 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 5 or amino acids 161-245 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 32), and the Ig-like C2-type domain 3 of a wt FGFR3 polypeptide (e.g., amino acids 247-10 335 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 5 or amino acids 268-310 of a wt FGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 32).  
15 An ECD of a FGFR3 can also include a fragment of the wildtype FGFR3 Ig-like C2-type domain 3, for instance, aa 247-288 of SEQ ID NO: 1, which can further include, e.g., an amino acid substitution of a cysteine residue with a serine residue or another conservative amino acid substitution (e.g., alanine, glycine, proline, or threonine) at position 253 of SEQ ID NO: 1 (e.g., aa 247-288 of SEQ ID NO: 2). Additionally, an ECD can include an Ig-like C2-type domain 3 of,  
20 e.g., aa 247-335 of SEQ ID NO: 4. Thus, exemplary ECDs of FGFR3 polypeptides include, e.g., those polypeptides having the amino acid sequence of aa 1-288 of SEQ ID NOs: 1 and 2 or aa 1-335 of SEQ ID NOs: 4 and 33. In particular, the ECD of a FGFR3 polypeptide includes aa 1-335 of SEQ ID NO: 33.

The term “FGFR3-related skeletal disease,” as used herein, refers to a skeletal disease 25 that is caused by an abnormal increase in the activation of FGFR3, such as by expression of a gain-of-function mutant of the FGFR3. The phrase “gain-of-function mutant of the FGFR3” refers to a mutant of the FGFR3 exhibiting a biological activity, such as triggering downstream signaling, which is higher than the biological activity of the corresponding wild-type FGFR3 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5) in the presence of a FGF 30 ligand. FGFR3-related skeletal diseases can include an inherited or a sporadic disease. Exemplary FGFR3-related skeletal diseases include, but are not limited to, achondroplasia, thanatophoric dysplasia type I (TDI), thanatophoric dysplasia type II (TDII), severe achondroplasia with developmental delay and acanthosis nigricans (SADDAN), hypochondroplasia, a craniosynostosis syndrome (e.g., Muenke syndrome, Crouzon syndrome, 35 and Crouzonodermoskeletal syndrome), and camptodactyly, tall stature, and hearing loss syndrome (CATSHL).

The terms “fibroblast growth factor” and “FGF” refer to a member of the FGF family, which includes structurally related signaling molecules involved in various metabolic processes,

including endocrine signaling pathways, development, wound healing, and angiogenesis. FGFs play key roles in the proliferation and differentiation of a wide range of cell and tissue types. The term preferably refers to FGF1, FGF2, FGF9, FGF18, FGF19, and FGF21, which have been shown to bind FGFR3. For instance, FGFs can include human FGF1 (e.g., a polypeptide 5 having the amino acid sequence of SEQ ID NO: 13), human FGF2 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 14), human FGF9 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 15), human FGF18 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 16), human FGF19 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 38), and human FGF21 (e.g., a polypeptide having the amino acid 10 sequence of SEQ ID NO: 39).

The terms “fibroblast growth factor receptor 3,” “FGFR3,” or “FGFR3 receptor,” as used herein, refer to a polypeptide that specifically binds one or more FGFs (e.g., FGF1, FGF2, FGF9, FGF18, FGF19, and/or FGF21). The human *FGFR3* gene, which is located on the distal short arm of chromosome 4, encodes an 806 amino acid protein precursor (fibroblast growth 15 factor receptor 3 isoform 1 precursor), which contains 19 exons, and includes a signal peptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 6 or 35). Mutations in the FGFR3 amino acid sequence that lead to skeletal growth disorders, (e.g., achondroplasia), include, e.g., the substitution of a glycine residue at position 380 with an arginine residue (i.e., G380R). The naturally occurring human FGFR3 gene has a nucleotide sequence as shown in 20 Genbank Accession number NM\_000142.4 and the naturally occurring human FGFR3 protein has an amino acid sequence as shown in Genbank Accession number NP\_000133, herein represented by SEQ ID NO: 5. The wildtype FGFR3 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5) consists of an extracellular immunoglobulin-like membrane domain including Ig-like C2-type domains 1-3 (amino acid residues 1 to 335), a transmembrane domain 25 (amino acid residues 345 to 377), and an intracellular domain (amino acid residues 378 to 784). FGFR3s can include fragments and/or variants (e.g., splice variants, such as splice variants utilizing alternate exon 8 rather than exon 9) of the full-length, wild-type FGFR3 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5).

The terms “fragment” and “portion” refer to a part of a whole, such as a polypeptide or 30 nucleic acid molecule that contains, 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 or portion 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, 35 370, 380, 390, 400, 500, 600, 700, or more amino acid residues, up to the entire length of the reference polypeptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5 or 32). For example, a FGFR3 fragment can include any polypeptide having at least 200, 205, 210, 215, 220, 225, 235, 230, 240, 245, 250, 255, 260, 265, 275, 280, 285, 290, or 300

consecutive amino acids of SEQ ID NO: 1 or 2. Additionally, a FGFR3 fragment can include any polypeptide having at least 200, 205, 210, 215, 220, 225, 235, 230, 240, 245, 250, 255, 260, 265, 275, 280, 285, 290, 300, 305, 310, 315, 320, 325, 330, 335, 345, or 345 consecutive amino acids of SEQ ID NO: 4 or 33.

5 As used herein, the term “host cell” refers to a vehicle that includes the necessary cellular components, e.g., organelles, needed to express an sFGFR3 polypeptide from a corresponding polynucleotide. The nucleic acid sequence of the polynucleotide is typically included in a nucleic acid vector (e.g., a plasmid, an artificial chromosome, a viral vector, or a phage vector) that can be introduced into the host cell by conventional techniques known in the 10 art (e.g., transformation, transfection, electroporation, calcium phosphate precipitation, and direct microinjection). A host cell may be a prokaryotic cell, e.g., a bacterial or an archaeal cell, or a eukaryotic cell, e.g., a mammalian cell (e.g., a Chinese Hamster Ovary (CHO) cell or a Human Embryonic Kidney 293 (HEK 293)). Preferably, the host cell is a mammalian cell, such as a CHO cell.

15 By “isolated” is meant separated, recovered, or purified from its natural environment. For example, an isolated sFGFR3 polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2 or 4) can be characterized by a certain degree of purity after isolating the sFGFR3 polypeptide from, e.g., cell culture media. An isolated sFGFR3 polypeptide can be at least 75% pure, such that the 20 sFGFR3 polynucleotide constitutes at least 75% by weight of the total material (e.g., polypeptides, polynucleotides, cellular debris, and environmental contaminants) present in the preparation (e.g., at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 99%, or at least 99.5% by weight of the total material present in the preparation). Likewise, an isolated polynucleotide encoding an sFGFR3 polypeptide (e.g., a polynucleotide 25 having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37), or an isolated host cell (e.g., CHO cell, a HEK 293 cell, L cell, C127 cell, 3T3 cell, BHK cell, or COS-7 cell) can be at least 75% pure, such that the polynucleotide or host cell constitutes at least 75% by weight of the total material (e.g., polypeptides, polynucleotides, cellular debris, and environmental 30 contaminants) present in the preparation (e.g., at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 99%, or at least 99.5% by weight of the total material present in the preparation).

“Polynucleotide” and “nucleic acid molecule,” as used interchangeably herein, refer to polymers of nucleotides of any length, and include DNA and RNA. The nucleotides can be deoxyribonucleotides, ribonucleotides, modified nucleotides or bases, and/or analogs thereof, or 35 any substrate that can be incorporated into a polymer by DNA or RNA polymerase or by a synthetic reaction. A polynucleotide can include modified nucleotides, such as methylated nucleotides and analogs thereof. If present, modification to the nucleotide structure can be imparted before or after assembly of the polymer. The sequence of nucleotides can be

interrupted by non-nucleotide components. A polynucleotide can be further modified after synthesis, such as by conjugation with a label.

The terms “patient” and “subject” refer to a mammal, including, but not limited to, a human (e.g., a human having a skeletal growth retardation disorder, such as achondroplasia) or a non-human mammal (e.g., a non-human mammal having a skeletal growth retardation disorder, such as achondroplasia), such as a bovine, equine, canine, ovine, or feline. Preferably, the patient is a human having a skeletal growth retardation disorder (e.g., achondroplasia), particularly an infant, a child, or an adolescent having a skeletal growth retardation disorder (e.g., achondroplasia).

The terms “parenteral administration,” “administered parenterally,” and other grammatically equivalent phrases, as used herein, refer to a mode of administration of compositions including an sFGFR3 polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34) other than enteral and topical administration, usually by injection, and include, without limitation, subcutaneous, intradermal, intravenous, intranasal, intraocular, pulmonary, intramuscular, intra-arterial, intrathecal, intracapsular, intraorbital, intracardiac, intradermal, intrapulmonary, intraperitoneal, transtracheal, subcuticular, intraarticular, subcapsular, subarachnoid, intraspinal, epidural, intracerebral, intracranial, intracarotid, and intrasternal injection and infusion.

By “pharmaceutically acceptable diluent, excipient, carrier, or adjuvant” is meant a diluent, excipient, carrier, or adjuvant, respectively that is physiologically acceptable to the subject (e.g., a human) while retaining the therapeutic properties of the pharmaceutical composition (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34) with which it is administered. One exemplary pharmaceutically acceptable carrier is physiological saline. Other physiologically acceptable diluents, excipients, carriers, or adjuvants and their formulations are known to one skilled in the art.

By “pharmaceutical composition” is meant a composition containing an active agent, such as an sFGFR3 (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34), formulated with at least one pharmaceutically acceptable excipient, carrier, or diluent. The pharmaceutical composition may be manufactured or sold with the approval of a governmental regulatory agency as part of a therapeutic regimen for the treatment of a disease or event (e.g., a skeletal growth retardation disorder, such as achondroplasia) in a patient (e.g., a patient having a skeletal growth retardation disorder, such as a patient having achondroplasia). Pharmaceutical

compositions can be formulated, e.g., for parenteral administration, such as for subcutaneous administration (e.g. by subcutaneous injection) or intravenous administration (e.g., as a sterile solution free of particulate emboli and in a solvent system suitable for intravenous use), or for oral administration (e.g., as a tablet, capsule, caplet, gelcap, or syrup).

5 As used herein, the term “sequence identity” refers to the percentage of amino acid (or nucleic acid) residues of a candidate sequence, e.g., an FGFR3 polypeptide, that are identical to the amino acid (or nucleic acid) residues of a reference sequence, e.g., a wild-type sFGFR3 polypeptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5 or 32) or an sFGFR3 polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide 10 having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34), after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent identity (e.g., gaps can be introduced in one or both of the candidate and reference sequences for optimal alignment and non-homologous sequences can be disregarded for 15 comparison purposes). Alignment for purposes of determining percent identity can be achieved in various ways that are within the skill in the art, for instance, using publicly available computer software, such as BLAST, BLAST-2, BLAST-P, BLAST-N, BLAST-X, WU-BLAST-2, ALIGN, ALIGN-2, CLUSTAL, or Megalign (DNASTAR) software. Those skilled in the art can determine appropriate parameters for measuring alignment, including any algorithms needed to achieve 20 maximal alignment over the full length of the sequences being compared. For instance, the percent amino acid (or nucleic acid) sequence identity of a given candidate sequence to, with, or against a given reference sequence (which can alternatively be phrased as a given candidate sequence that has or includes a certain percent amino acid (or nucleic acid) sequence identity to, with, or against a given reference sequence) is calculated as follows:

25 
$$100 \times (\text{fraction of A/B})$$

where A is the number of amino acid (or nucleic acid) residues scored as identical in the alignment of the candidate sequence and the reference sequence, and where B is the total number of amino acid (or nucleic acid) residues in the reference sequence. In particular, a reference sequence aligned for comparison with a candidate sequence can show that the 30 candidate sequence exhibits from, e.g., 50% to 100% identity across the full length of the candidate sequence or a selected portion of contiguous amino acid (or nucleic acid) residues of the candidate sequence. The length of the candidate sequence aligned for comparison purpose is at least 30%, e.g., at least 40%, e.g., at least 50%, 60%, 70%, 80%, 90%, or 100% of the length of the reference sequence. When a position in the candidate sequence is occupied by 35 the same amino acid (or nucleic acid) residue as the corresponding position in the reference sequence, then the molecules are identical at that position.

By “signal peptide” is meant a short peptide (e.g., 5-30 amino acids in length, such as 22 amino acids in length) at the N-terminus of a polypeptide that directs a polypeptide towards the

secretory pathway (e.g., the extracellular space). The signal peptide is typically cleaved during secretion of the polypeptide. The signal sequence may direct the polypeptide to an intracellular compartment or organelle, e.g., the Golgi apparatus. A signal sequence may be identified by homology, or biological activity, to a peptide 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 peptide 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, Wis. 53705, BLAST, or PILEUP/Prettybox programs). A signal peptide can be one that is, for example, substantially identical to the amino acid sequence of SEQ ID NO: 6 or 10 35.

The term "skeletal growth retardation disorder," as used herein, refers to a skeletal disease characterized by deformities and/or malformations of the bones. These disorders include, but are not limiting to, skeletal growth retardation disorders caused by growth plate (physeal) fractures, idiopathic skeletal growth retardation disorders, or FGFR3-related skeletal 15 diseases. In particular, a patient having a skeletal growth retardation disorder (e.g., achondroplasia) may have bones that are shorter than the bones of a healthy patient. For example, the skeletal growth retardation disorder may include a skeletal dysplasia, e.g., achondroplasia, homozygous achondroplasia, heterozygous achondroplasia, achondrogenesis, acrodysostosis, acromesomelic dysplasia, atelosteogenesis, camptomelic dysplasia, 20 chondrodysplasia 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, 25 fibrochondrogenesis, fibrous dysplasia, hereditary multiple exostoses, 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, 30 neurofibromatosis, osteoarthritis, osteochondrodysplasia, osteogenesis imperfecta, 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, and spondyloepimetaphyseal dysplasia.

The terms "soluble fibroblast growth factor receptor 3," "soluble FGFR3," and "sFGFR3" 35 refer to a FGFR3 that is characterized by the absence or functional disruption of all or a substantial part of the transmembrane domain and any polypeptide portion that would anchor the FGFR3 polypeptide to a cell membrane (e.g., a tyrosine kinase domain). An sFGFR3 polypeptide is a non-membrane bound form of an FGFR3 polypeptide. In particular, the

transmembrane domain of FGFR3 extends from amino acid residues 345 to 377 of the wild-type FGFR3 sequence (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5) or amino acid residues 367 to 399 of the wild-type FGFR3 sequence including a signal peptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 32). Thus, the sFGFR3 polypeptide can include a deletion of a portion or all of amino acid residues 345 to 377 of the wild-type FGFR3 polypeptide sequence (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5) or amino acid residues 367 to 399 of the wild-type FGFR3 sequence including a signal peptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 32). The sFGFR3 polypeptide can further include deletions of the cytoplasmic domain of the wild-type FGFR3 polypeptide sequence (amino acid residues 378 to 784 of SEQ ID NO: 5) or the wild-type FGFR3 polypeptide sequence including a signal peptide sequence (amino acid residues 378 to 806 of SEQ ID NO: 32).

Exemplary sFGFR3 polypeptides can include, but are not limited to, at least amino acids 1 to 100, 1 to 125, 1 to 150, 1 to 175, 1 to 200, 1 to 205, 1 to 210, 1 to 215, 1 to 220, 1 to 225, 1 to 230, 1 to 235, 1 to 240, 1 to 245, 1 to 250, 1 to 252, 1 to 255, 1 to 260, 1 to 265, 1 to 270, 1 to 275, 1 to 280, 1 to 285, 1 to 290, 1 to 295, or 1 to 300, or 1 to 301 of SEQ ID NOs: 1 or 2. sFGFR3 polypeptides can 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 any of these sFGFR3 polypeptides of SEQ ID NO: 1 or 2. Additionally, exemplary sFGFR3 polypeptides can include, but are not limited to, at least amino acids 1 to 100, 1 to 125, 1 to 150, 1 to 175, 1 to 200, 1 to 205, 1 to 210, 1 to 215, 1 to 220, 1 to 225, 1 to 230, 1 to 235, 1 to 240, 1 to 245, 1 to 250, 1 to 255, 1 to 260, 1 to 265, 1 to 270, 1 to 275, 1 to 280, 1 to 285, 1 to 290, 1 to 295, 1 to 300, 1 to 305, 1 to 310, 1 to 315, 1 to 320, 1 to 325, 1 to 330, 1 to 335, 1 to 340, 1 to 345, or 1 to 348 of SEQ ID NO: 4 or 33. sFGFR3 polypeptides can 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 any of these sFGFR3 polypeptides having the amino acid sequence of SEQ ID NO: 4 or 33. Any of the above sFGFR3 polypeptides or variants thereof can optionally include a signal peptide at the N-terminal position, such as amino acids 1 to 22 of SEQ ID NO: 6 (MGAPACALALCVAVAIVAGASS) or amino acids 1 to 19 of SEQ ID NO: 35 (e.g., MMSFVSLLLVGILFHATQA).

By "treating" and "treatment" is meant a reduction (e.g., by at least about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 60%, about 70%, about 80%, about 90%, about 95%, about 99%, or even 100%) in the progression or severity of a skeletal growth retardation disorder (e.g., achondroplasia), or in the progression, severity, or frequency of one or more symptoms of a skeletal growth retardation disorder (e.g., achondroplasia) in a patient (e.g., a human, such as an infant, a child, or an adolescent). Treatment can occur for a treatment period, in which an sFGFR3 polypeptide is

administered for a period of time (e.g., days, months, years, or longer) to treat a patient (e.g., a human, such as an infant, a child, or an adolescent) having a skeletal growth retardation disorder, such as achondroplasia. Exemplary symptoms of achondroplasia that can be treated with an sFGFR3 (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34) include, but are not limited to, short stature, a long trunk, shortened limbs, an adult height of between about 42 to about 56 inches, a relatively large head, a forehead that is prominent, underdeveloped portions of the face, genu valgum (e.g., "knock-knee"), a waddling gait, short and stubby fingers, short and stubby toes, limited ability to straighten the arm at the elbow, an excessive curve of the lower back, dental problems (e.g. from overcrowding of teeth), weight control problems, neurological problems, respiratory problems, and/or pain and numbness in the lower back and/or spine.

The term "variant," with respect to a polypeptide, refers to a polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34) that differs by one or more changes in the amino acid sequence from the polypeptide from which the variant is derived (e.g., the parent polypeptide, such a polypeptide having the amino acid sequence of SEQ ID NO: 1 or 7). The term "variant," with respect to a polynucleotide, refers to a polynucleotide (e.g., a polynucleotide encoding a sFGFR3 polypeptide, such as a polynucleotide having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37) that differs by one or more changes in the nucleic acid sequence from the polynucleotide from which the variant is derived (e.g., the parent polynucleotide). The changes in the amino acid or nucleic acid sequence of the variant can be, e.g., amino acid or nucleic acid substitutions, insertions, deletions, N-terminal truncations, or C-terminal truncations, or any combination thereof. In particular, the amino acid substitutions may be conservative and/or non-conservative substitutions. A variant can be characterized by amino acid sequence identity or nucleic acid sequence identity to the parent polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34) or parent polynucleotide (e.g., a polynucleotide encoding a sFGFR3 polypeptide, such as a polynucleotide having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37), respectively. For example, a variant can 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 a polypeptide having the amino acid sequence of SEQ ID NO: 1, 2, 4, or 33. A variant can also include any polynucleotide 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 a polynucleotide having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37.

By “vector” is meant a DNA construct that includes one or more polynucleotides, or fragments thereof, encoding an sFGFR3 polypeptide (e.g., an sFGFR3 polypeptide or variant thereof, such as a polypeptide having the amino acid sequence of SEQ ID NO: 2, 4, or 33, or a sFGFR3 polypeptide including a signal peptide, such as a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34). The vector can be used to infect a cell (e.g., a host cell or a cell of a patient having a human skeletal growth retardation disorder, such as achondroplasia), which results in the translation of the polynucleotides of the vector into a sFGFR3 polypeptide.

10 One type of vector is a “plasmid,” which refers to a circular double stranded DNA loop into which additional DNA segments may be ligated. Certain vectors are capable of autonomous replication in a host cell into which they are introduced (e.g., bacterial vectors having a bacterial origin of replication and episomal mammalian vectors). Other vectors (e.g., non-episomal mammalian vectors) can be integrated into the genome of a host cell upon introduction into the

15 host cell, and thereby are replicated along with the host genome. Moreover, certain vectors are capable of directing the expression of genes to which they are operatively linked. In general, expression vectors of utility in recombinant DNA techniques are often in the form of plasmids.

The term “unit dosage form(s)” refers to physically discrete unit(s) suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect, in association with any suitable pharmaceutical excipient, carrier, or diluent.

The recitation herein of numerical ranges by endpoints is intended to include all numbers subsumed within that range (e.g., a recitation of 1 to 5 includes 1, 1.5, 2, 2.75, 3, 3.80, 4, and 5).

25 Other features and advantages of the invention will be apparent from the following Detailed Description and from the claims.

#### BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

**FIGS. 1A-1D** are graphs showing sensorgrams of the sFGFR3 polypeptides.

30 Sensorgrams are shown for sFGFR3\_Del1 (SEQ ID NO: 7), sFGFR3\_Del4 (SEQ ID NO: 1), and sFGFR3\_Del4-LK1-LK2 (SEQ ID NO: 10; Fig. 1A); sFGFR3\_Del1 (SEQ ID NO: 7) and sFGFR3\_Del1-D3 (SEQ ID NO: 9; Fig. 1B); sFGFR3\_Del4-LK1-LK2 (SEQ ID NO: 10), sFGFR3\_Del4-LK1-LK2-C253S (SEQ ID NO: 11), and sFGFR3\_Del4-LK1-LK2-D3 (SEQ ID NO: 12; Fig. 1C); and sFGFR3\_Del4 (SEQ ID NO: 1), sFGFR3\_Del4-C253S (SEQ ID NO: 2),

35 and sFGFR3\_Del4-D3 (SEQ ID NO: 33; Fig. 1D).

**FIGS. 2A-2C** are images of Western blots of the sFGFR3 polypeptides. Western blots under reducing (R) and non-reducing (NR) conditions are shown for sFGFR3\_Del1, sFGFR3\_Del1-C253S (SEQ ID NO: 8), and sFGFR3\_Del1-D3 (Fig. 2A); sFGFR3\_Del4-LK1-

LK2, sFGFR3\_Del4-LK1-LK2-C253S, and sFGFR3\_Del4-LK1-LK2-D3 (Fig. 2B); and sFGFR3\_Del4, sFGFR3\_Del4-C253S, and sFGFR3\_Del4-D3 (Fig. 2C).

**FIGS. 3A-3B** are graphs showing a sensorgram (Fig. 3A) and proliferation assays of sFGFR3\_Del4, sFGFR3\_Del4-C253S, and sFGFR3\_Del4-D3 (Fig. 3B) using *Fgfr3<sup>ach/+</sup>* chondrocyte cells in the presence of FGF2.

**FIG. 4** is a graph showing luciferase signaling in Serum Response Element-Luciferase (SRE-Luc) HEK cells expressing FGFR3<sup>G380R</sup> incubated with sFGFR3\_Del4-D3 at 0 nM, 70 nM, and 280nM with or without 1 ng/mL of hFGF2 (\* indicates p value < 0.05; \*\*\* indicates a p value < 0.001 compared to sFGFR3\_Del4-D3 at 0 nM).

**FIG. 5** is a graph showing the percentage of living animals (wild type (wt) and *Fgfr3<sup>ach/+</sup>* mice) after 3 days of treatment with a low dose (0.25 mg/kg) of sFGFR3\_Del4-D3 relative to age (days). The percentage of living wt mice receiving vehicle (PBS) is also shown.

**FIG. 6** is an image showing the amino acid residues corresponding to the Ig-like C2-type domains 1 (IgI), 2 (IgII), and 3 (IgIII) of wildtype FGFR3 polypeptide (SEQ ID NO: 5 or 32), sFGFR3\_Del4-C253S (SEQ ID NO: 2), and a variant of sFGFR3\_Del4-D3 (SEQ ID NO:33). sFGFR3\_Del4-C253S includes an amino acid substitution of a cysteine residue with a serine residue at position 253 of SEQ ID NO: 1.

**FIGS. 7A-7B** are images of Western blots of the sFGFR3 polypeptides. Western blots under reducing (R) and non-reducing (NR) conditions are shown for 2.3 mg/ml and 23 mg/ml sFGFR3\_Del1-D3 (Fig. 7A) and 1.5 mg/ml and 15 mg/ml sFGFR3\_Del1-C253S (Fig. 7B).

**FIGS. 8A-8B** are graphs showing the melting temperature ( $T_m$ ) of sFGFR3\_Del4-C253S in 20 mM phosphate, 40mM NaCl, pH 7.5 buffer and 40 mM citrate, 40mM NaCl, pH 6.5 buffer (Fig. 8A) and the  $T_m$  of sFGFR3\_Del4-D3 in 20 mM phosphate, 40mM NaCl, pH 7.5 buffer and 20 mM citrate, 40mM NaCl, pH 6.5 buffer (Fig. 8B).

**FIGS. 9A-9C** are graphs showing the fast protein liquid chromatography (FPLC) elution profiles of sFGFR3\_Del4-D3. FPLC elution profiles are shown for Fig. 9A: sFGFR3\_Del4-D3 at 0 minutes, 2 hours, and 24 hours in cpm/fraction (Fig. 9A); Fig. 9B: sFGFR3\_Del4-D3 administered by intravenous bolus at 1 minute, 15 minutes, 30 minute, 2 hours, and 24 hours in cpm/fraction and as normalized to the highest peak (shown in Fig. 9B cont.); Fig 9C: sFGFR3\_Del4-D3 administered by subcutaneous injection at 30 minutes, 2 hours, 4 hours, and 24 hours in cpm/fraction and as normalized to the highest peak (shown in Fig. 9C cont.).

**FIGS. 10A-10B** are graphs showing the percentage (%) of proliferation of *Fgfr3<sup>ach/+</sup>* chondrocyte cells in the presence of the sFGFR3 polypeptides. *Fgfr3<sup>ach/+</sup>* chondrocyte proliferation is shown for 1 ug/ml, 10 ug/ml, and 50 ug/ml of sFGFR3\_Del4-D3 (Fig. 10A) and for 1 ug/ml, 10 ug/ml, and 50 ug/ml of sFGFR3\_Del4-C253S (Fig. 10B).

**FIG. 11** is a graph showing the PK profiles of 2.5 mg/kg sFGFR3\_Del4-D3 administered subcutaneously and 2.5 mg/kg sFGFR3\_Del4-D3 administered intravenously.

**FIG. 12** is a graph showing the concentration of  $^{125}\text{I}$ - sFGFR3\_Del4-D3 in kidney, liver, spleen, lung, and heart tissue at 30 minutes, 120 minutes, and 1440 minutes after intravenous administration. The concentration is expressed as the percent of injected dose per gram (%ID/g).

5 **FIG. 13** is a graph showing the concentration of  $^{125}\text{I}$ - sFGFR3\_Del4-D3 in kidney, liver, spleen, lung, and heart tissue at 30 minutes, 120 minutes, 240 minutes, 480 minutes, and 1440 minutes after subcutaneous administration. The concentration is expressed as %ID/g.

10 **FIG. 14A-14B** are graphs showing the concentration (c) and volume of distribution ( $V_d$ ) of  $^{125}\text{I}$ -sFGFR3\_Del4-D3 in brain tissue. Shown is the c of  $^{125}\text{I}$ -sFGFR3\_Del4-D3 before and 15 after correction for vascular content and degradation at 30 minutes, 2 hours, and 24 hours after intravenous bolus (Fig. 14A) and the  $V_d$  of  $^{125}\text{I}$ -sFGFR3\_Del4-D3 and RSA at 30 minutes, 2 hours, and 24 hours after intravenous bolus (Fig. 14B).

15 **FIG. 15** is a graph showing the percentage of surviving  $Fgfr3^{ach/+}$  mice administered sFGFR3\_Del4-D3. Shown are the surviving wild type mice,  $Fgfr3^{ach/+}$  mice administered PBS as vehicle,  $Fgfr3^{ach/+}$  mice administered 2.5 mg/kg sFGFR3\_Del4-D3 once weekly,  $Fgfr3^{ach/+}$  mice administered 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and  $Fgfr3^{ach/+}$  mice administered 10 mg/kg sFGFR3\_Del4-D3 twice weekly over 22 days.

20 **FIG. 16** is a graph showing the percentage (%) of locomotor and abdominal breathing complications in  $Fgfr3^{ach/+}$  mice administered PBS as vehicle, 2.5 mg/kg sFGFR3\_Del4-D3 once weekly, 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and 10 mg/kg sFGFR3\_Del4-D3 twice weekly.

25 **FIGS. 17A-17D** are graphs and an x-ray radiograph showing the length of  $Fgfr3^{ach/+}$  mice administered sFGFR3\_Del4-D3. Shown are the axial length (Fig. 17A), tail length (Fig. 17B), and tibia length (Fig. 17C) of wild type mice administered PBS as vehicle, and  $Fgfr3^{ach/+}$  mice administered PBS as vehicle, 2.5 mg/kg sFGFR3\_Del4-D3 once weekly, 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and 10 mg/kg sFGFR3\_Del4-D3 twice weekly. Also shown is the x-ray radiograph (Fig. 17D) of wild type mice administered PBS as vehicle and  $Fgfr3^{ach/+}$  mice administered PBS as vehicle, 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and 10 mg/kg sFGFR3\_Del4-D3 twice weekly. All measurements are in millimeters (mm).

30 **FIGS. 18A-18B** are a graph showing the cranium ratio and an x-ray radiograph showing the skulls of  $Fgfr3^{ach/+}$  mice administered sFGFR3\_Del4-D3, respectively. Shown in the graph (Fig. 18A) is the cranium ratio (L/W) of wild type mice administered PBS as vehicle and  $Fgfr3^{ach/+}$  mice administered PBS as vehicle, 2.5 mg/kg sFGFR3\_Del4-D3 once weekly, 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and 10 mg/kg sFGFR3\_Del4-D3 twice weekly. Shown 35 in the x-ray radiograph (Fig. 18B) is the skulls of wild type mice administered PBS as vehicle,  $Fgfr3^{ach/+}$  mice administered PBS as vehicle, wild type mice administered 10 mg/kg sFGFR3\_Del4-D3 twice weekly, and  $Fgfr3^{ach/+}$  mice administered 10 mg/kg sFGFR3\_Del4-D3 twice weekly.

**FIGS. 19A-19E** are graphs showing the kinetic profile for the binding of different concentrations of hFGF1, FGF2, hFGF9, hFGF18, hFGF19, and hFGF21 to immobilized SFGFR3\_DEL4-D3 in real time. Shown are the kinetic profiles for binding of hFGF1 at concentrations of 0.5 nM to 12 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19A); hFGF2 at 5 concentrations of 2 nM to 10 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19B); hFGF9 at concentrations of 1 nM to 5 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19C); hFGF18 at concentrations of 1 nM to 10 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19D); hFGF19 at concentrations of 2 nM to 20 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19E); and hFGF21 at 10 concentrations of 100 nM to 10000 nM to immobilized SFGFR3\_DEL4-D3 (FIG. 19F). The darker, overlapping lines represent the 2:1 binding model used to generate the  $K_d$  values.

**FIG. 20** is an image of a Western blot of non-induced wild type ATDC5 and retrovirally infected ATDC5 cells expressing FGFR3<sup>G380R</sup>.

**FIG 21** is a graph showing the induction of proliferation of ATDC5 FGFR3<sup>G380R</sup> cells in the presence of SFGFR3\_DEL4-D3 for three experiments. Untreated ATDC5 FGFR3<sup>G380R</sup> cells 15 were used as a control.

#### DETAILED DESCRIPTION OF THE INVENTION

We have discovered that soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptides and variants thereof can be used to treat skeletal growth retardation disorders, 20 such as achondroplasia, in a patient (e.g., a human, particularly an infant, a child, or an adolescent). In particular, sFGFR3 polypeptides of the invention feature a deletion of, e.g., amino acids 289 to 400 of SEQ ID NO: 5 or amino acids 311 to 422 of SEQ ID NO: 32, to provide the following exemplary sFGFR3 polypeptides: sFGFR3\_Del4 including an amino acid 25 substitution of a cysteine residue with a serine residue at position 253 (sFGFR3\_Del4-C253S; SEQ ID NO: 2) and sFGFR3\_Del4 including an extended Ig-like C2-type domain 3 (sFGFR3\_Del4-D3; SEQ ID NO: 33) and variants thereof, such as a sFGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 4. Additionally, the sFGFR3 polypeptides may 30 include a signal peptide, such as a sFGFR3 polypeptide having the amino acid sequence of SEQ ID NO: 18 or 34. See U.S. Provisional Application No. 62/276,222 and International Application No. PCT/US16/12553 for a description of sFGFR3\_Del4 (SEQ ID NO: 1), each of which is hereby incorporated herein by reference in their entirety.

For example, sFGFR3 polypeptides and variants thereof having at least 85% sequence identity (e.g., at least 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 NO: 1 can include 35 an amino acid substitution that removes a cysteine residue at position 253 of SEQ ID NO: 1 (e.g. sFGFR3\_Del4-C253S; a polypeptide having the amino acid sequence of SEQ ID NO: 2). In particular, an sFGFR3 polypeptide of the invention can include a substitution of a cysteine residue at position 253 of SEQ ID NO: 1 with, e.g., a serine residue. For example, the cysteine

residue at position 253 is substituted with a serine residue or, e.g., another conservative amino acid substitution, such as alanine, glycine, proline, or threonine.

The sFGFR3 polypeptides can also include a polypeptide sequence having at least 90% (e.g., at least 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity)

5 amino acid sequence identity to amino acid residues 23 to 357 of SEQ ID NO: 32, in which the polypeptide lacks a signal peptide and a transmembrane domain of FGFR3 and (i) is less than 500 amino acids in length; (ii) comprises 200 consecutive amino acids or fewer of an intracellular domain of FGFR3; and/or (iii) lacks a tyrosine kinase domain of FGFR3 (e.g., sFGFR3\_Del4-D3; a polypeptide having the amino acid sequence of SEQ ID NO: 33). Methods 10 for administering the sFGFR3 polypeptides of the invention to treat skeletal growth retardation disorders (e.g., achondroplasia) in a patient (e.g., a human, particularly an infant, a child, or an adolescent) are also described.

The sFGFR3 polypeptides, methods of production, methods of treatment, compositions, and kits of the invention are described herein.

15

### **Soluble Fibroblast Growth Factor Receptor 3 (sFGFR3) Polypeptides**

The invention features sFGFR3 polypeptides and variants thereof formulated for the treatment of skeletal growth retardation disorders (e.g., achondroplasia). In particular, the sFGFR3 polypeptides can have at least 85% sequence identity (e.g., 86%, 87%, 88%, 89%, 20 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity) to the amino acid sequence of SEQ ID NO: 1, in which the sFGFR3 polypeptide includes an amino acid substitution that removes a cysteine residue at position 253 of SEQ ID NO: 1 (e.g., sFGFR3\_Del4-C253S; a polypeptide having the amino acid sequence of SEQ ID NO: 2). For example, the cysteine residue at position 253 of SEQ ID NO: 1 is substituted with a serine 25 residue or a conservative amino acid substitution, such as alanine, glycine, proline, or threonine.

The sFGFR3 polypeptides and variants thereof can also include fragments of the amino acid sequence of SEQ ID NO: 2 (e.g., at least amino acids 1 to 200, 1 to 205, 1 to 210, 1 to 215, 1 to 220, 1 to 225, 1 to 235, 1 to 230, 1 to 240, 1 to 245, 1 to 250, 1 to 253, 1 to 255, 1 to 260, 1 to 265, 1 to 275, 1 to 280, 1 to 285, 1 to 290, or 1 to 300, of SEQ ID NO: 2) having at least 50% 30 sequence identity (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 SEQ ID NO: 2. Additionally, sFGFR3 polypeptides can include amino acids 1 to 301 of SEQ ID NO: 1, in which the sFGFR3 polypeptide includes an amino acid substitution of a cysteine residue with a serine residue at position 253 of SEQ ID NO: 1 (e.g., a polypeptide having the amino acid 35 sequence of SEQ ID NO: 2).

The sFGFR3 polypeptides and variants thereof can also include fragments of the amino acid sequence of SEQ ID NO: 33 (e.g., at least amino acids 1 to 200, 1 to 210, 1 to 220, 1 to 230, 1 to 240, 1 to 250, 1 to 260, 1 to 270, 1 to 280, 1 to 290, 1 to 300, 1 to 310, 1 to 320, 1 to

330, 1 to 340, 1 to 340, or 1 to 345 of SEQ ID NO: 33) having at least 50% sequence identity (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 SEQ ID NO: 33. In addition, the cysteine residue at position 253 of SEQ ID NO: 4 or 33 and/or position 316 of SEQ ID NO: 5 4, if present, can be substituted with a serine residue or a conservative amino acid substitution, such as alanine, glycine, proline, or threonine.

Given the results described herein, the invention is not limited to a particular sFGFR3 polypeptide or variants thereof. In addition to the exemplary sFGFR3 polypeptides and variants thereof discussed above, any polypeptide that binds one or more FGFs (e.g., FGF1 (e.g., a 10 polypeptide having the amino acid sequence of SEQ ID NO: 13), FGF2 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 14), FGF9 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 15), FGF18 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 16), FGF19 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 18 or 15 38), and/or FGF21 (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 39)) with similar binding affinity as the sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be used in the methods, such as for treating a skeletal growth retardation disorder, e.g., achondroplasia. The sFGFR3 polypeptides can be, for example, fragments of FGFR3 20 isoform 2 lacking exons 8 and 9 encoding the C-terminal half of the Ig-like C2-type domain 3 and exon 10 including the transmembrane domain (e.g., fragments of the amino acid sequence of SEQ ID NO: 5 or 32), corresponding to fragments of FGFR3 transcript variant 2 (Accession No. NM\_022965).

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), 25 sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4)) can include a signal peptide at the N-terminal position. An exemplary signal peptide can include, but is not limited to, amino acids 1 to 22 of SEQ ID NO: 6 (e.g., MGAPACALALCVAVAIIVAGASS) or amino acids 1 to 19 of SEQ ID NO: 35 (e.g., MMSFVSLLLVGILFHATQA). Accordingly, the sFGFR3 polypeptides include both secreted forms, which lack the N-terminal signal peptide, and non- 30 secreted forms, which include the N-terminal signal peptide. For instance, a secreted sFGFR3 polypeptide can include the amino acid sequence of SEQ ID NOs: 2, 4, or 33. Alternatively, the sFGFR3 polypeptide does include a signal peptide, such the amino acid sequence of SEQ ID NOs: 18, 19, or 34. One skilled in the art will appreciate that the position of the N-terminal signal peptide will vary in different sFGFR3 polypeptides and can include, for example, the first 35 5, 8, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 27, 30, or more amino acid residues on the N-terminus of the polypeptide. 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 [cbs.dtu.dk/services/SignalP/](http://cbs.dtu.dk/services/SignalP/).

Additionally, sFGFR3 polypeptides (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) of the invention can be glycosylated. In particular, a sFGFR3 polypeptide can be altered to increase or decrease the extent to which the sFGFR3 polypeptide is glycosylated. Addition or deletion of glycosylation sites to an sFGFR3 polypeptide can be accomplished by altering the amino acid sequence such that one or more glycosylation sites is created or removed. For example, N-linked glycosylation, in which an oligosaccharide is attached to the amide nitrogen of an asparagine residue, can occur at position Asn76, Asn148, Asn169, Asn 203, Asn240, Asn272, and/or Asn 294 of the amino acid sequence of sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 4 or 33), and variants thereof. One or more of these Asn residues can also be substituted to remove the glycosylation site. For instance, O-linked glycosylation, in which an oligosaccharide is attached to an oxygen atom of an amino acid residue, can occur at position Ser109, Thr126, Ser199, Ser274, Thr281, Ser298, Ser299, and/or Thr301 of the amino acid sequence of sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), variants thereof (SEQ ID NO: 4), and sFGFR3 polypeptides including a signal peptide (SEQ ID NO: 18 or 34). Additionally, O-linked glycosylation can occur at position Ser310 and/or Ser321 of sFGFR3\_Del4-D3 (SEQ ID NO: 33) and variants thereof (SEQ ID NO: 4). One or more of these Ser or Thr residues can also be substituted to remove the glycosylation site.

#### *sFGFR3 Fusion Polypeptides*

sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be fused to a functional domain from a heterologous polypeptide (e.g., a fragment crystallizable region of an immunoglobulin (Fc region; such as a polypeptide having the amino acid sequence of SEQ ID NOS: 25 and 26) or human serum albumin (HSA; such as a polypeptide having the amino acid sequence of SEQ ID NO: 27)) to provide a sFGFR3 fusion polypeptide. Optionally, a flexible linker, can be included between the sFGFR3 polypeptide and the heterologous polypeptide (e.g., an Fc region or HSA), such as a serine or glycine-rich sequence (e.g., a poly-glycine or a poly-glycine/serine linker, such as SEQ ID NOS: 28 and 29).

For example, the sFGFR3 polypeptides (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be a fusion polypeptide including, e.g., an Fc region of an immunoglobulin at the N-terminal or C-terminal domain. In particular, useful Fc regions can 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) from any mammal (e.g., a human). For instance, the Fc fragment human IgG-1 (SEQ ID NO: 25) or a variant of human IgG-1, such as a variant including a substitution of asparagine at position 297 of SEQ ID NO: 25 with alanine (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 26). The Fc fragments of the invention can include, for example, the CH2 and CH3 domains of the heavy chain and any portion of the hinge region. The sFGFR3 fusion polypeptides of the invention can also include, e.g., a monomeric Fc, such as a CH2 or CH3 domain. The Fc region may optionally be glycosylated at any appropriate one or more amino acid residues known to those skilled in the art. 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.

Additionally, the sFGFR3 polypeptides (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be conjugated to other molecules at the N-terminal or C-terminal domain for the purpose of improving the solubility and stability of the protein in aqueous solution. Examples of such molecules include human serum albumin (HSA), PEG, PSA, and bovine serum albumin (BSA). For instance, the sFGFR3 polypeptides can be conjugated to human HSA (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 27) or a fragment thereof.

The sFGFR3 fusion polypeptides can include a peptide linker region between the sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) and the heterologous polypeptide (e.g., an Fc region or HSA). The linker region may be of any sequence and length that allows the sFGFR3 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. For instance, linkers include or consist of flexible portions, e.g., regions without significant fixed secondary or tertiary structure. Preferred ranges are 5 to 25 and 10 to 20 amino acids in length. Such flexibility is generally increased if the amino acids are small and do not have bulky side chains that impede rotation or bending of the amino acid chain. Thus, preferably the peptide linker of the present invention has an increased content of small amino acids, in particular of glycines, alanines, serines, threonines, leucines and isoleucines.

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-rich linkers, e.g., containing at least 50%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or even 100% serine residues. In some cases, the amino acid sequence of a

linker consists only of glycine and serine residues. For example, the linker can be the amino acid sequence of GGGGAGGGG (SEQ ID NO: 28) or GGGGSGGGGGSGGGGS (SEQ ID NO: 29). A linker can optionally be glycosylated at any appropriate one or more amino acid residues. The linker can also be absent, in which the FGFR3 polypeptide and the heterologous polypeptide (e.g., an Fc region or HSA) are fused together directly, with no intervening residues.

#### *Polynucleotides encoding the sFGFR3 Polypeptides*

The invention further includes polynucleotides encoding the sFGFR3 polypeptides (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) that can be used to treat skeletal growth retardation disorders (e.g., achondroplasia) in a patient (e.g., a human, such as an infant, a child, or an adolescent), such as SEQ ID NOs: 20, 21, 36, or 37. For example, the polynucleotide can be the nucleic acid sequence of SEQ ID NO: 20 or 36, which encode sFGFR3\_Del4-C253S (SEQ ID NO: 2), or a variant having at least 85% sequence identity (e.g., 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity) to the nucleic acid sequence of SEQ ID NO: 20 or 36. Additionally, the polynucleotide can be the nucleic acid sequence of SEQ ID NO: 21 or 37, which encodes sFGFR3\_Del4-D3 (SEQ ID NO: 33), having at least 85% sequence identity (e.g., 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more sequence identity) to the nucleic acid sequence of SEQ ID NO: 21 or 37. The invention also includes polynucleotides encoding sFGFR3 fusion polypeptides (e.g., a sFGFR3 polypeptide fused to a heterologous polypeptide, such as a Fc region or HSA) and polynucleotides encoding sFGFR3 polypeptides without a signal peptide (e.g., polypeptides having the amino acid sequence of SEQ ID NOs: 2, 4, and 33) or with a signal peptide (e.g., polypeptides having the amino acid sequence of SEQ ID NOs: 18, 19, and 34). Additionally, the invention includes polynucleotides include one or more mutations to alter any of the glycosylation sites described herein.

Optionally, the sFGFR3 polynucleotides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be codon optimized to alter the codons in the nucleic acid, in particular to reflect the typical codon usage of the host organism (e.g., a human) without altering the sFGFR3 polypeptide encoded by the nucleic acid sequence of the polynucleotide. Codon-optimized polynucleotides (e.g., a polynucleotide having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37) can, e.g., facilitate genetic manipulations by decreasing the GC content and/or for expression in a host cell (e.g., a HEK 293 cell or a CHO cell). Codon-optimization can be performed by the skilled person, e.g. by using online tools such as the JAVA Codon Adaption Tool ([www.jcat.de](http://www.jcat.de)) or Integrated DNA Technologies Tool ([www.eu.idtdna.com/CodonOpt](http://www.eu.idtdna.com/CodonOpt)) by simply entering the nucleic acid

sequence of the polynucleotide and the host organism for which the codons are to be optimized. The codon usage of different organisms is available in online databases, for example, [www.kazusa.or.jp/codon](http://www.kazusa.or.jp/codon).

5 *Host cells for expression of the sFGFR3 polypeptides*

Mammalian cells can be used as host cells for expression of the sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Exemplary mammalian cell types useful in the methods include, but 10 are not limited to, human embryonic kidney (HEK; e.g., HEK 293) cells, Chinese Hamster Ovary (CHO) cells, L cells, C127 cells, 3T3 cells, BHK cells, COS-7 cells, HeLa cells, PC3 cells, Vero cells, MC3T3 cells, NS0 cells, Sp2/0 cells, VERY cells, BHK, MDCK cells, W138 cells, BT483 cells, Hs578T cells, HTB2 cells, BT20 cells, T47D cells, NS0 cells, CRL7030 cells, and Hs78Bst cells, or any other suitable mammalian host cell known in the art. Alternatively, *E.* 15 *coli* cells can be used as host cells for expression of the sFGFR3 polypeptides. Examples of *E. coli* strains include, but are not limited to, *E. coli* 294 (ATCC® 31,446), *E. coli* λ 1776 (ATCC® 31,537, *E. coli* BL21 (DE3) (ATCC® BAA-1025), *E. coli* RV308 (ATCC® 31,608), or any other suitable *E. coli* strain known in the art.

20 *Vectors including polynucleotides encoding the sFGFR3 polypeptides*

The invention also features recombinant vectors including any one or more of the polynucleotides described above. The vectors of the invention can be used to deliver a polynucleotide encoding a sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a 25 sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), which can include mammalian, viral, and bacterial expression vectors. For example, the vectors can be plasmids, artificial chromosomes (e.g. BAG, PAC, and YAC), and virus or phage vectors, and may optionally include a promoter, enhancer, or regulator for the expression of the polynucleotide. The vectors can also contain one or more selectable marker genes, such as an ampicillin, 30 neomycin, and/or kanamycin resistance gene in the case of a bacterial plasmid or a resistance gene for a fungal vector. Vectors can be used *in vitro* for the production of DNA or RNA or used to transfect or transform a host cell, such as a mammalian host cell for the production of a sFGFR3 polypeptide encoded by the vector. The vectors can also be adapted to be used *in vivo* in a method of gene therapy.

35 Exemplary viral vectors that can be used to deliver a polynucleotide encoding a sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) include a retrovirus, adenovirus (e.g., Ad2, Ad5, Ad11,

Ad12, Ad24, Ad26, Ad34, Ad35, Ad40, Ad48, Ad49, Ad50, and Pan9 (also known as AdC68)), parvovirus (e.g., adeno-associated viruses), coronavirus, negative strand RNA viruses such as orthomyxovirus (e.g., influenza virus), rhabdovirus (e.g., rabies and vesicular stomatitis virus), paramyxovirus (e.g. measles and Sendai), positive strand RNA viruses, such as picornavirus 5 and alphavirus, and double stranded DNA viruses including adenovirus, herpesvirus (e.g., Herpes Simplex virus types 1 and 2, Epstein-Barr virus, cytomegalovirus), and poxvirus (e.g., vaccinia, modified vaccinia Ankara (MVA), fowlpox and canarypox). Other viruses useful for delivering polynucleotides encoding sFGFR3 polypeptides include Norwalk virus, togavirus, 10 flavivirus, reoviruses, papovavirus, hepadnavirus, and hepatitis virus. Examples of retroviruses include avian leukosis-sarcoma, mammalian C-type, B-type viruses, D-type viruses, HTLV-BLV group, lentivirus, and spumavirus (Coffin, J. M., *Retroviridae: The viruses and their replication*, *In Fundamental Virology*, Third Edition, B. N. Fields, et al., Eds., Lippincott-Raven Publishers, Philadelphia, 1996).

