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(54) **TAT-UTROPHIN AS A PROTEIN THERAPY FOR DYSTROPHINOPATHIES**

(75) Inventors: **James M. Ervasti**, Shoreview, MN (US); **Kevin J. Sonnemann**, Minneapolis, MN (US)

(73) Assignee: **Wisconsin Alumni Research Foundation**, Madison, WI (US)

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(51) **Int. Cl.**
C07K 19/00 (2006.01)
C12N 15/62 (2006.01)

(52) **U.S. Cl.** **435/69.7**; 514/2; 530/350; 536/23.4

(58) **Field of Classification Search** None
See application file for complete search history.

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Primary Examiner—John D Ulm

(74) *Attorney, Agent, or Firm*—Joseph T. Leone, Esq.; DeWitt Ross & Stevens S.C.

(57) **ABSTRACT**

Disclosed is a fusion protein including a full-length TAT-utrophin or an anti-dystrophinopathic fragment thereof, a method of treating dystrophinopathies (including Duchenne muscular dystrophy) using the fusion protein, a pharmaceutical composition for treating dystrophinopathies in mammals comprising the fusion protein, and nucleic acid constructs for expressing the fusion protein.

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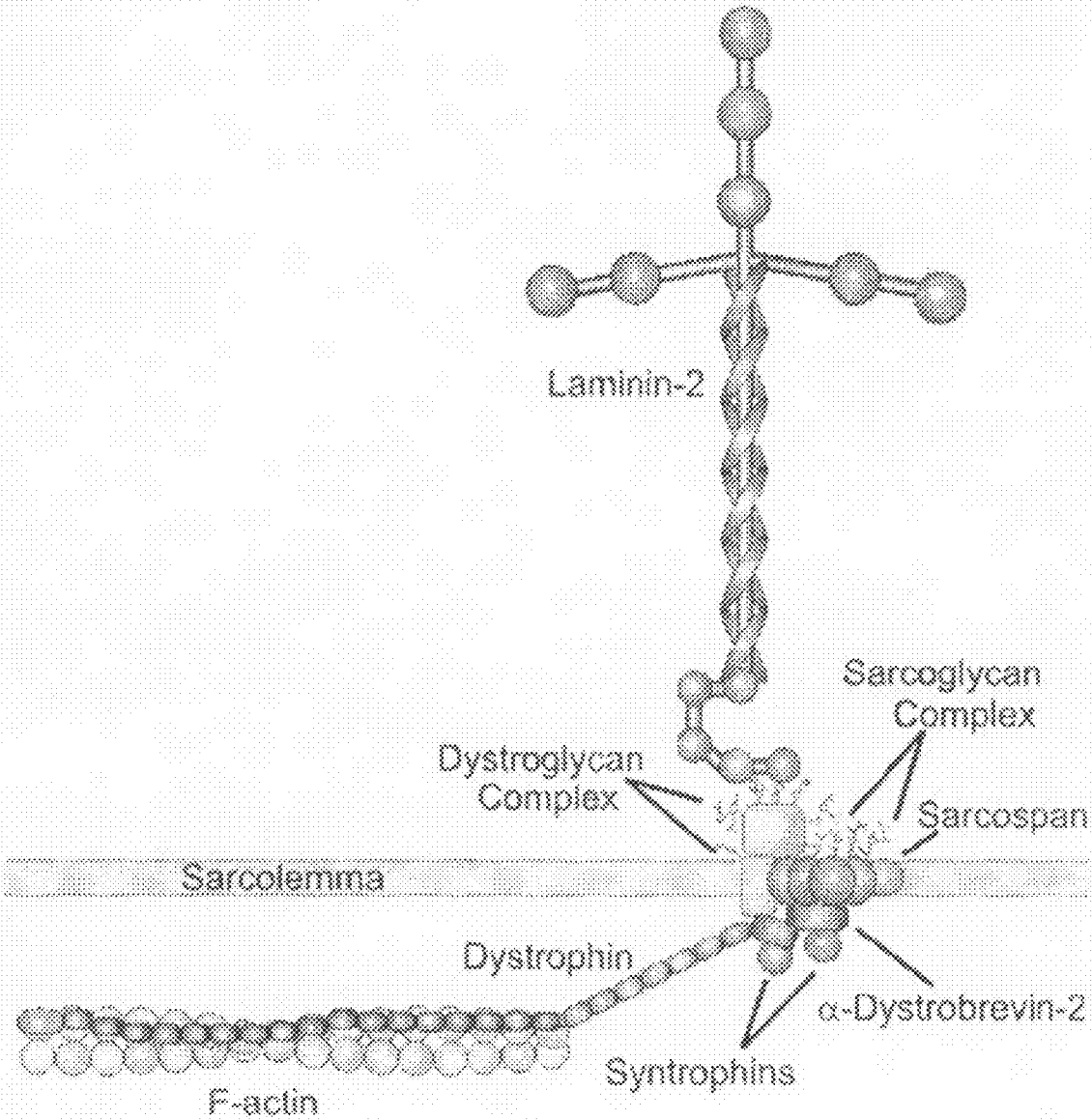


FIG. 1

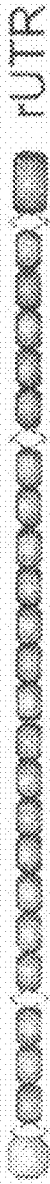
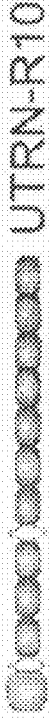
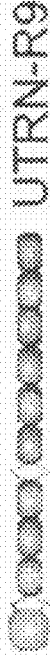
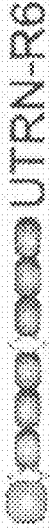
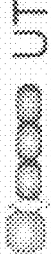
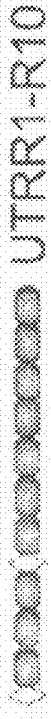
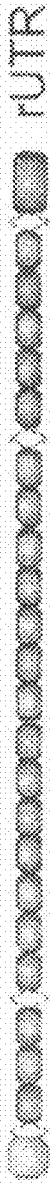
K_d (μ M)	B_{max}	Protect?	Protein
0.2	1:14	Yes	 UTRN-R10
0.6	1:12	Yes	 UTRN-R9
1.4	1:10	Yes	 UTRN-R6
1.5	1:5	Partial	 UTRN-R3
2	1:3	No	 UTR261
16	1:1	No	 UTRR1-R10
No Binding Activity			 rUTR

FIG. 2

FIG. 3A

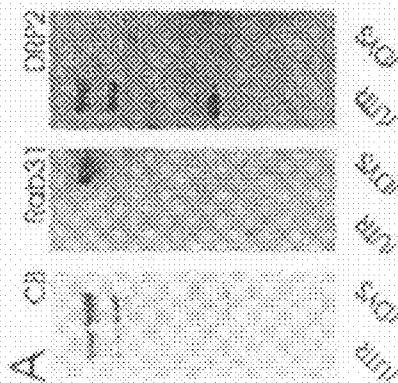


FIG. 3D

K _d (μM)	B _{max}	Protect?	Protein
0.6	1:24	Yes	(GPC) DGC
0.4	1:24	Yes	rDYS
0.2	1:14	Yes	rUTR

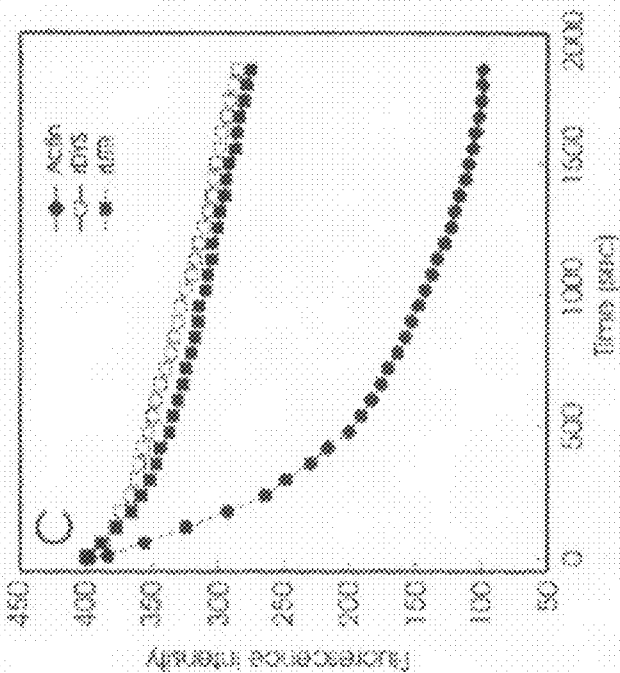


FIG. 3B

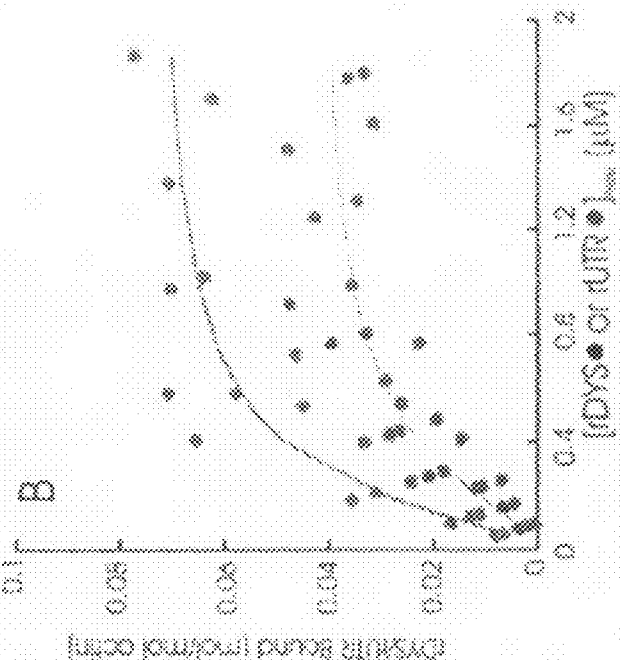


FIG. 3C

Line/Protein	% Total Protein	% Dys _{wt}
WT/Dystrophin	0.02	100
WT/Utrophin	0.0006	3
<i>mdx</i> /Utrophin	0.0013	7
Fiona/Utrophin	0.014	70

FIG. 4

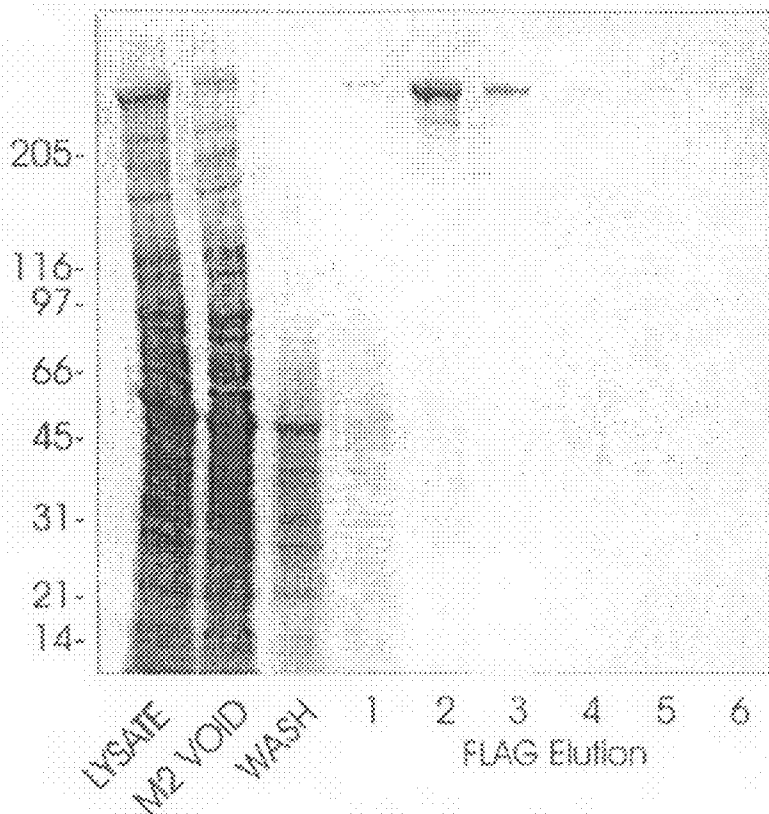


FIG. 5

FIG. 6A

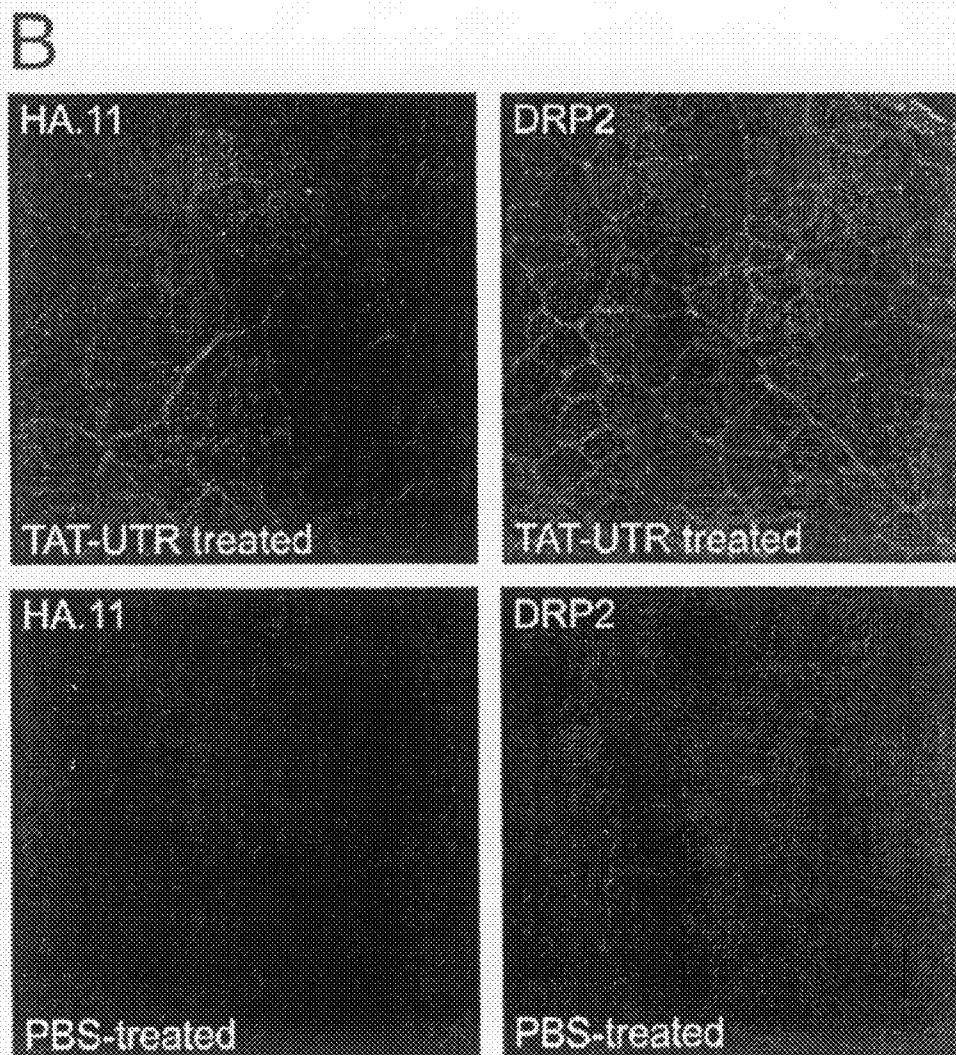
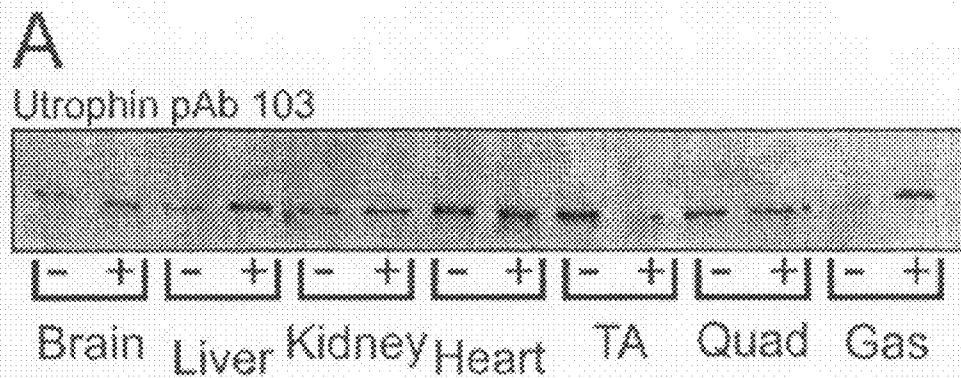


FIG. 6B

FIG. 7A

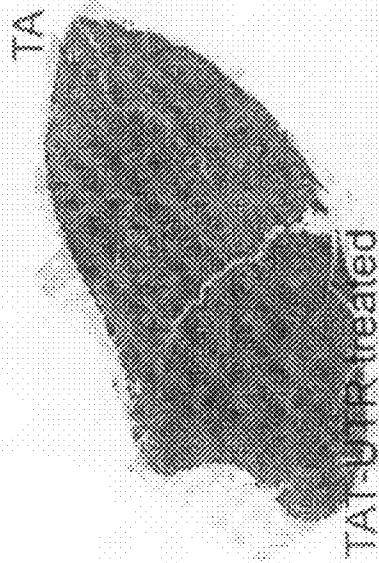


FIG. 7B

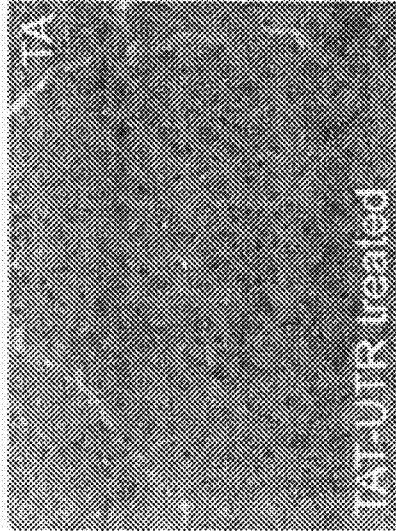


FIG. 7C

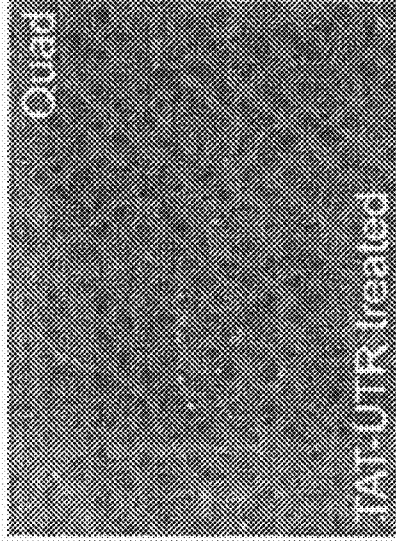


FIG. 7D

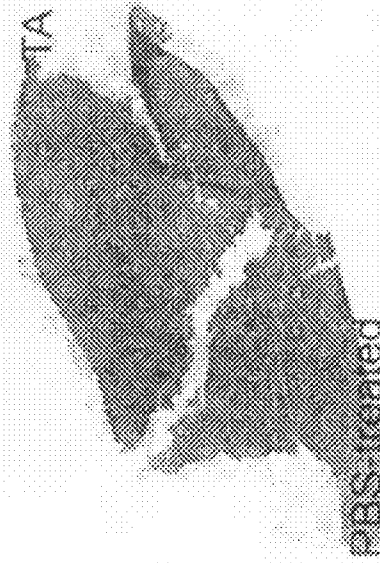


FIG. 7E

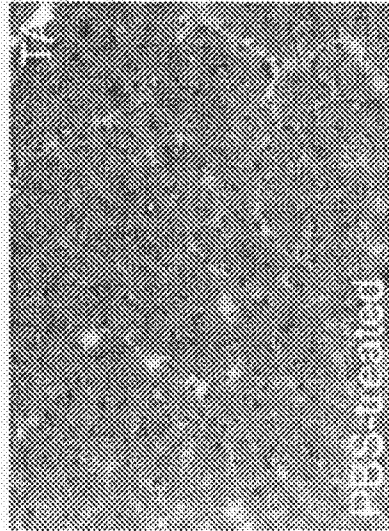


FIG. 7F

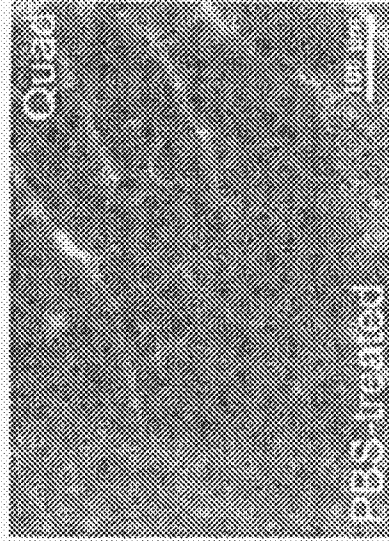


FIG. 7D

FIG. 7E

FIG. 7F

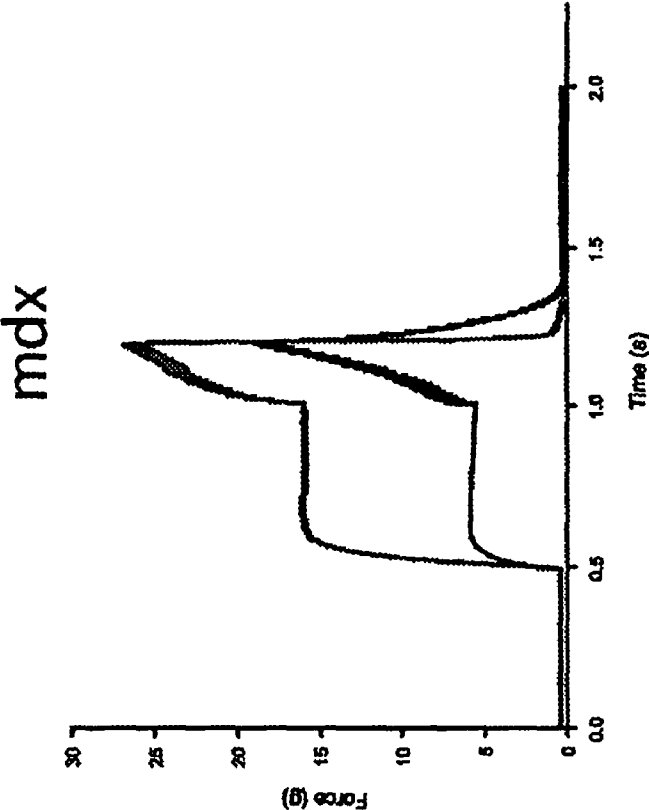


FIG. 8B

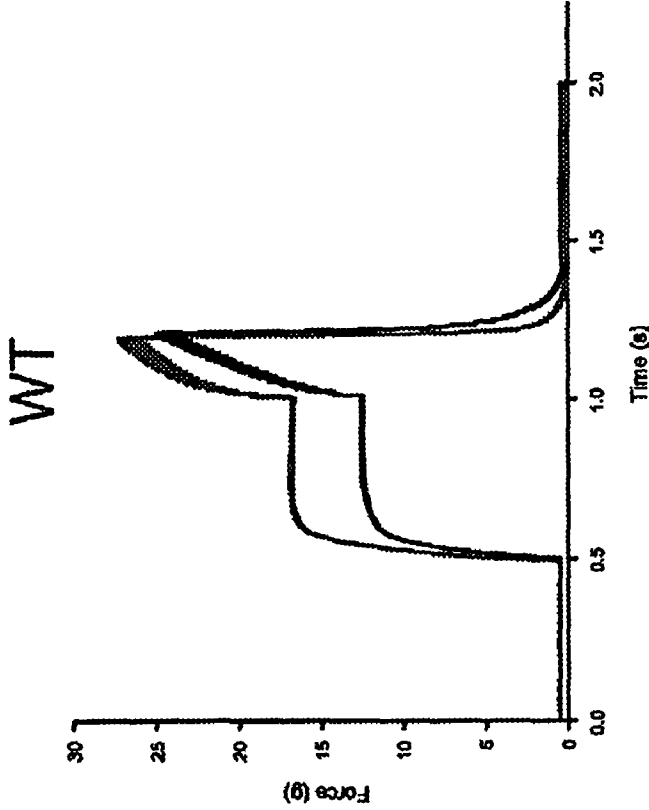


FIG. 8A

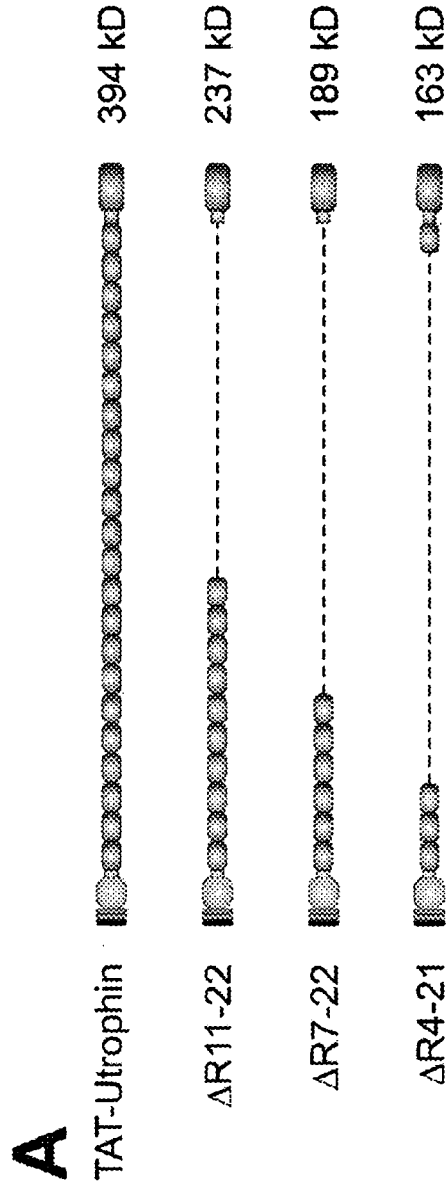


FIG. 9A

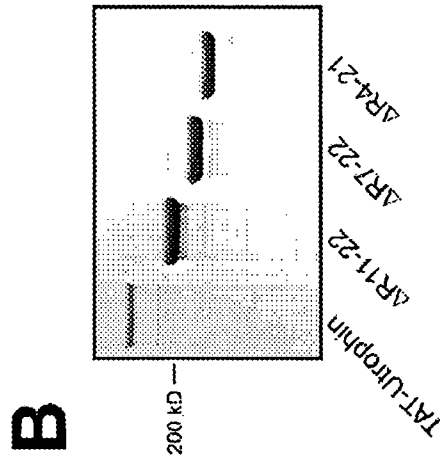


FIG. 9B

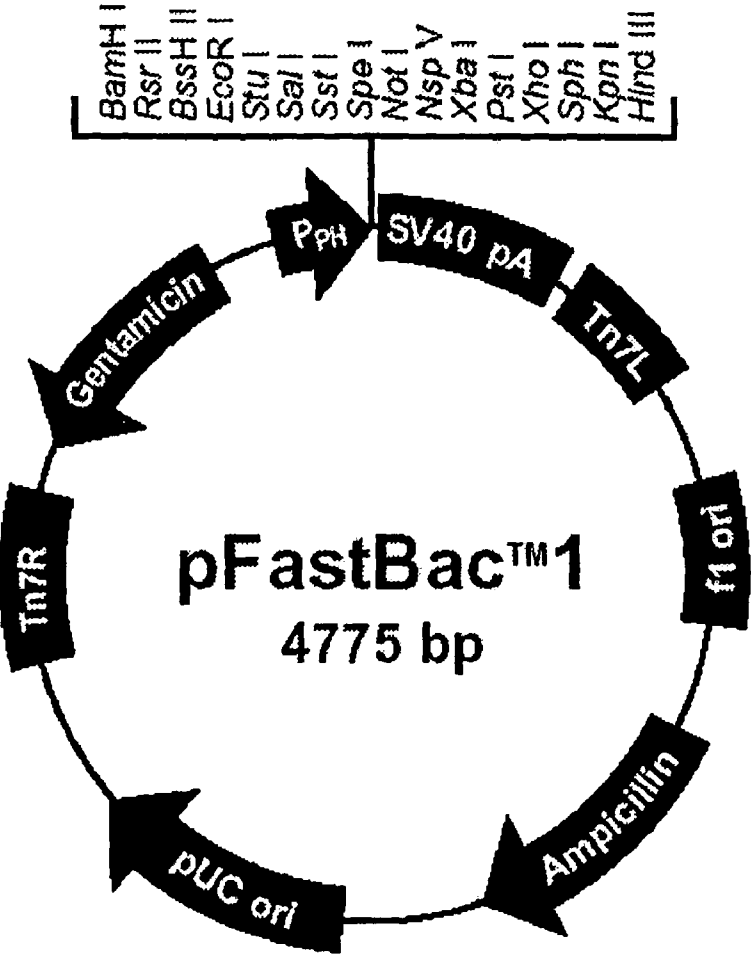


FIG. 10

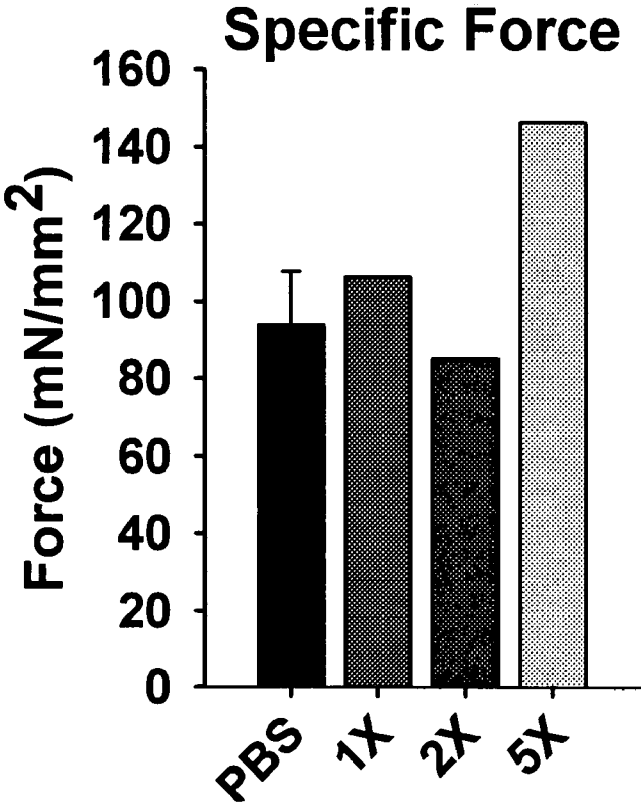


FIG. 11

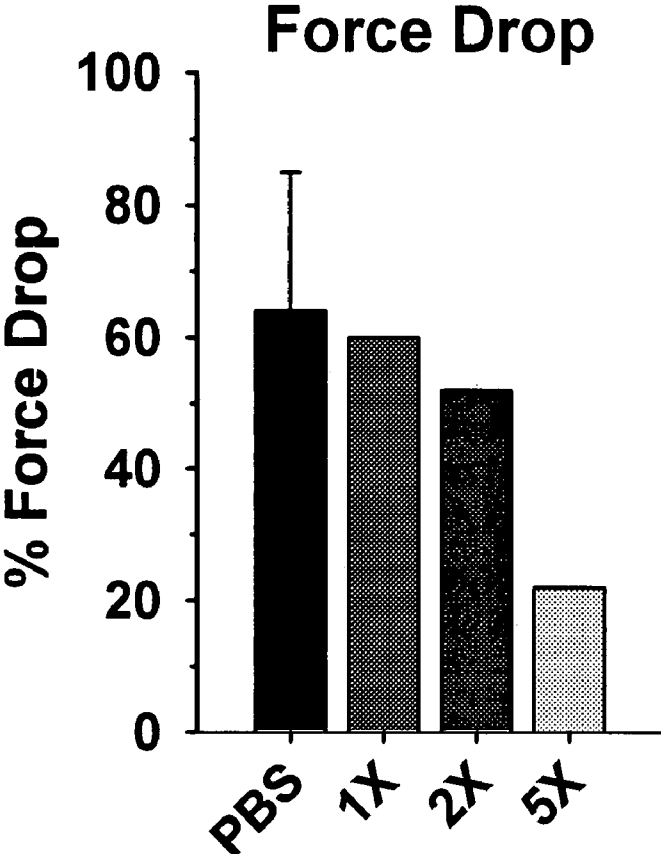
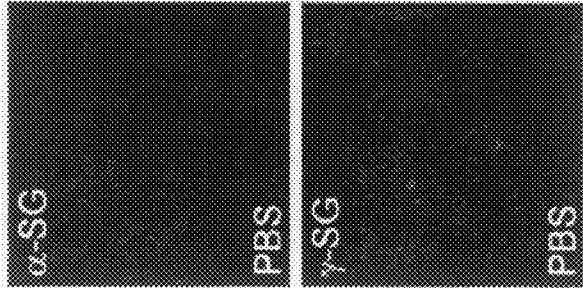
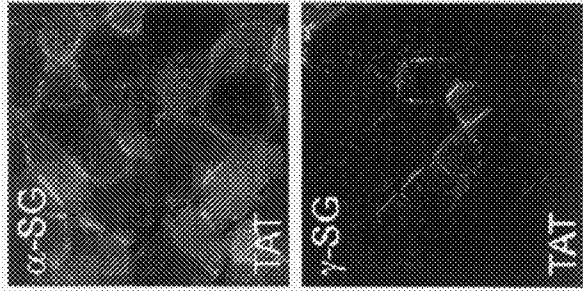


FIG. 12

B



A

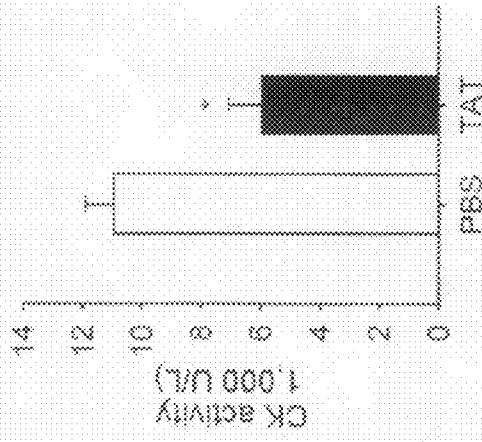
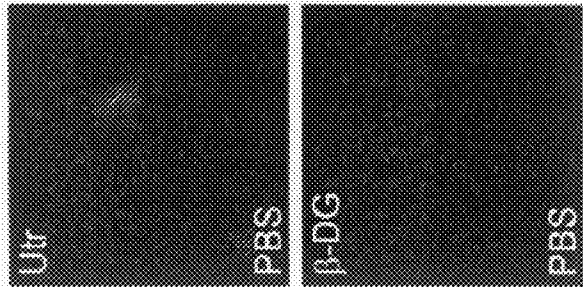
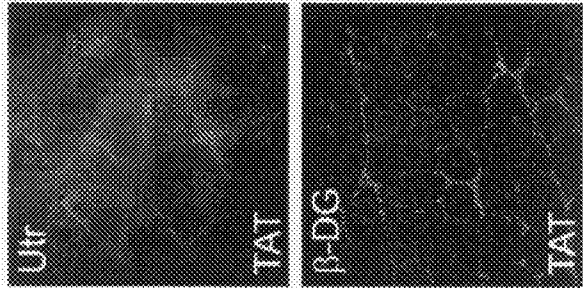


FIG. 13A

FIG. 13B

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TAT-UTROPHIN AS A PROTEIN THERAPY FOR DYSTROPHINOPATHIES

CROSS-REFERENCE TO RELATED APPLICATIONS

Priority is hereby claimed to provisional application Ser. No. 60/868,119, filed Dec. 1, 2006, which is incorporated herein by reference.

FEDERAL FUNDING STATEMENT

This invention was made with United States government support awarded by the following agency: NIH Grant AR042423. The United States government has certain rights in this invention.

FIELD OF THE INVENTION

The invention is directed to a fusion protein comprising a full-length TAT-utrophin or an anti-dystrophinopathic fragment thereof, a method of treating dystrophinopathies (including Duchenne muscular dystrophy) using the fusion protein and a pharmaceutical composition for treating dystrophinopathies in mammals comprising the fusion protein.

BACKGROUND

Duchenne muscular dystrophy (DMD) is the most prevalent and severe form of human muscular dystrophy. DMD occurs with an incidence of 1 in 4000 male births. Onset of DMD is typically between 3 and 6 years of age with skeletal muscle weakness preferentially affecting the large proximal muscle groups. The disease is invariably progressive, leading to loss of ambulation by 11 to 13 years, and death typically in the 20's. Significant laboratory findings include grossly elevated serum CK-MM levels. Skeletal muscle biopsy samples reveal a dystrophic pattern of muscle degeneration and regeneration with fiber-size variation, increased central nuclei, and progressive interstitial fibrosis.

Becker muscular dystrophy (BMD) was long considered to be a potentially allelic disorder because of its clinical similarities to DMD and a common pattern of X-linked inheritance. The shared genetic basis for DMD and BMD was confirmed after the identification of the protein dystrophin; both DMD and BMD patients were shown to have dystrophin gene mutations. Typically, patients with DMD lack any detectable dystrophin expression in their skeletal muscles, and this is correlated with deletion mutations that disrupt the translational reading frame or point mutations that create stop codons. In contrast, muscle from patients with BMD contains mutated dystrophins having an altered size and/or reduced abundance secondary to deletion mutations that maintain the reading frame.

While clinical descriptions of DMD date back to the 1850's, over 100 years passed before evidence suggested that the muscle cell plasma membrane, or sarcolemma, is compromised in DMD muscle. The molecular basis for DMD and its associated sarcolemmal instability became more clear with landmark studies published in the mid-to-late 1980's which identified the gene encoding dystrophin as being defective in DMD (O'Brien and Kunkel, 2001). The DMD locus spans over 2.5 million bases distinguishing it as the largest gene in the human genome. The array of transcripts expressed from the DMD gene is complex due to the presence of multiple promoters and alternative splicing. The largest

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transcripts encode dystrophin, a four-domain protein with a predicted molecular weight of 427,000. Dystrophin is the predominant DMD transcript expressed in striated muscle. DMD gene mutations, deletions, or duplications most frequently result in a loss of dystrophin expression in muscle of patients afflicted with DMD. Based on its localization to the cytoplasmic face of the sarcolemma, and its sequence similarity with domains/motifs common to proteins of the actin-based cytoskeleton, dystrophin was hypothesized early on to play a mechanical role in anchoring the sarcolemma to the underlying cytoskeleton. It has also been hypothesized that dystrophin plays a role in protecting the sarcolemma against stress imposed during muscle contraction or stretch.

Biochemical studies aimed at confirming the hypothesized structure and function of dystrophin revealed its tight association with a multi-subunit complex, the so-named dystrophin-glycoprotein complex. See FIG. 1, which is a schematic representation showing the sarcolemma and the interaction of dystrophin with the other elements of the dystrophin-glycoprotein complex. Through its cysteine-rich and C-terminal domains, dystrophin in striated muscle interacts with the integral membrane dystroglycan sub-complex and the sarcoglycan/sarcospan sub-complex, as well as the subsarcolemmal dystrobrevins and syntrophins (Cohn and Campbell, 2000; Blake et al., 2002). The N-terminal domain and a portion of middle rod domain of dystrophin act in concert to effect an extensive lateral association with actin filaments in vitro (Rybakova et al., 1996) and in vivo (Rybakova et al., 2000; Warner et al., 2002; Rybakova and Ervasti, 1997; Amann et al., 1998; Amann et al., 1999).

Utrophin is a widely expressed autosomal gene product with high sequence similarity to dystrophin (Tinsley et al., 1992). Utrophin is distributed throughout the sarcolemma in fetal and regenerating muscle, but is down-regulated in normal adult muscle and is restricted to the myotendinous and neuromuscular junctions (Blake et al., 1996). Because utrophin and dystrophin bind the same complement of proteins (Matsumura et al., 1992; Kramarcy et al., 1994; Winder et al., 1995), it was hypothesized that utrophin may be capable of compensating for dystrophin deficiency. Indeed, continued utrophin expression in adult mdx mice partially attenuates the phenotype associated with dystrophin deficiency. In short, mice lacking both dystrophin and utrophin exhibit a more severe phenotype similar to that seen in human DMD patients (Deconinck et al., 1997a; Grady et al., 1997). Moreover, transgenic overexpression of full-length utrophin completely rescued the dystrophic phenotype in mdx mice (Tinsley et al., 1998).

Methods to express and purify full-length utrophin using a baculovirus system has been demonstrated (Rybakova et al., 2002 and 2006). It has also been shown that purified recombinant utrophin is a soluble, rod-shaped monomer with the expected molecular weight of 400,000 Da. Recombinant utrophin-bound actin filaments display an affinity ($K_d=0.2$ μ M) similar to that measured for purified dystrophin-glycoprotein complex (Rybakova et al., 2002). Recombinant utrophin-bound F-actin displays a stoichiometry of 1 utrophin per 14 actin monomers, which implies a more extensive lateral association with actin filaments than anticipated from studies with isolated fragments, but a less extensive lateral association than the 1 per 24 stoichiometry measured for purified recombinant dystrophin (Rybakova et al., 2006). Like the dystrophin-glycoprotein complex, recombinant utrophin protected actin filaments from forced depolymerization in a concentration-dependent manner that saturated at molar ratios equal to or greater than 1 utrophin per 14 actin monomers. Also different from purified dystrophin-glycoprotein com-

plex, the binding of recombinant utrophin to actin filaments was completely insensitive to increasing ionic strength up to 0.8 M. These results (Rybakova et al., 2002) (Rybakova et al., 2006) indicate that dystrophin and utrophin both bind laterally alongside actin filaments through contributions by the spectrin-like repeats of the rod domain, but that the rod domain epitopes involved differ between the two proteins. Utrophin appears to bind laterally along actin filaments through a contribution of the first 10 acidic spectrin-like repeats (Rybakova et al., 2002) rather than a cluster of basic repeats as employed by dystrophin (Rybakova et al., 1996; Amann et al., 1998); (Rybakova et al., 2006).

Most viruses, including the human immunodeficiency viruses (HIV), encode proteins for regulating genome transcription. In HIV, the *tat* gene plays a role in driving the transcription of the HIV genetic code. The *tat* gene encodes a small nuclear protein of from 86 to 101 amino acids, depending upon the viral strain. Both the *tat* gene and its encoded protein, TAT, are known. The protein itself is designated TAT, for "transactivator protein." The typical HIV-1 laboratory strains HXB2 and NL4-3 express an 86 amino acid-long TAT protein, while other HIV strains express a 101 amino acid-long TAT protein. See, for example, Kuppaswamy et al., 1989.

Despite all that is now known, and despite continuing efforts by many laboratories around the world (Gregorevic and Chamberlain, 2003), there is presently no cure or effective treatment to alleviate the devastating progression of DMD.

SUMMARY OF THE INVENTION

The primary object of the present invention is a method of treating dystrophinopathies in mammals, including humans. The method comprises administering an anti-dystrophinopathic-effective amount of a chimeric protein (i.e., a fusion protein) encoding TAT-utrophin. The chimeric protein is administered in an amount effective to transduce skeletal muscle cells and thereby to correct the pathologies associated with dystrophin deficiency. The chimeric protein may comprise a full-length TAT protein (e.g., 86 amino acids long or 101 amino acids long) or a fragment thereof, such as the HIV-1 TAT protein transduction sequence (see SEQ. ID. NO: 5). Similarly, the chimeric protein may comprise a full-length utrophin protein or an anti-dystrophinopathic-effective fragment thereof. (For purposes of brevity, both full-length and fragmented versions of the chimeric protein will be referred to herein as the "TAT-utrophin chimeric (or fusion) protein.") Utrophin fragments can be evaluated for their anti-dystrophinopathic effects by transgenically over-expressing the putative anti-dystrophinopathic fragment in mdx mice in the same fashion as Tinsley et al., 1998 and observing whether the dystrophic phenotype in the mdx mice is ameliorated or eliminated. Alternatively, the TAT-utrophin chimeras can be tested on mdx mice as described herein below for their anti-dystrophinopathic efficacy.

The invention is also directed to a baculovirus construct that drives the expression of the TAT-utrophin chimeric protein, the chimeric protein encoding TAT-utrophin itself, as well as a pharmaceutical composition for treating dystrophinopathies that comprises an anti-dystrophinopathic amount of the TAT-utrophin chimeric protein in combination with a pharmaceutically suitable carrier.

Thus, one version of the invention is directed to a fusion protein comprising a first protein region which is effective to transduce the fusion protein into mammalian muscle cells. The first protein region preferably comprises an HIV TAT

protein or a transduction-effective fragment thereof. The first protein region is operationally linked to a second protein region comprising a full-length utrophin protein or an anti-dystrophinopathic fragment thereof. Also included within the invention are pharmaceutically suitable salts of the fusion proteins.

Another version of the invention is directed to a nucleic acid construct (vector) that drives the expression of the above-noted fusion protein when the construct is transformed into a suitable host or disposed in a suitable cell-free expression system. Many cell-free expression systems are commercially available. For example, Promega (Madison, Wis.) markets a suitable cell-free expression system under the registered trademark "TNT." Promega's "TNT"®-brand systems are single-tube, coupled transcription/translation reactions for eukaryotic cell-free protein expression. To use these systems, 0.2 to 2.0 µg of circular plasmid DNA containing a T7, T3 or SP6 promoter, or a PCR-generated fragment containing a T7 promoter, is added to an aliquot of the "TNT"®-brand Quick Master Mix and incubated in a 50 µl reaction volume for 60 minutes at 30° C. Other cell-free systems are offered commercially by Qiagen (Valencia, Calif.), Invitrogen (Carlsbad, Calif.), and others.

The transformed host itself is also encompassed within the scope of the present invention.

Another version of the invention is directed to a pharmaceutical composition for treating dystrophinopathies in mammals, including humans. The pharmaceutical composition comprises a fusion protein as noted previously, or a pharmaceutically suitable salt thereof, in an anti-dystrophinopathic amount, in combination with a pharmaceutically suitable carrier.

Yet another version of the invention is directed to a method of treating dystrophinopathies, including DMD, in mammals. The method comprises administering to a mammalian subject in need thereof an anti-dystrophinopathic amount of an isolated fusion protein or a pharmaceutically suitable salt thereof, wherein the fusion protein comprises a first region which is effective to transduce the fusion protein into mammalian muscle cells. The first region is operationally linked to a second region comprising a full-length utrophin protein or an anti-dystrophinopathic fragment thereof.

