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(54) **RNA-GUIDED GENE EDITING AND GENE REGULATION**

(71) Applicant: **Duke University**, Durham, NC (US)

(72) Inventors: **Charles A. Gersbach**, Durham, NC (US); **Isaac B. Hilton**, Durham, NC (US); **Pablo Perez-Pinera**, Lynn, MA (US); **Ami M. Kabadi**, Durham, NC (US); **Pratiksha I. Thakore**, Durham, NC (US); **David G. Ousterout**, Atlanta, GA (US); **Joshua B. Black**, Durham, NC (US)

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Related U.S. Application Data

(63) Continuation of application No. 14/895,316, filed on Dec. 2, 2015, now Pat. No. 10,704,060, filed as application No. PCT/US14/41190 on Jun. 5, 2014.

(60) Provisional application No. 61/831,481, filed on Jun. 5, 2013, provisional application No. 61/839,127, filed on Jun. 25, 2013, provisional application No. 61/904,911, filed on Nov. 15, 2013, provisional application No. 61/967,466, filed on Mar. 19, 2014, provisional application No. 61/981,575, filed on Apr. 18, 2014.

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A61K 48/00	(2006.01)
C12N 9/96	(2006.01)
C07K 14/47	(2006.01)
C12N 9/22	(2006.01)
C12N 15/90	(2006.01)

(52) **U.S. Cl.**

CPC ***C12N 15/85*** (2013.01); ***A61K 38/465*** (2013.01); ***A61K 48/0058*** (2013.01); ***C12N 9/96*** (2013.01); ***C07K 14/4708*** (2013.01); ***C12N 9/22*** (2013.01); *C12N 2840/20* (2013.01); *C12Y 301/00* (2013.01); *A61K 48/005* (2013.01); *C07K 2319/71* (2013.01); *C12N 2740/16043* (2013.01); *C12N 2800/40* (2013.01); ***C12N 15/907*** (2013.01)

(57) **ABSTRACT**

Disclosed herein are Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR)/CRISPR-associated (Cas) 9-based system related compositions and methods of using said CRISPR/Cas9-based system related compositions for altering gene expression and genome engineering. Also disclosed herein are compositions and methods of using said compositions for altering gene expression and genome engineering in muscle, such as skeletal muscle and cardiac muscle.

Specification includes a Sequence Listing.

Fig. 1A

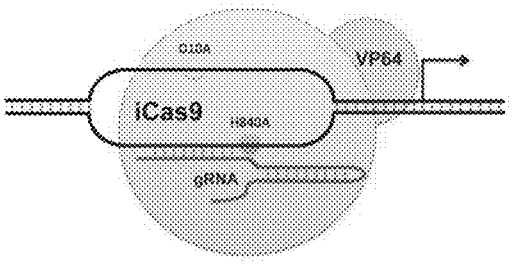


Fig. 1B

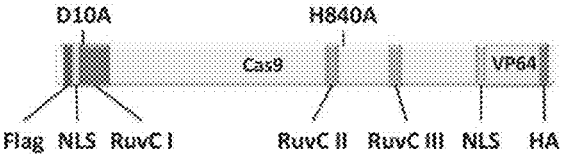


Fig. 1C

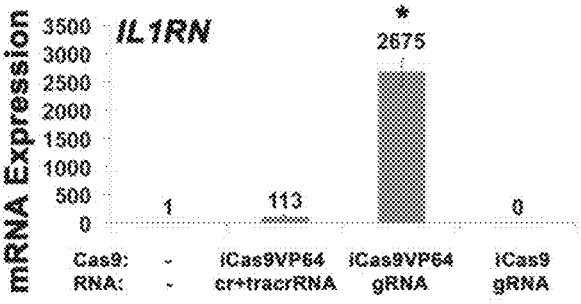


Fig. 1D

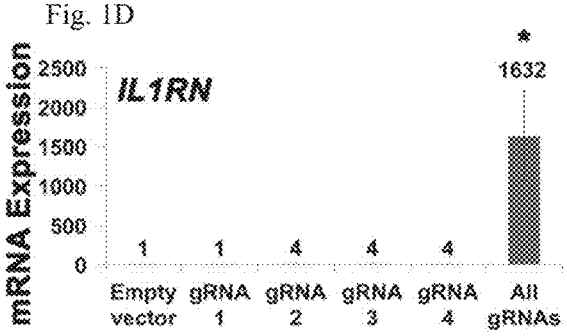


Fig. 1E

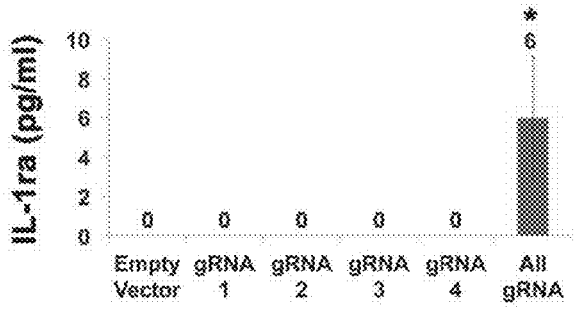
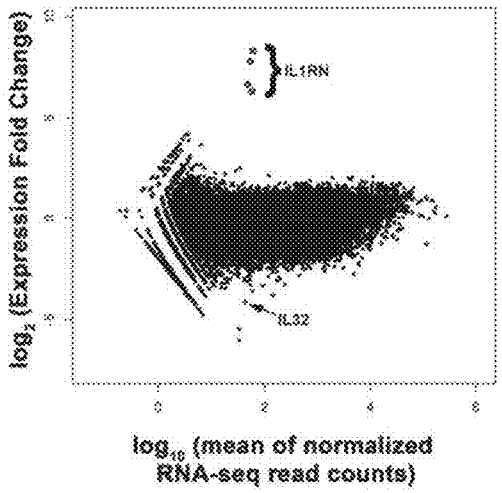


Fig. 1F



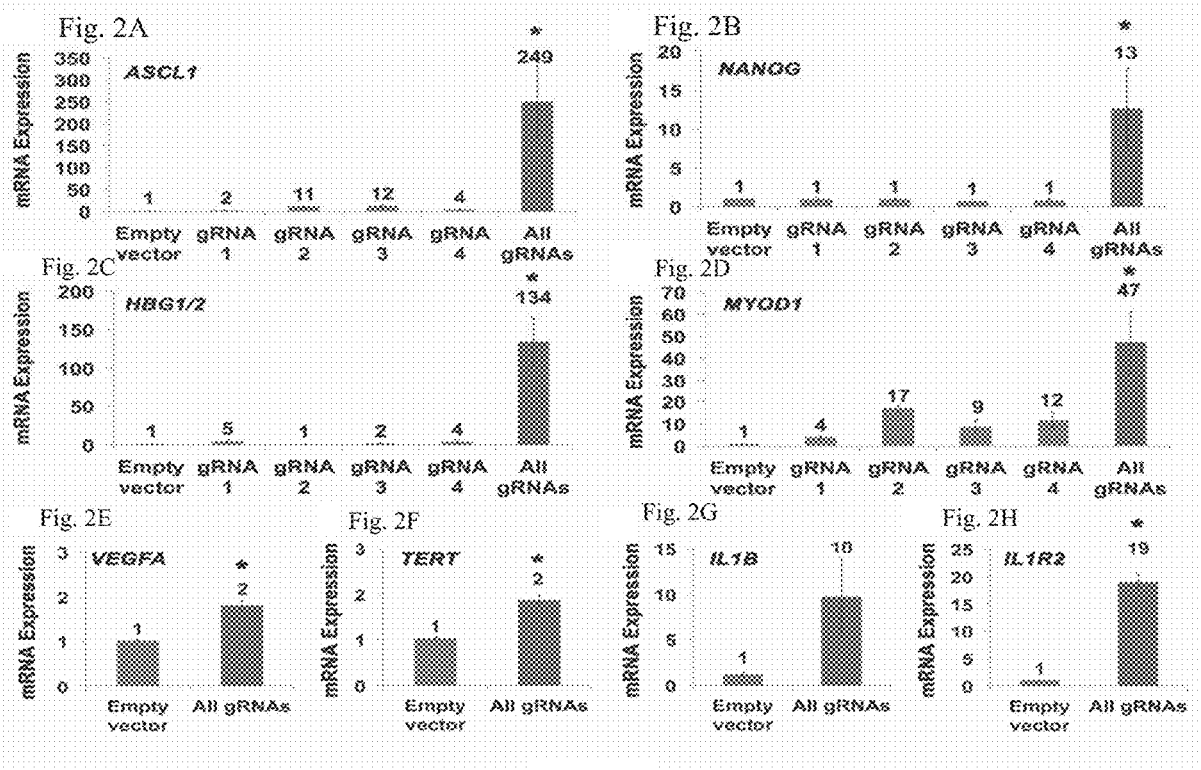


Fig. 2A
ASCL1
mRNA Expression
350
300
250
200
150
100
50
0
1
2
11
12
4
249
*
Empty vector
gRNA 1
gRNA 2
gRNA 3
gRNA 4
All gRNAs
Fig. 2B
NANOG
20
15
10
5
0
1
1
1
1
1
13
Fig. 2C
HBG1/2
200
150
100
50
0
1
5
1
2
4
134
Fig. 2D
MYOD1
70
60
50
40
30
20
10
0
1
4
17
9
12
47
Fig. 2E
VEGFA
3
2
1
0
1
2
Empty vector
All gRNAs
Fig. 2F
TERT
1
2
Fig. 2G
IL1B
15
10
5
0
1
18
Fig. 2H
IL1R2
25
20
15
10
5
0
1
19

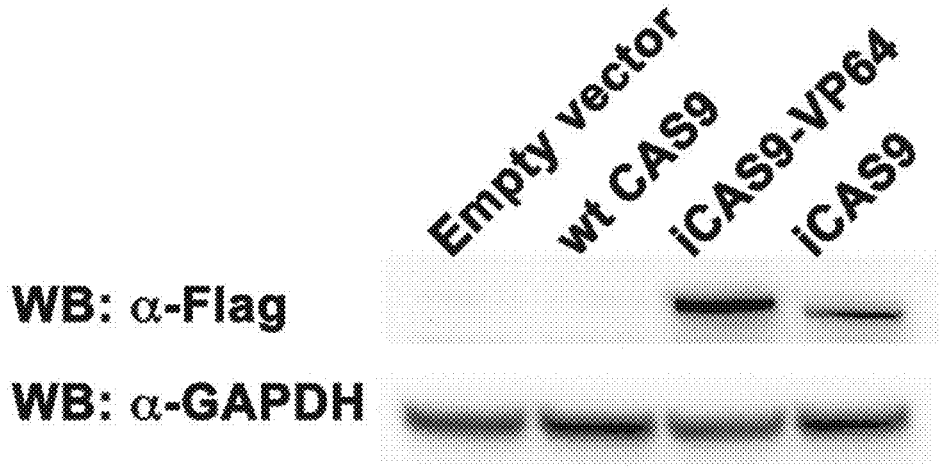

Empty vector
wt CAS9
iCAS9-VP64
iCAS9
WB: α-Flag
WB: α-GAPDH

Fig. 3

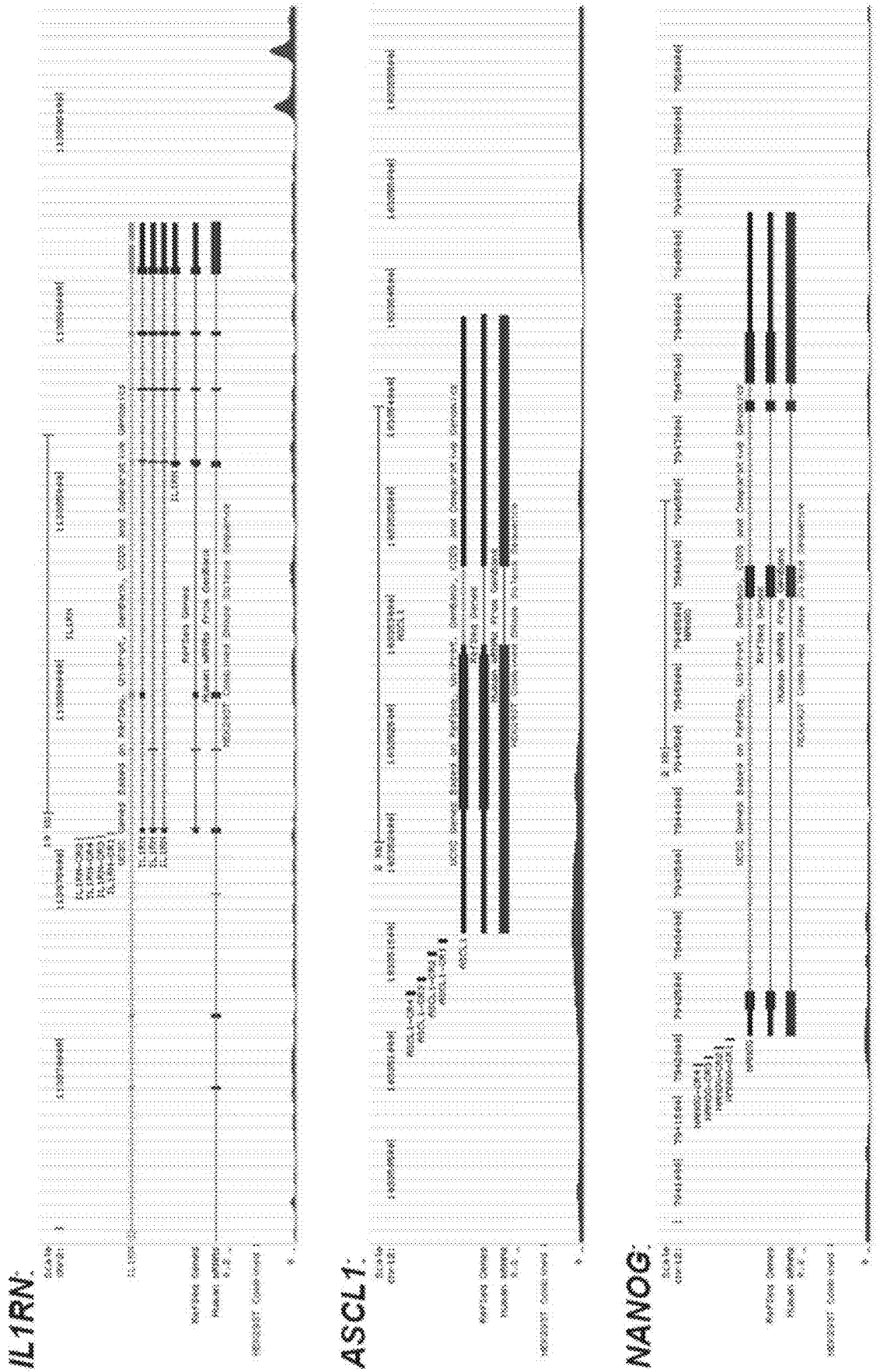


Fig. 4

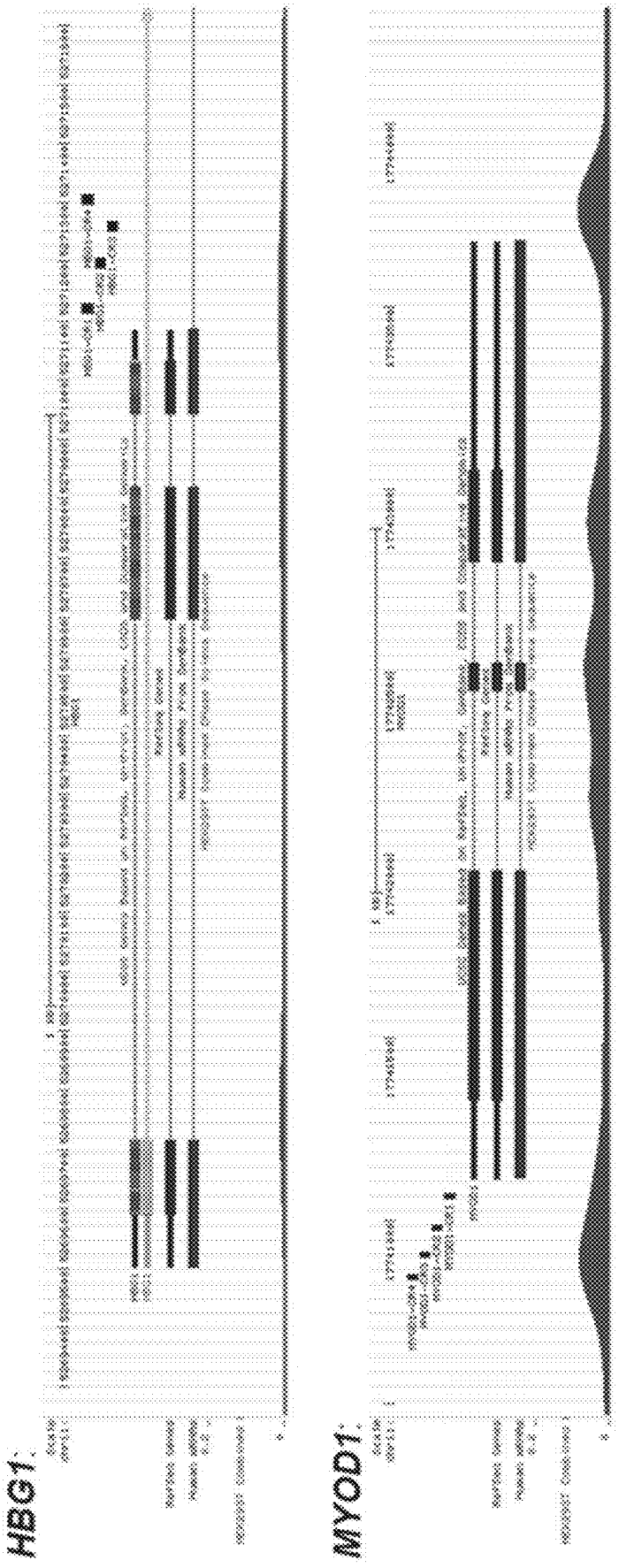


Fig. 4 (Continued)

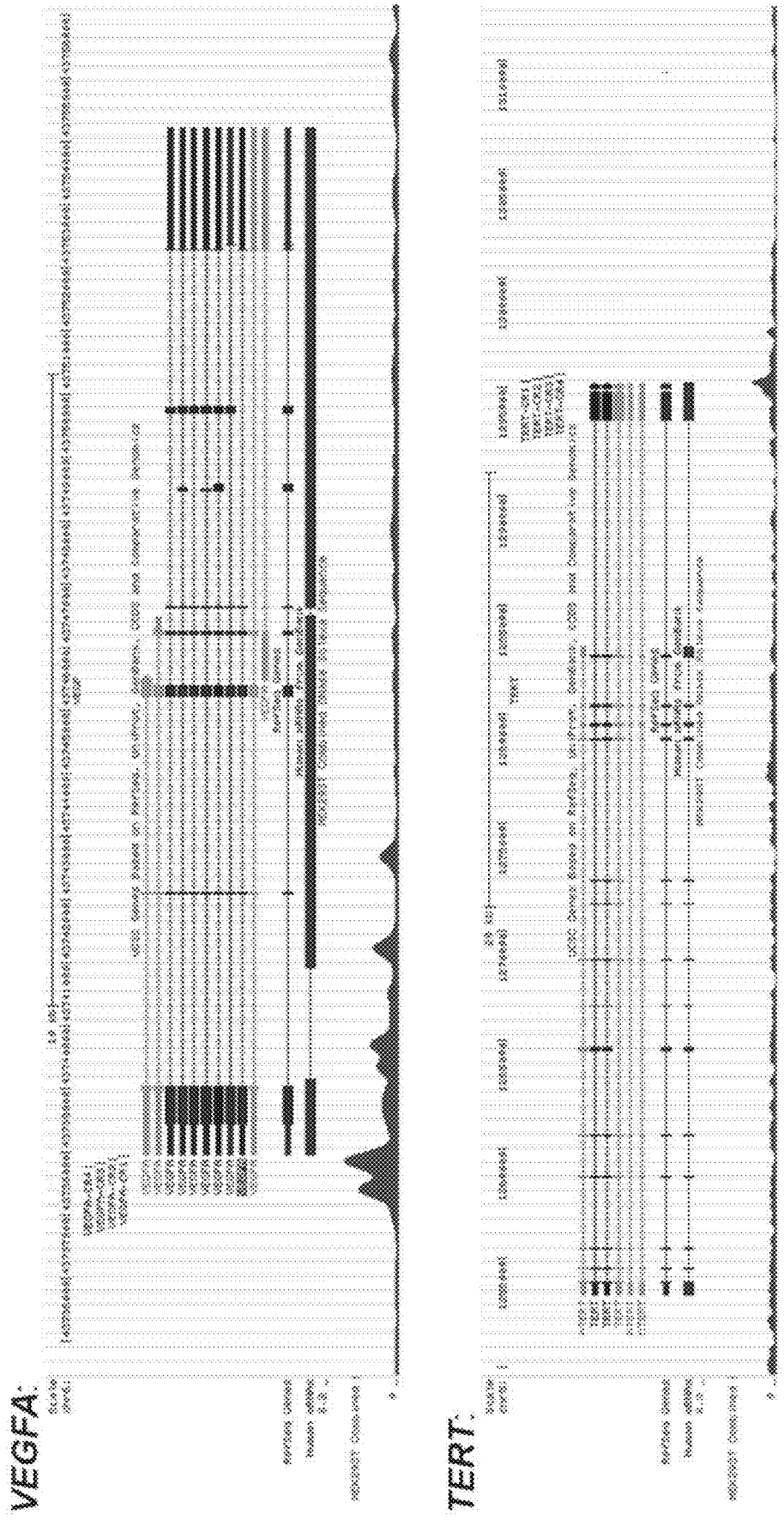


Fig. 4 (Continued)

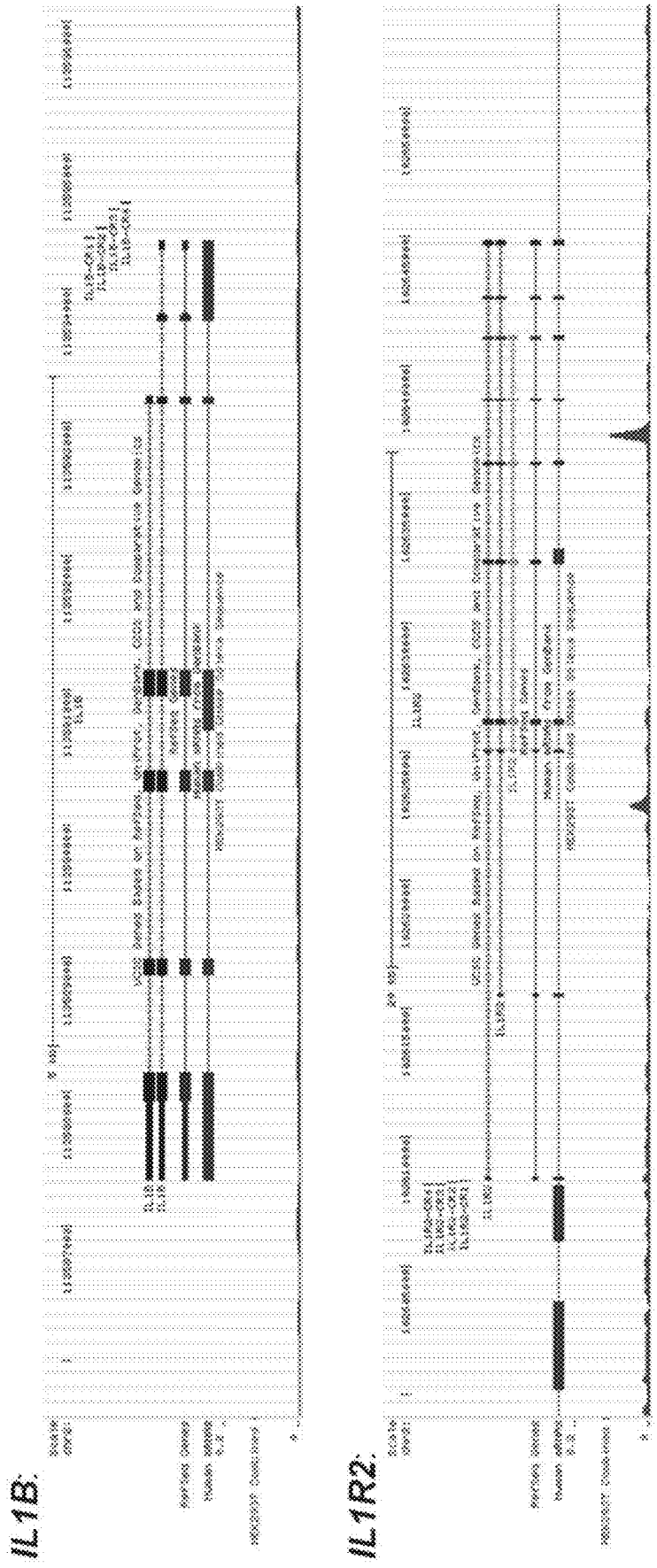


Fig. 4 (Continued)

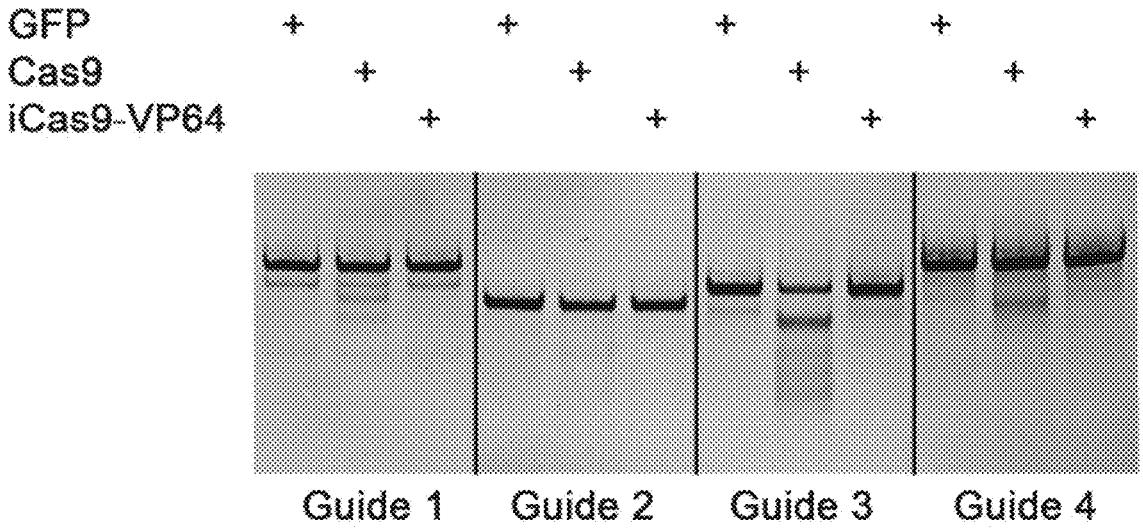

GFP
Cas9
iCas9-VP64
Guide 1
Guide 2
Guide 3
Guide 4

Fig. 5

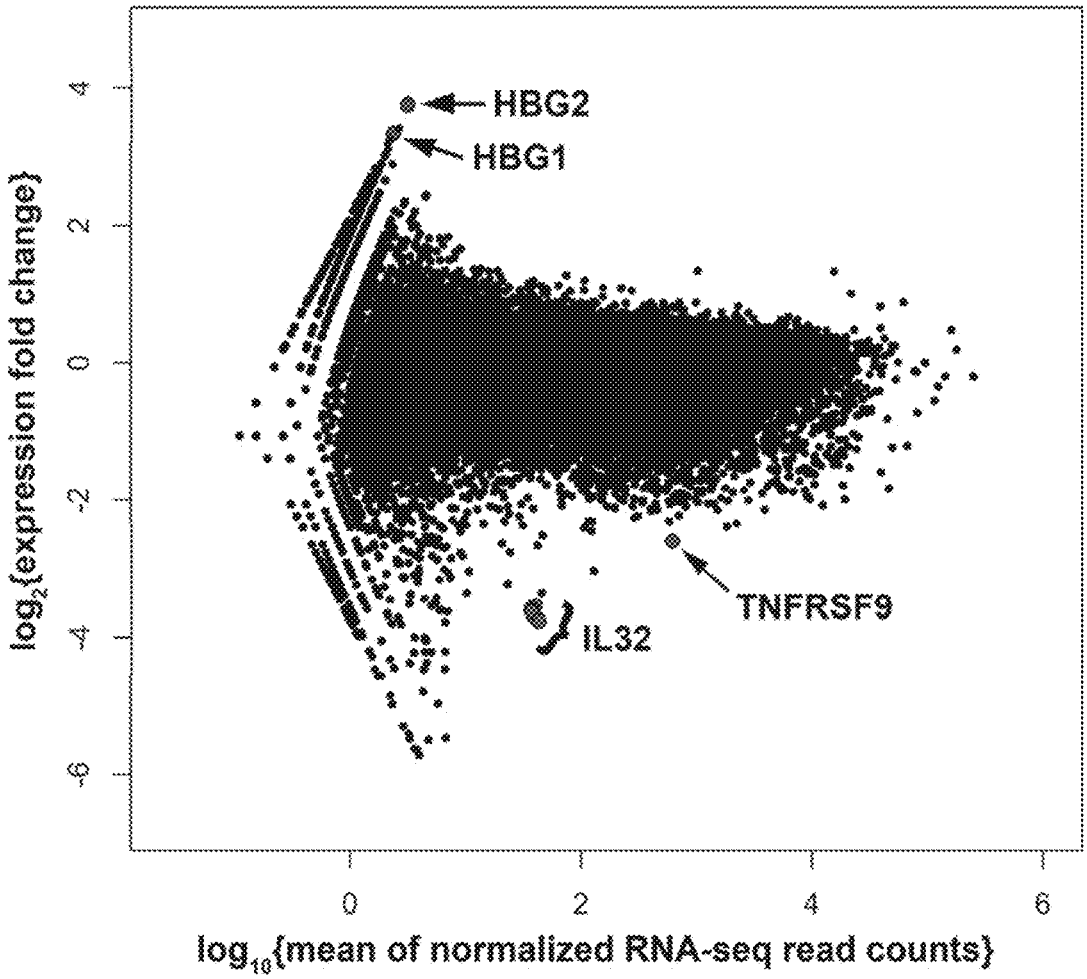

HBG2
HBG1
TNFRSF9
IL32
log2{expression fold change}
log10{mean of normalized RNA-seq read counts}
4
2
0
-2
-4
-6
0
2
4
6

Fig. 6

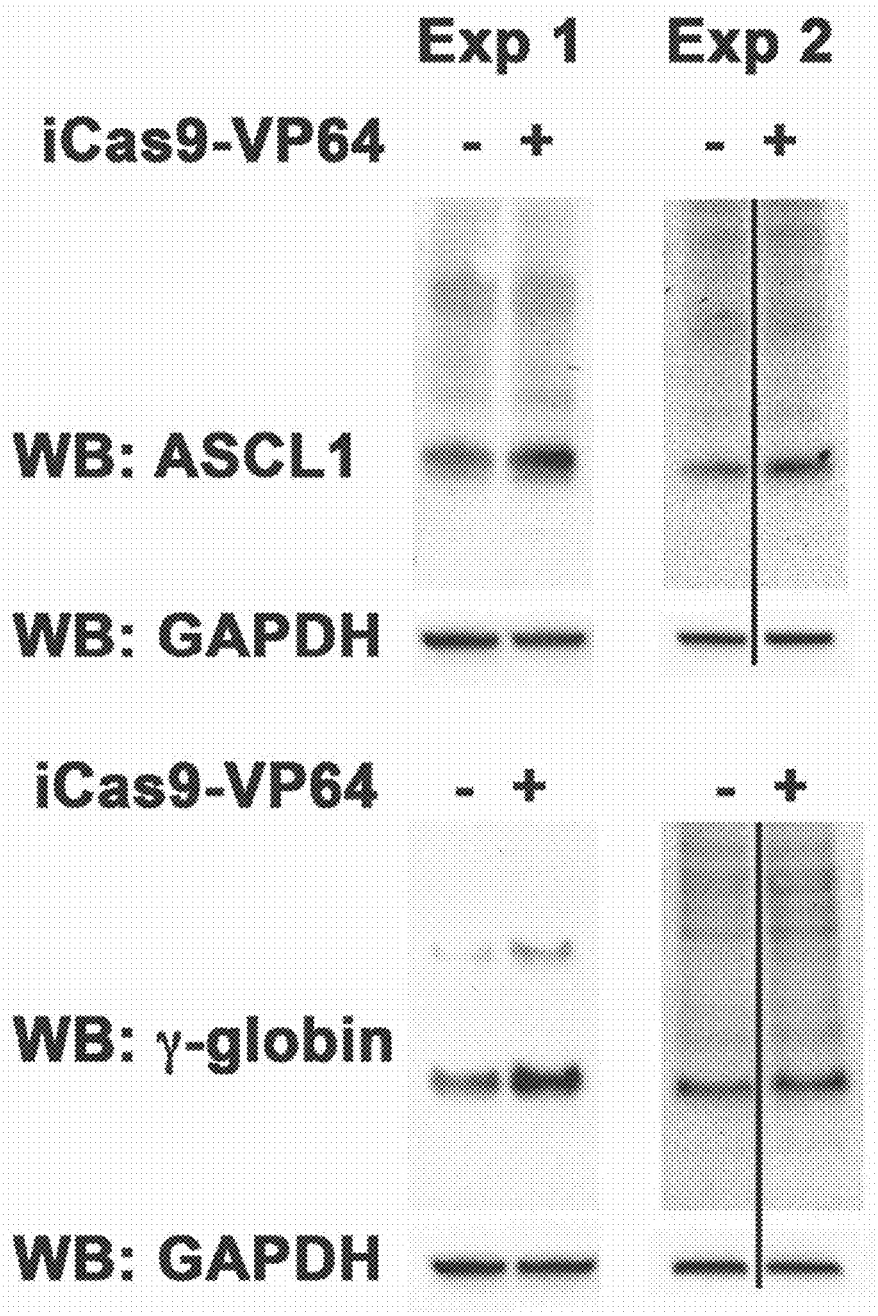

Exp 1
Exp 2
iCas9-VP64
- + - +
WB: ASCL1
WB: GAPDH
iCas9-VP64
- + - +
WB: γ-globin
WB: GAPDH

Fig. 7

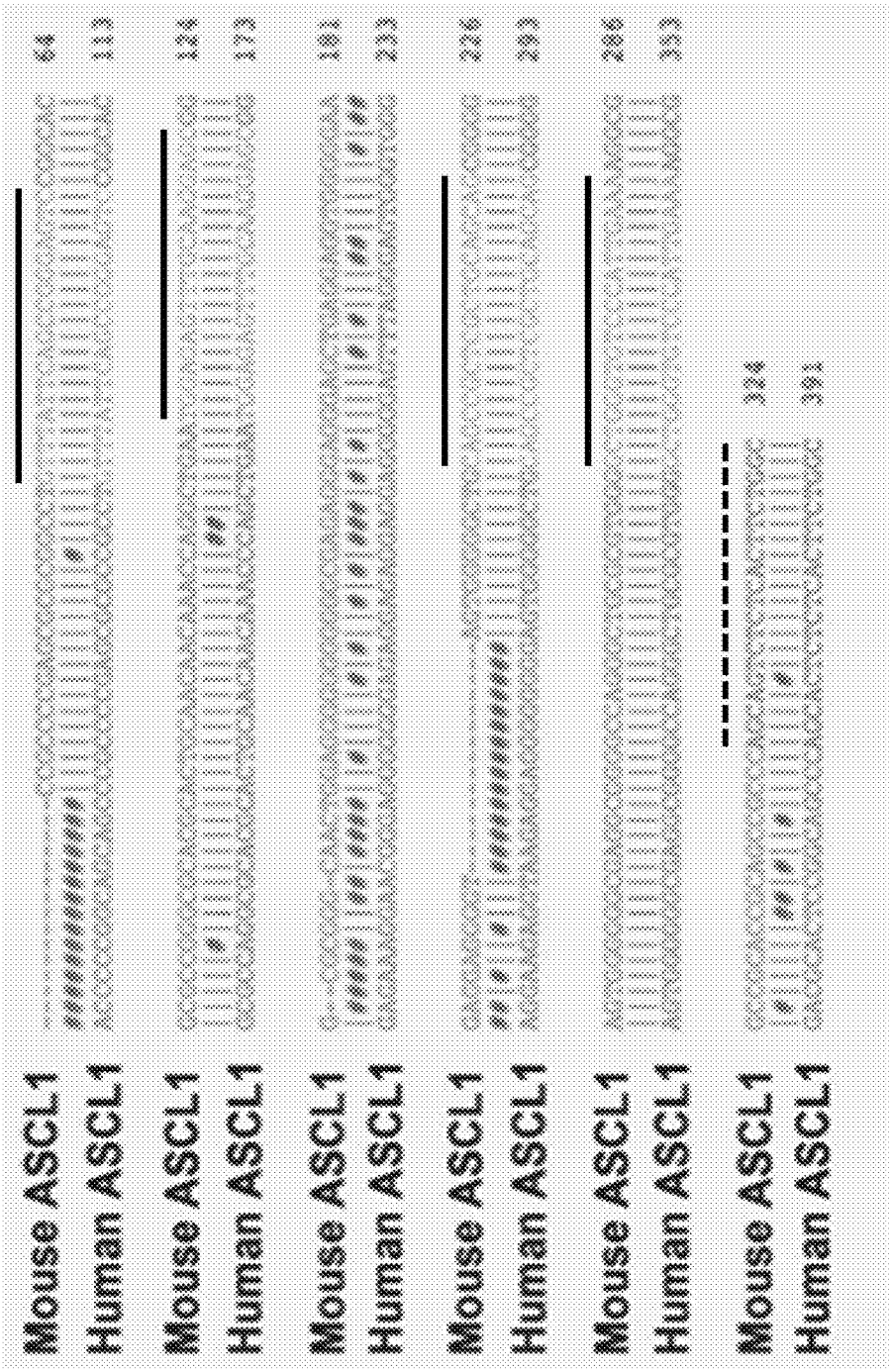

Mouse ASCL1
Human ASCL1
Mouse ASCL1
Human ASCL1
Mouse ASCL1
Human ASCL1
Mouse ASCL1
Human ASCL1
Mouse ASCL1
Human ASCL1
Mouse ASCL1
Human ASCL1

Fig. 8A

Fig. 8B

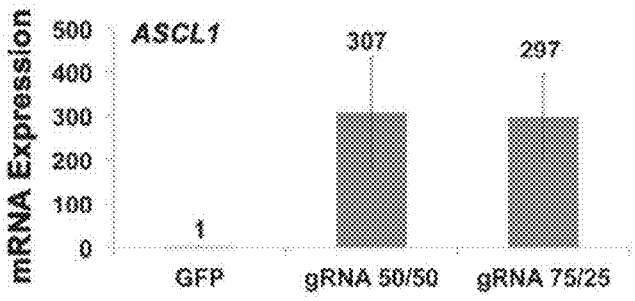


Fig. 8C

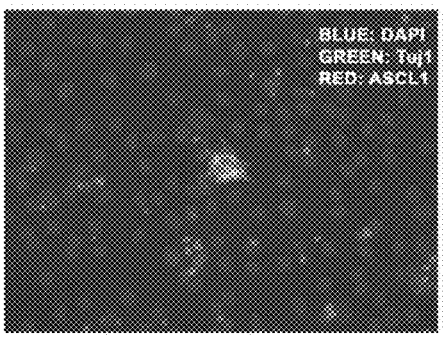


Fig. 8D

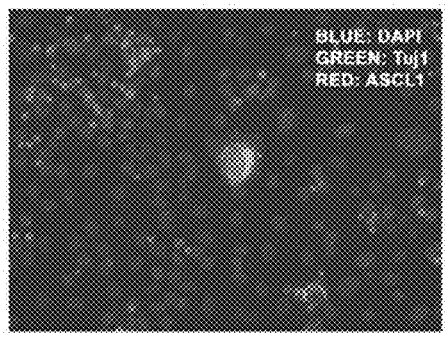
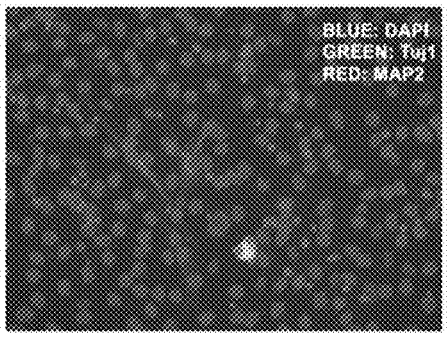


Fig. 8E



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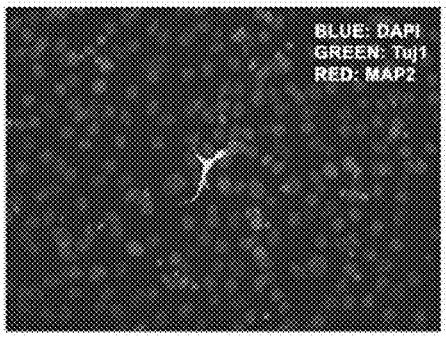


Fig. 8G

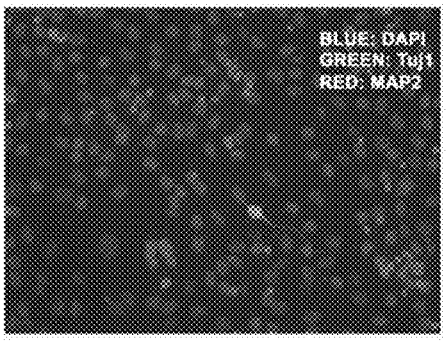
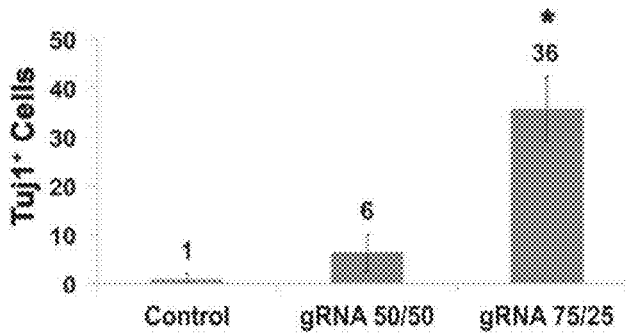


Fig. 8H



M*DYKDHDGDYKDHDIDYKDDDDK*MA**PKKKRKV**GRGMDKKYSIGLAIGTNSVGWAVITD
EYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQ
EIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVD
STDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASG
VDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQL
SKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEH
HQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEE
LLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPY
YVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVL
PKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDY
FKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDRE
MIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFAN
RNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKV
MGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNE
KLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSD
NVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQIT
KHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAY
LNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTE
ITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESI
LPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIME
RSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSK
YVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLS
AYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITG
LYETRIDLSQLGGDPIAGSKAS**PKKKRKV**GRA***DALDDFDLDMLGSDALDDFDLDMLGS***
DALDDFDLDMLGSDALDDFDLDMLINY*PYDVPDYA*S (SEQ ID NO: 1)

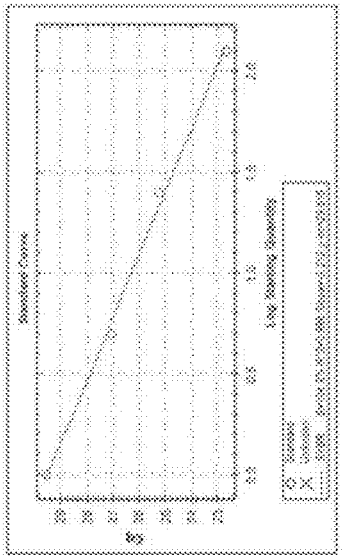
FLAG epitope tag = italicized
Nuclear localization sequence = bold
Streptococcus pyogenes Cas9 = underlined
VP64 (4x minimal VP16 domain) = italicized and bold
HA epitope tag = italicized and underlined

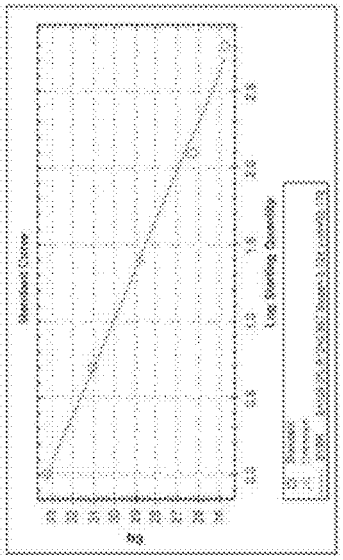
Fig. 9A

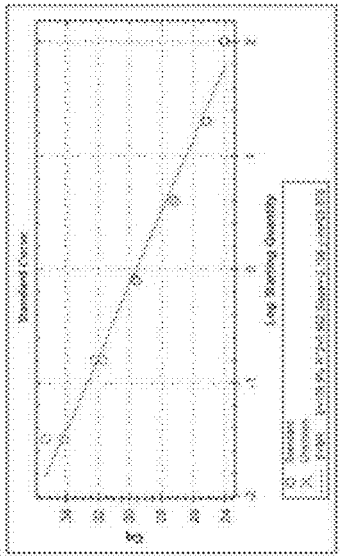
GAGGGCCTATTTCCCATGATTCCTTCATATTTGCATATACGATACAAGGCTG
TTAGAGAGATAATTGGAATTAATTTGACTGTAAACACAAAGATATTAGTACA
AAATACGTGACGTAGAAAGTAATAATTTCTTGGGTAGTTTGCAGTTTTAAAA
TTATGTTTTAAAATGGACTATCATATGCTTATCGTAACTTGAAAGTATTTCGA
TTTCTTGGCTTTATATATCTTGTGGAAAGGACGAAACACCG*GGTCTTC*GA*GA*
*AGAC*CT*GTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGCTAGTCCGTTATC*
*AACTTGAAAAAGTGGCACCGAGTCGGTGC***TTTTTTT** (SEQ ID NO:2)

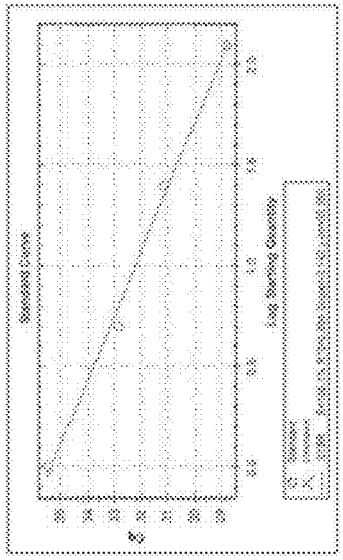
U6 promoter = bold
+1 transcription start site = underlined
BbsI restriction sites to clone in guide RNA = italicized and underlined
Chimeric guide RNA sequence = italicized
Poly-T terminator sequence = bold and underlined

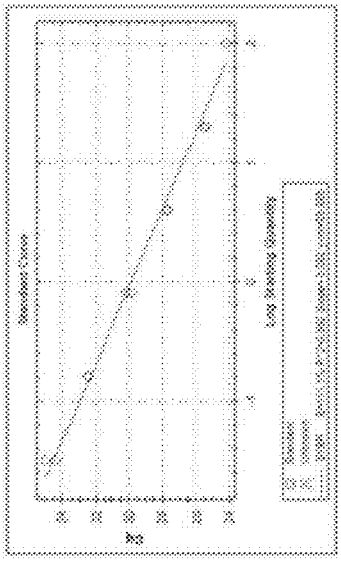
Fig. 9B











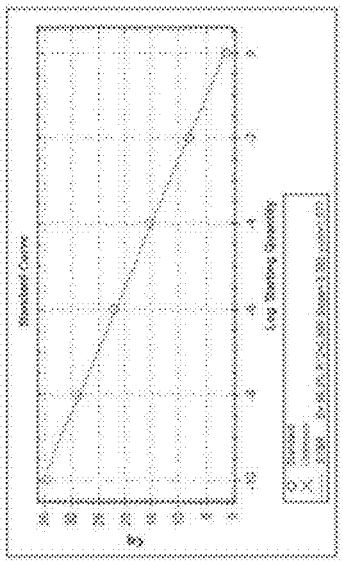
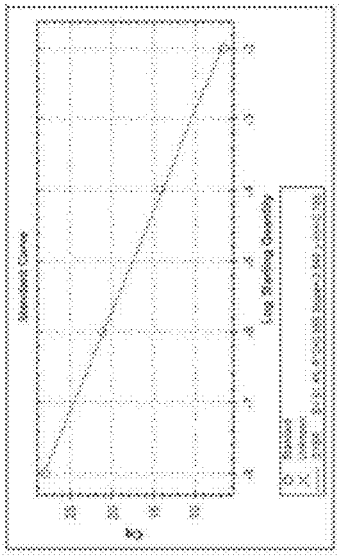
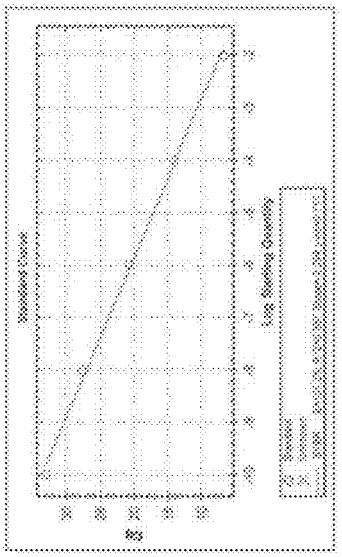


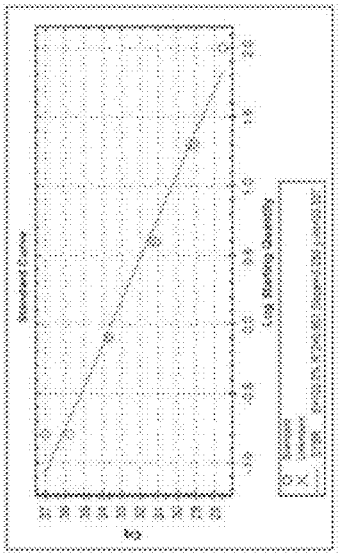
Fig. 10



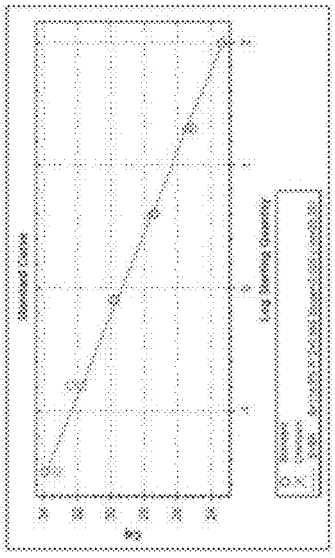
MYOD1



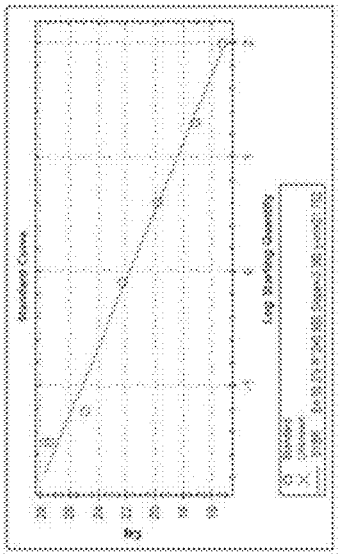
HGB1



mASCL1



IL1RN



GAPDH

Fig. 10 (Continued)

Fig. 11A

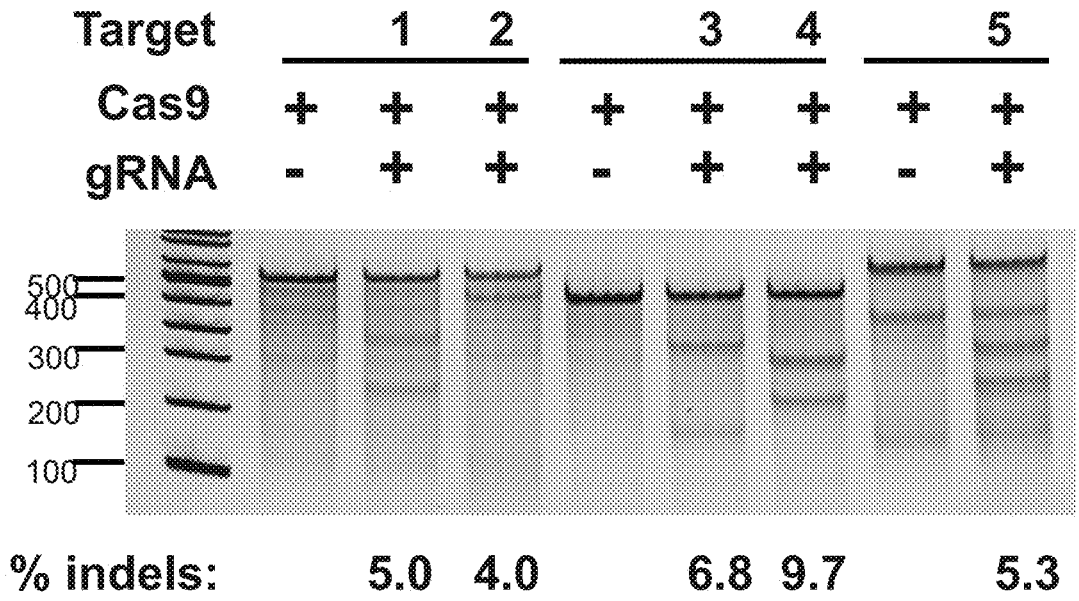


Fig. 11B

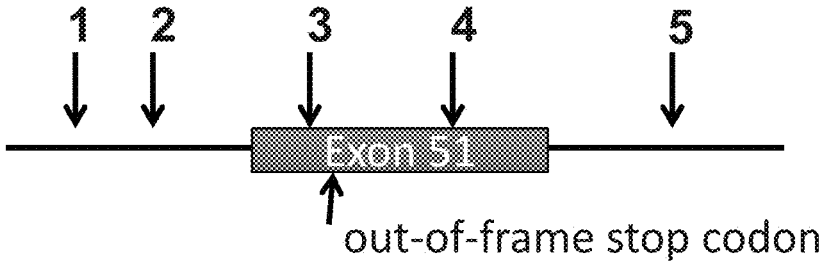


Fig. 11C

Expected cleavage sizes
CR1=352/237
CR2=461/128
CR3=301/150
CR4=272/179
CR5=280/203

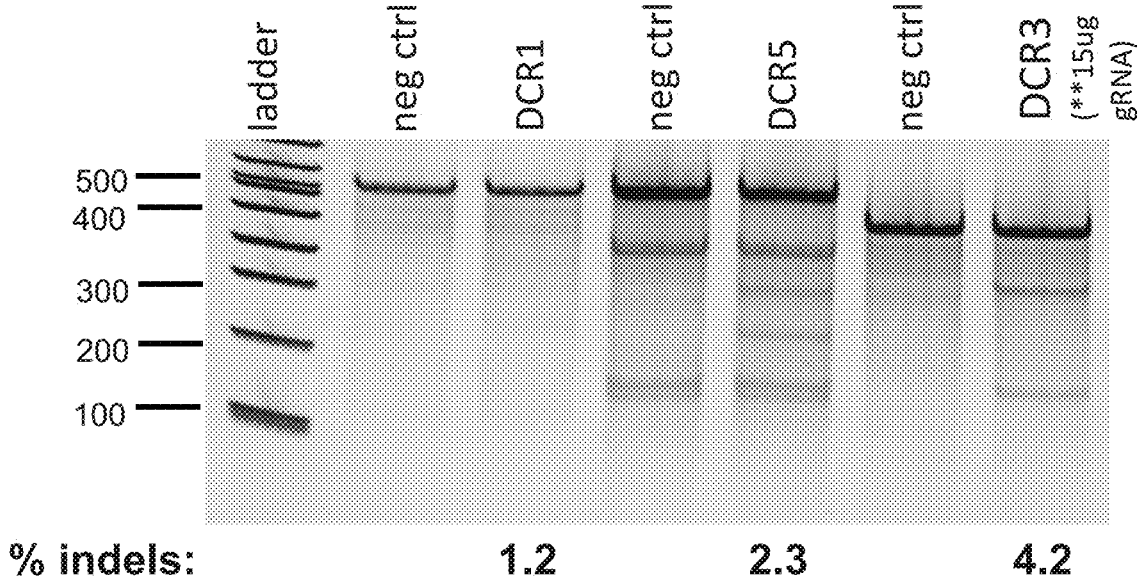

ladder
neg ctrl
DCR1
neg ctrl
DCR5
neg ctrl
DCR3
(**15ug gRNA)
500
400
300
200
100
% indels:
1.2
2.3
4.2