## 15 **Methods of Production**

Polynucleotides encoding sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be produced by any method known in the art. For instance, a polynucleotide is generated using molecular 20 cloning methods and is placed within a vector, such as a plasmid, an artificial chromosome, a viral vector, or a phage vector. The vector is used to transform the polynucleotide into a host cell appropriate for the expression of the sFGFR3 polypeptide.

### *Nucleic acid vector construction and host cells*

25 The sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be produced from a host cell. The polynucleotides (e.g., polynucleotides having the nucleic acid sequence of SEQ ID NO: 20, 21, 36, or 37 and variants thereof) encoding sFGFR3 polypeptides can be included in vectors 30 that can be introduced into the host cell by conventional techniques known in the art (e.g., transformation, transfection, electroporation, calcium phosphate precipitation, direct microinjection, or infection). The choice of vector depends in part on the host cells to be used. Generally, host cells are of either prokaryotic (e.g., bacterial) or eukaryotic (e.g., mammalian) origin.

35 A polynucleotide encoding an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be prepared by a variety of methods known in the art. These methods include, but are not limited to,

oligonucleotide-mediated (or site-directed) mutagenesis and PCR mutagenesis. A polynucleotide encoding an sFGFR3 polypeptide can be obtained using standard techniques, e.g., gene synthesis. Alternatively, a polynucleotide encoding a wild-type sFGFR3 polypeptide (e.g., a polypeptide having the amino acid sequence of SEQ ID NO: 5 or 32) can be mutated to 5 contain specific amino acid substitutions (e.g., an amino acid substitution of a cysteine residue with a serine residue or a conservative amino acid substitution, such as alanine, glycine, proline, or threonine, at position 253 of SEQ ID NO: 33 and/or position 316 of SEQ ID NO: 4) using standard techniques in the art, e.g., QuikChange<sup>TM</sup> mutagenesis. Polynucleotides encoding an sFGFR3 polypeptide can be synthesized using, e.g., a nucleotide synthesizer or PCR 10 techniques.

Polynucleotides encoding sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be inserted into a vector capable of replicating and expressing the polynucleotide in prokaryotic or 15 eukaryotic host cells. Exemplary vectors useful in the methods can include, but are not limited to, a plasmid, an artificial chromosome, a viral vector, and a phage vector. For example, a viral vector can include the viral vectors described above, such as a retroviral vector, adenoviral vector, or poxviral vector (e.g., vaccinia viral vector, such as Modified Vaccinia Ankara (MVA)), 20 adeno-associated viral vector, and alphaviral vector)) containing the nucleic acid sequence of a polynucleotide encoding the sFGFR3 polypeptide. Each vector can contain various components that may be adjusted and optimized for compatibility with the particular host cell. For example, the vector components may include, but are not limited to, an origin of replication, 25 a selection marker gene, a promoter, a ribosome binding site, a signal sequence, the nucleic acid sequence of the polynucleotide encoding the sFGFR3 polypeptide, and/or a transcription termination sequence.

The above-described vectors may be introduced into appropriate host cells (e.g., HEK 293 cells or CHO cells) using conventional techniques in the art, e.g., transformation, transfection, electroporation, calcium phosphate precipitation, and direct microinjection. Once the vectors are introduced into host cells for the production of an sFGFR3 polypeptide of the 30 invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), host cells are cultured in conventional nutrient media modified as appropriate for inducing promoters, selecting transformants, or amplifying the polynucleotides (e.g. SEQ ID NOs: 20 and 21 and variants thereof) encoding the sFGFR3 polypeptide. Methods for 35 expression of therapeutic proteins, such as sFGFR3 polypeptides, are known in the art, see, for example, Paulina Balbas, Argelia Lorence (eds.) *Recombinant Gene Expression: Reviews and Protocols (Methods in Molecular Biology)*, Humana Press 2nd ed. 2004 (July 20, 2004) and Vladimir Voynov and Justin A. Caravella (eds.) *Therapeutic Proteins: Methods and Protocols*

(*Methods in Molecular Biology*) Humana Press 2nd ed. 2012 (June 28, 2012), each of which is hereby incorporated by reference in its entirety.

*sFGFR3 polypeptide production, recovery, and purification*

5 Host cells (e.g., HEK 293 cells or CHO cells) used to produce the sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be grown in media known in the art and suitable for culturing of the selected host cells. Examples of suitable media for mammalian host cells include Minimal  
10 Essential Medium (MEM), Dulbecco's Modified Eagle's Medium (DMEM), Expi293™ Expression Medium, DMEM with supplemented fetal bovine serum (FBS), and RPMI-1640. Examples of suitable media for bacterial host cells include Luria broth (LB) plus necessary supplements, such as a selection agent, e.g., ampicillin. Host cells are cultured at suitable temperatures, such as from about 20 °C to about 39 °C, e.g., from 25 °C to about 37 °C, preferably 37 °C, and  
15 CO<sub>2</sub> levels, such as 5 to 10% (preferably 8%). The pH of the medium is generally from about 6.8 to 7.4, e.g., 7.0, depending mainly on the host organism. If an inducible promoter is used in the expression vector, sFGFR3 polypeptide expression is induced under conditions suitable for the activation of the promoter.

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2),  
20 sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be recovered from the supernatant of the host cell. Alternatively, the sFGFR3 polypeptide can be recovered by disrupting the host cell (e.g., using osmotic shock, sonication, or lysis), followed by centrifugation or filtration to remove the sFGFR3 polypeptide. Upon recovery of the sFGFR3  
25 polypeptide, the sFGFR3 polypeptide can then be further purified. An sFGFR3 polypeptide can be purified by any method known in the art of protein purification, such as protein A affinity, other chromatography (e.g., ion exchange, affinity, and size-exclusion column chromatography), centrifugation, differential solubility, or by any other standard technique for the purification of proteins (see *Process Scale Purification of Antibodies*, Uwe Gottschalk (ed.) John Wiley &  
30 Sons, Inc., 2009, hereby incorporated by reference in its entirety).

35 Optionally, the sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be conjugated to a detectable label for purification. Examples of suitable labels for use in purification of the sFGFR3 polypeptides include, but are not limited to, a protein tag, a fluorophore, a chromophore, a radiolabel, a metal colloid, an enzyme, or a chemiluminescent, or bioluminescent molecule. In particular, protein tags that are useful for purification of the sFGFR3 polypeptides can include, but are not limited to, chromatography tags (e.g., peptide

tags consisting of polyanionic amino acids, such as a FLAG-tag, or a hemagglutinin "HA" tag), affinity tags (e.g., a poly(His) tag, chitin binding protein (CBP), maltose binding protein (MBP), or glutathione-S-transferase (GST)), solubilization tags (e.g., thioredoxin (TRX) and poly(NANP)), epitope tags (e.g., V5-tag, Myc-tag, and HA-tag), or fluorescence tags (e.g., GFP, GFP variants, 5 RFP, and RFP variants).

### Methods of Treatment

Provided herein are methods for treating a skeletal growth retardation disorder in a patient, such as a patient having achondroplasia (e.g., a human having achondroplasia). In 10 particular, the patient is one that exhibits or is likely to develop one or more symptoms of a skeletal growth retardation disorder (e.g., achondroplasia). The method involves administering an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) to the patient having a skeletal 15 growth retardation disorder, such as a patient having achondroplasia (e.g., a human having achondroplasia). In particular, the method involves administering sFGFR3\_Del4-C253S (SEQ ID NO: 2) or sFGFR3\_Del4-D3 (SEQ ID NO: 33) to the patient having a skeletal growth retardation disorder, such as a patient having achondroplasia (e.g., a human having achondroplasia). For example, the patient is an infant or child having a skeletal growth 20 retardation disorder, such as an infant, a child, or an adolescent having achondroplasia (e.g., a human having achondroplasia).

The patient (e.g., a human) can be treated before symptoms of a skeletal growth retardation disorder (e.g., achondroplasia) appear or after symptoms of a skeletal growth retardation disorder (e.g., achondroplasia) develop. In particular, patients that can be treated 25 with a sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) are those exhibiting symptoms including, but not limited to, short limbs, short trunk, bowlegs, a waddling gait, skull 30 malformations, cloverleaf skull, craniosynostosis, wormian bones, anomalies of the hands, anomalies of the feet, hitchhiker thumb, and/or chest anomalies. Furthermore, treatment with an sFGFR3 polypeptide can result in an improvement in one or more of the aforementioned symptoms of a skeletal growth retardation disorder (e.g., relative to an untreated patient), such as achondroplasia.

The patient (e.g., a human) can be diagnosed with a skeletal growth retardation disorder, 35 such as achondroplasia, before administration of an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or

34)). Additionally, the patient having a skeletal growth retardation disorder, such as achondroplasia, can be one that has not previously been treated with an sFGFR3 polypeptide.

#### *Skeletal Growth Retardation Disorders*

5       Skeletal growth retardation disorders can be treated by administering an sFGFR3 polypeptide as described herein to a patient (e.g., a human) in need thereof. The method involves administering to the patient (e.g., a human) having the skeletal growth retardation disorder an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 10 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Skeletal growth retardation disorders that can be treated with the sFGFR3 polypeptides are characterized by deformities and/or malformations of the bones and can include, but are not limited to, FGFR3-related skeletal diseases. In particular, the patient is treated with sFGFR3\_Del4-C253S (SEQ ID NO: 2) or sFGFR3\_Del4-D3 (SEQ ID NO: 33).

15       Administration of an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can treat a skeletal growth retardation disorder including, but not limited to, achondroplasia, achondrogenesis, acrodysostosis, acromesomelic dysplasia, atelosteogenesis, camptomelic dysplasia, 20 chondrodysplasia punctata, rhizomelic type of chondrodysplasia punctata, cleidocranial dysostosis, congenital short femur, Crouzon syndrome, Apert syndrome, Jackson-Weiss syndrome, Pfeiffer syndrome, Crouzonodermoskeletal syndrome, dactyly, brachydactyly, camptodactyly, polydactyly, syndactyly, diastrophic dysplasia, dwarfism, dyssegmental dysplasia, enchondromatosis, fibrochondrogenesis, fibrous dysplasia, hereditary multiple 25 exostoses, 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 (such as type 1 (e.g., with bone manifestations or without bone 30 manifestations), type 2, or schwannomatosis), osteoarthritis, osteochondrodysplasia, osteogenesis imperfecta, 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, and spondyloepimetaphyseal dysplasia.

35       For instance, the sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be used to treat symptoms associated with a skeletal growth retardation disorder, including the disorders

described above, such as achondroplasia. Non-limiting examples of symptoms of skeletal growth retardation disorders that can be treated with the sFGFR3 polypeptides, include short limbs and trunk, bowlegs, a waddling gait, skull malformations (e.g., a large head), cloverleaf skull, craniosynostosis (e.g., premature fusion of the bones in the skull), wormian bones (e.g., 5 abnormal thread-like connections between the bones in the skull), anomalies of the hands and feet (e.g., polydactyly or extra fingers), "hitchhiker" thumbs and abnormal fingernails and toenails, and chest anomalies (e.g., pear-shaped chest or narrow thorax). Additional symptoms that can be treated by administering sFGFR3 polypeptides can also include non-skeletal 10 abnormalities in patients having skeletal growth retardation disorders, such as anomalies of the eyes, mouth, and ears, such as congenital cataracts, myopia, cleft palate, or deafness; brain malformations, such as hydrocephaly, 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 disabilities.

15 Treatment with the sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can also increase survival of patients (e.g., humans) with skeletal growth retardation disorders (e.g., achondroplasia). For example, the survival rate of patients treated with the sFGFR3 20 polypeptides can increase by at least 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, or more relative to, e.g., an untreated patient with a skeletal growth retardation disorder (e.g., achondroplasia), over a treatment period of days, months, years, or longer. In particular, administration of sFGFR3\_Del4-D3 can increase survival of patients (e.g., humans) with skeletal growth retardation disorders (e.g., relative to an untreated patient), such as achondroplasia.

25 Any skeletal growth retardation disorder that is a FGFR3-related skeletal disease (e.g., caused by or associated with overactivation of FGFR3 as result of a gain-of-function FGFR3 mutation) can be treated by administering an sFGFR3 polypeptide of the invention ((e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) to a patient (e.g., a human). For example, FGFR3-related skeletal diseases can include, 30 but are not limited to, achondroplasia, thanatophoric dysplasia type I (TDI), thanatophoric dysplasia type II (TDII), severe achondroplasia with developmental delay and acanthosis nigricans (SADDAN), hypochondroplasia, and craniosynostosis (e.g., Muenke syndrome, Crouzon syndrome, and Crouzonodermoskeletal syndrome).

35 Patients (e.g., humans) with mutations in the *FGFR3* gene associated with different FGFR3-related skeletal disorders, such as achondroplasia, hypochondroplasia, SADDAN, TDI, and TDII, can be treated with sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a

sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). For example, the sFGFR3 polypeptides can be administered to treat achondroplasia resulting from the G380R, G375C, G346E or S279C mutations of the *FGFR3* gene. Administration of the sFGFR3 polypeptides can be used to treat the following exemplary FGFR3-related skeletal disorders:

5 hypochondroplasia resulting from the G375C, G346E or S279C mutations of the *FGFR3* gene; TDI resulting from the R248C, S248C, G370C, S371C, Y373C, X807R, X807C, X807G, X807S, X807W and K650M mutations of the *FGFR3* gene; TDII resulting from the K650E mutation of the *FGFR3* gene; and SADDAN resulting from the K650M mutation of the *FGFR3* gene.

Any of the aforementioned mutations in the *FGFR3* gene (e.g., the G380R mutation of the *FGFR3* gene) can be detected in a sample from the patient (e.g., a human with achondroplasia, hypochondroplasia, SADDAN, TDI, and TDII) prior to or after treatment with an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Additionally, the parents of the patient and/or fetal samples (e.g., fetal nucleic acid obtained from maternal blood, placental, or fetal samples) can be tested by methods known in the art for the mutation in the *FGFR3* gene to determine their need for treatment.

#### *Achondroplasia*

20 Achondroplasia is the most common cause of dwarfism in humans and can be treated by administering sFGFR3 polypeptides as described herein. In particular, achondroplasia can be treated by administering an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Accordingly, 25 administration of the sFGFR3 polypeptides can result in an improvement in symptoms including, but not limited to, growth retardation, skull deformities, orthodontic defects, 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, respiratory problems, fatigue, 30 pain, numbness in the lower back and/or spine, and/or obesity.

Patients treated using the sFGFR3 polypeptides of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can include infants, children, and adults with achondroplasia. In particular, infants are often diagnosed with 35 achondroplasia at birth, and thus, treatment with the sFGFR3 polypeptides can begin as early as possible in the patient's life, e.g., shortly after birth, or prior to birth (*in utero*).

Symptoms of achondroplasia in patients (e.g., humans) can also be monitored prior to or after a patient is treated with an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-

C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). For instance, symptoms of achondroplasia can be monitored prior to treatment to assess the severity of achondroplasia and condition of the patient prior to performing the methods.

5 The methods can include diagnosis of achondroplasia in a patient and monitoring the patient for changes in the symptoms of achondroplasia, such as changes in body weight and skull size (e.g., skull length and/or skull width) of the patient. Changes in body weight and skull size can be monitored over a period of time, e.g., 1, 2, 3, 4 or more times per month or per year or approximately every 1, 2, 3, 4, 5, 6, 7, 8, 12 or 16 weeks over the course of treatment with  
10 the sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Body weight and/or skull size of the patient having achondroplasia can also be determined at treatment specific events, such as before and/or after administration of the sFGFR3 polypeptide.

15 For example, body weight and/or skull size can be measured in response to administration of the sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)). Body weight can be measured by weighing the patient having achondroplasia on a scale, preferably in a standardized manner,  
20 such as with the same or no clothes or at a certain time of the day, preferably in a fasting state (e.g., in the morning before breakfast or after at least 1, 2, 3, 4, 5 or more hours of fasting).  
Skull size can be represented by length, height, width, and/or circumference of the skull. Measurements can be performed using any known or self-devised standardized method. For a human subject, the measurement of skull circumference is preferred, which can be measured  
25 using a flexible and non-stretchable material, such as a tape, wrapped around the widest possible circumference of the head (e.g. from the most prominent part of the forehead around to the widest part of the back of the head). The height of the skull of the subject (e.g., human) can also be determined from the underside of the chin to the uppermost point of the head.  
Preferably, any measurement is performed more than once, e.g. at least 2, 3, 4, 5, 6, 7, 8, 9, 10,  
30 or more times.

#### *Administration of sFGFR3 Polypeptides*

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be administered by any route known in the art, such as by parenteral administration, enteral administration, or topical administration. In particular, the sFGFR3 polypeptide can be administered to the patient having a skeletal growth retardation disorder (e.g., achondroplasia) subcutaneously (e.g., by

subcutaneous injection), intravenously, intramuscularly, intra-arterially, intrathecally, or intraperitoneally.

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be administered to a patient (e.g., a human) at a predetermined dosage, such as in an effective amount to treat a skeletal growth retardation disorder (e.g., achondroplasia), without inducing significant toxicity. For example, sFGFR3 polypeptides can be administered to a patient having skeletal growth retardation disorders (e.g., achondroplasia) in individual doses ranging from about 0.002 mg/kg to about 50 mg/kg (e.g., from 2.5 mg/kg to 30 mg/kg, from 0.002 mg/kg to 20 mg/kg, from 0.01 mg/kg to 2 mg/kg, from .2 mg/kg to 20 mg/kg, from 0.01 mg/kg to 10 mg/kg, from 10 mg/kg to 100 mg/kg, from 0.1 mg/kg to 50 mg/kg, 0.5 mg/kg to 20 mg/kg, 1.0 mg/kg to 10 mg/kg, 1.5 mg/kg to 5 mg/kg, or 0.2 mg/kg to 3 mg/kg). In particular, the sFGFR3 polypeptide can be administered in individual doses of, e.g., 0.001 mg/kg to 50 mg/kg, such as 2.5 mg/kg to about 10 mg/kg.

Exemplary doses of an sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) for administration to a patient (e.g., a human) having a skeletal growth retardation disorder (e.g., achondroplasia) include, e.g., 0.005, 0.01, 0.02, 0.05, 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, 1, 2, 2.5, 3, 3.5, 4, 4.5, 5, 5.5, 6, 6.5, 7, 7.5, 8, 8.5, 9, 9.5, 10, 11, 12, 13, 14, 15, 20, 25, 30, 35, 40, 45, or 50 mg/kg. These doses can be administered one or more times (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 or more times) per day, week, month, or year. For example, an sFGFR3 polypeptide can be administered to patients in a weekly dosage ranging, e.g., from about 0.0014 mg/kg/week to about 140 mg/kg/week, e.g., about 0.14 mg/kg/week to about 105 mg/kg/week, or, e.g., about 1.4 mg/kg/week to about 70 mg/kg/week (e.g., 2.5 mg/kg/week, 5 mg/kg/week, 10 mg/kg/week, 20 mg/kg/week, 30 mg/kg/week, 40 mg/kg/week, or 50 mg/kg/week).

### Gene Therapy

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can also be delivered through gene therapy, where a polynucleotide encoding the sFGFR3 polypeptide 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 Miner. Res. 26: 135-142, 2011), each of which is hereby incorporated by reference.

An sFGFR3 polypeptide of the invention (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)) can be produced by the cells of a patient (e.g., a human) having a skeletal growth retardation disorder (e.g., achondroplasia) by 5 administrating a vector (e.g., a plasmid, an artificial chromosome (e.g. BAG, PAC, and YAC), or a viral vector) containing the nucleic acid sequence of a polynucleotide encoding the sFGFR3 polypeptide. For example, a viral vector can be a retroviral vector, adenoviral vector, or poxviral vector (e.g., vaccinia viral vector, such as Modified Vaccinia Ankara (MVA)), adeno-associated viral vector, or alphaviral vector. The vector, once inside a cell of the patient (e.g., a human) 10 having a skeletal growth retardation disorder (e.g., achondroplasia), by, e.g., transformation, transfection, electroporation, calcium phosphate precipitation, or direct microinjection, will promote expression of the sFGFR3 polypeptide, which is then secreted from the cell. The invention further includes cell-based therapies, in which the patient (e.g., a human) is administered a cell expressing the sFGFR3 polypeptide.

15

### Pharmaceutical Compositions

Pharmaceutical compositions of the invention can include an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants 20 thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide, vector, and/or host cell of the invention. Compositions including an sFGFR3 polypeptide, polynucleotide, vector, and/or host cell can be formulated at a range of dosages, in a variety of formulations, and in combination with pharmaceutically acceptable excipients, carriers, or diluents.

25 A pharmaceutical composition including an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide, vector, and/or host cell of the invention can be formulated at a specific dosage, such as a dosage that is effective for treating a patient (e.g., a human) skeletal growth retardation disorder 30 (e.g., achondroplasia), without inducing significant toxicity. For example, the compositions can be formulated to include between about 1 mg/mL and about 500 mg/mL of the sFGFR3 polypeptide (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 of the sFGFR3 polypeptide).

35 The pharmaceutical compositions including an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide, vector, and/or host cell of the invention can be prepared in a variety of forms, such as a liquid

solution, dispersion or suspension, powder, or other ordered structure suitable for stable storage. For example, compositions including an sFGFR3 polypeptide intended for systemic or local delivery can be in the form of injectable or infusible solutions, such as for parenteral administration (e.g., subcutaneous, intravenous, intramuscular, intra-arterial, intrathecal, or 5 intraperitoneal administration). sFGFR3 compositions for injection (e.g., subcutaneous or intravenous injection) can be formulated using a sterile solution or any pharmaceutically acceptable liquid as a vehicle. Pharmaceutically acceptable vehicles include, but are not limited to, sterile water, physiological saline, and cell culture media (e.g., Dulbecco's Modified Eagle Medium (DMEM),  $\alpha$ -Modified Eagles Medium ( $\alpha$ -MEM), F-12 medium). Formulation methods 10 are known in the art, see e.g., Banga (ed.) *Therapeutic Peptides and Proteins: Formulation, Processing and Delivery Systems* (2nd ed.) Taylor & Francis Group, CRC Press (2006), which is hereby incorporated by reference in its entirety.

Compositions including an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 15 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide, vector, and/or host cell of the invention can be provided to patients (e.g., humans) having skeletal growth retardation disorders (e.g. achondroplasia) in combination with pharmaceutically acceptable excipients, carriers, or diluents. Acceptable excipients, carriers, or diluents can include buffers, 20 antioxidants, preservatives, polymers, amino acids, and carbohydrates. Aqueous excipients, carriers, or diluents can include water, water-alcohol solutions, emulsions or suspensions including saline, buffered medical parenteral vehicles including sodium chloride solution, Ringer's dextrose solution, dextrose plus sodium chloride solution, Ringer's solution containing lactose, and fixed oils. Examples of non-aqueous excipients, carriers, or diluents are propylene glycol, polyethylene glycol, vegetable oil, fish oil, and injectable organic esters.

25 Pharmaceutically acceptable salts can also be included in the compositions including an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide, vector, and/or host cell of the invention. Exemplary pharmaceutically acceptable salts can include mineral acid salts (e.g., 30 hydrochlorides, hydrobromides, phosphates, and sulfates) and salts of organic acids (e.g., acetates, propionates, malonates, and benzoates). Additionally, auxiliary substances, such as wetting or emulsifying agents and pH buffering substances, can be present. A thorough discussion of pharmaceutically acceptable excipients, carriers, and diluents is available in *Remington: The Science and Practice of Pharmacy*, 22<sup>nd</sup> Ed., Allen (2012), which is hereby 35 incorporated by reference in its entirety.

Pharmaceutical compositions including an sFGFR3 polypeptide (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotide,

vector, and/or host cell of the invention can also be formulated with a carrier that will protect the sFGFR3 polypeptide against rapid release, such as a controlled release formulation, including implants and microencapsulated delivery systems. For example, the sFGFR3 composition can be entrapped in microcapsules prepared by coacervation techniques or by interfacial

5 polymerization, such as hydroxymethylcellulose, gelatin, or poly-(methylmethacrylate) microcapsules; colloidal drug delivery systems (e.g., liposomes, albumin microspheres, microemulsions, nano-particles, or nanocapsules); or macroemulsions. Additionally, an sFGFR3 composition can be formulated as a sustained-release composition. For example, sustained-release compositions can include semi-permeable matrices of solid hydrophobic

10 polymers containing the sFGFR3 polypeptides, polynucleotides, vectors, or host cells of the invention, in which the matrices are in the form of shaped articles, such as films or microcapsules.

## Kits

15 Kits of the invention can include one or more sFGFR3 polypeptides (e.g. sFGFR3\_Del4-C253S (SEQ ID NO: 2), sFGFR3\_Del4-D3 (SEQ ID NO: 33), and variants thereof (SEQ ID NO: 4) or a sFGFR3 polypeptide including a signal peptide (SEQ ID NO: 18 or 34)), polynucleotides, vectors, and/or cells of the invention as described herein. For example, the sFGFR3 polypeptide, polynucleotide, vector, and/or cell can be present in a container (e.g., a glass vial)

20 in liquid form (e.g., in water or a buffered salt solution, such as, 2 mM to 20 mM of sodium phosphate, pH 6.5 or 7.0, and 25 mM to 250 mM sodium chloride). Alternatively, the sFGFR3 polypeptide, polynucleotide, and/or vector is present in a container (e.g., a glass vial) in lyophilized form, which can optionally include a diluent (e.g., water or a buffered salt solution) for reconstitution of the lyophilized sFGFR3 polypeptide, polynucleotide, vector, and/or cell into

25 liquid form prior to administration. The sFGFR3 polypeptide, polynucleotide, vector, and/or cell can also be present in a kit in another formulation as described herein. The kit components can be provided in dosage form to facilitate administration, and optionally, can include materials required for administration and/or instructions for patient treatment consistent with the methods. For example, the kit can include instructions for use, which guides the user (e.g., the physician)

30 with respect to the administration of the sFGFR3 polypeptide, polynucleotide, vector, and/or cell.

## EXAMPLES

The following examples are intended to illustrate, rather than limit, the disclosure. These studies feature the administration of the sFGFR3 polypeptides of sFGFR3\_Del4-C253S (SEQ ID NO: 2) and sFGFR3\_Del4-D3 (SEQ ID NO: 33) to patients (e.g., humans) having achondroplasia, to treat achondroplasia and symptoms associated therewith.

### Example 1: Production of sFGFR3 Polypeptides

sFGFR3\_Del4-C253S (SEQ ID NO: 2) and sFGFR3\_Del4-D3 (SEQ ID NO: 33) were produced by transient transfection in three different suspension cell types: HEK 293 freestyle, CHO-S freestyle cells and Expi CHO-S cells. For production in HEK 293 freestyle and CHO-S freestyle cells, transfection was performed using polyethylenimine (PEIpro® - Polyplus-transfection), 5 according to the manufacturer's directions. Proteins were harvested after three days. For sFGFR3 polypeptide production in Expi CHO-S cells, transfection was performed using Expifectamine as described by the manufacturer using the High Titer production protocol. A time course was performed and sFGFR3 polypeptides were optimally harvested after 12 days. Western blots were then performed using 50 ng of sFGFR3 polypeptide. Classical western blot 10 protocols were used with B9 as a primary antibody (anti FGFR3, sc-13121, Santa Cruz) diluted 1:2000 in blocking buffer and anti-mouse IgG secondary antibody (Anti-mouse IgG, #7076, Cell signaling) diluted 1:5000 in blocking buffer.

### **Example 2: Purification of sFGFR3 Polypeptides**

15 sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 were each purified using a two-step purification process including ion exchange chromatography and size exclusion chromatography.

For ion exchange chromatography, 300 mL of culture supernatant was purified by cross flow filtration (ÄKTA™ flux, GE Healthcare) using 5 µm and 0.2 µm capsules (KGF-A0504 TT 20 and KMP-HEC 9204 TT, GE Healthcare, respectively). The purified sample including sFGFR3\_Del4-C253S or sFGFR3\_Del4-D3 was then loaded on an equilibrated column at 20 mL/min, after adjusting the sample's conductivity to 14 mS/cm (ÄKTA™ pure 25 (GE Healthcare)). Columns used were HiPrep Q FF 26/10 (GE Healthcare) with a bed volume of 53 mL. The binding buffer was 1X PBS and the elution buffer was PBS 1X + 1 M NaCl. The 25 column was washed with four column volumes of 1X PBS. Elution of sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 was performed by two steps of 5% NaCl and 10% NaCl using four column volumes of each. Both 5% NaCl and 10% NaCl were pooled and concentrated by cross flow filtration (ÄKTA™ flux, GE Healthcare). The remaining volume was then concentrated on a 30 kDa filter by centrifugation at 4°C, 3,900 g for 10 min (MILLIPORE® UFC903024 AMICON® 30 Ultra-15 Centrifugal Filter Concentrator). For size exclusion chromatography, the remaining volume was loaded on a HiLoad 26/600 SUPERDEX™ 200 prep grade (28-9893-36, GE Healthcare) with a bed volume of 320 mL. Loading volume did not exceed 12.8 mL. Elution was performed in 1X PBS.

35 **Example 3: Kinetic Assays and Dissociation Constant (K<sub>d</sub>) Measurements of sFGFR3 Polypeptides**

Calibration Free Concentration Analysis and kinetic assays of sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 were performed with a Sensor Chip CM5 (GE Healthcare). Human FGF2

(hFGF2) was covalently immobilized to the Sensor Chip CM5 at a level of about 5000 RU by amine coupling. To achieve 5000 RU, hFGF2 was immobilized for 420 seconds at a flow rate 10  $\mu$ l/min and a concentration 25  $\mu$ g/ml. Running buffer was HBS-EP+ Buffer (GE Healthcare). Regeneration buffer was 100mM sodium acetate with 2M sodium chloride pH 4.5. FGF binding, 5 dissociation constant ( $K_d$ ) measurements, and kinetic parameters were determined by Surface Plasmon Resonance using a BIACORE™ T200 (GE Healthcare). The model used for kinetic assays and  $K_d$  determination was a 1:1 binding algorithm.

#### **Example 4: Proliferation Assays of sFGFR3 Polypeptides**

10 Both ATDC5 and ATDC5 FGFR3<sup>G380R</sup> cell lines were seeded at a density of 25,000 cells/cm<sup>2</sup> in NUNC™ MICROWELL™ 96-Well Optical-Bottom Plates with Polymer Base (ThermoFisher Scientific, Catalog No. 165305). After a 24 hour incubation period, cells were depleted for 48 hour in 0.5 % BSA and then stimulated for 72 hour with sFGFR3\_Del4-C253S or sFGFR3\_Del4-D3 with and without hFGF2 (Peprotech). Cell proliferation was then measured 15 using the CyQUANT® Direct Cell Proliferation Assay (Molecular Probes, Catalog No. C35012). After stimulation, 10 $\mu$ L of CyQUANT® Direct Cell Proliferation (Invitrogen; 1mL 1X PBS, 250 $\mu$ L background suppressor, and 50 $\mu$ L nuclear stain) was added per well. ATDC5 and ATDC5 FGFR3<sup>G380R</sup> cells were then incubated at room temperature in the dark for 2 hours. Fluorescence was read using the VARIOSKAN™ LUX multimode microplate reader 20 (ThermoFisher Scientific).

#### **Example 5: Luciferase Assays of sFGFR3 Polypeptides**

Serum Response Element-Luciferase (SRE-Luc) HEK cells expressing FGFR3<sup>G380R</sup> were seeded at a density of 100,000 cells/cm<sup>2</sup> in a standard culture 96 well plate. Cells were then 25 depleted for 24 hours with 0.5% heat inactivated Fetal Bovine Serum (hiFBS), before being treated with sFGFR3\_Del4-D3 at concentrations of 0 nm, 70 nm, and 280 nm with or without 1 ng/ml of hFGF2 for 24h. The culture plate was equilibrated to room temperature for 15 minutes prior to adding 100 $\mu$ L per well of Firefly Luc One-Step Glow Assay Working Solution (ThermoFisher Scientific, Catalog No. 16197), then shaken at 600 rpm for 3 minutes. The plate 30 was incubated at room temperature for 10 minutes and each cell lysate was transferred to a white opaque 96 well plate to increase luminescence signal and decrease cross contamination. The luminescence signal was read using the VARIOSKAN™ LUX multimode microplate reader (ThermoFisher Scientific).

35 **Example 6: *In vivo* Efficacy Study of sFGFR3 Polypeptides**

Experiments were performed on transgenic *Fgfr3*<sup>ach/+</sup> animals in which expression of the mutant FGFR3 is driven by the Col2a1 promoter/enhancer. Mice were exposed to a 12 hour light/dark cycle and had free access to standard laboratory food and water. Genotypes were

verified by PCR of genomic DNA using the primers 5'-AGGTGGCCTTGACACCTACCAGG-3' (SEQ ID NO: 30) and 5'-TCTGTTGTTCCTCCCTGTTGG-3' (SEQ ID NO: 31), which amplify 360 bp of the FGFR3 transgene.

5 sFGFR3\_Del4-D3 produced using CHO cells was evaluated at a subcutaneous dose of 0.25 mg/kg twice weekly. At day 3, all newborn mice from a single litter received the same dose. Control litters received 10  $\mu$ l of PBS (vehicle). Thereafter, subcutaneous injections of sFGFR3\_Del4-D3 (0.25 mg/kg) were administered twice a week for three weeks, alternatively on the left and right sides of the back. Mice were observed daily with particular attention to locomotion and urination alterations. Breeding was performed to generate litters with half wild 10 type and half heterozygous *Fgfr3<sup>ach/+</sup>* mice. To avoid bias due to phenotype penetrance variations, experiments were performed on at least two litters (one treated and one control) from the same breeders. Previous data indicated there was no statistical difference between males and females, and thus, males and females were considered one group for all analyses.

15 At day 22, all animals were sacrificed by lethal injection of pentobarbital, and gender was determined. All subsequent measurements and analyses were performed without knowledge of mice genotype to avoid investigator bias. Genotyping was performed at the end of the study to reveal the correspondence of data with a specific genotype. Since achondroplasia is a disease with phenotypic variability, all animals were included in the study. Animals dead before day 22 were used to investigate the impact of treatment on premature death. Surviving animals at day 20 22 were used for all analyses. All experiments and data measurements were performed by blinded experimenters at all time points.

25 Following sacrifice at day 22, body weights were measured. Cadavers were carefully skinned, eviscerated, and skeletal measurements were performed based on X-rays. Organs were harvested, weighed, and stored in 10% formalin for further histological analysis using standard paraffin-embedded techniques. Organs were then observed for macroscopic abnormalities, such as modification of color or texture and presence of nodules. The Principles of Laboratory Animal Care (NIH publication no. 85-23, revised 1985; <http://grants1.nih.gov/grants/olaw/references/phspol.htm>) and the European commission guidelines for the protection of animals used for scientific purposes (30 [http://ec.europa.eu/environment/chemicals/lab\\_animals/legislation\\_en.htm](http://ec.europa.eu/environment/chemicals/lab_animals/legislation_en.htm)) were followed during all animal experiments. All procedures were approved by the Institutional Ethic Committee for the use of Laboratory Animals (CIEPAL Azur) (approval # NCE-2012-52).

#### **Example 7: The Cell Line used to produce sFGFR3 Polypeptides did not impact Activity**

35 The FGF2 binding activity, Kd, and effect on cellular signaling of sFGFR3\_Del1 (SEQ ID NO: 7), sFGFR3\_Del4 (SEQ ID NO: 1), and sFGFR3\_Del4-LK1-LK2 (SEQ ID NO: 10) produced in suspension HEK 293 cells or CHO cells were compared. HEK 293 cells or CHO cells differ in post-translation modification of proteins. Expression of the sFGFR3 polypeptides in different

cell lines did not impact Kd, binding activity, or the effect of the sFGFR3 polypeptides on intracellular signaling inhibition (FIGS. 1A-1D).

**Example 8: Improved Production of sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3**

5 The sFGFR3 polypeptides of sFGFR3\_Del1 (SEQ ID NO: 7), sFGFR3\_Del4 (SEQ ID NO: 1), and sFGFR3\_Del4-LK1-LK2 (SEQ ID NO: 10) were each modified to include either an amino acid substitution of a cysteine residue with a serine residue at position 253 or an extended Ig-like C2-type domain 3 (SEQ ID NO: 33). These modifications of sFGFR3\_Del1 and sFGFR3\_Del4-LK1-LK2 had no or minimal effect on production of the sFGFR3 polypeptides, 10 since aggregation was still visible (FIGS. 2A and 2B, respectively). Surprisingly, modification of sFGFR3\_Del4 to include either an amino acid substitution of a cysteine residue with a serine residue at position 253 (sFGFR3\_Del4-C253S) or an extended Ig-like C2-type domain 3 (SEQ ID NO: 33)) improved production of the sFGFR3 polypeptides. In particular, there was minimal aggregation of sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 under both reducing and non- 15 reducing conditions (FIG. 2C). The inclusion of C253S or D3 also resulted in a relative increase in production compared to sFGFR3\_Del4, a two-fold increase in sFGFR3\_Del4-C253S production and a 3-fold increase in sFGFR3\_Del4-D3 production.

Additionally, sFGFR3\_Del4, sFGFR3\_Del4-C253S, and sFGFR3\_Del4-D3 exhibited similar Kd and were not affected by cell type specific changes in post translational modifications.

20 In Expi CHO cells, the Kd of sFGFR3\_Del4 was 0.8 nM, the Kd of sFGFR3\_Del4-C253S was 0.6 nM, and the Kd of sFGFR3\_Del4-D3 was 0.7 nM (FIG. 3A and Table 1).

**Table 1. Dissociation constant (Kd) of sFGFR3 polypeptides.**

| sFGFR3 Polypeptide | Kd (nM) |
|--------------------|---------|
| sFGFR3_Del4        | 0.8     |
| sFGFR3_Del4-C253S  | 0.6     |
| sFGFR3_Del4-D3     | 0.7     |

25 **Example 9: sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 are Equally Active *In Vitro***

sFGFR3\_Del4, sFGFR3\_Del4-C253S, and sFGFR3\_Del4-D3 restored proliferation of ATDC5 cells genetically modified to overexpress the FGFR3<sup>ach</sup> mutation (ATDC5 FGFR3<sup>G380R</sup> cell lines). At a dose of 36 nM, sFGFR3\_Del4 produced using HEK 293 cells increased proliferation to 115.5%, sFGFR3\_Del4 produced using CHO-S cells increased proliferation to 116%, sFGFR3\_Del4-C253S produced using CHO-S cells increased proliferation to 114.4%, and sFGFR3\_Del4-D3 using CHO-S cells increased proliferation to 120.1% (FIG. 3B).

30 sFGFR3\_Del4-D3 was also tested in the FGFR3<sup>G380R</sup> expressing SRE(-Luc) HEK cell line at doses of 0 nM, 70 nM, and 280nM with or without 1 ng/ml of hFGF2 (FIG. 4; n=8). Data shown in FIG. 4 are the mean +/- standard error of the mean (SEM). These data followed a 35 normal law and have equal variance based on the D'Agostino- Pearson omnibus normality test.

Statistical comparisons with and without sFGFR3\_Del4-D3 were performed using a student t-test. As shown in FIG. 4, sFGFR3\_Del4-D3 decreases luciferase signalling in the SRE cell line.

**Example 10: sFGFR3\_Del4-D3 restores Bone Growth, prevents Mortality, and restores  
5 Foramen Magnum Shape in Mice with Achondroplasia**

An in vivo efficacy study was performed as in Example 6 using a low dose (0.25 mg/kg) of sFGFR3\_Del4-D3. A total of 60 mice were included in the vehicle group, with 32 wild type (wt) mice and 28 *Fgfr3<sup>ach/+</sup>* mice. The treated group included 40 mice, with 19 wt mice and 21 *Fgfr3<sup>ach/+</sup>* mice. Surprisingly, the low dose of sFGFR3\_Del4-D3 almost completely prevented 10 the premature death of mice with achondroplasia (FIG. 5). In the control group, 53.6% of the *Fgfr3<sup>ach/+</sup>* mice died before weaning, whereas only 4.8% of mice in the treated group died before day 22 and 20% of mice died following treatment with sFGFR3\_Del1 at 0.25 mg/kg (Table 2; see also Garcia et al. *Sci. Transl. Med.* 5:203ra124, 2013, hereby incorporated by reference in its entirety).

15 sFGFR3\_Del4-D3 also partially restored bone growth with correction of the initial discrepancy between wt and *Fgfr3<sup>ach/+</sup>* mice on the axial and appendicular skeleton (Table 2). In contrast to prior results of treatment with a low dose of sFGFR3\_Del1, treatment with low dose of sFGFR3\_Del4-D3 restored normal foramen magnum shape.

20 **Table 2. In vivo results of administering a high dose of sFGFR3\_Del1, a low dose of sFGFR3\_Del1, and a low dose of sFGFR3\_Del4-D3 to mice with achondroplasia**

|   | 2.5 mg/kg<br>sFGFR3_Del1<br>(Garcia et al.) | 0.25 mg/kg<br>sFGFR3_Del1<br>(Garcia et al.) | 0.25 mg/kg<br>sFGFR3_Del4-D3 |
|---|---|--|------------------------------|
| Mortality                               | 12%   | 20%  | 4.8%                         |
| Axial correction                        | 77%   | 24%  | 10%                          |
| Appendicular<br>correction              | 150-215%                                    | 18-42%                                       | 11-42%                       |
| Foramen shape<br>correction (ratio W/H) | Not determined                              | Not determined                               | 111%                         |

**Example 11: Treatment of Achondroplasia by Administration of sFGFR3\_Del4-C253S**

25 A human patient (e.g., an infant, child, adolescent, or adult) suffering from achondroplasia can be treated by administering sFGFR3\_Del4-C253S (FIG. 6; SEQ ID NO: 2) by an appropriate route (e.g., by subcutaneous injection) at a particular dosage (e.g., between 0.0002 mg/kg/day to about 20 mg/kg/day, such as 0.001 mg/kg/day to 7 mg/kg/day) over a course of days, weeks, months, or years. The progression of achondroplasia that is treated with sFGFR3\_Del4-C253S 30 can be monitored by one or more of several established methods. A physician can monitor the patient by direct observation in order to evaluate how the symptoms of achondroplasia exhibited by the patient have changed in response to treatment. For instance, a physician may monitor changes in body weight, skull length, and/or skull width of the patient over a period of time, e.g.,

1, 2, 3, 4 or more times per month or per year or approximately every 1, 2, 3, 4, 5, 6, 7, 8, 12, or 16 weeks over the course of treatment with sFGFR3\_Del4-C253S. Body weight and/or skull size of the patient or changes thereof can also be determined at treatment specific events, e.g. before and/or after administration of sFGFR3\_Del4-C253S. For example, body weight and/or skull size are measured in response to administration of sFGFR3\_Del4-C253S.

**Example 12: Treatment of Achondroplasia by Administration of sFGFR3\_Del4-D3**

Additionally, a human patient (e.g., an infant, child, adolescent, or adult) suffering from achondroplasia can be treated by administering the sFGFR3 polypeptide of sFGFR3\_Del4-D3 (SEQ ID NO: 33) by an appropriate route (e.g., by subcutaneous injection) at a particular dosage (e.g., between 0.0002 mg/kg/day to about 20 mg/kg/day, such as 0.001 mg/kg/day to 7 mg/kg/day) over a course of days, weeks, months, or years. The progression of achondroplasia that is treated with sFGFR3\_Del4-D3 can be monitored by one or more of several established methods. A physician can monitor the patient by direct observation in order to evaluate how the symptoms of achondroplasia exhibited by the patient have changed in response to treatment. For instance, a physician may monitor changes in body weight, skull length, and/or skull width of the patient over a period of time, e.g., 1, 2, 3, 4 or more times per month or per year or approximately every 1, 2, 3, 4, 5, 6, 7, 8, 12, or 16 weeks over the course of treatment with sFGFR3\_Del4-D3. Body weight and/or skull size of the patient or changes thereof can also be determined at treatment specific events, e.g. before and/or after administration of sFGFR3\_Del4-D3. For example, body weight and/or skull size are measured in response to administration of sFGFR3\_Del4-D3.

**Example 13: Production of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S**

The sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S polypeptides were purified as described in Example 2. Modification of sFGFR3\_Del4 to include either an extended Ig-like C2-type domain 3 (FGFR3\_Del4-D3) or an amino acid substitution of a cysteine residue with a serine residue at position 253 (sFGFR3\_Del4-C253S) improved production of the sFGFR3 polypeptides. In particular, there was less than about 2% aggregation of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S (as observed upon loading using a concentration of 2.3 mg/ml or 23 mg/ml for FGFR3\_Del4-D3 and 1.5 mg/ml and 15 mg/ml of sFGFR3\_Del4-C253S) under both reducing and non-reducing conditions using sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE; FIGS. 7A and 7B, respectively). Following production of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S in fed-batch cultures, the top five clones were separated using capillary electrophoresis to yield 0.93 to 1.0 g/L and 0.98 to 1.1 g/L of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S, respectively. Viral filtration using ion-exchange chromatography resulted in a yield of greater than 60% for both sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S.

**Example 14: Pharmacokinetics and Tissue Distribution of sFGFR3\_Del4-D3 *in vivo***

*In vivo* studies were performed to investigate the pharmacokinetic parameters of sFGFR3\_Del4-D3, the uptake of sFGFR3\_Del4-D3 across the blood brain barrier, and the tissue distribution of sFGFR3\_Del4-D3 in kidney, liver, spleen, lung, and heart. The studies 5 described herein included four arms with five groups of C57BL/6J mice per arm and a total of four mice (n=4) per group (Table 3). Mice were male and weighed 25 to 30 grams.

**Table 3. Overview of mice used in studies of sFGFR3\_Del4-D3.**

| Arm | sFGFR3_Del4-D3 (mg/kg) | Route | PK  | BBB | Tissue distribution |
|-----|------------------------|-------|-----|-----|---------------------|
| 1   | 0.25                   | SC    | yes | no  | no                  |
| 2   | 2.5                    | SC    | yes | no  | yes                 |
| 3   | 2.5                    | IV    | yes | Yes | yes                 |
| 4   | 10                     | SC    | yes | no  | no                  |

10 Group 1 was sampled at 1 minute, 15 minutes, and 30 minutes; group 2 was sampled at 4 hours; group 3 was sampled at 24 hours; group 4 was sampled at 36 hours; and group 5 was sampled at 48 hours. For Group 1, an indwelling intra-arterial catheter (PE-10) was inserted into one common carotid artery under isoflurane anesthesia and used for repeated blood sampling at the 30 minute final sampling time point. For intravenous injection, <sup>125</sup>I-sFGFR3\_Del4-D3 was injected intravenously into the jugular vein, which was exposed by skin incision under isoflurane anesthesia. Group 1 mice remained anesthetized throughout the experiments. Repeated blood samples (2 x ~50µL) were drawn from the arterial catheter at 1 minute and 15 minutes after intravenous injection. For groups 2 to 5, after injection of <sup>125</sup>I-sFGFR3\_Del4-D3, the skin was closed with a surgical clip, and the mice were allowed to wake 15 up and returned to the cage. At 5 minutes before termination time for group 3, mice were re-anesthetized and received an intravenous bolus of <sup>3</sup>H-albumin into the jugular vein. The <sup>3</sup>H tracer dose was targeted to yield a ratio of <sup>125</sup>I to <sup>3</sup>H in blood, which is suitable for double isotope labeling with a lower dose at later sampling times. At the terminal sampling time (2 hours, 3 hours, 24 hours, 36 hours, and 48 hours), a blood sample was collected, and the 20 animal was euthanized. The brain was sampled for homogenization and determination of tissue concentration of tracers. Endpoints of the studies included pharmacokinetic parameters for sFGFR3\_Del4-D3 (terminal half life), uptake of sFGFR3\_Del4-D3 across the blood brain barrier, and the tissue distribution of sFGFR3\_Del4-D3 in kidney, liver, spleen, lung, and heart.

25

**30 Example 15: Thermal and Plasma Stability of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S**

The thermal stability of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S in mouse plasma was investigated using differential scanning colorimetry. For sFGFR3\_Del4-D3, two buffers (20 mM phosphate, 40mM NaCl, pH 7.5, and 20 mM citrate, 40mM NaCl, pH 6.5) were added to polypeptide samples. For sFGFR3\_Del4-C253S, two buffers (20 mM phosphate, 40mM NaCl,

pH 7.5, and 40 mM citrate, 40mM NaCl, pH 6.5) were added to polypeptide samples. The melting temperature ( $T_m$ ) for sFGFR3\_Del4-C253S in the 20 mM phosphate, 40mM NaCl, pH 7.5 buffer was 52°C and 56°C, and the  $T_m$  for sFGFR3\_Del4-C253S in the 40 mM citrate, 40mM NaCl, pH 6.5 buffer was 55°C and 60°C (FIG. 8A). For sFGFR3\_Del4-D3, two buffers (20 mM phosphate, 40mM NaCl, pH 7.5, and 20 mM citrate, 40mM NaCl, pH 6.5) were added to polypeptide samples. The  $T_m$  for sFGFR3\_Del4-D3 in the 20 mM phosphate, 40mM NaCl, pH 7.5 buffer was 50°C and 54°C, and the  $T_m$  for sFGFR3\_Del4-D3 in the 20 mM citrate, 40mM NaCl, pH 6.5 buffer was 53°C and 58°C (FIG. 8B). These results indicate that both sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S show two domains of polypeptide stability and unfolding.

