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(54) **Title**: VIRAL VECTORS ENCODING RECOMBINANT FVIII VARIANTS WITH INCREASED EXPRESSION FOR GENE THERAPY OF HEMOPHILIA A

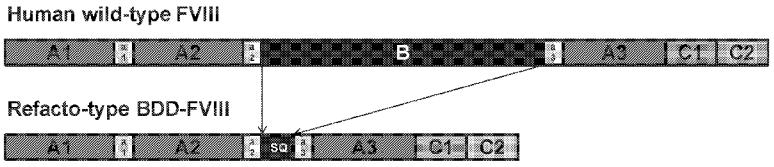

Figure 1

(57) **Abstract**: The present disclosure provides, among other aspects, codon-altered polynucleotides encoding Factor VIII variants for expression in mammalian cells. In some embodiments, the disclosure also provides mammalian gene therapy vectors and methods for treating hemophilia A.

WO 2017/083762 A1

VIRAL VECTORS ENCODING RECOMBINANT FVIII VARIANTS WITH INCREASED EXPRESSION FOR GENE THERAPY OF HEMOPHILIA A

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to United States Provisional Patent Application No. 62/255,317, filed November 13, 2015, the content of which is hereby incorporated by reference in its entirety for all purposes.

SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on November 9, 2016, is named 008073_5107_WO_Sequence_Listing.txt and is 353,479 bytes in size.

BACKGROUND OF THE DISCLOSURE

[0003] Blood coagulation proceeds through a complex and dynamic biological pathway of interdependent biochemical reactions, referred to as the coagulation cascade. Coagulation Factor VIII (FVIII) is a key component in the cascade. Factor VIII is recruited to bleeding sites, and forms a Xase complex with activated Factor IX (FIXa) and Factor X (FX). The Xase complex activates FX, which in turn activates prothrombin to thrombin, which then activates other components in the coagulation cascade to generate a stable clot (reviewed in Saenko *et al.*, *Trends Cardiovasc. Med.*, 9:185-192 (1999); Lenting *et al.*, *Blood*, 92:3983-3996 (1998)).

[0004] Hemophilia A is a congenital X-linked bleeding disorder characterized by a deficiency in Factor VIII activity. Diminished Factor VIII activity inhibits a positive feedback loop in the coagulation cascade. This causes incomplete coagulation, which manifests as bleeding episodes with increased duration, extensive bruising, spontaneous oral and nasal bleeding, joint stiffness and chronic pain, and possibly internal bleeding and anemia in severe cases (Zhang *et al.*, *Clinic. Rev. Allerg. Immunol.*, 37:114-124 (2009)).

[0005] Conventionally, hemophilia A is treated by Factor VIII replacement therapy, which consists of administering Factor VIII protein (e.g., plasma-derived or recombinantly-produced Factor VIII) to an individual with hemophilia A. Factor VIII is administered prophylactically to prevent or reduce frequency of bleeding episodes, in response to an acute bleeding episode, and/or perioperatively to manage bleeding during surgery. However, there are several undesirable features of Factor VIII replacement therapy.

[0006] First, Factor VIII replacement therapy is used to treat or manage hemophilia A, but does not cure the underlying Factor VIII deficiency. Because of this, individuals with hemophilia A require Factor VIII replacement therapy for the duration of their lives. Continuous treatment is expensive and requires the individual to maintain strict compliance, as missing only a few prophylactic doses can have serious consequences for individuals with severe hemophilia A.

[0007] Second, because Factor VIII has a relatively short half-life in vivo, conventional prophylactic Factor VIII replacement therapy requires administration every second or third day. This places a burden on the individual to maintain compliance throughout their life. While third generation "long-acting" Factor VIII drugs may reduce the frequency of administration, prophylactic Factor FVIII replacement therapy with these drugs still requires monthly, weekly, or more frequent administration in perpetuity. For example, prophylactic treatment with ELOCTATE™ [Antihemophilic Factor (Recombinant), Fc Fusion Protein] requires administration every three to five days (ELOCTATE™ Prescribing Information, Biogen Idec Inc., (2015)). Moreover, the long-term effects of chemically modified biologics (e.g., pegylated polypeptides) are not yet fully understood.

[0008] Third, between 15% and 30% of all individuals receiving Factor VIII replacement therapy form anti-Factor VIII inhibitor antibodies, rendering the therapy inefficient. Factor VIII bypass therapy (e.g., administration of plasma-derived or recombinantly-produced prothrombin complex concentrates) can be used to treat hemophilia in individuals that form inhibitor antibodies. However, Factor VIII bypass therapy is less effective than Factor VIII replacement therapy (Mannucci P.M., J Thromb Haemost., 1(7):1349-55 (2003)) and may be associated with an increased risk of cardiovascular complication (Luu and Ewenstein, Haemophilia, 10 Suppl. 2:10-16 (2004)).

[0009] Somatic gene therapy holds great promise for the treatment of hemophilia A because it would remedy the underlying under-expression functional Factor VIII activity (e.g., due to missense or nonsense mutations), rather than provide a one-time dose of Factor VIII activity to the individual. Because of this difference in the mechanism of action, as compared to Factor VIII replacement therapy, one-time administration of a Factor VIII gene therapy vector may provide an individual with Factor VIII for several years, reducing the cost of treatment and eliminating the need for continued patient compliance.

[0010] Coagulation Factor IX (FIX) gene therapy has been used effectively to treat individuals with hemophilia B, a related blood coagulation condition characterized by diminished Factor IX activity (Manno C.S., et al., Nat Med., 12(3):342-47 (2006)). However, Factor VIII gene therapy presents several unique challenges. For example, the full-length, wild-type Factor VIII polypeptide (2351 amino acids; UniProt accession number P00451) is five times larger than the full-length, wild-type Factor IX polypeptide (461 amino acids; UniProt accession number P00740). As such, the coding sequence of wild-type Factor VIII is 7053 base pairs, which is too large to be packaged in conventional AAV gene therapy vectors. Further, reported recombinant expression of B-domain deleted variants of Factor VIII (BDD-FVIII) has been poor. As such, several groups have attempted to alter the codon usage of BDD-FVIII constructs, with limited success.

BRIEF SUMMARY OF DISCLOSURE

[0011] Accordingly, there is a need for Factor VIII variants whose coding sequences are more efficiently packaged into, and delivered via, gene therapy vectors. There is also a need for synthetic, codon-altered nucleic acids which express Factor VIII more efficiently. Such Factor VIII variants and codon-altered nucleic acids allow for improved treatment of Factor VIII deficiencies (e.g., hemophilia A). The above deficiencies and other problems associated with the treatment of Factor VIII deficiencies (e.g., hemophilia A) are reduced or eliminated by the disclosed codon-altered Factor VIII variants.

[0012] In accordance with some embodiments, the present disclosure provides nucleic acids encoding Factor VIII variants that have high sequence identity to the disclosed codon-altered sequences of the Factor VIII heavy chain (e.g., CS01-HC-NA, CS04-HC-NA, or CS23-HC-NA)

and light chain (CS01-LC-NA, CS04-LC-NA, or CS23-LC-NA). In some embodiments, these nucleic acids further include a sequence encoding a linker sequence that replaces the native Factor VIII B-domain (e.g., a linker sequences comprising a furin cleavage site), between the sequences coding for the Factor VIII heavy and light chains.

[0013] In one aspect, the disclosure provides a polynucleotide including a nucleotide sequence encoding a Factor VIII polypeptide. The Factor VIII polypeptide includes a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS04-HC-NA (SEQ ID NO: 3). The light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS04-LC-NA (SEQ ID NO: 4). The polypeptide linker comprises a furin cleavage site.

[0014] In one embodiment of the polynucleotides described above, the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO04 (SEQ ID NO: 6).

[0015] In one aspect, the disclosure provides a polynucleotide including a nucleotide sequence encoding a Factor VIII polypeptide. The Factor VIII polypeptide includes a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS01-HC-NA (SEQ ID NO: 24). The light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS01-LC-NA (SEQ ID NO: 25). The polypeptide linker comprises a furin cleavage site.

[0016] In one embodiment of the polynucleotides described above, the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO01 (SEQ ID NO: 5).

[0017] In one aspect, the disclosure provides a polynucleotide including a nucleotide sequence encoding a Factor VIII polypeptide. The Factor VIII polypeptide includes a light chain, a heavy

chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS23-HC-NA (SEQ ID NO: 22). The light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS23-LC-NA (SEQ ID NO: 23). The polypeptide linker comprises a furin cleavage site.

[0018] In one embodiment of the polynucleotides described above, the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO23 (SEQ ID NO: 7).

[0019] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 96% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 96% identity to the respective light chain sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0020] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 97% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 97% identity to the respective light chain sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0021] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 98% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 98% identity to the respective light chain

sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0022] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 99% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 99% identity to the respective light chain sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0023] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 99.5% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 99.5% identity to the respective light chain sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0024] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide has at least 99.9% identity to the respective heavy chain sequence (e.g., CS04-HC-NA (SEQ ID NO: 3), CS01-HC-NA (SEQ ID NO: 24), or CS23-HC-NA (SEQ ID NO: 22)), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide has at least 99.9% identity to the respective light chain sequence (e.g., CS04-LC-NA (SEQ ID NO: 4), CS01-LC-NA (SEQ ID NO: 25), or CS23-LC-NA (SEQ ID NO: 23)).

[0025] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide is CS04-HC-NA (SEQ ID NO: 3), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide is CS04-LC-NA (SEQ ID NO: 4).

[0026] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide is CS01-HC-NA (SEQ ID NO: 24), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide is CS01-LC-NA (SEQ ID NO: 25).

[0027] In one embodiment of the polynucleotides described above, the first nucleotide sequence encoding the heavy chain of the Factor VIII polypeptide is CS23-HC-NA (SEQ ID NO: 22), and the second nucleotide sequence encoding the light chain of the Factor FVIII polypeptide is CS23-LC-NA (SEQ ID NO: 23).

[0028] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-FL-NA, wherein the polynucleotide encodes a Factor VIII polypeptide.

[0029] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-FL-NA, wherein the polynucleotide encodes a Factor VIII polypeptide.

[0030] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-FL-NA, wherein the polynucleotide encodes a Factor VIII polypeptide.

[0031] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 96% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0032] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 97% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0033] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 98% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0034] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0035] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.5% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0036] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.9% identity to the respective full-length polynucleotide sequence (e.g., CS04-FL-NA (SEQ ID NO: 1), CS01-FL-NA (SEQ ID NO: 13), or CS23-FL-NA (SEQ ID NO: 20)).

[0037] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS04-FL-NA (SEQ ID NO: 1).

[0038] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS01-FL-NA (SEQ ID NO: 13).

[0039] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS23-FL-NA (SEQ ID NO: 20).

[0040] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 95% identity to CS04-FL-AA (SEQ ID NO: 2).

[0041] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 96% identity to CS04-FL-AA (SEQ ID NO: 2).

[0042] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 97% identity to CS04-FL-AA (SEQ ID NO: 2).

[0043] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 98% identity to CS04-FL-AA (SEQ ID NO: 2).

[0044] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 99% identity to CS04-FL-AA (SEQ ID NO: 2).

[0045] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 99.5% identity to CS04-FL-AA (SEQ ID NO: 2).

[0046] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 99.9% identity to CS04-FL-AA (SEQ ID NO: 2).

[0047] In one embodiment of the polynucleotides described above, the polynucleotide encodes a Factor VIII polypeptide comprising the amino acid sequence of CS04-FL-AA (SEQ ID NO: 2).

[0048] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-SC1-NA (SEQ ID NO: 9), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0049] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-SC2-NA (SEQ ID NO: 11), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0050] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-SC1-NA (SEQ ID NO: 26), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0051] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-SC2-NA (SEQ ID NO: 27), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0052] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-SC1-NA (SEQ ID NO: 28), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0053] In one aspect, the disclosure provides a polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-SC2-NA (SEQ ID NO: 29), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

[0054] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 96% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0055] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 97% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0056] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 98% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0057] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0058] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.5% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0059] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.9% identity to the respective full-length polynucleotide sequence (e.g., CS04-SC1-NA (SEQ ID NO: 9), CS04-SC2-NA (SEQ ID NO: 11), CS01-SC1-NA (SEQ ID NO: 26), CS01-SC2-NA (SEQ ID NO: 27), CS23-SC1-NA (SEQ ID NO: 28), or CS23-SC2-NA (SEQ ID NO: 29)).

[0060] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS04-SC1-NA (SEQ ID NO: 9).

[0061] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS04-SC2-NA (SEQ ID NO: 11).

[0062] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS01-SC1-NA (SEQ ID NO: 26).

[0063] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS01-SC2-NA (SEQ ID NO: 27).

[0064] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS23-SC1-NA (SEQ ID NO: 28).

[0065] In one embodiment of the polynucleotides described above, the nucleotide sequence is CS23-SC2-NA (SEQ ID NO: 29).

[0066] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 95% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA,

CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0067] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 96% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-

NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0068] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 97% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA,

CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0069] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 98% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-

NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0070] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0071] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.5% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0072] In one embodiment of the polynucleotides described above, the nucleotide sequence has at least 99.5% identity to a sequence selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA,

CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0073] In one embodiment of the polynucleotides described above, the nucleotide sequence is selected from the group consisting of CS01-FL-NA, CS01-HC-NA, CS01-LC-NA, CS04-FL-NA, CS04-HC-NA, CS04-LC-NA, CS23-FL-NA, CS23-HC-NA, CS23-LC-NA, CS01m1-FL-NA, CS01m2-FL-NA, CS01m3-FL-NA, CS01m4-FL-NA, CS01m12-FL-NA, CS01m13-FL-NA, CS01m23-FL-NA, CS01m24-FL-NA, CS01m34-FL-NA, CS01m123-FL-NA, CS01m234-FL-NA, CS04m1-FL-NA, CS04m2-FL-NA, CS04m3-FL-NA, CS04m4-FL-NA, CS04m12-FL-NA, CS04m13-FL-NA, CS04m23-FL-NA, CS04m24-FL-NA, CS04m34-FL-NA, CS04m123-FL-NA, CS04m234-FL-NA, CS23m1-FL-NA, CS23m2-FL-NA, CS23m3-FL-NA, CS23m4-FL-

NA, CS23m12-FL-NA, CS23m13-FL-NA, CS23m23-FL-NA, CS23m24-FL-NA, CS23m34-FL-NA, CS23m123-FL-NA, CS23m234-FL-NA, CS01-SC1-NA, CS04-SC1-NA, CS23-SC1-NA, CS01m1-SC1-NA, CS01m2-SC1-NA, CS01m3-SC1-NA, CS01m4-SC1-NA, CS01m12-SC1-NA, CS01m13-SC1-NA, CS01m23-SC1-NA, CS01m24-SC1-NA, CS01m34-SC1-NA, CS01m123-SC1-NA, CS01m234-SC1-NA, CS04m1-SC1-NA, CS04m2-SC1-NA, CS04m3-SC1-NA, CS04m4-SC1-NA, CS04m12-SC1-NA, CS04m13-SC1-NA, CS04m23-SC1-NA, CS04m24-SC1-NA, CS04m34-SC1-NA, CS04m123-SC1-NA, CS04m234-SC1-NA, CS23m1-SC1-NA, CS23m2-SC1-NA, CS23m3-SC1-NA, CS23m4-SC1-NA, CS23m12-SC1-NA, CS23m13-SC1-NA, CS23m23-SC1-NA, CS23m24-SC1-NA, CS23m34-SC1-NA, CS23m123-SC1-NA, CS23m234-SC1-NA, CS01-SC2-NA, CS04-SC2-NA, CS23-SC2-NA, CS01m1-SC2-NA, CS01m2-SC2-NA, CS01m3-SC2-NA, CS01m4-SC2-NA, CS01m12-SC2-NA, CS01m13-SC2-NA, CS01m23-SC2-NA, CS01m24-SC2-NA, CS01m34-SC2-NA, CS01m123-SC2-NA, CS01m234-SC2-NA, CS04m1-SC2-NA, CS04m2-SC2-NA, CS04m3-SC2-NA, CS04m4-SC2-NA, CS04m12-SC2-NA, CS04m13-SC2-NA, CS04m23-SC2-NA, CS04m24-SC2-NA, CS04m34-SC2-NA, CS04m123-SC2-NA, CS04m234-SC2-NA, CS23m1-SC2-NA, CS23m2-SC2-NA, CS23m3-SC2-NA, CS23m4-SC2-NA, CS23m12-SC2-NA, CS23m13-SC2-NA, CS23m23-SC2-NA, CS23m24-SC2-NA, CS23m34-SC2-NA, CS23m123-SC2-NA, and CS23m234-SC2-NA.

[0074] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide comprises a glycosylation polypeptide positioned between two consecutive amino acids.

[0075] In one embodiment of the polynucleotides described above, the encoded polypeptide linker includes a glycosylation peptide with an amino acid sequence having at least 92% identity to a glycosylation peptide selected from the group consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

[0076] In one embodiment of the polynucleotides described above, the encoded polypeptide linker comprises a glycosylation peptide with an amino acid sequence selected from the group

consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

[0077] In one embodiment of the polynucleotides described above, the glycosylation peptide is encoded by a polynucleotide with a nucleotide sequence having at least 95% identity to a sequence selected from the group consisting of NG1-NA, NG4-NA, NG5-NA, NG6-NA, NG7-NA, NG9-NA, NG10-NA, NG16-NA, NG17-NA, NG18-NA, NG19-NA, NG20-NA, NG21-NA and NGV-NA.

[0078] In one embodiment of the polynucleotides described above, the glycosylation peptide is encoded by a polynucleotide with a nucleotide sequence selected from one of NG1-NA, NG4-NA, NG5-NA, NG6-NA, NG7-NA, NG9-NA, NG10-NA, NG16-NA, NG17-NA, NG18-NA, NG19-NA, NG20-NA, NG21-NA and NGV-NA.

[0079] In one embodiment of the polynucleotides described above, the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to a sequence selected from the group consisting of BDLNG1-NA, BDLNG3-NA, BDLNG5-NA, BDLNG6-NA, BDLNG9-NA, BDLNG10-NA, BDLNG16-NA, BDLNG17-NA, BDLNG18-NA, BDLNG19-NA, BDLNG20-NA and BDLNG21-NA.

[0080] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes an F328S (SPI, F309S SPE) amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0081] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes I105V, A127S, G151K, M166T, and L171P (SPI; I86V, A108S, G132K, M147T, and L152P, SPE, respectively) amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0082] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and b) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19). In some embodiments (e.g., where the encoded FVIII molecule includes a portion of the N-terminal region of the wild-type B-domain), the

encoded Factor VIII polypeptide also includes a deletion of amino acids SF760-761, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0083] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) an F328S (SPI; F309S SPE) amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), and b) C1918G and C1922G (SPI; C1899G and C1903 SPE, respectively) amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0084] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) an F328S (SPI; F309S SPE) amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), and b) I105V, A127S, G151K, M166T, and L171P (SPI; I86V, A108S, G132K, M147T, and L152P, SPE, respectively) amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0085] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), b) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and c) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19). In some embodiments (e.g., where the encoded FVIII molecule includes a portion of the N-terminal region of the wild-type B-domain), the encoded Factor VIII polypeptide also includes a deletion of amino acids SF760-761, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0086] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19), b) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and c) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19). In some embodiments (e.g., where the encoded FVIII molecule includes a portion of the N-terminal region of the wild-type B-domain), the encoded Factor VIII polypeptide also includes a deletion of amino acids SF760-761, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0087] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), b) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19), and c) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0088] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), b) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19), c) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and d) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19). In some embodiments (e.g., where the encoded FVIII molecule includes a portion of the N-terminal region of the wild-type B-domain), the encoded Factor VIII polypeptide also includes a deletion of amino acids SF760-761, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0089] In one embodiment of the polynucleotides described above, the encoded Factor VIII polypeptide includes a) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19), b) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19), c) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19), d) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and e) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19). In some embodiments (e.g., where the encoded FVIII molecule includes a portion of the N-terminal region of the wild-type B-domain), the encoded Factor VIII polypeptide also includes a deletion of amino acids SF760-761, relative to FVIII-FL-AA (SEQ ID NO: 19).

[0090] In one embodiment of the polynucleotides described above, the polynucleotide also includes a promoter element operably linked to the polynucleotide encoding the Factor VIII polypeptide.

[0091] In one embodiment of the polynucleotides described above, the polynucleotide also includes an enhancer element operably linked to the polynucleotide encoding the Factor VIII polypeptide.

[0092] In one embodiment of the polynucleotides described above, the polynucleotide also includes a polyadenylation element operably linked to the polynucleotide encoding the Factor VIII polypeptide.

[0093] In one embodiment of the polynucleotides described above, the polynucleotide also includes an intron operatively linked to the nucleotide sequence encoding the Factor VIII polypeptide.

[0094] In one embodiment of the polynucleotides described above, the intron is positioned between a promoter element and the translation initiation site (e.g., the first coding ATG) of the nucleotide sequence encoding a Factor VIII polypeptide.

[0095] In another aspect, the disclosure provides a mammalian gene therapy vector including a polynucleotide as described above.

[0096] In one embodiment of the mammalian gene therapy vector described above, the mammalian gene therapy vector is an adeno-associated virus (AAV) vector.

[0097] In one embodiment of the mammalian gene therapy vector described above, the AAV vector is an AAV-8 vector.

[0098] In another aspect, the disclosure provides a method for treating hemophilia A including administering, to a patient in need thereof, a mammalian gene therapy vector as described above.

[0099] In another aspect, the disclosure provides a mammalian gene therapy vector as described above for treating hemophilia A.

[00100] In another aspect, the disclosure provides the use of a mammalian gene therapy vector as described above for the manufacture of a medicament for treating hemophilia A.

[00101] In another aspect, the disclosure provides a Factor VIII polypeptide including a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to

the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-HC-AAm23. The light chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-LC-AAm23. The polypeptide linker of the Factor VIII polypeptide includes a furin cleavage site. The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

[00102] In another aspect, the disclosure provides a Factor VIII polypeptide including a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-HC-AAm123. The light chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-LC-AAm123. The polypeptide linker of the Factor VIII polypeptide includes a furin cleavage site. The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19), and iv) an F328S amino acid substitution.

[00103] In another aspect, the disclosure provides a Factor VIII polypeptide including a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-HC-AAm234. The light chain of the Factor VIII polypeptide has a sequence at least 95% identical to the sequence CS01-LC-AAm234. The polypeptide linker of the Factor VIII polypeptide includes a furin cleavage site. The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19), and iv) F328S/C1918G/C1922G amino acid substitutions.

[00104] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence at least 96% identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the Factor FVIII polypeptide has a sequence at least 96% identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00105] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence at least 97% identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the Factor FVIII polypeptide has a sequence at least 97% identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00106] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence at least 98% identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the Factor FVIII polypeptide has a sequence at least 98% identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00107] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence at least 99% identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the Factor FVIII polypeptide has a sequence at least 99% identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00108] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence at least 99.5% identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the Factor FVIII polypeptide has a sequence at least 99.5% identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00109] In one embodiment of the Factor VIII polypeptides described, the heavy chain of the Factor VIII polypeptide has a sequence identical to the respective heavy chain sequence (e.g., CS01-HC-AAm23, CS01-HC-AAm123, or CS01-HC-AAm234), and the light chain of the

Factor FVIII polypeptide has a sequence identical to the respective light chain sequence (e.g., CS01-LC-AAm23, CS01-LC-AAm123, or CS01-LC-AAm234).

[00110] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker has at least 95% identity to BDL-SQ-AA (SEQ ID NO: 30).

[00111] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker has the amino acid sequence of BDL-SQ-AA (SEQ ID NO: 30).

[00112] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker includes a glycosylation peptide with an amino acid sequence having at least 92% identity to a glycosylation peptide selected from the group consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

[00113] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker includes a glycosylation peptide selected from the group consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

[00114] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker has an amino acid sequence having at least 95% identity to a sequence selected from the group consisting of BDLNG1-AA, BDLNG3-AA, BDLNG5-AA, BDLNG6-AA, BDLNG9-AA, BDLNG10-AA, BDLNG16-AA, BDLNG17-AA, BDLNG18-AA, BDLNG19-AA, BDLNG20-AA and BDLNG21-AA.

[00115] In one embodiment of the Factor VIII polypeptides described above, the polypeptide linker has an amino acid sequence selected from the group consisting of BDLNG1-AA, BDLNG3-AA, BDLNG5-AA, BDLNG6-AA, BDLNG9-AA, BDLNG10-NA, BDLNG16-AA, BDLNG17-AA, BDLNG18-AA, BDLNG19-AA, BDLNG20-AA and BDLNG21-AA.

[00116] In another aspect, the disclosure provides a Factor VIII polypeptide having an amino acid sequence with at least 95% identity to CS40-FL-AAm23 (SEQ ID NO: 104). The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid

substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), and iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

[00117] In another aspect, the disclosure provides a Factor VIII polypeptide having an amino acid sequence with at least 95% identity to CS40-FL-AAm123. The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19), and iv) an F328S amino acid substitution.

[00118] In another aspect, the disclosure provides a Factor VIII polypeptide having an amino acid sequence with at least 95% identity to CS40-FL-AAm234. The Factor VIII polypeptide includes i) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, ii) a deletion of amino acids AIEPR755-759, relative to FVIII-FL-AA (SEQ ID NO: 19), iii) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19), and iv) F328S/C1918G/C1922G amino acid substitutions.

[00119] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence at least 96% identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

[00120] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence at least 97% identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

[00121] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence at least 98% identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

[00122] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence at least 99% identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

[00123] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence at least 99.5% identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

[00124] In one embodiment of the Factor VIII polypeptides described, the Factor VIII polypeptide has a sequence identical to the respective full-length sequence (e.g., CS40-FL-AAm23 (SEQ ID NO: 104), CS40-FL-AAm123, or CS40-FL-AAm234).

BRIEF DESCRIPTION OF DRAWINGS

[00125] Figure 1 shows schematic illustrations of the wild-type and ReFacto-type human Factor VIII protein constructs.

[00126] Figures 2A and 2B show the CS04 codon-altered nucleotide sequence (SEQ ID NO: 1) encoding a Factor VIII variant in accordance with some embodiments ("CS04-FL-NA" for full-length coding sequence).

[00127] Figure 3 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 2) encoded by the CS04 codon-altered nucleotide sequence in accordance with some embodiments ("CS04-FL-AA" for full-length amino acid sequence).

[00128] Figure 4 shows the portion of the CS04 codon-altered nucleotide sequence (SEQ ID NO: 3) encoding the heavy chain of a Factor VIII variant in accordance with some embodiments ("CS04-HC-NA").

[00129] Figure 5 shows the portion of the CS04 codon-altered nucleotide sequence (SEQ ID NO: 4) encoding the light chain of a Factor VIII variant in accordance with some embodiments ("CS04-LC-NA").

[00130] Figure 6 shows exemplary coding sequences (SEQ ID NOS 5-7 and 36-48, respectively, in order of appearance) for B-domain substituted linkers in accordance with some embodiments. BDLO01 (SEQ ID NO: 5), BDLO04 (SEQ ID NO: 6), and BDLO23 (SEQ ID NO: 7) are the respective portions of the CS01, CS04, and CS23 codon-altered nucleotide sequences that encode a B-domain substituted linker, respectively.

[00131] Figures 7A, 7B, and 7C show an AAV vector sequence (SEQ ID NO: 8) containing an CS04 codon-altered nucleotide sequence in accordance with some embodiments ("CS04-AV-NA").

[00132] Figures 8A and 8B show the CS01m1 codon-altered nucleotide sequence (SEQ ID NO: 49) encoding a Factor VIII variant with an F328S amino acid substitution in accordance with some embodiments ("CS01m1-FL-NA").

[00133] Figures 9A and 9B show the CS04Δ(760-1667) (SPI; CS04Δ(741-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 9) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS04-SC1-NA").

[00134] Figure 10 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 10) encoded by the CS01Δ(760-1667) (SPI; CS01Δ(741-1648), SPE), CS04Δ(760-1667) (SPI; CS04Δ(741-1648), SPE), and CS23Δ(760-1667) (SPI; CS23Δ(741-1648), SPE) codon-altered nucleotide sequences in accordance with some embodiments ("CS01-SC1-AA," "CS04-SC1-AA," and "CS23-SC1-AA," respectively).

[00135] Figures 11A and 11B show the CS04Δ(772-1667) (SPI; CS04Δ (753-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 11) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS04-SC2-NA").

[00136] Figure 12 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 12) encoded by the CS01Δ(772-1667) (SPI; CS01Δ(753-1648), SPE), CS04Δ(772-1667) (SPI; CS04Δ(753-1648), SPE), and CS23Δ(772-1667) (SPI; CS23Δ(753-1648), SPE) codon-altered nucleotide sequence in accordance with some embodiments ("CS01-SC2-AA," "CS04-SC2-AA," and "CS23-SC2-AA," respectively).

[00137] Figure 13A and 13B show amino acid and nucleotide sequences for exemplary glycosylation peptides that are inserted into the B-domain substituted linker in accordance with some embodiments. "NG1" or NG1-AA" is the code for the amino acid sequence, shown in the top line. "NG1-NA" is the code for the nucleic acid sequence, shown in the bottom line for each set. Figure 13A and 13B disclose the amino acid sequences as SEQ ID NOS 51, 53, 55, 57, 59,

61, 63, 65, 67, 69, 71, 73, and 75, and the nucleotide sequences as SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, all respectively, in order of appearance.

[00138] Figure 14 shows the results of in silico prediction of in vivo N-glycosylation of the wild-type Factor VIII B-domain. Figure discloses SEQ ID NOS 76 and 76-82, respectively, in order of appearance.

[00139] Figure 15 shows the results of in silico prediction of in vivo N-glycosylation of the V3 peptide linker. Figure discloses SEQ ID NOS 83 and 83-89, respectively, in order of appearance.

[00140] Figures 16A and 16B show the CS01 codon-altered nucleotide sequence (SEQ ID NO: 13) encoding a Factor VIII variant in accordance with some embodiments ("CS01-FL-NA").

[00141] Figures 17A and 17B show the CS08 codon-altered nucleotide sequence (SEQ ID NO: 14) encoding a Factor VIII variant in accordance with some embodiments ("CS08-FL-NA").

[00142] Figures 18A and 18B show the CS10 codon-altered nucleotide sequence (SEQ ID NO: 15) encoding a Factor VIII variant in accordance with some embodiments ("CS10-FL-NA").

[00143] Figures 19A and 19B show the CS11 codon-altered nucleotide sequence (SEQ ID NO: 16) encoding a Factor VIII variant in accordance with some embodiments ("CS11-FL-NA").

[00144] Figures 20A and 20B show the CS40 wild-type ReFacto coding sequence (SEQ ID NO: 17), in accordance with some embodiments ("CS40-FL-NA").

[00145] Figures 21A and 21B show the CH25 codon-altered nucleotide sequence (SEQ ID NO: 18) encoding a Factor VIII variant in accordance with some embodiments ("CH25-FL-NA").

[00146] Figure 22 shows a wild-type human Factor VIII amino acid sequence (SEQ ID NO: 19), in accordance with some embodiments ("FVIII-FL-AA").

[00147] Figure 23 illustrates the scheme for cloning the pCS40, pCS01, pCS04, pCS08, pCS10, pCS11, and pCh25 constructs, by inserting synthetic Refacto-type BDD-FVIII DNA sequences into the vector backbone pCh-BB01 via AscI and NotI restriction sites.

[00148] Figure 24 shows the integrity of AAV vector genome preparations, as analyzed by agarose gel electrophoresis. Lane 1, DNA marker; lane 2, vCS40; lane 3, vCS01; lane 4, vCS04. The AAV vectors have all the same-sized genomes, migrating at approximately 5 kb (arrow, right side). The scale on the left side indicates size of the DNA fragments in kilobases (kb).

[00149] Figure 25 shows the protein analysis of AAV vector preparations by PAGE and silver staining. Lane 1, protein marker (M); lane 2, vCS40, lane 3, vCS01; and lane 4, vCS04. The constructs all have the same AAV8 capsids consisting of VP1, VP2, and VP3 (arrows right side). The scale on the left side indicates size of the protein marker in kilodaltons (kDa).

[00150] Figure 26A and 26B show the CS23 codon-altered nucleotide sequence (SEQ ID NO: 20) encoding a Factor VIII variant in accordance with some embodiments ("CS23-FL-NA").

[00151] Figure 27 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 21) encoded by the CS23 codon-altered nucleotide sequence in accordance with some embodiments ("CS23-FL-AA").

[00152] Figure 28 shows the portion of the CS23 codon-altered nucleotide sequence (SEQ ID NO: 22) encoding the heavy chain of a Factor VIII variant in accordance with some embodiments ("CS23-HC-NA").

[00153] Figure 29 shows the portion of the CS23 codon-altered nucleotide sequence (SEQ ID NO: 23) encoding the light chain of a Factor VIII variant in accordance with some embodiments("CS23-LC-NA").

[00154] Figures 30A and 30B show the CS01m13 codon-altered nucleotide sequence (SEQ ID NO: 90) encoding a Factor VIII variant with m1 (F328S) and m3 amino acid substitutions in accordance with some embodiments ("CS01-FL-NA-m13").

[00155] Figures 31A and 31B show the CS01m23 codon-altered nucleotide sequence (SEQ ID NO: 91) encoding a Factor VIII variant with the m2 and m3 mutation sets in accordance with some embodiments ("CS01-FL-NA-m23").

[00156] Figures 32A and 32B show the CS01m3 codon-altered nucleotide sequence (SEQ ID NO: 92) encoding a Factor VIII variant with m3 amino acid substitutions in accordance with some embodiments ("CS01-FL-NA-m3").

[00157] Figures 33A and 33B show the CS01m2 codon-altered nucleotide sequence (SEQ ID NO: 93) encoding a Factor VIII variant with the m2 mutation set (I105V/A127S/G151K/M166T/L171P (SPI)) amino acid substitutions in accordance with some embodiments ("CS01-FL-NA-m2").

[00158] Figures 34A and 34B show the CS04m2 codon-altered nucleotide sequence (SEQ ID NO: 94) encoding a Factor VIII variant with the m2 mutants (I105V/A127S/G151K/M166T/L171P (SPI)) amino acid substitutions in accordance with some embodiments ("CS01-FL-NA-m2").

[00159] Figures 35A and 35B show the CS04m3 codon-altered nucleotide sequence (SEQ ID NO: 95) encoding a Factor VIII variant with m3 amino acid substitutions in accordance with some embodiments ("CS04-FL-NA-m3").

[00160] Figures 36A and 36B show the CS04m23 codon-altered nucleotide sequence (SEQ ID NO: 96) encoding a Factor VIII variant with the m2 mutant set (I105V/A127S/G151K/M166T/L171P (SPI)) and m3 amino acid substitutions in accordance with some embodiments ("CS04-FL-NA-m23").

[00161] Figures 37A and 37B show the CS04m1 codon-altered nucleotide sequence (SEQ ID NO: 97) encoding a Factor VIII variant with an m1 (F328S) amino acid substitution in accordance with some embodiments ("CS04-FL-NA-m1").

[00162] Figures 38A and 38B show the CS04m13 codon-altered nucleotide sequence (SEQ ID NO: 98) encoding a Factor VIII variant with m1 and m3 amino acid substitutions in accordance with some embodiments ("CS04-FL-NA-m13")

[00163] Figures 39A and 39B show the CS23m13 codon-altered nucleotide sequence (SEQ ID NO: 99) encoding a Factor VIII variant with m1 and m3 amino acid substitutions in accordance with some embodiments ("CS23m13-FL-NA")

[00164] Figures 40A and 40B show the CS23m3 codon-altered nucleotide sequence (SEQ ID NO: 100) encoding a Factor VIII variant with m3 amino acid substitutions in accordance with some embodiments ("CS23-FL-NA-m3")

[00165] Figures 41A and 41B show the CS23m2 codon-altered nucleotide sequence (SEQ ID NO: 101) encoding a Factor VIII variant with the m2 mutant set (I105V/A127S/G151K/M166T/L171P amino acid substitutions) in accordance with some embodiments ("CS23-FL-NA-m2").

[00166] Figures 42A and 42B show the CS23m1 codon-altered nucleotide sequence (SEQ ID NO: 102) encoding a Factor VIII variant with an m1 (F328S) amino acid substitution in accordance with some embodiments ("CS23-FL-NA-m1").

[00167] Figures 43A and 43B show the CS23m23 codon-altered nucleotide sequence (SEQ ID NO: 103) encoding a Factor VIII variant with the m2 mutant set (I105V/A127S/G151K/M166T/L171P) and m3 amino acid substitutions in accordance with some embodiments ("CS23-FL-NA-m23").

[00168] Figure 44 depicts cloning of the pCS constructs, done by inserting synthetic Refacto-type BDD-FVIII carrying different mutations (see inserted table) into the vector backbone pCh-BB01 via AscI and NotI restriction sites.

[00169] Figure 45 depicts the protein analysis of AAV vector preparations by PAGE and silver staining. Lane 1, protein marker (M); lane 2, vCS01, lane 3, vCS17; lane 4, vCS19; lane 5, vCS20; lane 6, vCS40; lane 7, vCS04; lane 8, vCS17; lane 9, vCS24 construct. The constructs have all the same AAV8 capsids consisting of VP1, VP2 and VP3 (arrows right side). The scale on the left side indicates size of the protein marker in kilo Daltons (kDa).

[00170] Figure 46 depicts the integrity of AAV vector genome preparations analyzed by agarose gel electrophoresis. Lane 1, DNA marker (M); lane 2, vCS04, lane 3, vCS17; lane 4,

vCS20; lane 5, vCS24; lane 6, vCS16; lane 7, vCS40 construct. Vector load is 1.5E10 vg per lane. The AAV vectors have the same-sized genomes, migrating at approximately 5 kb (arrow, right side). The scale on the left side indicates size of the DNA fragments in kilobases (kb).

[00171] Figure 47 shows the portion of the CS01 codon-altered nucleotide sequence (SEQ ID NO: 24) encoding the heavy chain of a Factor VIII variant in accordance with some embodiments ("CS01-HC-NA").

[00172] Figure 48 shows the portion of the CS01 codon-altered nucleotide sequence (SEQ ID NO: 25) encoding the light chain of a Factor VIII variant in accordance with some embodiments("CS01-LC-NA").

[00173] Figures 49A and 49B show the CS01Δ(760-1667) (SPI; CS01Δ(741-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 26) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS01-SC1-NA").

[00174] Figures 50A and 50B show the CS01Δ(772-1667) (SPI; CS01Δ(753-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 27) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS01-SC2-NA").

[00175] Figures 51A and 51B show the CS23Δ(760-1667) (SPI; CS23Δ(741-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 28) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS23-SC1-NA").

[00176] Figures 52A and 52B show the CS23Δ(772-1667) (SPI; CS23Δ (753-1648), SPE) codon-altered nucleotide sequence (SEQ ID NO: 29) encoding a single-chain Factor VIII variant in accordance with some embodiments ("CS23-SC2-NA").

[00177] Figure 53 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 104) encoded by the CS01m23 codon-altered nucleotide sequence in accordance with some embodiments ("CS01m23-FL-AA").

[00178] Figure 54 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 105) encoded by the CS04m3 codon-altered nucleotide sequence in accordance with some embodiments ("CS01m23-FL-AA").

[00179] Figure 55 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 106) encoded by the CS01m12 codon-altered nucleotide sequence in accordance with some embodiments ("CS01m12-FL-AA").

[00180] Figure 56 shows the Factor VIII variant amino acid sequence (SEQ ID NO: 107) encoded by the CS04m12 codon-altered nucleotide sequence in accordance with some embodiments ("CS04m12-FL-AA").

[00181] Figures 57A and 57B show the CS01m12 codon-altered nucleotide sequence (SEQ ID NO: 108) encoding a Factor VIII variant with m1 (F328S) and m2 amino acid substitutions in accordance with some embodiments ("CS01-FL-NAm12").

[00182] Figures 57A and 57B show the CS04m12 codon-altered nucleotide sequence (SEQ ID NO: 109) encoding a Factor VIII variant with m1 (F328S) and m2 amino acid substitutions in accordance with some embodiments ("CS04-FL-NAm12").

DETAILED DESCRIPTION OF DISCLOSURE

I. Introduction

[00183] AAV-based gene therapy holds great promise for the treatment of hemophiliacs. For hemophilia B, first clinical data are encouraging in that FIX levels of about 10% can be maintained in at least some patients for more than 1 year. For hemophilia A however, achieving therapeutic expression levels of 5-10% with AAV vectors remains challenging for various reasons. First, the Factor VIII coding sequence is too large for conventional AAV-based vectors. Second, engineered B-domain deleted or truncated Factor VIII constructs suffer from poor expression in vivo, even when codon-optimized. Third, these B-domain deleted or truncated Factor VIII variant constructs have short half-lives in vivo, exacerbating the effects of poor expression. Fourth, even when expressed, FVIII is not efficiently secreted from cells, as are other coagulation factors, such as Factor IX.

[00184] Moreover, these challenges cannot be addressed by simply administering higher doses of the gene therapy construct. According to current knowledge, the vector dose of an AAV-based gene therapy vector should be increased above $2x10^{12}$ vg/kg bodyweight. This is

because at such high doses a T cell immune response is triggered, which destroys transduced cells and, as a consequence, transgene expression is reduced or even eliminated. Therefore, strategies to improve the expression of FVIII are needed to make FVIII gene therapy a viable therapeutic option for hemophilia A patients.

[00185] The present disclosure relates to the discovery of codon-altered Factor VIII variant coding sequences that solve these and other problems associated with Factor VIII gene therapy. For example, the polynucleotides disclosed herein provide markedly improved expression in mammalian cells, and display improved virion packaging due to stabilized packing interactions. In some implementations, these advantages are realized by using coding sequences for the heavy and light chains of Factor VIII with high sequence identity to the codon altered CS01, CS04, and CS23 constructs (e.g., with high sequence identity to one of the CS01-HC, CS04-HC, and CS23-HC heavy chain coding sequences and high sequence identity to one of the CS01-LC, CS04-LC, and CS23-LC light chain coding sequences).

[00186] In some implementations, the Factor VIII molecules encoded by the polynucleotides described herein have been shortened by truncating, deleting, or replacing the wild-type B-domain. As such, the polynucleotides are better suited for expressing Factor VIII via conventional gene therapy vectors, which inefficiently express larger polypeptides, such as the wild-type Factor VIII.

[00187] Advantageously, it is shown herein that the CS01, CS04, and CS23 codon-altered Factor VIII variant coding sequences provide superior expression of a B-domain deleted Factor VIII construct in vivo. For example, it is demonstrated in Example 2 and Example 4 that intravenous administration of AAV-based gene therapy vectors having the CS01 (SEQ ID NO: 13), CS04 (SEQ ID NO: 1), and CS23 (SEQ ID NO: 20) coding sequence provide 18-fold, 74-fold, and 30-fold increases in Factor VIII expression, relative to the corresponding CS40 construct encoded with the wild-type polynucleotide sequence (SEQ ID NO: 17), in Factor VIII knock-out mice (**Table 4** and **Table 7**).

[00188] Further, it also shown herein that the CS01 and CS04 codon-altered Factor VIII variant coding sequences provide superior virion packaging and virus production. For example, it is demonstrated in Example 1 that AAV vector constructs containing the CS01 and CS04

constructs provided 5 to 7-fold greater viral yield, relative to the corresponding CS40 construct encoded with the wild-type polynucleotide sequence, when isolated from the same amount of cell pellet.