Fig. 12

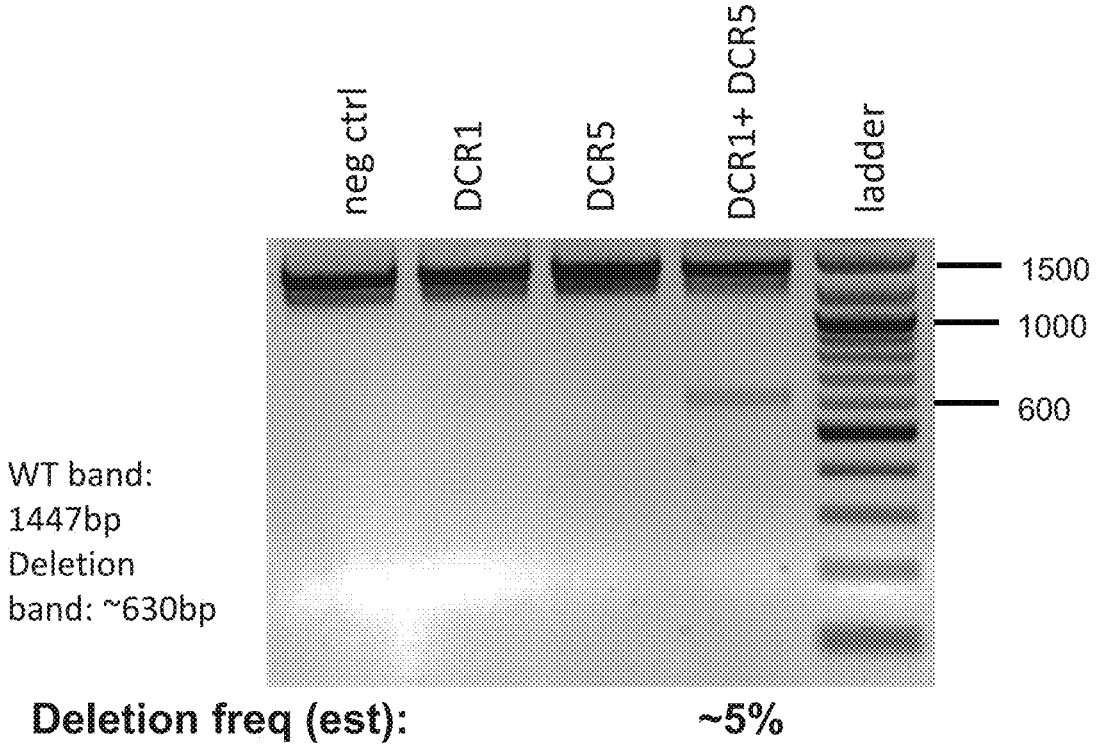

neg ctrl
DCR1
DCR5
DCR1+ DCR5
ladder
1500
1000
600
WT band:
1447bp
Deletion
band: ~630bp
Deletion freq (est):
~5%

Fig. 13

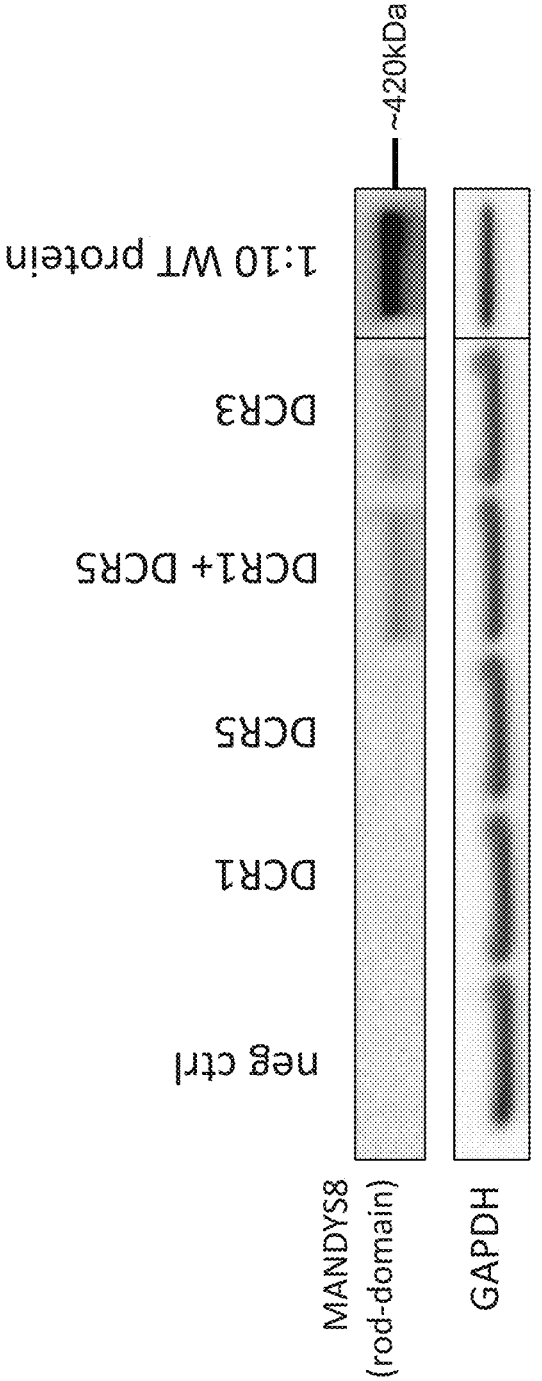

~420kDa
1:10 WT protein
DCR3
DCR1+ DCR5
DCR5
DCR1
neg ctrl
MANDYS8
(rod-domain)
GAPDH

Fig. 14

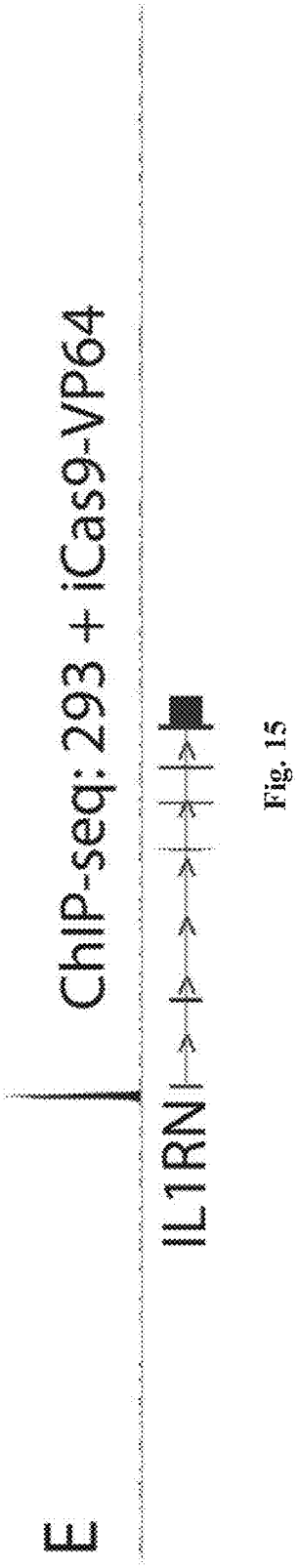

E
ChIP-seq: 293 + iCas9-VP64
IL1RN

Fig. 15

Fig. 16A

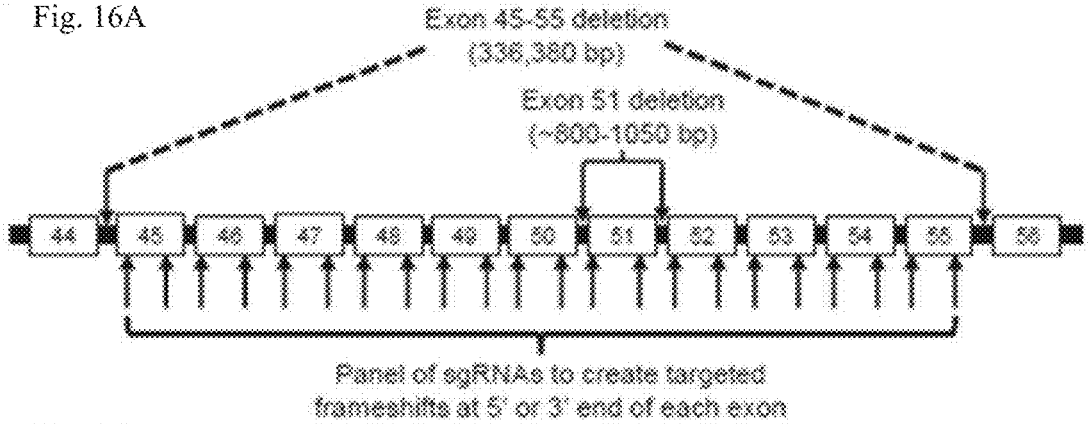


Fig. 16B

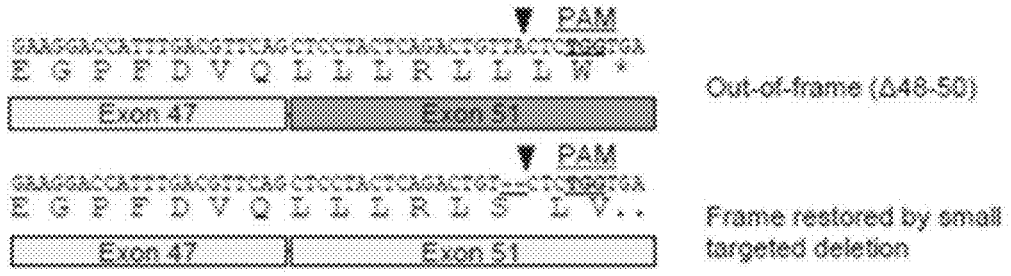


Fig. 16C

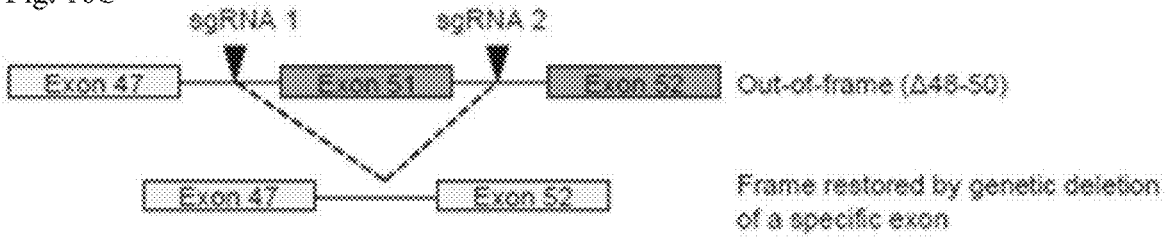
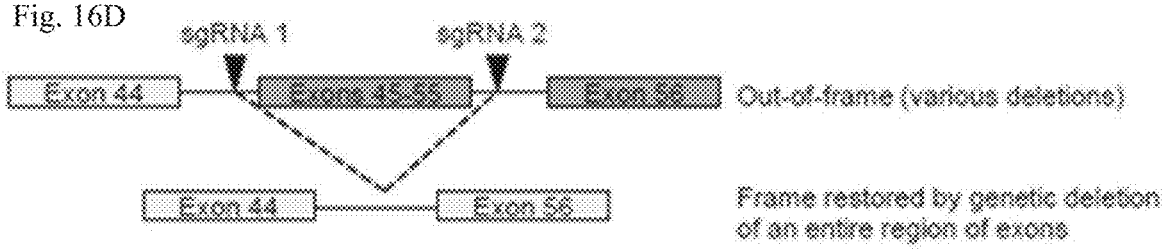


Fig. 16D



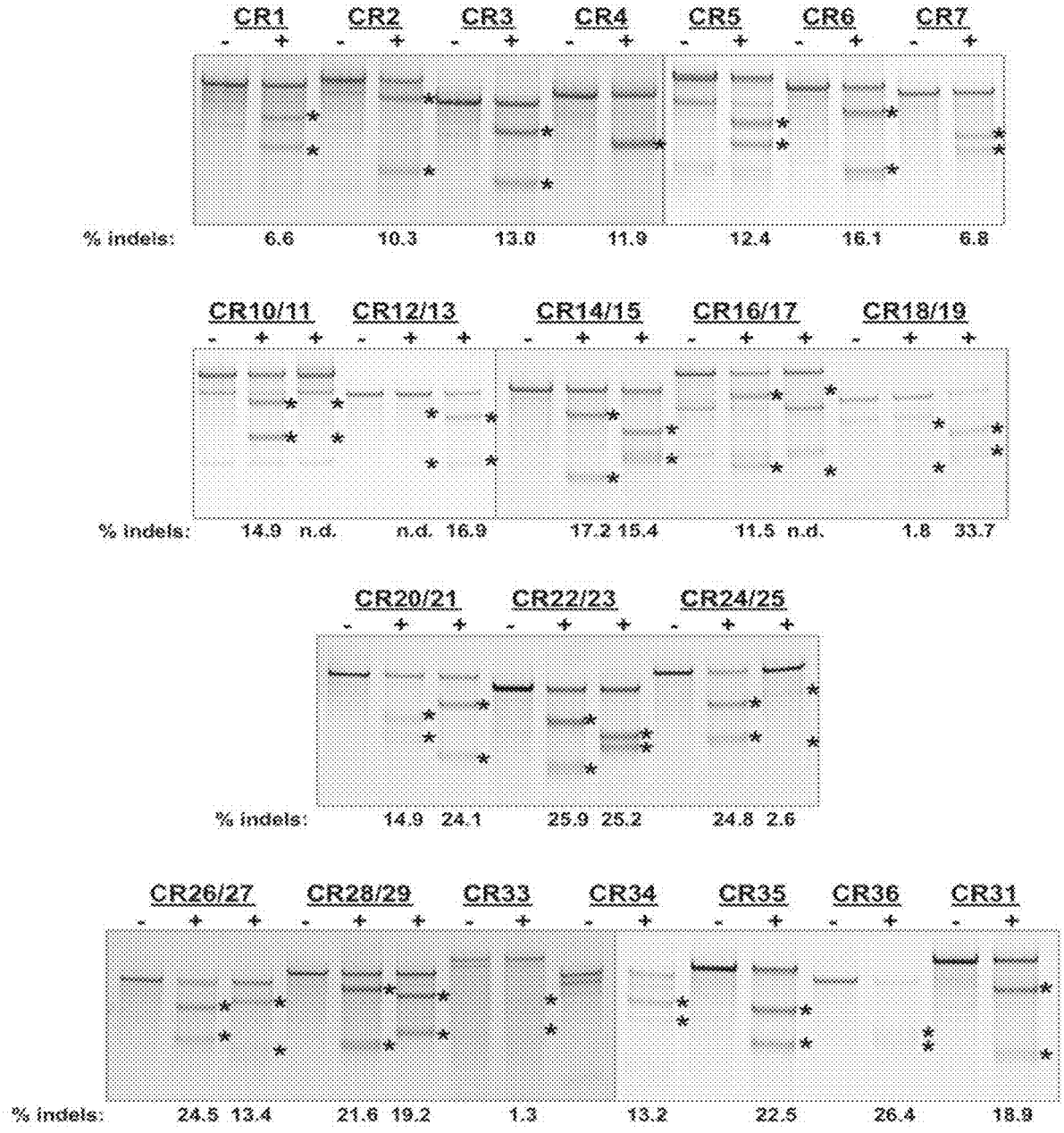

CR1
CR2
CR3
CR4
CR5
CR6
CR7
% indels: 6.6 10.3 13.0 11.9 12.4 16.1 6.8
CR10/11
CR12/13
CR14/15
CR16/17
CR18/19
% indels: 14.9 n.d. n.d. 16.9 17.2 15.4 11.5 n.d. 1.8 33.7
CR20/21
CR22/23
CR24/25
% indels: 14.9 24.1 25.9 25.2 24.8 2.6
CR26/27
CR28/29
CR33
CR34
CR35
CR36
CR31
% indels: 24.5 13.4 21.6 19.2 1.3 13.2 22.5 26.4 18.8

Fig. 17

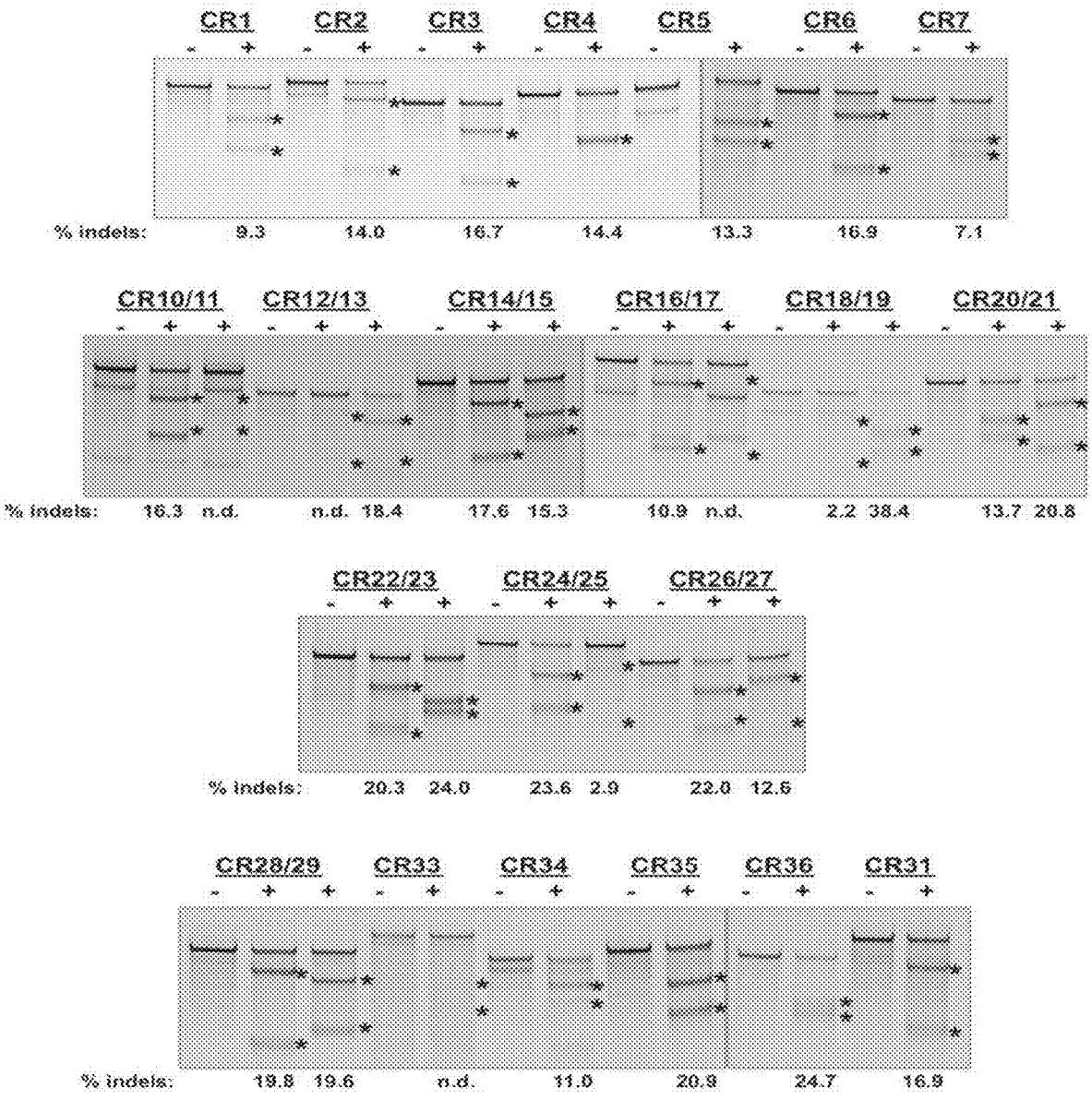

CR1
CR2
CR3
CR4
CR5
CR6
CR7
% indels: 9.3 14.0 16.7 14.4 13.3 16.9 7.1
CR10/11
CR12/13
CR14/15
CR16/17
CR18/19
CR20/21
% indels: 16.3 n.d. n.d. 18.4 17.6 15.3 10.9 n.d. 2.2 38.4 13.7 20.8
CR22/23
CR24/25
CR26/27
% indels: 20.3 24.0 23.6 2.9 22.0 12.6
CR28/29
CR33
CR34
CR35
CR36
CR31
% indels: 19.8 19.6 n.d. 11.0 20.9 24.7 16.9

Fig. 18

Fig. 19A

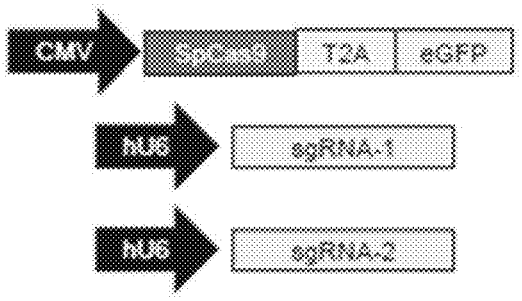


Fig. 19B

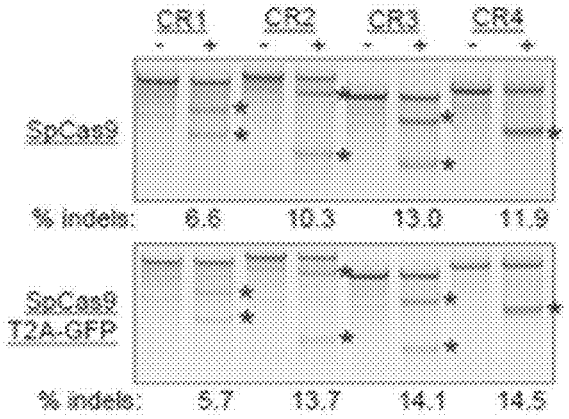


Fig. 19C

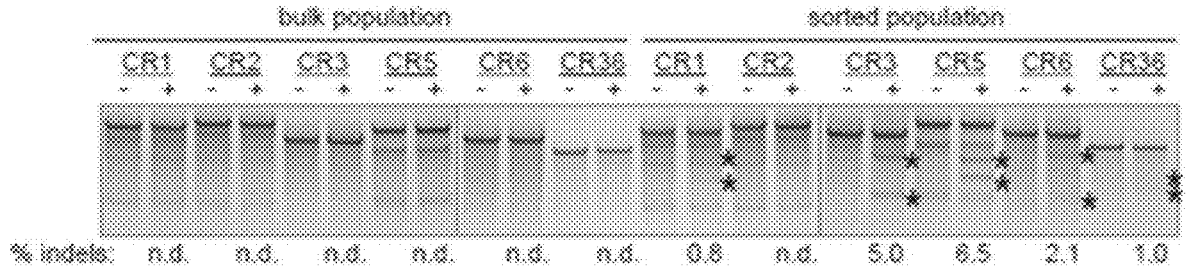


Fig. 19D

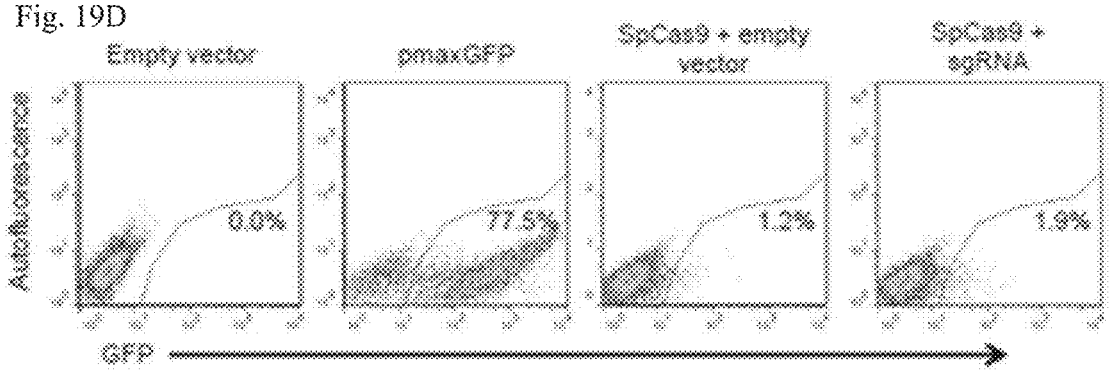


Fig. 20A

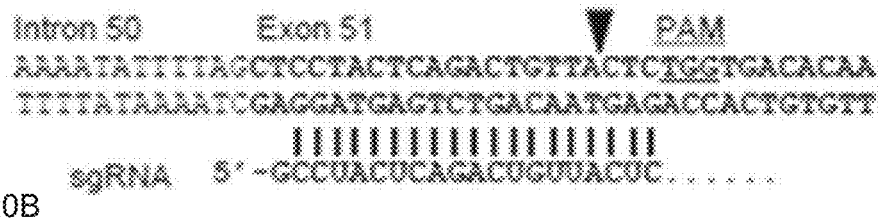


Fig. 20B

Fig. 20C

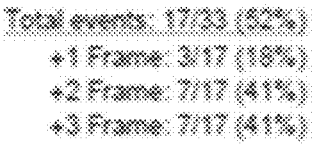


Fig. 20D

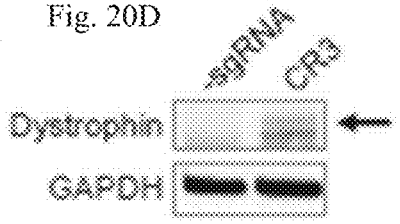


Fig. 21A

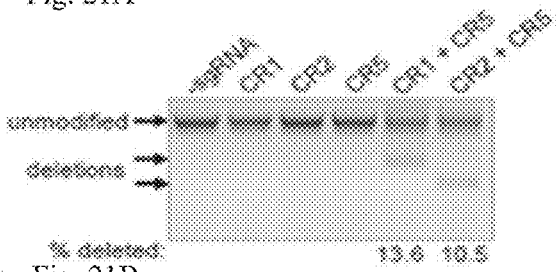


Fig. 21B

Fig. 21C

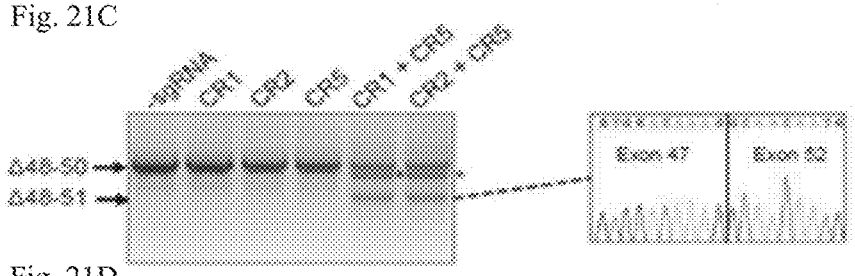


Fig. 21D

Fig. 22A

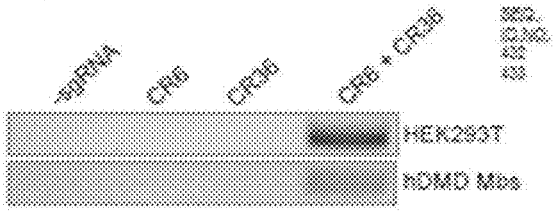


Fig. 22B

Fig. 22C

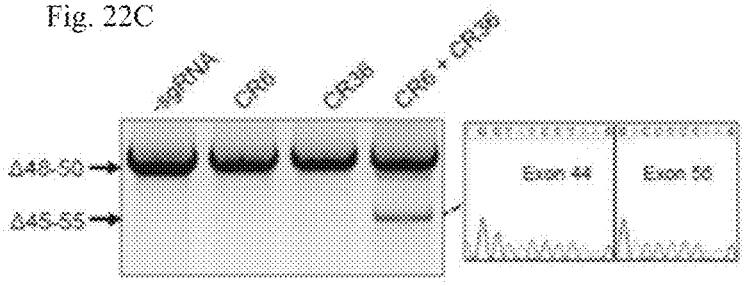
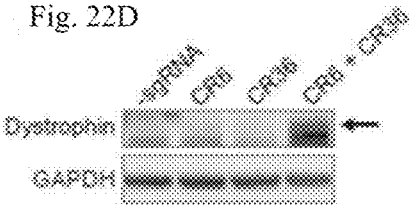


Fig. 22D



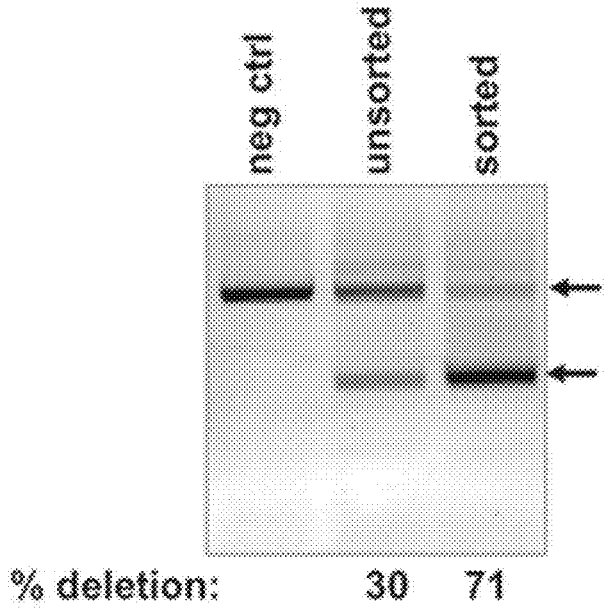


Fig. 23

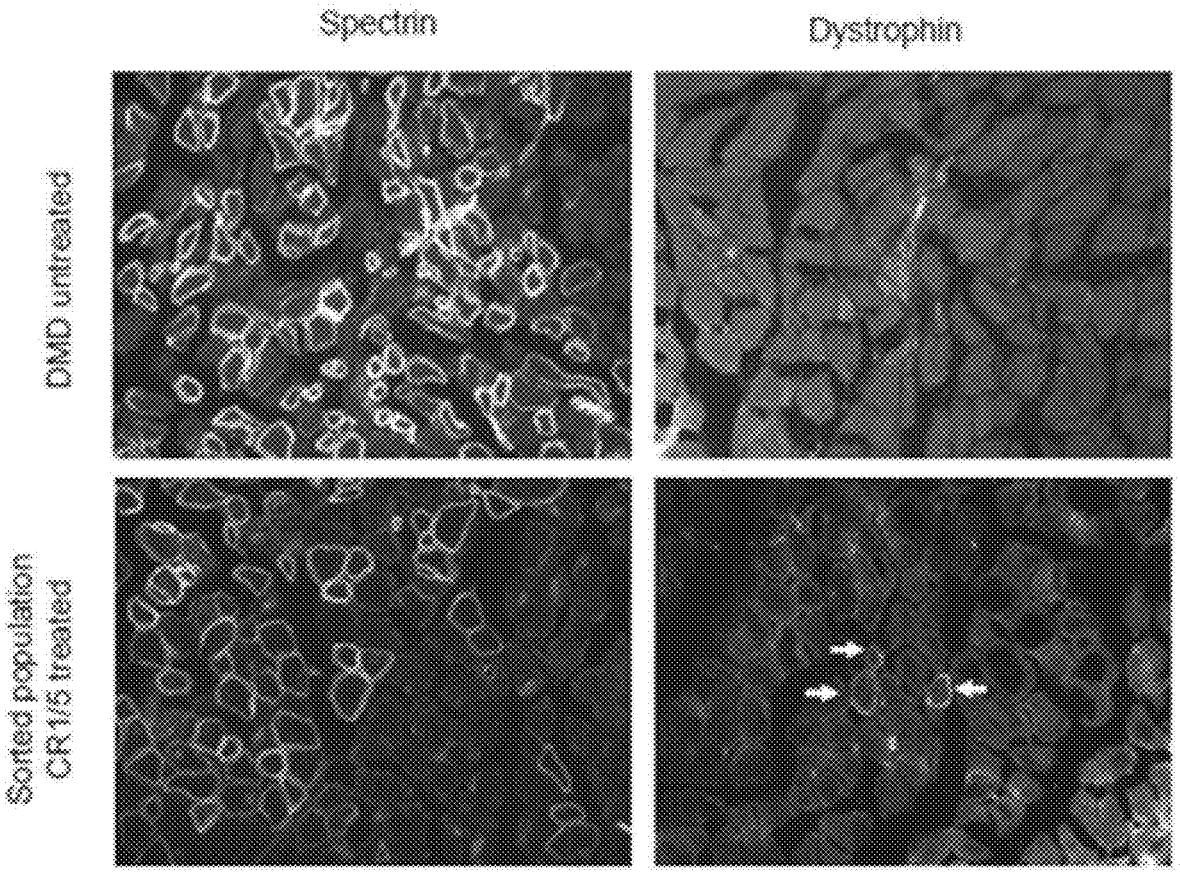

Spectrin
Dystrophin
DMD untreated
Sorted population
CR1/6 treated

Fig. 24

Fig. 25A

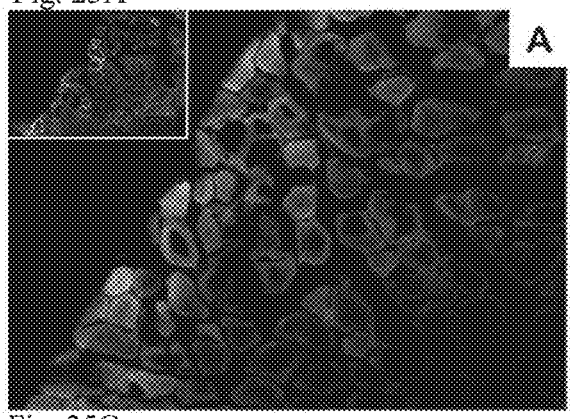


Fig. 25B

Fig. 25C

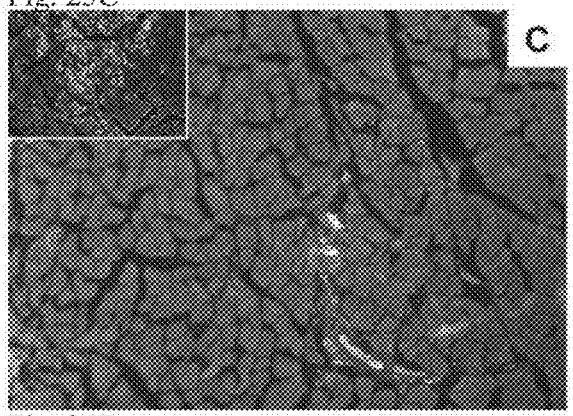


Fig. 25D

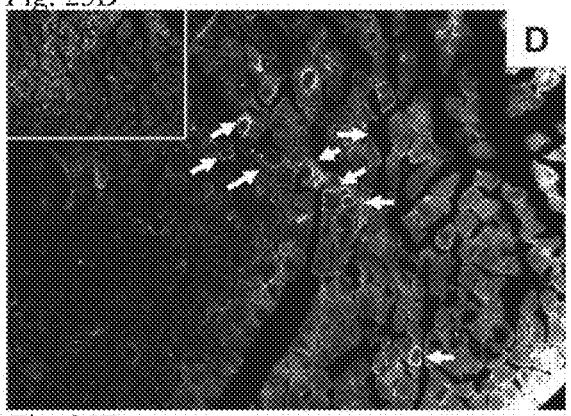


Fig. 25E

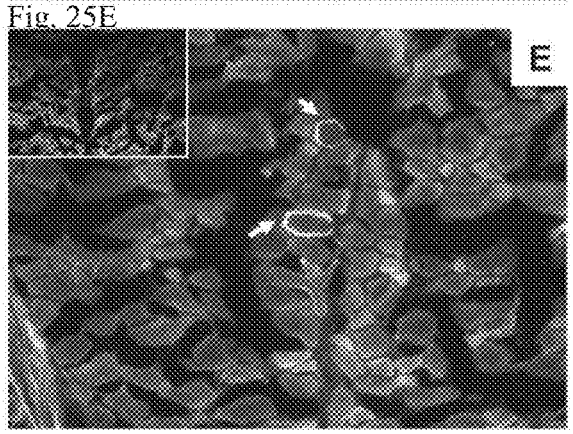


Fig. 25F

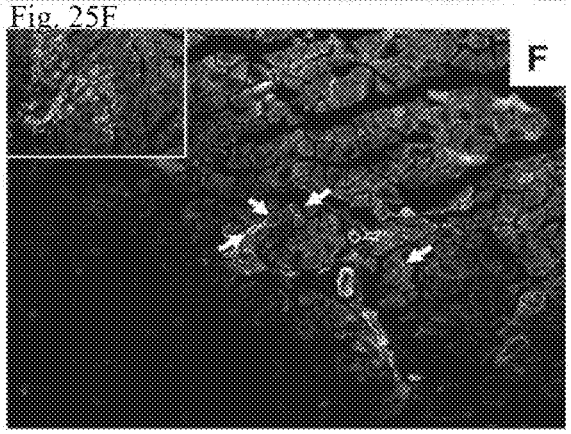
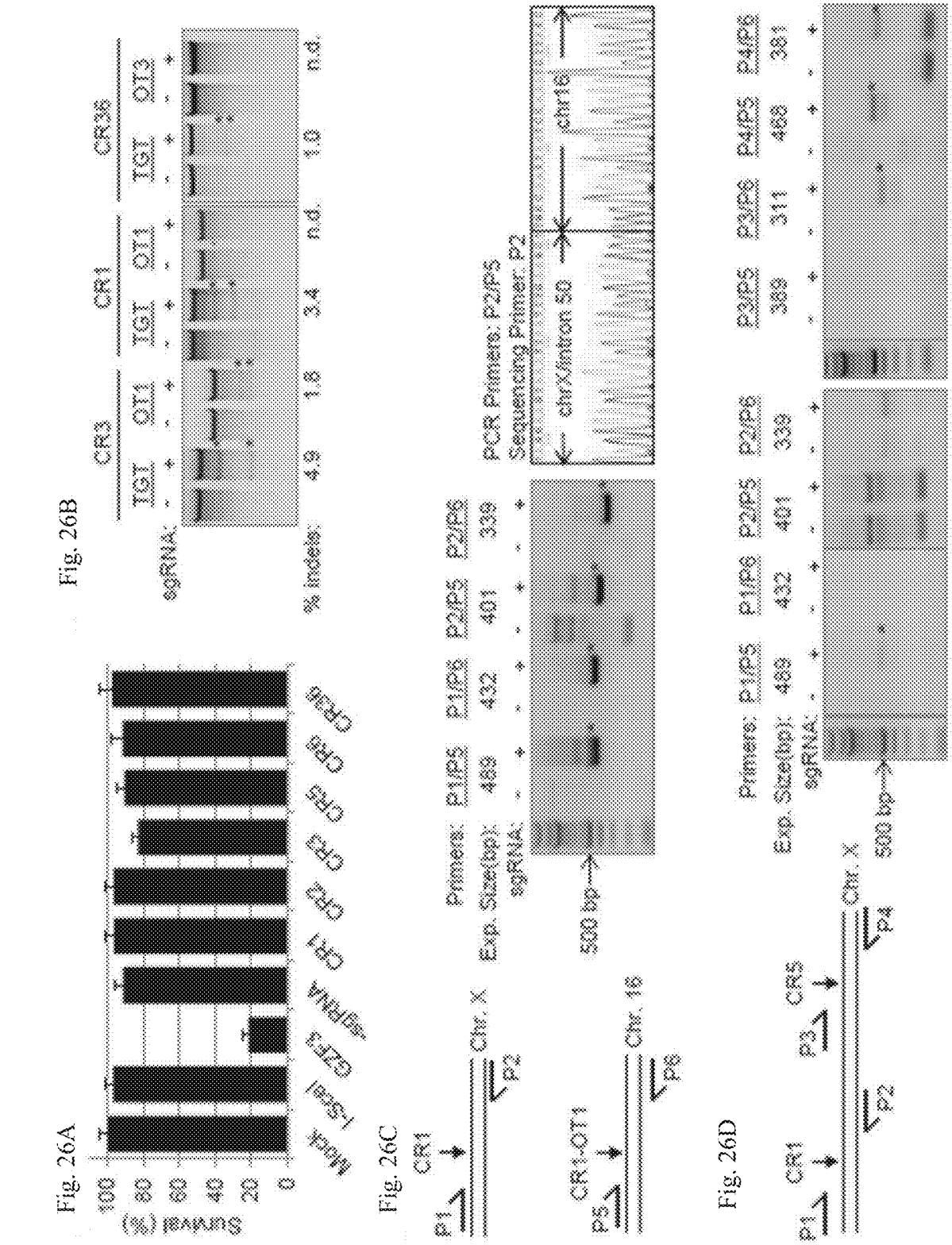


Fig. 26A

Fig. 26B

Fig. 26C

Fig. 26D



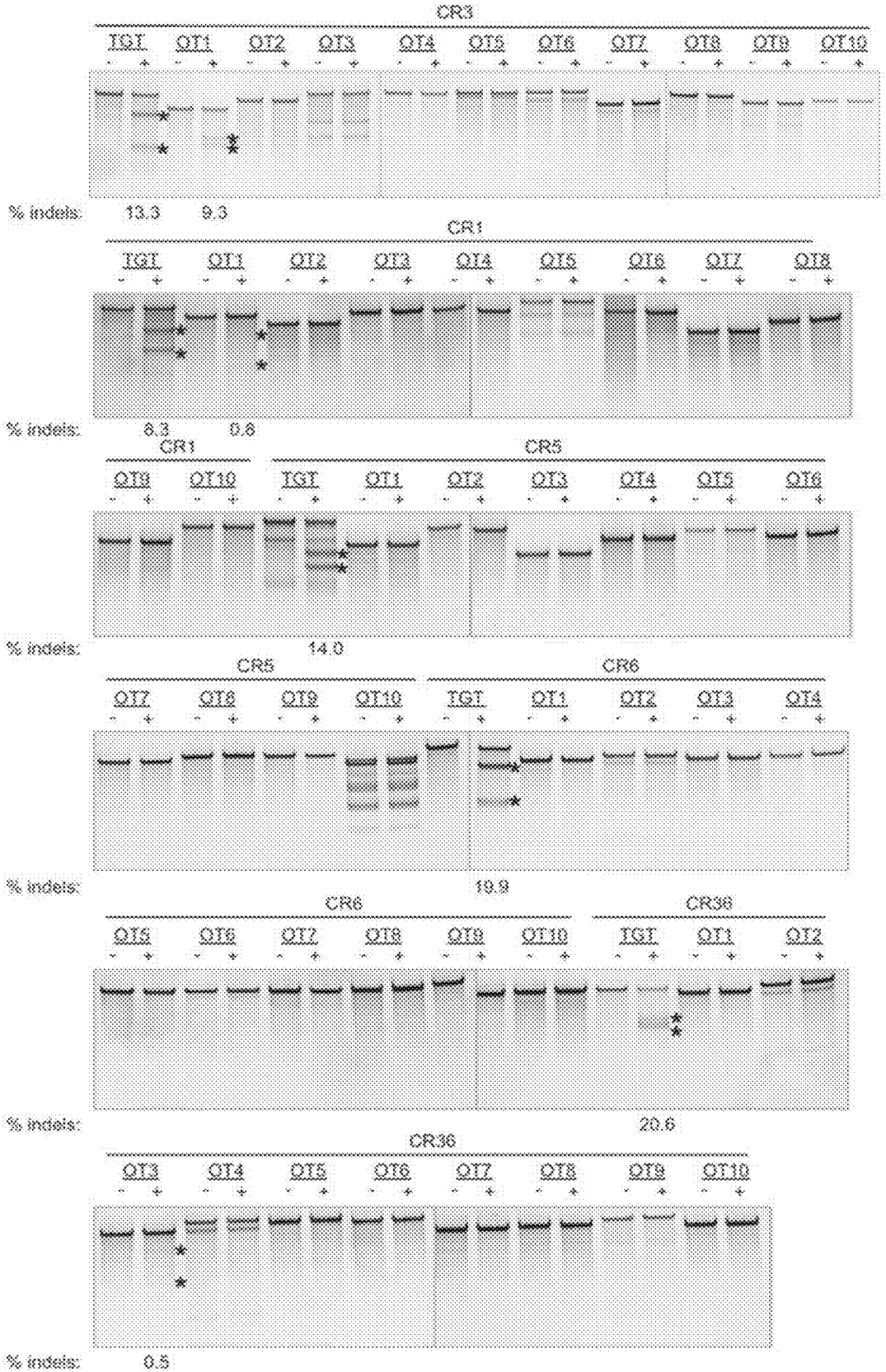

CR3
TGT OT1 OT2 OT3 OT4 OT5 OT6 OT7 OT8 OT9 OT10
% indels: 13.3 9.3
CR1
TGT OT1 OT2 OT3 OT4 OT5 OT6 OT7 OT8
% indels: 8.3 0.8
CR1
CR5
OT9 OT10 TGT OT1 OT2 OT3 OT4 OT5 OT6
% indels: 14.0
CR5
CR6
OT7 OT8 OT9 OT10 TGT OT1 OT2 OT3 OT4
% indels: 19.9
CR6
CR36
OT5 OT6 OT7 OT8 OT9 OT10 TGT OT1 OT2
% indels: 20.6
CR36
OT3 OT4 OT5 OT6 OT7 OT8 OT9 OT10
% indels: 0.5

Fig. 27

Fig. 28A

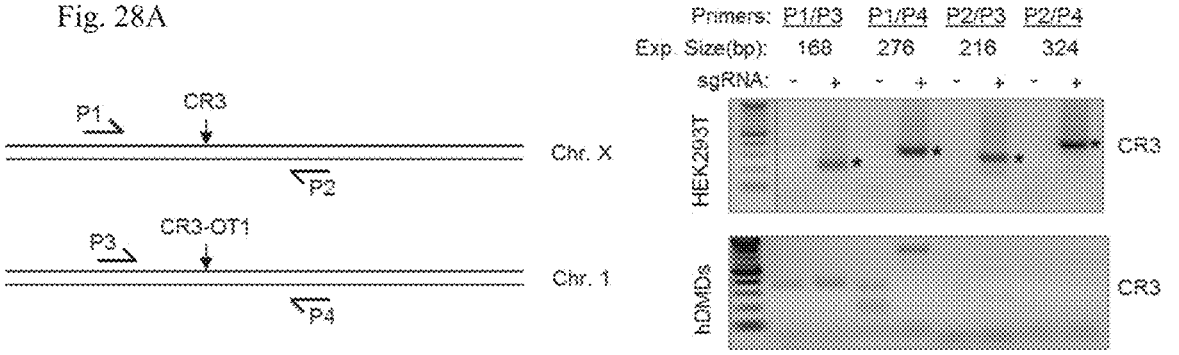


Fig. 28B

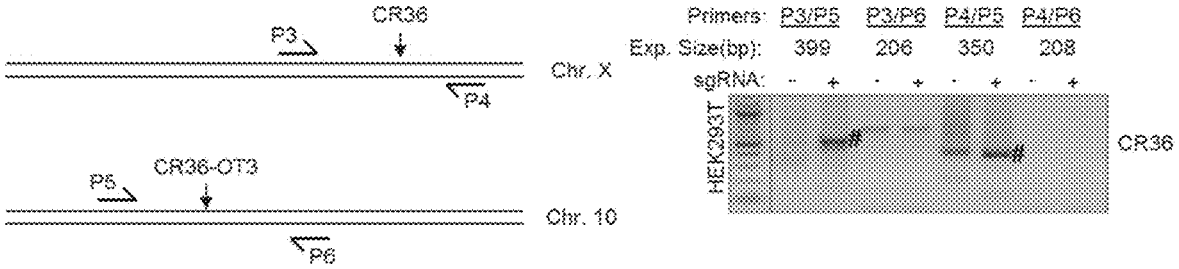
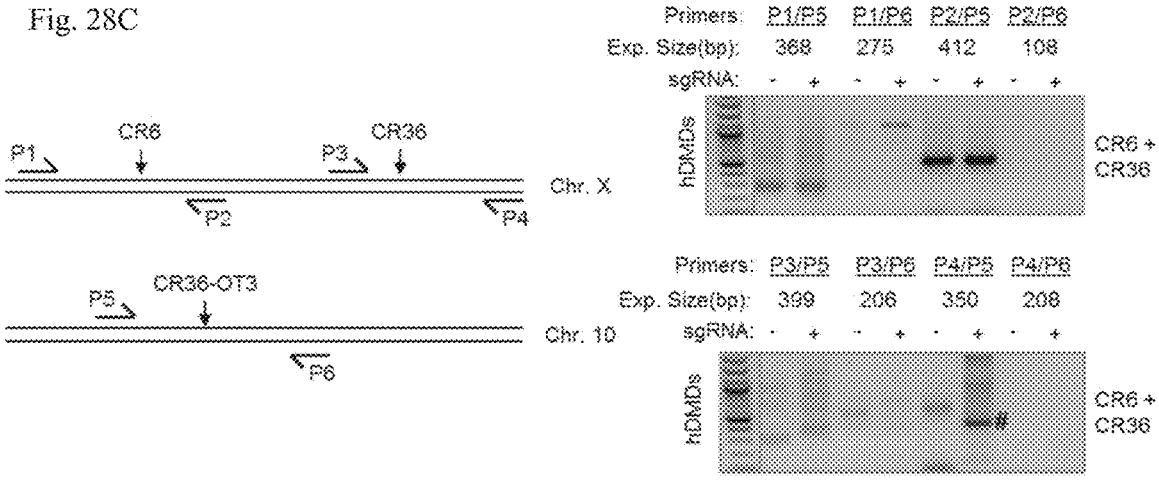


Fig. 28C



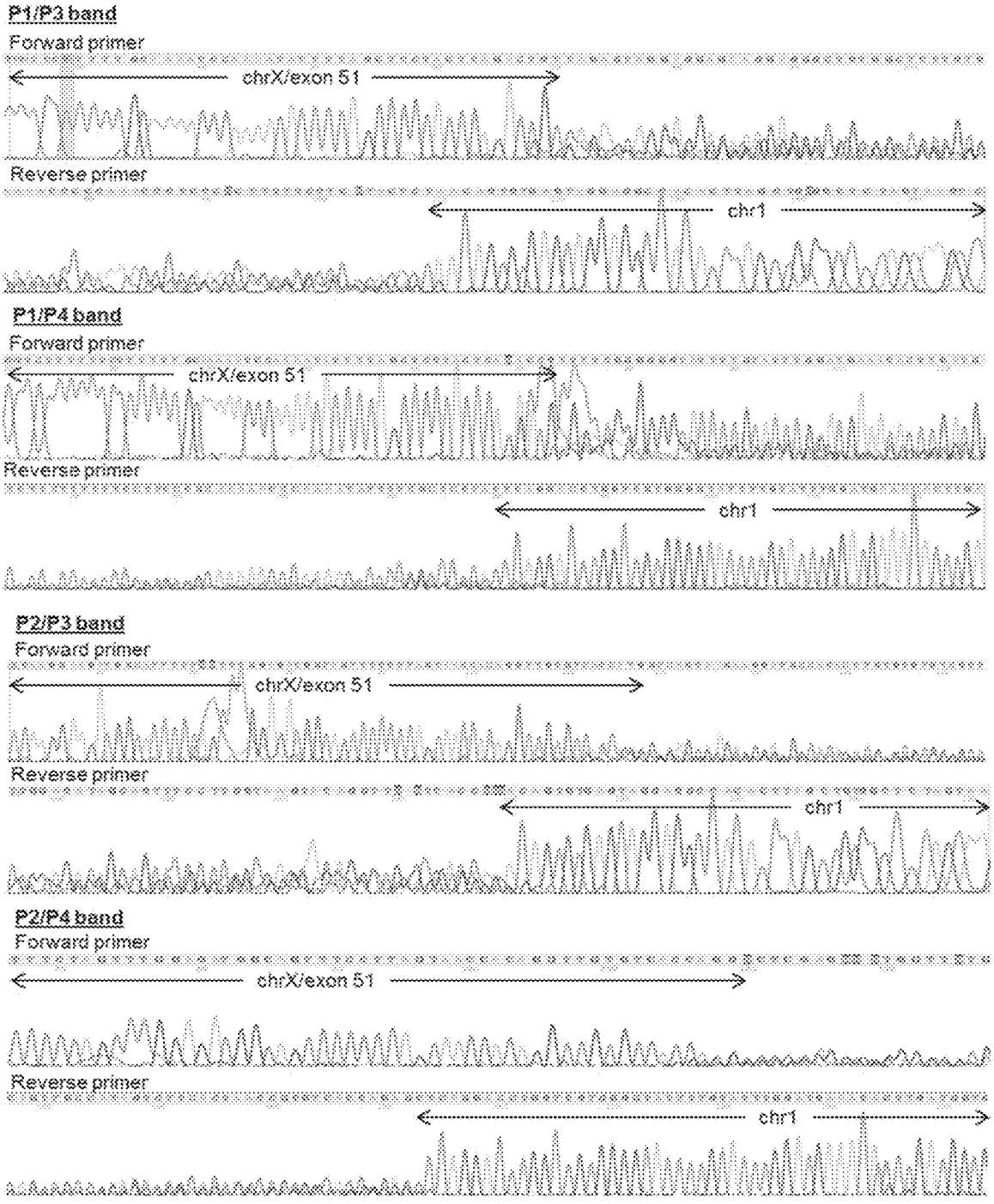

P1/P3 band
Forward primer
chrX/exon 51
Reverse primer
chr1
P1/P4 band
Forward primer
chrX/exon 51
Reverse primer
chr1
P2/P3 band
Forward primer
chrX/exon 51
Reverse primer
chr1
P2/P4 band
Forward primer
chrX/exon 51
Reverse primer
chr1

Fig. 29

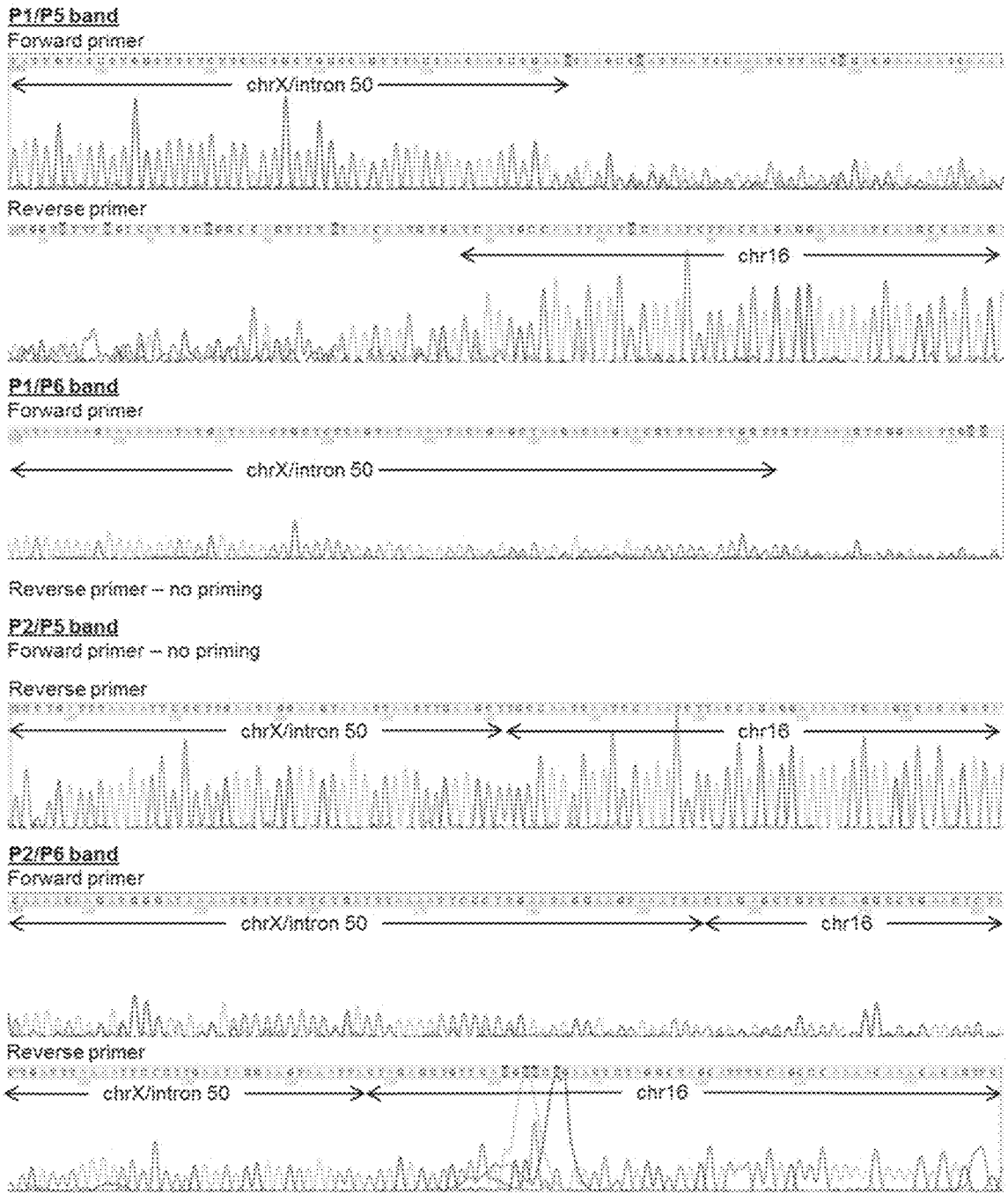

P1/P5 band
Forward primer
chrX/intron 50
Reverse primer
chr16
P1/P6 band
Forward primer
chrX/intron 50
Reverse primer -- no priming
P2/P5 band
Forward primer -- no priming
Reverse primer
chrX/intron 50
chr16
P2/P6 band
Forward primer
chrX/intron 50
chr16
Reverse primer
chrX/intron 50
chr16