As described herein, the present inventors have expressed full-length utrophin in a baculovirus system and have shown that the expressed protein can be purified as a highly soluble monomer. The monomer has actin-binding activities similar to those measured for recombinant dystrophin and purified dystrophin glycoprotein complex. The invention also encompasses a baculovirus expression construct (i.e. a "bacmid") that encodes full-length mouse utrophin fused with an amino-terminal peptide corresponding to the protein transduction domain of the HIV TAT protein. TAT-utrophin expresses to high levels in insect cells, is fully soluble, and can be rapidly purified by affinity chromatography.

Transduction of TAT-utrophin into the skeletal muscle of dystrophin-deficient mdx mice corrects the dystrophic phenotype displayed by the mdx mice.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a schematic diagram of the dystrophin-glycoprotein complex.

FIG. 2 is a graph depicting the relative lengths and actin-binding properties (K_d and B_{max}) of serially-deleted constructs of utrophin.

FIGS. 3A, 3B, 3C, and 3D compare in various terms the actin-binding properties of recombinant dystrophin versus

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the actin-binding properties of utrophin. FIG. 3A shows parallel gels containing (moving from left to right) a Coomassie blue-stained gel loaded with recombinant utrophin (rUTR) and recombinant dystrophin (rDYS), western blots stained with rabbit 31 antibodies (Rab31) specific for dystrophin, and DRP2 antibodies specific against utrophin. FIG. 3B is a graph depicting F-actin co-sedimentation data for rDYS (lower trace) and rUTR (upper trace); the X-axis plots concentration in μM , the Y-axis plots bound rDYS and rUTR (mol/mol actin). FIG. 3C is a graph depicting the effect of dystrophin/utrophin on depolymerization of actin filaments containing PRODAN-labeled monomers (\bullet -actin, \square -rDYS, \blacksquare -rUTR). FIG. 3D is a graph depicting the relative lengths and actin-binding properties (K_d and B_{max}) of the serially-deleted constructs.

FIG. 4 is a graph depicting the quantitation of dystrophin and utrophin levels in skeletal muscle. The abundance of dystrophin and utrophin was measured in wild-type, mdx, and "Fiona" transgenic mdx mice overexpressing full-length utrophin by quantitative western blot analysis using recombinant dystrophin and utrophin as standards. Values are expressed as percent of total muscle protein and percent of dystrophin abundance in wild-type muscle.

FIG. 5 is a gel showing the expression and purification of TAT-utrophin in the baculovirus system. See the examples for lane assignments.

FIGS. 6A and 6B depict uptake and membrane localization of TAT-utrophin in mdx muscle. FIG. 6A is a western blot that shows increased utrophin immuno-reactivity in several tissues of an mdx mouse after 6 intraperitoneal injections of TAT-UTR (+) compared to PBS-injected controls (-). FIG. 6B depicts the results of immunofluorescence analysis, which shows increased sarcolemmal HA-tag and DRP2 immunoreactivity in the TAT-UTR-treated animal (upper-left and upper-right panels, respectively) as compared to the sarcolemmal HA-tag and DRP2 immunoreactivity in PBS-injected controls (lower-left and lower-right panels, respectively).

FIGS. 7A, 7B, 7C, 7D, 7E, and 7F are a series of photographs showing greatly reduced histopathology in TAT-utrophin-treated mdx muscle versus controls. FIGS. 7A, 7B, and 7C depict TAT-utrophin-treated mdx muscle (TA and quadriceps), while FIGS. 7D, 7E, and 7F depict PBS-treated mdx muscle (TA and quadriceps). Haematoxylin and eosin stained sections revealed decreased numbers of centrally nucleated fibers and less fibrosis in TAT-UTR treated compared to PBS-injected mdx muscle.

FIGS. 8A and 8B are graphs depicting the increased susceptibility of mdx muscles to eccentric contraction. Shown are tracings of maximal force versus time obtained during the first (upper trace) and fifth (lower trace) eccentric contraction imposed on isolated EDL muscle from wild-type (WT) and dystrophin-deficient mdx mice. Note the greater force drop in mdx muscle versus WT muscle as previously reported by Petrof et al. (1993) and Moens et al. (1993).

FIG. 9A is a schematic representation of mini- and micro-TAT-utrophin constructs according to the present invention. FIG. 9B is a Coomassie Blue-stained protein gel of the truncated TAT-utrophin constructs depicted schematically in FIG. 9A.

FIG. 10 is a map of the pFastBac1-brand plasmid, available commercially from Invitrogen.

FIG. 11 is a histogram depicting the dose-dependent ability of TAT-utrophin to increase the specific force of muscle tissue in mdx mice treated with the TAT-utrophin.

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FIG. 12 is a histogram depicting the dose-dependent ability of TAT-utrophin to decrease contraction-induced injury of muscle tissue in mdx mice treated with the TAT-utrophin.

FIG. 13A depicts the results of immunofluorescence analysis on $10\ \mu\text{m}$ thick cryosections from PBS- or TAT-utrophin-injected quadriceps from mdx mice. Primary antibodies to utrophin (NCL-DRP2; Utr), β -dystroglycan (NCL-b-DG; β -DG), α -sarcoglycan (NCL-a-SARC; α -SG), and γ -sarcoglycan (NCL-g-SARC; γ -SG) demonstrated peripherally localized dystrophin complex members in the TAT-utrophin-treated mice. FIG. 3B is a histogram depicting serum activity levels of the muscle enzyme creatine kinase from PBS- or TAT-utrophin-injected quadriceps from mdx mice. Creatine kinase levels were reduced 50% in 38 day-old TAT-utrophin-treated mice as compared to PBS-injected controls. (*) denotes $p < 0.05$.

DETAILED DESCRIPTION OF THE INVENTION

20 Definitions and Abbreviations:

The following abbreviations and definitions are used throughout the specification and claims. Any terms not explicitly defined herein are to be given their accepted meanings in the fields of molecular biology, physiology, and/or biochemistry.

Affinity tag: Any moiety (typically a small oligopeptide) that can be affixed to a protein (by any means) which allows the resulting fused entity to be isolated by affinity chromatography.

30 Anti-dystrophinopathic fragment: a fragment of a full-length utrophin protein that functions to ameliorate dystrophinopathic symptoms when administered as part of the fusion protein described herein. Explicitly included within this definition are the utrophin fragments shown in SEQ. ID. NOS: 10-25 in the attached Sequence List. (The "delta" nomenclature used in the Sequence List reflects the number of deleted repeats. Thus, the construct "murine TAT-UTR delta 4-21" encodes a murine TAT-utrophin fusion protein deleted for repeats 4-21.) It is preferred that the fragment be no more than 75% of the mass of the full-length utrophin protein, more preferred that the fragment be no more than 50% of the mass of the full-length utrophin protein, and still more preferred that the fragment be no more than 25% of the mass of the full-length utrophin protein.

45 Bacmid: baculovirus shuttle vector.

BMD: Becker muscular dystrophy.

DMD: Duchenne muscular dystrophy.

Dystrophinopathy: All pathological conditions in mammals, including humans, due in full or in part to mutations in the gene(s) encoding the protein dystrophin (both now known or discovered in the future). Explicitly included within the definition of "dystrophinopathy" are BMD, DMD, EDMD, SBMA, XLDCM, elevated serum creatine kinase, and the like.

55 EDL: extensor digitorum longus muscle.

EDMD: Emery-Dreifuss muscular dystrophy.

"FLAG"-brand polypeptide: Generally, any polypeptide having the sequence DYKDDDDK (SEQ. ID. NO: 1), or a fragment thereof, such as the tetrapeptide DYKD (SEQ. ID. NO. 1), which can be used for isolating fusion proteins via affinity chromatography. The terms "FLAG" and "ANTI-FLAG" are registered trademarks of Sigma-Aldrich Biotechnology LP (St. Louis, Mo.). "FLAG"-brand polypeptides are available commercially from Sigma-Aldrich. See also Chubet & Brizzard (1996) "Vectors for expression and secretion of FLAG epitope-tagged proteins in mammalian cells," *Bio-techniques* 20(1):136-141.

HIV-TAT or TAT: Human immunodeficiency virus transactivator protein. "Tat" is short for "transactivator," a regulatory gene that accelerates the production of more HIV virus. "TAT" designates the protein, while "tat" designates the corresponding gene that encodes the TAT protein. In its native milieu, the TAT protein binds to the start of a new HIV RNA strand. Once bound, TAT encourages the transcription of the remainder of the HIV genetic code. TAT from HIV is a protein containing from 86 to 101 amino acids, depending upon the strain of HIV. The 86 amino acid-long sequence of HIV-1 TAT is shown in SEQ. ID. NO: 2. The entire genomic sequence of the HIV-1 virus, including the tat gene (at nts 5377-5591 and 7925-7970), is shown in SEQ. ID. NO: 3. See Gaynor, R. B. (1995) Regulation of HIV-1 gene expression by the transactivator protein Tat. *Curr Top Microbiol Immunol* 193, 51-77. See also GenBank Accession No. AF033819 for a fully annotated version of the HIV-1 genomic sequence.

LGMD: Limb-Girdle muscular dystrophy.

mdx Mice: A strain of mice arising from a spontaneous mutation (mdx) in inbred C57BL mice. The mutation is X chromosome-linked and produces viable homozygous animals that lack the muscle protein dystrophin. Mdx mice have high serum levels of muscle enzymes, and possess histological lesions similar to human muscular dystrophy. The histological features, linkage, and map position of mdx make these mice a widely utilized animal model for Duchenne muscular dystrophy. Mdx mice can be purchased from several commercial suppliers, including The Jackson Laboratory, Bar Harbor, Me. (sold under the registered trademark "JAX").

Operationally linked: when referring to two or more regions of a protein or a nucleotide sequence, "operationally linked" means the two regions are physically linked either directly or indirectly via intervening amino acid residues, nucleotide bases, or any other type of linking moiety.

PBS: phosphate-buffered saline.

PCR: polymerase chain reaction.

Pharmaceutically-suitable salt: any acid or base addition salt whose counter-ions are non-toxic to the patient in pharmaceutical doses of the salts so that the beneficial inhibitory effects inherent in the free base or free acid are not vitiated by side effects ascribable to the counter-ions. A host of pharmaceutically-suitable salts are well known in the art. For basic active ingredients, all acid addition salts are useful as sources of the free base form even if the particular salt, per se, is desired only as an intermediate product as, for example, when the salt is formed only for purposes of purification, and identification, or when it is used as intermediate in preparing a pharmaceutically-suitable salt by ion exchange procedures. Pharmaceutically-suitable salts include, without limitation, those derived from mineral acids and organic acids, explicitly including hydrohalides, e.g., hydrochlorides and hydrobromides, sulphates, phosphates, nitrates, sulphamates, acetates, citrates, lactates, tartrates, malonates, oxalates, salicylates, propionates, succinates, fumarates, maleates, methylene bis-hydroxynaphthoates, gentisates, isethionates, di-p-toluoyl-tartrates, methane sulphonates, ethanesulphonates, benzene-sulphonates, p-toluenesulphonates, cyclohexylsulphamates, quinates, and the like. Base addition salts include those derived from alkali or alkaline earth metal bases or conventional organic bases, such as triethylamine, pyridine, piperidine, morpholine, N methylmorpholine, and the like.

SBMA: Spinal bulbar muscular atrophy (also known as Kennedy's disease).

TA: tibialis anterior muscle.

Transduction: in general, the transfer of DNA from one cell to another; typically transduction is mediated via a bacteriophage, but any means of transferring the DNA from its

original source to its ultimate destination are included within the term "transduction" as used herein.

UTR or UTRN: utrophin. The nucleotide sequence for the human utrophin gene and the corresponding amino acid sequence for the encoded human utrophin protein are shown in SEQ ID NOS: 6 and 7, respectively; the nucleotide sequence for the murine utrophin gene and the corresponding amino acid sequence for the murine utrophin protein are shown in SEQ. ID. NOS: 8 and 9, respectively.

WT: wild-type.

XLDCM: X-linked dilated cardiomyopathy.

A first version of the invention is directed to a TAT-utrophin fusion protein (TAT-UTR), and use of the TAT-UTR fusion protein to treat dystrophinopathies in mammals, including humans. To demonstrate the efficacy of the TAT-UTR to treat dystrophinopathies in mammals, the mdx mouse is used as a model to demonstrate that TAT-UTR is imported into striated muscle cells and that the TAT-UTR fusion protein eliminates or significantly reduces the dystrophic phenotype in mdx mice.

Thus, in this first version of the invention, purified TAT-utrophin is injected into dystrophin-deficient mdx mice in an anti-dystrophic-effective amount. The mdx mouse model serves to demonstrate efficacy in all mammals, including humans. Measurements are then taken to assess the extent to which the TAT-utrophin is transduced into striated muscle cells in vivo. The localization of the TAT-utrophin is then assessed to determine how much of the TAT-utrophin is localized to the sarcolemma. (As shown in FIG. 1, natural dystrophin exerts its biological effect in close conjunction with the sarcolemma.) Measurements are also taken to determine whether the TAT-UTR fusion protein becomes stably associated with other dystrophin-associated proteins. The progress of mdx mice treated with the TAT-UTR is then followed to measure the improvement of several well-established parameters of the dystrophic phenotype, such as specific force and force drop in the muscles of the treated mice versus the control mice.

A second version of the invention is directed to mini- and micro-TAT-UTR constructs and methods of using these constructs to treat dystrophinopathies in mammals, including humans. Thus, the invention also encompasses truncated mini- and micro-TAT-utrophin constructs and the use of these truncated versions of the protein to treat dystrophinopathies. Reducing the physical size of the fusion protein results in improved protein transduction in vivo. Two representative truncated constructs are described herein. These truncated fusion proteins are designed to retain full activity for all known binding partners of utrophin, but with a 40 to 50% reduction in the mass of the protein. A third construct is designed to mimic the structure of the most extensively truncated, fully-functional dystrophin micro-gene.

Using TAT-UTR as a protein-based therapy for treating dystrophinopathies is a relatively low-cost, low-risk, but high-return approach to treating these currently intractable and fatal conditions. At present, there simply is no effective treatment available to treat prevalent dystrophinopathies such as DMD.

The present invention includes a series of utrophin constructs encoding the amino-terminal, actin-binding domain alone (UTRN), or the amino-terminal domain plus 4, 7, 10, or 11 spectrin-like repeats. FIG. 2 depicts the relative lengths of these constructs and their binding characteristics. As shown in FIG. 2, the constructs are designated herein as UTRN-R3, UTRN-R6, UTRN-R9, and UTRN-R10, respectively. Interestingly, the UTRN-R10 protein bound to actin filaments with essentially the same properties as full-length recombinant

utrophin (rUTR), which suggests UTRN-R10 encodes the complete actin-binding region of utrophin (see FIG. 2). The UTRN-R9, UTRN-R6, and UTRN-R3 proteins each bound to actin filaments with progressively lower affinity and stoichiometry as compared to full-length utrophin and UTRN-R10. (See FIG. 2.) These results demonstrate that the first ten (10) spectrin-like repeats of utrophin dramatically enhance the F-actin binding affinity and lateral association of the amino-terminal domain and provide a molecular basis for the greater effectiveness of full-length utrophin in rescuing dystrophin-deficient muscle as compared to a utrophin mini-gene deleted for spectrin-like repeats 4-19.

The present inventors have also expressed and characterized full-length mouse dystrophin. Recombinant dystrophin binds to actin filaments with a K_d of 0.4 μ M and B_{max} of 1 dystrophin molecule per 24 actin monomers (see FIG. 3D, second construct), which is remarkably close to the actin-binding properties of purified dystrophin-glycoprotein complex (Rybakova et al., 1996). In direct comparisons (see FIGS. 3A, 3B, and 3C), dystrophin and utrophin differed only in their extent of lateral association with actin filaments (1-to-24 for dystrophin and 1-to-14 for utrophin), and in the effect of increasing ionic strength on actin filament binding. These results strongly suggest that dystrophin and utrophin are both actin-binding proteins, but that the molecular epitopes important for filament binding differ between the two proteins.

While transgenic utrophin overexpression rescued all known phenotypes associated with dystrophin-deficiency in mdx mice (Tinsley et al., 1998), there remains a widespread perception that utrophin levels must greatly exceed the amount of dystrophin expressed in normal muscle in order to cause full rescue from the dystrophinopathic phenotype exhibited by mdx mice. This perception is based, at least in part, on an early quantitative estimate (Hoffman et al., 1987) of dystrophin abundance in normal muscle (0.002% of total muscle protein) and the present inventors' own measurements of utrophin expression (Rybakova et al., 2002) in normal (0.0006%) and mdx muscle (0.0013%), as well as in the Fiona line of transgenic mdx mice that overexpress utrophin to levels (0.014%) that fully corrected the mdx phenotype. (See Tinsley et al., 1998). From these measurements, it can reasonably be concluded that up to 7-fold greater levels of utrophin (0.014%/0.002%) may be necessary to compensate for dystrophin deficiency.

However, the early measurements of dystrophin levels in normal muscle used a relatively small recombinant protein fragment (Hoffman et al., 1987). While state-of-the-art at that time, the much smaller protein fragment used as the standard likely transferred to nitrocellulose more efficiently than the full-length dystrophin protein. Thus, it is possible that the previous measurements (Hoffman et al., 1987) may have significantly underestimated the abundance of dystrophin in normal muscle. Therefore, the abundance of dystrophin in normal skeletal muscle has now been measured by quantitative western blotting using full-length recombinant mouse dystrophin as the standard and iodinated secondary antibody as previously described for utrophin (Rybakova et al., 2002). The measurements (see the table shown in FIG. 4) suggest that the abundance of dystrophin in normal muscle is 10-times greater ($0.021 \pm 0.003\%$, $n=7$) (Rybakova et al., 2006) than previously reported (Hoffman et al., 1987). The new measurements more closely agree with the measured abundance of dystrophin (Ohlendeck et al., 1991) in highly purified sarcolemma vesicles (2% of sarcolemmal protein) and with quantitative estimates that sarcolemmal proteins comprise 1% of total muscle protein based on the density of sodium channels in total homogenates (0.09 pmol/mg total

protein) and in purified sarcolemmal vesicles (8 pmol/mg sarcolemmal protein) from rat skeletal muscle (Barchi and Weigele, 1979).

Most importantly, however, these data indicate that utrophin can fully rescue the mdx phenotype (Tinsley et al., 1998) when expressed to levels approaching that of dystrophin in normal muscle (0.014%/0.02%=70%).

The present invention is thus a method of using recombinant utrophin as a protein-based therapy for treating dystrophinopathies in general and DMD in particular. The present method uses TAT-utrophin chimeric (i.e., fusion) proteins. The TAT portion of the chimeric protein serves to mobilize the protein (i.e., transduce the protein) into muscle cells. The UTR portion of the chimeric protein serves to ameliorate or to eliminate the dystrophic condition.

One distinct benefit of the invention is that utrophin itself is not toxic. Therefore, the TAT-UTR fusion proteins can be administered in relatively high doses, thereby making it easier to transduce therapeutically effective amounts of the TAT-UTR fusion protein into muscle cells. Ubiquitous transgenic over-expression of utrophin itself caused no toxicity in a broad range of tissues (Fisher et al., 2001). Thus, in the present invention, an 11 kb full-length mammalian utrophin cDNA (mouse) (Guo et al., 1996) was cloned in-frame into the bacterial expression vector pTAT (Nagahara et al., 1998), which was kindly provided by Dr. Steven Dowdy (University of California, San Diego). A Kozak consensus sequence and a "FLAG"-brand type epitope were engineered in-frame 5' to TAT-utrophin by PCR.

The FLAG-TAT-utrophin construct was inserted into the pFastBac 1 donor plasmid (purchased commercially from Invitrogen, Carlsbad, Calif.). A map of the pFastBac1 donor plasmid is shown in FIG. 10 and the complete sequence of pFastBac1 is presented in SEQ. ID. NO: 4. Subsequent trans-formation into DHIOBac cells (purchased commercially from Invitrogen, catalog no. 18290-015) allowed for site-specific transposition into bMON14272 bacmid DNA. (The bMON14272 bacmid, along with the helper plasmid pMON7124, are included with the DHIOBac cells sold by Invitrogen. See Invitrogen's catalog no. 10359-016, and the product literature for Invitrogen's "BAC-TO-BAC"[®]-brand baculovirus expression system.)

Colonies containing recombinant bacmid DNA were identified by blue/white screening and high titer viral stocks were used to infect Sf21 insect cells (*Spodoptera frugiperda*) for protein expression. (Sf21 cells are available commercially from a number of international suppliers, including Orbigen Inc., San Diego Calif., and Gentaur, Brussels, Belgium.) Infected Sf21 cells were harvested 72 h post-infection and TAT-utrophin was purified from cell lysates using "ANTI-FLAG"-brand M2 affinity resin (obtained commercially from Sigma-Aldrich, St. Louis, Mo.). The gel depicted in FIG. 5 shows that FLAG-TAT-utrophin is expressed as a fully soluble protein and can be easily purified by "ANTI-FLAG" M2 affinity chromatography. Thus, sufficient TAT-utrophin can easily be prepared to perform a host of experiments. Moving from left-to-right, the lanes of the gel in FIG. 5 depict the cell lysate prior to chromatography, the M2 column void volume, and the M2 column wash. The lanes numbered 1-6 then depict the elution of the M2 column to obtain the resulting fusion protein.

To assess whether TAT-utrophin is measurably transduced into skeletal muscle, a 2.5 week-old mdx mouse received six intraperitoneal injections of TAT-utrophin (20 mg/kg in sterile PBS) administered biweekly. As a control, a littermate mdx mouse was sham-injected with sterile PBS in parallel. At age six weeks, both mice were euthanized, perfused with

PBS, and muscle tissue was excised for western blot, immunofluorescence and histological analyses. Western blot analysis of lysates from several tissues showed increased utrophin immunoreactivity in the TAT-utrophin-treated mdx mouse compared to the PBS-injected animal. See FIG. 6A, which is a gel depicting the utrophin immunoreactivity of the treated mouse versus the untreated mouse in several different tissue types. Importantly, immunofluorescence analysis of muscle cryosections revealed both increased HA-tag and DRP2 immunoreactivity localized to the sarcolemma of muscle from the animal treated with TAT-utrophin. See FIG. 6B, where the two upper panels depict immunoreactivity in the treated mouse and the two lower panels depict immunoreactivity in the untreated mouse.

Most strikingly, light microscopic analysis of haematoxylin and eosin-stained muscle cryosections showed dramatically decreased fibrosis and numbers of centrally nucleated myofibers in the TAT-utrophin treated animal compared to PBS-injected control. Compare FIGS. 7A, 7B, and 7C (which are photos of tibialis anterior ("TA") and quadriceps ("QUAD") muscle fibers from treated mice) to FIGS. 7D, 7E, and 7F (which are corresponding photos from untreated mice). In the quadriceps, the percentage of centrally nucleated fibers was 48% in the PBS-injected control, but only 24% in the TAT-utrophin-treated animal. The combined data of FIGS. 6A, 6B, 7A, 7B, 7C, 7D, 7E, and 7F show that TAT-utrophin effectively transduced skeletal muscle cells *in vivo*, correctly localized to the sarcolemma, and improved the histopathology of dystrophin-deficient mdx muscle.

Of course, recovery of muscle function is the ultimate criterion for evaluating the efficacy of any therapy for dystrophinopathies. Several studies have demonstrated that specific force production by mdx muscle is significantly decreased. It has also been shown that mdx muscle is hypersensitive to lengthening and eccentric contraction (Petrof et al., 1993; Moens et al., 1993). Therefore, these parameters were measured in sham- and TAT-utrophin treated mdx mice. (Kind thanks are extended to Dr. Richard L. Moss for his aid in conducting these tests.) FIGS. 8A and 8B provide data demonstrating that the eccentric contraction protocol described in Petrof et al. (1993) and Moens et al. (1993) can be performed and that these tests performed by the present inventors reproduced the key findings of Petrof et al. (1993) and Moens et al. (1993).

Regarding the key utility of the present invention, the Examples presented below clearly demonstrate that dystrophinopathic mammals treated according to the present invention show a significantly increased specific force produced by their muscles as compared to untreated dystrophinopathic mammals, as well as a significantly decreased force drop. See Example 3 and FIGS. 11 and 12. Thus, the utility of the compounds, compositions, and methods of the present invention is to ameliorate the disabling effects of dystrophinopathic conditions in mammals, including DMD in humans.

As indicated above, the invention includes pharmaceutical compositions comprising the fusion protein(s) described herein and/or a pharmaceutically acceptable salt thereof together with a pharmaceutically acceptable carrier. The compositions may also include other therapeutically active substances in addition to the fusion protein and/or salt thereof. The pharmaceutical compositions of the invention comprise an amount of the fusion protein and/or a pharmaceutically suitable salt thereof that is effective to ameliorate dystrophinopathic symptoms in a mammal suffering from a dystrophinopathy. In a pharmaceutical composition of the invention, the carrier must be pharmaceutically acceptable in the sense of being compatible with other ingredients in the

particular composition and not deleterious to the recipient thereof. The compositions include those suitable for oral, topical, rectal or parenteral (including subcutaneous, intramuscular, intraperitoneal, intradermal and intravenous) administration. Parenteral administration, either via the intramuscular or the intraperitoneal routes, is preferred.

In a particular version of the invention, the pharmaceutical compositions comprise the active ingredient (the fusion protein or a salt thereof) presented in a unit dosage form. The term "unit dosage" or "unit dose" is denoted to mean a predetermined amount of the active ingredient sufficient to be effective for treating dystrophinopathy. Preferred unit dosage formulations are those containing a daily dose, daily sub-dose, or an appropriate fraction thereof, of the administered active ingredient.

The pharmaceutical compositions may be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing the active ingredient into association with a carrier which constitutes one or more accessory ingredients. In general, the compositions are prepared by uniformly and intimately bringing the active ingredient into association with a liquid or solid carrier and then, if necessary, shaping the product into the desired unit dosage form.

Compositions of the present invention suitable for oral administration may be presented as discrete unit dosages, e.g., as capsules, cachets, tablets, boluses, lozenges and the like, each containing a predetermined amount of the active ingredient; as a powder or granules; or in liquid form, e.g., as a collyrium, suspension, solution, syrup, elixir, emulsion, dispersion and the like.

A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active compound in a free-flowing form, e.g., a powder or granules, optionally mixed with accessory ingredients or excipients, e.g., binders, lubricants, inert diluents, surface active or dispersing agents. Molded tablets may be made by molding in a suitable machine, a mixture of the powdered active compound with any suitable carrier.

Compositions suitable for parenteral administration conveniently comprise a sterile injectable preparation of the active ingredient in, for example, a solution which is preferably isotonic with the blood of the recipient. Useful formulations also comprise concentrated solutions or solids containing the active ingredient which upon dilution with an appropriate solvent give a solution suitable for parenteral administration. The parenteral compositions include aqueous and non-aqueous formulations which may contain conventional adjuvants such as buffers, bacteriostats, sugars, thickening agents and the like. The compositions may be presented in unit dose or multi-dose containers, for example, sealed ampules and vials.

Compositions suitable for topical or local application (including ophthalmological administration) comprise the active ingredient formulated into pharmaceutically-acceptable topical vehicles by conventional methodologies. Common formulations include drops, collyriums, aerosol sprays, lotions, gels, ointments, plasters, shampoos, transferosomes, liposomes and the like.

Compositions suitable for inhalation administration, wherein the carrier is a solid, include a micronized powder or liquid formulation having a particle size in the range of from about 5 μm or less to about 500 μm , for rapid inhalation through the nasal or oral passage from a conventional inhalation squeeze or spray container. Suitable liquid nasal com-

positions include conventional nasal sprays, nasal drops and the like, of aqueous solutions of the active ingredient and optional adjuvants.

In addition to the aforementioned ingredients, the compositions of this invention may further include one or more optional accessory ingredients(s) utilized in the art of pharmaceutical formulations, e.g., diluents, buffers, flavoring agents, colorants, binders, surfactants, thickeners, lubricants, suspending agents, preservatives (including antioxidants), and the like.

The amount of active ingredient required to be effective for any specific dystrophinopathy in any specific patient will, of course, vary with the individual mammal being treated and is ultimately at the discretion of the medical or veterinary practitioner. The factors to be considered include the species and sex of the mammal, the dystrophinopathic condition being treated, the route of administration, the nature of the formulation, the mammal's body weight, surface area, age and general condition, and the particular compound to be administered.

In general, the pharmaceutical compositions of this invention contain from about 0.5 to about 500 mg and, preferably, from about 5 to about 350 mg of the active ingredient, preferably in a unit dosage form, for each of the indicated activities. However, a suitable effective dose is in the range of about 0.1 to about 200 mg/kg body weight per day, preferably in the range of about 1 to about 100 mg/kg per day, calculated as the non-salt form of the fusion protein. The total daily dose may be given as a single dose, multiple doses, e.g., two to six times per day, or by intravenous or parenteral infusion for a selected duration. Dosages above or below the range cited above are within the scope of the present invention and may be administered to the individual patient if desired and necessary. In topical formulations, the subject compounds are preferably utilized at concentrations of from about 0.1% to about 5.0% by weight.

EXAMPLES

The following Examples are presented solely to provide a more complete description of the invention disclosed and claimed herein. The Examples do not limit the scope of the invention claimed herein in any fashion.

Example 1

—Expression, Purification of TAT-Utrophin; General Protocols:

1.a. Expression and Purification of TAT-Utrophin. High titer stocks of recombinant baculovirus encoding the "FLAG"-tagged TAT-utrophin chimera were used to infect Sf21 insect cells for protein expression by a shaker culture procedure described in the manufacturer's instructions. Infected Sf21 cells were harvested 72 h post-infection and resuspended in 10 ml of 50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1% Triton X-100, and a cocktail of protease inhibitors. The soluble lysate was circulated over a 2 ml "ANTI-FLAG" M2 agarose column (Sigma-Aldrich). The column was washed extensively with 10 mM Tris-HCl, pH 7.4, 150 mM NaCl and bound protein eluted with the same buffer containing 100 µg/ml "FLAG"-brand peptide (Sigma-Aldrich). Purified protein was concentrated in a Centricon 100 column (Amicon) and quantified with the Bio-Rad DC Protein Assay Kit using BSA as standard. The typical yield of pure utrophin was 700 µg when only five 177 cm² plates of cell monolayer were used as a starting material. The protocols can be easily scaled up as needed.

Quality control analysis. The data indicate that TAT-utrophin is abundantly expressed in a highly soluble form that can be readily purified by "ANTI-FLAG" affinity chromatography (see FIG. 5). It is critical to note that including the TAT sequence within the fusion protein has no adverse effect on utrophin structure/function. The purified TAT-utrophin is to be analyzed by gel permeation chromatography (Rybakova and Ervasti, 1997), velocity sedimentation analysis (Ervasti et al., 1991) and electron microscopy after rotary shadowing (Rybakova et al., 2002). These analyses yield quantitative measures for the native molecular weight, dimensions, shape, oligomeric/aggregative state as well as an assessment of proper folding.

The F-actin binding properties of TAT-utrophin are measured using the established high-speed co-sedimentation assay (see FIG. 3B) and binding data is analyzed by nonlinear regression analysis. These experiments will yield both the apparent K_d and B_{max} of recombinant protein binding to F-actin. See FIG. 3D. The ability of different proteins to protect actin filaments from depolymerization is measured by monitoring the time-dependent decay in fluorescence of preformed filaments seeded with PRODAN-labeled (i.e., 6-propionyl-2-(N,N-dimethyl)aminonaphthalene-labeled) monomers at Cys374 (Marriott et al., 1988; Miyata et al., 1997) as shown in FIG. 3C. All data is compared to those measured for recombinant utrophin performed in parallel.

More specifically, an 11 kb full-length murine utrophin cDNA was subcloned in-frame into the bacterial expression vector pTAT to generate PTAT-Utr. To prepare for eventual expression and purification of TAT-Utrophin in Sf21 insect cells using a baculovirus expression system, a Kozak consensus sequence and FLAG-epitope were engineered in-frame at the extreme 5' end of TAT-Utr using PCR primers KJS36 (5' gcggccgcacacatggactacaagga-
caacgatgacaaggctacggccgaagaac-3') (SEQ. ID. NO: 26) (FLAG-epitope is underlined) and KJS32 (5'-ggagatgcacagcaacagtttcaggacttagg-3') (SEQ. ID. NO: 27). This FLAG-TAT-utrophin construct was inserted into the bacmid donor plasmid pFastBac1 (Invitrogen, Carlsbad, Calif.) and subsequently transformed into DH10BAC (Invitrogen) bacterial cells to allow for site-specific transposition into bacmid DNA. Recombinant bacmid DNA was purified and used to transfect Sf21 cell monolayers in order to generate recombinant baculovirus. Recombinant virus infection of Sf21 monolayers and recombinant protein purification using anti-FLAG M2 affinity resin (Sigma, St. Louis, Mo.) was performed as per the manufacturer's instructions.

Elution fractions were pooled, dialyzed against phosphate buffered saline (PBS) overnight, and concentrated using a Centricon 100 (Millipore, Concord, Mass.). The purified protein was sterilized for injection by passage through a 0.22 µm filter and injected into the intraperitoneal cavity of mdx mice at a concentration of 0.5 to 1.0 mg/ml. The pure protein was stable for up to 4 days when kept on wet ice at 4° C. (assessed by a lack of degradation on Coomassie blue stained SDS-polyacrylamide gels), so a single protein preparation was utilized for up to 2 injections when possible. Otherwise, protein was prepared fresh for each injection.

1.b. Treatment Time Course. Pairs of female C57Bl/10ScSn-Dmdmdx/J (The Jackson Laboratory, Bar Harbor, Me.) littermates were treated in parallel, one of which received a dose of 20 µg TAT-utrophin/g body weight while the control mouse received equal volume injections of sterile PBS. A total of 6 biweekly injections were administered over three weeks, beginning at 18 days and culminating at 35 days of age. Three days after the final injection, serum and tissue were collected for creatine kinase, western blot, immunofluo-

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rescence, histological, and physiological analyses. Animals were housed and treated in accordance with the standards set by the University of Wisconsin Institutional Animal and Care and Use Committee.

1.c. Protein extracts and Western Blotting. Tissues were dissected from freshly killed mice and snap frozen in liquid nitrogen. Frozen tissue was pulverized with a liquid nitrogen-cooled mortar and pestle and solubilized in 1% SDS, 5 mM EGTA, and a cocktail of protease inhibitors. Samples were incubated for 2 minutes at 100° C. and centrifuged 2 min at 12000×g. The supernatant protein concentration was determined with the Bio-Rad DC protein assay kit using bovine serum albumin as standard. Equal amounts of protein was separated by SDS-PAGE and transferred to nitrocellulose. Western blot analysis of utrophin levels was performed with rabbit polyclonal antibody 103 raised against the carboxyl-terminus of utrophin (generously provided by Dr. Stanley Froehner, University of Washington) diluted 1:250 in BLOTTO (i.e., bovine lacto transfer technique optimizer, a blocking reagent made from nonfat dry milk and PBS) (5% milk in PBS, pH 7.5) and anti-FLAG monoclonal antibody M2 (Sigma) diluted 1:1000 in BLOTTO. (BLOTTO blocking reagents are also commercially available from, for example, Thermo-Fisher Scientific, Waltham, Mass., catalog no. PI-37530.)

1.d. Histological and Morphometric Analysis. Individual muscles were dissected from freshly killed mice, coated with "O.C.T." matrix solution ("TissueTek"®-brand, Sakura Finetek, Torrance, Calif.; O.C.T. refers to "optimum cutting temperature," a specimen matrix formulation comprising water-soluble glycols and resins for cryostat sectioning at temperatures of -10° C. and below), and rapidly frozen in liquid nitrogen-cooled isopentane. Ten (10) µm thick cryosections were cut on a Leica CM3050 cryostat, allowed to dry, and stained with hematoxylin and eosin-phloxine. Sections cut from the mid-belly of both the tibialis anterior and quadriceps were selected for histological assessment. Images were collected on a Zeiss Axiovert 25 microscope and compiled into montages of entire sections in CorelDraw 10 and exported to Scion Image (Scion Corporation, Frederick, Maryland) for morphometric analyses. The percentage of centrally nucleated fibers and fiber diameters were determined from one muscle of each mouse, with every fiber scored for CNF analysis and ~700 fiber diameters measured per muscle section. A Student's t-test was used to compare average CNF values and average fiber diameter. To determine statistical significance of fiber diameter variability, a student's t-test was performed on the standard deviations of individual muscle sections.

1.e. Immunofluorescence. 10 µm thick cryosections were fixed in 4% paraformaldehyde for 10 minutes, washed 3×10 minutes in PBS, and blocked in 5% goat serum for 30 minutes. Primary antibodies were applied in 5% goat serum overnight at 4° C. and washed off 3×10 minutes in PBS. "ALEXA"®-brand 488- or 568-conjugated secondary antibodies (Invitrogen) were incubated for 30 min before a final 3×10 minute wash cycle. Coverslips were applied with a drop of Anti-Fade Reagent (Molecular Probes) and confocal images obtained using a Bio-Rad MRC 1000 scan head mounted transversely to an inverted Nikon Diaphot 200 microscope at the Keck Center for Biological Imaging. Primary monoclonal antibodies used were anti-HA tag HA. 11 (BABC0, Berkeley, Calif.) 1:1000; anti-utrophin DRP2 (Novacastra, Newcastle upon Tyne, UK) 1: 10; anti-β-dystroglycan b-DG (Novacastra) 1:1000; anti-α-sarcoglycan (NCL-a-SARC; α-SG), (Novacastra) 1:1000; and anti-γ-sarcoglycan g-SARC (Novacastra) 1:1000.

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1.f. Contractile Properties. All mechanical properties were adapted from Petrof et al. After rapid PBS perfusion, the extensor digitorum longus (EDL) muscles were quickly dissected tendon to tendon and immersed in an O₂-saturated Ringer's solution (135 mM NaCl, 4 mM KCl, 1 mM MgCl₂, 10 mM HEPES, 10 mM glucose, and 1.8 mM CaCl₂, pH 7.4) at 25° C. Suture silk (4-0) was used to attach one tendon to a rigid support and the other to a dual lever force transducer (Aurora Scientific, Ontario, Canada) and the entire apparatus was immersed in oxygenated Ringer's solution and allowed to equilibrate for 5 minutes. Muscles were stimulated through two platinum plate electrodes on either side of the muscle. A range of twitch stimulations were performed to determine L_o, the muscle length at which maximal twitch force was produced. After 5 minutes of recovery, the EDL was maximally activated to determine maximal tetanic tension. Data were normalized against cross-sectional area of each individual muscle.

Protection against mechanical injury was assessed by subjecting the muscle to a series of five eccentric contractions (ECC). Each ECC consisted of maximally activating the muscle for 700 ms, with a stretch of 0.5 L_o/s over the final 200 ms to result in a total stretch of 0.1 L_o. Five minutes of recovery time was allowed between each ECC. Force drop was calculated as (ECC1-ECC5)/ECC1. Data were compared using ANOVA followed by a Tukey post hoc test.

1.g. Serum CK Analysis. Retro-orbital bleeds were performed on anesthetized mice using heparinized capillary tubes. Approximately 100 µl of blood was obtained per mouse, centrifuged at 5000 rpm and the serum layer removed and stored at -80° C. for analysis. Creatine kinase levels were determined using Vitros CK DT slides (Ortho-Clinical Diagnostics, Raritan, N.J.) and analyzed using a Kodak Ektachem DT60 Analyzer as per the manufacturer's instructions. Data were collected in Units/ml and compared using a Student's T-test.

Example 2

—Effect of TAT-Utrophin on the Dystrophic Phenotype of mdx Mice:

In this Example, purified TAT-utrophin is injected into dystrophin-deficient mdx mice. The mice are then examined to assess the extent to which the TAT-utrophin is transduced into striated muscle cells in vivo. The extent of uptake is measured, and the amount of TAT-utrophin localized to the sarcolemma is determined. Optionally, it may also be determined whether the TAT-utrophin becomes stably associated with other dystrophin-associated proteins. The quantitative improvement of several well-established parameters of the dystrophic phenotype is then measured in mdx mice treated with TAT-utrophin and compared to untreated controls and placebo groups.

Administration of TAT-utrophin—Purified TAT-utrophin is dialyzed against phosphate-buffered saline and sterilized by passage through a Millex-GP 0.22 µm filter. Assuming 100% protein transduction specifically into skeletal muscle, a minimal dose of 11 µg TAT-utrophin per gram body weight is believed to compensate for dystrophin deficiency. Of course, it is likely that TAT-utrophin will distribute to all tissues and transduction efficiency will almost certainly be less than complete. Therefore, TAT-utrophin is preferably administered via intraperitoneal injection at several different concentrations ranging from 1-5 mg/ml and total injection volumes of 0.1-0.5 ml.

Measurement of TAT-Utrophin Uptake and Cellular Location - TAT-utrophin uptake into skeletal muscle and cellular localization is assessed by two methods. In the first method, mice are deeply anesthetized with avertin, the chest wall is opened, and the animals are infused for 20 minutes with phosphate-buffered saline through the left ventricle with an outflow path from the right atrium. Skeletal muscles are then dissected and used in the preparation of KCl-washed skeletal muscle membranes (Ohlendieck et al., 1991), or immediately snap-frozen in liquid nitrogen to prepare SDS total protein lysates (Rybakova et al., 2002). Both preparations are analyzed for TAT-utrophin content by quantitative western blot analysis using "ANTI-FLAG"-brand M2 antibody (Sigma-Aldrich) detected with ¹²⁵I-goat anti-mouse IgG and the signals quantitated by phosphor autoradiography. Analysis of total protein lysates and KCl-washed membranes provides a measure of the fraction of TAT-utrophin stably associated with the sarcolemma. The absolute utrophin content in SDS muscle lysates of TAT-utrophin-treated mice is also quantitatively compared to the utrophin content of sham-treated mdx mice and transgenic mdx mice expressing full-length utrophin (Fiona) to levels that rescue all known phenotypes of mdx mice. These comparisons provide a quantitative assessment of the TAT-utrophin uptake relative to a fully-rescued transgenic animal model.

In the second method, anesthetized animals are infused for 2 minutes with PBS followed by a 20 minute infusion of 2% paraformaldehyde in PBS. Various skeletal muscles are dissected, post-fixed for 5 minutes in 2% paraformaldehyde, and frozen in liquid nitrogen-cooled isopentane. From 8 μm cryosections, both the uptake and cellular location of TAT-utrophin is assessed using confocal immunofluorescence microscopy.

The KCl-washed membranes, SDS lysates and cryosections prepared from TAT-utrophin-treated mdx mice are also used to detect alterations in the abundance and sarcolemmal localization of other proteins within the dystrophin-glycoprotein complex including α- and β-dystroglycan, α-, β-, γ- and δ-sarcoglycans, syntrophin and α-dystrobrevins. Relative protein abundance can be assessed by quantitative western blot analysis of total muscle SDS extracts (Rybakova et al., 2002), while cellular localization and organization can be assessed by immunofluorescence analysis of both longitudinal and transverse cryosections and mechanically peeled sarcolemma (Rybakova et al., 2000).

Assessment of costamere structure and function—To assess whether TAT-utrophin treatment can restore mechanical coupling between the sarcolemma and costameric γ-actin, confocal immunofluorescence microscopy analysis is performed on mechanically peeled sarcolemma (Rybakova et al., 2000) from sham and TAT-utrophin-treated mdx mice. Paraformaldehyde-fixed sarcolemma are blocked for 2 h at 4° C. with 5% serum in PBS and incubated with the appropriate primary antibodies overnight at 4° C. The specimens are washed with PBS, incubated with fluorescent secondary antibody for 30 min at 37° C., rinsed and sealed under coverslips in an anti-fade solution.

Assessment of Dystrophic Phenotype—Skeletal and cardiac muscle of dystrophin-deficient mdx mice exhibits several histologic and physiologic defects in common with patients suffering from Duchenne muscular dystrophy. Most notable (and easily measured) are a dramatic elevation in centrally nucleated fibers of irregular size resulting from muscle fiber necrosis/regeneration and elevated serum creatine kinase levels due to sarcolemmal instability.

For histologic analysis, 8 μm cryosections of skeletal muscle from control, sham-injected, and TAT-utrophin-in-

jected mdx mice are stained with haematoxylin and eosin and the percentage of central nuclei and mean fiber diameter measured. Histological analyses are also performed on several different muscles to compare the effects of TAT-utrophin on different fiber types, and muscles experiencing different work loads and activities. Measurement of these parameters in C57BL/10 control and sham-injected mdx mice provides a baseline and elevated values for normal and dystrophic muscle, respectively. While the number of centrally-nucleated fibers is already quite high (~40%) in 4 week-old mdx mice (Warner et al., 2002), this parameter doubles yet again by 10-12 weeks of age (Warner et al., 2002). Therefore, it is possible to measure a decrease in the percentage of centrally nucleated fibers in mdx mice treated for 2 months with TAT-utrophin compared to sham-treated mice.

To assess for sarcolemmal damage, quantitative colorimetric analysis of serum creatine kinase levels is performed using CK DT slides (Ortho-Clinical Diagnostics) measured with a Kodak Ektachem DT 60 Analyzer. A minimum of 5 animals in each treatment regime are measured at several time points post-injection. Evans blue infiltration is also assessed, which has been shown to accumulate significantly in dystrophin deficient mdx cardiac and skeletal muscle (Straub et al., 1997). Evans blue dye in sterile PBS is injected into the tail veins of control and knockout littermates and the animals sacrificed 3-6 h after dye administration. After skinning, the animals are visually inspected for macroscopic dye uptake by a blue coloration of limb muscles. 100% of mdx mice and 0% of control mice exhibit indication of membrane damage by this technique (Straub et al., 1997). In addition, 8 μm cryosections are examined by immunofluorescence microscopy to quantitate the fraction of muscle cells infiltrated by Evans blue (Straub et al., 1997).