[00189] Advantageously, Applicants also found that the improved Factor VIII activity generated from the CS01, CS04, and CS23 codon altered sequences could be further enhanced by introducing mutations into the underlying Factor VIII polypeptide sequence. For example, as demonstrated in Example 4, the F328S, X5, and X1 mutations, alone and in combination with one another, further increased FVIII activity when expressed in vivo in the CS01 or CS04 codon altered background 2 to 7-fold, relative to the wild type, codon altered constructs (**Table 7**). More strikingly, these codon altered sequences, encoding the mutant Factor VIII mutants, provided up to 246-fold greater increase as compared to the corresponding CS40 construct encoded with the wild-type polynucleotide sequence (**Table 7**).

II. Definitions

[00190] As used herein, the following terms have the meanings ascribed to them unless specified otherwise.

[00191] As used herein, the terms "Factor VIII" and "FVIII" are used interchangeably, and refer to any protein with Factor VIII activity (e.g., active FVIII, often referred to as FVIIIa) or protein precursor (e.g., pro-protein or pre-pro-protein) of a protein with Factor VIII activity, particularly Factor IXa cofactor activity. In an exemplary embodiment, a Factor VIII polypeptide refers to a polypeptide that has sequences with high sequence identity (e.g., at least 70%, 75%, 80%, 85%, 90%, 95%, 99%, or more) to the heavy and light chains of a wild type Factor VIII polypeptide. In some embodiments, the B-domain of a Factor VIII polypeptide is deleted, truncated, or replaced with a linker polypeptide to reduce the size of the polynucleotide encoding the Factor VIII polypeptide. In an exemplary embodiment, amino acids 20-1457 of SEQ ID NO: 2 constitute a Factor VIII polypeptide.

[00192] Non-limiting examples of wild type Factor VIII polypeptides include human pre-pro-Factor VIII (e.g., GenBank accession nos. AAA52485, CAA25619, AAA58466, AAA52484, AAA52420, AAV85964, BAF82636, BAG36452, CAI41660, CAI41666,

CAI41672, CAI43241, CAO03404, EAW72645, AAH22513, AAH64380, AAH98389, AAI11968, AAI11970, or AAB61261), corresponding pro-Factor VIII, and natural variants thereof; porcine pre-pro-Factor VIII (e.g., UniProt accession nos. F1RZ36 or K7GSZ5), corresponding pro-Factor VIII, and natural variants thereof; mouse pre-pro-Factor VIII (e.g., GenBank accession nos. AAA37385, CAM15581, CAM26492, or EDL29229), corresponding pro-Factor VIII, and natural variants thereof; rat pre-pro-Factor VIII (e.g., GenBank accession no. AAQ21580), corresponding pro-Factor VIII, and natural variants thereof; rat pre-pro-Factor VIII; and other mammalian Factor VIII homologues (e.g., monkey, ape, hamster, guinea pig, etc.).

[00193] As used herein, a Factor VIII polypeptide includes natural variants and artificial constructs with Factor IX cofactor activity. As used in the present disclosure, Factor VIII encompasses any natural variants, alternative sequences, isoforms, or mutant proteins that retain some basal Factor IX cofactor activity (e.g., at least 5%, 10%, 25%, 50%, 75%, or more of the corresponding wild type activity). Examples of Factor VIII amino acid variations (relative to FVIII-FL-AA (SEQ ID NO: 19)) found in the human population include, without limitation, S19R, R22T, Y24C, Y25C, L26P/R, E30V, W33G, Y35C/H, G41C, R48C/K, K67E/N, L69P, E72K, D75E/V/Y, P83R, G89D/V, G92A/V, A97P, E98K, V99D, D101G/H/V, V104D, K108T, M110V, A111T/V, H113R/Y, L117F/R, G121S, E129V, G130R, E132D, Y133C, D135G/Y, T137A/I, S138R, E141K, D145H, V147D, Y155H, V159A, N163K, G164D/V, P165S, C172W, S176P, S179P, V181E/M, K185T, D186G/N/Y, S189L, L191F, G193R, L195P, C198G, S202N/R, F214V, L217H, A219D/T, V220G, D222V, E223K, G224W, T252I, V253F, N254I, G255V, L261P, P262L, G263S, G266F, C267Y, W274C, H275L, G278R, G280D, E284K, V285G, E291G/K, T294I, F295L, V297A, N299I, R301C/H/L, A303E/P, I307S, S308L, F312S, T314A/I, A315V, G323E, L326P, L327P/V, C329F, I331V, M339T, E340K, V345A/L, C348R/S/Y, Y365C, R391C/H/P, S392L/P, A394S, W401G, I405F/S, E409G, W412G/R, K427I, L431F/S, R437P/W, I438F, G439D/S/V, Y442C, K444R, Y450D/N, T454I, F455C, G466E, P470L/R/T, G474E/R/V, E475K, G477V, D478N, T479R, F484C, A488G, R490G, Y492C/H, Y492H, I494T, P496R, G498R, R503H, G513S/V, I522Y, K529E, W532G, P540T, T541S, D544N, R546W, R550C/G/H, S553P, S554C/G, V556D, R560T, D561G/H/Y, I567T, P569R, S577F, V578A, D579A/H, N583S, Q584H/K/R, I585R/T, M586V, D588G/Y, L594Q,

S596P, N601D/K, R602G, S603I/R, W604C, Y605H/S, N609I, R612C, N631K/S, M633I, S635N, N637D/I/S, Y639C, L644V, L650F, V653A/M, L659P, A663V, Q664P, F677L, M681I, V682F, Y683C/N, T686R, F698L, M699T/V, M701I, G705V, G710W, N713I, R717L/W, G720D/S, M721I/L, A723T, L725Q, V727F, E739K, Y742C, R795G, P947R, V1012L, E1057K, H1066Y, D1260E, K1289Q, Q1336K, N1460K, L1481P, A1610S, I1698T, Y1699C/F, E1701K, Q1705H, R1708C/H, T1714S, R1715G, A1720V, E1723K, D1727V, Y1728C, R1740G, K1751Q, F1762L, R1768H, G1769R, L1771P, L1775F/V, L1777P, G1779E/R, P1780L, I1782R, D1788H, M1791T, A1798P, S1799H, R1800C/G/H, P1801A, Y1802C, S1803Y, F1804S, L1808F, M1842I, P1844S, T1845P, E1848G, A1853T/V, S1858C, K1864E, D1865N/Y, H1867P/R, G1869D/V, G1872E, P1873R, L1875P, V1876L, C1877R/Y, L1882P, R1888I, E1894G, I1901F, E1904D/K, S1907C/R, W1908L, Y1909C, A1939T/V, N1941D/S, G1942A, M1945V, L1951F, R1960L/Q, L1963P, S1965I, M1966I/V, G1967D, S1968R, N1971T, H1973L, G1979V, H1980P/Y, F1982I, R1985Q, L1994P, Y1998C, G2000A, T2004R, M2007I, G2013R, W2015C, R2016P/W, E2018G, G2022D, G2028R, S2030N, V2035A, Y2036C, N2038S, 2040Y, G2045E/V, I2051S, I2056N, A2058P, W2065R, P2067L, A2070V, S2082N, S2088F, D2093G/Y, H2101D, T2105N, Q2106E/P/R, G2107S, R2109C, I2117F/S, Q2119R, F2120C/L, Y2124C, R2135P, S2138Y, T2141N, M2143V, F2145C, N2148S, N2157D, P2162L, R2169C/H, P2172L/Q/R, T2173A/I, H2174D, R2178C/H/L, R2182C/H/P, M2183R/V, L2185S/W, S2192I, C2193G, P2196R, G2198V, E2200D, I2204T, I2209N, A2211P, A2220P, P2224L, R2228G/L/P/Q, L2229F, V2242M, W2248C/S, V2251A/E, M2257V, T2264A, Q2265R, F2279C/I, I2281T, D2286G, W2290L, G2304V, D2307A, P2319L/S, R2323C/G/H/L, R2326G/L/P/Q, Q2330P, W2332R, I2336F, R2339T, G2344C/D/S, and C2345S/Y. Factor VIII proteins also include polypeptides containing post-translational modifications.

[00194] Generally, polynucleotides encoding Factor VIII encode for an inactive single-chain polypeptide (e.g., a pre-pro-protein) that undergoes post-translational processing to form an active Factor VIII protein (e.g., FVIIIa). For example, referring to Figure 1, the wild type human Factor VIII pre-pro-protein is first cleaved to release the encoded signal peptide (not shown), forming a first single-chain pro-protein (shown as "human wild-type FVIII). The pro-protein is then cleaved between the B and A3 domains to form a first polypeptide that includes

the Factor VIII heavy chain (e.g., the A1 and A2 domains) and B-domain, and a second polypeptide that includes the Factor VIII light chain (e.g., including the A3, C1, and C3 domains). The first polypeptide is further cleaved to remove the B-domain, and also to separate the A1 and A2 domains, which remain associated with the Factor VIII light chain in the mature Factor VIIIa protein. For review of the Factor VIII maturation process, see Graw et al., Nat Rev Genet., 6(6):488-501 (2005), the content of which is incorporated herein by reference in its entirety for all purposes.

[00195] However, in some embodiments, the Factor VIII polypeptide is a single-chain Factor VIII polypeptide. Single-chain Factor VIII polypeptides are engineered to remove natural cleavage sites, and optionally remove, truncate, or replace the B-domain of Factor VIII. As such, they are not matured by cleavage (other than cleavage of an optional signal and/or leader peptide), and are active as a single chain. Non-limiting examples of single-chain Factor VIII polypeptides are described in Zollner et al. (Thromb Res, 134(1):125-31 (2014)) and Donath et al. (Biochem J., 312(1):49-55 (1995)), the disclosures of which are hereby incorporated by reference in their entireties for all purposes.

[00196] As used herein, the terms "Factor VIII heavy chain," or simply "heavy chain," refers to the aggregate of the A1 and A2 domains of a Factor VIII polypeptide. In an exemplary embodiment, amino acids 20-759 of CS04-FL-AA (SEQ ID NO: 2) constitute a Factor VIII heavy chain.

[00197] As used herein, the term "Factor VIII light chain," or simply "light chain," refers to the aggregate of the A3, C1, and C2 domains of a Factor VIII polypeptide. In an exemplary embodiment, amino acids 774-1457 CS04-FL-AA (SEQ ID NO: 2) constitute a Factor VIII light chain. In some embodiments, a Factor VIII light chain excludes the acidic a3 peptide, which is released during maturation in vivo.

[00198] Generally, Factor VIII heavy and light chains are expressed as a single polypeptide chain, e.g., along with an optional B-domain or B-domain substituted linker. However, in some embodiments, a Factor VIII heavy chain and Factor VIII light chain are expressed as separate polypeptide chains (e.g., co-expressed), and reconstituted to form a Factor VIII protein (e.g., in vivo or in vitro).

[00199] As used herein, the terms "B-domain substituted linker" and "Factor VIII linker" are used interchangeably, and refer to truncated versions of a wild type Factor VIII B-domain (e.g., amino acids 760-1667 of FVIII-FL-AA (SEQ ID NO: 19)) or peptides engineered to replace the B-domain of a Factor VIII polypeptide. As used herein, a Factor VIII linker is positioned between the C-terminus of a Factor VIII heavy chain and the N-terminus of a Factor VIII light chain in a Factor VIII variant polypeptide in accordance with some embodiments. Non-limiting examples of B-domain substituted linkers are disclosed in U.S. Patent Nos. 4,868,112, 5,112,950, 5,171,844, 5,543,502, 5,595,886, 5,610,278, 5,789,203, 5,972,885, 6,048,720, 6,060,447, 6,114,148, 6,228,620, 6,316,226, 6,346,513, 6,458,563, 6,924,365, 7,041,635, and 7,943,374; U.S. Patent Application Publication Nos. 2013/024960, 2015/0071883, and 2015/0158930; and PCT Publication Nos. WO 2014/064277 and WO 2014/127215, the disclosures of which are hereby incorporated by reference, in their entireties, for all purposes.

[00200] Unless otherwise specified herein, the numbering of Factor VIII amino acids refers to the corresponding amino acid in the full-length, wild-type human Factor VIII sequence (FVIII-FL-AA), presented as SEQ ID NO: 19 in Figure 22. As such, when referring to an amino acid substitution in a Factor VIII variant protein disclosed herein, the recited amino acid number refers to the analogous (e.g., structurally or functionally equivalent) and/or homologous (e.g., evolutionarily conserved in the primary amino acid sequence) amino acid in the full-length, wild-type Factor VIII sequence. For example, a T2105N amino acid substitution refers to a T to N substitution at position 2105 of the full-length, wild-type human Factor VIII sequence (FVIII-FL-AA; SEQ ID NO: 19), a T to N substitution at position 1211 of the Factor VIII variant protein encoded by CS04 (CS04-FL-AA; SEQ ID NO: 2), and a T to N substitution at position 1212 of the Factor VIII variant encoded by CS04m3 (CS04m3-FL-AA; SEQ ID NO: 105).

[00201] As described herein, the Factor VIII amino acid numbering system is dependent on whether the Factor VIII signal peptide (e.g., amino acids 1-19 of the full-length, wild-type human Factor VIII sequence) is included. Where the signal peptide is included, the numbering is referred to as "signal peptide inclusive" or "SPI". Where the signal peptide is not included, the numbering is referred to as "signal peptide exclusive" or "SPE." For example, F328S is SPI numbering for the same amino acid as F309S, in SPE numbering. Unless otherwise indicated, all

amino acid numbering refers to the corresponding amino acid in the full-length, wild-type human Factor VIII sequence (FVIII-FL-AA), presented as SEQ ID NO: 19 in Figure 22.

[00202] As described herein, the codon-altered polynucleotides provide increased expression of transgenic Factor VIII in vivo (e.g., when administered as part of a gene therapy vector), as compared to the level of Factor VIII expression provided by a natively-coded Factor VIII construct (e.g., a polynucleotide encoding the same Factor VIII construct using the wild-type human codons). As used herein, the term "increased expression" refers to an increased level of transgenic Factor VIII activity in the blood of an animal administered the codon-altered polynucleotide encoding Factor VIII, as compared to the level of transgenic Factor VIII activity in the blood of an animal administered a natively-coded Factor VIII construct. The activity levels can be measured using any Factor VIII activity known in the art. An exemplary assay for determining Factor VIII activity is the Technochrome FVIII assay (Technoclone, Vienna, Austria).

[00203] In some embodiments, increased expression refers to at least 25% greater transgenic Factor VIII activity in the blood of an animal administered the codon-altered Factor VIII polynucleotide, as compared to the level of transgenic Factor VIII activity in the blood of an animal administered a natively coded Factor VIII polynucleotide. In some embodiments, increased expression refers to at least 50% greater, at least 75% greater, at least 100% greater, at least 3-fold greater, at least 4-fold greater, at least 5-fold greater, at least 6-fold greater, at least 7-fold greater, at least 8-fold greater, at least 9-fold greater, at least 10-fold greater, at least 15-fold greater, at least 20-fold greater, at least 25-fold greater, at least 30-fold greater, at least 40-fold greater, at least 50-fold greater, at least 60-fold greater, at least 70-fold greater, at least 80-fold greater, at least 90-fold greater, at least 100-fold greater, at least 125-fold greater, at least 150-fold greater, at least 175-fold greater, at least 200-fold greater, at least 225-fold greater, or at least 250-fold greater transgenic Factor VIII activity in the blood of an animal administered the codon-altered Factor VIII polynucleotide, as compared to the level of transgenic Factor VIII activity in the blood of an animal administered a natively coded Factor VIII polynucleotide.

[00204] As described herein, the codon-altered polynucleotides provide increased vector production, as compared to the level of vector production provided by a natively-coded Factor

VIII construct (e.g., a polynucleotide encoding the same Factor VIII construct using the wild-type human codons). As used herein, the term "increased virus production" refers to an increased vector yield in cell culture (e.g., titer per liter culture) inoculated with the codon-altered polynucleotide encoding Factor VIII, as compared to the vector yield in cell culture inoculated with a natively-coded Factor VIII construct. The vector yields can be measured using any vector titer assay known in the art. An exemplary assay for determining vector yield (e.g., of an AAV vector) is qPCR targeting the AAV2 inverted terminal repeats (Aurnhammer, Human Gene Therapy Methods: Part B 23:18–28 (2012)).

[00205] In some embodiments, increased virus production refers to at least 25% greater codon-altered vector yield, as compared to the yield of a natively-coded Factor VIII construct in the same type of culture. In some embodiments, increased vector production refers to at least 50% greater, at least 75% greater, at least 100% greater, at least 3-fold greater, at least 4-fold greater, at least 5-fold greater, at least 6-fold greater, at least 7-fold greater, at least 8-fold greater, at least 9-fold greater, at least 10-fold greater, at least 15-fold greater, or at least 20-fold greater codon-altered vector yield, as compared to the yield of a natively-coded Factor VIII construct in the same type of culture.

[00206] As used herein, the term "hemophilia" refers to a group of disease states broadly characterized by reduced blood clotting or coagulation. Hemophilia may refer to Type A, Type B, or Type C hemophilia, or to the composite of all three diseases types. Type A hemophilia (hemophilia A) is caused by a reduction or loss of factor VIII (FVIII) activity and is the most prominent of the hemophilia subtypes. Type B hemophilia (hemophilia B) results from the loss or reduction of factor IX (FIX) clotting function. Type C hemophilia (hemophilia C) is a consequence of the loss or reduction in factor XI (FXI) clotting activity. Hemophilia A and B are X-linked diseases, while hemophilia C is autosomal. Conventional treatments for hemophilia include both prophylactic and on-demand administration of clotting factors, such as FVIII, FIX, including Bebulin®–VH, and FXI, as well as FEIBA-VH, desmopressin, and plasma infusions.

[00207] As used herein, the term "FVIII gene therapy" includes any therapeutic approach of providing a nucleic acid encoding Factor VIII to a patient to relieve, diminish, or prevent the reoccurrence of one or more symptoms (e.g., clinical factors) associated with hemophilia. The

term encompasses administering any compound, drug, procedure, or regimen comprising a nucleic acid encoding a Factor VIII molecule, including any modified form of Factor VIII (e.g., Factor VIII variant), for maintaining or improving the health of an individual with hemophilia. One skilled in the art will appreciate that either the course of FVIII therapy or the dose of a FVIII therapeutic agent can be changed, e.g., based upon the results obtained in accordance with the present disclosure.

[00208] As used herein, the term "bypass therapy" includes any therapeutic approach of providing non-Factor VIII hemostatic agents, compounds or coagulation factors to a patient to relieve, diminish, or prevent the reoccurrence of one or more symptoms (e.g., clinical factors) associated with hemophilia. Non-Factor VIII compounds and coagulation factors include, but are not limited to, Factor VIII Inhibitor Bypass Activity (FEIBA), recombinant activated factor VII (FVIIa), prothrombin complex concentrates, and activated prothrombin complex concentrates. These non-Factor VIII compounds and coagulation factors may be recombinant or plasma-derived. One skilled in the art will appreciate that either the course of bypass therapy or the dose of bypass therapy can be changed, e.g., based upon the results obtained in accordance with the present disclosure.

[00209] As used herein, a “combination therapy” including administration of a nucleic acid encoding a Factor VIII molecule and a conventional hemophilia A therapeutic agent includes any therapeutic approach of providing both a nucleic acid encoding a Factor VIII molecule and a Factor VIII molecule and/or non-Factor VIII hemostatic agent (e.g., bypass therapeutic agent) to a patient to relieve, diminish, or prevent the reoccurrence of one or more symptoms (e.g., clinical factors) associated with hemophilia. The term encompasses administering any compound, drug, procedure, or regimen including a nucleic acid encoding a Factor VIII molecule, including any modified form of factor VIII, which is useful for maintaining or improving the health of an individual with hemophilia and includes any of the therapeutic agents described herein.

[00210] The terms "therapeutically effective amount or dose" or "therapeutically sufficient amount or dose" or "effective or sufficient amount or dose" refer to a dose that produces therapeutic effects for which it is administered. For example, a therapeutically effective amount

of a drug useful for treating hemophilia can be the amount that is capable of preventing or relieving one or more symptoms associated with hemophilia. The exact dose will depend on the purpose of the treatment, and will be ascertainable by one skilled in the art using known techniques (*see*, *e.g.*, Lieberman, *Pharmaceutical Dosage Forms* (vols. 1-3, 1992); Lloyd, *The Art, Science and Technology of Pharmaceutical Compounding* (1999); Pickar, *Dosage Calculations* (1999); and *Remington: The Science and Practice of Pharmacy*, 20th Edition, 2003, Gennaro, Ed., Lippincott, Williams & Wilkins).

[00211] As used herein, the term "gene" refers to the segment of a DNA molecule that codes for a polypeptide chain (e.g., the coding region). In some embodiments, a gene is positioned by regions immediately preceding, following, and/or intervening the coding region that are involved in producing the polypeptide chain (e.g., regulatory elements such as a promoter, enhancer, polyadenylation sequence, 5'-untranslated region, 3'-untranslated region, or intron).

[00212] As used herein, the term "regulatory elements" refers to nucleotide sequences, such as promoters, enhancers, terminators, polyadenylation sequences, introns, etc, that provide for the expression of a coding sequence in a cell.

[00213] As used herein, the term "promoter element" refers to a nucleotide sequence that assists with controlling expression of a coding sequence. Generally, promoter elements are located 5' of the translation start site of a gene. However, in certain embodiments, a promoter element may be located within an intron sequence, or 3' of the coding sequence. In some embodiments, a promoter useful for a gene therapy vector is derived from the native gene of the target protein (e.g., a Factor VIII promoter). In some embodiments, a promoter useful for a gene therapy vector is specific for expression in a particular cell or tissue of the target organism (e.g., a liver-specific promoter). In yet other embodiments, one of a plurality of well characterized promoter elements is used in a gene therapy vector described herein. Non-limiting examples of well-characterized promoter elements include the CMV early promoter, the β-actin promoter, and the methyl CpG binding protein 2 (MeCP2) promoter. In some embodiments, the promoter is a constitutive promoter, which drives substantially constant expression of the target protein. In other embodiments, the promoter is an inducible promoter, which drives expression of the target

protein in response to a particular stimulus (e.g., exposure to a particular treatment or agent). For a review of designing promoters for AAV-mediated gene therapy, see Gray et al. (Human Gene Therapy 22:1143-53 (2011)), the contents of which are expressly incorporated by reference in their entirety for all purposes.

[00214] As used herein, the term "vector" refers to any vehicle used to transfer a nucleic acid (e.g., encoding a Factor VIII gene therapy construct) into a host cell. In some embodiments, a vector includes a replicon, which functions to replicate the vehicle, along with the target nucleic acid. Non-limiting examples of vectors useful for gene therapy include plasmids, phages, cosmids, artificial chromosomes, and viruses, which function as autonomous units of replication in vivo. In some embodiments, a vector is a viral vehicle for introducing a target nucleic acid (e.g., a codon-altered polynucleotide encoding a Factor VIII variant). Many modified eukaryotic viruses useful for gene therapy are known in the art. For example, adeno-associated viruses (AAVs) are particularly well suited for use in human gene therapy because humans are a natural host for the virus, the native viruses are not known to contribute to any diseases, and the viruses illicit a mild immune response.

[00215] As used herein, the term “CpG island” refers to a region within a polynucleotide having a statistically elevated density of CpG dinucleotides. As used herein, a region of a polynucleotide (e.g., a polynucleotide encoding a codon-altered Factor VIII protein) is a CpG island if, over a 200-base pair window: (i) the region has GC content of greater than 50%, and (ii) the ratio of observed CpG dinucleotides per expected CpG dinucleotides is at least 0.6, as defined by the relationship:

$$\frac{N[CpG]*N[length\ of\ window]}{N[C]*N\,[G]} \geq 0.6.$$

For additional information on methods for identifying CpG islands, *see* Gardiner-Garden M. et al., J Mol Biol., 196(2):261-82 (1987), the content of which is expressly incorporated herein by reference, in its entirety, for all purposes.

[00216] As used herein, the term "nucleic acid" refers to deoxyribonucleotides or ribonucleotides and polymers thereof in either single- or double-stranded form, and complements thereof. The term encompasses nucleic acids containing known nucleotide analogs or modified

backbone residues or linkages, which are synthetic, naturally occurring, and non-naturally occurring, which have similar binding properties as the reference nucleic acid, and which are metabolized in a manner similar to the reference nucleotides. Examples of such analogs include, without limitation, phosphorothioates, phosphoramidates, methyl phosphonates, chiral-methyl phosphonates, 2-O-methyl ribonucleotides, and peptide-nucleic acids (PNAs).

[00217] The term "amino acid" refers to naturally occurring and non-natural amino acids, including amino acid analogs and amino acid mimetics that function in a manner similar to the naturally occurring amino acids. Naturally occurring amino acids include those encoded by the genetic code, as well as those amino acids that are later modified, e.g., hydroxyproline, y-carboxyglutamate, and O-phosphoserine. Naturally occurring amino acids can include, e.g., D- and L-amino acids. The amino acids used herein can also include non-natural amino acids. Amino acid analogs refer to compounds that have the same basic chemical structure as a naturally occurring amino acid, i.e., any carbon that is bound to a hydrogen, a carboxyl group, an amino group, and an R group, e.g., homoserine, norleucine, methionine sulfoxide, or methionine methyl sulfonium. Such analogs have modified R groups (e.g., norleucine) or modified peptide backbones, but retain the same basic chemical structure as a naturally occurring amino acid. Amino acid mimetics refer to chemical compounds that have a structure that is different from the general chemical structure of an amino acid, but that function in a manner similar to a naturally occurring amino acid. Amino acids may be referred to herein by either their commonly known three letter symbols or by the one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Nucleotides, likewise, may be referred to by their commonly accepted single-letter codes.

[00218] The nucleotide sequences that encode the mutant Factor VIII constructs herein may be identical to the coding sequence provided herein or may be a different coding sequence, which sequence, as a result of the redundancy or degeneracy of the genetic code, encodes the same polypeptides as the coding sequences provided herein. One of ordinary skill in the art will recognize that each codon in a nucleic acid (except AUG, which is ordinarily the only codon for methionine, and TGG, which is ordinarily the only codon for tryptophan) can be modified to yield a functionally identical molecule. Accordingly, each variation of a nucleic acid which

encodes a same polypeptide is implicit in each described sequence with respect to the expression product, but not with respect to actual gene therapy constructs.

[00219] As to amino acid sequences, one of ordinary skill in the art will recognize that individual substitutions, deletions or additions to a nucleic acid or peptide sequence that alters, adds or deletes a single amino acid or a small percentage of amino acids in the encoded sequence is a "conservatively modified variant" where the alteration results in the substitution of an amino acid with a chemically similar amino acid. Conservative substitution tables providing functionally similar amino acids are well known in the art. Such conservatively modified variants are in addition to and do not exclude polymorphic variants, interspecies homologs, and alleles of the disclosure.

[00220] Conservative amino acid substitutions providing functionally similar amino acids are well known in the art. Dependent on the functionality of the particular amino acid, e.g., catalytic, structural, or sterically important amino acids, different groupings of amino acid may be considered conservative substitutions for each other. **Table 1** provides groupings of amino acids that are considered conservative substitutions based on the charge and polarity of the amino acid, the hydrophobicity of the amino acid, the surface exposure/structural nature of the amino acid, and the secondary structure propensity of the amino acid.

Table 1. Groupings of conservative amino acid substitutions based on the functionality of the residue in the protein.

Important Feature	Conservative Groupings
Charge/Polarity	1. H, R, and K 2. D and E 3. C, T, S, G, N, Q, and Y 4. A, P, M, L, I, V, F, and W
Hydrophobicity	1. D, E, N, Q, R, and K 2. C, S, T, P, G, H, and Y 3. A, M, I, L, V, F, and W

Structural/Surface Exposure	1. D, E, N, Q, H, R, and K 2. C, S, T, P, A, G, W, and Y 3. M, I, L, V, and F
Secondary Structure Propensity	1. A, E, Q, H, K, M, L, and R 2. C, T, I, V, F, Y, and W 3. S, G, P, D, and N
Evolutionary Conservation	1. D and E 2. H, K, and R 3. N and Q 4. S and T 5. L, I, and V 6. F, Y, and W 7. A and G 8. M and C

[00221] The terms "identical" or percent "identity," in the context of two or more nucleic acids or peptide sequences, refer to two or more sequences or subsequences that are the same or have a specified percentage of amino acid residues or nucleotides that are the same (i.e., about 60% identity, preferably 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or higher identity over a specified region, when compared and aligned for maximum correspondence over a comparison window or designated region) as measured using a BLAST or BLAST 2.0 sequence comparison algorithms with default parameters described below, or by manual alignment and visual inspection.

[00222] As is known in the art, a number of different programs may be used to identify whether a protein (or nucleic acid as discussed below) has sequence identity or similarity to a known sequence. Sequence identity and/or similarity is determined using standard techniques known in the art, including, but not limited to, the local sequence identity algorithm of Smith &

Waterman, Adv. Appl. Math., 2:482 (1981), by the sequence identity alignment algorithm of Needleman & Wunsch, J. Mol. Biol., 48:443 (1970), by the search for similarity method of Pearson & Lipman, Proc. Natl. Acad. Sci. U.S.A., 85:2444 (1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Drive, Madison, WI), the Best Fit sequence program described by Devereux et al., Nucl. Acid Res., 12:387-395 (1984), preferably using the default settings, or by inspection. Preferably, percent identity is calculated by FastDB based upon the following parameters: mismatch penalty of 1; gap penalty of 1; gap size penalty of 0.33; and joining penalty of 30, "Current Methods in Sequence Comparison and Analysis," Macromolecule Sequencing and Synthesis, Selected Methods and Applications, pp 127-149 (1988), Alan R. Liss, Inc, all of which are incorporated by reference.

[00223] An example of a useful algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using progressive, pair wise alignments. It may also plot a tree showing the clustering relationships used to create the alignment. PILEUP uses a simplification of the progressive alignment method of Feng & Doolittle, J. Mol. Evol. 35:351-360 (1987); the method is similar to that described by Higgins & Sharp CABIOS 5:151-153 (1989), both incorporated by reference. Useful PILEUP parameters including a default gap weight of 3.00, a default gap length weight of 0.10, and weighted end gaps.

[00224] Another example of a useful algorithm is the BLAST algorithm, described in: Altschul et al., J. Mol. Biol. 215, 403-410, (1990); Altschul et al., Nucleic Acids Res. 25:3389-3402 (1997); and Karlin et al., Proc. Natl. Acad. Sci. U.S.A. 90:5873-5787 (1993), both incorporated by reference. A particularly useful BLAST program is the WU-BLAST-2 program which was obtained from Altschul et al., Methods in Enzymology, 266:460-480 (1996); http://blast.wustl/edu/blast/ README.html]. WU-BLAST-2 uses several search parameters, most of which are set to the default values. The adjustable parameters are set with the following values: overlap span =1, overlap fraction = 0.125, word threshold (T) = 11. The HSP S and HSP S2 parameters are dynamic values and are established by the program itself depending upon the composition of the particular sequence and composition of the particular database against which the sequence of interest is being searched; however, the values may be adjusted to increase sensitivity.

[00225] An additional useful algorithm is gapped BLAST, as reported by Altschul et al., Nucl. Acids Res., 25:3389-3402, incorporated by reference. Gapped BLAST uses BLOSUM-62 substitution scores; threshold T parameter set to 9; the two-hit method to trigger ungapped extensions; charges gap lengths of k a cost of 10+k; Xu set to 16, and Xg set to 40 for database search stage and to 67 for the output stage of the algorithms. Gapped alignments are triggered by a score corresponding to ~22 bits.

[00226] A % amino acid sequence identity value is determined by the number of matching identical residues divided by the total number of residues of the "longer" sequence in the aligned region. The "longer" sequence is the one having the most actual residues in the aligned region (gaps introduced by WU-Blast-2 to maximize the alignment score are ignored). In a similar manner, "percent (%) nucleic acid sequence identity" with respect to the coding sequence of the polypeptides identified is defined as the percentage of nucleotide residues in a candidate sequence that are identical with the nucleotide residues in the coding sequence of the cell cycle protein. A preferred method utilizes the BLASTN module of WU-BLAST-2 set to the default parameters, with overlap span and overlap fraction set to 1 and 0.125, respectively.

[00227] The alignment may include the introduction of gaps in the sequences to be aligned. In addition, for sequences which contain either more or fewer amino acids than the protein encoded by the sequence of Figure 2 (SEQ ID NO:1), it is understood that in one embodiment, the percentage of sequence identity will be determined based on the number of identical amino acids or nucleotides in relation to the total number of amino acids or nucleotides. Thus, for example, sequence identity of sequences shorter than that shown in Figure 2 (SEQ ID NO:1), as discussed below, will be determined using the number of nucleotides in the shorter sequence, in one embodiment. In percent identity calculations relative weight is not assigned to various manifestations of sequence variation, such as, insertions, deletions, substitutions, etc.

[00228] In one embodiment, only identities are scored positively (+1) and all forms of sequence variation including gaps are assigned a value of "0", which obviates the need for a weighted scale or parameters as described below for sequence similarity calculations. Percent sequence identity may be calculated, for example, by dividing the number of matching identical residues by the total number of residues of the "shorter" sequence in the aligned region and

multiplying by 100. The "longer" sequence is the one having the most actual residues in the aligned region.

[00229] The term "allelic variants" refers to polymorphic forms of a gene at a particular genetic locus, as well as cDNAs derived from mRNA transcripts of the genes, and the polypeptides encoded by them. The term "preferred mammalian codon" refers a subset of codons from among the set of codons encoding an amino acid that are most frequently used in proteins expressed in mammalian cells as chosen from the following list: Gly (GGC, GGG); Glu (GAG); Asp (GAC); Val (GTG, GTC); Ala (GCC, GCT); Ser (AGC, TCC); Lys (AAG); Asn (AAC); Met (ATG); Ile (ATC); Thr (ACC); Trp (TGG); Cys (TGC); Tyr (TAT, TAC); Leu (CTG); Phe (TTC); Arg (CGC, AGG, AGA); Gln (CAG); His (CAC); and Pro (CCC).

[00230] As used herein, the term codon-altered refers to a polynucleotide sequence encoding a polypeptide (e.g., a Factor VIII variant protein), where at least one codon of the native polynucleotide encoding the polypeptide has been changed to improve a property of the polynucleotide sequence. In some embodiments, the improved property promotes increased transcription of mRNA coding for the polypeptide, increased stability of the mRNA (e.g., improved mRNA half-life), increased translation of the polypeptide, and/or increased packaging of the polynucleotide within the vector. Non-limiting examples of alterations that can be used to achieve the improved properties include changing the usage and/or distribution of codons for particular amino acids, adjusting global and/or local GC content, removing AT-rich sequences, removing repeated sequence elements, adjusting global and/or local CpG dinucleotide content, removing cryptic regulatory elements (e.g., TATA box and CCAAT box elements), removing of intron/exon splice sites, improving regulatory sequences (e.g., introduction of a Kozak consensus sequence), and removing sequence elements capable of forming secondary structure (e.g., stem-loops) in the transcribed mRNA.

[00231] As discussed herein, there are various nomenclatures to refer to components of the disclosure herein. "CS-number" (e.g. "CS04", "CS01", "CS23", etc.) refer to codon altered polynucleotides encoding FVIII polypeptides and/or the encoded polypeptides, including variants. For example, CS01-FL refers to the **F**ull **L**ength codon altered CS01 polynucleotide sequence or amino acid sequence (sometimes referred to herein as "CS01-FL-AA" for the

Amino **A**cid sequence and "CS01-FL-NA" for the **N**ucleic **A**cid sequence) encoded by the CS01 polynucleotide sequence. Similarly, "CS01-LC" refers to either the codon altered nucleic acid sequence ("CS01-LC-NA") encoding the light chain of a FVIII polypeptide or the amino acid sequence (also sometimes referred to herein as "CS01-LC-AA") of the FVIII light chain encoded by the CS01 polynucleotide sequence. Likewise, CS01-HC, CS01-HC-AA and CS01-HC-NA are the same for the FVIII heavy chain. As will be appreciated by those in the art, for constructs such as CS01, CS04, CS23, etc., that are only codon-altered (e.g. they do not contain additional amino acid substitutions as compared to Refacto), the amino acid sequences will be identical, as the amino acid sequences are not altered by the codon optimization. Thus, sequence constructs of the disclosure include, but are not limited to, CS01-FL-NA, CS01-FL-AA, CS01-LC-NA, CS01-LC-AA, CS01-HC-AA, CS01-HC-NA, CS04-FL-NA, CS04-FL-AA, CS04-LC-NA, CS04-LC-AA, CS04-HC-AA, CS04-HC-NA, CS23-FL-NA, CS23-FL-AA, CS23-LC-NA, CS23-LC-AA, CS23-HC-AA and CS23-HC-NA.

[00232] This nomenclature also applies to glycosylation peptides as shown in Figure 13, such that "NGA1-AA" refers to the amino acid sequence and NGA1-NA refers to the nucleic acid sequence.

[00233] The disclosure also includes additional new Factor VIII variants, as described below, with the appropriate nomenclature.

III. Codon-Altered Factor VIII Variants

[00234] In some embodiments, the present disclosure provides codon-altered polynucleotides encoding Factor VIII variants. These codon-altered polynucleotides provide markedly improved expression of Factor VIII when administered in an AAV-based gene therapy construct. The codon-altered polynucleotides also demonstrate improved AAV-virion packaging, as compared to conventionally codon-optimized constructs. As demonstrated in Example 2 and Example 4, Applicants have achieve these advantages through the discovery of three codon-altered polynucleotides (CS01-FL-NA, CS04-FL-NA, and CS23-FL-NA) encoding a Factor VIII polypeptide with human wild-type Factor VIII heavy and light chains, and a short, 14 amino acid, B-domain substituted linker (the "SQ" linker) containing a furin cleavage site to facilitate maturation of an active FVIIIa protein in vivo. As further demonstrated in Example 4,

incorporation of various combinations of the F328S, X5, and X1 amino acid mutations into the encoded Factor VIII molecule further increased the in vivo expression of Factor VIII activity.

[00235] In one embodiment, a codon-altered polynucleotide provided herein has nucleotide sequences with high sequence identity to at least the sequences within CS01, CS04, or CS23 (SEQ ID NOS 13, 1, and 20, respectively) encoding the Factor VIII heavy chain and Factor VIII light chains. As known in the art, the B-domain of Factor VIII is dispensable for activity in vivo. Thus, in some embodiments, the codon-altered polynucleotides provided herein completely lack a Factor VIII B-domain. In some embodiments, the native Factor VIII B-domain is replaced with a short amino acid linker containing a furin cleavage site, e.g., the "SQ" linker consisting of amino acids 760-773 of the CS01, CS04, or CS23 (SEQ ID NOS 2, 2, and 21, respectively) constructs. The "SQ" linker is also referred to as BDLO04, (-AA for the amino acid sequence and -NA for the nucleotide sequence shown in Figure 6).

[00236] In one embodiment, the Factor VIII heavy and light chains encoded by the codon-altered polynucleotide are human Factor VIII heavy and light chains, respectively. In other embodiments, the Factor VIII heavy and light chains encoded by the codon-altered polynucleotide are heavy and light chain sequences from another mammal (e.g., porcine Factor VIII). In yet other embodiments, the Factor VIII heavy and light chains are chimeric heavy and light chains (e.g., a combination of human and a second mammalian sequence). In yet other embodiments, the Factor VIII heavy and light chains are humanized version of the heavy and light chains from another mammal, e.g., heavy and light chain sequences from another mammal in which human residues are substituted at select positions to reduce the immunogenicity of the resulting peptide when administered to a human.

[00237] The GC content of human genes varies widely, from less than 25% to greater than 90%. However, in general, human genes with higher GC contents are expressed at higher levels. For example, Kudla et al. (PLoS Biol., 4(6):80 (2006)) demonstrate that increasing a gene's GC content increases expression of the encoded polypeptide, primarily by increasing transcription and effecting a higher steady state level of the mRNA transcript. Generally, the desired GC content of a codon-optimized gene construct is equal or greater than 60%. However, native AAV genomes have GC contents of around 56%.

[00238] Accordingly, in some embodiments, the codon-altered polynucleotides provided herein have a CG content that more closely matches the GC content of native AAV virions (e.g., around 56% GC), which is lower than the preferred CG contents of polynucleotides that are conventionally codon-optimized for expression in mammalian cells (e.g., at or above 60% GC). As outlined in Example 1, CS04-FL-NA (SEQ ID NO: 1), which has a GC content of about 56%, has improved virion packaging as compared to similarly codon-altered coding sequences with higher GC content.

[00239] Thus, in some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is less than 60%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is less than 59%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is less than 58%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is less than 57%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is no more than 56%.

[00240] In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 54% to 59%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 55% to 59%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 56% to 59%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 54% to 58%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 55% to 58%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 56% to 58%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 54% to 57%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 55% to 57%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 56% to 57%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 54% to 56%. In some

embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is from 55% to 56%.

[00241] In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56±0.5%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56±0.4%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56±0.3%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56±0.2%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56±0.1%. In some embodiments, the overall GC content of a codon-altered polynucleotide encoding a Factor VIII polypeptide is 56%.

A. Factor VIII Amino Acid Substitutions

[00242] To further increase the efficiency of AAV-vector based expression of the Factor VIII constructs described herein, amino acid substitutions know to improve secretion, increase specific activity, and/or enhanced the stability of Factor VIII are further incorporated, in some implementations. A number of potential variants were identified that increase the plasma levels of FVIII activity at a given vector dose. These variants include those with a more efficient signal peptide, amino acid substitutions that prevent BiP interactions, amino acid substitutions resembling more efficiently secreted Factor VIII orthologs (e.g., porcine Factor VIII), single-chain Factor VIII variants, and amino acid substitutions that stabilize Factor VIII and/or reduce subunit dissociation.

[00243] Mutation of residues A108, R121, and L2302 (SPE), located at the interface between the A1 and C2 domains, increases the stability of Factor VIII. For example, the A108I amino acid substitution introduces a hydrophobic residue that better fills the inter-domain space, stabilizing the interaction. Likewise, an R121C/L2302C (SPE) double amino acid substitution introduces a disulfide bond spanning the A1-C2 domains, further stabilizing the interaction. Taken together, all three amino acid substitutions increase the thermal stability of Factor VIII by 3 to 4-fold. For review, see Wakabayashi et al., J Biol Chem. 286(29):25748-55 (2011) and Wakabayashi et al., Thromb Haemost. 10(3):492-95 (2012). Accordingly, in some

embodiments, the encoded Factor VIII polypeptide includes A108I and/or R121C/L2302C amino acid substitutions.

[00244] Mutation of E113 (SPE), located within the calcium binding domain of Factor VIII, increases the specific FVIII clotting activity. For example, E113A appears to increase FXase formation through increased FVIII affinity for Factor IXa. Specifically, the E113A amino acid substitution increases specific FVIII clotting activity two-fold and increases affinity for Factor IXa by four-fold (Biochemistry, 41:8485 (2002); J. Biol. Chem., 279:12677 (2004); and Biochemistry, 44:10298 (2005)). Accordingly, in some embodiments, the encoded Factor VIII polypeptides include an E113A amino acid substitution.

[00245] Substitution of one or more amino acid residues surrounding the Factor VIII APC cleavage site (residues 331-341 (SPE)) reduce Factor VIIIa inactivation by activated protein C, without affecting FVIII activity. For example PQL333-335VDQ (SPE) amino acid substitutions reduce Factor VIII inactivation by 16-fold. Likewise, MKN336-339GNQ amino acid substitutions reduce Factor VIII inactivation by 9-fold. When combined, the two triple amino acid substitutions (e.g., PQLRMKN333-339VDQRGNQ) (SEQ ID NOS 34 and 35, respectively) reduce Factor VIII inactivation by 100-fold (J. Biol. Chem., 282:20264 (2007). Accordingly, in some embodiments, the encoded Factor VIII polypeptide include PQL333-335VDQ and/or MKN337-339GNQ (SPE) amino acid substitutions.