Fig. 30

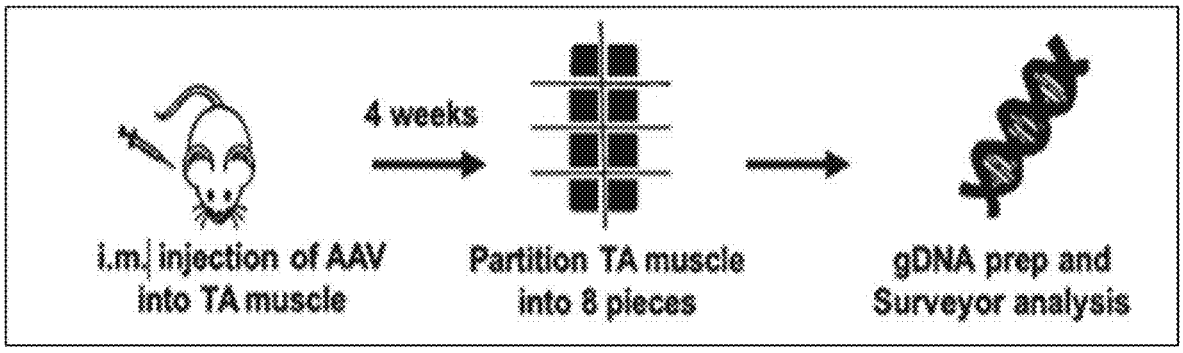

4 weeks
i.m. injection of AAV into TA muscle
Partition TA muscle into 8 pieces
gDNA prep and Surveyor analysis

Fig. 31

Fig. 32A

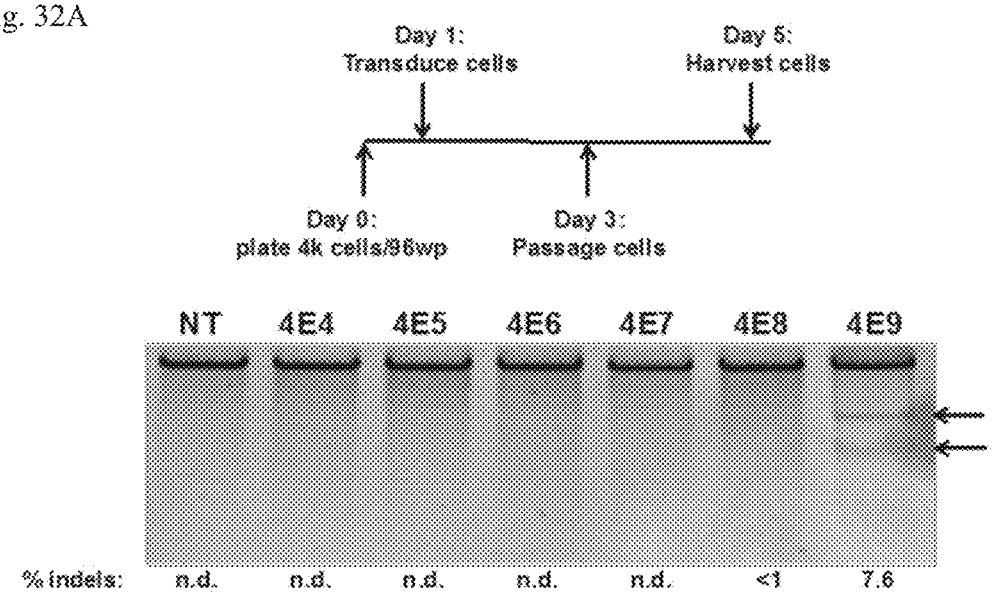


Fig. 32B

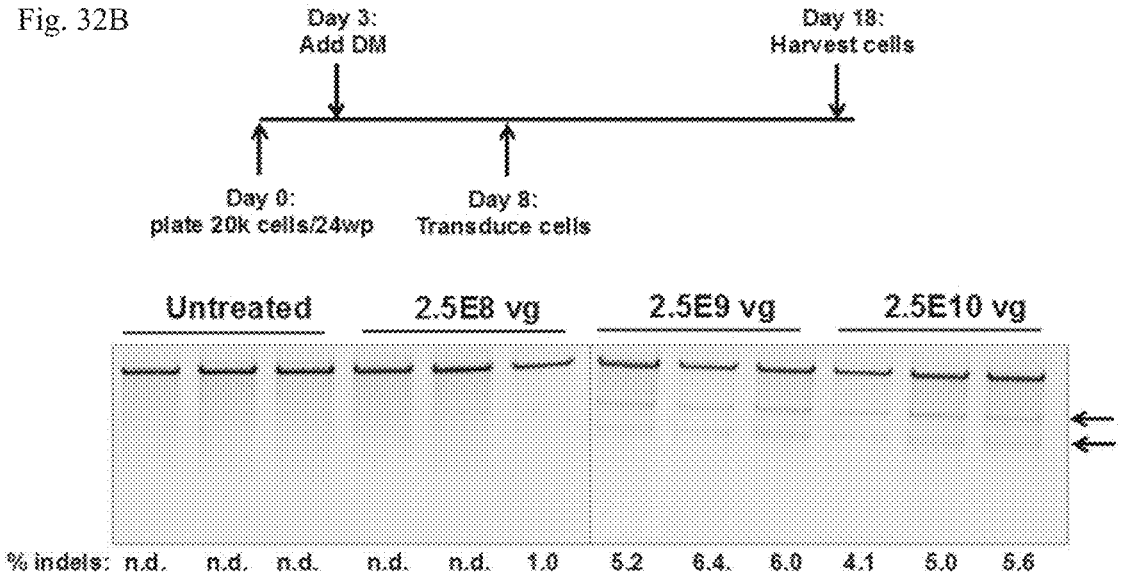
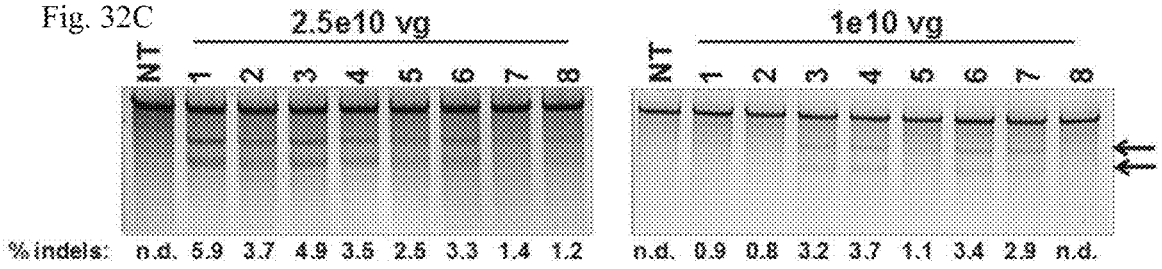


Fig. 32C



Rosa T2A opt DNA sequence (SEQ ID NO: 434)

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TCTCCGACCCTGGGGCGCTCGTCCGGCACCAAAGGACGCATACAGGGGAAAAGCCGTATAAGTGCCCCGAGTGTGGA
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GTGCGGGAAATCGTTTAGCAGATCCGATGACTTGGTGCGCCACCAGCGGACACACAGGGTGAAAAGCCCTACAAAT
GCCCGGAGTGTGGGAAGTCGTTTTCAAGGTCGGATCATCTGACTACCCATCAGCGCACCCATACGGGAGCggcggcc
cgcgccctGGTGAAGAGCGAGCTGGAGGAGAAGAAGTCCGAGCTGCGGCACAAGCTGAAGTACGTGCCCCACGAGTA
CATCGAGCTGATCGAGATCGCCAGGAACCCCACCCAGGACCGCATCCTGGAGATGAAGGTGATGGAGTTCTTCATGA
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CGCAAAACCAACTGCAATGGCGCCGTGCTGAGCGTGGAGGAGCTGCTGATCGGCGGCGAGATGATCAAAGCCGGCAC
CCTGACACTGGAGGAGGTGCGGCGCAAGTTCAACAACGGCGAGATCAACTTCGAGGGCAGAGGAAGTCTTCTAACAT
GCGGTGACGTGGAGGAGAATCCCGGCCCTAgatcTGACTACAAAGACCATGACGGTGATTATAAAGATCATGACATC
GATTACAAGGATGACGATGACAAGATGGCCCCCAAGAAGAAGaggaaggtgggcctcgagccGGGAGAGAAGCCGTA
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GACCGCATCCTGGAGATGAAGGTGATGGAGTTCTTCATGAAGGTGTACGGCTACAGGGGAGAGCACCTGGGCGGAAG
CAGAAAGCCTGACGGCGCCATCTATACAGTGGGCAGCCCCATCGATTACGGCGTGATCGTGGACACAAAGGCCTACA
GCGGCGGCTACAATCTGCCTATCGGCCAGGCCGACGAGATGGAGAGATACGTGGAGGAGAACCAGACACGGGATAAG
CACCTCAACCCCAACGAGTGGTGGAAGGTGTACCCTAGCAGCGTGACCGAGTTCAAGTTCCTGTTCGTGAGCGGCCA
CTTCAAGGGCAACTACAAGGCCCAGCTGACCAGGCTGAACCACATCACCAACTGCAATGGCGCCGTGCTGAGCGTGG
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Rosa T2A opt protein sequence (SEQ ID NO: 435)

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KSFSQRARLERHQRTHTGEKPYKCPECGKSFSESDDLVRHQRTHTGEKPYKCPECGKSFSRSDHLTTHQRTHTGAAA
KALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSP
IDYGVIVDTKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLN
HITNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF

EGRGSLLTCGDVEENPGPRSDYKDHDGDYKDHDI
DYKDDDDKMAPKKKRKVGLEPGEKPYKCPECGKSFSQSGDLRRHQRTHTGEKPYKCPECGKSFSTSGSLVRHQRTHT
GEKPYKCPECGKSFSQRAHLERHQRTHTGEKPYKCPECGKSFSTHLDLIRRQRTHTGEKPYKCPECGKSFSRKDNLK
NHQRTHTGEKPYKCPECGKSFSQLAHLRAHQRTHTGAAARALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQ
DRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVDTKAYSGGYNLPIGQADEMERTVEENQTRDK
HLNPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNHITNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNN
GEINF

Fig. 33

SASTG capsid sequence (SEQ ID NO: 436)

tggccactccctctatgcgcactgctcgctcggtgggcctggcgaccaaaggtgccagacggacgtgctttgca
cgtccggccccaccgagcgagcgagtgcgcatagagggagtggccaactccatcactagaggtatggcagtgacgta
acgcgaagcgcgcgaagcgagaccacgcctaccagctgcgtcagcagtcaggtgaccctttgcgacagtttgcgac
accacgtggccgctgagggtatatatttctcgagtgagcgaaccaggagctccatttgacgcgaaatttgaacgag
cagcagccatgccgggttctacgagattgtcctgaaggtcccgagtgacctggacgagcacctgccgggcatttct
aactcgtttgttaactgggtggccgagaaggaatgggagctgccgccggattctgacatggatccgaatctgattga
gcaggcacccctgaccgtggccgaaaagcttcagcgcgagttcctggtggagtggcgccgcgtgagtaaggccccgg
aggccctctttttttgtccagttcgaaaaggggagacctacttccacctgcacgtgctgattgagaccatcgggtc
aaatccatggtggtcggccgctacgtgagccagattaaagagaagctggtgaccgcatctaccgcgggtcgagcc
gcagcttccgaactggttcgcggtgaccaaaacgcgaaatggcgccgggggcgggaacaaggtggtggacgactgct
acatccccaactacctgctccccaagacccagccgagctccagtgggcgtggactaacatggaccagtatttaagc
gcctgtttgaatctgcggagcgtaaacggctggtggcgcagcatctgacgcacgtgtcgcagacgcaggagcagaa
caaagagaatcagaaccccaatttctgacgccgcggtcatcaggtcaaaaacctcagccaggtacatggagctggtcg
ggtggctggtggaccgcgggatcacgtcagaaaagcaatggattcaggaggaccaggcctcgtacatctccttcaac
gccgcctccaactcgcggtcccagatcaaggccgcgctggacaatgcctccaagatcatgagcctgacaaagacggc
tccggactacctggtgggcagcaacccgccggaggacattaccaaaaatcggatctaccaaatcctggagctgaacg
ggtacgatccgcagtacgcggcctcgtcttcctgggctgggcgcaaaagaagttcgggaagaggaacaccatctgg
ctctttgggccggccacgacgggtaaaaccaacatcgccggaagccatcgcccacgccgtgcccttctacggctgcgt
aaactggaccaatgagaacttcctcattcaacgatigcgtcgacaagatggtgatctggtgggaggagggcaagatga
cggccaaggtcgtggagagcgccaaggccattctgggcggaagcaaggtgcgcgtggaccaaaagtgcaagtcatcg
gcccagatcgaacccactcccgtgatcgtcacctccaacaccaacatgtgcgccgtgattgacgggaacagcaccac
cttcgagcatcagcagccgctgcaggaccggatgtttaaatttgaacttacccgccgttggaccatgactttggga
aggtcaccaaacaggaagtaaaggacttttccggtgggcttccgatcacgtgactgacgtggctcatgagttctac
gtcagaaagggtggagctaagaaacgcccccgcctccaatgacgcggatgtaagcgagccaaaacggcagtgcacgtc
acttgcgcagccgacaacgtcagacgcggaagcaccggcggactacgcggacaggtaccaaaacaaatgtttctcgtc
acgtgggcatgaatctgatgctttttcccctgtaaaacatgcgagagaatgaatcaaatttccaatgtctgttttacg
catggtcaaagagactgtgggaatgcttcccctggaatgtcagaatctcaacccgtttctgtcgtcaaaaagaagac
ttatcagaaactgtgtccaattcatcatatcctgggaagggcacccgagattgcctgttcggcctgcgatttggcca
atgtggacttggatgactgtgtttctgagcaataaatgacttaaaccaggtatggctgctgacggttatcttccaga
ttggctcgaggacaaccttttctgaaggcattcgtgagtggtgggctctgaaacctggagtccctcaacccaaagcga
accaacaacaccaggacaacgtcgcgggtcttgtgcttccgggttacaaatacctcggacccggtaacggactcgac
aaaggagagccggtcaacgaggcggacgcggcagccctcgaacacgacaaagcttacgaccagcagctcaaggccgg
tgacaacccgtacctcaagtacaaccacgccgacgccgagtttcaggagcgtcttcaagaagatacgtctttgggg
gcaaccttggcagaqcagtcttccaggccaaaaagaggatccttgagcctcttggtctggttgaggaagcagctaaa
acggctcctggaaagaagaggcctgtagatcagtctcctcaggaaccggactcatcatctggtgttggcaaatcggg
caaacagcctgccagaaaaagactaaatttcggtcagactggcgactcagagtcagtcccagaccctcaacctctcg
gagaaccaccagcagccccccacaagtttgtyatctaatacaatggcttcaggcggtggcgcaccaatggcagacaat
aacgagggtgccgatggagtgggtaattcctcaggaaattggcattgcgattcccaatggctgggcgacagagtcat
caccaccagcaccagaacctgggccctgcccacttacaacaaccatctctacaagcaaatctccagcGCCTtcaACGg
gagcttcaaacgacaaccactactttggctacagcacccctttgggggtattttgactttaacagattccactgccac
ttctcaccacgtgactggcagcgactcattaacaacaactgggqattccggcccaagaaactcagcttcaagctctt
caacatccaagttaaagaggtcacgcagaacgatggcacgacgactattgccaataacctaccagcacggttcaag
tgtttacggactcggagtatcagctcccgtacgtgctcgggtcggcgcaccaaggctgtctcccgccgtttccagcg
gacgtcttcatggtcccctcagtatggatacctcaccctgaacaacggaagtcaagcggtgggacgctcatcctttta
ctgcctggagtacttcccttcgcagatgctaaggactggaaataacttccaattcagctataccttcgaggatgtac
cttttcacagcagctacgctcacagccagagtttggatcgcttgatgaatcctcttattgatcagtatctgtactac
ctgaacagaacgcaaggaacaacctctggaacaaccaaccaatcacggctgcttttagccaggctgggcctcagtc
tatgtctttgcaggccagaaattggctacctgggccctgctaccggcaacagagactttcaaagactgctaacgaca
acaacaacagtaacttttccttggacagcggccagcaaatatcatctcaatggccgcgactcgctggtgaatccagga
ccagctatggccagtcacaaggacgatgaagaaaaattttccctatgcaoggcaatctaatatttggcaaagaagg
gacaacggcaagtaacgcagaattagataatgtaatgattacggatgaagaagagattcgtaccaccaatcctgtgg
caacagagcagtatggaactgtggcaaataacttgcagagctcaaatacagctcccacgactagaactgtcaatgat
caggggcccttacctggcatggtgtggcaagatcgtgacgtgtaccttcaaggacctatctgggcaaagattcctca
cacggatggacactttcatcctttctcctctgatgggaggctttggactgaaacatccgcctcctcaaatcatgatca
aaaatactccggtaccggcaaatcctccgacgactttcagcccggccaagtttgcttcatttatcactcagtactcc

Fig. 34

actggacaggtcagcgtggaaattgagtgggagctacagaaagaaaacagcaaacgttggaatccagagattcagta
cacttccaactacaacaagtctgttaatgtggactttactgtagacactaatggtgtttatagtgaacctcgcccta
ttggaacccggtatctcacacgaaacttgtaatcctggttaatcaataaacogtttaattcgtttcagttgaacttt
ggctcttgtgcaacttcttatcttatcttgtttccatggctactgcgtagataagcagcggcctgcggcgcttgcgct
tcgcggtttacaactgctggttaatatttaactctcgccatacctctagtgatggagttggccactccctctatgcg
cactcgctcgctcggtggggcccggacgtgcaaagcacgtccgtctggcgaccttggtcgccaggcccccaccgagcg
agcgagtgcgcatagagggagtggccaa

SASTG capsid peptide sequence (SEQ ID NO: 437)

MAADGYLPDWLEDNLSEGIREWWALKPGVPQPKANQQHQDNRRGLVLPGYKYLGPGNGLDKGEPVNEADAAALEHDK
AYDQQLKAGDNPYLKYNHADAEFQERLQEDTSFGGNLGRAVFQAKKRILEPLGLVEEAAKTAPGKKRPVIQSPQEPD
SSSGVGKSGKQPARKRLNFGQTGDSESVPDPQPLGEPPAAPTSLGSNTMASGGGAPMADNNEGADGVGNSSGNWHCD
SQWLGDRVITTSTRTWALPTYNNHLYKQISSASTGASNDNHYFGYSTPWGYFDFNRFHCHFSPRDWQRLINNNWGFR
PKKLSFKLFNIQVKEVTQNDGTTTIANNLTSTVQVFTDSEYQLPYVLGSAHQGCLPPFPADVFMVPQYGYLTLNNGS
QAVGRSSFYCLEYFPSQMLRTGNNFQFSYTFEDVPFHSSYAHSQSLDRLMNPLIDQYLYYLNRTQGTTSGTTNQSRL
LFSQAGPQSMSLQARNWLPGPCYRQQRLSKTANDNNNSNFPWTAASKYHLNGRDSLVNPGPAMASHKDDEEKFFPMH
GNLIFGKEGTTASNAELDNVMITDEEEIRTTNPVATEQYGTVANNLQSSNTAPTTRTVNDQGALPGMVWQDRDVYLQ
GPIWAKIPHTDGHFHPSPLMGGFGLKHPPPQIMIKNTPVPANPPTTFSPAKFASFITQYSTGQVSVEIEWELQKENS
KRWNPEIQYTSNYNKSVNVDFTVDTNGVYSEPRPIGTRYLTRNL

Fig. 34 (continued)

DZF16 ZFN target site (SEQ ID NO:442):

5'-CAAACTAGAAATGCCATCTTCCTTGATGTTGGAGGTACCTGC-3'

3'-GTTTGATCTTTACGGTAGAAGGAACTACAACCTCCATGGACG-5'

DZF16-L6 left full amino acid sequence (SEQ ID NO:443)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGKSFSPKDALRGHQRTHTGEKPY
KCPECGKSFSHKTTLTNHQPTHTGEKPYKCPECGKSFSQRNALAGHQRTHTGEKPYKCPECGKSFSHKNA
LQNHQRTHTGEKPYKCPECGKSFSDPGHLVRHQRTHTGEKPYKCPECGKSFSTSGNLVRHQPTHTGAAAR
ALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAI
YTVGSPIDYGVIVDTKAYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFKFLFVSGH
FKGNYKAQLTRLNHITNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

DZF16-R6 right full amino acid sequence (SEQ ID NO:444)

MDYKDHDGDYKDHDIDYKDDDDRMAPKKKRKVGRLEPGEKPYKCPECGKSFSQQRSLVGHQRTHTGEKPY
KCPECGKSFSDKKDLTRHQRTHTGEKPYKCPECGKSFSTSGHLVRHQRTHTGEKPYKCPECGKSFSQRAH
LERHQRTHTGEKPYKCPECGKSFSTSGSLVRHQRTHTGEKPYKCPECGKSFSTSGNLVRHQRTHTGAAAR
ALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAI
YTVGSPIDYGVIVDTKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGH
FKGNYKAQLTRLNRKTNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

Fig. 35

E51C3 target site (SEQ ID NO:445):

5'-(t)ATCTGCCCATGACTGGCGCAGGG(a)-3'

3'-(a)TAGACGGGTACTGACCGCGTCCC(t)-5'

E51C-3L left full amino acid sequence (SEQ ID NO:446)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPFQCRICMRNFSSKQALAVHTRTHTGEKPF
QCRICMRNFSQSTTLKRHLRTHTGEKPFQCRICMRNFSRSDHLSLHLKTHLRGSQLVKSELEEKKSELRH
KLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVDTK
AYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNRKT
NCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

E51C-3R right full amino acid sequence (SEQ ID NO:447)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPFQCRICMRNFSRRAHLQNHTRTHTGEKPF
QCRICMRNFSQSTTLKRHLRTHTGEKPFQCRICMRNFSDGGHLTRHLKTHLRGSQLVKSELEEKKSELRH
KLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVDTK
AYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNHIT
NCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

Fig. 36

DZF15 target site (SEQ ID NO:448):

5'-ACTAGAAATGCCATCTTCCTTGATGTTGGAGGTACCTGCTCT-3'
3'-TGATCTTTACGGTAGAAGGAACTACAACCTCCATGGACGAGA-5'

DZF15-L6 left full amino acid sequence (SEQ ID NO:449)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGKSFSHRTTLTNHQRTHTGEKPY
KCPECGKSFSQHNALAGHQRTHTGEKPYKCPECGKSFSHENALQNHQRTHTGEKPYKCPECGKSFSDPGH
LVRHQRTHTGEKPYKCPECGKSFSTSGNLVRHQRTHTGERPYKCPECGKSFSQSSNLVRHQRTHTGAAAR
ALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAI
YTVGSPIDYGVIVDTKAYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFKFLFVSGH
FKGNYKAQLTRLNHITNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

DZF15-R6 right full amino acid sequence (SEQ ID NO:450)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGKSFSQRNALAGHQRTHTGEKPY
KCPECGKSFSQQRSLVGHQRTHTGEKPYKCPECGKSFSDKKDLTRHQRTHTGEKPYKCPECGKSFSTSGH
LVRHQRTHTGERPYKCPECGKSFSQRAHLERHQRTHTGERPYKCPECGKSFSTSGSLVRHQRTHTGAAAR
ALVKSELEEKKSELRHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAI
YTVGSPIDYGVIVDTKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGH
FKGNYKAQLTRLNRKTNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

DZF15-L5 left full amino acid sequence (SEQ ID NO:451)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGKSFSQRNALAGHQRTHTGEKPY
KCPECGKSFSHENALQNHQRTHTGEKPYKCPECGKSFSDFGHLVRHQRTHTGEKPYKCPECGKSFSTSGN
LVRHQRTHTGEKPYKCPECGKSFSQSSNLVRHQRTHTGAAARALVKSELEEKKSELRHKLKYVPHEYIEL
IEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVDTKAYSGGYNLPIGQ
ADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNHITNCNGAVLSVEEL
LIGGEMIKAGTLTLEEVRRKFNNGEINF*

DZF15-R5 right full amino acid sequence (SEQ ID NO:452)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGKSFSQQRSLVGHQRTHTGEKPY
KCPECGKSFSDKKDLTRHQRTHTGEKPYKCPECGKSFSTSGHLVRHQRTHTGEKPYKCPECGKSFSQRAH
LERHQRTHTGEKPYKCPECGKSFSTSGSLVRHQRTHTGAAARALVKSELEEKKSELRHKLKYVPHEYIEL
IEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVDTKAYSGGYNLPIGQ
ADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNRKTNCNGAVLSVEEL
LIGGEMIKAGTLTLEEVRRKFNNGEINF*

Fig. 37

E51C4 target site (SEQ ID NO:453):

5'-(t)GCCATCTTCCTTGATGTTGGAGGT(a)-3'

3'-(a)CGGTAGAAGGAACTACAACCTCCA(t)-5'

E51C-4L left full amino acid sequence (SEQ ID NO:454)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGERPFQCRICMRNFSSPSKLARHTRTHTGEKPF
QCRICMRNFSVRHNLTRHLRTHTGEKPFQCRICMRNFSQRNNLGRHLKTHTGAAARALVKSELEEKKSEL
RHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVD
TKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNH
KTNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

E51C-4R right full amino acid sequence (SEQ ID NO:455)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGERPFQCRICMRNFSIPNHLARHTRTHTGEKPF
QCRICMRNFSQSASLKRHLRTHTGEKPFQCRICMRNFSHHNSLTRHLKTHTGAAARALVKSELEEKKSEL
RHKLKYVPHEYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVD
TKAYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNH
ITNCNGAVLSVEELLIGGEMIKAGTLTLEEVRRKFNNGEINF*

Fig. 38

"Single vector, multiplex CRISPR system"

"Dual vector, multiplex CRISPR system"

"Single vector, single gRNA system"

Fig. 39

SaCas9-NLS (NLS underlined) (SEQ ID NO: 459)

atgaaaaggaactacattctgggctggacatcgggattacaagcgtgggtatgggattattgactatgaaacaag
ggacgtgatcgacgcaggcgtcagactgttcaaggaggccaacgtggaaaacaatgagggacggagaagcaagaggg
gagccaggcgcctgaaacgacggagaaggcacagaatccagagggtgaagaaactgctgttcgattacaacctgctg
accgaccattctgagctgagtggaattaatccttatgaagccagggtgaaaggcctgagtcagaagctgtcagagga
agagtttttccgcagctctgctgcacctggctaagcgccgaggagtgcataacgtcaatgaggtggaagaggacaccg
gcaacgagctgtctacaaaggaacagatctcacgcaatagcaaagctctggaagagaagtatgtcgcagagctgcag
ctggaacggctgaagaaagatggcgaggtgagagggtcaattaataggttcaagacaagcgactacgtcaaagaagc
caagcagctgctgaaagtgcagaaggcttaccaccagctggatcagagcttcatcgatacttatatcgacctgctgg
agactcggagaacctactatgagggaccaggagaagggagccccttcggatggaaagacatcaaggaatggtacgag
atgctgatgggacattgcacctatttttccagaagagctgagaagcgtcaagtacgcttataacgcagatctgtacaa
cgccctgaatgacctgaacaacctggtcatcaccagggatgaaaacgagaaactggaatactatgagaagttccaga
tcatcgaaaacgtgtttaagcagaagaaaaagcctacactgaaacagattgctaaggagatcctggtcaacgaagag
gacatcaagggctaccgggtgacaagcactggaaaaccagagttcaccaatctgaaagtgtatcacgatattaagga
catcacagcacggaaagaaatcattgagaacgccgaactgctggatcagattgctaagatcctgactatctaccaga
gctccgaggacatccaggaagagctgactaacctgaacagcgagctgacccaggaagagatcgaacagattagtaat
ctgaaggggtacaccggaacacacaacctgtccctgaaagctatcaatctgattctggatgagctgtggcatacaaa
cgacaatcagattgcaatctttaaccggctgaagctggtcccaaaaaaggtggacctgagtcagcagaaagagatcc
caaccacactggtggacgatttcattctgtcacccgtggtcaagcggagcttcatccagagcatcaaagtgatcaac
gccatcatcaagaagtacggcctgcccaatgatatcattatcgagctggctagggagaagaacagcaaggacgcaca
gaagatgatcaatgagatgcagaaacgaaaccggcagaccaatgaacgcattgaagagattatccgaactaccggga
aagagaacgcaaagtacctgattgaaaaaatcaagctgcacgatatgcaggagggaaagtgctgtattctctggag
gccatccccctggaggacctgctgaacaatccattcaactacgaggtcgatcatattatccccagaagcgtgtcctt
cgacaattcctttaacaacaaggtgctggtcaagcaggaagagaactctaaaaagggcaataggactcctttccagt
acctgtctagttcagattccaagatctcttacgaaaccttaaaaagcacattctgaatctggccaaaggaaagggc
cgcatcagcaagaccaaaaaggagtacctgctggaagagcgggacatcaacagattctccgtccagaaggattttat
taacggaatctggtggacacaagataccgtactcgcggcctgatgaatctgctgcgatcctatttccgggtgaaca
atctggatgtgaaagtcaagtccatcaacgggcgttcacatctttctgaggcgcaaatggaagtttaaaaaggag
cgcaacaaagggtacaagcaccatgccgaagatgctctgattatcgcaaatgccgacttcatctttaaggagtggaa
aaagctggacaaagccaagaaagtgatggagaaccagatgtttcgaagagaagcaggccgaatctatgcccgaaatcg
agacagaacaggagtacaaggagatttttcatcactcctcaccagatcaagcatatcaaggatttcaaggactacaag
tactctcaccgggtggataaaaagcccaacagagagctgatcaatgacaccctgtatagtacaagaaaagacgataa
ggggaataccctgattgtgaacaatctgaacggactgtacgacaaagataatgacaagctgaaaaagctgatcaaca
aaagtcccgagaagctgctgatgtaccaccatgatcctcagacatatcagaaactgaagctgattatggagcagtac
ggcgacgagaagaacccactgtataagtactatgaagagactgggaactacctgaccaagtatagcaaaaaggataa
tggccccgtgatcaagaagatcaagtactatgggaacaagctgaatgcccatctggacatcacagacgattaccctа
acagtcgcaacaaggtggtcaagctgtcactgaagccatacagattcgatgtctatctggacaacggcgtgtataaa
tttgtgactgtcaagaatctggatgtcatcaaaaaggagaactactatgaagtgaatagcaagtgctacgaagaggc
taaaaagctgaaaaagattagcaaccaggcagagttcatcgcctcctttacaacaacgacctgattaagatcaatg
gcgaactgtatagggtcatcgggtgaacaatgatctgctgaaccgcattgaagtgaatatgattgacatcacttac
cgagagtatctggaaaacatgaatgataagcgccccctcgaattatcaaaacaattgcctctaagactcagagtat
caaaaagtactcaaccgacattctgggaaacctgtatgaggtgaagagcaaaaagcaccctcagattatcaaaaagg
gcagcggaggcaagcgtcctgctgctactaagaaagctggtcaagctaagaaaaagaaaaggatcctacccatacgat
gttccagattacgcttaa

SaCas9 gRNA (SEQ ID NO: 460)

[protospacer]gttttagtactctggaaacagaatctactaaaacaaggcaaaatgccgtgtttatctcgtcaac
ttgttggcgagatttttt

Fig. 40

NmCas9 (NLS 1 underlined NLS 2 underlined/bold, HA tag bold)(SEQ ID NO: 451)

atggtgcctaagaagaagagaaaggtggctgccttcaaacctaattcaatcaactacatcctcggcctcgatatcgg
catcgcatccgtcggctgggcgatggtagaaattgacgaagaagaaaacccccatccgcctgatttgatttgggcgtgc
gcgtatttgagcgtgccgaagtaccgaaaacaggcgactccttgccatggcaaggcgtttggcgccgcagtgttcgc
cgcctgacccgccgtcgcgcccacccgcctgcttcggacccgccgcctattgaaacgcgaaggcgtattacaagccgc
caattttgacgaaaacggcttgattaaatccttaccgaatacaccatggcaacttcgcgcagccgcattagaccgca
aactgacgcctttagagtggtcggcagtcttgttgcatttaatcaaacatcgcggctatttatcgcaacggaaaaac
gagggcgaaactgccgataaggagcttggcgctttgcttaaaggcgtagccggcaatgcccatgccttacagacagg
cgatttccgcacaccggccgaattggctttaaataaatttgagaaagaaagcggccatatccgcaatcagcgcagcg
attattcgcatacgttcagccgcaaagatttacaggcggagctgattttgctgtttgaaaaacaaaaagaatttggc
aatccgcatgtttcaggcggccttaaagaaggtattgaaaccctactgatgacgcaacgccctgccctgtccggcga
tgccgttcaaaaaatgttggggcattgcaccttcgaacggcagagccgaaagccgctaaaaacacctacacagccg
aacgtttcatctggctgaccaagctgaacaacctgcgtatttagagcaaggcagcgagcggccattgaccgatacc
gaacgcgccacgcttatggacgagccatacagaaaatccaaactgacttacgcacaagcccgtaagctgctgggttt
agaagataccgccttttttcaaaggcttgcgctatggtaaagacaatgccgaagcctcaacattgatggaaatgaagg
cctaccatgccatcagccgtgcactggaaaaagaaggattgaaagacaaaaaatcccccattaaacctttctcccgaa
ttacaagacgaaatcggcacggcattctccctgttcaaaaccgatgaagacattacaggccgtctgaaagaccgtat
acagcccgaaatcttagaagcgctgttgaaacacatcagcttcgataagttcgtccaaatttccttgaaagcattgc
gccgaattgtgcctctaatggaacaaggcaaacgttacgatgaagcctgcgccgaaatctacggagaccattacggc
aagaagaatacggaagaaaagatttatctgccgccgattcccgccgacgaaatccgcaacccccgtcgtcttgcgcgc
cttatctcaagcacgtaaggtcattaacggcgtggtacgccgttacggctcccccagctcgtatccatattgaaactg
caagggaagtaggtaaatcgtttaaagaccgcaaagaaattgagaaacgccaagaagaaaaccgcaaagaccgggaa
aaagccgccgccaaattccgagagtatttcccccaatttgtcggagaacccaaatccaaagatattctgaaactgcg
cctgtacgagcaacaacacggcaaatgcctgtattcgggcaaagaaatcaacttagggcgtctgaacgaaaaaggct
atgtcgaaatcgaccatgccctgccgttctcgcgcacatgggacgacagtttcaacaataaagtactggtattgggc
agcgaaaaccaaaacaaaggcaatcaaaccccttacgaatacttcaacggcaaagacaacagccgcgaatggcagga
atttaaagcgcgtgtcgaaaccagccgtttcccgcgcagtaaaaaacaacggattctgctgcaaaaattcgatgaag
acggctttaaagaacgcaatctgaacgacacgcgctacgtcaaccgtttcctgtgtcaatttgttgccgaccgtatg
cggctgacaggtaaaggcaagaaacgtgtctttgcatccaacggacaaattaccaatctgttgcgcggcttttgggg
attgcgcaaagtgcgtgcggaaaacgaccgccatcacgccttggacgccgtcgtcgttgcctgctcgaccgttgcca
tgcagcagaaaattaccgttttgtacgctataaagagatgaacgcgtttgacggtaaaaccatagacaaagaaaca
ggagaagtgctgcatcaaaaaacacacttcccacaacottgggaatttttcgcacaagaagtcatgattcgcgtctt
cggcaaaccggacggcaaacccgaattcgaagaagccgataccctagaaaaactgcgcacgttgcttgccgaaaaat
tatcatctcgccccgaagccgtacacgaatacgttacgccactgtttgtttcacgcgcgcccaatcggaagatgagc
gggcaagggcatatggagaccgtcaaatccgccaaacgactggacgaaggcgtcagcgtgttgcgcgtaccgctgac
acagttaaaactgaaagacttggaaaaaatggtcaatcgggagcgcgaacctaagctatacgaagcactgaaagcac
ggctggaagcacataaagacgatcctgccaaagcctttgccgagccgttttacaaatacgataaagcaggcaacgc
acccaacaggtaaaagccgtacgcgtagagcaagtacagaaaaccggcgtatgggtgcgcaaccataacggtattgc
cgacaacgcaaccatggtgcgcgtagatgtgtttgagaaaggcgacaagtattatctggtaccgatttacagttggc
aggtagcgaaagggatttgccggatagggctgttgtacaaggaaaagatgaagaagattggcaacttattgatgat
agtttcaactttaaattctcattacaccctaatgatttagtcgaggttataacaaaaaaagctagaatgtttggtta
ctttgccagctgccatcgaggcacaggtaatatcaatatacgcattcatgatcttgatcataaaattggcaaaaatg
gaatactggaaggtatcggcgtcaaaaccgccctttcattccaaaaatacccaaattgacgaactgggcaaagaaatc
agaccatgccgtctgaaaaaacgcccgcctgtccgttacccatacgatgttccagattacgctgcagctccagcagc
gaagaaaaagaagctggattaa

NmCas9 short hairpin from Thomson PNAS 2013 (SEQ ID NO: 462)

[protospacer]GTTGTAGCTCCCTTTCTCATTTCGGAAACGAAATGAGAACCGTTGCTACAATAAGGCCGTCTGA
AAAGATGTGCCGCAACGCTCTGCCCCTTAAAGCTTCTGCTTTAAGGGGCTTTTTTT

NmCas9 long hairpin from Church Nature Biotech 2013 (SEQ ID NO: 463)

[protospacer]GTTGTAGCTCCCTTTCTCATTTCGCAGTGCTACAATGAAAATTGTCGCACTGCGAAATGAGAACC
GTTGCTACAATAAGGCCGTCTGAAAAGATGTGCCGCAACGCTCTGCCCCTTAAAGCTTCTGCTTTAAGGGGCTTTTT
TT

Fig. 41

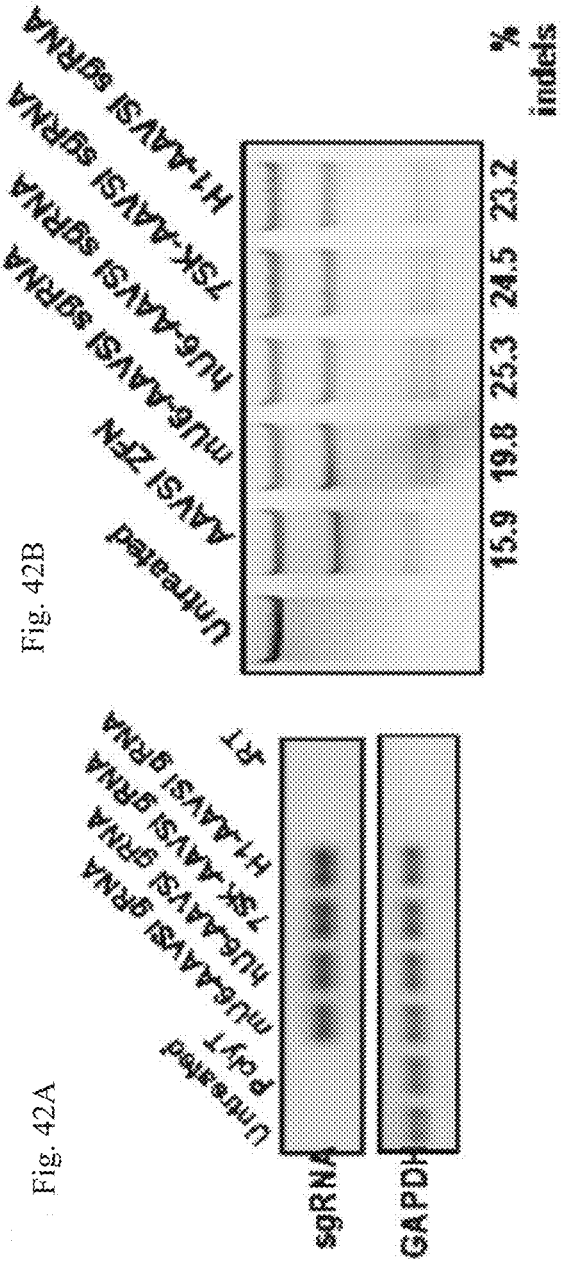


Fig. 42A

Fig. 42B

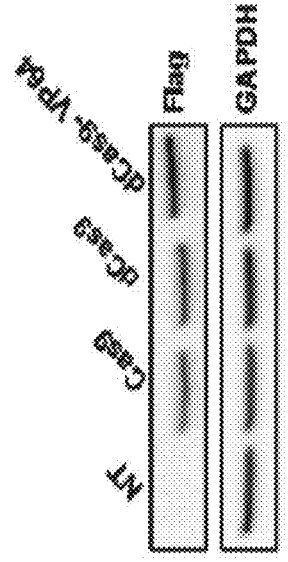


Fig. 42C

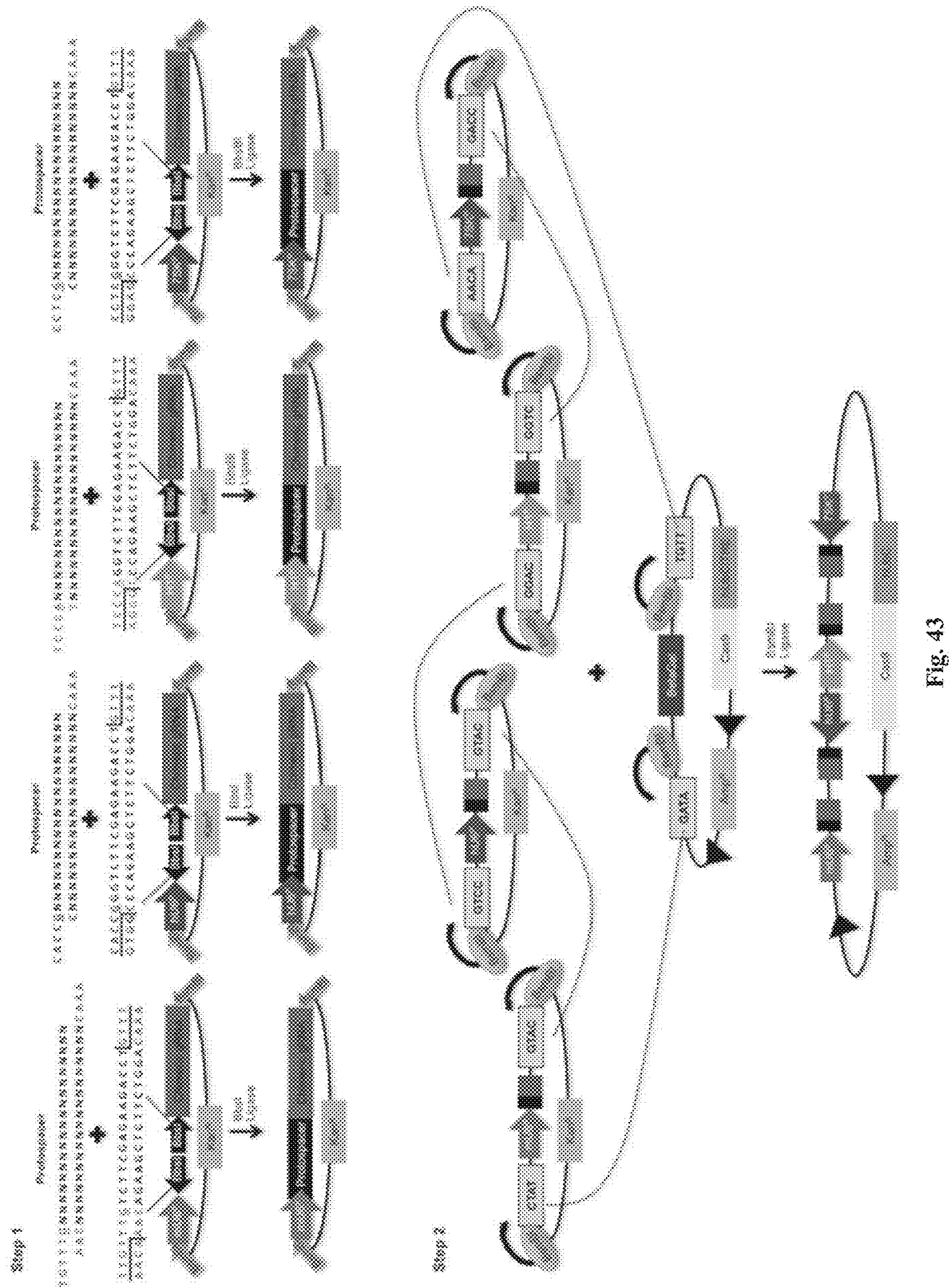


Fig. 43

Fig. 44A

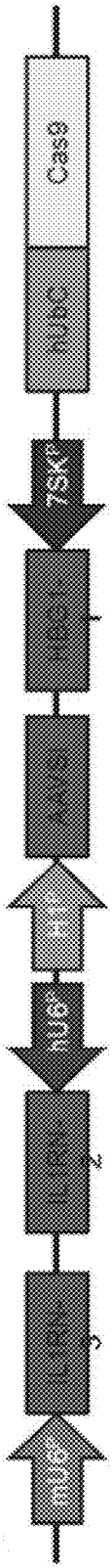
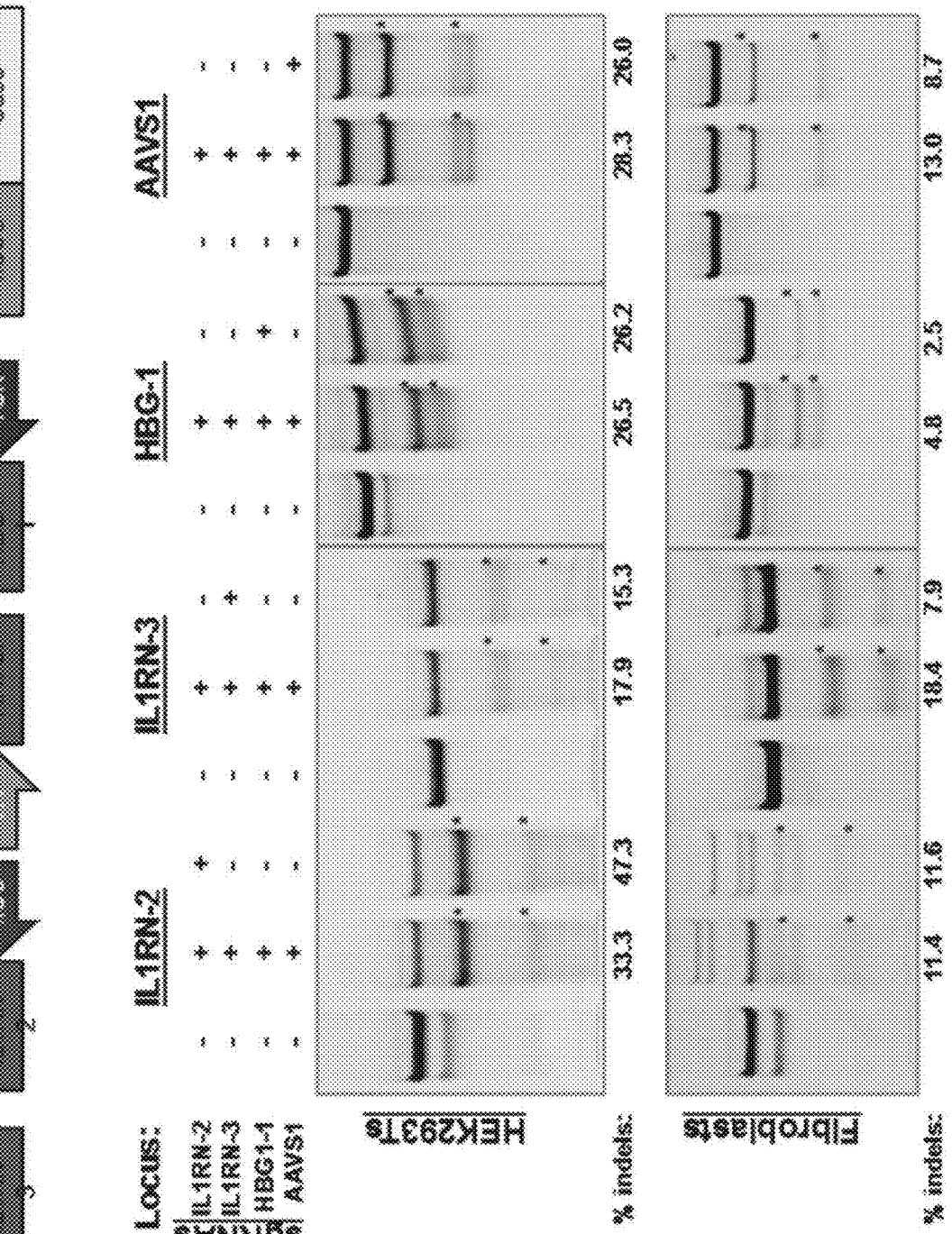


Fig. 44B



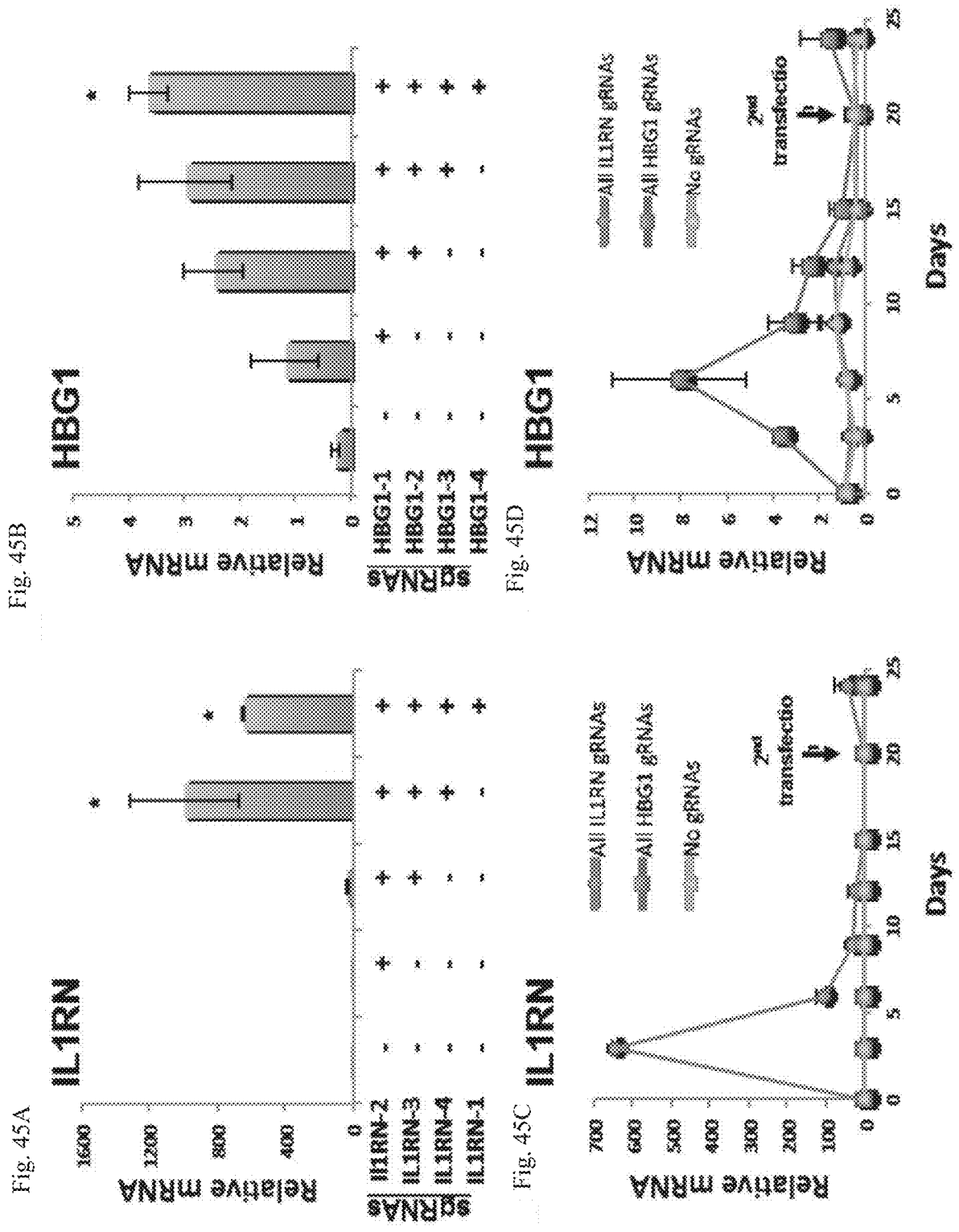


Fig. 45A

Fig. 45B

Fig. 45C

Fig. 45D

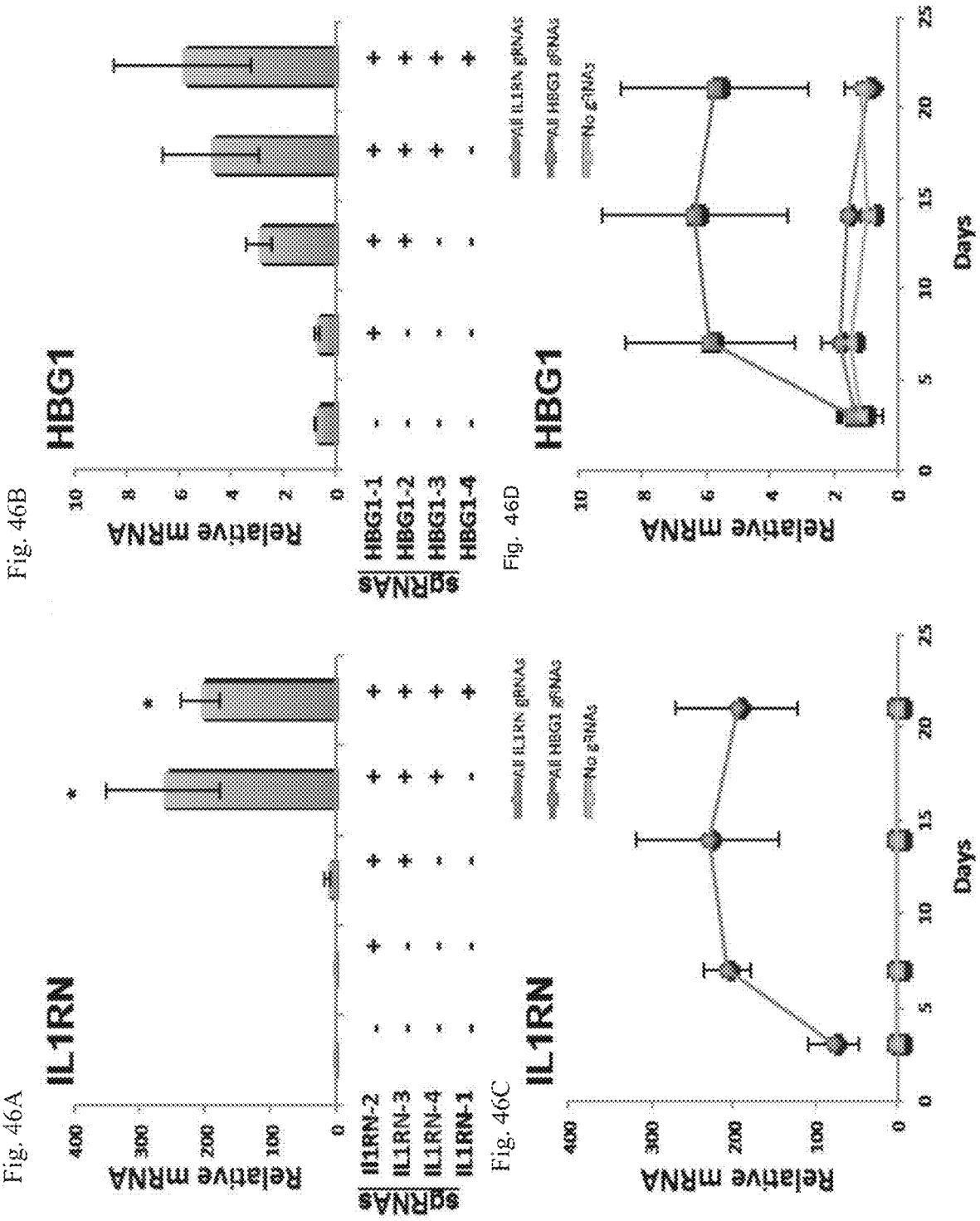


Fig. 46A

Fig. 46B

Fig. 46C

Fig. 46D

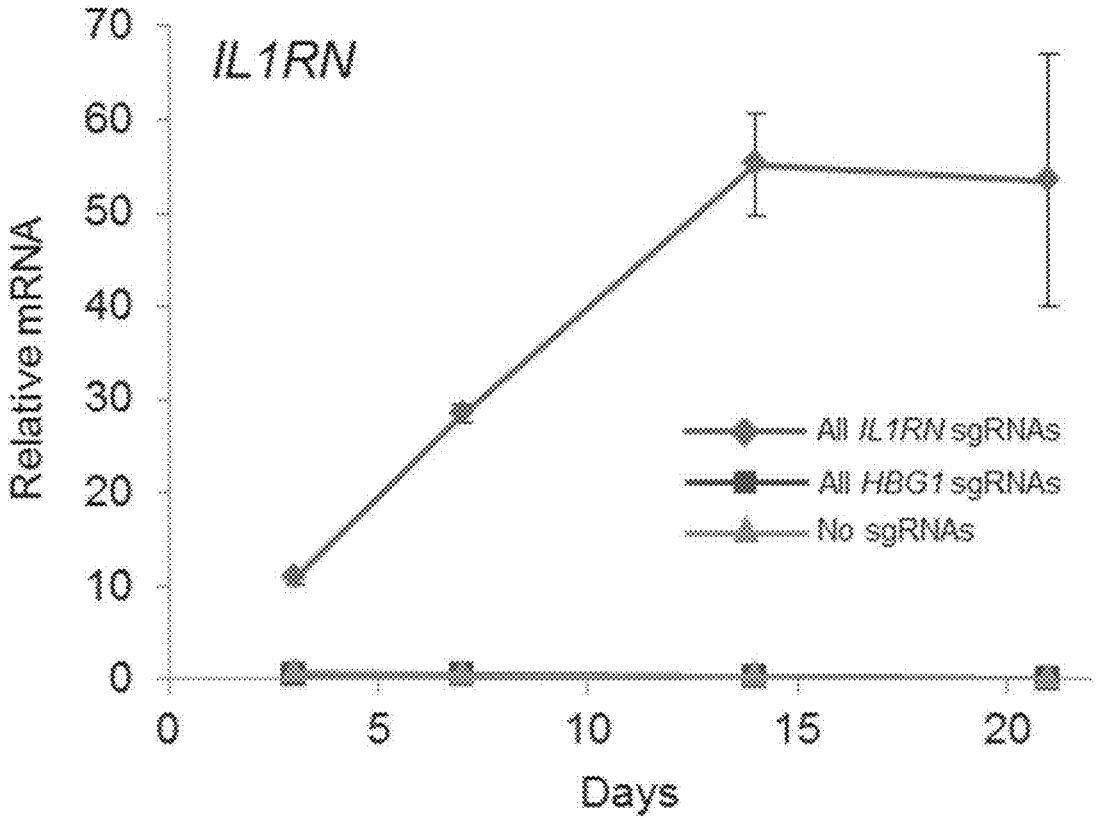

IL1RN
Relative mRNA
70
60
50
40
30
20
10
0
0
5
10
15
20
Days
All IL1RN sgRNAs
All HBG1 sgRNAs
No sgRNAs