The *ex vivo* plasma stability of sFGFR3\_Del4-D3 with a Histidine tag was determined by labeling purified sFGFR3\_Del4-D3 with  $^{125}\text{I}$ -tracer using the Bolton-Hunter method, followed by purification on PD-10 (Sephadex® G-25) columns. The trichloroacetic acid (TCA) precipitability of peak fractions was also determined to confirm stability of the  $^{125}\text{I}$ -tracer. Mouse plasma (n = 4) pre-warmed to 37°C was spiked with the  $^{125}\text{I}$ -sFGFR3\_Del4-D3 to a concentration of ~10 cpm/mL and then vortexed. The plasma samples were incubated with the  $^{125}\text{I}$ -sFGFR3\_Del4-D3 in an Eppendorf ThermoMixer® under gentle rotation (300 rpm). Aliquots were then collected for TCA precipitation (10  $\mu\text{l}$  sample and 100  $\mu\text{l}$  2% BSA) and for injection onto an Fast Performance Liquid Chromatography (FPLC) column (20  $\mu\text{l}$  sample and 150  $\mu\text{l}$  10 mM PBS, pH 7.4) at intervals of 0, 30, 60, 120, 180, and 360 minutes. Aliquots were stored on ice until TCA precipitation or FPLC injection was performed.

For TCA precipitation, 1 mL ice cold 10% TCA was added to plasma samples, incubated for 10 minutes on ice, centrifuged at 4,000g for 5 minutes, and then the supernatant and pellet were separated and both were counted in a gamma counter. For evaluation of the *ex vivo* plasma stability, 100  $\mu\text{l}$  of the sample was injected on an FPLC column (Superdex® 200 10/300 GL) and eluted at a rate of 0.75 ml/min for 1.5 column volumes. Fractions of 1 ml were collected from the column and then measured in a gamma counter. The plasma stability of sFGFR3\_Del4-D3 at 37°C was determined to be 95% at 0 minutes, 95% at 2 hours, and ~92% at 24 hours with only minor aggregation (FIG. 9A).

The *in vivo* stability of sFGFR3\_Del4-D3 in plasma after administration by intravenous and subcutaneous injection was also determined. sFGFR3\_Del4-D3 was labeled with  $^{125}\text{I}$ -tracer using the Bolton-Hunter method, followed by purification on PD-10 (Sephadex® G-25) columns. The  $^{125}\text{I}$ -labeled sFGFR3\_Del4-D3 (10  $\mu\text{Ci}$  in ~50  $\mu\text{L}$  PBS) was administered by intravenous or subcutaneous injection into anesthetized C57Bl/6 mice. The  $^{125}\text{I}$ -tracer protein dose (approximately 0.1 mg/kg) was complemented with unlabeled protein to a total dose of 2.5 mg/kg. Rat serum albumin used as a vascular marker was labeled with [ $^3\text{H}$ ]-NSP (N-succinimidyl[2,3- $^3\text{H}$ ]Propionate; Perkin Elmer) and purified on PD-10 (Sephadex® G25) columns.

For the stability of sFGFR3\_Del4-D3 in plasma after intravenous bolus injection, FPLC elution profiles showed no degradation products in plasma up to 15 minutes (FIG. 9B). At 30 minutes after administration of sFGFR3\_Del4-D3, a small amount of low molecular weight degradation products appeared, which increased by 2 hours, but largely disappeared by 24 hours. For the stability of sFGFR3\_Del4-D3 in plasma after subcutaneous injection, FPLC elution profiles showed some degradation products in plasma at 30 minutes, with increased degradation by 2 hours and 4 hours (FIG. 9C). The low amount of tracer left in plasma after 24 hours appears largely as the intact sFGFR3\_Del4-D3 polypeptide. The lower panel chromatograms for FIGS. 9B and 9C are presented as normalized to the highest peak in each individual run for easier comparison of the elution patterns.

#### **Example 16: Ligand Binding Activity of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S**

Experiments were performed to characterize the binding affinity of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S for human FGF2. The dissociation constant (Kd) of sFGFR3\_Del4-D3 and Kd of sFGFR3\_Del4-C253S for FGF2 were determined as described in Example 3 with a regeneration buffer of 20mM phosphate, 40mM NaCl, pH 7.5. Concentrations of 13 nM, 6.5 nM, 3.25 nM, and 1.75 nM were tested for both sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S. The Kd of sFGFR3\_Del4-D3 was determined to be ~3.6 nm, and the Kd of sFGFR3\_Del4-C253S was determined to be ~6.9 nm. These results indicate that sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S have binding activity for FGF2 in the low nM range.

#### **Example 17: sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S Exhibit Functional Activity *in vitro***

Functional activity of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S was tested using a proliferation assay. Proliferation assays using ATDC5 cells genetically modified to overexpress the FGFR3<sup>ach</sup> mutation (ATDC5 FGFR3<sup>G380R</sup> cell lines) were performed as described in Example 4 with concentrations of 1 ug/ml, 10 ug/ml, and 50 ug/ml for sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S. At each of these concentrations, sFGFR3\_Del4-C253S and sFGFR3\_Del4-D3 restored proliferation of the FGFR3<sup>G380R</sup> cells (FIG. 10A and 10B). The EC50 was determined to be about 10 nM for both sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S based on a concentration of 1 ug/ml. These results indicate that sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S are biologically active in the low nM range.

#### **Example 18: Pharmacokinetic Profile of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S**

The pharmacokinetic (PK) profile of sFGFR3\_Del4-D3 administered subcutaneously or intravenously at a dose of 2.5 mg/kg was used to determine the terminal elimination half-life of sFGFR3\_Del4-D3 (FIG. 11). Samples were collected at 30 minutes, 2 hours, 4 hours, 8 hours, 24 hours, 36 hours, and 48 hours for mice administered sFGFR3\_Del4-D3 subcutaneously.

Samples were collected at 1 minute, 15 minutes, 30 minutes, 2 hours, 24 hours, and 36 hours for mice administered sFGFR3\_Del4-D3 intravenously. The subcutaneous terminal elimination half-life of 2.5 mg/kg sFGFR3\_Del4-D3 was ~20 hours, while the intravenous terminal elimination half-life of 2.5 mg/kg sFGFR3\_Del4-D3 was ~7 hours. From the PK profile, the  $T_{max}$  5 was ~8 hours, the  $C_{max}$  was ~ 4.5 nM, and the estimated bioavailability was ~30% for 2.5 mg/kg sFGFR3\_Del4-D3 administered subcutaneously. There was rapid clearance of sFGFR3\_Del4-D3 administered intravenously during the  $\alpha$  phase followed by a slower  $\beta$  phase clearance, with a similar intravenous PK profile for sFGFR3\_Del4-C253S.

10 **Example 19: The Kidney and Liver are the Main Clearance Routes of sFGFR3\_Del4-D3**

Clearance of sFGFR3\_Del4-D3 was evaluated in kidney, liver, spleen, lung, and heart tissue after 30 minutes, 120 minutes, and 1440 minutes following intravenous administration of 2.5 mg/kg sFGFR3\_Del4-D3 and after 30 minutes, 120 minutes, 240 minutes, 480 minutes, and 1440 minutes following subcutaneous administration of 2.5 mg/kg sFGFR3\_Del4-D3. The liver 15 and kidney were the major route of sFGFR3\_Del4-D3 clearance for intravenous administration (FIG. 12). The kidney was the major route of sFGFR3\_Del4-D3 clearance for subcutaneous administration (FIG. 13).

**Example 20: sFGFR3\_Del4-D3 does not Cross the Blood Brain Barrier**

Pharmacokinetic studies were also performed to determine the uptake of sFGFR3\_Del4-20 D3 across the blood brain barrier in wild-type mice. After intravenous bolus injection, brain tissue uptake of sFGFR3\_Del4-D3 was measured at three time points (30 minutes, 2 hours, and 24 hours). sFGFR3\_Del4-D3 was injected as radiolabeled tracer ( $^{125}I$ - sFGFR3\_Del4-D3) with 2.5 mg/kg unlabeled sFGFR3\_Del4-D3. The injected dose of  $^{125}I$ - sFGFR3\_Del4-D3 was about 25 10  $\mu$ Ci per animal, which corresponds to less than 0.1 mg/kg. After euthanizing the mice at 30 minutes, 2 hours, and 24 hours, the concentration of  $^{125}I$ - sFGFR3\_Del4-D3 in organs and plasma was measured by liquid scintillation counting.

The  $^{125}I$ - sFGFR3\_Del4-D3 concentration was corrected for metabolism in plasma and in 30 brain samples by measuring the fraction of trichloroacetic acid (TCA) precipitable material (e.g., intact tracer). The validity of the TCA correction was also confirmed by injecting samples on a size exclusion fast protein liquid chromatography (FPLC) column. The organ concentration of  $^{125}I$ - sFGFR3\_Del4-D3 was corrected for intravascular content ( $V_0$ ) by injecting radiolabeled 35 albumin ( $^3H$ -RSA) shortly before sacrificing the animal. The apparent organ volume of distribution of RSA represents  $V_0$ . The dose of albumin was negligible (on the order of 1% of the physiological concentration). For all organs other than the brain, the concentrations were calculated by subtracting the vascular content and taking into account the TCA precipitable fraction in plasma. However, no correction was made for the uptake of degraded material into these organs other than the brain because no TCA precipitation was performed.

The brain concentrations were calculated by the following formula:  $C_{\text{brain}(\text{corr.})} = [V_d(\text{sFGFR3_Del4-D3}) - V_0] \times C_{\text{plasma}(\text{terminal})}$ , in which  $V_d(\text{sFGFR3_Del4-D3})$  is the volume of distribution of sFGFR3\_Del4-D3 in brain (calculated as  $C_{\text{brain}} / C_{\text{plasma}}$ ),  $V_0$  is the volume of albumin distributed in the brain, and  $C_{\text{plasma}(\text{terminal})}$  is the plasma concentration of sFGFR3\_Del4-D3 at the terminal sampling time. All concentrations were expressed as the percent of injected dose per gram or ml (%ID/g or %ID/mL), respectively, and the dose of the intravenous bolus equals 100%. These values can be converted to [mg/g] or [mg/mL] by multiplication with the injected dose: (body weight in g /1000 g) x 2.5 mg. All body weights were in the range of 25 g – 30 g.

There was no detectable brain uptake of  $^{125}\text{I}$ - sFGFR3\_Del4-D3, as indicated by corrected brain concentrations (after correction for vascular content and degradation (TCA precipitability)) at any of the measured time points (FIG. 14A). Additionally, the  $V_d$  of RSA ( $=V_0$ ) and  $^{125}\text{I}$ - sFGFR3\_Del4-D3 was not significantly different at any of the measured time points (30 minutes, 2 hours, and 24 hours) as determined by a paired t-test (FIG. 14B). In conclusion, there is no measurable uptake of sFGFR3\_Del4-D3 into brain tissue of mice at 30 minutes, 2 hours, and 24 hours at a dose of 2.5 mg/kg injected as an intravenous bolus.

#### **Example 21: *In Vivo* Efficacy of sFGFR3\_Del4-D3 for the Treatment of Achondroplasia**

sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S were each evaluated at a subcutaneous dose of 2.5 mg/kg once or twice weekly or 10 mg/kg twice weekly. Breeding was performed to generate 30 litters with half wild type and half heterozygous  $Fgfr3^{ach/+}$  mice (Table 4).

**Table 4. Subcutaneous administration of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S to wild type (WT) and  $Fgfr3^{ach/+}$  mice.**

|                          |                 | PBS (pooled) | 2.5mg 1X week | 2.5mg 2X week | 10mg 2X week        |
|--------------------------|-----------------|--------------|---------------|---------------|---------------------|
| <b>sFGFR3_Del4-D3</b>    |                 |              |               |               |                     |
|                          | WT              | 65           | 26            | 22            | 23                  |
|                          | $Fgfr3^{ach/+}$ | 43           | 26            | 25            | 30                  |
|                          |                 |              |               |               | <b>total N= 260</b> |
| <b>sFGFR3_Del4-C253S</b> |                 |              |               |               |                     |
|                          | WT              | 65           | 26            | 22            | 23                  |
|                          | $Fgfr3^{ach/+}$ | 27           | 22            | 18            | 28                  |
|                          |                 |              |               |               | <b>total N= 231</b> |
|                          | % survival      | 62.8         | 84.6          | 72.0          | 93.3                |

|  |             |      |      |      |     |
|--|-------------|------|------|------|-----|
|  | % mortality | 37.2 | 15.4 | 28.0 | 6.7 |
|--|-------------|------|------|------|-----|

At day 3, all newborn mice from a single litter received the same dose. Control litters received 10  $\mu$ l of PBS (vehicle). Thereafter, subcutaneous injections of sFGFR3\_Del4-D3 and sFGFR3\_Del4-C253S were administered at doses of 2.5 mg/kg once or twice weekly or 10 mg/kg twice a week for three weeks, alternatively on the left and right sides of the back. Mice were observed daily with particular attention to locomotion and urination alterations and weighed on days of injection. Mice with complications were observed twice a day for surveillance. Previous data indicated there was no statistical difference between males and females, and thus, males and females were considered one group for all analyses.

At day 22, all animals were sacrificed by lethal injection of pentobarbital, and gender was determined. All subsequent measurements and analyses were performed without knowledge of mice genotype to avoid investigator bias. Genotyping was performed at the end of the study to reveal the correspondence of data with a specific genotype. Since achondroplasia is a disease with phenotypic variability, all animals were included in the study. Animals dead before day 22 were used to investigate the impact of treatment on premature death. Surviving animals at day 22 were used for all analyses. All experiments and data measurements were performed by blinded experimenters at all time points.

Subcutaneous administration of sFGFR3\_Del4-D3 at 2.5 mg/kg once or twice weekly or 10 mg/kg twice weekly increased survival of *Fgfr3<sup>ach/+</sup>* mice relative to *Fgfr3<sup>ach/+</sup>* mice receiving PBS (FIG. 15 and Table 4). In particular, administration of 10 mg/kg sFGFR3\_Del4-D3 twice weekly resulted in 93% survival of *Fgfr3<sup>ach/+</sup>* mice, administration of 2.5 mg/kg sFGFR3\_Del4-D3 once weekly resulted in 84% survival in *Fgfr3<sup>ach/+</sup>* mice, and administration of 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly resulted in 72% survival in *Fgfr3<sup>ach/+</sup>* mice, while the survival of *Fgfr3<sup>ach/+</sup>* mice receiving PBS was 62.8%. The mortality of *Fgfr3<sup>ach/+</sup>* mice administered 10 mg/kg sFGFR3\_Del4-D3 twice weekly was 6.7%, the mortality of *Fgfr3<sup>ach/+</sup>* mice administered 2.5 mg/kg sFGFR3\_Del4-D3 once weekly was 15.4%, the mortality of *Fgfr3<sup>ach/+</sup>* mice administered 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly was 28.0%, and the mortality of *Fgfr3<sup>ach/+</sup>* mice administered PBS was 37.2%. Statistical analysis of *Fgfr3<sup>ach/+</sup>* mice survival following treatment with sFGFR3\_Del4-D3 was performed using the Agostino and Pearson omnibus normality test following a t-test. All investigated groups passed the normality tests. The P-values from these analyses are shown below, in which \* represent a P-value of <0.05 and \*\*\* represents a P-value of <0.001 (Table 5).

**Table 5. P-values for subcutaneous administration of sFGFR3\_Del4-D3 to wild type (WT) and *Fgfr3<sup>ach/+</sup>* mice.**

| Group Comparison | P Value |
|------------------|---------|
|------------------|---------|

|  |     |
|--|-----|
| Wt vs ach  | *** |
| <i>Fgfr3<sup>ach/+</sup></i> PBS vs <i>Fgfr3<sup>ach/+</sup></i> 2.5 mg/kg, 1x | *** |
| <i>Fgfr3<sup>ach/+</sup></i> PBS vs <i>Fgfr3<sup>ach/+</sup></i> 2.5 mg/kg, 2x | *   |
| <i>Fgfr3<sup>ach/+</sup></i> PBS vs <i>Fgfr3<sup>ach/+</sup></i> 10 mg/kg, 2x  | *** |
| Wt PBS vs <i>Fgfr3<sup>ach/+</sup></i> 10 mg/kg, 2x                            | ns  |

Subcutaneous administration of sFGFR3\_Del4-D3 at 2.5 mg/kg once or twice weekly or 10 mg/kg twice weekly also decreased the severity and frequency of locomotor problems and complications in abdominal breathing in *Fgfr3<sup>ach/+</sup>* mice relative to *Fgfr3<sup>ach/+</sup>* mice receiving PBS

5 (FIG. 16). In particular, locomotor problems decreased the most in *Fgfr3<sup>ach/+</sup>* mice administered subcutaneously 10 mg/kg sFGFR3\_Del4-D3 twice weekly followed by mice administered sFGFR3\_Del4-D3 2.5 mg/kg twice weekly and mice administered sFGFR3\_Del4-D3 2.5 mg/kg once weekly. Complications in abdominal breathing decreased the most in *Fgfr3<sup>ach/+</sup>* mice administered subcutaneously 10 mg/kg sFGFR3\_Del4-D3 twice weekly followed by mice 10 administered sFGFR3\_Del4-D3 2.5 mg/kg once weekly and then mice administered sFGFR3\_Del4-D3 2.5 mg/kg twice weekly. These results show that sFGFR3\_Del4-D3 reduces symptoms of achondroplasia in *Fgfr3<sup>ach/+</sup>* mice.

Subcutaneous administration of sFGFR3\_Del4-D3 also significantly increased total body length, including axial length and tail length, and long bones (p = 0.07) in *Fgfr3<sup>ach/+</sup>* mice 15 receiving 2.5 mg/kg sFGFR3\_Del4-D3 once or twice weekly or 10 mg/kg sFGFR3\_Del4-D3 twice weekly relative to *Fgfr3<sup>ach/+</sup>* mice receiving PBS (FIGS. 17A-17C). Tail and body length (axial length) were measured using the same digital caliper on whole skeletons. Tibia length was measured on digital X-rays. Administration of 10 mg/kg sFGFR3\_Del4-D3 twice weekly 20 resulted in 51% axial correction (body and tail length) of *Fgfr3<sup>ach/+</sup>* mice, followed by 43% axial correction in *Fgfr3<sup>ach/+</sup>* receiving 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, and 39% axial correction in *Fgfr3<sup>ach/+</sup>* mice receiving 2.5 mg/kg sFGFR3\_Del4-D3 once weekly. Increases in bone and body length were also evident from x-ray radiographs of *Fgfr3<sup>ach/+</sup>* mice administered 2.5 mg/kg or 10 mg/kg sFGFR3\_Del4-D3 twice weekly relative to *Fgfr3<sup>ach/+</sup>* mice receiving PBS 25 (FIG. 17D). Administration of 10 mg/kg sFGFR3\_Del4-D3 twice weekly resulted in 86% appendicular correction (tibia and femur length) of *Fgfr3<sup>ach/+</sup>* mice, followed by 68% appendicular correction in *Fgfr3<sup>ach/+</sup>* receiving 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly and 54% appendicular correction in *Fgfr3<sup>ach/+</sup>* mice receiving 2.5 mg/kg sFGFR3\_Del4-D3 once weekly.

Subcutaneous administration of sFGFR3\_Del4-D3 also resulted in a dose-dependent improvement in cranial ratio (length/width (L/W)) in *Fgfr3<sup>ach/+</sup>* mice relative to *Fgfr3<sup>ach/+</sup>* mice 30 receiving PBS (FIG. 18A). *Fgfr3<sup>ach/+</sup>* mice subcutaneously administered 10 mg/kg

sFGFR3\_Del4-D3 twice weekly exhibited the greatest improvement in the cranium ratio (L/W), followed by *Fgfr3<sup>ach/+</sup>* mice administered 2 mg/kg sFGFR3\_Del4-D3 twice weekly and *Fgfr3<sup>ach/+</sup>* mice administered 2 mg/kg sFGFR3\_Del4-D3 once weekly. In particular, administration of 10 mg/kg sFGFR3\_Del4-D3 twice weekly resulted in 37% skull shape correction (L/W ratio) of 5 *Fgfr3<sup>ach/+</sup>* mice, followed by 29% skull shape correction in *Fgfr3<sup>ach/+</sup>* receiving 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly and 19% skull shape correction in *Fgfr3<sup>ach/+</sup>* mice receiving 2.5 mg/kg sFGFR3\_Del4-D3 once weekly. Improvements in the cranial ratio were also evident from 10 x-ray radiographs of *Fgfr3<sup>ach/+</sup>* mice administered 10 mg/kg sFGFR3\_Del4-D3 relative to *Fgfr3<sup>ach/+</sup>* mice receiving PBS (FIG. 18B). Bone measurements (presented in mm and mean  $\pm$  SEM) for body length, tail, femur, tibia, and cranial ratio are shown below (Table 6). These results indicate the dose-dependent *in vivo* efficacy of sFGFR3\_Del4-D3 as demonstrated by increased survival, reduced number of complications, increased bone growth, and improvements in skeletal proportions of *Fgfr3<sup>ach/+</sup>* mice.

15 **Table 6. Bone measurements (presented in mm and mean  $\pm$  SEM) for body length, tail, femur, tibia, and cranial ratio of WT and *Fgfr3<sup>ach/+</sup>* mice administered subcutaneously sFGFR3\_Del4-D3.**

|               | <i>Efficacy of sFGFR3_Del4-D3</i> |  |                          |                           |                          |
|---------------|-----------------------------------|--|--------------------------|---------------------------|--------------------------|
|               | WT                                | PBS in<br><i>Fgfr3<sup>ach/+</sup></i><br>mice | 2.5 mg/kg<br>once weekly | 2.5 mg/kg<br>twice weekly | 10 mg/kg<br>twice weekly |
| Body length   | 144.8 $\pm$ 0.53                  | 129.2 $\pm$ 1.98                               | 135 $\pm$ 1.48           | 135.5 $\pm$ 1.75          | 135.2 $\pm$ 1.58         |
| Tail          | 77.65 $\pm$ 0.39                  | 70.25 $\pm$ 1.1                                | 73.37 $\pm$ 1.66         | 73.69 $\pm$ 1.5           | 74.95 $\pm$ 0.91         |
| Femur         | 10.94 $\pm$ 0.05                  | 10.14 $\pm$ 0.13                               | 10.47 $\pm$ 0.08         | 10.58 $\pm$ 0.09          | 10.63 $\pm$ 0.10         |
| Tibia         | 14.19 $\pm$ 0.05                  | 13.67 $\pm$ 0.14                               | 14.02 $\pm$ 0.10         | 14.09 $\pm$ 0.12          | 14.25 $\pm$ 0.12         |
| Cranial ratio | 1.99 $\pm$ 0.01                   | 1.79 $\pm$ 0.01                                | 1.83 $\pm$ 0.02          | 1.85 $\pm$ 0.01           | 1.86 $\pm$ 0.02          |

20 Additionally, comparison of the bone measurements for *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del1 at a dosage of 2.5 mg/kg twice weekly show that administration sFGFR3\_Del4-D3 at a dosage of 2.5 mg/kg twice weekly was comparable to or more effective in increasing the bone, tail, femur, and tibia length and improving the cranial ratio of *Fgfr3<sup>ach/+</sup>* mice (Table 7). In particular, the body length of *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del4-D3 improved to 135.5  $\pm$  1.75 mm relative to 134.4  $\pm$  1.17 mm for *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del1; the tail length of *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del4-D3 improved to 73.69  $\pm$  1.5 mm relative to 71.58  $\pm$  0.86 mm for *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del1; the femur length of *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del4-D3 improved to 10.58  $\pm$  0.09 mm relative to 10.01  $\pm$  0.06 mm for *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del1; the tibia length of *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del4-D3 improved to 14.09  $\pm$  0.12 mm relative to 13.27  $\pm$  0.31 mm for *Fgfr3<sup>ach/+</sup>* mice 25 30

administered sFGFR3\_Del1; and the cranial ratio of *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del4-D3 improved to  $1.85 \pm 0.01$  mm relative to  $1.81 \pm 0.02$  mm for *Fgfr3<sup>ach/+</sup>* mice administered sFGFR3\_Del1.

5 **Table 7. Bone measurements (presented in mm and mean  $\pm$  SEM) for body length, tail, femur, tibia, and cranial ratio of WT and *Fgfr3<sup>ach/+</sup>* mice administered subcutaneously sFGFR3\_Del1 (data described in Garcia et al. *Sci. Transl. Med.* 5:203ra124, 2013).**

| <b><i>Efficacy of sFGFR3_Del1</i></b> |                  |  |                         |                        |
|---------------------------------------|------------------|--|-------------------------|------------------------|
|                                       | WT               | PBS in <i>Fgfr3<sup>ach/+</sup></i> mice | 0.25 mg/kg twice weekly | 2.5 mg/kg twice weekly |
| body length                           | $133.9 \pm 0.8$  | $118.5 \pm 1.76$                         | $132.4 \pm 1.26$        | $134.4 \pm 1.17$       |
| tail                                  | $71.9 \pm 0.49$  | $64.48 \pm 1.1$                          | $71.05 \pm 0.99$        | $71.58 \pm 0.86$       |
| femur                                 | $10.05 \pm 0.17$ | $9.67 \pm 0.16$                          | $9.85 \pm 0.10$         | $10.01 \pm 0.06$       |
| tibia                                 | $13.43 \pm 0.19$ | $12.62 \pm 0.18$                         | $12.87 \pm 0.14$        | $13.27 \pm 0.31$       |
| cranial ratio                         | $1.94 \pm 0.01$  | $1.75 \pm 0.01$                          | $1.77 \pm 0.02$         | $1.81 \pm 0.02$        |

10 **Example 22: No Organ Toxicity Associated with Administration of sFGFR3\_Del4-D3**

Histopathological studies were performed to characterize organ toxicity associated with sFGFR3\_Del4-D3 administration. Wild type mice (6 males and 6 females per dose) were administered PBS, 2.5 mg/kg sFGFR3\_Del4-D3 once weekly, 2.5 mg/kg sFGFR3\_Del4-D3 twice weekly, or 10 mg/kg sFGFR3\_Del4-D3 twice weekly. Organs investigated included the kidney, 15 skin, salivary glands, mandibular lymph nodes, gall bladder, spleen, pancreas, lungs, heart, aorta, jejunum, colon, and liver. There were no histopathological results indicating organ toxicity in wild-type mice administered any of the doses of sFGFR3\_Del4-D3. These results indicate that there was no toxicity associated with administration of sFGFR3\_Del4-D3 up to 10 mg/kg twice weekly.

20 **Example 23: Determination of Binding Affinity of sFGFR3\_Del4-D3 to Fibroblast Growth Factors**

We determined that sFGFR3\_Del4-D3 binds to Fibroblast Growth Factors (FGF) ligands and acts as a decoy to prevent the binding of FGFs to the membrane bound FGFR3. Surface Plasmon Resonance was performed using a BIACORE™ T200 (GE Healthcare) to determine 25 the  $K_d$  values for different human FGFs (hFGFs) binding to immobilized sFGFR3\_Del4-D3. In particular,  $K_d$  values for the paracrine hFGFs of hFGF1 (FIG. 19A), hFGF2 (FIG. 19B), hFGF9 (FIG. 19C), and hFGF18 (FIG. 19D) and the endocrine hFGFs of hFGF19 (FIG. 19E) and hFGF21 (FIG. 19F) were determined. All four paracrine FGF ligands bound sFGFR3\_Del4-D3 with nanomolar (nM) affinity (Table 8).

| Paracrine FGFs | Binding                    | $k_{a1}$ (1/Ms)  | $k_{a2}$ (1/Ms) | $k_{d1}$ (1/s)   | $k_{d2}$ (1/s)  | $K_D$ (M) Kinetic                               | $\text{Chi}^2$ ( $\text{RU}^2$ ) average | $K_D$ (M) Steady state                       | $\text{Chi}^2$ ( $\text{RU}^2$ ) average |
|----------------|----------------------------|------------------|-----------------|------------------|-----------------|---|--|--|--|
| FGF1           | 2:1 binding & steady state | $2.0^* 10^{+11}$ | $1.2^* 10^{-3}$ | 1610             | $6.4^* 10^{-4}$ | $2.6^* 10^{-9}$ (+/- $1.9^* 10^{-9}$ , n = 3)   | 0.138                                    | $5.7^* 10^{-9}$ (+/- $2.1^* 10^{-9}$ , n=3)  | 0.247                                    |
| FGF2           | 1:1 binding                | $9.0^* 10^{+5}$  |                 | $4.75^* 10^{-4}$ |                 | $6.1^* 10^{-10}$ (+/- $1.7^* 10^{-10}$ , n = 3) | 13.6                                     |  |  |
| FGF9           | 2:1 binding & steady state | $2.3^* 10^{+6}$  | $3.0^* 10^{-2}$ | $2.6^* 10^{-2}$  | $3.6^* 10^{-3}$ | $1.8^* 10^{-9}$ (+/- $0.25^* 10^{-9}$ , n = 3)  | 0.14                                     | $3.6^* 10^{-9}$ (n = 1)                      | 0.25                                     |
| FGF18          | 1:1 binding & steady state | $2.0^* 10^{+5}$  |                 | $9.1^* 10^{-3}$  |                 | $4.5^* 10^{-9}$ (+/- $2.5^* 10^{-9}$ , n = 3)   | 9.7                                      | $6.4^* 10^{-9}$ (+/- $0.89^* 10^{-9}$ , n=4) | 11.8                                     |
| Endocrine FGFs |                            |                  |                 |                  |                 |   |  |  |  |

|        |             |                       |                       |                       |                       |   |      |  |  |
|--------|-------------|-----------------------|-----------------------|-----------------------|-----------------------|---|------|--|--|
| hFGF19 | 2:1 binding | 5.4* 10 <sup>+4</sup> | 7.3* 10 <sup>-3</sup> | 1.5* 10 <sup>-1</sup> | 3.6* 10 <sup>-3</sup> | <b>4.8* 10<sup>-7</sup></b><br>(+/- 3.2*10 <sup>-7</sup> , n = 3) | 0.05 |  |  |
| hFGF21 | 2:1 binding | 258                   | 1.8* 10 <sup>-2</sup> | 5.5* 10 <sup>-3</sup> | 1.4* 10 <sup>-3</sup> | <b>2.8* 10<sup>-5</sup></b><br>(n = 2)                            | 0.56 |  |  |

**Table 8. Summary of  $K_d$  determination and values for human, paracrine FGFs (hFGF1, hFGF2, hFGF9, and hFGF18) and human, endocrine FGFs (hFGF19 and hFGF21).**

5

For FGF2 and FGF18, a good fit was achieved with a 1:1 binding model, which is the most direct model of binding affinity. This model describes a 1:1 binding interaction at the surface of the chip with immobilized sFGFR3\_DEL4-D3 binding different FGFs: A + B = AB with single on- and off rate. The 2:1 model also describes a 1:1 interaction of FGF binding to SFGFR3\_DEL4-D3, but also assumes a conformational change that stabilizes the complex: A + B = AB = AB\* and represents two on- and off-rates. This model assumes that the conformationally changed complex (sFGFR3\_DEL4-D3 bound to FGF) can only dissociate by reversing the conformational change. The experimental data for hFGF1, hFGF9, hFGF19, and hFGF21 were determined to fit the 2:1 model very well, and thus,  $K_d$  for hFGF1, hFGF9, hFGF19, and hFGF21 were derived from the 2:1 model.

Despite hFGF1, hFGF9, hFGF19, and hFGF21 all having a  $K_d$  in the low nM range, the kinetic profiles of these hFGFs differed significantly. For example, FGF1 binds sFGFR3\_Del4-D3 with a very fast on-rate and off-rate, while FGF2 does not bind sFGFR3\_Del4-D3 with as fast of an on-rate or off-rate as FGF1, resulting in an overall smaller  $K_d$  for FGF2 compared to FGF1 (Table 8). A significantly lower affinity was measured between sFGFR3\_Del4-D3 and hFGF19 or hFGF21, which are members of the endocrine FGF15/FGF19 subfamily, relative to the paracrine hFGFs (Table 8 and FIGS. 19D and 19E). The FGF15/FGF19 subfamily uses Klotho instead of proteoglycans as a co-factor and has evolved into endocrine-acting growth factors, which are important for the systemic regulation of metabolic parameters, such as phosphate, bile acid, carbohydrate, and lipid metabolism.

These results demonstrate that there was a high affinity interaction of sFGFR3\_Del4-D3 with hFGF1, hFGF2, hFGF9, and hFGF18, while there was a low affinity interaction of sFGFR3\_Del4-D3 with FGF19 and FGF21. The low affinity of sFGFR3\_Del4-D3 for FGF19 and FGF21 is advantageous as sFGFR3\_Del4-D3 will have a low probability of interfering with the function of these FGFs *in vivo*.

#### **Example 24: *In Vitro* Proliferation Assay of sFGFR3\_Del4-D3**

Following binding of FGFs, FGFR3 dimerizes to initiate signaling cascades. Several downstream signaling pathways are associated with FGF signaling. In chondrocytes, dimerized

FGFR3 results in an anti-proliferative signal/early differentiation signal into the chondrocyte, which eventually leads to inhibition of bone growth. For example, the RAS/MAPK pathway propagates signals to negatively affect proliferation, terminal differentiation, and post-mitotic matrix synthesis, and the STAT1 pathway mediates the inhibition of chondrocyte proliferation in concert with the cell cycle regulators p107 and 130 and cell cycle inhibitor p21Waf/Cip1. Gene expression studies suggest a number of other pathways are also involved in down-regulation of growth-promoting molecules or induction of anti-proliferative functions.

To study FGFR3-decoy induced inhibition of FGFR3<sup>G380R</sup> in a chondrocytic cell model, studies were performed to determine the effect of sFGFR3\_Del4-D3 on the proliferation of ATDC5 cells genetically modified to overexpress the FGFR3<sup>ach</sup> mutation (ATDC5 FGFR3<sup>G380R</sup> cells). The chondrocytic cell line ATDC5 cell, which was first isolated from the differentiating teratocarcinoma stem cell line AT805, is commonly used as a model for *in vitro* chondrocyte research. ATDC5 cells were first infected with a retroviral expression vector and a stable cell line expressing FGFR3<sup>G380R</sup> was generated. The expression of FGFR3<sup>G380R</sup> in the ATDC5 cell line was determined via Western blot (FIG. 20). Extracts of ATDC5 cells expressing FGFR3<sup>G380R</sup> at passage one (G380R #1) and two (G380R #2) after resistant cell selection and extracts of control ATDC5 cells were blotted and detected with antibodies for total phosphorylation of FGFR3 (pFGFR3), the specific phosphotyrosine 724 in FGFR3 (pFGFR3 Y724), and total FGFR3 expression (FGFR3). Total extracellular signal-related kinase expression was used as loading control (ERK). Addition of SFGFR3\_DEL4-D3 to the ATDC5 FGFR3<sup>G380R</sup> cells dose-dependently increased the proliferation index of the ATDC5 FGFR3<sup>G380R</sup> cells by two-fold with an EC<sub>50</sub> of 1.25 +/- 0.27 nM (FIG. 21). These results demonstrate that addition of SFGFR3\_DEL4-D3 to ATDC5 FGFR3<sup>G380R</sup> cells overcomes the negative growth signal mediated by FGFR3<sup>G380R</sup> in a cellular model of achondroplasia and are in line with the anti-proliferative signal mediated by FGFR3 in chondrocytes, which is more pronounced when the chondrocytes express a FGFR3 including the G380R mutation.

## OTHER EMBODIMENTS

All publications, patents, and patent applications mentioned in the above specification are hereby incorporated by reference to the same extent as if each individual publication, patent or patent application was specifically and individually indicated to be incorporated by reference in its entirety. Various modifications and variations of the described methods, pharmaceutical compositions, and kits 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 will be understood that it is capable of further modifications and 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. This application is intended to cover any variations, uses, or adaptations of the invention following, in general, the principles of the invention and including such departures from the present disclosure come within known customary practice within the art to which the invention pertains and may be applied to the essential features herein before set forth.

5

## CLAIMS

1. A soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide comprising an amino acid sequence of SEQ ID NO: 33.
2. A soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide encodable by nucleotides 52-1098 of SEQ ID NO: 21.
3. A soluble fibroblast growth factor receptor 3 (sFGFR3) polypeptide encoded by nucleotides 52-1098 of SEQ ID NO: 21.
4. The sFGFR3 polypeptide of any one of claims 1-3, wherein the sFGFR3 polypeptide binds to at least one of fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), or fibroblast growth factor 18 (FGF18).
5. The sFGFR3 polypeptide of any one of claims 1-4, wherein the sFGFR3 polypeptide binds to fibroblast growth factor 1 (FGF1), fibroblast growth factor 2 (FGF2), fibroblast growth factor 9 (FGF9), and fibroblast growth factor 18 (FGF18).
6. The sFGFR3 polypeptide of claim 4 or claim 5, wherein the binding is characterized by an equilibrium dissociation constant (Kd) of from about 0.2 nM to about 20 nM.
7. The sFGFR3 polypeptide of any one of claims 4-6, wherein the binding is characterized by a Kd of from about 1 nM to about 10 nM.
8. A nucleic acid encoding the sFGFR3 polypeptide of any one of claims 1-7.
9. A vector comprising the nucleic acid of claim 8.
10. The vector of claim 9, wherein the vector is a plasmid, an artificial chromosome, a viral vector, or a phage vector.
11. A host cell comprising the nucleic acid of claim 8 or the vector of claim 9 or claim 10.
12. The host cell of claim 11, wherein the host cell is a HEK293 cell.
13. The host cell of claim 11, wherein the host cell is a CHO cell.

14. A method of producing the sFGFR3 polypeptide of any one of claims 1-7, the method comprising:

- (i) culturing the host cell of any one of claims 11-13 in a culture medium under conditions suitable for expression of the sFGFR3 polypeptide; and
- (ii) recovering the sFGFR3 polypeptide from the culture medium.

15. The method of claim 14, further comprising a step of (iii) purifying the sFGFR3 polypeptide by ion exchange chromatography and size exclusion chromatography.

16. The method of claim 14 or claim 15, wherein the sFGFR3 polypeptide is less than 2% aggregated.

17. The method of any one of claims 14 to 16, wherein the recovering step yields a concentration of sFGFR3 polypeptide of from about 0.9 g/L to about 1.1 g/L.

18. A sFGFR3 polypeptide when produced by the method of any one of claims 14 to 17.

19. A composition comprising:

the sFGFR3 polypeptide of any one of claims 1-7 or 18; and  
a pharmaceutically acceptable excipient.

20. The composition of claim 19, wherein the composition comprises less than 2% aggregation of the sFGFR3 polypeptide.

21. A method comprising administering to a subject in need thereof the sFGFR3 polypeptide of any one of claims 1-7 or 18 or the composition of claim 19 or claim 20.

22. A method for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof, the method comprising administering to the subject the sFGFR3 polypeptide of 30 any one of claims 1-7 or 18 or the composition of claim 19 or claim 20.

23. Use of the sFGFR3 polypeptide of any one of claims 1-7 or 18 or the composition of claim 19 or claim 20 for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof.

35

24. Use of the sFGFR3 polypeptide of any one of claims 1-7 or 18 in the manufacture of a medicament for treating an FGFR3-related skeletal growth retardation disorder in a subject in need thereof.

25. The method of claim 22 or the use of claim 23 or claim 24, wherein the FGFR3-related skeletal growth retardation disorder is achondroplasia, thanatophoric dysplasia type I (TDI), thanatophoric dysplasia type II (TDII), severe achondroplasia with developmental delay and acanthosis nigricans (SADDAN), hypochondroplasia, a craniostenosis syndrome, or camptodactyly, tall stature, and hearing loss syndrome (CATSHL).
26. The method or use of any one of claims 22-25, wherein the FGFR3-related skeletal growth retardation disorder is achondroplasia.
27. The method or use of any one of claims 22-25, wherein the FGFR3-related skeletal growth retardation disorder is hypochondroplasia.
28. A method for treating achondroplasia in a human subject in need thereof, the method comprising administering to the human subject the sFGFR3 polypeptide of any one of claims 1-7 or 18 or the composition of claim 19 or claim 20.
29. Use of the sFGFR3 polypeptide of any one of claims 1-7 or 18 or the composition of claim 19 or claim 20 for treating achondroplasia in a human subject in need thereof.
30. Use of the sFGFR3 polypeptide of any one of claims 1-7 or 18 in the manufacture of a medicament for treating achondroplasia in a human subject in need thereof.
31. The method or use of any one of claims 22-30, wherein the subject is administered a dose of the sFGFR3 polypeptide of about 0.0002 mg/kg to about 30 mg/kg.
32. The method or use of any one of claim 22-31, wherein the subject is administered a dose of the sFGFR3 polypeptide of about 0.2 mg/kg to 3 mg/kg.
33. The method or use of any one of claims 22-32, wherein the sFGFR3 polypeptide or the composition improves one or more symptoms in the subject.
34. The method or use of any one of claims 22-33, wherein administration of the sFGFR3 polypeptide or the composition:
  35. a) increases survival of the subject;
  - b) improves locomotion of the subject;
  - c) improves abdominal breathing in the subject;
  - d) increases body and/or bone length of the subject; or

- e) improves the cranial ratio and/or restores foramen magnum shape in the subject.

35. The method or use of any one of claims 22-34, wherein the sFGFR3 polypeptide or composition is administered subcutaneously.

36. The method or use of any one of claims 22-34, wherein the sFGFR3 polypeptide or composition is administered intravenously.