Assessment of contractile function—Several studies have demonstrated that specific force production by mdx muscle is significantly decreased and hypersensitive to lengthening, or eccentric contraction (Petrof et al., 1993; Moens et al., 1993). Thus, the measure isometric twitch and tetanic tension in intact muscles from sham- and TAT-utrophin treated mdx mice are measured. The EDL muscle is dissected tendon to tendon and allowed to equilibrate in oxygenated mammalian Ringers' solution (Eddinger et al., 1986), and then tied into a dual mode force transducer (Aurora Scientific). The muscle length (L_o) at which maximal twitch tension is obtained is determined with a single pulse at a stimulation frequency of 2500 Hz at increasing muscle lengths. After a 10 minute wait, the muscle undergoes a series of 5 eccentric contractions (ECC) with the maximal tetanic tension measured for each round. The ECC protocol involves stimulation at 150 Hz at L_o for 500 msec followed by lengthening the muscle by 0.5 L_o /sec for 200 msec before relaxing at a rate of 0.5 L_o /sec for 200 msec. This protocol results in a stretch equal to 10% L_o . There is a 5 minute wait in between each ECC to allow the muscle to recover. All measurements are recorded and analyzed using Dynamic Muscle Control and Analysis Software (Aurora Scientific).

Example 3

—Generation of Mini- and Micro-TAT-Utrophin Constructs:

In parallel with the experiments described in Example 1, the invention also encompasses fusion proteins wherein the utrophin portion of the fusion protein has been truncated (to lower the molecular weight of the fusion protein), without deleteriously impacting the anti-dystrophinopathic activity of

the fusion protein. Thus, the invention encompasses truncated, but fully functional mini- and micro-TAT-utrophin constructs. It is hoped that reducing the size of the chimera leads to improved protein uptake.

Bacmid construction—Mini- and micro-TAT-utrophin constructs are generated with the “BAC-TO-BAC”-brand expression system (anvitrogen), which has been used to express full-length mouse utrophin (Rybakova et al., 2002), dystrophin (see FIG. 3A), and numerous truncation constructs (see FIG. 2). Briefly, all expression constructs are PCR-amplified from the TAT-utrophin construct using PfuUltra high-fidelity DNA polymerase (Stratagene) to incorporate an amino-terminal “FLAG”-brand type purification tag (DYKDDDDK) (SEQ. ID. NO: 1) followed by the HIV TAT protein transduction sequence (YGRKKRRQRRR) (SEQ. ID. NO: 5). The HIV TAT protein transduction sequence is preferred. However, any sequence that functions to transduce the fusion protein into mammalian muscle cells may be used in its place. The mini- and micro-constructs planned or actually made are shown schematically in FIG. 9. Preferably, the constructs all contain intact cysteine-rich and carboxy-terminal domains to ensure optimal β -dystroglycan binding activity (Ishikawa-Sakurai et al., 2004).

Based on actin-binding studies of serially-truncated utrophin constructs performed by the present inventors (data not shown), it is expected that TAT-UTRAR11-22 should have near-optimal actin filament binding activity, but with a 40% reduction in molecular weight (237,000) compared to full-length utrophin (394,000). TAT-UTRAR7-22, which is less than half the molecular weight of full-length utrophin (189,000) will also be evaluated, but at the expense of diminished actin-binding activity. TAT-UTRAR4-21 will also be generated and tested. This construct is expected to bind actin with the lowest affinity. It is an attractive compound for incorporation into a pharmaceutical composition because based on its small size (42% of full-length utrophin), and in light of the success of the analogous dystrophin micro-gene to rescue the mdx phenotype (Harper et al., 2002).

pFASTBAC1 donor plasmids carrying each new TAT construct is transformed into DHIOBAC for site-specific transposition into bMON14272 bacmid DNA. Colonies containing recombinant bacmid DNA are identified by blue-white screening and high titer viral stocks produced for infection of Sf21 insect cells for protein expression. Protein purification, quality control and transduction efficacy are performed as described earlier.

Example 4

—Dose-Dependent Amelioration of Dystrophin-Deficient Phenotype:

To determine whether the ability of TAT-utrophin to ameliorate the dystrophin-deficient phenotype is dose-dependent, the protective effects of increased dosages of TAT-utrophin were assessed on the dystrophin-deficient mdx mouse. An initial dosage of 20 μ g protein/g mouse body weight was arbitrarily designated as a dosage of “1 \times .” A study was then performed in which littermate mdx mice were injected with 1 \times (20 μ g protein/g body weight), 2 \times (40 μ g protein/g body weight), and 5 \times (100 μ g protein/g body weight) levels of TAT-utrophin. The timeline of treatment was consistent with the original 1 \times studies (see above) in which 2.5 week-old mdx mice received six intraperitoneal injections at the indicated dosage. The doses were administered bi-weekly. As controls, littermate mdx mice received sterile PBS injections in parallel. At six weeks of age, treated and control mice were eutha-

nized and assessed for several functional and histological parameters of dystrophin deficiency.

Of note, the 5 \times -treated mdx mouse demonstrated an approximately 45% increase in specific force generation over PBS-injected mice (see FIG. 11, which depicts the results for PBS-treated versus TAT-utrophin-treated mice). Specific force is an index of maximal force generated by a muscle normalized against the cross-sectional area of the muscle; mdx muscle typically generates approximately 25-30% less specific force than wild-type mice (Petrof et al., 1993).

Additionally, the treated mdx mice exhibited a dose-dependent improvement in protection against contraction-induced injury (see FIG. 12, which depicts the results for PBS-treated versus TAT-utrophin-treated mice). Contraction-induced injury is a parameter quantified by the drop in maximal force generation after five (5) consecutive damaging eccentric contractions. Wild-type force drop values are typically 15-25%, while the corresponding mdx values range from 60-80% (Petrof et al., 1993). As shown in FIG. 12, the 5 \times -treated mdx mice had a force drop value in the range of 20%, which is well within the range for non-mdx, wild-type mice. In contrast, the PBS-treated mdx mice had a force drop value typical of mdx mice, an approximately 65% drop.

Example 5

—Reduction of Serum Creatine Kinase in TAT-Utrophin-treated Mice:

To assess whether the protective effects of TAT-utrophin were mitigated through restoration of dystrophin complex members to the sarcolemma, immunofluorescence analyses were carried out on cryosections from TAT-utrophin and PBS-injected quadriceps. While no signal was observed on cryosections from PBS-treated muscle stained for the transmembrane glycoproteins β -dystroglycan, α -sarcoglycan, and γ -sarcoglycan, each antibody probe revealed intense staining along the periphery of muscle cells from TAT-utrophin-treated mice (FIG. 13A). Sarcolemmal integrity was also assessed by measuring serum levels of the muscle-specific enzyme creatine kinase (CK), which are typically elevated ~20 fold in mdx mice. TAT-utrophin-treated mice demonstrated a 50% reduction in serum CK activity compared to PBS-injected controls. See FIG. 13B. These results strongly indicate that TAT-utrophin not only restored dystrophin complex members to the sarcolemma but also partially protected against membrane instability.

The significance of these Examples is that they show that the TAT-utrophin constructs function to ameliorate dystrophinopathy in a dose-dependent fashion. The Examples also show the now best-known combination of transduction efficiency, size, and pharmacological activity to rescue phenotypically dystrophic mammals.

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<210> SEQ ID NO 7

<211> LENGTH: 3433

<212> TYPE: PRT

<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 7

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35        40        45
Gly Lys Pro Pro Ile Asn Asp Met Phe Thr Asp Leu Lys Asp Gly Arg
50        55        60
Lys Leu Leu Asp Leu Leu Glu Gly Leu Thr Gly Thr Ser Leu Pro Lys
65        70        75        80
Glu Arg Gly Ser Thr Arg Val His Ala Leu Asn Asn Val Asn Arg Val
85        90        95
Leu Gln Val Leu His Gln Asn Asn Val Glu Leu Val Asn Ile Gly Gly
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Thr Asp Ile Val Asp Gly Asn His Lys Leu Thr Leu Gly Leu Leu Trp
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Ile Ala Val Ile His Glu Lys	Gln Pro Asp Val Ile	Leu Glu Ala		
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Tyr Arg His Gln Leu Asp Glu	Ile Ile Cys Trp Leu	Thr Lys Ala		
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<210> SEQ ID NO 9

<211> LENGTH: 3430

<212> TYPE: PRT

<213> ORGANISM: Mus musculus

<400> SEQUENCE: 9

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Gln Lys Lys Thr Phe Thr Lys Trp Ile Asn Ala Arg Phe Ser Lys Ser
 35             40             45
Gly Lys Pro Pro Ile Ser Asp Met Phe Ser Asp Leu Lys Asp Gly Arg
 50             55             60
Lys Leu Leu Asp Leu Leu Glu Gly Leu Thr Gly Thr Ser Leu Pro Lys
 65             70             75             80
Glu Arg Gly Ser Thr Arg Val His Ala Leu Asn Asn Val Asn Arg Val
 85             90             95
Leu Gln Val Leu His Gln Asn Asn Val Asp Leu Val Asn Ile Gly Gly
 100            105            110
Thr Asp Ile Val Asp Gly Asn Pro Lys Leu Thr Leu Gly Leu Leu Trp
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Arg	Gln	Thr	Thr	Arg	Pro	Tyr	Ser	Gln	Val	Asn	Val	Leu	Asn	Phe	Thr
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Thr	Ser	Trp	Thr	Asp	Gly	Leu	Ala	Phe	Asn	Ala	Val	Leu	His	Arg	His
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Lys	Pro	Asp	Leu	Phe	Ser	Trp	Asp	Arg	Val	Val	Lys	Met	Ser	Pro	Ile
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Glu	Arg	Leu	Glu	His	Ala	Phe	Ser	Lys	Ala	His	Thr	Tyr	Leu	Gly	Ile
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Glu	Lys	Leu	Leu	Asp	Pro	Glu	Asp	Val	Ala	Val	His	Leu	Pro	Asp	Lys
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Lys	Ser	Ile	Ile	Met	Tyr	Leu	Thr	Ser	Leu	Phe	Glu	Val	Leu	Pro	Gln
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Gln	Val	Thr	Ile	Asp	Ala	Ile	Arg	Glu	Val	Glu	Thr	Leu	Pro	Arg	Lys
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Val	Leu	Thr	Trp	Leu	Leu	Ser	Ala	Glu	Asp	Thr	Phe	Gln	Glu	Gln	Asp
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Asp	Ile	Ser	Asp	Asp	Val	Glu	Glu	Val	Lys	Glu	Gln	Phe	Ala	Thr	His
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Glu	Thr	Phe	Met	Met	Glu	Leu	Thr	Ala	His	Gln	Ser	Ser	Val	Gly	Ser
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Asp	Ala	Leu	Met	Glu	Leu	Gln	Lys	Lys	Gln	Leu	Gln	Gln	Leu	Ser	Ser
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Trp	Leu	Ala	Leu	Thr	Glu	Glu	Arg	Ile	Gln	Lys	Met	Glu	Ser	Leu	Pro
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Leu	Gly	Asp	Asp	Leu	Pro	Ser	Leu	Gln	Lys	Leu	Leu	Gln	Glu	His	Lys
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Ser	Leu	Gln	Asn	Asp	Leu	Glu	Ala	Glu	Gln	Val	Lys	Val	Asn	Ser	Leu
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Thr	His	Met	Val	Val	Ile	Val	Asp	Glu	Asn	Ser	Gly	Glu	Ser	Ala	Thr
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Ala	Leu	Leu	Glu	Asp	Gln	Leu	Gln	Lys	Leu	Gly	Glu	Arg	Trp	Thr	Ala
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Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp	Asn	Arg	Leu	Gln	Glu	Ile	Ser
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		995					1000						1005		
His	Leu	Leu	Glu	Glu	Ile	Thr	Pro	Arg	Leu	Arg	Asp	Phe	Glu	Ala	
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His Val Leu Asp Val Arg	Asp Val Asp Pro Asp Val	Ile Gln Ala		
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His Leu Asp Lys Cys Met	Lys Leu Tyr Lys Thr Leu	Ser Glu Val		
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1685	1690	1695		
Asp Ala Ser Leu Gln Val	Glu Asn Val Arg Glu Gln	Ala Ile Ile		
1700	1705	1710		
Leu Val Asn Ala Arg Gly	Ser Ala Ser Arg Glu Leu	Val Glu Pro		
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Lys Leu Ala Glu Leu Ser	Arg Asn Phe Glu Lys Val	Ser Gln His		
1730	1735	1740		
Ile Lys Ser Ala Arg Met	Leu Ile Gly Gln Asp Pro	Ser Ser Tyr		
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Gln Gly Leu Asp Pro Ala	Gly Thr Val Gln Ala Ala	Glu Ser Phe		
1760	1765	1770		

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Ser	Asp	Leu	Glu	Asn	Leu	Glu	Gln	Asp	Ile	Glu	Asn	Met	Leu	Lys
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Val	Val	Glu	Lys	His	Leu	Asp	Pro	Asn	Asn	Asp	Glu	Lys	Met	Asp
1790						1795						1800		
Glu	Glu	Gln	Ala	Gln	Ile	Glu	Glu	Val	Leu	Gln	Arg	Gly	Glu	His
1805						1810						1815		
Leu	Leu	His	Glu	Pro	Met	Glu	Asp	Ser	Lys	Lys	Glu	Lys	Ile	Arg
1820						1825						1830		
Leu	Gln	Leu	Leu	Leu	Leu	His	Thr	Arg	Tyr	Asn	Lys	Ile	Lys	Thr
1835						1840						1845		
Ile	Pro	Ile	Gln	Gln	Arg	Lys	Thr	Ile	Pro	Val	Ser	Ser	Gly	Ile
1850						1855						1860		
Thr	Ser	Ser	Ala	Leu	Pro	Ala	Asp	Tyr	Leu	Val	Glu	Ile	Asn	Lys
1865						1870						1875		
Ile	Leu	Leu	Thr	Leu	Asp	Asp	Ile	Glu	Leu	Ser	Leu	Asn	Met	Pro
1880						1885						1890		
Glu	Leu	Asn	Thr	Thr	Val	Tyr	Lys	Asp	Phe	Ser	Phe	Gln	Glu	Asp
1895						1900						1905		
Ser	Leu	Lys	Ser	Ile	Lys	Gly	Gln	Leu	Asp	Arg	Leu	Gly	Glu	Gln
1910						1915						1920		
Ile	Ala	Val	Val	His	Glu	Lys	Gln	Pro	Asp	Val	Ile	Val	Glu	Ala
1925						1930						1935		
Ser	Gly	Pro	Glu	Ala	Ile	Gln	Ile	Arg	Asp	Met	Leu	Ala	Gln	Leu
1940						1945						1950		
Asn	Ala	Lys	Trp	Asp	Arg	Val	Asn	Arg	Val	Tyr	Ser	Asp	Arg	Arg
1955						1960						1965		
Gly	Ser	Phe	Ala	Arg	Ala	Val	Glu	Glu	Trp	Arg	Gln	Phe	His	His
1970						1975						1980		
Asp	Leu	Asp	Asp	Leu	Thr	Gln	Trp	Leu	Ser	Glu	Ala	Glu	Asp	Leu
1985						1990						1995		
Leu	Val	Asp	Thr	Cys	Ala	Pro	Asp	Gly	Ser	Leu	Asp	Leu	Glu	Lys
2000						2005						2010		
Ala	Arg	Ala	Gln	Gln	Leu	Glu	Leu	Glu	Glu	Gly	Leu	Ser	Ser	His
2015						2020						2025		
Gln	Pro	Ser	Leu	Ile	Lys	Val	Asn	Arg	Lys	Gly	Glu	Asp	Leu	Val
2030						2035						2040		
Gln	Arg	Leu	Arg	Pro	Ser	Glu	Ala	Ser	Phe	Leu	Lys	Glu	Lys	Leu
2045						2050						2055		
Ala	Gly	Phe	Asn	Gln	Arg	Trp	Ser	Thr	Leu	Val	Ala	Glu	Val	Glu
2060						2065						2070		
Ala	Leu	Gln	Pro	Arg	Leu	Lys	Gly	Glu	Ser	Gln	Gln	Val	Leu	Gly
2075						2080						2085		
Tyr	Lys	Arg	Arg	Leu	Asp	Glu	Val	Thr	Cys	Trp	Leu	Thr	Lys	Val
2090						2095						2100		
Glu	Ser	Ala	Val	Gln	Lys	Arg	Ser	Thr	Pro	Asp	Pro	Glu	Glu	Ser
2105						2110						2115		
Pro	Gln	Glu	Leu	Thr	Asp	Leu	Ala	Gln	Glu	Thr	Glu	Val	Gln	Ala
2120						2125						2130		
Glu	Asn	Ile	Lys	Trp	Leu	Asn	Arg	Ala	Glu	Leu	Glu	Met	Leu	Ser
2135						2140						2145		
Asp	Lys	Asn	Leu	Ser	Leu	Arg	Glu	Arg	Glu	Lys	Leu	Ser	Glu	Ser
2150						2155						2160		

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Leu	Arg	Asn	Val	Asn	Thr	Thr	Trp	Thr	Lys	Val	Cys	Arg	Glu	Val
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Pro	Ser	Leu	Leu	Lys	Thr	Arg	Thr	Gln	Asp	Pro	Cys	Ser	Ala	Pro
2180						2185					2190			
Gln	Met	Arg	Met	Ala	Ala	His	Pro	Asn	Val	Gln	Lys	Val	Val	Leu
2195						2200					2205			
Val	Ser	Ser	Ala	Ser	Asp	Ala	Pro	Leu	Arg	Gly	Gly	Leu	Glu	Ile
2210						2215					2220			
Ser	Val	Pro	Ala	Asp	Leu	Asp	Lys	Thr	Ile	Thr	Glu	Leu	Ala	Asp
2225						2230					2235			
Trp	Leu	Val	Leu	Ile	Asp	Gln	Met	Leu	Lys	Ser	Asn	Ile	Val	Thr
2240						2245					2250			
Val	Gly	Asp	Val	Lys	Glu	Ile	Asn	Lys	Thr	Val	Ser	Arg	Met	Lys
2255						2260					2265			
Ile	Thr	Lys	Ala	Asp	Leu	Glu	Gln	Arg	His	Pro	Gln	Leu	Asp	Cys
2270						2275					2280			
Val	Phe	Thr	Leu	Ala	Gln	Asn	Leu	Lys	Asn	Lys	Ala	Ser	Ser	Ser
2285						2290					2295			
Asp	Val	Arg	Thr	Ala	Ile	Thr	Glu	Lys	Leu	Glu	Lys	Leu	Lys	Thr
2300						2305					2310			
Gln	Trp	Glu	Ser	Thr	Gln	His	Gly	Val	Glu	Leu	Arg	Arg	Gln	Gln
2315						2320					2325			
Leu	Glu	Asp	Met	Val	Val	Asp	Ser	Leu	Gln	Trp	Asp	Asp	His	Arg
2330						2335					2340			
Glu	Glu	Thr	Glu	Glu	Leu	Met	Arg	Lys	Tyr	Glu	Ala	Arg	Phe	Tyr
2345						2350					2355			
Met	Leu	Gln	Gln	Ala	Arg	Arg	Asp	Pro	Leu	Ser	Lys	Gln	Val	Ser
2360						2365					2370			
Asp	Asn	Gln	Leu	Leu	Leu	Gln	Glu	Leu	Gly	Ser	Gly	Asp	Gly	Val
2375						2380					2385			
Ile	Met	Ala	Phe	Asp	Asn	Val	Leu	Gln	Lys	Leu	Leu	Glu	Glu	Tyr
2390						2395					2400			
Ser	Gly	Asp	Asp	Thr	Arg	Asn	Val	Glu	Glu	Thr	Thr	Glu	Tyr	Leu
2405						2410					2415			
Lys	Thr	Ser	Trp	Val	Asn	Leu	Lys	Gln	Ser	Ile	Ala	Asp	Arg	Gln
2420						2425					2430			
Ser	Ala	Leu	Glu	Ala	Glu	Leu	Gln	Thr	Val	Gln	Thr	Ser	Arg	Arg
2435						2440					2445			
Asp	Leu	Glu	Asn	Phe	Val	Lys	Trp	Leu	Gln	Glu	Ala	Glu	Thr	Thr
2450						2455					2460			
Ala	Asn	Val	Leu	Ala	Asp	Ala	Ser	Gln	Arg	Glu	Asn	Ala	Leu	Gln
2465						2470					2475			
Asp	Ser	Val	Leu	Ala	Arg	Gln	Leu	Arg	Gln	Gln	Met	Leu	Asp	Ile
2480						2485					2490			
Gln	Ala	Glu	Ile	Asp	Ala	His	Asn	Asp	Ile	Phe	Lys	Ser	Ile	Asp
2495						2500					2505			
Gly	Asn	Arg	Gln	Lys	Met	Val	Lys	Ala	Leu	Gly	Asn	Ser	Glu	Glu
2510						2515					2520			
Ala	Thr	Met	Leu	Gln	His	Arg	Leu	Asp	Asp	Met	Asn	Gln	Arg	Trp
2525						2530					2535			
Asn	Asp	Leu	Lys	Ala	Lys	Ser	Ala	Ser	Ile	Arg	Ala	His	Leu	Glu
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Ala	Ser	Ala	Glu	Lys	Trp	Asn	Arg	Leu	Leu	Ala	Ser	Leu	Glu	Glu

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2555	2560	2565
Leu Ile Lys Trp Leu Asn Met Lys Asp Glu Glu Leu Lys Lys Gln 2570 2575 2580		
Met Pro Ile Gly Gly Asp Val Pro Ala Leu Gln Leu Gln Tyr Asp 2585 2590 2595		
His Cys Lys Val Leu Arg Arg Glu Leu Lys Glu Lys Glu Tyr Ser 2600 2605 2610		
Val Leu Asn Ala Val Asp Gln Ala Arg Val Phe Leu Ala Asp Gln 2615 2620 2625		
Pro Ile Glu Ala Pro Glu Glu Pro Arg Arg Asn Pro Gln Ser Lys 2630 2635 2640		
Thr Glu Leu Thr Pro Glu Glu Arg Ala Gln Lys Ile Ala Lys Ala 2645 2650 2655		
Met Arg Lys Gln Ser Ser Glu Val Arg Glu Lys Trp Glu Asn Leu 2660 2665 2670		
Asn Ala Val Thr Ser Asn Trp Gln Lys Gln Val Gly Lys Ala Leu 2675 2680 2685		
Glu Lys Leu Arg Asp Leu Gln Gly Ala Met Asp Asp Leu Asp Ala 2690 2695 2700		
Asp Met Lys Glu Val Glu Ala Val Arg Asn Gly Trp Lys Pro Val 2705 2710 2715		
Gly Asp Leu Leu Ile Asp Ser Leu Gln Asp His Ile Glu Lys Thr 2720 2725 2730		
Leu Ala Phe Arg Glu Glu Ile Ala Pro Ile Asn Leu Lys Val Lys 2735 2740 2745		
Thr Met Asn Asp Leu Ser Ser Gln Leu Ser Pro Leu Asp Leu His 2750 2755 2760		
Pro Ser Leu Lys Met Ser Arg Gln Leu Asp Asp Leu Asn Met Arg 2765 2770 2775		
Trp Lys Leu Leu Gln Val Ser Val Asp Asp Arg Leu Lys Gln Leu 2780 2785 2790		
Gln Glu Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu 2795 2800 2805		
Ser Thr Ser Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn 2810 2815 2820		
Lys Val Pro Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp 2825 2830 2835		
Asp His Pro Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu 2840 2845 2850		
Asn Asn Val Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg 2855 2860 2865		
Arg Leu Gln Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Asn Thr 2870 2875 2880		
Thr Asn Glu Val Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln 2885 2890 2895		
Leu Leu Ser Val Pro Asp Val Ile Asn Cys Leu Thr Thr Thr Tyr 2900 2905 2910		
Asp Gly Leu Glu Gln Leu His Lys Asp Leu Val Asn Val Pro Leu 2915 2920 2925		
Cys Val Asp Met Cys Leu Asn Trp Leu Leu Asn Val Tyr Asp Thr 2930 2935 2940		
Gly Arg Thr Gly Lys Ile Arg Val Gln Ser Leu Lys Ile Gly Leu 2945 2950 2955		

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Met	Ser	Leu	Ser	Lys	Gly	Leu	Leu	Glu	Glu	Lys	Tyr	Arg	Cys	Leu
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Phe	Lys	Glu	Val	Ala	Gly	Pro	Thr	Glu	Met	Cys	Asp	Gln	Arg	Gln
2975						2980						2985		
Leu	Gly	Leu	Leu	Leu	His	Asp	Ala	Ile	Gln	Ile	Pro	Arg	Gln	Leu
2990						2995						3000		
Gly	Glu	Val	Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile	Glu	Pro	Ser	Val
3005						3010						3015		
Arg	Ser	Cys	Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro	Glu	Ile	Ser	Val
3020						3025						3030		
Lys	Glu	Phe	Ile	Asp	Trp	Met	His	Leu	Glu	Pro	Gln	Ser	Met	Val
3035						3040						3045		
Trp	Leu	Pro	Val	Leu	His	Arg	Val	Ala	Ala	Ala	Glu	Thr	Ala	Lys
3050						3055						3060		
His	Gln	Ala	Lys	Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly
3065						3070						3075		
Phe	Arg	Tyr	Arg	Ser	Leu	Lys	His	Phe	Asn	Tyr	Asp	Val	Cys	Gln
3080						3085						3090		
Ser	Cys	Phe	Phe	Ser	Gly	Arg	Thr	Ala	Lys	Gly	His	Lys	Leu	His
3095						3100						3105		
Tyr	Pro	Met	Val	Glu	Tyr	Cys	Ile	Pro	Thr	Thr	Ser	Gly	Glu	Asp
3110						3115						3120		
Val	Arg	Asp	Phe	Thr	Lys	Val	Leu	Lys	Asn	Lys	Phe	Arg	Ser	Lys
3125						3130						3135		
Lys	Tyr	Phe	Ala	Lys	His	Pro	Arg	Leu	Gly	Tyr	Leu	Pro	Val	Gln
3140						3145						3150		
Thr	Val	Leu	Glu	Gly	Asp	Asn	Leu	Glu	Thr	Pro	Ile	Thr	Leu	Ile
3155						3160						3165		
Ser	Met	Trp	Pro	Glu	His	Tyr	Asp	Pro	Ser	Gln	Ser	Pro	Gln	Leu
3170						3175						3180		
Phe	His	Asp	Asp	Thr	His	Ser	Arg	Ile	Glu	Gln	Tyr	Ala	Thr	Arg
3185						3190						3195		
Leu	Ala	Gln	Met	Glu	Arg	Thr	Asn	Gly	Ser	Phe	Leu	Thr	Asp	Ser
3200						3205						3210		
Ser	Ser	Thr	Thr	Gly	Ser	Val	Glu	Asp	Glu	His	Ala	Leu	Ile	Gln
3215						3220						3225		
Gln	Tyr	Cys	Gln	Thr	Leu	Gly	Gly	Glu	Ser	Pro	Val	Ser	Gln	Pro
3230						3235						3240		
Gln	Ser	Pro	Ala	Gln	Ile	Leu	Lys	Ser	Val	Glu	Arg	Glu	Glu	Arg
3245						3250						3255		
Gly	Glu	Leu	Glu	Arg	Ile	Ile	Ala	Asp	Leu	Glu	Glu	Glu	Gln	Arg
3260						3265						3270		
Asn	Leu	Gln	Val	Glu	Tyr	Glu	Gln	Leu	Lys	Glu	Gln	His	Leu	Arg
3275						3280						3285		
Arg	Gly	Leu	Pro	Val	Gly	Ser	Pro	Pro	Asp	Ser	Ile	Val	Ser	Pro
3290						3295						3300		
His	His	Thr	Ser	Glu	Asp	Ser	Glu	Leu	Ile	Ala	Glu	Ala	Lys	Leu
3305						3310						3315		
Leu	Arg	Gln	His	Lys	Gly	Arg	Leu	Glu	Ala	Arg	Met	Gln	Ile	Leu
3320						3325						3330		
Glu	Asp	His	Asn	Lys	Gln	Leu	Glu	Ser	Gln	Leu	His	Arg	Leu	Arg
3335						3340						3345		

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Gln Leu Leu Glu Gln Pro Asp Ser Asp Ser Arg Ile Asn Gly Val
 3350 3355 3360

Ser Pro Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu
 3365 3370 3375

Asp Thr Asp Pro Gly Pro Gln Phe His Gln Ala Ala Ser Glu Asp
 3380 3385 3390

Leu Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Asp Val
 3395 3400 3405

Met Glu Gln Ile Asn Ser Thr Phe Pro Ser Cys Ser Ser Asn Val
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Pro Ser Arg Pro Gln Ala Met
 3425 3430

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 <222> LOCATION: (1)..(4083)
 <220> FEATURE:
 <221> NAME/KEY: misc_feature
 <222> LOCATION: (1)..(117)
 <223> OTHER INFORMATION: TAT and epitope tag coding region

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cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac 96
 Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20 25 30

gtc cca gac tat gct ggc tcc atg gcc aag tat gga gaa cat gaa gcc 144
 Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
 35 40 45

agt cct gac aat ggg cag aac gaa ttc agt gat atc att aag tcc aga 192
 Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50 55 60

tct gat gaa cac aat gac gta cag aag aaa acc ttt acc aaa tgg ata 240
 Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65 70 75 80

aat gct cga ttt tca aag agt ggg aaa cca ccc atc aat gat atg ttc 288
 Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
 85 90 95

aca gac ctc aaa gat gga agg aag cta ttg gat ctt cta gaa ggc ctc 336
 Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100 105 110

aca gga aca tca ctg cca aag gaa cgt ggt tcc aca agg gta cat gcc 384
 Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115 120 125

tta aat aac gtc aac aga gtg ctg cag gtt tta cat cag aac aat gtg 432
 Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
 130 135 140

gaa tta gtg aat ata ggg gga act gac att gtg gat gga aat cac aaa 480
 Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
 145 150 155 160

ctg act ttg ggg tta ctt tgg agc atc att ttg cac tgg cag gtg aaa 528
 Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
 165 170 175

gat gtc atg aag gat gtc atg tcg gac ctg cag cag acg aac agt gag 576
 Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu

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Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu		
			500					505				510					
cag	gtg	aaa	gta	aat	tca	cta	act	cac	atg	gtg	gtc	att	gtt	gat	gaa	1584	
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu		
		515					520					525					
aac	agt	ggt	gag	agt	gct	aca	gct	atc	cta	gaa	gac	cag	tta	cag	aaa	1632	
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys		
		530					535					540					
ctt	ggt	gag	cgc	tgg	aca	gca	gta	tgc	cgt	tgg	act	gaa	gaa	cgc	tgg	1680	
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp		
		545			550					555					560		
aat	agg	tta	caa	gaa	atc	aat	ata	ttg	tgg	cag	gaa	tta	ttg	gaa	gaa	1728	
Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu		
			565						570						575		
cag	tgc	ttg	ttg	aaa	gct	tgg	tta	acc	gaa	aaa	gaa	gag	gct	tta	aat	1776	
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn		
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aaa	gtc	cag	aca	agc	aac	ttc	aaa	gac	caa	aag	gaa	cta	agt	gtc	agt	1824	
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser		
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gtt	cga	cgt	ctg	gct	att	ttg	aag	gaa	gac	atg	gaa	atg	aag	cgt	caa	1872	
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln		
		610			615					620							
aca	ttg	gat	cag	ctg	agt	gag	att	ggc	cag	gat	gtg	gga	caa	tta	ctt	1920	
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu		
		625			630					635					640		
gat	aat	tcc	aag	gca	tct	aag	aag	atc	aac	agt	gac	tca	gag	gaa	ctg	1968	
Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu		
			645					650							655		
act	caa	aga	tgg	gat	tct	ttg	gtt	cag	aga	cta	gaa	gat	tcc	tcc	aac	2016	
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn		
			660					665							670		
cag	gtg	act	cag	gct	gta	gca	aag	ctg	ggg	atg	tct	cag	att	cct	cag	2064	
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln		
		675					680								685		
aag	gac	ctt	ttg	gag	act	gtt	cgt	gta	aga	gaa	caa	gca	att	aca	aaa	2112	
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys		
		690				695					700						
aaa	tct	aag	cag	gaa	ctg	cct	cct	cct	cct	ccc	cca	aag	aag	aga	cag	2160	
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln		
		705			710					715					720		
atc	cat	gtg	gat	gcc	cac	aga	gat	ttt	gga	cca	tcc	tct	cag	cat	ttt	2208	
Ile	His	Val	Asp	Ala	His	Arg	Asp	Phe	Gly	Pro	Ser	Ser	Gln	His	Phe		
			725					730							735		
ctc	tct	acg	tca	gtc	cag	ctg	cgg	tgg	caa	aga	tcc	att	tca	cat	aat	2256	
Leu	Ser	Thr	Ser	Val	Gln	Leu	Pro	Trp	Gln	Arg	Ser	Ile	Ser	His	Asn		
			740					745							750		
aaa	gtg	ccc	tat	tac	atc	aac	cat	caa	aca	cag	acc	acc	tgt	tgg	gac	2304	
Lys	Val	Pro	Tyr	Tyr	Ile	Asn	His	Gln	Thr	Gln	Thr	Thr	Cys	Trp	Asp		
		755				760									765		
cat	cct	aaa	atg	acc	gaa	ctc	ttt	caa	tcc	ctt	gct	gac	ctg	aat	aat	2352	
His	Pro	Lys	Met	Thr	Glu	Leu	Phe	Gln	Ser	Leu	Ala	Asp	Leu	Asn	Asn		
		770				775									780		
gta	cgt	ttt	tct	gcc	tac	cgt	aca	gca	atc	aaa	atc	cga	aga	cta	caa	2400	
Val	Arg	Phe	Ser	Ala	Tyr	Arg	Thr	Ala	Ile	Lys	Ile	Arg	Arg	Leu	Gln		
		785			790					795					800		
aaa	gca	cta	tgt	ttg	gat	ctc	tta	gag	ttg	agt	aca	aca	aat	gaa	att	2448	
Lys	Ala	Leu	Cys	Leu	Asp	Leu	Leu	Glu	Leu	Ser	Thr	Thr	Asn	Glu	Ile		
				805					810						815		

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ttc aaa cag cac aag ttg aac caa aat gac cag ctc ctc agt gtt cca	2496
Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu Ser Val Pro	
820 825 830	
gat gtc atc aac tgt ctg aca aca act tat gat gga ctt gag caa atg	2544
Asp Val Ile Asn Cys Leu Thr Thr Tyr Asp Gly Leu Glu Gln Met	
835 840 845	
cat aag gac ctg gtc aac gtt cca ctc tgt gtt gat atg tgt ctc aat	2592
His Lys Asp Leu Val Asn Val Pro Leu Cys Val Asp Met Cys Leu Asn	
850 855 860	
tgg ttg ctc aat gtc tat gac acg ggt cga act gga aaa att aga gtg	2640
Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg Thr Gly Lys Ile Arg Val	
865 870 875 880	
cag agt ctg aag att gga tta atg tct ctc tcc aaa ggt ctc ttg gaa	2688
Gln Ser Leu Lys Ile Gly Leu Met Ser Leu Ser Lys Gly Leu Leu Glu	
885 890 895	
gaa aaa tac aga tat ctc ttt aag gaa gtt gca ggg cca aca gaa atg	2736
Glu Lys Tyr Arg Tyr Leu Phe Lys Glu Val Ala Gly Pro Thr Glu Met	
900 905 910	
tgt gac cag agg cag ctg ggc ctg tta ctt cat gat gcc atc cag atc	2784
Cys Asp Gln Arg Gln Leu Gly Leu Leu Leu His Asp Ala Ile Gln Ile	
915 920 925	
ccc cgg cag cta ggt gaa gta gca gct ttt gga ggc agt aat att gag	2832
Pro Arg Gln Leu Gly Glu Val Ala Ala Phe Gly Gly Ser Asn Ile Glu	
930 935 940	
cct agt gtt cgc agc tgc ttc caa cag aat aac aat aaa cca gaa ata	2880
Pro Ser Val Arg Ser Cys Phe Gln Gln Asn Asn Asn Lys Pro Glu Ile	
945 950 955 960	
agt gtg aaa gag ttt ata gat tgg atg cat ttg gaa cca cag tcc atg	2928
Ser Val Lys Glu Phe Ile Asp Trp Met His Leu Glu Pro Gln Ser Met	
965 970 975	
gtt tgg ctc cca gtt tta cat cga gtg gca gca gcg gag act gca aaa	2976
Val Trp Leu Pro Val Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys	
980 985 990	
cat cag gcc aaa tgc aac atc tgt aaa gaa tgt cca att gtc ggg ttc	3024
His Gln Ala Lys Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe	
995 1000 1005	
agg tat aga agc ctt aag cat ttt aac tat gat gtc tgc cag agt	3069
Arg Tyr Arg Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser	
1010 1015 1020	
tgt ttc ttt tcg ggt cga aca gca aaa ggt cac aaa tta cat tac	3114
Cys Phe Phe Ser Gly Arg Thr Ala Lys Gly His Lys Leu His Tyr	
1025 1030 1035	
cca atg gtg gaa tat tgt ata cct aca aca tct ggg gaa gat gta	3159
Pro Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val	
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cga gac ttc aca aag gta ctt aag aac aag ttc agg tcg aag aag	3204
Arg Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys	
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tac ttt gcc aaa cac cct cga ctt ggt tac ctg cct gtc cag aca	3249
Tyr Phe Ala Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr	
1070 1075 1080	
gtt ctt gaa ggt gac aac tta gag act cct atc aca ctc atc agt	3294
Val Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser	
1085 1090 1095	
atg tgg cca gag cac tat gac ccc tca caa tct cct caa ctg ttt	3339
Met Trp Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe	
1100 1105 1110	
cat gat gac acc cat tca aga ata gaa caa tat gcc aca cga ctg	3384
His Asp Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu	
1115 1120 1125	

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gcc cag atg gaa agg act aat ggg tct ttt ctc act gat agc agc 3429
Ala Gln Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser
1130 1135 1140

tcc acc aca gga agt gtg gaa gac gag cac gcc ctc atc cag cag 3474
Ser Thr Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln
1145 1150 1155

tat tgc caa aca ctc gga gga gag tcc cca gtg agc cag ccg cag 3519
Tyr Cys Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln
1160 1165 1170

agc cca gct cag atc ctg aag tca gta gag agg gaa gaa cgt gga 3564
Ser Pro Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly
1175 1180 1185

gaa ctg gag agg atc att gct gac ctg gag gaa gaa caa aga aat 3609
Glu Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn
1190 1195 1200

cta cag gtg gag tat gag cag ctg aag gac cag cac ctc cga agg 3654
Leu Gln Val Glu Tyr Glu Gln Leu Lys Asp Gln His Leu Arg Arg
1205 1210 1215

ggg ctc cct gtc ggt tca ccg cca gag tcg att ata tct ccc cat 3699
Gly Leu Pro Val Gly Ser Pro Glu Ser Ile Ile Ser Pro His
1220 1225 1230

cac acg tct gag gat tca gaa ctt ata gca gaa gca aaa ctc ctc 3744
His Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu
1235 1240 1245

agg cag cac aaa ggt cgg ctg gag gct agg atg cag att tta gaa 3789
Arg Gln His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu
1250 1255 1260

gat cac aat aaa cag ctg gag tct cag ctc cac cgc ctc cga cag 3834
Asp His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln
1265 1270 1275

ctg ctg gag cag cct gaa tct gat tcc cga atc aat ggt gtt tcc 3879
Leu Leu Glu Gln Pro Glu Ser Asp Ser Arg Ile Asn Gly Val Ser
1280 1285 1290

cca tgg gct tct cct cag cat tct gca ctg agc tac tcg ctt gat 3924
Pro Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp
1295 1300 1305

cca gat gcc tcc ggc cca cag ttc cac cag gca gcg gga gag gac 3969
Pro Asp Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp
1310 1315 1320

ctg ctg gcc cca ccg cac gac acc agc acg gat ctc acg gag gtc 4014
Leu Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val
1325 1330 1335

atg gag cag att cac agc acg ttt cca tct tgc tgc cca aat gtt 4059
Met Glu Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val
1340 1345 1350

ccc agc agg cca cag gca atg tga 4083
Pro Ser Arg Pro Gln Ala Met
1355 1360

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<210> SEQ ID NO 11
<211> LENGTH: 1360
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

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<400> SEQUENCE: 11

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1 5 10 15
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
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Val	Pro	Asp	Tyr	Ala	Gly	Ser	Met	Ala	Lys	Tyr	Gly	Glu	His	Glu	Ala
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Ser	Pro	Asp	Asn	Gly	Gln	Asn	Glu	Phe	Ser	Asp	Ile	Ile	Lys	Ser	Arg
	50					55					60				
Ser	Asp	Glu	His	Asn	Asp	Val	Gln	Lys	Lys	Thr	Phe	Thr	Lys	Trp	Ile
65				70						75					80
Asn	Ala	Arg	Phe	Ser	Lys	Ser	Gly	Lys	Pro	Pro	Ile	Asn	Asp	Met	Phe
			85						90					95	
Thr	Asp	Leu	Lys	Asp	Gly	Arg	Lys	Leu	Leu	Asp	Leu	Leu	Glu	Gly	Leu
		100						105					110		
Thr	Gly	Thr	Ser	Leu	Pro	Lys	Glu	Arg	Gly	Ser	Thr	Arg	Val	His	Ala
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Leu	Asn	Asn	Val	Asn	Arg	Val	Leu	Gln	Val	Leu	His	Gln	Asn	Asn	Val
	130					135					140				
Glu	Leu	Val	Asn	Ile	Gly	Gly	Thr	Asp	Ile	Val	Asp	Gly	Asn	His	Lys
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Leu	Thr	Leu	Gly	Leu	Leu	Trp	Ser	Ile	Ile	Leu	His	Trp	Gln	Val	Lys
			165						170					175	
Asp	Val	Met	Lys	Asp	Val	Met	Ser	Asp	Leu	Gln	Gln	Thr	Asn	Ser	Glu
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Lys	Ile	Leu	Leu	Ser	Trp	Val	Arg	Gln	Thr	Thr	Arg	Pro	Tyr	Ser	Gln
		195					200					205			
Val	Asn	Val	Leu	Asn	Phe	Thr	Thr	Ser	Trp	Thr	Asp	Gly	Leu	Ala	Phe
	210					215					220				
Asn	Ala	Val	Leu	His	Arg	His	Lys	Pro	Asp	Leu	Phe	Ser	Trp	Asp	Lys
225					230					235					240
Val	Val	Lys	Met	Ser	Pro	Ile	Glu	Arg	Leu	Glu	His	Ala	Phe	Ser	Lys
			245						250					255	
Ala	Gln	Thr	Tyr	Leu	Gly	Ile	Glu	Lys	Leu	Leu	Asp	Pro	Glu	Asp	Val
			260						265				270		
Ala	Val	Gln	Leu	Pro	Asp	Lys	Lys	Ser	Ile	Ile	Met	Tyr	Leu	Thr	Ser
		275					280					285			
Leu	Phe	Glu	Val	Leu	Pro	Gln	Gln	Val	Thr	Ile	Asp	Ala	Ile	Arg	Glu
	290					295					300				
Val	Glu	Thr	Leu	Pro	Arg	Lys	Tyr	Lys	Lys	Glu	Cys	Glu	Glu	Glu	Ala
305					310					315					320
Ile	Asn	Ile	Gln	Ser	Thr	Ala	Pro	Glu	Glu	Glu	His	Glu	Ser	Pro	Arg
			325						330					335	
Ala	Glu	Thr	Pro	Ser	Thr	Val	Thr	Glu	Val	Asp	Met	Asp	Leu	Asp	Ser
			340					345					350		
Tyr	Gln	Ile	Ala	Leu	Glu	Glu	Val	Leu	Thr	Trp	Leu	Leu	Ser	Ala	Glu
		355					360						365		
Asp	Thr	Phe	Gln	Glu	Gln	Asp	Asp	Ile	Ser	Asp	Asp	Val	Glu	Glu	Val
	370					375					380				
Lys	Asp	Gln	Phe	Ala	Thr	His	Glu	Ala	Phe	Met	Met	Glu	Leu	Thr	Ala
385					390					395					400
His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Ile
			405						410					415	
Thr	Gln	Gly	Thr	Leu	Ser	Asp	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln
			420					425					430		
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met
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Gln	Leu	Gln	Gln	Leu	Ser	Ala	Trp	Leu	Thr	Leu	Thr	Glu	Glu	Arg	Ile
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Gln	Lys	Met	Glu	Thr	Cys	Pro	Leu	Asp	Asp	Asp	Val	Lys	Ser	Leu	Gln
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Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu
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Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu
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Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys
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Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp
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Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu
			565					570						575	
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn
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Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser
		595					600					605			
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln
	610					615					620				
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu
625				630						635					640
Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu
			645					650						655	
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn
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Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln
		675					680					685			
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys
	690					695				700					
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln
705				710						715					720
Ile	His	Val	Asp	Ala	His	Arg	Asp	Phe	Gly	Pro	Ser	Ser	Gln	His	Phe
			725					730						735	
Leu	Ser	Thr	Ser	Val	Gln	Leu	Pro	Trp	Gln	Arg	Ser	Ile	Ser	His	Asn
			740					745				750			
Lys	Val	Pro	Tyr	Tyr	Ile	Asn	His	Gln	Thr	Gln	Thr	Thr	Cys	Trp	Asp
		755					760					765			
His	Pro	Lys	Met	Thr	Glu	Leu	Phe	Gln	Ser	Leu	Ala	Asp	Leu	Asn	Asn
	770					775					780				
Val	Arg	Phe	Ser	Ala	Tyr	Arg	Thr	Ala	Ile	Lys	Ile	Arg	Arg	Leu	Gln
785				790						795					800
Lys	Ala	Leu	Cys	Leu	Asp	Leu	Leu	Glu	Leu	Ser	Thr	Thr	Asn	Glu	Ile
			805					810						815	
Phe	Lys	Gln	His	Lys	Leu	Asn	Gln	Asn	Asp	Gln	Leu	Leu	Ser	Val	Pro
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Asp	Val	Ile	Asn	Cys	Leu	Thr	Thr	Thr	Tyr	Asp	Gly	Leu	Glu	Gln	Met
		835					840					845			
His	Lys	Asp	Leu	Val	Asn	Val	Pro	Leu	Cys	Val	Asp	Met	Cys	Leu	Asn
	850					855					860				
Trp	Leu	Leu	Asn	Val	Tyr	Asp	Thr	Gly	Arg	Thr	Gly	Lys	Ile	Arg	Val
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Gln Ser Leu Lys Ile Gly Leu Met Ser Leu Ser Lys Gly Leu Leu Glu
 885 890 895