Fig. 47

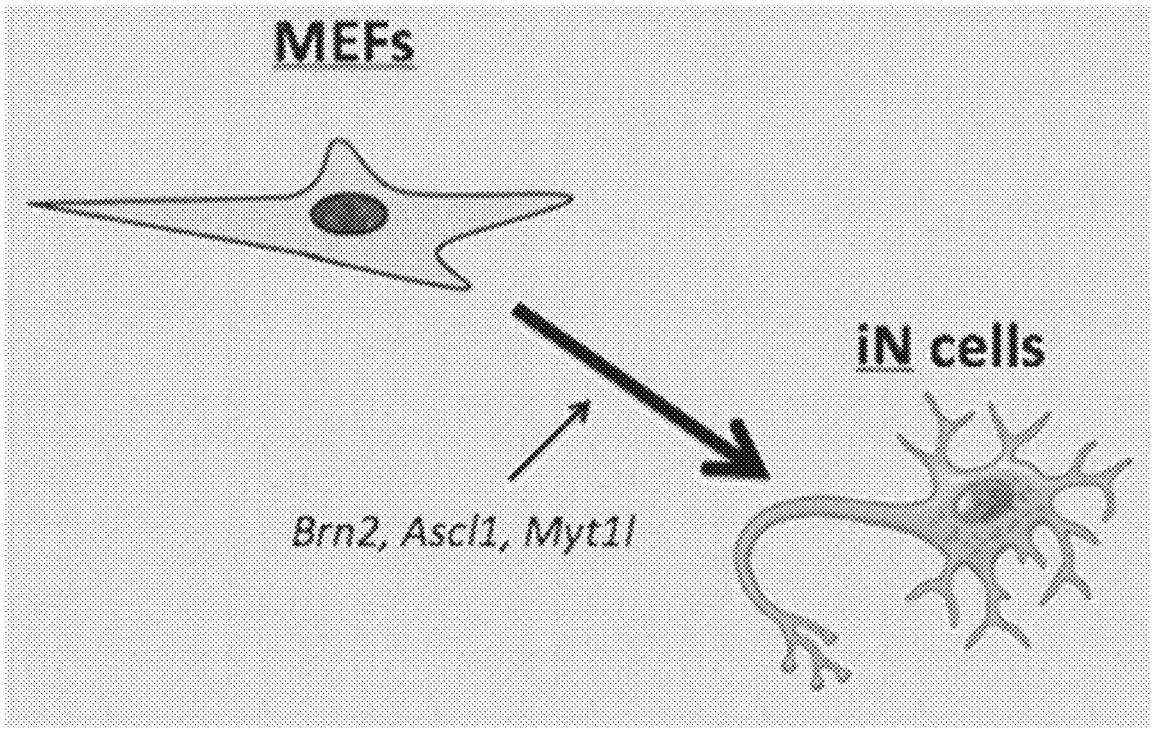


Fig. 48

Fig. 49A

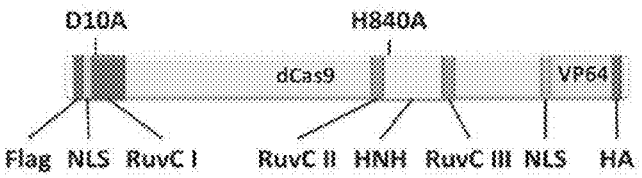


Fig. 49B

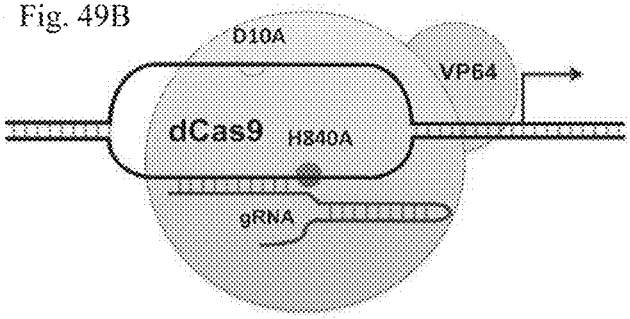


Fig. 49C

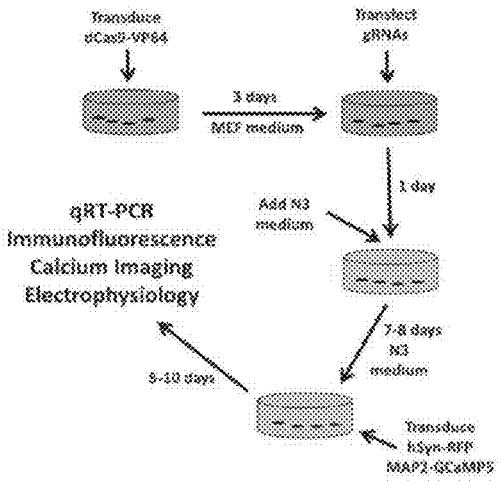


Fig. 50A

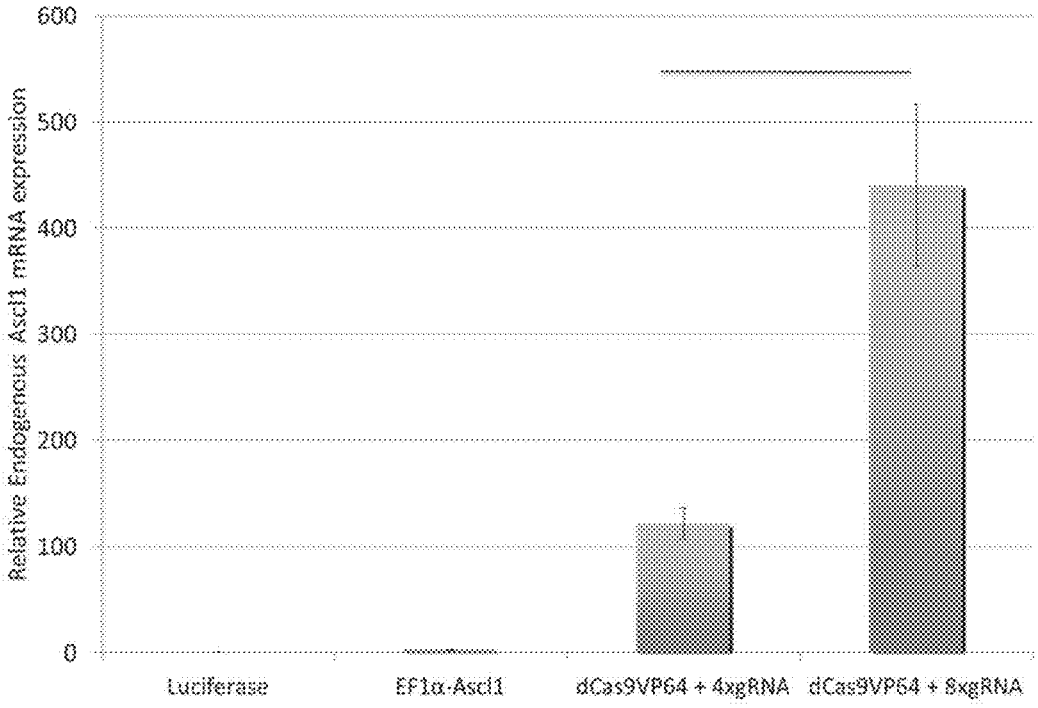


Fig. 50B

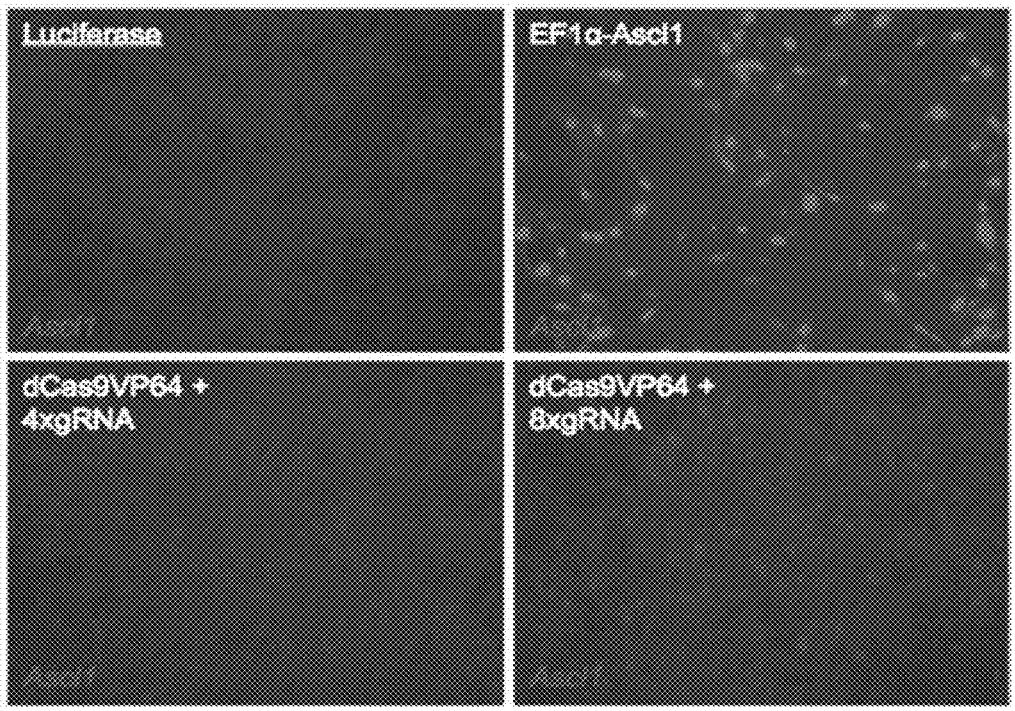


Fig. 51A

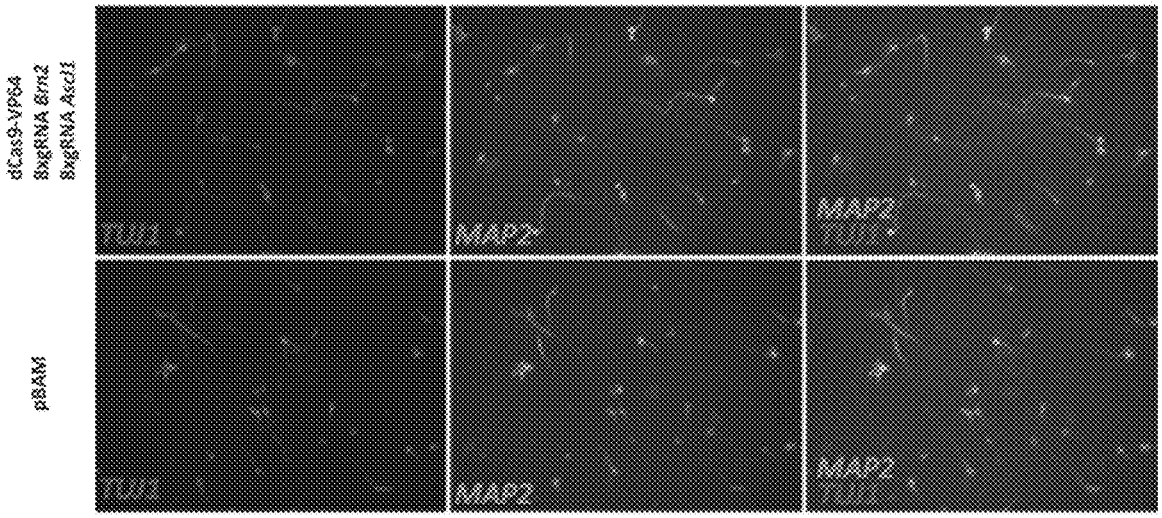


Fig. 51B

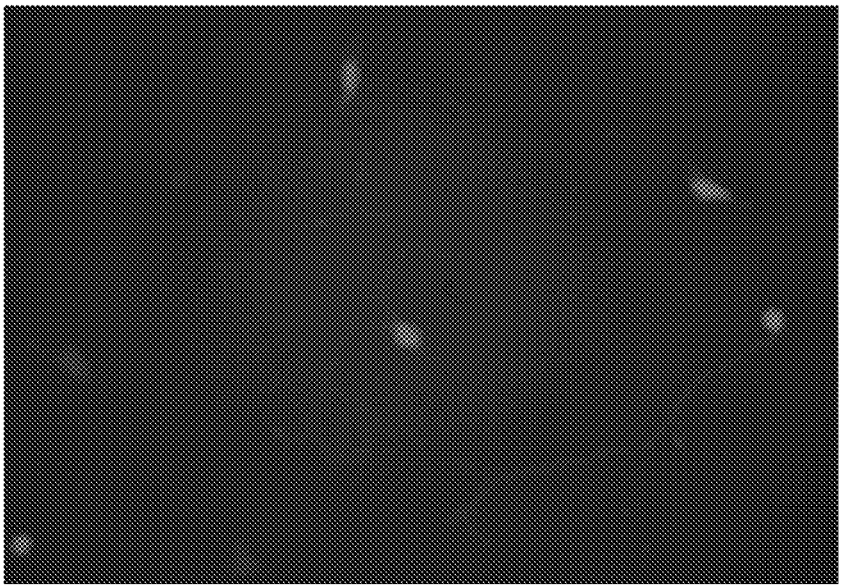


Fig. 52A

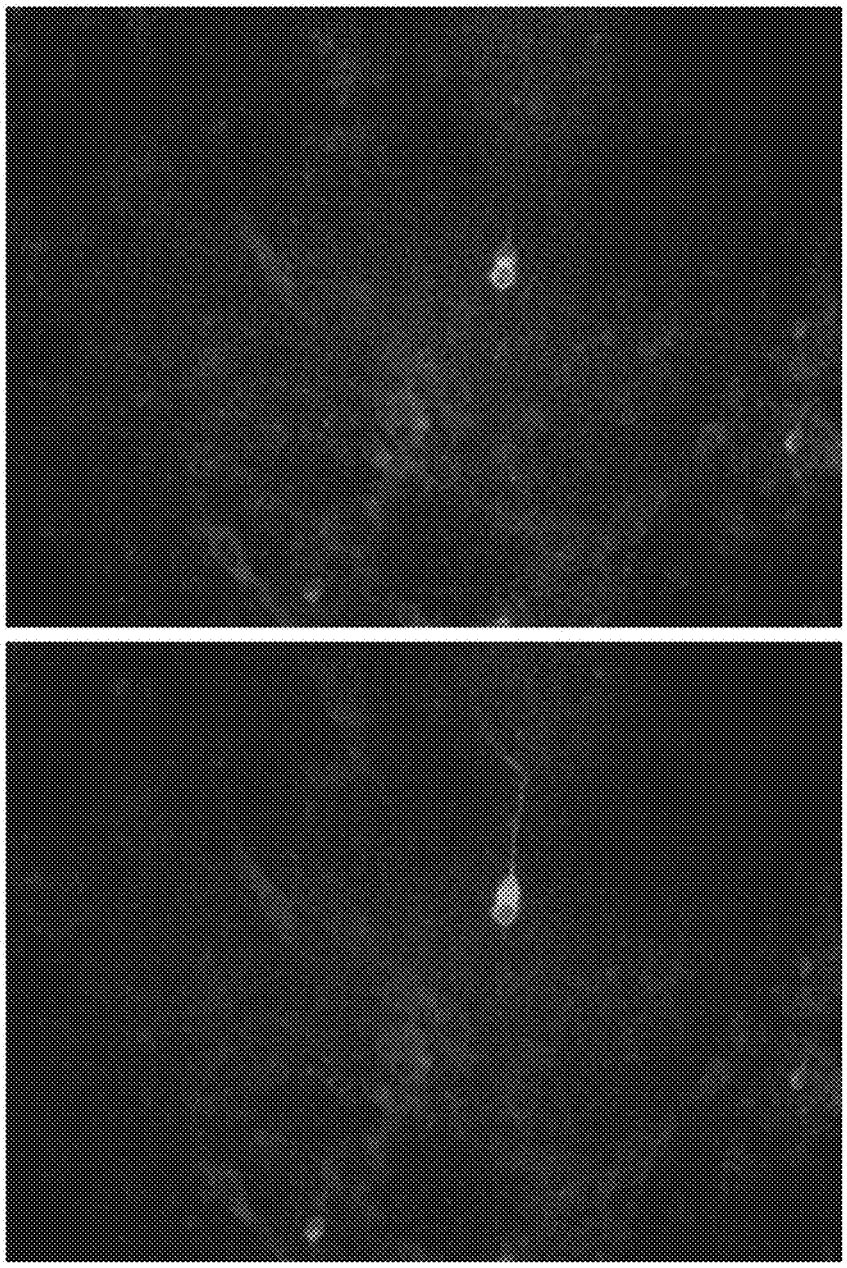


Fig. 52B

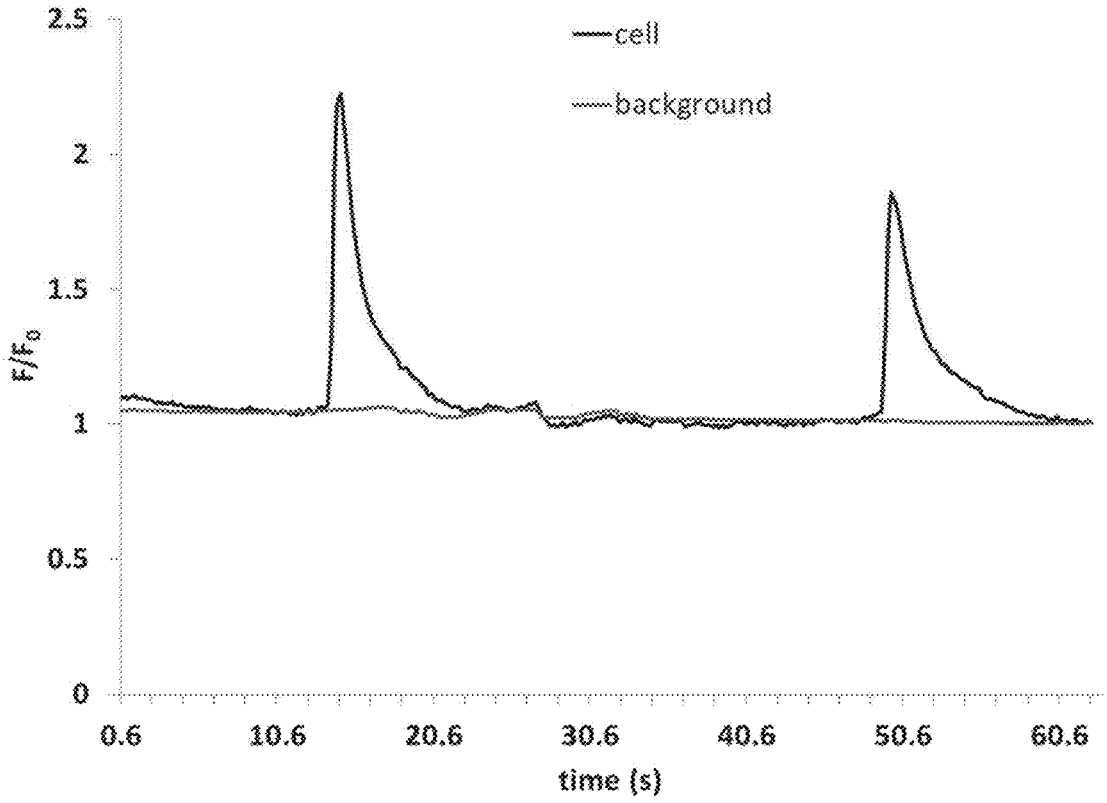

2.5
2
1.5
1
0.5
0
F/F0
cell
background
0.6
10.6
20.6
30.6
40.6
50.6
60.6
time (s)

Fig. 53A

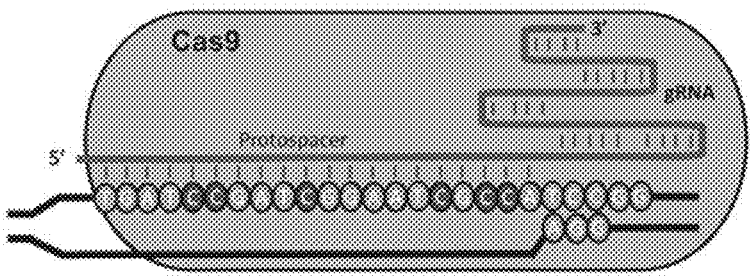


Fig. 53B

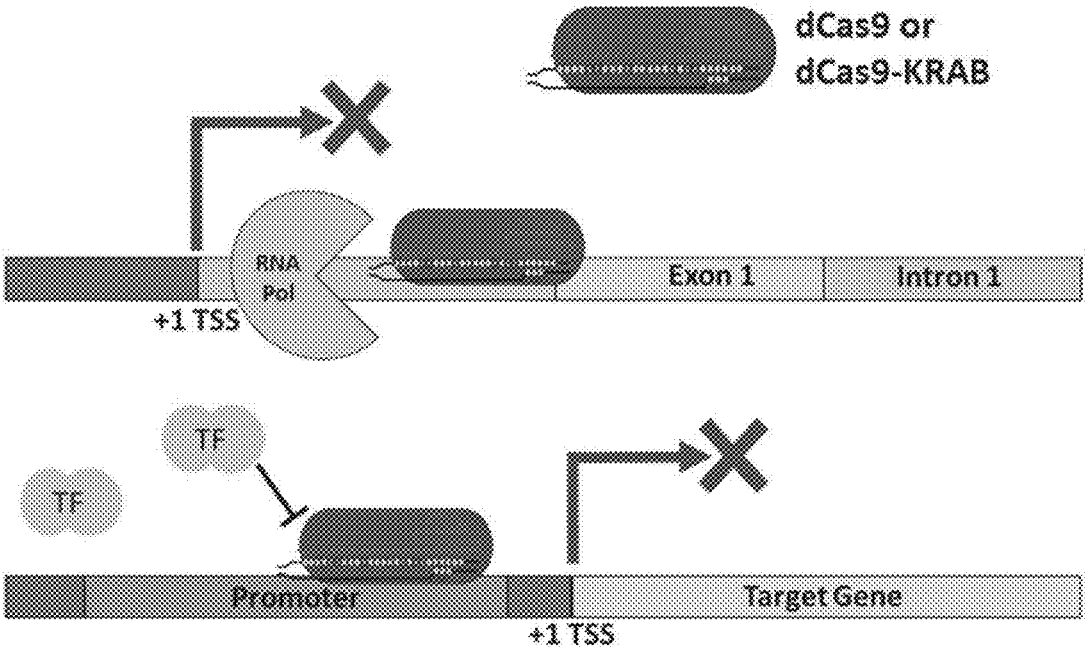
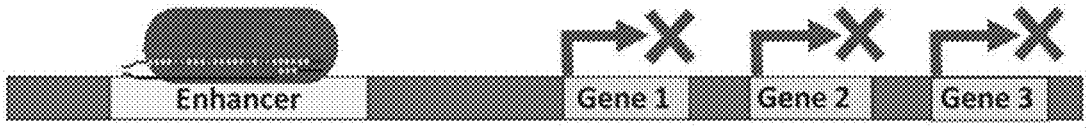


Fig. 53C



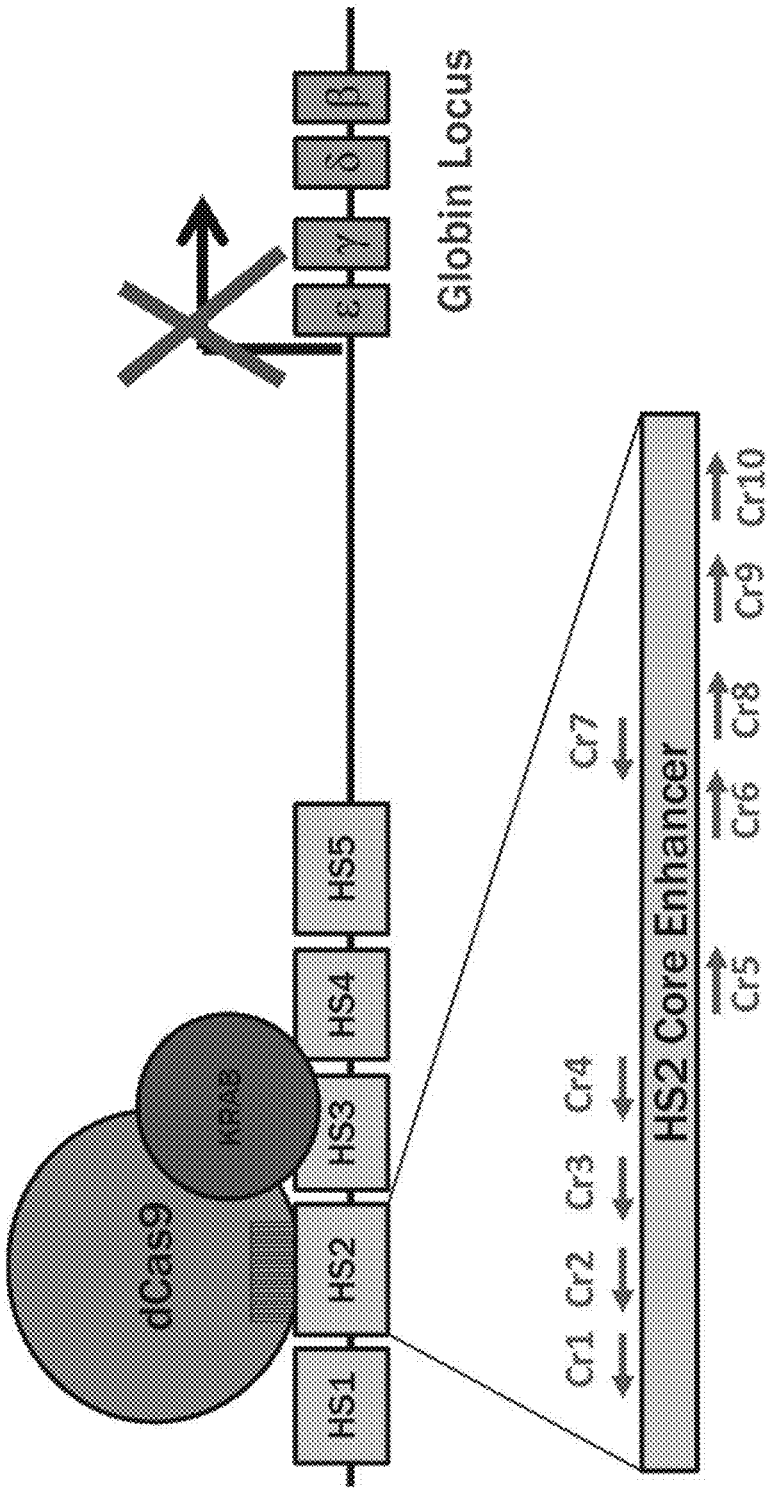

Globin Locus
ε
Gγ
Aγ
δ
β
HS5
HS4
HS3
HS2
HS1
dCas9
KRAB
Cr1
Cr2
Cr3
Cr4
Cr5
Cr6
Cr7
Cr8
Cr9
Cr10
HS2 Core Enhancer

Fig. 54

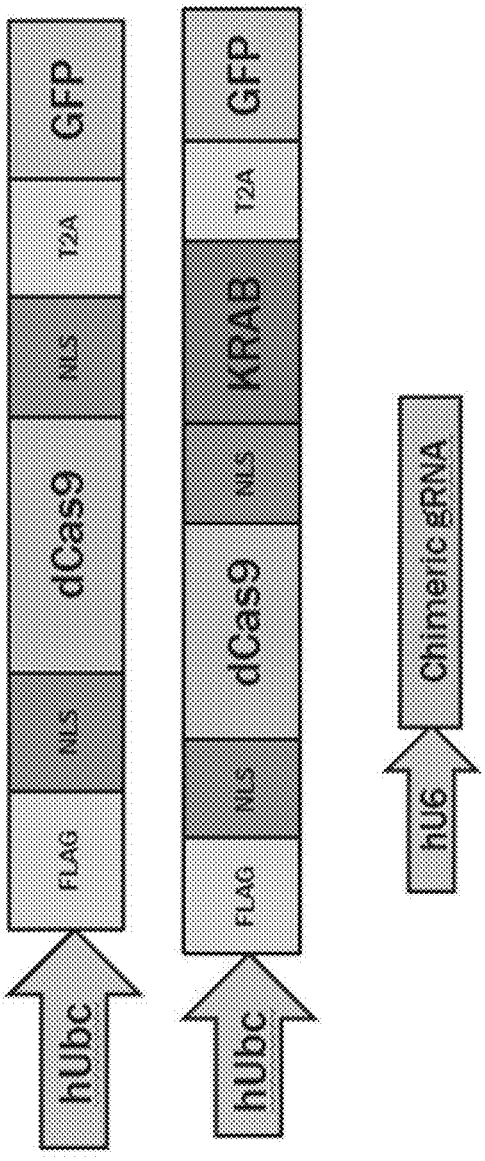

hUbc
FLAG
NLS
dCas9
NLS
T2A
GFP
hUbc
FLAG
NLS
dCas9
NLS
KRAB
T2A
GFP
hU6
Chimeric gRNA

Fig. 55A

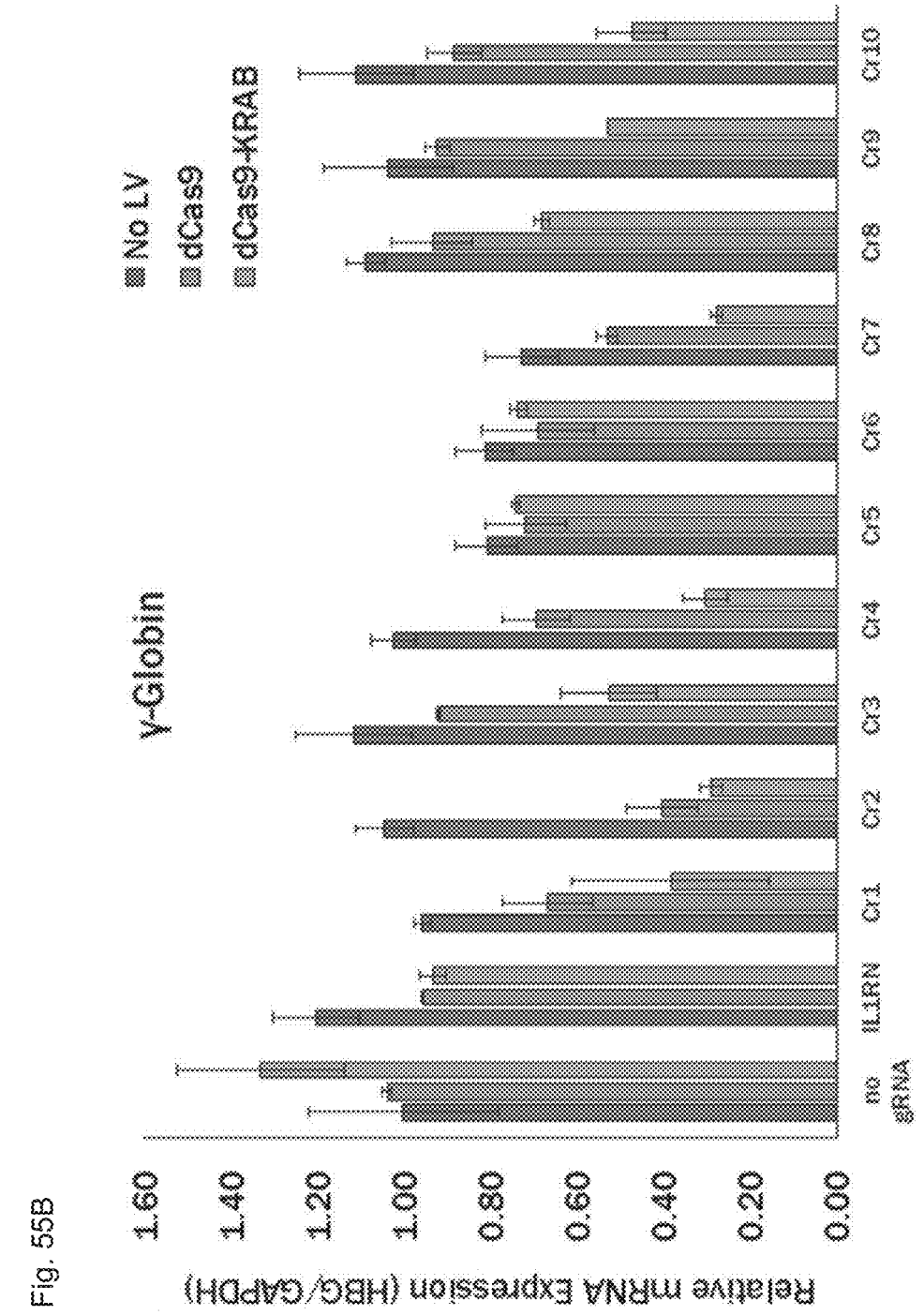

γ-Globin
No LV
dCas9
dCas9-KRAB
Relative mRNA Expression (HBG/GAPDH)
1.60
1.40
1.20
1.00
0.80
0.60
0.40
0.20
0.00
no gRNA
IL1RN
Cr1
Cr2
Cr3
Cr4
Cr5
Cr6
Cr7
Cr8
Cr9
Cr10

Fig. 55B

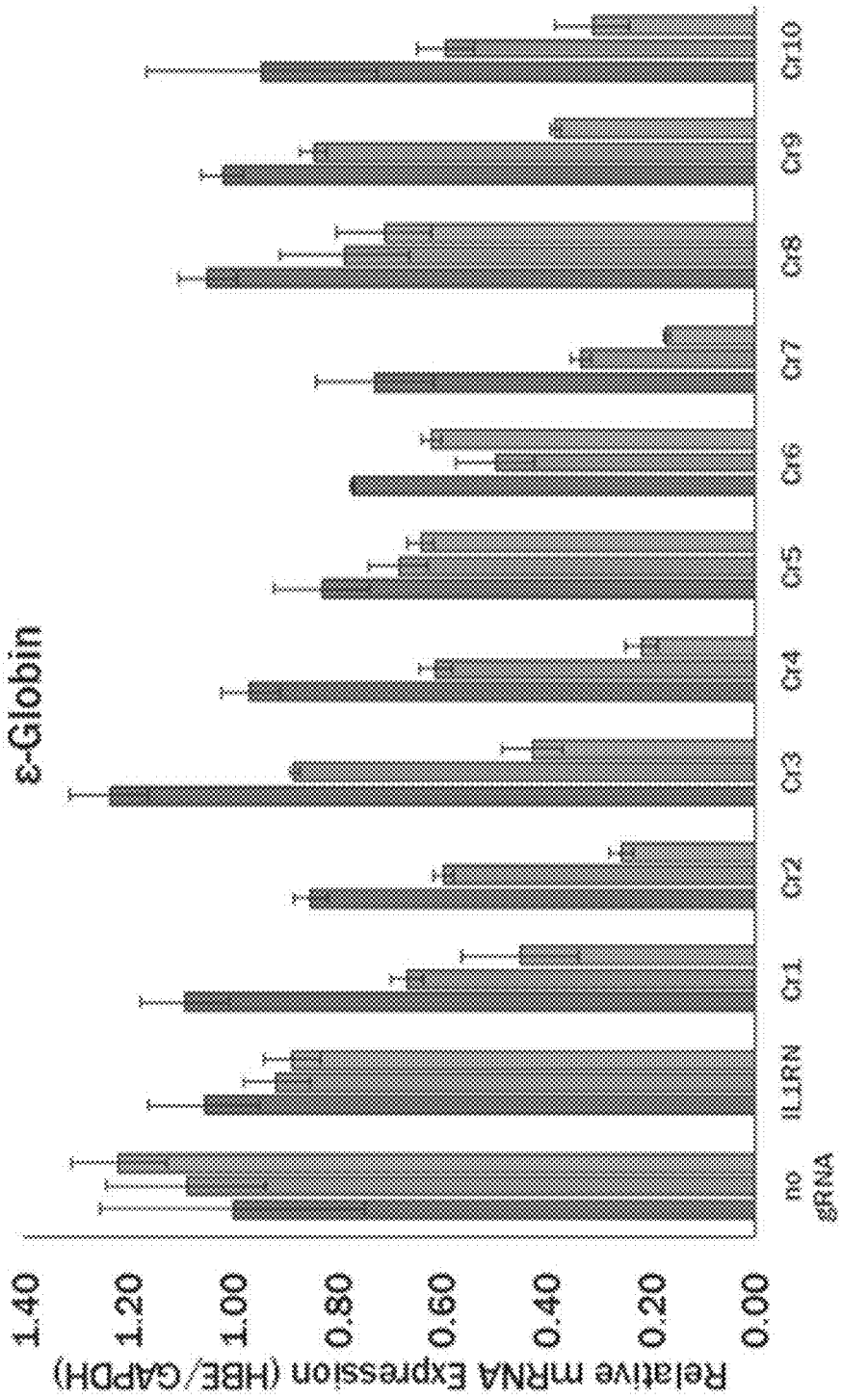

ε-Globin
Relative mRNA Expression (HBE/GAPDH)
1.40
1.20
1.00
0.80
0.60
0.40
0.20
0.00
no gRNA
IL1RN
Cr1
Cr2
Cr3
Cr4
Cr5
Cr6
Cr7
Cr8
Cr9
Cr10

Fig. 55C

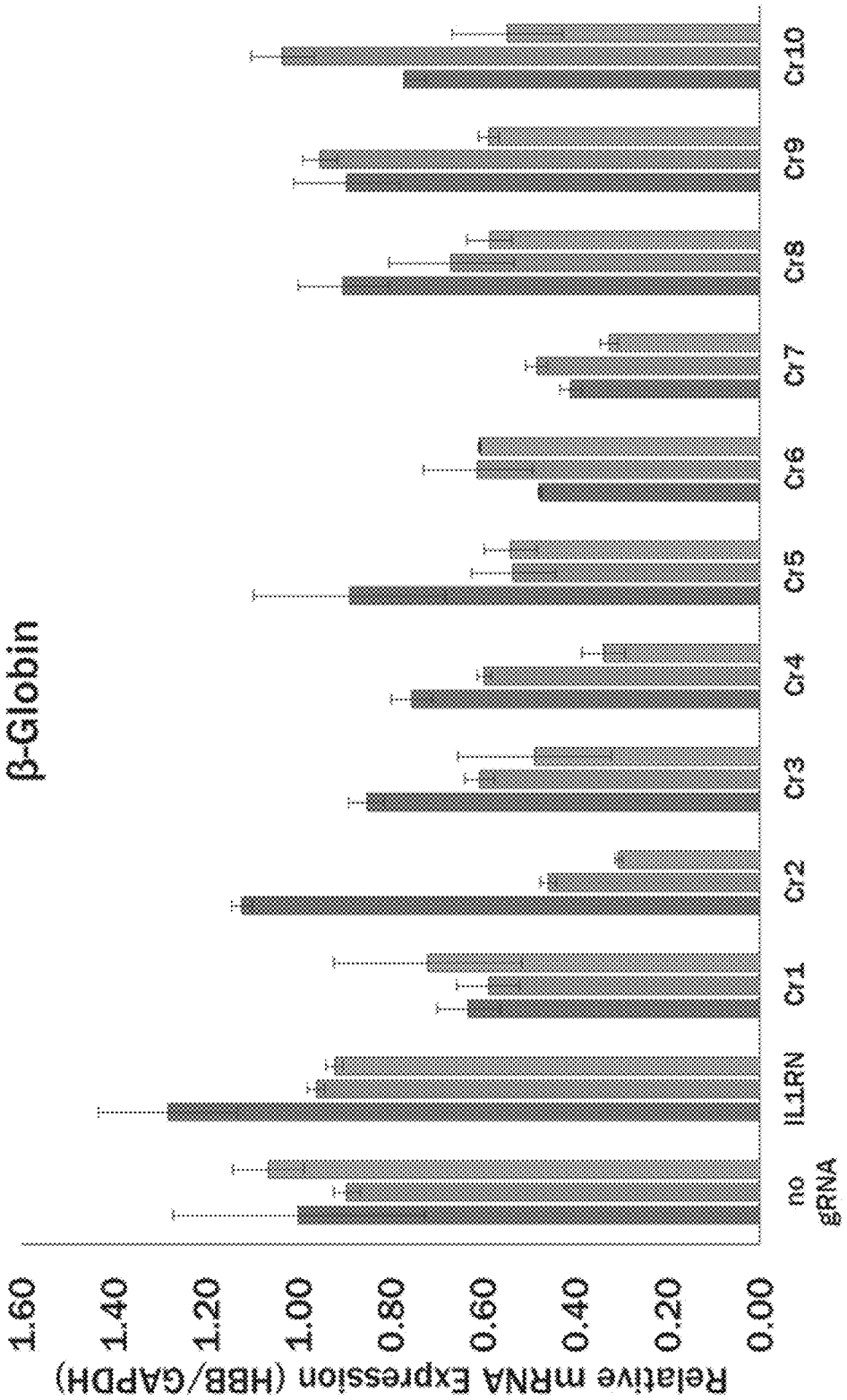

β-Globin
Relative mRNA Expression (HBB/GAPDH)
1.60
1.40
1.20
1.00
0.80
0.60
0.40
0.20
0.00
no gRNA
IL1RN
Cr1
Cr2
Cr3
Cr4
Cr5
Cr6
Cr7
Cr8
Cr9
Cr10

Fig. 55D

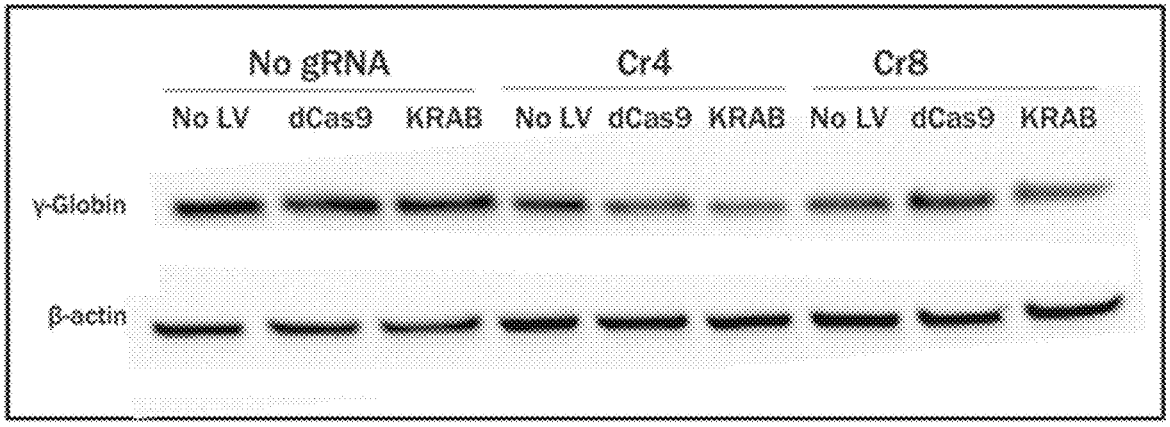

No gRNA
Cr4
Cr8
No LV
dCas9
KRAB
No LV
dCas9
KRAB
No LV
dCas9
KRAB
γ-Globin
β-actin

Fig. 55E

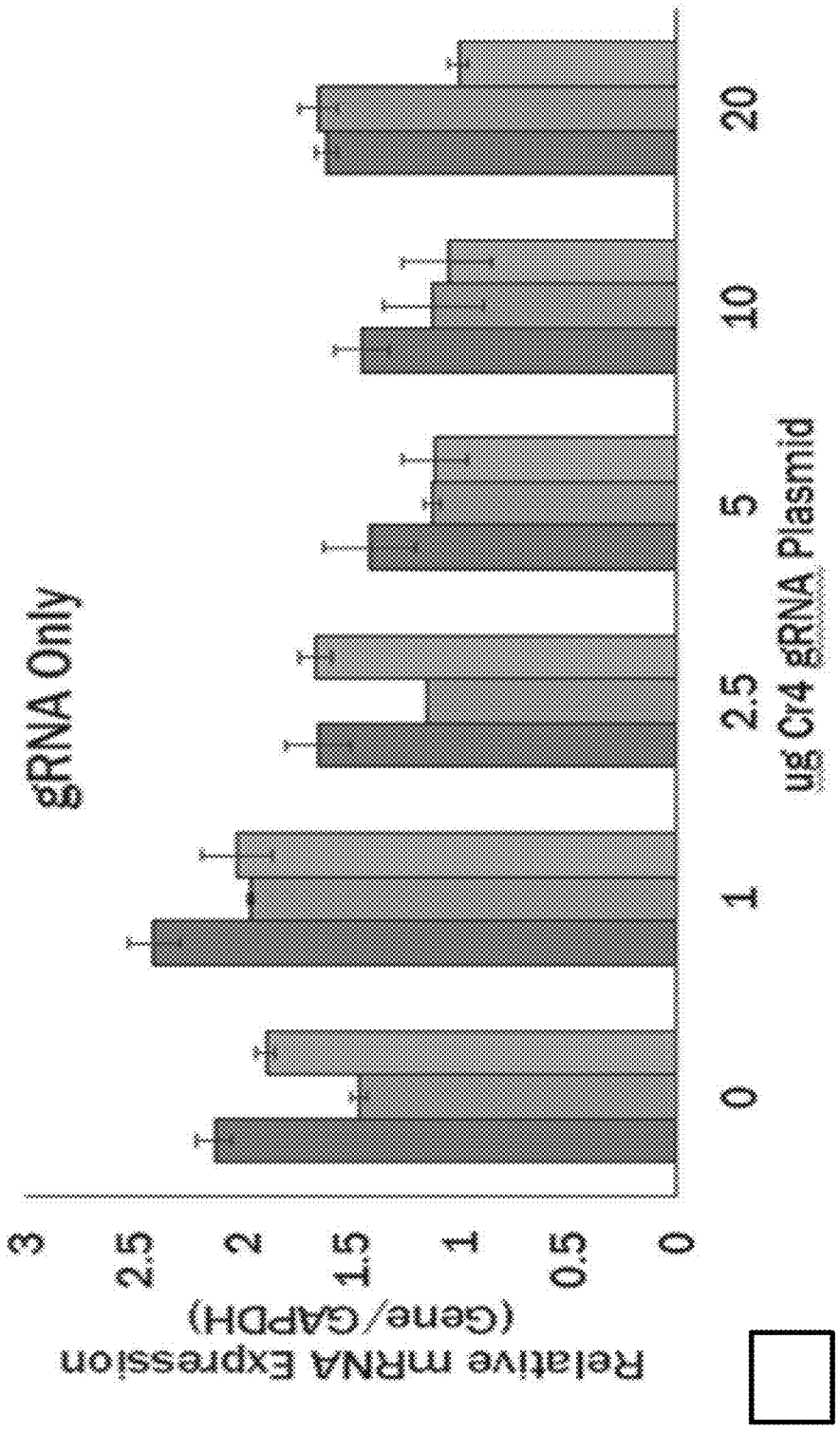

Relative mRNA Expression
(Gene/GAPDH)
3
2.5
2
1.5
1
0.5
0
0
1
2.5
5
10
20
ug Cr4 gRNA Plasmid
gRNA Only

Fig. 56A

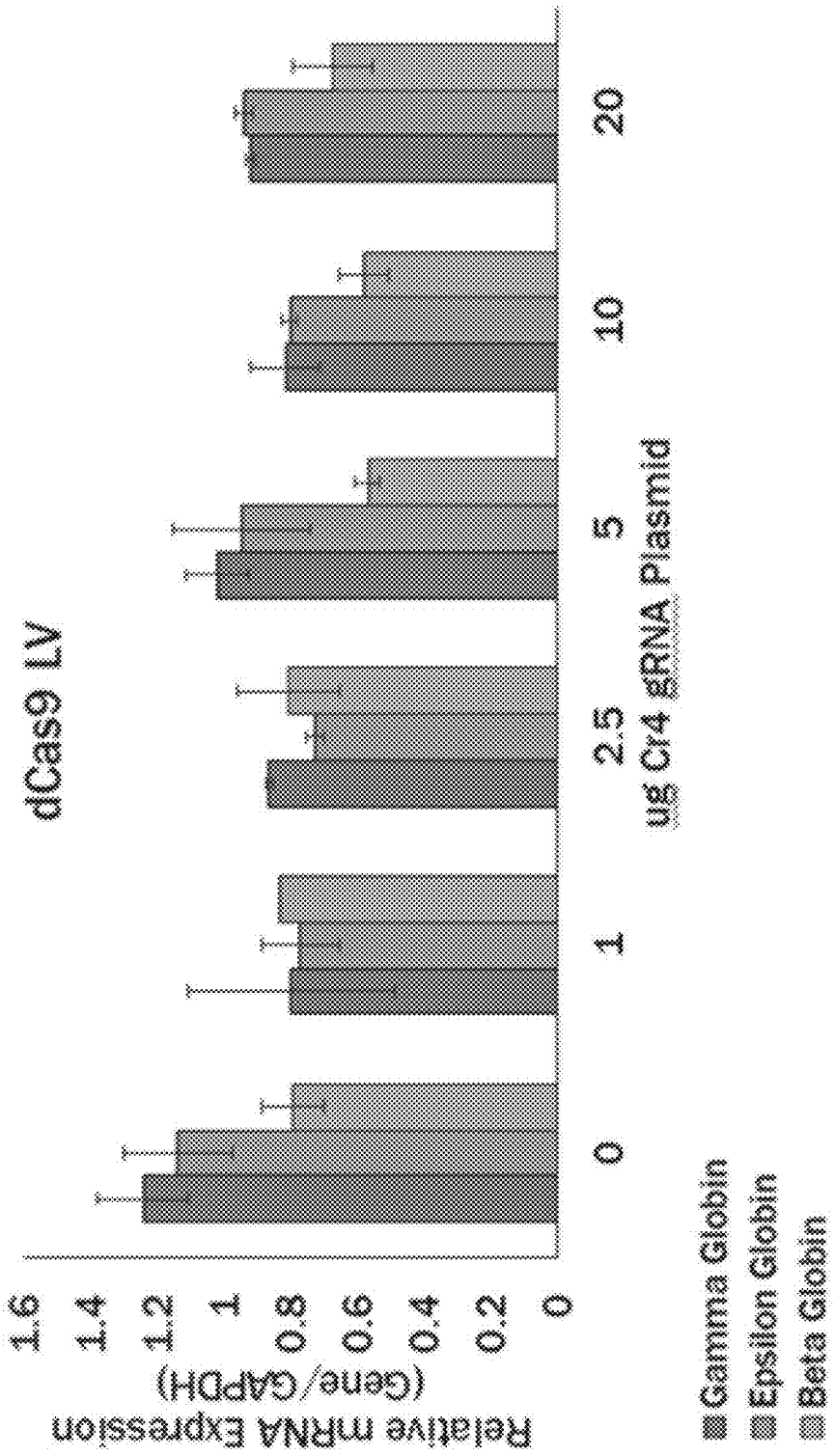

dCas9 LV
Relative mRNA Expression
(Gene/GAPDH)
1.6
1.4
1.2
1
0.8
0.6
0.4
0.2
0
0
1
2.5
5
10
20
ug Cr4 gRNA Plasmid
Gamma Globin
Epsilon Globin
Beta Globin

Fig. 56B

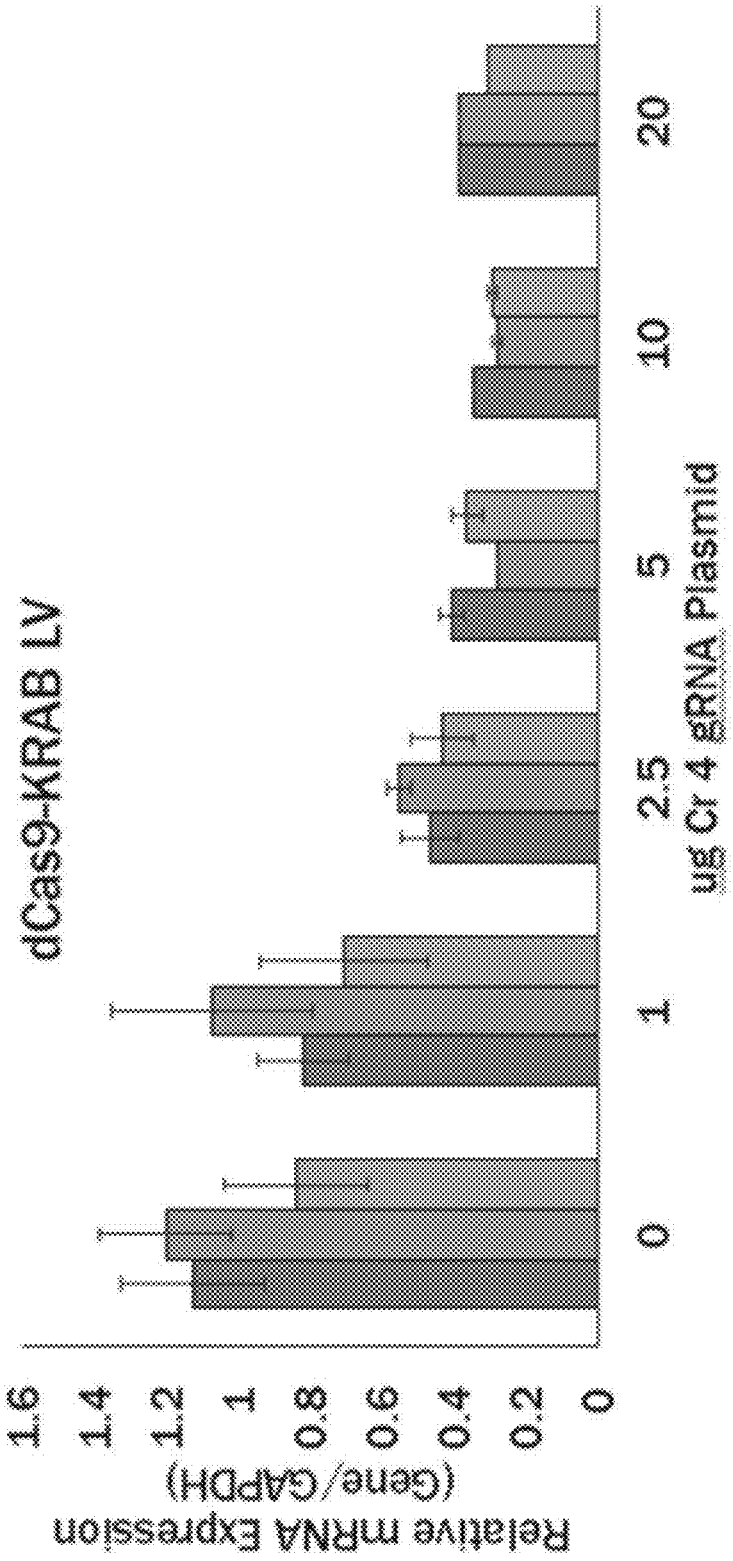

dCas9-KRAB LV
Relative mRNA Expression
(Gene/GAPDH)
1.6
1.4
1.2
1
0.8
0.6
0.4
0.2
0
0
1
2.5
5
10
20
ug Cr 4 gRNA Plasmid