FIG. 1A

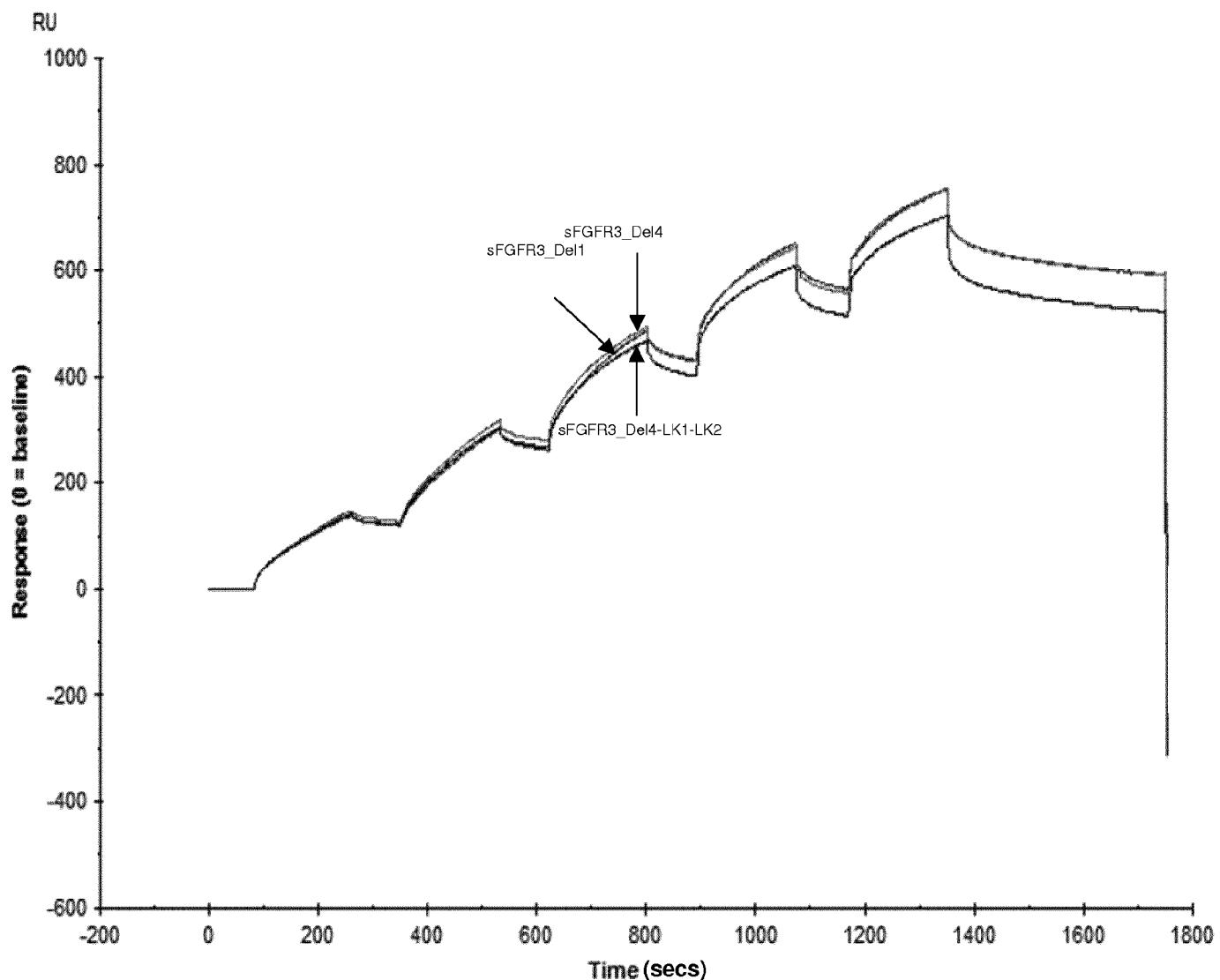


FIG. 1B

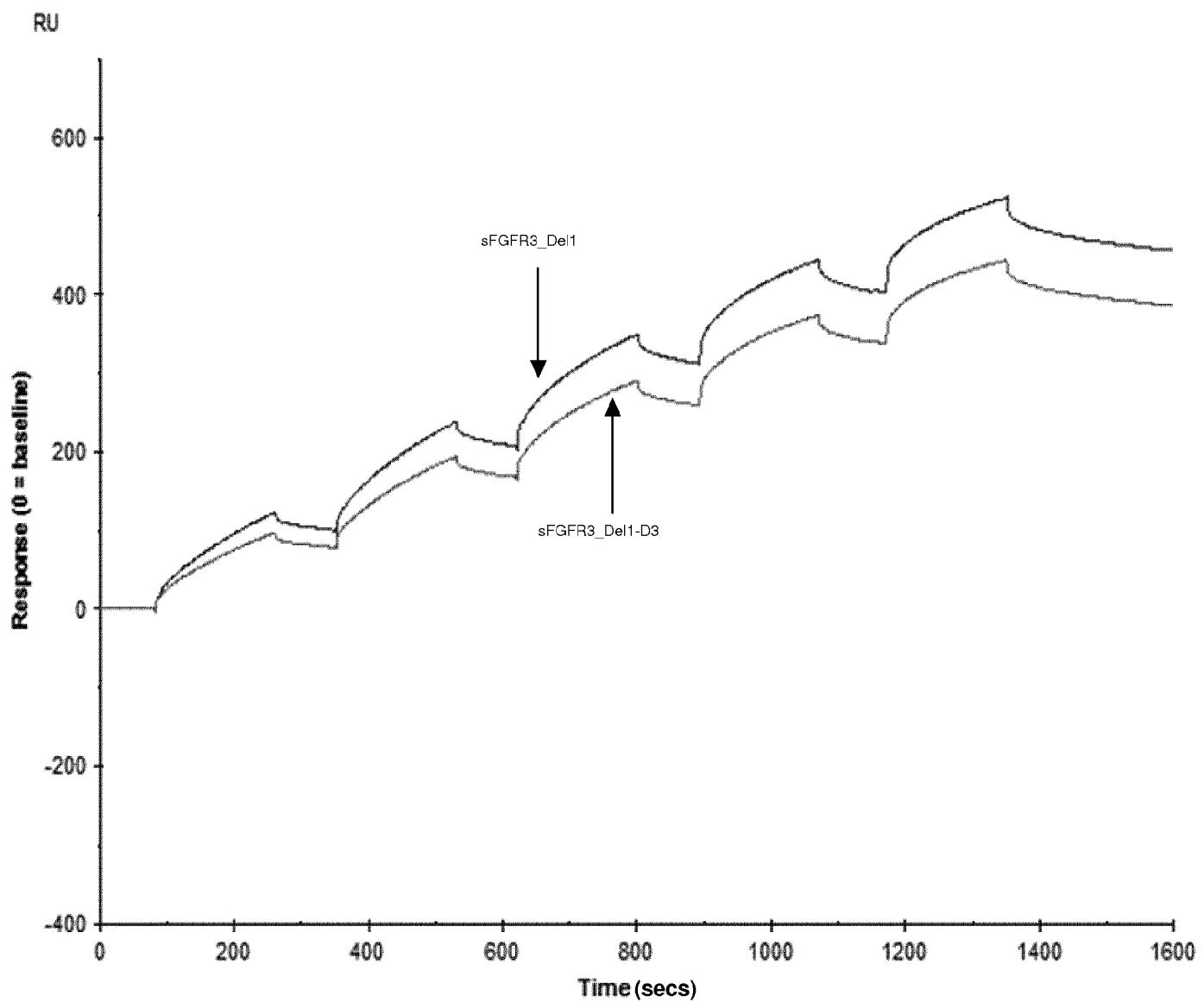


FIG. 1C

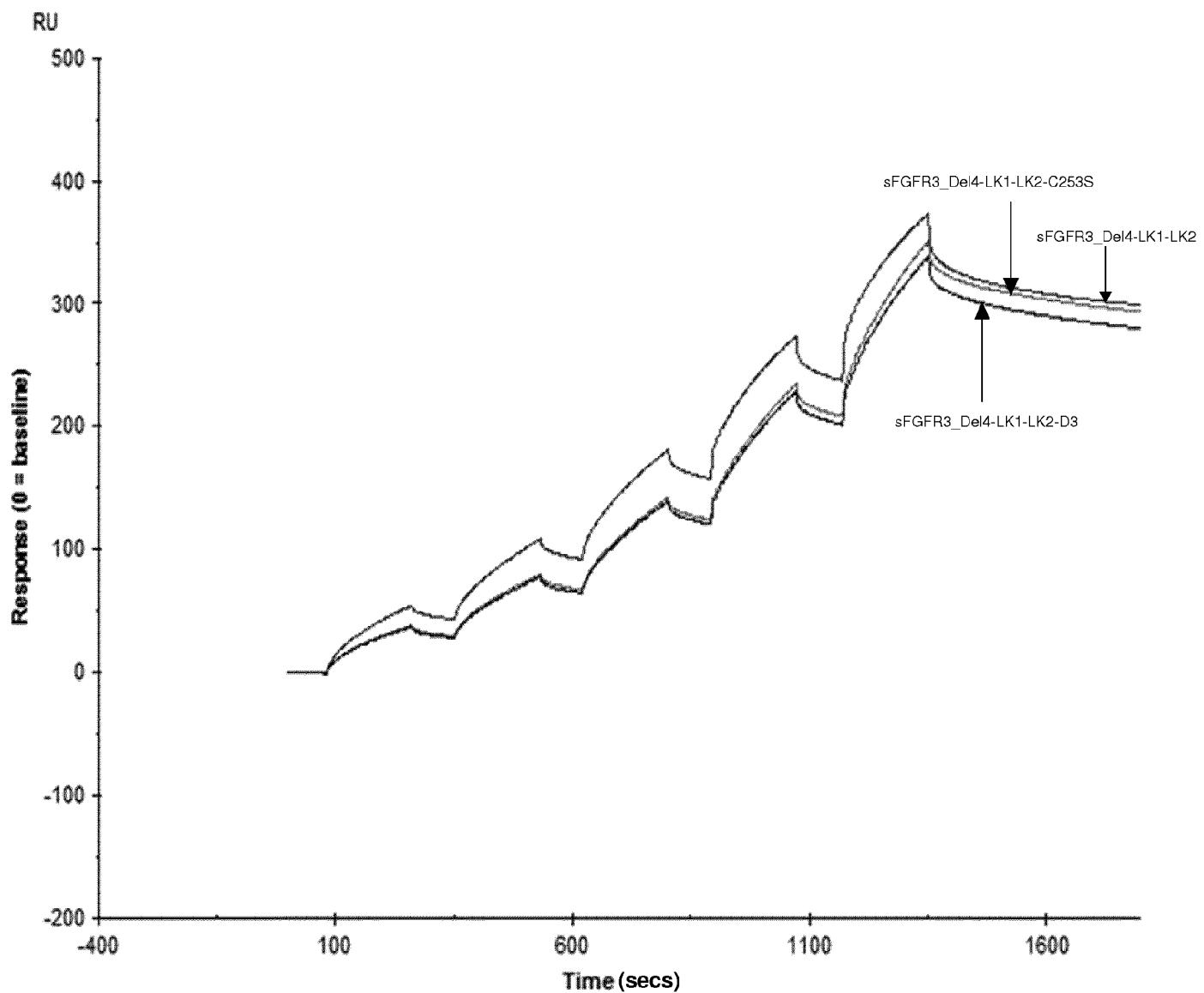


FIG. 1D

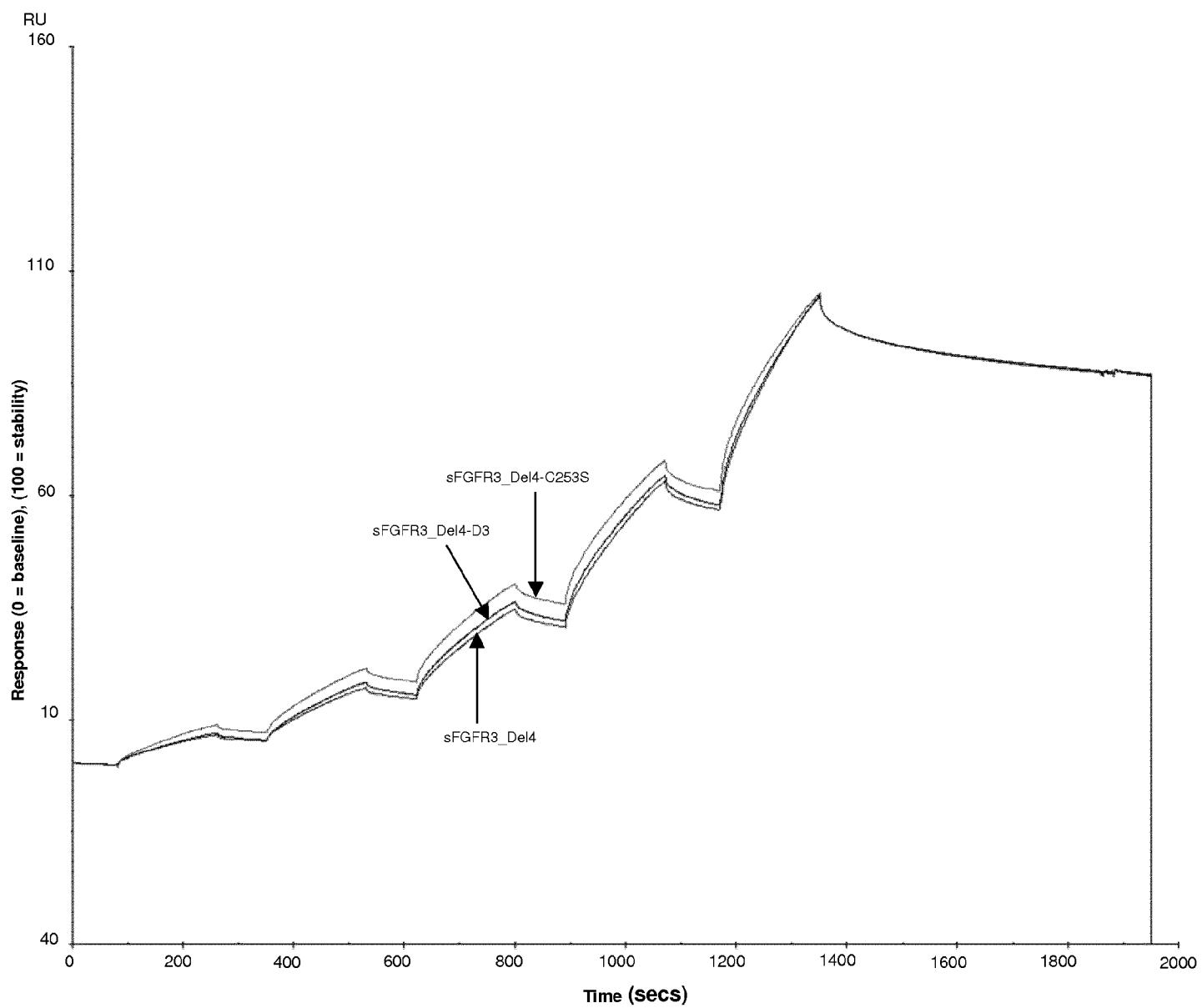
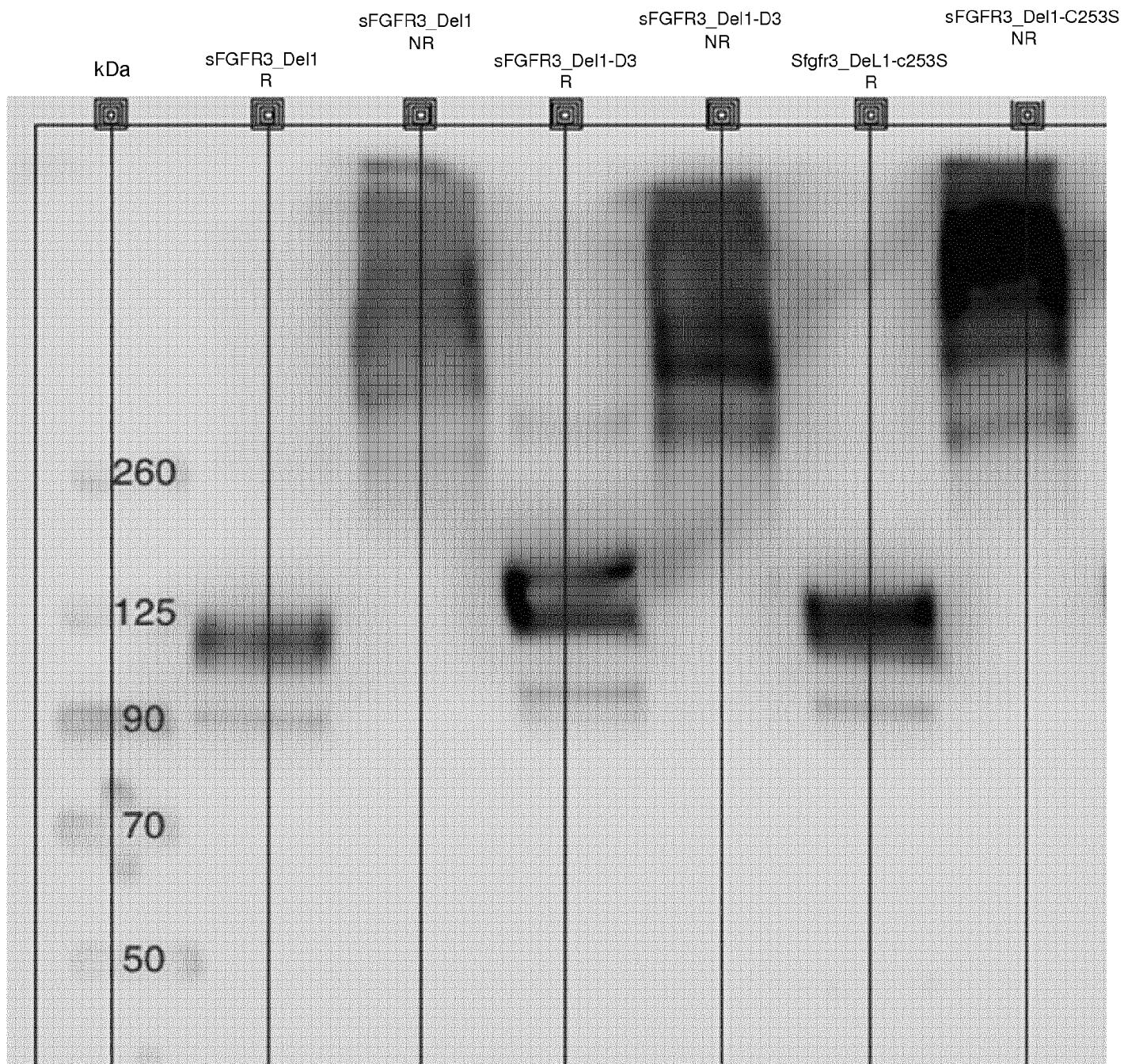
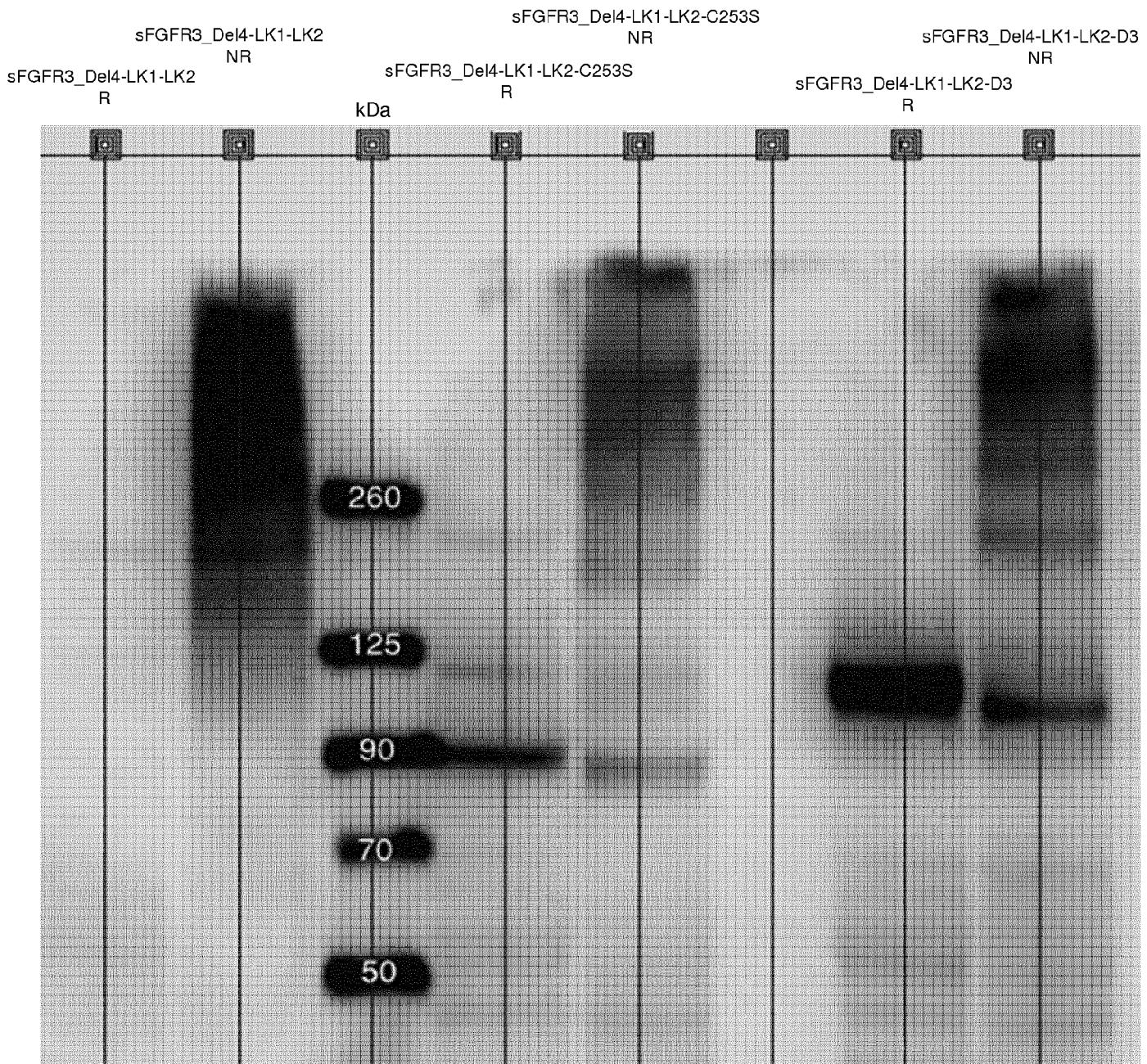


FIG. 2A



## FIG. 2B



## FIG. 2C

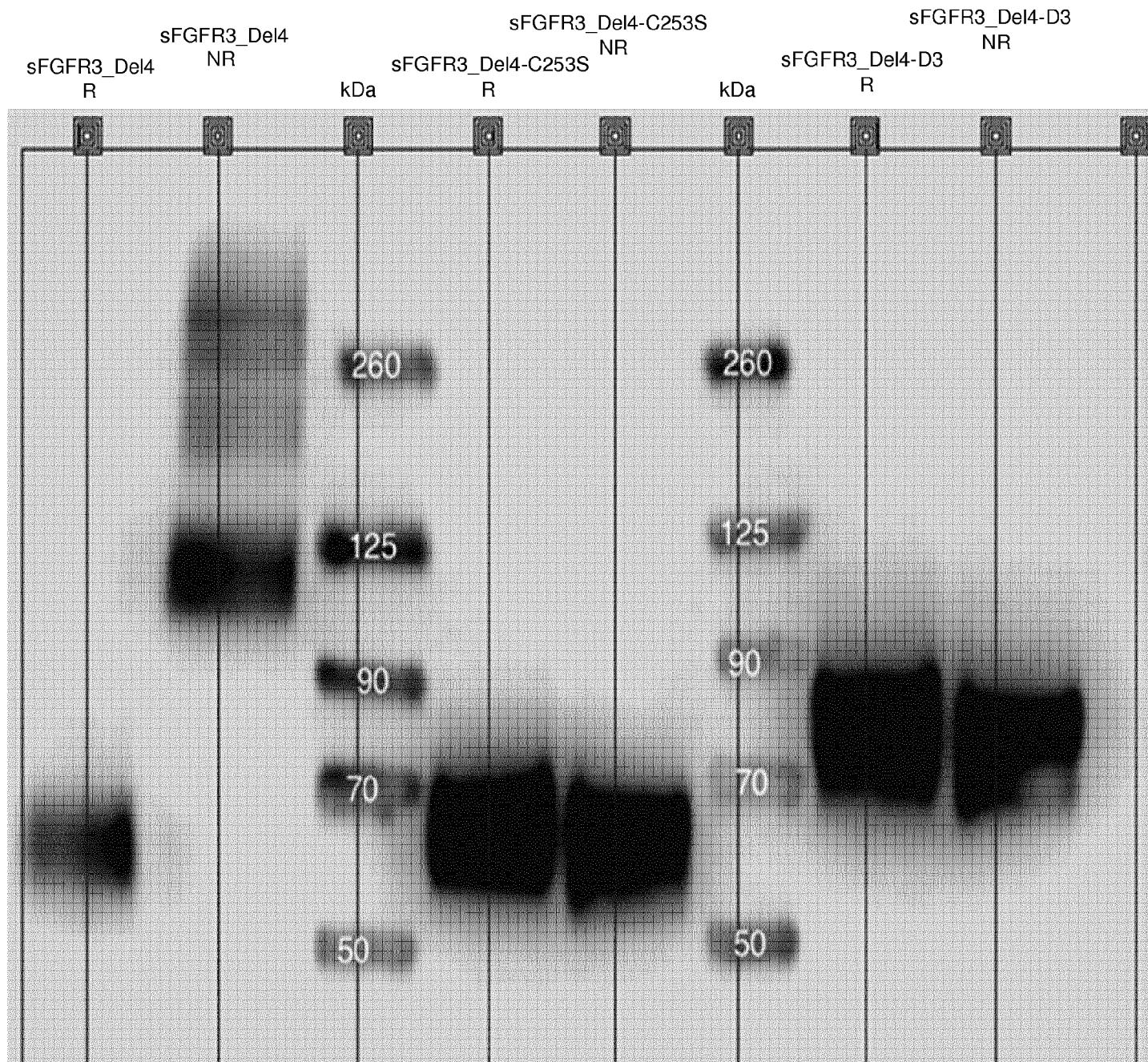


FIG. 3A

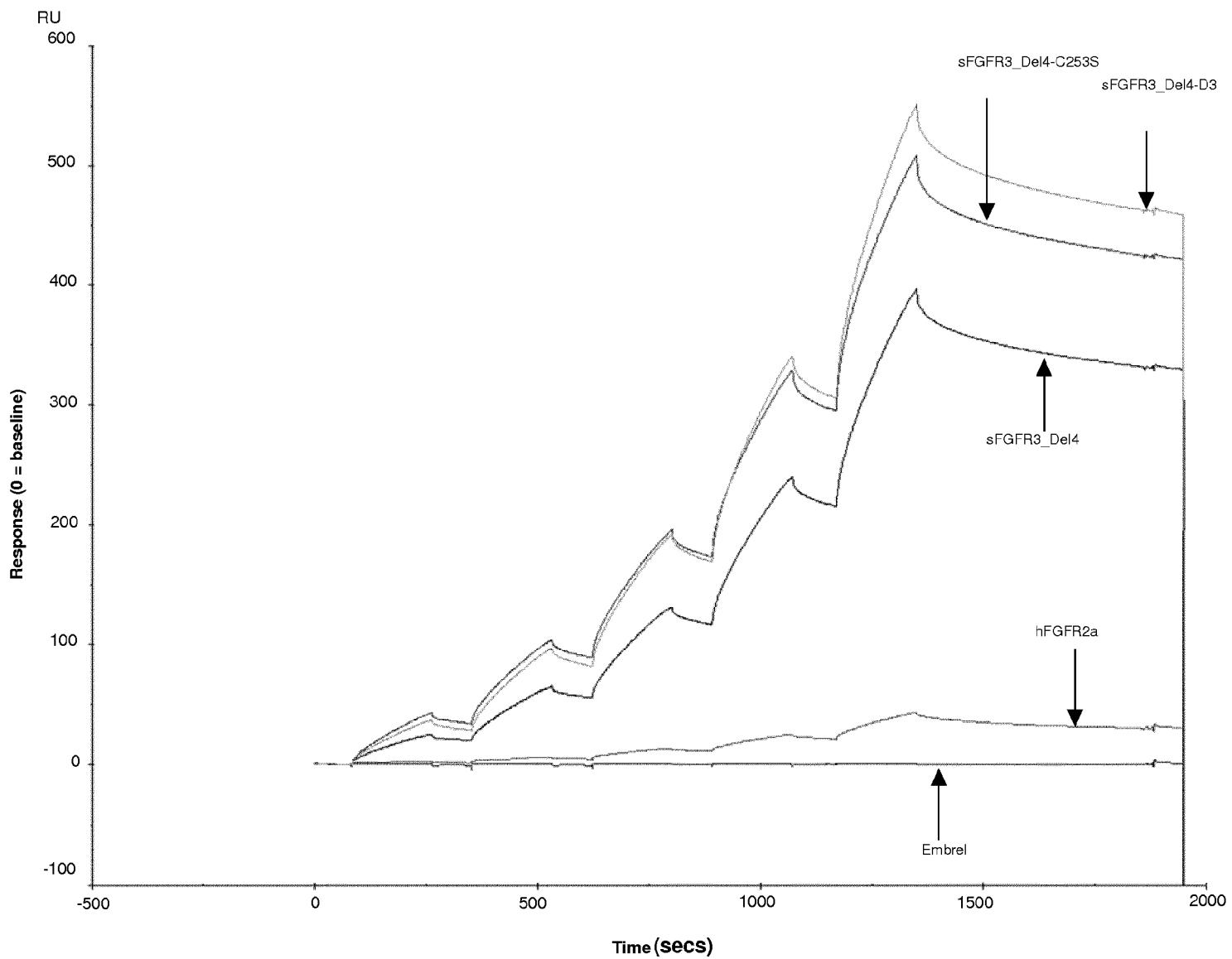


FIG. 3B

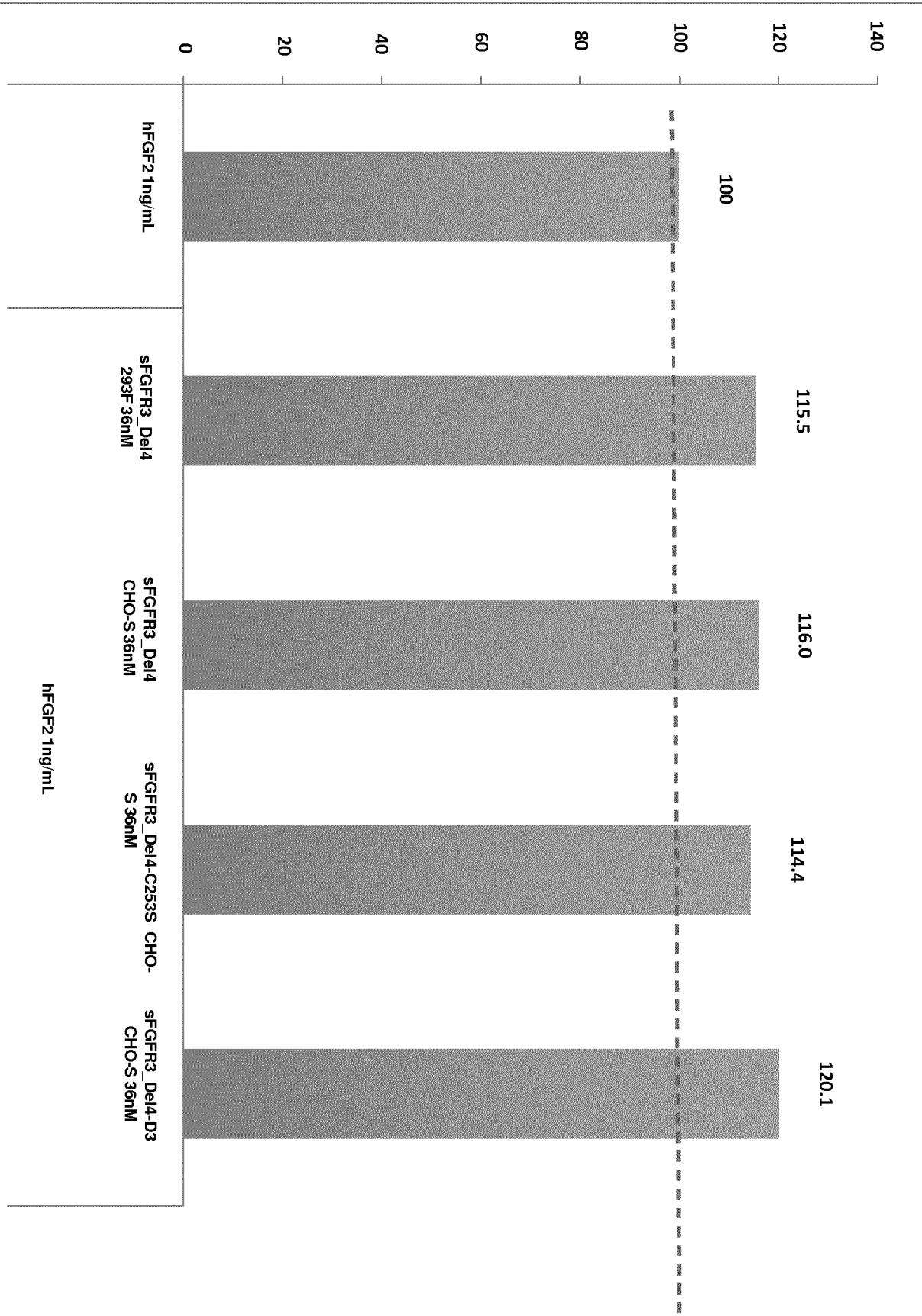


FIG. 4

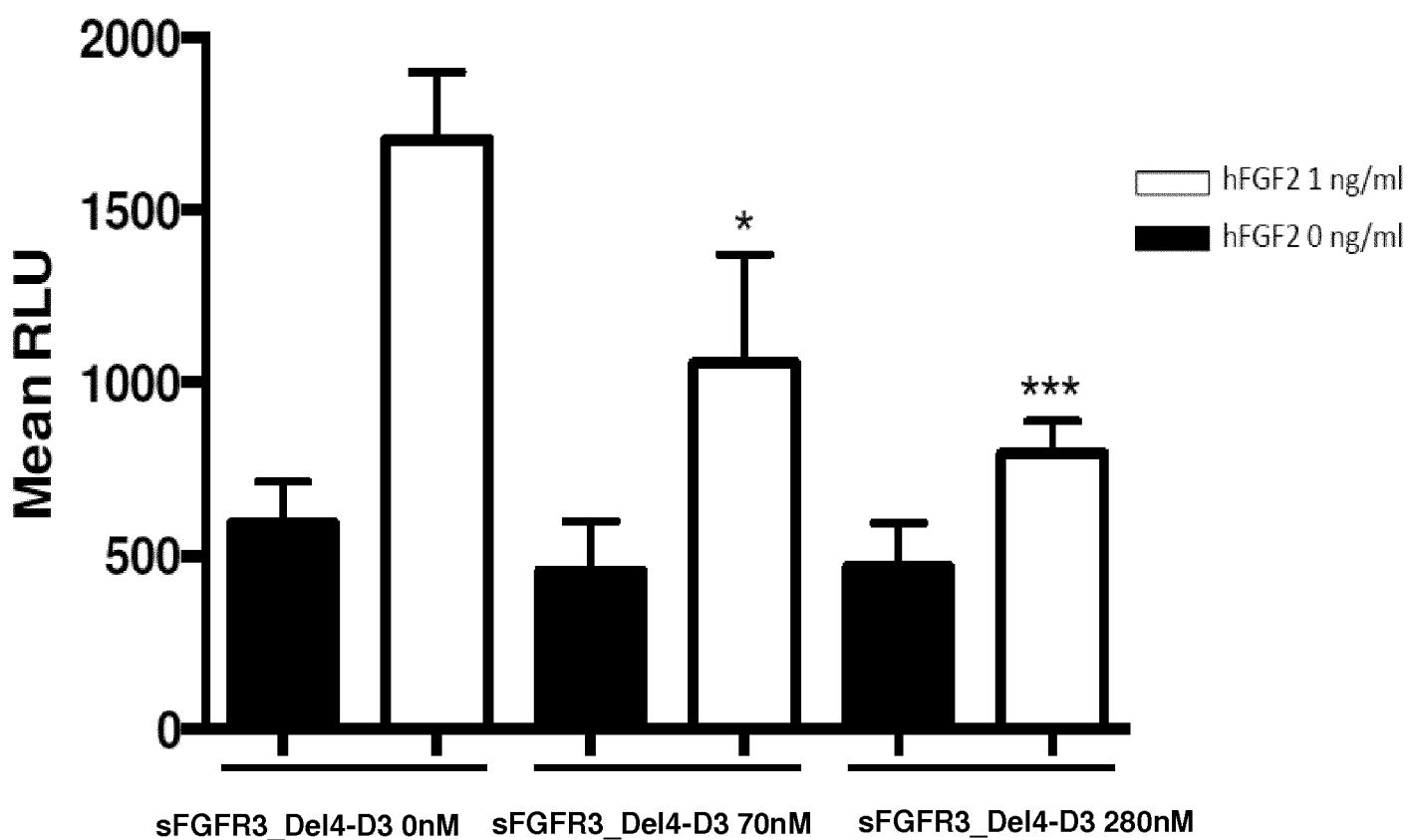
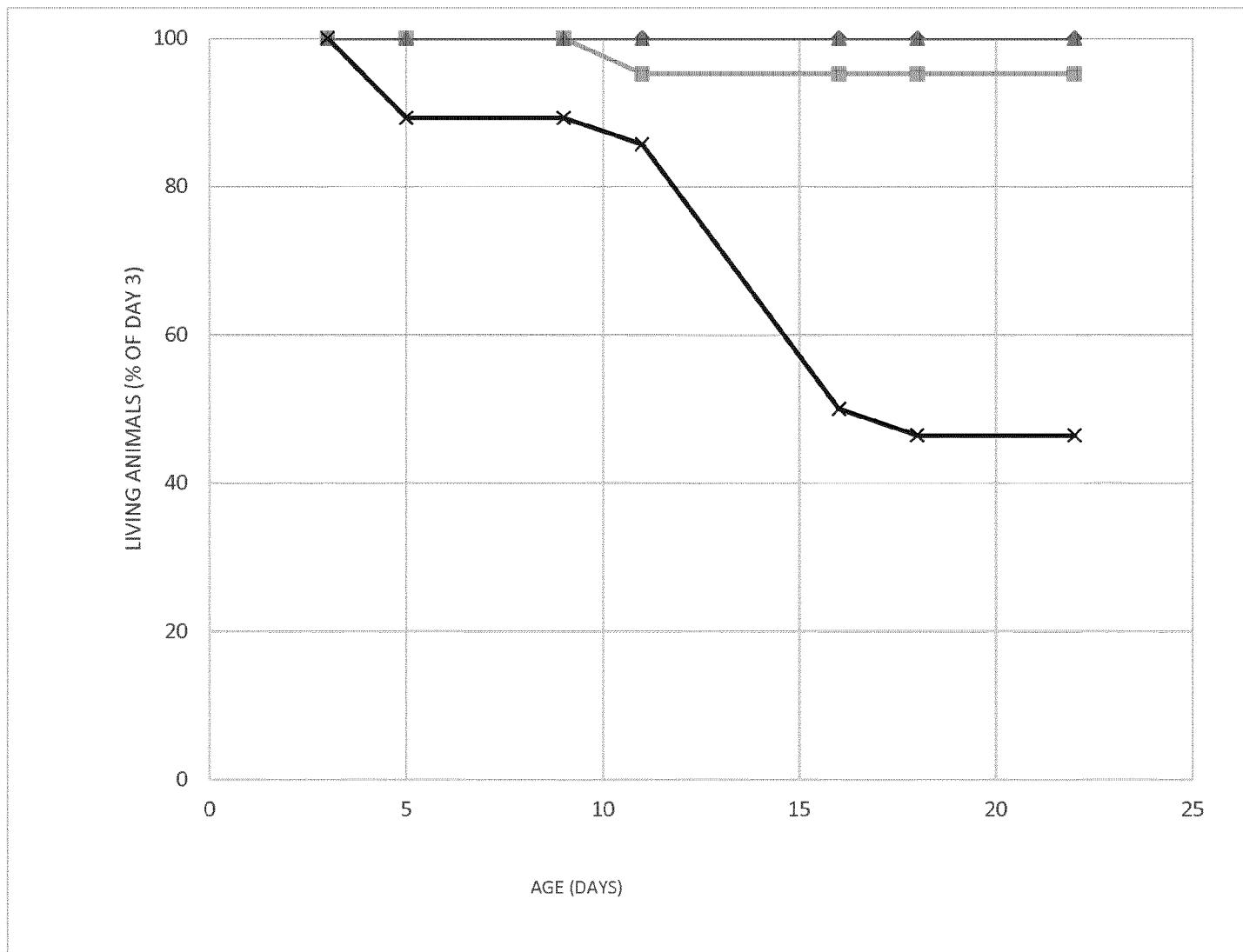
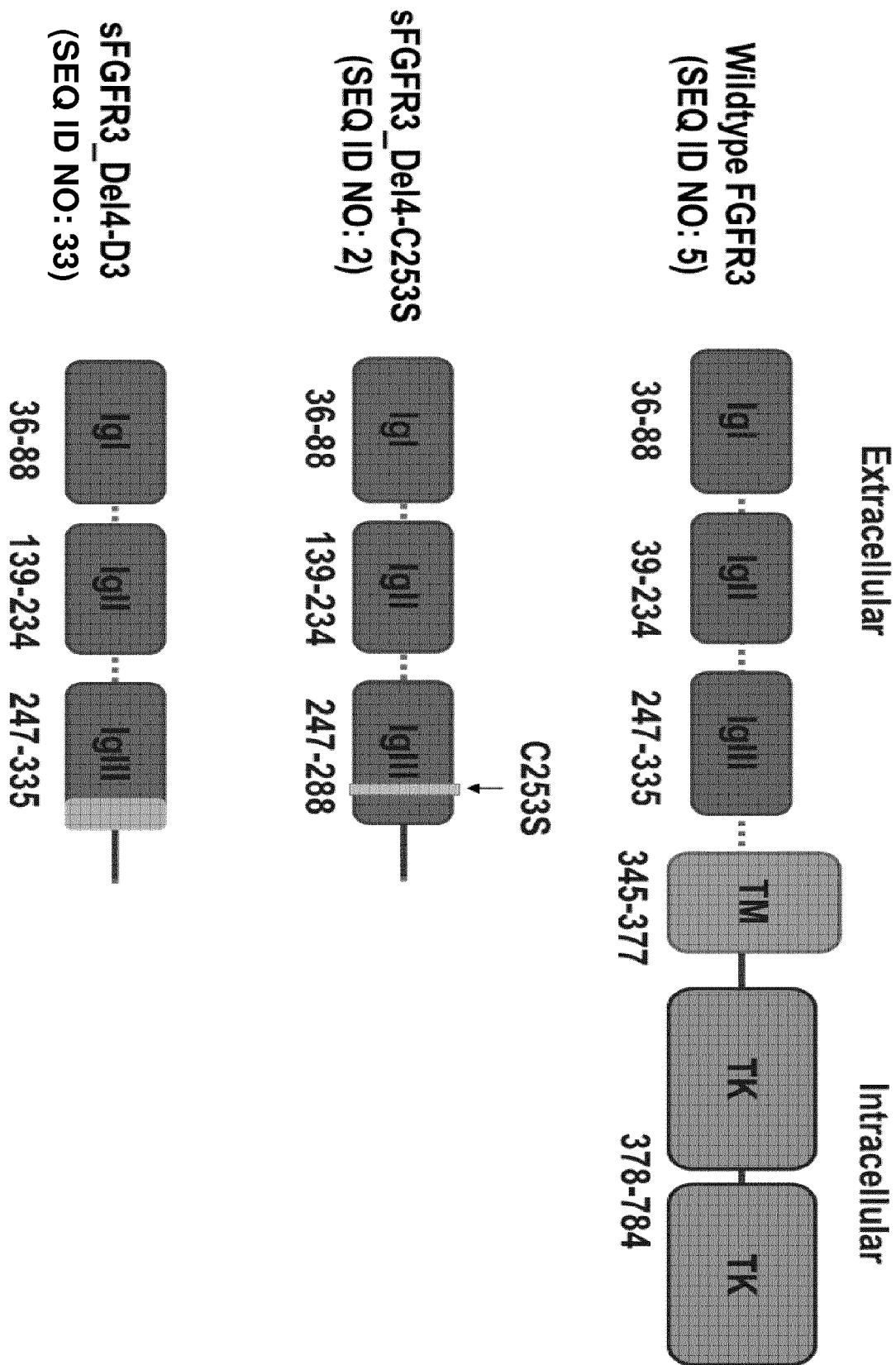