Glu Lys Tyr Arg Tyr Leu Phe Lys Glu Val Ala Gly Pro Thr Glu Met
 900 905 910

Cys Asp Gln Arg Gln Leu Gly Leu Leu Leu His Asp Ala Ile Gln Ile
 915 920 925

Pro Arg Gln Leu Gly Glu Val Ala Ala Phe Gly Gly Ser Asn Ile Glu
 930 935 940

Pro Ser Val Arg Ser Cys Phe Gln Gln Asn Asn Asn Lys Pro Glu Ile
 945 950 955 960

Ser Val Lys Glu Phe Ile Asp Trp Met His Leu Glu Pro Gln Ser Met
 965 970 975

Val Trp Leu Pro Val Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys
 980 985 990

His Gln Ala Lys Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe
 995 1000 1005

Arg Tyr Arg Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser
 1010 1015 1020

Cys Phe Phe Ser Gly Arg Thr Ala Lys Gly His Lys Leu His Tyr
 1025 1030 1035

Pro Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val
 1040 1045 1050

Arg Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys
 1055 1060 1065

Tyr Phe Ala Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr
 1070 1075 1080

Val Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser
 1085 1090 1095

Met Trp Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe
 1100 1105 1110

His Asp Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu
 1115 1120 1125

Ala Gln Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser
 1130 1135 1140

Ser Thr Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln
 1145 1150 1155

Tyr Cys Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln
 1160 1165 1170

Ser Pro Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly
 1175 1180 1185

Glu Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn
 1190 1195 1200

Leu Gln Val Glu Tyr Glu Gln Leu Lys Asp Gln His Leu Arg Arg
 1205 1210 1215

Gly Leu Pro Val Gly Ser Pro Pro Glu Ser Ile Ile Ser Pro His
 1220 1225 1230

His Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu
 1235 1240 1245

Arg Gln His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu
 1250 1255 1260

Asp His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln
 1265 1270 1275

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Leu Leu Glu Gln Pro Glu Ser Asp Ser Arg Ile Asn Gly Val Ser
 1280 1285 1290

Pro Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp
 1295 1300 1305

Pro Asp Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp
 1310 1315 1320

Leu Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val
 1325 1330 1335

Met Glu Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val
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Pro Ser Arg Pro Gln Ala Met
 1355 1360

<210> SEQ ID NO 12
 <211> LENGTH: 5070
 <212> TYPE: DNA
 <213> ORGANISM: Homo sapiens
 <220> FEATURE:
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 <222> LOCATION: (1)..(5070)
 <220> FEATURE:
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 <222> LOCATION: (1)..(117)
 <223> OTHER INFORMATION: TAT and epitope tag coding sequence

<400> SEQUENCE: 12

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 Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20 25 30

gtc cca gac tat gct ggc tcc atg gcc aag tat gga gaa cat gaa gcc 144
 Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
 35 40 45

agt cct gac aat ggg cag aac gaa ttc agt gat atc att aag tcc aga 192
 Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50 55 60

tct gat gaa cac aat gac gta cag aag aaa acc ttt acc aaa tgg ata 240
 Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65 70 75 80

aat gct cga ttt tca aag agt ggg aaa cca ccc atc aat gat atg ttc 288
 Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
 85 90 95

aca gac ctc aaa gat gga agg aag cta ttg gat ctt cta gaa ggc ctc 336
 Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100 105 110

aca gga aca tca ctg cca aag gaa cgt ggt tcc aca agg gta cat gcc 384
 Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115 120 125

tta aat aac gtc aac aga gtg ctg cag gtt tta cat cag aac aat gtg 432
 Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
 130 135 140

gaa tta gtg aat ata ggg gga act gac att gtg gat gga aat cac aaa 480
 Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
 145 150 155 160

ctg act ttg ggg tta ctt tgg agc atc att ttg cac tgg cag gtg aaa 528
 Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
 165 170 175

gat gtc atg aag gat gtc atg tcg gac ctg cag cag acg aac agt gag 576
 Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu

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Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu		
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Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu		
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aac	agt	ggg	gag	agt	gct	aca	gct	atc	cta	gaa	gac	cag	tta	cag	aaa	1632	
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys		
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Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp		
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Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu		
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cag	tgc	ttg	ttg	aaa	gct	tgg	tta	acc	gaa	aaa	gaa	gag	gct	tta	aat	1776	
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn		
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Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser		
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gtt	cga	cgt	ctg	gct	att	ttg	aag	gaa	gac	atg	gaa	atg	aag	cgt	caa	1872	
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln		
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Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu		
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Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu		
			645					650						655			
act	caa	aga	tgg	gat	tct	ttg	ggt	cag	aga	cta	gaa	gat	tcc	tcc	aac	2016	
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn		
			660					665					670				
cag	gtg	act	cag	gct	gta	gca	aag	ctg	ggg	atg	tct	cag	att	cct	cag	2064	
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln		
		675					680					685					
aag	gac	ctt	ttg	gag	act	gtt	cgt	gta	aga	gaa	caa	gca	att	aca	aaa	2112	
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys		
	690					695					700						
aaa	tct	aag	cag	gaa	ctg	cct	cct	cct	cct	ccc	cca	aag	aag	aga	cag	2160	
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln		
	705				710					715					720		
atc	cat	gtg	gat	att	gaa	gct	aag	aaa	aag	ttt	gat	gct	ata	agt	gca	2208	
Ile	His	Val	Asp	Ile	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Ala		
			725							730					735		
gag	ctg	ttg	aac	tgg	att	ttg	aaa	tgg	aaa	act	gcc	att	cag	acc	aca	2256	
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Trp	Lys	Thr	Ala	Ile	Gln	Thr	Thr		
			740					745					750				
gag	ata	aaa	gag	tat	atg	aag	atg	caa	gac	act	tcc	gaa	atg	aaa	aag	2304	
Glu	Ile	Lys	Glu	Tyr	Met	Lys	Met	Gln	Asp	Thr	Ser	Glu	Met	Lys	Lys		
		755					760					765					
aag	ttg	aag	gca	tta	gaa	aaa	gaa	cag	aga	gaa	aga	atc	ccc	aga	gca	2352	
Lys	Leu	Lys	Ala	Leu	Glu	Lys	Glu	Gln	Arg	Glu	Arg	Ile	Pro	Arg	Ala		
		770				775						780					
gat	gaa	tta	aac	caa	act	gga	caa	atc	ctt	gtg	gag	caa	atg	gga	aaa	2400	
Asp	Glu	Leu	Asn	Gln	Thr	Gly	Gln	Ile	Leu	Val	Glu	Gln	Met	Gly	Lys		
	785				790					795					800		
gaa	ggc	ctt	cct	act	gaa	gaa	ata	aaa	aat	gtt	ctg	gag	aag	gtt	tca	2448	
Glu	Gly	Leu	Pro	Thr	Glu	Glu	Ile	Lys	Asn	Val	Leu	Glu	Lys	Val	Ser		
				805						810					815		

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tca gaa tgg aag aat gta tct caa cat ttg gaa gat cta gaa aga aag	2496
Ser Glu Trp Lys Asn Val Ser Gln His Leu Glu Asp Leu Glu Arg Lys	
820 825 830	
att cag cta cag gaa gat ata aat gct tat ttc aag cag ctt gat gag	2544
Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Glu	
835 840 845	
ctt gaa aag gtc atc aag aca aag gag gag tgg gta aaa cac act tcc	2592
Leu Glu Lys Val Ile Lys Thr Lys Glu Glu Trp Val Lys His Thr Ser	
850 855 860	
att tct gaa tct tcc cgg cag tcc ttg cca agc ttg aag gat tcc tgt	2640
Ile Ser Glu Ser Ser Arg Gln Ser Leu Pro Ser Leu Lys Asp Ser Cys	
865 870 875 880	
cag cgg gaa ttg aca aat ctt ctt ggc ctt cac ccc aaa att gaa atg	2688
Gln Arg Glu Leu Thr Asn Leu Leu Gly Leu His Pro Lys Ile Glu Met	
885 890 895	
gct cgt gca agc tgc tcg gcc ctg atg tct cag cct tct gcc cca gat	2736
Ala Arg Ala Ser Cys Ser Ala Leu Met Ser Gln Pro Ser Ala Pro Asp	
900 905 910	
ttt gtc cag cgg ggc ttc gat agc ttt ctg ggc cgc tac caa gct gta	2784
Phe Val Gln Arg Gly Phe Asp Ser Phe Leu Gly Arg Tyr Gln Ala Val	
915 920 925	
caa gag gct gta gag gat cgt caa caa cat cta gag aat gaa ctg aag	2832
Gln Glu Ala Val Glu Asp Arg Gln Gln His Leu Glu Asn Glu Leu Lys	
930 935 940	
ggc caa cct gga cat gca tat ctg gaa aca ttg aaa aca ctg aaa gat	2880
Gly Gln Pro Gly His Ala Tyr Leu Glu Thr Leu Lys Thr Leu Lys Asp	
945 950 955 960	
gtg cta aat gat tca gaa aat aag gcc cag gtg tct ctg aat gtc ctt	2928
Val Leu Asn Asp Ser Glu Asn Lys Ala Gln Val Ser Leu Asn Val Leu	
965 970 975	
aat gat ctt gcc aag gtg gag aag gcc ctg caa gaa aaa aag acc ctt	2976
Asn Asp Leu Ala Lys Val Glu Lys Ala Leu Gln Glu Lys Lys Thr Leu	
980 985 990	
gat gaa atc ctt gag aat cag aaa cct gca tta cat aaa ctt gca gaa	3024
Asp Glu Ile Leu Glu Asn Gln Lys Pro Ala Leu His Lys Leu Ala Glu	
995 1000 1005	
gaa aca aag gct ctg gag aaa aat gtt cat cct gat gta gaa aaa	3069
Glu Thr Lys Ala Leu Glu Lys Asn Val His Pro Asp Val Glu Lys	
1010 1015 1020	
tta tat aag caa gaa ttt gat gat gtg caa gga aag tgg aac aag	3114
Leu Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Lys Trp Asn Lys	
1025 1030 1035	
cta aag gtc ttg gtt tcc aaa gat cta cat ttg ctt gag gaa att	3159
Leu Lys Val Leu Val Ser Lys Asp Leu His Leu Leu Glu Glu Ile	
1040 1045 1050	
gcc cac aga gat ttt gga cca tcc tct cag cat ttt ctc tct acg	3204
Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr	
1055 1060 1065	
tca gtc cag ctg cgg tgg caa aga tcc att tca cat aat aaa gtg	3249
Ser Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn Lys Val	
1070 1075 1080	
ccc tat tac atc aac cat caa aca cag acc acc tgt tgg gac cat	3294
Pro Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp His	
1085 1090 1095	
cct aaa atg acc gaa ctc ttt caa tcc ctt gct gac ctg aat aat	3339
Pro Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn	
1100 1105 1110	
gta cgt ttt tct gcc tac cgt aca gca atc aaa atc cga aga cta	3384
Val Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu	
1115 1120 1125	

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caa aaa gca cta tgt ttg gat ctc tta gag ttg agt aca aca aat	3429
Gln Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Ser Thr Thr Asn	
1130 1135 1140	
gaa att ttc aaa cag cac aag ttg aac caa aat gac cag ctc ctc	3474
Glu Ile Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu	
1145 1150 1155	
agt gtt cca gat gtc atc aac tgt ctg aca aca act tat gat gga	3519
Ser Val Pro Asp Val Ile Asn Cys Leu Thr Thr Thr Tyr Asp Gly	
1160 1165 1170	
ctt gag caa atg cat aag gac ctg gtc aac gtt cca ctc tgt gtt	3564
Leu Glu Gln Met His Lys Asp Leu Val Asn Val Pro Leu Cys Val	
1175 1180 1185	
gat atg tgt ctc aat tgg ttg ctc aat gtc tat gac acg ggt cga	3609
Asp Met Cys Leu Asn Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg	
1190 1195 1200	
act gga aaa att aga gtg cag agt ctg aag att gga tta atg tct	3654
Thr Gly Lys Ile Arg Val Gln Ser Leu Lys Ile Gly Leu Met Ser	
1205 1210 1215	
ctc tcc aaa ggt ctc ttg gaa gaa aaa tac aga tat ctc ttt aag	3699
Leu Ser Lys Gly Leu Leu Glu Glu Lys Tyr Arg Tyr Leu Phe Lys	
1220 1225 1230	
gaa gtt gca ggg cca aca gaa atg tgt gac cag agg cag ctg ggc	3744
Glu Val Ala Gly Pro Thr Glu Met Cys Asp Gln Arg Gln Leu Gly	
1235 1240 1245	
ctg tta ctt cat gat gcc atc cag atc ccc cgg cag cta ggt gaa	3789
Leu Leu Leu His Asp Ala Ile Gln Ile Pro Arg Gln Leu Gly Glu	
1250 1255 1260	
gta gca gct ttt gga ggc agt aat att gag cct agt gtt cgc agc	3834
Val Ala Ala Phe Gly Gly Ser Asn Ile Glu Pro Ser Val Arg Ser	
1265 1270 1275	
tgc ttc caa cag aat aac aat aaa cca gaa ata agt gtg aaa gag	3879
Cys Phe Gln Gln Asn Asn Asn Lys Pro Glu Ile Ser Val Lys Glu	
1280 1285 1290	
ttt ata gat tgg atg cat ttg gaa cca cag tcc atg gtt tgg ctc	3924
Phe Ile Asp Trp Met His Leu Glu Pro Gln Ser Met Val Trp Leu	
1295 1300 1305	
cca gtt tta cat cga gtg gca gca gcg gag act gca aaa cat cag	3969
Pro Val Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys His Gln	
1310 1315 1320	
gcc aaa tgc aac atc tgt aaa gaa tgt cca att gtc ggg ttc agg	4014
Ala Lys Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe Arg	
1325 1330 1335	
tat aga agc ctt aag cat ttt aac tat gat gtc tgc cag agt tgt	4059
Tyr Arg Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser Cys	
1340 1345 1350	
ttc ttt tcg ggt cga aca gca aaa ggt cac aaa tta cat tac cca	4104
Phe Phe Ser Gly Arg Thr Ala Lys Gly His Lys Leu His Tyr Pro	
1355 1360 1365	
atg gtg gaa tat tgt ata cct aca aca tct ggg gaa gat gta cga	4149
Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val Arg	
1370 1375 1380	
gac ttc aca aag gta ctt aag aac aag ttc agg tcg aag aag tac	4194
Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys Tyr	
1385 1390 1395	
ttt gcc aaa cac cct cga ctt ggt tac ctg cct gtc cag aca gtt	4239
Phe Ala Lys His Pro Arg Leu Leu Gly Tyr Leu Pro Val Gln Thr Val	
1400 1405 1410	
ctt gaa ggt gac aac tta gag act cct atc aca ctc atc agt atg	4284
Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser Met	

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1415	1420	1425	
tgg cca gag cac tat gac ccc tca caa tct cct caa ctg ttt cat			4329
Trp Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe His			
1430	1435	1440	
gat gac acc cat tca aga ata gaa caa tat gcc aca cga ctg gcc			4374
Asp Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu Ala			
1445	1450	1455	
cag atg gaa agg act aat ggg tct ttt ctc act gat agc agc tcc			4419
Gln Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser Ser			
1460	1465	1470	
acc aca gga agt gtg gaa gac gag cac gcc ctc atc cag cag tat			4464
Thr Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln Tyr			
1475	1480	1485	
tgc caa aca ctc gga gga gag tcc cca gtg agc cag ccg cag agc			4509
Cys Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln Ser			
1490	1495	1500	
cca gct cag atc ctg aag tca gta gag agg gaa gaa cgt gga gaa			4554
Pro Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly Glu			
1505	1510	1515	
ctg gag agg atc att gct gac ctg gag gaa gaa caa aga aat cta			4599
Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn Leu			
1520	1525	1530	
cag gtg gag tat gag cag ctg aag gac cag cac ctc cga agg ggg			4644
Gln Val Glu Tyr Glu Gln Leu Lys Asp Gln His Leu Arg Arg Gly			
1535	1540	1545	
ctc cct gtc ggt tca ccg cca gag tcg att ata tct ccc cat cac			4689
Leu Pro Val Gly Ser Pro Pro Glu Ser Ile Ile Ser Pro His His			
1550	1555	1560	
acg tct gag gat tca gaa ctt ata gca gaa gca aaa ctc ctc agg			4734
Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg			
1565	1570	1575	
cag cac aaa ggt cgg ctg gag gct agg atg cag att tta gaa gat			4779
Gln His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp			
1580	1585	1590	
cac aat aaa cag ctg gag tct cag ctc cac cgc ctc cga cag ctg			4824
His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu			
1595	1600	1605	
ctg gag cag cct gaa tct gat tcc cga atc aat ggt gtt tcc cca			4869
Leu Glu Gln Pro Glu Ser Asp Ser Arg Ile Asn Gly Val Ser Pro			
1610	1615	1620	
tgg gct tct cct cag cat tct gca ctg agc tac tcg ctt gat cca			4914
Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Pro			
1625	1630	1635	
gat gcc tcc ggc cca cag ttc cac cag gca gcg gga gag gac ctg			4959
Asp Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp Leu			
1640	1645	1650	
ctg gcc cca ccg cac gac acc agc acg gat ctc acg gag gtc atg			5004
Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val Met			
1655	1660	1665	
gag cag att cac agc acg ttt cca tct tgc tgc cca aat gtt ccc			5049
Glu Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val Pro			
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agc agg cca cag gca atg tga			5070
Ser Arg Pro Gln Ala Met			
1685			

<210> SEQ ID NO 13

<211> LENGTH: 1689

<212> TYPE: PRT

<213> ORGANISM: Homo sapiens

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<400> SEQUENCE: 13

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Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
20 25 30
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
35 40 45
Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
50 55 60
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
65 70 75 80
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
85 90 95
Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
100 105 110
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
115 120 125
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
130 135 140
Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
145 150 155 160
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
165 170 175
Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu
180 185 190
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln
195 200 205
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe
210 215 220
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Ser Trp Asp Lys
225 230 235 240
Val Val Lys Met Ser Pro Ile Glu Arg Leu Glu His Ala Phe Ser Lys
245 250 255
Ala Gln Thr Tyr Leu Gly Ile Glu Lys Leu Leu Asp Pro Glu Asp Val
260 265 270
Ala Val Gln Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser
275 280 285
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu
290 295 300
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Ala
305 310 315 320
Ile Asn Ile Gln Ser Thr Ala Pro Glu Glu Glu His Glu Ser Pro Arg
325 330 335
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser
340 345 350
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu
355 360 365
Asp Thr Phe Gln Glu Gln Asp Asp Ile Ser Asp Asp Val Glu Glu Val
370 375 380
Lys Asp Gln Phe Ala Thr His Glu Ala Phe Met Met Glu Leu Thr Ala
385 390 395 400
His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Ile

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405				410				415							
Thr	Gln	Gly	Thr	Leu	Ser	Asp	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln
			420								425				430
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met
		435					440							445	
Asp	Arg	Gln	Ser	Arg	Leu	His	Asp	Val	Leu	Met	Glu	Leu	Gln	Lys	Lys
	450					455					460				
Gln	Leu	Gln	Gln	Leu	Ser	Ala	Trp	Leu	Thr	Leu	Thr	Glu	Glu	Arg	Ile
465					470						475				480
Gln	Lys	Met	Glu	Thr	Cys	Pro	Leu	Asp	Asp	Asp	Val	Lys	Ser	Leu	Gln
					485						490				495
Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu
			500						505					510	
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu
		515					520							525	
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys
	530					535								540	
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp
545					550					555					560
Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu
					565					570					575
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn
			580						585					590	
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser
		595					600							605	
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln
	610					615					620				
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu
625					630					635					640
Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu
					645					650					655
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn
			660						665					670	
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln
		675					680							685	
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys
	690					695					700				
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln
705					710					715					720
Ile	His	Val	Asp	Ile	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Ala
					725					730					735
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Trp	Lys	Thr	Ala	Ile	Gln	Thr	Thr
			740						745					750	
Glu	Ile	Lys	Glu	Tyr	Met	Lys	Met	Gln	Asp	Thr	Ser	Glu	Met	Lys	Lys
		755					760							765	
Lys	Leu	Lys	Ala	Leu	Glu	Lys	Glu	Gln	Arg	Glu	Arg	Ile	Pro	Arg	Ala
	770					775					780				
Asp	Glu	Leu	Asn	Gln	Thr	Gly	Gln	Ile	Leu	Val	Glu	Gln	Met	Gly	Lys
785					790					795					800
Glu	Gly	Leu	Pro	Thr	Glu	Glu	Ile	Lys	Asn	Val	Leu	Glu	Lys	Val	Ser
					805					810					815
Ser	Glu	Trp	Lys	Asn	Val	Ser	Gln	His	Leu	Glu	Asp	Leu	Glu	Arg	Lys
					820					825					830

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Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Glu
 835 840 845
 Leu Glu Lys Val Ile Lys Thr Lys Glu Glu Trp Val Lys His Thr Ser
 850 855 860
 Ile Ser Glu Ser Ser Arg Gln Ser Leu Pro Ser Leu Lys Asp Ser Cys
 865 870 875 880
 Gln Arg Glu Leu Thr Asn Leu Leu Gly Leu His Pro Lys Ile Glu Met
 885 890 895
 Ala Arg Ala Ser Cys Ser Ala Leu Met Ser Gln Pro Ser Ala Pro Asp
 900 905 910
 Phe Val Gln Arg Gly Phe Asp Ser Phe Leu Gly Arg Tyr Gln Ala Val
 915 920 925
 Gln Glu Ala Val Glu Asp Arg Gln Gln His Leu Glu Asn Glu Leu Lys
 930 935 940
 Gly Gln Pro Gly His Ala Tyr Leu Glu Thr Leu Lys Thr Leu Lys Asp
 945 950 955 960
 Val Leu Asn Asp Ser Glu Asn Lys Ala Gln Val Ser Leu Asn Val Leu
 965 970 975
 Asn Asp Leu Ala Lys Val Glu Lys Ala Leu Gln Glu Lys Lys Thr Leu
 980 985 990
 Asp Glu Ile Leu Glu Asn Gln Lys Pro Ala Leu His Lys Leu Ala Glu
 995 1000 1005
 Glu Thr Lys Ala Leu Glu Lys Asn Val His Pro Asp Val Glu Lys
 1010 1015 1020
 Leu Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Lys Trp Asn Lys
 1025 1030 1035
 Leu Lys Val Leu Val Ser Lys Asp Leu His Leu Leu Glu Glu Ile
 1040 1045 1050
 Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr
 1055 1060 1065
 Ser Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn Lys Val
 1070 1075 1080
 Pro Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp His
 1085 1090 1095
 Pro Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn
 1100 1105 1110
 Val Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu
 1115 1120 1125
 Gln Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Ser Thr Thr Asn
 1130 1135 1140
 Glu Ile Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu
 1145 1150 1155
 Ser Val Pro Asp Val Ile Asn Cys Leu Thr Thr Thr Tyr Asp Gly
 1160 1165 1170
 Leu Glu Gln Met His Lys Asp Leu Val Asn Val Pro Leu Cys Val
 1175 1180 1185
 Asp Met Cys Leu Asn Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg
 1190 1195 1200
 Thr Gly Lys Ile Arg Val Gln Ser Leu Lys Ile Gly Leu Met Ser
 1205 1210 1215
 Leu Ser Lys Gly Leu Leu Glu Glu Lys Tyr Arg Tyr Leu Phe Lys
 1220 1225 1230

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Glu	Val	Ala	Gly	Pro	Thr	Glu	Met	Cys	Asp	Gln	Arg	Gln	Leu	Gly
1235						1240					1245			
Leu	Leu	Leu	His	Asp	Ala	Ile	Gln	Ile	Pro	Arg	Gln	Leu	Gly	Glu
1250						1255					1260			
Val	Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile	Glu	Pro	Ser	Val	Arg	Ser
1265						1270					1275			
Cys	Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro	Glu	Ile	Ser	Val	Lys	Glu
1280						1285					1290			
Phe	Ile	Asp	Trp	Met	His	Leu	Glu	Pro	Gln	Ser	Met	Val	Trp	Leu
1295						1300					1305			
Pro	Val	Leu	His	Arg	Val	Ala	Ala	Ala	Glu	Thr	Ala	Lys	His	Gln
1310						1315					1320			
Ala	Lys	Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly	Phe	Arg
1325						1330					1335			
Tyr	Arg	Ser	Leu	Lys	His	Phe	Asn	Tyr	Asp	Val	Cys	Gln	Ser	Cys
1340						1345					1350			
Phe	Phe	Ser	Gly	Arg	Thr	Ala	Lys	Gly	His	Lys	Leu	His	Tyr	Pro
1355						1360					1365			
Met	Val	Glu	Tyr	Cys	Ile	Pro	Thr	Thr	Ser	Gly	Glu	Asp	Val	Arg
1370						1375					1380			
Asp	Phe	Thr	Lys	Val	Leu	Lys	Asn	Lys	Phe	Arg	Ser	Lys	Lys	Tyr
1385						1390					1395			
Phe	Ala	Lys	His	Pro	Arg	Leu	Gly	Tyr	Leu	Pro	Val	Gln	Thr	Val
1400						1405					1410			
Leu	Glu	Gly	Asp	Asn	Leu	Glu	Thr	Pro	Ile	Thr	Leu	Ile	Ser	Met
1415						1420					1425			
Trp	Pro	Glu	His	Tyr	Asp	Pro	Ser	Gln	Ser	Pro	Gln	Leu	Phe	His
1430						1435					1440			
Asp	Asp	Thr	His	Ser	Arg	Ile	Glu	Gln	Tyr	Ala	Thr	Arg	Leu	Ala
1445						1450					1455			
Gln	Met	Glu	Arg	Thr	Asn	Gly	Ser	Phe	Leu	Thr	Asp	Ser	Ser	Ser
1460						1465					1470			
Thr	Thr	Gly	Ser	Val	Glu	Asp	Glu	His	Ala	Leu	Ile	Gln	Gln	Tyr
1475						1480					1485			
Cys	Gln	Thr	Leu	Gly	Gly	Glu	Ser	Pro	Val	Ser	Gln	Pro	Gln	Ser
1490						1495					1500			
Pro	Ala	Gln	Ile	Leu	Lys	Ser	Val	Glu	Arg	Glu	Glu	Arg	Gly	Glu
1505						1510					1515			
Leu	Glu	Arg	Ile	Ile	Ala	Asp	Leu	Glu	Glu	Glu	Gln	Arg	Asn	Leu
1520						1525					1530			
Gln	Val	Glu	Tyr	Glu	Gln	Leu	Lys	Asp	Gln	His	Leu	Arg	Arg	Gly
1535						1540					1545			
Leu	Pro	Val	Gly	Ser	Pro	Pro	Glu	Ser	Ile	Ile	Ser	Pro	His	His
1550						1555					1560			
Thr	Ser	Glu	Asp	Ser	Glu	Leu	Ile	Ala	Glu	Ala	Lys	Leu	Leu	Arg
1565						1570					1575			
Gln	His	Lys	Gly	Arg	Leu	Glu	Ala	Arg	Met	Gln	Ile	Leu	Glu	Asp
1580						1585					1590			
His	Asn	Lys	Gln	Leu	Glu	Ser	Gln	Leu	His	Arg	Leu	Arg	Gln	Leu
1595						1600					1605			
Leu	Glu	Gln	Pro	Glu	Ser	Asp	Ser	Arg	Ile	Asn	Gly	Val	Ser	Pro
1610						1615					1620			
Trp	Ala	Ser	Pro	Gln	His	Ser	Ala	Leu	Ser	Tyr	Ser	Leu	Asp	Pro

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1625	1630	1635	
Asp Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp Leu			
1640	1645	1650	
Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val Met			
1655	1660	1665	
Glu Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val Pro			
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Ser Arg Pro Gln Ala Met			
1685			
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Met Asp Tyr Lys Asp Asp Asp Asp Lys Gly Tyr Gly Arg Lys Lys Arg			
1 5 10 15			
cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac			96
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp			
20 25 30			
gtc cca gac tat gct ggc tcc atg gcc aag tat gga gaa cat gaa gcc			144
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala			
35 40 45			
agt cct gac aat ggg cag aac gaa ttc agt gat atc att aag tcc aga			192
Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg			
50 55 60			
tct gat gaa cac aat gac gta cag aag aaa acc ttt acc aaa tgg ata			240
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile			
65 70 75 80			
aat gct cga ttt tca aag agt ggg aaa cca ccc atc aat gat atg ttc			288
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe			
85 90 95			
aca gac ctc aaa gat gga agg aag cta ttg gat ctt cta gaa ggc ctc			336
Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu			
100 105 110			
aca gga aca tca ctg cca aag gaa cgt ggt tcc aca agg gta cat gcc			384
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala			
115 120 125			
tta aat aac gtc aac aga gtg ctg cag gtt tta cat cag aac aat gtg			432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val			
130 135 140			
gaa tta gtg aat ata ggg gga act gac att gtg gat gga aat cac aaa			480
Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys			
145 150 155 160			
ctg act ttg ggg tta ctt tgg agc atc att ttg cac tgg cag gtg aaa			528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys			
165 170 175			
gat gtc atg aag gat gtc atg tcg gac ctg cag cag acg aac agt gag			576
Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu			
180 185 190			
aag atc ctg ctc agc tgg gtg cgt cag acc acc agg ccc tac agc caa			624
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln			

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gtc aac gtc ctc aac ttc acc acc agc tgg aca gat gga ctc gcc ttt						672
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe						
210		215		220		
aat gct gtc ctc cac cga cat aaa cct gat ctc ttc agc tgg gat aaa						720
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Ser Trp Asp Lys						
225		230		235		240
gtt gtc aaa atg tca cca att gag aga ctt gaa cat gcc ttc agc aag						768
Val Val Lys Met Ser Pro Ile Glu Arg Leu Glu His Ala Phe Ser Lys						
		245		250		255
gct caa act tat ttg gga att gaa aag ctg tta gat cct gaa gat gtt						816
Ala Gln Thr Tyr Leu Gly Ile Glu Lys Leu Leu Asp Pro Glu Asp Val						
		260		265		270
gcc gtt cag ctt cct gac aag aaa tcc ata att atg tat tta aca tct						864
Ala Val Gln Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser						
		275		280		285
ttg ttt gag gtg cta cct cag caa gtc acc ata gac gcc atc cgt gag						912
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu						
		290		295		300
gta gag aca ctc cca agg aaa tat aaa aaa gaa tgt gaa gaa gag gca						960
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Ala						
		305		310		315
att aat ata cag agt aca gcg cct gag gag gag cat gag agt ccc cga						1008
Ile Asn Ile Gln Ser Thr Ala Pro Glu Glu Glu His Glu Ser Pro Arg						
		325		330		335
gct gaa act ccc agc act gtc act gag gtt gac atg gat ctg gac agc						1056
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser						
		340		345		350
tat cag att gcg ttg gag gaa gtg ctg acc tgg ttg ctt tct gct gag						1104
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu						
		355		360		365
gac act ttc cag gag cag gat gat att tct gat gat gtt gaa gaa gtc						1152
Asp Thr Phe Gln Glu Gln Asp Asp Ile Ser Asp Asp Val Glu Glu Val						
		370		375		380
aaa gac cag ttt gca acc cat gaa gct ttt atg atg gaa ctg act gca						1200
Lys Asp Gln Phe Ala Thr His Glu Ala Phe Met Met Glu Leu Thr Ala						
		385		390		395
cac cag agc agt gtg ggc agc gtc ctg cag gca ggc aac caa ctg ata						1248
His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Ile						
		405		410		415
aca caa gga act ctg tca gac gaa gaa gaa ttt gag att cag gaa cag						1296
Thr Gln Gly Thr Leu Ser Asp Glu Glu Glu Phe Glu Ile Gln Glu Gln						
		420		425		430
atg acc ctg ctg aat gct aga tgg gag gct ctt agg gtg gag agt atg						1344
Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met						
		435		440		445
gac aga cag tcc cgg ctg cac gat gtg ctg atg gaa ctg cag aag aag						1392
Asp Arg Gln Ser Arg Leu His Asp Val Leu Met Glu Leu Gln Lys Lys						
		450		455		460
caa ctg cag cag ctc tcc gcc tgg tta aca ctc aca gag gag cgc att						1440
Gln Leu Gln Gln Leu Ser Ala Trp Leu Thr Leu Thr Glu Glu Arg Ile						
		465		470		475
cag aag atg gaa act tgc ccc ctg gat gat gat gta aaa tct cta caa						1488
Gln Lys Met Glu Thr Cys Pro Leu Asp Asp Asp Val Lys Ser Leu Gln						
		485		490		495
aag ctg cta gaa gaa cat aaa agt ttg caa agt gat ctt gag gct gaa						1536
Lys Leu Leu Glu Glu His Lys Ser Leu Gln Ser Asp Leu Glu Ala Glu						
		500		505		510
cag gtg aaa gta aat tca cta act cac atg gtg gtc att gtt gat gaa						1584

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Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu	
		515					520					525				
aac	agt	ggt	gag	agt	gct	aca	gct	atc	cta	gaa	gac	cag	tta	cag	aaa	1632
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys	
	530				535					540						
ctt	ggt	gag	cgc	tgg	aca	gca	gta	tgc	cgt	tgg	act	gaa	gaa	cgc	tgg	1680
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp	
545				550					555						560	
aat	agg	tta	caa	gaa	atc	aat	ata	ttg	tgg	cag	gaa	tta	ttg	gaa	gaa	1728
Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu	
				565					570					575		
cag	tgc	ttg	ttg	aaa	gct	tgg	tta	acc	gaa	aaa	gaa	gag	gct	tta	aat	1776
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn	
		580						585						590		
aaa	gtc	cag	aca	agc	aac	ttc	aaa	gac	caa	aag	gaa	cta	agt	gtc	agt	1824
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser	
		595				600						605				
gtt	cga	cgt	ctg	gct	att	ttg	aag	gaa	gac	atg	gaa	atg	aag	cgt	caa	1872
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln	
	610					615					620					
aca	ttg	gat	cag	ctg	agt	gag	att	ggc	cag	gat	gtg	gga	caa	tta	ctt	1920
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu	
	625				630					635					640	
gat	aat	tcc	aag	gca	tct	aag	aag	atc	aac	agt	gac	tca	gag	gaa	ctg	1968
Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu	
				645					650					655		
act	caa	aga	tgg	gat	tct	ttg	gtt	cag	aga	cta	gaa	gat	tcc	tcc	aac	2016
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn	
			660					665					670			
cag	gtg	act	cag	gct	gta	gca	aag	ctg	ggg	atg	tct	cag	att	cct	cag	2064
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln	
		675					680							685		
aag	gac	ctt	ttg	gag	act	gtt	cgt	gta	aga	gaa	caa	gca	att	aca	aaa	2112
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys	
		690				695					700					
aaa	tct	aag	cag	gaa	ctg	cct	cct	cct	cct	ccc	cca	aag	aag	aga	cag	2160
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln	
	705				710					715					720	
atc	cat	gtg	gat	att	gaa	gct	aag	aaa	aag	ttt	gat	gct	ata	agt	gca	2208
Ile	His	Val	Asp	Ile	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Ala	
				725					730					735		
gag	ctg	ttg	aac	tgg	att	ttg	aaa	tgg	aaa	act	gcc	att	cag	acc	aca	2256
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Trp	Lys	Thr	Ala	Ile	Gln	Thr	Thr	
			740					745					750			
gag	ata	aaa	gag	tat	atg	aag	atg	caa	gac	act	tcc	gaa	atg	aaa	aag	2304
Glu	Ile	Lys	Glu	Tyr	Met	Lys	Met	Gln	Asp	Thr	Ser	Glu	Met	Lys	Lys	
		755					760					765				
aag	ttg	aag	gca	tta	gaa	aaa	gaa	cag	aga	gaa	aga	atc	ccc	aga	gca	2352
Lys	Leu	Lys	Ala	Leu	Glu	Lys	Glu	Gln	Arg	Glu	Arg	Ile	Pro	Arg	Ala	
		770				775						780				
gat	gaa	tta	aac	caa	act	gga	caa	atc	ctt	gtg	gag	caa	atg	gga	aaa	2400
Asp	Glu	Leu	Asn	Gln	Thr	Gly	Gln	Ile	Leu	Val	Glu	Gln	Met	Gly	Lys	
				785		790				795					800	
gaa	ggc	ctt	cct	act	gaa	gaa	ata	aaa	aat	gtt	ctg	gag	aag	gtt	tca	2448
Glu	Gly	Leu	Pro	Thr	Glu	Glu	Ile	Lys	Asn	Val	Leu	Glu	Lys	Val	Ser	
				805					810					815		
tca	gaa	tgg	aag	aat	gta	tct	caa	cat	ttg	gaa	gat	cta	gaa	aga	aag	2496
Ser	Glu	Trp	Lys	Asn	Val	Ser	Gln	His	Leu	Glu	Asp	Leu	Glu	Arg	Lys	
				820				825						830		

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ctt gaa aag gtc atc aag aca aag gag gag tgg gta aaa cac act tcc Leu Glu Lys Val Ile Lys Thr Lys Glu Glu Trp Val Lys His Thr Ser 850 855 860	2592
att tct gaa tct tcc cgg cag tcc ttg cca agc ttg aag gat tcc tgt Ile Ser Glu Ser Ser Arg Gln Ser Leu Pro Ser Leu Lys Asp Ser Cys 865 870 875 880	2640
cag cgg gaa ttg aca aat ctt ctt ggc ctt cac ccc aaa att gaa atg Gln Arg Glu Leu Thr Asn Leu Leu Gly Leu His Pro Lys Ile Glu Met 885 890 895	2688
gct cgt gca agc tgc tgc gcc ctg atg tct cag cct tct gcc cca gat Ala Arg Ala Ser Cys Ser Ala Leu Met Ser Gln Pro Ser Ala Pro Asp 900 905 910	2736
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ggc caa cct gga cat gca tat ctg gaa aca ttg aaa aca ctg aaa gat Gly Gln Pro Gly His Ala Tyr Leu Glu Thr Leu Lys Thr Leu Lys Asp 945 950 955 960	2880
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gaa aca aag gct ctg gag aaa aat gtt cat cct gat gta gaa aaa Glu Thr Lys Ala Leu Glu Lys Asn Val His Pro Asp Val Glu Lys 1010 1015 1020	3069
tta tat aag caa gaa ttt gat gat gtg caa gga aag tgg aac aag Leu Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Lys Trp Asn Lys 1025 1030 1035	3114
cta aag gtc ttg gtt tcc aaa gat cta cat ttg ctt gag gaa att Leu Lys Val Leu Val Ser Lys Asp Leu His Leu Leu Glu Glu Ile 1040 1045 1050	3159
gct ctc aca ctc aga gct ttt gag gcc gat tca aca gtc att gag Ala Leu Thr Leu Arg Ala Phe Glu Ala Asp Ser Thr Val Ile Glu 1055 1060 1065	3204
aag tgg atg gat ggc gtg aaa gac ttc tta atg aaa cag cag gct Lys Trp Met Asp Gly Val Lys Asp Phe Leu Met Lys Gln Gln Ala 1070 1075 1080	3249
gcc caa gga gac gac gca ggt cta cag agg cag tta gac cag tgc Ala Gln Gly Asp Asp Ala Gly Leu Gln Arg Gln Leu Asp Gln Cys 1085 1090 1095	3294
tct gca ttt gtt aat gaa ata gaa aca att gaa tca tct ctg aaa Ser Ala Phe Val Asn Glu Ile Glu Thr Ile Glu Ser Ser Leu Lys 1100 1105 1110	3339
aac atg aag gaa ata gag act aat ctt cga agt ggt cca gtt gct Asn Met Lys Glu Ile Glu Thr Asn Leu Arg Ser Gly Pro Val Ala 1115 1120 1125	3384
gga ata aaa act tgg gtg cag aca aga cta ggt gac tac caa act Gly Ile Lys Thr Trp Val Gln Thr Arg Leu Gly Asp Tyr Gln Thr 1130 1135 1140	3429

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gtg aga gtg aag att ctc aag gac aac atc aag tta tta gct gcc 3699 Val Arg Val Lys Ile Leu Lys Asp Asn Ile Lys Leu Leu Ala Ala 1220 1225 1230
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ctt gct gac ctg aat aat gta cgt ttt tct gcc tac cgt aca gca 4329 Leu Ala Asp Leu Asn Asn Val Arg Phe Ser Ala Tyr Arg Thr Ala

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gag ttg agt aca aca aat gaa att ttc aaa cag cac aag ttg aac Glu Leu Ser Thr Thr Asn Glu Ile Phe Lys Gln His Lys Leu Asn 1460 1465 1470			4419
caa aat gac cag ctc ctc agt gtt cca gat gtc atc aac tgt ctg Gln Asn Asp Gln Leu Leu Ser Val Pro Asp Val Ile Asn Cys Leu 1475 1480 1485			4464
aca aca act tat gat gga ctt gag caa atg cat aag gac ctg gtc Thr Thr Thr Tyr Asp Gly Leu Glu Gln Met His Lys Asp Leu Val 1490 1495 1500			4509
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gtc tat gac acg ggt cga act gga aaa att aga gtg cag agt ctg Val Tyr Asp Thr Gly Arg Thr Gly Lys Ile Arg Val Gln Ser Leu 1520 1525 1530			4599
aag att gga tta atg tct ctc tcc aaa ggt ctc ttg gaa gaa aaa Lys Ile Gly Leu Met Ser Leu Ser Lys Gly Leu Leu Glu Glu Lys 1535 1540 1545			4644
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ccc cgg cag cta ggt gaa gta gca gct ttt gga ggc agt aat att Pro Arg Gln Leu Gly Glu Val Ala Ala Phe Gly Gly Ser Asn Ile 1580 1585 1590			4779
gag cct agt gtt cgc agc tgc ttc caa cag aat aac aat aaa cca Glu Pro Ser Val Arg Ser Cys Phe Gln Gln Asn Asn Asn Lys Pro 1595 1600 1605			4824
gaa ata agt gtg aaa gag ttt ata gat tgg atg cat ttg gaa cca Glu Ile Ser Val Lys Glu Phe Ile Asp Trp Met His Leu Glu Pro 1610 1615 1620			4869
cag tcc atg gtt tgg ctc cca gtt tta cat cga gtg gca gca gcg Gln Ser Met Val Trp Leu Pro Val Leu His Arg Val Ala Ala Ala 1625 1630 1635			4914
gag act gca aaa cat cag gcc aaa tgc aac atc tgt aaa gaa tgt Glu Thr Ala Lys His Gln Ala Lys Cys Asn Ile Cys Lys Glu Cys 1640 1645 1650			4959
cca att gtc ggg ttc agg tat aga agc ctt aag cat ttt aac tat Pro Ile Val Gly Phe Arg Tyr Arg Ser Leu Lys His Phe Asn Tyr 1655 1660 1665			5004
gat gtc tgc cag agt tgt ttc ttt tgc ggt cga aca gca aaa ggt Asp Val Cys Gln Ser Cys Phe Phe Ser Gly Arg Thr Ala Lys Gly 1670 1675 1680			5049
cac aaa tta cat tac cca atg gtg gaa tat tgt ata cct aca aca His Lys Leu His Tyr Pro Met Val Glu Tyr Cys Ile Pro Thr Thr 1685 1690 1695			5094
tct ggg gaa gat gta cga gac ttc aca aag gta ctt aag aac aag Ser Gly Glu Asp Val Arg Asp Phe Thr Lys Val Leu Lys Asn Lys 1700 1705 1710			5139
ttc agg tgc aag aag tac ttt gcc aaa cac cct cga ctt ggt tac Phe Arg Ser Lys Lys Tyr Phe Ala Lys His Pro Arg Leu Gly Tyr 1715 1720 1725			5184
ctg cct gtc cag aca gtt ctt gaa ggt gac aac tta gag act cct			5229