Fig. 56C

Fig. 57A

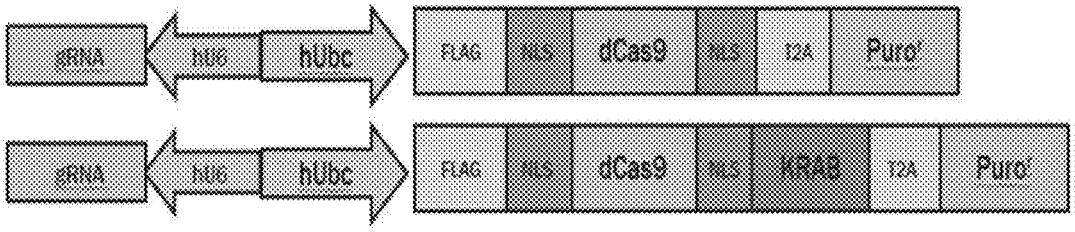


Fig. 57B

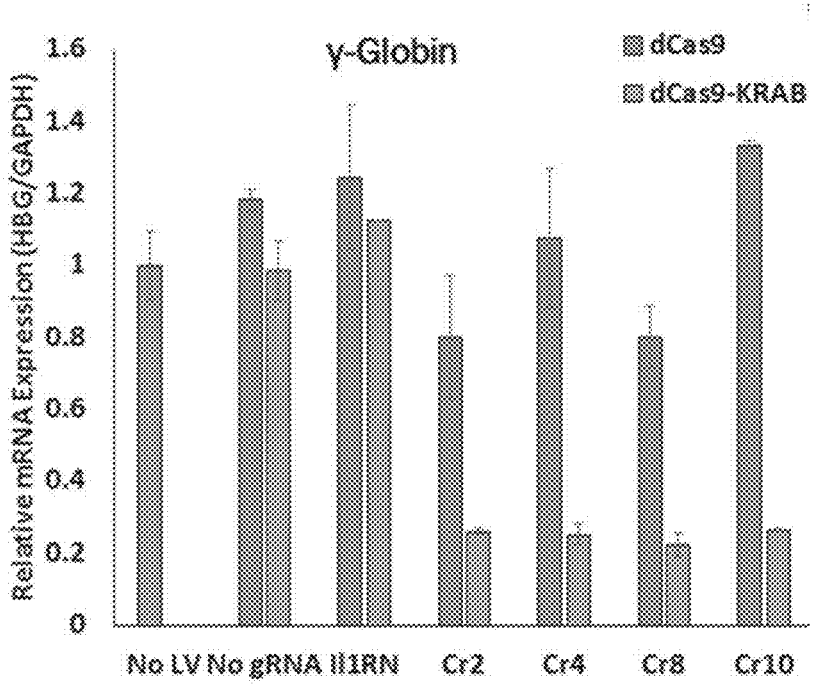


Fig. 57C

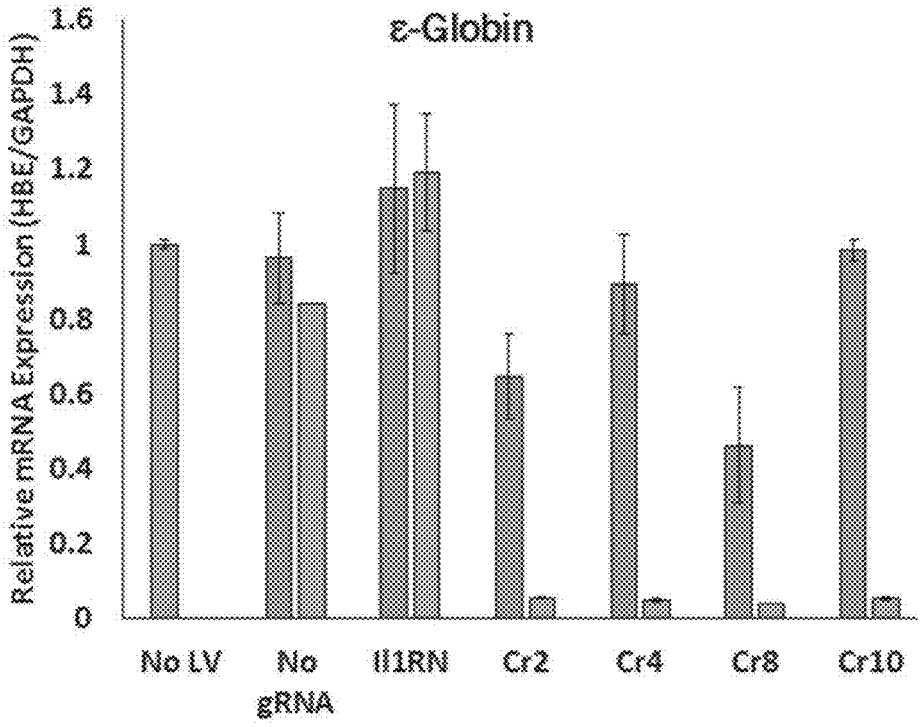
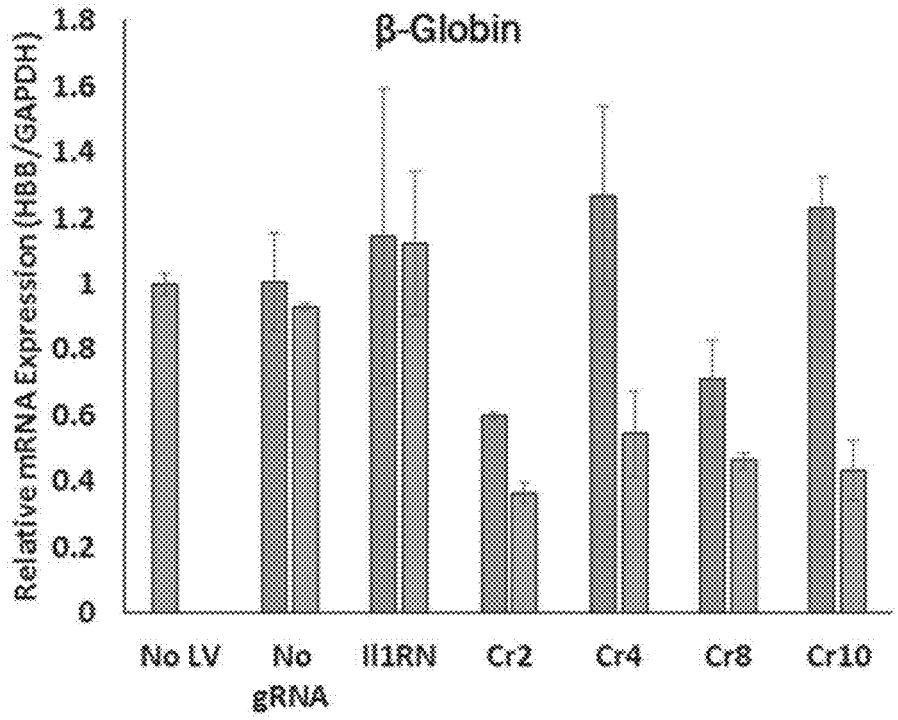


Fig. 57D



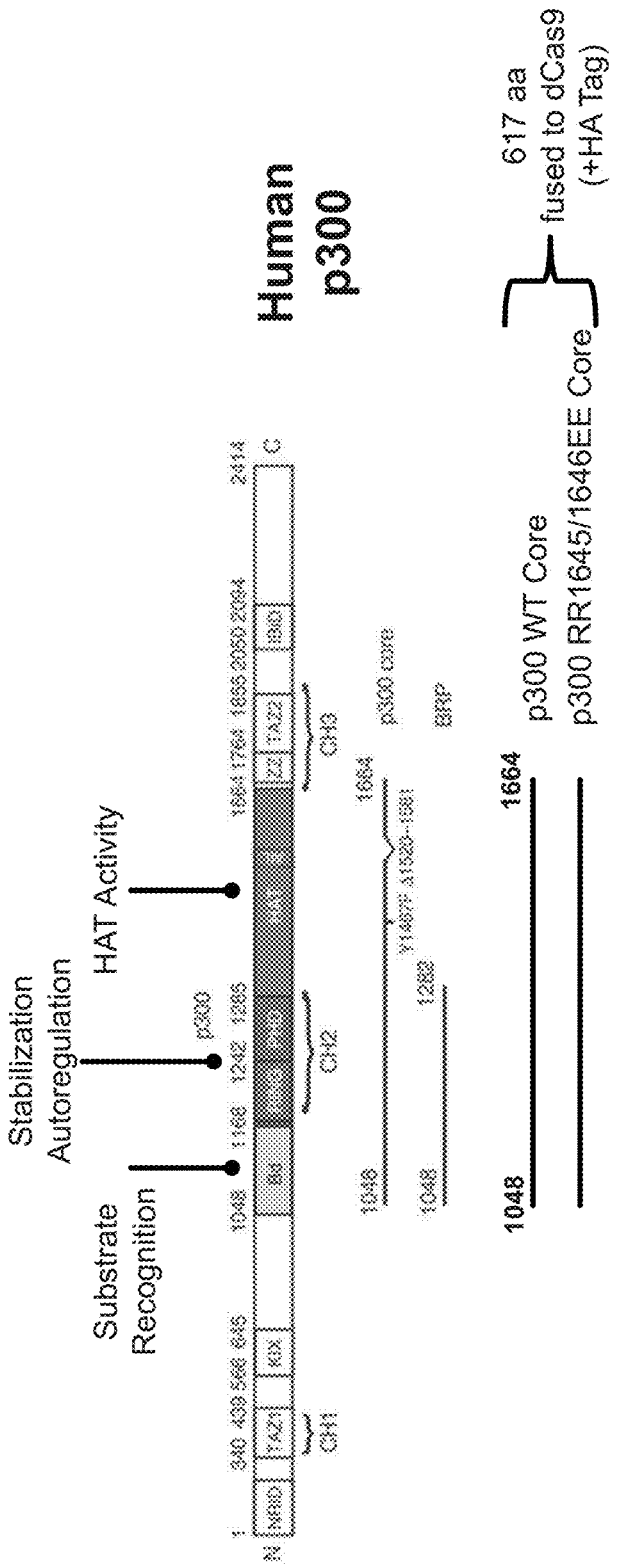

Human
p300
617 aa
fused to dCas9
(+HA Tag)
p300 WT Core
p300 RR1645/1646EE Core
1664
1048
Stabilization
Autoregulation
HAT Activity
Substrate
Recognition
p300
CH1
CH2
CH3
p300 core
BRP
N
C

Fig. 58

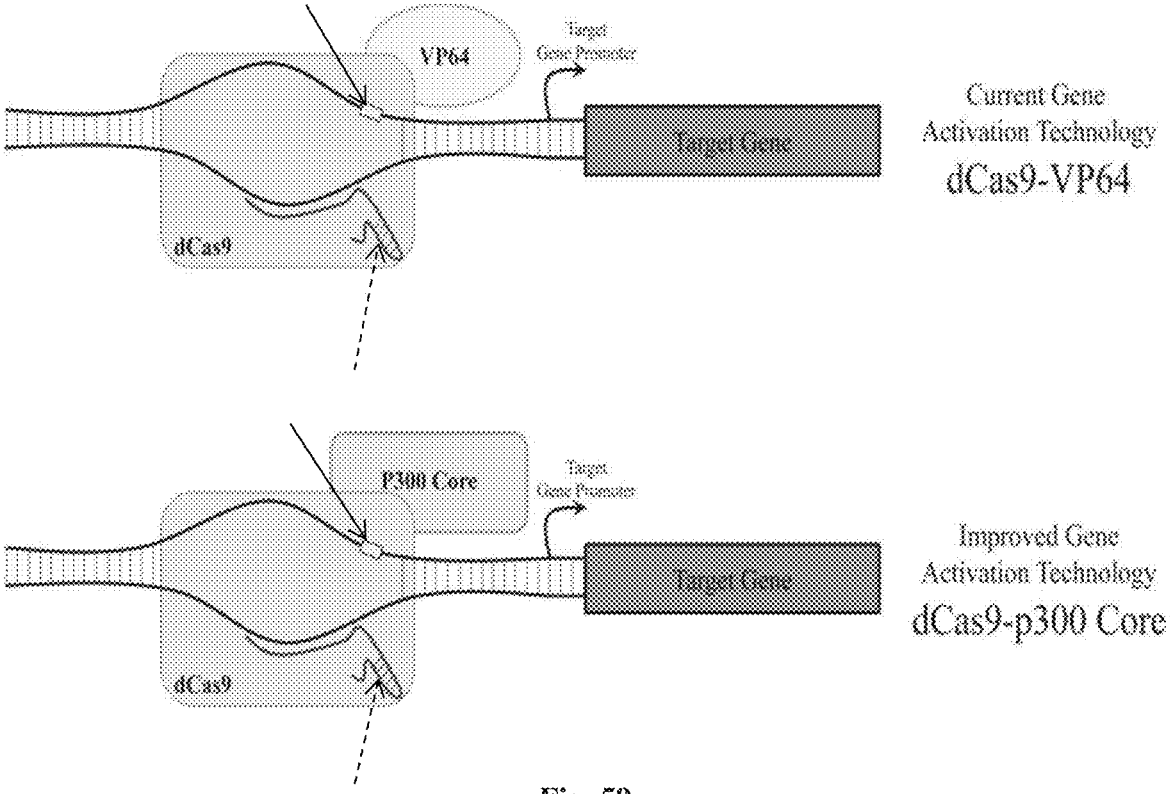


Fig. 59

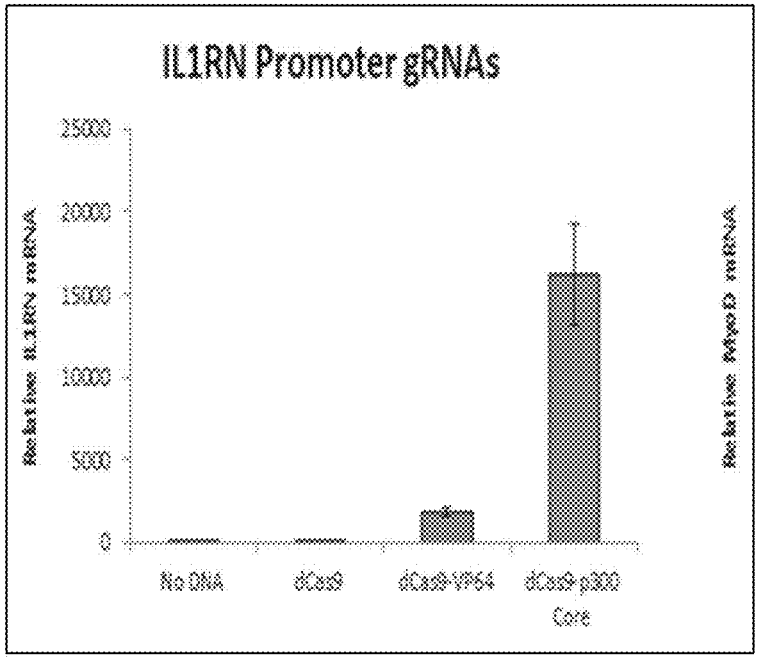


Fig. 60A

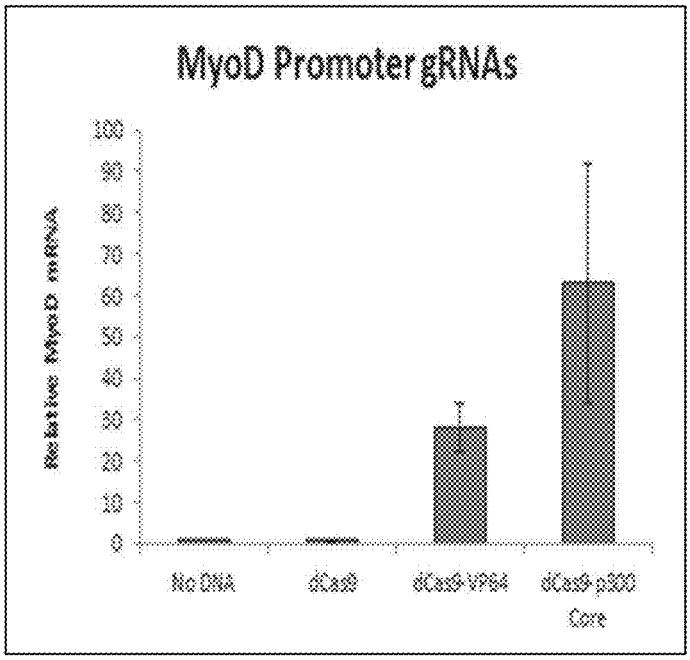


Fig. 60B

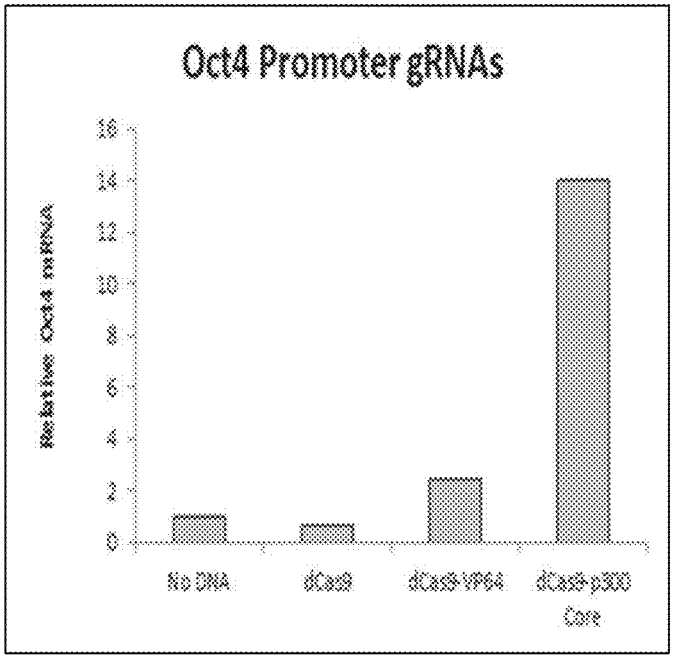


Fig. 60C

i. dCas9

MDYKDHDGDYKDHDIDYKDDDDMAPKKKRKVGRGMDKKYSIGLAIGTNSVGWAVITDEY
KVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEI
FSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDS
TDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASG
VDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQ
LSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRY
DEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDG
TEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILT
FRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNE
KVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLK
EDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTL
FEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKS
DGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVD
ELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQ
LQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRG
KSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETR
QITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAH
DAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNF
FKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGF
SKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLG
ITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNE
LALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADA
NLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDA
TLIHQSITGLYETRIDLSQLGGDPKKKRKVG

Amino Acid Legend:

"Flag" Epitope
Nuclear Localization Sequence
Streptococcus pyogenes Cas9 (D10A, H840A)
VP64 Effector
p300 Core Effector
"HA" Epitope

Fig. 61A

ii. dCas9^{VP64}

MDYKDHDGDYKDHDIDYKDDDDMAPKKKRKVGRGMDKKYSIGLAIGTNSVGWAVITDEY
KVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEI
FSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDS
TDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASG
VDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQ
LSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRY
DEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDG
TEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILT
FRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNE
KVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLK
EDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTL
FEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKS
DGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVD
ELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQ
LQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRG
KSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETR
QITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAH
DAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNF
FKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGF
SKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLG
ITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNE
LALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADA
NLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDA
TLIHQSITGLYETRIDLSQLGGDPIAGSKASPKKKRKVGRADALDDFDLDMLGSDALDDF
DLDMLGSDALDDFDLDMLGSDALDDFDLDMLINYPYDVPDYAS

Fig. 61B

iii. dCas9$^{p300\ Core}$

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRGMDKKYSIGLAIGTNSVGWAVITDEY
KVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEI
FSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDS
TDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASG
VDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQ
LSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRY
DEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDG
TEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILT
FRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNE
KVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLK
EDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTL
FEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKS
DGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVD
ELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQ
LQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDENRG
KSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETR
QITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAH
DAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNF
FKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGF
SKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLG
ITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNE
LALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADA
NLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDA
TLIHQSITGLYETRIDLSQLGGDPIAGSKASPKKKRKVGRAIFKPEELRQALMPTLEALY
RQDPESLPFRQPVDPQLLGIPDYFDIVKSPMDLSTIKRKLDTGQYQEPWQYVDDIWLMFN
NAWLYNRKTSRVYKYCSKLSEVFEQEIDPVMQSLGYCCGRKLEFSPQTLCCYGKQLCTIP
RDATYYSYQNRYHFCEKCFNEIQGESVSLGDDPSQPQTTINKEQFSKRKNDTLDPELFVE
CTECGRKMHQICVLHHEIIWPAGFVCDGCLKKSARTRKENKFSAKRLPSTRLGTFLENRV
NDFLRRQNHPESGEVTVRVVHASDKTVEVKPGMKARFVDSGEMAESFPYRTKALFAFEEI
DGVDLCFFGMHVQEYGSDCPPPNQRRVYISYLDSVHFFRPKCLRTAVYHEILIGYLEYVK
KLGYTTGHIWACPPSEGDDYIFHCHPPDQKIPKPKRLQEWYKKMLDKAVSERIVHDYKDI
FKQATEDRLTSAKELPYFEGDFWPNVLEESIKELEQEEERKREENTSNESTDVTKGDSK
NAKKQNNKKTSKNKSSLSRGNKKKPGMPNVSNDLSQKLYATMEKHKEVFFVIRLIAGPAA
NSLPPIVDPDPLIPCDLMDGRDAFLTLARDKHLEFSSLRRAQWSTMCMLVELHTQSQDYP
YDVPDYAS

Fig. 61C

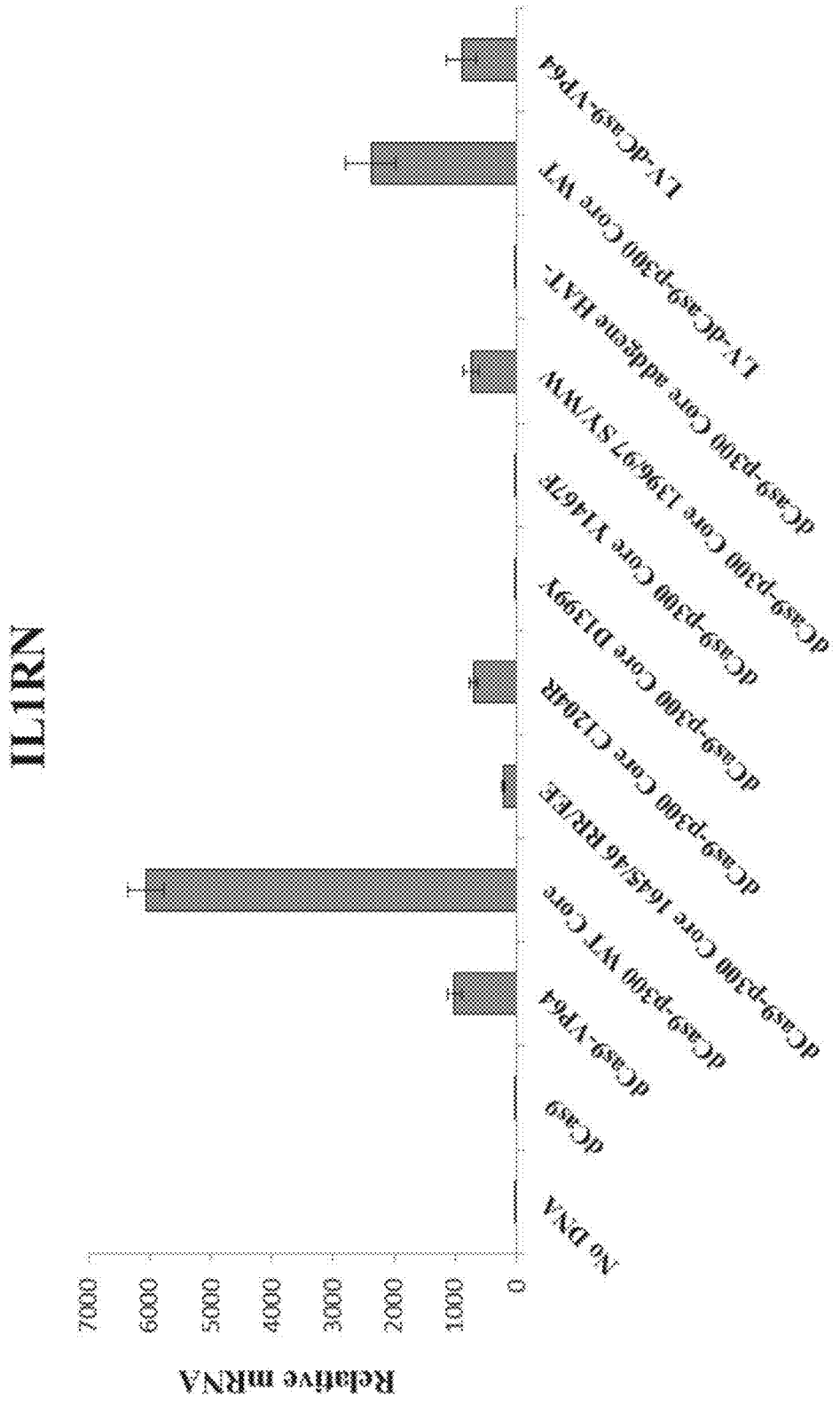

IL1RN
Relative mRNA
7000
6000
5000
4000
3000
2000
1000
0
No DNA
dCas9
dCas9-VP64
dCas9-p300 WT Core
dCas9-p300 Core 1645/46 RRPFE
dCas9-p300 Core C1208R
dCas9-p300 Core D1399Y
dCas9-p300 Core Y1467F
dCas9-p300 Core 1396/97 SV/WW
dCas9-p300 Core additional HAT-
LV-dCas9-p300 Core WT
LV-dCas9-VP64

Fig. 62

Fig. 63A

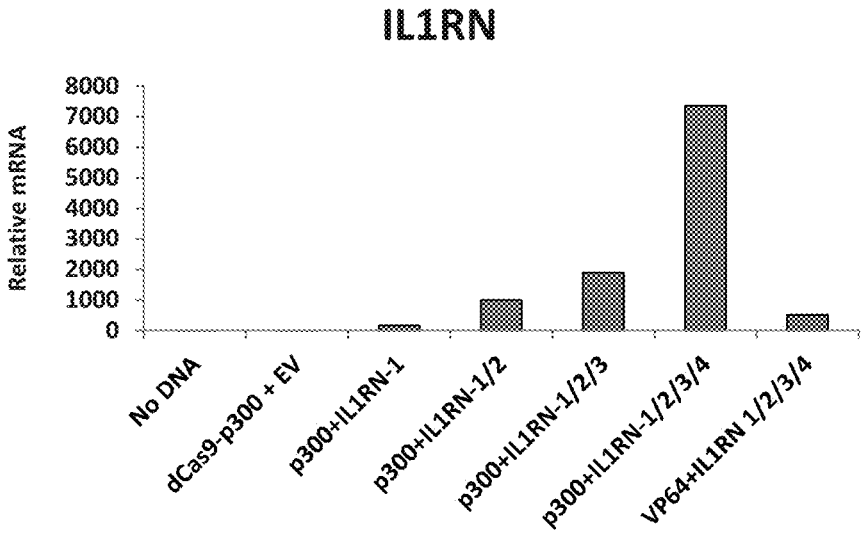


Fig. 63B

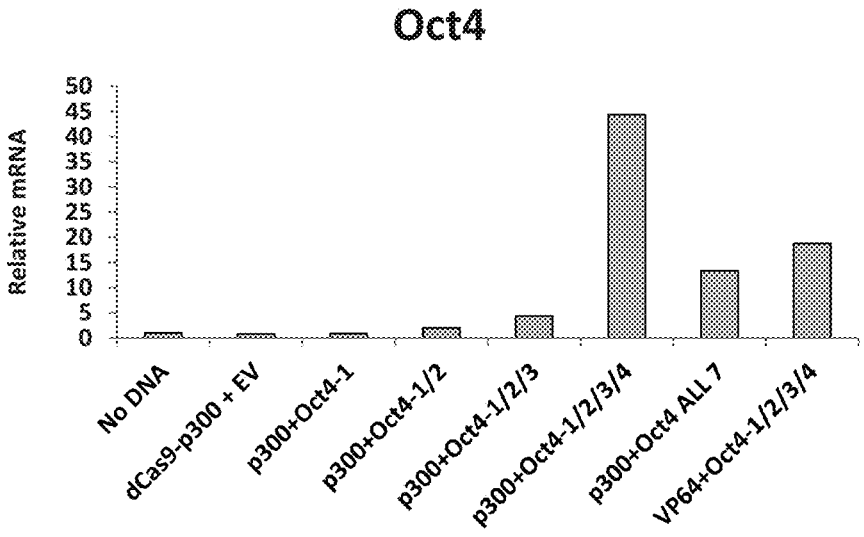


Fig. 64A

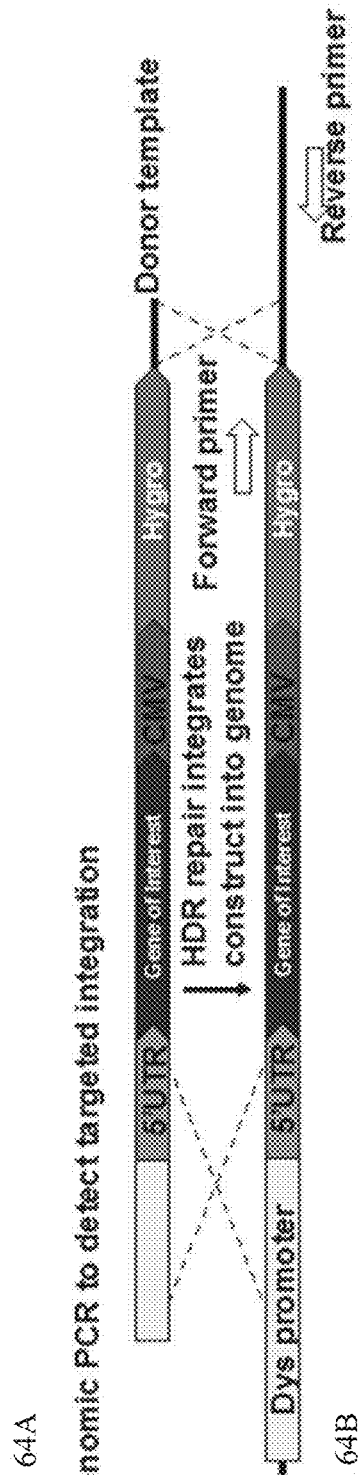


Fig. 64B

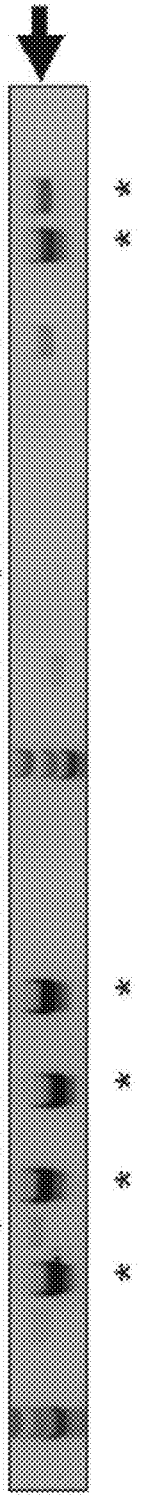
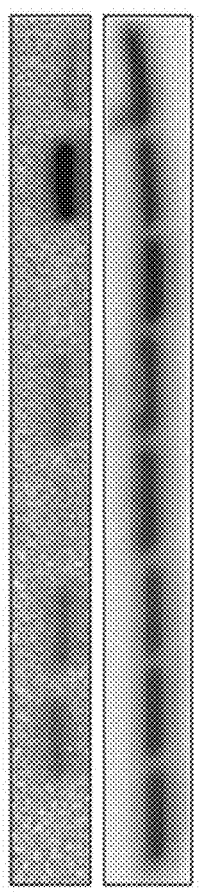


Fig. 64C



RNA-GUIDED GENE EDITING AND GENE REGULATION

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application is a continuation of U.S. patent application Ser. No. 14/895,316, filed Dec. 2, 2015, which is the national stage filing under 35 U.S.C. 371 of International Patent Application No. PCT/US2014/041190, filed Jun. 5, 2014, which claims the benefit of priority to U.S. Provisional Application No. 61/831,481, filed Jun. 5, 2013, U.S. Provisional Application No. 61/839,127, filed Jun. 25, 2013, U.S. Provisional Application No. 61/904,911, filed Nov. 15, 2013, U.S. Provisional Application No. 61/967,466, filed Mar. 19, 2014, and U.S. Provisional Application No. 61/981,575, filed Apr. 18, 2014, the entire contents of each of which are hereby incorporated by reference.

STATEMENT OF GOVERNMENT INTEREST

[0002] This invention was made with government support under federal grant numbers DP2-OD008586 and R01DA036865 awarded by NIH and CBET-1151035 awarded by the National Science Foundation. The U.S. Government has certain rights to this invention.

SEQUENCE LISTING

[0003] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on Feb. 27, 2018, is named 028193-9164-US03_Subt_Seq_List.txt and is 331,614 bytes in size.

TECHNICAL FIELD

[0004] The present disclosure relates to the field of gene expression alteration, genome engineering and genomic alteration of genes using Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR)/CRISPR-associated (Cas) 9-based systems and viral delivery systems. The present disclosure also relates to the field of genome engineering and genomic alteration of genes in muscle, such as skeletal muscle and cardiac muscle.

BACKGROUND

[0005] Synthetic transcription factors have been engineered to control gene expression for many different medical and scientific applications in mammalian systems, including stimulating tissue regeneration, drug screening, compensating for genetic defects, activating silenced tumor suppressors, controlling stem cell differentiation, performing genetic screens, and creating synthetic gene circuits. These transcription factors can target promoters or enhancers of endogenous genes, or be purposefully designed to recognize sequences orthogonal to mammalian genomes for transgene regulation. The most common strategies for engineering novel transcription factors targeted to user-defined sequences have been based on the programmable DNA-binding domains of zinc finger proteins and transcription-activator like effectors (TALEs). Both of these approaches involve applying the principles of protein-DNA interactions of these domains to engineer new proteins with unique DNA-binding specificity. Although these methods have been widely successful for many applications, the protein engineering necessary for manipulating protein-DNA interactions can be laborious and require specialized expertise.

[0006] Additionally, these new proteins are not always effective. The reasons for this are not yet known but may be related to the effects of epigenetic modifications and chromatin state on protein binding to the genomic target site. In addition, there are challenges in ensuring that these new proteins, as well as other components, are delivered to each cell. Existing methods for delivering these new proteins and their multiple components include delivery to cells on separate plasmids or vectors which leads to highly variable expression levels in each cell due to differences in copy number. Additionally, gene activation following transfection is transient due to dilution of plasmid DNA, and temporary gene expression may not be sufficient for inducing therapeutic effects. Furthermore, this approach is not amenable to cell types that are not easily transfected. Thus another limitation of these new proteins is the potency of transcriptional activation.

[0007] Site-specific nucleases can be used to introduce site-specific double strand breaks at targeted genomic loci. This DNA cleavage stimulates the natural DNA-repair machinery, leading to one of two possible repair pathways. In the absence of a donor template, the break will be repaired by non-homologous end joining (NHEJ), an error-prone repair pathway that leads to small insertions or deletions of DNA. This method can be used to intentionally disrupt, delete, or alter the reading frame of targeted gene sequences. However, if a donor template is provided along with the nucleases, then the cellular machinery will repair the break by homologous recombination, which is enhanced several orders of magnitude in the presence of DNA cleavage. This method can be used to introduce specific changes in the DNA sequence at target sites. Engineered nucleases have been used for gene editing in a variety of human stem cells and cell lines, and for gene editing in the mouse liver. However, the major hurdle for implementation of these technologies is delivery to particular tissues in vivo in a way that is effective, efficient, and facilitates successful genome modification.

[0008] Hereditary genetic diseases have devastating effects on children in the United States. These diseases currently have no cure and can only be managed by attempts to alleviate the symptoms. For decades, the field of gene therapy has promised a cure to these diseases. However technical hurdles regarding the safe and efficient delivery of therapeutic genes to cells and patients have limited this approach. Duchenne Muscular Dystrophy (DMD) is the most common hereditary monogenic disease and occurs in 1 in 3500 males. DMD is the result of inherited or spontaneous mutations in the dystrophin gene. Dystrophin is a key component of a protein complex that is responsible for regulating muscle cell integrity and function. DMD patients typically lose the ability to physically support themselves during childhood, become progressively weaker during the teenage years, and die in their twenties. Current experimental gene therapy strategies for DMD require repeated administration of transient gene delivery vehicles or rely on permanent integration of foreign genetic material into the genomic DNA. Both of these methods have serious safety concerns. Furthermore, these strategies have been limited by an inability to deliver the large and complex dystrophin gene sequence.

SUMMARY

[0009] The present invention is directed to a fusion protein comprising two heterologous polypeptide domains. The first polypeptide domain comprises a Clustered Regularly Interspaced Short Palindromic Repeats associated (Cas) protein and the second polypeptide domain has an activity selected from the group consisting of transcription activation activity, transcription repression activity, transcription release factor activity, histone modification activity, nuclease activity, nucleic acid association activity, methylase activity, and demethylase activity. The Cas protein may comprise Cas9. The Cas9 may comprise at least one amino acid mutation which knocks out nuclease activity of Cas9. The at least one amino acid mutation may be at least one of D10A and H840A. The Cas protein may comprise iCas9 (amino acids 36-1403 of SEQ ID NO: 1). The second polypeptide domain may have transcription activation activity. The second polypeptide domain may comprise at least one VP16 transcription activation domain repeat. The second polypeptide domain may comprise a VP16 tetramer ("VP64") or a p65 activation domain. The fusion protein may further comprise a linker connecting the first polypeptide domain to the second polypeptide domain. The fusion protein may comprise iCas9-VP64.

[0010] The present invention is directed to a DNA targeting system comprising said fusion protein and at least one guide RNA (gRNA). The at least one gRNA may comprise a 12-22 base pair complementary polynucleotide sequence of the target DNA sequence followed by a protospacer-adjacent motif. The at least one gRNA may target a promoter region of a gene, an enhancer region of a gene, or a transcribed region of a gene. The at least one gRNA may target an intron of a gene. The at least one gRNA may target an exon of a gene. The at least one gRNA may target a the promoter region of a gene selected from the group consisting of ASCL1, BRN2, MYTIL, NANOG, VEGFA, TERT, IL1B, IL1R2, IL1RN, HBG1, HBG2, and MYOD1. The at least one gRNA may comprise at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625.

[0011] The present invention is directed to a DNA targeting system that binds to a dystrophin gene comprising Cas9 and at least one guide RNA (gRNA). The at least one gRNA may target an intron of the dystrophin gene. The at least one gRNA may target an exon of the dystrophin gene. The at least one guide RNA may comprise at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625. The DNA targeting system may comprise between one and ten different gRNAs.

[0012] The present invention is directed to an isolated polynucleotide encoding said fusion protein or said DNA targeting system.

[0013] The present invention is directed to a vector comprising said isolated polynucleotide.

[0014] The present invention is directed to a cell comprising said isolated polynucleotide or said vector.

[0015] The present invention is directed to a method of modulating mammalian gene expression in a cell. The method comprises contacting the cell with said fusion protein, said DNA targeting system, said isolated polynucleotide, or said vector. The gene expression may be induced.

[0016] The present invention is directed to a method of transdifferentiating or inducing differentiation of a cell. The method comprises contacting the cell with said fusion protein, said DNA targeting system, said isolated polynucleotide, or said vector. The cell may be a fibroblast cell or an induced pluripotent stem cells. The fibroblast cell may be trandifferentiated into a neuronal cell or a myogenic cell. The DNA targeting system may be contacted with the cell and at least one gRNA targets a promoter region of at least one gene selected from the group consisting of ASCL1, BRN2, MYOD1, and MYT1L. The DNA targeting system may comprise at least one gRNA that targets the promoter region of the ASCL1 gene and at least one gRNA that targets the promoter region of the BRN2 gene. The DNA targeting system may comprise between one and twenty different gRNAs. The DNA targeting system may comprise 8 or 16 different gRNAs. The DNA targeting system may comprise dCas9-VP64. The DNA targeting system may be delivered to the cell virally or non-virally.

[0017] The present invention is directed to a method of correcting a mutant gene in a cell. The method comprises administering to a cell containing said DNA targeting system, said isolated polynucleotide, or said vector. The correction of the mutant gene may comprise homology-directed repair. The method may further comprise administering to the cell a donor DNA. The mutant gene may comprise a frameshift mutation which causes a premature stop codon and a truncated gene product. The correction of the mutant gene may comprise nuclease mediated non-homologous end joining. The correction of the mutant gene may comprise a deletion of a premature stop codon, a disruption of a splice acceptor site, a deletion of one or more exons, or disruption of a splice donor sequence. The deletion of one or more exons may result in the correction of the reading frame.

[0018] The present invention is directed to a method of treating a subject in need thereof having a mutant dystrophin gene. The method comprises administering to the subject said DNA targeting system, said isolated polynucleotide, or said vector. The subject may be suffering from Duchenne muscular dystrophy.

[0019] The present invention is directed to a method of correcting a mutant dystrophin gene in a cell. The method comprises administering to a cell containing a mutant dystrophin gene said DNA targeting system, said isolated polynucleotide, said vector, or said cell. The mutant dystrophin gene may comprise a premature stop codon, disrupted reading frame via gene deletion, an aberrant splice acceptor site, or an aberrant splice donor site, and wherein the target region is upstream or downstream of the premature stop codon, disrupted reading frame, aberrant splice acceptor site, or the aberrant splice donor site. The correction of the mutant dystrophin gene may comprise homology-directed repair. The method may further comprise administering to the cell a donor DNA. The mutant dystrophin gene may comprise a frameshift mutation which causes a premature stop codon and a truncated gene product. The correction of the mutant dystrophin gene may comprise nuclease mediated non-homologous end joining. The correction of the mutant dystrophin gene may comprise a deletion of a premature stop codon, correction of a disrupted reading frame, or modulation of splicing by disruption of a splice acceptor site or disruption of a splice donor sequence. The correction of the mutant dystrophin gene may comprise a deletion of exons 45-55 or exon 51.

[0020] The present invention is directed to a kit comprising said fusion protein, said DNA targeting system, said isolated polynucleotide, said vector, or said cell.

[0021] The present invention is directed to a method of modulating mammalian gene expression in a cell. The method comprises contacting the cell with a polynucleotide encoding a DNA targeting system. The DNA targeting system comprises said fusion protein and at least one guide RNA (gRNA). The DNA targeting system may comprise between one and ten different gRNAs. The different gRNAs may bind to different target regions within the target gene. The target regions may be separated by at least one nucleotide. The target regions may be separated by about 15 to about 700 base pairs. Each of the different gRNAs may bind to at least one different target genes. The different target genes may be located on same chromosome. The different target genes may be located on different chromosomes. The at least one target region may be within a non-open chromatin region, an open chromatin region, a promoter region of the target gene, an enhancer region of the target gene, a transcribed region of the target gene, or a region upstream of a transcription start site of the target gene. The at least one target region may be located between about 1 to about 1000 base pairs upstream of a transcription start site of a target gene. The at least one target region may be located between about 1 to about 600 base pairs upstream of a transcription start site of a target gene. The gene expression may be induced. The DNA targeting system may comprise two different gRNAs, three different gRNAs, four different gRNAs, five different gRNAs, six different gRNAs, seven different gRNAs, eight different gRNAs, nine different gRNAs, or ten different gRNAs. The at least one guide RNA may target a promoter region of a gene selected from the group consisting of ASCL1, BRN2, MYT1L, NANOG, VEGFA, TERT, IL1B, IL1R2, IL1RN, HBG1, HBG2, and MYOD1. The at least one guide RNA may comprise at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625. The at least one target region may be within an intron or an exon of a target gene.

[0022] The present invention is directed to a composition for inducing mammalian gene expression in a cell. The composition comprises said fusion protein and at least one guide RNA (gRNA).

[0023] The present invention is directed to a composition for inducing mammalian gene expression in a cell. The composition comprises an isolated polynucleotide sequence encoding said fusion protein and at least one guide RNA (gRNA). The at least one guide RNA may target a promoter region of a gene selected from the group consisting of ASCL1, BRN2, MYT1L, NANOG, VEGFA, TERT, IL1B, IL1R2, IL1RN, HBG1, HBG2, and MYOD1. The at least one guide RNA may comprise at least one of SEQ ID NOs 5-40, 65-144, 492-515, 540-563, and 585-625.

[0024] The present invention is directed to a cell comprising said composition for inducing mammalian gene expression in a cell.

[0025] The present invention is directed to a kit comprising said composition for inducing mammalian gene expression in a cell or said cell comprising said composition for inducing mammalian gene expression in a cell.

[0026] The present invention is directed to a kit for inducing mammalian gene expression in a cell. The kit comprises said composition for inducing mammalian gene expression in a cell or said cell comprising said composition for inducing mammalian gene expression in a cell.

[0027] The present invention is directed to a composition for genome editing in a muscle of a subject. The composition comprises a modified adeno-associated virus (AAV) vector and a nucleotide sequence encoding a site-specific nuclease. The muscle is skeletal muscle or cardiac muscle. The modified AAV vector may have enhanced cardiac and skeletal muscle tissue tropism. The site-specific nuclease may comprise a zinc finger nuclease, a TAL effector nuclease, or a CRISPR/Cas9 system. The site-specific nuclease may bind a gene or locus in the cell of the muscle. The gene or locus may be dystrophin gene. The composition may further comprise a donor DNA or transgene.

[0028] The present invention is directed to a kit comprising said composition for genome editing in a muscle of a subject.

[0029] The present invention is directed to a method of genome editing in a muscle of a subject. The method comprises administering to the muscle said composition for genome editing in a muscle of a subject, wherein the muscle is skeletal muscle or cardiac muscle. The genome editing may comprise correcting a mutant gene or inserting a transgene. Correcting a mutant gene may comprise deleting, rearranging, or replacing the mutant gene. Correcting the mutant gene may comprise nuclease-mediated non-homologous end joining or homology-directed repair.

[0030] The present invention is directed to a method of treating a subject. The method comprises administering said composition for genome editing in a muscle of a subject to a muscle of the subject, wherein the muscle is skeletal muscle or cardiac muscle. The subject may be suffering from a skeletal muscle condition or a genetic disease. The subject may be suffering from Duchenne muscular dystrophy.

[0031] The present invention is directed to a method of correcting a mutant gene in a subject, the method comprises administering said composition for genome editing in a muscle of a subject. The muscle is skeletal muscle or cardiac muscle. The composition may be injected into the skeletal muscle of the subject. The composition may be injected systemically to the subject. The skeletal muscle may be tibialis anterior muscle.

[0032] The present invention is directed to a modified lentiviral vector for genome editing in a subject comprising a first polynucleotide sequence encoding said fusion protein and a second polynucleotide sequence encoding at least one sgRNA. The first polynucleotide sequence may be operably linked to a first promoter. The first promoter may be a constitutive promoter, an inducible promoter, a repressible promoter, or a regulatable promoter. The second polynucleotide sequence may encode between one and ten different sgRNAs. The second polynucleotide sequence may encode two different sgRNAs, three different sgRNAs, four different sgRNAs, five different sgRNAs, six different sgRNAs, seven different sgRNAs, eight different sgRNAs, nine different sgRNAs, or ten different sgRNAs. Each of the polynucleotide sequences encoding the different sgRNAs may be operably linked to a promoter. Each of the promoters operably linked to the different sgRNAs may be the same promoter. Each of the promoters operably linked to the different sgRNAs may be different promoters. The promoter may be a constitutive promoter, an inducible promoter, a repressible promoter, or a regulatable promoter. The sgRNA may bind to a target gene. Each of the sgRNA may bind to a different target region within one target loci. Each of the sgRNA may bind to a different target region within different gene loci. The fusion protein may comprise Cas9 protein or iCas9-VP64 protein. The fusion protein may comprise a

VP64 domain, a p300 domain, or a KRAB domain. The two or more endogenous genes may be transcriptionally activated. The two or more endogenous genes may be repressed.

[0033] The present invention is directed to a method of activating an endogenous gene in a cell. The method comprises contacting a cell with said modified lentiviral vector. The endogenous gene may be transiently activated. The endogenous gene may be stably activated. The endogenous gene may be transiently repressed. The endogenous gene may be stably repressed. The fusion protein may be expressed at similar levels to the sgRNAs. The fusion protein may be expressed at different levels to the sgRNAs. The cell may be a primary human cell.

[0034] The present invention is directed to a method of multiplex gene editing in a cell. The method comprises contacting a cell with said modified lentiviral vector. The multiplex gene editing may comprise correcting at least one mutant gene or inserting a transgene. Correcting a mutant gene may comprise deleting, rearranging, or replacing the at least one mutant gene. Correcting the at least one mutant gene may comprise nuclease-mediated non-homologous end joining or homology-directed repair. The multiplex gene editing may comprise deleting at least one gene, wherein the gene is an endogenous normal gene or a mutant gene. The multiplex gene editing may comprise deleting at least two genes. The multiplex gene editing may comprise deleting between two and ten genes.

[0035] The present invention is directed to a method of modulating gene expression of at least one target gene in a cell. The method comprises contacting a cell with said modified lentiviral vector. The gene expression of at least two genes may be modulated. The gene expression of between two genes and ten genes may be modulated. The gene expression of the at least one target gene may be modulated when gene expression levels of the at least one target gene are increased or decreased compared to normal gene expression levels for the at least one target gene.

BRIEF DESCRIPTION OF THE DRAWINGS

[0036] FIGS. **1**A-**1**F show RNA-guided activation of the human IL1RN gene by iCas9-VP64. (FIG. **1**A, FIG. **1**B) An RNA-guided transcriptional activator was created by fusing the inactivated Cas9 (iCas9, D10A/H840A) to the VP64 transactivation domain. iCas9-VP64 recognizes genomic target sites through the hybridization of a guide RNA (gRNA) to a 20 bp target sequence. (FIG. **1**C) Expression plasmids for four gRNAs or crRNA/tracrRNAs targeted to sequences in the IL1RN promoter were co-transfected with the iCas9-VP64 expression plasmid into HEK293T cells. Activation of IL1RN expression was assessed by qRT-PCR. (FIG. **1**D) The four gRNA expression plasmids were co-transfected with iCas9-VP64 individually or in combination. Robust gene activation was observed by qRT-PCR only in response to the combination of gRNAs. (FIG. **1**E) Activation of IL1RN expression was confirmed by assessing secretion of the IL-1ra gene product into the media by ELISA. IL-1ra was only detected in three of the six samples treated with the combination of gRNAs. For FIGS. **1**C-**1**E, data are shown as the mean±s.e.m. (n=3 independent experiments). Treatment with the combination of gRNAs was statistically different than all other treatments (*P≤0.02) by Tukey's test. (FIG. **1**F) RNA-seq was performed on samples treated with empty expression vector (n=2) or co-transfected with the expression plasmids for iCas9-VP64 and the four gRNAs targeting IL1RN (n=2). The only statistically significant changes in gene expression between these treatments were an increase in the four IL1RN isoforms (false discovery rate $\leq 3\times10-4$) and a decrease in IL32 (false discovery rate=0.03).

[0037] FIGS. **2**A-**2**H show RNA-guided activation of human genes relevant to cell and gene therapy, genetic reprogramming, and regenerative medicine. HEK293T cells were transfected with the iCas9-VP64 expression plasmid and four gRNAs individually or in combination. Target gene expression was measured by qRT-PCR and normalized to GAPDH mRNA levels. Data are shown as the mean±s.e.m. (n=3 independent experiments). Treatment with the combination of gRNAs was statistically different than all other treatments (*P<0.05) by Tukey's test.

[0038] FIG. **3** shows expression of iCas9-VP64. Expression of iCas9-VP64 in transfected HEK293 cells was confirmed by western blot for the N-terminal Flag epitope tag. The wt Cas9 expression plasmid does not contain the epitope tag.

[0039] FIG. **4** shows positions of gRNA target sites and DNAse hypersensitivity of human target genes. The four gRNA target sites for each locus are designated as custom tracks above each gene and DNase-seq data indicating DNAse-hypersensitive open chromatin regions is shown below each gene. DNase-seq was performed in HEK293T cells to identify DNase hypersensitive regions, as previously described (Song et al., Cold Spring Harbor protocols 2010, pdb prot5384 (2010); Song et al. *Genome Res* 21, 1757-1767 (2011)). The results show that open chromatin was not a requirement for gene activation by combinations of gRNAs with iCas9-VP64.

[0040] FIG. **5** shows the absence of nuclease activity by iCas9-VP64. Wild-type Cas9 or inactivated (D10A, H840A) iCas9-VP64 expression plasmids were co-transfected with expression plasmids for four different guide RNAs targeting the IL1RN promoter. Nuclease activity was determined by the Surveyor assay (Guschin et al., *Methods Mol Biol* 649, 247-256 (2010)). The lower molecular weight bands indicative of nuclease activity and DNA repair by non-homologous end joining are only present following treatment with wild-type Cas9, supporting abrogation of nuclease activity by iCas9-VP64.

[0041] FIG. **6** shows RNA-seq for samples treated with gRNAs targeting HBG1 and HBG2. RNA-seq was performed on samples treated with a control empty expression vector (n=3) or cotransfected with the expression plasmids for iCas9-VP64 and the four gRNAs targeting HBG1 (n=2). Three of these gRNAs also target HBG2. Increases in both HBG1 and HBG2 relative to control were observed but were not statistically significant due to low expression levels. The only statistically significant changes in gene expression between these treatments were decreases in IL32 (false discovery rate=0.0007) and TNFRS9 (false discovery rate=0.002).

[0042] FIG. **7** shows upregulation of Ascl1 and γ-globin by iCas9-VP64. HEK293T cells were transfected with iCas9-VP64 and four gRNAs targeting the ASCL1 or HBG1 promoter. Levels of corresponding Ascl1 and γ-globin protein production were assessed by western blot. Low levels of these proteins were detectable in HEK293T cells and increases in expression were detectable following iCas9-VP64 treatment in two independent experiments.