FIG. 5



- ▲ Fgfr3ach/+ sFGFR3\_Del4-D3 0.25 mg/kg
- WT PBS
- ✖ Fgfr3ach/+ PBS

FIG. 6



## FIG. 7A

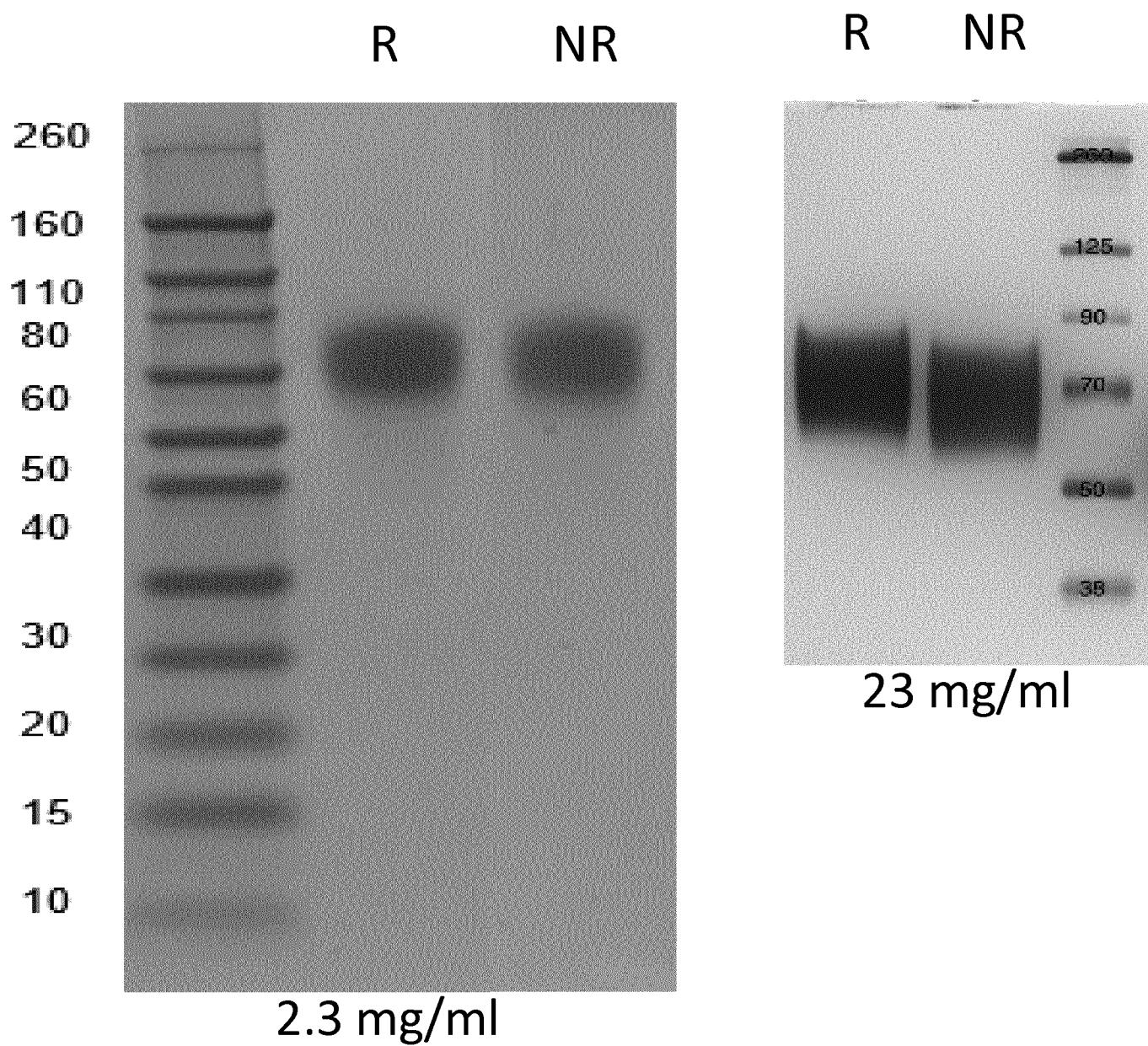


FIG. 7B

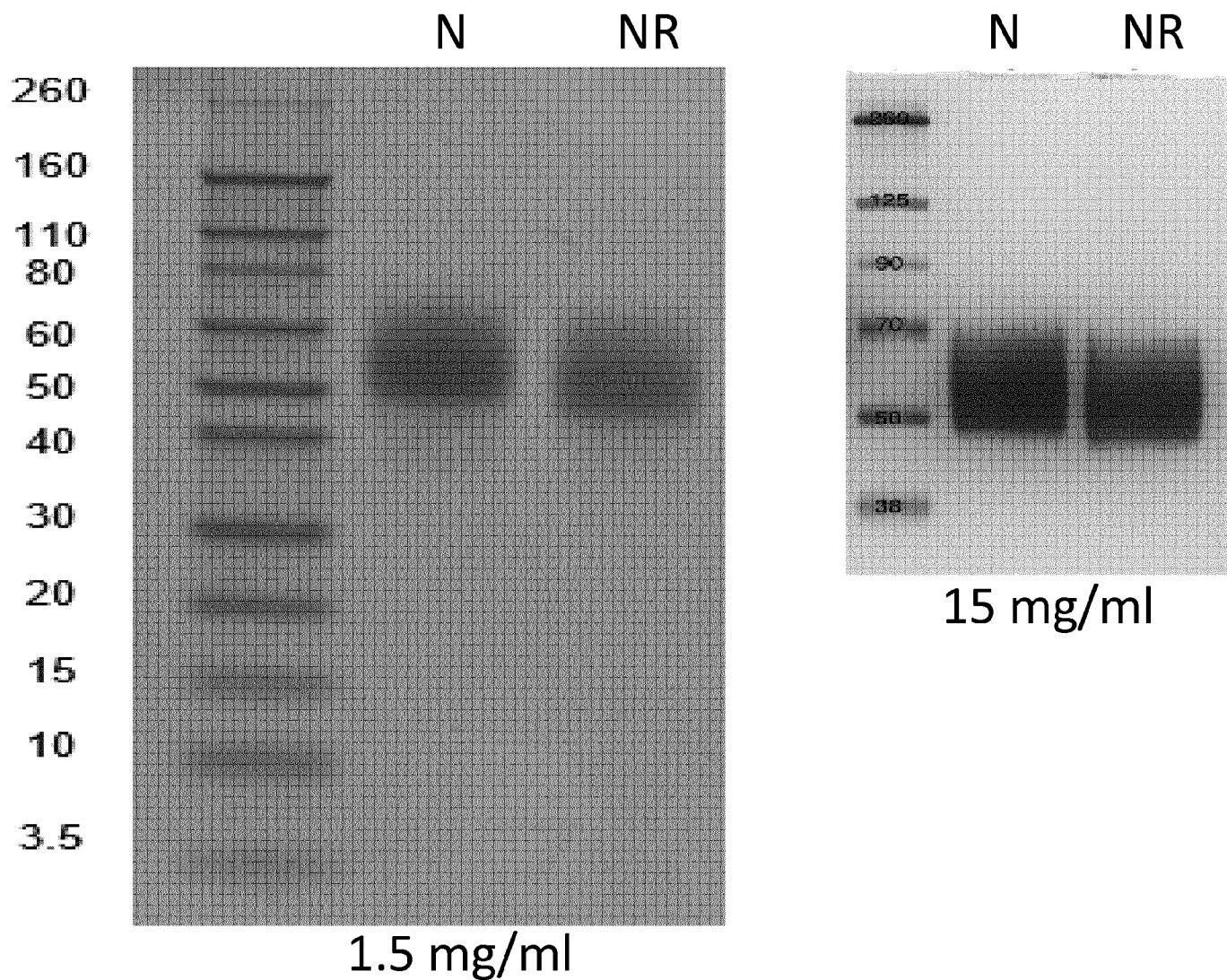


FIG. 8A

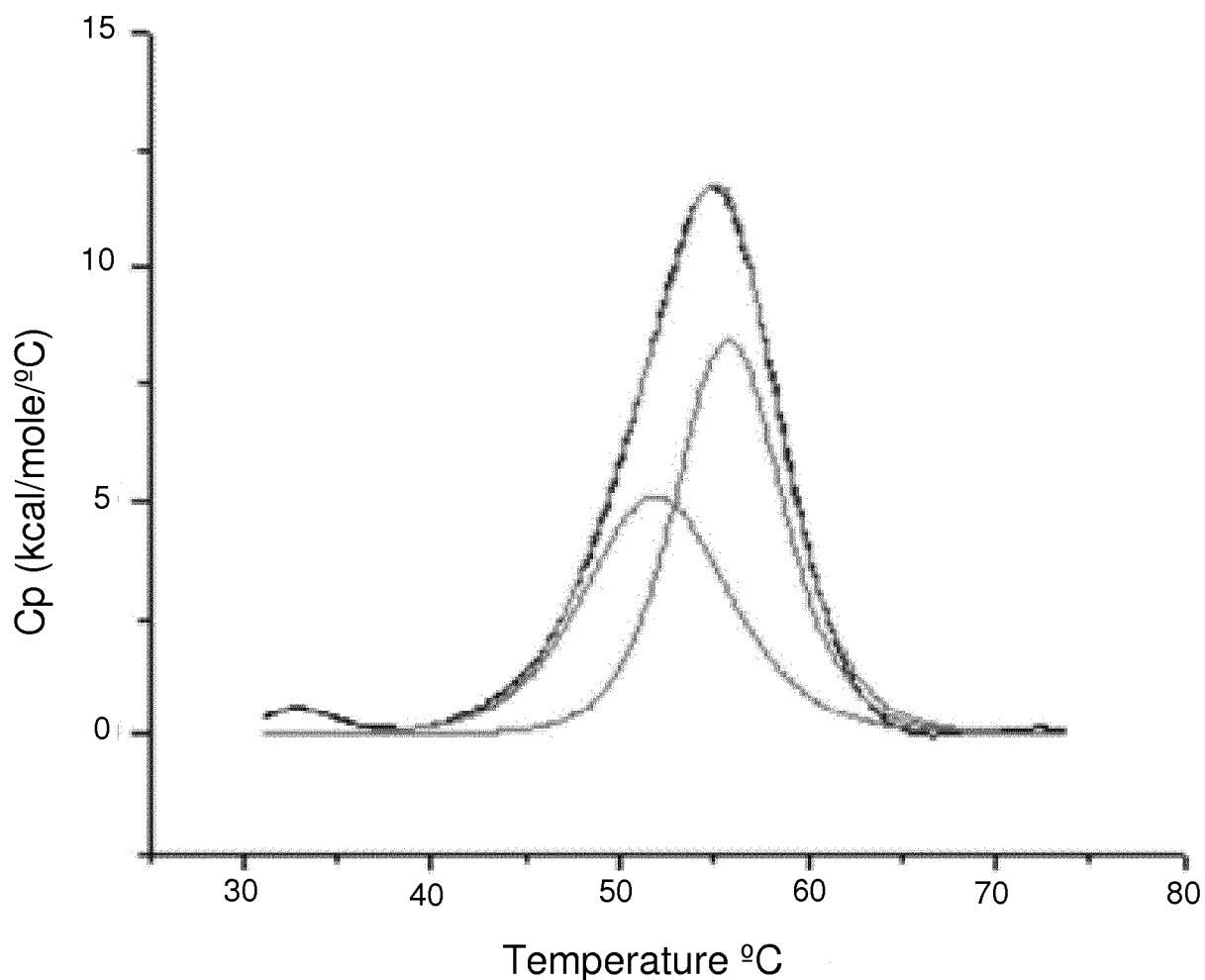
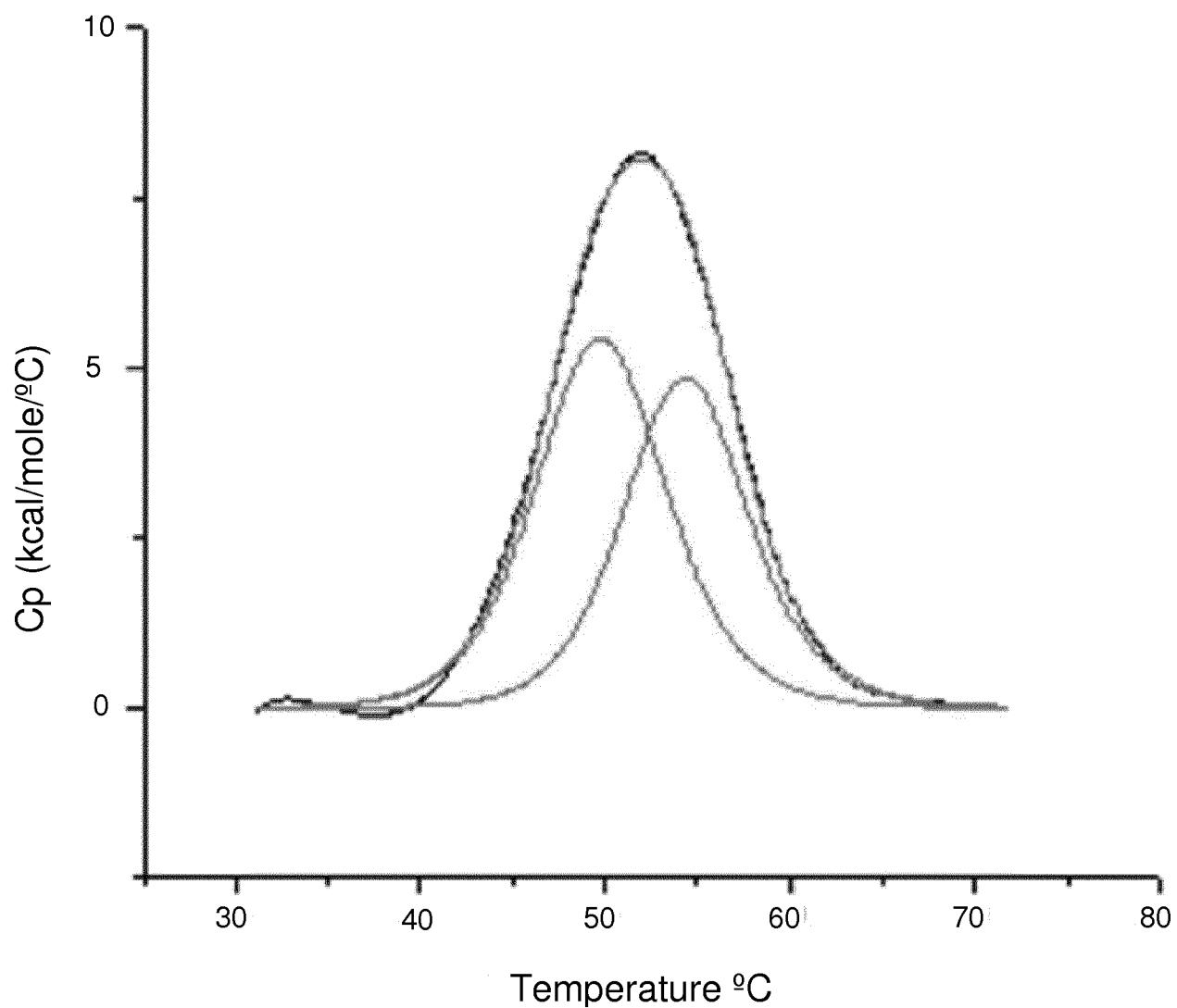
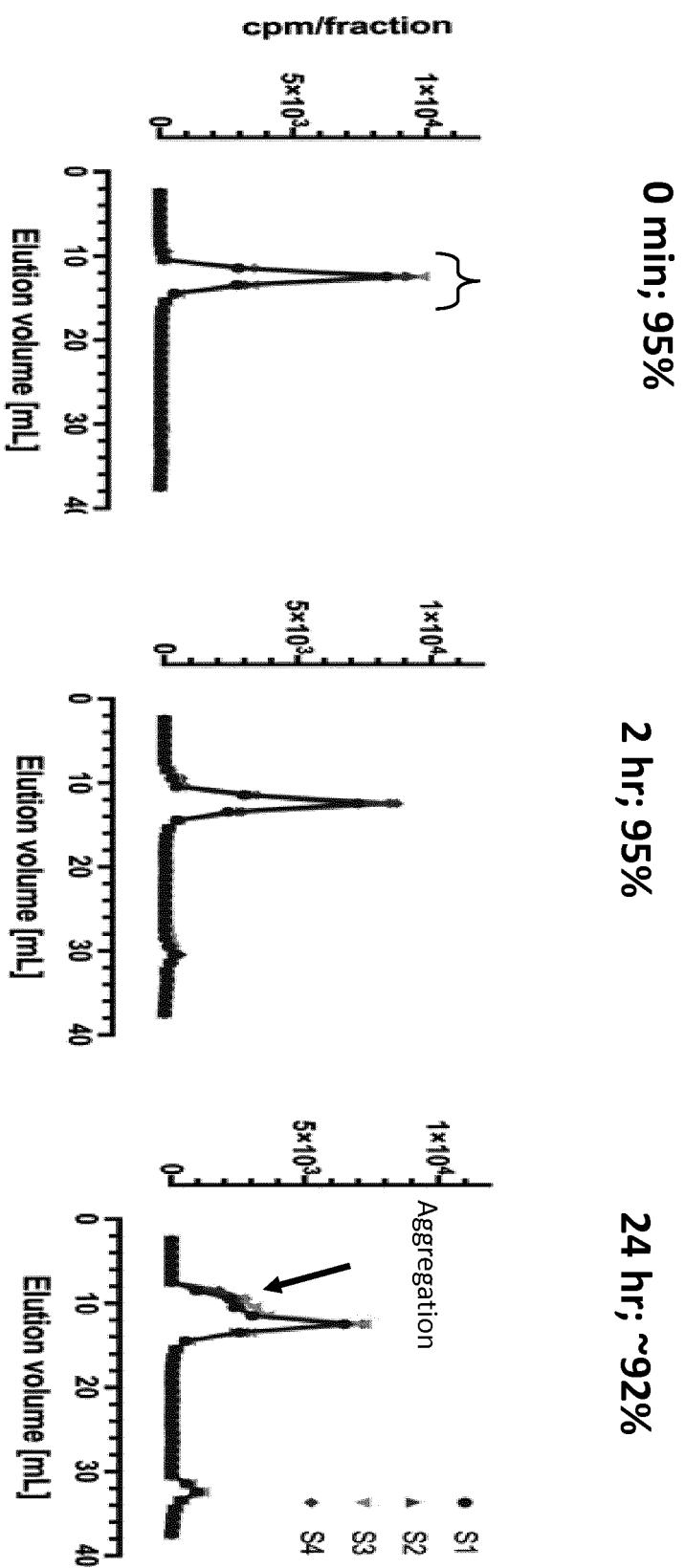
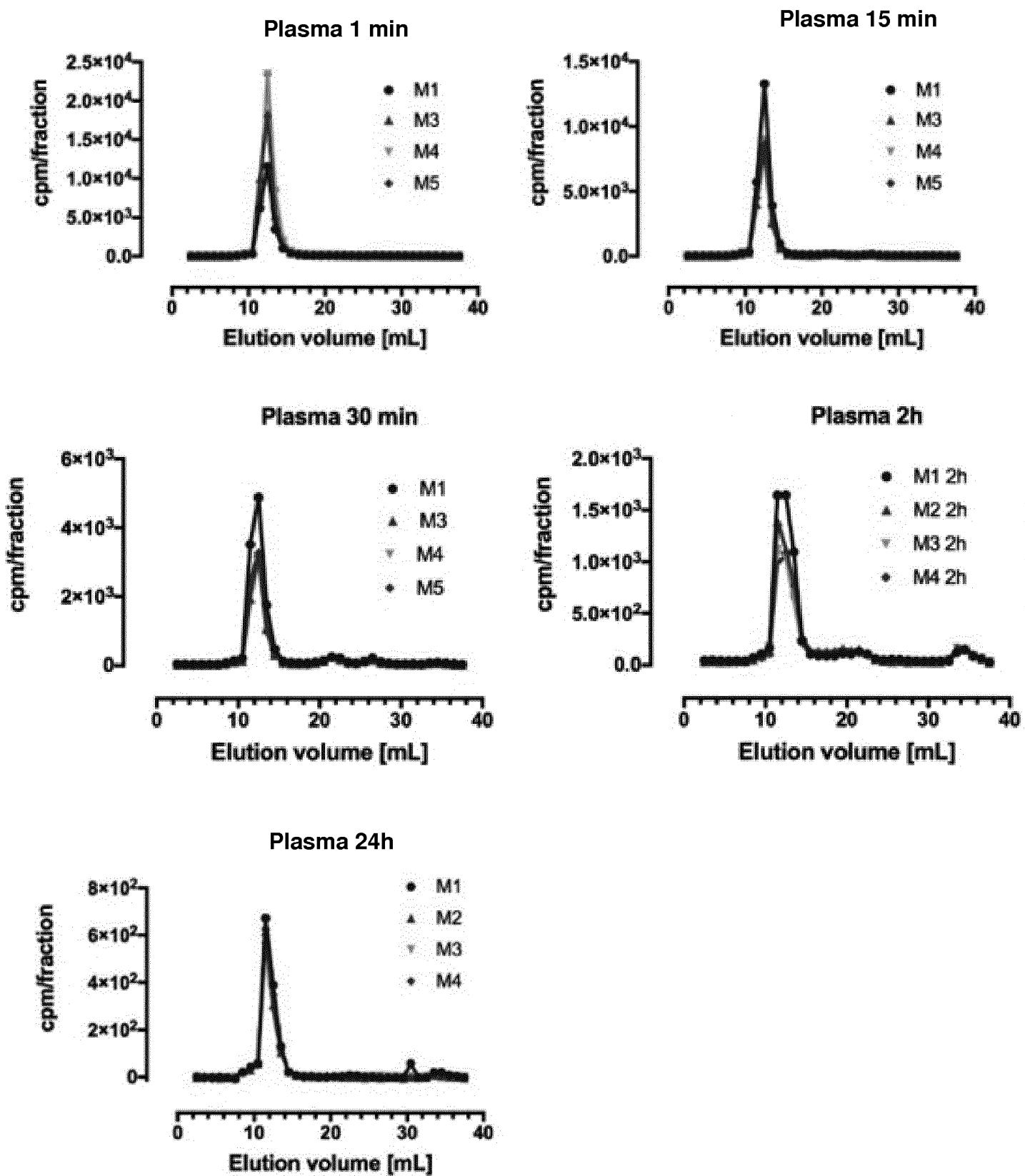


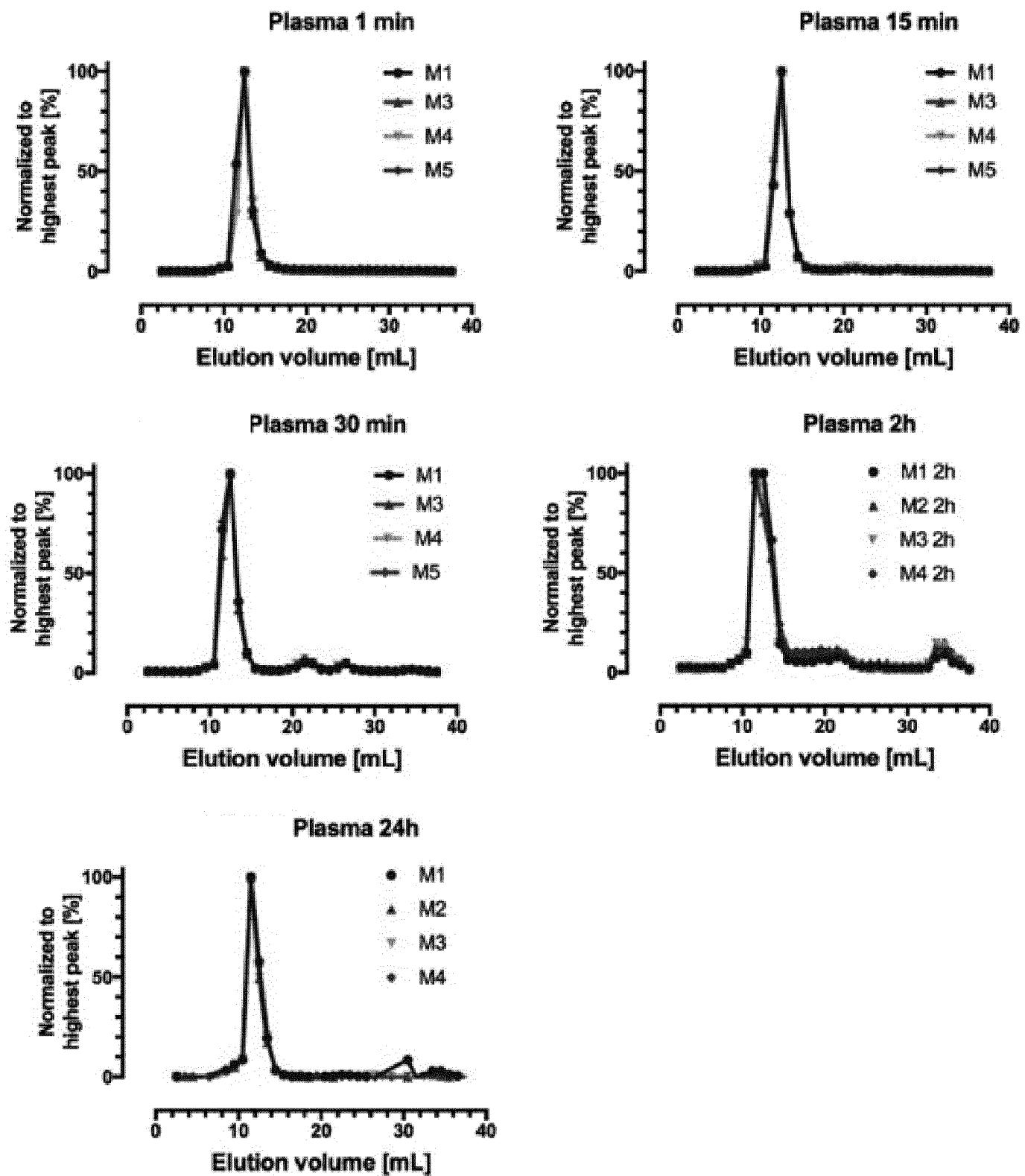
FIG. 8B







## FIG. 9B cont.



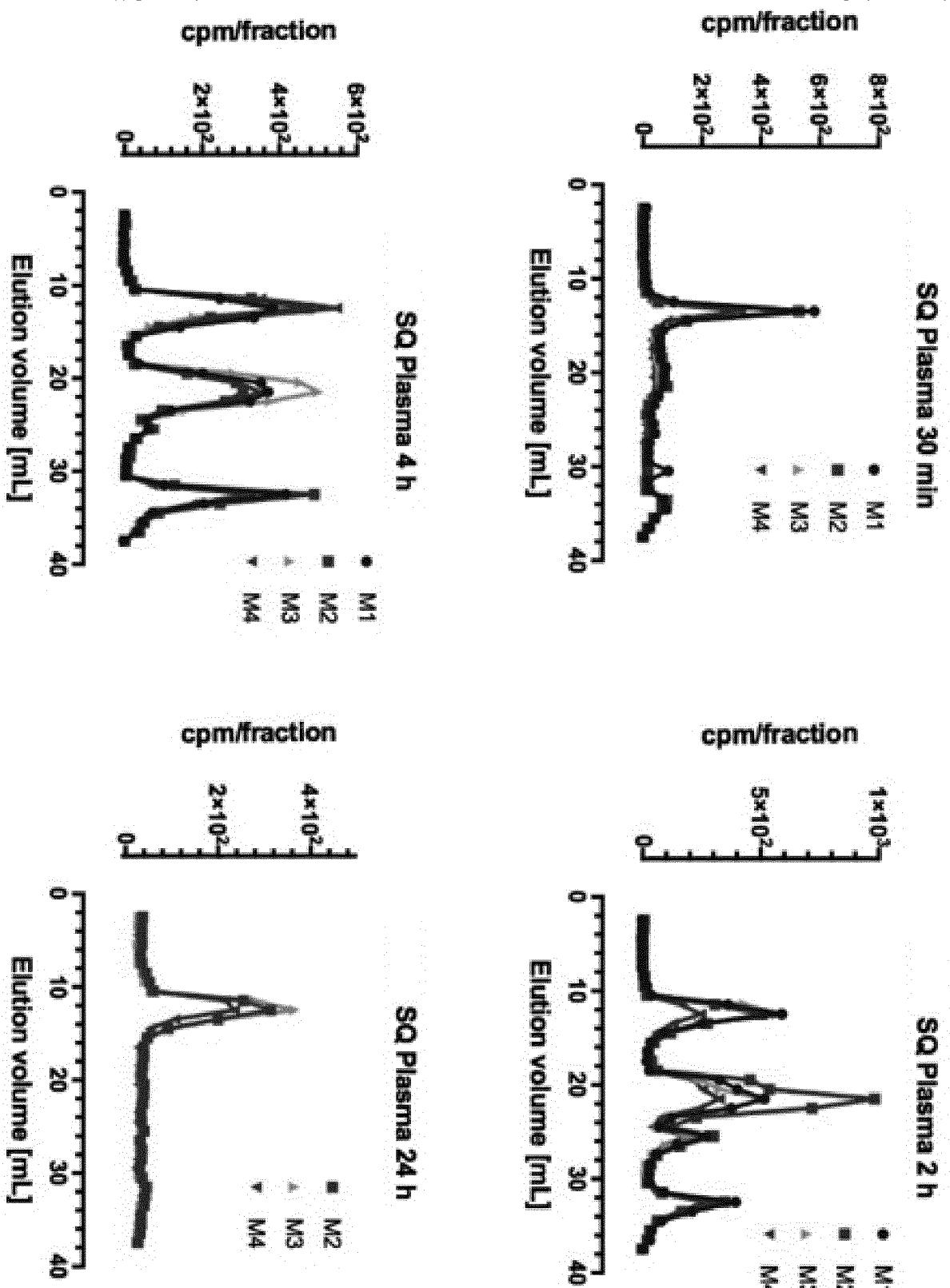


FIG. 9C cont.

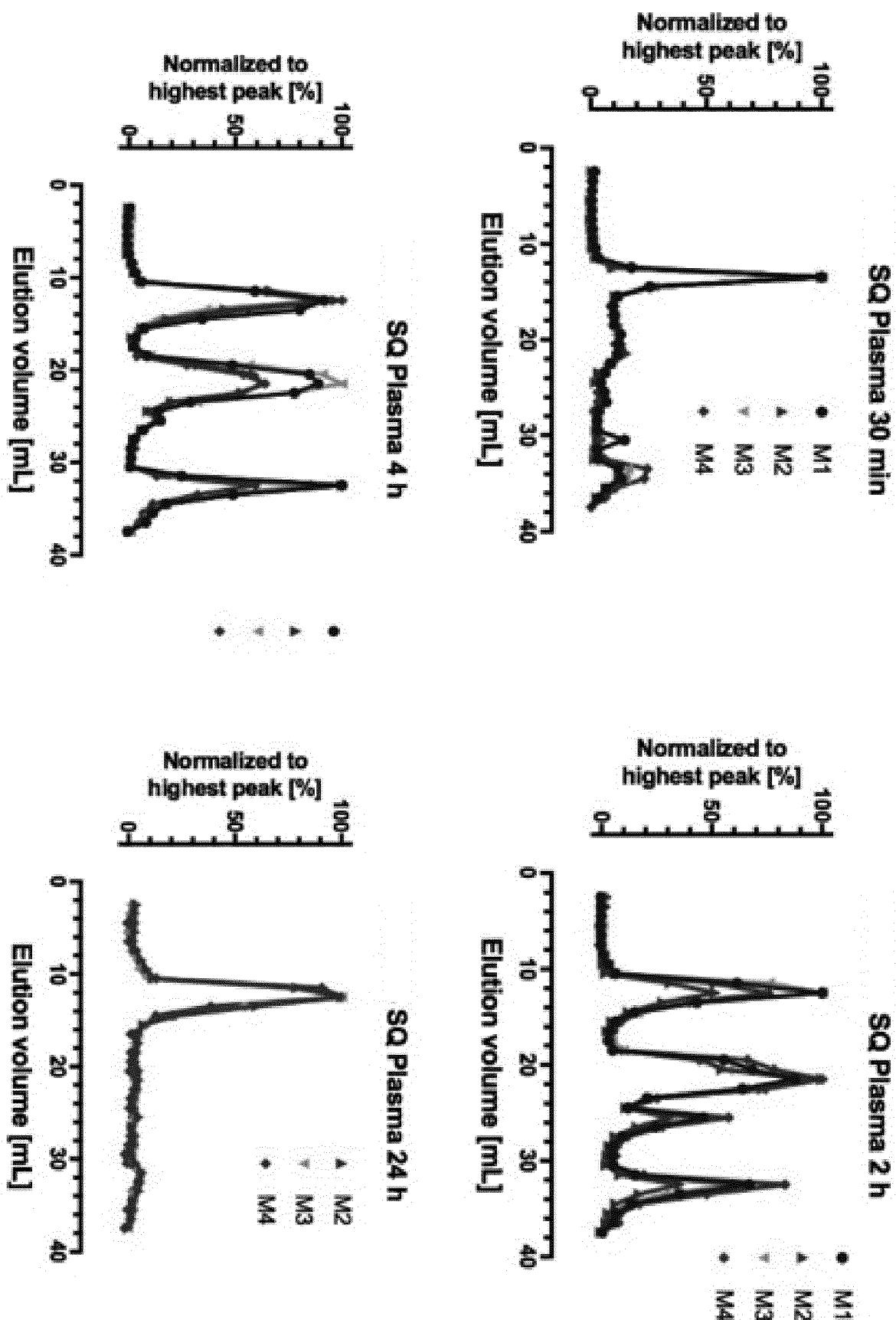


FIG. 10A

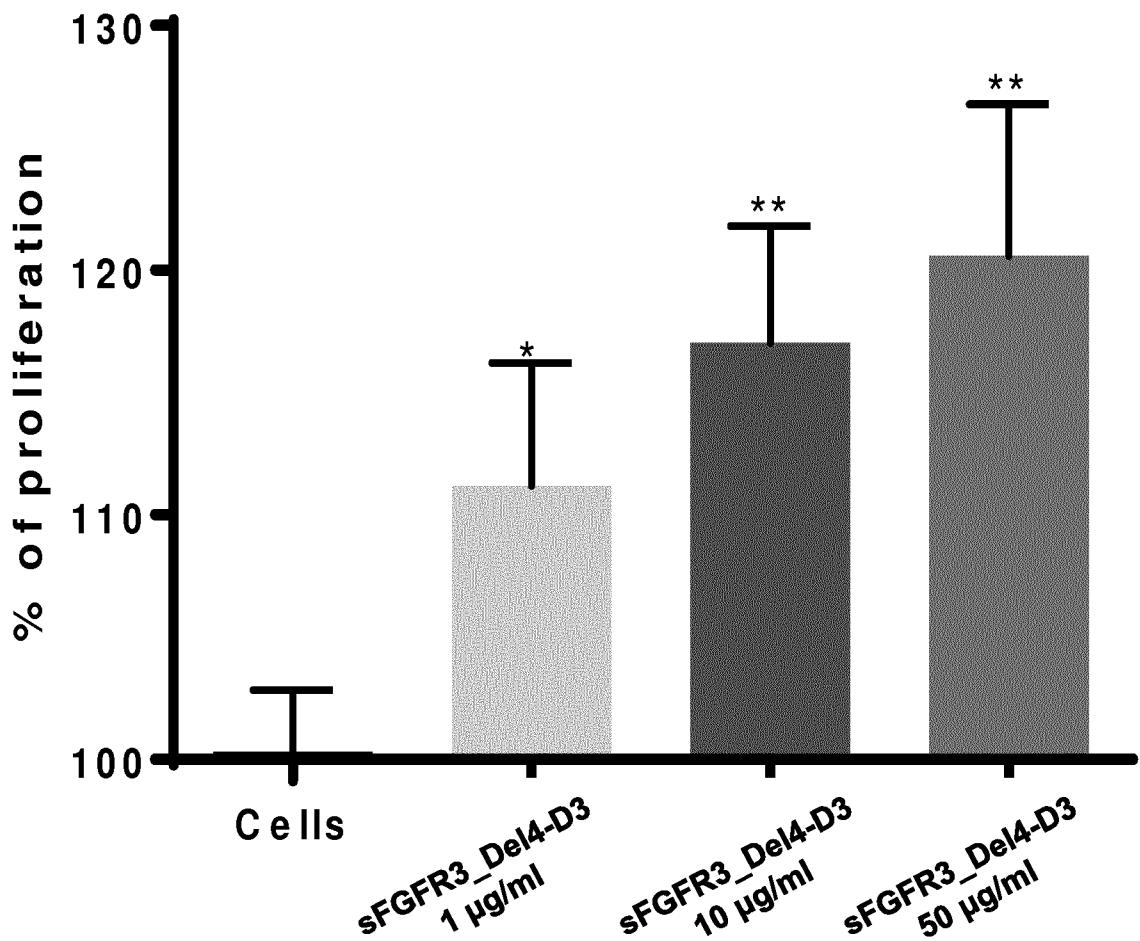


FIG. 10B

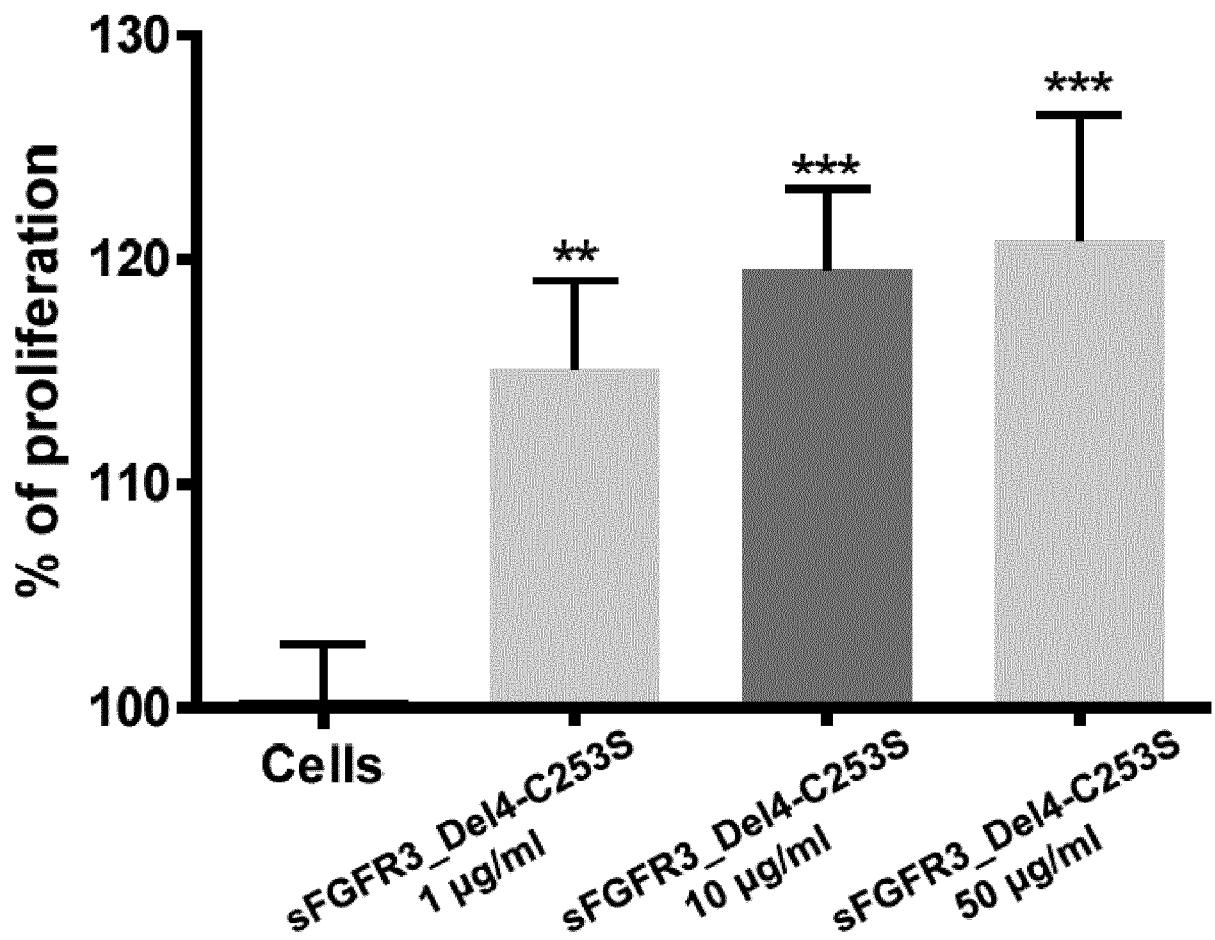


FIG. 11

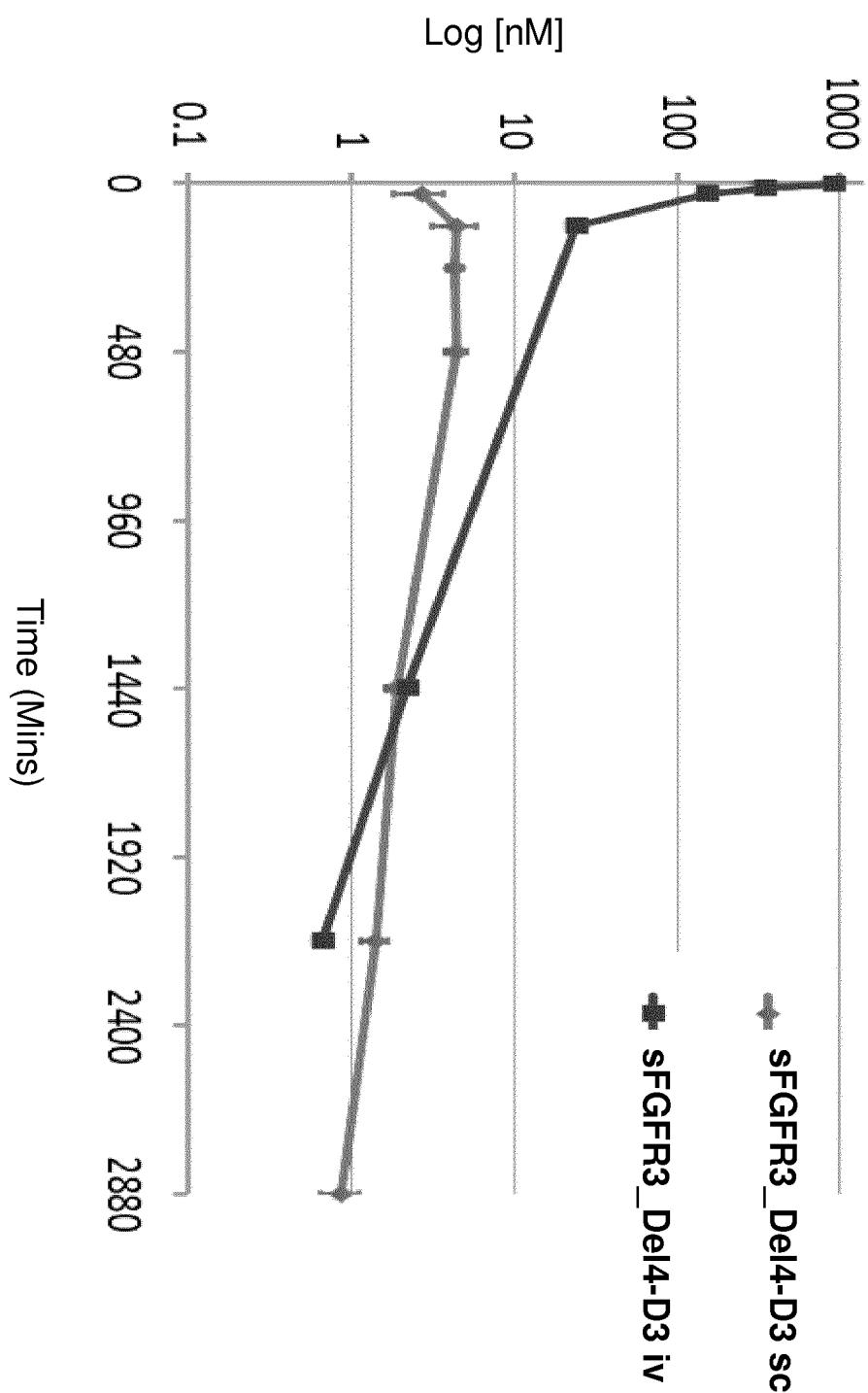


FIG. 12

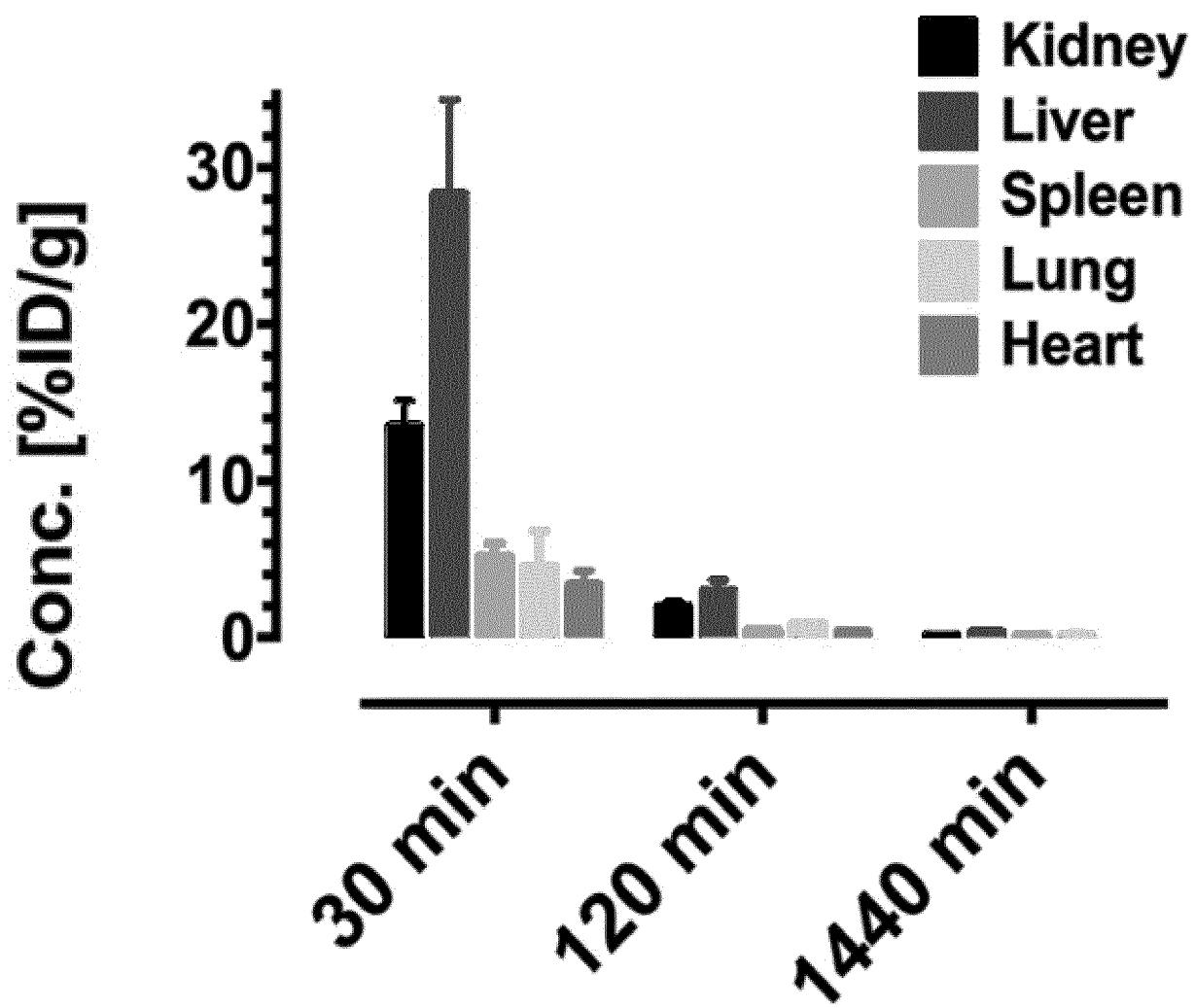
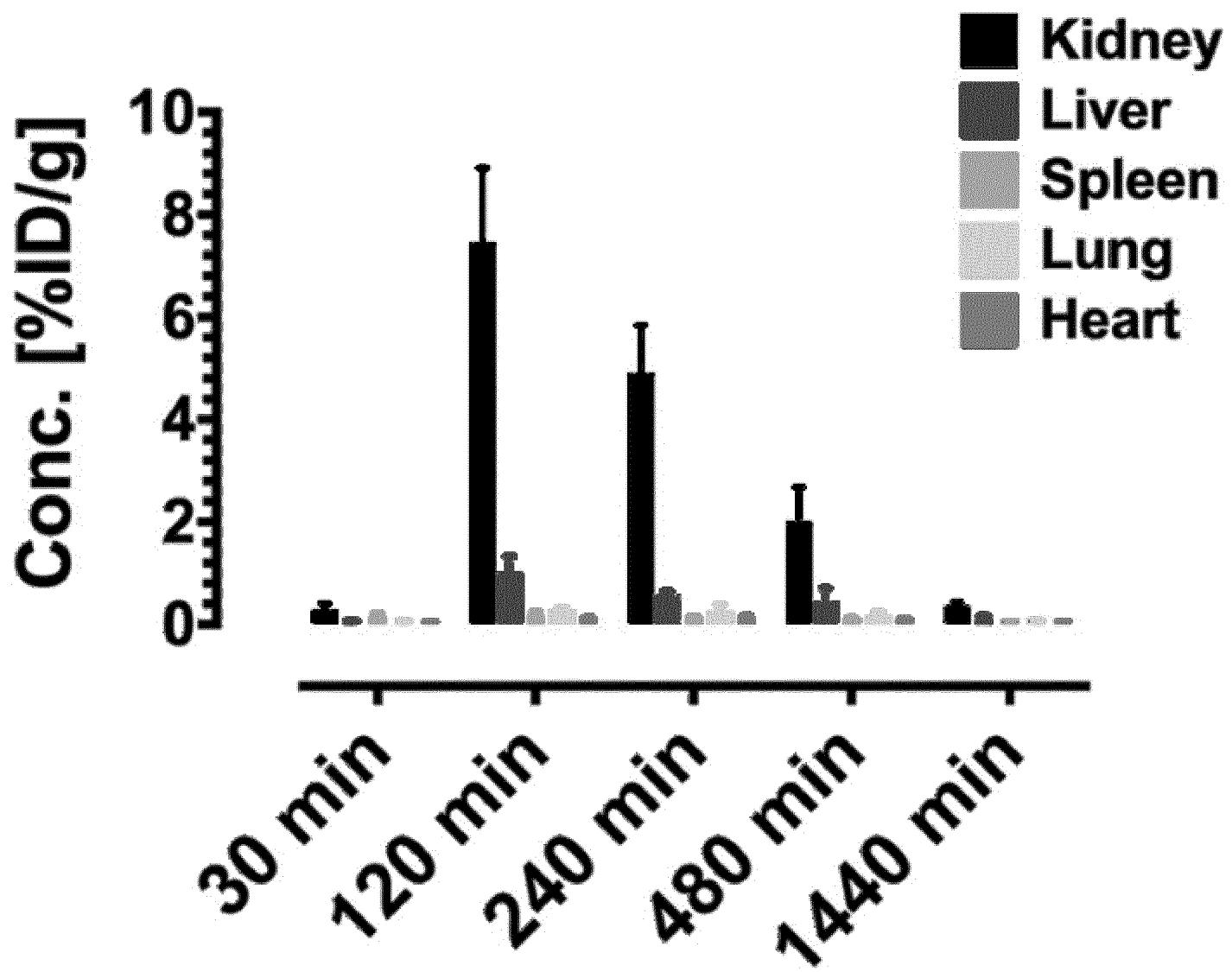


FIG. 13



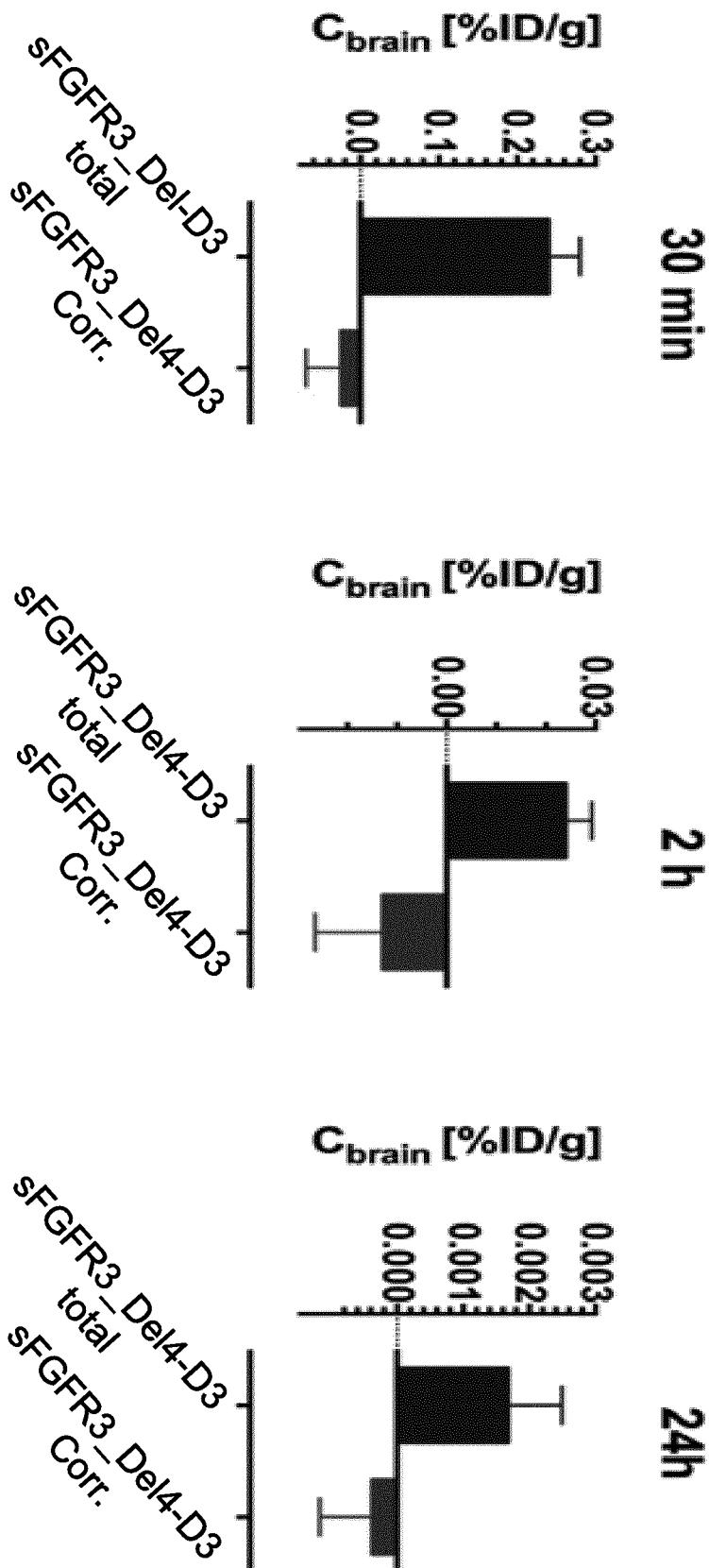


FIG. 14B



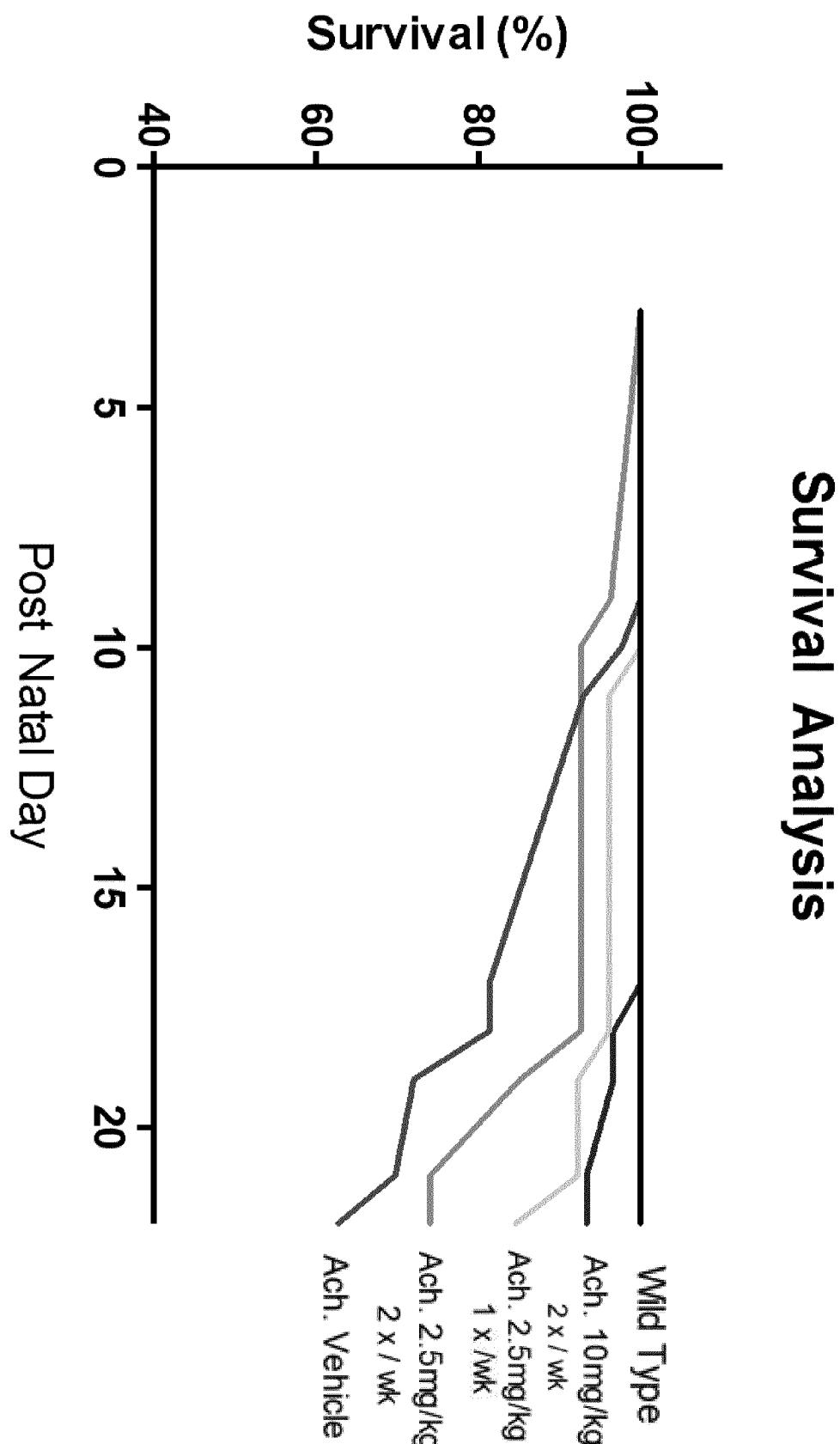
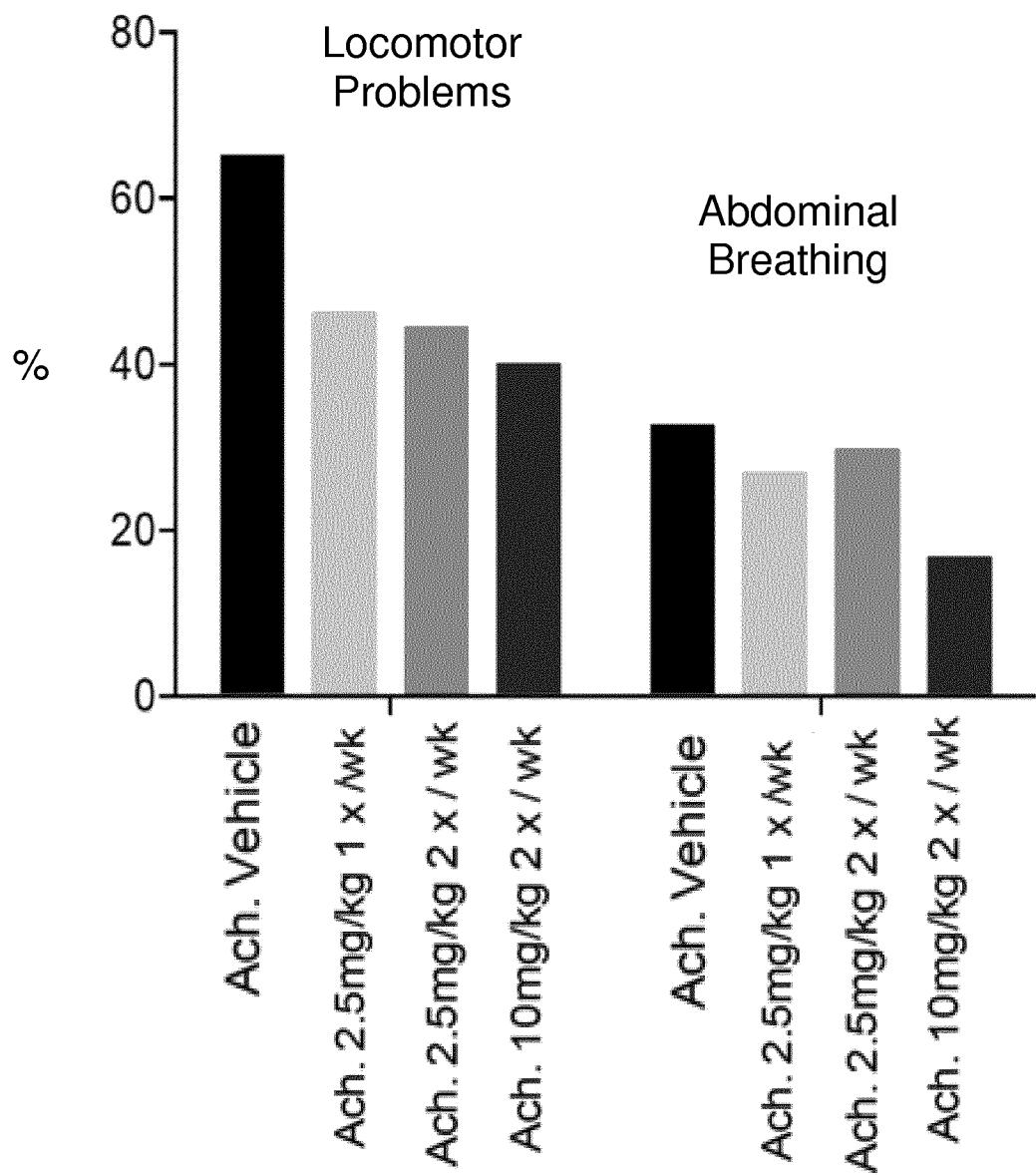