[00246] Mutations within the A2 domain interface also increase Factor VIII stability. Specifically, mutating charged residues in the A1-A2 and A2-A3 domain interfaces increases stability and retention of the A2 subunit in Factor VIIIa. For example, mutation of D519, E665, and E1984 to V or A yields up to 2-fold increased stability in Factor VIII and up to 5-fold stability in Factor VIIIa. Specifically, D519A/E665V amino acid substitutions provide a 3-fold increase in stability; D519V/E665V amino acid substitutions provide a 2-fold increase in stability, an 8-fold decrease in A2 dissociation, and a 2-4-fold increase in thrombin generation potential; D519V/E1984A amino acid substitutions provide a 2-fold increase in stability; and D519V/E665V/E1984A amino acid substitution provide a 2-fold increase in stability (Blood 112:2761-69 (2008); J. Thromb. Haemost., 7:438-44 (2009)). Accordingly, in some

embodiments, the encoded Factor VIII polypeptides include one or more of D519A/V, E665A/V, and E1984A/V amino acid substitutions.

[00247] Of particular relevance to the present disclosure are a number of specific mutations that can be included separately or in combinations with other variants described herein. These variants are coded as sets herein as follows: "m1" refers to a single amino acid change, "m2" is a set of 5 amino acid variants, "m3" is a combination of a deletion of 7 amino acids and an insertion of six amino acids that span the junction between the polypeptide linker and the heavy chain, "m4" is a combination of the m1 single mutation and the m5 double mutation, and "m5" is a set of two cysteine ablations. These mutations are described below. These can be included in any particular construct alone or in combination with other variants, and they are coded accordingly. For example, "m23" is a combination of the m2 and m3 variants onto a particular scaffold, as outlined herein; thus "CS01m23-FL-NA" or "CS01-FL-NAm23" refers to the CS01 codon-altered polynucleotide sequence with the nucleotides encoding the m2 and m3 mutations included, and "CS01m23-FL-AA" or "CS01-FL-AAm23" refers to the amino acid sequence. As CS01 is codon-altered but does not change the amino acid sequence of Refacto, these can be thought of on the amino acid level as mutations as compared to the Refacto amino acid sequence of CS01-FL-AA (SEQ ID NO: 2).

[00248] In many embodiments, the polypeptides of the disclosure are made with the "m1" variant included. Mutations within an 11 amino acid hydrophobic β-sheet in the A1 domain, which interacts with BiP, increase secretion of Factor VIII. For example, an F328S (SPI, F309S SPE) amino acid substitution within the pocket increased Factor VIII secretion 3-fold. The F328S variant is referred to herein as the "m1" mutation and is within the heavy chain. Again, as described herein, the number of the variants can be done inclusive of the signal peptide, "**S**ignal **P**eptide **I**nclusive", or "SPI", or starting from the processed final protein sequence, "**S**ignal **P**eptide **E**xclusive", or "SPE". Thus, using SPI numbering, the mutation F328S is the same as the F309 SPE mutant. Generally the specification uses the SPI numbering, but as will be appreciated by those in the art, either numbering system results in the same mutation(s).

[00249] Accordingly, included in the present disclosure are polypeptides that include the m1 mutation, including CS01-FL-AAm1, CS01-HC-AAm1, CS04-FL-AAm1, CS04-HC-AAm1

CS23-FL-AAm1, CS23-HC-AAm1, CS40-FL-AAm1 and CS40-HC-AAm1 (all of which encode the same corresponding protein sequences).

[00250] In addition, included in the present disclosure are not only polypeptide sequences that include the m1 mutation, but also those codon-altered polynucleotide sequences that encode proteins with the m1 mutation, such as CS01-FL-NAm1, CS01-HC-NAm1, CS04-FL-NAm1, CS04-HC-NAm1, CS23-FL-NAm1, CS23-HC-NA-m1, CS40-FL-NAm1 and CS40-HC-NAm1.

[00251] In many embodiments, the polypeptides of the disclosure are made with the "m2" variant set included, which is the I105V/A127S/G151K/M166T/L171P mutations (SPI numbering; (SPE numbering is V86I/S108A/K132G/T147M/P152L, respectively). The m2 mutation set is based on the fact that substitution of porcine amino acids 82-176 for the corresponding human amino acids in a B-domain deleted gene therapy construct increased Factor VIII activity when expressed in HEK293 cells (W. Xiao, communication). *Id.* Back-mutation of single porcine amino acids into the human BDD-FVIII construct identified five amino acids within the A1 domain that contribute to this phenomenon: I105V, A127S, G151K, M166T, and L171P (SPI). Introduction of the combination of these mutations into the human construct recapitulated the improved activity of the larger porcine substitution. *Id.* Accordingly, in some embodiments, the encoded Factor VIII polypeptides include one or more amino acid substitutions selected from I105V, A127S, G151K, M166T, and L171P, with the entire 5 amino acid set, m2, finding particular use in many embodiments. As for the m1 mutation, the m2 variants are in the heavy chain, and thus the present disclosure includes polypeptides that include the m2 mutation, including CS01-FL-AAm2, CS01-HC-AAm2, CS04-FL-AAm2, CS04-HC-AAm2, CS23-FL-AAm2, CS23-HC-AAm2, CS40-FL-AAm2 and CS40-HC-AAm2 (all of which encode the same corresponding protein sequences).

[00252] In addition, included in the present disclosure are not only polypeptide sequences that include the m2 mutation, but also those codon-altered polynucleotide sequences that encode proteins with the m2 mutation, such as CS01-FL-NAm2, CS01-HC-NAm2, CS04-FL-NAm2, CS04-HC-NAm2, CS23-FL-NAm2, CS23-HC-NA-m2, CS40-FL-NAm2 and CS40-HC-NAm2.

[00253] In additional embodiments, the polypeptides and polynucleotides of the disclosure include m3 mutations. m3 is the substitution of seven amino acids for six across the HC-B

domain interface that introduces an additional glycosylation site introduced close to the interface. Accordingly, in some embodiments, m3 is the deletion of amino acids AIEPRSF755-761 and the insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19) (e.g., AIEPRSF755-761TTYVNRSL) ("TTYVNRSL" disclosed as SEQ ID NO: 33). Residues AIEPR755-759, relative to SEQ ID NO: 19, fall within the end of the heavy chain, while residues S760 and F761 fall within the B-domain. In some embodiments, where the FVIII B-domain is deleted, truncated, or replaced, residues S760 and F761 may not be present in the underlying amino acid sequence being mutated. Accordingly, in some embodiments, m3 is the deletion of amino acids AIEPR755-759 and the insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19) (e.g., AIEPR755-759TTYVNRSL ("TTYVNRSL" disclosed as SEQ ID NO: 33)

[00254] The m3 variants are in the junction between the heavy chain and the B domain, and thus the present disclosure includes polypeptides that include the m3 mutation, including CS01-FL-AAm3, CS01-HC-AAm3, CS04-FL-AAm3, CS04-HC-AAm3, CS23-FL-AAm3, CS23-HC-AAm3, CS40-FL-AAm3 and CS40-HC-AAm3 (all of which encode the same corresponding protein sequences).

[00255] In addition, included in the present disclosure are not only polypeptide sequences that include the m3 mutation, but also those codon-altered polynucleotide sequences that encode proteins with the m3 mutations, such as CS01-FL-NAm3, CS01-HC-NAm3, CS04-FL-NAm3, CS04-HC-NAm3, CS23-FL-NAm3, CS23-HC-NA-m3, CS40-FL-NAm3 and CS40-HC-NAm3.

[00256] In additional embodiments, the polypeptides and polynucleotides of the disclosure include m4 mutations. Elimination of the C1899-C1903 disulfide bond in Factor VIII also increased secretion. Moreover, the increases in Factor VIII secretion are additive for the combination of F328S (SPI, F309S SPE) and C1918G/C1922G amino acid substitutions (Miao et al., Blood, 103:3412-19 (2004); Selvaraj et al., J. Thromb. Haemost., 10:107-15 (2012)). Accordingly, in some embodiments, the encoded Factor VIII polypeptides include m4 mutations, which is the F328S (SPI, F309S SPE) and C1918G/C1922G (SPI) amino acid substitutions. As the F328S variant is in the heavy chain and the two cysteine variants are in the light chain, polypeptide sequences that include m4 mutations are CS01-FL-AAm4, CS01-HC-AAm4, CS01-

LC-AAm4, CS04-FL-AAm4, CS04-HC-AAm4, CS04-LC-AAm4, CS23-FL-AAm4, CS23-HC-AAm4 and CS23-LC-AAm4.

[00257] In addition, included in the present disclosure are not only polypeptide sequences that include the m4 mutation, but also those codon-altered polynucleotide sequences that encode proteins with the m4 mutations, such as CS01-FL-NAm4, CS01-HC-NAm4, CS01-LC-NAm4, CS04-FL-NAm4, CS04-HC-NAm4, CS04-LC-NAm4, CS23-FL-NAm4, CS23-HC-NAm4, CS23-LC-NAm4, CS40-FL-NA-m4, CS40-HC-NA-m4 and CS40-LC-NA-m4.

[00258] In additional embodiments, the polypeptides and polynucleotides of the disclosure include m5 mutations. As above, elimination of the C1899-C1903 disulfide bond in Factor VIII also increased secretion. C1918G/C1922G (SPI) amino acid substitutions, contained within the light chain, referred to herein as the m5 mutation set.

[00259] The m5 variants are in the light chain, and thus the present disclosure includes polypeptides that include the m5 mutation, including CS01-FL-AAm5, CS01-LC-AAm5, CS04-FL-AAm5, CS04-LC-AAm5, CS23-FL-AAm5, CS23-LC-AAm5, CS40-FL-AAm5 and CS40-LC-AAm5 (all of which encode the same corresponding protein sequences).

[00260] In addition, included in the present disclosure are not only polypeptide sequences that include the m5 mutation, but also those codon-altered polynucleotide sequences that encode proteins with the m5 mutations, such as CS01-FL-NAm5, CS01-LC-NAm5, CS04-FL-NAm5, CS04-LC-NAm5, CS23-FL-NA-m5, CS23-LC-NA-m5, CS40-FL-NA-m5 and CS40-LC-NA-m5.

[00261] In addition to specific constructs (both amino acid and nucleic acid) that include m1, m2, m3, m4 and m5 individually, combinations of mutation sets can be made as outlined herein. As noted herein, these are noted as “m12”, which is the combination of m1 and m2 sets, or “m123” which is the combination of m1, m2 and m3 sets. Thus, included in the disclosure are dual combinations including m12, m13, m14, m15, m23, m24, m25, m34, m35 and m45. Also included are triple combinations, m123, m124, m125, m234, m235 and m345. Further included are quad combinations, m1234, m1235, m1345 and the m12345 combination.

Of particular interest in some embodiments are the following mutation sets: m1, m2, m3 and m4, m23, m123, and m234.

B. Factor VIII B-domain Substituted Linkers

[00262] In some embodiments, the linkage between the FVIII heavy chain and the light chain (e.g., the B-domain in wild-type Factor VIII) is further altered. Due to size constraints of AAV packaging capacity, B-domain deleted, truncated, and or linker substituted variants should improve the efficacy of the FVIII gene therapy construct. The most conventionally used B-domain substituted linker is that of SQ FVIII, which retains only 14 amino acids of the B domain as linker sequence. Another variant of porcine VIII ("OBI-1," described in U.S. Patent No. 6,458,563) is well expressed in CHO cells, and has a slightly longer linker of 24 amino acids. In some embodiments, the Factor VIII constructs encoded by the codon-altered polynucleotides described herein include an SQ-type B-domain linker sequence. In other embodiments, the Factor VIII constructs encoded by the codon-altered polynucleotides described herein include an OBI-1-type B-domain linker sequence.

[00263] In some embodiments, the encoded Factor VIII polypeptides described herein include an SQ-type B-domain linker, including amino acids 760-762/1657-1667 of the wild-type human Factor VIII B-domain (FVIII-FL-AA; SEQ ID NO: 19) (Sandberg et al. Thromb. Haemost. 85:93 (2001)). In some embodiments, the SQ-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments, the SQ-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the SQ-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00264] In some embodiments, the encoded Factor VIII polypeptides described herein include a Greengene-type B-domain linker, including amino acids 760/1582-1667 of the wild-type human Factor VIII B-domain (FVIII-FL-AA; SEQ ID NO: 19) (Oh et al., Biotechnol. Prog., 17:1999 (2001)). In some embodiments, the Greengene-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments,

the Greengene-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the Greengene-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00265] In some embodiments, the encoded Factor VIII polypeptides described herein include an extended SQ-type B-domain linker (SFSQNPPVLKRHQR; BDL-SQ-AA; SEQ ID NO: 30), including amino acids 760-769/1657-1667 of the wild-type human Factor VIII B-domain (FVIII-FL-AA; SEQ ID NO: 19) (Thim et al., Haemophilia, 16:349 (2010)). In some embodiments, the extended SQ-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments, the extended SQ-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the extended SQ-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00266] In some embodiments, the encoded Factor VIII polypeptides described herein include a porcine OBI-1-type B-domain linker, including the amino acids SFAQNSRPPSASAPKPPVLRRHQR (SEQ ID NO: 31) from the wild-type porcine Factor VIII B-domain (Toschi et al., Curr. Opin. Mol. Ther. 12:517 (2010)). In some embodiments, the porcine OBI-1-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments, the porcine OBI-1-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the porcine OBI-1-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00267] In some embodiments, the encoded Factor VIII polypeptides described herein include a human OBI-1-type B-domain linker, including amino acids 760-772/1655-1667 of the

wild-type human Factor VIII B-domain (FVIII-FL-AA; SEQ ID NO: 19). In some embodiments, the human OBI-1-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments, the human OBI-1-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the human OBI-1-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00268] In some embodiments, the encoded Factor VIII polypeptides described herein include an O8-type B-domain linker, including the amino acids SFSQNSRHQAYRYRRG (SEQ ID NO: 32) from the wild-type porcine Factor VIII B-domain (Toschi et al., Curr. Opin. Mol. Ther. 12:517 (2010)). In some embodiments, the porcine OBI-1-type B-domain linker has one amino acid substitution relative to the corresponding wild-type sequence. In some embodiments, the porcine OBI-1-type B-domain linker has two amino acid substitutions relative to the corresponding wild-type sequence. In some embodiments, a glycosylation peptide is inserted into the porcine OBI-1-type B-domain linker. In some embodiments, the glycosylation peptide is selected from those shown in Figure 13 (SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance).

[00269] Removal of the B-domain from Factor VIII constructs does not appear to affect the activity of the activated enzyme (e.g., FVIIIa), presumably because the B-domain is removed during activation. However, the B-domain of Factor VIII contains several residues that are post-translationally modified, e.g., by N- or O-linked glycosylation. In silico analysis (Prediction of N-glycosylation sites in human proteins, R. Gupta, E. Jung and S. Brunak, in preparation (2004)) of the wild-type Factor VIII B-domain predicts that at least four of these sites are glycosylated in vivo (Figure 14). It is thought that these modifications within the B-domain contribute to the post-translational regulation and/or half-life of Factor VIII in vivo.

[00270] While the Factor VIII B-domain is absent in mature Factor VIIIa protein, glycosylation within the B-domain of the precursor Factor VIII molecule may increase the circulating half-life of the protein prior to activation. Thus, in some embodiments, the

polypeptide linker of the encoded Factor VIII constructs described herein includes one or more glycosylation sequences, to allow for glycosylation in vivo. In some embodiments, the polypeptide linker includes at least one consensus glycosylation sequence (e.g., an N- or O-linked glycosylation consensus sequence). In some embodiments, the polypeptide linker includes at least two consensus glycosylation sequences. In some embodiments, the polypeptide linker includes at least three consensus glycosylation sequences. In some embodiments, the polypeptide linker includes at least four consensus glycosylation sequences. In some embodiments, the polypeptide linker includes at least five consensus glycosylation sequences. In some embodiments, the polypeptide linker includes at least 6, 7, 8, 9, 10, or more consensus glycosylation sequences.

[00271] In some embodiments, the polypeptide linker contains at least one N-linked glycosylation sequence N-X-S/T, where X is any amino acid other than P, S, or T. In some embodiments, the polypeptide linker contains at least two N-linked glycosylation sequences N-X-S/T, where X is any amino acid other than P, S, or T. In some embodiments, the polypeptide linker contains at least three N-linked glycosylation sequences N-X-S/T, where X is any amino acid other than P, S, or T. In some embodiments, the polypeptide linker contains at least four N-linked glycosylation sequences N-X-S/T, where X is any amino acid other than P, S, or T. In some embodiments, the polypeptide linker contains at least five N-linked glycosylation sequences N-X-S/T, where X is any amino acid other than P, S, or T. In some embodiments, the polypeptide linker contains at least 6, 7, 8, 9, 10, or more N-linked glycosylation sequences N-X-S/T, where X is any amino acid other than P, S, or T.

[00272] In some embodiments, the polypeptide linker includes a glycosylation peptide with high sequence identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation polypeptide has at least 92% identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than two amino acid substitutions relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than one amino acid substitution

relative to any of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has an amino acid sequence selected from any of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B.

[00273] In some embodiments, the glycosylation peptide has at least 92% identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 90% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has at least 92% identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 95% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has at least 92% identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 98% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B.

[00274] In some embodiments, the glycosylation peptide has no more than two amino acid substitutions relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, and is encoded by a polynucleotide sequence having at least 90% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than two amino acid substitutions relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, and is encoded by a polynucleotide sequence having at least 95% identity to

a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than two amino acid substitutions relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, and is encoded by a polynucleotide sequence having at least 98% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B.

[00275] In some embodiments, the glycosylation peptide has no more than one amino acid substitution relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 90% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than one amino acid substitution relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 95% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has no more than one amino acid substitution relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 98% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B.

[00276] In some embodiments, the glycosylation peptide has a sequence selected from any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 90% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as

shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has a sequence selected from any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 95% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide has a sequence selected from any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B and is encoded by a polynucleotide sequence having at least 98% identity to a corresponding nucleotide sequence selected from SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance, as shown in Figures 13A-13B.

[00277] In some embodiments, the Factor VIII polypeptide encoded by a codon-altered polynucleotide described herein has a B-domain substituted linker in which a glycosylation peptide is inserted into the SQ linker sequence (amino acids 760-773 of CS04-FL-AA; SEQ ID NO: 2). In a specific embodiment, the glycosylation peptide is selected from selected from any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, a glycosylation peptide having at least 92% identity to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, a glycosylation peptide having no more than two amino acid substitutions relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B, and a glycosylation peptide having no more than one amino acid substitution relative to any one of SEQ ID NOS 51, 53, 55, 57, 59, 61, 63, 65, 67, 69, 71, 73, and 75, respectively, in order of appearance, as shown in Figures 13A-13B. In some embodiments, the glycosylation peptide is inserted in the SQ peptide between residues N768 and P769 (relative to CS04-FL-AA; SEQ ID NO: 2).

[00278] In some embodiments, the polypeptide linker of the Factor VIII construct is encoded by a third nucleotide sequence having high sequence identity to any one of those shown in Figure 6 (SEQ ID NOS 5-7 and 36-48, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 95% identity to any one of those shown

in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 96% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 97% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 98% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 99% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 99.5% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence has at least 99.9% identity to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance). In some embodiments, the third nucleotide sequence is identical to any one of those shown in Figure 13 (SEQ ID NOS 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, and 74, respectively, in order of appearance).

C. Codon-altered Polynucleotides Encoding a Factor VIII Variant with a Cleavable Linker

CS04 Codon Altered Polynucleotides

[00279] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a Factor VIII variant polypeptide with a linker that is cleavable in vivo. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS04-HC-NA (SEQ ID NO: 3), which is the portion of CS04-FL-NA (SEQ ID NO: 1) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to

CS04-LC-NA (SEQ ID NO: 4), which is the portion of CS04-FL-NA (SEQ ID NO: 1) encoding for a Factor VIII light chain. The polypeptide linker includes a furin cleavage site, which allows for maturation in vivo (e.g., after expression in vivo or administration of the precursor polypeptide).

[00280] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively.

[00281] In some embodiments, the polypeptide linker of the Factor VIII construct is encoded by a third nucleotide sequence having high sequence identity to BDLO04 (SEQ ID NO: 6), which encodes the 14-amino acid linker corresponding to amino acids 760-773 of CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the third nucleotide sequence has at least 95% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 96% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 97% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 98% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence is identical to BDLO04 (SEQ ID NO: 6).

[00282] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 95% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 96% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 97% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 98% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 99% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS04-FL-NA (SEQ ID NO: 1). In some embodiments, the nucleotide sequence is identical to CS04-FL-NA (SEQ ID NO: 1).

[00283] In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 97% identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 98% identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99% identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99.5% identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99.9% identity to CS04-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence is identical to CS04-FL-AA (SEQ ID NO: 2).

[00284] In some embodiments, the Factor VIII variant encoded by the CS04 polynucleotide, having high sequence homology to CS04-FL-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00285] In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide

comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m5 amino acid substitution.

[00286] In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m23 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m35 amino acid substitutions.

[00287] In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m125 amino acid substitutions.

CS01 Codon Altered Polynucleotides

[00288] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a Factor VIII variant polypeptide with a linker that is cleavable in vivo. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS01-HC-NA (SEQ ID NO: 24), which is the portion of CS01-FL-NA (SEQ ID NO: 13) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to CS01-LC-NA (SEQ ID NO: 25), which is the portion of CS01-FL-NA (SEQ ID NO: 13) encoding for a Factor VIII light chain. The polypeptide linker includes a furin cleavage site,

which allows for maturation in vivo (e.g., after expression in vivo or administration of the precursor polypeptide).

[00289] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively.

[00290] In some embodiments, the polypeptide linker of the Factor VIII construct is encoded by a third nucleotide sequence having high sequence identity to BDLO04 (SEQ ID NO: 6), which encodes the 14-amino acid linker corresponding to amino acids 760-773 of CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the third nucleotide sequence has at least 95% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 96% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 97% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 98% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence is identical to BDLO04 (SEQ ID NO: 6).

[00291] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments,

the nucleotide sequence has at least 95% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 96% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 97% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 98% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 99% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS01-FL-NA (SEQ ID NO: 13). In some embodiments, the nucleotide sequence is identical to CS01-FL-NA (SEQ ID NO: 13).

[00292] In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 97% identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 98% identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99% identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99.5% identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence has at least 99.9% identity to CS01-FL-AA (SEQ ID NO: 2). In some embodiments, the amino acid sequence is identical to CS01-FL-AA (SEQ ID NO: 2).

[00293] In some embodiments, the Factor VIII variant encoded by the CS01 polynucleotide, having high sequence homology to CS01-FL-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00294] In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m5 amino acid substitution.

[00295] In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m23 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m35 amino acid substitutions.

[00296] In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m125 amino acid substitutions.

CS23 Codon Altered Polynucleotides

[00297] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a Factor VIII variant polypeptide with a linker that is cleavable in vivo. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS23-HC-NA (SEQ ID NO: 22), which is the portion of CS23-FL-NA (SEQ ID NO: 20) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to CS23-LC-NA (SEQ ID NO: 23), which is the portion of CS23-FL-NA (SEQ ID NO: 20) encoding for a Factor VIII light chain. The polypeptide linker includes a furin cleavage site, which allows for maturation in vivo (e.g., after expression in vivo or administration of the precursor polypeptide).

[00298] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively.

[00299] In some embodiments, the polypeptide linker of the Factor VIII construct is encoded by a third nucleotide sequence having high sequence identity to BDLO04 (SEQ ID NO: 6), which encodes the 14-amino acid linker corresponding to amino acids 760-773 of CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the third nucleotide sequence has at least 95% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 96% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 97% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence has at least 98% identity to BDLO04 (SEQ ID NO: 6). In some embodiments, the third nucleotide sequence is identical to BDLO04 (SEQ ID NO: 6).

[00300] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 95% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 96% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 97% identity to CS23-FL-NA

(SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 98% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 99% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS23-FL-NA (SEQ ID NO: 20). In some embodiments, the nucleotide sequence is identical to CS23-FL-NA (SEQ ID NO: 20).

[00301] In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence has at least 97% identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence has at least 98% identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence has at least 99% identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence has at least 99.5% identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence has at least 99.9% identity to CS23-FL-AA (SEQ ID NO: 21). In some embodiments, the amino acid sequence is identical to CS23-FL-AA (SEQ ID NO: 21).

[00302] In some embodiments, the Factor VIII variant encoded by the CS23 polynucleotide, having high sequence homology to CS23-FL-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00303] In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m5 amino acid substitution.

[00304] In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the

Factor VIII variant encoded by the CS23 polynucleotide comprises m23 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m35 amino acid substitutions.

[00305] In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m125 amino acid substitutions.

D. Codon-altered Polynucleotides Encoding a Single-chain Factor VIII Protein

[00306] Factor VIII constructs in which the furin cleavage site located at the C-terminal end of the B-domain is removed retain activity as a single chain polypeptide, despite that normal maturation of the Factor VIII molecule cannot occur (Leyte et al. (1991)). Similarly, a B-domain deleted Factor VIII construct with an attenuated furin site (containing an R1664H amino acid substitution) is more biologically active than the corresponding Factor VIII construct with a wild-type furin cleavage site (Siner et al. (2013)). Accordingly, in some embodiments, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a single-chain Factor VIII variant polypeptide. The single-chain Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The polypeptide linker does not include a furin cleavage site.

Single-chain CS04 Codon Altered Polynucleotides

[00307] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a single-chain Factor VIII variant polypeptide. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and an optional

polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS04-HC-NA (SEQ ID NO: 3), which is the portion of CS04-FL-NA (SEQ ID NO: 1) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to CS04-LC-NA (SEQ ID NO: 4), which is the portion of CS04-FL-NA (SEQ ID NO: 1) encoding for a Factor VIII light chain. The optional polypeptide linker does not include a furin cleavage site.

[00308] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS04-HC-NA and CS04-LC-NA (SEQ ID NOS 3 and 4), respectively.

[00309] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 95% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 96% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 97% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 98% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 99% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence

has at least 99.5% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS04-SC1-NA (SEQ ID NO: 9). In some embodiments, the nucleotide sequence is identical to CS04-SC1-NA (SEQ ID NO: 9).

[00310] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 95% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 96% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 97% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 98% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 99% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS04-SC2-NA (SEQ ID NO: 11). In some embodiments, the nucleotide sequence is identical to CS04-SC2-NA (SEQ ID NO: 11).

[00311] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS04-SC1-AA (SEQ ID NO: 10; human Factor VIIIΔ(760-1667) (SPI; *Hs*FVIIIΔ(741-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 97% identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 98% identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99% identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.5% identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.9% identity to CS04-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence is identical to CS04-SC1-AA (SEQ ID NO: 10).

[00312] In some embodiments, the Factor VIII variant encoded by the CS04-SC1 polynucleotide, having high sequence homology to CS04-SC1-AA (e.g., at least 95%, 96%,

97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00313] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS04-SC2-AA (SEQ ID NO: 12; human Factor VIIIΔ(772-1667) (SPI; *Hs*FVIIIΔ(753-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 97% identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 98% identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99% identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99.5% identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99.9% identity to CS04-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence is identical to CS04-SC2-AA (SEQ ID NO: 12).

[00314] In some embodiments, the single-chain Factor VIII variant encoded by the CS04-SC2 polynucleotide, having high sequence homology to CS04-SC2-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00315] In one embodiment, the single-chain Factor VIII variant encoded by the CS04 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises an m5 amino acid substitution.

[00316] In one embodiment, the single-chain Factor VIII variant encoded by the CS04 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m23 amino

acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m35 amino acid substitutions.

[00317] In one embodiment, the single-chain Factor VIII variant encoded by the CS04 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS04 polynucleotide comprises m125 amino acid substitutions.

Single-chain CS01 Codon Altered Polynucleotides

[00318] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a single-chain Factor VIII variant polypeptide. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and an optional polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS01-HC-NA (SEQ ID NO: 24), which is the portion of CS01-FL-NA (SEQ ID NO: 13) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to CS01-LC-NA (SEQ ID NO: 25), which is the portion of CS01-FL-NA (SEQ ID NO: 13) encoding for a Factor VIII light chain. The optional polypeptide linker does not include a furin cleavage site.

[00319] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to

CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS01-HC-NA and CS01-LC-NA (SEQ ID NOS 24 and 25), respectively.

[00320] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 95% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 96% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 97% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 98% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 99% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS01-SC1-NA (SEQ ID NO: 26). In some embodiments, the nucleotide sequence is identical to CS01-SC1-NA (SEQ ID NO: 26).

[00321] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 95% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 96% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 97% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 98% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 99% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS01-SC2-NA (SEQ ID NO: 27). In some

embodiments, the nucleotide sequence has at least 99.9% identity to CS01-SC2-NA (SEQ ID NO: 27). In some embodiments, the nucleotide sequence is identical to CS01-SC2-NA (SEQ ID NO: 27).

[00322] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS01-SC1-AA (SEQ ID NO: 10; human Factor VIIIΔ(760-1667) (SPI; *Hs*FVIIIΔ(741-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 97% identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 98% identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99% identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.5% identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.9% identity to CS01-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence is identical to CS01-SC1-AA (SEQ ID NO: 10).

[00323] In some embodiments, the Factor VIII variant encoded by the CS01-SC1 polynucleotide, having high sequence homology to CS01-SC1-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00324] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS01-SC2-AA (SEQ ID NO: 12; human Factor VIIIΔ(772-1667) (SPI; *Hs*FVIIIΔ(753-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 97% identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 98% identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99% identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99.5% identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid

sequence has at least 99.9% identity to CS01-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence is identical to CS01-SC2-AA (SEQ ID NO: 12).

[00325] In some embodiments, the single-chain Factor VIII variant encoded by the CS01-SC2 polynucleotide, having high sequence homology to CS01-SC2-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00326] In one embodiment, the single-chain Factor VIII variant encoded by the CS01 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises an m5 amino acid substitution.

[00327] In one embodiment, the single-chain Factor VIII variant encoded by the CS01 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m23 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m35 amino acid substitutions.

[00328] In one embodiment, the single-chain Factor VIII variant encoded by the CS01 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS01 polynucleotide comprises m125 amino acid substitutions.

Single-chain CS23 Codon Altered Polynucleotides

[00329] In one embodiment, the codon-altered polynucleotides provided herein include a nucleotide sequence encoding a single-chain Factor VIII variant polypeptide. The Factor VIII polypeptide includes a Factor VIII light chain, a Factor VIII heavy chain, and an optional polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain. The heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having high sequence identity to CS23-HC-NA (SEQ ID NO: 22), which is the portion of CS23-FL-NA (SEQ ID NO: 20) encoding for a Factor VIII heavy chain. The light chain of the Factor VIII polypeptide is encoded by a second nucleotide sequence with high sequence identity to CS23-LC-NA (SEQ ID NO: 23), which is the portion of CS23-FL-NA (SEQ ID NO: 20) encoding for a Factor VIII light chain. The optional polypeptide linker does not include a furin cleavage site.

[00330] In some embodiments, the first and second nucleotide sequences have at least 95% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 96% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 97% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 98% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.5% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences have at least 99.9% sequence identity to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively. In some embodiments, the first and second nucleotide sequences are identical to CS23-HC-NA and CS23-LC-NA (SEQ ID NOS 22 and 23), respectively.

[00331] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS23-SC1-NA (SEQ ID NO: 28). In some

embodiments, the nucleotide sequence has at least 95% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 96% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 97% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 98% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 99% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS23-SC1-NA (SEQ ID NO: 28). In some embodiments, the nucleotide sequence is identical to CS23-SC1-NA (SEQ ID NO: 28).

[00332] In some embodiments, the codon-altered polynucleotide has a nucleotide sequence with high sequence identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 95% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 96% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 97% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 98% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 99% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 99.5% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence has at least 99.9% identity to CS23-SC2-NA (SEQ ID NO: 29). In some embodiments, the nucleotide sequence is identical to CS23-SC2-NA (SEQ ID NO: 29).

[00333] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS23-SC1-AA (SEQ ID NO: 10; human Factor VIIIΔ(760-1667) (SPI; CS04Δ(741-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 97% identity to CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 98% identity to CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99% identity to

CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.5% identity to CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence has at least 99.9% identity to CS23-SC1-AA (SEQ ID NO: 10). In some embodiments, the amino acid sequence is identical to CS23-SC1-AA (SEQ ID NO: 10).

[00334] In some embodiments, the Factor VIII variant encoded by the CS23-SC1 polynucleotide, having high sequence homology to CS23-SC1-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00335] In some embodiments, the single-chain Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS23-SC2-AA (SEQ ID NO: 12; human Factor VIIIΔ(772-1667) (SPI; *Hs*FVIIIΔ(753-1648), SPE)). In some embodiments, the Factor VIII variant encoded by the codon-altered polynucleotide has an amino acid sequence with high sequence identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 97% identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 98% identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99% identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99.5% identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence has at least 99.9% identity to CS23-SC2-AA (SEQ ID NO: 12). In some embodiments, the amino acid sequence is identical to CS23-SC2-AA (SEQ ID NO: 12).

[00336] In some embodiments, the single-chain Factor VIII variant encoded by the CS23-SC2 polynucleotide, having high sequence homology to CS23-SC2-AA (e.g., at least 95%, 96%, 97%, 98%, 99%, 99.5%, or 99.9% identity), comprises one or more amino acid substitutions selected from m1, m2, m3, m4, and m5.

[00337] In one embodiment, the single-chain Factor VIII variant encoded by the CS23 polynucleotide comprises an m1 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m2 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m3 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23

polynucleotide comprises an m4 amino acid substitution. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises an m5 amino acid substitution.

[00338] In one embodiment, the single-chain Factor VIII variant encoded by the CS23 polynucleotide comprises m12 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m13 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m23 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m24 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m25 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m34 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m35 amino acid substitutions.

[00339] In one embodiment, the single-chain Factor VIII variant encoded by the CS23 polynucleotide comprises m123 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m234 amino acid substitutions. In one embodiment, the Factor VIII variant encoded by the CS23 polynucleotide comprises m125 amino acid substitutions.

E. Factor VIII Expression Vectors

[00340] In some embodiments, the codon-altered polynucleotides described herein are integrated into expression vectors. Non-limiting examples of expression vectors include viral vectors (e.g., vectors suitable for gene therapy), plasmid vectors, bacteriophage vectors, cosmids, phagemids, artificial chromosomes, and the like.

[00341] Non-limiting examples of viral vectors include: retrovirus, e.g., Moloney murine leukemia virus (MMLV), Harvey murine sarcoma virus, murine mammary tumor virus, and Rous sarcoma virus; adenoviruses, adeno-associated viruses; SV40-type viruses; polyomaviruses; Epstein-Barr viruses; papilloma viruses; herpes viruses; vaccinia viruses; and polio viruses.

[00342] In some embodiments, the codon-altered polynucleotides described herein are integrated into a gene therapy vector. In some embodiments, the gene therapy vector is a retrovirus, and particularly a replication-deficient retrovirus. Protocols for the production of replication-deficient retroviruses are known in the art. For review, see Kriegler, M., Gene Transfer and Expression, A Laboratory Manual, W.H. Freeman Co., New York (1990) and Murry, E. J., Methods in Molecular Biology, Vol. 7, Humana Press, Inc., Cliffton, N.J. (1991).

[00343] In one embodiment, the gene therapy vector is an adeno-associated virus (AAV) based gene therapy vector. AAV systems have been described previously and are generally well known in the art (Kelleher and Vos, *Biotechniques*, 17(6):1110-17 (1994); Cotten et al., *Proc Natl Acad Sci USA*, 89(13):6094-98 (1992); Curiel, *Nat Immun*, 13(2-3):141-64 (1994); Muzyczka, *Curr Top Microbiol Immunol*, 158:97-129 (1992); and Asokan A, et al., Mol. Ther., 20(4):699-708 (2012), each incorporated herein by reference in their entireties for all purposes). Details concerning the generation and use of rAAV vectors are described, for example, in U.S. Patent Nos. 5,139,941 and 4,797,368, each incorporated herein by reference in their entireties for all purposes. In a particular embodiment, the AAV vector is an AAV-8 vector.

[00344] In some embodiments, the codon-altered polynucleotides described herein are integrated into a retroviral expression vector. These systems have been described previously, and are generally well known in the art (Mann *et al.*, *Cell*, 33:153-159, 1983; Nicolas and Rubinstein, In: Vectors: A survey of molecular cloning vectors and their uses, Rodriguez and Denhardt, eds., Stoneham: Butterworth, pp. 494-513, 1988; Temin, In: Gene Transfer, Kucherlapati (ed.), New York: Plenum Press, pp. 149-188, 1986). In a specific embodiment, the retroviral vector is a lentiviral vector (*see*, for example, Naldini *et al.*, *Science*, 272(5259):263-267, 1996; Zufferey *et al.*, *Nat Biotechnol*, 15(9):871-875, 1997; Blomer *et al.*, *J Virol.*, 71(9):6641-6649, 1997; U.S. Pat. Nos. 6,013,516 and 5,994,136).

[00345] A wide variety of vectors can be used for the expression of a Factor VIII polypeptide from a codon-altered polypeptide in cell culture, including eukaryotic and prokaryotic expression vectors. In certain embodiments, a plasmid vector is contemplated for use in expressing a Factor VIII polypeptide in cell culture. In general, plasmid vectors containing replicon and control sequences which are derived from species compatible with the

host cell are used in connection with these hosts. The vector can carry a replication site, as well as marking sequences which are capable of providing phenotypic selection in transformed cells. The plasmid will include the codon-altered polynucleotide encoding the Factor VIII polypeptide, operably linked to one or more control sequences, for example, a promoter.

[00346] Non-limiting examples of vectors for prokaryotic expression include plasmids such as pRSET, pET, pBAD, etc., wherein the promoters used in prokaryotic expression vectors include lac, trc, trp, recA, araBAD, etc. Examples of vectors for eukaryotic expression include: (i) for expression in yeast, vectors such as pAO, pPIC, pYES, pMET, using promoters such as AOX1, GAP, GAL1, AUG1, etc; (ii) for expression in insect cells, vectors such as pMT, pAc5, pIB, pMIB, pBAC, etc., using promoters such as PH, p10, MT, Ac5, OpIE2, gp64, polh, etc., and (iii) for expression in mammalian cells, vectors such as pSVL, pCMV, pRc/RSV, pcDNA3, pBPV, etc., and vectors derived from viral systems such as vaccinia virus, adeno-associated viruses, herpes viruses, retroviruses, etc., using promoters such as CMV, SV40, EF-1, UbC, RSV, ADV, BPV, and β-actin.

IV. Examples

Example 1 – Construction of a Codon-Altered Factor VIII Variant Expression Sequence

[00347] Two hurdles had to be overcome in order to create a Factor VIII coding sequence that is effective for gene therapy of hemophilia A. First, because of the genomic size limitations of conventional gene therapy delivery vectors (e.g., AAV virions), the encoded Factor VIII polypeptide had to be shortened considerably. Second, the coding sequence had to be altered to: (i) stabilize packaging interactions within the delivery vector, (ii) stabilize the mRNA intermediary, and (iii) improve the robustness of transcription/translation of the mRNA.

[00348] To achieve the first objective, Applicants started with a B-domain deleted Factor VIII variant construct, referred to herein as "FVIII-BDD-SQ." In this construct, the B-domain is replaced with a fourteen amino acid sequence referred to as the "SQ" sequence. Recombinant FVIII-BDD-SQ is sold under the trade name REFACTO®, and has been shown to be effective for the management of hemophilia A. However, the native coding sequence for FVIII-BDD-SQ,

which includes human wild-type nucleic acid sequences for the Factor VIII heavy and light chains, is ineffectively expressed in gene therapy vectors.

[00349] To address the poor expression of the native FVIII-BDD-SQ, the codon optimization algorithm described in Fath et al. (PLoS ONE, 6:e17596 (2011)), modified as described in Ward et al. (Blood, 117:798 (2011)) and in McIntosh et al. (Blood, 121, 3335-3344 (2013)), was applied to the FVIII-BDD-SQ sequence to create first intermediate coding sequence CS04a. However, Applicants recognized that the CS04a sequence created using the modified algorithm could be improved by further modifying the sequence. Accordingly, Applicants re-introduced CpG dinucleotides, re-introduced the CGC codon for arginine, changed the leucine and serine codon distributions, re-introduced highly conserved codon pairs, and removed cryptic TATA box, CCAAT box, and splice site elements, while avoiding CpG islands and local overrepresentation of AT-rich and GC-rich stretches.

[00350] First, the modified algorithm systematically replaces codons containing CpG-dinucleotides (e.g., arginine codons) with non-CpG-dinucleotide codons, and eliminates/avoids CpG-dinucleotides created by neighboring codons. This strict avoidance of CpG dinucleotides is usually done to prevent TLR-induced immunity after intramuscular injection of DNA vaccines. However, doing so limits the codon optimization possibilities. For example, the modified algorithm excludes use of the complete set of CGX arginine codons. This is particularly disruptive in the coding of genes for expression in human cells, because CGC is the most frequently used arginine codon in highly expressed human genes. Additionally, avoiding the creation of CpGs by neighboring codons further limits the optimization possibilities (e.g., limits the number of codon pairs that may be used together).

[00351] Because TLR-induced immunity is not expected to be a problem associated with liver-directed, AAV-based gene therapy, codons including CpGs, and neighboring codons creating CpGs, were re-introduced into intermediate coding sequence CS04a, preferentially in the sequence coding for the Factor VIII light chain (e.g., at the 3' end of the FVIII-BDD-SQ coding sequence). This allowed for more frequent use of preferred human codons, particularly those for arginine. Care was taken, however, to avoid creation of CpG islands, which are regions of coding sequence having a high frequency of CpG sites. This is contrary to the teachings of

Krinner et al. (Nucleic Acids Res., 42(6):3551-64 (2014)), which suggests that CpG domains downstream of transcriptional start sites promote high levels of gene expression.

[00352] Second, the modified algorithm applies certain codons exclusively, such as CTG for leucine, GTG for valine, and CAG for glutamine. However, this offends the principles of balanced codon use, for example, as proposed in Haas et al. (Current Biology, 6(3):315-24 (1996)). To account for the overuse of preferred codons by the modified algorithm, alternate leucine codons were re-introduced where allowed by the other rules applied to the codon alteration (e.g., CpG frequency and GC content).

[00353] Third, the modified algorithm replaces codon pairs without regard to how conserved they are in nature, when certain criteria (e.g., the presence of CG-dinucleotides) are met. To account for beneficial properties which may have been conserved by evolution, the most conserved codon pairs that were replaced by the algorithm and the most conserved preferred codon pairs, e.g., as described in Tats et al. (BMC Genomics 9:463 (2008)), were analyzed and adjusted where allowed by the other rules applied to the codon alteration (e.g., CpG frequency and GC content).

[00354] Fourth, serine codons used in the intermediate coding sequence were also re-engineered. Specifically, AGC, TCC, and TCT serine codons were introduced into the modified coding sequence with higher frequency, to better match overall for human codon usage (Haas et al., supra).