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Leu	Pro	Val	Gln	Thr	Val	Leu	Glu	Gly	Asp	Asn	Leu	Glu	Thr	Pro			
	1730					1735					1740						
atc	aca	ctc	atc	agt	atg	tgg	cca	gag	cac	tat	gac	ccc	tca	caa		5274	
Ile	Thr	Leu	Ile	Ser	Met	Trp	Pro	Glu	His	Tyr	Asp	Pro	Ser	Gln			
	1745					1750					1755						
tct	cct	caa	ctg	ttt	cat	gat	gac	acc	cat	tca	aga	ata	gaa	caa		5319	
Ser	Pro	Gln	Leu	Phe	His	Asp	Asp	Thr	His	Ser	Arg	Ile	Glu	Gln			
	1760					1765					1770						
tat	gcc	aca	cga	ctg	gcc	cag	atg	gaa	agg	act	aat	ggg	tct	ttt		5364	
Tyr	Ala	Thr	Arg	Leu	Ala	Gln	Met	Glu	Arg	Thr	Asn	Gly	Ser	Phe			
	1775					1780					1785						
ctc	act	gat	agc	agc	tcc	acc	aca	gga	agt	gtg	gaa	gac	gag	cac		5409	
Leu	Thr	Asp	Ser	Ser	Ser	Thr	Thr	Gly	Ser	Val	Glu	Asp	Glu	His			
	1790					1795					1800						
gcc	ctc	atc	cag	cag	tat	tgc	caa	aca	ctc	gga	gga	gag	tcc	cca		5454	
Ala	Leu	Ile	Gln	Gln	Tyr	Cys	Gln	Thr	Leu	Gly	Gly	Glu	Ser	Pro			
	1805					1810					1815						
gtg	agc	cag	ccg	cag	agc	cca	gct	cag	atc	ctg	aag	tca	gta	gag		5499	
Val	Ser	Gln	Pro	Gln	Ser	Pro	Ala	Gln	Ile	Leu	Lys	Ser	Val	Glu			
	1820					1825					1830						
agg	gaa	gaa	cgt	gga	gaa	ctg	gag	agg	atc	att	gct	gac	ctg	gag		5544	
Arg	Glu	Glu	Arg	Gly	Glu	Leu	Glu	Arg	Ile	Ile	Ala	Asp	Leu	Glu			
	1835					1840					1845						
gaa	gaa	caa	aga	aat	cta	cag	gtg	gag	tat	gag	cag	ctg	aag	gac		5589	
Glu	Glu	Gln	Arg	Asn	Leu	Gln	Val	Glu	Tyr	Glu	Gln	Leu	Lys	Asp			
	1850					1855					1860						
cag	cac	ctc	cga	agg	ggg	ctc	cct	gtc	ggc	tca	ccg	cca	gag	tcg		5634	
Gln	His	Leu	Arg	Arg	Gly	Leu	Pro	Val	Gly	Ser	Pro	Pro	Glu	Ser			
	1865					1870					1875						
att	ata	tct	ccc	cat	cac	acg	tct	gag	gat	tca	gaa	ctt	ata	gca		5679	
Ile	Ile	Ser	Pro	His	His	Thr	Ser	Glu	Asp	Ser	Glu	Leu	Ile	Ala			
	1880					1885					1890						
gaa	gca	aaa	ctc	ctc	agg	cag	cac	aaa	ggc	cgg	ctg	gag	gct	agg		5724	
Glu	Ala	Lys	Leu	Leu	Arg	Gln	His	Lys	Gly	Arg	Leu	Glu	Ala	Arg			
	1895					1900					1905						
atg	cag	att	tta	gaa	gat	cac	aat	aaa	cag	ctg	gag	tct	cag	ctc		5769	
Met	Gln	Ile	Leu	Glu	Asp	His	Asn	Lys	Gln	Leu	Glu	Ser	Gln	Leu			
	1910					1915					1920						
cac	cgc	ctc	cga	cag	ctg	ctg	gag	cag	cct	gaa	tct	gat	tcc	cga		5814	
His	Arg	Leu	Arg	Gln	Leu	Leu	Glu	Gln	Pro	Glu	Ser	Asp	Ser	Arg			
	1925					1930					1935						
atc	aat	ggc	gtt	tcc	cca	tgg	gct	tct	cct	cag	cat	tct	gca	ctg		5859	
Ile	Asn	Gly	Val	Ser	Pro	Trp	Ala	Ser	Pro	Gln	His	Ser	Ala	Leu			
	1940					1945					1950						
agc	tac	tcg	ctt	gat	cca	gat	gcc	tcc	ggc	cca	cag	ttc	cac	cag		5904	
Ser	Tyr	Ser	Leu	Asp	Pro	Asp	Ala	Ser	Gly	Pro	Gln	Phe	His	Gln			
	1955					1960					1965						
gca	gcg	gga	gag	gac	ctg	ctg	gcc	cca	ccg	cac	gac	acc	agc	acg		5949	
Ala	Ala	Gly	Glu	Asp	Leu	Leu	Ala	Pro	Pro	His	Asp	Thr	Ser	Thr			
	1970					1975					1980						
gat	ctc	acg	gag	gtc	atg	gag	cag	att	cac	agc	acg	ttt	cca	tct		5994	
Asp	Leu	Thr	Glu	Val	Met	Glu	Gln	Ile	His	Ser	Thr	Phe	Pro	Ser			
	1985					1990					1995						
tgc	tgc	cca	aat	gtt	ccc	agc	agg	cca	cag	gca	atg	tga				6033	
Cys	Cys	Pro	Asn	Val	Pro	Ser	Arg	Pro	Gln	Ala	Met						
	2000					2005					2010						

<210> SEQ ID NO 15

<211> LENGTH: 2010

<212> TYPE: PRT

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<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 15

Met Asp Tyr Lys Asp Asp Asp Lys Gly Tyr Gly Arg Lys Lys Arg
 1 5 10 15
 Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20 25 30
 Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
 35 40 45
 Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50 55 60
 Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65 70 75 80
 Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
 85 90 95
 Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100 105 110
 Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115 120 125
 Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
 130 135 140
 Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
 145 150 155 160
 Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
 165 170 175
 Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu
 180 185 190
 Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln
 195 200 205
 Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe
 210 215 220
 Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Ser Trp Asp Lys
 225 230 235 240
 Val Val Lys Met Ser Pro Ile Glu Arg Leu Glu His Ala Phe Ser Lys
 245 250 255
 Ala Gln Thr Tyr Leu Gly Ile Glu Lys Leu Leu Asp Pro Glu Asp Val
 260 265 270
 Ala Val Gln Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser
 275 280 285
 Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu
 290 295 300
 Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Ala
 305 310 315 320
 Ile Asn Ile Gln Ser Thr Ala Pro Glu Glu Glu His Glu Ser Pro Arg
 325 330 335
 Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser
 340 345 350
 Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu
 355 360 365
 Asp Thr Phe Gln Glu Gln Asp Asp Ile Ser Asp Asp Val Glu Glu Val
 370 375 380
 Lys Asp Gln Phe Ala Thr His Glu Ala Phe Met Met Glu Leu Thr Ala
 385 390 395 400

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His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Ile
				405					410					415	
Thr	Gln	Gly	Thr	Leu	Ser	Asp	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln
			420					425					430		
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met
		435					440					445			
Asp	Arg	Gln	Ser	Arg	Leu	His	Asp	Val	Leu	Met	Glu	Leu	Gln	Lys	Lys
	450					455					460				
Gln	Leu	Gln	Gln	Leu	Ser	Ala	Trp	Leu	Thr	Leu	Thr	Glu	Glu	Arg	Ile
465					470					475					480
Gln	Lys	Met	Glu	Thr	Cys	Pro	Leu	Asp	Asp	Asp	Val	Lys	Ser	Leu	Gln
				485					490					495	
Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu
			500					505					510		
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu
		515					520					525			
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys
		530				535					540				
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp
545					550					555					560
Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu
				565					570					575	
Gln	Cys	Leu	Leu	Lys	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asn
			580				585						590		
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser
		595				600						605			
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln
	610					615					620				
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu
625					630					635					640
Asp	Asn	Ser	Lys	Ala	Ser	Lys	Lys	Ile	Asn	Ser	Asp	Ser	Glu	Glu	Leu
				645					650					655	
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn
			660					665						670	
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln
		675					680					685			
Lys	Asp	Leu	Leu	Glu	Thr	Val	Arg	Val	Arg	Glu	Gln	Ala	Ile	Thr	Lys
	690					695					700				
Lys	Ser	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln
705					710					715					720
Ile	His	Val	Asp	Ile	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Ala
				725					730					735	
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Trp	Lys	Thr	Ala	Ile	Gln	Thr	Thr
			740					745					750		
Glu	Ile	Lys	Glu	Tyr	Met	Lys	Met	Gln	Asp	Thr	Ser	Glu	Met	Lys	Lys
		755					760					765			
Lys	Leu	Lys	Ala	Leu	Glu	Lys	Glu	Gln	Arg	Glu	Arg	Ile	Pro	Arg	Ala
	770					775						780			
Asp	Glu	Leu	Asn	Gln	Thr	Gly	Gln	Ile	Leu	Val	Glu	Gln	Met	Gly	Lys
785					790					795					800
Glu	Gly	Leu	Pro	Thr	Glu	Glu	Ile	Lys	Asn	Val	Leu	Glu	Lys	Val	Ser
				805					810					815	
Ser	Glu	Trp	Lys	Asn	Val	Ser	Gln	His	Leu	Glu	Asp	Leu	Glu	Arg	Lys

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Lys	Val	Pro	Ser	Gly	Gly	Gln	Glu	Leu	Thr	Ser	Glu	Leu	Asn	Val
1235						1240					1245			
Val	Leu	Glu	Asn	Tyr	Gln	Leu	Leu	Cys	Asn	Arg	Ile	Arg	Gly	Lys
1250						1255					1260			
Cys	His	Thr	Leu	Glu	Glu	Val	Trp	Ser	Cys	Trp	Ile	Glu	Leu	Leu
1265						1270					1275			
His	Tyr	Leu	Asp	Leu	Glu	Thr	Thr	Trp	Leu	Asn	Thr	Leu	Glu	Glu
1280						1285					1290			
Arg	Met	Lys	Ser	Thr	Glu	Val	Leu	Pro	Glu	Lys	Thr	Asp	Ala	Val
1295						1300					1305			
Asn	Glu	Ala	Leu	Glu	Ser	Leu	Glu	Ser	Val	Leu	Arg	His	Pro	Ala
1310						1315					1320			
Asp	Asn	Arg	Thr	Gln	Ile	Arg	Glu	Leu	Gly	Gln	Thr	Leu	Ile	Asp
1325						1330					1335			
Gly	Gly	Ile	Leu	Asp	Asp	Ile	Ile	Ser	Glu	Lys	Leu	Glu	Ala	Phe
1340						1345					1350			
Asn	Ser	Arg	Tyr	Glu	Asp	Leu	Ser	His	Leu	Ala	Glu	Ser	Lys	Gln
1355						1360					1365			
Ile	Ser	Leu	Glu	Lys	Gln	Ala	His	Arg	Asp	Phe	Gly	Pro	Ser	Ser
1370						1375					1380			
Gln	His	Phe	Leu	Ser	Thr	Ser	Val	Gln	Leu	Pro	Trp	Gln	Arg	Ser
1385						1390					1395			
Ile	Ser	His	Asn	Lys	Val	Pro	Tyr	Tyr	Ile	Asn	His	Gln	Thr	Gln
1400						1405					1410			
Thr	Thr	Cys	Trp	Asp	His	Pro	Lys	Met	Thr	Glu	Leu	Phe	Gln	Ser
1415						1420					1425			
Leu	Ala	Asp	Leu	Asn	Asn	Val	Arg	Phe	Ser	Ala	Tyr	Arg	Thr	Ala
1430						1435					1440			
Ile	Lys	Ile	Arg	Arg	Leu	Gln	Lys	Ala	Leu	Cys	Leu	Asp	Leu	Leu
1445						1450					1455			
Glu	Leu	Ser	Thr	Thr	Asn	Glu	Ile	Phe	Lys	Gln	His	Lys	Leu	Asn
1460						1465					1470			
Gln	Asn	Asp	Gln	Leu	Leu	Ser	Val	Pro	Asp	Val	Ile	Asn	Cys	Leu
1475						1480					1485			
Thr	Thr	Thr	Tyr	Asp	Gly	Leu	Glu	Gln	Met	His	Lys	Asp	Leu	Val
1490						1495					1500			
Asn	Val	Pro	Leu	Cys	Val	Asp	Met	Cys	Leu	Asn	Trp	Leu	Leu	Asn
1505						1510					1515			
Val	Tyr	Asp	Thr	Gly	Arg	Thr	Gly	Lys	Ile	Arg	Val	Gln	Ser	Leu
1520						1525					1530			
Lys	Ile	Gly	Leu	Met	Ser	Leu	Ser	Lys	Gly	Leu	Leu	Glu	Glu	Lys
1535						1540					1545			
Tyr	Arg	Tyr	Leu	Phe	Lys	Glu	Val	Ala	Gly	Pro	Thr	Glu	Met	Cys
1550						1555					1560			
Asp	Gln	Arg	Gln	Leu	Gly	Leu	Leu	Leu	His	Asp	Ala	Ile	Gln	Ile
1565						1570					1575			
Pro	Arg	Gln	Leu	Gly	Glu	Val	Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile
1580						1585					1590			
Glu	Pro	Ser	Val	Arg	Ser	Cys	Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro
1595						1600					1605			
Glu	Ile	Ser	Val	Lys	Glu	Phe	Ile	Asp	Trp	Met	His	Leu	Glu	Pro
1610						1615					1620			

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Gln Ser	Met Val Trp	Leu Pro	Val Leu His Arg	Val	Ala Ala Ala
1625		1630		1635	
Glu Thr	Ala Lys His	Gln Ala	Lys Cys Asn Ile	Cys	Lys Glu Cys
1640		1645		1650	
Pro Ile	Val Gly Phe	Arg Tyr	Arg Ser Leu Lys	His	Phe Asn Tyr
1655		1660		1665	
Asp Val	Cys Gln Ser	Cys Phe	Phe Ser Gly Arg	Thr	Ala Lys Gly
1670		1675		1680	
His Lys	Leu His Tyr	Pro Met	Val Glu Tyr Cys	Ile	Pro Thr Thr
1685		1690		1695	
Ser Gly	Glu Asp Val	Arg Asp	Phe Thr Lys Val	Leu	Lys Asn Lys
1700		1705		1710	
Phe Arg	Ser Lys Lys	Tyr Phe	Ala Lys His Pro	Arg	Leu Gly Tyr
1715		1720		1725	
Leu Pro	Val Gln Thr	Val Leu	Glu Gly Asp Asn	Leu	Glu Thr Pro
1730		1735		1740	
Ile Thr	Leu Ile Ser	Met Trp	Pro Glu His Tyr	Asp	Pro Ser Gln
1745		1750		1755	
Ser Pro	Gln Leu Phe	His Asp	Asp Thr His Ser	Arg	Ile Glu Gln
1760		1765		1770	
Tyr Ala	Thr Arg Leu	Ala Gln	Met Glu Arg Thr	Asn	Gly Ser Phe
1775		1780		1785	
Leu Thr	Asp Ser Ser	Ser Thr	Thr Gly Ser Val	Glu	Asp Glu His
1790		1795		1800	
Ala Leu	Ile Gln Gln	Tyr Cys	Gln Thr Leu Gly	Gly	Glu Ser Pro
1805		1810		1815	
Val Ser	Gln Pro Gln	Ser Pro	Ala Gln Ile Leu	Lys	Ser Val Glu
1820		1825		1830	
Arg Glu	Glu Arg Gly	Glu Leu	Glu Arg Ile Ile	Ala	Asp Leu Glu
1835		1840		1845	
Glu Glu	Gln Arg Asn	Leu Gln	Val Glu Tyr Glu	Gln	Leu Lys Asp
1850		1855		1860	
Gln His	Leu Arg Arg	Gly Leu	Pro Val Gly Ser	Pro	Pro Glu Ser
1865		1870		1875	
Ile Ile	Ser Pro His	His Thr	Ser Glu Asp Ser	Glu	Leu Ile Ala
1880		1885		1890	
Glu Ala	Lys Leu Leu	Arg Gln	His Lys Gly Arg	Leu	Glu Ala Arg
1895		1900		1905	
Met Gln	Ile Leu Glu	Asp His	Asn Lys Gln Leu	Glu	Ser Gln Leu
1910		1915		1920	
His Arg	Leu Arg Gln	Leu Leu	Glu Gln Pro Glu	Ser	Asp Ser Arg
1925		1930		1935	
Ile Asn	Gly Val Ser	Pro Trp	Ala Ser Pro Gln	His	Ser Ala Leu
1940		1945		1950	
Ser Tyr	Ser Leu Asp	Pro Asp	Ala Ser Gly Pro	Gln	Phe His Gln
1955		1960		1965	
Ala Ala	Gly Glu Asp	Leu Leu	Ala Pro Pro His	Asp	Thr Ser Thr
1970		1975		1980	
Asp Leu	Thr Glu Val	Met Glu	Gln Ile His Ser	Thr	Phe Pro Ser
1985		1990		1995	
Cys Cys	Pro Asn Val	Pro Ser	Arg Pro Gln Ala	Met	
2000		2005		2010	

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<210> SEQ ID NO 16
<211> LENGTH: 6327
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<220> FEATURE:
<221> NAME/KEY: CDS
<222> LOCATION: (1)..(6327)
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (1)..(117)
<223> OTHER INFORMATION: TAT and epitope tag coding sequence

<400> SEQUENCE: 16

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Met Asp Tyr Lys Asp Asp Asp Lys Gly Tyr Gly Arg Lys Lys Arg
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cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac      96
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
20         25         30

gtc cca gac tat gct ggc tcc atg gcc aag tat gga gaa cat gaa gcc      144
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
35         40         45

agt cct gac aat ggg cag aac gaa ttc agt gat atc att aag tcc aga      192
Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
50         55         60

tct gat gaa cac aat gac gta cag aag aaa acc ttt acc aaa tgg ata      240
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
65         70         75         80

aat gct cga ttt tca aag agt ggg aaa cca ccc atc aat gat atg ttc      288
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
85         90         95

aca gac ctc aaa gat gga agg aag cta ttg gat ctt cta gaa ggc ctc      336
Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
100        105        110

aca gga aca tca ctg cca aag gaa cgt ggt tcc aca agg gta cat gcc      384
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
115        120        125

tta aat aac gtc aac aga gtg ctg cag gtt tta cat cag aac aat gtg      432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
130        135        140

gaa tta gtg aat ata ggg gga act gac att gtg gat gga aat cac aaa      480
Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
145        150        155        160

ctg act ttg ggg tta ctt tgg agc atc att ttg cac tgg cag gtg aaa      528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
165        170        175

gat gtc atg aag gat gtc atg tgc gac ctg cag cag acg aac agt gag      576
Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu
180        185        190

aag atc ctg ctc agc tgg gtg cgt cag acc acc agg ccc tac agc caa      624
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln
195        200        205

gtc aac gtc ctc aac ttc acc acc agc tgg aca gat gga ctc gcc ttt      672
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe
210        215        220

aat gct gtc ctc cac cga cat aaa cct gat ctc ttc agc tgg gat aaa      720
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Ser Trp Asp Lys
225        230        235        240

gtt gtc aaa atg tca cca att gag aga ctt gaa cat gcc ttc agc aag      768
Val Val Lys Met Ser Pro Ile Glu Arg Leu Glu His Ala Phe Ser Lys
245        250        255

gct caa act tat ttg gga att gaa aag ctg tta gat cct gaa gat gtt      816

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Ala	Gln	Thr	Tyr	Leu	Gly	Ile	Glu	Lys	Leu	Leu	Asp	Pro	Glu	Asp	Val			
			260					265					270					
gcc	ggt	cag	ctt	cct	gac	aag	aaa	tcc	ata	att	atg	tat	tta	aca	tct	864		
Ala	Val	Gln	Leu	Pro	Asp	Lys	Lys	Ser	Ile	Ile	Met	Tyr	Leu	Thr	Ser			
		275					280					285						
ttg	ttt	gag	gtg	cta	cct	cag	caa	gtc	acc	ata	gac	gcc	atc	cgt	gag	912		
Leu	Phe	Glu	Val	Leu	Pro	Gln	Gln	Val	Thr	Ile	Asp	Ala	Ile	Arg	Glu			
		290				295					300							
gta	gag	aca	ctc	cca	agg	aaa	tat	aaa	aaa	gaa	tgt	gaa	gaa	gag	gca	960		
Val	Glu	Thr	Leu	Pro	Arg	Lys	Tyr	Lys	Lys	Glu	Cys	Glu	Glu	Glu	Ala			
305					310					315					320			
att	aat	ata	cag	agt	aca	gcg	cct	gag	gag	gag	cat	gag	agt	ccc	cga	1008		
Ile	Asn	Ile	Gln	Ser	Thr	Ala	Pro	Glu	Glu	Glu	His	Glu	Ser	Pro	Arg			
					325					330					335			
gct	gaa	act	ccc	agc	act	gtc	act	gag	ggt	gac	atg	gat	ctg	gac	agc	1056		
Ala	Glu	Thr	Pro	Ser	Thr	Val	Thr	Glu	Val	Asp	Met	Asp	Leu	Asp	Ser			
			340					345					350					
tat	cag	att	gcg	ttg	gag	gaa	gtg	ctg	acc	tgg	ttg	ctt	tct	gct	gag	1104		
Tyr	Gln	Ile	Ala	Leu	Glu	Glu	Val	Leu	Thr	Trp	Leu	Leu	Ser	Ala	Glu			
		355					360						365					
gac	act	ttc	cag	gag	cag	gat	gat	att	tct	gat	gat	ggt	gaa	gaa	gtc	1152		
Asp	Thr	Phe	Gln	Glu	Gln	Asp	Asp	Ile	Ser	Asp	Asp	Val	Glu	Glu	Val			
		370				375					380							
aaa	gac	cag	ttt	gca	acc	cat	gaa	gct	ttt	atg	atg	gaa	ctg	act	gca	1200		
Lys	Asp	Gln	Phe	Ala	Thr	His	Glu	Ala	Phe	Met	Met	Glu	Leu	Thr	Ala			
385					390					395					400			
cac	cag	agc	agt	gtg	ggc	agc	gtc	ctg	cag	gca	ggc	aac	caa	ctg	ata	1248		
His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Ile			
				405				410						415				
aca	caa	gga	act	ctg	tca	gac	gaa	gaa	gaa	ttt	gag	att	cag	gaa	cag	1296		
Thr	Gln	Gly	Thr	Leu	Ser	Asp	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln			
			420					425						430				
atg	acc	ctg	ctg	aat	gct	aga	tgg	gag	gct	ctt	agg	gtg	gag	agt	atg	1344		
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met			
			435				440						445					
gac	aga	cag	tcc	cgg	ctg	cac	gat	gtg	ctg	atg	gaa	ctg	cag	aag	aag	1392		
Asp	Arg	Gln	Ser	Arg	Leu	His	Asp	Val	Leu	Met	Glu	Leu	Gln	Lys	Lys			
		450				455					460							
caa	ctg	cag	cag	ctc	tcc	gcc	tgg	tta	aca	ctc	aca	gag	gag	cgc	att	1440		
Gln	Leu	Gln	Gln	Leu	Ser	Ala	Trp	Leu	Thr	Leu	Thr	Glu	Glu	Arg	Ile			
465				470						475					480			
cag	aag	atg	gaa	act	tgc	ccc	ctg	gat	gat	gat	gta	aaa	tct	cta	caa	1488		
Gln	Lys	Met	Glu	Thr	Cys	Pro	Leu	Asp	Asp	Asp	Val	Lys	Ser	Leu	Gln			
				485						490					495			
aag	ctg	cta	gaa	gaa	cat	aaa	agt	ttg	caa	agt	gat	ctt	gag	gct	gaa	1536		
Lys	Leu	Leu	Glu	Glu	His	Lys	Ser	Leu	Gln	Ser	Asp	Leu	Glu	Ala	Glu			
			500					505						510				
cag	gtg	aaa	gta	aat	tca	cta	act	cac	atg	gtg	gtc	att	gtt	gat	gaa	1584		
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu			
		515					520						525					
aac	agt	ggg	gag	agt	gct	aca	gct	atc	cta	gaa	gac	cag	tta	cag	aaa	1632		
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Ile	Leu	Glu	Asp	Gln	Leu	Gln	Lys			
		530				535					540							
ctt	ggt	gag	cgc	tgg	aca	gca	gta	tgc	cgt	tgg	act	gaa	gaa	cgc	tgg	1680		
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp			
545				550						555					560			
aat	agg	tta	caa	gaa	atc	aat	ata	ttg	tgg	cag	gaa	tta	ttg	gaa	gaa	1728		
Asn	Arg	Leu	Gln	Glu	Ile	Asn	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu			
				565					570						575			

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cag tgc ttg ttg aaa gct tgg tta acc gaa aaa gaa gag gct tta aat	1776
Gln Cys Leu Leu Lys Ala Trp Leu Thr Glu Lys Glu Glu Ala Leu Asn	
580 585 590	
aaa gtc cag aca agc aac ttc aaa gac caa aag gaa cta agt gtc agt	1824
Lys Val Gln Thr Ser Asn Phe Lys Asp Gln Lys Glu Leu Ser Val Ser	
595 600 605	
gtt cga cgt ctg gct att ttg aag gaa gac atg gaa atg aag cgt caa	1872
Val Arg Arg Leu Ala Ile Leu Lys Glu Asp Met Glu Met Lys Arg Gln	
610 615 620	
aca ttg gat cag ctg agt gag att ggc cag gat gtg gga caa tta ctt	1920
Thr Leu Asp Gln Leu Ser Glu Ile Gly Gln Asp Val Gly Gln Leu Leu	
625 630 635 640	
gat aat tcc aag gca tct aag aag atc aac agt gac tca gag gaa ctg	1968
Asp Asn Ser Lys Ala Ser Lys Lys Ile Asn Ser Asp Ser Glu Glu Leu	
645 650 655	
act caa aga tgg gat tct ttg gtt cag aga cta gaa gat tcc tcc aac	2016
Thr Gln Arg Trp Asp Ser Leu Val Gln Arg Leu Glu Asp Ser Ser Asn	
660 665 670	
cag gtg act cag gct gta gca aag ctg ggg atg tct cag att cct cag	2064
Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln	
675 680 685	
aag gac ctt ttg gag act gtt cgt gta aga gaa caa gca att aca aaa	2112
Lys Asp Leu Leu Glu Thr Val Arg Val Arg Glu Gln Ala Ile Thr Lys	
690 695 700	
aaa tct aag cag gaa ctg cct cct cct ccc cca aag aag aga cag	2160
Lys Ser Lys Gln Glu Leu Pro Pro Pro Pro Pro Pro Lys Lys Arg Gln	
705 710 715 720	
atc cat gtg gat att gaa gct aag aaa aag ttt gat gct ata agt gca	2208
Ile His Val Asp Ile Glu Ala Lys Lys Lys Phe Asp Ala Ile Ser Ala	
725 730 735	
gag ctg ttg aac tgg att ttg aaa tgg aaa act gcc att cag acc aca	2256
Glu Leu Leu Asn Trp Ile Leu Lys Trp Lys Thr Ala Ile Gln Thr Thr	
740 745 750	
gag ata aaa gag tat atg aag atg caa gac act tcc gaa atg aaa aag	2304
Glu Ile Lys Glu Tyr Met Lys Met Gln Asp Thr Ser Glu Met Lys Lys	
755 760 765	
aag ttg aag gca tta gaa aaa gaa cag aga gaa aga atc ccc aga gca	2352
Lys Leu Lys Ala Leu Glu Lys Glu Gln Arg Glu Arg Ile Pro Arg Ala	
770 775 780	
gat gaa tta aac caa act gga caa atc ctt gtg gag caa atg gga aaa	2400
Asp Glu Leu Asn Gln Thr Gly Gln Ile Leu Val Glu Gln Met Gly Lys	
785 790 795 800	
gaa ggc ctt cct act gaa gaa ata aaa aat gtt ctg gag aag gtt tca	2448
Glu Gly Leu Pro Thr Glu Glu Ile Lys Asn Val Leu Glu Lys Val Ser	
805 810 815	
tca gaa tgg aag aat gta tct caa cat ttg gaa gat cta gaa aga aag	2496
Ser Glu Trp Lys Asn Val Ser Gln His Leu Glu Asp Leu Glu Arg Lys	
820 825 830	
att cag cta cag gaa gat ata aat gct tat ttc aag cag ctt gat gag	2544
Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Glu	
835 840 845	
ctt gaa aag gtc atc aag aca aag gag gag tgg gta aaa cac act tcc	2592
Leu Glu Lys Val Ile Lys Thr Lys Glu Glu Trp Val Lys His Thr Ser	
850 855 860	
att tct gaa tct tcc cgg cag tcc ttg cca agc ttg aag gat tcc tgt	2640
Ile Ser Glu Ser Ser Arg Gln Ser Leu Pro Ser Leu Lys Asp Ser Cys	
865 870 875 880	
cag cgg gaa ttg aca aat ctt ctt ggc ctt cac ccc aaa att gaa atg	2688
Gln Arg Glu Leu Thr Asn Leu Leu Gly Leu His Pro Lys Ile Glu Met	
885 890 895	

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gct cgt gca agc tgc tgc gcc ctg atg tct cag cct tct gcc cca gat	2736
Ala Arg Ala Ser Cys Ser Ala Leu Met Ser Gln Pro Ser Ala Pro Asp	
900 905 910	
ttt gtc cag cgg ggc ttc gat agc ttt ctg ggc cgc tac caa gct gta	2784
Phe Val Gln Arg Gly Phe Asp Ser Phe Leu Gly Arg Tyr Gln Ala Val	
915 920 925	
caa gag gct gta gag gat cgt caa caa cat cta gag aat gaa ctg aag	2832
Gln Glu Ala Val Glu Asp Arg Gln Gln His Leu Glu Asn Glu Leu Lys	
930 935 940	
ggc caa cct gga cat gca tat ctg gaa aca ttg aaa aca ctg aaa gat	2880
Gly Gln Pro Gly His Ala Tyr Leu Glu Thr Leu Lys Thr Leu Lys Asp	
945 950 955 960	
gtg cta aat gat tca gaa aat aag gcc cag gtg tct ctg aat gtc ctt	2928
Val Leu Asn Asp Ser Glu Asn Lys Ala Gln Val Ser Leu Asn Val Leu	
965 970 975	
aat gat ctt gcc aag gtg gag aag gcc ctg caa gaa aaa aag acc ctt	2976
Asn Asp Leu Ala Lys Val Glu Lys Ala Leu Gln Glu Lys Lys Thr Leu	
980 985 990	
gat gaa atc ctt gag aat cag aaa cct gca tta cat aaa ctt gca gaa	3024
Asp Glu Ile Leu Glu Asn Gln Lys Pro Ala Leu His Lys Leu Ala Glu	
995 1000 1005	
gaa aca aag gct ctg gag aaa aat gtt cat cct gat gta gaa aaa	3069
Glu Thr Lys Ala Leu Glu Lys Asn Val His Pro Asp Val Glu Lys	
1010 1015 1020	
tta tat aag caa gaa ttt gat gat gtg caa gga aag tgg aac aag	3114
Leu Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Lys Trp Asn Lys	
1025 1030 1035	
cta aag gtc ttg gtt tcc aaa gat cta cat ttg ctt gag gaa att	3159
Leu Lys Val Leu Val Ser Lys Asp Leu His Leu Leu Glu Glu Ile	
1040 1045 1050	
gct ctc aca ctc aga gct ttt gag gcc gat tca aca gtc att gag	3204
Ala Leu Thr Leu Arg Ala Phe Glu Ala Asp Ser Thr Val Ile Glu	
1055 1060 1065	
aag tgg atg gat ggc gtg aaa gac ttc tta atg aaa cag cag gct	3249
Lys Trp Met Asp Gly Val Lys Asp Phe Leu Met Lys Gln Gln Ala	
1070 1075 1080	
gcc caa gga gac gac gca ggt cta cag agg cag tta gac cag tgc	3294
Ala Gln Gly Asp Asp Ala Gly Leu Gln Arg Gln Leu Asp Gln Cys	
1085 1090 1095	
tct gca ttt gtt aat gaa ata gaa aca att gaa tca tct ctg aaa	3339
Ser Ala Phe Val Asn Glu Ile Glu Thr Ile Glu Ser Ser Leu Lys	
1100 1105 1110	
aac atg aag gaa ata gag act aat ctt cga agt ggt cca gtt gct	3384
Asn Met Lys Glu Ile Glu Thr Asn Leu Arg Ser Gly Pro Val Ala	
1115 1120 1125	
gga ata aaa act tgg gtg cag aca aga cta ggt gac tac caa act	3429
Gly Ile Lys Thr Trp Val Gln Thr Arg Leu Gly Asp Tyr Gln Thr	
1130 1135 1140	
caa ctg gag aaa ctt agc aag gag atc gct act caa aaa agt agg	3474
Gln Leu Glu Lys Leu Ser Lys Glu Ile Ala Thr Gln Lys Ser Arg	
1145 1150 1155	
ttg tct gaa agt caa gaa aaa gct gcg aac ctg aag aaa gac ttg	3519
Leu Ser Glu Ser Gln Glu Lys Ala Ala Asn Leu Lys Lys Asp Leu	
1160 1165 1170	
gca gag atg cag gaa tgg atg acc cag gcc gag gaa gaa tat ttg	3564
Ala Glu Met Gln Glu Trp Met Thr Gln Ala Glu Glu Glu Tyr Leu	
1175 1180 1185	
gag cgg gat ttt gag tac aag tca cca gaa gag ctt gag agt gct	3609
Glu Arg Asp Phe Glu Tyr Lys Ser Pro Glu Glu Leu Glu Ser Ala	

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1190	1195	1200	
gtg gaa gag atg aag agg gca Val Glu Glu Met Lys Arg Ala 1205	aaa gag gat Lys Glu Asp 1210	gtg ttg cag aag gag Val Leu Gln Lys Glu 1215	3654
gtg aga gtg aag att ctc aag Val Arg Val Lys Ile Leu Lys 1220	gac aac atc aag tta Asp Asn Ile Lys Leu 1225	tta gct gcc Leu Ala Ala 1230	3699
aag gtg ccc tct ggt ggc cag Lys Val Pro Ser Gly Gly Gln 1235	gag ttg acg tct gag Glu Leu Thr Ser Glu 1240	ctg aat gtt Leu Asn Val 1245	3744
gtg ctg gag aat tac caa ctt Val Leu Glu Asn Tyr Gln Leu 1250	ctt tgt aat aga att Leu Cys Asn Arg Ile 1255	cga gga aag Arg Gly Lys 1260	3789
tgc cac acg cta gag gag gtc Cys His Thr Leu Glu Glu Val 1265	tgg tct tgt tgg att Trp Ser Cys Trp Ile 1270	gaa ctg ctt Glu Leu Leu 1275	3834
cac tat ttg gat ctt gaa act His Tyr Leu Asp Leu Glu Thr 1280	acc tgg tta aac act Thr Trp Leu Asn Thr 1285	ttg gaa gag Leu Glu Glu 1290	3879
cgg atg aag agc aca gag gtc Arg Met Lys Ser Thr Glu Val 1295	ctg cct gag aag acg Leu Pro Glu Lys Thr 1300	gat gct gtc Asp Ala Val 1305	3924
aac gaa gcc ctg gag tct ctg Asn Glu Ala Leu Glu Ser Leu 1310	gaa tct gtt ctg cgc Glu Ser Val Leu Arg 1315	cac ccg gca His Pro Ala 1320	3969
gat aat cgc acc cag att cga Asp Asn Arg Thr Gln Ile Arg 1325	gag ctt ggc cag act Glu Leu Gly Gln Thr 1330	ctg att gat Leu Ile Asp 1335	4014
ggg ggg atc ctg gat gat ata Gly Gly Ile Leu Asp Asp Ile 1340	atc agt gag aaa ctg Ile Ser Glu Lys Leu 1345	gag gct ttc Glu Ala Phe 1350	4059
aac agc cga tat gaa gat cta Asn Ser Arg Tyr Glu Asp Leu 1355	agt cac ctg gca gag Ser His Leu Ala Glu 1360	agc aag cag Ser Lys Gln 1365	4104
att tct ttg gaa aag caa ctc Ile Ser Leu Glu Lys Gln Leu 1370	cag gtg ctg cgg gaa Gln Val Leu Arg Glu 1375	act gac cag Thr Asp Gln 1380	4149
atg ctt caa gtc ttg caa gag Met Leu Gln Val Leu Gln Glu 1385	agc ttg ggg gag ctg Ser Leu Gly Glu Leu 1390	gac aaa cag Asp Lys Gln 1395	4194
ctc acc aca tac ctg act gac Leu Thr Thr Tyr Leu Thr Asp 1400	agg ata gat gct ttc Arg Ile Asp Ala Phe 1405	caa gtt cca Gln Val Pro 1410	4239
cag gaa gct cag aaa atc caa Gln Glu Ala Gln Lys Ile Gln 1415	gca gag atc tca gcc Ala Glu Ile Ser Ala 1420	cat gag cta His Glu Leu 1425	4284
acc cta gag gag ttg aga aga Thr Leu Glu Glu Leu Arg Arg 1430	aat atg cgt tct cag Asn Met Arg Ser Gln 1435	ccc ctg acc Pro Leu Thr 1440	4329
tcc cca gag agt agg act gcc Ser Pro Glu Ser Arg Thr Ala 1445	aga gga gga agt cag Arg Gly Gly Ser Gln 1450	atg gat gtg Met Asp Val 1455	4374
cta cag agg aaa ctc cga gag Leu Gln Arg Lys Leu Arg Glu 1460	gtg tcc aca aag ttc Val Ser Thr Lys Phe 1465	cag ctt gcc Gln Leu Ala 1470	4419
cac aga gat ttt gga cca tcc His Arg Asp Phe Gly Pro Ser 1475	tct cag cat ttt ctc Ser Gln His Phe Leu 1480	tct acg tca Ser Thr Ser 1485	4464
gtc cag ctg ccg tgg caa aga 1490	tcc att tca cat aat 1495	aaa gtg ccc 1500	4509

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Val	Gln	Leu	Pro	Trp	Gln	Arg	Ser	Ile	Ser	His	Asn	Lys	Val	Pro				
	1490					1495					1500							
tat	tac	atc	aac	cat	caa	aca	cag	acc	acc	tgt	tgg	gac	cat	cct			4554	
Tyr	Tyr	Ile	Asn	His	Gln	Thr	Gln	Thr	Thr	Cys	Trp	Asp	His	Pro				
	1505					1510					1515							
aaa	atg	acc	gaa	ctc	ttt	caa	tcc	ctt	gct	gac	ctg	aat	aat	gta			4599	
Lys	Met	Thr	Glu	Leu	Phe	Gln	Ser	Leu	Ala	Asp	Leu	Asn	Asn	Val				
	1520					1525					1530							
cgt	ttt	tct	gcc	tac	cgt	aca	gca	atc	aaa	atc	cga	aga	cta	caa			4644	
Arg	Phe	Ser	Ala	Tyr	Arg	Thr	Ala	Ile	Lys	Ile	Arg	Arg	Leu	Gln				
	1535					1540					1545							
aaa	gca	cta	tgt	ttg	gat	ctc	tta	gag	ttg	agt	aca	aca	aat	gaa			4689	
Lys	Ala	Leu	Cys	Leu	Asp	Leu	Leu	Glu	Leu	Ser	Thr	Thr	Asn	Glu				
	1550					1555					1560							
att	ttc	aaa	cag	cac	aag	ttg	aac	caa	aat	gac	cag	ctc	ctc	agt			4734	
Ile	Phe	Lys	Gln	His	Lys	Leu	Asn	Gln	Asn	Asp	Gln	Leu	Leu	Ser				
	1565					1570					1575							
gtt	cca	gat	gtc	atc	aac	tgt	ctg	aca	aca	act	tat	gat	gga	ctt			4779	
Val	Pro	Asp	Val	Ile	Asn	Cys	Leu	Thr	Thr	Thr	Tyr	Asp	Gly	Leu				
	1580					1585					1590							
gag	caa	atg	cat	aag	gac	ctg	gtc	aac	ggt	cca	ctc	tgt	ggt	gat			4824	
Glu	Gln	Met	His	Lys	Asp	Leu	Val	Asn	Val	Pro	Leu	Cys	Val	Asp				
	1595					1600					1605							
atg	tgt	ctc	aat	tgg	ttg	ctc	aat	gtc	tat	gac	acg	ggt	cga	act			4869	
Met	Cys	Leu	Asn	Trp	Leu	Leu	Asn	Val	Tyr	Asp	Thr	Gly	Arg	Thr				
	1610					1615					1620							
gga	aaa	att	aga	gtg	cag	agt	ctg	aag	att	gga	tta	atg	tct	ctc			4914	
Gly	Lys	Ile	Arg	Val	Gln	Ser	Leu	Lys	Ile	Gly	Leu	Met	Ser	Leu				
	1625					1630					1635							
tcc	aaa	ggt	ctc	ttg	gaa	gaa	aaa	tac	aga	tat	ctc	ttt	aag	gaa			4959	
Ser	Lys	Gly	Leu	Leu	Glu	Glu	Lys	Tyr	Arg	Tyr	Leu	Phe	Lys	Glu				
	1640					1645					1650							
gtt	gca	ggg	cca	aca	gaa	atg	tgt	gac	cag	agg	cag	ctg	ggc	ctg			5004	
Val	Ala	Gly	Pro	Thr	Glu	Met	Cys	Asp	Gln	Arg	Gln	Leu	Gly	Leu				
	1655					1660					1665							
tta	ctt	cat	gat	gcc	atc	cag	atc	ccc	cgg	cag	cta	ggt	gaa	gta			5049	
Leu	Leu	His	Asp	Ala	Ile	Gln	Ile	Pro	Arg	Gln	Leu	Gly	Glu	Val				
	1670					1675					1680							
gca	gct	ttt	gga	ggc	agt	aat	att	gag	cct	agt	ggt	cgc	agc	tgc			5094	
Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile	Glu	Pro	Ser	Val	Arg	Ser	Cys				
	1685					1690					1695							
ttc	caa	cag	aat	aac	aat	aaa	cca	gaa	ata	agt	gtg	aaa	gag	ttt			5139	
Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro	Glu	Ile	Ser	Val	Lys	Glu	Phe				
	1700					1705					1710							
ata	gat	tgg	atg	cat	ttg	gaa	cca	cag	tcc	atg	ggt	tgg	ctc	cca			5184	
Ile	Asp	Trp	Met	His	Leu	Glu	Pro	Gln	Ser	Met	Val	Trp	Leu	Pro				
	1715					1720					1725							
gtt	tta	cat	cga	gtg	gca	gca	gcg	gag	act	gca	aaa	cat	cag	gcc			5229	
Val	Leu	His	Arg	Val	Ala	Ala	Ala	Glu	Thr	Ala	Lys	His	Gln	Ala				
	1730					1735					1740							
aaa	tgc	aac	atc	tgt	aaa	gaa	tgt	cca	att	gtc	ggg	ttc	agg	tat			5274	
Lys	Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly	Phe	Arg	Tyr				
	1745					1750					1755							
aga	agc	ctt	aag	cat	ttt	aac	tat	gat	gtc	tgc	cag	agt	tgt	ttc			5319	
Arg	Ser	Leu	Lys	His	Phe	Asn	Tyr	Asp	Val	Cys	Gln	Ser	Cys	Phe				
	1760					1765					1770							
ttt	tcg	ggt	cga	aca	gca	aaa	ggt	cac	aaa	tta	cat	tac	cca	atg			5364	
Phe	Ser	Gly	Arg	Thr	Ala	Lys	Gly	His	Lys	Leu	His	Tyr	Pro	Met				
	1775					1780					1785							

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gtg gaa tat tgt ata cct aca aca tct ggg gaa gat gta cga gac	5409
Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val Arg Asp	
1790 1795 1800	
ttc aca aag gta ctt aag aac aag ttc agg tcg aag aag tac ttt	5454
Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys Tyr Phe	
1805 1810 1815	
gcc aaa cac cct cga ctt ggt tac ctg cct gtc cag aca gtt ctt	5499
Ala Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr Val Leu	
1820 1825 1830	
gaa ggt gac aac tta gag act cct atc aca ctc atc agt atg tgg	5544
Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser Met Trp	
1835 1840 1845	
cca gag cac tat gac ccc tca caa tct cct caa ctg ttt cat gat	5589
Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe His Asp	
1850 1855 1860	
gac acc cat tca aga ata gaa caa tat gcc aca cga ctg gcc cag	5634
Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu Ala Gln	
1865 1870 1875	
atg gaa agg act aat ggg tct ttt ctc act gat agc agc tcc acc	5679
Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser Thr	
1880 1885 1890	
aca gga agt gtg gaa gac gag cac gcc ctc atc cag cag tat tgc	5724
Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln Tyr Cys	
1895 1900 1905	
caa aca ctc gga gga gag tcc cca gtg agc cag ccg cag agc cca	5769
Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln Ser Pro	
1910 1915 1920	
gct cag atc ctg aag tca gta gag agg gaa gaa cgt gga gaa ctg	5814
Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly Glu Leu	
1925 1930 1935	
gag agg atc att gct gac ctg gag gaa gaa caa aga aat cta cag	5859
Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn Leu Gln	
1940 1945 1950	
gtg gag tat gag cag ctg aag gac cag cac ctc cga agg ggg ctc	5904
Val Glu Tyr Glu Gln Leu Lys Asp Gln His Leu Arg Arg Gly Leu	
1955 1960 1965	
cct gtc ggt tca ccg cca gag tcg att ata tct ccc cat cac acg	5949
Pro Val Gly Ser Pro Pro Glu Ser Ile Ile Ser Pro His His Thr	
1970 1975 1980	
tct gag gat tca gaa ctt ata gca gaa gca aaa ctc ctc agg cag	5994
Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg Gln	
1985 1990 1995	
cac aaa ggt cgg ctg gag gct agg atg cag att tta gaa gat cac	6039
His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp His	
2000 2005 2010	
aat aaa cag ctg gag tct cag ctc cac cgc ctc cga cag ctg ctg	6084
Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu Leu	
2015 2020 2025	
gag cag cct gaa tct gat tcc cga atc aat ggt gtt tcc cca tgg	6129
Glu Gln Pro Glu Ser Asp Ser Arg Ile Asn Gly Val Ser Pro Trp	
2030 2035 2040	
gct tct cct cag cat tct gca ctg agc tac tcg ctt gat cca gat	6174
Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Pro Asp	
2045 2050 2055	
gcc tcc ggc cca cag ttc cac cag gca gcg gga gag gac ctg ctg	6219
Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp Leu Leu	
2060 2065 2070	
gcc cca ccg cac gac acc agc acg gat ctc acg gag gtc atg gag	6264
Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val Met Glu	
2075 2080 2085	

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cag att cac agc acg ttt cca tct tgc tgc cca aat gtt ccc agc 6309
 Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val Pro Ser
 2090 2095 2100

agg cca cag gca atg tga 6327
 Arg Pro Gln Ala Met
 2105

<210> SEQ ID NO 17
 <211> LENGTH: 2108
 <212> TYPE: PRT
 <213> ORGANISM: Homo sapiens

<400> SEQUENCE: 17

Met Asp Tyr Lys Asp Asp Asp Asp Lys Gly Tyr Gly Arg Lys Lys Arg
 1 5 10 15

Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20 25 30

Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Glu His Glu Ala
 35 40 45

Ser Pro Asp Asn Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50 55 60

Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65 70 75 80

Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Asn Asp Met Phe
 85 90 95

Thr Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100 105 110

Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115 120 125

Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
 130 135 140

Glu Leu Val Asn Ile Gly Gly Thr Asp Ile Val Asp Gly Asn His Lys
 145 150 155 160

Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
 165 170 175

Asp Val Met Lys Asp Val Met Ser Asp Leu Gln Gln Thr Asn Ser Glu
 180 185 190

Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln
 195 200 205

Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe
 210 215 220

Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Ser Trp Asp Lys
 225 230 235 240

Val Val Lys Met Ser Pro Ile Glu Arg Leu Glu His Ala Phe Ser Lys
 245 250 255

Ala Gln Thr Tyr Leu Gly Ile Glu Lys Leu Leu Asp Pro Glu Asp Val
 260 265 270

Ala Val Gln Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser
 275 280 285

Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu
 290 295 300

Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Ala
 305 310 315 320

Ile Asn Ile Gln Ser Thr Ala Pro Glu Glu Glu His Glu Ser Pro Arg
 325 330 335

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Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser
340 345 350

Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu
355 360 365

Asp Thr Phe Gln Glu Gln Asp Asp Ile Ser Asp Asp Val Glu Glu Val
370 375 380

Lys Asp Gln Phe Ala Thr His Glu Ala Phe Met Met Glu Leu Thr Ala
385 390 395 400

His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Ile
405 410 415

Thr Gln Gly Thr Leu Ser Asp Glu Glu Glu Phe Glu Ile Gln Glu Gln
420 425 430

Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met
435 440 445

Asp Arg Gln Ser Arg Leu His Asp Val Leu Met Glu Leu Gln Lys Lys
450 455 460

Gln Leu Gln Gln Leu Ser Ala Trp Leu Thr Leu Thr Glu Glu Arg Ile
465 470 475 480

Gln Lys Met Glu Thr Cys Pro Leu Asp Asp Asp Val Lys Ser Leu Gln
485 490 495

Lys Leu Leu Glu Glu His Lys Ser Leu Gln Ser Asp Leu Glu Ala Glu
500 505 510

Gln Val Lys Val Asn Ser Leu Thr His Met Val Val Ile Val Asp Glu
515 520 525

Asn Ser Gly Glu Ser Ala Thr Ala Ile Leu Glu Asp Gln Leu Gln Lys
530 535 540

Leu Gly Glu Arg Trp Thr Ala Val Cys Arg Trp Thr Glu Glu Arg Trp
545 550 555 560

Asn Arg Leu Gln Glu Ile Asn Ile Leu Trp Gln Glu Leu Leu Glu Glu
565 570 575

Gln Cys Leu Leu Lys Ala Trp Leu Thr Glu Lys Glu Glu Ala Leu Asn
580 585 590

Lys Val Gln Thr Ser Asn Phe Lys Asp Gln Lys Glu Leu Ser Val Ser
595 600 605

Val Arg Arg Leu Ala Ile Leu Lys Glu Asp Met Glu Met Lys Arg Gln
610 615 620

Thr Leu Asp Gln Leu Ser Glu Ile Gly Gln Asp Val Gly Gln Leu Leu
625 630 635 640

Asp Asn Ser Lys Ala Ser Lys Lys Ile Asn Ser Asp Ser Glu Glu Leu
645 650 655

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

Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln
675 680 685

Lys Asp Leu Leu Glu Thr Val Arg Val Arg Glu Gln Ala Ile Thr Lys
690 695 700

Lys Ser Lys Gln Glu Leu Pro Pro Pro Pro Pro Lys Lys Arg Gln
705 710 715 720

Ile His Val Asp Ile Glu Ala Lys Lys Lys Phe Asp Ala Ile Ser Ala
725 730 735

Glu Leu Leu Asn Trp Ile Leu Lys Trp Lys Thr Ala Ile Gln Thr Thr
740 745 750

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Glu Ile Lys Glu Tyr Met Lys Met Gln Asp Thr Ser Glu Met Lys Lys
 755 760 765
 Lys Leu Lys Ala Leu Glu Lys Glu Gln Arg Glu Arg Ile Pro Arg Ala
 770 775 780
 Asp Glu Leu Asn Gln Thr Gly Gln Ile Leu Val Glu Gln Met Gly Lys
 785 790 795 800
 Glu Gly Leu Pro Thr Glu Glu Ile Lys Asn Val Leu Glu Lys Val Ser
 805 810 815
 Ser Glu Trp Lys Asn Val Ser Gln His Leu Glu Asp Leu Glu Arg Lys
 820 825 830
 Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Glu
 835 840 845
 Leu Glu Lys Val Ile Lys Thr Lys Glu Glu Trp Val Lys His Thr Ser
 850 855 860
 Ile Ser Glu Ser Ser Arg Gln Ser Leu Pro Ser Leu Lys Asp Ser Cys
 865 870 875 880
 Gln Arg Glu Leu Thr Asn Leu Leu Gly Leu His Pro Lys Ile Glu Met
 885 890 895
 Ala Arg Ala Ser Cys Ser Ala Leu Met Ser Gln Pro Ser Ala Pro Asp
 900 905 910
 Phe Val Gln Arg Gly Phe Asp Ser Phe Leu Gly Arg Tyr Gln Ala Val
 915 920 925
 Gln Glu Ala Val Glu Asp Arg Gln Gln His Leu Glu Asn Glu Leu Lys
 930 935 940
 Gly Gln Pro Gly His Ala Tyr Leu Glu Thr Leu Lys Thr Leu Lys Asp
 945 950 955 960
 Val Leu Asn Asp Ser Glu Asn Lys Ala Gln Val Ser Leu Asn Val Leu
 965 970 975
 Asn Asp Leu Ala Lys Val Glu Lys Ala Leu Gln Glu Lys Lys Thr Leu
 980 985 990
 Asp Glu Ile Leu Glu Asn Gln Lys Pro Ala Leu His Lys Leu Ala Glu
 995 1000 1005
 Glu Thr Lys Ala Leu Glu Lys Asn Val His Pro Asp Val Glu Lys
 1010 1015 1020
 Leu Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Lys Trp Asn Lys
 1025 1030 1035
 Leu Lys Val Leu Val Ser Lys Asp Leu His Leu Leu Glu Glu Ile
 1040 1045 1050
 Ala Leu Thr Leu Arg Ala Phe Glu Ala Asp Ser Thr Val Ile Glu
 1055 1060 1065
 Lys Trp Met Asp Gly Val Lys Asp Phe Leu Met Lys Gln Gln Ala
 1070 1075 1080
 Ala Gln Gly Asp Asp Ala Gly Leu Gln Arg Gln Leu Asp Gln Cys
 1085 1090 1095
 Ser Ala Phe Val Asn Glu Ile Glu Thr Ile Glu Ser Ser Leu Lys
 1100 1105 1110
 Asn Met Lys Glu Ile Glu Thr Asn Leu Arg Ser Gly Pro Val Ala
 1115 1120 1125
 Gly Ile Lys Thr Trp Val Gln Thr Arg Leu Gly Asp Tyr Gln Thr
 1130 1135 1140
 Gln Leu Glu Lys Leu Ser Lys Glu Ile Ala Thr Gln Lys Ser Arg
 1145 1150 1155
 Leu Ser Glu Ser Gln Glu Lys Ala Ala Asn Leu Lys Lys Asp Leu

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1160	1165	1170
Ala Glu Met Gln Glu Trp Met Thr Gln Ala Glu Glu Glu Tyr Leu 1175 1180 1185		
Glu Arg Asp Phe Glu Tyr Lys Ser Pro Glu Glu Leu Glu Ser Ala 1190 1195 1200		
Val Glu Glu Met Lys Arg Ala Lys Glu Asp Val Leu Gln Lys Glu 1205 1210 1215		
Val Arg Val Lys Ile Leu Lys Asp Asn Ile Lys Leu Leu Ala Ala 1220 1225 1230		
Lys Val Pro Ser Gly Gly Gln Glu Leu Thr Ser Glu Leu Asn Val 1235 1240 1245		
Val Leu Glu Asn Tyr Gln Leu Leu Cys Asn Arg Ile Arg Gly Lys 1250 1255 1260		
Cys His Thr Leu Glu Glu Val Trp Ser Cys Trp Ile Glu Leu Leu 1265 1270 1275		
His Tyr Leu Asp Leu Glu Thr Thr Trp Leu Asn Thr Leu Glu Glu 1280 1285 1290		
Arg Met Lys Ser Thr Glu Val Leu Pro Glu Lys Thr Asp Ala Val 1295 1300 1305		
Asn Glu Ala Leu Glu Ser Leu Glu Ser Val Leu Arg His Pro Ala 1310 1315 1320		
Asp Asn Arg Thr Gln Ile Arg Glu Leu Gly Gln Thr Leu Ile Asp 1325 1330 1335		
Gly Gly Ile Leu Asp Asp Ile Ile Ser Glu Lys Leu Glu Ala Phe 1340 1345 1350		
Asn Ser Arg Tyr Glu Asp Leu Ser His Leu Ala Glu Ser Lys Gln 1355 1360 1365		
Ile Ser Leu Glu Lys Gln Leu Gln Val Leu Arg Glu Thr Asp Gln 1370 1375 1380		
Met Leu Gln Val Leu Gln Glu Ser Leu Gly Glu Leu Asp Lys Gln 1385 1390 1395		
Leu Thr Thr Tyr Leu Thr Asp Arg Ile Asp Ala Phe Gln Val Pro 1400 1405 1410		
Gln Glu Ala Gln Lys Ile Gln Ala Glu Ile Ser Ala His Glu Leu 1415 1420 1425		
Thr Leu Glu Glu Leu Arg Arg Asn Met Arg Ser Gln Pro Leu Thr 1430 1435 1440		
Ser Pro Glu Ser Arg Thr Ala Arg Gly Gly Ser Gln Met Asp Val 1445 1450 1455		
Leu Gln Arg Lys Leu Arg Glu Val Ser Thr Lys Phe Gln Leu Ala 1460 1465 1470		
His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr Ser 1475 1480 1485		
Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn Lys Val Pro 1490 1495 1500		
Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp His Pro 1505 1510 1515		
Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn Val 1520 1525 1530		
Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu Gln 1535 1540 1545		
Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Ser Thr Thr Asn Glu 1550 1555 1560		

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Ile	Phe	Lys	Gln	His	Lys	Leu	Asn	Gln	Asn	Asp	Gln	Leu	Leu	Ser
1565						1570					1575			
Val	Pro	Asp	Val	Ile	Asn	Cys	Leu	Thr	Thr	Thr	Tyr	Asp	Gly	Leu
1580						1585					1590			
Glu	Gln	Met	His	Lys	Asp	Leu	Val	Asn	Val	Pro	Leu	Cys	Val	Asp
1595						1600					1605			
Met	Cys	Leu	Asn	Trp	Leu	Leu	Asn	Val	Tyr	Asp	Thr	Gly	Arg	Thr
1610						1615					1620			
Gly	Lys	Ile	Arg	Val	Gln	Ser	Leu	Lys	Ile	Gly	Leu	Met	Ser	Leu
1625						1630					1635			
Ser	Lys	Gly	Leu	Leu	Glu	Glu	Lys	Tyr	Arg	Tyr	Leu	Phe	Lys	Glu
1640						1645					1650			
Val	Ala	Gly	Pro	Thr	Glu	Met	Cys	Asp	Gln	Arg	Gln	Leu	Gly	Leu
1655						1660					1665			
Leu	Leu	His	Asp	Ala	Ile	Gln	Ile	Pro	Arg	Gln	Leu	Gly	Glu	Val
1670						1675					1680			
Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile	Glu	Pro	Ser	Val	Arg	Ser	Cys
1685						1690					1695			
Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro	Glu	Ile	Ser	Val	Lys	Glu	Phe
1700						1705					1710			
Ile	Asp	Trp	Met	His	Leu	Glu	Pro	Gln	Ser	Met	Val	Trp	Leu	Pro
1715						1720					1725			
Val	Leu	His	Arg	Val	Ala	Ala	Ala	Glu	Thr	Ala	Lys	His	Gln	Ala
1730						1735					1740			
Lys	Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly	Phe	Arg	Tyr
1745						1750					1755			
Arg	Ser	Leu	Lys	His	Phe	Asn	Tyr	Asp	Val	Cys	Gln	Ser	Cys	Phe
1760						1765					1770			
Phe	Ser	Gly	Arg	Thr	Ala	Lys	Gly	His	Lys	Leu	His	Tyr	Pro	Met
1775						1780					1785			
Val	Glu	Tyr	Cys	Ile	Pro	Thr	Thr	Ser	Gly	Glu	Asp	Val	Arg	Asp
1790						1795					1800			
Phe	Thr	Lys	Val	Leu	Lys	Asn	Lys	Phe	Arg	Ser	Lys	Lys	Tyr	Phe
1805						1810					1815			
Ala	Lys	His	Pro	Arg	Leu	Gly	Tyr	Leu	Pro	Val	Gln	Thr	Val	Leu
1820						1825					1830			
Glu	Gly	Asp	Asn	Leu	Glu	Thr	Pro	Ile	Thr	Leu	Ile	Ser	Met	Trp
1835						1840					1845			
Pro	Glu	His	Tyr	Asp	Pro	Ser	Gln	Ser	Pro	Gln	Leu	Phe	His	Asp
1850						1855					1860			
Asp	Thr	His	Ser	Arg	Ile	Glu	Gln	Tyr	Ala	Thr	Arg	Leu	Ala	Gln
1865						1870					1875			
Met	Glu	Arg	Thr	Asn	Gly	Ser	Phe	Leu	Thr	Asp	Ser	Ser	Ser	Thr
1880						1885					1890			
Thr	Gly	Ser	Val	Glu	Asp	Glu	His	Ala	Leu	Ile	Gln	Gln	Tyr	Cys
1895						1900					1905			
Gln	Thr	Leu	Gly	Gly	Glu	Ser	Pro	Val	Ser	Gln	Pro	Gln	Ser	Pro
1910						1915					1920			
Ala	Gln	Ile	Leu	Lys	Ser	Val	Glu	Arg	Glu	Glu	Arg	Gly	Glu	Leu
1925						1930					1935			
Glu	Arg	Ile	Ile	Ala	Asp	Leu	Glu	Glu	Glu	Gln	Arg	Asn	Leu	Gln
1940						1945					1950			

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Val Glu Tyr Glu Gln Leu Lys Asp Gln His Leu Arg Arg Gly Leu
 1955 1960 1965

Pro Val Gly Ser Pro Pro Glu Ser Ile Ile Ser Pro His His Thr
 1970 1975 1980

Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg Gln
 1985 1990 1995

His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp His
 2000 2005 2010

Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu Leu
 2015 2020 2025

Glu Gln Pro Glu Ser Asp Ser Arg Ile Asn Gly Val Ser Pro Trp
 2030 2035 2040

Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Pro Asp
 2045 2050 2055

Ala Ser Gly Pro Gln Phe His Gln Ala Ala Gly Glu Asp Leu Leu
 2060 2065 2070

Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Glu Val Met Glu
 2075 2080 2085

Gln Ile His Ser Thr Phe Pro Ser Cys Cys Pro Asn Val Pro Ser
 2090 2095 2100

Arg Pro Gln Ala Met
 2105

<210> SEQ ID NO 18
 <211> LENGTH: 4080
 <212> TYPE: DNA
 <213> ORGANISM: Mus musculus
 <220> FEATURE:
 <221> NAME/KEY: CDS
 <222> LOCATION: (1)..(4080)
 <220> FEATURE:
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 <223> OTHER INFORMATION: TAT and epitope tag coding sequence

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cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac	96
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp	
20 25 30	
gtc cca gac tat gct ggc tcc atg gcc aag tat ggg gac ctt gaa gcc	144
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala	
35 40 45	
agg cct gat gat ggg cag aac gaa ttc agt gac atc att aag tcc aga	192
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg	
50 55 60	
tct gat gaa cac aat gat gta cag aag aaa acc ttt acc aaa tgg ata	240
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile	
65 70 75 80	
aac gct cga ttt tcc aag agt ggg aaa cca ccc atc agt gat atg ttc	288
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe	
85 90 95	
tca gac ctc aaa gat ggg aga aag ctc ttg gat ctt ctc gaa ggc ctc	336
Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu	
100 105 110	
aca gga aca tca ttg cca aag gaa cgt ggt tcc aca agg gtg cat gcc	384
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala	
115 120 125	

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tta aac aat gtc aac cga gtg cta cag gtt tta cat cag aac aat gtg	432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val	
130 135 140	
gac ttg gtg aat att gga ggc acg gac att gtg gct gga aat ccc aag	480
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys	
145 150 155 160	
ctg act tta ggg tta ctc tgg agc atc att ctg cac tgg cag gtg aag	528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys	
165 170 175	
gat gtc atg aaa gat atc atg tca gac ctg cag cag aca aac agc gag	576
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu	
180 185 190	
aag atc ctg ctg agc tgg gtg cgg cag acc acc agg ccc tac agt caa	624
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln	
195 200 205	
gtc aac gtc ctc aac ttc acc acc agc tgg acc gat gga ctc gcg ttc	672
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe	
210 215 220	
aac gcc gtg ctc cac cgg cac aaa cca gat ctc ttc gac tgg gac gag	720
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu	
225 230 235 240	
atg gtc aaa atg tcc cca att gag aga ctt gac cat gct ttt gac aag	768
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys	
245 250 255	
gcc cac act tct ttg gga att gaa aag ctc cta agt cct gaa act gtt	816
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val	
260 265 270	
gct gtg cat ctc cct gac aag aaa tcc ata att atg tat tta acg tct	864
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser	
275 280 285	
ctg ttt gag gtg ctt cct cag caa gtc acg ata gat gcc atc cga gag	912
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu	
290 295 300	
gtg gag act ctc cca agg aag tat aag aaa gaa tgt gaa gag gaa gaa	960
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu	
305 310 315 320	
att cat atc cag agt gca gtg ctg gca gag gaa ggc cag agt ccc cga	1008
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg	
325 330 335	
gct gag acc cct agc acc gtc act gaa gtg gac atg gat ttg gac agc	1056
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser	
340 345 350	
tac cag ata gcg cta gag gaa gtg ctg acg tgg ctg ctg tcc gcg gag	1104
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu	
355 360 365	
gac acg ttc cag gag caa cat gac att tct gat gat gtc gaa gaa gtc	1152
Asp Thr Phe Gln Glu Gln His Asp Ile Ser Asp Asp Val Glu Glu Val	
370 375 380	
aaa gag cag ttt gct acc cat gaa act ttt atg atg gag ctg aca gca	1200
Lys Glu Gln Phe Ala Thr His Glu Thr Phe Met Met Glu Leu Thr Ala	
385 390 395 400	
cac cag agc agc gtg ggg agc gtc ctg cag gct ggc aac cag ctg atg	1248
His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Met	
405 410 415	
aca caa ggg act ctg tcc aga gag gag gag ttt gag atc cag gaa cag	1296
Thr Gln Gly Thr Leu Ser Arg Glu Glu Glu Phe Glu Ile Gln Glu Gln	
420 425 430	
atg acc ttg ctg aat gca agg tgg gag gcg ctc cgg gtg gag agc atg	1344
Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met	

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gag agg cag tcc cgg ctg cac gac gct ctg atg gag ctg cag aag aaa	1392
Glu Arg Gln Ser Arg Leu His Asp Ala Leu Met Glu Leu Gln Lys Lys	
450 455 460	
cag ctg cag cag ctc tca agc tgg ctg gcc ctc aca gaa gag cgc att	1440
Gln Leu Gln Gln Leu Ser Ser Trp Leu Ala Leu Thr Glu Glu Arg Ile	
465 470 475 480	
cag aag atg gag agc ctc ccg ctg ggt gat gac ctg ccc tcc ctg cag	1488
Gln Lys Met Glu Ser Leu Pro Leu Gly Asp Asp Leu Pro Ser Leu Gln	
485 490 495	
aag ctg ctt caa gaa cat aaa agt ttg caa aat gac ctt gaa gct gaa	1536
Lys Leu Leu Gln Glu His Lys Ser Leu Gln Asn Asp Leu Glu Ala Glu	
500 505 510	
cag gtg aag gta aat tcc tta act cac atg gtg gtg att gtg gat gaa	1584
Gln Val Lys Val Asn Ser Leu Thr His Met Val Val Ile Val Asp Glu	
515 520 525	
aac agt ggg gag agt gcc aca gct ctt ctg gaa gat cag tta cag aaa	1632
Asn Ser Gly Glu Ser Ala Thr Ala Leu Leu Glu Asp Gln Leu Gln Lys	
530 535 540	
ctg ggt gag cgc tgg aca gct gta tgc cgc tgg act gaa gaa cgt tgg	1680
Leu Gly Glu Arg Trp Thr Ala Val Cys Arg Trp Thr Glu Glu Arg Trp	
545 550 555 560	
aac agg ttg caa gaa atc agt att ctg tgg cag gaa tta ttg gaa gag	1728
Asn Arg Leu Gln Glu Ile Ser Ile Leu Trp Gln Glu Leu Leu Glu Glu	
565 570 575	
cag tgt ctg ttg gag gct tgg ctc acc gaa aag gaa gag gct ttg gat	1776
Gln Cys Leu Leu Glu Ala Trp Leu Thr Glu Lys Glu Glu Ala Leu Asp	
580 585 590	
aaa gtt caa acc agc aac ttt aaa gac cag aag gaa cta agt gtc agt	1824
Lys Val Gln Thr Ser Asn Phe Lys Asp Gln Lys Glu Leu Ser Val Ser	
595 600 605	
gtc cgg cgt ctg gct ata ttg aag gaa gac atg gaa atg aag agg cag	1872
Val Arg Arg Leu Ala Ile Leu Lys Glu Asp Met Glu Met Lys Arg Gln	
610 615 620	
act ctg gat caa ctg agt gag att ggc cag gat gtg ggc caa tta ctc	1920
Thr Leu Asp Gln Leu Ser Glu Ile Gly Gln Asp Val Gly Gln Leu Leu	
625 630 635 640	
agt aat ccc aag gca tct aag aag atg aac agt gac tct gag gag cta	1968
Ser Asn Pro Lys Ala Ser Lys Lys Met Asn Ser Asp Ser Glu Glu Leu	
645 650 655	
aca cag aga tgg gat tct ctg gtt cag aga ctc gaa gac tct tct aac	2016
Thr Gln Arg Trp Asp Ser Leu Val Gln Arg Leu Glu Asp Ser Ser Asn	
660 665 670	
cag gtg act cag gcg gta gcg aag ctc ggc atg tcc cag att cca cag	2064
Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln	
675 680 685	
aag gac cta ttg gag acc gtt cat gtg aga gaa caa ggg atg gtg aag	2112
Lys Asp Leu Leu Glu Thr Val His Val Arg Glu Gln Gly Met Val Lys	
690 695 700	
aag ccc aag cag gaa ctg cct cct cct ccc cca cca aag aag aga cag	2160
Lys Pro Lys Gln Glu Leu Pro Pro Pro Pro Pro Pro Lys Lys Arg Gln	
705 710 715 720	
att cac gtg gac gcc cac aga gat ttt ggg cca tct tct caa cac ttt	2208
Ile His Val Asp Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe	
725 730 735	
ctg tcc act tca gtc cag ctg ccg tgg cag aga tcc att tca cat aat	2256
Leu Ser Thr Ser Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn	
740 745 750	
aaa gtg ccc tat tac atc aac cat caa aca cag aca acc tgt tgg gat	2304

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Lys Val	Pro Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp	
	755 760 765	
cat cct aaa atg act gag ctc ttc caa tcc ctt gct gat ctg aat aat		2352
His Pro Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn		
770 775 780		
gta cgt ttc tct gcc tac cgc aca gca atc aaa att cga agg ctg caa		2400
Val Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu Gln		
785 790 795 800		
aaa gca tta tgt ctg gat ctc tta gag ctg aat acg acg aat gaa gtt		2448
Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Asn Thr Thr Asn Glu Val		
805 810 815		
ttc aag cag cac aaa ctg aac caa aat gat cag ctc ctg agt gtc cca		2496
Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu Ser Val Pro		
820 825 830		
gac gtc atc aac tgt ctg acc acc act tac gat ggg ctt gag cag ctg		2544
Asp Val Ile Asn Cys Leu Thr Thr Tyr Asp Gly Leu Glu Gln Leu		
835 840 845		
cac aag gac ttg gtc aat gtt cca ctc tgc gtc gat atg tgt ctc aac		2592
His Lys Asp Leu Val Asn Val Pro Leu Cys Val Asp Met Cys Leu Asn		
850 855 860		
tgg ctg ctc aac gta tac gac acg ggc cgg act gga aaa att cgg gta		2640
Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg Thr Gly Lys Ile Arg Val		
865 870 875 880		
cag agt ctg aag att gga ttg atg tct ctc tcc aaa ggc ctc tta gaa		2688
Gln Ser Leu Lys Ile Gly Leu Met Ser Leu Ser Lys Gly Leu Leu Glu		
885 890 895		
gag aaa tac aga tgt ctc ttt aag gag gtg gca ggg cca act gag atg		2736
Glu Lys Tyr Arg Cys Leu Phe Lys Glu Val Ala Gly Pro Thr Glu Met		
900 905 910		
tgt gac cag cgg cag ctt ggc ctg cta ctt cac gat gcc atc cag atc		2784
Cys Asp Gln Arg Gln Leu Gly Leu Leu Leu His Asp Ala Ile Gln Ile		
915 920 925		
cct agg cag ctg ggg gaa gta gca gcc ttt ggg ggc agt aac att gag		2832
Pro Arg Gln Leu Gly Glu Val Ala Ala Phe Gly Gly Ser Asn Ile Glu		
930 935 940		
ccc agt gtc cgc agc tgc ttc cag cag aat aac aac aag cca gaa atc		2880
Pro Ser Val Arg Ser Cys Phe Gln Gln Asn Asn Asn Lys Pro Glu Ile		
945 950 955 960		
agt gtg aag gag ttt ata gac tgg atg cat ttg gaa ccc cag tcc atg		2928
Ser Val Lys Glu Phe Ile Asp Trp Met His Leu Glu Pro Gln Ser Met		
965 970 975		
gtg tgg ttg ccg gtt ctg cat cgg gtc gca gct gct gag act gca aaa		2976
Val Trp Leu Pro Val Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys		
980 985 990		
cat cag gcc aaa tgc aac atc tgc aaa gaa tgc ccg att gtt ggg ttc		3024
His Gln Ala Lys Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe		
995 1000 1005		
aga tac agg agc cta aag cat ttt aat tat gat gtc tgc cag agt		3069
Arg Tyr Arg Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser		
1010 1015 1020		
tgc ttc ttt tct gga aga aca gca aag ggc cac aag tta cat tac		3114
Cys Phe Phe Ser Gly Arg Thr Ala Lys Gly His Lys Leu His Tyr		
1025 1030 1035		
ccg atg gta gaa tac tgc ata ccg aca aca tct ggg gaa gat gtg		3159
Pro Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val		
1040 1045 1050		
aga gat ttc act aag gtg ctg aag aac aag ttc agg tcc aag aaa		3204
Arg Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys		
1055 1060 1065		

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<210> SEQ ID NO 19
<211> LENGTH: 1359
<212> TYPE: PRT
<213> ORGANISM: Mus musculus

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Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20          25          30
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala
 35          40          45
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50          55          60
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65          70          75          80
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe
 85          90          95
Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100         105         110
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115         120         125
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val
 130         135         140
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys
 145         150         155         160
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys
 165         170         175
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu
 180         185         190
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln
 195         200         205
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe
 210         215         220
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu
 225         230         235         240
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys
 245         250         255
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val
 260         265         270
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser
 275         280         285
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu
 290         295         300
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu
 305         310         315         320
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg
 325         330         335
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser
 340         345         350
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu
 355         360         365
Asp Thr Phe Gln Glu Gln His Asp Ile Ser Asp Asp Val Glu Glu Val

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370					375					380					
Lys	Glu	Gln	Phe	Ala	Thr	His	Glu	Thr	Phe	Met	Met	Glu	Leu	Thr	Ala
385					390					395					400
His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Met
				405						410					415
Thr	Gln	Gly	Thr	Leu	Ser	Arg	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln
			420							425					430
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met
		435					440					445			
Glu	Arg	Gln	Ser	Arg	Leu	His	Asp	Ala	Leu	Met	Glu	Leu	Gln	Lys	Lys
	450					455					460				
Gln	Leu	Gln	Gln	Leu	Ser	Ser	Trp	Leu	Ala	Leu	Thr	Glu	Glu	Arg	Ile
465				470							475				480
Gln	Lys	Met	Glu	Ser	Leu	Pro	Leu	Gly	Asp	Asp	Leu	Pro	Ser	Leu	Gln
				485					490						495
Lys	Leu	Leu	Gln	Glu	His	Lys	Ser	Leu	Gln	Asn	Asp	Leu	Glu	Ala	Glu
			500						505					510	
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu
		515					520					525			
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Leu	Leu	Glu	Asp	Gln	Leu	Gln	Lys
	530					535					540				
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp
545				550						555					560
Asn	Arg	Leu	Gln	Glu	Ile	Ser	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu
				565					570					575	
Gln	Cys	Leu	Leu	Glu	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asp
			580					585						590	
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser
		595					600					605			
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln
	610					615					620				
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu
625				630						635					640
Ser	Asn	Pro	Lys	Ala	Ser	Lys	Lys	Met	Asn	Ser	Asp	Ser	Glu	Glu	Leu
				645					650					655	
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn
			660						665					670	
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln
		675					680					685			
Lys	Asp	Leu	Leu	Glu	Thr	Val	His	Val	Arg	Glu	Gln	Gly	Met	Val	Lys
	690					695					700				
Lys	Pro	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln
705				710						715					720
Ile	His	Val	Asp	Ala	His	Arg	Asp	Phe	Gly	Pro	Ser	Ser	Gln	His	Phe
				725					730					735	
Leu	Ser	Thr	Ser	Val	Gln	Leu	Pro	Trp	Gln	Arg	Ser	Ile	Ser	His	Asn
			740					745					750		
Lys	Val	Pro	Tyr	Tyr	Ile	Asn	His	Gln	Thr	Gln	Thr	Thr	Cys	Trp	Asp
		755					760					765			
His	Pro	Lys	Met	Thr	Glu	Leu	Phe	Gln	Ser	Leu	Ala	Asp	Leu	Asn	Asn
	770					775					780				
Val	Arg	Phe	Ser	Ala	Tyr	Arg	Thr	Ala	Ile	Lys	Ile	Arg	Arg	Leu	Gln
785				790						795					800

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Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Asn Thr Thr Asn Glu Val
 805 810 815

Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu Ser Val Pro
 820 825 830

Asp Val Ile Asn Cys Leu Thr Thr Thr Tyr Asp Gly Leu Glu Gln Leu
 835 840 845

His Lys Asp Leu Val Asn Val Pro Leu Cys Val Asp Met Cys Leu Asn
 850 855 860

Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg Thr Gly Lys Ile Arg Val
 865 870 875 880

Gln Ser Leu Lys Ile Gly Leu Met Ser Leu Ser Lys Gly Leu Leu Glu
 885 890 895

Glu Lys Tyr Arg Cys Leu Phe Lys Glu Val Ala Gly Pro Thr Glu Met
 900 905 910

Cys Asp Gln Arg Gln Leu Gly Leu Leu Leu His Asp Ala Ile Gln Ile
 915 920 925

Pro Arg Gln Leu Gly Glu Val Ala Ala Phe Gly Gly Ser Asn Ile Glu
 930 935 940

Pro Ser Val Arg Ser Cys Phe Gln Gln Asn Asn Asn Lys Pro Glu Ile
 945 950 955 960

Ser Val Lys Glu Phe Ile Asp Trp Met His Leu Glu Pro Gln Ser Met
 965 970 975

Val Trp Leu Pro Val Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys
 980 985 990

His Gln Ala Lys Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe
 995 1000 1005

Arg Tyr Arg Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser
 1010 1015 1020

Cys Phe Phe Ser Gly Arg Thr Ala Lys Gly His Lys Leu His Tyr
 1025 1030 1035

Pro Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val
 1040 1045 1050

Arg Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys
 1055 1060 1065

Tyr Phe Ala Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr
 1070 1075 1080

Val Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser
 1085 1090 1095

Met Trp Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe
 1100 1105 1110

His Asp Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu
 1115 1120 1125

Ala Gln Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser
 1130 1135 1140

Ser Thr Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln
 1145 1150 1155

Tyr Cys Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln
 1160 1165 1170

Ser Pro Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly
 1175 1180 1185

Glu Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn
 1190 1195 1200

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Leu Gln Val Glu Tyr Glu Gln Leu Lys Glu Gln His Leu Arg Arg
 1205 1210 1215
 Gly Leu Pro Val Gly Ser Pro Pro Asp Ser Ile Val Ser Pro His
 1220 1225 1230
 His Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu
 1235 1240 1245
 Arg Gln His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu
 1250 1255 1260
 Asp His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln
 1265 1270 1275
 Leu Leu Glu Gln Pro Asp Ser Asp Ser Arg Ile Asn Gly Val Ser
 1280 1285 1290
 Pro Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp
 1295 1300 1305
 Thr Asp Pro Gly Pro Gln Phe His Gln Ala Ala Ser Glu Asp Leu
 1310 1315 1320
 Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Asp Val Met
 1325 1330 1335
 Glu Gln Ile Asn Ser Thr Phe Pro Ser Cys Ser Ser Asn Val Pro
 1340 1345 1350
 Ser Arg Pro Gln Ala Met
 1355

<210> SEQ ID NO 20
 <211> LENGTH: 5067
 <212> TYPE: DNA
 <213> ORGANISM: Mus musculus
 <220> FEATURE:
 <221> NAME/KEY: CDS
 <222> LOCATION: (1)..(5067)
 <220> FEATURE:
 <221> NAME/KEY: misc_feature
 <222> LOCATION: (1)..(117)
 <223> OTHER INFORMATION: TAT and epitope tag coding sequence

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 cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac 96
 Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp
 20 25 30
 gtc cca gac tat gct ggc tcc atg gcc aag tat ggg gac ctt gaa gcc 144
 Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala
 35 40 45
 agg cct gat gat ggg cag aac gaa ttc agt gac atc att aag tcc aga 192
 Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg
 50 55 60
 tct gat gaa cac aat gat gta cag aag aaa acc ttt acc aaa tgg ata 240
 Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile
 65 70 75 80
 aac gct cga ttt tcc aag agt ggg aaa cca ccc atc agt gat atg ttc 288
 Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe
 85 90 95
 tca gac ctc aaa gat ggg aga aag ctc ttg gat ctt ctc gaa ggc ctc 336
 Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu
 100 105 110
 aca gga aca tca ttg cca aag gaa cgt ggt tcc aca agg gtg cat gcc 384
 Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala
 115 120 125

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tta aac aat gtc aac cga gtg cta cag gtt tta cat cag aac aat gtg	432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val	
130 135 140	
gac ttg gtg aat att gga ggc acg gac att gtg gct gga aat ccc aag	480
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys	
145 150 155 160	
ctg act tta ggg tta ctc tgg agc atc att ctg cac tgg cag gtg aag	528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys	
165 170 175	
gat gtc atg aaa gat atc atg tca gac ctg cag cag aca aac agc gag	576
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu	
180 185 190	
aag atc ctg ctg agc tgg gtg cgg cag acc acc agg ccc tac agt caa	624
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln	
195 200 205	
gtc aac gtc ctc aac ttc acc acc agc tgg acc gat gga ctc gcg ttc	672
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe	
210 215 220	
aac gcc gtg ctc cac cgg cac aaa cca gat ctc ttc gac tgg gac gag	720
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu	
225 230 235 240	
atg gtc aaa atg tcc cca att gag aga ctt gac cat gct ttt gac aag	768
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys	
245 250 255	
gcc cac act tct ttg gga att gaa aag ctc cta agt cct gaa act gtt	816
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val	
260 265 270	
gct gtg cat ctc cct gac aag aaa tcc ata att atg tat tta acg tct	864
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser	
275 280 285	
ctg ttt gag gtg ctt cct cag caa gtc acg ata gat gcc atc cga gag	912
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu	
290 295 300	
gtg gag act ctc cca agg aag tat aag aaa gaa tgt gaa gag gaa gaa	960
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu	
305 310 315 320	
att cat atc cag agt gca gtg ctg gca gag gaa ggc cag agt ccc cga	1008
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg	
325 330 335	
gct gag acc cct agc acc gtc act gaa gtg gac atg gat ttg gac agc	1056
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser	
340 345 350	
tac cag ata gcg cta gag gaa gtg ctg acg tgg ctg ctg tcc gcg gag	1104
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu	
355 360 365	
gac acg ttc cag gag caa cat gac att tct gat gat gtc gaa gaa gtc	1152
Asp Thr Phe Gln Glu Gln His Asp Ile Ser Asp Asp Val Glu Glu Val	
370 375 380	
aaa gag cag ttt gct acc cat gaa act ttt atg atg gag ctg aca gca	1200
Lys Glu Gln Phe Ala Thr His Glu Thr Phe Met Met Glu Leu Thr Ala	
385 390 395 400	
cac cag agc agc gtg ggg agc gtc ctg cag gct ggc aac cag ctg atg	1248
His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Met	
405 410 415	
aca caa ggg act ctg tcc aga gag gag gag ttt gag atc cag gaa cag	1296
Thr Gln Gly Thr Leu Ser Arg Glu Glu Glu Phe Glu Ile Gln Glu Gln	
420 425 430	
atg acc ttg ctg aat gca agg tgg gag gcg ctc cgg gtg gag agc atg	1344
Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met	

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cag ctg cag cag ctc tca agc tgg ctg gcc ctc aca gaa gag cgc att Gln Leu Gln Gln Leu Ser Ser Trp Leu Ala Leu Thr Glu Glu Arg Ile 465 470 475 480	1440
cag aag atg gag agc ctc ccg ctg ggt gat gac ctg ccc tcc ctg cag Gln Lys Met Glu Ser Leu Pro Leu Gly Asp Asp Leu Pro Ser Leu Gln 485 490 495	1488
aag ctg ctt caa gaa cat aaa agt ttg caa aat gac ctt gaa gct gaa Lys Leu Leu Gln Glu His Lys Ser Leu Gln Asn Asp Leu Glu Ala Glu 500 505 510	1536
cag gtg aag gta aat tcc tta act cac atg gtg gtg att gtg gat gaa Gln Val Lys Val Asn Ser Leu Thr His Met Val Val Ile Val Asp Glu 515 520 525	1584
aac agt ggg gag agt gcc aca gct ctt ctg gaa gat cag tta cag aaa Asn Ser Gly Glu Ser Ala Thr Ala Leu Leu Glu Asp Gln Leu Gln Lys 530 535 540	1632
ctg ggt gag cgc tgg aca gct gta tgc cgc tgg act gaa gaa cgt tgg Leu Gly Glu Arg Trp Thr Ala Val Cys Arg Trp Thr Glu Glu Arg Trp 545 550 555 560	1680
aac agg ttg caa gaa atc agt att ctg tgg cag gaa tta ttg gaa gag Asn Arg Leu Gln Glu Ile Ser Ile Leu Trp Gln Glu Leu Leu Glu Glu 565 570 575	1728
cag tgt ctg ttg gag gct tgg ctc acc gaa aag gaa gag gct ttg gat Gln Cys Leu Leu Glu Ala Trp Leu Thr Glu Lys Glu Glu Ala Leu Asp 580 585 590	1776
aaa gtt caa acc agc aac ttt aaa gac cag aag gaa cta agt gtc agt Lys Val Gln Thr Ser Asn Phe Lys Asp Gln Lys Glu Leu Ser Val Ser 595 600 605	1824
gtc cgg cgt ctg gct ata ttg aag gaa gac atg gaa atg aag agg cag Val Arg Arg Leu Ala Ile Leu Lys Glu Asp Met Glu Met Lys Arg Gln 610 615 620	1872
act ctg gat caa ctg agt gag att ggc cag gat gtg ggc caa tta ctc Thr Leu Asp Gln Leu Ser Glu Ile Gly Gln Asp Val Gly Gln Leu Leu 625 630 635 640	1920
agt aat ccc aag gca tct aag aag atg aac agt gac tct gag gag cta Ser Asn Pro Lys Ala Ser Lys Lys Met Asn Ser Asp Ser Glu Glu Leu 645 650 655	1968
aca cag aga tgg gat tct ctg gtt cag aga ctc gaa gac tct tct aac Thr Gln Arg Trp Asp Ser Leu Val Gln Arg Leu Glu Asp Ser Ser Asn 660 665 670	2016
cag gtg act cag gcg gta gcg aag ctc ggc atg tcc cag att cca cag Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln 675 680 685	2064
aag gac cta ttg gag acc gtt cat gtg aga gaa caa ggg atg gtg aag Lys Asp Leu Leu Glu Thr Val His Val Arg Glu Gln Gly Met Val Lys 690 695 700	2112
aag ccc aag cag gaa ctg cct cct cct ccc cca cca aag aag aga cag Lys Pro Lys Gln Glu Leu Pro Pro Pro Pro Pro Pro Lys Lys Arg Gln 705 710 715 720	2160
att cac gtg gac gtg gag gcc aag aaa aag ttt gat gct ata agt aca Ile His Val Asp Val Glu Ala Lys Lys Lys Phe Asp Ala Ile Ser Thr 725 730 735	2208
gag ctg ctg aac tgg att ttg aaa tca aag act gcc att cag aac aca Glu Leu Leu Asn Trp Ile Leu Lys Ser Lys Thr Ala Ile Gln Asn Thr 740 745 750	2256
gag atg aaa gaa tat aag aag tcg cag gag acc tca gga atg aaa aag	2304

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Glu Met	Lys Glu Tyr Lys Lys Ser Gln Glu Thr Ser Gly Met Lys Lys	
	755 760 765	
aaa ttg aag gga tta gag aaa gaa cag aag gaa aat ctg ccc cga ctg		2352
Lys Leu Lys Gly Leu Glu Lys Glu Gln Lys Glu Asn Leu Pro Arg Leu		
	770 775 780	
gac gaa ctg aat caa acc gga caa acc ctc cgg gag caa atg gga aaa		2400
Asp Glu Leu Asn Gln Thr Gly Gln Thr Leu Arg Glu Gln Met Gly Lys		
	785 790 795 800	
gaa ggc ctt cca ctg aaa gaa gta aac gat gtt ctg gaa agg gtt tcg		2448
Glu Gly Leu Pro Leu Lys Glu Val Asn Asp Val Leu Glu Arg Val Ser		
	805 810 815	
ttg gag tgg aag atg ata tct cag cag cta gaa gat ctg gga agg aag		2496
Leu Glu Trp Lys Met Ile Ser Gln Gln Leu Glu Asp Leu Gly Arg Lys		
	820 825 830	
atc cag ctg cag gaa gat ata aat gct tat ttt aag cag ctt gat gcc		2544
Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Ala		
	835 840 845	
att gag gag acc atc aag gag aag gaa gag tgg ctg agg ggc aca ccc		2592
Ile Glu Glu Thr Ile Lys Glu Lys Glu Glu Trp Leu Arg Gly Thr Pro		
	850 855 860	
att tct gaa tcg ccc cgg cag ccc ttg cca ggc tta aag gat tct tgc		2640
Ile Ser Glu Ser Pro Arg Gln Pro Leu Pro Gly Leu Lys Asp Ser Cys		
	865 870 875 880	
cag agg gaa ctg aca gat ctc ctt ggc ctt cac ccc aga att gag acg		2688
Gln Arg Glu Leu Thr Asp Leu Leu Gly Leu His Pro Arg Ile Glu Thr		
	885 890 895	
ctg tgt gca agc tgt tca gcc ctg aag tct cag ccc tgt gtc cca ggt		2736
Leu Cys Ala Ser Cys Ser Ala Leu Lys Ser Gln Pro Cys Val Pro Gly		
	900 905 910	
ttt gtc cag cag ggt ttt gac gac ctt cga cat cat tac cag gct gtt		2784
Phe Val Gln Gln Gly Phe Asp Asp Leu Arg His His Tyr Gln Ala Val		
	915 920 925	
gcg aag gct tta gag gaa tac caa caa caa cta gaa aat gag ctg aag		2832
Ala Lys Ala Leu Glu Glu Tyr Gln Gln Gln Leu Glu Asn Glu Leu Lys		
	930 935 940	
agc cag cct gga ccc gag tat ttg gac aca ctg aat acc ctg aaa aaa		2880
Ser Gln Pro Gly Pro Glu Tyr Leu Asp Thr Leu Asn Thr Leu Lys Lys		
	945 950 955 960	
atg cta agc gag tca gaa aag gcg gcc cag gcc tct ctg aat gcc ctg		2928
Met Leu Ser Glu Ser Glu Lys Ala Ala Gln Ala Ser Leu Asn Ala Leu		
	965 970 975	
aac gat ccc ata gcg gtg gag cag gcc ctg cag gag aaa aag gcc ctt		2976
Asn Asp Pro Ile Ala Val Glu Gln Ala Leu Gln Glu Lys Lys Ala Leu		
	980 985 990	
gat gaa acc ctt gag aat cag aaa cat acg tta cat aag ctt tca gaa		3024
Asp Glu Thr Leu Glu Asn Gln Lys His Thr Leu His Lys Leu Ser Glu		
	995 1000 1005	
gaa acg aag act ttg gag aaa aat atg ctt cct gat gtg ggg aaa		3069
Glu Thr Lys Thr Leu Glu Lys Asn Met Leu Pro Asp Val Gly Lys		
	1010 1015 1020	
atg tat aaa caa gaa ttt gat gat gtc caa ggc aga tgg aat aaa		3114
Met Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Arg Trp Asn Lys		
	1025 1030 1035	
gta aag acc aag gtt tcc aga gac tta cac ttg ctc gag gaa atc		3159
Val Lys Thr Lys Val Ser Arg Asp Leu His Leu Leu Glu Glu Ile		
	1040 1045 1050	
gcc cac aga gat ttt ggg cca tct tct caa cac ttt ctg tcc act		3204
Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr		
	1055 1060 1065	