FIG. 15

FIG. 16



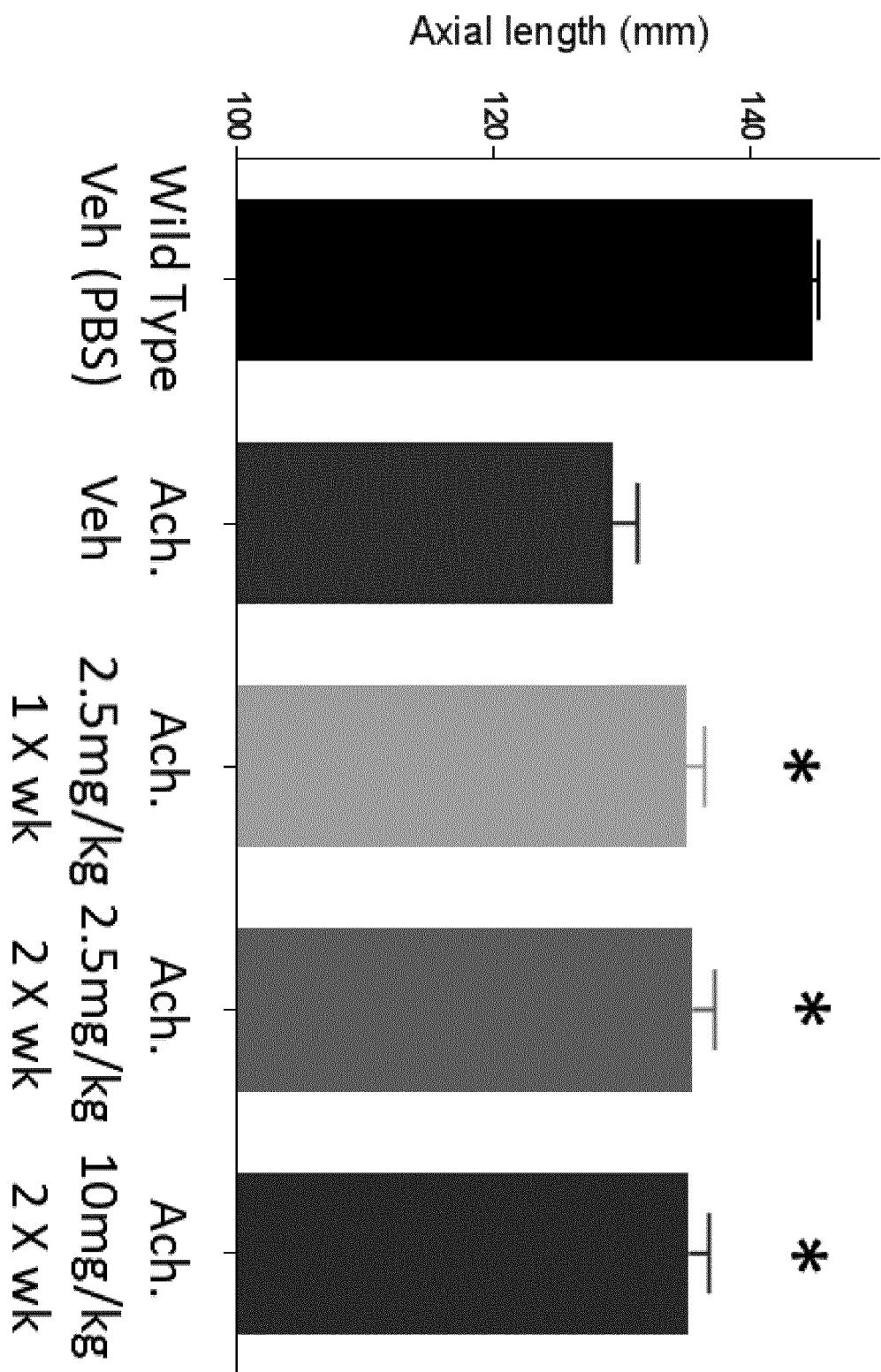


FIG. 17A

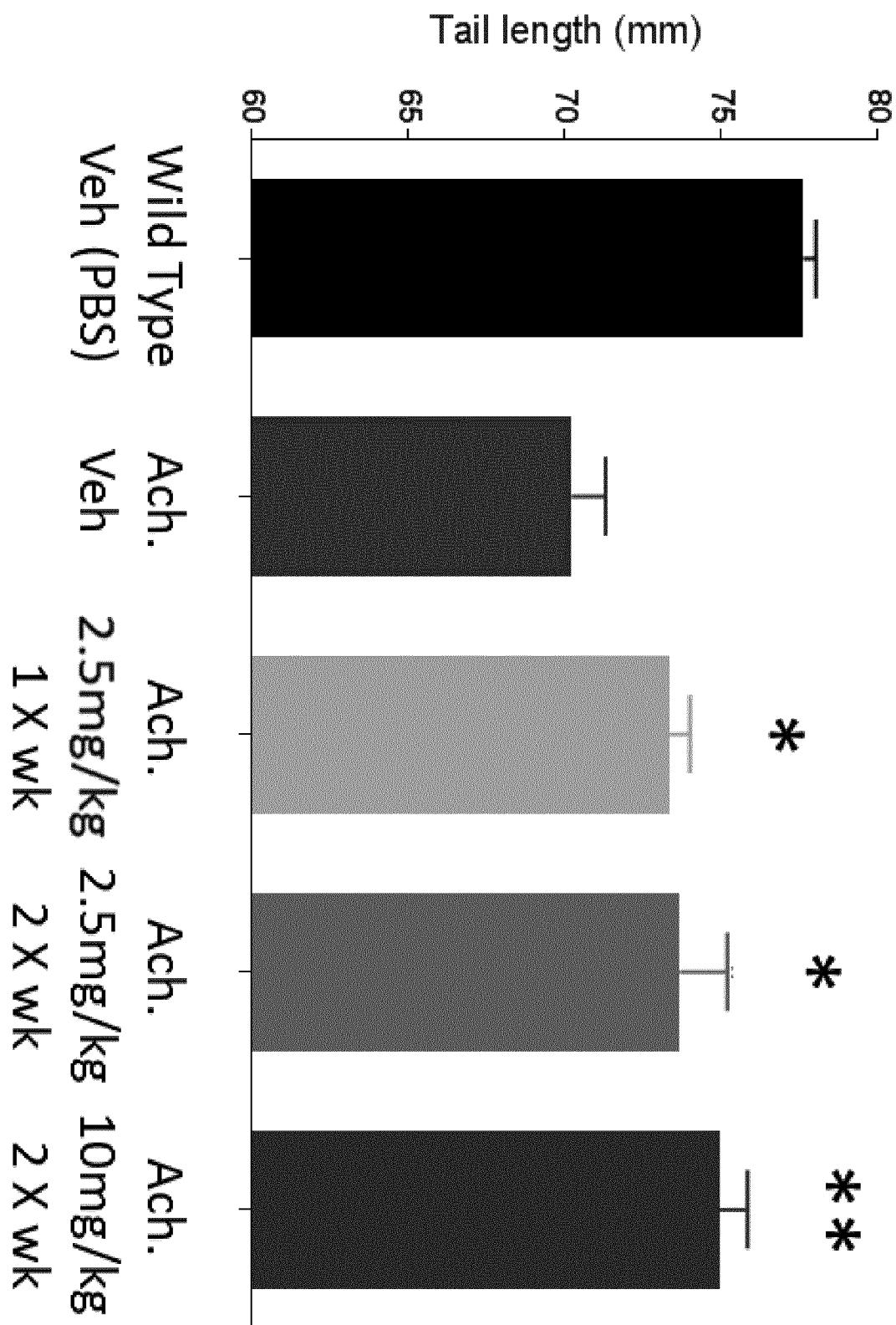


FIG. 17B

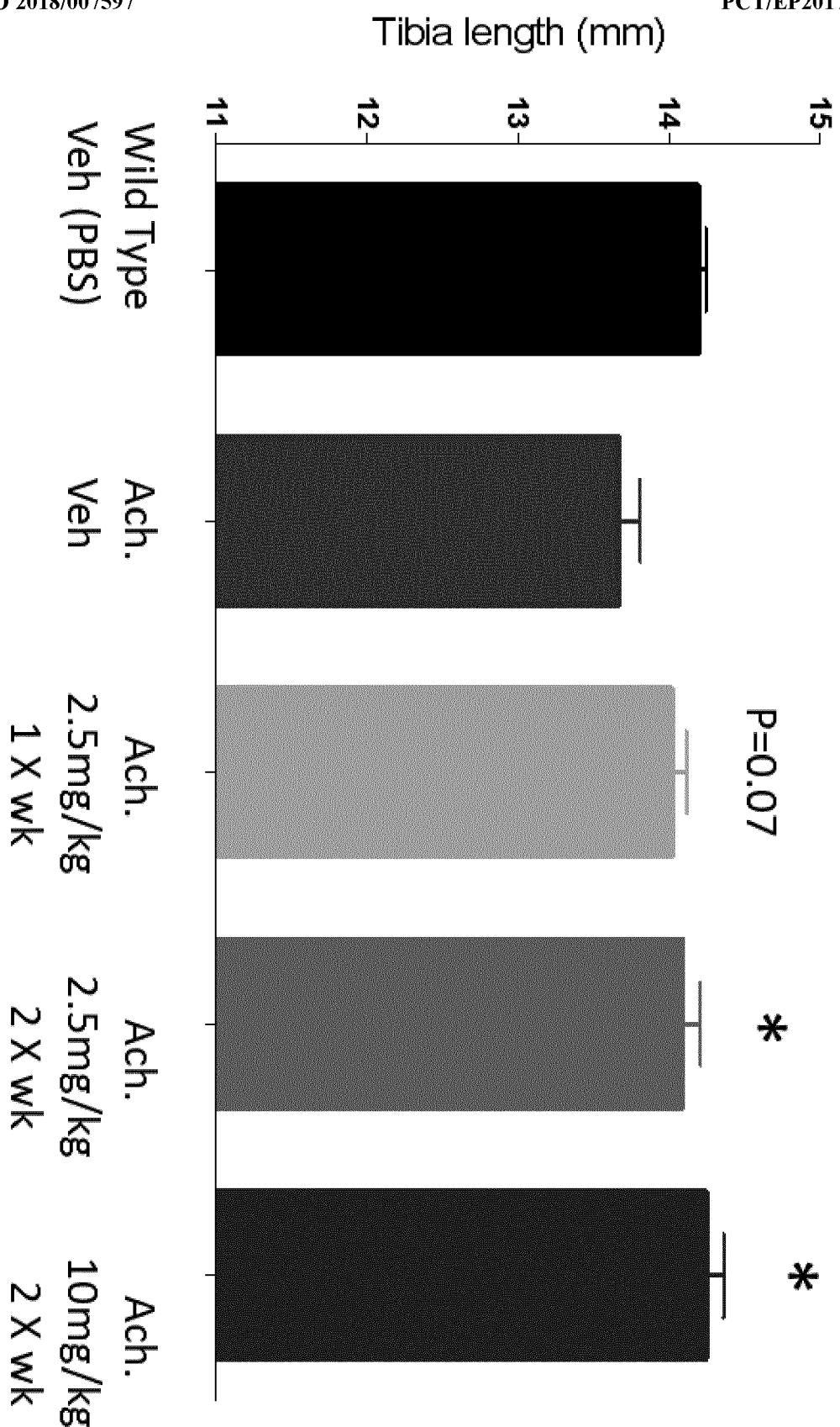
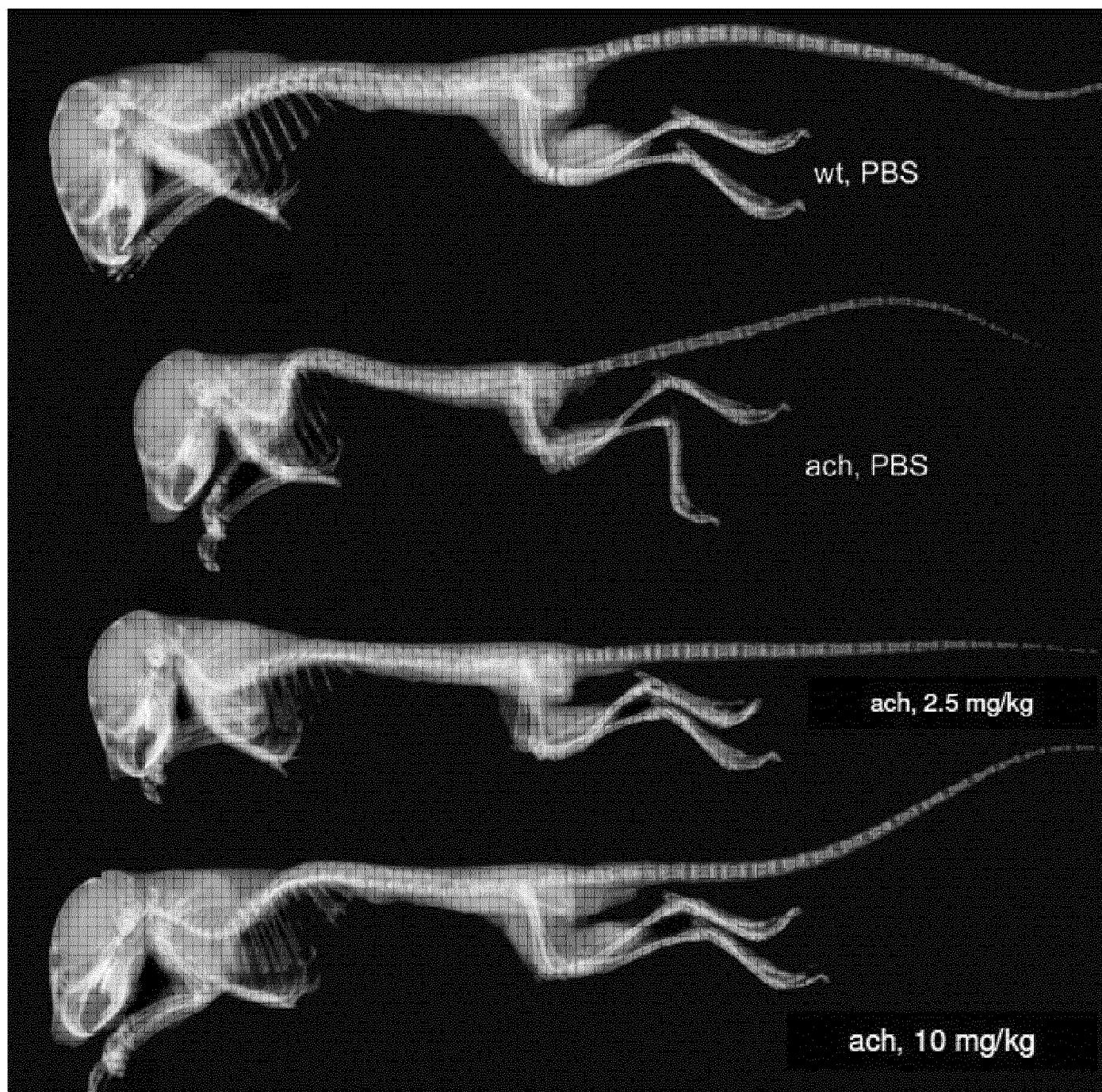
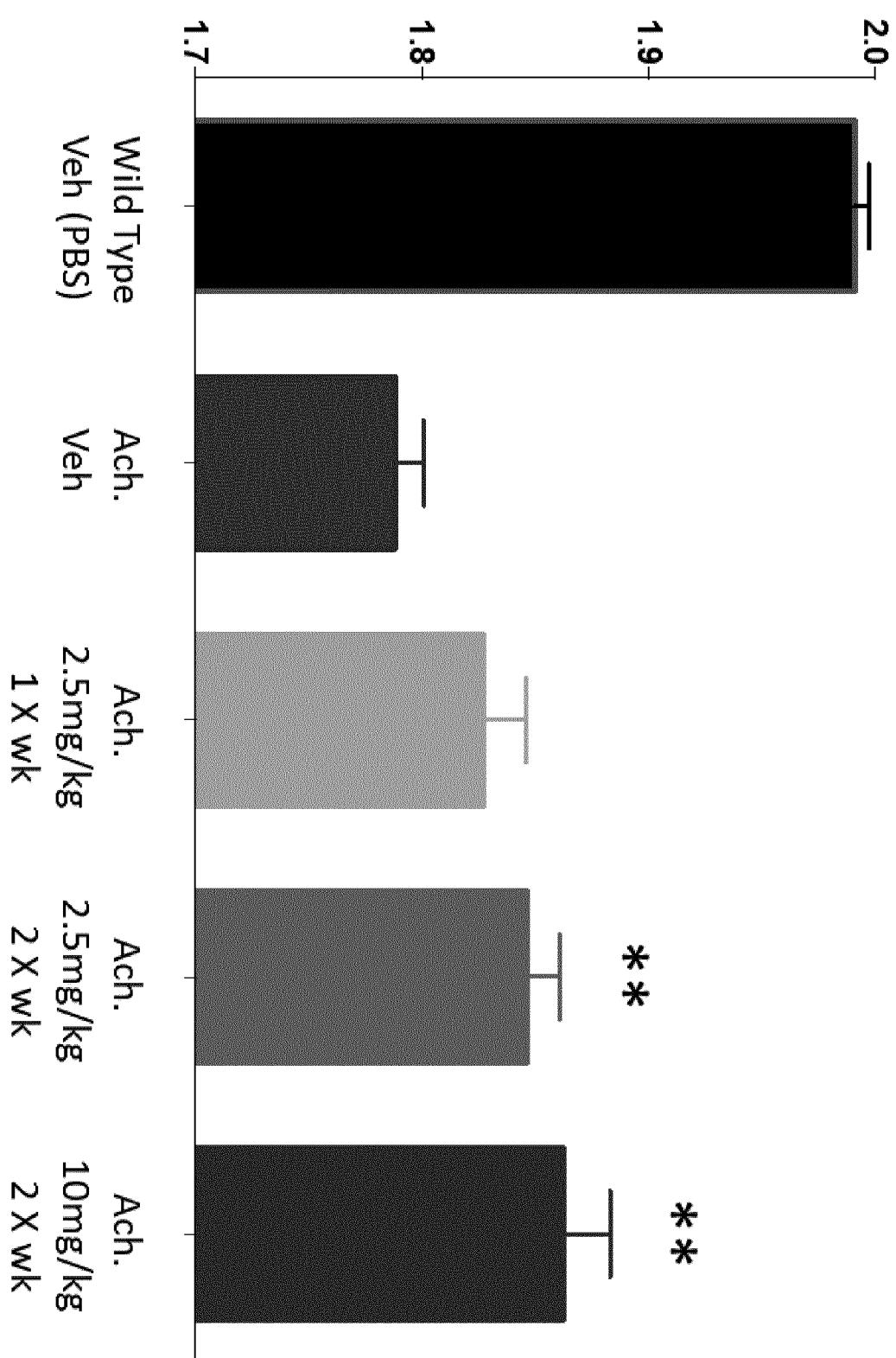


FIG. 17C

FIG. 17D





## FIG. 18B



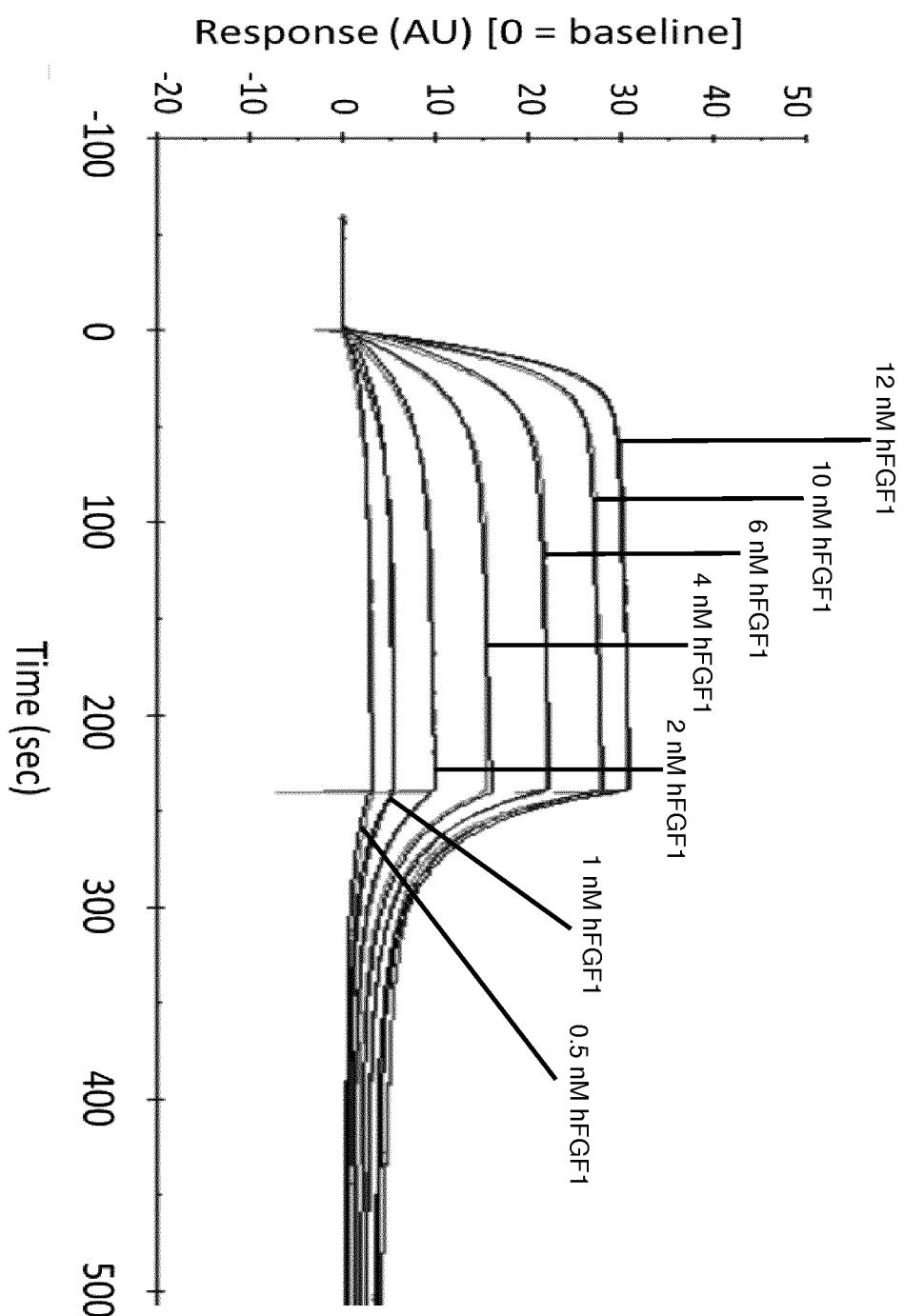


FIG. 19A

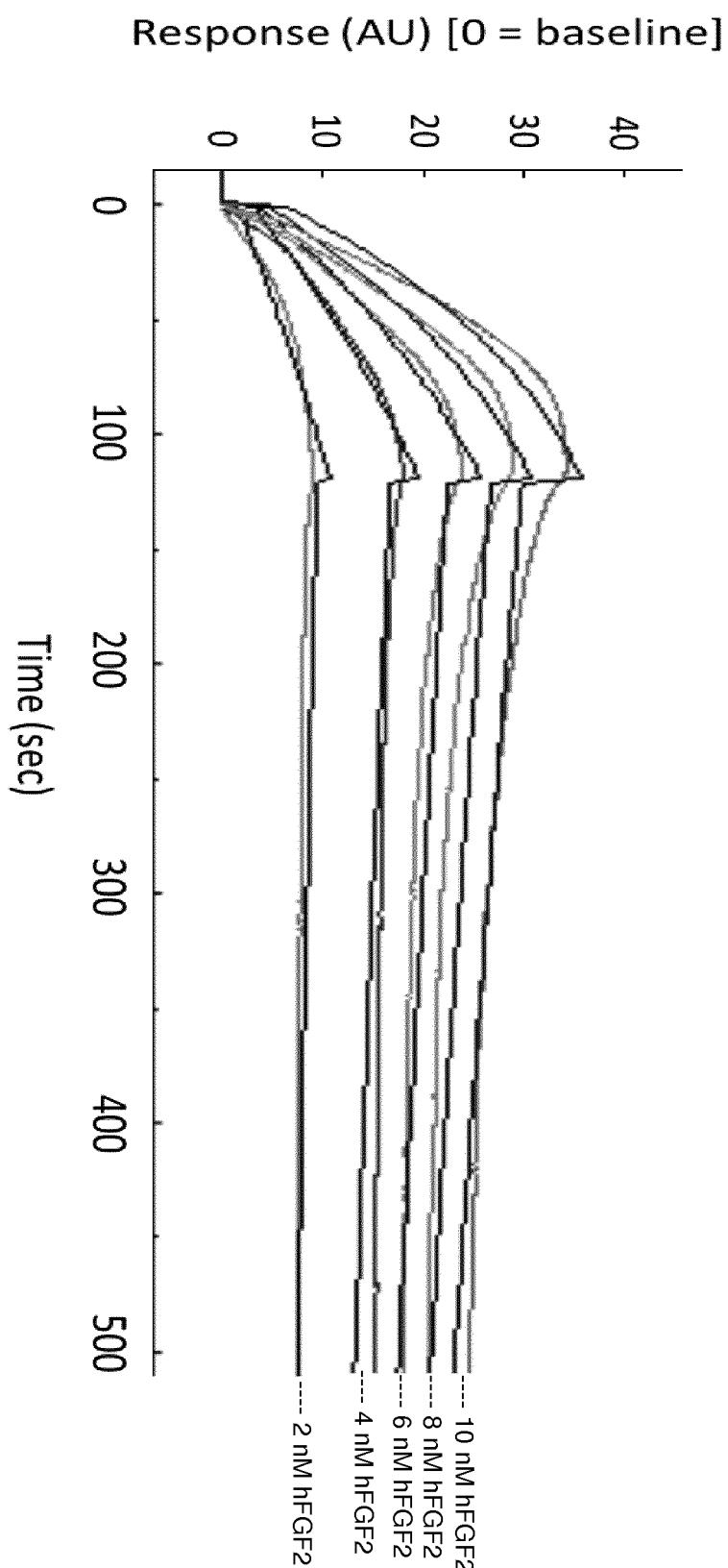


FIG. 19B

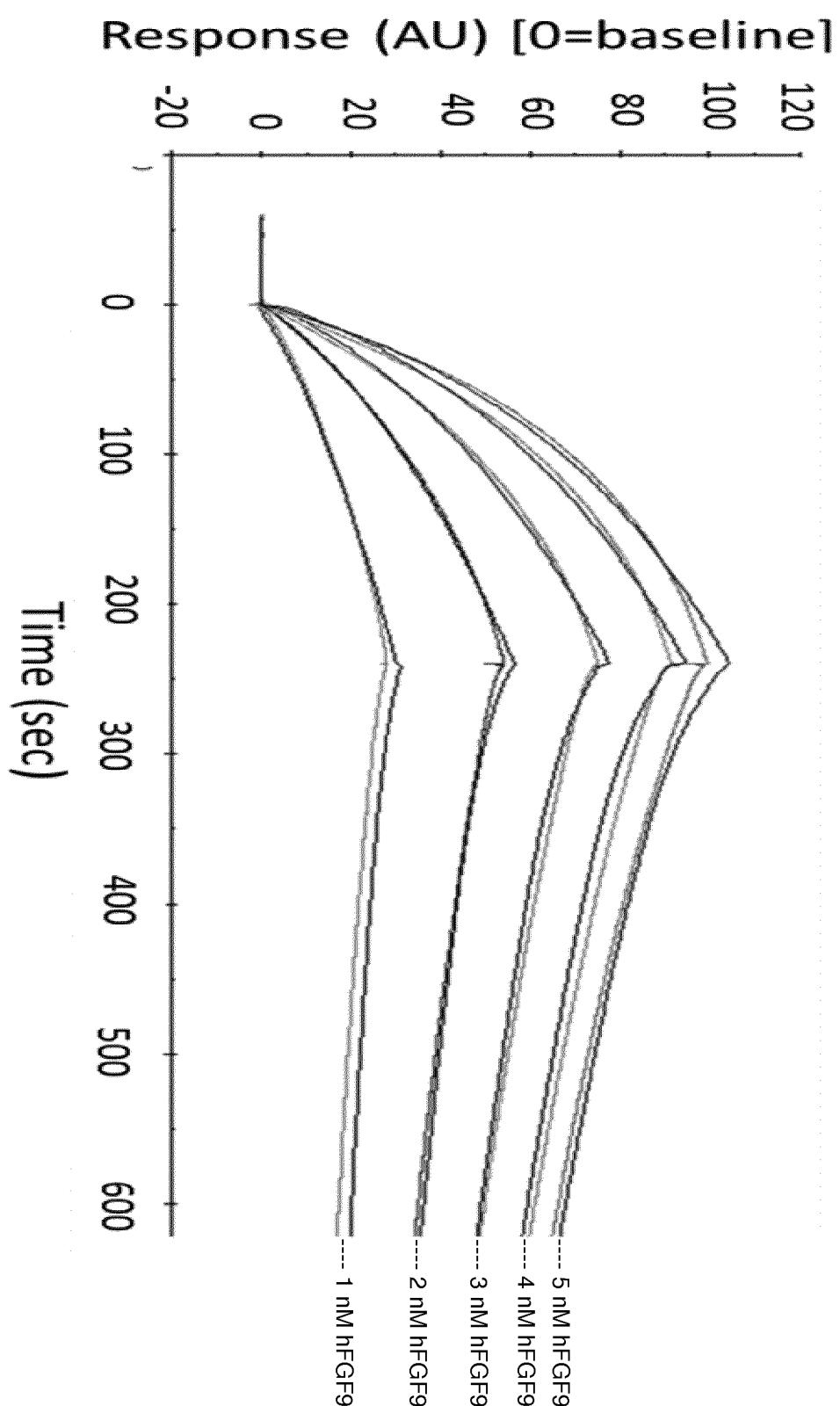


FIG. 19C

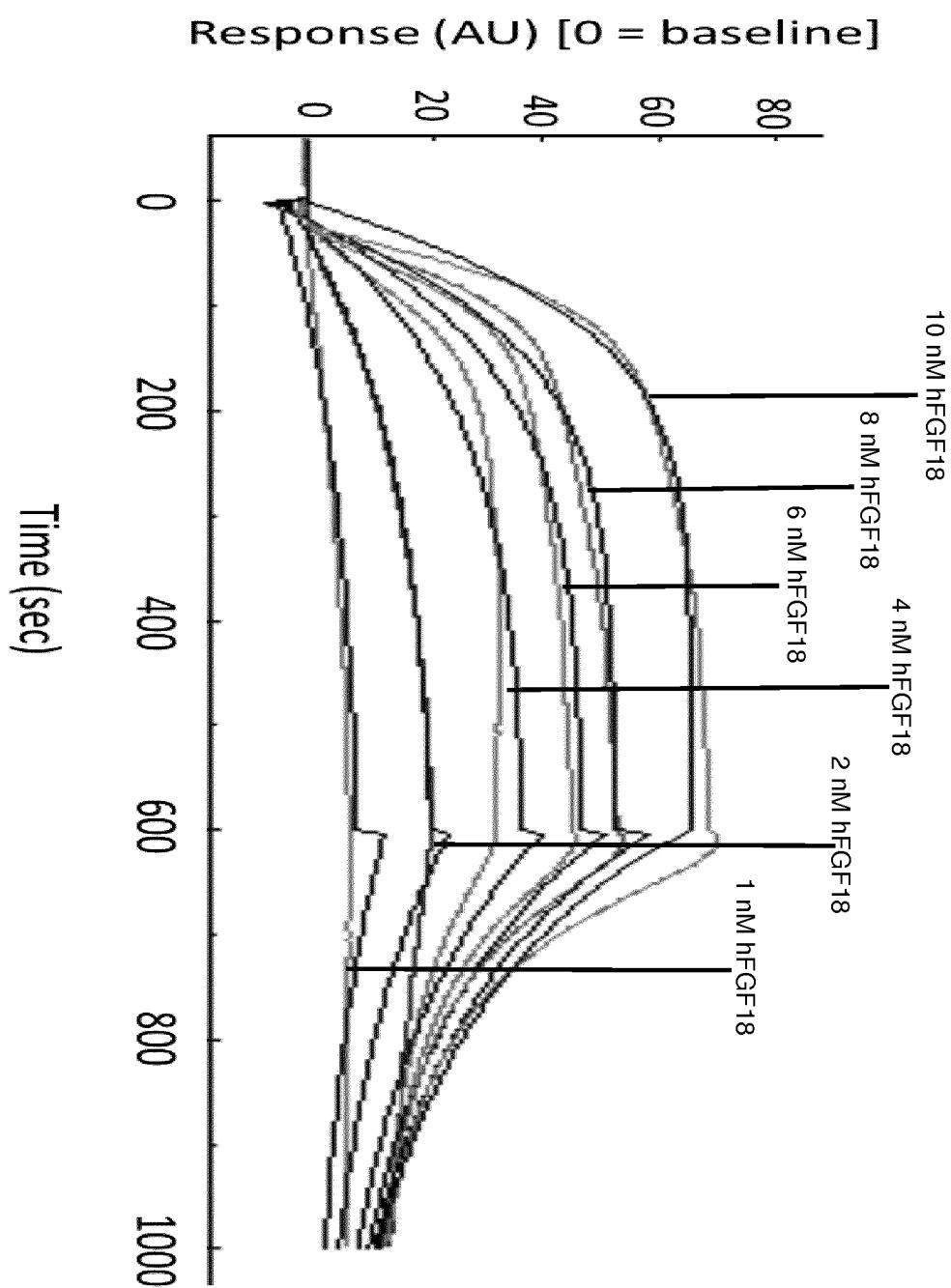


FIG. 19D

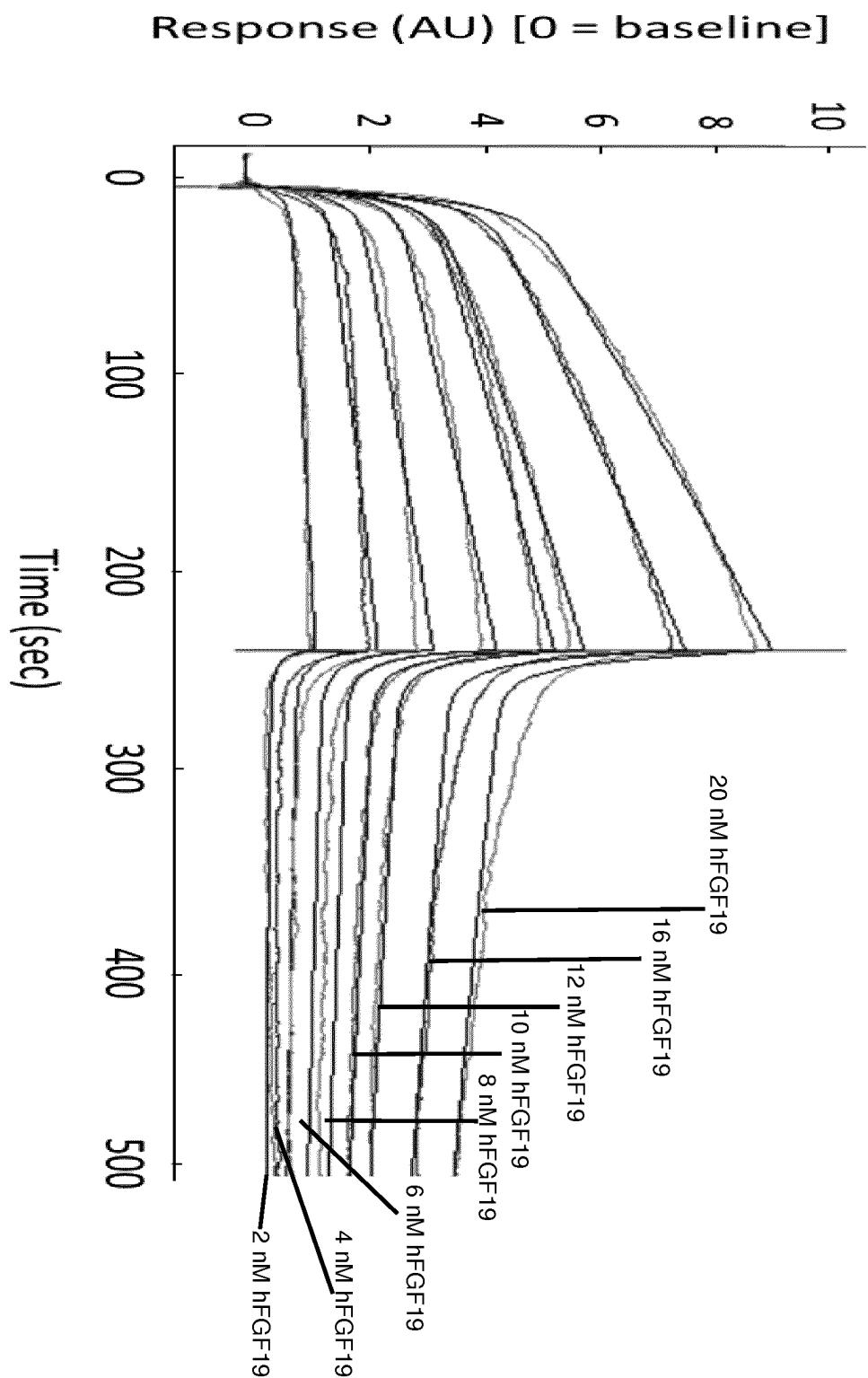


FIG. 19E

FIG. 19F

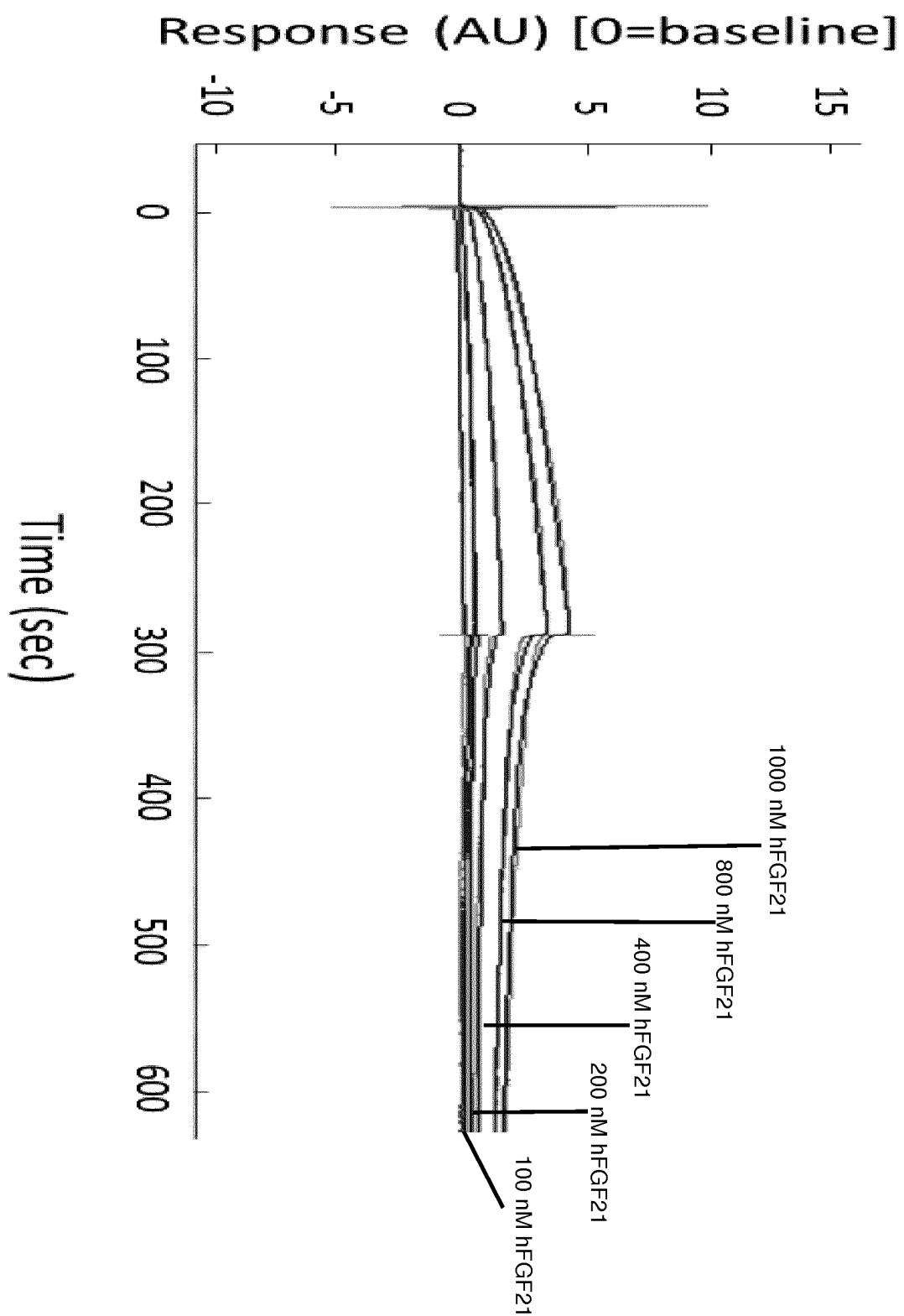


FIG. 20

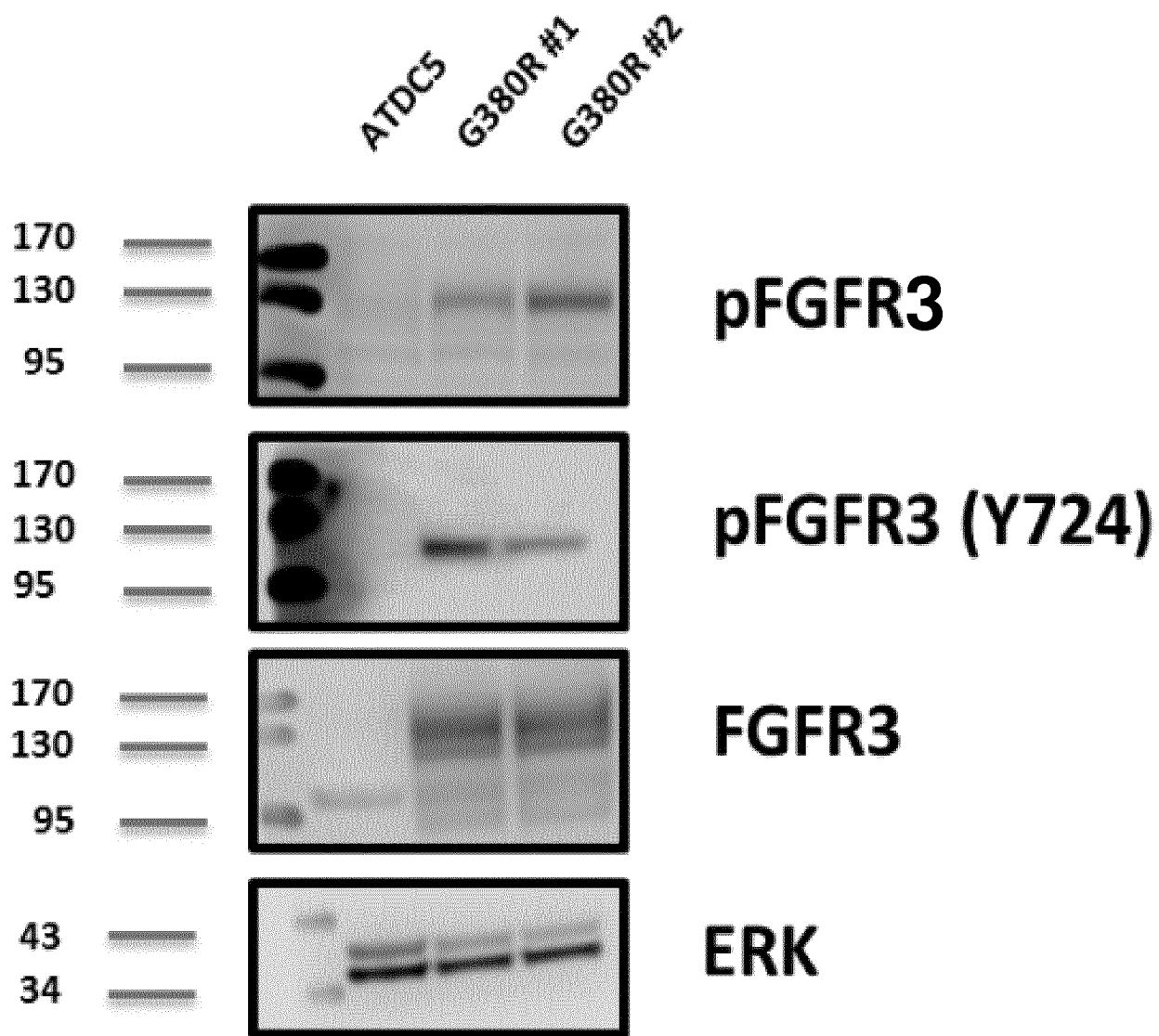
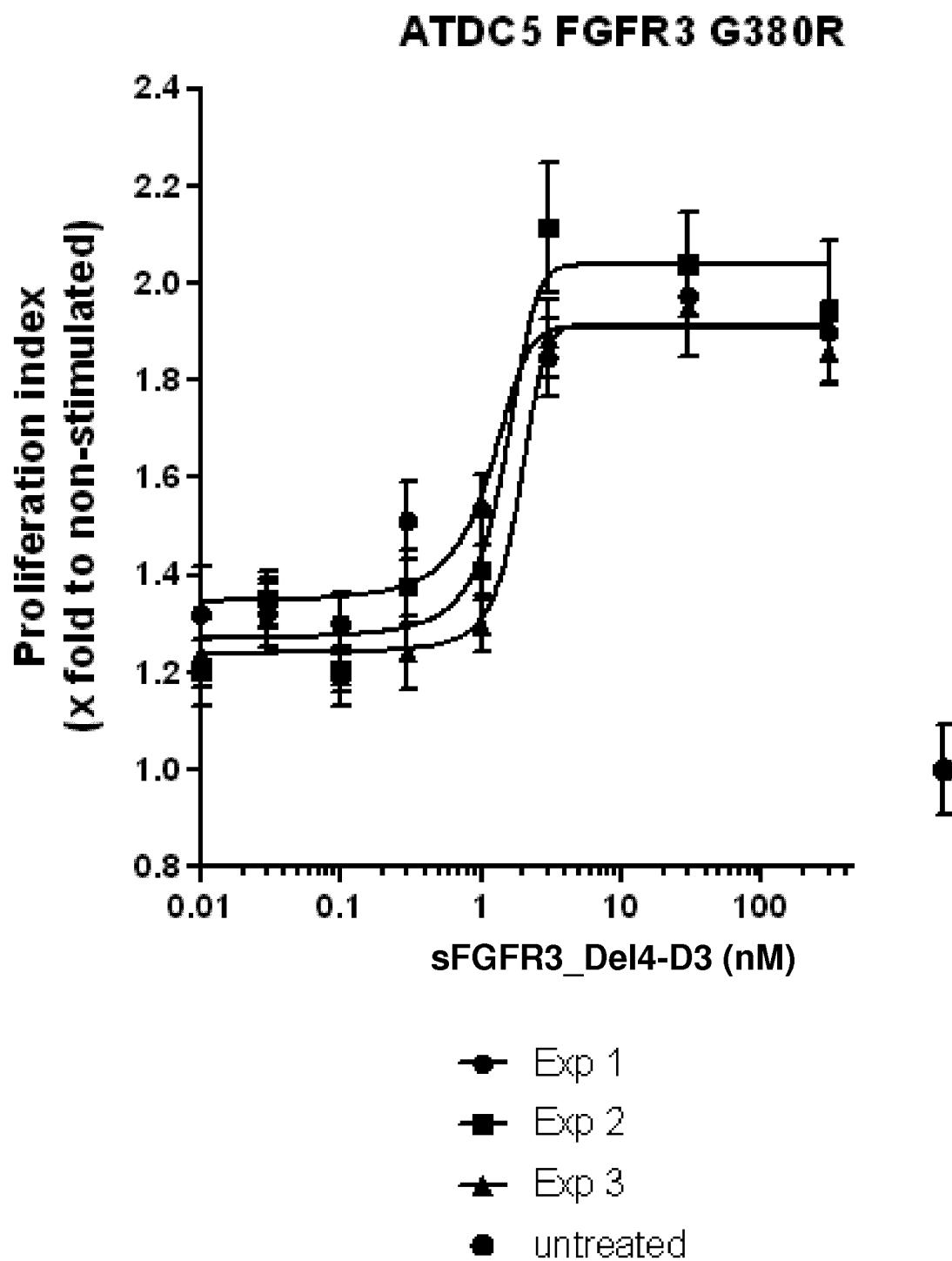