[0043] FIGS. **8**A-**8**H show activation of downstream targets of Ascl1 in iCas9-VP64-treated murine embryonic

fibroblasts. Mouse embryonic fibroblasts (MEFs) were transfected with a control GFP expression plasmid or the iCas9-VP64 expression plasmid and a combination of four gRNA expression plasmids targeting ASCL1 at a ratio of 50:50 or 75:25. (FIG. **8**A) The gRNA target sites in the human ASCL1 promoter (SEQ ID NO: 3) are conserved in the mouse ASCL1 promoter (SEQ ID NO: 4). Target sites are indicated by solid lines and the transcribed region is indicated by dashed line. (FIG. **8**B) ASCL1 expression in MEFs increased at two days after iCas9-VP64/gRNA treatment as determined by qRT-PCR. (FIGS. **8**C-**8**H) After 10 days in neural induction media, cells were stained for Ascl1 and Tuj1, an early marker of neuronal differentiation (FIGS. **8**C-**8**D), or for Tuj1 and MAP2, a marker of more mature neuronal differentiation (FIGS. **8**D-**8**F). Some Tuj1-positive cells adopted neuronal morphologies (FIGS. **8**F-**8**G) and a single cell was found to be positive for Tuj1 and MAP2 (FIG. **8**G). (FIG. **8**H) Tuj1-positive cells were readily identified in the iCas9-VP64/gRNA-treated cultures (~0.05%) but were absent in controls. n=3 independent samples and data are represented as mean standard error of the mean. gRNA 75/25 is significantly different than gRNA 50/50 and control (*P<0.01, Tukey's test).

[0044] FIG. **9**A shows the iCas9-VP64 protein sequence (SEQ ID NO: 1) and FIG. **9**B shows the sequence of the gRNA expression cassette with U6 promoter (SEQ ID NO: 2).

[0045] FIG. **10** shows the standard curves for qRT-PCR. For each gene, the experimental sample with the highest expression level was diluted to create a standard curve that was assayed by qRT-PCR to ensure efficient amplification over an appropriate dynamic range. The efficiencies of all amplification reactions were within 90-115%.

[0046] FIGS. **11**A-**11**C show the validation of RNA-guided repair. FIG. **11**A shows the Surveyor assay results of genomic DNA harvested from HEK 293T cells two days after Cas9 was co-transfected into the cells with empty vector (negative control) or gRNA. FIG. **11**B shows the location of the gRNA target. FIG. **11**C shows the expected cleavage sizes for each gRNA.

[0047] FIG. **12** shows RNA-guided repair in DMD 8036 (del48-50) cells as shown by Surveyor assay.

[0048] FIG. **13** shows RNA-guided repair in DMD 8036 (del48-50) cells as shown by PCR across the entire locus. The PCR of a wild-type dystrophin gene generates a fragment of 1447 bp in size, whereas PCR of the mutant gene in the DMD 8036 cell line shows a deletion of approximately 817 bp. The deletion band after introduction of the CRISPR/Cas9-based system was approximately 630 bp.

[0049] FIG. **14** shows RNA-guided repair in DMD 8036 (del48-50) cells as shown by Western blot with MANDYS8 (anti-dystrophin antibody) and GAPDH antibody (positive control).

[0050] FIG. **15** shows ChIP sequencing data illustrating the specific binding of iCas9-VP64 targeting the IL1RN promoter. HEK 293T cells were transfected with iCas9-VP64 targeting the IL1RN promoter.

[0051] FIGS. **16**A-**16**D CRISPR/Cas9 targeting of the dystrophin gene. (FIG. **16**A) sgRNA sequences were designed to bind sequences in the exon 45-55 mutational hotspot region of the dystrophin gene, such that gene editing could restore dystrophin expression from a wide variety of patient-specific mutations. Arrows within introns indicate sgRNA targets designed to delete entire exons from the genome. Arrows within exons indicate sgRNA targets designed to create targeted frameshifts in the dystrophin gene. (FIG. **16**B) Example of frame correction following introduction of small insertions or deletions by NHEJ DNA repair in exon 51 using the CR3 sgRNA. FIG. **16**B discloses SEQ ID NOS 633-636, respectively, in order of appearance. (FIG. **16**C) Schematic of multiplex sgRNA targets designed to delete exon 51 and restore the dystrophin reading frame in a patient mutation with the deletion of exons 48-50. (FIG. **16**D) Schematic of multiplex sgRNA targets designed to delete the entire exon 45-55 region to address a variety of DMD patient mutations.

[0052] FIG. **17** shows images of TBE-PAGE gels used to quantify Surveyor assay results to measure day 3 gene modification in Table 7. Asterisks mark expected sizes of bands indicative of nuclease activity.

[0053] FIG. **18** shows images of TBE-PAGE gels used to quantify Surveyor assay results to measure day 10 gene modification in Table 7. Asterisks mark expected sizes of bands indicative of nuclease activity.

[0054] FIGS. **19**A-**19**D show fluorescence-activated flow sorting to enrich genetically modified DMD myoblasts. (FIG. **19**A) A plasmid expressing a human-codon optimized SpCas9 protein linked to a GFP marker using a T2A ribosomal skipping peptide sequence was co-electroporated into human DMD myoblasts with one or two plasmids carrying sgRNA expression cassettes. (FIG. **19**B) The indicated sgRNA expression cassettes were independently co-transfected into HEK293 Ts with a separate plasmid expressing SpCas9 with (bottom) or without (top) a GFP marker linked to SpCas9 by a T2A ribosomal skipping peptide sequence. Gene modification frequencies were assessed at 3 days post-transfection by the Surveyor assay. (FIG. **19**C) DMD myoblasts with deletions of exons 48-50 in the dystrophin gene were treated with sgRNAs that correct the dystrophin reading frame in these patient cells. Gene modification was assessed at 20 days post-electroporation in unsorted (bulk) or GFP+ sorted cells. (FIG. **19**D) GFP expression in DMD myoblasts 3 days after electroporation with indicated expression plasmids. Transfection efficiencies and sorted cell populations are indicated by the gated region.

[0055] FIGS. **20**A-**20**D show targeted frameshifts to restore the dystrophin reading frame using CRISPR/Cas9. (FIG. **20**A) The 5' region of exon 51 was targeted using a sgRNA (SEQ ID NO: 637), CR3, that binds immediately upstream of the first out-of-frame stop codon. PAM: protospacer-adjacent motif. (FIG. **20**B) The exon 51 locus was PCR amplified from HEK293T cells treated with SpCas9 and CR3 expression cassettes. Sequences of individual clones were determined by Sanger sequencing. The top sequence (bolded, exon in red) is the native, unmodified sequence. The number of clones for each sequence is indicated in parentheses. (FIG. **20**C) Summary of total gene editing efficiency and reading frame conversions resulting from gene modification shown in FIG. **20**B. (FIG. **20**D) Western blot for dystrophin expression in human DMD myoblasts treated with SpCas9 and the CR3 sgRNA expression cassette (FIG. **19**C) to create targeted frameshifts to restore the dystrophin reading frame. Dystrophin expression was probed using an antibody against the rod-domain of the dystrophin protein after 6 days of differentiation.

[0056] FIGS. **21**A-**21**D show deletion of exon 51 from the human genome using multiplex CRISPR/Cas9 gene editing.

(FIG. **21**A) End-point genomic PCR across the exon 51 locus in human DMD myoblasts with a deletion of exons 48-50. The top arrow indicates the expected position of full-length PCR amplicons and the two lower arrows indicate the expected position of PCR amplicons with deletions caused by the indicated sgRNA combinations. (FIG. **21**B) PCR products from FIG. **21**A were cloned and individual clones were sequenced to determine insertions and deletions present at the targeted locus (SEQ ID NOS 424, 638, 425-428, 639 and 429-431, respectively, in order of appearance). The top row shows the wild-type unmodified sequence and the triangles indicate SpCas9 cleavage sites. At the right are representative chromatograms showing the sequences of the expected deletion junctions (SEQ ID NOS 640-642, respectively, in order of appearance). (FIG. **21**C) End-point RT-PCR analysis of dystrophin mRNA transcripts in CRISPR/Cas9-modified human Δ48-50 DMD myoblasts treated with the indicated sgRNAs. A representative chromatogram of the expected deletion PCR product is shown at the right. Asterisk: band resulting from hybridization of the deletion product strand to the unmodified strand. (FIG. **21**D) Rescue of dystrophin protein expression by CRISPR/Cas9 genome editing was assessed by western blot for the dystrophin protein with GAPDH as a loading control. The arrow indicates the expected restored dystrophin protein band.

[0057] FIGS. **22**A-**22**D show deletion of the entire exon 45-55 region in human DMD myoblasts by multiplex CRISPR/Cas9 gene editing. (FIG. **22**A) End-point genomic PCR of genomic DNA to detect deletion of the region between intron 44 and intron 55 after treating HEK293 Ts or DMD myoblasts with the indicated sgRNAs. (FIG. **22**B) Individual clones of PCR products (SEQ ID NOS 432, 643 and 433, respectively, in order of appearance) of the expected size for the deletions from DMD myoblasts in FIG. **22**A were analyzed by Sanger sequencing to determine the sequences of genomic deletions present at the targeted locus. Below is a representative chromatograms showing the sequence of the expected deletion junctions (SEQ ID NO: 644). (FIG. **22**C) End-point RT-PCR analysis of dystrophin mRNA transcripts in CRISPR/Cas9-modified human Δ48-50 DMD myoblasts treated with the indicated sgRNAs. A representative chromatogram of the expected deletion PCR product is shown at the right (SEQ ID NO: 645). (FIG. **22**D) Analysis of restored dystrophin protein expression by western blot following electroporation of DMD myoblasts with sgRNAs targeted to intron 44 and/or intron 55.

[0058] FIG. **23** shows verification of flow cytometry-based enrichment of gene-modified DMD myoblasts used for in vivo cell transplantation experiment. DMD myoblasts were treated with Cas9 with or without sgRNA expression vectors for CR1 and CR5 and sorted for GFP+ cells by flow cytometry. Deletions at the exon 51 locus were detected by end-point PCR using primers flanking the locus. Neg ctrl: DMD myoblasts treated with Cas9 only and sorted for GFP+ cells.

[0059] FIG. **24** shows expression of restored human dystrophin in vivo following transplantation of CRISPR/Cas9-treated human DMD myoblasts into immunodeficient mice. Human Δ48-50 DMD myoblasts were treated with SpCas9, CR1, and CR5 to delete exon 51 and sorted for GFP expression as shown in FIGS. **19**A-**19**D. These sorted cells and untreated control cells were injected into the hind limbs of immunodeficient mice and assessed for human-specific protein expression in muscle fibers after 4 weeks post-transplantation. Cryosections were stained with anti-human spectrin, which is expressed by both uncorrected and corrected myoblasts that have fused into mouse myofibers, or anti-human dystrophin antibodies as indicated. White arrows indicate muscle fibers positive for human dystrophin.

[0060] FIGS. **25**A-**25**F show additional immunofluorescence images probing human dystrophin expression. Serial sections from regions stained with anti-human spectrin are shown inset in top left. (FIG. **25**A-FIG. **25**C) Sections from muscles injected with untreated human DMD myoblasts. (FIG. **25**D-FIG. **25**F) Sections from muscles injected with CR1/5 treated human DMD myoblasts enriched by flow cytometry. White arrows indicate dystrophin positive fibers.

[0061] FIGS. **26**A-**26**D shows evaluation of CRISPR/Cas9 toxicity and off-target effects for CR1/CR5-mediated deletion of exon 51 in human cells. (FIG. **26**A) Results of a cytotoxicity assay in HEK293T cells treated with human-optimized SpCas9 and the indicated sgRNA constructs. Cytotoxicity is based on survival of GFP-positive cells that are co-transfected with the indicated nuclease. I-SceI is a well-characterized non-toxic meganuclease and GZF3 is a known toxic zinc finger nuclease. (FIG. **26**B) Surveyor analysis at off-target sites in sorted hDMD cells treated with expression cassettes encoding Cas9 the indicated sgRNAs. These three off-target sites tested in hDMD cells were identified from a panel of 50 predicted sites tested in HEK293T cells (FIG. **27** and Table 4). TGT: on-target locus for indicated sgRNA. OT:off-target locus. (FIG. **26**C, FIG. **26**D) End-point nested PCR to detect chromosomal translocations in (FIG. **26**C) HEK293T cells treated with Cas9 and CR1 or (FIG. **26**D) sorted hDMD cells treated with Cas9, CR1, and CR5. The schematic depicts the relative location of nested primer pairs customized for each translocation event. The expected size of each band was estimated based on the primer size and the location of the predicted sgRNA cut site at each locus. Asterisks indicate bands detected at the expected size. The identities of the bands in FIG. **26**C were verified by Sanger sequencing from each end (FIG. **30**). A representative chromatogram for the P2/P5 translocation in HEK293T cells is shown (SEQ ID NO: 646).

[0062] FIG. **27** shows images of TBE-PAGE gels used to quantify Surveyor assay results to measure on-target and off-target gene modification in Table 4. Asterisks mark expected sizes of bands indicative of nuclease activity.

[0063] FIGS. **28**A-**28**C show end-point nested PCR to detect chromosomal translocations caused by CRISPR/Cas9 off-target activity for CR3 and CR6/CR36 in human cells. Nested end-point PCR analysis was used to detect translocations in (FIG. **28**A) HEK293T or sorted hDMD cells treated with Cas9 and CR3 as indicated, (FIG. **28**B) HEK293T cells treated with Cas9 and CR36 alone, or (FIG. **28**C) sorted hDMD cells treated with Cas9, CR6, and CR36 expression cassettes. The second nested PCR reaction for translocation was amplified using custom primers for each predicted translocation locus to maximize specificity (See Table 4). The schematic depicts the relative location of nested primer pairs used to probe for the presence of translocations. Each possible translocation event was first amplified from genomic DNA isolated from cells treated with or without the indicated sgRNA(s). A second nested PCR reaction was performed using primers within the predicted PCR amplicons that would result from translocations. Expected size was estimated based on the indicated

primer binding site and the predicted sgRNA cut site at each locus. *indicates bands detected at the expected size and verified by Sanger sequencing from each end. # indicates amplicons in which Sanger sequencing showed sequences other than the predicted translocation, likely a result of mispriming during the nested PCR.

[0064] FIG. **29** shows Sanger sequencing chromatograms (SEQ ID NOS 647-654, respectively, in order of appearance) for bands detected in FIGS. **28**A-**28**C resulting from translocations between CR3 and CR3-OT1, on chromosomes X and 1, respectively, in HEK293T cells treated with Cas9 and CR3 gene cassettes. Arrows show regions of homology to the indicated chromosome nearby the expected break points caused by the appropriate sgRNAs. Note that sequencing reads become out of phase near the break point due to the error-prone nature of DNA repair by non-homologous end-joining.

[0065] FIG. **30** shows Sanger sequencing chromatograms (SEQ ID NOS 655-660, respectively, in order of appearance) for bands detected in FIG. **26**C resulting from translocations between CR1 and CR1-OT1, on chromosomes X and 16, respectively, in HEK293T cells treated with Cas9 and CR1 gene cassettes. Arrows show regions of homology to the indicated chromosome nearby the expected break points caused by the appropriate sgRNAs. Note that sequencing reads become out of phase near the break point due to the error-prone nature of DNA repair by non-homologous end-joining.

[0066] FIG. **31** shows an overview of in vivo AAV injections and tissue harvest.

[0067] FIGS. **32**A-**32**C show Surveyor analysis of Rosa26 ZFN activities in skeletal muscle in vitro and in vivo following delivery of AAV-SASTG-ROSA. Arrows indicate expected bands resulting from Surveyor cleavage. n.d.: not detected. (FIG. **32**A) Proliferating C2C12s were transduced with the indicated amount of virus and harvested at 4 days post-infection. Arrows indicate expected bands sizes resulting from Surveyor cleavage. (FIG. **32**B) C2C12s were incubated in differentiation medium for 5 days and then transduced with the indicated amount of AAV-SASTG-ROSA virus in 24 well plates. Samples were collected at 10 days post-transduction. (FIG. **32**C) The indicated amount of AAV-SASTG-ROSA was injected directly into the tibialis anterior of C57BL/6J mice and muscles were harvested 4 weeks post-infection. The harvested TA muscles were partitioned into 8 separate pieces for genomic DNA analysis, each shown in a separate lane.

[0068] FIG. **33** shows Rosa T2A opt DNA sequence (SEQ ID NO: 434) and Rosa T2A opt protein sequence (SEQ ID NO: 435).

[0069] FIG. **34** shows SASTG capsid DNA sequence (SEQ ID NO:436) and SASTG capsid peptide sequence (SEQ ID NO: 437).

[0070] FIG. **35** shows DZF16 ZFN target site sequence (SEQ ID NO: 442), DZF16-L6 left full amino acid sequence (SEQ ID NO: 443) and DZF16-R6 right full amino acid sequence (SEQ ID NO: 444).

[0071] FIG. **36** shows E51C3 ZFN target site sequence (SEQ ID NO: 445), E51C3-3L left full amino acid sequence (SEQ ID NO: 446) and E51C3-3R right full amino acid sequence (SEQ ID NO: 447).

[0072] FIG. **37** shows DZF15 ZFN target site sequence (SEQ ID NO: 448), DZF15-L6 left full amino acid sequence (SEQ ID NO: 449), DZF15-R6 right full amino acid sequence (SEQ ID NO: 450), DZF15-L5 left full amino acid sequence (SEQ ID NO: 451), DZF15-R5 right full amino acid sequence (SEQ ID NO: 452).

[0073] FIG. **38** shows E51C4 ZFN target site sequence (SEQ ID NO: 453), E51C4-4L left full amino acid sequence (SEQ ID NO: 454) and E51C4-4R right full amino acid sequence (SEQ ID NO: 455).

[0074] FIG. **39** shows schematic diagrams of a "Single vector, multiplex CRISPR system," Dual vector, multiplex CRISPR system," and "Single vector, single gRNA system."

[0075] FIG. **40** shows the nucleotide sequences of SaCas9-NLS (with the NLS underlined) (SEQ ID NO: 459) and SaCas9 gRNA (SEQ ID NO: 460).

[0076] FIG. **41** shows the nucleotide sequences of NmCas9 (with the NLS 1 underlined, the NLS 2 underlined and bolded, and the HA tag bolded), NmCas9 short hairpin from Thomson *PNAS* 2013 (SEQ ID NO: 462), and NmCas9 long hairpin from Church Nature Biotech 2013 (SEQ ID NO: 463).

[0077] FIGS. **42**A-**42**C show validation of sgRNA and lentiviral Cas9 expression constructs. (FIG. **42**A) Constructs encoding unique Pol III promoters expressing sgRNAs targeting the AAVS1 locus or a construct containing the hU6 promoter immediately followed by poly-thymidine to terminate expression ("PolyT") were transfected into HEK293T cells. End-point RT-PCR was used to probe for expression of each indicated promoter/sgRNA construct two days post-transfection. -RT: no reverse transcriptase control. (FIG. **42**B) HEK293 Ts were transfected with expression vectors encoding the AAVS1 zinc-finger nuclease or Cas9-T2A-GFP and the indicated promoter/sgRNA expression cassettes and assessed for gene modification levels 3 days post-transfection using the Surveyor assay. (FIG. **42**C) HEK293T cells were transduced with lentiviral constructs encoding the indicated Cas9-T2A-GFP constructs without sgRNAs and assessed for Cas9 expression by western blot 7 days post-transduction by probing for a FLAG epitope tag on the N-terminus of the Cas9 protein.

[0078] FIG. **43** shows Golden Gate assembly of single lentiviral CRISPR/Cas9 expression cassettes. FIG. **43** discloses SEQ ID NOS 661-672, respectively, in order of appearance.

[0079] FIGS. **44**A-**44**B show single lentiviral delivery of a multiplex CRISPR/Cas9 system. (FIG. **44**A) Four sgRNAs targeting distinct genomic loci were cloned into a lentiviral vector expressing the active Cas9 nuclease. (FIG. **44**B) HEK293 Ts and primary human dermal fibroblasts were transduced with lentivirus expressing the indicated sgRNAs and assayed for cleavage events using the Surveyor assay. HEK293 Ts were assayed 7 days post transduction. The human fibroblasts were assayed 10 days post transduction.

[0080] FIGS. **45**A-**45**D show transient gene activation in HEK293 Ts stably expressing dCas9-VP64. HEK293 Ts were transduced with lentivirus to stably express dCas9-VP64 and were subsequently transfected with plasmid expressing the indicated sgRNA combinations. By varying the number of sgRNAs delivered, tunable endogenous gene activation of the endogenous IL1RN (FIG. **45**A) and HBG1 (FIG. **45**B) loci was achieved 3 days post transfection. Peak levels of endogenous IL1RN (FIG. **45**C) and HBG1 (FIG. **45**D) were observed 3-6 days post transfection and the level of activation returned to background levels between days 15-20. Importantly, the cell lines were able to reactive

following a second transfection on day 20 albeit at a lower level than previously observed.

[0081] FIGS. **46**A-**46**D show stable gene activation in HEK293 Ts using a single lentiviral multiplex dCas9-VP64 vector. HEK293 Ts were transduced with lentivirus to stably express dCas9-VP64 and the indicated combinations of gRNAs. By varying the number of sgRNAs delivered, tunable endogenous gene activation of the endogenous IL1RN (FIG. **46**A) and HBG1 (FIG. **46**B) loci was achieved 7 days post transduction. Peak levels of endogenous IL1RN (FIG. **46**C) and HBG1 (FIG. **46**D) were observed 6 days post transduction and the level of activation was sustained out to day 21.

[0082] FIG. **47** shows IL1RN mRNA expression levels.

[0083] FIG. **48** shows a schematic representing the direct conversion of fibroblasts to neurons through ectopic expression of the BAM neuronal transcription factors.

[0084] FIGS. **49**A-**49**C show (FIG. **49**A) Schematic of the dCas9-VP64 construct. dCas9-VP64 is a catalytically inactive form of the Cas9 protein fused to a tetramer of the VP16 transcriptional activation domain. (FIG. **49**B) Schematic showing the mechanism of RNA-guided recruitment of dCas9-VP64 to a genomic target. (FIG. **49**C) Schematic of the experimental protocol to generate iNs with CRISPR/Cas9 transcription factors.

[0085] FIGS. **50**A-**50**B show endogenous ASCL1 expression at day 3 determined by (FIG. **50**A) qRT-PCR or total ASCL1 protein detected by (FIG. **50**B) immunofluorescence in MEFs transduced with dCas9-VP64 and transfected with either gRNAs targeted to the ASCL1 promoter, ASCL1 cDNA, or luciferase. Asterisk (*) indicates significant ($p<0.05$) increase in ASCL1 expression with the co-delivery of 8 gRNAs compared to 4 gRNAs. Ectopic expression of ASCL1 produced more protein than induced by dCas9-VP64 and 8 gRNAs targeted to the Ascl1 promoter, but did not activate the endogenous locus by day 3 in culture.

[0086] FIG. **51**A shows TUJ1 and MAP2-positive cells generated by ectopic BAM factors or by dCas9-VP64 and gRNAs targeted to the BRN2 and ASCL1 promoters. FIG. **51**B shows cells with neuronal morphology expressing a hSyn-RFP reporter at day 11 in N3 medium.

[0087] FIG. **52**A shows a cell with neuronal morphology positive for the GCaMP5 calcium indicator in the presence (bottom) or absence (top) of KCl in the culture medium. FIG. **52**B shows a trace of normalized fluorescent intensity over time showing depolarization of the cell in response to KCl addition.

[0088] FIGS. **53**A-**53**C show the CRISPR/Cas9 platform for control of mammalian gene regulation. (FIG. **53**A) Cas9-based effectors bind genomic sequences in the presence of a chimeric gRNA molecule consisting of a constant region that complexes with Cas9 preceded by an exchangeable 20 bp protospacer that confers target site specificity. FIG. **53**A discloses SEQ ID NO: 673. (FIG. **53**B) Cas9-based synthetic transcription factors repress transcription of a target gene by interfering with RNA polymerase activity or by binding within the promoter and blocking the binding sites of endogenous transcription factors. (FIG. **53**C) Targeting regulatory elements such as enhancers could also potentially block the expression of multiple distal genes.

[0089] FIG. **54** shows targeting the HS2 enhancer using CRISPR/dCas9-KRAB. The HS2 region is a potent enhancer that distally regulates the expression of globin genes >10 kb downstream. A panel of single gRNAs was designed to target sites along the enhancer region.

[0090] FIGS. **55**A-**55**E show that single gRNAs targeting the HS2 enhancer effect potent transcriptional repression of globin genes. (FIG. **55**A) dCas9 and dCas9-KRAB repressors were delivered on a lentiviral vector. Single gRNAs were transiently transfected for screening. When assayed by quantitative RT-PCR at 3 days post-transfection, K562s expressing dCas9-KRAB achieve up to 80% repression of (FIG. **55**B) γ-globin, (FIG. **55**C) ε-globin, and (FIG. **55**D) β-globin genes, as compared to control cells that received no gRNA treatment. (FIG. **55**E) Protein expression in cells expressing dCas9 or dCas9-KRAB and treated with Cr4 or Cr8 show mild repression of γ-globin expression at day 3, compared to β-actin controls.

[0091] FIGS. **56**A-**56**C show expression of globin locus genes with varying doses of gRNA plasmid delivered to cells treated with (FIG. **56**A) no lentivirus, (FIG. **56**B) dCas9 lentivirus, or (FIG. **56**C) dCas9-KRAB lentivirus. Increasing the dose of Cr4 gRNA plasmid delivered enhanced repression in dCas9-KRAB treated cells, indicating that both the dCas9-KRAB effector and targeted gRNA play a role in achieving repression.

[0092] FIGS. **57**A-**57**D show that stably delivering single gRNAs with dCas9-KRAB silences expression of the globin genes. (FIG. **57**A) dCas9 and dCas9-KRAB repressors were co-expressed on a lentiviral vector with single gRNAs. When assayed by quantitative RT-PCR at 7 days post-transduction, K562s expressing dCas9-KRAB achieve up to 95% repression of (FIG. **57**B) γ-globin, (FIG. **57**C) ε-globin, and (FIG. **57**D) β-globin genes, as compared to control cells that received no lentiviral treatment.

[0093] FIG. **58** shows that isolating the p300 HAT "Core" for targeted epigenetic modification of histones only via dCas9 fusion.

[0094] FIG. **59** shows a simplified schematic of *S. pyogenes* dCas9-VP64 fusion (top) and dCas9-p300 core fusion (bottom). The Protospacer Adjacent Motifs (PAM) are shown with arrows at target gene loci and synthetic guide RNA (gRNA) is shown with hatched arrows.

[0095] FIGS. **60**A-**60**C show representative data at three human loci demonstrating the efficacy of activation using dCas9-p300 in relation to dCas9-VP64 and dCas9 without any fused effector domain in the human 293T cell culture line.

[0096] FIGS. **61**A-**61**C show the amino acid sequences of the dCas9 constructs. The legend for all FIGS. **61**A-**61**C is shown in FIG. **61**A. FIGS. **61**A-**61**C disclose SEQ ID NOS 674-676, respectively, in order of appearance.

[0097] FIG. **62** shows that HAT-dCase9-p300 fusion proteins fail to activate gene expression.

[0098] FIGS. **63**A-**63**B show that gRNA's also act synergistically with dCas9-p300 Core.

[0099] FIGS. **64**A-**64**C show TALEN mediated integration of minidystrophin at the 5'UTR of the Dp427m skeletal muscle isoform of dystrophin in skeletal myoblast cell lines derived from human DMD patients carrying different deletions in the dystrophin gene. DMD patient cells were electroporated with constructs encoding a TALEN pair active at the 5'UTR locus and a donor template carrying the minidystrophin gene. (FIG. **64**A) Schematic showing how minidystrophin is integrated into the 5'UTR. (FIG. **64**B) Hygromycin-resistant clonal cell lines were isolated and screened by PCR for successful site-specific integrations at the 5'UTR

using the primers shown in FIG. **64**A. Asterisks indicate clones selected for further analysis in FIG. **64**C. (FIG. **64**C) Clonally isolated DMD myoblasts with detected integration events were differentiated for 6 days and assessed for expression of an HA tag fused to the C terminus of minidystrophin.

DETAILED DESCRIPTION

[0100] As described herein, certain methods and engineered CRISPR/CRISPR-associated (Cas) 9-based system compositions have been discovered to be useful for altering the expression of genes, genome engineering, and correcting or reducing the effects of mutations in genes involved in genetic diseases. The CRISPR/Cas9-based system involves a Cas9 protein and at least one guide RNA, which provide the DNA targeting specificity for the system. In particular, the present disclosure describes a Cas9 fusion protein that combines the DNA sequence targeting function of the CRISPR/Cas9-based system with an additional activity, thus allowing changes in gene expression and/or epigenetic status. The system may also be used in genome engineering and correcting or reducing the effects of gene mutations.

[0101] The present disclosure also provides certain compositions and methods for delivering CRISPR/CRISPR-associated (Cas) 9-based system and multiple gRNAs to target one or more endogenous genes. Co-transfection of multiple sgRNAs targeted to a single promoter allow for synergistic activation, however, co-transfection of multiple plasmids leads to variable expression levels in each cell due to differences in copy number. Additionally, gene activation following transfection is transient due to dilution of plasmid DNA over time. Moreover, many cell types are not easily transfected and transient gene expression may not be sufficient for inducing a therapeutic effect. To address these limitations, a single lentiviral system was developed to express Cas9 and up to four sgRNAs from independent promoters. A platform is disclosed that expresses Cas9 or dCas9 fusion proteins and up to four gRNAs from a single lentiviral vector. The lentiviral vector expresses a constitutive or inducible Cas9 or dCas9-VP64 in addition to one, two, three, or four gRNAs expressed from independent promoters. This system enables control of both the magnitude and timing of CRISPR/Cas9-based gene regulation. Furthermore, the lentiviral platform provides the potent and sustained levels of gene expression that will facilitate therapeutic applications of the CRISPR/Cas9 system in primary cells. Finally, this system may be used for editing multiple genes simultaneously, such as the concurrent knockout of several oncogenes.

[0102] The present disclosure also provides certain compositions and methods for delivering site-specific nucleases to skeletal muscle and cardiac muscle using modified adeno-associated virus (AAV) vectors. The site-specific nucleases, which may be engineered, are useful for altering the expression of genes, genome engineering, correcting or reducing the effects of mutations in genes involved in genetic diseases, or manipulating genes involved in other conditions affecting skeletal muscle or cardiac muscle or muscle regeneration. The engineered site-specific nucleases may include a zinc finger nuclease (ZFN), a TAL effector nuclease (TALEN), and/or a CRISPR/Cas9 system for genome editing. As described herein, genes in skeletal muscle tissue were successfully edited in vivo using this unique delivery system. The disclosed invention provides a means to rewrite the human genome for therapeutic applications and target model species for basic science applications.

[0103] Gene editing is highly dependent on cell cycle and complex DNA repair pathways that vary from tissue to tissue. Skeletal muscle is a very complex environment, consisting of large myofibers with more than 100 nuclei per cell. Gene therapy and biologics in general have been limited for decades by in vivo delivery hurdles. These challenges include stability of the carrier in vivo, targeting the right tissue, getting sufficient gene expression and active gene product, and avoiding toxicity that might overcome activity, which is common with gene editing tools. Other delivery vehicles, such as direct injection of plasmid DNA, work to express genes in skeletal muscle and cardiac muscle in other contexts, but do not work well with these site-specific nucleases for achieving detectable levels of genome editing.

[0104] While many gene sequences are unstable in AAV vectors and therefore undeliverable, these site-specific nucleases are surprisingly stable in the AAV vectors. When these site-specific nucleases are delivered and expressed, they remained active in the skeletal muscle tissue. The protein stability and activity of the site-specific nucleases are highly tissue type- and cell type-dependent. These active and stable nucleases are able to modify gene sequences in the complex environment of skeletal muscle. The current invention describes a way to deliver active forms of this class of therapeutics to skeletal muscle or cardiac muscle that is effective, efficient and facilitates successful genome modification.

[0105] The present disclosure also provides certain fusion epigenetic effector molecules, a dCas9-p300 fusion protein, which provides a robust and potentially more widely applicable tool for synthetic transcriptional modulation compared to the dCas9-VP64 fusion. The activated target genes to a substantially greater extent than the dCas9-VP64 fusion protein at all loci tested. In addition, the p300 has intrinsic endogenous activity at enhancers within the human genome. The dCas9-p300 fusion protein may be able to activate endogenous target gene promoters and enhancer regions.

[0106] The dCas9-p300 fusion protein can be used in human tissue culture cell lines to activate gene expression. This fusion protein may be used to direct the epigenetic state of target loci within human cells with precision and predictability in order to control differentiation, modulate cellular regulation, and apply innovative potential therapies. Current technologies are limited in the strength of activation and the extent and sustainability of epigenetic modulation; obstacles which may be obviated via utilization of this new fusion protein.

[0107] Section headings as used in this section and the entire disclosure herein are merely for organizational purposes and are not intended to be limiting.

1. DEFINITIONS

[0108] Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art. In case of conflict, the present document, including definitions, will control. Preferred methods and materials are described below, although methods and materials similar or equivalent to those described herein can be used in practice or testing of the present invention. All publications, patent applications, patents and other references mentioned herein are

incorporated by reference in their entirety. The materials, methods, and examples disclosed herein are illustrative only and not intended to be limiting.

[0109] The terms "comprise(s)," "include(s)," "having," "has," "can," "contain(s)," and variants thereof, as used herein, are intended to be open-ended transitional phrases, terms, or words that do not preclude the possibility of additional acts or structures. The singular forms "a," "and" and "the" include plural references unless the context clearly dictates otherwise. The present disclosure also contemplates other embodiments "comprising," "consisting of" and "consisting essentially of," the embodiments or elements presented herein, whether explicitly set forth or not.

[0110] For the recitation of numeric ranges herein, each intervening number there between with the same degree of precision is explicitly contemplated. For example, for the range of 6-9, the numbers 7 and 8 are contemplated in addition to 6 and 9, and for the range 6.0-7.0, the number 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, and 7.0 are explicitly contemplated.

[0111] "Adeno-associated virus" or "AAV" as used interchangeably herein refers to a small virus belonging to the genus Dependovirus of the Parvoviridae family that infects humans and some other primate species. AAV is not currently known to cause disease and consequently the virus causes a very mild immune response.

[0112] "Binding region" as used herein refers to the region within a nuclease target region that is recognized and bound by the nuclease.

[0113] "Cardiac muscle" or "heart muscle" as used interchangeably herein means a type of involuntary striated muscle found in the walls and histological foundation of the heart, the myocardium. Cardiac muscle is made of cardiomyocytes or myocardiocytes. Myocardiocytes show striations similar to those on skeletal muscle cells but contain only one, unique nucleus, unlike the multinucleated skeletal cells.

[0114] "Cardiac muscle condition" as used herein refers to a condition related to the cardiac muscle, such as cardiomyopathy, heart failure, arrhythmia, and inflammatory heart disease.

[0115] "Coding sequence" or "encoding nucleic acid" as used herein means the nucleic acids (RNA or DNA molecule) that comprise a nucleotide sequence which encodes a protein. The coding sequence can further include initiation and termination signals operably linked to regulatory elements including a promoter and polyadenylation signal capable of directing expression in the cells of an individual or mammal to which the nucleic acid is administered. The coding sequence may be codon optimize.

[0116] "Complement" or "complementary" as used herein means a nucleic acid can mean Watson-Crick (e.g., A-T/U and C-G) or Hoogsteen base pairing between nucleotides or nucleotide analogs of nucleic acid molecules. "Complementarity" refers to a property shared between two nucleic acid sequences, such that when they are aligned antiparallel to each other, the nucleotide bases at each position will be complementary.

[0117] "Correcting", "genome editing" and "restoring" as used herein refers to changing a mutant gene that encodes a truncated protein or no protein at all, such that a full-length functional or partially full-length functional protein expression is obtained. Correcting or restoring a mutant gene may include replacing the region of the gene that has the mutation or replacing the entire mutant gene with a copy of the gene that does not have the mutation with a repair mechanism such as homology-directed repair (HDR). Correcting or restoring a mutant gene may also include repairing a frameshift mutation that causes a premature stop codon, an aberrant splice acceptor site or an aberrant splice donor site, by generating a double stranded break in the gene that is then repaired using non-homologous end joining (NHEJ). NHEJ may add or delete at least one base pair during repair which may restore the proper reading frame and eliminate the premature stop codon. Correcting or restoring a mutant gene may also include disrupting an aberrant splice acceptor site or splice donor sequence. Correcting or restoring a mutant gene may also include deleting a non-essential gene segment by the simultaneous action of two nucleases on the same DNA strand in order to restore the proper reading frame by removing the DNA between the two nuclease target sites and repairing the DNA break by NHEJ.

[0118] "Donor DNA", "donor template" and "repair template" as used interchangeably herein refers to a double-stranded DNA fragment or molecule that includes at least a portion of the gene of interest. The donor DNA may encode a full-functional protein or a partially-functional protein.

[0119] "Duchenne Muscular Dystrophy" or "DMD" as used interchangeably herein refers to a recessive, fatal, X-linked disorder that results in muscle degeneration and eventual death. DMD is a common hereditary monogenic disease and occurs in 1 in 3500 males. DMD is the result of inherited or spontaneous mutations that cause nonsense or frame shift mutations in the dystrophin gene. The majority of dystrophin mutations that cause DMD are deletions of exons that disrupt the reading frame and cause premature translation termination in the dystrophin gene. DMD patients typically lose the ability to physically support themselves during childhood, become progressively weaker during the teenage years, and die in their twenties.

[0120] "Dystrophin" as used herein refers to a rod-shaped cytoplasmic protein which is a part of a protein complex that connects the cytoskeleton of a muscle fiber to the surrounding extracellular matrix through the cell membrane. Dystrophin provides structural stability to the dystroglycan complex of the cell membrane that is responsible for regulating muscle cell integrity and function. The dystrophin gene or "DMD gene" as used interchangeably herein is 2.2 megabases at locus Xp21. The primary transcription measures about 2,400 kb with the mature mRNA being about 14 kb. 79 exons code for the protein which is over 3500 amino acids.

[0121] "Exon 51" as used herein refers to the 51st exon of the dystrophin gene. Exon 51 is frequently adjacent to frame-disrupting deletions in DMD patients and has been targeted in clinical trials for oligonucleotide-based exon skipping. A clinical trial for the exon 51 skipping compound eteplirsen recently reported a significant functional benefit across 48 weeks, with an average of 47% dystrophin positive fibers compared to baseline. Mutations in exon 51 are ideally suited for permanent correction by NHEJ-based genome editing.

[0122] "Frameshift" or "frameshift mutation" as used interchangeably herein refers to a type of gene mutation wherein the addition or deletion of one or more nucleotides causes a shift in the reading frame of the codons in the mRNA. The shift in reading frame may lead to the alteration

in the amino acid sequence at protein translation, such as a missense mutation or a premature stop codon.

[0123] "Functional" and "full-functional" as used herein describes protein that has biological activity. A "functional gene" refers to a gene transcribed to mRNA, which is translated to a functional protein.

[0124] "Fusion protein" as used herein refers to a chimeric protein created through the joining of two or more genes that originally coded for separate proteins. The translation of the fusion gene results in a single polypeptide with functional properties derived from each of the original proteins.

[0125] "Genetic construct" as used herein refers to the DNA or RNA molecules that comprise a nucleotide sequence that encodes a protein. The coding sequence includes initiation and termination signals operably linked to regulatory elements including a promoter and polyadenylation signal capable of directing expression in the cells of the individual to whom the nucleic acid molecule is administered. As used herein, the term "expressible form" refers to gene constructs that contain the necessary regulatory elements operable linked to a coding sequence that encodes a protein such that when present in the cell of the individual, the coding sequence will be expressed.

[0126] "Genetic disease" as used herein refers to a disease, partially or completely, directly or indirectly, caused by one or more abnormalities in the genome, especially a condition that is present from birth. The abnormality may be a mutation, an insertion or a deletion. The abnormality may affect the coding sequence of the gene or its regulatory sequence. The genetic disease may be, but not limited to DMD, hemophilia, cystic fibrosis, Huntington's chorea, familial hypercholesterolemia (LDL receptor defect), hepatoblastoma, Wilson's disease, congenital hepatic porphyria, inherited disorders of hepatic metabolism, Lesch Nyhan syndrome, sickle cell anemia, thalassaemias, xeroderma pigmentosum, Fanconi's anemia, retinitis pigmentosa, ataxia telangiectasia, Bloom's syndrome, retinoblastoma, and Tay-Sachs disease.

[0127] "Homology-directed repair" or "HDR" as used interchangeably herein refers to a mechanism in cells to repair double strand DNA lesions when a homologous piece of DNA is present in the nucleus, mostly in G2 and S phase of the cell cycle. HDR uses a donor DNA template to guide repair and may be used to create specific sequence changes to the genome, including the targeted addition of whole genes. If a donor template is provided along with the site specific nuclease, such as with a CRISPR/Cas9-based systems, then the cellular machinery will repair the break by homologous recombination, which is enhanced several orders of magnitude in the presence of DNA cleavage. When the homologous DNA piece is absent, non-homologous end joining may take place instead.

[0128] "Genome editing" as used herein refers to changing a gene. Genome editing may include correcting or restoring a mutant gene. Genome editing may include knocking out a gene, such as a mutant gene or a normal gene. Genome editing may be used to treat disease or enhance muscle repair by changing the gene of interest.

[0129] "Identical" or "identity" as used herein in the context of two or more nucleic acids or polypeptide sequences means that the sequences have a specified percentage of residues that are the same over a specified region. The percentage may be calculated by optimally aligning the two sequences, comparing the two sequences over the specified region, determining the number of positions at which the identical residue occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the specified region, and multiplying the result by 100 to yield the percentage of sequence identity. In cases where the two sequences are of different lengths or the alignment produces one or more staggered ends and the specified region of comparison includes only a single sequence, the residues of single sequence are included in the denominator but not the numerator of the calculation. When comparing DNA and RNA, thymine (T) and uracil (U) may be considered equivalent. Identity may be performed manually or by using a computer sequence algorithm such as BLAST or BLAST 2.0.

[0130] "Mutant gene" or "mutated gene" as used interchangeably herein refers to a gene that has undergone a detectable mutation. A mutant gene has undergone a change, such as the loss, gain, or exchange of genetic material, which affects the normal transmission and expression of the gene. A "disrupted gene" as used herein refers to a mutant gene that has a mutation that causes a premature stop codon. The disrupted gene product is truncated relative to a full-length undisrupted gene product.

[0131] "Non-homologous end joining (NHEJ) pathway" as used herein refers to a pathway that repairs double-strand breaks in DNA by directly ligating the break ends without the need for a homologous template. The template-independent re-ligation of DNA ends by NHEJ is a stochastic, error-prone repair process that introduces random micro-insertions and micro-deletions (indels) at the DNA breakpoint. This method may be used to intentionally disrupt, delete, or alter the reading frame of targeted gene sequences. NHEJ typically uses short homologous DNA sequences called microhomologies to guide repair. These microhomologies are often present in single-stranded overhangs on the end of double-strand breaks. When the overhangs are perfectly compatible, NHEJ usually repairs the break accurately, yet imprecise repair leading to loss of nucleotides may also occur, but is much more common when the overhangs are not compatible.

[0132] "Normal gene" as used herein refers to a gene that has not undergone a change, such as a loss, gain, or exchange of genetic material. The normal gene undergoes normal gene transmission and gene expression.

[0133] "Nuclease mediated NHEJ" as used herein refers to NHEJ that is initiated after a nuclease, such as a cas9, cuts double stranded DNA.

[0134] "Nucleic acid" or "oligonucleotide" or "polynucleotide" as used herein means at least two nucleotides covalently linked together. The depiction of a single strand also defines the sequence of the complementary strand. Thus, a nucleic acid also encompasses the complementary strand of a depicted single strand. Many variants of a nucleic acid may be used for the same purpose as a given nucleic acid. Thus, a nucleic acid also encompasses substantially identical nucleic acids and complements thereof. A single strand provides a probe that may hybridize to a target sequence under stringent hybridization conditions. Thus, a nucleic acid also encompasses a probe that hybridizes under stringent hybridization conditions.

[0135] Nucleic acids may be single stranded or double stranded, or may contain portions of both double stranded and single stranded sequence. The nucleic acid may be

DNA, both genomic and cDNA, RNA, or a hybrid, where the nucleic acid may contain combinations of deoxyribo- and ribo-nucleotides, and combinations of bases including uracil, adenine, thymine, cytosine, guanine, inosine, xanthine hypoxanthine, isocytosine and isoguanine. Nucleic acids may be obtained by chemical synthesis methods or by recombinant methods.

[0136] "Operably linked" as used herein means that expression of a gene is under the control of a promoter with which it is spatially connected. A promoter may be positioned 5' (upstream) or 3' (downstream) of a gene under its control. The distance between the promoter and a gene may be approximately the same as the distance between that promoter and the gene it controls in the gene from which the promoter is derived. As is known in the art, variation in this distance may be accommodated without loss of promoter function.

[0137] "Partially-functional" as used herein describes a protein that is encoded by a mutant gene and has less biological activity than a functional protein but more than a non-functional protein.

[0138] "Premature stop codon" or "out-of-frame stop codon" as used interchangeably herein refers to nonsense mutation in a sequence of DNA, which results in a stop codon at location not normally found in the wild-type gene. A premature stop codon may cause a protein to be truncated or shorter compared to the full-length version of the protein.

[0139] "Promoter" as used herein means a synthetic or naturally-derived molecule which is capable of conferring, activating or enhancing expression of a nucleic acid in a cell. A promoter may comprise one or more specific transcriptional regulatory sequences to further enhance expression and/or to alter the spatial expression and/or temporal expression of same. A promoter may also comprise distal enhancer or repressor elements, which may be located as much as several thousand base pairs from the start site of transcription. A promoter may be derived from sources including viral, bacterial, fungal, plants, insects, and animals. A promoter may regulate the expression of a gene component constitutively, or differentially with respect to cell, the tissue or organ in which expression occurs or, with respect to the developmental stage at which expression occurs, or in response to external stimuli such as physiological stresses, pathogens, metal ions, or inducing agents. Representative examples of promoters include the bacteriophage T7 promoter, bacteriophage T3 promoter, SP6 promoter, lac operator-promoter, tac promoter, SV40 late promoter, SV40 early promoter, RSV-LTR promoter, CMV IE promoter, SV40 early promoter or SV40 late promoter and the CMV IE promoter.

[0140] "Repeat variable diresidue" or "RVD" as used interchangeably herein refers to a pair of adjacent amino acid residues within a DNA recognition motif (also known as "RVD module"), which includes 33-35 amino acids, of a TALE DNA-binding domain. The RVD determines the nucleotide specificity of the RVD module. RVD modules may be combined to produce an RVD array. The "RVD array length" as used herein refers to the number of RVD modules that corresponds to the length of the nucleotide sequence within the TALEN target region that is recognized by a TALEN, i.e., the binding region.

[0141] "Site-specific nuclease" as used herein refers to an enzyme capable of specifically recognizing and cleaving DNA sequences. The site-specific nuclease may be engineered. Examples of engineered site-specific nucleases include zinc finger nucleases (ZFNs), TAL effector nucleases (TALENs), and CRISPR/Cas9-based systems.

[0142] "Skeletal muscle" as used herein refers to a type of striated muscle, which is under the control of the somatic nervous system and attached to bones by bundles of collagen fibers known as tendons. Skeletal muscle is made up of individual components known as myocytes, or "muscle cells", sometimes colloquially called "muscle fibers." Myocytes are formed from the fusion of developmental myoblasts (a type of embryonic progenitor cell that gives rise to a muscle cell) in a process known as myogenesis. These long, cylindrical, multinucleated cells are also called myofibers.

[0143] "Skeletal muscle condition" as used herein refers to a condition related to the skeletal muscle, such as muscular dystrophies, aging, muscle degeneration, wound healing, and muscle weakness or atrophy.

[0144] "Spacers" and "spacer region" as used interchangeably herein refers to the region within a TALEN or ZFN target region that is between, but not a part of, the binding regions for two TALENs or ZFNs.

[0145] "Subject" and "patient" as used herein interchangeably refers to any vertebrate, including, but not limited to, a mammal (e.g., cow, pig, camel, llama, horse, goat, rabbit, sheep, hamsters, guinea pig, cat, dog, rat, and mouse, a non-human primate (for example, a monkey, such as a cynomolgous or rhesus monkey, chimpanzee, etc.) and a human). In some embodiments, the subject may be a human or a non-human. The subject or patient may be undergoing other forms of treatment.

[0146] "Target gene" as used herein refers to any nucleotide sequence encoding a known or putative gene product. The target gene may be a mutated gene involved in a genetic disease.

[0147] "Target region" as used herein refers to the region of the target gene to which the site-specific nuclease is designed to bind and cleave.

[0148] "Transcription activator-like effector" or "TALE" as used herein refers to a protein structure that recognizes and binds to a particular DNA sequence. The "TALE DNA-binding domain" refers to a DNA-binding domain that includes an array of tandem 33-35 amino acid repeats, also known as RVD modules, each of which specifically recognizes a single base pair of DNA. RVD modules may be arranged in any order to assemble an array that recognizes a defined sequence.

[0149] A binding specificity of a TALE DNA-binding domain is determined by the RVD array followed by a single truncated repeat of 20 amino acids. A TALE DNA-binding domain may have 12 to 27 RVD modules, each of which contains an RVD and recognizes a single base pair of DNA. Specific RVDs have been identified that recognize each of the four possible DNA nucleotides (A, T, C, and G). Because the TALE DNA-binding domains are modular, repeats that recognize the four different DNA nucleotides may be linked together to recognize any particular DNA sequence. These targeted DNA-binding domains may then be combined with catalytic domains to create functional enzymes, including artificial transcription factors, methyltransferases, integrases, nucleases, and recombinases.

[0150] "Transcription activator-like effector nucleases" or "TALENs" as used interchangeably herein refers to engineered fusion proteins of the catalytic domain of a nuclease,

such as endonuclease FokI, and a designed TALE DNA-binding domain that may be targeted to a custom DNA sequence. A "TALEN monomer" refers to an engineered fusion protein with a catalytic nuclease domain and a designed TALE DNA-binding domain. Two TALEN monomers may be designed to target and cleave a TALEN target region.

[0151] "Transgene" as used herein refers to a gene or genetic material containing a gene sequence that has been isolated from one organism and is introduced into a different organism. This non-native segment of DNA may retain the ability to produce RNA or protein in the transgenic organism, or it may alter the normal function of the transgenic organism's genetic code. The introduction of a transgene has the potential to change the phenotype of an organism.

[0152] "Variant" used herein with respect to a nucleic acid means (i) a portion or fragment of a referenced nucleotide sequence; (ii) the complement of a referenced nucleotide sequence or portion thereof; (iii) a nucleic acid that is substantially identical to a referenced nucleic acid or the complement thereof; or (iv) a nucleic acid that hybridizes under stringent conditions to the referenced nucleic acid, complement thereof, or a sequences substantially identical thereto.

[0153] "Variant" with respect to a peptide or polypeptide that differs in amino acid sequence by the insertion, deletion, or conservative substitution of amino acids, but retain at least one biological activity. Variant may also mean a protein with an amino acid sequence that is substantially identical to a referenced protein with an amino acid sequence that retains at least one biological activity. A conservative substitution of an amino acid, i.e., replacing an amino acid with a different amino acid of similar properties (e.g., hydrophilicity, degree and distribution of charged regions) is recognized in the art as typically involving a minor change. These minor changes may be identified, in part, by considering the hydropathic index of amino acids, as understood in the art. Kyte et al., *J. Mol. Biol.* 157:105-132 (1982). The hydropathic index of an amino acid is based on a consideration of its hydrophobicity and charge. It is known in the art that amino acids of similar hydropathic indexes may be substituted and still retain protein function. In one aspect, amino acids having hydropathic indexes of ±2 are substituted. The hydrophilicity of amino acids may also be used to reveal substitutions that would result in proteins retaining biological function. A consideration of the hydrophilicity of amino acids in the context of a peptide permits calculation of the greatest local average hydrophilicity of that peptide. Substitutions may be performed with amino acids having hydrophilicity values within ±2 of each other. Both the hydrophobicity index and the hydrophilicity value of amino acids are influenced by the particular side chain of that amino acid. Consistent with that observation, amino acid substitutions that are compatible with biological function are understood to depend on the relative similarity of the amino acids, and particularly the side chains of those amino acids, as revealed by the hydrophobicity, hydrophilicity, charge, size, and other properties.

[0154] "Vector" as used herein means a nucleic acid sequence containing an origin of replication. A vector may be a viral vector, bacteriophage, bacterial artificial chromosome or yeast artificial chromosome. A vector may be a DNA or RNA vector. A vector may be a self-replicating extrachromosomal vector, and preferably, is a DNA plasmid. For example, the vector may encode an iCas9-VP64 fusion protein comprising the amino acid sequence of SEQ ID NO: 1 or at least one gRNA nucleotide sequence of any one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625. Alternatively, the vector may encode Cas9 and at least one gRNA nucleotide sequence of any one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625.

[0155] "Zinc finger" as used herein refers to a protein structure that recognizes and binds to DNA sequences. The zinc finger domain is the most common DNA-binding motif in the human proteome. A single zinc finger contains approximately 30 amino acids and the domain typically functions by binding 3 consecutive base pairs of DNA via interactions of a single amino acid side chain per base pair.

[0156] "Zinc finger nuclease" or "ZFN" as used interchangeably herein refers to a chimeric protein molecule comprising at least one zinc finger DNA binding domain effectively linked to at least one nuclease or part of a nuclease capable of cleaving DNA when fully assembled.

[0157] Unless otherwise defined herein, scientific and technical terms used in connection with the present disclosure shall have the meanings that are commonly understood by those of ordinary skill in the art. For example, any nomenclatures used in connection with, and techniques of, cell and tissue culture, molecular biology, immunology, microbiology, genetics and protein and nucleic acid chemistry and hybridization described herein are those that are well known and commonly used in the art. The meaning and scope of the terms should be clear; in the event however of any latent ambiguity, definitions provided herein take precedent over any dictionary or extrinsic definition. Further, unless otherwise required by context, singular terms shall include pluralities and plural terms shall include the singular.

2. COMPOSITIONS FOR GENOME EDITING

[0158] The present invention is directed to compositions for genome editing, genomic alteration or altering gene expression of a target gene. The compositions may include a may include viral vector and fusion protein such as a site-specific nuclease or CRISPR/Cas9-system with at least one gRNA.

[0159] a. Compositions for Genome Editing in Muscle

[0160] The present invention is directed to a composition for genome editing a target gene in skeletal muscle or cardiac muscle of a subject. The composition includes a modified AAV vector and a nucleotide sequence encoding a site-specific nuclease. The composition delivers active forms of site-specific nucleases to skeletal muscle or cardiac muscle. The composition may further comprise a donor DNA or a transgene. These compositions may be used in genome editing, genome engineering, and correcting or reducing the effects of mutations in genes involved in genetic diseases and/or other skeletal or cardiac muscle conditions.

[0161] The target gene may be involved in differentiation of a cell or any other process in which activation, repression, or disruption of a gene may be desired, or may have a mutation such as a deletion, frameshift mutation, or a nonsense mutation. If the target gene has a mutation that causes a premature stop codon, an aberrant splice acceptor site or an aberrant splice donor site, the site-specific nucleases may be designed to recognize and bind a nucleotide sequence upstream or downstream from the premature stop

codon, the aberrant splice acceptor site or the aberrant splice donor site. The site-specific nucleases may also be used to disrupt normal gene splicing by targeting splice acceptors and donors to induce skipping of premature stop codons or restore a disrupted reading frame. The site-specific nucleases may or may not mediate off-target changes to protein-coding regions of the genome.

3. CRISPR SYSTEM

[0162] "Clustered Regularly Interspaced Short Palindromic Repeats" and "CRISPRs", as used interchangeably herein refers to loci containing multiple short direct repeats that are found in the genomes of approximately 40% of sequenced bacteria and 90% of sequenced archaea. The CRISPR system is a microbial nuclease system involved in defense against invading phages and plasmids that provides a form of acquired immunity. The CRISPR loci in microbial hosts contain a combination of CRISPR-associated (Cas) genes as well as non-coding RNA elements capable of programming the specificity of the CRISPR-mediated nucleic acid cleavage. Short segments of foreign DNA, called spacers, are incorporated into the genome between CRISPR repeats, and serve as a 'memory' of past exposures. Cas9 forms a complex with the 3' end of the sgRNA, and the protein-RNA pair recognizes its genomic target by complementary base pairing between the 5' end of the sgRNA sequence and a predefined 20 bp DNA sequence, known as the protospacer. This complex is directed to homologous loci of pathogen DNA via regions encoded within the crRNA, i.e., the protospacers, and protospacer-adjacent motifs (PAMs) within the pathogen genome. The non-coding CRISPR array is transcribed and cleaved within direct repeats into short crRNAs containing individual spacer sequences, which direct Cas nucleases to the target site (protospacer). By simply exchanging the 20 bp recognition sequence of the expressed sgRNA, the Cas9 nuclease can be directed to new genomic targets. CRISPR spacers are used to recognize and silence exogenous genetic elements in a manner analogous to RNAi in eukaryotic organisms.