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tca gtc cag ctg ccg tgg cag	aga tcc att tca cat	aat aaa gtg	3249
Ser Val Gln Leu Pro Trp Gln	Arg Ser Ile Ser His	Asn Lys Val	
1070	1075	1080	
ccc tat tac atc aac cat caa	aca cag aca acc tgt	tgg gat cat	3294
Pro Tyr Tyr Ile Asn His Gln	Thr Gln Thr Thr Cys	Trp Asp His	
1085	1090	1095	
cct aaa atg act gag ctc ttc	caa tcc ctt gct gat	ctg aat aat	3339
Pro Lys Met Thr Glu Leu Phe	Gln Ser Leu Ala Asp	Leu Asn Asn	
1100	1105	1110	
gta cgt ttc tct gcc tac cgc	aca gca atc aaa att	cga agg ctg	3384
Val Arg Phe Ser Ala Tyr Arg	Thr Ala Ile Lys Ile	Arg Arg Leu	
1115	1120	1125	
caa aaa gca tta tgt ctg gat	ctc tta gag ctg aat	acg acg aat	3429
Gln Lys Ala Leu Cys Leu Asp	Leu Leu Glu Leu Asn	Thr Thr Asn	
1130	1135	1140	
gaa gtt ttc aag cag cac aaa	ctg aac caa aat gat	cag ctc ctg	3474
Glu Val Phe Lys Gln His Lys	Leu Asn Gln Asn Asp	Gln Leu Leu	
1145	1150	1155	
agt gtc cca gac gtc atc aac	tgt ctg acc acc act	tac gat ggg	3519
Ser Val Pro Asp Val Ile Asn	Cys Leu Thr Thr Thr	Tyr Asp Gly	
1160	1165	1170	
ctt gag cag ctg cac aag gac	ttg gtc aat gtt cca	ctc tgc gtc	3564
Leu Glu Gln Leu His Lys Asp	Leu Val Asn Val Pro	Leu Cys Val	
1175	1180	1185	
gat atg tgt ctc aac tgg ctg	ctc aac gta tac gac	acg ggc cgg	3609
Asp Met Cys Leu Asn Trp Leu	Leu Asn Val Tyr Asp	Thr Gly Arg	
1190	1195	1200	
act gga aaa att cgg gta cag	agt ctg aag att gga	ttg atg tct	3654
Thr Gly Lys Ile Arg Val Gln	Ser Leu Lys Ile Gly	Leu Met Ser	
1205	1210	1215	
ctc tcc aaa ggc ctc tta gaa	gag aaa tac aga tgt	ctc ttt aag	3699
Leu Ser Lys Gly Leu Leu Glu	Glu Lys Tyr Arg Cys	Leu Phe Lys	
1220	1225	1230	
gag gtg gca ggg cca act gag	atg tgt gac cag cgg	cag ctt ggc	3744
Glu Val Ala Gly Pro Thr Glu	Met Cys Asp Gln Arg	Gln Leu Gly	
1235	1240	1245	
ctg cta ctt cac gat gcc atc	cag atc cct agg cag	ctg ggg gaa	3789
Leu Leu Leu His Asp Ala Ile	Gln Ile Pro Arg Gln	Leu Gly Glu	
1250	1255	1260	
gta gca gcc ttt ggg ggc agt	aac att gag ccc agt	gtc cgc agc	3834
Val Ala Ala Phe Gly Gly Ser	Asn Ile Glu Pro Ser	Val Arg Ser	
1265	1270	1275	
tgc ttc cag cag aat aac aac	aag cca gaa atc agt	gtg aag gag	3879
Cys Phe Gln Gln Asn Asn Asn	Lys Pro Glu Ile Ser	Val Lys Glu	
1280	1285	1290	
ttt ata gac tgg atg cat ttg	gaa ccc cag tcc atg	gtg tgg ttg	3924
Phe Ile Asp Trp Met His Leu	Glu Pro Gln Ser Met	Val Trp Leu	
1295	1300	1305	
ccg gtt ctg cat cgg gtc gca	gct gct gag act gca	aaa cat cag	3969
Pro Val Leu His Arg Val Ala	Ala Ala Glu Thr Ala	Lys His Gln	
1310	1315	1320	
gcc aaa tgc aac atc tgc aaa	gaa tgc ccg att gtt	ggg ttc aga	4014
Ala Lys Cys Asn Ile Cys Lys	Glu Cys Pro Ile Val	Gly Phe Arg	
1325	1330	1335	
tac agg agc cta aag cat ttt	aat tat gat gtc tgc	cag agt tgc	4059
Tyr Arg Ser Leu Lys His Phe	Asn Tyr Asp Val Cys	Gln Ser Cys	
1340	1345	1350	
ttc ttt tct gga aga aca gca	aag ggc cac aag tta	cat tac ccg	4104
Phe Phe Ser Gly Arg Thr Ala	Lys Gly His Lys Leu	His Tyr Pro	
1355	1360	1365	

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atg gta gaa tac tgc ata ccg aca aca tct ggg gaa gat gtg aga	4149
Met Val Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val Arg	
1370 1375 1380	
gat ttc act aag gtg ctg aag aac aag ttc agg tcc aag aaa tat	4194
Asp Phe Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys Tyr	
1385 1390 1395	
ttt gcc aaa cat cct cgg ctt ggc tac ctg cct gtc cag acc gtg	4239
Phe Ala Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr Val	
1400 1405 1410	
ctg gaa ggg gac aac tta gaa act cct atc acg ctc atc agt atg	4284
Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser Met	
1415 1420 1425	
tgg cca gag cac tat gac ccc tcc cag tcc cct cag ctg ttt cat	4329
Trp Pro Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe His	
1430 1435 1440	
gat gac acc cac tca aga ata gag caa tac gct aca cga ctg gcc	4374
Asp Asp Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu Ala	
1445 1450 1455	
cag atg gaa agg aca aac ggg tcc ttc cta act gat agc agc tct	4419
Gln Met Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser Ser	
1460 1465 1470	
aca aca gga agc gtg gag gat gag cat gcc ctc atc cag cag tac	4464
Thr Thr Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln Tyr	
1475 1480 1485	
tgc cag acc ctg ggc ggg gag tca cct gtg agt cag ccg cag agt	4509
Cys Gln Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln Ser	
1490 1495 1500	
cca gct cag atc ctg aag tcc gtg gag agg gaa gag cgt ggg gaa	4554
Pro Ala Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly Glu	
1505 1510 1515	
ctg gag cgg atc att gct gac ttg gag gaa gag caa aga aat ctg	4599
Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn Leu	
1520 1525 1530	
cag gtg gag tat gag cag ctg aag gag cag cac cta aga agg ggt	4644
Gln Val Glu Tyr Glu Gln Leu Lys Glu Gln His Leu Arg Arg Gly	
1535 1540 1545	
ctc cct gtg ggc tcc cct cca gac tcc atc gta tct cct cac cac	4689
Leu Pro Val Gly Ser Pro Pro Asp Ser Ile Val Ser Pro His His	
1550 1555 1560	
aca tct gag gac tca gaa ctt ata gca gaa gct aaa ctc ctg cgg	4734
Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg	
1565 1570 1575	
cag cac aaa ggg cgg ctg gag gcg agg atg caa att ttg gaa gat	4779
Gln His Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp	
1580 1585 1590	
cac aat aaa cag ctg gag tct cag ctg cac cgc ctc aga cag ctc	4824
His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu	
1595 1600 1605	
ctg gag cag cct gac tct gac tcc cgc atc aat ggt gtc tcc ccc	4869
Leu Glu Gln Pro Asp Ser Asp Ser Arg Ile Asn Gly Val Ser Pro	
1610 1615 1620	
tgg gct tcc cca cag cat tct gca ttg agc tac tca ctt gac act	4914
Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Thr	
1625 1630 1635	
gac cca ggc cca cag ttc cac cag gca gca tct gag gac ctg ctg	4959
Asp Pro Gly Pro Gln Phe His Gln Ala Ala Ser Glu Asp Leu Leu	
1640 1645 1650	
gcc cca cct cac gac act agc acg gac ctc acg gac gtg atg gag	5004
Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Asp Val Met Glu	

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1655	1660	1665	
cag atc aac agc acg ttt ccc tct tgc agc tca aat gtc ccc agc			5049
Gln Ile Asn Ser Thr Phe Pro Ser Cys Ser Ser Asn Val Pro Ser			
1670	1675	1680	
agg cca cag gca atg tga			5067
Arg Pro Gln Ala Met			
1685			
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<212> TYPE: PRT			
<213> ORGANISM: Mus musculus			
<400> SEQUENCE: 21			
Met Asp Tyr Lys Asp Asp Asp Asp Lys Gly Tyr Gly Arg Lys Lys Arg			
1	5	10	15
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp			
	20	25	30
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala			
	35	40	45
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg			
	50	55	60
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile			
65	70	75	80
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe			
	85	90	95
Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu			
	100	105	110
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala			
	115	120	125
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val			
	130	135	140
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys			
145	150	155	160
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys			
	165	170	175
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu			
	180	185	190
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln			
	195	200	205
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe			
	210	215	220
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu			
225	230	235	240
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys			
	245	250	255
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val			
	260	265	270
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser			
	275	280	285
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu			
	290	295	300
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu			
305	310	315	320
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg			

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325						330						335					
Ala	Glu	Thr	Pro	Ser	Thr	Val	Thr	Glu	Val	Asp	Met	Asp	Leu	Asp	Ser		
340						345						350					
Tyr	Gln	Ile	Ala	Leu	Glu	Glu	Val	Leu	Thr	Trp	Leu	Leu	Ser	Ala	Glu		
355						360						365					
Asp	Thr	Phe	Gln	Glu	Gln	His	Asp	Ile	Ser	Asp	Asp	Val	Glu	Glu	Val		
370						375						380					
Lys	Glu	Gln	Phe	Ala	Thr	His	Glu	Thr	Phe	Met	Met	Glu	Leu	Thr	Ala		
385						390						395					
His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Met		
405						410						415					
Thr	Gln	Gly	Thr	Leu	Ser	Arg	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln		
420						425						430					
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met		
435						440						445					
Glu	Arg	Gln	Ser	Arg	Leu	His	Asp	Ala	Leu	Met	Glu	Leu	Gln	Lys	Lys		
450						455						460					
Gln	Leu	Gln	Gln	Leu	Ser	Ser	Trp	Leu	Ala	Leu	Thr	Glu	Glu	Arg	Ile		
465						470						475					
Gln	Lys	Met	Glu	Ser	Leu	Pro	Leu	Gly	Asp	Asp	Leu	Pro	Ser	Leu	Gln		
485						490						495					
Lys	Leu	Leu	Gln	Glu	His	Lys	Ser	Leu	Gln	Asn	Asp	Leu	Glu	Ala	Glu		
500						505						510					
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu		
515						520						525					
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Leu	Leu	Glu	Asp	Gln	Leu	Gln	Lys		
530						535						540					
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp		
545						550						555					
Asn	Arg	Leu	Gln	Glu	Ile	Ser	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu		
565						570						575					
Gln	Cys	Leu	Leu	Glu	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asp		
580						585						590					
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser		
595						600						605					
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln		
610						615						620					
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu		
625						630						635					
Ser	Asn	Pro	Lys	Ala	Ser	Lys	Lys	Met	Asn	Ser	Asp	Ser	Glu	Glu	Leu		
645						650						655					
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn		
660						665						670					
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln		
675						680						685					
Lys	Asp	Leu	Leu	Glu	Thr	Val	His	Val	Arg	Glu	Gln	Gly	Met	Val	Lys		
690						695						700					
Lys	Pro	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln		
705						710						715					
Ile	His	Val	Asp	Val	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Thr		
725						730						735					
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Ser	Lys	Thr	Ala	Ile	Gln	Asn	Thr		
740						745						750					

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Glu Met Lys Glu Tyr Lys Lys Ser Gln Glu Thr Ser Gly Met Lys Lys
 755 760 765
 Lys Leu Lys Gly Leu Glu Lys Glu Gln Lys Glu Asn Leu Pro Arg Leu
 770 775 780
 Asp Glu Leu Asn Gln Thr Gly Gln Thr Leu Arg Glu Gln Met Gly Lys
 785 790 795 800
 Glu Gly Leu Pro Leu Lys Glu Val Asn Asp Val Leu Glu Arg Val Ser
 805 810 815
 Leu Glu Trp Lys Met Ile Ser Gln Gln Leu Glu Asp Leu Gly Arg Lys
 820 825 830
 Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Ala
 835 840 845
 Ile Glu Glu Thr Ile Lys Glu Lys Glu Glu Trp Leu Arg Gly Thr Pro
 850 855 860
 Ile Ser Glu Ser Pro Arg Gln Pro Leu Pro Gly Leu Lys Asp Ser Cys
 865 870 875 880
 Gln Arg Glu Leu Thr Asp Leu Leu Gly Leu His Pro Arg Ile Glu Thr
 885 890 895
 Leu Cys Ala Ser Cys Ser Ala Leu Lys Ser Gln Pro Cys Val Pro Gly
 900 905 910
 Phe Val Gln Gln Gly Phe Asp Asp Leu Arg His His Tyr Gln Ala Val
 915 920 925
 Ala Lys Ala Leu Glu Glu Tyr Gln Gln Gln Leu Glu Asn Glu Leu Lys
 930 935 940
 Ser Gln Pro Gly Pro Glu Tyr Leu Asp Thr Leu Asn Thr Leu Lys Lys
 945 950 955 960
 Met Leu Ser Glu Ser Glu Lys Ala Ala Gln Ala Ser Leu Asn Ala Leu
 965 970 975
 Asn Asp Pro Ile Ala Val Glu Gln Ala Leu Gln Glu Lys Lys Ala Leu
 980 985 990
 Asp Glu Thr Leu Glu Asn Gln Lys His Thr Leu His Lys Leu Ser Glu
 995 1000 1005
 Glu Thr Lys Thr Leu Glu Lys Asn Met Leu Pro Asp Val Gly Lys
 1010 1015 1020
 Met Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Arg Trp Asn Lys
 1025 1030 1035
 Val Lys Thr Lys Val Ser Arg Asp Leu His Leu Leu Glu Glu Ile
 1040 1045 1050
 Ala His Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr
 1055 1060 1065
 Ser Val Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn Lys Val
 1070 1075 1080
 Pro Tyr Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp His
 1085 1090 1095
 Pro Lys Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn
 1100 1105 1110
 Val Arg Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu
 1115 1120 1125
 Gln Lys Ala Leu Cys Leu Asp Leu Leu Glu Leu Asn Thr Thr Asn
 1130 1135 1140
 Glu Val Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu
 1145 1150 1155

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Ser	Val	Pro	Asp	Val	Ile	Asn	Cys	Leu	Thr	Thr	Thr	Tyr	Asp	Gly
1160						1165						1170		
Leu	Glu	Gln	Leu	His	Lys	Asp	Leu	Val	Asn	Val	Pro	Leu	Cys	Val
1175						1180					1185			
Asp	Met	Cys	Leu	Asn	Trp	Leu	Leu	Asn	Val	Tyr	Asp	Thr	Gly	Arg
1190						1195					1200			
Thr	Gly	Lys	Ile	Arg	Val	Gln	Ser	Leu	Lys	Ile	Gly	Leu	Met	Ser
1205						1210					1215			
Leu	Ser	Lys	Gly	Leu	Leu	Glu	Glu	Lys	Tyr	Arg	Cys	Leu	Phe	Lys
1220						1225					1230			
Glu	Val	Ala	Gly	Pro	Thr	Glu	Met	Cys	Asp	Gln	Arg	Gln	Leu	Gly
1235						1240					1245			
Leu	Leu	Leu	His	Asp	Ala	Ile	Gln	Ile	Pro	Arg	Gln	Leu	Gly	Glu
1250						1255					1260			
Val	Ala	Ala	Phe	Gly	Gly	Ser	Asn	Ile	Glu	Pro	Ser	Val	Arg	Ser
1265						1270					1275			
Cys	Phe	Gln	Gln	Asn	Asn	Asn	Lys	Pro	Glu	Ile	Ser	Val	Lys	Glu
1280						1285					1290			
Phe	Ile	Asp	Trp	Met	His	Leu	Glu	Pro	Gln	Ser	Met	Val	Trp	Leu
1295						1300					1305			
Pro	Val	Leu	His	Arg	Val	Ala	Ala	Ala	Glu	Thr	Ala	Lys	His	Gln
1310						1315					1320			
Ala	Lys	Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly	Phe	Arg
1325						1330					1335			
Tyr	Arg	Ser	Leu	Lys	His	Phe	Asn	Tyr	Asp	Val	Cys	Gln	Ser	Cys
1340						1345					1350			
Phe	Phe	Ser	Gly	Arg	Thr	Ala	Lys	Gly	His	Lys	Leu	His	Tyr	Pro
1355						1360					1365			
Met	Val	Glu	Tyr	Cys	Ile	Pro	Thr	Thr	Ser	Gly	Glu	Asp	Val	Arg
1370						1375					1380			
Asp	Phe	Thr	Lys	Val	Leu	Lys	Asn	Lys	Phe	Arg	Ser	Lys	Lys	Tyr
1385						1390					1395			
Phe	Ala	Lys	His	Pro	Arg	Leu	Gly	Tyr	Leu	Pro	Val	Gln	Thr	Val
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Leu	Glu	Gly	Asp	Asn	Leu	Glu	Thr	Pro	Ile	Thr	Leu	Ile	Ser	Met
1415						1420					1425			
Trp	Pro	Glu	His	Tyr	Asp	Pro	Ser	Gln	Ser	Pro	Gln	Leu	Phe	His
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1445						1450					1455			
Gln	Met	Glu	Arg	Thr	Asn	Gly	Ser	Phe	Leu	Thr	Asp	Ser	Ser	Ser
1460						1465					1470			
Thr	Thr	Gly	Ser	Val	Glu	Asp	Glu	His	Ala	Leu	Ile	Gln	Gln	Tyr
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Cys	Gln	Thr	Leu	Gly	Gly	Glu	Ser	Pro	Val	Ser	Gln	Pro	Gln	Ser
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Leu	Glu	Arg	Ile	Ile	Ala	Asp	Leu	Glu	Glu	Glu	Gln	Arg	Asn	Leu
1520						1525					1530			
Gln	Val	Glu	Tyr	Glu	Gln	Leu	Lys	Glu	Gln	His	Leu	Arg	Arg	Gly
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Leu	Pro	Val	Gly	Ser	Pro	Pro	Asp	Ser	Ile	Val	Ser	Pro	His	His

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His Asn Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu			
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Trp Ala Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Thr			
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Ala Pro Pro His Asp Thr Ser Thr Asp Leu Thr Asp Val Met Glu			
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Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp			
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gtc cca gac tat gct ggc tcc atg gcc aag tat ggg gac ctt gaa gcc			144
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala			
35 40 45			
agg cct gat gat ggg cag aac gaa ttc agt gac atc att aag tcc aga			192
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg			
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tct gat gaa cac aat gat gta cag aag aaa acc ttt acc aaa tgg ata			240
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile			
65 70 75 80			
aac gct cga ttt tcc aag agt ggg aaa cca ccc atc agt gat atg ttc			288
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe			
85 90 95			
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Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu			
100 105 110			
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Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala			
115 120 125			
tta aac aat gtc aac cga gtg cta cag gtt tta cat cag aac aat gtg			432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val			
130 135 140			

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Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys	
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ctg act tta ggg tta ctc tgg agc atc att ctg cac tgg cag gtg aag	528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys	
165 170 175	
gat gtc atg aaa gat atc atg tca gac ctg cag cag aca aac agc gag	576
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu	
180 185 190	
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Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln	
195 200 205	
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Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe	
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Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu	
225 230 235 240	
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Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys	
245 250 255	
gcc cac act tct ttg gga att gaa aag ctc cta agt cct gaa act gtt	816
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val	
260 265 270	
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Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser	
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Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu	
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Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu	
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Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg	
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Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser	
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Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu	
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Asp Thr Phe Gln Glu Gln His Asp Ile Ser Asp Asp Val Glu Glu Val	
370 375 380	
aaa gag cag ttt gct acc cat gaa act ttt atg atg gag ctg aca gca	1200
Lys Glu Gln Phe Ala Thr His Glu Thr Phe Met Met Glu Leu Thr Ala	
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His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Met	
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aca caa ggg act ctg tcc aga gag gag gag ttt gag atc cag gaa cag	1296
Thr Gln Gly Thr Leu Ser Arg Glu Glu Glu Phe Glu Ile Gln Glu Gln	
420 425 430	
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Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met	
435 440 445	
gag agg cag tcc cgg ctg cac gac gct ctg atg gag ctg cag aag aaa	1392
Glu Arg Gln Ser Arg Leu His Asp Ala Leu Met Glu Leu Gln Lys Lys	
450 455 460	

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aag ctg ctt caa gaa cat aaa agt ttg caa aat gac ctt gaa gct gaa Lys Leu Leu Gln Glu His Lys Ser Leu Gln Asn Asp Leu Glu Ala Glu 500 505 510	1536
cag gtg aag gta aat tcc tta act cac atg gtg gtg att gtg gat gaa Gln Val Lys Val Asn Ser Leu Thr His Met Val Val Ile Val Asp Glu 515 520 525	1584
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cag gtg act cag gcg gta gcg aag ctc ggc atg tcc cag att cca cag Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln 675 680 685	2064
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att cac gtg gac gtg gag gcc aag aaa aag ttt gat gct ata agt aca Ile His Val Asp Val Glu Ala Lys Lys Lys Phe Asp Ala Ile Ser Thr 725 730 735	2208
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gag atg aaa gaa tat aag aag tcg cag gag acc tca gga atg aaa aag Glu Met Lys Glu Tyr Lys Lys Ser Gln Glu Thr Ser Gly Met Lys Lys 755 760 765	2304
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ttg gag tgg aag atg ata tct cag cag cta gaa gat ctg gga agg aag Leu Glu Trp Lys Met Ile Ser Gln Gln Leu Glu Asp Leu Gly Arg Lys 820 825 830			2496
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att gag gag acc atc aag gag aag gaa gag tgg ctg agg ggc aca ccc Ile Glu Glu Thr Ile Lys Glu Lys Glu Glu Trp Leu Arg Gly Thr Pro 850 855 860			2592
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cag agg gaa ctg aca gat ctc ctt ggc ctt cac ccc aga att gag acg Gln Arg Glu Leu Thr Asp Leu Leu Gly Leu His Pro Arg Ile Glu Thr 885 890 895			2688
ctg tgt gca agc tgt tca gcc ctg aag tct cag ccc tgt gtc cca ggt Leu Cys Ala Ser Cys Ser Ala Leu Lys Ser Gln Pro Cys Val Pro Gly 900 905 910			2736
ttt gtc cag cag ggt ttt gac gac ctt cga cat cat tac cag gct gtt Phe Val Gln Gln Gly Phe Asp Asp Leu Arg His His Tyr Gln Ala Val 915 920 925			2784
gcg aag gct tta gag gaa tac caa caa caa cta gaa aat gag ctg aag Ala Lys Ala Leu Glu Glu Tyr Gln Gln Gln Leu Glu Asn Glu Leu Lys 930 935 940			2832
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acc ccc aga ctc cga gat ttt gag gct gat tca gaa gtc att gag Thr Pro Arg Leu Arg Asp Phe Glu Ala Asp Ser Glu Val Ile Glu 1055 1060 1065			3204
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Thr	Phe	Ala	Asn	Glu	Ile	Glu	Thr	Ile	Glu	Ser	Ser	Leu	Lys	Asn			
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Met	Arg	Glu	Val	Glu	Thr	Ser	Leu	Gln	Arg	Cys	Pro	Val	Thr	Gly			
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Leu	Asp	Ser	Gln	Glu	Lys	Ala	Leu	Asn	Leu	Lys	Lys	Asp	Leu	Ala			
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Arg	Asp	Phe	Glu	Tyr	Lys	Ser	Pro	Glu	Glu	Leu	Glu	Ser	Ala	Val			
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Glu	Glu	Met	Lys	Arg	Ala	Lys	Glu	Asp	Val	Leu	Gln	Lys	Glu	Val			
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Arg	Val	Lys	Ile	Leu	Lys	Asp	Ser	Ile	Lys	Leu	Val	Ala	Ala	Lys			
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Val	Pro	Ser	Gly	Gly	Gln	Glu	Leu	Thr	Ser	Glu	Phe	Asn	Glu	Val			
1235						1240					1245						
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Leu	Glu	Ser	Tyr	Gln	Leu	Leu	Cys	Asn	Arg	Ile	Arg	Gly	Lys	Cys			
1250						1255					1260						
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His	Thr	Leu	Glu	Glu	Val	Trp	Ser	Cys	Trp	Val	Glu	Leu	Leu	His			
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1310						1315					1320						
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Asn	Arg	Thr	Gln	Ile	Arg	Glu	Leu	Gly	Gln	Thr	Leu	Ile	Asp	Gly			
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Ser His	Asn Lys	Val Pro	Tyr	Tyr Ile	Asn His	Gln	Thr Gln	Thr	
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Thr Cys	Trp Asp	His Pro	Lys	Met Thr	Glu Leu	Phe	Gln Ser	Leu	
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Ala Asp	Leu Asn	Asn Val	Arg	Phe Ser	Ala Tyr	Arg	Thr Ala	Ile	
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Lys Ile	Arg Arg	Leu Gln	Lys	Ala Leu	Cys Leu	Asp	Leu Leu	Glu	
1445			1450			1455			
ctg aat	acg acg	aat gaa	gtt	ttc aag	cag cac	aaa	ctg aac	caa	4419
Leu Asn	Thr Thr	Asn Glu	Val	Phe Lys	Gln His	Lys	Leu Asn	Gln	
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Asn Asp	Gln Leu	Leu Ser	Val	Pro Asp	Val Ile	Asn	Cys Leu	Thr	
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Val Pro	Leu Cys	Val Asp	Met	Cys Leu	Asn Trp	Leu	Leu Asn	Val	
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Tyr Asp	Thr Gly	Arg Thr	Gly	Lys Ile	Arg Val	Gln	Ser Leu	Lys	
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Ile Gly	Leu Met	Ser Leu	Ser	Lys Gly	Leu Leu	Glu	Glu Lys	Tyr	
1535			1540			1545			
aga tgt	ctc ttt	aag gag	gtg	gca ggg	cca act	gag	atg tgt	gac	4689
Arg Cys	Leu Phe	Lys Glu	Val	Ala Gly	Pro Thr	Glu	Met Cys	Asp	
1550			1555			1560			
cag cgg	cag ctt	ggc ctg	cta	ctt cac	gat gcc	atc	cag atc	cct	4734
Gln Arg	Gln Leu	Gly Leu	Leu	Leu His	Asp Ala	Ile	Gln Ile	Pro	
1565			1570			1575			
agg cag	ctg ggg	gaa gta	gca	gcc ttt	ggg ggc	agt	aac att	gag	4779
Arg Gln	Leu Gly	Glu Val	Ala	Ala Phe	Gly Gly	Ser	Asn Ile	Glu	
1580			1585			1590			
ccc agt	gtc cgc	agc tgc	ttc	cag cag	aat aac	aac	aag cca	gaa	4824
Pro Ser	Val Arg	Ser Cys	Phe	Gln Gln	Asn Asn	Asn	Lys Pro	Glu	
1595			1600			1605			
atc agt	gtg aag	gag ttt	ata	gac tgg	atg cat	ttg	gaa ccc	cag	4869
Ile Ser	Val Lys	Glu Phe	Ile	Asp Trp	Met His	Leu	Glu Pro	Gln	
1610			1615			1620			
tcc atg	gtg tgg	ttg ccg	gtt	ctg cat	cgg gtc	gca	gct gct	gag	4914
Ser Met	Val Trp	Leu Pro	Val	Leu His	Arg Val	Ala	Ala Ala	Glu	
1625			1630			1635			
act gca	aaa cat	cag gcc	aaa	tgc aac	atc tgc	aaa	gaa tgc	ccg	4959
Thr Ala	Lys His	Gln Ala	Lys	Cys Asn	Ile Cys	Lys	Glu Cys	Pro	
1640			1645			1650			
att gtt	ggg ttc	aga tac	agg	agc cta	aag cat	ttt	aat tat	gat	5004
Ile Val	Gly Phe	Arg Tyr	Arg	Ser Leu	Lys His	Phe	Asn Tyr	Asp	
1655			1660			1665			
gtc tgc	cag agt	tgc ttc	ttt	tct gga	aga aca	gca	aag ggc	cac	5049
Val Cys	Gln Ser	Cys Phe	Phe	Ser Gly	Arg Thr	Ala	Lys Gly	His	
1670			1675			1680			

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aag tta cat tac ccg atg gta gaa tac tgc ata ccg aca aca tct	5094
Lys Leu His Tyr Pro Met Val Glu Tyr Cys Ile Pro Thr Thr Ser	
1685 1690 1695	
ggg gaa gat gtg aga gat ttc act aag gtg ctg aag aac aag ttc	5139
Gly Glu Asp Val Arg Asp Phe Thr Lys Val Leu Lys Asn Lys Phe	
1700 1705 1710	
agg tcc aag aaa tat ttt gcc aaa cat cct cgg ctt ggc tac ctg	5184
Arg Ser Lys Lys Tyr Phe Ala Lys His Pro Arg Leu Gly Tyr Leu	
1715 1720 1725	
cct gtc cag acc gtg ctg gaa ggg gac aac tta gaa act cct atc	5229
Pro Val Gln Thr Val Leu Glu Gly Asp Asn Leu Glu Thr Pro Ile	
1730 1735 1740	
acg ctc atc agt atg tgg cca gag cac tat gac ccc tcc cag tcc	5274
Thr Leu Ile Ser Met Trp Pro Glu His Tyr Asp Pro Ser Gln Ser	
1745 1750 1755	
cct cag ctg ttt cat gat gac acc cac tca aga ata gag caa tac	5319
Pro Gln Leu Phe His Asp Asp Thr His Ser Arg Ile Glu Gln Tyr	
1760 1765 1770	
gct aca cga ctg gcc cag atg gaa agg aca aac ggg tcc ttc cta	5364
Ala Thr Arg Leu Ala Gln Met Glu Arg Thr Asn Gly Ser Phe Leu	
1775 1780 1785	
act gat agc agc tct aca aca gga agc gtg gag gat gag cat gcc	5409
Thr Asp Ser Ser Ser Thr Thr Gly Ser Val Glu Asp Glu His Ala	
1790 1795 1800	
ctc atc cag cag tac tgc cag acc ctg ggc ggg gag tca cct gtg	5454
Leu Ile Gln Gln Tyr Cys Gln Thr Leu Gly Gly Glu Ser Pro Val	
1805 1810 1815	
agt cag ccg cag agt cca gct cag atc ctg aag tcc gtg gag agg	5499
Ser Gln Pro Gln Ser Pro Ala Gln Ile Leu Lys Ser Val Glu Arg	
1820 1825 1830	
gaa gag cgt ggg gaa ctg gag cgg atc att gct gac ttg gag gaa	5544
Glu Glu Arg Gly Glu Leu Glu Arg Ile Ile Ala Asp Leu Glu Glu	
1835 1840 1845	
gag caa aga aat ctg cag gtg gag tat gag cag ctg aag gag cag	5589
Glu Gln Arg Asn Leu Gln Val Glu Tyr Glu Gln Leu Lys Glu Gln	
1850 1855 1860	
cac cta aga agg ggt ctc cct gtg ggc tcc cct cca gac tcc atc	5634
His Leu Arg Arg Gly Leu Pro Val Gly Ser Pro Pro Asp Ser Ile	
1865 1870 1875	
gta tct cct cac cac aca tct gag gac tca gaa ctt ata gca gaa	5679
Val Ser Pro His His Thr Ser Glu Asp Ser Glu Leu Ile Ala Glu	
1880 1885 1890	
gct aaa ctc ctg cgg cag cac aaa ggg cgg ctg gag gcg agg atg	5724
Ala Lys Leu Leu Arg Gln His Lys Gly Arg Leu Glu Ala Arg Met	
1895 1900 1905	
caa att ttg gaa gat cac aat aaa cag ctg gag tct cag ctg cac	5769
Gln Ile Leu Glu Asp His Asn Lys Gln Leu Glu Ser Gln Leu His	
1910 1915 1920	
cgc ctc aga cag ctc ctg gag cag cct gac tct gac tcc cgc atc	5814
Arg Leu Arg Gln Leu Leu Glu Gln Pro Asp Ser Asp Ser Arg Ile	
1925 1930 1935	
aat ggt gtc tcc ccc tgg gct tcc cca cag cat tct gca ttg agc	5859
Asn Gly Val Ser Pro Trp Ala Ser Pro Gln His Ser Ala Leu Ser	
1940 1945 1950	
tac tca ctt gac act gac cca ggc cca cag ttc cac cag gca gca	5904
Tyr Ser Leu Asp Thr Asp Pro Gly Pro Gln Phe His Gln Ala Ala	
1955 1960 1965	
tct gag gac ctg ctg gcc cca cct cac gac act agc acg gac ctc	5949
Ser Glu Asp Leu Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu	

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1970	1975	1980	
acg gac gtg atg gag cag atc aac agc acg ttt ccc tct tgc agc			5994
Thr Asp Val Met Glu Gln Ile Asn Ser Thr Phe Pro Ser Cys Ser			
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2000	2005		
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	20	25	30
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala			
	35	40	45
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg			
	50	55	60
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile			
65	70	75	80
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe			
	85	90	95
Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu			
	100	105	110
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala			
	115	120	125
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val			
	130	135	140
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys			
145	150	155	160
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys			
	165	170	175
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu			
	180	185	190
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln			
	195	200	205
Val Asn Val Leu Asn Phe Thr Thr Ser Trp Thr Asp Gly Leu Ala Phe			
	210	215	220
Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu			
225	230	235	240
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys			
	245	250	255
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val			
	260	265	270
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser			
	275	280	285
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu			
	290	295	300
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu			
305	310	315	320
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg			

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325						330						335					
Ala	Glu	Thr	Pro	Ser	Thr	Val	Thr	Glu	Val	Asp	Met	Asp	Leu	Asp	Ser		
340						345						350					
Tyr	Gln	Ile	Ala	Leu	Glu	Glu	Val	Leu	Thr	Trp	Leu	Leu	Ser	Ala	Glu		
355						360						365					
Asp	Thr	Phe	Gln	Glu	Gln	His	Asp	Ile	Ser	Asp	Asp	Val	Glu	Glu	Val		
370						375						380					
Lys	Glu	Gln	Phe	Ala	Thr	His	Glu	Thr	Phe	Met	Met	Glu	Leu	Thr	Ala		
385						390						395					
His	Gln	Ser	Ser	Val	Gly	Ser	Val	Leu	Gln	Ala	Gly	Asn	Gln	Leu	Met		
405						410						415					
Thr	Gln	Gly	Thr	Leu	Ser	Arg	Glu	Glu	Glu	Phe	Glu	Ile	Gln	Glu	Gln		
420						425						430					
Met	Thr	Leu	Leu	Asn	Ala	Arg	Trp	Glu	Ala	Leu	Arg	Val	Glu	Ser	Met		
435						440						445					
Glu	Arg	Gln	Ser	Arg	Leu	His	Asp	Ala	Leu	Met	Glu	Leu	Gln	Lys	Lys		
450						455						460					
Gln	Leu	Gln	Gln	Leu	Ser	Ser	Trp	Leu	Ala	Leu	Thr	Glu	Glu	Arg	Ile		
465						470						475					
Gln	Lys	Met	Glu	Ser	Leu	Pro	Leu	Gly	Asp	Asp	Leu	Pro	Ser	Leu	Gln		
485						490						495					
Lys	Leu	Leu	Gln	Glu	His	Lys	Ser	Leu	Gln	Asn	Asp	Leu	Glu	Ala	Glu		
500						505						510					
Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu		
515						520						525					
Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Leu	Leu	Glu	Asp	Gln	Leu	Gln	Lys		
530						535						540					
Leu	Gly	Glu	Arg	Trp	Thr	Ala	Val	Cys	Arg	Trp	Thr	Glu	Glu	Arg	Trp		
545						550						555					
Asn	Arg	Leu	Gln	Glu	Ile	Ser	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu		
565						570						575					
Gln	Cys	Leu	Leu	Glu	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asp		
580						585						590					
Lys	Val	Gln	Thr	Ser	Asn	Phe	Lys	Asp	Gln	Lys	Glu	Leu	Ser	Val	Ser		
595						600						605					
Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln		
610						615						620					
Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu		
625						630						635					
Ser	Asn	Pro	Lys	Ala	Ser	Lys	Lys	Met	Asn	Ser	Asp	Ser	Glu	Glu	Leu		
645						650						655					
Thr	Gln	Arg	Trp	Asp	Ser	Leu	Val	Gln	Arg	Leu	Glu	Asp	Ser	Ser	Asn		
660						665						670					
Gln	Val	Thr	Gln	Ala	Val	Ala	Lys	Leu	Gly	Met	Ser	Gln	Ile	Pro	Gln		
675						680						685					
Lys	Asp	Leu	Leu	Glu	Thr	Val	His	Val	Arg	Glu	Gln	Gly	Met	Val	Lys		
690						695						700					
Lys	Pro	Lys	Gln	Glu	Leu	Pro	Pro	Pro	Pro	Pro	Pro	Lys	Lys	Arg	Gln		
705						710						715					
Ile	His	Val	Asp	Val	Glu	Ala	Lys	Lys	Lys	Phe	Asp	Ala	Ile	Ser	Thr		
725						730						735					
Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Ser	Lys	Thr	Ala	Ile	Gln	Asn	Thr		
740						745						750					

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Glu Met Lys Glu Tyr Lys Lys Ser Gln Glu Thr Ser Gly Met Lys Lys
 755 760 765
 Lys Leu Lys Gly Leu Glu Lys Glu Gln Lys Glu Asn Leu Pro Arg Leu
 770 775 780
 Asp Glu Leu Asn Gln Thr Gly Gln Thr Leu Arg Glu Gln Met Gly Lys
 785 790 795 800
 Glu Gly Leu Pro Leu Lys Glu Val Asn Asp Val Leu Glu Arg Val Ser
 805 810 815
 Leu Glu Trp Lys Met Ile Ser Gln Gln Leu Glu Asp Leu Gly Arg Lys
 820 825 830
 Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Ala
 835 840 845
 Ile Glu Glu Thr Ile Lys Glu Lys Glu Glu Trp Leu Arg Gly Thr Pro
 850 855 860
 Ile Ser Glu Ser Pro Arg Gln Pro Leu Pro Gly Leu Lys Asp Ser Cys
 865 870 875 880
 Gln Arg Glu Leu Thr Asp Leu Leu Gly Leu His Pro Arg Ile Glu Thr
 885 890 895
 Leu Cys Ala Ser Cys Ser Ala Leu Lys Ser Gln Pro Cys Val Pro Gly
 900 905 910
 Phe Val Gln Gln Gly Phe Asp Asp Leu Arg His His Tyr Gln Ala Val
 915 920 925
 Ala Lys Ala Leu Glu Glu Tyr Gln Gln Gln Leu Glu Asn Glu Leu Lys
 930 935 940
 Ser Gln Pro Gly Pro Glu Tyr Leu Asp Thr Leu Asn Thr Leu Lys Lys
 945 950 955 960
 Met Leu Ser Glu Ser Glu Lys Ala Ala Gln Ala Ser Leu Asn Ala Leu
 965 970 975
 Asn Asp Pro Ile Ala Val Glu Gln Ala Leu Gln Glu Lys Lys Ala Leu
 980 985 990
 Asp Glu Thr Leu Glu Asn Gln Lys His Thr Leu His Lys Leu Ser Glu
 995 1000 1005
 Glu Thr Lys Thr Leu Glu Lys Asn Met Leu Pro Asp Val Gly Lys
 1010 1015 1020
 Met Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Arg Trp Asn Lys
 1025 1030 1035
 Val Lys Thr Lys Val Ser Arg Asp Leu His Leu Leu Glu Glu Ile
 1040 1045 1050
 Thr Pro Arg Leu Arg Asp Phe Glu Ala Asp Ser Glu Val Ile Glu
 1055 1060 1065
 Lys Trp Val Ser Gly Ile Lys Asp Phe Leu Met Lys Glu Gln Ala
 1070 1075 1080
 Ala Gln Gly Asp Ala Ala Gln Ser Gln Leu Asp Gln Cys Ala
 1085 1090 1095
 Thr Phe Ala Asn Glu Ile Glu Thr Ile Glu Ser Ser Leu Lys Asn
 1100 1105 1110
 Met Arg Glu Val Glu Thr Ser Leu Gln Arg Cys Pro Val Thr Gly
 1115 1120 1125
 Val Lys Thr Trp Val Gln Ala Arg Leu Val Asp Tyr Gln Ser Gln
 1130 1135 1140
 Leu Glu Lys Phe Ser Lys Glu Ile Ala Ile Gln Lys Ser Arg Leu
 1145 1150 1155

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Leu Asp 1160	Ser Gln Glu Lys 1165	Ala	Leu Asn Leu Lys Lys 1170	Asp Leu Ala
Glu Met 1175	Gln Glu Trp Met 1180	Ala	Gln Ala Glu Glu Asp 1185	Tyr Leu Glu
Arg Asp 1190	Phe Glu Tyr Lys 1195	Ser	Pro Glu Glu Leu Glu 1200	Ser Ala Val
Glu Glu 1205	Met Lys Arg Ala 1210	Lys	Glu Asp Val Leu Gln 1215	Lys Glu Val
Arg Val 1220	Lys Ile Leu Lys 1225	Asp	Ser Ile Lys Leu Val 1230	Ala Ala Lys
Val Pro 1235	Ser Gly Gly Gln 1240	Glu	Leu Thr Ser Glu Phe 1245	Asn Glu Val
Leu Glu 1250	Ser Tyr Gln Leu 1255	Leu	Cys Asn Arg Ile Arg 1260	Gly Lys Cys
His Thr 1265	Leu Glu Glu Val 1270	Trp	Ser Cys Trp Val Glu 1275	Leu Leu His
Tyr Leu 1280	Asp Leu Glu Thr 1285	Thr	Trp Leu Asn Thr Leu 1290	Glu Glu Arg
Val Arg 1295	Ser Thr Glu Ala 1300	Leu	Pro Glu Arg Ala Glu 1305	Ala Val His
Glu Ala 1310	Leu Glu Ser Leu 1315	Glu	Ser Val Leu Arg His 1320	Pro Ala Asp
Asn Arg 1325	Thr Gln Ile Arg 1330	Glu	Leu Gly Gln Thr Leu 1335	Ile Asp Gly
Gly Ile 1340	Leu Asp Asp Ile 1345	Ile	Ser Glu Lys Leu Glu 1350	Ala Phe Asn
Ser Arg 1355	Tyr Glu Glu Leu 1360	Ser	His Leu Ala Glu Ser 1365	Lys Gln Ile
Ser Leu 1370	Glu Lys Gln Ala 1375	His	Arg Asp Phe Gly Pro 1380	Ser Ser Gln
His Phe 1385	Leu Ser Thr Ser 1390	Val	Gln Leu Pro Trp Gln 1395	Arg Ser Ile
Ser His 1400	Asn Lys Val Pro 1405	Tyr	Tyr Ile Asn His Gln 1410	Thr Gln Thr
Thr Cys 1415	Trp Asp His Pro 1420	Lys	Met Thr Glu Leu Phe 1425	Gln Ser Leu
Ala Asp 1430	Leu Asn Asn Val 1435	Arg	Phe Ser Ala Tyr Arg 1440	Thr Ala Ile
Lys Ile 1445	Arg Arg Leu Gln 1450	Lys	Ala Leu Cys Leu Asp 1455	Leu Leu Glu
Leu Asn 1460	Thr Thr Asn Glu 1465	Val	Phe Lys Gln His Lys 1470	Leu Asn Gln
Asn Asp 1475	Gln Leu Leu Ser 1480	Val	Pro Asp Val Ile Asn 1485	Cys Leu Thr
Thr Thr 1490	Tyr Asp Gly Leu 1495	Glu	Gln Leu His Lys Asp 1500	Leu Val Asn
Val Pro 1505	Leu Cys Val Asp 1510	Met	Cys Leu Asn Trp Leu 1515	Leu Asn Val
Tyr Asp 1520	Thr Gly Arg Thr 1525	Gly	Lys Ile Arg Val Gln 1530	Ser Leu Lys
Ile Gly 1535	Leu Met Ser Leu 1540	Ser	Lys Gly Leu Leu Glu 1545	Glu Lys Tyr
Arg Cys	Leu Phe Lys Glu Val	Ala	Gly Pro Thr Glu	Met Cys Asp

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Gln Arg	Gln Leu Gly Leu	Leu Leu His Asp Ala	Ile	Gln Ile Pro	
1565		1570	1575		
Arg Gln	Leu Gly Glu Val	Ala Ala Phe Gly Gly Ser	Asn Ile Glu		
1580		1585	1590		
Pro Ser	Val Arg Ser Cys	Phe Gln Gln Asn Asn Asn	Lys Pro Glu		
1595		1600	1605		
Ile Ser	Val Lys Glu Phe	Ile Asp Trp Met His Leu	Glu Pro Gln		
1610		1615	1620		
Ser Met	Val Trp Leu Pro	Val Leu His Arg Val Ala	Ala Ala Glu		
1625		1630	1635		
Thr Ala	Lys His Gln Ala	Lys Cys Asn Ile Cys Lys	Glu Cys Pro		
1640		1645	1650		
Ile Val	Gly Phe Arg Tyr	Arg Ser Leu Lys His Phe	Asn Tyr Asp		
1655		1660	1665		
Val Cys	Gln Ser Cys Phe	Phe Ser Gly Arg Thr Ala	Lys Gly His		
1670		1675	1680		
Lys Leu	His Tyr Pro Met	Val Glu Tyr Cys Ile Pro	Thr Thr Ser		
1685		1690	1695		
Gly Glu	Asp Val Arg Asp	Phe Thr Lys Val Leu Lys	Asn Lys Phe		
1700		1705	1710		
Arg Ser	Lys Lys Tyr Phe	Ala Lys His Pro Arg Leu	Gly Tyr Leu		
1715		1720	1725		
Pro Val	Gln Thr Val Leu	Glu Gly Asp Asn Leu Glu	Thr Pro Ile		
1730		1735	1740		
Thr Leu	Ile Ser Met Trp	Pro Glu His Tyr Asp Pro	Ser Gln Ser		
1745		1750	1755		
Pro Gln	Leu Phe His Asp	Asp Thr His Ser Arg Ile	Glu Gln Tyr		
1760		1765	1770		
Ala Thr	Arg Leu Ala Gln	Met Glu Arg Thr Asn Gly	Ser Phe Leu		
1775		1780	1785		
Thr Asp	Ser Ser Ser Thr	Thr Gly Ser Val Glu Asp	Glu His Ala		
1790		1795	1800		
Leu Ile	Gln Gln Tyr Cys	Gln Thr Leu Gly Gly Glu	Ser Pro Val		
1805		1810	1815		
Ser Gln	Pro Gln Ser Pro	Ala Gln Ile Leu Lys Ser	Val Glu Arg		
1820		1825	1830		
Glu Glu	Arg Gly Glu Leu	Glu Arg Ile Ile Ala Asp	Leu Glu Glu		
1835		1840	1845		
Glu Gln	Arg Asn Leu Gln	Val Glu Tyr Glu Gln Leu	Lys Glu Gln		
1850		1855	1860		
His Leu	Arg Arg Gly Leu	Pro Val Gly Ser Pro Pro	Asp Ser Ile		
1865		1870	1875		
Val Ser	Pro His His Thr	Ser Glu Asp Ser Glu Leu	Ile Ala Glu		
1880		1885	1890		
Ala Lys	Leu Leu Arg Gln	His Lys Gly Arg Leu Glu	Ala Arg Met		
1895		1900	1905		
Gln Ile	Leu Glu Asp His	Asn Lys Gln Leu Glu Ser	Gln Leu His		
1910		1915	1920		
Arg Leu	Arg Gln Leu Leu	Glu Gln Pro Asp Ser Asp	Ser Arg Ile		
1925		1930	1935		
Asn Gly	Val Ser Pro Trp	Ala Ser Pro Gln His Ser	Ala Leu Ser		
1940		1945	1950		