FIG. 21



eof-seql.txt  
SEQUENCE LISTING

<110> THERACHON  
INSTITUT NATIONAL DE LA SANTE ET DE LA RECHERCHE MEDI CALE  
UNIVERSITE NICE SOPHIA ANTI POLIS  
<120> SOLUBLE FIBROBLAST GROWTH FACTOR RECEPTOR 3 (SFGFR3) POLYPEPTIDES  
AND USES THEREOF  
<130> 901-10 PCT  
<150> 62/467,478  
<151> 2017-03-06  
<150> 62/359,607  
<151> 2016-07-07  
<160> 39  
<170> Patent in version 3.5  
<210> 1  
<211> 301  
<212> PRT  
<213> Artificial Sequence  
<220>  
<223> Synthetic construct  
<400> 1

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Ala Ala Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Ala Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Ala Ser His Gl u  
65 70 75 80

Asp Ser Gl y Ala Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser Gl y Asp Asp  
100 105 110

Gl u Asp Gl y Gl u Asp Gl u Ala Gl u Asp Thr Gl y Val Asp Thr Gl y Ala  
115 120 125

eol f-seql.txt

Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys Lys Leu Leu Ala Val  
130 135 140

Pro Ala Ala Asn Thr Val Arg Phe Arg Cys Pro Ala Ala Gly Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Gln Gln Trp Ser Leu Val  
180 185 190

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Glu Asn Lys Phe Gly Ser Ile Arg Gln Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Glu Arg Ser Pro His Arg Pro Ile Leu Gln Ala Gly Leu Pro Ala Asn  
225 230 235 240

Gln Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Ala Gln Pro His Ile Gln Trp Leu Lys His Val Glu Val Asn  
260 265 270

Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr  
290 295 300

<210> 2

<211> 301

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 2

Glu Ser Leu Gly Thr Glu Gln Arg Val Val Gly Arg Ala Ala Glu Val  
1 5 10 15

Pro Gly Pro Glu Pro Gly Gln Gln Glu Gln Leu Val Phe Gly Ser Gly  
20 25 30

Asp Ala Val Glu Leu Ser Cys Pro Pro Pro Gly Gly Gly Pro Met Gly  
Page 2

eol f-seq1 . txt

| 35   | 40                             | 45              |     |
|--|--------------------------------|-----------------|-----|
| Pro Thr Val Trp Val Lys Asp Glu              | Thr Glu Leu Val                | Pro Ser Glu Arg |     |
| 50 55  | 60                             |                 |     |
| Val Leu Val Gly Pro Glu Arg Leu Glu          | Val Leu Asn Ala Ser His Glu    |                 |     |
| 65 70  | 75                             | 80              |     |
| Asp Ser Gly Ala Tyr Ser Cys Arg Glu          | Arg Leu Thr Glu Arg Val        | Leu             |     |
| 85   | 90                             | 95              |     |
| Cys His Phe Ser Val Arg Val Thr Asp          | Ala Pro Ser Ser Glu            | Asp Asp         |     |
| 100  | 105                            | 110             |     |
| Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp    | Thr Gl y Val Asp Thr Gl y Al a |                 |     |
| 115  | 120                            | 125             |     |
| Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys  | Lys Leu Leu Al a Val           |                 |     |
| 130  | 135                            | 140             |     |
| Pro Al a Al a Asn Thr Val Arg Phe Arg Cys    | Pro Al a Al a Gl y Asn Pro     |                 |     |
| 145  | 150                            | 155             | 160 |
| Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y     | Arg Gl u Phe Arg Gl y Gl u     |                 |     |
| 165  | 170                            | 175             |     |
| His Arg Ile Gl y Gl y Ile Lys Leu Arg        | His Gl n Gl n Trp Ser Leu Val  |                 |     |
| 180  | 185                            | 190             |     |
| Met Glu Ser Val Val Pro Ser Asp Arg Gl y Asn | Tyr Thr Cys Val Val            |                 |     |
| 195  | 200                            | 205             |     |
| Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr   | Tyr Thr Leu Asp Val Leu        |                 |     |
| 210  | 215                            | 220             |     |
| Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n    | Al a Gl y Leu Pro Al a Asn     |                 |     |
| 225  | 230                            | 235             | 240 |
| Gl n Thr Al a Val Leu Gl y Ser Asp Val       | Gl u Phe His Ser Lys Val       | Tyr             |     |
| 245  | 250                            | 255             |     |
| Ser Asp Al a Gl n Pro His Ile Gl n Trp       | Leu Lys His Val Gl u Val       | Asn             |     |
| 260  | 265                            | 270             | 275 |
| Gl y Ser Lys Val Gl y Pro Asp Gl y           | Thr Pro Tyr Val Thr Val        | Leu Lys         |     |
| 275  | 280                            | 285             |     |

eol f-seql . txt

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr  
290 295 300

<210> 3

<211> 47

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 3

Thr Ala Gly Ala Asn Thr Thr Asp Lys Glu Leu Glu Val Leu Ser Leu  
1 5 10 15

Asn Val Thr Phe Glu Asp Ala Gly Glu Tyr Thr Cys Leu Ala Gly Asn  
20 25 30

Ser Ile Gly Phe Ser His His Ser Ala Trp Leu Val Val Leu Pro  
35 40 45

<210> 4

<211> 348

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 4

Glu Ser Leu Gly Thr Glu Gln Arg Val Val Gly Arg Ala Ala Glu Val  
1 5 10 15

Pro Gly Pro Glu Pro Gly Gln Gln Glu Gln Leu Val Phe Gly Ser Gly  
20 25 30

Asp Ala Val Glu Leu Ser Cys Pro Pro Pro Gly Gly Gly Pro Met Gly  
35 40 45

Pro Thr Val Trp Val Lys Asp Gly Thr Gly Leu Val Pro Ser Glu Arg  
50 55 60

Val Leu Val Gly Pro Gln Arg Leu Gln Val Leu Asn Ala Ser His Glu  
65 70 75 80

Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg Leu Thr Gln Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser Gly Asp Asp  
100 105 110

eol f-seql . txt

Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp Thr Gl y Val Asp Thr Gl y Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gl y Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y Arg Gl u Phe Arg Gl y Gl u  
165 170 175

His Arg Ile Gl y Gl y Ile Lys Leu Arg His Gl n Gl n Trp Ser Leu Val  
180 185 190

Met Gl u Ser Val Val Pro Ser Asp Arg Gl y Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n Al a Gl y Leu Pro Al a Asn  
225 230 235 240

Gl n Thr Al a Val Leu Gl y Ser Asp Val Gl u Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Al a Gl n Pro His Ile Gl n Trp Leu Lys His Val Gl u Val Asn  
260 265 270

Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Thr Al a Gl y Al a Asn Thr Thr Asp Lys Gl u Leu Gl u Val Leu Ser Leu  
290 295 300

Asn Val Thr Phe Gl u Asp Al a Gl y Gl u Tyr Thr Cys Leu Al a Gl y Asn  
305 310 315 320

Ser Ile Gl y Phe Ser His His Ser Al a Trp Leu Val Val Leu Pro Val  
325 330 335

Ser Leu Gl u Ser Asn Al a Ser Met Ser Ser Asn Thr  
340 345

<210> 5  
<211> 784

eol f-seql . txt

<212> PRT

<213> Homo sapiens

<400> 5

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Al a Al a Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Al a Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Al a Ser His Gl u  
65 70 75 80

Asp Ser Gl y Al a Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Al a Pro Ser Ser Gl y Asp Asp  
100 105 110

Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp Thr Gl y Val Asp Thr Gl y Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gl y Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y Arg Gl u Phe Arg Gl y Gl u  
165 170 175

His Arg Ile Gl y Gl y Ile Lys Leu Arg His Gl n Gl n Trp Ser Leu Val  
180 185 190

Met Gl u Ser Val Val Pro Ser Asp Arg Gl y Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n Al a Gl y Leu Pro Al a Asn  
225 230 235 240

eol f-seql . txt

Gl n Thr Al a Val Leu Gl y Ser Asp Val Gl u Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Al a Gl n Pro His Ile Gl n Trp Leu Lys His Val Gl u Val Asn  
260 265 270

Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Thr Al a Gl y Al a Asn Thr Thr Asp Lys Gl u Leu Gl u Val Leu Ser Leu  
290 295 300

His Asn Val Thr Phe Gl u Asp Al a Gl y Gl u Tyr Thr Cys Leu Al a Gl y  
305 310 315 320

Asn Ser Ile Gl y Phe Ser His His Ser Al a Trp Leu Val Val Leu Pro  
325 330 335

Al a Gl u Gl u Gl u Leu Val Gl u Al a Asp Gl u Al a Gl y Ser Val Tyr Al a  
340 345 350

Gl y Ile Leu Ser Tyr Gl y Val Gl y Phe Phe Leu Phe Ile Leu Val Val  
355 360 365

Al a Al a Val Thr Leu Cys Arg Leu Arg Ser Pro Pro Lys Lys Gl y Leu  
370 375 380

Gl y Ser Pro Thr Val His Lys Ile Ser Arg Phe Pro Leu Lys Arg Gl n  
385 390 395 400

Val Ser Leu Gl u Ser Asn Al a Ser Met Ser Ser Asn Thr Pro Leu Val  
405 410 415

Arg Ile Al a Arg Leu Ser Ser Gl y Gl u Gl y Pro Thr Leu Al a Asn Val  
420 425 430

Ser Gl u Leu Gl u Leu Pro Al a Asp Pro Lys Trp Gl u Leu Ser Arg Al a  
435 440 445

Arg Leu Thr Leu Gl y Lys Pro Leu Gl y Gl u Gl y Cys Phe Gl y Gl n Val  
450 455 460 465

Val Met Al a Gl u Al a Ile Gl y Ile Asp Lys Asp Arg Al a Al a Lys Pro  
465 470 475 480

Val Thr Val Al a Val Lys Met Leu Lys Asp Asp Al a Thr Asp Lys Asp  
Page 7

## eol f-seql . txt

485

490

495

Leu Ser Asp Leu Val Ser Glu Met Glu Met Met Lys Met Ile Gly Lys  
 500 505 510

His Lys Asn Ile Ile Asn Leu Leu Gly Ala Cys Thr Gln Gly Gly Pro  
 515 520 525

Leu Tyr Val Leu Val Glu Tyr Ala Ala Lys Gly Asn Leu Arg Glu Phe  
 530 535 540

Leu Arg Ala Arg Arg Pro Pro Gly Leu Asp Tyr Ser Phe Asp Thr Cys  
 545 550 555 560

Lys Pro Pro Glu Glu Gln Leu Thr Phe Lys Asp Leu Val Ser Cys Ala  
 565 570 575

Tyr Gln Val Ala Arg Gly Met Glu Tyr Leu Ala Ser Gln Lys Cys Ile  
 580 585 590

His Arg Asp Leu Ala Ala Arg Asn Val Leu Val Thr Glu Asp Asn Val  
 595 600 605

Met Lys Ile Ala Asp Phe Gly Leu Ala Arg Asp Val His Asn Leu Asp  
 610 615 620

Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu Pro Val Lys Trp Met Ala  
 625 630 635 640

Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr His Gln Ser Asp Val Trp  
 645 650 655

Ser Phe Gly Val Leu Leu Trp Glu Ile Phe Thr Leu Gly Gly Ser Pro  
 660 665 670

Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe Lys Leu Leu Lys Glu Gly  
 675 680 685

His Arg Met Asp Lys Pro Ala Asn Cys Thr His Asp Leu Tyr Met Ile  
 690 695 700

Met Arg Glu Cys Trp His Ala Ala Pro Ser Gln Arg Pro Thr Phe Lys  
 705 710 715 720

Gln Leu Val Glu Asp Leu Asp Arg Val Leu Thr Val Thr Ser Thr Asp  
 725 730 735

eol f-seql.txt

Gl u Tyr Leu Asp Leu Ser Al a Pro Phe Gl u Gl n Tyr Ser Pro Gl y Gl y  
740 745 750

Gl n Asp Thr Pro Ser Ser Ser Ser Gl y Asp Asp Ser Val Phe Al a  
755 760 765

Hi s Asp Leu Leu Pro Pro Al a Pro Pro Ser Ser Gl y Gl y Ser Arg Thr  
770 775 780

<210> 6

<211> 22

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 6

Met Gl y Al a Pro Al a Cys Al a Leu Al a Leu Cys Val Al a Val Al a Ile  
1 5 10 15

Val Al a Gl y Al a Ser Ser  
20

<210> 7

<211> 672

<212> PRT

<213> Homo sapiens

<400> 7

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Al a Al a Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Al a Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Al a Ser His Gl u  
65 70 75 80

Asp Ser Gl y Al a Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Al a Pro Ser Ser Gl y Asp Asp  
100 105 110

eol f-seql . txt

Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp Thr Gl y Val Asp Thr Gl y Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gl y Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y Arg Gl u Phe Arg Gl y Gl u  
165 170 175

Hi s Arg Ile Gl y Gl y Ile Lys Leu Arg Hi s Gl n Gl n Trp Ser Leu Val  
180 185 190

Met Gl u Ser Val Val Pro Ser Asp Arg Gl y Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Gl u Arg Ser Pro Hi s Arg Pro Ile Leu Gl n Al a Gl y Leu Pro Al a Asn  
225 230 235 240

Gl n Thr Al a Val Leu Gl y Ser Asp Val Gl u Phe Hi s Cys Lys Val Tyr  
245 250 255

Ser Asp Al a Gl n Pro Hi s Ile Gl n Trp Leu Lys Hi s Val Gl u Val Asn  
260 265 270

Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Val Ser Leu Gl u Ser Asn Al a Ser Met Ser Ser Asn Thr Pro Leu Val  
290 295 300

Arg Ile Al a Arg Leu Ser Ser Gl y Gl u Gl y Pro Thr Leu Al a Asn Val  
305 310 315 320

Ser Gl u Leu Gl u Leu Pro Al a Asp Pro Lys Trp Gl u Leu Ser Arg Al a  
325 330 335

Arg Leu Thr Leu Gl y Lys Pro Leu Gl y Gl u Gl y Cys Phe Gl y Gl n Val  
340 345 350

Val Met Al a Gl u Al a Ile Gl y Ile Asp Lys Asp Arg Al a Al a Lys Pro  
Page 10

eol f-seql . txt

355 360 365

Val Thr Val Ala Val Lys Met Leu Lys Asp Asp Ala Thr Asp Lys Asp  
370 375 380

Leu Ser Asp Leu Val Ser Glu Met Glu Met Met Lys Met Ile Gly Lys  
385 390 395 400

His Lys Asn Ile Ile Asn Leu Leu Gly Ala Cys Thr Gln Gly Gly Pro  
405 410 415

Leu Tyr Val Leu Val Glu Tyr Ala Ala Lys Gly Asn Leu Arg Glu Phe  
420 425 430

Leu Arg Ala Arg Arg Pro Pro Gly Leu Asp Tyr Ser Phe Asp Thr Cys  
435 440 445

Lys Pro Pro Glu Glu Gln Leu Thr Phe Lys Asp Leu Val Ser Cys Ala  
450 455 460

Tyr Gln Val Ala Arg Gly Met Glu Tyr Leu Ala Ser Gln Lys Cys Ile  
465 470 475 480

His Arg Asp Leu Ala Ala Arg Asn Val Leu Val Thr Glu Asp Asn Val  
485 490 495

Met Lys Ile Ala Asp Phe Gly Leu Ala Arg Asp Val His Asn Leu Asp  
500 505 510

Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu Pro Val Lys Trp Met Ala  
515 520 525

Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr His Gln Ser Asp Val Trp  
530 535 540

Ser Phe Gly Val Leu Leu Trp Glu Ile Phe Thr Leu Gly Gly Ser Pro  
545 550 555 560

Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe Lys Leu Leu Lys Glu Gly  
565 570 575

His Arg Met Asp Lys Pro Ala Asn Cys Thr His Asp Leu Tyr Met Ile  
580 585 590

Met Arg Glu Cys Trp His Ala Ala Pro Ser Gln Arg Pro Thr Phe Lys  
595 600 605

eol f-seql.txt

Gl n Leu Val Gl u Asp Leu Asp Arg Val Leu Thr Val Thr Ser Thr Asp  
610 615 620

Gl u Tyr Leu Asp Leu Ser Al a Pro Phe Gl u Gl n Tyr Ser Pro Gl y Gl y  
625 630 635 640

Gl n Asp Thr Pro Ser Ser Ser Gl y Asp Asp Ser Val Phe Al a  
645 650 655

Hi s Asp Leu Leu Pro Pro Al a Pro Pro Ser Ser Gl y Gl y Ser Arg Thr  
660 665 670

<210> 8  
<211> 672

<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 8

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Al a Al a Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Al a Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Al a Ser His Gl u  
65 70 75 80

Asp Ser Gl y Al a Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Al a Pro Ser Ser Gl y Asp Asp  
100 105 110

Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp Thr Gl y Val Asp Thr Gl y Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gl y Asn Pro

eol f-seql . txt

145

150

155

160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Glu Glu Trp Ser Leu Val  
180 185 190

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Glu Asn Lys Phe Gly Ser Ile Arg Glu Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Glu Arg Ser Pro His Arg Pro Ile Leu Glu Ala Gly Leu Pro Ala Asn  
225 230 235 240

Gln Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Ser Lys Val Tyr  
245 250 255

Ser Asp Ala Gln Pro His Ile Gln Trp Leu Lys His Val Glu Val Asn  
260 265 270

Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr Pro Leu Val  
290 295 300

Arg Ile Ala Arg Leu Ser Ser Gly Glu Gly Pro Thr Leu Ala Asn Val  
305 310 315 320

Ser Glu Leu Glu Leu Pro Ala Asp Pro Lys Trp Glu Leu Ser Arg Ala  
325 330 335

Arg Leu Thr Leu Gly Lys Pro Leu Gly Glu Gly Cys Phe Gly Gln Val  
340 345 350

Val Met Ala Glu Ala Ile Gly Ile Asp Lys Asp Arg Ala Ala Lys Pro  
355 360 365

Val Thr Val Ala Val Lys Met Leu Lys Asp Asp Ala Thr Asp Lys Asp  
370 375 380

Leu Ser Asp Leu Val Ser Glu Met Glu Met Met Lys Met Ile Gly Lys  
385 390 395 400

eol f-seql.txt

His Lys Asn Ile Ile Asn Leu Leu Gly Ala Cys Thr Gln Gly Gly Pro  
405 410 415

Leu Tyr Val Leu Val Glu Tyr Ala Ala Lys Gly Asn Leu Arg Glu Phe  
420 425 430

Leu Arg Ala Arg Arg Pro Pro Gly Leu Asp Tyr Ser Phe Asp Thr Cys  
435 440 445

Lys Pro Pro Glu Glu Gln Leu Thr Phe Lys Asp Leu Val Ser Cys Ala  
450 455 460

Tyr Gln Val Ala Arg Gly Met Glu Tyr Leu Ala Ser Gln Lys Cys Ile  
465 470 475 480

His Arg Asp Leu Ala Ala Arg Asn Val Leu Val Thr Glu Asp Asn Val  
485 490 495

Met Lys Ile Ala Asp Phe Gly Leu Ala Arg Asp Val His Asn Leu Asp  
500 505 510

Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu Pro Val Lys Trp Met Ala  
515 520 525

Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr His Gln Ser Asp Val Trp  
530 535 540

Ser Phe Gly Val Leu Leu Trp Glu Ile Phe Thr Leu Gly Gly Ser Pro  
545 550 555 560

Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe Lys Leu Leu Lys Glu Gly  
565 570 575

His Arg Met Asp Lys Pro Ala Asn Cys Thr His Asp Leu Tyr Met Ile  
580 585 590

Met Arg Glu Cys Trp His Ala Ala Pro Ser Gln Arg Pro Thr Phe Lys  
595 600 605

Gln Leu Val Glu Asp Leu Asp Arg Val Leu Thr Val Thr Ser Thr Asp  
610 615 620

Glu Tyr Leu Asp Leu Ser Ala Pro Phe Glu Gln Tyr Ser Pro Glu Gly  
625 630 635 640

Gln Asp Thr Pro Ser Ser Ser Ser Gly Asp Asp Ser Val Phe Ala  
645 650 655

eol f-seql . txt

Hi s Asp Leu Leu Pro Pro Al a Pro Pro Ser Ser Gly Gly Ser Arg Thr  
660 665 670

<210> 9  
<211> 720

<212> PRT

<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 9

Glu Ser Leu Gly Thr Glu Gln Arg Val Val Gly Arg Ala Ala Glu Val  
1 5 10 15

Pro Gly Pro Glu Pro Gly Gln Gln Glu Gln Leu Val Phe Gly Ser Gly  
20 25 30

Asp Ala Val Glu Leu Ser Cys Pro Pro Pro Gly Gly Gly Pro Met Gly  
35 40 45

Pro Thr Val Trp Val Lys Asp Gly Thr Gly Leu Val Pro Ser Glu Arg  
50 55 60

Val Leu Val Gly Pro Gln Arg Leu Gln Val Leu Asn Ala Ser His Glu  
65 70 75 80

Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg Leu Thr Gln Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Al a Pro Ser Ser Gly Asp Asp  
100 105 110

Glu Asp Gly Glu Asp Glu Al a Glu Asp Thr Gly Val Asp Thr Gly Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gly Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Gln Gln Trp Ser Leu Val  
180 185 190

eol f-seql.txt

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n Ala Gl y Leu Pro Ala Asn  
225 230 235 240

Gl n Thr Ala Val Leu Gl y Ser Asp Val Gl u Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Ala Gl n Pro His Ile Gl n Trp Leu Lys His Val Gl u Val Asn  
260 265 270

Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Thr Ala Gl y Ala Asn Thr Thr Asp Lys Gl u Leu Gl u Val Leu Ser Leu  
290 295 300

His Asn Val Thr Phe Gl u Asp Ala Gl y Gl u Tyr Thr Cys Leu Ala Gl y  
305 310 315 320

Asn Ser Ile Gl y Phe Ser His His Ser Ala Trp Leu Val Val Leu Pro  
325 330 335

Val Ser Leu Gl u Ser Asn Ala Ser Met Ser Ser Asn Thr Pro Leu Val  
340 345 350

Arg Ile Ala Arg Leu Ser Ser Gl y Gl u Gl y Pro Thr Leu Ala Asn Val  
355 360 365

Ser Gl u Leu Gl u Leu Pro Ala Asp Pro Lys Trp Gl u Leu Ser Arg Ala  
370 375 380

Arg Leu Thr Leu Gl y Lys Pro Leu Gl y Gl u Gl y Cys Phe Gl y Gl n Val  
385 390 395 400

Val Met Ala Gl u Ala Ile Gl y Ile Asp Lys Asp Arg Ala Ala Lys Pro  
405 410 415

Val Thr Val Ala Val Lys Met Leu Lys Asp Asp Ala Thr Asp Lys Asp  
420 425 430

Leu Ser Asp Leu Val Ser Gl u Met Gl u Met Met Lys Met Ile Gl y Lys  
435 440 445

eol f-seql . txt

His Lys Asn Ile Ile Asn Leu Leu Gly Ala Cys Thr Glu Gly Gly Pro  
450 455 460

Leu Tyr Val Leu Val Glu Tyr Ala Ala Lys Gly Asn Leu Arg Glu Phe  
465 470 475 480

Leu Arg Ala Arg Arg Pro Pro Gly Leu Asp Tyr Ser Phe Asp Thr Cys  
485 490 495

Lys Pro Pro Glu Glu Gln Leu Thr Phe Lys Asp Leu Val Ser Cys Ala  
500 505 510

Tyr Gln Val Ala Arg Gly Met Glu Tyr Leu Ala Ser Gln Lys Cys Ile  
515 520 525

His Arg Asp Leu Ala Ala Arg Asn Val Leu Val Thr Glu Asp Asn Val  
530 535 540

Met Lys Ile Ala Asp Phe Gly Leu Ala Arg Asp Val His Asn Leu Asp  
545 550 555 560

Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu Pro Val Lys Trp Met Ala  
565 570 575

Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr His Gln Ser Asp Val Trp  
580 585 590

Ser Phe Gly Val Leu Leu Trp Glu Ile Phe Thr Leu Gly Gly Ser Pro  
595 600 605

Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe Lys Leu Leu Lys Glu Gly  
610 615 620

His Arg Met Asp Lys Pro Ala Asn Cys Thr His Asp Leu Tyr Met Ile  
625 630 635 640

Met Arg Glu Cys Trp His Ala Ala Pro Ser Gln Arg Pro Thr Phe Lys  
645 650 655

Gln Leu Val Glu Asp Leu Asp Arg Val Leu Thr Val Thr Ser Thr Asp  
660 665 670

Glu Tyr Leu Asp Leu Ser Ala Pro Phe Glu Gln Tyr Ser Pro Gly Gly  
675 680 685

Gln Asp Thr Pro Ser Ser Ser Ser Ser Gly Asp Asp Ser Val Phe Ala  
690 695 700

eol f-seql . txt

His Asp Leu Leu Pro Pro Ala Pro Pro Ser Ser Gly Gly Ser Arg Thr  
705 710 715 720

<210> 10  
<211> 512  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 10

Glu Ser Leu Gly Thr Glu Gln Arg Val Val Gly Arg Ala Ala Glu Val  
1 5 10 15

Pro Gly Pro Glu Pro Gly Gln Gln Glu Gln Leu Val Phe Gly Ser Gly  
20 25 30

Asp Ala Val Glu Leu Ser Cys Pro Pro Pro Gly Gly Gly Pro Met Gly  
35 40 45

Pro Thr Val Trp Val Lys Asp Gly Thr Gly Leu Val Pro Ser Glu Arg  
50 55 60

Val Leu Val Gly Pro Gln Arg Leu Gln Val Leu Asn Ala Ser His Glu  
65 70 75 80

Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg Leu Thr Gln Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser Gly Asp Asp  
100 105 110

Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr Gly Val Asp Thr Gly Ala  
115 120 125

Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys Lys Leu Leu Ala Val  
130 135 140

Pro Ala Ala Asn Thr Val Arg Phe Arg Cys Pro Ala Ala Gly Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Gln Gln Trp Ser Leu Val  
180 185 190

eol f-seql . txt

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Glu Asn Lys Phe Gly Ser Ile Arg Gln Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Glu Arg Ser Pro His Arg Pro Ile Leu Gln Ala Gly Leu Pro Ala Asn  
225 230 235 240

Gln Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Ala Gln Pro His Ile Gln Trp Leu Lys His Val Glu Val Asn  
260 265 270

Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr Ser Gly Ser  
290 295 300

Gly Ser Gly Ser Gly Ser Gly Ser Gly Ser Val Val Phe Pro  
305 310 315 320

Tyr Phe Pro Arg Leu Gly Arg Tyr Asn Leu Asn Phe His Glu Ala Gln  
325 330 335

Gln Ala Cys Leu Asp Gln Asp Ala Val Ile Ala Ser Phe Asp Gln Leu  
340 345 350

Tyr Asp Ala Trp Arg Gly Gly Leu Asp Trp Cys Asn Ala Gly Trp Leu  
355 360 365

Ser Asp Gly Ser Val Gln Tyr Pro Ile Thr Lys Pro Arg Glu Pro Cys  
370 375 380

Gly Gly Gln Asn Thr Val Pro Gly Val Arg Asn Tyr Gly Phe Trp Asp  
385 390 395 400

Lys Asp Lys Ser Arg Tyr Asp Val Phe Cys Phe Thr Ser Asn Phe Asn  
405 410 415

Gly Arg Phe Tyr Tyr Leu Ile His Pro Thr Lys Leu Thr Tyr Asp Glu  
420 425 430

Ala Val Gln Ala Cys Leu Asn Asp Gly Ala Gln Ile Ala Lys Val Gly  
435 440 445

eol f-seql . txt

Gln Ile Phe Ala Ala Trp Lys Ile Leu Gly Tyr Asp Arg Cys Asp Ala  
450 455 460

Gly Trp Leu Ala Asp Gly Ser Val Arg Tyr Pro Ile Ser Arg Pro Arg  
465 470 475 480

Arg Arg Cys Ser Pro Thr Glu Ala Ala Val Arg Phe Val Gly Phe Pro  
485 490 495

Asp Lys Lys His Lys Leu Tyr Gly Val Tyr Cys Phe Arg Ala Tyr Asn  
500 505 510

<210> 11

<211> 512

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 11

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Ala Ala Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Ala Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Ala Ser His Gl u  
65 70 75 80

Asp Ser Gl y Ala Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser Gl y Asp Asp  
100 105 110

Gl u Asp Gl y Gl u Asp Gl u Ala Gl u Asp Thr Gl y Val Asp Thr Gl y Ala  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Ala Val  
130 135 140

eol f-seql . txt

Pro Ala Ala Asn Thr Val Arg Phe Arg Cys Pro Ala Ala Gly Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Gln Gln Trp Ser Leu Val  
180 185 190

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gly Ser Ile Arg Gln Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Gl u Arg Ser Pro His Arg Pro Ile Leu Gln Ala Gly Leu Pro Ala Asn  
225 230 235 240

Gln Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Ser Lys Val Tyr  
245 250 255

Ser Asp Ala Gln Pro His Ile Gln Trp Leu Lys His Val Glu Val Asn  
260 265 270

Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr Ser Gly Ser  
290 295 300

Gly Ser Gly Ser Gly Ser Gly Ser Gly Ser Val Val Phe Pro  
305 310 315 320

Tyr Phe Pro Arg Leu Gly Arg Tyr Asn Leu Asn Phe His Glu Ala Gln  
325 330 335

Gln Ala Cys Leu Asp Gln Asp Ala Val Ile Ala Ser Phe Asp Gln Leu  
340 345 350

Tyr Asp Ala Trp Arg Gly Gly Leu Asp Trp Cys Asn Ala Gly Trp Leu  
355 360 365

Ser Asp Gly Ser Val Gln Tyr Pro Ile Thr Lys Pro Arg Glu Pro Cys  
370 375 380

Gly Gly Gln Asn Thr Val Pro Gly Val Arg Asn Tyr Gly Phe Trp Asp  
385 390 395 400

eol f-seql . txt

Lys Asp Lys Ser Arg Tyr Asp Val Phe Cys Phe Thr Ser Asn Phe Asn  
405 410 415

Gly Arg Phe Tyr Tyr Leu Ile His Pro Thr Lys Leu Thr Tyr Asp Glu  
420 425 430

Ala Val Gln Ala Cys Leu Asn Asp Gly Ala Gln Ile Ala Lys Val Gly  
435 440 445

Gln Ile Phe Ala Ala Trp Lys Ile Leu Gly Tyr Asp Arg Cys Asp Ala  
450 455 460

Gly Trp Leu Ala Asp Gly Ser Val Arg Tyr Pro Ile Ser Arg Pro Arg  
465 470 475 480

Arg Arg Cys Ser Pro Thr Glu Ala Ala Val Arg Phe Val Gly Phe Pro  
485 490 495

Asp Lys Lys His Lys Leu Tyr Gly Val Tyr Cys Phe Arg Ala Tyr Asn  
500 505 510

<210> 12

<211> 560

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 12

Glu Ser Leu Gly Thr Glu Gln Arg Val Val Gly Arg Ala Ala Glu Val  
1 5 10 15

Pro Gly Pro Glu Pro Gly Gln Gln Glu Gln Leu Val Phe Gly Ser Gly  
20 25 30

Asp Ala Val Glu Leu Ser Cys Pro Pro Pro Gly Gly Pro Met Gly  
35 40 45

Pro Thr Val Trp Val Lys Asp Gly Thr Gly Leu Val Pro Ser Glu Arg  
50 55 60

Val Leu Val Gly Pro Gln Arg Leu Gln Val Leu Asn Ala Ser His Glu  
65 70 75 80

Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg Leu Thr Gln Arg Val Leu  
85 90 95

eol f-seql . txt

Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser Gly Asp Asp  
100 105 110

Glu Asp Glu Glu Asp Glu Ala Glu Asp Thr Gly Val Asp Thr Gly Ala  
115 120 125

Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys Lys Leu Leu Ala Val  
130 135 140

Pro Ala Ala Asn Thr Val Arg Phe Arg Cys Pro Ala Ala Gly Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe Arg Gly Glu  
165 170 175

His Arg Ile Gly Gly Ile Lys Leu Arg His Gln Gln Trp Ser Leu Val  
180 185 190

Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr Cys Val Val  
195 200 205

Glu Asn Lys Phe Gly Ser Ile Arg Gln Thr Tyr Thr Leu Asp Val Leu  
210 215 220

Glu Arg Ser Pro His Arg Pro Ile Leu Gln Ala Gly Leu Pro Ala Asn  
225 230 235 240

Gln Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Ala Gln Pro His Ile Gln Trp Leu Lys His Val Glu Val Asn  
260 265 270

Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Thr Ala Gly Ala Asn Thr Thr Asp Lys Glu Leu Glu Val Leu Ser Leu  
290 295 300

His Asn Val Thr Phe Glu Asp Ala Gly Glu Tyr Thr Cys Leu Ala Gly  
305 310 315 320

Asn Ser Ile Gly Phe Ser His His Ser Ala Trp Leu Val Val Leu Pro  
325 330 335

Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr Ser Gly Ser  
340 345 350

eol f-seql . txt

Gly Ser Gly Ser Gly Ser Gly Ser Gly Ser Val Val Phe Pro  
355 360 365

Tyr Phe Pro Arg Leu Gly Arg Tyr Asn Leu Asn Phe His Glu Ala Glu  
370 375 380

Gln Ala Cys Leu Asp Gln Asp Ala Val Ile Ala Ser Phe Asp Gln Leu  
385 390 395 400

Tyr Asp Ala Trp Arg Gly Gly Leu Asp Trp Cys Asn Ala Gly Trp Leu  
405 410 415

Ser Asp Gly Ser Val Gln Tyr Pro Ile Thr Lys Pro Arg Glu Pro Cys  
420 425 430

Gly Gly Gln Asn Thr Val Pro Gly Val Arg Asn Tyr Gly Phe Trp Asp  
435 440 445

Lys Asp Lys Ser Arg Tyr Asp Val Phe Cys Phe Thr Ser Asn Phe Asn  
450 455 460

Gly Arg Phe Tyr Tyr Leu Ile His Pro Thr Lys Leu Thr Tyr Asp Glu  
465 470 475 480

Ala Val Gln Ala Cys Leu Asn Asp Gly Ala Gln Ile Ala Lys Val Gly  
485 490 495

Gln Ile Phe Ala Ala Trp Lys Ile Leu Gly Tyr Asp Arg Cys Asp Ala  
500 505 510

Gly Trp Leu Ala Asp Gly Ser Val Arg Tyr Pro Ile Ser Arg Pro Arg  
515 520 525

Arg Arg Cys Ser Pro Thr Glu Ala Ala Val Arg Phe Val Gly Phe Pro  
530 535 540

Asp Lys Lys His Lys Leu Tyr Gly Val Tyr Cys Phe Arg Ala Tyr Asn  
545 550 555 560

<210> 13  
<211> 155

<212> PRT

<213> Homo sapiens

<400> 13

Met Ala Glu Gly Glu Ile Thr Thr Phe Thr Ala Leu Thr Glu Lys Phe  
1 5 10 15

eol f-seql . txt

Asn Leu Pro Pro Gly Asn Tyr Lys Lys Pro Lys Leu Leu Tyr Cys Ser  
20 25 30

Asn Gly Gly His Phe Leu Arg Ile Leu Pro Asp Gly Thr Val Asp Gly  
35 40 45

Thr Arg Asp Arg Ser Asp Gln His Ile Gln Leu Gln Leu Ser Ala Glu  
50 55 60

Ser Val Gly Glu Val Tyr Ile Lys Ser Thr Glu Thr Gly Gln Tyr Leu  
65 70 75 80

Ala Met Asp Thr Asp Gly Leu Leu Tyr Gly Ser Gln Thr Pro Asn Glu  
85 90 95

Gl u Cys Leu Phe Leu Gl u Arg Leu Gl u Gl u Asn His Tyr Asn Thr Tyr  
100 105 110

Ile Ser Lys Lys His Ala Glu Lys Asn Trp Phe Val Gly Leu Lys Lys  
115 120 125

Asn Gly Ser Cys Lys Arg Gly Pro Arg Thr His Tyr Gly Gln Lys Ala  
130 135 140

Ile Leu Phe Leu Pro Leu Pro Val Ser Ser Asp  
145 150 155

<210> 14

<211> 288

<212> PRT

<213> Homo sapiens

<400> 14

Met Val Gly Val Gly Gly Asp Val Gl u Asp Val Thr Pro Arg Pro  
1 5 10 15

Gly Gly Cys Gln Ile Ser Gly Arg Gly Ala Arg Gly Cys Asn Gly Ile  
20 25 30

Pro Gly Ala Ala Ala Trp Gl u Ala Ala Leu Pro Arg Arg Arg Pro Arg  
35 40 45

Arg His Pro Ser Val Asn Pro Arg Ser Arg Ala Ala Gly Ser Pro Arg  
50 55 60

Thr Arg Gly Arg Arg Thr Gl u Gl u Arg Pro Ser Gly Ser Arg Leu Gly  
65 70 75 80

eol f-seql . txt

Asp Arg Gly Arg Gly Arg Ala Leu Pro Gly Gly Arg Leu Gly Gly Arg  
85 90 95

Gly Arg Gly Arg Ala Pro Glu Arg Val Gly Gly Arg Gly Arg Gly Arg  
100 105 110

Gly Thr Ala Ala Pro Arg Ala Ala Pro Ala Ala Arg Gly Ser Arg Pro  
115 120 125

Gly Pro Ala Gly Thr Met Ala Ala Gly Ser Ile Thr Thr Leu Pro Ala  
130 135 140

Leu Pro Glu Asp Gly Gly Ser Gly Ala Phe Pro Pro Gly His Phe Lys  
145 150 155 160

Asp Pro Lys Arg Leu Tyr Cys Lys Asn Gly Gly Phe Phe Leu Arg Ile  
165 170 175

His Pro Asp Gly Arg Val Asp Gly Val Arg Glu Lys Ser Asp Pro His  
180 185 190

Ile Lys Leu Gln Leu Gln Ala Glu Glu Arg Gly Val Val Ser Ile Lys  
195 200 205

Gly Val Cys Ala Asn Arg Tyr Leu Ala Met Lys Glu Asp Gly Arg Leu  
210 215 220

Leu Ala Ser Lys Cys Val Thr Asp Glu Cys Phe Phe Glu Arg Leu  
225 230 235 240

Glu Ser Asn Asn Tyr Asn Thr Tyr Arg Ser Arg Lys Tyr Thr Ser Trp  
245 250 255

Tyr Val Ala Leu Lys Arg Thr Gly Gln Tyr Lys Leu Gly Ser Lys Thr  
260 265 270

Gly Pro Gly Gln Lys Ala Ile Leu Phe Leu Pro Met Ser Ala Lys Ser  
275 280 285

<210> 15  
<211> 208  
<212> PRT  
<213> Homo sapiens

<400> 15

Met Ala Pro Leu Gly Glu Val Gly Asn Tyr Phe Gly Val Gln Asp Ala  
1 5 10 15

eol f-seql . txt

Val Pro Phe Gl y Asn Val Pro Val 20 Leu Pro Val Asp Ser Pro Val Leu 25 30

Leu Ser Asp His Leu Gl y Gl n Ser Gl u Al a Gl y Gl y Leu Pro Arg Gl y 35 40 45

Pro Al a Val Thr Asp Leu Asp His Leu Lys Gl y Ile Leu Arg Arg Arg 50 55 60

Gl n Leu Tyr Cys Arg Thr Gl y Phe His Leu Gl u Ile Phe Pro Asn Gl y 65 70 75 80

Thr Ile Gl n Gl y Thr Arg Lys Asp His Ser Arg Phe Gl y Ile Leu Gl u 85 90 95

Phe Ile Ser Ile Al a Val Gl y Leu Val 100 Ser Ile Arg Gl y Val 105 Asp Ser 110

Gl y Leu Tyr Leu Gl y Met Asn Gl u Lys Gl y Gl u Leu Tyr Gl y Ser Gl u 115 120 125

Lys Leu Thr Gl n Gl u Cys Val Phe Arg Gl u Gl n Phe Gl u Gl u Asn Trp 130 135 140

Tyr Asn Thr Tyr Ser Ser Asn Leu Tyr Lys His Val Asp Thr Gl y Arg 145 150 155 160

Arg Tyr Tyr Val Al a Leu Asn Lys Asp Gl y Thr Pro Arg Gl u Gl y Thr 165 170 175

Arg Thr Lys Arg His Gl n Lys Phe Thr His Phe Leu Pro Arg Pro Val 180 185 190

Asp Pro Asp Lys Val Pro Gl u Leu Tyr Lys Asp Ile Leu Ser Gl n Ser 195 200 205

<210> 16

<211> 207

<212> PRT

<213> Homo sapiens

<400> 16

Met Tyr Ser Al a Pro Ser Al a Cys Thr Cys Leu Cys Leu His Phe Leu 1 5 10 15

Leu Leu Cys Phe Gl n Val Gl n Val 20 Leu Val Al a Gl u Gl u Asn Val Asp 25 30

eol f-seql . txt

Phe Arg Ile His Val Glu Asn Gln Thr Arg Ala Arg Asp Asp Val Ser  
35 40 45

Arg Lys Gln Leu Arg Leu Tyr Gln Leu Tyr Ser Arg Thr Ser Gly Lys  
50 55 60

His Ile Gln Val Leu Gly Arg Arg Ile Ser Ala Arg Gly Glu Asp Gly  
65 70 75 80

Asp Lys Tyr Ala Gln Leu Leu Val Glu Thr Asp Thr Phe Gly Ser Gln  
85 90 95

Val Arg Ile Lys Gly Lys Glu Thr Glu Phe Tyr Leu Cys Met Asn Arg  
100 105 110

Lys Gly Lys Leu Val Gly Lys Pro Asp Gly Thr Ser Lys Glu Cys Val  
115 120 125

Phe Ile Glu Lys Val Leu Glu Asn Asn Tyr Thr Ala Leu Met Ser Ala  
130 135 140

Lys Tyr Ser Gly Trp Tyr Val Gly Phe Thr Lys Lys Gly Arg Pro Arg  
145 150 155 160

Lys Gly Pro Lys Thr Arg Glu Asn Gln Gln Asp Val His Phe Met Lys  
165 170 175

Arg Tyr Pro Lys Gly Gln Pro Glu Leu Gln Lys Pro Phe Lys Tyr Thr  
180 185 190

Thr Val Thr Lys Arg Ser Arg Arg Ile Arg Pro Thr His Pro Ala  
195 200 205

<210> 17

<211> 323

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 17

Met Gly Ala Pro Ala Cys Ala Leu Ala Leu Cys Val Ala Val Ala Ile  
1 5 10 15

Val Ala Gly Ala Ser Ser Glu Ser Leu Gly Thr Glu Gln Arg Val Val  
20 25 30

eol f-seql . txt

Gly Arg Ala Ala Glu Val Pro Gly Pro Glu Pro Gly Glu Glu Glu  
35 40 45

Leu Val Phe Gly Ser Gly Asp Ala Val Glu Leu Ser Cys Pro Pro Pro  
50 55 60

Gly Gly Gly Pro Met Gly Pro Thr Val Trp Val Lys Asp Gly Thr Gly  
65 70 75 80

Leu Val Pro Ser Glu Arg Val Leu Val Gly Pro Glu Arg Leu Glu Val  
85 90 95

Leu Asn Ala Ser His Glu Asp Ser Gly Ala Tyr Ser Cys Arg Glu Arg  
100 105 110

Leu Thr Glu Arg Val Leu Cys His Phe Ser Val Arg Val Thr Asp Ala  
115 120 125

Pro Ser Ser Gly Asp Asp Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr  
130 135 140

Gly Val Asp Thr Gly Ala Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp  
145 150 155 160

Lys Lys Leu Leu Ala Val Pro Ala Ala Asn Thr Val Arg Phe Arg Cys  
165 170 175

Pro Ala Ala Gly Asn Pro Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly  
180 185 190

Arg Glu Phe Arg Gly Glu His Arg Ile Gly Gly Ile Lys Leu Arg His  
195 200 205

Gln Gln Trp Ser Leu Val Met Glu Ser Val Val Pro Ser Asp Arg Gly  
210 215 220

Asn Tyr Thr Cys Val Val Glu Asn Lys Phe Gly Ser Ile Arg Glu Thr  
225 230 235 240

Tyr Thr Leu Asp Val Leu Glu Arg Ser Pro His Arg Pro Ile Leu Glu  
245 250 255

Ala Gly Leu Pro Ala Asn Glu Thr Ala Val Leu Gly Ser Asp Val Glu  
260 265 270

Phe His Cys Lys Val Tyr Ser Asp Ala Glu Pro His Ile Glu Trp Leu  
275 280 285

eol f-seql . txt

Lys His Val Glu Val Asn Gly Ser Lys Val Gly Pro Asp Gly Thr Pro  
290 295 300

Tyr Val Thr Val Leu Lys Val Ser Leu Glu Ser Asn Ala Ser Met Ser  
305 310 315 320

Ser Asn Thr

<210> 18

<211> 323

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 18

Met Gly Ala Pro Ala Cys Ala Leu Ala Leu Cys Val Ala Val Ala Ile  
1 5 10 15

Val Ala Gly Ala Ser Ser Glu Ser Leu Gly Thr Glu Glu Arg Val Val  
20 25 30

Gly Arg Ala Ala Glu Val Pro Gly Pro Glu Pro Gly Glu Glu Glu Glu  
35 40 45

Leu Val Phe Gly Ser Gly Asp Ala Val Glu Leu Ser Cys Pro Pro Pro  
50 55 60

Gly Gly Gly Pro Met Gly Pro Thr Val Trp Val Lys Asp Gly Thr Gly  
65 70 75 80

Leu Val Pro Ser Glu Arg Val Leu Val Gly Pro Glu Arg Leu Glu Val  
85 90 95

Leu Asn Ala Ser His Glu Asp Ser Gly Ala Tyr Ser Cys Arg Glu Arg  
100 105 110

Leu Thr Glu Arg Val Leu Cys His Phe Ser Val Arg Val Thr Asp Ala  
115 120 125

Pro Ser Ser Gly Asp Asp Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr  
130 135 140

Gly Val Asp Thr Gly Ala Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp  
145 150 155 160

eol f-seql . txt

Lys Lys Leu Leu Al a Val Pro Al a Al a Asn Thr Val Arg Phe Arg Cys  
165 170 175

Pro Al a Al a Gl y Asn Pro Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y  
180 185 190

Arg Gl u Phe Arg Gl y Gl u His Arg Ile Gl y Gl y Ile Lys Leu Arg His  
195 200 205

Gl n Gl n Trp Ser Leu Val Met Gl u Ser Val Val Pro Ser Asp Arg Gl y  
210 215 220

Asn Tyr Thr Cys Val Val Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr  
225 230 235 240

Tyr Thr Leu Asp Val Leu Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n  
245 250 255

Al a Gl y Leu Pro Al a Asn Gl n Thr Al a Val Leu Gl y Ser Asp Val Gl u  
260 265 270

Phe His Cys Lys Val Tyr Ser Asp Al a Gl n Pro His Ile Gl n Trp Leu  
275 280 285

Lys His Val Gl u Val Asn Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro  
290 295 300