[0163] Three classes of CRISPR systems (Types I, II and III effector systems) are known. The Type II effector system carries out targeted DNA double-strand break in four sequential steps, using a single effector enzyme, Cas9, to cleave dsDNA. Compared to the Type I and Type III effector systems, which require multiple distinct effectors acting as a complex, the Type II effector system may function in alternative contexts such as eukaryotic cells. The Type II effector system consists of a long pre-crRNA, which is transcribed from the spacer-containing CRISPR locus, the Cas9 protein, and a tracrRNA, which is involved in pre-crRNA processing. The tracrRNAs hybridize to the repeat regions separating the spacers of the pre-crRNA, thus initiating dsRNA cleavage by endogenous RNase III. This cleavage is followed by a second cleavage event within each spacer by Cas9, producing mature crRNAs that remain associated with the tracrRNA and Cas9, forming a Cas9:crRNA-tracrRNA complex.

[0164] The Cas9:crRNA-tracrRNA complex unwinds the DNA duplex and searches for sequences matching the crRNA to cleave. Target recognition occurs upon detection of complementarity between a "protospacer" sequence in the target DNA and the remaining spacer sequence in the crRNA. Cas9 mediates cleavage of target DNA if a correct protospacer-adjacent motif (PAM) is also present at the 3' end of the protospacer. For protospacer targeting, the sequence must be immediately followed by the protospacer-adjacent motif (PAM), a short sequence recognized by the Cas9 nuclease that is required for DNA cleavage. Different Type II systems have differing PAM requirements. The *S. pyogenes* CRISPR system may have the PAM sequence for this Cas9 (SpCas9) as 5'-NRG-3', where R is either A or G, and characterized the specificity of this system in human cells. A unique capability of the CRISPR/Cas9 system is the straightforward ability to simultaneously target multiple distinct genomic loci by co-expressing a single Cas9 protein with two or more sgRNAs. For example, the Streptococcuspyogenes Type II system naturally prefers to use an "NGG" sequence, where "N" can be any nucleotide, but also accepts other PAM sequences, such as "NAG" in engineered systems (Hsu et al., *Nature Biotechnology* (2013) doi:10.1038/nbt.2647). Similarly, the Cas9 derived from *Neisseria meningitidis* (NmCas9) normally has a native PAM of NNNNGATT, but has activity across a variety of PAMs, including a highly degenerate NNNNGNNN PAM (Esvelt et al. *Nature Methods* (2013) doi:10.1038/nmeth.2681).

4. CRISPR/CAS9-BASED SYSTEM

[0165] An engineered form of the Type II effector system of Streptococcuspyogenes was shown to function in human cells for genome engineering. In this system, the Cas9 protein was directed to genomic target sites by a synthetically reconstituted "guide RNA" ("gRNA", also used interchangeably herein as a chimeric single guide RNA ("sgRNA")), which is a crRNA-tracrRNA fusion that obviates the need for RNase III and crRNA processing in general (see FIG. **53**A). Provided herein are CRISPR/Cas9-based engineered systems for use in genome editing and treating genetic diseases. The CRISPR/Cas9-based engineered systems may be designed to target any gene, including genes involved in a genetic disease, aging, tissue regeneration, or wound healing. The CRISPR/Cas9-based systems may include a Cas9 protein or Cas9 fusion protein and at least one gRNA. The Cas9 fusion protein may, for example, include a domain that has a different activity that what is endogenous to Cas9, such as a transactivation domain.

[0166] The target gene may be involved in differentiation of a cell or any other process in which activation of a gene may be desired, or may have a mutation such as a frameshift mutation or a nonsense mutation. If the target gene has a mutation that causes a premature stop codon, an aberrant splice acceptor site or an aberrant splice donor site, the CRISPR/Cas9-based system may be designed to recognize and bind a nucleotide sequence upstream or downstream from the premature stop codon, the aberrant splice acceptor site or the aberrant splice donor site. The CRISPR-Cas9-based system may also be used to disrupt normal gene splicing by targeting splice acceptors and donors to induce skipping of premature stop codons or restore a disrupted reading frame. The CRISPR/Cas9-based system may or may not mediate off-target changes to protein-coding regions of the genome.

[0167] a. Cas9

[0168] The CRISPR/Cas9-based system may include a Cas9 protein or a Cas9 fusion protein. Cas9 protein is an endonuclease that cleaves nucleic acid and is encoded by the CRISPR loci and is involved in the Type II CRISPR system. The Cas9 protein may be from any bacterial or archaea species, such as Streptococcuspyogenes. The Cas9 protein

may be mutated so that the nuclease activity is inactivated. An inactivated Cas9 protein from Streptococcuspyogenes (iCas9, also referred to as "dCas9") with no endonuclease activity has been recently targeted to genes in bacteria, yeast, and human cells by gRNAs to silence gene expression through steric hindrance. As used herein, "iCas9" and "dCas9" both refer to a Cas9 protein that has the amino acid substitutions D10A and H840A and has its nuclease activity inactivated. For example, the CRISPR/Cas9-based system may include a Cas9 of SEQ ID NO: 459 or 461.

[0169] b. Cas9 Fusion Protein

[0170] The CRISPR/Cas9-based system may include a fusion protein. The fusion protein may comprise two heterologous polypeptide domains, wherein the first polypeptide domain comprises a Cas protein and the second polypeptide domain has an activity such as transcription activation activity, transcription repression activity, transcription release factor activity, histone modification activity, nuclease activity, nucleic acid association activity, methylase activity, or demethylase activity. The fusion protein may include a Cas9 protein or a mutated Cas9 protein, as described above, fused to a second polypeptide domain that has an activity such as transcription activation activity, transcription repression activity, transcription release factor activity, histone modification activity, nuclease activity, nucleic acid association activity, methylase activity, or demethylase activity.

[0171] (1) Transcription Activation Activity

[0172] The second polypeptide domain may have transcription activation activity, i.e., a transactivation domain. For example, gene expression of endogenous mammalian genes, such as human genes, may be achieved by targeting a fusion protein of iCas9 and a transactivation domain to mammalian promoters via combinations of gRNAs. The transactivation domain may include a VP16 protein, multiple VP16 proteins, such as a VP48 domain or VP64 domain, or p65 domain of NF kappa B transcription activator activity. For example, the fusion protein may be iCas9-VP64.

[0173] (2) Transcription Repression Activity

[0174] The second polypeptide domain may have transcription repression activity. The second polypeptide domain may have a Kruppel associated box activity, such as a KRAB domain, ERF repressor domain activity, Mxi1 repressor domain activity, SID4X repressor domain activity, Mad-SID repressor domain activity or TATA box binding protein activity. For example, the fusion protein may be dCas9-KRAB.

[0175] (3) Transcription Release Factor Activity

[0176] The second polypeptide domain may have transcription release factor activity. The second polypeptide domain may have eukaryotic release factor 1 (ERF1) activity or eukaryotic release factor 3 (ERF3) activity.

[0177] (4) Histone Modification Activity

[0178] The second polypeptide domain may have histone modification activity. The second polypeptide domain may have histone deacetylase, histone acetyltransferase, histone demethylase, or histone methyltransferase activity. The histone acetyltransferase may be p300 or CREB-binding protein (CBP) protein, or fragments thereof. For example, the fusion protein may be dCas9-p300.

[0179] (5) Nuclease Activity

[0180] The second polypeptide domain may have nuclease activity that is different from the nuclease activity of the Cas9 protein. A nuclease, or a protein having nuclease activity, is an enzyme capable of cleaving the phosphodiester bonds between the nucleotide subunits of nucleic acids. Nucleases are usually further divided into endonucleases and exonucleases, although some of the enzymes may fall in both categories. Well known nucleases are deoxyribonuclease and ribonuclease.

[0181] (6) Nucleic Acid Association Activity

[0182] The second polypeptide domain may have nucleic acid association activity or nucleic acid binding protein-DNA-binding domain (DBD) is an independently folded protein domain that contains at least one motif that recognizes double- or single-stranded DNA. A DBD can recognize a specific DNA sequence (a recognition sequence) or have a general affinity to DNA. nucleic acid association region selected from the group consisting of helix-turn-helix region, leucine zipper region, winged helix region, winged helix-turn-helix region, helix-loop-helix region, immunoglobulin fold, B3 domain, Zinc finger, HMG-box, Wor3 domain, TAL effector DNA-binding domain.

[0183] (7) Methylase Activity

[0184] The second polypeptide domain may have methylase activity, which involves transferring a methyl group to DNA, RNA, protein, small molecule, cytosine or adenine. The second polypeptide domain may include a DNA methyltransferase.

[0185] (8) Demethylase Activity

[0186] The second polypeptide domain may have demethylase activity. The second polypeptide domain may include an enzyme that remove methyl (CH3-) groups from nucleic acids, proteins (in particular histones), and other molecules. Alternatively, the second polypeptide may covert the methyl group to hydroxymethylcytosine in a mechanism for demethylating DNA. The second polypeptide may catalyze this reaction. For example, the second polypeptide that catalyzes this reaction may be Tet1.

[0187] c. gRNA

[0188] The gRNA provides the targeting of the CRISPR/Cas9-based system. The gRNA is a fusion of two noncoding RNAs: a crRNA and a tracrRNA. The sgRNA may target any desired DNA sequence by exchanging the sequence encoding a 20 bp protospacer which confers targeting specificity through complementary base pairing with the desired DNA target. gRNA mimics the naturally occurring crRNA: tracrRNA duplex involved in the Type II Effector system. This duplex, which may include, for example, a 42-nucleotide crRNA and a 75-nucleotide tracrRNA, acts as a guide for the Cas9 to cleave the target nucleic acid. The "target region", "target sequence" or "protospacer" as used interchangeably herein refers to the region of the target gene to which the CRISPR/Cas9-based system targets. The CRISPR/Cas9-based system may include at least one gRNA, wherein the gRNAs target different DNA sequences. The target DNA sequences may be overlapping. The target sequence or protospacer is followed by a PAM sequence at the 3' end of the protospacer. Different Type II systems have differing PAM requirements. For example, the *Streptococcus pyogenes* Type II system uses an "NGG" sequence, where "N" can be any nucleotide.

[0189] The number of gRNA administered to the cell may be at least 1 gRNA, at least 2 different gRNA, at least 3 different gRNA at least 4 different gRNA, at least 5 different gRNA, at least 6 different gRNA, at least 7 different gRNA, at least 8 different gRNA, at least 9 different gRNA, at least

10 different gRNAs, at least 11 different gRNAs, at least 12 different gRNAs, at least 13 different gRNAs, at least 14 different gRNAs, at least 15 different gRNAs, at least 16 different gRNAs, at least 17 different gRNAs, at least 18 different gRNAs, at least 18 different gRNAs, at least 20 different gRNAs, at least 25 different gRNAs, at least 30 different gRNAs, at least 35 different gRNAs, at least 40 different gRNAs, at least 45 different gRNAs, or at least 50 different gRNAs. The number of gRNA administered to the cell may be between at least 1 gRNA to at least 50 different gRNAs, at least 1 gRNA to at least 45 different gRNAs, at least 1 gRNA to at least 40 different gRNAs, at least 1 gRNA to at least 35 different gRNAs, at least 1 gRNA to at least 30 different gRNAs, at least 1 gRNA to at least 25 different gRNAs, at least 1 gRNA to at least 20 different gRNAs, at least 1 gRNA to at least 16 different gRNAs, at least 1 gRNA to at least 12 different gRNAs, at least 1 gRNA to at least 8 different gRNAs, at least 1 gRNA to at least 4 different gRNAs, at least 4 gRNAs to at least 50 different gRNAs, at least 4 different gRNAs to at least 45 different gRNAs, at least 4 different gRNAs to at least 40 different gRNAs, at least 4 different gRNAs to at least 35 different gRNAs, at least 4 different gRNAs to at least 30 different gRNAs, at least 4 different gRNAs to at least 25 different gRNAs, at least 4 different gRNAs to at least 20 different gRNAs, at least 4 different gRNAs to at least 16 different gRNAs, at least 4 different gRNAs to at least 12 different gRNAs, at least 4 different gRNAs to at least 8 different gRNAs, at least 8 different gRNAs to at least 50 different gRNAs, at least 8 different gRNAs to at least 45 different gRNAs, at least 8 different gRNAs to at least 40 different gRNAs, at least 8 different gRNAs to at least 35 different gRNAs, 8 different gRNAs to at least 30 different gRNAs, at least 8 different gRNAs to at least 25 different gRNAs, 8 different gRNAs to at least 20 different gRNAs, at least 8 different gRNAs to at least 16 different gRNAs, or 8 different gRNAs to at least 12 different gRNAs.

[0190] The gRNA may comprise a complementary polynucleotide sequence of the target DNA sequence followed by a PAM sequence. The gRNA may comprise a "G" at the 5' end of the complementary polynucleotide sequence. The gRNA may comprise at least a 10 base pair, at least a 11 base pair, at least a 12 base pair, at least a 13 base pair, at least a 14 base pair, at least a 15 base pair, at least a 16 base pair, at least a 17 base pair, at least a 18 base pair, at least a 19 base pair, at least a 20 base pair, at least a 21 base pair, at least a 22 base pair, at least a 23 base pair, at least a 24 base pair, at least a 25 base pair, at least a 30 base pair, or at least a 35 base pair complementary polynucleotide sequence of the target DNA sequence followed by a PAM sequence. The PAM sequence may be "NGG", where "N" can be any nucleotide. The gRNA may target at least one of the promoter region, the enhancer region or the transcribed region of the target gene. The gRNA may include a nucleic acid sequence of at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, 585-625, 462 (FIG. **40**), 464 (FIG. **41**), and 465 (FIG. **41**).

[0191] The gRNA may target any nucleic acid sequence. The nucleic acid sequence target may be DNA. The DNA may be any gene. For example, the gRNA may target a gene, such as BRN2, MYT1L, ASCL1, NANOG, VEGFA, TERT, IL1B, IL1R2, IL1RN, HBG1, HBG2, MYOD1, OCT4, and DMD.

[0192] (1) Dystrophin

[0193] Dystrophin is a rod-shaped cytoplasmic protein which is a part of a protein complex that connects the cytoskeleton of a muscle fiber to the surrounding extracellular matrix through the cell membrane. Dystrophin provides structural stability to the dystroglycan complex of the cell membrane. The dystrophin gene is 2.2 megabases at locus Xp21. The primary transcription measures about 2,400 kb with the mature mRNA being about 14 kb. 79 exons code for the protein which is over 3500 amino acids. Normal skeleton muscle tissue contains only small amounts of dystrophin but its absence of abnormal expression leads to the development of severe and incurable symptoms. Some mutations in the dystrophin gene lead to the production of defective dystrophin and severe dystrophic phenotype in affected patients. Some mutations in the dystrophin gene lead to partially-functional dystrophin protein and a much milder dystrophic phenotype in affected patients.

[0194] DMD is the result of inherited or spontaneous mutations that cause nonsense or frame shift mutations in the dystrophin gene. Naturally occurring mutations and their consequences are relatively well understood for DMD. It is known that in-frame deletions that occur in the exon 45-55 region contained within the rod domain can produce highly functional dystrophin proteins, and many carriers are asymptomatic or display mild symptoms. Furthermore, more than 60% of patients may theoretically be treated by targeting exons in this region of the dystrophin gene. Efforts have been made to restore the disrupted dystrophin reading frame in DMD patients by skipping non-essential exons during mRNA splicing to produce internally deleted but functional dystrophin proteins. The deletion of internal dystrophin exons retain the proper reading frame but cause the less severe Becker muscular dystrophy.

[0195] (2) CRISPR/Cas9-Based System for Targeting Dystrophin

[0196] A CRISPR/Cas9-based system specific for dystrophin gene are disclosed herein. The CRISPR/Cas9-based system may include Cas9 and at least one gRNA to target the dystrophin gene. The CRISPR/Cas9-based system may bind and recognize a target region. The target regions may be chosen immediately upstream of possible out-of-frame stop codons such that insertions or deletions during the repair process restore the dystrophin reading frame by frame conversion. Target regions may also be splice acceptor sites or splice donor sites, such that insertions or deletions during the repair process disrupt splicing and restore the dystrophin reading frame by splice site disruption and exon exclusion. Target regions may also be aberrant stop codons such that insertions or deletions during the repair process restore the dystrophin reading frame by eliminating or disrupting the stop codon.

[0197] Single or multiplexed sgRNAs may be designed to restore the dystrophin reading frame by targeting the mutational hotspot at exons 45-55 and introducing either intra-exonic small insertions and deletions, or large deletions of one or more exons. Following treatment with Cas9 and one or more sgRNAs, dystrophin expression may be restored in Duchenne patient muscle cells in vitro. Human dystrophin was detected in vivo following transplantation of genetically corrected patient cells into immunodeficient mice. Significantly, the unique multiplex gene editing capabilities of the CRISPR/Cas9 system enable efficiently generating large deletions of this mutational hotspot region that can correct

up to 62% of patient mutations by universal or patient-specific gene editing approaches.

[0198] The CRISPR/Cas9-based system may use gRNA of varying sequences and lengths. Examples of gRNAs may be found in Table 6. The CRISPR/Cas9-based system may target a nucleic acid sequence of SEQ ID NOs: 65-144, or a complement thereof. The gRNA may include a nucleotide sequence selected from the group consisting of SEQ ID NO: 65-144, or a complement thereof. For example, the disclosed CRISPR/Cas9-based systems were engineered to mediate highly efficient gene editing at exon 51 of the dystrophin gene. These CRISPR/Cas9-based systems restored dystrophin protein expression in cells from DMD patients.

[0199] (a) Exons 51 and 45-55

[0200] Exon 51 is frequently adjacent to frame-disrupting deletions in DMD. Elimination of exon 51 from the dystrophin transcript by exon skipping can be used to treat approximately 15% of all DMD patients. This class of DMD mutations is ideally suited for permanent correction by NIEJ-based genome editing and HDR. The CRISPR/Cas9-based systems described herein have been developed for targeted modification of exon 51 in the human dystrophin gene. These CRISPR/Cas9-based systems were transfected into human DMD cells and mediated efficient gene modification and conversion to the correct reading frame. Protein restoration was concomitant with frame restoration and detected in a bulk population of CRISPR/Cas9-based system-treated cells. Similarly, the elimination of exons 45-55 of the dystrophin transcript can be used to treat approximately 62% of all DMD patients.

[0201] (3) AAV/CRISPR Constructs

[0202] AAV may be used to deliver CRISPRs using various construct configurations (see FIG. **39**). For example, AAV may deliver Cas9 and gRNA expression cassettes on separate vectors. Alternatively, if the small Cas9 proteins, derived from species such as *Staphylococcus aureus* or *Neisseria meningitidis*, are used then both the Cas9 and up to two gRNA expression cassettes may be combined in a single AAV vector within the 4.7 kb packaging limit (see FIG. **39**).

5. MULTIPLEX CRISPR/CAS9-BASED SYSTEM

[0203] The present disclosure is directed to a multiplex CRISPR/Cas9-Based System which includes a CRISPR/CRISPR-associated (Cas) 9-based system, such as Cas9 or dCas9, and multiple gRNAs to target one or more endogenous genes. This platform utilizes a convenient Golden Gate cloning method to rapidly incorporate up to four independent sgRNA expression cassettes into a single lentiviral vector. Each sgRNA was efficiently expressed and could mediate multiplex gene editing at diverse loci in immortalized and primary human cell lines. Transient transcriptional activation in cell lines stably expressing dCas9-VP64 was demonstrated to be tunable by synergistic activation with one to four sgRNAs. Furthermore, the single lentiviral vector can induce sustained and long-term endogenous gene expression in immortalized and primary human cells. This system allows for rapid assembly of a single lentiviral vector that enables efficient multiplex gene editing or activation in model and primary cell lines.

[0204] The multiplex CRISPR/Cas9-Based System provides potency of transcriptional activation and tunable induction of transcriptional activation. Readily generated by Golden Gate assembly, the final vector expresses a constitutive Cas9 or dCas9-VP64 in addition to one, two, three, or four sgRNAs expressed from independent promoters. Each promoter is capable of efficiently expressing sgRNAs that direct similar levels of Cas9 nuclease activity. Furthermore, lentiviral delivery of a single vector expressing Cas9 and four sgRNAs targeting independent loci resulted in simultaneous multiplex gene editing of all four loci. Tunable transcriptional activation at two endogenous genes in both transient and stable contexts was achieved using lentiviral delivery of Cas9 with or without sgRNAs. Highly efficient and long-term gene activation in primary human cells is accomplished. This system is therefore an attractive and efficient method to generate multiplex gene editing and long-term transcriptional activation in human cells.

[0205] The multiplex CRISPR/Cas9-Based System allows efficient multiplex gene editing for simultaneously inactivating multiple genes. The CRISPR/Cas9 system can simultaneously target multiple distinct genomic loci by co-expressing a single Cas9 protein with two or more sgRNAs, making this system uniquely suited for multiplex gene editing or synergistic activation applications. The CRISPR/Cas9 system greatly expedites the process of molecular targeting to new sites by simply modifying the expressed sgRNA molecule. The single lentiviral vector may be combined with methods for achieving inducible control of these components, either by chemical or optogenetic regulation, to facilitate investigation of the dynamics of gene regulation in both time and space.

[0206] The multiplex CRISPR/Cas9-based systems may transcriptionally activate two or more endogenous genes. The multiplex CRISPR/Cas9-based systems may transcriptionally repress two or more endogenous genes. For example, at least two endogenous genes, at least three endogenous genes, at least four endogenous genes, at least five endogenous genes, or at least ten endogenous genes may be activated or repressed by the multiplex CRISPR/Cas9-based system. Between two and fifteen genes, between two and ten genes, between two and five genes, between five and fifteen genes, or between five and ten genes may be activated or repressed by the multiplex CRISPR/Cas9-based system.

[0207] (1) Modified Lentiviral Vector

[0208] The multiplex CRISPR/Cas9-based system includes a modified lentiviral vector. The modified lentiviral vector includes a first polynucleotide sequence encoding a fusion protein and a second polynucleotide sequence encoding at least one sgRNA. The fusion protein may be the fusion protein of the CRISPR/Cas9-based system, as described above. The first polynucleotide sequence may be operably linked to a promoter. The promoter may be a constitutive promoter, an inducible promoter, a repressible promoter, or a regulatable promoter.

[0209] The second polynucleotide sequence encodes at least 1 sgRNA. For example, the second polynucleotide sequence may encode at least 1 sgRNA, at least 2 sgRNAs, at least 3 sgRNAs, at least 4 sgRNAs, at least 5 sgRNAs, at least 6 sgRNAs, at least 7 sgRNAs, at least 8 sgRNAs, at least 9 sgRNAs, at least 10 sgRNAs, at least 11 sgRNA, at least 12 sgRNAs, at least 13 sgRNAs, at least 14 sgRNAs, at least 15 sgRNAs, at least 16 sgRNAs, at least 17 sgRNAs, at least 18 sgRNAs, at least 19 sgRNAs, at least 20 sgRNAs, at least 25 sgRNA, at least 30 sgRNAs, at least 35 sgRNAs, at least 40 sgRNAs, at least 45 sgRNAs, or at least 50 sgRNAs. The second polynucleotide sequence may encode between 1 sgRNA and 50 sgRNAs, between 1 sgRNA and

45 sgRNAs, between 1 sgRNA and 40 sgRNAs, between 1 sgRNA and 35 sgRNAs, between 1 sgRNA and 30 sgRNAs, between 1 sgRNA and 25 different sgRNAs, between 1 sgRNA and 20 sgRNAs, between 1 sgRNA and 16 sgRNAs, between 1 sgRNA and 8 different sgRNAs, between 4 different sgRNAs and 50 different sgRNAs, between 4 different sgRNAs and 45 different sgRNAs, between 4 different sgRNAs and 40 different sgRNAs, between 4 different sgRNAs and 35 different sgRNAs, between 4 different sgRNAs and 30 different sgRNAs, between 4 different sgRNAs and 25 different sgRNAs, between 4 different sgRNAs and 20 different sgRNAs, between 4 different sgRNAs and 16 different sgRNAs, between 4 different sgRNAs and 8 different sgRNAs, between 8 different sgRNAs and 50 different sgRNAs, between 8 different sgRNAs and 45 different sgRNAs, between 8 different sgRNAs and 40 different sgRNAs, between 8 different sgRNAs and 35 different sgRNAs, between 8 different sgRNAs and 30 different sgRNAs, between 8 different sgRNAs and 25 different sgRNAs, between 8 different sgRNAs and 20 different sgRNAs, between 8 different sgRNAs and 16 different sgRNAs, between 16 different sgRNAs and 50 different sgRNAs, between 16 different sgRNAs and 45 different sgRNAs, between 16 different sgRNAs and 40 different sgRNAs, between 16 different sgRNAs and 35 different sgRNAs, between 16 different sgRNAs and 30 different sgRNAs, between 16 different sgRNAs and 25 different sgRNAs, or between 16 different sgRNAs and 20 different sgRNAs. Each of the polynucleotide sequences encoding the different sgRNAs may be operably linked to a promoter. The promoters that are operably linked to the different sgRNAs may be the same promoter. The promoters that are operably linked to the different sgRNAs may be different promoters. The promoter may be a constitutive promoter, an inducible promoter, a repressible promoter, or a regulatable promoter.

[0210] The at least one sgRNA may bind to a target gene or loci. If more than one sgRNA is included, each of the sgRNAs binds to a different target region within one target loci or each of the sgRNA binds to a different target region within different gene loci. The fusion protein may include Cas9 protein or iCas9-VP64 protein. The fusion protein may include a VP64 domain, a p300 domain, or a KRAB domain.

6. SITE-SPECIFIC NUCLEASES

[0211] The composition, as described above, includes a nucleotide sequence encoding a site-specific nuclease that binds and cleaves a target region. The site-specific nuclease may be engineered. For example, an engineered site-specific nuclease may be a CRISPR/Cas9-based system, a ZFN, or a TALEN. The site-specific nuclease may bind and cleave a gene or locus in the genome of a cell in the skeletal muscle or cardiac muscle. For example, the gene or locus may be the Rosa26 locus or the dystrophin gene.

[0212] a. CRISPR/Cas9-based system

[0213] The CRISPR/Cas9-based system, as described above, may be used to introduce site-specific double strand breaks at targeted genomic loci.

[0214] b. Zinc Finger Nucleases (ZFN)

[0215] The site-specific nuclease may be a ZFN. A single zinc finger contains approximately 30 amino acids and the domain functions by binding 3 consecutive base pairs of DNA via interactions of a single amino acid side chain per base pair. The modular structure of the zinc finger motif permits the conjunction of several domains in series, allowing for the recognition and targeting of extended sequences in multiples of 3 nucleotides. These targeted DNA-binding domains can be combined with a nuclease domain, such as FokI, to generate a site-specific nuclease, called a"zinc finger nuclease" (ZFNs) that can be used to introduce site-specific double strand breaks at targeted genomic loci. This DNA cleavage stimulates the natural DNA-repair machinery, leading to one of two possible repair pathways, NTEJ andHFIDR. For example, the ZFN may target the Rosa26 locus (Perez-Pinera et al. *Nucleic Acids Research* (2012) 40:3741-3752) or a dystrophin gene. Examples of ZFNs are shown in Table 1and FIGS. **35-38**. In Table 1, the DNA recognition helices are underlined and "Fok ELD-S" and "Fok KKR-S" refers to the FokI nuclease domain that is fused to the zinc finger protein DNA-binding domains. In FIGS. **35-38**, the target DNA sequence in the target sites (i.e., in SEQ ID NOs: 442, 445, 448, and 453) and the DNA recognition helices in the ZFN amino acid sequences (i.e., in SEQ ID NOs: 443, 444, 446, 447, 449-452, 454, and 455) are underlined, respectively.

TABLE 1

Full amino acid sequences of identified ZFNs.

ZFN B left Fok ELD-S full amino acid sequence (SEQ ID NO: 438)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGK

SFSRKDALRGHQRTHTGEKPYKCPECGKSFSHRTTLTNHQRTHTGEKPY

KCPECGKSFSQRNALAGHQRTHTGEKPYKCPECGKSFSHKNALQNHQRT

HTGEKPYKCPECGKSFSDPGHLVRHQRTHTGEKPYKCPECGKSFSTSGN

LVRHQRTHTGAAARALVKSELEEKKSELRHKLKYVPHEYIELIEIARNP

TQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVD

TKAYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEWWKVYPSSVTEFK

FLFVSGHFKGNYKAQLTRLNHITNCNGAVLSVEELLIGGEMIKAGTLTL

EEVRRKFNNGEINF*

ZFN B right Fok KKR-S full amino acid sequence (SEQ ID NO: 439)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGEKPYKCPECGK

SFSQQRSLVGHQRTHTGEKPYKCPECGKSFSDKKDLTRHQRTHTGEKPY

KCPECGKSFSTSGHLVRHQRTHTGEKPYKCPECGKSFSQRAHLERHQRT

HTGEKPYKCPECGKSFSTSGSLVRHQRTHTGEKPYKCPECGKSFSTSGN

LVRHQRTHTGAAARALVKSELEEKKSELRHKLKYVPHEYIELIEIARNP

TQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYTVGSPIDYGVIVD

TKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEWWKVYPSSVTEFK

FLFVSGHFKGNYKAQLTRLNRKTNCNGAVLSVEELLIGGEMIKAGTLTL

EEVRRKFNNGEINF*

ZFN J left Fok KKR-S full amino acid sequence (SEQ ID NO: 440)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGERPFQCRICMR

NFSSKQALAVHTRTHTGEKPFQCRICMRNFSQSTTLKRHLRTHTGEKPF

QCRICMRNFSRSDHLSLHLKTHLRGSQLVKSELEEKKSELRHKLKYVPH

TABLE 1-continued

Full amino acid sequences of identified ZFNs.

EYIELIEIARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYT

VGSPIDYGVIVDTKAYSGGYNLPIGQADEMQRYVKENQTRNKHINPNEW

WKVYPSSVTEFKFLFVSGHFKGNYKAQLTRLNRKTNCNGAVLSVEELLI

GGEMIKAGTLTLEEVRRKFNNGEINF*

ZFN J right Fok ELD-S full amino acid sequence (SEQ ID NO: 441)

MDYKDHDGDYKDHDIDYKDDDDKMAPKKKRKVGRLEPGERPEQCRICMR

NESRRAHLQNHTRTHTGEKPEQCRICMRNESQSTTLKRHLRTHTGEKPE

QCRICMRNESDGGHLTRHLKTHLRGSQLVKSELEEKKSELRHKLKYVPH

EYIELIETARNPTQDRILEMKVMEFFMKVYGYRGEHLGGSRKPDGAIYT

VGSPIDYGVIVDTKAYSGGYNLPIGQADEMERYVEENQTRDKHLNPNEW

WKVYPSSVTEFKFLFVSGHEKGNYKAQLTRLNHITNCNGAVLSVEELLI

GGEMIKAGTLTLEEVRRKFNNGEINF*

[0216] c. TAL Effector Nucleases (TALENs)

[0217] TALENs may be used to introduce site-specific double strand breaks at targeted genomic loci. Site-specific double-strand breaks are created when two independent TALENs bind to nearby DNA sequences, thereby permitting dimerization of FokI and cleavage of the target DNA. TALENs have advanced genome editing due to their high rate of successful and efficient genetic modification. This DNA cleavage may stimulate the natural DNA-repair machinery, leading to one of two possible repair pathways: homology-directed repair (HDR) or the non-homologous end joining (NHEJ) pathway. The TALENs may be designed to target any gene involved in a genetic disease.

[0218] The TALENs may include a nuclease and a TALE DNA-binding domain that binds to the target gene in a TALEN target region. The target gene may have a mutation such as a frameshift mutation or a nonsense mutation. If the target gene has a mutation that causes a premature stop codon, the TALEN may be designed to recognize and bind a nucleotide sequence upstream or downstream from the premature stop codon. A "TALEN target region" includes the binding regions for two TALENs and the spacer region, which occurs between the binding regions. The two TALENs bind to different binding regions within the TALEN target region, after which the TALEN target region is cleaved. Examples of TALENs are described in International Patent Application No. PCT/US2013/038536, which is incorporated by reference in its entirety.

7. TRANSCRIPTIONAL ACTIVATORS

[0219] The composition, as described above, includes a nucleotide sequence encoding a transcriptional activator that activates a target gene. The transcriptional activator may be engineered. For example, an engineered transcriptional activator may be a CRISPR/Cas9-based system, a zinc finger fusion protein, or a TALE fusion protein.

[0220] a. CRISPR/Cas9-Based System

[0221] The CRISPR/Cas9-based system, as described above, may be used to activate transcription of a target gene with RNA. The CRISPR/Cas9-based system may include a fusion protein, as described above, wherein the second polypeptide domain has transcription activation activity or histone modification activity. For example, the second polypeptide domain may include VP64 or p300.

[0222] b. Zinc Finger Fusion Proteins

[0223] The transcriptional activator may be a zinc finger fusion protein. The zinc finger targeted DNA-binding domains, as described above, can be combined with a domain that has transcription activation activity or histone modification activity. For example, the domain may include VP64 or p300.

[0224] c. TALE Fusion Proteins

[0225] TALE fusion proteins may be used to activate transcription of a target gene. The TALE fusion protein may include a TALE DNA-binding domain and a domain that has transcription activation activity or histone modification activity. For example, the domain may include VP64 or p300.

8. COMPOSITIONS

[0226] The present invention is directed to a composition for altering gene expression and engineering or altering genomic DNA in a cell or subject. The composition may also include a viral delivery system.

[0227] a. Compositions for Genome Editing in Muscle

[0228] The present invention is directed to a composition for genome editing a target gene in skeletal muscle or cardiac muscle of a subject. The composition includes a modified AAV vector and a nucleotide sequence encoding a site-specific nuclease. The composition delivers active forms of site-specific nucleases to skeletal muscle or cardiac muscle. The composition may further comprise a donor DNA or a transgene. These compositions may be used in genome editing, genome engineering, and correcting or reducing the effects of mutations in genes involved in genetic diseases and/or other skeletal or cardiac muscle conditions.

[0229] The target gene may be involved in differentiation of a cell or any other process in which activation, repression, or disruption of a gene may be desired, or may have a mutation such as a deletion, frameshift mutation, or a nonsense mutation. If the target gene has a mutation that causes a premature stop codon, an aberrant splice acceptor site or an aberrant splice donor site, the site-specific nucleases may be designed to recognize and bind a nucleotide sequence upstream or downstream from the premature stop codon, the aberrant splice acceptor site or the aberrant splice donor site. The site-specific nucleases may also be used to disrupt normal gene splicing by targeting splice acceptors and donors to induce skipping of premature stop codons or restore a disrupted reading frame. The site-specific nucleases may or may not mediate off-target changes to protein-coding regions of the genome.

[0230] b. Adeno-Associated Virus Vectors

[0231] The composition, as described above, includes a modified adeno-associated virus (AAV) vector. The modified AAV vector may have enhanced cardiac and skeletal muscle tissue tropism. The modified AAV vector may be capable of delivering and expressing the site-specific nuclease in the cell of a mammal. For example, the modified AAV vector may be an AAV-SASTG vector (Piacentino et al. (2012) Human Gene Therapy 23:635-646). The modified AAV vector may deliver nucleases to skeletal and cardiac muscle in vivo. The modified AAV vector may be based on

one or more of several capsid types, including AAV1, AAV2, AAV5, AAV6, AAV8, and AAV9. The modified AAV vector may be based on AAV2 pseudotype with alternative muscle-tropic AAV capsids, such as AAV2/1, AAV2/6, AAV2/7, AAV2/8, AAV2/9, AAV2.5 and AAV/SASTG vectors that efficiently transduce skeletal muscle or cardiac muscle by systemic and local delivery (Seto et al. Current Gene Therapy (2012) 12:139-151).

[0232] c. CRISPR/Cas9-Based System

[0233] The present disclosure also provides DNA targeting systems or compositions of at least one CRISPR/Cas9-based system, as described above. These compositions may be used in genome editing, genome engineering, and correcting or reducing the effects of mutations in genes involved in genetic diseases. The composition includes a CRISPR/Cas9-based system that includes a Cas9 protein or Cas9 fusion protein, as described above. The CRISPR/Cas9-based system may also include at least one gRNA, as described above.

[0234] d. Multiplex CRISPR/Cas9-Based System

[0235] The present disclosure also provides multiplex CRISPR/Cas9-based systems, as described above. These compositions may be used in genome editing, genome engineering, and correcting or reducing the effects of mutations in genes involved in genetic diseases. These compositions may be used to target more than one gene. The composition includes a modified lentiviral vector comprising a CRISPR/Cas9-based system that includes a Cas9 protein or Cas9 fusion protein, as described above and more than one gRNA, as described above.

9. METHODS OF USES

[0236] Potential applications of the compositions are diverse across many areas of science and biotechnology. The disclosed compositions may be used to repair genetic mutations that cause disease. The disclosed compositions may be used to disrupt genes such that gene disruption leads to increases in muscle regeneration or muscle strength, or decreases in muscle aging. The disclosed compositions may be used to introduce therapeutic genes to be expressed systemically from skeletal muscle or cardiac muscle, such as clotting factors or monoclonal antibodies. The disclosed compositions may be used to modulate mammalian gene expression. The disclosed compositions may be used to transdifferentiate or induce the differentiation of a cell or correct a mutant gene in a cell. Examples of activation of genes related to cell and gene therapy, genetic reprogramming, and regenerative medicine are provided. RNA-guided transcriptional activators may be used to reprogram cell lineage specification. Activation of endogenous genes encoding the key regulators of cell fate, rather than forced overexpression of these factors, may potentially lead to more rapid, efficient, stable, or specific methods for genetic reprogramming, transdifferentiation, and/or induced differentiation.

10. METHODS OF GENOME EDITING IN MUSCLE

[0237] The present disclosure is directed to a method of genome editing in a skeletal muscle or cardiac muscle of a subject. The method comprises administering to the skeletal muscle or cardiac muscle of the subject the composition for genome editing in skeletal muscle or cardiac muscle, as described above. The genome editing may include correcting a mutant gene or inserting a transgene. Correcting the mutant gene may include deleting, rearranging, or replacing the mutant gene. Correcting the mutant gene may include nuclease-mediated NHEJ or HDR.

11. METHODS OF USING CRISPR/CAS9-BASED SYSTEM

[0238] Potential applications of the CRISPR/Cas9-based system are diverse across many areas of science and biotechnology. The disclosed CRISPR/Cas9-based systems may be used to modulate mammalian gene expression. The disclosed CRISPR/Cas9-based systems may be used to transdifferentiate or induce differentiation of a cell or correct a mutant gene in a cell. Examples of activation of genes related to cell and gene therapy, genetic reprogramming, and regenerative medicine are provided. RNA-guided transcriptional activators may be used to reprogram cell lineage specification. Although reprogramming was incomplete and inefficient in these experiments, there are many ways by which this method could be improved, including repeated transfections of iCas9-VP64/gRNA combinations, stable expression of these factors, and targeting multiple genes, such as Brn2 and Mytl1 in addition to Ascl1 for transdifferentiation into a neuronal phenotype. Activation of endogenous genes encoding the key regulators of cell fate, rather than forced overexpression of these factors, may potentially lead to more rapid, efficient, stable, or specific methods for genetic reprogramming and transdifferentiation or induced differentiation of a cell. Finally, Cas9 fusions to other domains, including repressive and epigenetic-modifying domains, could provide a greater diversity of RNA-guided transcriptional regulators to complement other RNA-based tools for mammalian cell engineering.

[0239] a. Methods of Activating Gene Expression

[0240] The present disclosure provides a mechanism for activating the expression of endogenous genes, such as mammalian genes, based on targeting a transcriptional activator to promoters via RNA using a CRISPR/Cas9 based system, as described above. This is fundamentally different from previously described methods based on engineering sequence-specific DNA-binding proteins and may provide opportunities for targeted gene regulation. Because the generation of gRNA expression plasmids simply involves synthesizing two short custom oligonucleotides and one cloning step, it is possible to generate many new gene activators quickly and economically. The gRNAs can also be transfected directly to cells following in vitro transcription. Multiple gRNAs targeted to single promoters were shown, but simultaneous targeting of multiple promoters could also be possible. Recognition of genomic target sites with RNAs, rather than proteins, may also circumvent limitations of targeting epigenetically modified sites, such as methylated DNA.

[0241] In contrast to current methods based on engineering DNA-binding proteins, Cas9 fused to a transcriptional activation domain can be targeted by combinations of guide RNA molecules to induce the expression of endogenous human genes. This straightforward and versatile approach for targeted gene activation circumvents the need for engineering new proteins and allows for widespread synthetic gene regulation.

[0242] The method may include administering to a cell or subject a CRISPR/Cas9-based system, a polynucleotide or

vector encoding said CRISPR/Cas9-based system, or DNA targeting systems or compositions of at least one CRISPR/Cas9-based system, as described above. The method may include administering a CRISPR/Cas9-based system, such as administering a Cas9 fusion protein containing transcription activation domain or a nucleotide sequence encoding said Cas9 fusion protein. The Cas9 fusion protein may include a transcription activation domain such as aVP16 protein or a transcription co-activator such as a p300 protein.

[0243] (1) dCas9-VP16

[0244] The Cas9 fusion protein may include a transcription activation domain such as aVP16 protein. The transcription activation domain may contain at least 1 VP16 protein, at least 2 VP16 proteins, at least 3 VP16 proteins, at least 4 VP16 proteins (i.e., a VP64 activator domain), at least 5 VP16 proteins, at least 6 VP16 proteins, at least 6 VP16 proteins, or at least 10 VP16 proteins. The Cas9 protein may be a Cas9 protein in which the nuclease activity is inactivated. For example, the Cas9 protein in the fusion protein may be iCas9 (amino acids 36-1403 of SEQ ID NO: 1), which includes the amino acid substitutions of D10A and H840A. The Cas9 fusion protein may be iCas9-VP64.

[0245] (2) dCas9-p300

[0246] The Cas9 fusion protein may include a transcription co-activation domain such as a p300 protein. The p300 protein (also known as EP300 or E1A binding protein p300) is encoded by the EP300 gene and regulates the activity of many genes in tissues throughout the body. The p300 protein plays a role in regulating cell growth and division, prompting cells to mature and assume specialized functions (differentiate) and preventing the growth of cancerous tumors. The p300 protein activates transcription by connecting transcription factors with a complex of proteins that carry out transcription in the cell's nucleus. The p300 interaction with transcription factors is managed by one or more of p300 domains: the nuclear receptor interaction domain (RID), the CREB and MYB interaction domain (KIX), the cysteine/histidine regions (TAZ1/CH and TAZ2/CH3) and the interferon response binding domain (IBiD). The last four domains, KIX, TAZ1, TAZ2 and IBiD of p300, each bind tightly to a sequence spanning both transactivation domains 9aaTADs of transcription factor p53. The protein functions as histone acetyltransferase that regulates transcription via chromatin remodeling, and is important in the processes of cell proliferation and differentiation. It mediates cAMP-gene regulation by binding specifically to phosphorylated CREB protein.

[0247] The p300 protein may activate Mothers against decapentaplegic homolog 7, MAF, TSG101, Peroxisome proliferator-activated receptor alpha, NPAS2, PAX6, DDX5, MYBL2, Mothers against decapentaplegic homolog 1, Mothers against decapentaplegic homolog 2, Lymphoid enhancer-binding factor 1, SNIP1, TRERF1, STAT3, EID1, RAR-related orphan receptor alpha, ELK1, HIF1A, ING5, Peroxisome proliferator-activated receptor gamma, SS18, TCF3, Zif268, Estrogen receptor alpha, GPS2, MyoD, YY1, ING4, PROX1, CITED1, HNF1A, MEF2C, MEF2D, MAML1, Twist transcription factor, PTMA, IRF2, DTX1, Flap structure-specific endonuclease 1, Myocyte-specific enhancer factor 2A, CDX2, BRCA1, HNRPU, STAT6, CITED2, RELA, TGS1, CEBPB, Mdm2, NCOA6, NFATC2, Thyroid hormone receptor alpha, BCL3, TFAP2A, PCNA, P53 and TAL1.

[0248] The transcription co-activation domain may include a human p300 protein or a fragment thereof. The transcription co-activation domain may include a wild-type human p300 protein or a mutant human p300 protein, or fragments thereof. The transcription co-activation domain may include the core lysine-acetyltranserase domain of the human p300 protein, i.e., the p300 HAT Core (also known as "p300 WT Core"; see FIG. **58**). The Cas9 protein may be a Cas9 protein in which the nuclease activity is inactivated. For example, the Cas9 protein in the fusion protein may be iCas9 (amino acids 36-1403 of SEQ ID NO: 1), which includes the amino acid substitutions of D10A and H840A. The Cas9 fusion protein may be iCas9-p300 WT Core.

[0249] (3) gRNA

[0250] The method may also include administering to a cell or subject a CRISPR/Cas9-based system at least one gRNA, wherein the gRNAs target different DNA sequences. The target DNA sequences may be overlapping. The number of gRNA administered to the cell may be at least 1 gRNA, at least 2 different gRNAs, at least 3 different gRNAs at least 4 different gRNAs, at least 5 different gRNAs, at least 6 different gRNAs, at least 7 different gRNAs, at least 8 different gRNAs, at least 9 different gRNAs, at least 10 different gRNAs, at least 11 different gRNAs, at least 12 different gRNAs, at least 13 different gRNAs, at least 14 different gRNAs, at least 15 different gRNAs, at least 16 different gRNAs, at least 17 different gRNAs, at least 18 different gRNAs, at least 18 different gRNAs, at least 20 different gRNAs, at least 25 different gRNAs, at least 30 different gRNAs, at least 35 different gRNAs, at least 40 different gRNAs, at least 45 different gRNAs, or at least 50 different gRNAs. The number of gRNA administered to the cell may be between at least 1 gRNA to at least 50 different gRNAs, at least 1 gRNA to at least 45 different gRNAs, at least 1 gRNA to at least 40 different gRNAs, at least 1 gRNA to at least 35 different gRNAs, at least 1 gRNA to at least 30 different gRNAs, at least 1 gRNA to at least 25 different gRNAs, at least 1 gRNA to at least 20 different gRNAs, at least 1 gRNA to at least 16 different gRNAs, at least 1 gRNA to at least 12 different gRNAs, at least 1 gRNA to at least 8 different gRNAs, at least 1 gRNA to at least 4 different gRNAs, at least 4 gRNAs to at least 50 different gRNAs, at least 4 different gRNAs to at least 45 different gRNAs, at least 4 different gRNAs to at least 40 different gRNAs, at least 4 different gRNAs to at least 35 different gRNAs, at least 4 different gRNAs to at least 30 different gRNAs, at least 4 different gRNAs to at least 25 different gRNAs, at least 4 different gRNAs to at least 20 different gRNAs, at least 4 different gRNAs to at least 16 different gRNAs, at least 4 different gRNAs to at least 12 different gRNAs, at least 4 different gRNAs to at least 8 different gRNAs, at least 8 different gRNAs to at least 50 different gRNAs, at least 8 different gRNAs to at least 45 different gRNAs, at least 8 different gRNAs to at least 40 different gRNAs, at least 8 different gRNAs to at least 35 different gRNAs, 8 different gRNAs to at least 30 different gRNAs, at least 8 different gRNAs to at least 25 different gRNAs, 8 different gRNAs to at least 20 different gRNAs, at least 8 different gRNAs to at least 16 different gRNAs, 8 different gRNAs to at least 12 different gRNAs, at least 8 different gRNAs to at least 8 different gRNAs,

[0251] The gRNA may comprise a complementary polynucleotide sequence of the target DNA sequence followed by NGG. The gRNA may comprise a "G" at the 5' end of the

complementary polynucleotide sequence. The gRNA may comprise at least a 10 base pair, at least a 11 base pair, at least a 12 base pair, at least a 13 base pair, at least a 14 base pair, at least a 15 base pair, at least a 16 base pair, at least a 17 base pair, at least a 18 base pair, at least a 19 base pair, at least a 20 base pair, at least a 21 base pair, at least a 22 base pair, at least a 23 base pair, at least a 24 base pair, at least a 25 base pair, at least a 30 base pair, or at least a 35 base pair complementary polynucleotide sequence of the target DNA sequence followed by NGG. The gRNA may target at least one of the promoter region, the enhancer region or the transcribed region of the target gene. The gRNA may include a nucleic acid sequence of at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625.

[0252] b. Methods of Repressing Gene Expression

[0253] The present disclosure provides a mechanism for repressing the expression of endogenous genes, such as mammalian genes, based on targeting genomic regulatory elements, such as distal enhancers, via RNA using a CRISPR/Cas9 based system, as described above. The Cas9 fusion protein may include a transcriptional repressor, such as the KRAB repressor. The Cas9 fusion protein may be dCas9-KRAB. The dCas9-KRAB may additionally affect epigenetic gene regulation by recruiting heterochromatin-forming factors to the targeted locus. The CRISPR/dCas9-KRAB system may be used to repress the transcription of genes, but can also be used to target genomic regulatory elements which were previously inaccessible by traditional repression methods such as RNA interference (FIG. **53**B). Delivering dCas9-KRAB with gRNAs targeted to a distal enhancer may disrupt expression of multiple genes regulated by the targeted enhancer (see FIG. **53**C). The targeted enhancer may be any enhancer for a gene such as the HS2 enhancer.

[0254] a. Methods of Transdifferentiation or Induced Differentiation

[0255] The present disclosure provides a mechanism for transdifferentiating or inducing differentiation of cells by activating endogenous genes via RNA using a CRISPR/Cas9-based system, as described above.

[0256] (1) Transdifferentiation

[0257] The CRISPR/Cas9-based system may be used to transdifferentiate cells. Transdifferentiation, also known as lineage reprogramming or direct conversion, is a process where cells convert from one differentiated cell type to another without undergoing an intermediate pluripotent state or progenitor cell type. It is a type of metaplasia, which includes all cell fate switches, including the interconversion of stem cells. Transdifferentiation of cells has potential uses for disease modeling, drug discovery, gene therapy and regenerative medicine. Activation of endogenous genes, such as BRN2, MYT1L, ASCL1, NANOG, and/or MYOD1, using the CRISPR/Cas9 based system described above may lead to transdifferentiation of several cell types, such as fibroblasts, cardiomyocytes, hepatocytes, chondrocytes, mesenchymal progenitor cells, hematopoetic stem cells, or smooth muscle cells, into neuronal and myogenic phenotypes, respectively.

[0258] (2) Inducing Differentiation

[0259] The CRISPR/Cas9-based system may be used to induce differentiation of cells, such as stem cells, cardiomyocytes, hepatocytes, chondrocytes, mesenchymal progenitor cells, hematopoetic stem cells, or smooth muscle cells. For example, stem cells, such as embryonic stem cells or pluripotent stem cells, may be induced to differentiate into muscle cells or vascular endothelial cell, i.e., induce neuronal or myogenic differentiation.

12. USES OF MULTIPLEX CRISPR/CAS9-BASED SYSTEM

[0260] The multiplex CRISPR/Cas9-Based System takes advantage of the simplicity and low cost of sgRNA design and may be helpful in exploiting advances in high-throughput genomic research using CRISPR/Cas9 technology. For example, the single lentiviral vectors described here are useful in expressing Cas9 and numerous sgRNAs in difficult cell lines, such as primary fibroblasts described here (FIG. **47**). The multiplex CRISPR/Cas9-Based System may be used in the same ways as the CRISPR/Cas9-Based System described above.

[0261] In addition to the described transcriptional activation and nuclease functionality, this system will be useful for expressing other novel Cas9-based effectors that control epigenetic modifications for diverse purposes, including interrogation of genome architecture and pathways of endogenous gene regulation. As endogenous gene regulation is a delicate balance between multiple enzymes, multiplexing Cas9 systems with different functionalities will allow for examining the complex interplay among different regulatory signals. The vector described here should be compatible with aptamer-modified sgRNAs and orthogonal Cas9s to enable independent genetic manipulations using a single set of sgRNAs.

[0262] The multiplex CRISPR/Cas9-Based System may be used to activate at least one endogenous gene in a cell. The method includes contacting a cell with the modified lentiviral vector. The endogenous gene may be transiently activated or stably activated. The endogenous gene may be transiently repressed or stably repressed. The fusion protein may be expressed at similar levels to the sgRNAs. The fusion protein may be expressed at different levels compared to the sgRNAs. The cell may be a primary human cell.

[0263] The multiplex CRISPR/Cas9-Based System may be used in a method of multiplex gene editing in a cell. The method includes contacting a cell with the modified lentiviral vector. The multiplex gene editing may include correcting a mutant gene or inserting a transgene. Correcting a mutant gene may include deleting, rearranging or replacing the mutant gene. Correcting the mutant gene may include nuclease-mediated non-homologous end joining or homology-directed repair. The multiplex gene editing may include deleting or correcting at least one gene, wherein the gene is an endogenous normal gene or a mutant gene. The multiplex gene editing may include deleting or correcting at least two genes. For example, at least two genes, at least three genes, at least four genes, at least five genes, at least six genes, at least seven genes, at least eight genes, at least nine genes, or at least ten genes may be deleted or corrected.

[0264] The multiplex CRISPR/Cas9-Based System may be used in a method of multiplex modulation of gene expression in a cell. The method includes contacting a cell with the modified lentiviral vector. The method may include modulating the gene expression levels of at least one gene. The gene expression of the at least one target gene is modulated when gene expression levels of the at least one target gene are increased or decreased compared to normal

gene expression levels for the at least one target gene. The gene expression levels may be RNA or protein levels.

13. METHODS OF CORRECTING A MUTANT GENE AND TREATING A SUBJECT

[0265] The present disclosure is also directed to a method of correcting a mutant gene in a subject. The method comprises administering to the skeletal muscle or cardiac muscle of the subject the composition for genome editing in skeletal muscle or cardiac muscle, as described above. Use of the composition to deliver the site-specific nuclease to the skeletal muscle or cardiac muscle may restore the expression of a full-functional or partially-functional protein with a repair template or donor DNA, which can replace the entire gene or the region containing the mutation. The site-specific nuclease may be used to introduce site-specific double strand breaks at targeted genomic loci. Site-specific double-strand breaks are created when the site-specific nuclease binds to a target DNA sequences, thereby permitting cleavage of the target DNA. This DNA cleavage may stimulate the natural DNA-repair machinery, leading to one of two possible repair pathways: homology-directed repair (HDR) or the non-homologous end joining (NHEJ) pathway.

[0266] The present disclosure is directed to genome editing with a site-specific nuclease without a repair template, which can efficiently correct the reading frame and restore the expression of a functional protein involved in a genetic disease. The disclosed site-specific nucleases may involve using homology-directed repair or nuclease-mediated non-homologous end joining (NHEJ)-based correction approaches, which enable efficient correction in proliferation-limited primary cell lines that may not be amenable to homologous recombination or selection-based gene correction. This strategy integrates the rapid and robust assembly of active site-specific nucleases with an efficient gene editing method for the treatment of genetic diseases caused by mutations in nonessential coding regions that cause frameshifts, premature stop codons, aberrant splice donor sites or aberrant splice acceptor sites.

[0267] a. Nuclease Mediated Non-Homologous End Joining

[0268] Restoration of protein expression from an endogenous mutated gene may be through template-free NHEJ-mediated DNA repair. In contrast to a transient method targeting the target gene RNA, the correction of the target gene reading frame in the genome by a transiently expressed site-specific nuclease may lead to permanently restored target gene expression by each modified cell and all of its progeny.

[0269] Nuclease mediated NHEJ gene correction may correct the mutated target gene and offers several potential advantages over the HDR pathway. For example, NHEJ does not require a donor template, which may cause non-specific insertional mutagenesis. In contrast to HDR, NHEJ operates efficiently in all stages of the cell cycle and therefore may be effectively exploited in both cycling and post-mitotic cells, such as muscle fibers. This provides a robust, permanent gene restoration alternative to oligonucleotide-based exon skipping or pharmacologic forced read-through of stop codons and could theoretically require as few as one drug treatment. NHEJ-based gene correction using a CRISPR/Cas9-based system, as well as other engineered nucleases including meganucleases and zinc finger nucleases, may be combined with other existing ex vivo and in vivo platforms for cell- and gene-based therapies, in addition to the plasmid electroporation approach described here. For example, delivery of a CRISPR/Cas9-based system by mRNA-based gene transfer or as purified cell permeable proteins could enable a DNA-free genome editing approach that would circumvent any possibility of insertional mutagenesis.

[0270] b. Homology-Directed Repair

[0271] Restoration of protein expression from an endogenous mutated gene may involve homology-directed repair. The method as described above further includes administrating a donor template to the cell. The donor template may include a nucleotide sequence encoding a full-functional protein or a partially-functional protein. For example, the donor template may include a miniaturized dystrophin construct, termed minidystrophin ("minidys"), a full-functional dystrophin construct for restoring a mutant dystrophin gene, or a fragment of the dystrophin gene that after homology-directed repair leads to restoration of the mutant dystrophin gene.