[00355] Fifth, TATA box, CCAAT box elements, and intron/exon splice sites were screened and removed from the modified coding sequence. When modifying the coding sequence, care was taken to avoid local overrepresentation of AT-rich or GC rich stretches.

[00356] Finally, in addition to optimizing the codon usage within the coding sequence, the structural requirements of the underlying AAV virion were considered when further refining the intermediate coding sequence CS04a. AAV vectors (e.g., the nucleic acid portion of an AAV virion) are packaged as single stranded DNA molecules into their capsids (for review, see, Daya and Berns, Clin. Microbiol Rev., 21(4):583-93 (2008)). The GC content of the vector is therefore likely to influence packaging of the genome and, thus, vector yields during production.

Like many algorithms, the modified algorithm used here creates an optimized gene sequence with a GC content of at least 60% (*see*, Fath et al., PLoS One, 6(3):e17596 (2011) (erratum in: PLoS One, (6)3 (2011)). However, the AAV8 capsid protein is encoded by a nucleotide sequence having a lower GC content of about 56%. Thus, to better mimic the native AAV8 capsid protein coding sequence, the GC content of the intermediate coding sequence CS04a was reduced to 56%.

[00357] The resulting CS04 coding sequence, shown in Figure 2, has an overall GC content of 56%. The CpG-dinucleotide content of the sequence is moderate. However, CpG dinucleotides are predominantly present in the downstream portion of the coding sequence, e.g., the portion coding for the Factor VIII light chain. The CS04 sequence has 79.77% nucleotide sequence identity to the corresponding coding sequences in wild-type Factor VIII (Genbank accession M14113).

[00358] For comparison purposes, several other codon-optimized, ReFacto constructs were prepared. CS01 was constructed by applying the codon-optimization algorithm of Fath et al., as modified by Ward et al., as done for CS04. However, unlike CS04, the CS01 construct does not contain any CpG islands. The CS08 ReFacto construct was codon-optimized as described in Radcliff P.M. et al., Gene Therapy, 15:289-97 (2008), the content of which is hereby expressly incorporated by reference herein, in its entirety, for all purposes. The CS10 codon-optimized ReFacto construct was obtained from Eurofins Genomics (Ebersberg, Germany). The CS11 codon-optimized ReFacto construct was obtained from Integrated DNA Technologies, Inc. (Coralville, USA). The CH25 codon-optimized ReFacto construct was obtained from ThermoFischer Scientific's GeneArt services (Regensburg, Germany). The CS40 ReFacto construct consists of the wild type Factor VIII coding sequence. The algorithm used to construct CS23 is based on the JCAT tool (www.jcat.de), an on-line tool for codon-optimizations (Grote et al., 2005; Nucl. Acids Res. W526-31). The sequence was further modified to more reflect the codon usage of the albumin superfamily (Mirsafian et al. 2014: Sc. Word Journal 2014, ID 639682). The sequence identities shared between each of the ReFacto coding sequences is shown in **Table 2**, below.

Table 2 – Percent identity matrix for codon-altered Factor VIII constructs.

	CS01	**CS04**	**CS08**	**CS10**	**CS11**	**CS40**	**CH25**	**CS23**
CS01	100%							
CS04	93.0%	100%						
CS08	80.7%	82.2.%	100%					
CS10	79.1%	79.4%	78.4%	100%				
CS11	78.3%	78.3%	78.1%	77.5%	100%			
CS40	79.6%	79.8%	76.7%	77.6%	75.4%	100%		
CH25	81.3%	85.1%	85.0%	79.9%	79.4%	75.8%	100%	
CS23	84.3%	89.2%	85.1%	80.3%	79.9	76.5%	93.2%	100%

[00359] Plasmids of each construct were constructed by cloning different synthetic DNA fragments into the same vector backbone plasmid (pCh-BB01). DNA synthesis of the Refacto-type BDD-FVIII fragments with flanking AscI and NotI enzyme restriction sites were done by ThermoFischer Scientific (Regensburg, Germany). The vector backbone contains two flanking AAV2-derived inverted terminal repeats (ITRs) that encompass a promoter/ enhancer sequence derived from the liver-specific murine transthyretin gene, AscI and NotI enzyme restriction sites for insertion of the respective Refacto-type BDD-FVIII and a synthetic polyA site. After ligation of the prepared vector backbone and inserts via the AscI and NotI sites, the resulting plasmids were amplified in milligram scale. The Refacto-type BDD-FVIII sequences of the constructs were verified by direct sequencing (Microsynth, Balgach, Switzerland). The cloning resulted in seven different plasmid constructs named pCS40, pCS01, pCS04, pCS08, pCS10, pCS11, and pCh25 (Figure 23). The constructs have the same vector backbone and encode the same B-domain deleted FVIII protein (Refacto-type BDD-FVIII), but differ in their FVIII coding sequence.

[00360] AAV8-based vectors were prepared by the three plasmid transfection method, as described in Grieger JC, et al. (Virus Vectors Using Suspension HEK293 Cells and Continuous Harvest of Vector From the Culture Media for GMP FIX and FLT1 Clinical Vector, Mol Ther., Oct 6. (2015) doi: 10.1038/mt.2015.187. [Epub ahead of print]), the content of which is hereby expressly incorporated by reference herein, in its entirety, for all purposes. HEK293 suspensions cells were used for plasmid transfections using the corresponding FVIII vector plasmid, the

helper plasmid pXX6-80 (carrying adenoviral helper genes), and the packaging plasmid pGSK2/8 (contributing the rep2 and cap8 genes). To isolate the AAV8 constructs, the cell pellets of one liter cultures were processed using iodixanol gradients, as described in Grieger et al. (2015, Supra). The procedure resulted in vector preparations called vCS01, vCS04, vCS08, vCS10, vCS11, and vCH25. Vectors were quantified by qPCR using the universal qPCR procedure targeting the AAV2 inverted terminal repeats (Aurnhammer, Human Gene Therapy Methods: Part B 23:18–28 (2012)). A control vector plasmid carrying AAV2 inverted terminal repeats served for preparing the standard curve. The resulting vCS04 construct is presented as SEQ ID NO: 8 in Figures 7A-7C.

[00361] The integrity of the vector genomes was analyzed by AAV agarose gel electrophoresis. The electrophoresis was performed as described in Fagone et al., Human Gene Therapy Methods 23:1-7 (2012). Briefly, AAV vector preparations were incubated at 75 °C for 10 minutes in the presence of 0.5% SDS and then cooled down to room temperature. Approximately 1.5E10 vector genomes (vg) were loaded per lane on a 1% 1xTAE agarose gel and electrophoresed for 60 min at 7 V/cm of gel length. The gel was then stained in 2x GelRed (Biotium Cat# 41003) solution and imaged by ChemiDocTMMP (Biorad). The results shown in Figure 24 demonstrate that the vCS01, vCS04, and vCS40 viral vectors have the same-sized genome, indicated by a distinct band in the 5kb range (Figure 24, lanes 2-4). Despite a vector size of approx. 5.2 kb, the genome is a homogenous band confirming correct packaging of the somewhat oversized genome (relative to an AAV wild-type genome of 4.7 kb). All other vCS vector preparations show the same genomic size (data not shown).

[00362] In order to confirm the expected pattern of capsid proteins, SDS PAGE followed by silver staining was performed with the vectors vCS01, vCS04, and vCS40 (Figure 25). As shown in the figure, the downstream purification procedure resulted in highly purified material displaying the expected protein pattern of VP1, VP2 and VP3 (Figure 25, lanes 2-4). The same pattern was seen with all other viral preparations (not shown). The SDS-PAGE procedure of AAV preparations was done according to standard procedures. Each lane contained 1E10 vg of the respective viral construct, and were separated on a 4-12% Bis-Tris (NuPAGE® Novex, Life Technologies) gel as per manufacturer's instructions. Silver staining was performed with a SilverQuestTM kit (Novex, Life Technologies) according to the manufacturer's instructions.

[00363] Surprisingly, AAV vectors vCS01 and vCS04 had higher virion packaging, measured by higher yields in AAV virus production, as compared to the vCS40 wild-type coding construct and the other codon-optimized constructs. As shown in **Table 3**, the vCS01 and vCS04 vectors replicated substantially better than vCS40, providing a 5-7 fold yield increase in AAV titer.

Table 3 – Yields per liter cell culture obtained with AAV vector constructs vCS01, vCS04, and vCD40, as purified from cell pellets.

Construct	**Vector concentration [vg/ml] x10E12**	**Yields [vg /liter] x10E12**	**Fold increase vs wt**
vCS40	2.0	11.0	-
vCS01	9.2	51.4	4.7
vCS04 – Sample 1	17.6	79.2	7.2
vCS04 – Sample 2	15.9	58.8	5.4

Example 2 – In Vivo Expression of Codon-Altered Factor VIII Variant Expression Sequences

[00364] To test the biological potency of the codon-altered Factor VIII variant sequences, the ReFacto-type FVIII constructs described in Example 1 were administered to mice lacking Factor VIII. Briefly, the assays were performed in C57Bl/6 FVIII knock-out (ko) mice (with 6-8 animals per group) by tail vein injection of 4E12 vector genomes (vg) per kilogram body weight of mouse. Blood was drawn 14 days after injection by retroorbital puncture and plasma was prepared and frozen using standard procedures. Expression levels at day 14 were chosen because there is minimal influence of inhibitory antibodies at this time, which are seen in some animals of this mouse model at later times. FVIII activity in the mouse plasma was determined using the Technochrome FVIII assay performed, with only minor modifications, as suggested by the manufacture (Technoclone, Vienna, Austria). For the assay, the plasma samples were appropriately diluted and mixed with assay reagents, containing thrombin, activated factor IX (FIXa), phospholipids, factor X and calcium. Following FVIII activation by thrombin a complex with FIXa, phospholipids and calcium is formed. This complex activates FX to activated FX (FXa) which in turn cleaves para-nitroanilide (pNA) from the chromogenic substrate. The

kinetics of pNA formation is measured at 405 nm. The rate is directly proportional to the FVIII concentration in the sample. FVIII concentrations are read from a reference curve and results are given in IU FVIII/milliliter.

[00365] The results, presented in **Table 4** below, demonstrate that the codon-altered sequences designed using commercial algorithms (CS10, CS11, and CH25) provided only a modest increase in BDD-Factor VIII (3-4 fold) as compared to the wild-type BDD-Factor VIII construct (CS40). Similarly, the codon-altered BDD-Factor VIII construct prepared as described in Radcliffe et al. (CS08), only provided a 3-4 fold increase in BDD-FVIII expression. This result is consistent with the results reported in Radcliff et al. Surprisingly, the CS01, CS04, and CS23 constructs provided much higher BDD-FVIII expression in the in-vivo biopotency assays (18-, 74-, and -30-fold increases, respectively).

Table 4 – Expression of FVIII in the plasma of FVIII-knock-out mice induced by the different AAV vector constructs.

Construct	**Codon Algorithm**	**Average FVIII Expression at Day 14 [IU/ml]**	**Standard deviation**	**Number of mice**	**Fold increase vs wt**
vCS40	Human wild-type	0.03	0.03	12	-
vCS01	Applicants'	0.55	0.28	22	18.3
vCS04	Applicants'	2.21	1.20	55	73.7
vCS08	Radcliffe et al.	0.11	0.01	6	3.6
vCS10	Eurofins	0.09	0.01	7	3.0
vCS11	IDT	0.08	0.02	8	2.7
vCH25	GeneArt	0.13	0.12	18	4.3
vCS23	Applicants'	0.91	0.32	5	30.3

Example 3 – Design of Glycosylation Peptides for the B-domain Substituted Linker

[00366] Others have shown that inclusion of a small peptide (the "V3 peptide") containing six putative N-linked glycosylation sites from the wild-type Factor VIII B-domain, into a B-domain deleted gene therapy construct, increased Factor VIII levels in the plasma of mice

(McIntosh et al., Blood 121(17):3335-44 (2013)). However, in order to maintain the small size of the B-domain substituted linker, the glycosylation sites were taken out of the context of the wild-type B-domain. In silico prediction (Gupta et al., Supra) of the linker containing the V3 peptide suggests that only two of these glycosylation sites in the V3 peptide will be modified in vivo (Figure 15).

[00367] Thus, Applicants attempted to identify alternative glycosylation peptides that would support higher levels of glycosylation in vivo, which matched wild type glycosylation more closely than the V3 peptide. Applicants designed and tested several alternative glycosylation peptides, in silico. Several of these peptides, shown in Figures 13A-13B, were predicted to have equal or greater glycosylation in vivo than the V3 peptide, when placed between amino acids N768 and P769 of the B-domain substituted linker in SEQ ID NO:2. The results of the in silico predictions are shown in **Table 5**, below. **Table 5** also reports the results of expression experiments performed for several constructs encoding a ReFacto-type Factor VIII protein with a glycosylation peptide incorporated into the B-domain substituted linker, in a CS01 codon-optimized background.

Table 5 – Prediction of N-glycosylation in B-domain substituted linker peptides and performance of AAV vector constructs in vivo.

Sequence	Number of Predicted N-glycosylation sites	Day 28 expression [IU/ml]	SD	Number of mice [n]	Fold expression
vCS01	0	0.74	0.52	5	21
vNG1/CS01	4	n.d.	-	-	-
vNG4/CS01	**3**	**1.93**	**0.57**	**6**	**55**
vNG5/CS01	2	n.d.	-	-	-
vNG6/CS01	1	0.80	0.67	5	23
vNG9/CS01	1	n.d.	-	-	-
vNG10/CS01	**2**	**2.66**	**0.52**	**6**	**76**
vNG16/CS01	**2**	**1.59**	**0.57**	**6**	**45**
vNG17/CS01	2	n.d.	-	-	-
vNG18/CS01	2	n.d.	-	-	-
vNG19/CS01	2	0.88	0.25	5	25
vNG20/CS01	2	n.d.	-	-	-

vNG21/CS01	2	n.d.	-	-	-
vCS40	0	0.035	0.030	12	1

[00368] AAV vectors containing the NG variants were constructed as described in Example 1 and tested in FVIII knock-out mice as described in Example 2. All virus vectors (except the control vector vCS40) shown in **Table 5** are based on the algorithm as used in vCS01. A parallel set of constructs using the algorithm of vCS04 was also prepared (vNG/CS04 series) and is tested in the mouse model. Results were compared to the expression levels achieved with the wild-type vCS40 construct. The day 28 expression levels were chosen in this example, because expression levels of the majority of construct reached the highest levels at this time point. Three AAV vectors achieved greater than 40-fold FVIII expression levels including vNG4/CS01, vNG10/CS01 and vNG16/CS01 (**Table 5**). The corrresponding constructs vNG4/CS04, vNG10/CS04 and vNG16/CS04 are expected to show even higher expression because they are based on the superior vCS04 algorithm.

[00369] Surprisingly, the AAV vectors of the vNG/CS01 series had higher virion packaging, measured by higher yields in AAV virus production, as compared to the vCS40 wild-type coding construct. As shown in **Table 6**, the vNG/CS01-based vectors replicated substantially better than vCS40, providing an approximately 3- fold yield increase in AAV titer.

[00370] **Table 6** – Yields per liter cell culture obtained with AAV vector constructs as purified from cell pellets.

Sequence	**Vector conc. [vg/ml] x10^{12}**	**Yields [vg/liter] x10^{12}**	**Fold increase vs wild-type**
vCS01	9.17	51.35	4.7
vNG1/CS01	2.13	17.04	1.5
vNG4/CS01	5.74	33.01	3.0
vNG5/CS01	6.91	27.29	2.5
vNG6/CS01	7.01	40.66	3.7
vNG9/CS01	6.39	29.39	2.7
vNG10/CS01	8.57	37.71	3.4
vNG16/CS01	5.3	28.36	2.6

vNG17/CS01	4.24	32.22	2.9
vNG18/CS01	6.11	37.88	3.4
vNG19/CS01	9.42	39.56	3.6
vNG20/CS01	4.09	30.27	2.8
vNG21/CS01	n.d	-	-
vCS40	2.03	11	1.0

Example 4 – Construction of Mutant BDD-FVIII constructs

[00371] Numerous different mutated Refacto-type BDD-FVIII constructs, carrying amino acid mutations within the Factor VIII heavy chain and/or B-domain substituted linker, were cloned and screened. The corresponding vectors, as referred to herein as the "vCS" series of vectors, encode BDD-FVIII variants in the CS01, CS04, and CS23 codon-altered backgrounds. The method used to construct the CS01 and CS04 backgrounds is described in Example 1. The method used to construct CS23 was based on the JCAT tool (www.jcat.de), an on-line tool for codon-optimizations (Grote et al., 2005; Nucl. Acids Res. W526-31). The sequence was further modified to better reflect the codon usage of the albumin superfamily (Mirsafian et al., Sc. Word Journal, ID 639682 (2014)), the content of which is hereby expressly incorporated by reference, in its entirety, for all purposes.

[00372] Combinations of three types of mutations were included in the FVIII sequences of the vCS series of constructs. The first amino acid change introduced into the FVIII sequence is the X1 mutation (TTYVNRSL (SEQ ID NO: 33); X. Xiao), which introduces an additional glycosylation site near the B-domain substituted linker. The X1 mutation is also referred to herein as the "m3" mutation. The second amino acid change made in the FVIII sequence includes the F328S (SPI, F309S SPE) mutation, an amino acid change known to improve secretion of FVIII (Swaaroop, J. Biol. Chem., 272:24121–24 (1997)). This mutation is also referred to herein as the "m1" mutation. The third change is the so-called X5 mutation, which is a combination of five amino acid changes in the A1 domain of the heavy chain that improves specific activity and secretion of BDD-FVIII (Cao et al., 2014; ASGCT abstract #460; details of mutations disclosed in oral presentation). The X5 mutation is also referred to herein as the "m2" mutation. Next, combinations of X1 and F328S (SPI, F309S SPE) were made, followed by

combinations of X1 and X5, also referred to as "X6," and yet other combinations of X5 and F328S (SPI, F309S SPE) were made (**Table 7**).

[00373] **Gene synthesis and cloning of the vector plasmids.** The plasmids were constructed by cloning different synthetic DNA fragments into the same vector backbone plasmid (pCh-BB01). DNA synthesis of the Refacto-type BDD-FVIII fragments with flanking *Asc*I and *Not*I enzyme restriction sites were done by ThermoFischer Scientific (Regensburg, Germany). The vector backbone contains two flanking AAV2-derived inverted terminal repeats (ITRs) that encompass a promoter/enhancer sequence derived from the liver-specific murine transthyretin gene, *Asc*I and *Not*I enzyme restriction sites for insertion of the respective Refacto-type BDD-FVIII, and a synthetic polyA site. After ligation of the prepared vector backbone and insertions via the *Asc*I and *Not*I sites, the resulting plasmids were amplified in milligram scale. The Refacto-type BDD-FVIII sequences of the constructs were verified by direct sequencing (Microsynth, Balgach, Switzerland). The cloning resulted in different plasmid constructs, as shown in Figure 44.

[00374] **Small scale vector preparations and quantification by quantitative PCR (qPCR).** AAV8-based vectors were prepared by the three plasmid transfection method essentially as described in Grieger et al. (2015, Supra). HEK293 suspensions cells were used for plasmid transfections using the corresponding FVIII vector plasmid, the helper plasmid pXX6X80 (carrying adenoviral helper genes) and the packaging plasmid pGSK2/8 (contributing the rep2 and cap8 genes). In the downstream process the cell pellet of a one liter culture was processed using iodixanol gradients as described above. The procedure resulted in vector preparations as outlined in **Table 8**. Vectors were quantified by qPCR using the universal qPCR procedure targeting the AAV2 inverted terminal repeats (Aurnhammer, HUMAN GENE THERAPY METHODS: Part B 23:18–28 (2012)). An accurately quantified vector plasmid carrying AAV2 Inverted terminal repeats served for preparing the standard curve.

[00375] **AAV vector characterizations**. The integrity of the vector genome was analyzed by AAV agarose gel electrophoresis. The electrophoresis was done similar as described in Fagone et al. (Human Gene Therapy Methods, 23:1-7 (2012)). AAV vector preparations were incubated at 75 °C for 10 minutes in the presence of 0.5% SDS and then cooled down to room

temperature. Approximately 1.5E10 vector genomes (vg) were loaded per lane on a 1% 1xTAE agarose gel and electrophoresed for 60 min at 7 V/cm of gel length. The gel was then stained in 2x GelRed (Biotium Cat# 41003) solution and imaged by ChemiDoc™MP (Biorad). The results of a selection of vectors are shown in Figure 45. The viral vectors vCS04 (control), vCS17, vCS20, vCS24, vCS16 and vCS40 (control) show all the same-sized genome as a distinct band in the 5kb range (Figure 45, lanes 2-7; arrow right side). Despite a vector size of approx. 5.2 kb, the genome is a homogenous band confirming correct packaging of the somewhat oversized genome (relative to an AAV wild-type genome of 4.7 kb).

[00376] In order to confirm purity of the vector and the expected pattern of capsid proteins, SDS PAGE followed by silver staining was performed with the vectors, as shown in Figure 46. As shown in the figure, the downstream purification procedure resulted in highly purified material displaying the expected protein pattern of VP1, VP2 and VP3 (Figure 46 lanes 2-9; arrows right hand side). The SDS-PAGE procedure of AAV preparations was done according to standard procedures. The amounts of 1E10 vg per lane were separated on a 4-12% Bis-Tris (NuPAGE® Novex, Life Technologies) gel as per manufacturer's instructions. Silver staining was performed with a SilverQuest™ kit (Novex, Life Technologies) according to the instructions of the manufacturer.

[00377] **In-vivo biopotency screening of vectors.** The different Refacto-type BDD-FVIII constructs were screened in mice. The assay was performed in C57Bl/6 FVIII knock-out (ko) mice (with 6-8 animals per group) by tail vein injection of 4E12 vector genomes (vg) per kilogram body weight of mouse. Blood was drawn 14 days after injection by retroorbital puncture and plasma was prepared and frozen using standard procedures. FVIII activity in mouse plasma was determined with a chromogenic assay from Technoclone with minor modifications (Technochrome FVIII, Technoclone, Vienna, Austria). In brief, the plasma sample was appropriately diluted and mixed with assay reagents, containing thrombin, activated factor IX (FIXa), phospholipids, factor X and calcium. Following FVIII activation by thrombin a complex with FIXa, phospholipids and calcium is formed. This complex activates FX to activated FX (FXa) which in turn cleaves para-nitroanilide (pNA) from the chromogenic substrate. The kinetics of pNA formation is measured at 405 nm. The rate is directly

proportional to the FVIII concentration in the sample. FVIII concentrations are read from a reference curve and results are given in IU FVIII/milliliter.

[00378] The results of the mouse biopotency assay (day 14 expression data of FVIII in international units per milliliter [IU/ml] in mouse plasma and fold expression compared to the wild-type vCS40 control) are shown in **Table 7.** AAV vectors vCS19, vCS26 and vCS32 all contain the X1 glycosylation site in the CS01, CS04, and CS23 codon-altered backgrounds, respectively. As seen in **Table** 7, surprisingly high expression levels were obtained, as compared to the wild-type construct vCS40 (level defined as 1). vCS26, for instance, expressed 202-fold higher levels compared to the wild-type vCS40 vector. Another control construct for the X1-series of vectors, vCH111, that contains the X1 mutation in the Geneart codon context, showed a more modest increase in expression (12-fold).

[00379] **Vectors** vCS16, vCS28, and vCS34 all contain the F328S (SPI, F309S SPE) mutation enhancing secretion in the CS01, CS04, and CS23 codon-altered backgrounds, respectively. As seen in **Table** 7, high expression levels (45-93-fold higher than the wt vCS40 control) were obtained with vCS16 and vCS28.

[00380] **Vectors** vCS20, vCS24, and vCS33 contain the X5 mutation in the CS01, CS04, and CS23 codon-altered backgrounds, respectively. The best performing variant in the X5 series was vCS20, achieving levels of >3 units/ml after day 14 and a 121-fold increase over the wt vCS40 control.

[00381] **Vectors** vCS17, vCS29, and vCS31 contain the combination of the X1 and F328S (SPI, F309S SPE) mutations in the CS01, CS04, and CS23 codon-altered backgrounds, respectively (Table 6). The vCS17 and vCS29 constructs achieved very high expression levels in the mouse studies (115 to 246-fold increase over the vCS40 control). Remarkably, in the FVIII KO mouse model used, the majority of mice treated with the vCS17 construct did not develop neutralizing antibodies over time, evidenced by increasing levels of FVIII at later time points (e.g., day 28 and day 42; data not shown). This is an unexpected finding, because in some other constructs the expression levels began to decrease with time due to the formation of neutralizing antibodies. The CS01 background combined with the secretion-enhancing mutations F328S (SPI, F309S SPE) and X1 resulted in low immunogenicity induction.

[00382] **Vectors** vCS18, vCS27, and vCS35 contain the combination of the X1 and X5 mutations in the CS01, CS04, and CS23 codon-altered backgrounds, respectively. The combination of these two mutations was also very efficient. A 145-fold increase over the vCS40 control could be achieved with vCS18, for example (**Table** 7).

[00383] Vectors vCS48 and vCS49 contain the combination of the X5 and F328S (SPI, F309S SPE) mutations in the CS01 and CS04 codon-altered backgrounds, respectively. The combination of these two mutations was also very efficient. One of the largest increases of all mutants, a 239-fold increase over the vCS40 control, could be achieved with vCS49 confirming the special value of the combinations including the F328S (SPI, F309S SPE) mutation.

[00384] A further surprising observation was that the mutant AAV vectors grew substantially better than the vCS40 construct harboring the wild-type BDD-FVIII codons. Sequence optimization resulted in a several-fold yield increase in vector production. In some of the best expressing constructs (e.g., vCS29, vCS17, vCS20, and vCS26) the increase in yields due to codon-alteration and/or mutant sequence was approximately 3-5-fold higher, as compared to the wild-type vector (**Table 8**).

[00385] Expression of BDD-FVIII in the plasma of FVIII-knock-out mice induced by the different AAV vector constructs is shown in **Table** 7. The constructs have the same vector backbone, however, encode different types of mutated FVIII, including different codon optimization backgrounds. Expression levels at day 14 were chosen because at this time point there is minimal influence of inhibitory antibodies usually seen in some animals in the mouse model at later times. N.d., not determined.

Table 7 – In vivo biopotency data of vCS constructs.

#	Vector	Algorithm, mutations	Day 14 expression [IU/ml]	SD	Number of mice [n]	Fold expression
1	vCS19	CS01, X1	2.34	1.10	13	78
2	vCS26	CS04, X1	6.07	2.72	12	202
3	vCS32	CS23, X1	n.d.	-	-	-
4	vCS16	CS01, F328S	1.35	0.88	6	45
5	vCS28	CS04, F328S	2.78	0.92	7	93
6	vCS34	CS23, F328S	n.d.	-	-	-
7	vCS20	CS01, X5	3.62	1.96	21	121
8	vCS24	CS04, X5	0.79	0.89	18	26
9	vCS33	CS23, X5	n.d.	-	-	n.d.
10	vCS17	CS01, X1, F328S	3.44	1.92	20	115
11	vCS29	CS04, X1, F328S	7.39	2.64	9	246
12	vCS31	CS23, X1, F328S	n.d.			n.d.
13	vCS18	CS01, X1+X5 (X6)	4.34	2.50	6	145
14	vCS27	CS04, X1+X5 (X6)	8.03	3.97-	6-	268-
15	vCS35	CS23, X1+X5 (X6)	n.d.	-	-	-
19	vCS48	CS01, X5, F328S	2.54	0.72	8	85
20	vCS49	CS04, X5, F328S	7.17	1.30	7	239
	controls					
16	vCS40	Human wild-type	0.03	0.03	12	1
17	vCh25	Geneart	0.13	0.12	18	4
18	vCh111	Geneart +X1	0.37	0.21	17	12

Table 8 – Yields per liter cell culture (packaging efficiency) obtained with the different AAV vector constructs. The vectors were purified out of the cell pellets; n.d., not determined.

	construct	Algorithm, mutations	Vector conc. [vg/ml] $x10^{12}$	Yields [vg /liter] $x10^{12}$	Fold increase vs wt
1	vCS19	CS01, X1	9.71	36	3.22
2	**vCS26**	CS04, X1	5.93	32	2.87
3	vCS32	CS23, X1	n.d.	n.d.	n.d.
4	vCS16	CS01, F328S	6.51	29	2.56
5	vCS28	CS04, F328S	5.85	32	2.88
6	vCS34	CS23, F328S	n.d.	n.d.	n.d.
7	**vCS20**	CS01, X5	9.90	50	4.48
8	vCS24	CS04, X5	3.00	16	1.46
9	vCS33	CS23, X5	n.d.	n.d.	n.d.
10	**vCS17**	CS01, X1, F328S	8.94	37	3.34
11	**vCS29**	CS04, X1, F328S	7.42	53	4.72
12	vCS31	CS23, X1, F328S	n.d.	n.d.	n.d.
13	vCS18	CS01, X1+X5 (X6)	21.20	53	4.75
14	vCS27	CS04, X1+X5 (X6)	4.15	19	1.67
15	vCS35	CS23, X1+X5 (X6)	n.d.	n.d.	n.d.
16	vCS48	CS01, X5, F328S	7.14	42.1	3.77
17	vCS49	CS04, X5, F328S	8.27	37.2	3.33
18	vCS40	Human wild-type	2.03	11	1.00

[00386] It is understood that the examples and embodiments described herein are for illustrative purposes only and that various modifications or changes in light thereof will be suggested to persons skilled in the art and are to be included within the spirit and purview of this application and scope of the appended claims. All publications, patents, and patent applications cited herein are hereby incorporated by reference in their entirety for all purposes.

WHAT IS CLAIMED IS:

1. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-FL-NA (SEQ ID NO: 13), wherein the polynucleotide encodes a Factor VIII polypeptide.

2. The polynucleotide of claim 1, wherein the nucleotide sequence has at least 99% identity to CS01-FL-NA (SEQ ID NO: 13).

3. The polynucleotide of claim 1, wherein the nucleotide sequence is CS01-FL-NA (SEQ ID NO: 13).

4. A polynucleotide comprising a nucleotide sequence encoding a Factor VIII polypeptide, the Factor VIII polypeptide comprising a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain,

wherein the heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS01-HC-NA (SEQ ID NO: 24);

wherein the light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS01-LC-NA (SEQ ID NO: 25); and

wherein the polypeptide linker comprises a furin cleavage site.

5. The polynucleotide of claim 4, wherein the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO01 (SEQ ID NO: 5).

6. The polynucleotide according to claim 4 or 5, wherein:

the first nucleotide sequence has at least 99% identity to CS01-HC-NA (SEQ ID NO: 24); and

the second nucleotide sequence has at least 99% identity to CS01-LC-NA (SEQ ID NO: 25).

7. The polynucleotide according to claim 4 or 5, wherein:

the first nucleotide sequence is CS01-HC-NA (SEQ ID NO: 24); and

the second nucleotide sequence CS01-LC-NA (SEQ ID NO: 25).

8. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-FL-NA (SEQ ID NO:1), wherein the polynucleotide encodes a Factor VIII polypeptide.

9. The polynucleotide of claim 4, wherein the nucleotide sequence has at least 99% identity to CS04-FL-NA (SEQ ID NO: 1).

10. The polynucleotide of claim 4, wherein the nucleotide sequence is CS04-FL-NA (SEQ ID NO: 1).

11. A polynucleotide comprising a nucleotide sequence encoding a Factor VIII polypeptide, the Factor VIII polypeptide comprising a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain,

wherein the heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS04-HC-NA (SEQ ID NO: 3);

wherein the light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS04-LC-NA (SEQ ID NO: 4); and

wherein the polypeptide linker comprises a furin cleavage site.

12. The polynucleotide of claim 11, wherein the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO04 (SEQ ID NO: 6).

13. The polynucleotide according to claim 11 or 12, wherein:

the first nucleotide sequence has at least 99% identity to CS04-HC-NA (SEQ ID NO: 3); and

the second nucleotide sequence has at least 99% identity to CS04-LC-NA (SEQ ID NO: 4).

14. The polynucleotide according to claim 11 or 12, wherein:

the first nucleotide sequence is CS04-HC-NA (SEQ ID NO: 3); and

the second nucleotide sequence CS04-LC-NA (SEQ ID NO: 4).

15. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-FL-NA, wherein the polynucleotide encodes a Factor VIII polypeptide.

16. The polynucleotide of claim 15, wherein the nucleotide sequence has at least 99% identity to CS23-FL-NA (SEQ ID NO: 20).

17. The polynucleotide of claim 15, wherein the nucleotide sequence is CS23-FL-NA (SEQ ID NO: 20).

18. A polynucleotide comprising a nucleotide sequence encoding a Factor VIII polypeptide, the Factor VIII polypeptide comprising a light chain, a heavy chain, and a polypeptide linker joining the C-terminus of the heavy chain to the N-terminus of the light chain,

wherein the heavy chain of the Factor VIII polypeptide is encoded by a first nucleotide sequence having at least 95% identity to CS23-HC-NA (SEQ ID NO: 22);

wherein the light chain of the Factor FVIII polypeptide is encoded by a second nucleotide sequence having at least 95% identity to CS23-LC-NA (SEQ ID NO: 23); and

wherein the polypeptide linker comprises a furin cleavage site.

19. The polynucleotide of claim 18, wherein the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to BDLO23 (SEQ ID NO: 7).

20. The polynucleotide according to claim 18 or 19 wherein:

the first nucleotide sequence has at least 99% identity to CS23-HC-NA (SEQ ID NO: 22); and

the second nucleotide sequence has at least 99% identity to CS23-LC-NA (SEQ ID NO: 23).

21. The polynucleotide according to claim 18 or 19 wherein:

the first nucleotide sequence is CS23-HC-NA (SEQ ID NO: 22); and

the second nucleotide sequence CS23-LC-NA (SEQ ID NO: 23).

22. The polynucleotide according to any one of claims 4, 11, and 18, wherein the encoded polypeptide linker comprises a glycosylation peptide with an amino acid sequence having at least 92% identity to a glycosylation peptide selected from the group consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

23. The polynucleotide according to any one of claims 4, 11, and 18, wherein the encoded polypeptide linker comprises a glycosylation peptide selected from the group consisting of NG1-AA, NG4-AA, NG5-AA, NG6-AA, NG7-AA, NG9-AA, NG10-AA, NG16-AA, NG17-AA, NG18-AA, NG19-AA, NG20-AA, NG21-AA and NGV-AA.

24. The polynucleotide according to claim 22 or 23, wherein the glycosylation peptide is encoded by a polynucleotide with a nucleotide sequence having at least 95% identity to a sequence selected from the group consisting of NG1-NA, NG4-NA, NG5-NA, NG6-NA, NG7-NA, NG9-NA, NG10-NA, NG16-NA, NG17-NA, NG18-NA, NG19-NA, NG20-NA, NG21-NA and NGV-NA.

25. The polynucleotide according to claim 22 or 23, wherein the glycosylation peptide is encoded by a polynucleotide with a nucleotide sequence selected from one of NG1-NA, NG4-NA, NG5-NA, NG6-NA, NG7-NA, NG9-NA, NG10-NA, NG16-NA, NG17-NA, NG18-NA, NG19-NA, NG20-NA, NG21-NA and NGV-NA.

26. The polynucleotide according to any one of claims 4, 11, and 18, wherein the polypeptide linker is encoded by a third nucleotide sequence having at least 95% identity to a sequence selected from the group consisting of BDLNG1-NA, BDLNG3-NA, BDLNG5-NA, BDLNG6-NA, BDLNG9-NA, BDLNG10-NA, BDLNG16-NA, BDLNG17-NA, BDLNG18-NA, BDLNG19-NA, BDLNG20-NA and BDLNG21-NA.

27. The polynucleotide according to any one of claims 1 to 21, wherein the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 95% identity to CS04-FL-AA (SEQ ID NO: 2).

28. The polynucleotide according to any one of claims 1 to 21, wherein the polynucleotide encodes a Factor VIII polypeptide comprising the amino acid sequence of CS04-FL-AA (SEQ ID NO: 2).

29. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-SC1-NA (SEQ ID NO: 26), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

30. The polynucleotide of claim 29, wherein the nucleotide sequence has at least 99% identity to CS01-SC1-NA (SEQ ID NO: 26).

31. The polynucleotide of claim 29, wherein the nucleotide sequence is CS01-SC1-NA (SEQ ID NO: 26).

32. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-SC1-NA (SEQ ID NO: 9), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

33. The polynucleotide of claim 29, wherein the nucleotide sequence has at least 99% identity to CS04-SC1-NA (SEQ ID NO: 9).

34. The polynucleotide of claim 29, wherein the nucleotide sequence is CS04-SC1-NA (SEQ ID NO: 9).

35. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-SC1-NA (SEQ ID NO:28), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

36. The polynucleotide of claim 29, wherein the nucleotide sequence has at least 99% identity to CS23-SC1-NA (SEQ ID NO:28).

37. The polynucleotide of claim 29, wherein the nucleotide sequence is CS23-SC1-NA (SEQ ID NO:28).

38. The polynucleotide according to any one of claims 29 to 37, wherein the polynucleotide encodes a single-chain Factor VIII polypeptide comprising an amino acid sequence having at least 95% identity to CS01-SC1-AA (SEQ ID NO: 10).

39. The polynucleotide according to any one of claims 29 to 37, wherein the polynucleotide encodes a single-chain Factor VIII polypeptide comprising the amino acid sequence of CS01-SC1-AA (SEQ ID NO: 10).

40. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS01-SC2-NA (SEQ ID NO: 27), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

41. The polynucleotide of claim 40, wherein the nucleotide sequence has at least 99% identity to CS01-SC2-NA (SEQ ID NO: 27).

42. The polynucleotide of claim 40, wherein the nucleotide sequence is CS01-SC2-NA (SEQ ID NO: 27).

43. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS04-SC2-NA (SEQ ID NO: 11), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

44. The polynucleotide of claim 43, wherein the nucleotide sequence has at least 99% identity to CS04-SC2-NA (SEQ ID NO: 11).

45. The polynucleotide of claim 43, wherein the nucleotide sequence is CS04-SC2-NA (SEQ ID NO: 11).

46. A polynucleotide comprising a nucleotide sequence having at least 95% identity to CS23-SC2-NA (SEQ ID NO:29), wherein the polynucleotide encodes a single-chain Factor VIII polypeptide.

47. The polynucleotide of claim 43, wherein the nucleotide sequence has at least 99% identity to CS23-SC2-NA (SEQ ID NO:29).

48. The polynucleotide of claim 43, wherein the nucleotide sequence is CS23-SC2-NA (SEQ ID NO:29).

49. The polynucleotide according to any one of claims 40 to 48, wherein the polynucleotide encodes a single-chain Factor VIII polypeptide comprising an amino acid sequence having at least 95% identity to CS01-SC2-AA (SEQ ID NO: 12).

50. The polynucleotide according to any one of claims 40 to 48, wherein the polynucleotide encodes a single-chain Factor VIII polypeptide comprising the amino acid sequence of CS01-SC2-AA (SEQ ID NO: 12).

51. A polynucleotide comprising a sequence having at least 95% identity to a sequence selected from the group consisting of CS01-HC-NA, CS01-LC-NA, CS04-HC-NA, CS04-LC-NA, CS23-HC-NA, CS23-LC-NA.

52. A polynucleotide comprising a sequence having at least 99% identity to a sequence selected from the group consisting of CS01-HC-NA, CS01-LC-NA, CS04-HC-NA, CS04-LC-NA, CS23-HC-NA, CS23-LC-NA.

53. A polynucleotide comprising a sequence selected from the group consisting of CS01-HC-NA, CS01-LC-NA, CS04-HC-NA, CS04-LC-NA, CS23-HC-NA, CS23-LC-NA.

54. The polynucleotide according to any one of claims 51 to 53, wherein the polynucleotide encodes a Factor VIII polypeptide comprising an amino acid sequence having at least 95% identity to CS01-FL-AA (SEQ ID NO:2).

55. The polynucleotide according to any one of claims 51 to 53, wherein the polynucleotide encodes a Factor VIII polypeptide comprising the amino acid sequence of CS01-FL-AA (SEQ ID NO:2).

56. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19).

57. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

58. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) a deletion of amino acids AIEPRSF755-761, relative to FVIII-FL-AA (SEQ ID NO: 19); and

b) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

59. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19); and

b) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

60. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises I105V, A127S, G151K, M166T, L171P, and F328S amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

61. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19);

b) a deletion of amino acids AIEPRSF755-761, relative to FVIII-FL-AA (SEQ ID NO: 19); and

c) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

62. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19);

b) a deletion of amino acids AIEPRSF755-761, relative to FVIII-FL-AA (SEQ ID NO: 19); and

c) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

63. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19);

b) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19); and

c) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19).

64. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19);

b) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19);

c) a deletion of amino acids AIEPRSF755-761, relative to FVIII-FL-AA (SEQ ID NO: 19); and

d) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

65. The polynucleotide according to any one of claims 1 to 55, wherein the encoded Factor VIII polypeptide comprises:

a) I105V, A127S, G151K, M166T, and L171P amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19);

b) an F328S amino acid substitution, relative to FVIII-FL-AA (SEQ ID NO: 19);

c) C1918G and C1922G amino acid substitutions, relative to FVIII-FL-AA (SEQ ID NO: 19);

d) a deletion of amino acids AIEPRSF755-761, relative to FVIII-FL-AA (SEQ ID NO: 19); and

e) an insertion of amino acids TTYVNRSL (SEQ ID NO: 33) after N754, relative to FVIII-FL-AA (SEQ ID NO: 19).

66. The polynucleotide according to any one of claims 1 to 65, further comprising a promoter element operably linked to the polynucleotide encoding the Factor VIII polypeptide.

67. The polynucleotide of claim 66, wherein the promoter element is a liver-specific promoter sequence upstream of the nucleotide sequence encoding the Factor VIII polypeptide.

68. The polynucleotide of claim 67, further comprising an intron sequence positioned between the liver-specific promoter sequence and the nucleotide sequence encoding the Factor VIII polypeptide.

69. An adeno-associated virus (AAV) vector comprising a polynucleotide according to any one of claims 1 to 68.

70. An adeno-associated virus (AAV) particle comprising a polynucleotide according to any one of claims 1 to 68.

71. A host cell infected with an adeno-associated virus (AAV) particle comprising a polynucleotide according to any one of claims 1 to 68.

72. A method for producing an adeno-associated virus (AAV) particle comprising introducing a polynucleotide according to any one of claims 1 to 68 into a mammalian host cell, wherein the polynucleotide is competent for replication in the mammalian host cell.

73. A method for treating hemophilia A comprising administering, to a patient in need thereof, an adeno-associated virus (AAV) particle according to claim 70.

74. A method for transducing a host cell comprising contacting the host cell with an adeno-associated virus (AAV) particle according to claim 70.