Tyr Val Thr Val Leu Lys Val Ser Leu Gl u Ser Asn Al a Ser Met Ser  
305 310 315 320

Ser Asn Thr

<210> 19  
<211> 323  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 19

Met Gl y Al a Pro Al a Cys Al a Leu Al a Leu Cys Val Al a Val Al a Ile  
1 5 10 15

Val Al a Gl y Al a Ser Ser Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val  
20 25 30

eol f-seql.txt

Gly Arg Ala Ala Glu Val Pro Glu Pro Glu Pro Gly Gln Gln Glu Gln  
35 40 45

Leu Val Phe Gly Ser Gly Asp Ala Val Glu Leu Ser Cys Pro Pro Pro  
50 55 60

Gly Gly Gly Pro Met Gly Pro Thr Val Trp Val Lys Asp Gly Thr Gly  
65 70 75 80

Leu Val Pro Ser Glu Arg Val Leu Val Gly Pro Gln Arg Leu Gln Val  
85 90 95

Leu Asn Ala Ser His Glu Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg  
100 105 110

Leu Thr Gln Arg Val Leu Cys His Phe Ser Val Arg Val Thr Asp Ala  
115 120 125

Pro Ser Ser Gly Asp Asp Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr  
130 135 140

Gly Val Asp Thr Gly Ala Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp  
145 150 155 160

Lys Lys Leu Leu Ala Val Pro Ala Ala Asn Thr Val Arg Phe Arg Cys  
165 170 175

Pro Ala Ala Gly Asn Pro Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly  
180 185 190

Arg Glu Phe Arg Gly Glu His Arg Ile Gly Gly Ile Lys Leu Arg His  
195 200 205

Gln Gln Trp Ser Leu Val Met Glu Ser Val Val Pro Ser Asp Arg Gly  
210 215 220

Asn Tyr Thr Cys Val Val Glu Asn Lys Phe Gly Ser Ile Arg Gln Thr  
225 230 235 240

Tyr Thr Leu Asp Val Leu Glu Arg Ser Pro His Arg Pro Ile Leu Gln  
245 250 255

Ala Gly Leu Pro Ala Asn Gln Thr Ala Val Leu Gly Ser Asp Val Glu  
260 265 270

Phe His Ser Lys Val Tyr Ser Asp Ala Gln Pro His Ile Gln Trp Leu  
275 280 285

eol f-seql . txt

Lys His Val Glu Val Asn Glu Ser Lys Val Glu Pro Asp Glu Thr Pro  
290 295 300

Tyr Val Thr Val Leu Lys Val Ser Leu Glu Ser Asn Ala Ser Met Ser  
305 310 315 320

Ser Asn Thr

<210> 20  
<211> 987  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 20  
atgggagtga aggtgctgtt cgccctgatc tgtatcgccg tggccgaggc cgagtctctg 60  
ggcacagaac agagagtcgt gggcagagcc gccgaagtgc ctggacctga acctggccag 120  
caggaacagc tgggtttgg cagcggcgcac gccgtggAAC tgagctgtcc tccacctggc 180  
ggaggcccta tgggacctac cgtgtgggtc aaggatggca ccggactggc gcctagcgg 240  
agggtgctcg tgggacctca gagactgcag gtgctgaacg ccagccacga ggatagcggc 300  
gcctacagct gcagacagag actgacacag cgggtgctgt gccacttctc cgtcagagtg 360  
accgacgccc ctagctccgg cgacgatgag gatggcgaag atgaggccga ggacaccggc 420  
gtggacacag ggcgtccata ctggaccaga cccgagcggc tggacaagaa actgctggcc 480  
gtgcctgccc ccaacaccgt gcggttttaga tgtcctgccc cccgaaaccc caccggc 540  
atcagctggc tgaagaacgg cagagatTC cggggcggc acagaatcgg cggcatcaag 600  
ctgagacacc agcagtggc cctcgtatg gaaagcgtgg tgcccagcga ccggggcaac 660  
tacacctgtg tggtgaaaa caagttcggc agcatccggc agacctacac cctggacgtg 720  
ctggaaagaa gccccacag acccatcctg caggccggac tgcctgcaa tcagacagcc 780  
gtgctggca ggcgttgaa atttcacagc aaggtgtaca ggcgtccca gccccacatc 840  
cagtggctga aacacgtggc agtgaacggc agcaaagtgg gccccgacgg caccggc 900  
gtgaccgtgc tgaaggtgtc cctggaaagc aacgcccagca tgagcagcaa caccggactac 960  
aaggacgacg acgacaagtg aaagctt 987

<210> 21  
<211> 1131  
<212> DNA  
<213> Artificial Sequence

<220>

eol f-seql . txt

<223> Synthetic construct

<400> 21  
atgggagtga aggtgctgtt cgccctgatc tgtatcgccg tggccgaggc cgagtctctg 60  
ggcacagaac agagagtcgt gggcagagcc gccgaagtgc ctggacctga acctggccag 120  
caggaacagc tggtgtttgg cagcggcgac gccgtggaac ttagctgtcc tccacctggc 180  
ggaggcccta tgggacctac cgtgtgggtc aaggatggca ccggactggt gcctagcgag 240  
agggtgctcg tgggaccta gagactgcag gtgctgaacg ccagccacga ggatagcggc 300  
gcctacagct gcagacagag actgacacag cgggtgctgt gccacttctc cgtcagagtg 360  
accgacgccc ctagctccgg cgacgatgag gatggcgaag atgaggccga ggacaccggc 420  
gtggacacag gcgctccata ctggaccaga cccgagcggg tggacaagaa actgctggcc 480  
gtgcctgccc ccaacaccgt gcgggttaga tgtcctgccc ccggaaaccc caccccccagc 540  
atcagctggc tgaagaacgg cagagagttc cggggcggc acagaatcgg cggcatcaag 600  
ctgagacacc agcagtggtc cctcgtatg gaaagctgg tgcccagcga ccggggcaac 660  
tacacctgtg tggtgaaaaa caagttcggc agcatccggc agacctacac cctggacgtg 720  
ctggaaagaa gcccccacag acccatcctg caggccggac tgcctgccaa tcagacagcc 780  
gtgctggca gcgacgtgga atttcactgc aaggtgtaca gcgacgccc gcccccacatc 840  
cagtggctga aacacgtgga agtgaacggc agcaaagtgg gccccgacgg caccccttat 900  
gtgaccgtgc taaaaaccgc tggcgccaaat accaccgaca aagaactgga agtgcgtgagc 960  
ctgcacaacg tgacccctcgaa ggtgccggc gagtacacct gtctggccgg caacagcatc 1020  
ggcttcagcc accattctgc ctggctggtg gtgctgccc tgccctgga aagcaacgccc 1080  
agcatgagca gcaacaccga ctacaaggac gacgacgaca agtgaagact t 1131

<210> 22

<211> 2019

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 22  
atggagtctc tgggcacaga gcagagagtc gtgggcagag ccgctgaagt gcctggaccc 60  
gagcctggcc agcaggaaca gctggcttt ggctctggcg acgccgtgga actgagctgt 120  
cctccacctg gcggaggccc tatgggacct accgtgtggg tcaaggatgg caccggactg 180  
gtgcctagcg agagggtgct cgtggaccc cagagactgc aggtcctgaa cgccagccac 240  
gaggatagcg gcgcctacag ctgcagacag agactgaccc agcgggtgct gtgccacttc 300  
agcgtcagag tgaccgatgc ccccagcggc ggagatgacg aggtggcga ggtgaggcc 360

eol f-seql . txt

|            |            |            |             |             |            |      |
|------------|------------|------------|-------------|-------------|------------|------|
| gaggatacac | gcgtggacac | aggcgccct  | tactggacca  | gaccggagcg  | gatggacaag | 420  |
| aaactgctgg | ccgtgcctgc | cgcacacacc | gtgcggttta  | gatgcctgc   | cgcggaaac  | 480  |
| cccaccccca | gcatctcttg | gctgaagaac | ggcagagagt  | tccggggcga  | gcaccggatc | 540  |
| ggcggcatta | agctgagaca | ccagcagtgg | tccctggtca  | tggaaagcgt  | ggtgcccagc | 600  |
| gaccggggca | actacacctg | tgtggtgaa  | aacaagttcg  | gcagcatccg  | gcagacctac | 660  |
| accctggacg | tgctggaaag | aagccccac  | agacctatcc  | tgcaggccgg  | actgcctgcc | 720  |
| aatcagacag | ccgtgctggg | cagcgacgtg | gaatttcaca  | gcaaggtgta  | cagcgacgcc | 780  |
| cagccccaca | tccagtggct | gaagcacgtg | gaagtgaacg  | gcagcaaagt  | gggccccgac | 840  |
| ggcacccctt | acgtgaccgt | gctgaaagtg | tccctggaaa  | gcaacgcccag | catgagcagc | 900  |
| aacacccccc | tcgtgcggat | cgcacactg  | tctagcggag  | aggccctac   | cctggccaac | 960  |
| gtgtccgaac | tggaactgcc | cgcgcacccc | aagtgggagc  | tgagcagagc  | tagactgacc | 1020 |
| ctgggcaagc | ctctggcga  | gggctgtttt | ggacaggtgg  | tcatggccga  | ggccatcggc | 1080 |
| atcgacaagg | acagagccgc | caagcctgtg | accgtggccg  | tgaagatgct  | gaaggacgac | 1140 |
| gccaccgaca | aggacctgag | cgacctggtg | tccgagatgg  | aatgtatgaa  | gatgatcggc | 1200 |
| aagcacaaga | acatcatcaa | cctgctggc  | gcctgcaccc  | aggcggacc   | tctgtacgtg | 1260 |
| ctggtgaat  | acgcccaca  | ggcaacactg | agagagttcc  | tgagagccag  | aaggccccct | 1320 |
| ggcctggact | acagttcga  | tacctgcaag | ccccccgaag  | aacagctgac  | cttcaaggat | 1380 |
| ctggtgtcct | gcgcctatca | ggtggccaga | ggcatgaaat  | acctggccag  | ccagaagtgc | 1440 |
| atccaccggg | atctggccgc | cagaaacgtg | ctggtcacccg | aggacaacgt  | gatgaagatc | 1500 |
| gccgacttcg | gcctggcccg | ggacgtgcac | aacctggact  | actacaagaa  | aaccaccaac | 1560 |
| ggccggctgc | ccgtgaagtg | gatggccct  | gaggccctgt  | tcgacagagt  | gtacacccac | 1620 |
| cagagcgacg | tgtggtcctt | cggcgtgctg | ctgtgggaga  | tcttaccct   | ggcggcagc  | 1680 |
| ccttaccccg | gcatccctgt | ggaagaactg | ttcaagctgc  | tgaaagaggg  | ccacagaatg | 1740 |
| gacaagcccg | ccaactgcac | ccacgacctg | tacatgatca  | tgagagagt   | ctggcacgcc | 1800 |
| gctcccagcc | agaggcctac | ctttaagcag | ctggtggaaag | atctggaccg  | ggtgctgacc | 1860 |
| gtgaccagca | ccgacgagta | cctggatctg | agcgccttct  | tcgagcagta  | ctctcctggc | 1920 |
| ggccaggata | cccctagcag | cagctctagc | ggcgacgaca  | gcgtgttcgc  | ccacgatctg | 1980 |
| ctgcctccag | cccctcctag | ctctggcggc | tctagaacc   |             |            | 2019 |

<210> 23  
 <211> 1491  
 <212> DNA  
 <213> Artificial Sequence

<220>

eol f-seql . txt

<223> Synthetic construct

<400> 23

|             |             |             |            |             |             |      |
|-------------|-------------|-------------|------------|-------------|-------------|------|
| atggagagcc  | tggcacaga   | acagagagtc  | gtggcagag  | ccgccgaagt  | gcctggacct  | 60   |
| gaacctggcc  | agcaggaaca  | gctggcttt   | ggctctggcg | acgcccgtgg  | actgagctgt  | 120  |
| cctccacactg | gcggaggccc  | tatgggacct  | accgtgtggg | tcaaggatgg  | caccggactg  | 180  |
| gtgcctagcg  | agagggtgct  | cgtggacct   | cagagactgc | aggtcctgaa  | cgccagccac  | 240  |
| gaggatagcg  | gcccctacag  | ctgcagacag  | agactgaccc | agcgggtgct  | gtgccacttc  | 300  |
| agcgtcagag  | tgaccgatgc  | ccccagcagc  | ggagatgacg | aggatggcga  | ggatgaggcc  | 360  |
| gaggatacag  | gcgtggacac  | aggcgccct   | tactggacca | gaccggagcg  | gatggacaag  | 420  |
| aaactgctgg  | ccgtgcctgc  | cgccaaacacc | gtgcgggta  | gatgccctgc  | cgccggaaac  | 480  |
| cccacccca   | gcatctcttg  | gctgaagaac  | ggcagagagt | tccggggcga  | gcaccggatc  | 540  |
| ggcggcatta  | agctgagaca  | ccagcagtgg  | tccctggtca | tggaaagcgt  | ggtgccagc   | 600  |
| gaccggggca  | actacacctg  | tgtggtgaa   | aacaagttcg | gcagcatccg  | gcagacctac  | 660  |
| accctggacg  | tgctggaaag  | aagccccac   | agacctatcc | tgcaggccgg  | actgcctgcc  | 720  |
| aatcagacag  | ccgtgctggg  | cagcgtcgt   | gaatttact  | gcaaggtgta  | cagcgtcgtcc | 780  |
| cagccccaca  | tccagtggct  | gaagcacgtg  | gaagtgaacg | gcagcaaagt  | gggccccgac  | 840  |
| ggcacccctt  | acgtgaccgt  | gctgaaagtg  | tccctggaaa | gcaacgcccag | catgagcagc  | 900  |
| aacaccagcg  | gcagcggctc  | tggcagcgg   | tctggttctg | gctccggcag  | cgtgggttcc  | 960  |
| ccctacttcc  | cccgctggg   | ccggtacaac  | ctgaacttcc | atgaggccca  | gcaggcctgc  | 1020 |
| ctggaccagg  | atgccgtat   | cgccagcttc  | gaccagctgt | acgatgcttg  | gagaggcggc  | 1080 |
| ctggactggt  | gcaatgcccgg | ctggctgtct  | gacggcagcg | tgcagtaccc  | catcaccaag  | 1140 |
| cccagagagc  | cctgcggcgg  | acagaatacc  | gtgcccggcg | tgcggacta   | cggcttctgg  | 1200 |
| gacaaggaca  | agagcagata  | cgacgtttc   | tgcttcacca | gcaacttcaa  | cggccggttc  | 1260 |
| tactaccta   | tccacccac   | caagctgacc  | tacgacgagg | ccgtgcaggc  | ctgtctgaac  | 1320 |
| gatggcgtcc  | agatcgccaa  | agtggacag   | atcttcggcc | cctggaaagat | cctgggtctac | 1380 |
| gacagatgtg  | acgcccggat  | gctggccgac  | ggctccgtgc | ggtatccat   | cagccggcct  | 1440 |
| agaagaagat  | gcagccctac  | cgaggccgccc | gtcagattcg | tgggcttccc  | c           | 1491 |

<210> 24

<211> 1710

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 24

eol f-seql . txt

|            |            |             |             |             |            |      |
|------------|------------|-------------|-------------|-------------|------------|------|
| atgggagcac | cagcttgtgc | cctggctctg  | tgtgtggccg  | tggctattgt  | ggctggcgcc | 60   |
| tctagcgaga | gcctggcac  | agaacagaga  | gtcgtggca   | gagccgcca   | agtgcctgga | 120  |
| cctgaacctg | gccagcagga | acagctggc   | tttggctctg  | gcgacgcccgt | ggaactgagc | 180  |
| tgtccctcac | ctggcggagg | ccctatggga  | cctaccgtgt  | gggtcaagga  | tggcaccgga | 240  |
| ctgggccta  | gcgagaggg  | gctcgtggga  | cctcagagac  | tgcaggtcct  | gaacgcac   | 300  |
| cacgaggata | gcggcgcta  | cagctcaga   | cagagactga  | cccagcgggt  | gctgtgccac | 360  |
| ttcagcgtca | gagtgaccga | tgcccccagc  | agcggagatg  | acgaggatgg  | cgaggatgag | 420  |
| gccgaggata | caggcgtgga | cacaggcgcc  | ccttactgga  | ccagacccga  | gcggatggac | 480  |
| aagaaactgc | tggccgtgcc | tgccgccaac  | accgtgcgg   | ttagatgccc  | tgccgcccgg | 540  |
| aaccccaccc | ccagcatctc | ttggctgaag  | aacggcagag  | agttccgggg  | cgagcaccgg | 600  |
| atcggcggca | ttaagctgag | acaccagcag  | tggccctgg   | tcatggaaag  | cgtggtgccc | 660  |
| agcgaccggg | gcaactacac | ctgtgtgg    | gaaaacaagt  | tcggcagcat  | ccggcagacc | 720  |
| tacaccctgg | acgtgctgga | aagaagcccc  | cacagaccta  | tcctgcaggc  | cggactgcct | 780  |
| gccaatcaga | cagccgtgct | gggcagcgac  | gtggaaattc  | actgcaaggt  | gtacagcgac | 840  |
| gcccagcccc | acatccagtg | gctgaagcac  | gtggaaagtga | acggcagcaa  | agtggggccc | 900  |
| gacggcaccc | cttacgtgac | cgtgctgaaa  | accgctggcg  | ccaacaccac  | cgacaagaa  | 960  |
| ctggaagtgc | tgagcctgca | caacgtgacc  | ttcaggacg   | ccggcgagta  | cacctgtctg | 1020 |
| gccggcaata | gcatcggtt  | cagccaccac  | tctgcctggc  | tgggtgtct   | gccaggcgga | 1080 |
| ggctctgtgt | ccctggaaag | caacgccagc  | atgagcagca  | acaccagcg   | cagcggctct | 1140 |
| ggcagcgat  | ctggttctgg | ctccggcagc  | gtgggttcc   | cctacttccc  | ccggctgggc | 1200 |
| cgttacaacc | tgaactttca | tgaggcccag  | caggcctgcc  | tggaccagga  | tgccgtgatc | 1260 |
| gccagcttcg | accagctgta | cgatgttgg   | agaggcggcc  | tggactgg    | caatgccgc  | 1320 |
| tggctgtctg | acggcagcgt | gcagtacccc  | atcaccaagc  | ccagagagcc  | ctgcggcgga | 1380 |
| cagaataccg | tgcccggt   | gccaactac   | ggcttctgg   | acaaggacaa  | gagcagatac | 1440 |
| gacgtttct  | gcttaccac  | caacttcaac  | ggccggttct  | actacctgat  | ccacccacc  | 1500 |
| aagctgacct | acgacgaggc | cgtgcaggcc  | tgtctgaac   | atggcgccca  | gatgccaa   | 1560 |
| gtgggacaga | tcttcgcgc  | ctggaaagatc | ctgggctac   | acagatgtga  | cgccggatgg | 1620 |
| ctggccgacg | gctccgtg   | gtatccatc   | agccggccta  | gaagaagatg  | cagccctacc | 1680 |
| gaggccgccc | tcaattcgt  | gggcttcccc  |             |             |            | 1710 |

<210> 25  
 <211> 236  
 <212> PRT  
 <213> Homo sapiens

eol f-seql . txt

<400> 25

Gl y Gl y Gl y Gl y Al a Gl y Gl y Gl y Asp Lys Thr His Thr Cys Pro  
1 5 10 15

Pro Cys Pro Al a Pro Gl u Leu Leu Gl y Gl y Pro Ser Val Phe Leu Phe  
20 25 30

Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Gl u Val  
35 40 45

Thr Cys Val Val Val Asp Val Ser His Gl u Asp Pro Gl u Val Lys Phe  
50 55 60

Asn Trp Tyr Val Asp Gl y Val Gl u Val His Asn Al a Lys Thr Lys Pro  
65 70 75 80

Arg Gl u Gl u Gl n Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr  
85 90 95

Val Leu His Gl n Asp Trp Leu Asn Gl y Lys Gl u Tyr Lys Cys Lys Val  
100 105 110

Ser Asn Lys Al a Leu Pro Al a Pro Ile Gl u Lys Thr Ile Ser Lys Al a  
115 120 125

Lys Gl y Gl n Pro Arg Gl u Pro Gl n Val Tyr Thr Leu Pro Pro Ser Arg  
130 135 140

Asp Gl u Leu Thr Lys Asn Gl n Val Ser Leu Thr Cys Leu Val Lys Gl y  
145 150 155 160

Phe Tyr Pro Ser Asp Ile Al a Val Gl u Trp Gl u Ser Asn Gl y Gl n Pro  
165 170 175

Gl u Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gl y Ser  
180 185 190

Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gl n Gl n  
195 200 205

Gl y Asn Val Phe Ser Cys Ser Val Met His Gl u Al a Leu His Asn His  
210 215 220

Tyr Thr Gl n Lys Ser Leu Ser Leu Ser Pro Gl y Lys  
225 230 235

eol f-seql . txt

<210> 26  
<211> 236  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 26

Gly Gly Gly Gly Ala Gly Gly Gly Asp Lys Thr His Thr Cys Pro  
1 5 10 15

Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser Val Phe Leu Phe  
20 25 30

Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val  
35 40 45

Thr Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe  
50 55 60

Asn Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro  
65 70 75 80

Arg Glu Glu Gln Tyr Ala Ser Thr Tyr Arg Val Val Ser Val Leu Thr  
85 90 95

Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val  
100 105 110

Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala  
115 120 125

Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg  
130 135 140

Glu Glu Met Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly  
145 150 155 160

Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro  
165 170 175

Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser  
180 185 190

Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln  
195 200 205

Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His

eol f-seql . txt

210

215

220

Tyr Thr Glu Lys Ser Leu Ser Leu Ser Pro Glu Lys  
225 230 235

<210> 27

<211> 585

<212> PRT

<213> Homo sapiens

<400> 27

Asp Ala His Lys Ser Glu Val Ala His Arg Phe Lys Asp Leu Glu Glu  
1 5 10 15

Glu Asn Phe Lys Ala Leu Val Leu Ile Ala Phe Ala Glu Tyr Leu Glu  
20 25 30

Glu Cys Pro Phe Glu Asp His Val Lys Leu Val Asn Glu Val Thr Glu  
35 40 45

Phe Ala Lys Thr Cys Val Ala Asp Glu Ser Ala Glu Asn Cys Asp Lys  
50 55 60

Ser Leu His Thr Leu Phe Glu Asp Lys Leu Cys Thr Val Ala Thr Leu  
65 70 75 80

Arg Glu Thr Tyr Glu Glu Met Ala Asp Cys Cys Ala Lys Glu Glu Pro  
85 90 95

Glu Arg Asn Glu Cys Phe Leu Glu His Lys Asp Asp Asn Pro Asn Leu  
100 105 110

Pro Arg Leu Val Arg Pro Glu Val Asp Val Met Cys Thr Ala Phe His  
115 120 125

Asp Asn Glu Glu Thr Phe Leu Lys Lys Tyr Leu Tyr Glu Ile Ala Arg  
130 135 140

Arg His Pro Tyr Phe Tyr Ala Pro Glu Leu Leu Phe Phe Ala Lys Arg  
145 150 155 160

Tyr Lys Ala Ala Phe Thr Glu Cys Cys Glu Ala Ala Asp Lys Ala Ala  
165 170 175

Cys Leu Leu Pro Lys Leu Asp Glu Leu Arg Asp Glu Glu Lys Ala Ser  
180 185 190

Ser Ala Lys Glu Arg Leu Lys Cys Ala Ser Leu Glu Lys Phe Glu Glu

eol f-seql . txt

195 200 205  
Arg Ala Phe Lys Ala Trp Ala Val Ala Arg Leu Ser Glu Arg Phe Pro  
210 215 220  
Lys Ala Glu Phe Ala Glu Val Ser Lys Leu Val Thr Asp Leu Thr Lys  
225 230 235 240  
Val His Thr Glu Cys Cys His Glu Asp Leu Leu Glu Cys Ala Asp Asp  
245 250 255  
Arg Ala Asp Leu Ala Lys Tyr Ile Cys Glu Asn Glu Asp Ser Ile Ser  
260 265 270  
Ser Lys Leu Lys Glu Cys Cys Glu Lys Pro Leu Leu Glu Lys Ser His  
275 280 285  
Cys Ile Ala Glu Val Glu Asn Asp Glu Met Pro Ala Asp Leu Pro Ser  
290 295 300  
Leu Ala Ala Asp Phe Val Glu Ser Lys Asp Val Cys Lys Asn Tyr Ala  
305 310 315 320  
Glu Ala Lys Asp Val Phe Leu Glu Met Phe Leu Tyr Glu Tyr Ala Arg  
325 330 335  
Arg His Pro Asp Tyr Ser Val Val Leu Leu Leu Arg Leu Ala Lys Thr  
340 345 350  
Tyr Glu Thr Thr Leu Glu Lys Cys Cys Ala Ala Ala Asp Pro His Glu  
355 360 365  
Cys Tyr Ala Lys Val Phe Asp Glu Phe Lys Pro Leu Val Glu Glu Pro  
370 375 380  
Gln Asn Leu Ile Lys Gln Asn Cys Glu Leu Phe Glu Gln Leu Glu Glu  
385 390 395 400  
Tyr Lys Phe Gln Asn Ala Leu Leu Val Arg Tyr Thr Lys Lys Val Pro  
405 410 415  
Gln Val Ser Thr Pro Thr Leu Val Glu Val Ser Arg Asn Leu Glu Lys  
420 425 430  
Val Glu Ser Lys Cys Cys Lys His Pro Glu Ala Lys Arg Met Pro Cys  
435 440 445

eol f-seql.txt

Ala Glu Asp Tyr Leu Ser Val Val Leu Asn Glu Leu Cys Val Leu His  
450 455 460

Gl u Lys Thr Pro Val Ser Asp Arg Val Thr Lys Cys Cys Thr Gl u Ser  
465 470 475 480

Leu Val Asn Arg Arg Pro Cys Phe Ser Ala Leu Gl u Val Asp Gl u Thr  
485 490 495

Tyr Val Pro Lys Gl u Phe Asn Ala Gl u Thr Phe Thr Phe His Ala Asp  
500 505 510

Ile Cys Thr Leu Ser Gl u Lys Gl u Arg Gl n Ile Lys Lys Gl n Thr Ala  
515 520 525

Leu Val Gl u Leu Val Lys His Lys Pro Lys Ala Thr Lys Gl u Gl n Leu  
530 535 540

Lys Ala Val Met Asp Asp Phe Ala Ala Phe Val Gl u Lys Cys Cys Lys  
545 550 555 560

Ala Asp Asp Lys Gl u Thr Cys Phe Ala Gl u Gl u Gly Lys Lys Leu Val  
565 570 575

Ala Ala Ser Gl n Ala Ala Leu Gl y Leu  
580 585

<210> 28  
<211> 9  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 28

Gl y Gl y Gl y Gl y Ala Gl y Gl y Gl y Gl y  
1 5

<210> 29  
<211> 15  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 29

Gl y Gl y Gl y Gl y Ser Gl y Gl y Gl y Gl y Ser  
1 5 10 15

eol f-seql . txt

<210> 30  
<211> 24  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 30

Ala Gly Gly Thr Gly Gly Cys Cys Thr Thr Thr Gly Ala Cys Ala Cys  
1 5 10 15

Cys Thr Ala Cys Cys Ala Gly Gly  
20

<210> 31  
<211> 24  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 31

Thr Cys Thr Gly Thr Thr Gly Thr Gly Thr Thr Cys Cys Thr Cys  
1 5 10 15

Cys Cys Thr Gly Thr Thr Gly Gly  
20

<210> 32  
<211> 806  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 32

Met Gly Ala Pro Ala Cys Ala Leu Ala Leu Cys Val Ala Val Ala Ile  
1 5 10 15

Val Ala Gly Ala Ser Ser Glu Ser Leu Gly Thr Glu Glu Arg Val Val  
20 25 30

Gly Arg Ala Ala Glu Val Pro Gly Pro Glu Pro Gly Glu Glu Glu Glu  
35 40 45

Leu Val Phe Gly Ser Gly Asp Ala Val Glu Leu Ser Cys Pro Pro Pro  
50 55 60

eol f-seql.txt

Gly Gly Gly Pro Met Gly Pro Thr Val Trp Val Lys Asp Gly Thr Gly  
65 70 75 80

Leu Val Pro Ser Glu Arg Val Leu Val Gly Pro Gln Arg Leu Gln Val  
85 90 95

Leu Asn Ala Ser His Glu Asp Ser Gly Ala Tyr Ser Cys Arg Gln Arg  
100 105 110

Leu Thr Gln Arg Val Leu Cys His Phe Ser Val Arg Val Thr Asp Ala  
115 120 125

Pro Ser Ser Gly Asp Asp Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr  
130 135 140

Gly Val Asp Thr Gly Ala Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp  
145 150 155 160

Lys Lys Leu Leu Ala Val Pro Ala Ala Asn Thr Val Arg Phe Arg Cys  
165 170 175

Pro Ala Ala Gly Asn Pro Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly  
180 185 190

Arg Glu Phe Arg Gly Glu His Arg Ile Gly Gly Ile Lys Leu Arg His  
195 200 205

Gln Gln Trp Ser Leu Val Met Glu Ser Val Val Pro Ser Asp Arg Gly  
210 215 220

Asn Tyr Thr Cys Val Val Glu Asn Lys Phe Gly Ser Ile Arg Gln Thr  
225 230 235 240

Tyr Thr Leu Asp Val Leu Glu Arg Ser Pro His Arg Pro Ile Leu Gln  
245 250 255

Ala Gly Leu Pro Ala Asn Gln Thr Ala Val Leu Gly Ser Asp Val Glu  
260 265 270

Phe His Cys Lys Val Tyr Ser Asp Ala Gln Pro His Ile Gln Trp Leu  
275 280 285

Lys His Val Glu Val Asn Gly Ser Lys Val Gly Pro Asp Gly Thr Pro  
290 295 300

Tyr Val Thr Val Leu Lys Thr Ala Gly Ala Asn Thr Thr Asp Lys Glu  
305 310 315 320

eol f-seql . txt

Leu Glu Val Leu Ser Leu His Asn Val Thr Phe Glu Asp Ala Gly Glu  
325 330 335

Tyr Thr Cys Leu Ala Gly Asn Ser Ile Gly Phe Ser His His Ser Ala  
340 345 350

Trp Leu Val Val Leu Pro Ala Glu Glu Glu Leu Val Glu Ala Asp Glu  
355 360 365

Ala Gly Ser Val Tyr Ala Gly Ile Leu Ser Tyr Gly Val Gly Phe Phe  
370 375 380

Leu Phe Ile Leu Val Val Ala Ala Val Thr Leu Cys Arg Leu Arg Ser  
385 390 395 400

Pro Pro Lys Lys Gly Leu Gly Ser Pro Thr Val His Lys Ile Ser Arg  
405 410 415

Phe Pro Leu Lys Arg Gln Val Ser Leu Glu Ser Asn Ala Ser Met Ser  
420 425 430

Ser Asn Thr Pro Leu Val Arg Ile Ala Arg Leu Ser Ser Gly Glu Gly  
435 440 445

Pro Thr Leu Ala Asn Val Ser Glu Leu Glu Leu Pro Ala Asp Pro Lys  
450 455 460

Trp Glu Leu Ser Arg Ala Arg Leu Thr Leu Gly Lys Pro Leu Gly Glu  
465 470 475 480

Gly Cys Phe Gly Gln Val Val Met Ala Glu Ala Ile Gly Ile Asp Lys  
485 490 495

Asp Arg Ala Ala Lys Pro Val Thr Val Ala Val Lys Met Leu Lys Asp  
500 505 510

Asp Ala Thr Asp Lys Asp Leu Ser Asp Leu Val Ser Glu Met Glu Met  
515 520 525

Met Lys Met Ile Gly Lys His Lys Asn Ile Ile Asn Leu Leu Gly Ala  
530 535 540

Cys Thr Gln Gly Gly Pro Leu Tyr Val Leu Val Glu Tyr Ala Ala Lys  
545 550 555 560

Gly Asn Leu Arg Glu Phe Leu Arg Ala Arg Arg Pro Pro Gly Leu Asp  
565 570 575

eol f-seql . txt

Tyr Ser Phe Asp Thr Cys Lys Pro Pro Glu Glu Glu Leu Thr Phe Lys  
580 585 590

Asp Leu Val Ser Cys Ala Tyr Glu Val Ala Arg Gly Met Glu Tyr Leu  
595 600 605

Ala Ser Glu Lys Cys Ile His Arg Asp Leu Ala Ala Arg Asn Val Leu  
610 615 620

Val Thr Glu Asp Asn Val Met Lys Ile Ala Asp Phe Gly Leu Ala Arg  
625 630 635 640

Asp Val His Asn Leu Asp Tyr Tyr Lys Lys Thr Thr Asn Gly Arg Leu  
645 650 655

Pro Val Lys Trp Met Ala Pro Glu Ala Leu Phe Asp Arg Val Tyr Thr  
660 665 670

His Glu Ser Asp Val Trp Ser Phe Gly Val Leu Leu Trp Glu Ile Phe  
675 680 685

Thr Leu Gly Gly Ser Pro Tyr Pro Gly Ile Pro Val Glu Glu Leu Phe  
690 695 700

Lys Leu Leu Lys Glu Gly His Arg Met Asp Lys Pro Ala Asn Cys Thr  
705 710 715 720

His Asp Leu Tyr Met Ile Met Arg Glu Cys Trp His Ala Ala Pro Ser  
725 730 735

Glu Arg Pro Thr Phe Lys Glu Leu Val Glu Asp Leu Asp Arg Val Leu  
740 745 750

Thr Val Thr Ser Thr Asp Glu Tyr Leu Asp Leu Ser Ala Pro Phe Glu  
755 760 765

Glu Tyr Ser Pro Gly Gly Glu Asp Thr Pro Ser Ser Ser Ser Glu  
770 775 780 785

Asp Asp Ser Val Phe Ala His Asp Leu Leu Pro Pro Ala Pro Pro Ser  
790 795 800

Ser Gly Gly Ser Arg Thr  
805

eol f-seql . txt

<211> 349  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 33

Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Al a Al a Gl u Val  
1 5 10 15

Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe Gl y Ser Gl y  
20 25 30

Asp Al a Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y Pro Met Gl y  
35 40 45

Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro Ser Gl u Arg  
50 55 60

Val Leu Val Gl y Pro Gl n Arg Leu Gl n Val Leu Asn Al a Ser His Gl u  
65 70 75 80

Asp Ser Gl y Al a Tyr Ser Cys Arg Gl n Arg Leu Thr Gl n Arg Val Leu  
85 90 95

Cys His Phe Ser Val Arg Val Thr Asp Al a Pro Ser Ser Gl y Asp Asp  
100 105 110

Gl u Asp Gl y Gl u Asp Gl u Al a Gl u Asp Thr Gl y Val Asp Thr Gl y Al a  
115 120 125

Pro Tyr Trp Thr Arg Pro Gl u Arg Met Asp Lys Lys Leu Leu Al a Val  
130 135 140

Pro Al a Al a Asn Thr Val Arg Phe Arg Cys Pro Al a Al a Gl y Asn Pro  
145 150 155 160

Thr Pro Ser Ile Ser Trp Leu Lys Asn Gl y Arg Gl u Phe Arg Gl y Gl u  
165 170 175

His Arg Ile Gl y Gl y Ile Lys Leu Arg His Gl n Gl n Trp Ser Leu Val  
180 185 190

Met Gl u Ser Val Val Pro Ser Asp Arg Gl y Asn Tyr Thr Cys Val Val  
195 200 205

Gl u Asn Lys Phe Gl y Ser Ile Arg Gl n Thr Tyr Thr Leu Asp Val Leu  
210 215 220

eol f-seql . txt

Gl u Arg Ser Pro His Arg Pro Ile Leu Gl n Ala Gl y Leu Pro Ala Asn  
225 230 235 240

Gl n Thr Ala Val Leu Gl y Ser Asp Val Gl u Phe His Cys Lys Val Tyr  
245 250 255

Ser Asp Ala Gl n Pro His Ile Gl n Trp Leu Lys His Val Gl u Val Asn  
260 265 270

Gl y Ser Lys Val Gl y Pro Asp Gl y Thr Pro Tyr Val Thr Val Leu Lys  
275 280 285

Thr Ala Gl y Ala Asn Thr Thr Asp Lys Gl u Leu Gl u Val Leu Ser Leu  
290 295 300

His Asn Val Thr Phe Gl u Asp Ala Gl y Gl u Tyr Thr Cys Leu Ala Gl y  
305 310 315 320

Asn Ser Ile Gl y Phe Ser His His Ser Ala Trp Leu Val Val Leu Pro  
325 330 335

Val Ser Leu Gl u Ser Asn Ala Ser Met Ser Ser Asn Thr  
340 345

<210> 34

<211> 368

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetic construct

<400> 34

Met Met Ser Phe Val Ser Leu Leu Leu Val Gl y Ile Leu Phe His Ala  
1 5 10 15

Thr Gl n Ala Gl u Ser Leu Gl y Thr Gl u Gl n Arg Val Val Gl y Arg Ala  
20 25 30

Al a Gl u Val Pro Gl y Pro Gl u Pro Gl y Gl n Gl n Gl u Gl n Leu Val Phe  
35 40 45

Gl y Ser Gl y Asp Ala Val Gl u Leu Ser Cys Pro Pro Pro Gl y Gl y Gl y  
50 55 60

Pro Met Gl y Pro Thr Val Trp Val Lys Asp Gl y Thr Gl y Leu Val Pro  
65 70 75 80

eol f-seql . txt

Ser Glu Arg Val Leu Val Gly Pro Glu Arg Leu Glu Val Leu Asn Ala  
85 90 95

Ser His Glu Asp Ser Gly Ala Tyr Ser Cys Arg Glu Arg Leu Thr Glu  
100 105 110

Arg Val Leu Cys His Phe Ser Val Arg Val Thr Asp Ala Pro Ser Ser  
115 120 125

Gly Asp Asp Glu Asp Gly Glu Asp Glu Ala Glu Asp Thr Gly Val Asp  
130 135 140

Thr Gly Ala Pro Tyr Trp Thr Arg Pro Glu Arg Met Asp Lys Lys Leu  
145 150 155 160

Leu Ala Val Pro Ala Ala Asn Thr Val Arg Phe Arg Cys Pro Ala Ala  
165 170 175

Gly Asn Pro Thr Pro Ser Ile Ser Trp Leu Lys Asn Gly Arg Glu Phe  
180 185 190

Arg Gly Glu His Arg Ile Gly Gly Ile Lys Leu Arg His Glu Glu Trp  
195 200 205

Ser Leu Val Met Glu Ser Val Val Pro Ser Asp Arg Gly Asn Tyr Thr  
210 215 220

Cys Val Val Glu Asn Lys Phe Gly Ser Ile Arg Glu Thr Tyr Thr Leu  
225 230 235 240

Asp Val Leu Glu Arg Ser Pro His Arg Pro Ile Leu Glu Ala Gly Leu  
245 250 255

Pro Ala Asn Glu Thr Ala Val Leu Gly Ser Asp Val Glu Phe His Cys  
260 265 270

Lys Val Tyr Ser Asp Ala Glu Pro His Ile Glu Trp Leu Lys His Val  
275 280 285

Glu Val Asn Gly Ser Lys Val Gly Pro Asp Gly Thr Pro Tyr Val Thr  
290 295 300

Val Leu Lys Thr Ala Glu Ala Asn Thr Thr Asp Lys Glu Leu Glu Val  
305 310 315 320

Leu Ser Leu His Asn Val Thr Phe Glu Asp Ala Gly Glu Tyr Thr Cys  
325 330 335

eol f-seql . txt

Leu Ala Gly Asn Ser Ile Gly Phe Ser His His Ser Ala Trp Leu Val  
340 345 350

Val Leu Pro Val Ser Leu Glu Ser Asn Ala Ser Met Ser Ser Asn Thr  
355 360 365

<210> 35  
<211> 19  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 35

Met Met Ser Phe Val Ser Leu Leu Leu Val Gly Ile Leu Phe His Ala  
1 5 10 15

Thr Glu Ala

<210> 36  
<211> 963  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Synthetic construct

<400> 36  
atgatgtcct ttgtctctc gctcctggtt ggcatcctat tccatgccac ccaggccgag 60  
tccttgggaa cggagcagcg cgtcggtggg cgagcggcag aagtccggg cccagagccc 120  
ggccagcagg agcagtttgtt cttcggcagc gggatgctg tggagctgag ctgtccccg 180  
cccggggtg gtcccatggg gcccactgtc tgggtcaagg atggcacagg gctgggcc 240  
tcggagcgtg tcctgggtgg gccccagcgg ctgcaggtgc tgaatgcctc ccacgaggac 300  
tccggggcct acagctgccc gcagcggctc acgcagcgcg tactgtCCA cttcagtgtg 360  
cgggtgacag acgctccatc ctcggagat gacgaagacg gggaggacga ggctgaggac 420  
acaggtgtgg acacaggggc cccttactgg acacggccc agcggatgga caagaagctg 480  
ctggccgtgc cggccgccaa caccgtccgc ttccgctgcc cagccgctgg caacccact 540  
ccctccatct cctggctgaa gaacggcagg gagttccgcg gcgagcaccc cattggaggc 600  
atcaagctgc ggcattcagca gtggagcctg gtcatggaaa gcgtggtgcc ctggaccgc 660  
ggcaactaca cctgcgtcgt ggagaacaag tttggcagca tccggcagac gtacacgctg 720  
gacgtgctgg agcgctcccc gcaccggccc atcctgcagg cggggctgcc ggccaaccag 780

eol f-seql . txt

|             |                     |             |             |            |            |      |
|-------------|---------------------|-------------|-------------|------------|------------|------|
| acggcggtgc  | tgggcagcga          | cgtggagttc  | caactccaagg | tgtacagtga | cgcacagccc | 840  |
| cacatccagt  | ggctcaagca          | cgtggaggtg  | aatggcagca  | aggtgggccc | ggacggcaca | 900  |
| ccctacgtta  | ccgtgctcaa          | ggtgtccctg  | gagtccaacg  | cgtccatgag | ctccaacaca | 960  |
|             |                     |             | tga         |            |            | 963  |
| <210>       | 37                  |             |             |            |            |      |
| <211>       | 1107                |             |             |            |            |      |
| <212>       | DNA                 |             |             |            |            |      |
| <213>       | Artificial Sequence |             |             |            |            |      |
| <220>       |                     |             |             |            |            |      |
| <223>       | Synthetic construct |             |             |            |            |      |
| <400>       | 37                  |             |             |            |            |      |
| atgatgtcct  | ttgtctctt           | gctcctggtt  | ggcatcctat  | tccatgccac | ccaggccgag | 60   |
| tccttggga   | cggagcagcg          | cgtcggtggg  | cgagcggcag  | aagtcccggg | cccagagccc | 120  |
| ggccagcagg  | agcagtttgt          | cttcggcagc  | ggggatgctg  | tggagctgag | ctgtcccccg | 180  |
| cccgggggtg  | gtcccatggg          | gcccaactgtc | tgggtcaagg  | atggcacagg | gctgggtccc | 240  |
| tcggagcgtg  | tcctgggtgg          | gccccagcgg  | ctgcaggtgc  | tgaatgcctc | ccacgaggac | 300  |
| tccggggcct  | acagctgccc          | gcagcggctc  | acgcagcgcg  | tactgtgcca | ttcagtgtg  | 360  |
| cgggtgacag  | acgctccatc          | ctcgggagat  | gacgaagacg  | gggaggacga | ggctgaggac | 420  |
| acaggtgtgg  | acacaggggc          | cccttactgg  | acacggcccg  | agcggatgga | caagaagctg | 480  |
| ctggccgtgc  | cggccgccaa          | caccgtccgc  | ttccgctgcc  | cagccgctgg | caacccact  | 540  |
| ccctccatct  | cctggctgaa          | gaacggcagg  | gagttccgcg  | gcgagcaccc | cattggaggc | 600  |
| atcaagctgc  | ggcatcagca          | gtggagcctg  | gtcatggaaa  | gcgtggtgcc | ctcggaccgc | 660  |
| ggcaactaca  | cctgcgtcgt          | ggagaacaag  | tttggcagca  | tccggcagac | gtacacgctg | 720  |
| gacgtgctgg  | agcgctcccc          | gcaccggccc  | atccctgcagg | ctgggctgcc | tgctaaccag | 780  |
| acagcgggtgc | tgggcagcga          | cgtggagttc  | cactgcaagg  | tgtacagtga | cgcacagccc | 840  |
| cacatccagt  | ggctcaagca          | cgtggaggtg  | aatggcagca  | aggtgggccc | ggacggcaca | 900  |
| ccctacgtta  | ccgtgctcaa          | gacggcgggc  | gctaacacca  | ccgacaagga | gctagaggtt | 960  |
| ctctccttgc  | acaacgtcac          | ctttgaggac  | gccggggagt  | acacctgcct | ggcgggcaat | 1020 |
| tctattgggt  | tttctcatca          | ctctgcgtgg  | ctgggtggtgc | tgccagtgtc | cctggagtcc | 1080 |
| aacgcgtcca  | tgagctccaa          | cacatga     |             |            |            | 1107 |
| <210>       | 38                  |             |             |            |            |      |
| <211>       | 216                 |             |             |            |            |      |
| <212>       | PRT                 |             |             |            |            |      |
| <213>       | Homo sapiens        |             |             |            |            |      |
| <400>       | 38                  |             |             |            |            |      |

eol f-seql . txt

Met Arg Ser Gly Cys Val Val Val His Val Trp Ile Leu Ala Gly Leu  
1 5 10 15

Trp Leu Ala Val Ala Gly Arg Pro Leu Ala Phe Ser Asp Ala Gly Pro  
20 25 30

His Val His Tyr Gly Trp Gly Asp Pro Ile Arg Leu Arg His Leu Tyr  
35 40 45

Thr Ser Gly Pro His Gly Leu Ser Ser Cys Phe Leu Arg Ile Arg Ala  
50 55 60

Asp Gly Val Val Asp Cys Ala Arg Gly Gln Ser Ala His Ser Leu Leu  
65 70 75 80

Gl u Ile Lys Ala Val Ala Leu Arg Thr Val Ala Ile Lys Gly Val His  
85 90 95

Ser Val Arg Tyr Leu Cys Met Gly Ala Asp Gly Lys Met Gln Gly Leu  
100 105 110

Leu Gln Tyr Ser Glu Glu Asp Cys Ala Phe Glu Glu Glu Ile Arg Pro  
115 120 125

Asp Gly Tyr Asn Val Tyr Arg Ser Glu Lys His Arg Leu Pro Val Ser  
130 135 140

Leu Ser Ser Ala Lys Gln Arg Gln Leu Tyr Lys Asn Arg Gly Phe Leu  
145 150 155 160

Pro Leu Ser His Phe Leu Pro Met Leu Pro Met Val Pro Glu Glu Pro  
165 170 175

Gl u Asp Leu Arg Gly His Leu Glu Ser Asp Met Phe Ser Ser Pro Leu  
180 185 190

Gl u Thr Asp Ser Met Asp Pro Phe Gl y Leu Val Thr Gl y Leu Gl u Ala  
195 200 205

Val Arg Ser Pro Ser Phe Gl u Lys  
210 215

<210> 39

<211> 209

<212> PRT

<213> Homo sapiens

<400> 39

eol f-seql . txt

Met Asp Ser Asp Glu Thr Gly Phe Glu His Ser Gly Leu Trp Val Ser  
1 5 10 15

Val Leu Ala Gly Leu Leu Leu Gly Ala Cys Gln Ala His Pro Ile Pro  
20 25 30

Asp Ser Ser Pro Leu Leu Gln Phe Gly Gly Gln Val Arg Gln Arg Tyr  
35 40 45

Leu Tyr Thr Asp Asp Ala Gln Gln Thr Glu Ala His Leu Glu Ile Arg  
50 55 60

Glu Asp Gly Thr Val Gly Gly Ala Ala Asp Gln Ser Pro Glu Ser Leu  
65 70 75 80

Leu Gln Leu Lys Ala Leu Lys Pro Gly Val Ile Gln Ile Leu Gly Val  
85 90 95

Lys Thr Ser Arg Phe Leu Cys Gln Arg Pro Asp Gly Ala Leu Tyr Gly  
100 105 110

Ser Leu His Phe Asp Pro Glu Ala Cys Ser Phe Arg Glu Leu Leu Leu  
115 120 125

Glu Asp Gly Tyr Asn Val Tyr Gln Ser Glu Ala His Gly Leu Pro Leu  
130 135 140

His Leu Pro Gly Asn Lys Ser Pro His Arg Asp Pro Ala Pro Arg Gly  
145 150 155 160

Pro Ala Arg Phe Leu Pro Leu Pro Gly Leu Pro Pro Ala Leu Pro Glu  
165 170 175

Pro Pro Gly Ile Leu Ala Pro Gln Pro Pro Asp Val Gly Ser Ser Asp  
180 185 190

Pro Leu Ser Met Val Gly Pro Ser Gln Gly Arg Ser Pro Ser Tyr Ala  
195 200 205

Ser