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Tyr Ser Leu Asp Thr Asp Pro Gly Pro Gln Phe His Gln Ala Ala
 1955 1960 1965

Ser Glu Asp Leu Leu Ala Pro Pro His Asp Thr Ser Thr Asp Leu
 1970 1975 1980

Thr Asp Val Met Glu Gln Ile Asn Ser Thr Phe Pro Ser Cys Ser
 1985 1990 1995

Ser Asn Val Pro Ser Arg Pro Gln Ala Met
 2000 2005

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cgc cag cgc cgc cgc ggt gga tcc acc atg tcc ggc tat cca tat gac	96
Arg Gln Arg Arg Arg Gly Gly Ser Thr Met Ser Gly Tyr Pro Tyr Asp	
20 25 30	
gtc cca gac tat gct ggc tcc atg gcc aag tat ggg gac ctt gaa gcc	144
Val Pro Asp Tyr Ala Gly Ser Met Ala Lys Tyr Gly Asp Leu Glu Ala	
35 40 45	
agg cct gat gat ggg cag aac gaa ttc agt gac atc att aag tcc aga	192
Arg Pro Asp Asp Gly Gln Asn Glu Phe Ser Asp Ile Ile Lys Ser Arg	
50 55 60	
tct gat gaa cac aat gat gta cag aag aaa acc ttt acc aaa tgg ata	240
Ser Asp Glu His Asn Asp Val Gln Lys Lys Thr Phe Thr Lys Trp Ile	
65 70 75 80	
aac gct cga ttt tcc aag agt ggg aaa cca ccc atc agt gat atg ttc	288
Asn Ala Arg Phe Ser Lys Ser Gly Lys Pro Pro Ile Ser Asp Met Phe	
85 90 95	
tca gac ctc aaa gat ggg aga aag ctc ttg gat ctt ctc gaa ggc ctc	336
Ser Asp Leu Lys Asp Gly Arg Lys Leu Leu Asp Leu Leu Glu Gly Leu	
100 105 110	
aca gga aca tca ttg cca aag gaa cgt ggt tcc aca agg gtg cat gcc	384
Thr Gly Thr Ser Leu Pro Lys Glu Arg Gly Ser Thr Arg Val His Ala	
115 120 125	
tta aac aat gtc aac cga gtg cta cag gtt tta cat cag aac aat gtg	432
Leu Asn Asn Val Asn Arg Val Leu Gln Val Leu His Gln Asn Asn Val	
130 135 140	
gac ttg gtg aat att gga ggc acg gac att gtg gct gga aat ccc aag	480
Asp Leu Val Asn Ile Gly Gly Thr Asp Ile Val Ala Gly Asn Pro Lys	
145 150 155 160	
ctg act tta ggg tta ctc tgg agc atc att ctg cac tgg cag gtg aag	528
Leu Thr Leu Gly Leu Leu Trp Ser Ile Ile Leu His Trp Gln Val Lys	
165 170 175	
gat gtc atg aaa gat atc atg tca gac ctg cag cag aca aac agc gag	576
Asp Val Met Lys Asp Ile Met Ser Asp Leu Gln Gln Thr Asn Ser Glu	
180 185 190	
aag atc ctg ctg agc tgg gtg cgg cag acc acc agg ccc tac agt caa	624
Lys Ile Leu Leu Ser Trp Val Arg Gln Thr Thr Arg Pro Tyr Ser Gln	
195 200 205	

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Asn Ala Val Leu His Arg His Lys Pro Asp Leu Phe Asp Trp Asp Glu	
225 230 235 240	
atg gtc aaa atg tcc cca att gag aga ctt gac cat gct ttt gac aag	768
Met Val Lys Met Ser Pro Ile Glu Arg Leu Asp His Ala Phe Asp Lys	
245 250 255	
gcc cac act tct ttg gga att gaa aag ctc cta agt cct gaa act gtt	816
Ala His Thr Ser Leu Gly Ile Glu Lys Leu Leu Ser Pro Glu Thr Val	
260 265 270	
gct gtg cat ctc cct gac aag aaa tcc ata att atg tat tta acg tct	864
Ala Val His Leu Pro Asp Lys Lys Ser Ile Ile Met Tyr Leu Thr Ser	
275 280 285	
ctg ttt gag gtg ctt cct cag caa gtc acg ata gat gcc atc cga gag	912
Leu Phe Glu Val Leu Pro Gln Gln Val Thr Ile Asp Ala Ile Arg Glu	
290 295 300	
gtg gag act ctc cca agg aag tat aag aaa gaa tgt gaa gag gaa gaa	960
Val Glu Thr Leu Pro Arg Lys Tyr Lys Lys Glu Cys Glu Glu Glu Glu	
305 310 315 320	
att cat atc cag agt gca gtg ctg gca gag gaa ggc cag agt ccc cga	1008
Ile His Ile Gln Ser Ala Val Leu Ala Glu Glu Gly Gln Ser Pro Arg	
325 330 335	
gct gag acc cct agc acc gtc act gaa gtg gac atg gat ttg gac agc	1056
Ala Glu Thr Pro Ser Thr Val Thr Glu Val Asp Met Asp Leu Asp Ser	
340 345 350	
tac cag ata gcg cta gag gaa gtg ctg acg tgg ctg ctg tcc gcg gag	1104
Tyr Gln Ile Ala Leu Glu Glu Val Leu Thr Trp Leu Leu Ser Ala Glu	
355 360 365	
gac acg ttc cag gag caa cat gac att tct gat gat gtc gaa gaa gtc	1152
Asp Thr Phe Gln Glu Gln His Asp Ile Ser Asp Asp Val Glu Glu Val	
370 375 380	
aaa gag cag ttt gct acc cat gaa act ttt atg atg gag ctg aca gca	1200
Lys Glu Gln Phe Ala Thr His Glu Thr Phe Met Met Glu Leu Thr Ala	
385 390 395 400	
cac cag agc agc gtg ggg agc gtc ctg cag gct ggc aac cag ctg atg	1248
His Gln Ser Ser Val Gly Ser Val Leu Gln Ala Gly Asn Gln Leu Met	
405 410 415	
aca caa ggg act ctg tcc aga gag gag gag ttt gag atc cag gaa cag	1296
Thr Gln Gly Thr Leu Ser Arg Glu Glu Glu Phe Glu Ile Gln Glu Gln	
420 425 430	
atg acc ttg ctg aat gca agg tgg gag gcg ctc cgg gtg gag agc atg	1344
Met Thr Leu Leu Asn Ala Arg Trp Glu Ala Leu Arg Val Glu Ser Met	
435 440 445	
gag agg cag tcc cgg ctg cac gac gct ctg atg gag ctg cag aag aaa	1392
Glu Arg Gln Ser Arg Leu His Asp Ala Leu Met Glu Leu Gln Lys Lys	
450 455 460	
cag ctg cag cag ctc tca agc tgg ctg gcc ctc aca gaa gag gcg att	1440
Gln Leu Gln Gln Leu Ser Ser Trp Leu Ala Leu Thr Glu Glu Arg Ile	
465 470 475 480	
cag aag atg gag agc ctc ccg ctg ggt gat gac ctg ccc tcc ctg cag	1488
Gln Lys Met Glu Ser Leu Pro Leu Gly Asp Asp Leu Pro Ser Leu Gln	
485 490 495	
aag ctg ctt caa gaa cat aaa agt ttg caa aat gac ctt gaa gct gaa	1536
Lys Leu Leu Gln Glu His Lys Ser Leu Gln Asn Asp Leu Glu Ala Glu	
500 505 510	
cag gtg aag gta aat tcc tta act cac atg gtg gtg att gtg gat gaa	1584
Gln Val Lys Val Asn Ser Leu Thr His Met Val Val Ile Val Asp Glu	

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515		520		525		
aac agt ggg gag agt gcc aca gct ctt ctg gaa gat cag tta cag aaa						1632
Asn Ser Gly Glu Ser Ala Thr Ala Leu Leu Glu Asp Gln Leu Gln Lys						
530		535		540		
ctg ggt gag cgc tgg aca gct gta tgc cgc tgg act gaa gaa cgt tgg						1680
Leu Gly Glu Arg Trp Thr Ala Val Cys Arg Trp Thr Glu Glu Arg Trp						
545		550		555		560
aac agg ttg caa gaa atc agt att ctg tgg cag gaa tta ttg gaa gag						1728
Asn Arg Leu Gln Glu Ile Ser Ile Leu Trp Gln Glu Leu Leu Glu Glu						
	565			570		575
cag tgt ctg ttg gag gct tgg ctc acc gaa aag gaa gag gct ttg gat						1776
Gln Cys Leu Leu Glu Ala Trp Leu Thr Glu Lys Glu Glu Ala Leu Asp						
	580			585		590
aaa gtt caa acc agc aac ttt aaa gac cag aag gaa cta agt gtc agt						1824
Lys Val Gln Thr Ser Asn Phe Lys Asp Gln Lys Glu Leu Ser Val Ser						
	595			600		605
gtc cgg cgt ctg gct ata ttg aag gaa gac atg gaa atg aag agg cag						1872
Val Arg Arg Leu Ala Ile Leu Lys Glu Asp Met Glu Met Lys Arg Gln						
	610			615		620
act ctg gat caa ctg agt gag att ggc cag gat gtg ggc caa tta ctc						1920
Thr Leu Asp Gln Leu Ser Glu Ile Gly Gln Asp Val Gly Gln Leu Leu						
	625			630		635
agt aat ccc aag gca tct aag aag atg aac agt gac tct gag gag cta						1968
Ser Asn Pro Lys Ala Ser Lys Lys Met Asn Ser Asp Ser Glu Glu Leu						
	645			650		655
aca cag aga tgg gat tct ctg gtt cag aga ctc gaa gac tct tct aac						2016
Thr Gln Arg Trp Asp Ser Leu Val Gln Arg Leu Glu Asp Ser Ser Asn						
	660			665		670
cag gtg act cag gcg gta gcg aag ctc ggc atg tcc cag att cca cag						2064
Gln Val Thr Gln Ala Val Ala Lys Leu Gly Met Ser Gln Ile Pro Gln						
	675			680		685
aag gac cta ttg gag acc gtt cat gtg aga gaa caa ggg atg gtg aag						2112
Lys Asp Leu Leu Glu Thr Val His Val Arg Glu Gln Gly Met Val Lys						
	690			695		700
aag ccc aag cag gaa ctg cct cct cct ccc cca cca aag aag aga cag						2160
Lys Pro Lys Lys Glu Glu Leu Pro Pro Pro Pro Pro Lys Lys Arg Gln						
	705			710		715
att cac gtg gac gtg gag gcc aag aaa aag ttt gat gct ata agt aca						2208
Ile His Val Asp Val Glu Ala Lys Lys Lys Phe Asp Ala Ile Ser Thr						
	725			730		735
gag ctg ctg aac tgg att ttg aaa tca aag act gcc att cag aac aca						2256
Glu Leu Leu Asn Trp Ile Leu Lys Ser Lys Thr Ala Ile Gln Asn Thr						
	740			745		750
gag atg aaa gaa tat aag aag tcg cag gag acc tca gga atg aaa aag						2304
Glu Met Lys Lys Glu Tyr Lys Lys Ser Gln Glu Thr Ser Gly Met Lys Lys						
	755			760		765
aaa ttg aag gga tta gag aaa gaa cag aag gaa aat ctg ccc cga ctg						2352
Lys Leu Lys Gly Leu Glu Lys Glu Gln Lys Glu Asn Leu Pro Arg Leu						
	770			775		780
gac gaa ctg aat caa acc gga caa acc ctc cgg gag caa atg gga aaa						2400
Asp Glu Leu Asn Gln Thr Gly Gln Thr Leu Arg Glu Glu Gln Met Gly Lys						
	785			790		795
gaa ggc ctt cca ctg aaa gaa gta aac gat gtt ctg gaa agg gtt tcg						2448
Glu Gly Leu Pro Leu Lys Glu Val Asn Asp Val Leu Glu Arg Val Ser						
	805			810		815
ttg gag tgg aag atg ata tct cag cag cta gaa gat ctg gga agg aag						2496
Leu Glu Trp Lys Met Ile Ser Gln Gln Leu Glu Asp Leu Gly Arg Lys						
	820			825		830
atc cag ctg cag gaa gat ata aat gct tat ttt aag cag ctt gat gcc						2544

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Ile Gln Leu Gln Glu Asp Ile Asn Ala Tyr Phe Lys Gln Leu Asp Ala 835 840 845	
att gag gag acc atc aag gag aag gaa gag tgg ctg agg ggc aca ccc Ile Glu Glu Thr Ile Lys Glu Lys Glu Glu Trp Leu Arg Gly Thr Pro 850 855 860	2592
att tct gaa tcg ccc cgg cag ccc ttg cca ggc tta aag gat tct tgc Ile Ser Glu Ser Pro Arg Gln Pro Leu Pro Gly Leu Lys Asp Ser Cys 865 870 875 880	2640
cag agg gaa ctg aca gat ctc ctt ggc ctt cac ccc aga att gag acg Gln Arg Glu Leu Thr Asp Leu Leu Gly Leu His Pro Arg Ile Glu Thr 885 890 895	2688
ctg tgt gca agc tgt tca gcc ctg aag tct cag ccc tgt gtc cca ggt Leu Cys Ala Ser Cys Ser Ala Leu Lys Ser Gln Pro Cys Val Pro Gly 900 905 910	2736
ttt gtc cag cag ggt ttt gac gac ctt cga cat cat tac cag gct gtt Phe Val Gln Gln Gly Phe Asp Asp Leu Arg His His Tyr Gln Ala Val 915 920 925	2784
gcg aag gct tta gag gaa tac caa caa caa cta gaa aat gag ctg aag Ala Lys Ala Leu Glu Glu Tyr Gln Gln Gln Leu Glu Asn Glu Leu Lys 930 935 940	2832
agc cag cct gga ccc gag tat ttg gac aca ctg aat acc ctg aaa aaa Ser Gln Pro Gly Pro Glu Tyr Leu Asp Thr Leu Asn Thr Leu Lys Lys 945 950 955 960	2880
atg cta agc gag tca gaa aag gcg gcc cag gcc tct ctg aat gcc ctg Met Leu Ser Glu Ser Glu Lys Ala Ala Gln Ala Ser Leu Asn Ala Leu 965 970 975	2928
aac gat ccc ata gcg gtg gag cag gcc ctg cag gag aaa aag gcc ctt Asn Asp Pro Ile Ala Val Glu Gln Ala Leu Gln Glu Lys Lys Ala Leu 980 985 990	2976
gat gaa acc ctt gag aat cag aaa cat acg tta cat aag ctt tca gaa Asp Glu Thr Leu Glu Asn Gln Lys His Thr Leu His Lys Leu Ser Glu 995 1000 1005	3024
gaa acg aag act ttg gag aaa aat atg ctt cct gat gtg ggg aaa Glu Thr Lys Thr Leu Glu Lys Asn Met Leu Pro Asp Val Gly Lys 1010 1015 1020	3069
atg tat aaa caa gaa ttt gat gat gtc caa ggc aga tgg aat aaa Met Tyr Lys Gln Glu Phe Asp Asp Val Gln Gly Arg Trp Asn Lys 1025 1030 1035	3114
gta aag acc aag gtt tcc aga gac tta cac ttg ctc gag gaa atc Val Lys Thr Lys Val Ser Arg Asp Leu His Leu Leu Glu Glu Ile 1040 1045 1050	3159
acc ccc aga ctc cga gat ttt gag gct gat tca gaa gtc att gag Thr Pro Arg Leu Arg Asp Phe Glu Ala Asp Ser Glu Val Ile Glu 1055 1060 1065	3204
aag tgg gtg agt ggc atc aaa gac ttc ctc atg aaa gaa cag gct Lys Trp Val Ser Gly Ile Lys Asp Phe Leu Met Lys Glu Gln Ala 1070 1075 1080	3249
gcc caa gga gac gct gct gcg cag agc cag ctt gac caa tgt gct Ala Gln Gly Asp Ala Ala Ala Gln Ser Gln Leu Asp Gln Cys Ala 1085 1090 1095	3294
acg ttt gct aat gaa atc gaa acc atc gag tca tct ctg aag aac Thr Phe Ala Asn Glu Ile Glu Thr Ile Glu Ser Ser Leu Lys Asn 1100 1105 1110	3339
atg agg gaa gta gag act agc ctt cag agg tgt cca gtc act gga Met Arg Glu Val Glu Thr Ser Leu Gln Arg Cys Pro Val Thr Gly 1115 1120 1125	3384
gtc aag aca tgg gta cag gca aga cta gtg gat tac caa tcc caa Val Lys Thr Trp Val Gln Ala Arg Leu Val Asp Tyr Gln Ser Gln 1130 1135 1140	3429

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ctg gag	aaa ttc agc	aaa gag	att gct att	caa aaa	agc agg ctg	3474
Leu Glu	Lys Phe Ser	Lys Glu	Ile Ala Ile	Gln Lys	Ser Arg Leu	
1145		1150		1155		
tta gat	agt caa gaa	aaa gcc	ctg aac ttg	aaa aag	gat ttg gct	3519
Leu Asp	Ser Gln Glu	Lys Ala	Leu Asn Leu	Lys Lys	Asp Leu Ala	
1160		1165		1170		
gag atg	cag gag tgg	atg gca	cag gct gaa	gag gac	tac ctg gag	3564
Glu Met	Gln Glu Trp	Met Ala	Gln Ala Glu	Glu Asp	Tyr Leu Glu	
1175		1180		1185		
agg gac	ttc gag tac	aaa tct	cca gaa gaa	ctc gag	agt gcg gtg	3609
Arg Asp	Phe Glu Tyr	Lys Ser	Pro Glu Glu	Leu Glu	Ser Ala Val	
1190		1195		1200		
gag gaa	atg aag agg	gca aaa	gag gat gtg	ctg cag	aag gag gtg	3654
Glu Glu	Met Lys Arg	Ala Lys	Glu Asp Val	Leu Gln	Lys Glu Val	
1205		1210		1215		
agg gtg	aaa att ctg	aag gac	agc atc aag	ctg gtg	gct gcc aag	3699
Arg Val	Lys Ile Leu	Lys Asp	Ser Ile Lys	Leu Val	Ala Ala Lys	
1220		1225		1230		
gtg ccc	tct ggt ggc	cag gag	ttg acg tcg	gaa ttc	aac gag gtg	3744
Val Pro	Ser Gly Gly	Gln Glu	Leu Thr Ser	Glu Phe	Asn Glu Val	
1235		1240		1245		
ctg gag	agc tac cag	ctt ctg	tgc aat aga	att cga	ggg aag tgc	3789
Leu Glu	Ser Tyr Gln	Leu Leu	Cys Asn Arg	Ile Arg	Gly Lys Cys	
1250		1255		1260		
cac aca	ctg gag gag	gtc tgg	tct tgc tgg	gtg gag	ctg ctt cac	3834
His Thr	Leu Glu Glu	Val Trp	Ser Cys Trp	Val Glu	Leu Leu His	
1265		1270		1275		
tat ctg	gac ctg gag	acc acg	tgg ttg aac	acc ttg	gag gag cgc	3879
Tyr Leu	Asp Leu Glu	Thr Thr	Trp Leu Asn	Thr Leu	Glu Glu Arg	
1280		1285		1290		
gtg agg	agc acg gag	gcc ctg	cct gag agg	gca gaa	gct gtt cat	3924
Val Arg	Ser Thr Glu	Ala Leu	Pro Glu Arg	Ala Glu	Ala Val His	
1295		1300		1305		
gaa gct	ctg gag tct	ctt gag	tct gtt ttg	cgc cat	cca gcg gat	3969
Glu Ala	Leu Glu Ser	Leu Glu	Ser Val Leu	Arg His	Pro Ala Asp	
1310		1315		1320		
aat cgc	acc cag att	cgg gaa	ctt ggg cag	act ctg	att gat ggt	4014
Asn Arg	Thr Gln Ile	Arg Glu	Leu Gly Gln	Thr Leu	Ile Asp Gly	
1325		1330		1335		
gga atc	ctg gat gac	ata atc	agc gag aag	ctg gag	gct ttt aac	4059
Gly Ile	Leu Asp Asp	Ile Ile	Ser Glu Lys	Leu Glu	Ala Phe Asn	
1340		1345		1350		
agc cgc	tac gaa gag	ctg agt	cac ttg gcg	gag agc	aaa cag att	4104
Ser Arg	Tyr Glu Glu	Leu Ser	His Leu Ala	Glu Ser	Lys Gln Ile	
1355		1360		1365		
tct ttg	gag aag caa	ctc cag	gtc ctc cgc	gaa act	gac cac atg	4149
Ser Leu	Glu Lys Gln	Leu Gln	Val Leu Arg	Glu Thr	Asp His Met	
1370		1375		1380		
ctt cag	gtg ctg aag	gag agc	ctg ggg gag	ctg gac	aaa cag ctt	4194
Leu Gln	Val Leu Lys	Glu Ser	Leu Gly Glu	Leu Asp	Lys Gln Leu	
1385		1390		1395		
acc aca	tac ctg acg	gac agg	atc gat gcc	ttc caa	ctg cca cag	4239
Thr Thr	Tyr Leu Thr	Asp Arg	Ile Asp Ala	Phe Gln	Leu Pro Gln	
1400		1405		1410		
gaa gct	cag aag atc	caa gcc	gaa atc tca	gcc cat	gag ctc acc	4284
Glu Ala	Gln Lys Ile	Gln Ala	Glu Ile Ser	Ala His	Glu Leu Thr	
1415		1420		1425		
ctg gag	gag ctg agg	aag aat	gtg cgc tcc	cag ccc	ccg acg tcc	4329
Leu Glu	Glu Leu Arg	Lys Asn	Val Arg Ser	Gln Pro	Pro Thr Ser	
1430		1435		1440		

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cct gag ggc agg gcc acc aga gga gga agt cag atg gac atg cta	4374
Pro Glu Gly Arg Ala Thr Arg Gly Gly Ser Gln Met Asp Met Leu	
1445 1450 1455	
cag agg aaa ctt cga gag gtc tcc acc aaa ttc cag ctt gcc cac	4419
Gln Arg Lys Leu Arg Glu Val Ser Thr Lys Phe Gln Leu Ala His	
1460 1465 1470	
aga gat ttt ggg cca tct tct caa cac ttt ctg tcc act tca gtc	4464
Arg Asp Phe Gly Pro Ser Ser Gln His Phe Leu Ser Thr Ser Val	
1475 1480 1485	
cag ctg ccg tgg cag aga tcc att tca cat aat aaa gtg ccc tat	4509
Gln Leu Pro Trp Gln Arg Ser Ile Ser His Asn Lys Val Pro Tyr	
1490 1495 1500	
tac atc aac cat caa aca cag aca acc tgt tgg gat cat cct aaa	4554
Tyr Ile Asn His Gln Thr Gln Thr Thr Cys Trp Asp His Pro Lys	
1505 1510 1515	
atg act gag ctc ttc caa tcc ctt gct gat ctg aat aat gta cgt	4599
Met Thr Glu Leu Phe Gln Ser Leu Ala Asp Leu Asn Asn Val Arg	
1520 1525 1530	
ttc tct gcc tac cgc aca gca atc aaa att cga agg ctg caa aaa	4644
Phe Ser Ala Tyr Arg Thr Ala Ile Lys Ile Arg Arg Leu Gln Lys	
1535 1540 1545	
gca tta tgt ctg gat ctc tta gag ctg aat acg acg aat gaa gtt	4689
Ala Leu Cys Leu Asp Leu Leu Glu Leu Asn Thr Thr Asn Glu Val	
1550 1555 1560	
ttc aag cag cac aaa ctg aac caa aat gat cag ctc ctg agt gtc	4734
Phe Lys Gln His Lys Leu Asn Gln Asn Asp Gln Leu Leu Ser Val	
1565 1570 1575	
cca gac gtc atc aac tgt ctg acc acc act tac gat ggg ctt gag	4779
Pro Asp Val Ile Asn Cys Leu Thr Thr Thr Tyr Asp Gly Leu Glu	
1580 1585 1590	
cag ctg cac aag gac ttg gtc aat gtt cca ctc tgc gtc gat atg	4824
Gln Leu His Lys Asp Leu Val Asn Val Pro Leu Cys Val Asp Met	
1595 1600 1605	
tgt ctc aac tgg ctg ctc aac gta tac gac acg ggc cgg act gga	4869
Cys Leu Asn Trp Leu Leu Asn Val Tyr Asp Thr Gly Arg Thr Gly	
1610 1615 1620	
aaa att cgg gta cag agt ctg aag att gga ttg atg tct ctc tcc	4914
Lys Ile Arg Val Gln Ser Leu Lys Ile Gly Leu Met Ser Leu Ser	
1625 1630 1635	
aaa ggc ctc tta gaa gag aaa tac aga tgt ctc ttt aag gag gtg	4959
Lys Gly Leu Leu Glu Glu Lys Tyr Arg Cys Leu Phe Lys Glu Val	
1640 1645 1650	
gca ggg cca act gag atg tgt gac cag cgg cag ctt ggc ctg cta	5004
Ala Gly Pro Thr Glu Met Cys Asp Gln Arg Gln Leu Gly Leu Leu	
1655 1660 1665	
ctt cac gat gcc atc cag atc cct agg cag ctg ggg gaa gta gca	5049
Leu His Asp Ala Ile Gln Ile Pro Arg Gln Leu Gly Glu Val Ala	
1670 1675 1680	
gcc ttt ggg ggc agt aac att gag ccc agt gtc cgc agc tgc ttc	5094
Ala Phe Gly Gly Ser Asn Ile Glu Pro Ser Val Arg Ser Cys Phe	
1685 1690 1695	
cag cag aat aac aac aag cca gaa atc agt gtg aag gag ttt ata	5139
Gln Gln Asn Asn Asn Lys Pro Glu Ile Ser Val Lys Glu Phe Ile	
1700 1705 1710	
gac tgg atg cat ttg gaa ccc cag tcc atg gtg tgg ttg ccg gtt	5184
Asp Trp Met His Leu Glu Pro Gln Ser Met Val Trp Leu Pro Val	
1715 1720 1725	
ctg cat cgg gtc gca gct gct gag act gca aaa cat cag gcc aaa	5229
Leu His Arg Val Ala Ala Ala Glu Thr Ala Lys His Gln Ala Lys	

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1730	1735	1740	
tgc aac atc tgc aaa gaa tgc ccg att gtt ggg ttc aga tac agg Cys Asn Ile Cys Lys Glu Cys Pro Ile Val Gly Phe Arg Tyr Arg 1745 1750 1755			5274
agc cta aag cat ttt aat tat gat gtc tgc cag agt tgc ttc ttt Ser Leu Lys His Phe Asn Tyr Asp Val Cys Gln Ser Cys Phe Phe 1760 1765 1770			5319
tct gga aga aca gca aag ggc cac aag tta cat tac ccg atg gta Ser Gly Arg Thr Ala Lys Glu His Lys Leu His Tyr Pro Met Val 1775 1780 1785			5364
gaa tac tgc ata ccg aca aca tct ggg gaa gat gtg aga gat ttc Glu Tyr Cys Ile Pro Thr Thr Ser Gly Glu Asp Val Arg Asp Phe 1790 1795 1800			5409
act aag gtg ctg aag aac aag ttc agg tcc aag aaa tat ttt gcc Thr Lys Val Leu Lys Asn Lys Phe Arg Ser Lys Lys Tyr Phe Ala 1805 1810 1815			5454
aaa cat cct cgg ctt ggc tac ctg cct gtc cag acc gtg ctg gaa Lys His Pro Arg Leu Gly Tyr Leu Pro Val Gln Thr Val Leu Glu 1820 1825 1830			5499
ggg gac aac tta gaa act cct atc acg ctc atc agt atg tgg cca Gly Asp Asn Leu Glu Thr Pro Ile Thr Leu Ile Ser Met Trp Pro 1835 1840 1845			5544
gag cac tat gac ccc tcc cag tcc cct cag ctg ttt cat gat gac Glu His Tyr Asp Pro Ser Gln Ser Pro Gln Leu Phe His Asp Asp 1850 1855 1860			5589
acc cac tca aga ata gag caa tac gct aca cga ctg gcc cag atg Thr His Ser Arg Ile Glu Gln Tyr Ala Thr Arg Leu Ala Gln Met 1865 1870 1875			5634
gaa agg aca aac ggg tcc ttc cta act gat agc agc tct aca aca Glu Arg Thr Asn Gly Ser Phe Leu Thr Asp Ser Ser Ser Thr Thr 1880 1885 1890			5679
gga agc gtg gag gat gag cat gcc ctc atc cag cag tac tgc cag Gly Ser Val Glu Asp Glu His Ala Leu Ile Gln Gln Tyr Cys Gln 1895 1900 1905			5724
acc ctg ggc ggg gag tca cct gtg agt cag ccg cag agt cca gct Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln Ser Pro Ala 1910 1915 1920			5769
cag atc ctg aag tcc gtg gag agg gaa gag cgt ggg gaa ctg gag Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly Glu Leu Glu 1925 1930 1935			5814
cgg atc att gct gac ttg gag gaa gag caa aga aat ctg cag gtg Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn Leu Gln Val 1940 1945 1950			5859
gag tat gag cag ctg aag gag cag cac cta aga agg ggt ctc cct Glu Tyr Glu Gln Leu Lys Glu Gln His Leu Arg Arg Gly Leu Pro 1955 1960 1965			5904
gtg ggc tcc cct cca gac tcc atc gta tct cct cac cac aca tct Val Gly Ser Pro Pro Asp Ser Ile Val Ser Pro His His Thr Ser 1970 1975 1980			5949
gag gac tca gaa ctt ata gca gaa gct aaa ctc ctg cgg cag cac Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg Gln His 1985 1990 1995			5994
aaa ggg cgg ctg gag gcg agg atg caa att ttg gaa gat cac aat Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp His Asn 2000 2005 2010			6039
aaa cag ctg gag tct cag ctg cac cgc ctc aga cag ctc ctg gag Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu Leu Glu 2015 2020 2025			6084
cag cct gac tct gac tcc cgc atc aat ggt gtc tcc ccc tgg gct			6129

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Gln	Pro	Asp	Ser	Asp	Ser	Arg	Ile	Asn	Gly	Val	Ser	Pro	Trp	Ala			
	2030					2035					2040						
tcc	cca	cag	cat	tct	gca	ttg	agc	tac	tca	ctt	gac	act	gac	cca		6174	
Ser	Pro	Gln	His	Ser	Ala	Leu	Ser	Tyr	Ser	Leu	Asp	Thr	Asp	Pro			
	2045					2050					2055						
ggc	cca	cag	ttc	cac	cag	gca	gca	tct	gag	gac	ctg	ctg	gcc	cca		6219	
Gly	Pro	Gln	Phe	His	Gln	Ala	Ala	Ser	Glu	Asp	Leu	Leu	Ala	Pro			
	2060					2065					2070						
cct	cac	gac	act	agc	acg	gac	ctc	acg	gac	gtg	atg	gag	cag	atc		6264	
Pro	His	Asp	Thr	Ser	Thr	Asp	Leu	Thr	Asp	Val	Met	Glu	Gln	Ile			
	2075					2080					2085						
aac	agc	acg	ttt	ccc	tct	tgc	agc	tca	aat	gtc	ccc	agc	agg	cca		6309	
Asn	Ser	Thr	Phe	Pro	Ser	Cys	Ser	Ser	Asn	Val	Pro	Ser	Arg	Pro			
	2090					2095					2100						
cag	gca	atg	tga													6321	
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<210> SEQ ID NO 25

<211> LENGTH: 2106

<212> TYPE: PRT

<213> ORGANISM: Mus musculus

<400> SEQUENCE: 25

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Val	Pro	Asp	Tyr	Ala	Gly	Ser	Met	Ala	Lys	Tyr	Gly	Asp	Leu	Glu	Ala		
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Arg	Pro	Asp	Asp	Gly	Gln	Asn	Glu	Phe	Ser	Asp	Ile	Ile	Lys	Ser	Arg		
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Ser	Asp	Leu	Lys	Asp	Gly	Arg	Lys	Leu	Leu	Asp	Leu	Leu	Glu	Gly	Leu		
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Thr	Gly	Thr	Ser	Leu	Pro	Lys	Glu	Arg	Gly	Ser	Thr	Arg	Val	His	Ala		
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Leu	Asn	Asn	Val	Asn	Arg	Val	Leu	Gln	Val	Leu	His	Gln	Asn	Asn	Val		
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Asp	Leu	Val	Asn	Ile	Gly	Gly	Thr	Asp	Ile	Val	Ala	Gly	Asn	Pro	Lys		
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Leu	Thr	Leu	Gly	Leu	Leu	Trp	Ser	Ile	Ile	Leu	His	Trp	Gln	Val	Lys		
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Asp	Val	Met	Lys	Asp	Ile	Met	Ser	Asp	Leu	Gln	Gln	Thr	Asn	Ser	Glu		
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Lys	Ile	Leu	Leu	Ser	Trp	Val	Arg	Gln	Thr	Thr	Arg	Pro	Tyr	Ser	Gln		
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Val	Asn	Val	Leu	Asn	Phe	Thr	Thr	Ser	Trp	Thr	Asp	Gly	Leu	Ala	Phe		
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Asn	Ala	Val	Leu	His	Arg	His	Lys	Pro	Asp	Leu	Phe	Asp	Trp	Asp	Glu		
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Ala	Glu	Thr	Pro	Ser	Thr	Val	Thr	Glu	Val	Asp	Met	Asp	Leu	Asp	Ser
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Gln	Leu	Gln	Gln	Leu	Ser	Ser	Trp	Leu	Ala	Leu	Thr	Glu	Glu	Arg	Ile
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Gln	Lys	Met	Glu	Ser	Leu	Pro	Leu	Gly	Asp	Asp	Leu	Pro	Ser	Leu	Gln
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Lys	Leu	Leu	Gln	Glu	His	Lys	Ser	Leu	Gln	Asn	Asp	Leu	Glu	Ala	Glu
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Gln	Val	Lys	Val	Asn	Ser	Leu	Thr	His	Met	Val	Val	Ile	Val	Asp	Glu
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Asn	Ser	Gly	Glu	Ser	Ala	Thr	Ala	Leu	Leu	Glu	Asp	Gln	Leu	Gln	Lys
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Asn	Arg	Leu	Gln	Glu	Ile	Ser	Ile	Leu	Trp	Gln	Glu	Leu	Leu	Glu	Glu
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Gln	Cys	Leu	Leu	Glu	Ala	Trp	Leu	Thr	Glu	Lys	Glu	Glu	Ala	Leu	Asp
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Val	Arg	Arg	Leu	Ala	Ile	Leu	Lys	Glu	Asp	Met	Glu	Met	Lys	Arg	Gln
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Thr	Leu	Asp	Gln	Leu	Ser	Glu	Ile	Gly	Gln	Asp	Val	Gly	Gln	Leu	Leu
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Ser	Asn	Pro	Lys	Ala	Ser	Lys	Lys	Met	Asn	Ser	Asp	Ser	Glu	Glu	Leu
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			660					665					670		
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675					680					685					
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Glu	Leu	Leu	Asn	Trp	Ile	Leu	Lys	Ser	Lys	Thr	Ala	Ile	Gln	Asn	Thr
			740					745					750		
Glu	Met	Lys	Glu	Tyr	Lys	Lys	Ser	Gln	Glu	Thr	Ser	Gly	Met	Lys	Lys
		755					760					765			
Lys	Leu	Lys	Gly	Leu	Glu	Lys	Glu	Gln	Lys	Glu	Asn	Leu	Pro	Arg	Leu
	770					775					780				
Asp	Glu	Leu	Asn	Gln	Thr	Gly	Gln	Thr	Leu	Arg	Glu	Gln	Met	Gly	Lys
785				790						795					800
Glu	Gly	Leu	Pro	Leu	Lys	Glu	Val	Asn	Asp	Val	Leu	Glu	Arg	Val	Ser
				805					810					815	
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			820						825				830		
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Ile	Glu	Glu	Thr	Ile	Lys	Glu	Lys	Glu	Glu	Trp	Leu	Arg	Gly	Thr	Pro
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945				950						955					960
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			980						985					990	
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		995					1000					1005			
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	1070					1075					1080				
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Val	Lys	Thr	Trp	Val	Gln	Ala	Arg	Leu	Val	Asp	Tyr	Gln	Ser	Gln
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Leu	Glu	Lys	Phe	Ser	Lys	Glu	Ile	Ala	Ile	Gln	Lys	Ser	Arg	Leu
1145						1150						1155		
Leu	Asp	Ser	Gln	Glu	Lys	Ala	Leu	Asn	Leu	Lys	Lys	Asp	Leu	Ala
1160						1165						1170		
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1175						1180						1185		
Arg	Asp	Phe	Glu	Tyr	Lys	Ser	Pro	Glu	Glu	Leu	Glu	Ser	Ala	Val
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Arg	Val	Lys	Ile	Leu	Lys	Asp	Ser	Ile	Lys	Leu	Val	Ala	Ala	Lys
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1445						1450						1455		
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1475						1480						1485		

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Tyr	Ile	Asn	His	Gln	Thr	Gln	Thr	Thr	Cys	Trp	Asp	His	Pro	Lys
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Met	Thr	Glu	Leu	Phe	Gln	Ser	Leu	Ala	Asp	Leu	Asn	Asn	Val	Arg
1520						1525					1530			
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1580						1585					1590			
Gln	Leu	His	Lys	Asp	Leu	Val	Asn	Val	Pro	Leu	Cys	Val	Asp	Met
1595						1600					1605			
Cys	Leu	Asn	Trp	Leu	Leu	Asn	Val	Tyr	Asp	Thr	Gly	Arg	Thr	Gly
1610						1615					1620			
Lys	Ile	Arg	Val	Gln	Ser	Leu	Lys	Ile	Gly	Leu	Met	Ser	Leu	Ser
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Cys	Asn	Ile	Cys	Lys	Glu	Cys	Pro	Ile	Val	Gly	Phe	Arg	Tyr	Arg
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Glu	Tyr	Cys	Ile	Pro	Thr	Thr	Ser	Gly	Glu	Asp	Val	Arg	Asp	Phe
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Thr	Lys	Val	Leu	Lys	Asn	Lys	Phe	Arg	Ser	Lys	Lys	Tyr	Phe	Ala
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Glu	His	Tyr	Asp	Pro	Ser	Gln	Ser	Pro	Gln	Leu	Phe	His	Asp	Asp
1850						1855					1860			
Thr	His	Ser	Arg	Ile	Glu	Gln	Tyr	Ala	Thr	Arg	Leu	Ala	Gln	Met
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Thr Leu Gly Gly Glu Ser Pro Val Ser Gln Pro Gln Ser Pro Ala 1910	1915	1920
Gln Ile Leu Lys Ser Val Glu Arg Glu Glu Arg Gly Glu Leu Glu 1925	1930	1935
Arg Ile Ile Ala Asp Leu Glu Glu Glu Gln Arg Asn Leu Gln Val 1940	1945	1950
Glu Tyr Glu Gln Leu Lys Glu Gln His Leu Arg Arg Gly Leu Pro 1955	1960	1965
Val Gly Ser Pro Pro Asp Ser Ile Val Ser Pro His His Thr Ser 1970	1975	1980
Glu Asp Ser Glu Leu Ile Ala Glu Ala Lys Leu Leu Arg Gln His 1985	1990	1995
Lys Gly Arg Leu Glu Ala Arg Met Gln Ile Leu Glu Asp His Asn 2000	2005	2010
Lys Gln Leu Glu Ser Gln Leu His Arg Leu Arg Gln Leu Leu Glu 2015	2020	2025
Gln Pro Asp Ser Asp Ser Arg Ile Asn Gly Val Ser Pro Trp Ala 2030	2035	2040
Ser Pro Gln His Ser Ala Leu Ser Tyr Ser Leu Asp Thr Asp Pro 2045	2050	2055
Gly Pro Gln Phe His Gln Ala Ala Ser Glu Asp Leu Leu Ala Pro 2060	2065	2070
Pro His Asp Thr Ser Thr Asp Leu Thr Asp Val Met Glu Gln Ile 2075	2080	2085
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Gln Ala Met 2105		

<210> SEQ ID NO 26

<211> LENGTH: 59

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: PCR primer

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59

<210> SEQ ID NO 27

<211> LENGTH: 32

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: PCR primer

<400> SEQUENCE: 27

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32

<210> SEQ ID NO 28

<211> LENGTH: 4

<212> TYPE: PRT

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<213> ORGANISM: Artificial
 <220> FEATURE:
 <223> OTHER INFORMATION: Artificial polypeptide/FLAG fragment

<400> SEQUENCE: 28

Asp Tyr Lys Asp
 1

What is claimed is:

1. An isolated fusion protein comprising:
 - a first protein region comprising a human immunodeficiency virus transactivator protein (HIV-TAT) or a transduction-effective fragment thereof which is effective to transduce the fusion protein into mammalian muscle cells, operationally linked to;
 - a second protein region comprising a full-length utrophin protein or an anti-dystrophinopathic fragment thereof.
2. The isolated fusion protein of claim 1, further comprising an affinity tag operationally linked to the fusion protein.
3. The isolated fusion protein of claim 2, wherein the affinity tag comprises an amino acid sequence DYKDDDDK (SEQ. ID. NO: 1) or a fragment thereof.
4. The isolated fusion protein of claim 2, wherein the affinity tag comprises an amino acid sequence DYKD (SEQ. ID. NO: 28).
5. The isolated fusion protein of claim 1, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain domain plus 4, 7, 10, or 11 spectrin-like repeats.
6. The isolated fusion protein of claim 1, wherein the second protein region is an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.
7. The isolated fusion protein of claim 1, which is an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 11, 13, 15, 17, 19, 21, 23, and 25.
8. The isolated fusion protein of claim 1, wherein the first protein region is an amino acid sequence as shown in SEQ. ID. NO: 2.
9. The isolated fusion protein of claim 8, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain domain plus 4, 7, 10, or 11 spectrin-like repeats.
10. The isolated fusion protein of claim 8, wherein the second protein region is an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.
11. The isolated fusion protein of claim 1, wherein the first protein region is an amino acid sequence as shown in SEQ. ID. NO: 5: YGRKKRRQRRR.
12. The isolated fusion protein of claim 11, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain domain plus 4, 7, 10, or 11 spectrin-like repeats.
13. The isolated fusion protein of claim 11, wherein the second protein region is an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.
14. The isolated fusion protein of claim 1, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising no more than 75% of the mass of the full-length utrophin protein.
15. The isolated fusion protein of claim 1, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising no more than 50% of the mass of the full-length utrophin protein.
16. The isolated fusion protein of claim 1, wherein the second protein region is an anti-dystrophinopathic utrophin fragment comprising no more than 25% of the mass of the full-length utrophin protein.
17. Pharmaceutically suitable salts of the isolated fusion protein recited in claim 1.
18. A pharmaceutical composition for treating dystrophinopathies in mammals, including humans, comprising: an isolated fusion protein or a pharmaceutically suitable salt thereof as recited in any one of claims 1-7 and 8-17, in combination with a pharmaceutically suitable carrier.
19. A method of treating dystrophinopathies in mammals, the method comprising administering to a mammalian subject in need thereof an anti-dystrophinopathic amount of an isolated fusion protein or a pharmaceutically suitable salt thereof as recited in any one of claims 1-7 and 8-17.
20. An isolated nucleic acid expression construct encoding a fusion protein, the nucleic acid expression construct comprising:
 - a first nucleic acid region that encodes a first protein region of the fusion protein, wherein the first protein region comprises a human immunodeficiency virus transactivator protein (HIV-TAT) or a transduction-effective fragment thereof which is effective to transduce the fusion protein into mammalian muscle cells, operationally linked to;
 - a second nucleic acid region that encodes a second protein region of the fusion protein, wherein the second protein region comprises a full-length utrophin protein or an anti-dystrophinopathic fragment thereof;
 wherein the expression construct drives expression of the fusion protein when transformed into a suitable host cell or disposed into a suitable cell-free expression system.
21. The isolated nucleic acid expression construct of claim 2, further comprising a third nucleic acid region that encodes an affinity tag that is operationally linked to the fusion protein.
22. The isolated nucleic acid expression construct of claim 20, wherein the third nucleic acid region encodes an amino acid sequence DYKDDDDK (SEQ. ID. NO: 1) or a fragment thereof.
23. The isolated nucleic acid expression construct of claim 20, wherein the third nucleic acid region encodes an amino acid sequence DYKD (SEQ. ID. NO: 28).
24. The isolated nucleic acid expression construct of claim 20, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain domain plus 4, 7, 10, or 11 spectrin-like repeats.

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25. The isolated nucleic acid expression construct of claim 20, wherein the second nucleic acid region encodes an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.

26. The isolated nucleic acid expression construct of claim 20, which is a nucleic acid sequence selected from the group consisting of SEQ. ID. NOS: 10, 12, 14, 16, 18, 20, 22, and 24.

27. The isolated nucleic acid expression construct of claim 20, wherein the first nucleic acid region encodes an amino acid sequence as shown in SEQ. ID. NO: 2.

28. The isolated nucleic acid expression construct of claim 27, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain plus 4, 7, 10, or 11 spectrin-like repeats.

29. The isolated nucleic acid expression construct of claim 27, wherein the second nucleic acid region encodes an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.

30. The isolated nucleic acid expression construct of claim 20, wherein the first nucleic acid region encodes an amino acid sequence as shown in SEQ. ID. NO: 5: YGRKKRRQRRR.

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31. The isolated nucleic acid expression construct of claim 30, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising an amino-terminal actin-binding domain of the utrophin protein, or the amino-terminal actin-binding domain plus 4, 7, 10, or 11 spectrin-like repeats.

32. The isolated nucleic acid expression construct of claim 30, wherein the second nucleic acid region encodes an amino acid sequence selected from the group consisting of SEQ. ID. NOS: 7 and 9.

33. The isolated nucleic acid expression construct of claim 20, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising no more than 75% of the mass of the full-length utrophin protein.

34. The isolated nucleic acid expression construct of claim 20, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising no more than 50% of the mass of the full-length utrophin protein.

35. The isolated nucleic acid expression construct of claim 20, wherein the second nucleic acid region encodes an anti-dystrophinopathic utrophin fragment comprising no more than 25% of the mass of the full-length utrophin protein.

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