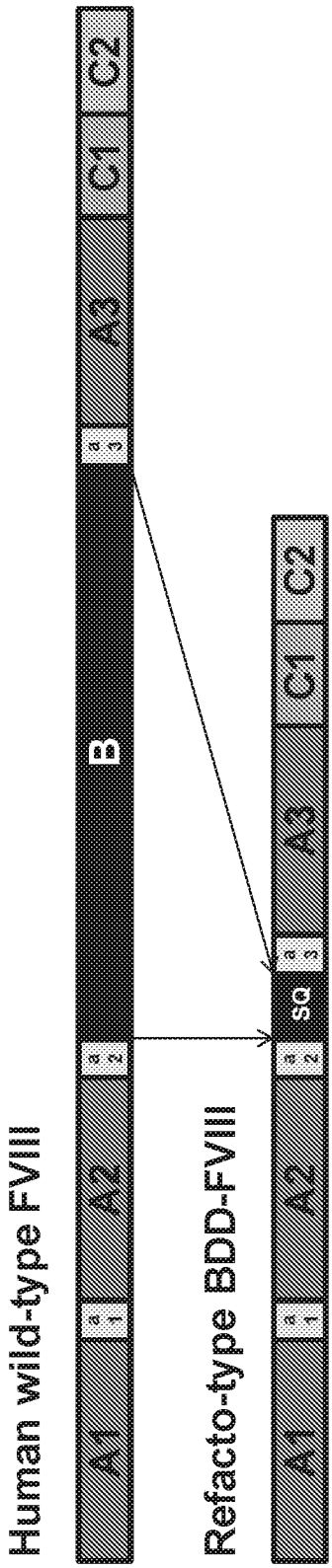
Human wild-type FVIII
A1
a1
A2
a2
B
a3
A3
C1
C2
Refacto-type BDD-FVIII
A1
a1
A2
a2
sq
a3
A3
C1
C2

Figure 1

CS04-FL-NA

atgcagattgagctgagcacctgcttcttcctgtgcctgctgaggttctgcttctctgccaccagga
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(Continued)

Figure 2A

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agtacaagatggccctgtacaacctctaccctggggtctttgagactgtggagatgctgccctccaa
agctggcatctggagggtggagtgcctcattggggagcacctgcatgctggcatgagcaccctgttc
ctggtctacagcaacaagtgccagacccccctgggaatggcctctggccacatcagggacttccaga
tcactgcctctggccagtatggccagtgggcccccaagctggccaggctccactactctggatccat
caatgcctggagcaccaaggagccattcagctggatcaaagtggacctgctggcccccatgatcatc
catggcatcaagacccaggggccaggcagaagttctccagcctgtacatcagccagttcatcatca
tgtacagcctggatggcaagaaatggcagacctacagaggcaactccactggaacactcatggtctt
ctttggcaatgtggacagctctggcatcaagcacaacatcttcaacccccccaatcatcgccagatac
atcaggctgcaccccacccactacagcatccgcagcaccctcaggatggagctgatgggctgtgacc
tgaactcctgcagcatgcccctgggcatggagagcaaggccatttctgatgcccagatcactgcctc
cagctacttcaccaacatgtttgccacctggagcccaagcaaggccaggctgcacctccagggaagg
agcaatgcctggaggccccaggtcaacaacccaaaggagtggctgcaggtggacttccagaagacca
tgaaggtcactggggtgaccacccaggggtcaagagcctgctcaccagcatgtatgtgaaggagtt
cctgatcagctccagccaggatggccaccagtggaccctcttcttccagaatggcaaggtcaaggtg
ttccagggcaaccaggacagcttcacCcctgtggtgaacagcctggacccccccctcctgaccagat
acctgaggattcacccccagagctgggtccaccagattgccctgaggatggaggtcctgggatgtga
ggcccaggacctgtactga (SEQ ID NO:1)

Figure 2B

CS04-FL-AA

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDAR'FPPRVPKSFPFNTSVVYK
KTLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVITLKNMASHPVSLHAVGVSYWKASEGAEY
DDQTSQREKEDDKVFPGGSHTYVWQVLKENGPMASDPLCLTYSYLSHVDLVKDLNSGLIGALLVCRE
GSLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRDAASARAWPKMHTVNGYVNRSLPGLIGC
HRKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNHRQASLEISPITFLTAQTLLMDLGQFLLFCHISSH
QHDGMEAYVKVDSCPEEPQLRMKNNEEAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKKHPKTW
VHYIAAEEEDWDYAPLVLAPDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQHESGIL
GPLLYGEVGDTLLIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIFKYKWTVT
VEDGPTKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVILFSVFDENR
SWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILSIGAQTDFLS
VFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRGMTALLKVSSCDKNTGD
YYEDSYEDISAYLLSKNNAIEPRSFSQNPPVLKRHQREITRTTLQSDQEEIDYDDTISVEMKKEDFD
IYDEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSSPHVLRNRAQSGSVPQFKKVVFQEFTDGSFTQ
PLYRGELNEHLGLLGPYIRAEVEDNIMVTFRNQASRPYSFYSSLISYEEDQRQGAEPRKNFVKPNET
KTYFWKVQHHMAPTKDEFDCKAWAYFSDVDLEKDVHSGLIGPLLVCHTNTLNPAHGRQVTVQEFALF
FTIFDETKSWYFTENMERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPGLVMAQDQRIRWYLLS
MGSNENIHSIHFSGHVFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWRVECLIGEHLHAGMSTL
FLVYSNKCQTPLGMASGHIRDFQITASGQYGQWAPKLARLHYSGSINAWSTKEPFSWIKVDLLAPMI
IHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGNVDSSGIKHNIFNPPIIAR
YIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASSYFTNMFATWSPSKARLHLQG
RSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTSMYVKEFLISSSQDGHQWTLFFQNGKVK
VFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWVHQIALRMEVLGCEAQDLY (SEQ ID NO:2)

Figure 3

CS04-HC-NA

gcc
accaggagat actacctggg ggctgtggag ctttcttggg actacatgca gtctgacctg
ggggagctgc ctgtggatgc caggttccca cccagagtgc ccaaatcctt cccattcaac
acctctgtgg tctacaagaa gaccctcttt gtggagttca ctgaccacct gttcaacatt
gccaaaccca ggccaccctg gatgggactc ctgggaccca ccattcaggc tgaggtgtat
gacactgtgg tcatcaccct caagaacatg gcctcccacc ctgtgagcct gcatgctgtg
ggggtcagct actggaaggc ctctgagggg gctgagtatg atgaccagac ctcccagagg
gagaaggagg atgacaaagt gttccctggg ggcagccaca cctatgtgtg gcaggtcctc
aaggagaatg gccccatggc ctctgaccca ctctgcctga cctactccta cctttctcat
gtggacctgg tcaaggacct caactctgga ctgattgggg ccctgctggt gtgcagggag
ggctccctgg ccaaagagaa gacccagacc ctgcacaagt tcattctcct gtttgctgtc
tttgatgagg gcaagagctg gcactctgaa accaagaact ccctgatgca ggacagggat
gctgcctctg ccagggcctg gcccaagatg cacactgtga atggctatgt gaacaggagc
ctgcctggac tcattggctg ccacaggaaa tctgtctact ggcatgtgat tggcatgggg
acaacccctg aggtgcactc cattttcctg gagggccaca ccttcctggt caggaaccac
agacaggcca gcctggagat cagccccatc accttcctca ctgcccagac cctgctgatg
gacctcggac agttcctgct gttctgccac atcagctccc accagcatga tggcatggag
gcctatgtca aggtggacag ctgccctgag gagccacagc tcaggatgaa gaacaatgag
gaggctgagg actatgatga tgacctgact gactctgaga tggatgtggt ccgctttgat
gatgacaaca gcccatcctt cattcagatc aggtctgtgg ccaagaaaca ccccaagacc
tgggtgcact acattgctgc tgaggaggag gactgggact atgccccact ggtcctggcc
cctgatgaca ggagctacaa gagccagtac ctcaacaatg gcccacagag gattggacgc
aagtacaaga aagtcaggtt catggcctac actgatgaaa ccttcaagac cagggaggcc
attcagcatg agtctggcat cctgggccca ctcctgtatg gggaggtggg ggacaccctg
ctcatcatct tcaagaacca ggcctccagg ccctacaaca tctacccaca tggcatcact
gatgtcaggc ccctgtacag ccgcaggctg ccaaaggggg tgaaacacct caaggacttc
cccattctgc ctggggagat cttcaagtac aagtggactg tcactgtgga ggatggacca
accaaatctg accccaggtg cctcaccaga tactactcca gctttgtgaa catggagagg
gacctggcct ctggcctgat tggcccactg ctcatctgct acaaggagtc tgtggaccag
aggggaaacc agatcatgtc tgacaagagg aatgtgattc tgttctctgt ctttgatgag
aacaggagct ggtacctgac tgagaacatt cagcgcttcc tgcccaaccc tgctggggtg
cagctggagg accctgagtt ccaggccagc aacatcatgc actccatcaa tggctatgtg
tttgacagcc tccagctttc tgtctgcctg catgaggtgg cctactggta cattctttct
attggggccc agactgactt cctttctgtc ttcttctctg gctacacctt caaacacaag
atggtgtatg aggacaccct gaccctcttc ccattctctg gggagactgt gttcatgagc
atggagaacc ctggcctgtg gattctggga tgccacaact ctgacttccg caacaggggc
atgactgccc tgctcaaagt ctcctcctgt gacaagaaca ctggggacta ctatgaggac
agctatgagg acatctctgc ctacctgctc agcaagaaca atgccattga gcccagg
(SEQ ID NO:3)

Figure 4

CS04-LC-NA

g agatcaccag gaccaccctc
cagtctgacc aggaggagat tgactatgat gacaccattt ctgtggagat gaagaaagag
gactttgaca tctatgacga ggacgagaac cagagcccaa ggagcttcca gaagaagacc
aggcactact tcattgctgc tgtggagcgc ctgtgggact atggcatgag ctccagcccc
catgtcctca ggaacagggc ccagtctggc tctgtgccac agttcaagaa agtggtcttc
caagagttca ctgatggcag cttcacccag cccctgtaca gaggggagct gaatgagcac
ctgggactcc tgggcccata catcagggct gaggtggagg acaacatcat ggtgaccttc
cgcaaccagg cctccaggcc ctacagcttc tacagctccc tcatcagcta tgaggaggac
cagaggcagg gggctgagcc acgcaagaac tttgtgaaac ccaatgaaac caagacctac
ttctggaaag tccagcacca catggccccc accaaggatg agtttgactg caaggcctgg
gcctacttct ctgatgtgga cctggagaag gatgtgcact ctggcctgat tggcccactc
ctggtctgcc acaccaacac cctgaaccct gcccatggaa ggcaagtgac tgtgcaggag
tttgccctct tcttcaccat ctttgatgaa accaagagct ggtacttcac tgagaacatg
gagcgcaact gcagggcccc atgcaacatt cagatggagg accccacctt caaagagaac
taccgcttcc atgccatcaa tggctacatc atggacaccc tgcctgggct tgtcatggcc
caggaccaga ggatcaggtg gtacctgctt tctatgggct ccaatgagaa cattcactcc
atccacttct ctgggcatgt cttcactgtg cgcaagaagg aggagtacaa gatggccctg
tacaacctct accctggggt ctttgagact gtggagatgc tgccctccaa agctggcatc
tggagggtgg agtgcctcat tggggagcac ctgcatgctg gcatgagcac cctgttcctg
gtctacagca acaagtgcca gacccccctg ggaatggcct ctggccacat cagggacttc
cagatcactg cctctggcca gtatggccag tgggccccca agctggccag gctccactac
tctggatcca tcaatgcctg gagcaccaag gagccattca gctggatcaa agtggacctg
ctggccccca tgatcatcca tggcatcaag acccaggggg ccaggcagaa gttctccagc
ctgtacatca gccagttcat catcatgtac agcctggatg gcaagaaatg gcagacctac
agaggcaact ccactggaac actcatggtc ttctttggca atgtggacag ctctggcatc
aagcacaaca tcttcaaccc cccaatcatc gccagataca tcaggctgca ccccacccac
tacagcatcc gcagcaccct caggatggag ctgatgggct gtgacctgaa ctcctgcagc
atgcccctgg gcatggagag caaggccatt tctgatgccc agatcactgc ctccagctac
ttcaccaaca tgtttgccac ctggagccca agcaaggcca ggctgcacct ccagggaagg
agcaatgcct ggaggcccca ggtcaacaac ccaaaggagt ggctgcaggt ggacttccag
aagaccatga aggtcactgg ggtgaccacc caggggtca agagcctgct caccagcatg
tatgtgaagg agttcctgat cagctccagc caggatggcc accagtggac cctcttcttc
cagaatggca aggtcaaggt gttccagggc aaccaggaca gcttcacccc tgtggtgaac
agcctggacc ccccctcct gaccagatac ctgaggattc acccccagag ctgggtccac
cagattgccc tgaggatgga ggtcctggga tgtgaggccc aggacctgta c
(SEQ ID NO:4)

Figure 5

BDLO01 - agc ttctctcaga atccacctgt cctgaagaga caccagaga (SEQ ID NO:5)

BDLO04 - agc ttcagccaga atccacctgt cctgaaacgc caccagagg (SEQ ID NO:6)

BDLO23 - agc ttcagccaga acccccccgt gctgaagagg caccagagg (SEQ ID NO:7)

BDLNG1 - agcttcagccagaatGTGAGCAACAATGTGAGCAACAATGCCACCAATAATGCTACCAACccacctgtcctgaaacgccaccagagg (SEQ ID NO:36)

BDLNG4 - agcttcagccagaatGTGAGCAACAATGCCACCAACAATGTGAGCAACccacctgtcctgaaacgccaccagagg (SEQ ID NO:37)

BDLNG5 - agcttcagccagaatGTGAGCAATAATGCCACCAACccacctgtcctgaaacgccaccagagg (SEQ ID NO:38)

BDLNG6 - agcttcagccagaatGTGAGCAATAATccacctgtcctgaaacgccaccagagg (SEQ ID NO:39)

BDLNG9 - agcttcagccagaatAGGAGCCTGccacctgtcctgaaacgccaccagagg (SEQ ID NO:40)

BDLNG10 - agcttcagccagaatGCCACTAATGTGTCTAACAACTCTGCTACCTCTGCTGACTCTGCTGTGAGCccacctgtcctgaaacgccaccagagg (SEQ ID NO:41)

BDLNG16 - agcttcagccagaatGCCACCAACTATGTGAACAGGAGCCTGccacctgtcctgaaacgccaccagagg (SEQ ID NO:42)

BDLNG17 - agcttcagccagaatGCCACCAACTATGTGAACAGGAGCCTGTCTGCCACCTCTGCTGACTCTGCTGTGAGCCAGAATccacctgtcctgaaacgccaccagagg(SEQ ID NO:43)

BDLNG18 - agcttcagccagaatGTGAGCAACAATGTGAGCAATGCTGTGTCTGCTGTGTCTGCTccacctgtcctgaaacgccaccagagg (SEQ ID NO:44)

BDLNG19 - agcttcagccagaatATCACTGTGGCCTCTGCCACCTCTAACATCACTGTGGCCTCTGCTGACccacctgtcctgaaacgccaccagagg (SEQ ID NO:45)

BDLNG20 - agcttcagccagaatATCACTGTGACCAACATCACTGTGACTGCCccacctgtcctgaaacgccaccagagg (SEQ ID NO:46)

BDLNG21 - agcttcagccagaatCAGACTGTGACCAACATCACTGTGACTGCCccacctgtcctgaaacgccaccagagg (SEQ ID NO:47)

BDLNGV - agcttcagccagaatGCCACTAATGTGTCTAACAACAGCAACACCAGCAATGACAGCAATGTGTCTccacctgtcctgaaacgccaccagagg (SEQ ID NO:48)

Figure 6

CS04-AV-NA

```
1    tcgcgcgttt cggtgatgac ggtgaaaacc tctgacacat gcagctcccg gagacggtca
61   cagcttgtct gtaagcggat gccgggagca gacaagcccg tcagggcgcg tcagcgggtg
121  ttggcgggtg tcggggctgg cttaactatg cggcatcaga gcagattgta ctgagagtgc
181  accatatgcg gtgtgaaata ccgcacagat gcgtaaggag aaaataccgc atcaggcgcc
241  attcgccatt caggctgcgc aactgttggg aagggcgatc ggtgcgggcc tcttcgctat
301  tacgccagct ggcgaaaggg ggatgtgctg caaggcgatt aagttgggta acgccagggt
361  tttcccagtc acgacgttgt aaaacgacgg ccagtgaatt cctcgagatt taaatgacgt
421  tggccactcc ctctctgcgc gctcgctcgc tcactgaggc cgggcgacca aaggtcgccc
481  gacgcccggg ctttgcccgg gcggcctcag tgagcgagcg agcgcgcaga gagggagtgg
541  ccaactccat cactaggggt tcctgagttt aaacttcgtc gacgattcga gcttgggctg
601  caggtcgagg gcactgggag gatgttgagt aagatggaaa actactgatg acccttgcag
661  agacagagta ttaggacatg tttgaacagg ggccgggcga tcagcaggta gctctagagg
721  atccccgtct gtctgcacat ttcgtagagc gagtgttccg atactctaat ctccctaggc
781  aaggttcata tttgtgtagg ttacttattc tccttttgtt gactaagtca ataatcagaa
841  tcagcaggtt tggagtcagc ttggcaggga tcagcagcct gggttggaag gagggggtat
901  aaaagccccc tcaccaggag aagccgtcac acagactagg cgcgccaccg ccaccatgca
961  gattgagctg agcacctgct tcttcctgtg cctgctgagg ttctgcttct ctgccaccag
1021 gagatactac ctgggggctg tggagctttc ttgggactac atgcagtctg acctgggggа
1081 gctgcctgtg gatgccaggt tcccacccag agtgcccaaa tccttcccat tcaacacctc
1141 tgtggtctac aagaagaccc tctttgtgga gttcactgac cacctgttca acattgccaa
1201 acccaggcca ccctggatgg gactcctggg acccaccatt caggctgagg tgtatgacac
1261 tgtggtcatc accctcaaga acatggcctc ccaccctgtg agcctgcatg ctgtgggggt
1321 cagctactgg aaggcctctg agggggctga gtatgatgac cagacctccc agagggagaa
1381 ggaggatgac aaagtgttcc ctgggggcag ccacacctat gtgtggcagg tcctcaagga
1441 gaatggcccc atggcctctg acccactctg cctgacctac tcctaccttt ctcatgtgga
1501 cctggtcaag gacctcaact ctggactgat tggggccctg ctggtgtgca gggagggctc
1561 cctggccaaa gagaagaccc agaccctgca caagttcatt ctcctgtttg ctgtctttga
1621 tgagggcaag agctggcact ctgaaaccaa gaactccctg atgcaggaca gggatgctgc
1681 ctctgccagg gcctggccca agatgcacac tgtgaatggc tatgtgaaca ggagcctgcc
1741 tggactcatt ggctgccaca ggaaatctgt ctactggcat gtgattggca tggggacaac
1801 ccctgaggtg cactccattt tcctggaggg ccacaccttc ctggtcagga accacagaca
1861 ggccagcctg gagatcagcc ccatcacctt cctcactgcc cagaccctgc tgatggacct
1921 cggacagttc ctgctgttct gccacatcag ctcccaccag catgatggca tggaggccta
1981 tgtcaaggtg gacagctgcc ctgaggagcc acagctcagg atgaagaaca atgaggaggc
2041 tgaggactat gatgatgacc tgactgactc tgagatggat gtggtccgct ttgatgatga
2101 caacagccca tccttcattc agatcaggtc tgtggccaag aaacacccca agacctgggt
2161 gcactacatt gctgctgagg aggaggactg ggactatgcc ccactggtcc tggcccctga
2221 tgacaggagc tacaagagcc agtacctcaa caatggccca cagaggattg gacgcaagta
2281 caagaaagtc aggttcatgg cctacactga tgaaaccttc aagaccaggg aggccattca
2341 gcatgagtct ggcatcctgg gcccactcct gtatggggag gtgggggaca ccctgctcat
2401 catcttcaag aaccaggcct ccaggcccta caacatctac ccacatggca tcactgatgt
2461 caggcccctg tacagccgca ggctgccaaa gggggtgaaa cacctcaagg acttccccat
```

Figure 7A

```
2521 tctgcctggg gagatcttca agtacaagtg gactgtcact gtggaggatg gaccaaccaa
2581 atctgacccc aggtgcctca ccagatacta ctccagcttt gtgaacatgg agagggacct
2641 ggcctctggc ctgattggcc cactgctcat ctgctacaag gagtctgtgg accagagggg
2701 aaaccagatc atgtctgaca agaggaatgt gattctgttc tctgtctttg atgagaacag
2761 gagctggtac ctgactgaga acattcagcg cttcctgccc aaccctgctg gggtgcagct
2821 ggaggaccct gagttccagg ccagcaacat catgcactcc atcaatggct atgtgtttga
2881 cagcctccag ctttctgtct gcctgcatga ggtggcctac tggtacattc tttctattgg
2941 ggcccagact gacttccttt ctgtcttctt ctctggctac accttcaaac acaagatggt
3001 gtatgaggac accctgaccc tcttcccatt ctctggggag actgtgttca tgagcatgga
3061 gaaccctggc ctgtggattc tgggatgcca caactctgac ttccgcaaca ggggcatgac
3121 tgccctgctc aaagtctcct cctgtgacaa gaacactggg gactactatg aggacagcta
3181 tgaggacatc tctgcctacc tgctcagcaa gaacaatgcc attgagccca ggagcttcag
3241 ccagaatcca cctgtcctga aacgccacca gagggagatc accaggacca ccctccagtc
3301 tgaccaggag gagattgact atgatgacac catttctgtg gagatgaaga aagaggactt
3361 tgacatctat gacgaggacg agaaccagag cccaaggagc ttccagaaga agaccaggca
3421 ctacttcatt gctgctgtgg agcgcctgtg ggactatggc atgagctcca gcccccatgt
3481 cctcaggaac agggcccagt ctggctctgt gccacagttc aagaaagtgg tcttccaaga
3541 gttcactgat ggcagcttca cccagcccct gtacagaggg gagctgaatg agcacctggg
3601 actcctgggc ccatacatca gggctgaggt ggaggacaac atcatggtga ccttccgcaa
3661 ccaggcctcc aggccctaca gcttctacag ctccctcatc agctatgagg aggaccagag
3721 gcaggggggct gagccacgca agaactttgt gaaacccaat gaaaccaaga cctacttctg
3781 gaaagtccag caccacatgg cccccaccaa ggatgagttt gactgcaagg cctgggccta
3841 cttctctgat gtggacctgg agaaggatgt gcactctggc ctgattggcc cactcctggt
3901 ctgccacacc aacaccctga accctgccca tggaaggcaa gtgactgtgc aggagtttgc
3961 cctcttcttc accatctttg atgaaaccaa gagctggtac ttcactgaga acatggagcg
4021 caactgcagg gccccatgca acattcagat ggaggacccc accttcaaag agaactaccg
4081 cttccatgcc atcaatggct acatcatgga caccctgcct gggcttgtca tggcccagga
4141 ccagaggatc aggtggtacc tgctttctat gggctccaat gagaacattc actccatcca
4201 cttctctggg catgtcttca ctgtgcgcaa gaaggaggag tacaagatgg ccctgtacaa
4261 cctctaccct ggggtctttg agactgtgga gatgctgccc tccaaagctg gcatctggag
4321 ggtggagtgc ctcattgggg agcacctgca tgctggcatg agcaccctgt tcctggtcta
4381 cagcaacaag tgccagaccc ccctgggaat ggcctctggc cacatcaggg acttccagat
4441 cactgcctct ggccagtatg gccagtgggc ccccaagctg gccaggctcc actactctgg
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4561 ccccatgatc atccatggca tcaagaccca gggggccagg cagaagttct ccagcctgta
4621 catcagccag ttcatcatca tgtacagcct ggatggcaag aaatggcaga cctacagagg
4681 caactccact ggaacactca tggtcttctt tggcaatgtg gacagctctg gcatcaagca
4741 caacatcttc aacccccccaa tcatcgccag atacatcagg ctgcacccca cccactacag
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4861 cctgggcatg gagagcaagg ccatttctga tgcccagatc actgcctcca gctacttcac
4921 caacatgttt gccacctgga gcccaagcaa ggccaggctg cacctccagg gaaggagcaa
4981 tgcctggagg ccccaggtca acaacccaaa ggagtggctg caggtggact tccagaagac
```

Figure 7B

5041 catgaaggtc actggggtga ccacccaggg ggtcaagagc ctgctcacca gcatgtatgt
5101 gaaggagttc ctgatcagct ccagccagga tggccaccag tggaccctct tcttccagaa
5161 tggcaaggtc aaggtgttcc agggcaacca ggacagcttc acccctgtgg tgaacagcct
5221 ggaccccccc ctcctgacca gatacctgag gattcacccc cagagctggg tccaccagat
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5341 cgctcttagt agcagtatcg ataataaaag atctttattt tcattagatc tgtgtgttgg
5401 ttttttgtgt gttaattaag ctcgcgaagg aacccctagt gatggagttg gccactccct
5461 ctctgcgcgc tcgctcgctc actgaggccg ggcgaccaaa ggtcgcccga cgcccgggct
5521 ttgcccgggc ggcctcagtg agcgagcgag cgcgcagaga gggagtggcc aagacgattt
5581 aaatgacaag cttggcgtaa tcatggtcat agctgtttcc tgtgtgaaat tgttatccgc
5641 tcacaattcc acacaacata cgagccggaa gcataaagtg taaagcctgg ggtgcctaat
5701 gagtgagcta actcacatta attgcgttgc gctcactgcc cgctttccag tcgggaaacc
5761 tgtcgtgcca gctgcattaa tgaatcggcc aacgcgcggg gagaggcggt ttgcgtattg
5821 ggcgctcttc cgcttcctcg ctcactgact cgctgcgctc ggtcgttcgg ctgcggcgag
5881 cggtatcagc tcactcaaag gcggtaatac ggttatccac agaatcaggg gataacgcag
5941 gaaagaacat gtgagcaaaa ggccagcaaa aggccaggaa ccgtaaaaag gccgcgttgc
6001 tggcgttttt ccataggctc cgcccccctg acgagcatca caaaaatcga cgctcaagtc
6061 agaggtggcg aaacccgaca ggactataaa gataccaggc gtttcccccc ggaagctccc
6121 tcgtgcgctc tcctgttccg accctgccgc ttaccggata cctgtccgcc tttctccctt
6181 cgggaagcgt ggcgctttct catagctcac gctgtaggta tctcagttcg gtgtaggtcg
6241 ttcgctccaa gctgggctgt gtgcacgaac ccccgttca gcccgaccgc tgcgccttat
6301 ccggtaacta tcgtcttgag tccaacccgg taagacacga cttatcgcca ctggcagcag
6361 ccactggtaa caggattagc agagcgaggt atgtaggcgg tgctacagag ttcttgaagt
6421 ggtggcctaa ctacggctac actagaagaa cagtatttgg tatctgcgct ctgctgaagc
6481 cagttacctt cggaaaaaga gttggtagct cttgatccgg caaacaaacc accgctggta
6541 gcggtggttt ttttgtttgc aagcagcaga ttacgcgcag aaaaaaagga tctcaagaag
6601 atcctttgat cttttctacg gggtctgacg ctcagtggaa cgaaaactca cgttaaggga
6661 ttttggtcat gagattatca aaaaggatct tcacctagat ccttttaaat taaaaatgaa
6721 gttttaaatc aatctaaagt atatatgagt aaacttggtc tgacagttac caatgcttaa
6781 tcagtgaggc acctatctca gcgatctgtc tatttcgttc atccatagtt gcctgactcc
6841 ccgtcgtgta gataactacg atacgggagg gcttaccatc tggccccagt gctgcaatga
6901 taccgcgaga cccacgctca ccggctccag atttatcagc aataaaccag ccagccggaa
6961 gggccgagcg cagaagtggt cctgcaactt tatccgcctc catccagtct attaattgtt
7021 gccgggaagc tagagtaagt agttcgccag ttaatagttt gcgcaacgtt gttgccattg
7081 ctacaggcat cgtggtgtca cgctcgtcgt ttggtatggc ttcattcagc tccggttccc
7141 aacgatcaag gcgagttaca tgatccccca tgttgtgcaa aaaagcggtt agctccttcg
7201 gtcctccgat cgttgtcaga agtaagttgg ccgcagtgtt atcactcatg gttatggcag
7261 cactgcataa ttctcttact gtcatgccat ccgtaagatg cttttctgtg actggtgagt
7321 actcaaccaa gtcattctga gaatagtgta tgcggcgacc gagttgctct tgcccggcgt
7381 caatacggga taataccgcg ccacatagca gaactttaaa agtgctcatc attggaaaac
7441 gttcttcggg gcgaaaactc tcaaggatct taccgctgtt gagatccagt tcgatgtaac
7501 ccactcgtgc acccaactga tcttcagcat cttttacttt caccagcgtt tctgggtgag
7561 caaaaacagg aaggcaaaat gccgcaaaaa agggaataag ggcgacacgg aaatgttgaa
7621 tactcatact cttccttttt caatattatt gaagcattta tcagggttat tgtctcatga
7681 gcggatacat atttgaatgt atttagaaaa ataaacaaat aggggttccg cgcacatttc
7741 cccgaaaagt gccacctgac gtctaagaaa ccattattat catgacatta acctataaaa
7801 ataggcgtat cacgaggccc tttcgtc (SEQ ID NO:8)

Figure 7C

CS01m1-FL-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGGA
GATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCTGT
GGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAGAAG
ACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATGGGAC
TCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATGGCATC
CCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAAGCCTCTGAAGGGGCTGAGTATGAT
GACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGGGATCTCACACCTATGTGT
GGCAAGTCCTCAAGGAGAATGGACCCATGGCATCTGACCCACTCTGCCTGACATACTCCTACCTTTC
TCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGCAGGGAAGGA
TCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTCTTTGATGAGG
GCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCCTCTGCCAGGGC
ATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTCATTGGCTGCCAC
AGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCACTCCATTTTCCTGG
AGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCTCCCATCACCTTCCT
CACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTCCTGCCACATCTCTTCCCACCAG
CATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCACAGCTCAGGATGAAGA
ACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATGGATGTGGTCAGATTTGA
TGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAACACCCCAAGACATGGGTG
CACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTCCTGGCCCCTGATGACAGGA
GCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGAAAGTACAAGAAAGTCAGATT
CATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAGCATGAGTCTGGCATTCTGGGA
CCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTCAAGAACCAGGCCTCCAGGCCCT
ACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTACAGCAGGAGACTGCCAAAAGGGGT
GAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTCAAGTACAAGTGGACTGTCACTGTG
GAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGATACTACTCCTCTTTTGTGAACATGG
AGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATCTGCTACAAGGAGTCTGTGGACCAGAG
AGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTGTTCTCTGTCTTTGATGAGAACAGATCA
TGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAACCCTGCTGGGGTGCAACTGGAAGACCCTG
AGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGCTATGTGTTTGACTCTCTCCAGCTTTCTGT
CTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCTATTGGGGCACAAACTGACTTCCTTTCTGTC
TTCTTCTCTGGATACACCTTCAAGCACAAGATGGTGTATGAGGACACCCTGACACTCTTCCCATTCT
CTGGGGAAACTGTGTTCATGAGCATGGAGAACCCTGGACTGTGGATTCTGGGATGCCACAACTCTGA
CTTCAGAAACAGGGGAATGACTGCACTGCTCAAAGTCTCCTCCTGTGACAAGAACACTGGGGACTAC
TATGAGGACTCTTATGAGGACATCTCTGCCTACCTGCTCAGCAAGAACAATGCCATTGAGCCCAGAA
GCTTCTCTCAGAATCCACCTGTCCTGAAGAGACACCAGAGAGAGATCACCAGGACAACCCTCCAGTC
TGACCAGGAAGAGATTGACTATGATGACACCATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATC
TATGATGAGGACGAGAACCAGTCTCCAAGATCATTCCAGAAGAAGACAAGACACTACTTCATTGCTG
CTGTGGAAAGACTGTGGGACTATGGCATGTCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTC
TGGCTCTGTGCCACAGTTCAAGAAAGTGGTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCC
CTGTACAGAGGGGAACTGAATGAGCACCTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAG
ACAACATCATGGTGACATTCAGAAACCAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAG
CTATGAGGAAGACCAGAGACAAGGGGCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAG
ACCTACTTCTGGAAAGTCCAGCACCACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGG

(Continued)

Figure 8A

CATACTTCTCTGATGTGGACCTGGAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTG
CCACACCAACACCCTGAACCCTGCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTC
ACCATCTTTGATGAAACCAAGTCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCAT
GCAACATTCAGATGGAAGACCCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACAT
CATGGACACCCTGCCTGGGCTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATG
GGATCCAATGAGAACATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGG
AATACAAGATGGCCCTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAA
AGCTGGCATCTGGAGGGTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTC
CTGGTCTACAGCAACAAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGA
TCACTGCCTCTGGCCAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCAT
CAATGCATGGTCAACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATT
CATGGCATCAAGACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCA
TGTACTCTCTGGATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTT
CTTTGGCAATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATAC
ATCAGGCTGCACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACC
TGAACTCCTGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATC
CTCTTACTTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGA
AGCAATGCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAA
TGAAAGTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTT
CCTGATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTG
TTCCAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGAT
ACCTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGA
GGCACAAGACCTGTACTGA (SEQ ID NO:49)

Figure 8B

CS04Δ(760-1667) - CS04-SC1-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGGAGATAC
TACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCTGTGGATGCCAGG
TTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAGAAGACCCTCTTTGTGGAG
TTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATGGGACTCCTGGGACCCACCATTCAG
GCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATGGCCTCCCACCCTGTGAGCCTGCATGCTGTG
GGGGTCAGCTACTGGAAGGCCTCTGAGGGGGCTGAGTATGATGACCAGACCTCCCAGAGGGAGAAGGAGGAT
GACAAAGTGTTCCCTGGGGGCAGCCACACCTATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCATGGCCTCT
GACCCACTCTGCCTGACCTACTCCTACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATT
GGGGCCCTGCTGGTGTGCAGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTC
CTGTTTGCTGTCTTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGAT
GCTGCCTCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCACTCCATT
TTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGCCCCATCACCTTC
CTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTTCTGCCACATCAGCTCCCACCAGCAT
GATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCACAGCTCAGGATGAAGAACAATGAG
GAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATGGATGTGGTCCGCTTTGATGATGACAACAGC
CCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAACACCCCAAGACCTGGGTGCACTACATTGCTGCTGAG
GAGGAGGACTGGGACTATGCCCCACTGGTCCTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAAC
AATGGCCCACAGAGGATTGGACGCAAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAG
ACCAGGGAGGCCATTCAGCATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTG
CTCATCATCTTCAAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCC
CTGTACAGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGATACTAC
TCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATCTGCTACAAGGAG
TCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTGTTCTCTGTCTTTGATGAG
AACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAACCCTGCTGGGGTGCAGCTGGAGGAC
CCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGCTATGTGTTTGACAGCCTCCAGCTTTCTGTC
TGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCTATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTC
TCTGGCTACACCTTCAAACACAAGATGGTGTATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACT
GTGTTCATGAGCATGGAGAACCCTGGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGC
ATGACTGCCCTGCTCAAAGTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGAC
ATCTCTGCCTACCTGCTCAGCAAGAACAATGCCATTGAGCCCAGGGAGATCACCAGGACCACCCTCCAGTCT
GACCAGGAGGAGATTGACTATGATGACACCATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGAC
GAGGACGAGAACCAGAGCCCAAGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGC
CTGTGGGACTATGGCATGAGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAG
TTCAAGAAAGTGGTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAAT
GAGCACCTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGCAAC
CAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAGGGGGCTGAG
CCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCACCACATGGCCCCC

(Continued)

Figure 9A

ACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTGGAGAAGGATGTGCACTCT
GGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCTGCCCATGGAAGGCAAGTGACTGTG
CAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAGAGCTGGTACTTCACTGAGAACATGGAGCGC
AACTGCAGGGCCCCATGCAACATTCAGATGGAGGACCCCACCTTCAAAGAGAACTACCGCTTCCATGCCATC
AATGGCTACATCATGGACACCCTGCCTGGGCTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTT
TCTATGGGCTCCAATGAGAACATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAG
GAGTACAAGATGGCCCTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCT
GGCATCTGGAGGGTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTAC
AGCAACAAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGGAGCACCAAG
GAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATCAAGACCCAGGGGGCC
AGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTGGATGGCAAGAAATGGCAG
ACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGCAATGTGGACAGCTCTGGCATCAAGCAC
AACATCTTCAACCCCCCAATCATCGCCAGATACATCAGGCTGCACCCCACCCACTACAGCATCCGCAGCACC
CTCAGGATGGAGCTGATGGGCTGTGACCTGAACTCCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATT
TCTGATGCCCAGATCACTGCCTCCAGCTACTTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGG
CTGCACCTCCAGGGAAGGAGCAATGCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGAC
TTCCAGAAGACCATGAAGGTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTG
AAGGAGTTCCTGATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAG
GTGTTCCAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGATAC
CTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGTGAGGCCCAG
GACCTGTACTGA (SEQ ID NO:9)

Figure 9B

CS04Δ(760-1667) - CS04-SC1-AA

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFNTSVVYKK
TLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVITLKNMASHPVSLHAVGVSYWKASEGAEYD
DQTSQREKEDDKVFPGGSHTYVWQVLKENGPMASDPLCLTYSYLSHVDLVKDLNSGLIGALLVCREG
SLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRDAASARAWPKMHTVNGYVNRSLPGLIGCH
RKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNHRQASLEISPITFLTAQTLLMDLGQFLLFCHISSHQ
HDGMEAYVKVDSCPEEPQLRMKNNEEAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKKHPKTWV
HYIAAEEEDWDYAPLVLAPDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQHESGILG
PLLYGEVGDTLLIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIFKYKWTVTV
EDGPTKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVILFSVFDENRS
WYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILSIGAQTDFLSV
FFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRGMTALLKVSSCDKNTGDY
YEDSYEDISAYLLSKNNAIEPREITRTTLQSDQEEIDYDDTISVEMKKEDFDIYDEDENQSPRSFQK
KTRHYFIAAVERLWDYGMSSSPHVLRNRAQSGSVPQFKKVVFQEFTDGSFTQPLYRGELNEHLGLLG
PYIRAEVEDNIMVTFRNQASRPYSFYSSLISYEEDQRQGAEPRKNFVKPNETKTYFWKVQHHMAPTK
DEFDCKAWAYFSDVDLEKDVHSGLIGPLLVCHTNTLNPAHGRQVTVQEFALFFTIFDETKSWYFTEN
MERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPGLVMAQDQRIRWYLLSMGSNENIHSIHFSGH
VFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWRVECLIGEHLHAGMSTLFLVYSNKCQTPLGMA
SGHIRDFQITASGQYGQWAPKLARLHYSGSINAWSTKEPFSWIKVDLLAPMIIHGIKTQGARQKFSS
LYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGNVDSSGIKHNIFNPPIIARYIRLHPTHYSIRSTL
RMELMGCDLNSCSMPLGMESKAISDAQITASSYFTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEW
LQVDFQKTMKVTGVTTQGVKSLLTSMYVKEFLISSSQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNS
LDPPLLTRYLRIHPQSWVHQIALRMEVLGCEAQDLY (SEQ ID NO:10)

Figure 10

CS04Δ(772-1667) - CS04-SC2-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGGCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGGGCAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCATGGCCTCTGACCCACTCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTTCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGGAGCTTCAGCCAGAATTCCAGACACCCCAGCACC
AGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGACACCATTTCT
GTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCAAGGAGCTTC

(Continued)

Figure 11A

CAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGCATGAGCTCC
AGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTGGTCTTC
CAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAGCACCTGGGA
CTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGCAACCAGGCC
TCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAGGGGGCTGAG
CCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCACCACATG
GCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTGGAGAAG
GATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCTGCCCAT
GGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAGAGCTGG
TACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAGGACCCCACC
TTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGGCTTGTC
ATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAACATTCACTCC
ATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCCCTGTACAAC
CTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGGGTGGAG
TGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGCAACAAGTGC
CAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGCCAGTAT
GGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGGAGCACCAAG
GAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATCAAGACCCAG
GGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTGGATGGC
AAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGCAATGTGGAC
AGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGGCTGCACCCC
ACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAACTCCTGCAGC
ATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGCTACTTCACC
AACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGCAATGCCTGG
AGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAGGTCACT
GGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTCCTGATCAGC
TCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTGTTCCAGGGC
AACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCCTCCTGACCAGATACCTGAGG
ATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGTGAGGCCCAG
GACCTGTACTGA (SEQ ID NO:11)

Figure 11B

CS04Δ(772-1667) - CS04-SC2-AA

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFNTSVVYKKTLFVEF
TDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVITLKNMASHPVSLHAVGVSYWKASEGAEYDDQTSQREKEDDK
VFPGGSHTYVWQVLKENGPMASDPLCLTYSYLSHVDLVKDLNSGLIGALLVCREGSLAKEKTQTLHKFILLFA
VFDEGKSWHSETKNSLMQDRDAASARAWPKMHTVNGYVNRSLPGLIGCHRKSVYWHVIGMGTTPEVHSIFLEG
HTFLVRNHRQASLEISPITFLTAQTLLMDLGQFLLFCHISSHQHDGMEAYVKVDSCPEEPQLRMKNNEEAEDY
DDDLTDSEMDVVRFDDDNSPSFIQIRSVAKKHPKTWVHYIAAEEEDWDYAPLVLAPDDRSYKSQYLNNGPQRI
GRKYKKVRFMAYTDETFKTREAIQHESGILGPLLYGEVGDTLLIIFKNQASRPYNIYPHGITDVRPLYSRRLP
KGVKHLKDFPILPGEIFKYKWTVTVEDGPTKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQ
IMSDKRNVILFSVFDENRSWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWY
ILSIGAQTDFLSVFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRGMTALLKVSSC
DKNTGDYYEDSYEDISAYLLSKNNAIEPRSFSQNSRHPSTREITRTTLQSDQEEIDYDDTISVEMKKEDFDIY
DEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSSPHVLRNRAQSGSVPQFKKVVFQEFTDGSFTQPLYRGELN
EHLGLLGPYIRAEVEDNIMVTFRNQASRPYSFYSSLISYEEDQRQGAEPRKNFVKPNETKTYFWKVQHHMAPT
KDEFDCKAWAYFSDVDLEKDVHSGLIGPLLVCHTNTLNPAHGRQVTVQEFALFFTIFDETKSWYFTENMERNC
RAPCNIQMEDPTFKENYRFHAINGYIMDTLPGLVMAQDQRIRWYLLSMGSNENIHSIHFSGHVFTVRKKEEYK
MALYNLYPGVFETVEMLPSKAGIWRVECLIGEHLHAGMSTLFLVYSNKCQTPLGMASGHIRDFQITASGQYGQ
WAPKLARLHYSGSINAWSTKEPFSWIKVDLLAPMIIHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGN
STGTLMVFFGNVDSSGIKHNIFNPPIIARYIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQIT
ASSYFTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTSMYVKEFLISS
SQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWVHQIALRMEVLGCEAQDLY
(SEQ ID NO:12)

Figure 12

NG1: V S N N V S N N A T N N A T N (SEQ ID NO:51)
GTG AGC AAC AAT GTG AGC AAC AAT GCC ACC AAT AAT GCT ACC AAC (SEQ ID NO:50)

NG4: V S N N A T N N V S N (SEQ ID NO:53)
GTG AGC AAC AAT GCC ACC AAC AAT GTG AGC AAC (SEQ ID NO:52)

NG5: V S N N A T N (SEQ ID NO:55)
GTG AGC AAT AAT GCC ACC AAC (SEQ ID NO:54)

NG6: V S N N (SEQ ID NO:57)
GTG AGC AAT AAT (SEQ ID NO:56)

NG9: R S L (SEQ ID NO:59)
AGG AGC CTG (SEQ ID NO:58)

NG10: A T N V S N N S A T S A D S A V S (SEQ ID NO:61)
GCC ACT AAT GTG TCT AAC AAC TCT GCT ACC TCT GCT GAC TCT GCT GTG AGC (SEQ ID NO:60)

NG16: A T N Y V N R S L (SEQ ID NO:63)
GCC ACC AAC TAT GTG AAC AGG AGC CTG (SEQ ID NO:62)

Figure 13A

NG17: A T N Y V N R S L S A T S A D S A V S Q N (SEQ ID NO:65)
GCC ACC AAC TAT GTG AAC AGG AGC CTG TCT GCC ACC TCT GCT GAC TCT GCT GTG AGC CAG AAT (SEQ ID NO:64)

NG18: V S N N V S N A V S A V S A (SEQ ID NO:67)
GTG AGC AAC AAT GTG AGC AAT GCT GTG TCT GCT GTG TCT GCT (SEQ ID NO:66)

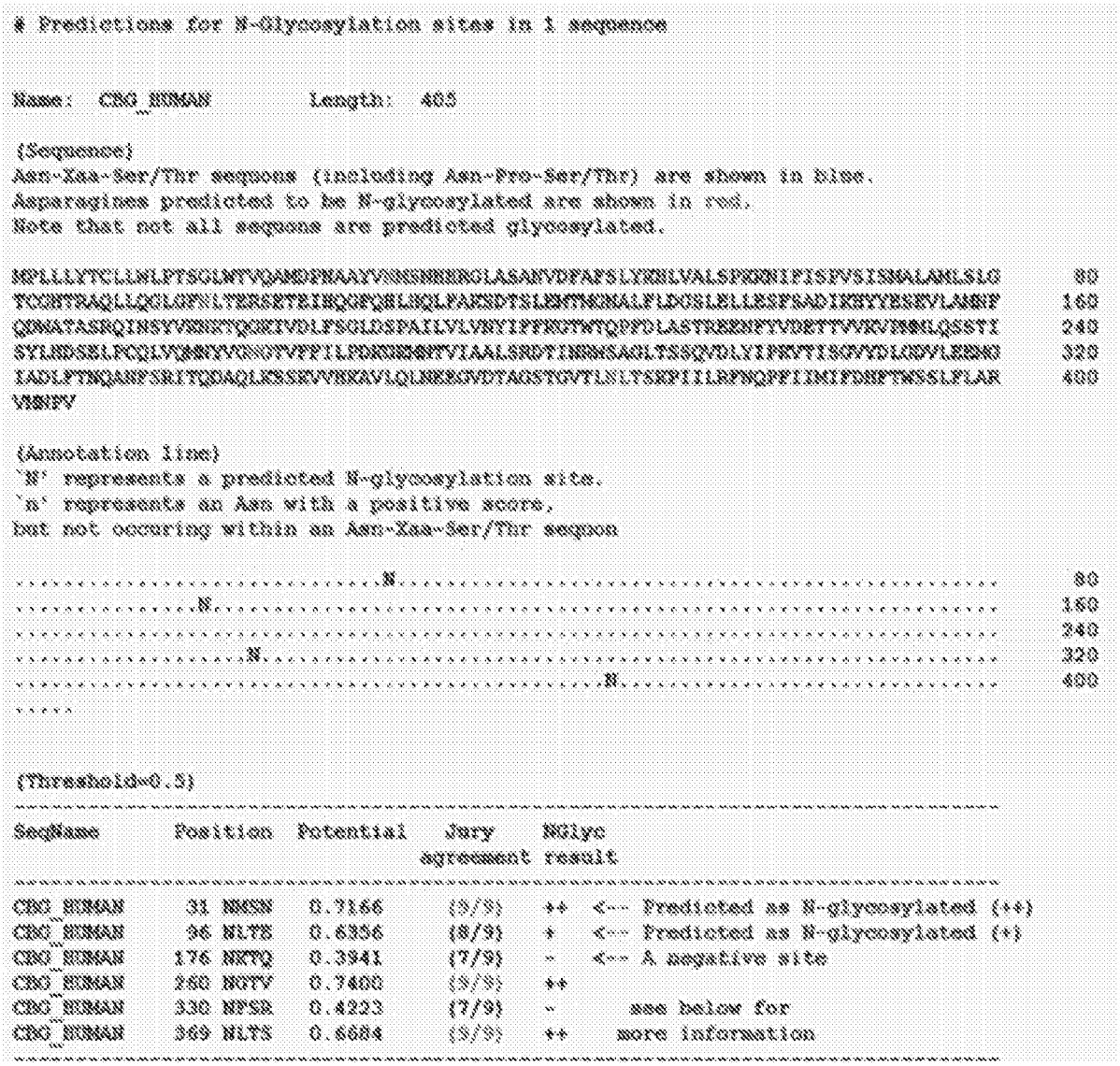
NG19: I T V A S A T S N I T V A S A D (SEQ ID NO:69)
ATC ACT GTG GCC TCT GCC ACC TCT AAC ATC ACT GTG GCC TCT GCT GAC (SEQ ID NO:68)

NG20: I T V T N I T V T A (SEQ ID NO:71)
ATC ACT GTG ACC AAC ATC ACT GTG ACT GCC (SEQ ID NO:70)

NG21: Q T V T N I T V T A (SEQ ID NO:73)
CAG ACT GTG ACC AAC ATC ACT GTG ACT GCC (SEQ ID NO:72)

NGV: A T N V S N N S N T S N D S N V S (SEQ ID NO:75)
GCC ACT AAT GTG TCT AAC AAC AGC AAC ACC AGC AAT GAC AGC AAT GTG TCT (SEQ ID NO:74)

Figure 13B



```
# Predictions for N-Glycosylation sites in 1 sequence

Name:  CBG_HUMAN        Length:  405

(Sequence)
Asn-Xaa-Ser/Thr sequons (including Asn-Pro-Ser/Thr) are shown in blue.
Asparagines predicted to be N-glycosylated are shown in red.
Note that not all sequons are predicted glycosylated.

MDLLLYTCLLWLPTSGLNTVQAMDPNAAIYVKSPRRSQLASAVVDFAFSLYRELVALSPDKNIFISPVSISMALAMLSLG      80
TQGHTRAQLLQGLGFNLTERSETEIHQGFQHLHQLFAKSDTSLEMTMGNALFLDGSLELLESFSADIKHYYESEVLAMNF     160
QDWATASRQINSYVSKGTQGKIVDLFSGLDSPAILVLVNYIFFKGTWTQPFDLASTRERNFYVDETTVVKVPMMLQSSTI     240
SYLHDSELPCQLVQMDYVGNGTVFFILPDRGKMQTVIAALSRDTINKWSAGLTSSQVDLYIPKVTISGVYDLGDVLEDMG     320
IADLFTNQANFSRITQDAQLKSSEVVHKAVLQLNEEGVDTAGSTGVTLNLTSKPIILRFNQPFIIMIFDNFTWSSLFLAR     400
VNSTV

(Annotation line)
`N' represents a predicted N-glycosylation site.
`n' represents an Asn with a positive score,
but not occuring within an Asn-Xaa-Ser/Thr sequon

.................................N.....................................................      80
.................N.....................................................................     160
.....................................................................................     240
.....................N...............................................................     320
.....................................................N...............................     400
.....

(Threshold=0.5)
-------------------------------------------------------------------------------
SeqName      Position  Potential   Jury    NGlyc
                                 agreement result
-------------------------------------------------------------------------------
CBG_HUMAN      31 NMSN   0.7166     (9/9)   ++  <-- Predicted as N-glycosylated (++)
CBG_HUMAN      96 NLTS   0.6356     (8/9)   +   <-- Predicted as N-glycosylated (+)
CBG_HUMAN     176 NKTQ   0.3941     (7/9)   -   <-- A negative site
CBG_HUMAN     260 NGTV   0.7400     (9/9)   ++
CBG_HUMAN     330 NFSR   0.4223     (7/9)   -      see below for
CBG_HUMAN     369 NLTS   0.6684     (9/9)   ++    more information
-------------------------------------------------------------------------------
```

Figure 14

```
Name: Sequence          Length: 41
LSKNNAIEPRSFSQNATNVSNNSNTSNDSNVSPPVLKRHQR
. . . . . N..N. . . . . .
(Threshold=0.5)
---------------------------------------------------------------
SeqName      Position  Potential   Jury    N-Glyc
                                 agreement result
---------------------------------------------------------------
Sequence       15 NATN   0.6224     (8/9)   +
Sequence       18 NVSN   0.6453     (9/9)   ++
Sequence       21 NNSN   0.4158     (8/9)   -
Sequence       24 NTSN   0.4158     (8/9)   -
Sequence       27 NDSN   0.3619     (8/9)   -
Sequence       30 NVSP   0.1149     (9/9)   ---
---------------------------------------------------------------
```

Figure 15

CS01-FL-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAAGCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGGGATCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCATGGCATCTGACCCACTCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGAAGCTTCTCTCAGAATCCACCTGTCCTGAAGAGA
CACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCAAGA
TCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGCATG
TCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG
GTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAGCAC

(Continued)

Figure 16A

CTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGAAAC
CAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAAGGG
GCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGACCTG
GAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
TCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAAGAC
CCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGGTCA
ACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATCAAG
ACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCTCTG
GATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGGCTG
CACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAACTCC
TGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCTTAC
TTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGCAAT
GCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATGAAA
GTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTCCTG
ATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTGTTC
CAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGATAC
CTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGAG
GCACAAGACCTGTACTGA (SEQ ID NO:13)

Figure 16B

CS08-FL-NA

ATGCAGATCGAACTGAGCACTTGCTTCTTCCTGTGTCTCCTGCGCTTTTGCTTCTCCGCCACAAGG
AGATACTATCTCGGTGCCGTGGAGCTCAGCTGGGACTACATGCAGAGCGACTTGGGTGAACTGCCT
GTGGACGCCAGGTTTCCACCCCGCGTGCCCAAGAGTTTCCCGTTCAACACCAGTGTCGTGTACAAG
AAAACCCTCTTCGTGGAATTCACCGACCACCTGTTCAACATCGCCAAACCGCGCCCTCCCTGGATG
GGGCTGCTCGGCCCGACGATCCAGGCTGAGGTCTATGACACGGTGGTGATTACCCTCAAGAACATG
GCTAGCCACCCGGTGAGCCTGCACGCCGTGGGCGTGTCCTATTGGAAAGCGTCCGAGGGTGCGGAG
TACGATGACCAGACTTCACAGCGGGAGAAGGAAGACGACAAAGTGTTCCCCGGGGGTTCCCACACC
TATGTCTGGCAGGTCCTGAAGGAGAATGGTCCTATGGCCTCCGACCCATTGTGCCTCACCTACTCT
TACCTAAGCCATGTGGATCTCGTCAAGGACCTGAACTCGGGGCTGATCGGCGCCCTGCTCGTGTGC
CGGGAGGGCTCACTGGCCAAGGAGAAGACCCAAACTCTGCACAAGTTCATCCTGCTGTTCGCGGTA
TTCGACGAGGGGAAGTCCTGGCACTCCGAGACCAAGAACAGCCTGATGCAGGACCGCGACGCAGCC
TCGGCCCGTGCGTGGCCAAAGATGCACACCGTGAACGGCTACGTTAACAGGAGCCTACCCGGCCTG
ATCGGCTGCCACCGCAAATCGGTCTACTGGCATGTGATCGGAATGGGCACAACGCCCGAGGTCCAC
AGTATCTTCCTCGAGGGCCACACTTTCCTGGTCCGGAATCACCGCCAGGCCAGCCTGGAGATCAGC
CCCATAACCTTTCTGACGGCGCAGACCTTACTCATGGATCTCGGCCAGTTCCTCCTGTTCTGCCAC
ATTTCGTCCCACCAGCACGATGGGATGGAAGCATATGTGAAAGTGGACTCCTGCCCCGAGGAACCC
CAGCTTAGGATGAAGAACAATGAGGAGGCCGAGGACTACGACGATGACCTTACCGATTCAGAAATG
GACGTAGTACGCTTTGACGACGACAACTCTCCATCCTTCATACAGATTCGCTCCGTCGCCAAGAAG
CACCCTAAGACTTGGGTGCACTACATCGCGGCCGAGGAGGAGGACTGGGATTATGCTCCCCTGGTG
CTGGCCCCCGACGACCGCAGCTACAAGAGCCAGTACCTGAATAACGGGCCCCAGCGCATCGGCCGG
AAGTACAAGAAAGTGCGGTTCATGGCTTACACGGACGAGACCTTCAAGACCCGGGAGGCTATCCAG
CATGAGAGCGGCATCTTGGGGCCCCTCCTGTACGGCGAAGTTGGAGACACACTGCTGATCATCTTC
AAGAACCAGGCGAGCAGGCCCTACAACATCTACCCCCACGGCATTACCGATGTCCGGCCGTTGTAC
AGCCGACGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTTCCGATCCTGCCGGGCGAGATCTTC
AAGTACAAGTGGACTGTGACCGTGGAGGATGGGCCGACCAAGAGCGATCCGCGCTGCCTGACCCGT
TACTACTCCAGCTTTGTCAATATGGAGCGCGACCTCGCTAGCGGCTTGATTGGCCCTCTGCTGATC
TGCTACAAGGAGTCCGTGGACCAGAGGGGGAATCAGATCATGAGTGACAAGAGGAACGTGATCCTG
TTCTCCGTGTTCGACGAAAACCGCAGCTGGTATCTCACCGAGAATATCCAGCGCTTCCTGCCCAAC
CCGGCCGGTGTGCAGCTGGAGGACCCCGAGTTTCAGGCCAGCAACATCATGCATTCTATCAACGGA
TATGTGTTTGATTCCCTGCAGCTCTCAGTGTGTCTGCACGAGGTCGCCTACTGGTATATCCTCAGC
ATTGGGGCACAGACCGACTTCCTGAGCGTGTTCTTCTCCGGGTATACCTTCAAGCACAAGATGGTG
TACGAGGATACCCTGACCCTGTTCCCCTTTAGCGGCGAAACCGTGTTTATGTCTATGGAGAACCCC
GGGCTCTGGATCCTTGGCTGCCATAACTCCGACTTCCGCAACCGCGGAATGACCGCGCTCCTGAAA
GTGTCGAGTTGTGACAAGAACACCGGCGACTATTACGAGGACAGTTACGAGGACATCTCTGCGTAC
CTCCTTAGCAAGAATAACGCCATCGAGCCAAGATCCTTCAGCCAGAACCCCCCAGTGCTGAAGAGG
CATCAGCGGGAGATCACCCGCACGACCCTGCAGTCGGATCAGGAGGAGATTGATTACGACGACACG
ATCAGTGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAAGATGAAAACCAGTCCCCTCGG
TCCTTCCAAAAGAAGACCCGGCACTACTTCATCGCCGCTGTGGAACGCCTGTGGGACTATGGAATG

(Continued)

Figure 17A

TCTTCTAGCCCTCACGTTTTGAGGAACCGCGCCCAGTCGGGCAGCGTGCCCCAGTTCAAGAAAGTG
GTGTTCCAGGAGTTCACCGACGGCTCCTTCACCCAGCCACTTTACCGGGGCGAGCTCAATGAACAT
CTGGGCCTGCTGGGACCCTACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACATTCCGGAAT
CAGGCCAGCAGACCATACAGTTTCTACAGTTCACTCATCTCCTACGAGGAGGACCAGCGCCAGGGG
GCTGAACCCCGTAAGAACTTCGTGAAGCCAAACGAAACAAAGACCTACTTCTGGAAGGTCCAGCAC
CACATGGCACCTACCAAGGACGAGTTCGATTGCAAGGCCTGGGCCTACTTCTCCGACGTGGACCTG
GAGAAAGATGTGCACAGCGGCCTGATTGGCCCTCTGCTGGTGTGTCACACGAACACACTCAACCCT
GCACACGGGCGGCAGGTCACTGTGCAGGAATTCGCCCTGTTCTTTACCATCTTTGATGAGACGAAG
TCCTGGTATTTCACCGAAAACATGGAGAGGAACTGCCGCGCACCCTGCAACATCCAGATGGAAGAT
CCGACATTCAAGGAGAACTACCGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGC
CTCGTGATGGCCCAAGACCAGCGTATCCGCTGGTATCTGCTGTCGATGGGCTCCAACGAGAACATC
CATAGTATCCACTTCAGCGGGCATGTCTTCACGGTGAGGAAAAAGGAGGAGTACAAGATGGCACTG
TACAACCTCTATCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCTCCAAGGCCGGCATCTGGAGA
GTGGAATGCCTGATCGGCGAGCACCTCCACGCTGGGATGTCCACGCTGTTCCTCGTTTACAGCAAT
AAGTGCCAGACCCCTCTGGGCATGGCGAGCGGCCACATCCGCGACTTCCAGATTACAGCCAGCGGC
CAGTACGGTCAGTGGGCTCCAAAGCTGGCCCGTCTGCACTACTCCGGATCCATCAACGCCTGGTCC
ACCAAGGAACCGTTCTCCTGGATCAAAGTAGACCTGCTAGCCCCCATGATCATTCACGGCATCAAG
ACACAAGGCGCCCGACAGAAGTTCTCGAGCCTCTATATCTCCCAGTTCATCATCATGTATAGCCTG
GACGGAAAGAAGTGGCAGACTTACCGCGGAAACTCGACAGGGACCCTGATGGTATTCTTCGGTAAC
GTGGACAGCTCCGGAATCAAGCACAACATCTTCAACCCACCCATTATCGCCCGCTACATCCGCCTG
CACCCCACTCACTATAGCATTAGGTCCACCCTGCGAATGGAGCTCATGGGCTGTGACCTGAACAGC
TGTAGCATGCCCCTCGGCATGGAGTCTAAGGCGATCTCCGACGCACAGATAACGGCATCATCCTAC
TTTACCAACATGTTCGCTACCTGGTCCCCCTCCAAGGCCCGACTCCACCTGCAAGGGAGATCCAAC
GCCTGGCGGCCACAGGTCAACAATCCCAAGGAGTGGCTGCAAGTGGACTTTCAGAAAACTATGAAA
GTCACCGGAGTGACCACACAGGGAGTGAAGTCTCTGCTGACCAGCATGTACGTGAAGGAGTTCCTC
ATCTCCAGTTCGCAGGATGGCCACCAGTGGACGTTGTTCTTCCAAAACGGTAAAGTCAAAGTCTTC
CAAGGGAACCAGGACAGCTTTACACCCGTCGTGAACTCCCTGGACCCCCCGCTTCTCACTAGATAC
CTCCGCATCCACCCTCAGAGCTGGGTGCACCAGATTGCCCTGCGCATGGAGGTTCTGGGGTGTGAA
GCCCAGGACCTGTACTAA (SEQ ID NO:14)

Figure 17B

CS10-FL-NA

ATGCAGATTGAGCTCTCCACCTGCTTCTTTCTCTGCCTTCTTCGCTTCTGCTTTTCTGCCACACGC
AGGTACTATTTGGGAGCAGTGGAACTGAGCTGGGATTACATGCAGAGTGACCTTGGTGAACTTCCT
GTGGACGCTCGTTTTCCACCTAGAGTTCCCAAGTCCTTCCCCTTCAACACCTCAGTGGTCTACAAG
AAAACGCTGTTTGTGGAGTTCACTGACCACCTCTTCAACATTGCCAAACCAAGACCCCCTTGGATG
GGATTGCTGGGACCCACAATACAAGCAGAAGTCTACGACACGGTGGTGATTACCCTGAAGAACATG
GCGTCACACCCTGTTTCACTTCACGCTGTTGGGGTCAGTTATTGGAAAGCCTCAGAGGGTGCGGAA
TACGATGATCAAACCAGCCAGAGGGAGAAGGAAGATGACAAGGTCTTTCCTGGGGGTAGCCATACC
TATGTTTGGCAGGTGCTGAAAGAGAATGGGCCTATGGCCTCTGATCCCTTGTGCCTCACATACTCT
TACCTGAGTCACGTCGACCTGGTGAAAGACCTGAATAGCGGTCTGATTGGTGCACTGCTTGTTTGT
AGAGAGGGGAGTTTGGCCAAGGAGAAAACTCAGACTCTCCACAAGTTTATCCTCCTGTTTGCTGTG
TTCGACGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAGGACAGAGATGCTGCA
TCTGCAAGGGCTTGGCCAAAAATGCACACAGTGAACGGCTATGTGAATCGATCACTGCCAGGACTG
ATAGGCTGTCATCGCAAGTCAGTGTATTGGCACGTTATCGGGATGGGAACAACTCCAGAAGTGCAC
AGCATCTTCCTTGAGGGCCACACTTTCCTGGTTCGGAATCATAGACAGGCCAGCCTTGAGATCAGC
CCAATCACCTTTCTGACTGCCCAAACCTTGCTGATGGATCTGGGACAGTTCCTCCTGTTTTGTCAC
ATCTCCTCCCACCAACATGACGGGATGGAGGCTTATGTGAAGGTCGATAGCTGTCCGGAGGAACCA
CAACTGAGGATGAAGAACAACGAAGAGGCAGAGGACTATGACGACGATCTGACTGACAGTGAAATG
GACGTGGTTCGGTTCGACGATGACAATTCTCCTTCATTTATCCAGATCCGTTCCGTGGCCAAGAAG
CACCCCAAGACTTGGGTTCATTACATCGCTGCTGAGGAGGAGGATTGGGACTACGCGCCCTTGGTG
TTGGCCCCAGACGATCGCTCATACAAGAGCCAGTACCTTAACAATGGTCCACAAAGGATCGGCCGG
AAGTACAAGAAGGTTAGATTTATGGCTTATACCGACGAGACTTTTAAAACTAGGGAAGCAATTCAG
CATGAAAGTGGCATTCTTGGACCCCTGCTGTATGGCGAGGTTGGCGACACCCTGCTGATTATCTTT
AAGAACCAGGCAAGCCGGCCCTACAACATCTACCCGCACGGCATAACCGATGTACGACCCCTGTAC
AGTCGCAGACTTCCTAAAGGGGTGAAACACCTGAAGGACTTCCCAATTCTGCCCGGGGAGATCTTC
AAGTATAAATGGACCGTGACGGTTGAGGATGGTCCCACAAAGTCCGATCCGAGATGCCTTACCCGA
TATTATTCCAGCTTCGTGAACATGGAAAGGGACCTGGCCAGCGGGCTGATTGGCCCACTGCTGATT
TGTTACAAGGAGTCTGTCGATCAAAGAGGAAACCAAATAATGAGCGACAAACGTAACGTCATCCTG
TTCAGCGTCTTTGATGAGAATAGAAGCTGGTACCTCACAGAAAATATTCAGCGGTTTCTGCCTAAC
CCCGCAGGCGTCCAGCTGGAAGATCCCGAGTTCCAAGCCTCAAACATCATGCATAGCATCAACGGA
TACGTATTCGATAGCCTGCAGCTGTCCGTCTGTCTCCATGAAGTGGCATATTGGTACATCCTGAGT
ATCGGGGCGCAGACCGACTTCCTGAGCGTGTTCTTTTCTGGATACACGTTCAAACACAAAATGGTC
TATGAAGATACCCTGACTCTGTTTCCATTCTCAGGAGAGACAGTCTTTATGAGTATGGAAAATCCT
GGACTGTGGATCCTGGGCTGTCACAATTCTGATTTTCGGAACAGAGGCATGACAGCCCTGCTTAAA
GTGAGCTCATGCGACAAGAACACCGGTGATTACTACGAAGATAGCTATGAGGACATCAGTGCGTAT
TTGCTCTCCAAGAACAACGCTATCGAGCCACGGTCTTTCAGTCAGAATCCTCCCGTTCTGAAGCGG
CATCAGCGCGAAATAACACGCACAACCCTTCAGTCAGACCAAGAGGAAATCGACTACGATGATACT
ATCTCTGTGGAGATGAAGAAGGAGGATTTCGACATTTACGACGAGGACGAGAATCAGTCCCCAAGG
AGCTTTCAGAAGAAAACAAGACACTATTTCATTGCCGCCGTGGAGCGACTGTGGGACTACGGCATG

(Continued)

Figure 18A

TCTAGCTCTCCGCATGTACTTAGAAATAGGGCACAAAGCGGATCCGTGCCTCAGTTTAAGAAAGTT
GTCTTTCAGGAGTTTACAGATGGCTCCTTCACCCAGCCCTTGTATCGCGGGGAACTCAATGAACAC
CTGGGCCTCCTGGGTCCTTATATTAGGGCCGAAGTCGAGGACAATATCATGGTGACCTTTAGGAAC
CAGGCATCTAGACCTTACTCTTTCTACTCCTCCCTGATATCCTATGAGGAGGACCAGCGGCAAGGC
GCTGAGCCTCGGAAGAACTTTGTGAAGCCAAATGAAACCAAAACATACTTTTGGAAAGTTCAGCAC
CACATGGCTCCCACGAAGGACGAATTTGACTGTAAAGCCTGGGCCTACTTCTCAGATGTAGATCTC
GAGAAAGACGTGCACTCAGGGCTCATTGGTCCCCTCCTGGTCTGTCATACTAATACCCTCAATCCA
GCACACGGACGTCAGGTAACCGTCCAGGAATTTGCCCTGTTCTTTACCATTTTCGATGAGACTAAA
TCCTGGTACTTTACCGAAAACATGGAGAGGAATTGCAGAGCCCCATGCAACATCCAGATGGAGGAC
CCTACCTTCAAAGAGAACTATCGCTTCCATGCCATTAACGGTTACATTATGGATACTCTCCCAGGA
CTTGTGATGGCACAGGATCAGCGGATAAGATGGTATCTGTTGAGCATGGGCTCCAACGAGAATATT
CACAGCATCCATTTCTCCGGTCACGTGTTTACAGTGAGAAAGAAAGAAGAGTACAAGATGGCTCTG
TATAATCTCTATCCAGGCGTATTCGAAACGGTGGAGATGTTGCCTAGCAAGGCCGGCATTTGGCGA
GTAGAATGCCTTATCGGGGAACATCTGCATGCCGGAATGAGCACGCTCTTCCTGGTGTATAGTAAC
AAGTGCCAGACTCCGCTGGGCATGGCATCTGGCCATATACGGGACTTTCAGATTACGGCTAGCGGG
CAGTATGGGCAGTGGGCACCCAAACTTGCGCGACTGCACTATTCAGGCTCTATCAATGCATGGTCC
ACCAAGGAACCCTTCTCTTGGATTAAGGTGGACCTTTTGGCGCCCATGATAATCCATGGGATCAAA
ACCCAGGGCGCTCGTCAGAAATTCTCATCACTCTACATCTCTCAGTTCATAATAATGTATTCACTG
GATGGGAAGAAATGGCAGACTTACAGAGGAAACAGCACCGGGACGCTGATGGTGTTCTTTGGCAAC
GTGGACAGCAGCGGCATCAAACACAACATCTTCAATCCTCCCATTATTGCCCGTTATATTAGACTG
CATCCCACTCACTACTCTATACGCAGCACACTTAGGATGGAGCTCATGGGATGCGACCTGAACAGT
TGTAGTATGCCCTTGGGGATGGAGTCCAAAGCTATAAGCGACGCACAAATTACAGCTAGCTCTTAC
TTTACGAATATGTTCGCCACGTGGAGCCCAAGCAAAGCCCGGCTGCATTTGCAGGGTCGGAGTAAT
GCTTGGCGCCCACAGGTGAATAACCCTAAGGAATGGTTGCAAGTAGATTTCCAGAAAACTATGAAG
GTAACCGGCGTCACTACACAGGGAGTCAAGTCCCTCTTGACCTCTATGTACGTCAAGGAGTTCCTG
ATTAGCAGCAGTCAGGATGGGCACCAATGGACACTGTTCTTCCAGAATGGGAAAGTTAAAGTATTT
CAGGGTAACCAGGACTCCTTTACACCTGTGGTGAATAGCCTCGACCCACCCCTGCTGACACGATAC
CTCCGCATCCACCCTCAGTCTTGGGTGCATCAAATTGCCCTGCGAATGGAGGTGTTGGGATGCGAA
GCTCAGGACCTCTACTGA (SEQ ID NO:15)

Figure 18B

CS11-FL-NA

ATGCAGATCGAACTCTCTACTTGCTTCTTCCTGTGCCTTCTGAGGTTCTGCTTCTCTGCCACTCGC
CGATATTACCTCGGGGCCGTGGAGTTGAGTTGGGACTACATGCAATCAGATCTGGGCGAACTCCCT
GTGGATGCCCGATTCCCACCGCGCGTGCCCAAGTCTTTCCCATTTAATACTTCTGTGGTGTACAAG
AAGACATTGTTTGTGGAGTTTACCGATCACCTGTTCAACATCGCCAAACCGCGGCCCCCATGGATG
GGTCTGCTTGGGCCCACCATTCAAGCGGAGGTCTATGATACAGTGGTGATAACGCTTAAGAACATG
GCGAGCCACCCAGTGTCTCTGCATGCCGTTGGTGTATCATATTGGAAGGCCAGCGAAGGAGCGGAG
TACGATGACCAGACCTCTCAGAGAGAGAAGGAAGACGATAAGGTTTTTCCTGGCGGAAGTCATACA
TATGTATGGCAGGTCCTGAAAGAGAATGGGCCGATGGCTTCTGACCCCCTTTGTCTTACCTATAGT
TATCTGAGCCACGTGGACCTGGTCAAGGACCTCAACAGTGGTCTGATTGGGGCTCTGCTTGTTTGT
AGAGAGGGTAGCTTGGCTAAGGAGAAAACCCAAACACTCCATAAGTTCATTTTGCTGTTCGCGGTG
TTCGACGAGGGAAAGAGTTGGCACAGCGAAACAAAGAATTCACTGATGCAAGACAGGGACGCCGCT
TCCGCAAGGGCTTGGCCTAAGATGCATACGGTGAATGGGTATGTGAACCGGAGCCTCCCGGGGCTG
ATCGGGTGCCATCGCAAGTCTGTTTACTGGCACGTCATTGGAATGGGGACAACGCCAGAGGTACAT
AGTATATTTCTTGAAGGCCACACGTTCCTCGTACGGAACCACCGACAGGCTTCCCTGGAGATAAGC
CCCATTACCTTTCTGACCGCTCAGACTCTGCTGATGGACCTTGGCCAGTTTCTCCTGTTCTGCCAT
ATTAGCAGCCACCAGCACGACGGTATGGAAGCATACGTGAAAGTCGATAGCTGTCCTGAGGAGCCT
CAGCTCAGAATGAAGAACAACGAGGAGGCCGAAGACTATGACGATGACCTTACAGATTCCGAGATG
GACGTGGTGCGCTTTGACGACGATAACAGTCCTAGTTTCATTCAAATCAGATCCGTAGCCAAAAAG
CATCCAAAGACATGGGTGCATTACATTGCAGCCGAAGAGGAGGATTGGGATTATGCGCCCCTTGTT
CTGGCTCCAGATGACAGGAGCTATAAGTCCCAGTACTTGAACAACGGGCCACAGCGAATCGGTAGA
AAATATAAGAAGGTAAGATTCATGGCCTACACTGACGAAACATTTAAAACCAGGGAAGCTATCCAA
CACGAATCTGGAATTCTCGGCCCTCTGCTCTACGGTGAGGTGGGGGACACCTTGCTGATCATTTTC
AAAAATCAGGCATCCAGGCCTTACAACATATACCCCCATGGCATCACCGATGTCCGCCCGCTGTAT
TCCAGAAGACTCCCCAAGGGAGTGAAACATCTGAAAGATTTTCCCATCCTGCCGGGCGAGATCTTT
AAATACAAATGGACTGTGACTGTAGAGGACGGGCCTACAAAATCAGACCCACGGTGCCTGACAAGG
TATTACAGTAGCTTCGTCAACATGGAACGCGACCTCGCCAGCGGACTCATTGGCCCACTGTTGATC
TGTTACAAAGAGTCAGTGGATCAGAGGGGAAATCAGATCATGAGCGATAAGAGAAACGTTATCCTG
TTTAGTGTCTTCGACGAGAACCGGTCTTGGTACCTTACTGAGAACATCCAGAGGTTCCTGCCGAAT
CCGGCTGGCGTTCAGCTCGAGGACCCAGAGTTCCAGGCCAGTAATATAATGCACTCAATCAACGGT
TATGTGTTCGATAGCCTGCAGCTGAGCGTCTGCCTCCACGAGGTAGCCTATTGGTACATATTGTCC
ATCGGGGCTCAGACCGATTTTCTGTCCGTGTTCTTTAGCGGGTATACCTTTAAACATAAAATGGTC
TATGAAGACACCCTGACCCTGTTCCCATTCTCCGGTGAGACTGTGTTCATGTCCATGGAGAACCCA
GGGCTGTGGATCCTGGGGTGTCACAATAGTGACTTTAGGAATCGGGGAATGACGGCACTGCTGAAG
GTGAGTTCTTGCGATAAAAATACAGGAGATTACTATGAGGATAGTTACGAGGATATCAGTGCCTAT
CTGCTTTCAAAAAACAACGCAATTGAGCCCCGGTCTTTCTCACAAAACCCCCCGGTGCTGAAGCGC
CACCAGCGCGAAATTACCCGGACAACCTTGCAGTCCGACCAGGAGGAAATCGATTATGACGATACT
ATCAGTGTAGAAATGAAAAAGGAGGATTTTGATATTTACGACGAAGACGAGAACCAGTCTCCGCGA

(Continued)

Figure 19A

AGTTTTCAGAAGAAAACGCGACACTACTTTATAGCTGCCGTGGAACGACTCTGGGATTATGGCATG
TCCTCCAGCCCTCATGTCCTTAGGAATCGAGCGCAGAGTGGCTCTGTGCCTCAGTTCAAAAAGGTT
GTGTTCCAGGAATTCACCGACGGCTCATTTACCCAGCCGCTGTACAGAGGCGAACTCAACGAACAC
CTTGGGCTGCTTGGGCCATATATTCGAGCAGAGGTGGAAGATAATATCATGGTAACCTTTAGAAAC
CAGGCGTCAAGACCCTATTCCTTCTACAGTTCTCTGATCAGCTACGAGGAGGACCAAAGACAGGGA
GCTGAACCCAGGAAGAACTTTGTGAAACCTAATGAGACCAAGACCTACTTCTGGAAGGTCCAGCAC
CATATGGCCCCAACTAAAGATGAATTCGATTGCAAGGCCTGGGCTTATTTCAGCGACGTGGATCTC
GAAAAGGATGTGCACAGCGGGTTGATCGGACCGCTTTTGGTGTGCCACACAAATACCCTCAATCCT
GCCCACGGGCGGCAGGTCACAGTTCAAGAGTTTGCACTCTTCTTTACAATATTTGACGAGACAAAG
TCATGGTATTTTACAGAGAATATGGAGAGAAATTGTCGCGCACCTTGCAACATTCAGATGGAGGAC
CCCACATTTAAGGAGAATTACAGATTTCATGCTATCAATGGGTACATTATGGATACTCTGCCTGGT
CTGGTCATGGCCCAGGATCAGCGCATAAGGTGGTACTTGCTGAGCATGGGATCTAATGAGAATATA
CACAGCATTCACTTCAGTGGCCACGTTTTTACTGTTAGAAAGAAGGAGGAGTACAAAATGGCGCTC
TACAACCTTTACCCGGGTGTGTTTGAGACAGTGGAGATGCTGCCAAGCAAGGCAGGCATCTGGAGG
GTTGAGTGTCTTATTGGGGAGCATCTGCATGCTGGAATGTCCACCCTCTTTCTTGTGTACAGCAAT
AAGTGCCAGACACCGCTTGGCATGGCCAGCGGCCACATTAGGGACTTTCAGATAACTGCCAGTGGA
CAGTACGGCCAGTGGGCTCCCAAGCTTGCAAGACTCCACTACTCCGGAAGCATAAACGCATGGAGC
ACCAAGGAACCCTTCTCTTGGATTAAGGTGGACCTGCTGGCGCCAATGATCATTCACGGCATAAAA
ACCCAAGGGGCACGACAGAAATTTTCATCTTTGTATATTAGTCAGTTTATCATCATGTACAGCTTG
GATGGAAAGAAGTGGCAGACGTACAGGGGCAATTCTACAGGAACACTTATGGTGTTTTTTGGGAAT
GTCGATTCCAGCGGGATCAAACATAACATCTTCAATCCTCCTATTATCGCCCGATATATCCGCCTG
CACCCTACGCATTACTCCATCAGGTCCACATTGAGAATGGAACTGATGGGGTGCGACCTGAATAGT
TGTAGTATGCCACTGGGCATGGAGTCTAAAGCCATCAGCGATGCACAGATCACTGCCAGCTCTTAC
TTCACCAACATGTTTGCAACTTGGTCCCCCTCTAAAGCTCGCCTGCATCTGCAGGGACGCTCAAAT
GCATGGCGACCACAGGTGAACAATCCAAAAGAGTGGCTCCAGGTCGACTTTCAGAAGACAATGAAG
GTAACAGGAGTGACAACCCAGGGTGTAAAAAGCCTCCTTACGAGTATGTACGTTAAGGAGTTTCTG
ATTTCTAGCTCCCAGGACGGACACCAGTGGACTCTGTTCTTCCAGAACGGCAAAGTGAAGGTATTT
CAGGGAAACCAGGATTCTTTTACCCCGGTAGTGAATAGCCTGGATCCACCGTTGCTGACCCGCTAT
CTGAGAATTCATCCACAATCCTGGGTGCATCAGATTGCCCTCCGGATGGAAGTGCTCGGCTGTGAA
GCTCAGGATCTGTATTAG (SEQ ID NO:16)

Figure 19B

CS40-FL-NA

ATGCAAATAGAGCTCTCCACCTGCTTCTTTCTGTGCCTTTTGCGATTCTGCTTTAGTGCCACCAGA
AGATACTACCTGGGTGCAGTGGAACTGTCATGGGACTATATGCAAAGTGATCTCGGTGAGCTGCCT
GTGGACGCAAGATTTCCTCCTAGAGTGCCAAAATCTTTTCCATTCAACACCTCAGTCGTGTACAAA
AAGACTCTGTTTGTAGAATTCACGGATCACCTTTTCAACATCGCTAAGCCAAGGCCACCCTGGATG
GGTCTGCTAGGTCCTACCATCCAGGCTGAGGTTTATGATACAGTGGTCATTACACTTAAGAACATG
GCTTCCCATCCTGTCAGTCTTCATGCTGTTGGTGTATCCTACTGGAAAGCTTCTGAGGGAGCTGAA
TATGATGATCAGACCAGTCAAAGGGAGAAAGAAGATGATAAAGTCTTCCCTGGTGGAAGCCATACA
TATGTCTGGCAGGTCCTGAAAGAGAATGGTCCAATGGCCTCTGACCCACTGTGCCTTACCTACTCA
TATCTTTCTCATGTGGACCTGGTAAAAGACTTGAATTCAGGCCTCATTGGAGCCCTACTAGTATGT
AGAGAAGGGAGTCTGGCCAAGGAAAAGACACAGACCTTGCACAAATTTATACTACTTTTTGCTGTA
TTTGATGAAGGGAAAAGTTGGCACTCAGAAACAAAGAACTCCTTGATGCAGGATAGGGATGCTGCA
TCTGCTCGGGCCTGGCCTAAAATGCACACAGTCAATGGTTATGTAAACAGGTCTCTGCCAGGTCTG
ATTGGATGCCACAGGAAATCAGTCTATTGGCATGTGATTGGAATGGGCACCACTCCTGAAGTGCAC
TCAATATTCCTCGAAGGTCACACATTTCTTGTGAGGAACCATCGCCAGGCGTCCTTGGAAATCTCG
CCAATAACTTTCCTTACTGCTCAAACACTCTTGATGGACCTTGGACAGTTTCTACTGTTTTGTCAT
ATCTCTTCCCACCAACATGATGGCATGGAAGCTTATGTCAAAGTAGACAGCTGTCCAGAGGAACCC
CAACTACGAATGAAAAATAATGAAGAAGCGGAAGACTATGATGATGATCTTACTGATTCTGAAATG
GATGTGGTCAGGTTTGATGATGACAACTCTCCTTCCTTTATCCAAATTCGCTCAGTTGCCAAGAAG
CATCCTAAAACTTGGGTACATTACATTGCTGCTGAAGAGGAGGACTGGGACTATGCTCCCTTAGTC
CTCGCCCCCGATGACAGAAGTTATAAAAGTCAATATTTGAACAATGGCCCTCAGCGGATTGGTAGG
AAGTACAAAAAAGTCCGATTTATGGCATACACAGATGAAACCTTTAAGACTCGTGAAGCTATTCAG
CATGAATCAGGAATCTTGGGACCTTTACTTTATGGGGAAGTTGGAGACACACTGTTGATTATATTT
AAGAATCAAGCAAGCAGACCATATAACATCTACCCTCACGGAATCACTGATGTCCGTCCTTTGTAT
TCAAGGAGATTACCAAAAGGTGTAAAACATTTGAAGGATTTTCCAATTCTGCCAGGAGAAATATTC
AAATATAAATGGACAGTGACTGTAGAAGATGGGCCAACTAAATCAGATCCTCGGTGCCTGACCCGC
TATTACTCTAGTTTCGTTAATATGGAGAGAGATCTAGCTTCAGGACTCATTGGCCCTCTCCTCATC
TGCTACAAAGAATCTGTAGATCAAAGAGGAAACCAGATAATGTCAGACAAGAGGAATGTCATCCTG
TTTTCTGTATTTGATGAGAACCGAAGCTGGTACCTCACAGAGAATATACAACGCTTTCTCCCCAAT
CCAGCTGGAGTGCAGCTTGAGGATCCAGAGTTCCAAGCCTCCAACATCATGCACAGCATCAATGGC
TATGTTTTTGATAGTTTGCAGTTGTCAGTTTGTTTGCATGAGGTGGCATACTGGTACATTCTAAGC
ATTGGAGCACAGACTGACTTCCTTTCTGTCTTCTTCTCTGGATATACCTTCAAACACAAAATGGTC
TATGAAGACACACTCACCCTATTCCCATTCTCAGGAGAAACTGTCTTCATGTCGATGGAAAACCCA
GGTCTATGGATTCTGGGGTGCCACAACTCAGACTTTCGGAACAGAGGCATGACCGCCTTACTGAAG
GTTTCTAGTTGTGACAAGAACACTGGTGATTATTACGAGGACAGTTATGAAGATATTTCAGCATAC
TTGCTGAGTAAAAACAATGCCATTGAACCAAGAAGCTTCTCCCAGAATCCACCAGTCTTGAAACGC
CATCAACGGGAAATAACTCGTACTACTCTTCAGTCAGATCAAGAGGAAATTGACTATGATGATACC
ATATCAGTTGAAATGAAGAAGGAAGATTTTGACATTTATGATGAGGATGAAAATCAGAGCCCCCGC
AGCTTTCAAAAGAAAACACGACACTATTTTATTGCTGCAGTGGAGAGGCTCTGGGATTATGGGATG

(Continued)

Figure 20A

AGTAGCTCCCCACATGTTCTAAGAAACAGGGCTCAGAGTGGCAGTGTCCCTCAGTTCAAGAAAGTT
GTTTTCCAGGAATTTACTGATGGCTCCTTTACTCAGCCCTTATACCGTGGAGAACTAAATGAACAT
TTGGGACTCCTGGGGCCATATATAAGAGCAGAAGTTGAAGATAATATCATGGTAACTTTCAGAAAT
CAGGCCTCTCGTCCCTATTCCTTCTATTCTAGCCTTATTTCTTATGAGGAAGATCAGAGGCAAGGA
GCAGAACCTAGAAAAAACTTTGTCAAGCCTAATGAAACCAAAACTTACTTTTGGAAAGTGCAACAT
CATATGGCACCCACTAAAGATGAGTTTGACTGCAAAGCCTGGGCTTATTTCTCTGATGTTGACCTG
GAAAAAGATGTGCACTCAGGCCTGATTGGACCCCTTCTGGTCTGCCACACTAACACACTGAACCCT
GCTCATGGGAGACAAGTGACAGTACAGGAATTTGCTCTGTTTTTCACCATCTTTGATGAGACCAAA
AGCTGGTACTTCACTGAAAATATGGAAAGAAACTGCAGGGCTCCCTGCAATATCCAGATGGAAGAT
CCCACTTTTAAAGAGAATTATCGCTTCCATGCAATCAATGGCTACATAATGGATACACTACCTGGC
TTAGTAATGGCTCAGGATCAAAGGATTCGATGGTATCTGCTCAGCATGGGCAGCAATGAAAACATC
CATTCTATTCATTTCAGTGGACATGTGTTCACTGTACGAAAAAAAGAGGAGTATAAAATGGCACTG
TACAATCTCTATCCAGGTGTTTTTGAGACAGTGGAAATGTTACCATCCAAAGCTGGAATTTGGCGG
GTGGAATGCCTTATTGGCGAGCATCTACATGCTGGGATGAGCACACTTTTTCTGGTGTACAGCAAT
AAGTGTCAGACTCCCCTGGGAATGGCTTCTGGACACATTAGAGATTTTCAGATTACAGCTTCAGGA
CAATATGGACAGTGGGCCCCAAAGCTGGCCAGACTTCATTATTCCGGATCAATCAATGCCTGGAGC
ACCAAGGAGCCCTTTTCTTGGATCAAGGTGGATCTGTTGGCACCAATGATTATTCACGGCATCAAG
ACCCAGGGTGCCCGTCAGAAGTTCTCCAGCCTCTACATCTCTCAGTTTATCATCATGTATAGTCTT
GATGGGAAGAAGTGGCAGACTTATCGAGGAAATTCCACTGGAACCTTAATGGTCTTCTTTGGCAAT
GTGGATTCATCTGGGATAAAACACAATATTTTTAACCCTCCAATTATTGCTCGATACATCCGTTTG
CACCCAACTCATTATAGCATTCGCAGCACTCTTCGCATGGAGTTGATGGGCTGTGATTTAAATAGT
TGCAGCATGCCATTGGGAATGGAGAGTAAAGCAATATCAGATGCACAGATTACTGCTTCATCCTAC
TTTACCAATATGTTTGCCACCTGGTCTCCTTCAAAAGCTCGACTTCACCTCCAAGGGAGGAGTAAT
GCCTGGAGACCTCAGGTGAATAATCCAAAAGAGTGGCTGCAAGTGGACTTCCAGAAGACAATGAAA
GTCACAGGAGTAACTACTCAGGGAGTAAAATCTCTGCTTACCAGCATGTATGTGAAGGAGTTCCTC
ATCTCCAGCAGTCAAGATGGCCATCAGTGGACTCTCTTTTTTCAGAATGGCAAAGTAAAGGTTTTT
CAGGGAAATCAAGACTCCTTCACACCTGTGGTGAACTCTCTAGACCCACCGTTACTGACTCGCTAC
CTTCGAATTCACCCCCAGAGTTGGGTGCACCAGATTGCCCTGAGGATGGAGGTTCTGGGCTGCGAG
GCACAGGACCTCTACTGA (SEQ ID NO:17)

Figure 20B

CH25-FL-NA

ATGCAGATCGAGCTGTCCACATGCTTTTTTCTGTGCCTGCTGCGGTTCTGCTTCAGCGCCACCCGG
CGGTACTACCTGGGCGCCGTGGAGCTGTCCTGGGACTACATGCAGAGCGACCTGGGCGAGCTGCCC
GTGGACGCCCGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCAGCGTGGTGTACAAG
AAAACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGGCCTCCGAGGGCGCCGAG
TACGACGACCAGACCAGCCAGCGGGAGAAAGAGGACGACAAAGTCTTTCCTGGCGGCAGCCACACC
TACGTGTGGCAGGTCCTGAAAGAAAACGGCCCCATGGCCTCCGACCCCCTGTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACAGCGGGCTGATTGGGGCCCTGCTGGTCTGC
CGGGAGGGCAGCCTGGCCAAAGAGAAAACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGACGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACCGGGACGCCGCC
TCTGCCAGAGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGAAGCCTGCCCGGCCTG
ATTGGCTGCCACCGGAAGAGCGTGTACTGGCACGTGATCGGCATGGGCACCACACCCGAGGTGCAC
AGCATCTTTCTGGAAGGGCACACCTTTCTGGTGCGGAACCACCGGCAGGCCAGCCTGGAAATCAGC
CCTATCACCTTCCTGACCGCCCAGACACTGCTGATGGACCTGGGCCAGTTCCTGCTGTTTTGCCAC
ATCAGCTCTCACCAGCACGACGGCATGGAAGCCTACGTGAAGGTGGACTCCTGCCCCGAGGAACCC
CAGCTGCGGATGAAGAACAACGAGGAAGCCGAGGACTACGACGACGACCTGACCGACAGCGAGATG
GACGTGGTGCGGTTCGACGACGACAACAGCCCCAGCTTCATCCAGATCAGAAGCGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAAGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGAAGCTACAAGAGCCAGTACCTGAACAATGGCCCCCAGCGGATCGGCCGG
AAGTACAAGAAAGTGCGGTTCATGGCCTACACCGACGAGACCTTCAAGACCCGGGAGGCCATCCAG
CACGAGAGCGGCATCCTGGGCCCCCTGCTGTACGGCGAAGTGGGCGACACACTGCTGATCATCTTC
AAGAACCAGGCCAGCCGGCCCTACAACATCTACCCCCACGGCATCACCGACGTGCGGCCCCTGTAC
AGCAGGCGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGACGGCCCCACCAAGAGCGACCCCAGATGCCTGACCCGG
TACTACAGCAGCTTCGTGAACATGGAACGGGACCTGGCCTCCGGGCTGATCGGACCTCTGCTGATC
TGCTACAAAGAAAGCGTGGACCAGCGGGGCAACCAGATCATGAGCGACAAGCGGAACGTGATCCTG
TTCAGCGTGTTCGATGAGAACCGGTCCTGGTATCTGACCGAGAACATCCAGCGGTTTCTGCCCAAC
CCTGCCGGGGTGCAGCTGGAAGATCCCGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TACGTGTTCGACAGCCTGCAGCTGTCCGTGTGTCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGAGCGTGTTCTTCAGCGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCTTTCAGCGGCGAGACCGTGTTCATGAGCATGGAAAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCCGGAACCGGGGCATGACCGCCCTGCTGAAG
GTGTCCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGATATCAGCGCCTAC
CTGCTGTCCAAGAACAACGCCATCGAGCCCAGAAGCTTCAGCCAGAACCCCCCTGTGCTGAAGCGG
CACCAGAGAGAGATCACCCGGACCACCCTGCAGTCCGACCAGGAAGAGATCGATTACGACGACACC

(Continued)

Figure 21A

ATCAGCGTGGAGATGAAAAAAGAAGATTTCGACATCTACGACGAGGACGAGAACCAGAGCCCCCGG
TCCTTCCAGAAGAAAACCCGGCACTACTTTATCGCCGCCGTGGAGCGGCTGTGGGACTACGGCATG
AGCAGCAGCCCCCACGTGCTGCGGAACCGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAAGTG
GTGTTCCAGGAATTCACCGACGGCAGCTTCACCCAGCCCCTGTACCGGGGCGAGCTGAACGAGCAC
CTGGGGCTGCTGGGGCCCTACATCAGGGCCGAAGTGGAGGACAACATCATGGTGACCTTCCGGAAT
CAGGCCAGCAGACCCTACTCCTTCTACAGCAGCCTGATCAGCTACGAAGAGGACCAGCGGCAGGGC
GCTGAACCCCGGAAGAACTTCGTGAAGCCCAATGAGACCAAGACCTACTTCTGGAAAGTGCAGCAC
CACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCAGCGACGTGGATCTG
GAAAAGGACGTGCACTCTGGACTGATTGGCCCTCTGCTGGTGTGCCACACCAACACCCTGAACCCC
GCCCACGGCCGGCAGGTGACCGTGCAGGAATTCGCCCTGTTCTTCACCATCTTCGACGAGACCAAG
TCCTGGTACTTCACCGAGAATATGGAACGGAACTGCAGAGCCCCCTGCAACATCCAGATGGAAGAT
CCTACCTTCAAAGAGAACTACCGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCTGGC
CTGGTGATGGCCCAGGACCAGAGGATCCGGTGGTATCTGCTGTCCATGGGCAGCAACGAGAATATC
CACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAAGAAGAGTACAAGATGGCCCTG
TACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGGCGG
GTGGAGTGTCTGATCGGCGAGCACCTGCATGCCGGGATGAGCACCCTGTTTCTGGTGTACAGCAAC
AAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCCGGGACTTCCAGATCACCGCCTCCGGC
CAGTACGGCCAGTGGGCCCCCAAGCTGGCCCGGCTGCACTACAGCGGCAGCATCAACGCCTGGTCC
ACCAAAGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCTATGATCATCCACGGCATTAAG
ACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GACGGCAAGAAGTGGCAGACCTACCGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGCAAC
GTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCCGGTACATCCGGCTG
CACCCCACCCACTACAGCATCAGATCCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAACTCC
TGCAGCATGCCTCTGGGCATGGAAAGCAAGGCCATCAGCGACGCCCAGATCACAGCCAGCAGCTAC
TTCACCAACATGTTCGCCACCTGGTCCCCCTCCAAGGCCAGGCTGCACCTGCAGGGCCGGTCCAAC
GCCTGGCGGCCTCAGGTGAACAACCCCAAAGAATGGCTGCAGGTGGACTTTCAGAAAACCATGAAG
GTGACCGGCGTGACCACCCAGGGCGTGAAAAGCCTGCTGACCAGCATGTACGTGAAAGAGTTTCTG
ATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTTCAGAACGGCAAGGTGAAAGTGTTC
CAGGGCAACCAGGACTCCTTCACCCCCGTGGTGAACTCCCTGGACCCCCCCCTGCTGACCCGCTAC
CTGCGGATCCACCCCCAGTCTTGGGTGCACCAGATCGCCCTGAGGATGGAAGTGCTGGGATGTGAG
GCCCAGGATCTGTACTGA (SEQ ID NO:18)

Figure 21B

FVIII-FL-AA

mqielstcff lcllrfcfsa trryylgave lswdymqsdl gelpvdarfp prvpksfpfn
tsvvykktlf veftdhlfni akprppwmgl lgptiqaevy dtvvitlknm ashpvslhav
gvsywkaseg aeyddqtsqr ekeddkvfpg gshtyvwqvl kengpmasdp lcltysylsh
vdlvkdlnsg ligallvcre gslakektqt lhkfillfav fdegkswhse tknslmqdrd
aasarawpkm htvngyvnrs lpgligchrk svywhvigmg ttpevhsifl eghtflvrnh
rqasleispi tfltaqtllm dlgqfllfch isshqhdgme ayvkvdscpe epqlrmknne
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pddrsyksqy lnngpqrigr kykkvrfmay tdetfktrea iqhesgilgp llygevgdtl
liifknqasr pyniyphgit dvrplysrrl pkgvkhlkdf pilpgeifky kwtvtvedgp
tksdprcltr yyssfvnmer dlasgligpl licykesvdq rgnqimsdkr nvilfsvfde
nrswylteni qrflpnpagv qledpefqas nimhsingyv fdslqlsvcl hevaywyils
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gkvpflrvat essaktpskl ldplawdnhy gtqipkeewk sqekspekta fkkkdtilsl
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Figure 22

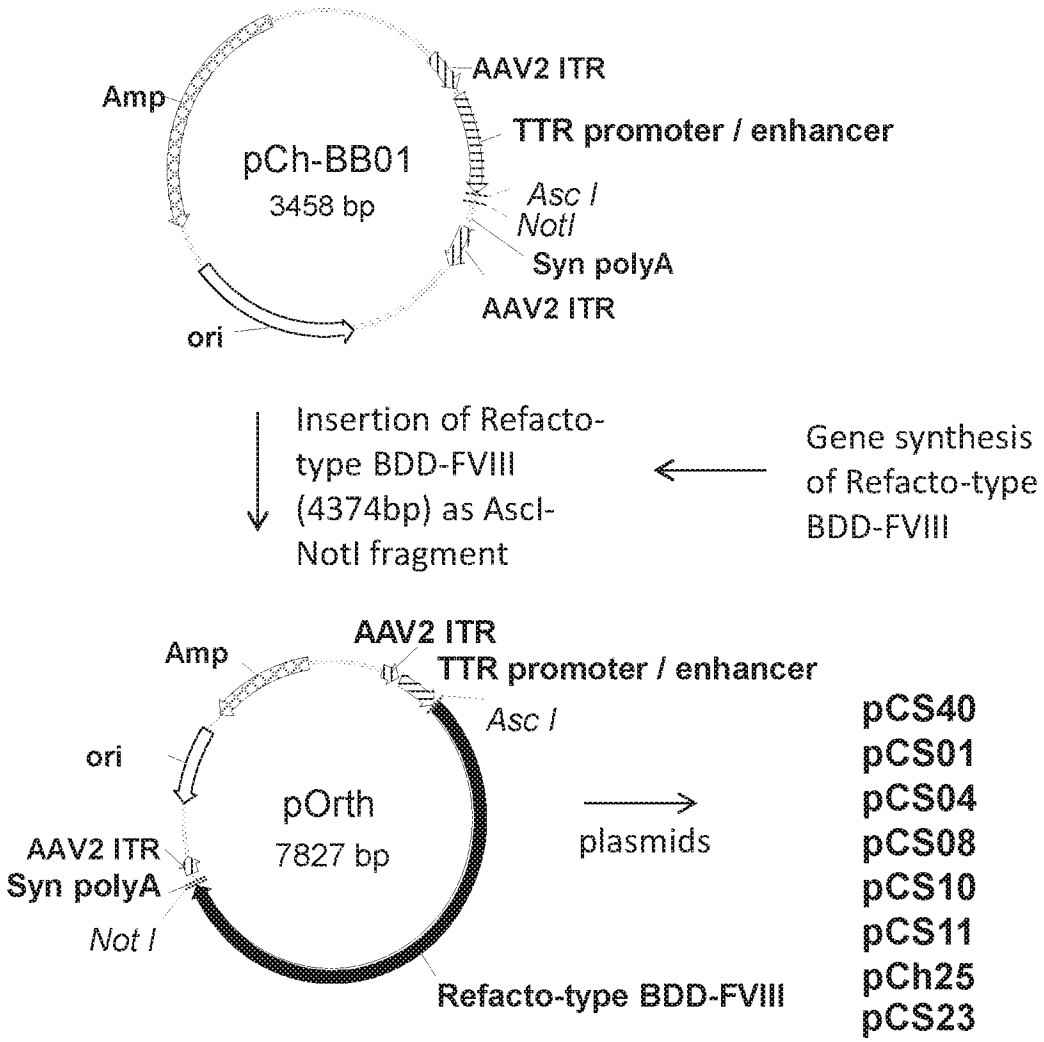
Amp
AAV2 ITR
TTR promoter / enhancer
pCh-BB01
3458 bp
Asc I
NotI
Syn polyA
AAV2 ITR
ori
Insertion of Refacto-type BDD-FVIII (4374bp) as AscI-NotI fragment
Gene synthesis of Refacto-type BDD-FVIII
Amp
AAV2 ITR
TTR promoter / enhancer
Asc I
ori
pOrth
7827 bp
AAV2 ITR
Syn polyA
Not I
plasmids
Refacto-type BDD-FVIII
pCS40
pCS01
pCS04
pCS08
pCS10
pCS11
pCh25
pCS23

Figure 23

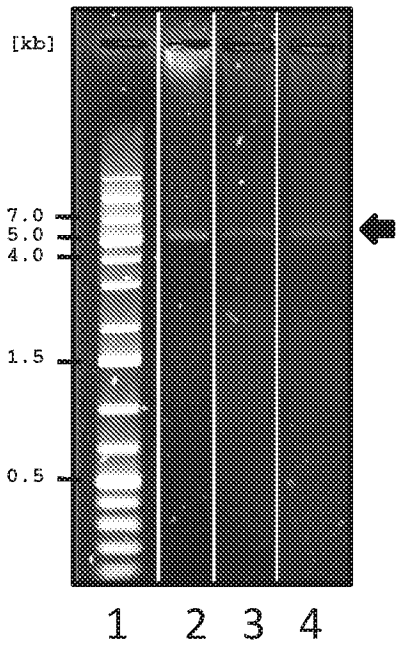

Figure 24

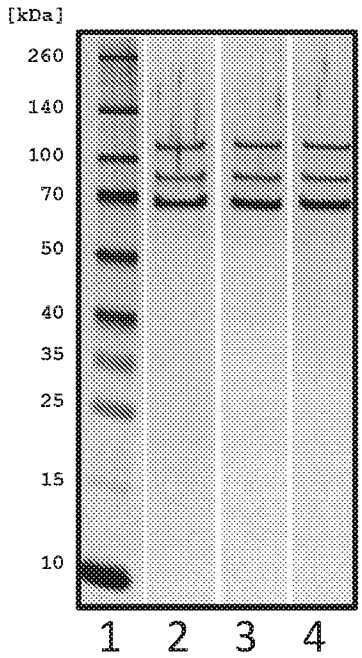

Figure 25

CS23-FL-NA

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cacgagtctggcatcctgggcccctgctgtacggcgaggtgggcgacaccctgctgatcatcttc
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(Continued)

Figure 26A

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Figure 26B

CS23-FL-AA

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YIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASSYFTNMFATWSPSKARLHLQG
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Figure 27

CS23-HC-NA

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(SEQ ID NO:22)

Figure 28

CS23-LC-NA

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(SEQ ID NO:23)

Figure 29

CS01m13-FL-NA

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ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGTCTCAGAATCCACCTGTCCTGAAG
AGACACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCA
AGATCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGC
ATGTCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 30A

GTGGTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAG
CACCTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGA
AACCAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAA
GGGGCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGAC
CTGGAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGTCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAA
GACCCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGG
TCAACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATC
AAGACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCT
CTGGATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGG
CTGCACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAAC
TCCTGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCT
TACTTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGC
AATGCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATG
AAAGTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTC
CTGATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTG
TTCCAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGA
TACCTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGT
GAGGCACAAGACCTGTACTGA (SEQ ID NO:90)

Figure 30B

CS01m23-FL-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAATCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGAAGTCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCACTGCATCTGACCCACCCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGTCTCAGAATCCACCTGTCCTGAAG
AGACACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCA
AGATCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGC
ATGTCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 31A

GTGGTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAG
CACCTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGA
AACCAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAA
GGGGCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGAC
CTGGAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGTCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAA
GACCCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGG
TCAACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATC
AAGACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCT
CTGGATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGG
CTGCACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAAC
TCCTGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCT
TACTTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGC
AATGCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATG
AAAGTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTC
CTGATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTG
TTCCAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGA
TACCTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGT
GAGGCACAAGACCTGTACTGA (SEQ ID NO:91)

Figure 31B

CS01m3-FL-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAAGCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGGGATCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCATGGCATCTGACCCACTCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGTCTCAGAATCCACCTGTCCTGAAG
AGACACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCA
AGATCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGC
ATGTCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 32A

GTGGTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAG
CACCTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGA
AACCAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAA
GGGGCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGAC
CTGGAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGTCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAA
GACCCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGG
TCAACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATC
AAGACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCT
CTGGATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGG
CTGCACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAAC
TCCTGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCT
TACTTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGC
AATGCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATG
AAAGTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTC
CTGATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTG
TTCCAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGA
TACCTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGT
GAGGCACAAGACCTGTACTGA (SEQ ID NO:92)

Figure 32B

CS01m2-FL-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAATCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGAAGTCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCACTGCATCTGACCCACCCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGAAGCTTCTCTCAGAATCCACCTGTCCTGAAGAGA
CACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCAAGA
TCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGCATG
TCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG

(Continued)

Figure 33A

GTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAGCAC
CTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGAAAC
CAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAAGGG
GCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGACCTG
GAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
TCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAAGAC
CCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGGTCA
ACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATCAAG
ACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCTCTG
GATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGGCTG
CACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAACTCC
TGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCTTAC
TTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGCAAT
GCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATGAAA
GTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTCCTG
ATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTGTTC
CAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGATAC
CTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGAG
GCACAAGACCTGTACTGA (SEQ ID NO:93)

Figure 33B

CS04m2-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGTCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGAAGAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCACTGCCTCTGACCCACCCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTTCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGGAGCTTCAGCCAGAATCCACCTGTCCTGAAACGC
CACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCAAGG
AGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGCATG
AGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG

(Continued)

Figure 34A

GTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAGCAC
CTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGCAAC
CAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAGGGG
GCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTG
GAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
AGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAGGAC
CCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGGAGC
ACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATCAAG
ACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGGCTG
CACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAACTCC
TGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGCTAC
TTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGCAAT
GCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAG
GTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTCCTG
ATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTGTTC
CAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGATAC
CTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGTGAG
GCCCAGGACCTGTACTGA (SEQ ID NO:94)

Figure 34B

CS04m3-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGGCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGGGCAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCATGGCCTCTGACCCACTCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTTCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGAGCCAGAATCCACCTGTCCTGAAA
CGCCACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCA
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGC
ATGAGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 35A

GTGGTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAG
CACCTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGC
AACCAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAG
GGGGCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGAGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAG
GACCCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGG
AGCACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATC
AAGACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGG
CTGCACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAAC
TCCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGC
TACTTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGC
AATGCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTC
CTGATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGA
TACCTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGT
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:95)

Figure 35B

CS04m23-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGTCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGAAGAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCACTGCCTCTGACCCACCCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTTCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGAGCCAGAATCCACCTGTCCTGAAA
CGCCACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCA
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGC
ATGAGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 36A

GTGGTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAG
CACCTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGC
AACCAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAG
GGGGCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGAGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAG
GACCCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGG
AGCACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATC
AAGACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGG
CTGCACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAAC
TCCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGC
TACTTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGC
AATGCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTC
CTGATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGA
TACCTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGT
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:96)

Figure 36B

CS04m1-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGGCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGGGCAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCATGGCCTCTGACCCACTCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTCCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGGAGCTTCAGCCAGAATCCACCTGTCCTGAAACGC
CACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCAAGG
AGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGCATG
AGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG

(Continued)

Figure 37A

GTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAGCAC
CTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGCAAC
CAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAGGGG
GCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTG
GAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
AGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAGGAC
CCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGGAGC
ACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATCAAG
ACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGGCTG
CACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAACTCC
TGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGCTAC
TTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGCAAT
GCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAG
GTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTCCTG
ATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTGTTC
CAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGATAC
CTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGTGAG
GCCCAGGACCTGTACTGA (SEQ ID NO:97)

Figure 37B

CS04m13-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGGCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGGGCAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCATGGCCTCTGACCCACTCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTCCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATACCACCTACGTGAACCGCTCCCTGAGCCAGAATCCACCTGTCCTGAAA
CGCCACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGAC
ACCATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCA
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGC
ATGAGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAA

(Continued)

Figure 38A

GTGGTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAG
CACCTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGC
AACCAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAG
GGGGCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAG
CACCACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAAC
CCTGCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACC
AAGAGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAG
GACCCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCT
GGGCTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAAC
ATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGG
AGGGTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGC
AACAAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCT
GGCCAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGG
AGCACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATC
AAGACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGC
AATGTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGG
CTGCACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAAC
TCCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGC
TACTTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGC
AATGCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTC
CTGATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGA
TACCTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGT
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:98)

Figure 38B

CS23m13-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGGCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCGGCAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCATGGCCAGCGACCCCCTGTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTCCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACACCACCTACGTGAACCGCTCCCTGAGCCAGAACCCCCCCGTGCTGAAG
AGGCACCAGAGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGAC
ACCATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCC
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGC
ATGAGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAG

(Continued)

Figure 39A

GTGGTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAG
CACCTGGGCCTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGG
AACCAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAG
GGCGCCGAGCCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAG
CACCACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAAC
CCCGCCCACGGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACC
AAGAGCTGGTACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAG
GACCCCACCTTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCC
GGCCTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAAC
ATCCACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGG
AGGGTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGC
AACAAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCT
GGCCAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGG
AGCACCAAGGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATC
AAGACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGACGGCAAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGC
AACGTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGG
CTGCACCCCACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAAC
AGCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGC
TACTTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGC
AACGCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTC
CTGATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCTGCTGACCAGG
TATCTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGC
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:99)

Figure 39B

CS23m3-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGGCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCGGCAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCATGGCCAGCGACCCCCTGTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTTCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACACCACCTACGTGAACCGCTCCCTGAGCCAGAACCCCCCCGTGCTGAAG
AGGCACCAGAGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGAC
ACCATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCC
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGC
ATGAGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAG

(Continued)

Figure 40A

GTGGTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAG
CACCTGGGCCTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGG
AACCAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAG
GGCGCCGAGCCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAG
CACCACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAAC
CCCGCCCACGGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACC
AAGAGCTGGTACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAG
GACCCCACCTTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCC
GGCCTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAAC
ATCCACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGG
AGGGTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGC
AACAAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCT
GGCCAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGG
AGCACCAAGGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATC
AAGACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGACGGCAAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGC
AACGTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGG
CTGCACCCCACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAAC
AGCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGC
TACTTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGC
AACGCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTC
CTGATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCTGCTGACCAGG
TATCTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGC
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:100)

Figure 40B

CS23m2-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGGTCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGTCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCAAGAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCACTGCCAGCGACCCCCCCTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTTCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACGCCATCGAGCCCAGGAGCTTCAGCCAGAACCCCCCCGTGCTGAAGAGG
CACCAGAGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGACACC
ATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCCAGG
AGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGCATG
AGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAGGTG

(Continued)

Figure 41A

GTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAGCAC
CTGGGCCTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGGAAC
CAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAGGGC
GCCGAGCCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAGCAC
CACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTG
GAGAAGGACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAACCCC
GCCCACGGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACCAAG
AGCTGGTACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAGGAC
CCCACCTTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCCGGC
CTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAACATC
CACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCCCTG
TACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGGAGG
GTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGCAAC
AAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCTGGC
CAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGGAGC
ACCAAGGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATCAAG
ACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GACGGCAAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGCAAC
GTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGGCTG
CACCCCACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAACAGC
TGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGCTAC
TTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGCAAC
GCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAG
GTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTCCTG
ATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTGTTC
CAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCCTGCTGACCAGGTAT
CTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGCGAG
GCCCAGGACCTGTACTGA (SEQ ID NO:101)

Figure 41B

CS23m1-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGGCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCGGCAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCATGGCCAGCGACCCCCTGTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTCCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACGCCATCGAGCCCAGGAGCTTCAGCCAGAACCCCCCCGTGCTGAAGAGG
CACCAGAGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGACACC
ATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCCAGG
AGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGCATG
AGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAGGTG

(Continued)

Figure 42A

GTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAGCAC
CTGGGCCTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGGAAC
CAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAGGGC
GCCGAGCCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAGCAC
CACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTG
GAGAAGGACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAACCCC
GCCCACGGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACCAAG
AGCTGGTACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAGGAC
CCCACCTTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCCGGC
CTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAACATC
CACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCCCTG
TACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGGAGG
GTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGCAAC
AAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCTGGC
CAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGGAGC
ACCAAGGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATCAAG
ACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GACGGCAAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGCAAC
GTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGGCTG
CACCCCACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAACAGC
TGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGCTAC
TTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGCAAC
GCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAG
GTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTCCTG
ATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTGTTC
CAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCCTGCTGACCAGGTAT
CTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGCGAG
GCCCAGGACCTGTACTGA (SEQ ID NO:102)

Figure 42B

CS23m23-FL-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGGTCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGTCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCAAGAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCACTGCCAGCGACCCCCCCTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTTCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACACCACCTACGTGAACCGCTCCCTGAGCCAGAACCCCCCCGTGCTGAAG
AGGCACCAGAGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGAC
ACCATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCC
AGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGC
ATGAGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAG

(Continued)

Figure 43A

GTGGTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAG
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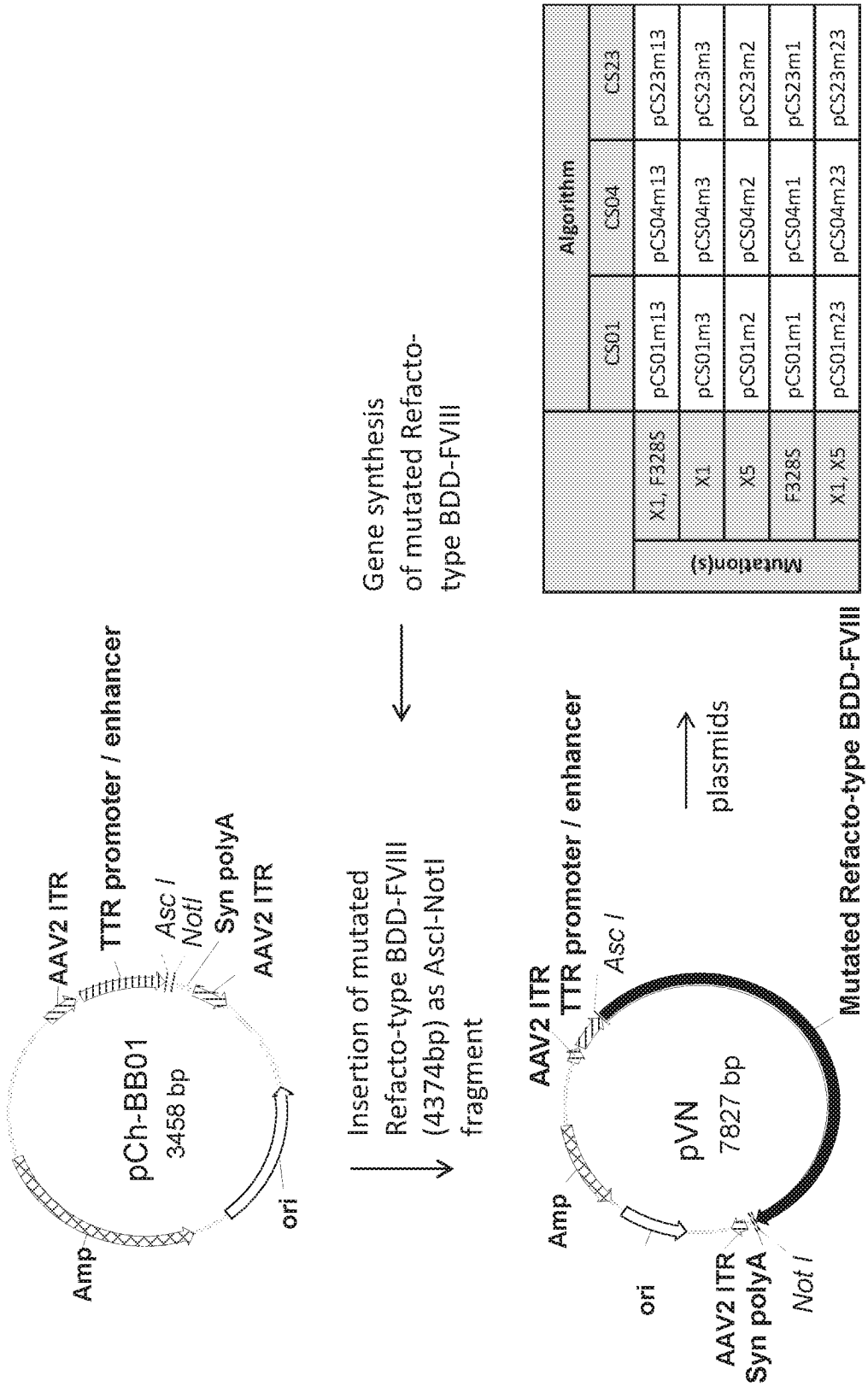
AACCAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAG
GGCGCCGAGCCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAG
CACCACATGGCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGAC
CTGGAGAAGGACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAAC
CCCGCCCACGGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACC
AAGAGCTGGTACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAG
GACCCCACCTTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCC
GGCCTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAAC
ATCCACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCC
CTGTACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGG
AGGGTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGC
AACAAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCT
GGCCAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGG
AGCACCAAGGAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATC
AAGACCCAGGGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGC
CTGGACGGCAAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGC
AACGTGGACAGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGG
CTGCACCCCACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAAC
AGCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGC
TACTTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGC
AACGCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATG
AAGGTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTC
CTGATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTG
TTCCAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCTGCTGACCAGG
TATCTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGC
GAGGCCCAGGACCTGTACTGA (SEQ ID NO:103)

Figure 43B



	Mutation(s)	Algorithm		
		CS01	CS04	CS23
	X1, F328S	pCS01m13	pCS04m13	pCS23m13
	X1	pCS01m3	pCS04m3	pCS23m3
	X5	pCS01m2	pCS04m2	pCS23m2
	F328S	pCS01m1	pCS04m1	pCS23m1
	X1, X5	pCS01m23	pCS04m23	pCS23m23

Figure 44

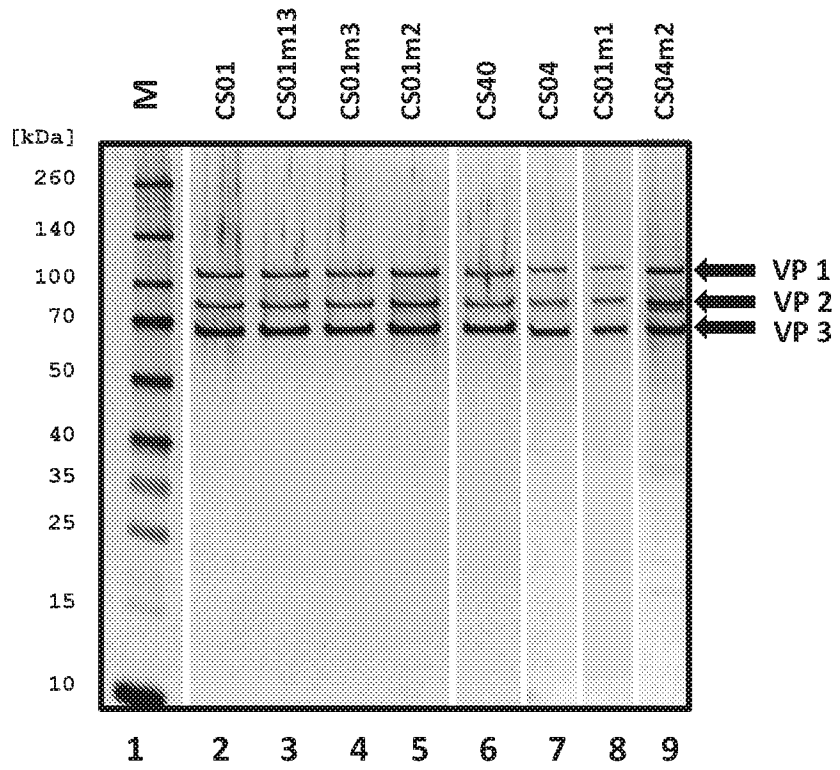


Figure 45

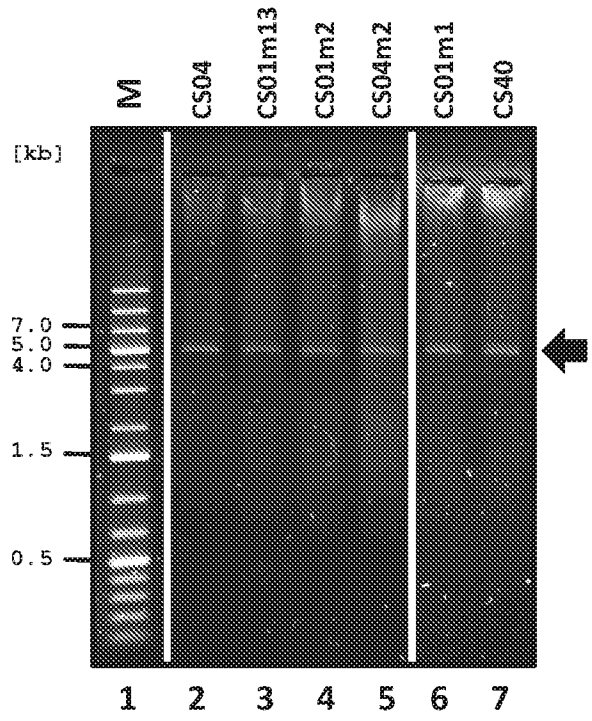


Figure 46

CS01-HC-NA

gcc
accaggagat actacctggg ggctgtggaa ctttcttggg actacatgca gtctgacctg
ggagagctgc ctgtggatgc caggttccca cccagagtgc ccaagtcctt cccattcaac
acctctgtgg tctacaagaa gacactcttt gtggaattca ctgaccacct gttcaacatt
gcaaaaccca gaccaccctg gatgggactc ctgggaccca ccattcaggc tgaggtgtat
gacactgtgg tcatcaccct caagaacatg gcatcccacc ctgtgtctct gcatgctgtg
ggagtctcat actggaaagc ctctgaaggg gctgagtatg atgaccagac atcccagaga
gagaaagagg atgacaaggt gttccctggg ggatctcaca cctatgtgtg gcaagtcctc
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gtggacctgg tcaaggacct caactctgga ctgattgggg cactgctggt gtgcagggaa
ggatccctgg ccaaggagaa aacccagaca ctgcacaagt tcattctcct gtttgctgtc
tttgatgagg gcaagtcttg gcactctgaa acaaagaact ccctgatgca agacagggat
gctgcctctg ccagggcatg gcccaagatg cacactgtga atggctatgt gaacagatca
ctgcctggac tcattggctg ccacaggaaa tctgtctact ggcatgtgat tggcatgggg
acaacccctg aagtgcactc cattttcctg gagggacaca ccttcctggt caggaaccac
agacaagcct ctctggagat ctctcccatc accttcctca ctgcacagac actgctgatg
gaccttggac agttcctgct gttctgccac atctcttccc accagcatga tggcatggaa
gcctatgtca aggtggactc atgccctgag gaaccacagc tcaggatgaa gaacaatgag
gaggctgagg actatgatga tgacctgact gactctgaga tggatgtggt cagatttgat
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tgggtgcact acattgctgc tgaggaagag gactgggact atgcaccact ggtcctggcc
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gatgtcaggc ccctgtacag caggagactg ccaaaagggg tgaaacacct caaggacttc
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acaaagtctg accccaggtg cctcaccaga tactactcct cttttgtgaa catggagaga
gacctggcat ctggactgat tggaccactg ctcatctgct acaaggagtc tgtggaccag
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aacagatcat ggtacctgac tgagaacatt cagagattcc tgcccaaccc tgctggggtg
caactggaag accctgagtt ccaggcaagc aacatcatgc actccatcaa tggctatgtg
tttgactctc tccagctttc tgtctgcctg catgaggtgg cctactggta cattctttct
attggggcac aaactgactt cctttctgtc ttcttctctg gatacacctt caagcacaag
atggtgtatg aggacaccct gacactcttc ccattctctg gggaaactgt gttcatgagc
atggagaacc ctggactgtg gattctggga tgccacaact ctgacttcag aaacagggga
atgactgcac tgctcaaagt ctcctcctgt gacaagaaca ctggggacta ctatgaggac
tcttatgagg acatctctgc ctacctgctc agcaagaaca atgccattga gcccaga
(SEQ ID NO:24X)

Figure 47

CS01-LC-NA

g agatcaccag gacaaccctc
cagtctgacc aggaagagat tgactatgat gacaccattt ctgtggagat gaagaaggag
gactttgaca tctatgatga ggacgagaac cagtctccaa gatcattcca gaagaagaca
agacactact tcattgctgc tgtggaaaga ctgtgggact atggcatgtc ttcctctccc
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caggagttca ctgatggctc attcacccag cccctgtaca gaggggaact gaatgagcac
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cagagacaag gggctgagcc aagaaagaac tttgtgaaac ccaatgaaac caagacctac
ttctggaaag tccagcacca catggcaccc accaaggatg agtttgactg caaggcctgg
gcatacttct ctgatgtgga cctggagaaa gatgtgcact ctggcctgat tggcccactc
ctggtctgcc acaccaacac cctgaaccct gcacatggaa ggcaagtgac tgtgcaggag
tttgccctct tcttcaccat ctttgatgaa accaagtcat ggtacttcac tgagaacatg
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tacaggttcc atgccatcaa tggctacatc atggacaccc tgcctgggct tgtcatggca
caggaccaga gaatcagatg gtacctgctt tctatgggat ccaatgagaa cattcactcc
atccacttct ctgggcatgt cttcactgtg agaaagaagg aggaatacaa gatggccctg
tacaacctct accctggggt ctttgagact gtggagatgc tgccctccaa agctggcatc
tggagggtgg aatgcctcat tggggagcac ctgcatgctg gcatgtcaac cctgttcctg
gtctacagca acaagtgcca gacacccctg ggaatggcct ctggccacat cagggacttc
cagatcactg cctctggcca gtatggccag tgggcaccca aactggccag gctccactac
tctggctcca tcaatgcatg gtcaaccaag gagccattct cttggatcaa ggtggacctg
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agaggcaact ccactggcac cctcatggtc ttctttggca atgtggacag ctctggcatc
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tactcaatca gatcaaccct caggatggaa ctgatgggat gtgacctgaa ctcctgctca
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aagacaatga aagtcactgg ggtgacaacc caggggtca agtctctgct cacctcaatg
tatgtgaagg agttcctgat ctcttcctca caggatggcc accagtggac actcttcttc
cagaatggca aagtcaaggt gttccagggc aaccaggact ctttcacacc tgtggtgaac
tcactggacc ccccctcct gacaagatac ctgagaattc acccccagtc ttgggtccac
cagattgccc tgagaatgga agtcctggga tgtgaggcac aagacctgta c
(SEQ ID NO:25)

Figure 48

CS01Δ(760-1667) - CS01-SC1-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGGAGATAC
TACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCTGTGGATGCCAGG
TTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAGAAGACACTCTTTGTGGAA
TTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATGGGACTCCTGGGACCCACCATTCAG
GCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATGGCATCCCACCCTGTGTCTCTGCATGCTGTG
GGAGTCTCATACTGGAAAGCCTCTGAAGGGGCTGAGTATGATGACCAGACATCCCAGAGAGAGAAAGAGGAT
GACAAGGTGTTCCCTGGGGGATCTCACACCTATGTGTGGCAAGTCCTCAAGGAGAATGGACCCATGGCATCT
GACCCACTCTGCCTGACATACTCCTACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATT
GGGGCACTGCTGGTGTGCAGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTC
CTGTTTGCTGTCTTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGAT
GCTGCCTCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCACTCCATT
TTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCTCCCATCACCTTC
CTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCACATCTCTTCCCACCAGCAT
GATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCACAGCTCAGGATGAAGAACAATGAG
GAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATGGATGTGGTCAGATTTGATGATGACAACTCT
CCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAACACCCCAAGACATGGGTGCACTACATTGCTGCTGAG
GAAGAGGACTGGGACTATGCACCACTGGTCCTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAAC
AATGGCCCACAAAGAATTGGAAGAAAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAG
ACAAGAGAAGCCATTCAGCATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTG
CTCATCATCTTCAAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCC
CTGTACAGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGATACTAC
TCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATCTGCTACAAGGAG
TCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTGTTCTCTGTCTTTGATGAG
AACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAACCCTGCTGGGGTGCAACTGGAAGAC
CCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGCTATGTGTTTGACTCTCTCCAGCTTTCTGTC
TGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCTATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTC
TCTGGATACACCTTCAAGCACAAGATGGTGTATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACT
GTGTTCATGAGCATGGAGAACCCTGGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGA
ATGACTGCACTGCTCAAAGTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGAC
ATCTCTGCCTACCTGCTCAGCAAGAACAATGCCATTGAGCCCAGAGAGATCACCAGGACAACCCTCCAGTCT
GACCAGGAAGAGATTGACTATGATGACACCATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGAT
GAGGACGAGAACCAGTCTCCAAGATCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGA
CTGTGGGACTATGGCATGTCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAG
TTCAAGAAAGTGGTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAAT
GAGCACCTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGAAAC
CAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAAGGGGCTGAG
CCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCACCACATGGCACCC

(Continued)

Figure 49A

ACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGACCTGGAGAAAGATGTGCACTCT
GGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCTGCACATGGAAGGCAAGTGACTGTG
CAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAGTCATGGTACTTCACTGAGAACATGGAGAGA
AACTGCAGAGCACCATGCAACATTCAGATGGAAGACCCCACCTTCAAGGAGAACTACAGGTTCCATGCCATC
AATGGCTACATCATGGACACCCTGCCTGGGCTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTT
TCTATGGGATCCAATGAGAACATTCACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAG
GAATACAAGATGGCCCTGTACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCT
GGCATCTGGAGGGTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTAC
AGCAACAAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGGTCAACCAAG
GAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATCAAGACACAGGGGGCA
AGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCTCTGGATGGCAAGAAGTGGCAG
ACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGCAATGTGGACAGCTCTGGCATCAAGCAC
AACATCTTCAACCCTCCCATCATTGCCAGATACATCAGGCTGCACCCCACCCACTACTCAATCAGATCAACC
CTCAGGATGGAACTGATGGGATGTGACCTGAACTCCTGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATT
TCTGATGCCCAGATCACTGCATCCTCTTACTTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGG
CTGCACCTCCAGGGAAGAAGCAATGCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGAC
TTCCAGAAGACAATGAAAGTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTG
AAGGAGTTCCTGATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAG
GTGTTCCAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGATAC
CTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGAGGCACAA
GACCTGTACTGA (SEQ ID NO:26)

Figure 49B

CS01Δ(772-1667) - CS01-SC2-NA

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCATCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAAGCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGGGATCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCATGGCATCTGACCCACTCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTTCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGAAGCTTCTCTCAGAATTCCAGACACCCCAGCACC
AGGGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGACACCATTTCT
GTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCAAGATCATTC

(Continued)

Figure 50A

CAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGCATGTCTTCC
TCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTGGTCTTC
CAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAGCACCTGGGA
CTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGAAACCAGGCC
TCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAAGGGGCTGAG
CCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCACCACATG
GCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGACCTGGAGAAA
GATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCTGCACAT
GGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAGTCATGG
TACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAAGACCCCACC
TTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGGCTTGTC
ATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAACATTCACTCC
ATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCCCTGTACAAC
CTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGGGTGGAA
TGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGCAACAAGTGC
CAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGCCAGTAT
GGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGGTCAACCAAG
GAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATCAAGACACAG
GGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCTCTGGATGGC
AAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGCAATGTGGAC
AGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGGCTGCACCCC
ACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAACTCCTGCTCA
ATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCTTACTTCACC
AACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGCAATGCCTGG
AGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATGAAAGTCACT
GGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTCCTGATCTCT
TCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTGTTCCAGGGC
AACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGATACCTGAGA
ATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGAGGCACAA
GACCTGTACTGA (SEQ ID NO:27)

Figure 50B

CS23Δ(760-1667) - CS23-SC1-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGGAGATAC
TACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCTGTGGACGCCAGG
TTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAGAAGACCCTGTTCGTGGAG
TTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATGGGCCTGCTGGGCCCCACCATCCAG
GCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATGGCCAGCCACCCCGTGAGCCTGCACGCCGTG
GGCGTGAGCTACTGGAAGGCCTCTGAGGGCGCCGAGTATGACGACCAGACCAGCCAGAGGGAGAAGGAGGAC
GACAAGGTGTTCCCCGGCGGCAGCCACACCTACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCATGGCCAGC
GACCCCCTGTGCCTGACCTACAGCTACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATC
GGCGCCCTGCTGGTGTGCAGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTG
CTGTTCGCCGTGTTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGAT
GCCGCCTCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCACAGCATC
TTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGCCCCATCACCTTC
CTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTTCTGCCACATCAGCAGCCACCAGCAC
GACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCCCAGCTGAGGATGAAGAACAACGAG
GAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATGGACGTGGTGAGGTTTGATGATGACAACAGC
CCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAGCACCCCAAGACCTGGGTGCACTACATCGCCGCCGAG
GAGGAGGACTGGGACTACGCCCCCCTGGTGCTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAAC
AACGGCCCCCAGAGGATCGGCAGGAAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAG
ACCAGGGAGGCCATCCAGCACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTG
CTGATCATCTTCAAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCC
CTGTACAGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGGTACTAC
AGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATCTGCTACAAGGAG
AGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTGTTCTCTGTGTTCGATGAG
AACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAACCCCGCCGGCGTGCAGCTGGAGGAC
CCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGCTACGTGTTCGACAGCCTGCAGCTGTCTGTG
TGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGCATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTC
TCTGGCTACACCTTCAAGCACAAGATGGTGTACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACC
GTGTTCATGAGCATGGAGAACCCCGGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGC
ATGACCGCCCTGCTGAAAGTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGAC
ATCAGCGCCTACCTGCTGAGCAAGAACAACGCCATCGAGCCCAGGGAGATCACCAGGACCACCCTGCAGAGC
GACCAGGAGGAGATCGACTATGATGACACCATCAGCGTGGAGATGAAGAAGGAGGACTTCGACATCTACGAC
GAGGACGAGAACCAGAGCCCCAGGAGCTTCCAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGG
CTGTGGGACTATGGCATGAGCAGCAGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAG
TTCAAGAAGGTGGTGTTCCAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAAC
GAGCACCTGGGCCTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGGAAC
CAGGCCAGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAGGGCGCCGAG

(Continued)

Figure 51A

CCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAGCACCACATGGCCCCC
ACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTGGAGAAGGACGTGCACAGC
GGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAACCCCGCCCACGGCAGGCAGGTGACCGTG
CAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACCAAGAGCTGGTACTTCACCGAGAACATGGAGAGG
AACTGCAGGGCCCCCTGCAACATCCAGATGGAGGACCCCACCTTCAAGGAGAACTACAGGTTCCACGCCATC
AACGGCTACATCATGGACACCCTGCCCGGCCTGGTGATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTG
AGCATGGGCAGCAACGAGAACATCCACAGCATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAG
GAGTACAAGATGGCCCTGTACAACCTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCC
GGCATCTGGAGGGTGGAGTGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTAC
AGCAACAAGTGCCAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCTGGC
CAGTACGGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGGAGCACCAAG
GAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATCAAGACCCAGGGCGCC
AGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTGGACGGCAAGAAGTGGCAG
ACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGCAACGTGGACAGCAGCGGCATCAAGCAC
AACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGGCTGCACCCCACCCACTACAGCATCAGGAGCACC
CTGCGGATGGAACTGATGGGCTGCGACCTGAACAGCTGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATC
TCTGACGCCCAGATCACCGCCAGCAGCTACTTCACCAACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGG
CTGCACCTGCAGGGCAGGAGCAACGCCTGGAGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGAC
TTCCAGAAGACCATGAAGGTGACCGGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTG
AAGGAGTTCCTGATCAGCAGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAG
GTGTTCCAGGGCAACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCTGCTGACCAGGTAT
CTGAGGATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGCGAGGCCCAG
GACCTGTACTGA (SEQ ID NO:28)

Figure 51B

CS23Δ(772-1667) - CS23-SC2-NA

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGCGCCGTGGAGCTGAGCTGGGACTACATGCAGTCTGACCTGGGCGAGCTGCCT
GTGGACGCCAGGTTCCCCCCCAGAGTGCCCAAGAGCTTCCCCTTCAACACCTCAGTGGTGTACAAG
AAGACCCTGTTCGTGGAGTTCACCGACCACCTGTTCAACATCGCCAAGCCCAGGCCCCCCTGGATG
GGCCTGCTGGGCCCCACCATCCAGGCCGAGGTGTACGACACCGTGGTGATCACCCTGAAGAACATG
GCCAGCCACCCCGTGAGCCTGCACGCCGTGGGCGTGAGCTACTGGAAGGCCTCTGAGGGCGCCGAG
TATGACGACCAGACCAGCCAGAGGGAGAAGGAGGACGACAAGGTGTTCCCCGGCGGCAGCCACACC
TACGTGTGGCAGGTGCTGAAGGAGAACGGCCCCATGGCCAGCGACCCCCTGTGCCTGACCTACAGC
TACCTGAGCCACGTGGACCTGGTGAAGGACCTGAACTCTGGCCTGATCGGCGCCCTGCTGGTGTGC
AGGGAGGGCAGCCTGGCCAAGGAGAAGACCCAGACCCTGCACAAGTTCATCCTGCTGTTCGCCGTG
TTCGATGAGGGCAAGAGCTGGCACAGCGAGACCAAGAACAGCCTGATGCAGGACAGGGATGCCGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACCGTGAACGGCTACGTGAACAGGAGCCTGCCCGGCCTG
ATCGGCTGCCACAGGAAGTCTGTGTACTGGCACGTGATCGGCATGGGCACCACCCCCGAGGTGCAC
AGCATCTTCCTGGAGGGCCACACCTTCCTGGTGAGGAACCACAGGCAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTGACCGCCCAGACCCTGCTGATGGACCTGGGCCAGTTCCTGCTGTTCTGCCAC
ATCAGCAGCCACCAGCACGACGGCATGGAGGCCTACGTGAAGGTGGACAGCTGCCCCGAGGAGCCC
CAGCTGAGGATGAAGAACAACGAGGAGGCCGAGGACTATGATGATGACCTGACCGACTCTGAGATG
GACGTGGTGAGGTTTGATGATGACAACAGCCCCAGCTTCATCCAGATCAGGTCTGTGGCCAAGAAG
CACCCCAAGACCTGGGTGCACTACATCGCCGCCGAGGAGGAGGACTGGGACTACGCCCCCCTGGTG
CTGGCCCCCGACGACAGGAGCTACAAGAGCCAGTACCTGAACAACGGCCCCCAGAGGATCGGCAGG
AAGTACAAGAAGGTCAGATTCATGGCCTACACCGACGAGACCTTCAAGACCAGGGAGGCCATCCAG
CACGAGTCTGGCATCCTGGGCCCCCTGCTGTACGGCGAGGTGGGCGACACCCTGCTGATCATCTTC
AAGAACCAGGCCAGCAGGCCCTACAACATCTACCCCCACGGCATCACCGATGTGAGGCCCCTGTAC
AGCAGGAGGCTGCCCAAGGGCGTGAAGCACCTGAAGGACTTCCCCATCCTGCCCGGCGAGATCTTC
AAGTACAAGTGGACCGTGACCGTGGAGGATGGCCCCACCAAGTCTGACCCCAGGTGCCTGACCAGG
TACTACAGCAGCTTCGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATCGGCCCCCTGCTGATC
TGCTACAAGGAGAGCGTGGACCAGAGGGGCAACCAGATCATGTCTGACAAGAGGAACGTGATCCTG
TTCTCTGTGTTCGATGAGAACAGGAGCTGGTATCTGACCGAGAACATCCAGAGGTTCCTGCCCAAC
CCCGCCGGCGTGCAGCTGGAGGACCCCGAGTTCCAGGCCAGCAACATCATGCACAGCATCAACGGC
TACGTGTTCGACAGCCTGCAGCTGTCTGTGTGCCTGCACGAGGTGGCCTACTGGTACATCCTGAGC
ATCGGCGCCCAGACCGACTTCCTGTCTGTGTTCTTCTCTGGCTACACCTTCAAGCACAAGATGGTG
TACGAGGACACCCTGACCCTGTTCCCCTTCAGCGGCGAGACCGTGTTCATGAGCATGGAGAACCCC
GGCCTGTGGATCCTGGGCTGCCACAACAGCGACTTCAGGAACAGGGGCATGACCGCCCTGCTGAAA
GTCAGCAGCTGCGACAAGAACACCGGCGACTACTACGAGGACAGCTACGAGGACATCAGCGCCTAC
CTGCTGAGCAAGAACAACGCCATCGAGCCCAGGAGCTTCAGCCAGAACTCCAGACACCCCAGCACC

(Continued)

Figure 52A

AGGGAGATCACCAGGACCACCCTGCAGAGCGACCAGGAGGAGATCGACTATGATGACACCATCAGC
GTGGAGATGAAGAAGGAGGACTTCGACATCTACGACGAGGACGAGAACCAGAGCCCCAGGAGCTTC
CAGAAGAAGACCAGGCACTACTTCATCGCCGCCGTGGAGAGGCTGTGGGACTATGGCATGAGCAGC
AGCCCCCACGTGCTGAGGAACAGGGCCCAGAGCGGCAGCGTGCCCCAGTTCAAGAAGGTGGTGTTC
CAGGAGTTCACCGACGGCAGCTTCACCCAGCCCCTGTACAGAGGCGAGCTGAACGAGCACCTGGGC
CTGCTGGGCCCCTACATCAGGGCCGAGGTGGAGGACAACATCATGGTGACCTTCAGGAACCAGGCC
AGCAGGCCCTACAGCTTCTACAGCAGCCTGATCAGCTACGAGGAGGACCAGAGGCAGGGCGCCGAG
CCCAGGAAGAACTTCGTGAAGCCCAACGAGACCAAGACCTACTTCTGGAAGGTGCAGCACCACATG
GCCCCCACCAAGGACGAGTTCGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTGGAGAAG
GACGTGCACAGCGGCCTGATCGGCCCCCTGCTGGTGTGCCACACCAACACCCTGAACCCCGCCCAC
GGCAGGCAGGTGACCGTGCAGGAGTTCGCCCTGTTCTTCACCATCTTCGACGAGACCAAGAGCTGG
TACTTCACCGAGAACATGGAGAGGAACTGCAGGGCCCCCTGCAACATCCAGATGGAGGACCCCACC
TTCAAGGAGAACTACAGGTTCCACGCCATCAACGGCTACATCATGGACACCCTGCCCGGCCTGGTG
ATGGCCCAGGACCAGAGGATCAGGTGGTATCTGCTGAGCATGGGCAGCAACGAGAACATCCACAGC
ATCCACTTCAGCGGCCACGTGTTCACCGTGAGGAAGAAGGAGGAGTACAAGATGGCCCTGTACAAC
CTGTACCCCGGCGTGTTCGAGACCGTGGAGATGCTGCCCAGCAAGGCCGGCATCTGGAGGGTGGAG
TGCCTGATCGGCGAGCACCTGCACGCCGGCATGAGCACCCTGTTCCTGGTGTACAGCAACAAGTGC
CAGACCCCCCTGGGCATGGCCAGCGGCCACATCAGGGACTTCCAGATCACCGCCTCTGGCCAGTAC
GGCCAGTGGGCCCCCAAGCTGGCCAGGCTGCACTACAGCGGCAGCATCAACGCCTGGAGCACCAAG
GAGCCCTTCAGCTGGATCAAGGTGGACCTGCTGGCCCCCATGATCATCCACGGCATCAAGACCCAG
GGCGCCAGGCAGAAGTTCAGCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTGGACGGC
AAGAAGTGGCAGACCTACAGGGGCAACAGCACCGGCACCCTGATGGTGTTCTTCGGCAACGTGGAC
AGCAGCGGCATCAAGCACAACATCTTCAACCCCCCCATCATCGCCAGGTACATCAGGCTGCACCCC
ACCCACTACAGCATCAGGAGCACCCTGCGGATGGAACTGATGGGCTGCGACCTGAACAGCTGCAGC
ATGCCCCTGGGCATGGAGAGCAAGGCCATCTCTGACGCCCAGATCACCGCCAGCAGCTACTTCACC
AACATGTTCGCCACCTGGAGCCCCAGCAAGGCCAGGCTGCACCTGCAGGGCAGGAGCAACGCCTGG
AGGCCCCAGGTGAACAACCCCAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAGGTGACC
GGCGTGACCACCCAGGGCGTGAAGAGCCTGCTGACCAGCATGTACGTGAAGGAGTTCCTGATCAGC
AGCAGCCAGGACGGCCACCAGTGGACCCTGTTCTTCCAGAACGGCAAAGTGAAGGTGTTCCAGGGC
AACCAGGACAGCTTCACCCCCGTGGTGAACAGCCTGGACCCCCCCCCTGCTGACCAGGTATCTGAGG
ATCCACCCCCAGAGCTGGGTGCACCAGATCGCCCTGAGAATGGAAGTGCTGGGATGCGAGGCCCAG
GACCTGTACTGA (SEQ ID NO:29)

Figure 52B

CS01m23-FL-AA (SEQ ID NO: 104)

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFN
TSVVYKKTLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVVTLKNMASHPVSLHAV
GVSYWKSSEGAEYDDQTSQREKEDDKVFPGKSHTYVWQVLKENGPTASDPPCLTYSYLSH
VDLVKDLNSGLIGALLVCREGSLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRD
AASARAWPKMHTVNGYVNRSLPGLIGCHRKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNH
RQASLEISPITFLTAQTLLMDLGQFLLFCHISSHQHDGMEAYVKVDSCPEEPQLRMKNNE
EAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKKHPKTWVHYIAAEEEDWDYAPLVLA
PDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQHESGILGPLLYGEVGDTL
LIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIFKYKWTVTVEDGP
TKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVILFSVFDE
NRSWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILS
IGAQTDFLSVFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRG
MTALLKVSSCDKNTGDYYEDSYEDISAYLLSKNNTTYVNRSLSQNPPVLKRHQREITRTT
LQSDQEEIDYDDTISVEMKKEDFDIYDEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSS
PHVLRNRAQSGSVPQFKKVVFQEFTDGSFTQPLYRGELNEHLGLLGPYIRAEVEDNIMVT
FRNQASRPYSFYSSLISYEEDQRQGAEPRKNFVKPNETKTYFWKVQHHMAPTKDEFDCKA
WAYFSDVDLEKDVHSGLIGPLLVCHTNTLNPAHGRQVTVQEFALFFTIFDETKSWYFTEN
MERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPGLVMAQDQRIRWYLLSMGSNENIH
SIHFSGHVFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWRVECLIGEHLHAGMSTLF
LVYSNKCQTPLGMASGHIRDFQITASGQYGQWAPKLARLHYSGSINAWSTKEPFSWIKVD
LLAPMIIHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGNVDSSG
IKHNIFNPPIIARYIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASS
YFTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTS
MYVKEFLISSSQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWV
HQIALRMEVLGCEAQDLY

Figure 53

CS04m3-FL-AA (SEQ ID NO: 105)

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFN
TSVVYKKTLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVITLKNMASHPVSLHAV
GVSYWKASEGAEYDDQTSQREKEDDKVFPGGSHTYVWQVLKENGPMASDPLCLTYSYLSH
VDLVKDLNSGLIGALLVCREGSLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRD
AASARAWPKMHTVNGYVNRSLPGLIGCHRKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNH
RQASLEISPITFLTAQTLLMDLGQFLLFCHISSHQHDGMEAYVKVDSCPEEPQLRMKNNE
EAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKKHPKTWVHYIAAEEEDWDYAPLVLA
PDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQHESGILGPLLYGEVGDTL
LIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIFKYKWTVTVEDGP
TKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVILFSVFDE
NRSWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILS
IGAQTDFLSVFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRG
MTALLKVSSCDKNTGDYYEDSYEDISAYLLSKNNTTYVNRSLSQNPPVLKRHQREITRTT
LQSDQEEIDYDDTISVEMKKEDFDIYDEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSS
PHVLRNRAQSGSVPQFKKVVFQEFTDGSFTQPLYRGELNEHLGLLGPYIRAEVEDNIMVT
FRNQASRPYSFYSSLISYEEDQRQGAEPRKNFVKPNETKTYFWKVQHHMAPTKDEFDCKA
WAYFSDVDLEKDVHSGLIGPLLVCHTNTLNPAHGRQVTVQEFALFFTIFDETKSWYFTEN
MERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPGLVMAQDQRIRWYLLSMGSNENIH
SIHFSGHVFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWRVECLIGEHLHAGMSTLF
LVYSNKCQTPLGMASGHIRDFQITASGQYGQWAPKLARLHYSGSINAWSTKEPFSWIKVD
LLAPMIIHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGNVDSSG
IKHNIFNPPIIARYIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASS
YFTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTS
MYVKEFLISSSQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWV
HQIALRMEVLGCEAQDLY

Figure 54

CS01-FL-AAm12 (SEQ ID NO: 106)

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFNTSVVYK
KTLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVVTLKNMASHPVSLHAVGVSYWKSSEGAE
YDDQTSQREKEDDKVFPGKSHTYVWQVLKENGPTASDPPCLTYSYLSHVDLVKDLNSGLIGALLVC
REGSLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRDAASARAWPKMHTVNGYVNRSLPGL
IGCHRKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNHRQASLEISPITFLTAQTLLMDLGQFLLSCH
ISSHQHDGMEAYVKVDSCPEEPQLRMKNNEEAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKK
HPKTWVHYIAAEEEDWDYAPLVLAPDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQ
HESGILGPLLYGEVGDTLLIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIF
KYKWTVTVEDGPTKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVIL
FSVFDENRSWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILS
IGAQTDFLSVFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRGMTALLK
VSSCDKNTGDYYEDSYEDISAYLLSKNNAIEPRSFSQNPPVLKRHQREITRTTLQSDQEEIDYDDT
ISVEMKKEDFDIYDEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSSPHVLRNRAQSGSVPQFKKV
VFQEFTDGSFTQPLYRGELNEHLGLLGPYIRAEVEDNIMVTFRNQASRPYSFYSSLISYEEDQRQG
AEPRKNFVKPNETKTYFWKVQHHMAPTKDEFDCKAWAYFSDVDLEKDVHSGLIGPLLVCHTNTLNP
AHGRQVTVQEFALFFTIFDETKSWYFTENMERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPG
LVMAQDQRIRWYLLSMGSNENIHSIHFSGHVFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWR
VECLIGEHLHAGMSTLFLVYSNKCQTPLGMASGHIRDFQITASGQYGQWAPKLARLHYSGSINAWS
TKEPFSWIKVDLLAPMIIHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGN
VDSSGIKHNIFNPPIIARYIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASSY
FTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTSMYVKEFL
ISSSQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWVHQIALRMEVLGCE
AQDLY

Figure 55

CS04-FL-AAm12 (SEQ ID NO: 107)

MQIELSTCFFLCLLRFCFSATRRYYLGAVELSWDYMQSDLGELPVDARFPPRVPKSFPFNTSVVYK
KTLFVEFTDHLFNIAKPRPPWMGLLGPTIQAEVYDTVVVTLKNMASHPVSLHAVGVSYWKSSEGAE
YDDQTSQREKEDDKVFPGKSHTYVWQVLKENGPTASDPPCLTYSYLSHVDLVKDLNSGLIGALLVC
REGSLAKEKTQTLHKFILLFAVFDEGKSWHSETKNSLMQDRDAASARAWPKMHTVNGYVNRSLPGL
IGCHRKSVYWHVIGMGTTPEVHSIFLEGHTFLVRNHRQASLEISPITFLTAQTLLMDLGQFLLSCH
ISSHQHDGMEAYVKVDSCPEEPQLRMKNNEEAEDYDDDLTDSEMDVVRFDDDNSPSFIQIRSVAKK
HPKTWVHYIAAEEEDWDYAPLVLAPDDRSYKSQYLNNGPQRIGRKYKKVRFMAYTDETFKTREAIQ
HESGILGPLLYGEVGDTLLIIFKNQASRPYNIYPHGITDVRPLYSRRLPKGVKHLKDFPILPGEIF
KYKWTVTVEDGPTKSDPRCLTRYYSSFVNMERDLASGLIGPLLICYKESVDQRGNQIMSDKRNVIL
FSVFDENRSWYLTENIQRFLPNPAGVQLEDPEFQASNIMHSINGYVFDSLQLSVCLHEVAYWYILS
IGAQTDFLSVFFSGYTFKHKMVYEDTLTLFPFSGETVFMSMENPGLWILGCHNSDFRNRGMTALLK
VSSCDKNTGDYYEDSYEDISAYLLSKNNAIEPRSFSQNPPVLKRHQREITRTTLQSDQEEIDYDDT
ISVEMKKEDFDIYDEDENQSPRSFQKKTRHYFIAAVERLWDYGMSSSPHVLRNRAQSGSVPQFKKV
VFQEFTDGSFTQPLYRGELNEHLGLLGPYIRAEVEDNIMVTFRNQASRPYSFYSSLISYEEDQRQG
AEPRKNFVKPNETKTYFWKVQHHMAPTKDEFDCKAWAYFSDVDLEKDVHSGLIGPLLVCHTNTLNP
AHGRQVTVQEFALFFTIFDETKSWYFTENMERNCRAPCNIQMEDPTFKENYRFHAINGYIMDTLPG
LVMAQDQRIRWYLLSMGSNENIHSIHFSGHVFTVRKKEEYKMALYNLYPGVFETVEMLPSKAGIWR
VECLIGEHLHAGMSTLFLVYSNKCQTPLGMASGHIRDFQITASGQYGQWAPKLARLHYSGSINAWS
TKEPFSWIKVDLLAPMIIHGIKTQGARQKFSSLYISQFIIMYSLDGKKWQTYRGNSTGTLMVFFGN
VDSSGIKHNIFNPPIIARYIRLHPTHYSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITASSY
FTNMFATWSPSKARLHLQGRSNAWRPQVNNPKEWLQVDFQKTMKVTGVTTQGVKSLLTSMYVKEFL
ISSSQDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPLLTRYLRIHPQSWVHQIALRMEVLGCE
AQDLY

Figure 56

CS01-FL-NAm12 (SEQ ID NO: 108)

ATGCAGATTGAGCTGTCCACCTGCTTCTTTCTGTGCCTGCTGAGATTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAACTTTCTTGGGACTACATGCAGTCTGACCTGGGAGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAGTCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACACTCTTTGTGGAATTCACTGACCACCTGTTCAACATTGCAAAACCCAGACCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCATCCCACCCTGTGTCTCTGCATGCTGTGGGAGTCTCATACTGGAAATCCTCTGAAGGGGCTGAG
TATGATGACCAGACATCCCAGAGAGAGAAAGAGGATGACAAGGTGTTCCCTGGGAAGTCTCACACC
TATGTGTGGCAAGTCCTCAAGGAGAATGGACCCACTGCATCTGACCCACCCTGCCTGACATACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCACTGCTGGTGTGC
AGGGAAGGATCCCTGGCCAAGGAGAAAACCCAGACACTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGTCTTGGCACTCTGAAACAAAGAACTCCCTGATGCAAGACAGGGATGCTGCC
TCTGCCAGGGCATGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGATCACTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAAGTGCAC
TCCATTTTCCTGGAGGGACACACCTTCCTGGTCAGGAACCACAGACAAGCCTCTCTGGAGATCTCT
CCCATCACCTTCCTCACTGCACAGACACTGCTGATGGACCTTGGACAGTTCCTGCTGTCCTGCCAC
ATCTCTTCCCACCAGCATGATGGCATGGAAGCCTATGTCAAGGTGGACTCATGCCCTGAGGAACCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCAGATTTGATGATGACAACTCTCCATCCTTCATTCAGATCAGGTCTGTGGCAAAGAAA
CACCCCAAGACATGGGTGCACTACATTGCTGCTGAGGAAGAGGACTGGGACTATGCACCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGTCTCAGTACCTCAACAATGGCCCACAAAGAATTGGAAGA
AAGTACAAGAAAGTCAGATTCATGGCCTACACTGATGAAACCTTCAAGACAAGAGAAGCCATTCAG
CATGAGTCTGGCATTCTGGGACCACTCCTGTATGGGGAAGTGGGAGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCAGGAGACTGCCAAAAGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGAGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACAAAGTCTGACCCCAGGTGCCTCACCAGA
TACTACTCCTCTTTTGTGAACATGGAGAGAGACCTGGCATCTGGACTGATTGGACCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGAGGCAACCAGATCATGTCTGACAAGAGAAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGATCATGGTACCTGACTGAGAACATTCAGAGATTCCTGCCCAAC
CCTGCTGGGGTGCAACTGGAAGACCCTGAGTTCCAGGCAAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACTCTCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCACAAACTGACTTCCTTTCTGTCTTCTTCTCTGGATACACCTTCAAGCACAAGATGGTG
TATGAGGACACCCTGACACTCTTCCCATTCTCTGGGGAAACTGTGTTCATGAGCATGGAGAACCCT
GGACTGTGGATTCTGGGATGCCACAACTCTGACTTCAGAAACAGGGGAATGACTGCACTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACTCTTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGAAGCTTCTCTCAGAATCCACCTGTCCTGAAGAGA
CACCAGAGAGAGATCACCAGGACAACCCTCCAGTCTGACCAGGAAGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAGGAGGACTTTGACATCTATGATGAGGACGAGAACCAGTCTCCAAGA
TCATTCCAGAAGAAGACAAGACACTACTTCATTGCTGCTGTGGAAAGACTGTGGGACTATGGCATG
TCTTCCTCTCCCCATGTCCTCAGGAACAGGGCACAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG

(Continued)

Figure 57A

GTCTTCCAGGAGTTCACTGATGGCTCATTCACCCAGCCCCTGTACAGAGGGGAACTGAATGAGCAC
CTGGGACTCCTGGGACCATACATCAGGGCTGAGGTGGAAGACAACATCATGGTGACATTCAGAAAC
CAGGCCTCCAGGCCCTACAGCTTCTACTCTTCCCTCATCAGCTATGAGGAAGACCAGAGACAAGGG
GCTGAGCCAAGAAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCACCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCATACTTCTCTGATGTGGACCTG
GAGAAAGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCACATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
TCATGGTACTTCACTGAGAACATGGAGAGAAACTGCAGAGCACCATGCAACATTCAGATGGAAGAC
CCCACCTTCAAGGAGAACTACAGGTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCACAGGACCAGAGAATCAGATGGTACCTGCTTTCTATGGGATCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGAGAAAGAAGGAGGAATACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAATGCCTCATTGGGGAGCACCTGCATGCTGGCATGTCAACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACACCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCACCCAAACTGGCCAGGCTCCACTACTCTGGCTCCATCAATGCATGGTCA
ACCAAGGAGCCATTCTCTTGGATCAAGGTGGACCTGCTGGCACCCATGATCATTCATGGCATCAAG
ACACAGGGGGCAAGACAGAAATTCTCCTCTCTGTACATCTCACAGTTCATCATCATGTACTCTCTG
GATGGCAAGAAGTGGCAGACATACAGAGGCAACTCCACTGGCACCCTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCTCCCATCATTGCCAGATACATCAGGCTG
CACCCCACCCACTACTCAATCAGATCAACCCTCAGGATGGAACTGATGGGATGTGACCTGAACTCC
TGCTCAATGCCCCTGGGAATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCATCCTCTTAC
TTCACCAACATGTTTGCCACCTGGTCACCATCAAAAGCCAGGCTGCACCTCCAGGGAAGAAGCAAT
GCCTGGAGACCCCAGGTCAACAACCCAAAGGAATGGCTGCAAGTGGACTTCCAGAAGACAATGAAA
GTCACTGGGGTGACAACCCAGGGGGTCAAGTCTCTGCTCACCTCAATGTATGTGAAGGAGTTCCTG
ATCTCTTCCTCACAGGATGGCCACCAGTGGACACTCTTCTTCCAGAATGGCAAAGTCAAGGTGTTC
CAGGGCAACCAGGACTCTTTCACACCTGTGGTGAACTCACTGGACCCCCCCCTCCTGACAAGATAC
CTGAGAATTCACCCCCAGTCTTGGGTCCACCAGATTGCCCTGAGAATGGAAGTCCTGGGATGTGAG
GCACAAGACCTGTACTGA

Figure 57B

CS04-FL-NAm12 (SEQ ID NO: 109)

ATGCAGATTGAGCTGAGCACCTGCTTCTTCCTGTGCCTGCTGAGGTTCTGCTTCTCTGCCACCAGG
AGATACTACCTGGGGGCTGTGGAGCTTTCTTGGGACTACATGCAGTCTGACCTGGGGGAGCTGCCT
GTGGATGCCAGGTTCCCACCCAGAGTGCCCAAATCCTTCCCATTCAACACCTCTGTGGTCTACAAG
AAGACCCTCTTTGTGGAGTTCACTGACCACCTGTTCAACATTGCCAAACCCAGGCCACCCTGGATG
GGACTCCTGGGACCCACCATTCAGGCTGAGGTGTATGACACTGTGGTCGTCACCCTCAAGAACATG
GCCTCCCACCCTGTGAGCCTGCATGCTGTGGGGGTCAGCTACTGGAAGTCCTCTGAGGGGGCTGAG
TATGATGACCAGACCTCCCAGAGGGAGAAGGAGGATGACAAAGTGTTCCCTGGGAAGAGCCACACC
TATGTGTGGCAGGTCCTCAAGGAGAATGGCCCCACTGCCTCTGACCCACCCTGCCTGACCTACTCC
TACCTTTCTCATGTGGACCTGGTCAAGGACCTCAACTCTGGACTGATTGGGGCCCTGCTGGTGTGC
AGGGAGGGCTCCCTGGCCAAAGAGAAGACCCAGACCCTGCACAAGTTCATTCTCCTGTTTGCTGTC
TTTGATGAGGGCAAGAGCTGGCACTCTGAAACCAAGAACTCCCTGATGCAGGACAGGGATGCTGCC
TCTGCCAGGGCCTGGCCCAAGATGCACACTGTGAATGGCTATGTGAACAGGAGCCTGCCTGGACTC
ATTGGCTGCCACAGGAAATCTGTCTACTGGCATGTGATTGGCATGGGGACAACCCCTGAGGTGCAC
TCCATTTTCCTGGAGGGCCACACCTTCCTGGTCAGGAACCACAGACAGGCCAGCCTGGAGATCAGC
CCCATCACCTTCCTCACTGCCCAGACCCTGCTGATGGACCTCGGACAGTTCCTGCTGTCCTGCCAC
ATCAGCTCCCACCAGCATGATGGCATGGAGGCCTATGTCAAGGTGGACAGCTGCCCTGAGGAGCCA
CAGCTCAGGATGAAGAACAATGAGGAGGCTGAGGACTATGATGATGACCTGACTGACTCTGAGATG
GATGTGGTCCGCTTTGATGATGACAACAGCCCATCCTTCATTCAGATCAGGTCTGTGGCCAAGAAA
CACCCCAAGACCTGGGTGCACTACATTGCTGCTGAGGAGGAGGACTGGGACTATGCCCCACTGGTC
CTGGCCCCTGATGACAGGAGCTACAAGAGCCAGTACCTCAACAATGGCCCACAGAGGATTGGACGC
AAGTACAAGAAAGTCAGGTTCATGGCCTACACTGATGAAACCTTCAAGACCAGGGAGGCCATTCAG
CATGAGTCTGGCATCCTGGGCCCACTCCTGTATGGGGAGGTGGGGGACACCCTGCTCATCATCTTC
AAGAACCAGGCCTCCAGGCCCTACAACATCTACCCACATGGCATCACTGATGTCAGGCCCCTGTAC
AGCCGCAGGCTGCCAAAGGGGGTGAAACACCTCAAGGACTTCCCCATTCTGCCTGGGGAGATCTTC
AAGTACAAGTGGACTGTCACTGTGGAGGATGGACCAACCAAATCTGACCCCAGGTGCCTCACCAGA
TACTACTCCAGCTTTGTGAACATGGAGAGGGACCTGGCCTCTGGCCTGATTGGCCCACTGCTCATC
TGCTACAAGGAGTCTGTGGACCAGAGGGGAAACCAGATCATGTCTGACAAGAGGAATGTGATTCTG
TTCTCTGTCTTTGATGAGAACAGGAGCTGGTACCTGACTGAGAACATTCAGCGCTTCCTGCCCAAC
CCTGCTGGGGTGCAGCTGGAGGACCCTGAGTTCCAGGCCAGCAACATCATGCACTCCATCAATGGC
TATGTGTTTGACAGCCTCCAGCTTTCTGTCTGCCTGCATGAGGTGGCCTACTGGTACATTCTTTCT
ATTGGGGCCCAGACTGACTTCCTTTCTGTCTTCTTCTCTGGCTACACCTTCAAACACAAGATGGTG
TATGAGGACACCCTGACCCTCTTCCCATTCTCTGGGGAGACTGTGTTCATGAGCATGGAGAACCCT
GGCCTGTGGATTCTGGGATGCCACAACTCTGACTTCCGCAACAGGGGCATGACTGCCCTGCTCAAA
GTCTCCTCCTGTGACAAGAACACTGGGGACTACTATGAGGACAGCTATGAGGACATCTCTGCCTAC
CTGCTCAGCAAGAACAATGCCATTGAGCCCAGGAGCTTCAGCCAGAATCCACCTGTCCTGAAACGC
CACCAGAGGGAGATCACCAGGACCACCCTCCAGTCTGACCAGGAGGAGATTGACTATGATGACACC
ATTTCTGTGGAGATGAAGAAAGAGGACTTTGACATCTATGACGAGGACGAGAACCAGAGCCCAAGG
AGCTTCCAGAAGAAGACCAGGCACTACTTCATTGCTGCTGTGGAGCGCCTGTGGGACTATGGCATG
AGCTCCAGCCCCCATGTCCTCAGGAACAGGGCCCAGTCTGGCTCTGTGCCACAGTTCAAGAAAGTG
GTCTTCCAAGAGTTCACTGATGGCAGCTTCACCCAGCCCCTGTACAGAGGGGAGCTGAATGAGCAC
CTGGGACTCCTGGGCCCATACATCAGGGCTGAGGTGGAGGACAACATCATGGTGACCTTCCGCAAC

(Continued)

Figure 58A

CAGGCCTCCAGGCCCTACAGCTTCTACAGCTCCCTCATCAGCTATGAGGAGGACCAGAGGCAGGGG
GCTGAGCCACGCAAGAACTTTGTGAAACCCAATGAAACCAAGACCTACTTCTGGAAAGTCCAGCAC
CACATGGCCCCCACCAAGGATGAGTTTGACTGCAAGGCCTGGGCCTACTTCTCTGATGTGGACCTG
GAGAAGGATGTGCACTCTGGCCTGATTGGCCCACTCCTGGTCTGCCACACCAACACCCTGAACCCT
GCCCATGGAAGGCAAGTGACTGTGCAGGAGTTTGCCCTCTTCTTCACCATCTTTGATGAAACCAAG
AGCTGGTACTTCACTGAGAACATGGAGCGCAACTGCAGGGCCCCATGCAACATTCAGATGGAGGAC
CCCACCTTCAAAGAGAACTACCGCTTCCATGCCATCAATGGCTACATCATGGACACCCTGCCTGGG
CTTGTCATGGCCCAGGACCAGAGGATCAGGTGGTACCTGCTTTCTATGGGCTCCAATGAGAACATT
CACTCCATCCACTTCTCTGGGCATGTCTTCACTGTGCGCAAGAAGGAGGAGTACAAGATGGCCCTG
TACAACCTCTACCCTGGGGTCTTTGAGACTGTGGAGATGCTGCCCTCCAAAGCTGGCATCTGGAGG
GTGGAGTGCCTCATTGGGGAGCACCTGCATGCTGGCATGAGCACCCTGTTCCTGGTCTACAGCAAC
AAGTGCCAGACCCCCCTGGGAATGGCCTCTGGCCACATCAGGGACTTCCAGATCACTGCCTCTGGC
CAGTATGGCCAGTGGGCCCCCAAGCTGGCCAGGCTCCACTACTCTGGATCCATCAATGCCTGGAGC
ACCAAGGAGCCATTCAGCTGGATCAAAGTGGACCTGCTGGCCCCCATGATCATCCATGGCATCAAG
ACCCAGGGGGCCAGGCAGAAGTTCTCCAGCCTGTACATCAGCCAGTTCATCATCATGTACAGCCTG
GATGGCAAGAAATGGCAGACCTACAGAGGCAACTCCACTGGAACACTCATGGTCTTCTTTGGCAAT
GTGGACAGCTCTGGCATCAAGCACAACATCTTCAACCCCCCAATCATCGCCAGATACATCAGGCTG
CACCCCACCCACTACAGCATCCGCAGCACCCTCAGGATGGAGCTGATGGGCTGTGACCTGAACTCC
TGCAGCATGCCCCTGGGCATGGAGAGCAAGGCCATTTCTGATGCCCAGATCACTGCCTCCAGCTAC
TTCACCAACATGTTTGCCACCTGGAGCCCAAGCAAGGCCAGGCTGCACCTCCAGGGAAGGAGCAAT
GCCTGGAGGCCCCAGGTCAACAACCCAAAGGAGTGGCTGCAGGTGGACTTCCAGAAGACCATGAAG
GTCACTGGGGTGACCACCCAGGGGGTCAAGAGCCTGCTCACCAGCATGTATGTGAAGGAGTTCCTG
ATCAGCTCCAGCCAGGATGGCCACCAGTGGACCCTCTTCTTCCAGAATGGCAAGGTCAAGGTGTTC
CAGGGCAACCAGGACAGCTTCACCCCTGTGGTGAACAGCCTGGACCCCCCCCTCCTGACCAGATAC
CTGAGGATTCACCCCCAGAGCTGGGTCCACCAGATTGCCCTGAGGATGGAGGTCCTGGGATGTGAG
GCCCAGGACCTGTACTGA

Figure 58B

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2016/061684

A. CLASSIFICATION OF SUBJECT MATTER
INV. C07K14/755 A61K48/00 C12N15/86
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07K A61K C12N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
EPO-Internal, Sequence Search

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2015/071883 A1 (COLOSI PETER CAMERON [US]) 12 March 2015 (2015-03-12) abstract; claims 1-14; sequence 45 paragraph [0060] paragraph [0130] -----	8,11,12, 14,18, 19,32, 43,51, 66-74
X	DATABASE Geneseq [Online] 5 December 2013 (2013-12-05), "Human diseases associated protein encoding optimized ORF, SEQ ID 1811.", XP002766200, retrieved from EBI accession no. GSN:BAW43417 Database accession no. BAW43417 sequence -/--	18,19,51

X Further documents are listed in the continuation of Box C.

X See patent family annex.

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"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

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Date of the actual completion of the international search	Date of mailing of the international search report
3 February 2017	15/02/2017

Name and mailing address of the ISA/	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Gurdjian, Didier

4

Form PCT/ISA/210 (second sheet) (April 2005)

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	& WO 2013/151666 A2 (MODERNA THERAPEUTICS [US]) 10 October 2013 (2013-10-10) abstract; claims 1,2; figure 1; sequence 1811	

A	US 2013/017997 A1 (SCHELLENBERGER VOLKER [US] ET AL) 17 January 2013 (2013-01-17) abstract; claims 1-26; figures 1-9; sequence 939	27

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/US2016/061684

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