[0272] c. Methods of Correcting a Mutant Gene and Treating a Subject Using CRISPR/Cas9

[0273] The present disclosure is also directed to genome editing with the CRISPR/Cas9-based system to restore the expression of a full-functional or partially-functional protein with a repair template or donor DNA, which can replace the entire gene or the region containing the mutation. The CRISPR/Cas9-based system may be used to introduce site-specific double strand breaks at targeted genomic loci. Site-specific double-strand breaks are created when the CRISPR/Cas9-based system binds to a target DNA sequences using the gRNA, thereby permitting cleavage of the target DNA. The CRISPR/Cas9-based system has the advantage of advanced genome editing due to their high rate of successful and efficient genetic modification. This DNA cleavage may stimulate the natural DNA-repair machinery, leading to one of two possible repair pathways: homology-directed repair (HDR) or the non-homologous end joining (NHEJ) pathway. For example, a CRISPR/Cas9-based system directed towards the dystrophin gene may include a gRNA having a nucleic acid sequence of any one of SEQ ID NOs: 65-115.

[0274] The present disclosure is directed to genome editing with CRISPR/Cas9-based system without a repair template, which can efficiently correct the reading frame and restore the expression of a functional protein involved in a genetic disease. The disclosed CRISPR/Cas9-based system and methods may involve using homology-directed repair or nuclease-mediated non-homologous end joining (NHEJ)-based correction approaches, which enable efficient correction in proliferation-limited primary cell lines that may not be amenable to homologous recombination or selection-based gene correction. This strategy integrates the rapid and robust assembly of active CRISPR/Cas9-based system with an efficient gene editing method for the treatment of genetic diseases caused by mutations in nonessential coding regions that cause frameshifts, premature stop codons, aberrant splice donor sites or aberrant splice acceptor sites.

[0275] The present disclosure provides methods of correcting a mutant gene in a cell and treating a subject suffering from a genetic disease, such as DMD. The method may include administering to a cell or subject a CRISPR/Cas9-based system, a polynucleotide or vector encoding said CRISPR/Cas9-based system, or composition of said

CRISPR/Cas9-based system as described above. The method may include administering a CRISPR/Cas9-based system, such as administering a Cas9 protein or Cas9 fusion protein containing a second domain having nuclease activity, a nucleotide sequence encoding said Cas9 protein or Cas9 fusion protein, and/or at least one gRNA, wherein the gRNAs target different DNA sequences. The target DNA sequences may be overlapping. The number of gRNA administered to the cell may be at least 1 gRNA, at least 2 different gRNA, at least 3 different gRNA at least 4 different gRNA, at least 5 different gRNA, at least 6 different gRNA, at least 7 different gRNA, at least 8 different gRNA, at least 9 different gRNA, at least 10 different gRNA, at least 15 different gRNA, at least 20 different gRNA, at least 30 different gRNA, or at least 50 different gRNA, as described above. The gRNA may include a nucleic acid sequence of at least one of SEQ ID NOs: 65-115. The method may involve homology-directed repair or non-homologous end joining.

14. METHODS OF TREATING A DISEASE

[0276] The present disclosure is directed to a method of treating a subject in need thereof. The method comprises administering to a tissue of a subject the composition for altering gene expression and engineering or altering genomic DNA in a cell or subject genome editing, as described above. The method may comprises administering to the skeletal muscle or cardiac muscle of the subject the composition for genome editing in skeletal muscle or cardiac muscle, as described above. The subject may be suffering from a skeletal muscle or cardiac muscle condition causing degeneration or weakness or a genetic disease. For example, the subject may be suffering from Duchenne muscular dystrophy, as described above.

[0277] a. Duchenne Muscular Dystrophy

[0278] The method, as described above, may be used for correcting the dystrophin gene and recovering full-functional or partially-functional protein expression of said mutated dystrophin gene. In some aspects and embodiments the disclosure provides a method for reducing the effects (e.g., clinical symptoms/indications) of DMD in a patient. In some aspects and embodiments the disclosure provides a method for treating DMD in a patient. In some aspects and embodiments the disclosure provides a method for preventing DMD in a patient. In some aspects and embodiments the disclosure provides a method for preventing further progression of DMD in a patient.

15. CONSTRUCTS AND PLASMIDS

[0279] The compositions, as described above, may comprise genetic constructs that encodes the CRISPR/Cas9-based system, as disclosed herein. The genetic construct, such as a plasmid, may comprise a nucleic acid that encodes the CRISPR/Cas9-based system, such as the Cas9 protein and Cas9 fusion proteins and/or at least one of the gRNAs. The compositions, as described above, may comprise genetic constructs that encodes the modified AAV vector and a nucleic acid sequence that encodes the site-specific nuclease, as disclosed herein. The genetic construct, such as a plasmid, may comprise a nucleic acid that encodes the site-specific nuclease. The compositions, as described above, may comprise genetic constructs that encodes the modified lentiviral vector, as disclosed herein. The genetic construct, such as a plasmid, may comprise a nucleic acid that encodes the Cas9-fusion protein and at least one sgRNA. The genetic construct may be present in the cell as a functioning extrachromosomal molecule. The genetic construct may be a linear minichromosome including centromere, telomeres or plasmids or cosmids.

[0280] The genetic construct may also be part of a genome of a recombinant viral vector, including recombinant lentivirus, recombinant adenovirus, and recombinant adenovirus associated virus. The genetic construct may be part of the genetic material in attenuated live microorganisms or recombinant microbial vectors which live in cells. The genetic constructs may comprise regulatory elements for gene expression of the coding sequences of the nucleic acid. The regulatory elements may be a promoter, an enhancer, an initiation codon, a stop codon, or a polyadenylation signal.

[0281] The nucleic acid sequences may make up a genetic construct that may be a vector. The vector may be capable of expressing the fusion protein, such as the Cas9-fusion protein or site-specific nuclease, in the cell of a mammal. The vector may be recombinant. The vector may comprise heterologous nucleic acid encoding the fusion protein, such as the Cas9-fusion protein or site-specific nuclease. The vector may be a plasmid. The vector may be useful for transfecting cells with nucleic acid encoding the Cas9-fusion protein or site-specific nuclease, which the transformed host cell is cultured and maintained under conditions wherein expression of the Cas9-fusion protein or the site-specific nuclease system takes place.

[0282] Coding sequences may be optimized for stability and high levels of expression. In some instances, codons are selected to reduce secondary structure formation of the RNA such as that formed due to intramolecular bonding.

[0283] The vector may comprise heterologous nucleic acid encoding the CRISPR/Cas9-based system or the site-specific nuclease and may further comprise an initiation codon, which may be upstream of the CRISPR/Cas9-based system or the site-specific nuclease coding sequence, and a stop codon, which may be downstream of the CRISPR/Cas9-based system or the site-specific nuclease coding sequence. The initiation and termination codon may be in frame with the CRISPR/Cas9-based system or the site-specific nuclease coding sequence. The vector may also comprise a promoter that is operably linked to the CRISPR/Cas9-based system or the site-specific nuclease coding sequence. The promoter operably linked to the CRISPR/Cas9-based system or the site-specific nuclease coding sequence may be a promoter from simian virus 40 (SV40), a mouse mammary tumor virus (MMTV) promoter, a human immunodeficiency virus (HIV) promoter such as the bovine immunodeficiency virus (BIV) long terminal repeat (LTR) promoter, a Moloney virus promoter, an avian leukosis virus (ALV) promoter, a cytomegalovirus (CMV) promoter such as the CMV immediate early promoter, Epstein Barr virus (EBV) promoter, or a Rous sarcoma virus (RSV) promoter. The promoter may also be a promoter from a human gene such as human ubiquitin C (hUbC), human actin, human myosin, human hemoglobin, human muscle creatine, or human metalothionein. The promoter may also be a tissue specific promoter, such as a muscle or skin specific promoter, natural or synthetic. Examples of such promoters are described in US Patent Application Publication No. US20040175727, the contents of which are incorporated herein in its entirety.

[0284] The vector may also comprise a polyadenylation signal, which may be downstream of the CRISPR/Cas9-based system or the site-specific nuclease. The polyadenylation signal may be a SV40 polyadenylation signal, LTR polyadenylation signal, bovine growth hormone (bGH) polyadenylation signal, human growth hormone (hGH) polyadenylation signal, or human β-globin polyadenylation signal. The SV40 polyadenylation signal may be a polyadenylation signal from a pCEP4 vector (Invitrogen, San Diego, Calif.).

[0285] The vector may also comprise an enhancer upstream of the CRISPR/Cas9-based system, i.e., the Cas9 protein or Cas9 fusion protein coding sequence or sgRNAs, or the site-specific nuclease. The enhancer may be necessary for DNA expression. The enhancer may be human actin, human myosin, human hemoglobin, human muscle creatine or a viral enhancer such as one from CMV, HA, RSV or EBV. Polynucleotide function enhancers are described in U.S. Pat. Nos. 5,593,972, 5,962,428, and WO94/016737, the contents of each are fully incorporated by reference. The vector may also comprise a mammalian origin of replication in order to maintain the vector extrachromosomally and produce multiple copies of the vector in a cell. The vector may also comprise a regulatory sequence, which may be well suited for gene expression in a mammalian or human cell into which the vector is administered. The vector may also comprise a reporter gene, such as green fluorescent protein ("GFP") and/or a selectable marker, such as hygromycin ("Hygro").

[0286] The vector may be expression vectors or systems to produce protein by routine techniques and readily available starting materials including Sambrook et al., Molecular Cloning and Laboratory Manual, Second Ed., Cold Spring Harbor (1989), which is incorporated fully by reference. In some embodiments the vector may comprise the nucleic acid sequence encoding the CRISPR/Cas9-based system, including the nucleic acid sequence encoding the Cas9 protein or Cas9 fusion protein and the nucleic acid sequence encoding the at least one gRNA comprising the nucleic acid sequence of at least one of SEQ ID NOs: 5-40, 65-144, 492-515, 540-563, and 585-625.

16. PHARMACEUTICAL COMPOSITIONS

[0287] The composition may be in a pharmaceutical composition. The pharmaceutical composition may comprise about 1 ng to about 10 mg of DNA encoding the CRISPR/Cas9-based system or CRISPR/Cas9-based system protein component, i.e., the Cas9 protein or Cas9 fusion protein. The pharmaceutical composition may comprise about 1 ng to about 10 mg of the DNA of the modified AAV vector and nucleotide sequence encoding the site-specific nuclease. The pharmaceutical composition may comprise about 1 ng to about 10 mg of the DNA of the modified lentiviral vector. The pharmaceutical compositions according to the present invention are formulated according to the mode of administration to be used. In cases where pharmaceutical compositions are injectable pharmaceutical compositions, they are sterile, pyrogen free and particulate free. An isotonic formulation is preferably used. Generally, additives for isotonicity may include sodium chloride, dextrose, mannitol, sorbitol and lactose. In some cases, isotonic solutions such as phosphate buffered saline are preferred. Stabilizers include gelatin and albumin. In some embodiments, a vasoconstriction agent is added to the formulation.

[0288] The composition may further comprise a pharmaceutically acceptable excipient. The pharmaceutically acceptable excipient may be functional molecules as vehicles, adjuvants, carriers, or diluents. The pharmaceutically acceptable excipient may be a transfection facilitating agent, which may include surface active agents, such as immune-stimulating complexes (ISCOMS), Freunds incomplete adjuvant, LPS analog including monophosphoryl lipid A, muramyl peptides, quinone analogs, vesicles such as squalene and squalene, hyaluronic acid, lipids, liposomes, calcium ions, viral proteins, polyanions, polycations, or nanoparticles, or other known transfection facilitating agents.

[0289] The transfection facilitating agent is a polyanion, polycation, including poly-L-glutamate (LGS), or lipid. The transfection facilitating agent is poly-L-glutamate, and more preferably, the poly-L-glutamate is present in the composition for genome editing in skeletal muscle or cardiac muscle at a concentration less than 6 mg/ml. The transfection facilitating agent may also include surface active agents such as immune-stimulating complexes (ISCOMS), Freunds incomplete adjuvant, LPS analog including monophosphoryl lipid A, muramyl peptides, quinone analogs and vesicles such as squalene and squalene, and hyaluronic acid may also be used administered in conjunction with the genetic construct. In some embodiments, the DNA vector encoding the composition may also include a transfection facilitating agent such as lipids, liposomes, including lecithin liposomes or other liposomes known in the art, as a DNA-liposome mixture (see for example WO9324640), calcium ions, viral proteins, polyanions, polycations, or nanoparticles, or other known transfection facilitating agents. Preferably, the transfection facilitating agent is a polyanion, polycation, including poly-L-glutamate (LGS), or lipid.

17. METHODS OF DELIVERY

[0290] Provided herein is a method for delivering the pharmaceutical formulations, preferably compositions described above, for providing genetic constructs. The delivery of the compositions may be the transfection or electroporation of the composition as a nucleic acid molecule that is expressed in the cell and delivered to the surface of the cell. The nucleic acid molecules may be electroporated using BioRad Gene Pulser Xcell or Amaxa Nucleofector IIb devices. Several different buffers may be used, including BioRad electroporation solution, Sigma phosphate-buffered saline product #D8537 (PBS), Invitrogen OptiMM I (OM), or Amaxa Nucleofector solution V (N.V.). Transfections may include a transfection reagent, such as Lipofectamine 2000.

[0291] Upon delivery of the composition to the tissue, and thereupon the vector into the cells of the mammal, the transfected cells will express the fusion protein, such as a CRISPR/Cas9-based system and/or a site-specific nuclease. The composition may be administered to a mammal to alter gene expression or to re-engineer or alter the genome. For example, the composition may be administered to a mammal to correct the dystrophin gene in a mammal. The mammal may be human, non-human primate, cow, pig, sheep, goat, antelope, bison, water buffalo, bovids, deer, hedgehogs, elephants, llama, alpaca, mice, rats, or chicken, and preferably human, cow, pig, or chicken.

[0292] a. CRISPR/Cas9-Based System

[0293] The vector encoding a CRISPR/Cas9-based system protein component, i.e., the Cas9 protein or Cas9 fusion protein, may be delivered to the mammal by DNA injection (also referred to as DNA vaccination) with and without in vivo electroporation, liposome mediated, nanoparticle facilitated, and/or recombinant vectors. The recombinant vector may be delivered by any viral mode. The viral mode may be recombinant lentivirus, recombinant adenovirus, and/or recombinant adeno-associated virus.

[0294] The nucleotide encoding a CRISPR/Cas9-based system protein component, i.e., the Cas9 protein or Cas9 fusion protein, may be introduced into a cell to genetically correct the target gene or alter gene expression of a gene, such as activate or repress endogenous genes. For example, a nucleotide encoding a CRISPR/Cas9-based system protein component, i.e., the Cas9 protein or Cas9 fusion protein, directed towards a mutant dystrophin gene by the gRNA may be introduced into a myoblast cell from a DMD patient. Alternatively, they may be introduced into a fibroblast cell from a DMD patient, and the genetically corrected fibroblast cell may be treated with MyoD to induce differentiation into myoblasts, which may be implanted into subjects, such as the damaged muscles of a subject to verify that the corrected dystrophin protein was functional and/or to treat the subject. The modified cells may also be stem cells, such as induced pluripotent stem cells, bone marrow-derived progenitors, skeletal muscle progenitors, human skeletal myoblasts from DMD patients, CD133+ cells, mesoangioblasts, and MyoD- or Pax7-transduced cells, or other myogenic progenitor cells. For example, the CRISPR/Cas9-based system may cause neuronal or myogenic differentiation of an induced pluripotent stem cell.

18. ROUTES OF ADMINISTRATION

[0295] The compositions may be administered to a subject by different routes including orally, parenterally, sublingually, transdermally, rectally, transmucosally, topically, via inhalation, via buccal administration, intrapleurally, intravenous, intraarterial, intraperitoneal, subcutaneous, intramuscular, intranasal intrathecal, and intraarticular or combinations thereof. For veterinary use, the composition may be administered as a suitably acceptable formulation in accordance with normal veterinary practice. The veterinarian may readily determine the dosing regimen and route of administration that is most appropriate for a particular animal. The compositions may be administered by traditional syringes, needleless injection devices, "microprojectile bombardment gone guns", or other physical methods such as electroporation ("EP"), "hydrodynamic method", or ultrasound.

[0296] The composition may be delivered to the mammal by several technologies including DNA injection (also referred to as DNA vaccination) with and without in vivo electroporation, liposome mediated, nanoparticle facilitated, recombinant vectors such as recombinant lentivirus, recombinant adenovirus, and recombinant adenovirus associated virus. The composition may be injected into the skeletal muscle or cardiac muscle. For example, the composition may be injected into the tibialis anterior muscle.

19. CELL TYPES

[0297] Any of these delivery methods and/or routes of administration could be utilized with a myriad of cell types, for example, those cell types currently under investigation for cell-based therapies. Cell types may be fibroblasts, pluripotent stem cells, cardiomyocytes, hepatocytes, chondrocytes, mesenchymal progenitor cells, hematopoetic stem cells, smooth muscle cells, or K562 human erythroid leukemia cell line.

[0298] a. DMD

[0299] Cell types currently under investigation for cell-based therapies of DMD include immortalized myoblast cells, such as wild-type and DMD patient derived lines, for example Δ48-50 DMD, DMD 8036 (del48-50), C25C14 and DMD-7796 cell lines, primal DMD dermal fibroblasts, induced pluripotent stem cells, bone marrow-derived progenitors, skeletal muscle progenitors, human skeletal myoblasts from DMD patients, CD133+ cells, mesoangioblasts, cardiomyocytes, hepatocytes, chondrocytes, mesenchymal progenitor cells, hematopoetic stem cells, smooth muscle cells, and MyoD- or Pax7-transduced cells, or other myogenic progenitor cells. Immortalization of human myogenic cells may be used for clonal derivation of genetically corrected myogenic cells. Cells may be modified ex vivo to isolate and expand clonal populations of immortalized DMD myoblasts that contain a genetically corrected dystrophin gene and are free of other nuclease-introduced mutations in protein coding regions of the genome. Alternatively, transient in vivo delivery of nucleases by non-viral or non-integrating viral gene transfer, or by direct delivery of purified proteins and gRNAs containing cell-penetrating motifs may enable highly specific correction in situ with minimal or no risk of exogenous DNA integration.

20. KITS

[0300] Provided herein is a kit, which may be used to edit a genome in skeletal muscle or cardiac muscle, such as correcting a mutant gene. The kit comprises a composition for genome editing in skeletal muscle or cardiac muscle, as described above, and instructions for using said composition. Instructions included in kits may be affixed to packaging material or may be included as a package insert. While the instructions are typically written or printed materials they are not limited to such. Any medium capable of storing such instructions and communicating them to an end user is contemplated by this disclosure. Such media include, but are not limited to, electronic storage media (e.g., magnetic discs, tapes, cartridges, chips), optical media (e.g., CD ROM), and the like. As used herein, the term "instructions" may include the address of an internet site that provides the instructions.

[0301] The composition for genome editing in skeletal muscle or cardiac muscle may include a modified AAV vector and a nucleotide sequence encoding a site-specific nuclease, as described above. The site-specific nuclease may include a ZFN, a TALEN, or CRISPR/Cas9-based system, as described above, that specifically binds and cleaves a mutated gene. The site-specific nuclease, as described above, may be included in the kit to specifically bind and target a particular region in the mutated gene. The site-specific nuclease may be specific for a mutated dystrophin gene, as described above. The kit may further include donor DNA, a gRNA, or a transgene, as described above.

[0302] a. CRISPR/Cas9-based system

[0303] Provided herein is a kit, which may be used to correct a mutated gene. The kit comprises at least one component for correcting a mutated gene and instructions for using the CRISPR/Cas9-based system. Instructions

included in kits may be affixed to packaging material or may be included as a package insert. While the instructions are typically written or printed materials they are not limited to such. Any medium capable of storing such instructions and communicating them to an end user is contemplated by this disclosure. Such media include, but are not limited to, electronic storage media (e.g., magnetic discs, tapes, cartridges, chips), optical media (e.g., CD ROM), and the like. As used herein, the term "instructions" may include the address of an internet site that provides the instructions.

[0304] At least one component may include at least one CRISPR/Cas9-based system, as described above, which specifically targets a gene. The kit may include a Cas9 protein or Cas9 fusion protein, a nucleotide sequence encoding said Cas9 protein or Cas9 fusion protein, and/or at least one gRNA. The CRISPR/Cas9-based system, as described above, may be included in the kit to specifically bind and target a particular target region upstream, within or downstream of the coding region of the target gene. For example, a CRISPR/Cas9-based system may be specific for a promoter region of a target gene or a CRISPR/Cas9-based system may be specific for a mutated gene, for example a mutated dystrophin gene, as described above. The kit may include donor DNA, as described above.

21. EXAMPLES

[0305] It will be readily apparent to those skilled in the art that other suitable modifications and adaptations of the methods of the present disclosure described herein are readily applicable and appreciable, and may be made using suitable equivalents without departing from the scope of the present disclosure or the aspects and embodiments disclosed herein. Having now described the present disclosure in detail, the same will be more clearly understood by reference to the following examples, which are merely intended only to illustrate some aspects and embodiments of the disclosure, and should not be viewed as limiting to the scope of the disclosure. The disclosures of all journal references, U.S. patents, and publications referred to herein are hereby incorporated by reference in their entireties.

[0306] The present invention has multiple aspects, illustrated by the following non-limiting examples.

Example 1

Materials and Methods

[0307] Cell culture and transfection. HEK293T cells were obtained from the American Tissue Collection Center (ATCC) through the Duke University Cancer Center Facilities and were maintained in DMEM supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin at 37° C. with 5% C02. HEK293T cells were transfected with Lipofectamine 2000 (Invitrogen) according to manufacturer's instructions. Transfection efficiencies were routinely higher than 80% as determined by fluorescence microscopy following delivery of a control eGFP expression plasmid. Cas9 expression plasmid was transfected at a mass ratio of 3:1 to either the individual gRNA expression plasmids or the identical amount of gRNA expression plasmid consisting of a mixture of equal amounts of the four gRNAs.

[0308] Primary mouse embryonic fibroblasts (PMEF-HL, Millipore, Billerica, Mass.) were seeded (75,000 per well) in 24-well TCPS plates (BD, Franklin Lakes, N.J.) and maintained at 37° C. and 5% CO_2 in complete MEF medium consisting of high glucose DMEM supplemented with 10% Premium Select FBS (Atlanta Biologicals, Lawrenceville, Ga.), 25 μg mL^{-1} gentamicin (Invitrogen), 1× GutaMAX, non-essential amino acids, sodium pyruvate, and β-mercaptoethanol (Invitrogen). MEF transfections were performed with a single 1 μg cm^{-2} dose of total plasmid DNA, delivered as cationic nanocomplexes following electrostatic condensation with poly(CBA-ABOL) in serum- and antibiotic-free OptiMEM, as described previously (Adler et al. *Molecular therapy. Nucleic acids* 1, e32 (2012)). OptiMEM was replaced with complete MEF medium 4 hrs after transfection. 48 hrs after transfection, MEFs were processed for qRT-PCR, or the complete MEF medium was replaced with N3 neural induction medium containing: DMEM/F-12 (Invitrogen), 1× N-2 Supplement (Invitrogen), 10 ng mL^{-1} human bFGF2 (Stemgent, Cambridge, Mass.), and 25 μg mL^{-1} gentamicin (Invitrogen). A GFP reporter vector (pmax-GFP, 3486 bp, Amaxa, Cologne, Germany) was used to optimize transfection conditions. Cas9 expression plasmid was transfected at a mass ratio of 3:1 or 1:1 to an equal mixture of four gRNA expression plasmids.

[0309] Plasmids. The plasmids encoding wild-type and H840A Cas9 were obtained from Addgene (Plasmid #39312 and Plasmid #39316; Jinek, et al. *Science* 337, 816-821 (2012)). H840A Cas9 was cloned into the vector pcDNA3.1 in frame with a FLAG epitope tag and a nuclear localization sequence (NLS) at the N-terminus with a primer pair that introduced the D10A mutation. The VP64 domain, an NLS, and an HA epitope tag were cloned in frame with the Cas9 ORF at the C-terminus (FIG. **1**A, FIG. **9**A). The tracrRNA and crRNA expression cassettes (Cong et al. *Science* 339, 819-823 (2013)) were ordered as gBlocks (Integrated DNA Technologies (IDT)) and cloned into a pZDonor plasmid (Sigma) with KpnI and SacII sites. A chimeric guide RNA expression cassette (Mali et al. *Science* 339, 823-826 (2013)) was also ordered as gBlocks with modifications to include a BbsI restriction site to facilitate rapid cloning of new guide RNA spacer sequences (FIG. **9**B). The oligonucleotides containing the target sequences were obtained from IDT, hybridized, phosphorylated, and cloned in the appropriate plasmids using BbsI sites. The target sequences are provided in Table 2.

TABLE 2

Target sequences of gRNAs

Target	Name	Sequence	SEQ ID NO
ASCL1	CR1	GCTGGGTGTCCCATTGAAA	5
	CR2	CAGCCGCTCGCTGCAGCAG	6
	CR3	TGGAGAGTTTGCAAGGAGC	7
	CR4	GTTTATTCAGCCGGGAGTC	8
NANOG	CR1	CGCCAGGAGGGGTGGGTCTA	9
	CR2	CCTTGGTGAGACTGGTAGA	10
	CR3	GTCTTCAGGTTCTGTTGCT	11
	CR4	ATATTCCTGATTTAAAAGT	12
VEGFA	CR1	TTAAAAGTCGGCTGGTAGC	13
	CR2	CGGGCCGGGGGCGGGGTCC	14
	CR3	GCCCGAGCCGCGTGTGGAA	15
	CR4	CCTTCATTGCGGCGGGCTG	16
TERT	CR1	CCGACCCCTCCCGGGTCCC	17
	CR2	CAGGACCGCGCTTCCCACG	18

TABLE 2-continued

Target sequences of qRNAs

Target	Name	Sequence	SEQ ID NO
	CR3	TGCACCCTGGGAGCGCGAG	19
	CR4	CCGCACGCACCTGTTCCCA	20
IL1B	CR1	AAAACAGCGAGGGAGAAAC	21
	CR2	TTAACTTGATTGTGAAATC	22
	CR3	AAAACAATGCATATTTGCA	23
	CR4	AAAATCCAGTATTTTAATG	24
IL1R2	CR1	ACCCAGCACTGCAGCCTGG	25
	CR2	AACTTATGCGGCGTTTCCT	26
	CR3	TCACTTTAAAACCACCTCT	27
	CR4	GCATCTTTTTCTCTTTAAT	28
IL1RN	CR1	TGTACTCTCTGAGGTGCTC	29
	CR2	ACGCAGATAAGAACCAGTT	30
	CR3	CATCAAGTCAGCCATCAGC	31
	CR4	GAGTCACCCTCCTGGAAAC	32
HBG1/2	CR1	GCTAGGGATGAAGAATAAA	33
	CR2	TTGACCAATAGCCTTGACA	34
	CR3	TGCAAATATCTGTCTGAAA	35
	CR4	AAATTAGCAGTATCCTCTT	36
MYOD1	CR1	CCTGGGCTCCGGGGCGTTT	37
	CR2	GGCCCCTGCGGCCACCCCG	38
	CR3	CTCCCTCCCTGCCCGGTAG	39
	CR4	AGGTTTGGAAAGGGCGTGC	40

[0310] Western blot. Cells were lysed in 50 mM Tris-Cl (pH 7.4), 150 mM NaCl, 0.50 Triton X-100, and 0.1% SDS. Lysates were mixed with loading buffer, boiled for 5 min, and equal volumes of protein were run in NuPAGE® Novex 0-2% or 10% Bis-Tris Gel polyacrylamide gels and transferred to nitrocellulose membranes. Non-specific antibody binding was blocked with 50 mM Tris/150 mM NaCl/0.1% Tween-20 (TBS-T) with 5% nonfat milk for 30 min. The membranes were incubated with primary antibodies (HRP-conjugated anti-Flag (Cell Signaling, Cat #2044) in 5% BSA in TBS-T diluted 1:1000 overnight; anti-GAPDH (Cell Signaling, clone 14C10) in 5% milk in TBS-T diluted 1:5000 for 30 min; anti-ASCL1 (Santa Cruz, clone sc-48449) in 5% BSA diluted 1:500; or anti-g-globin (Santa Cruz, clone 51-7) in 5% milk diluted 1:500 and the membranes were washed with TBS-T for 30 min. Membranes labeled with primary antibodies were incubated with anti-rabbit HRP-conjugated antibody (Sigma-Aldrich) diluted 1:5000 for 30 min, anti-goat (1:3000) or anti-mouse (1:5000) and washed with TBS-T for 30 min. Membranes were visualized using the Immun-Star WesternC™ Chemiluminescence Kit (Bio-Rad) and images were captured using a ChemiDoc™ XRS+ System and processed using Image-Lab software (Bio-Rad).

[0311] ELISA. Serum-free culture media (OPTI-MEM) was collected and frozen at −80° C. Human IL-1ra secretion into culture media was quantified via enzyme-linked immunosorbent assay (ELISA), according to the manufacturer's protocols (R&D Systems, Cat. No. DY280). The standard curve was prepared by diluting recombinant human IL-1ra in OPTI-MEM and the IL-1ra in culture media was measured undiluted. The samples were concentrated about 8 fold via centrifugation through 3 kDa MWCO filters for 20 min (Amicon Ultra, Cat # UFC500396). Reported values were corrected by the concentration factor for each sample.

[0312] Optical density was measured at 450 nm, with a wavelength correction at 540 nm. Each standard and sample was assayed in duplicate. The duplicate readings were averaged and normalized by subtracting the average zero standard optical density. A standard curve was generated by log-transforming the data and performing a linear regression of the IL-1ra concentration versus the optical density. Reported values are the mean and standard error of the mean from three independent experiments (n=3) that were performed on different days with technical duplicates that were averaged for each experiment.

[0313] qRT-PCR. Total RNA was isolated using the RNeasy Plus RNA isolation kit (Qiagen). cDNA synthesis was performed using the SuperScript® VILO™ cDNA Synthesis Kit (Invitrogen). Real-time PCR using PerfeCTa® SYBR® Green FastMix was performed with the CFX96 Real-Time PCR Detection System (Bio-Rad) with oligonucleotide primers reported in Table 3 that were designed using Primer3Plus software and purchased from IDT.

TABLE 3

Sequences of primers used for qRT-PCR.

Target	Forward Primer	SEQ ID NO	Reverse Primer	SEQ ID NO
hASCL1	GGAGCTTCTCGACT TCACCA	41	AACGCCACTGACAA GAAAGC	53
NANOG	GATTTGTGGGCCTG AAGAAA	42	CAGATCCATGGAGG AAGGAA	54
VEGFA	AAGGAGGAGGGCAG AATCAT	43	GGGTACTCCTGGAA GATGTCC	55
TERT	AAACCTTCCTCAGC TATGCCC	44	GTTTGCGACGCATG TTCCTC	56
IL1B	AGCTGATGGCCCTA AACAGA	45	AAGCCCTTGCTGTA GTGGTG	57
IL1R2	CAGGAGGACTCTGG CACCTA	46	CGGCAGGAAAGCAT CTGTAT	58
IL1RN	GGAATCCATGGAGG GAAGAT	47	TGTTCTCGCTCAGG TCAGTG	59
HBG1/2	GCTGAGTGAACTGC ACTGTGA	48	GAATTCTTTGCCGA AATGGA	60
MYOD1	CTCTCTGCTCCTTT GCCACA	49	GTGCTCTTCGGGTT TCAGGA	61
GAPDH	CAATGACCCCTTCA TTGACC	50	TTGATTTTGGAGGG ATCTCG	62
mASCL1	GGAACAAGAGCTGC TGGACT	51	GTTTTTCTGCCTCC CCATTT	63
mGAPDH	AACTTTGGCATTGT GGAAGG	52	GGATGCAGGGATGA TGTTCT	64

[0314] Primer specificity was confirmed by agarose gel electrophoresis and melting curve analysis. Reaction efficiencies over the appropriate dynamic range were calculated to ensure linearity of the standard curve (FIG. **10**). The results are expressed as fold-increase mRNA expression of the gene of interest normalized to GAPDH expression by the $\Delta\Delta C_T$ method. Reported values are the mean and standard error of the mean from three independent experiments

performed on different days (n=3) with technical duplicates that were averaged for each experiment.

[0315] RNA-Seq. RNA seq libraries were constructed. Briefly, first strand cDNA was synthesized from oligo dT Dynabead® (Invitrogen) captured mRNA using SuperScript® VILO™ cDNA Synthesis Kit (Invitrogen). Second strand cDNA was synthesized using DNA Polymerase I (New England Biolabs). cDNA was purified using Agencourt AMPure XP beads (Beckman Coulter) and Nextera transposase (Illumina; 5 min at 55° C.) was used to simultaneously fragment and insert sequencing primers into the double-stranded cDNA. Transposition reactions were halted using QG buffer (Qiagen) and fragmented cDNA was purified on AMPure XP beads. Indexed sequencing libraries were generated by 6 cycles of PCR.

[0316] Libraries were sequenced using 50-bp single end reads on two lanes of an Illumina HiSeq 2000 instrument, generating between 29 million and 74 million reads per library. Reads were aligned to human RefSeq transcripts using Bowtie (Langmead et al. Genome biology 10, R25 (2009)). The statistical significance of differential expression, including correction for multiple hypothesis testing, was calculated using DESeq (Anders et al. Genome biology 11, R106 (2010)). Raw RNA-seq reads and the number of reads aligned to each RefSeq transcript have been deposited for public access in the Gene Expression Omnibus (GEO), with accession number currently pending.

[0317] Immunofluorescence staining. For detection of Tuj1 and MAP2 expression, transfected MEFs were fixed at day 10 of culture in N3 medium with 4% PFA (EMS, Hatfield, Pa.) at room temperature (RT) for 20 min. Cells were then incubated in blocking buffer containing 0.2% Triton X-100, 3% w/v BSA, and 10% goat serum (Sigma-Aldrich, Saint Louis, Mo.) for two hrs at room temperature with rabbit anti-Tuj1 (Covance, Princeton, N.J., clone TUJ1 1-15-79, 1:500) and mouse anti-MAP2 (BD, clone Ap20, 1:500), or for an additional 24 hrs at 4° C. with mouse anti-Ascl1 (BD clone 24B72D11.1, 1:100). The cells were then washed three times with PBS, incubated for 1 hr at room temperature in blocking buffer with Alexa Fluor 488 goat anti-mouse IgG and Alexa Fluor 594 goat anti-rabbit IgG (Invitrogen, 1:200), and washed three times with PBS. Stained MEFs were then scanned with a Nikon Eclipse TE2000-U inverted fluorescence microscope with a ProScanII motorized stage (Prior Scientific, Rockland, Mass.) to produce large mosaic images of each complete culture area. These mosaics were processed with a FIJI macro to automatically and uniformly threshold each image according to local contrast, exclude small debris, and to count the number of $Tuj1^+$ cells in each well.

[0318] Statistics. Statistical analysis was performed by Tukey's test with alpha equal to 0.05 in JMP 10 Pro.

Example 2

Results

[0319] To create a CRISPR/Cas9-based transcriptional activation system, catalytic residues of Cas9 (D10A, H840A) were mutated to create iCas9 and genetically fused with a C-terminal VP64 acidic transactivation domain (FIG. **1**A, FIG. **1**). Robust expression of iCas9-VP64 was observed from the transfected plasmid in human embryonic kidney (HEK) 293T cells by western blot of the N-terminal Flag epitope tag (FIG. **3**). The CRISPR system recognizes its target through base pairing of a 20 bp sequence in the gRNA to a complementary DNA target, which is followed by the NGG protospacer-adjacent motif (PAM) sequence, where N is any base pair. Combinations of synthetic transcription factors targeted to endogenous human promoters result in synergistic and robust activation of gene expression. Therefore four gRNA target sites followed by the NGG PAM sequence were identified in the promoter of the IL1RN gene within 500 bp of the transcriptional start site (FIG. **4**, Table 2). To compare crRNA- and gRNA-based targeting strategies, the four target site sequences were introduced into crRNA and gRNA expression plasmids[17] and co-transfected with the iCas9-VP64 expression plasmid into HEK293T cells. Although substantial induction of IL1RN expression was observed by qRT-PCR in samples treated with the combination of crRNAs, much higher levels were achieved with the combination of gRNAs (FIG. **1**C). No changes to gene expression were observed in cells treated with gRNAs and an expression plasmid for iCas9 without VP64, demonstrating the critical role of the activation domain in modulating gene expression (FIG. **1**C). Nuclease activity at these target sites was confirmed to have been abrogated in the iCas9-VP64 system by performing the Surveyor assay to detect DNA repair events in samples treated with iCas9-VP64 and wild-type Cas9 (FIG. **5**). By transfecting each of the four gRNAs individually or in combination, targeting multiple sites in the promoter with combinations of gRNAs showed robust increases in gene expression (FIG. **1**D). High levels of IL1RN expression were observed only when the gRNA combinations were co-transfected with iCas9-VP64 (FIG. **1**D), as seen with other classes of engineered transcription factors. Similarly, production of the IL-1 receptor antagonist (IL-1ra) protein, encoded by the IL1RN gene, was only observed in three of the six samples treated with the combination of gRNAs across three different experiments, whereas it was never detected in samples treated with single gRNAs or control plasmid (FIG. **1**E). To examine the specificity of gene activation by iCas9-VP64, global gene expression of HEK293T cells treated with the combination of four gRNAs by RNA-seq was assessed (FIG. **1**F). Notably, the only genes with significantly increased expression relative to control (false discovery rate $\leq 3\times10^{-4}$) were the four isoforms expressed from the IL1RN locus (FIG. **4**), indicating a high level of specificity of gene activation.

[0320] To demonstrate the general applicability of this system, four gRNAs were designed to target each of the promoters of eight other genes relevant to medicine and biotechnology, including ASCL1, NANOG, HBG1/2, MYOD1, VEGFA, TERT, IL1B, and IL1R2 (FIG. **4**, Table 2). Forced expression of ASCL1 and MYOD1 leads to transdifferentiation of several cell types into neuronal and myogenic phenotypes, respectively. NANOG is a marker of pluripotency and that is also used in genetic reprogramming strategies. Activation of the homologs HBG1 and HBG2, which encode γ-globin during fetal development, can be used as a therapeutic strategy to compensate for β-globin mutations in sickle cell disease. Up-regulation of VEGFA by synthetic transcription factors has been explored as a strategy to enhance tissue regeneration and wound healing. The forced expression of telomerase, encoded by the TERT gene, can be used to immortalize cell lines. IL1B encodes the IL-1β cytokine that mediates inflammation and autoimmunity. IL-1β signaling can be blocked by expression of IL-1ra or the decoy receptor encoded by IL1R2. Expression of each

of these genes was enhanced by co-transfection of expression plasmids for iCas9-VP64 and the four gRNAs into HEK293T cells, as determined by qRT-PCR (FIGS. **2**A-**2**H). In some cases expression of a single gRNA was sufficient to induce gene expression, but in all cases co-transfection of the four gRNAs led to synergistic effects (FIGS. **2**A-**2**D). Notably, chromatin accessibility, as determined by DNase-seq, was not a predictor of successful gene activation (FIG. **4**). RNA-seq was performed on cells transfected with iCas9-VP64 and the four gRNAs targeting HBG1, three of which also target HBG2. This revealed specific and reproducible increases in expression of both HBG1 and HBG2, which cannot be distinguished by RNA-seq, although statistical significance was not achieved due to low total expression levels (FIG. **6**). Increases in protein expression of Asc1 and γ-globin following treatment with iCas9-VP64 and the four gRNAs were detected by western blot (FIG. **7**), corroborating higher mRNA levels observed by qRT-PCR (FIGS. **2**A-**2**H). Low baseline levels of Ascl1 and γ-globin protein expression were detectable in empty vector controls. As preliminary evidence that the activation of gene targets by iCas9-VP64 can lead to secondary changes in gene networks and cell phenotypes, expression plasmids for iCas9-VP64 and the four gRNAs targeting ASCL1 were co-transfected into murine embryonic fibroblasts (MEFs) (FIGS. **8**A-**8**H). Forced expression of Ascl1 in MEFs has been shown to partially activate the neuronal gene network, including the downstream target Tuj1. Because the gRNA target sites are conserved in the human and mouse ASCL1 promoters (FIG. **8**A), activation of ASCL1 expression was also observed in MEFs treated with plasmids encoding iCas9-VP64 and the four gRNAs (FIG. **8**B). Furthermore, cells expressing Ascl1 and the neuronal marker Tuj1 were readily detected by immunofluorescence staining 12 days after transfection in the iCas9-VP64/gRNA-treated samples (FIGS. **8**C-**8**H). No Tuj1-positive cells were observed in the cells treated with the control plasmid.

[0321] Thus far there has not been any comprehensive survey of the specificity of Cas9/CRISPR activity in mammalian cells. Using RNA-seq, targeted gene activation was shown to be exquisitely specific with no detectable off-target gene activation (FIG. **1**F, FIG. **6**). IL1RN and HBG12 were chosen for this specificity analysis as the gene products, IL-1ra and γ-globin, may not generate secondary effects on gene expression in HEK293T cells. Exploiting the synergistic activity of multiple weak transcriptional activators, in contrast to using a single strong activator, may increase specific gene regulation since it is unlikely that multiple adjacent off-target sites would exist at another locus. Interestingly, the IL32 gene was moderately downregulated (false discovery rate <0.03) in both the samples treated with iCas9-VP64 and either the IL1RN- or HBG1/2-targeted gRNAs compared to control samples treated with only an empty expression plasmid (FIG. **1**F, FIG. **6**). Because both the IL1RN and HBG1/2-targeted samples were similarly affected, it is unlikely that this is the result of off-target iCas9-VP64 activity related to the identity of the target sequences.

[0322] To evaluate the specificity with which iCas9-VP64 binds the genome, ChIP sequencing was performed using an anti-HA antibody on cells treated with iCas9-VP64 and four gRNAs targeting the IL1RN promoter. The experiment revealed that iCas9 targets the IL1RN promoter (FIG. **15**). Moreover, the experiment revealed an extremely high level of specificity. The iCas9 had only 10 potential off-target binding sites (FDR<5%). To further query the specificity, RNA sequencing experiments were performed with iCas9 EGEMs and found that only IL1RN gene isoforms increased in expression relative to control (FDR≤3×10.4).

Example 3

CRISPRs Targeting the Dystrophin Gene—Methods and Materials

[0323] Plasmid constructs. The expression cassettes for the *S. pyogenes* sgRNA and human codon optimized Cas9 (hCas9) nuclease were used, as previously described (Perez-Pinera et al., *Nat Methods* 10:973-976 (2013)). In order to create a fluorescent reporter system to enrich CRISPR/Cas9-modified cells, a GeneBlock (IDT) was synthesized containing a portion of the 3' end of the Cas9 coding sequence fused to a T2A skipping peptide immediately upstream of a multiple cloning site and subsequently cloned into the hCas9 expression vector. An eGFP reporter gene was then cloned into the T2A vector to allow co-translation of Cas9 and eGFP proteins from the same expression vector (hCas9-T2A-GFP, SEQ ID NO: 116).

[0324] Cell culture and transfection. HEK293T cells were obtained from the American Tissue Collection Center (ATCC) through the Duke Cell Culture Facility and were maintained in DMEM supplemented with 10% bovine calf serum and 1% penicillin/streptomycin. Immortalized myoblasts (Mamchaoui, K. et al. *Skelet Muscle* 1, 1-11 (2011)) (one from a wild-type donor, and two Δ48-50 DMD patient derived lines) were maintained in skeletal muscle media (PromoCell) supplemented with 20% bovine calf serum (Sigma), 50 μg/ml fetuin, 10 ng/ml human epidermal growth factor (Sigma), 1 ng/ml human basic fibroblast growth factor (Sigma), 10 μg/ml human insulin (Sigma), 1% GutaMAX (Invitrogen), and 1% penicillin/streptomycin (Invitrogen). Primary DMD dermal fibroblasts were obtained from the Coriell Cell repository (GM05162A, A46-50) and maintained in DMEM supplemented with 10% fetal bovine serum, 1 ng/mL human basic fibroblast growth factor, and 1% penicillin/streptomycin. All cell lines were maintained at 37° C. and 5% C02.

[0325] HEK293T cells were transfected with Lipofectamine 2000 (Invitrogen) with 400 ng of each expression vector according to the manufacturer's protocol in 24 well plates. Immortalized myoblasts and primary fibroblasts were transfected with 5 μg of each expression vector by electroporation using the Gene Pulser XCell (BioRad) with PBS as an electroporation buffer using optimized conditions for each line (FIGS. **1**A-**1**F) (Ousterout et al. Mol Ther 21:1718-1726 (2013)). Transfection efficiencies were measured by delivering an eGFP expression plasmid (pmaxGFP, Clontech) and using flow cytometry. These efficiencies were routinely ≥95% for HEK293T and ≥70% for the primary fibroblasts and immortalized myoblasts. For all experiments, the indicated mass of electroporated plasmid corresponds to the amount used for each CRISPR/Cas9-based system.

[0326] Cel-I Quantification of Endogenous Gene Modification (Surveyor Assay).

[0327] CRISPR/Cas9-based system-induced lesions at the endogenous target site were quantified using the Surveyor nuclease assay (Guschin, D. Y. et al. *Meth Mol Biol* 649, 247-256 (2010)), which can detect mutations characteristic of nuclease-mediated NHEJ. After transfection, cells were

incubated for 3 or 10 days at 37° C. and genomic DNA was extracted using the DNeasy Blood and Tissue kit (QIAGEN). The target locus was amplified by 35 cycles of PCR with the AccuPrime High Fidelity PCR kit (Invitrogen) using primers specific to each locus (see Table 4), such as 5'-GAGTTTGGCTCAAATTGTTACTCTT-3' (SEQ ID NO: 626) and 5'-GGGAAATGGTCTAGGAGAGTAAAGT-3' (SEQ ID NO: 627).

TABLE 4

Summary of top 10 off target sites predicted in silico and activity at each site as detected by the Surveyor assay in HEK293T cells transfected with Cas9 and the indicated sgRNA expression cassettes.

SEQ ID NO.	Guide	Target	Sequence	PAM	Score	Chr	Gene	Intron/ Exon	#MMs	% indels
67	CR3	Guide	GCCTACTCAGACTGTTACTC	–	–	–	–	–	–	–
150		Target	tCCTACTCAGACTGTTACTC	TGG	–	X	DMD	Exon	1	13.0
151		OT1	tCCTACTCAcACTGTTACTC	AGG	7.4	1	STRIP1	Intron	2	9.3
152		OT2	aCCTgCTCAcACTGTTACTC	CAG	2.5	2	ARHGAP25	Intron	3	n.d.
153		OT3	GCaTtCTCAaACTGTTACTC	AGG	2.4	13	None	None	3	n.d.
154		OT4	GgaTtCTCAcACTGTTACTC	GGG	1.3	14	PGPEP1	Exon	4	n.d.
155		OT5	aCaTACTtAtACTGTTACTC	TAG	1.3	19	MDGA2	Intron	4	n.d.
156		OT6	tatTcCTaAGACTGTTACTC	AAG	0.9	8	LPPR1	Intron	5	n.d.
157		OT7	aaggACTaAGACTGTTACTC	GGG	0.9	9	RNF122	Intron	5	n.d.
158		OT8	GagctCTCAtACTGTTACTC	TAG	0.8	3	DNMBP	Exon	5	n.d.
159		OT9	GCaaAaTgAGACTGTTACTC	CAG	0.8	5	SLC12A2	Intron	4	n.d.
160		OT10	cCtcAtTCAGACTGTTACTC	AAG	0.8	4	KCNIP4	Intron	4	n.d.
65	CR1	Guide	GATTGGCTTTGATTTCCCTA	–	–	–	–	–	–	–
161		Target	cATTGGCTTTGATTTCCCTA	GGG	–	X	DMD	Intron	1	8.3
162		OT1	aATTGGCATTGATTTCCCTA	GAG	7.1	16	None	None	2	0.8
163		OT2	cATTGGCTTTaATTTCCCTA	TAG	4.8	4	None	None	2	n.d.
164		OT3	GATaGGCTgTGATTTCCCTA	GAG	3.9	9	None	None	2	n.d.
165		OT4	GAaTaGCcTTGATTTCCCTA	AAG	2.4	1	None	None	3	n.d.
166		OT5	aATTtGCTTTGATTTCCCTg	AGG	1.5	1	TIMM17A	Intron	3	n.d.
167		OT6	GATgtGCTTTGATTTCCCTt	GGG	1.4	17	MYO1D	Intron	3	n.d.
168		OT7	aATTGGtTTTaATTTCCCTA	AAG	1.1	8	PIK1A	Intron	3	n.d.
169		OT8	aATTGGgTTTGATTTCCCTt	TGG	1.1	11	MS4A1	Intron	3	n.d.
170		OT9	GATgGGtTTTtATTTCCCTA	GAG	1.0	11	None	None	3	n.d.
171		OT10	GAaTGGtTTTGATTTCCCTg	GAG	1.0	11	None	None	3	n.d.
69	CR5	Guide	GCAGTTGCCTAAGAACTGGT	–	–	–	–	–	–	–
172		Target	aCAGTTGCCTAAGAACTGGT	GGG	–	X	DMD	Intron	1	14.0
173		OT1	cCAGTTGtCTAAGAACTGGg	GAG	1.5	5	NRG1	Intron	3	n.d.
174		OT2	GCAGTTGCCTgtGAACTGGT	AGG	1.4	X	None	None	2	n.d.
175		OT3	GCAGaTGCagAAGAACTGGT	GAG	1.4	19	SMIM7	Intron	3	n.d.
176		OT4	GCAGTTcCagAAGAACTGGT	GAG	0.9	11	GLB1L2	Intron	3	n.d.
177		OT5	caAcTTGCCTAtGAACTGGT	AGG	0.7	8	ASAP1	Intron	4	n.d.
178		OT6	aCAccTGCCTAAGAACTGGa	GGG	0.7	11	None	None	4	n.d.
179		OT7	tCAGgTGgCTAAGAACTGGg	TGG	0.7	14	NIN	Intron	4	n.d.
180		OT8	GaAGTTGgCcAAGAACTGGa	GAG	0.6	7	None	None	4	n.d.
181		OT9	GCtGcTGCCcAAGAACTGGc	AGG	0.6	11	AMOTL1	Intron	4	n.d.
182		OT10	tCAGcTGgCTAAGAACgGGT	AAG	0.6	7	ACTR3C	Intron	4	n.d.
70	CR6	Guide	GGGGCTCCACCCTCACGAGT	–	–	–	–	–	–	–
183		Target	aGGGCTCCACCCTCACGAGT	GGG	–	X	DMD	Intron	1	19.9
184		OT1	GcaGCTCagCCCTCACGAGT	CAG	0.8	3	None	None	4	n.d.
185		OT2	GGGGCTtCAgCaTCACGAGT	GAG	0.8	8	None	None	3	n.d.
186		OT3	GGGGCTCtcCCCTCACtAGT	GAG	0.6	8	None	None	3	n.d.
187		OT4	GGGGaTCCACCtTCACcAGT	CAG	0.6	2	None	None	3	n.d.
188		OT5	aGGGCTggACCCTCACaAGT	AAG	0.4	16	AXIN1	Intron	4	n.d.
189		OT6	tGGtCTCCtCCCcCACGAGT	GGG	0.4	2	None	None	4	n.d.
190		OT7	aGGGCTCCcaCCcCACGAGT	GAG	0.3	5	None	None	4	n.d.
191		OT8	GaGGCTCCAtaCTCACcAGT	GAG	0.3	11	None	None	4	n.d.
192		OT9	GGaGCTgCcCCtTCACGAGT	GGG	0.3	3	None	None	4	n.d.
193		OT10	atGaCTCCACCCTCAaGAGT	AAG	0.3	8	AGPAT5	None	4	n.d.
100	CR36	Guide	GCCTTCTTTATCCCCTATCG	–	–	–	–	–	–	–
194		Target	GCCTTCTTTATCCCCTATCG	AGG	–	X	DMD	Intron	0	20.6
195		OT1	GtCTgCTgTgTCCCCTATCG	GGG	1.3	21	None	None	4	n.d.
196		OT2	cCCTTCTcTATCCCCTgTCG	TGG	1.3	8	None	None	3	n.d.
197		OT3	GCCTTCTTTATCCCCTcTCt	TGG	0.9	10	None	None	2	0.5
198		OT4	GCgcTCTTTtTCCCCTATCt	TAG	0.6	16	None	None	4	n.d.
199		OT5	GCCcTCTgTcTCCCCTgTCG	CAG	0.5	1	NFASC	None	4	n.d.
200		OT6	tCCATCTtTgTCCCCTATtG	AGG	0.5	10	None	None	4	n.d.
201		OT7	aCCtTCTCTcTCCCCTATaG	AGG	0.5	5	LOC100996485	Intron	4	n.d.

TABLE 4-continued

Summary of top 10 off target sites predicted in silico and activity at each site as detected by the Surveyor assay in HEK293T cells transfected with Cas9 and the indicated sgRNA expression cassettes.

SEQ ID NO.	Guide	Target	Sequence	PAM	Score	Chr	Gene	Intron/ Exon	#MMs	% indels
202		OT8	GttTTCTTTtTCCCCTATgG	GAG	0.5	3	None	None	4	n.d.
203		OT9	tgCTTCTTaATCCCCTATCa	AAG	0.4	7	None	None	4	n.d.
204		OT10	aCCTTCTTacTCCCCTATCc	GGG	0.4	10	ADARB2	None	4	n.d.

n.d.: not detected.

[0328] The resulting PCR products were randomly melted and reannealed in a thermal cycler with the program: 95° C. for 240 s, followed by 85° C. for 60 s, 75° C. for 60 s, 65° C. for 60 s, 55° C. for 60 s, 45° C. for 60 s, 35° C. for 60 s, and 25° C. for 60 s with a −0.3° C./s rate between steps. Following reannealing, 8 μL of PCR product was mixed with 1 μL of Surveyor Nuclease S and 1 μL of Enhancer S (Transgenomic) and incubated at 42° C. for 1 hr. After incubation, 6 μL of digestion product was loaded onto a 10% TBE polyacrylamide gel and run at 200V for 30 min. The gels were stained with ethidium bromide and quantified using ImageLab (Bio-Rad) by densitometry as previously described (Guschin, et al. *Meth Mol Biol* 649, 247-256 (2010)).

[0329] Fluorescence-activated cell sorting of myoblasts. DMD myoblasts were electroporated with 5 micrograms each of hCas9-T2A-GFP and sgRNA expression vectors and incubated at 37° C. and 5% C02. Three days after electroporation, cells were trypsinized and collected for FACS sorting using a FACSvantage II sorting machine. GFP-positive cells were collected and grown for analysis.

[0330] PCR-based assay to detect genomic deletions. The exon 51 or exon 45-55 loci were amplified from genomic DNA by PCR (Invitrogen AccuPrime High Fidelity PCR kit) using primers flanking each locus. The flanking primers were CelI-CR1/2-F and CelI-CR5-R for exon 51 or CelI-CR6-F and CelI-CR36-R for exon 45-55 analysis (Table 4). PCR products were separated on TAE-agarose gels and stained with ethidium bromide for analysis.

[0331] PCR-based detection of translocations. Loci with predicted possible translocations were amplified by a two-step nested PCR (Invitrogen AccuPrime High Fidelity PCR kit for each step) of genomic DNA from cells transfected with Cas9 alone (control) or Cas9 with sgRNA. In the first step, translocations that may occur at each on-target and off-target sgRNA target site were amplified by 35 cycles of PCR using combinations of Surveyor primers for each locus that were modified to include restriction sites to facilitate cloning and sequencing analysis (Table 4). One microliter of each PCR reaction was subjected to a second round of amplification by 35 rounds of PCR using nested primer sets custom designed for each individual predicted translocation (Table 4). Each second nested PCR primer binds within the same approximate region within the primary amplicon; however, each pair was optimized using Primer3 online bioinformatics software to ensure specific detection of each translocation. PCR amplicons corresponding to the expected length of predicted translocations and only present in cells treated with sgRNA were purified (QIAGEN Gel Extraction kit) and analyzed by Sanger sequencing.

[0332] mRNA analysis. Immortalized myoblasts were differentiated into myofibers by replacing the growth medium with DMEM supplemented with 1% insulin-transferrin-selenium (Invitrogen #51500056) and 1% penicillin/streptomycin (Invitrogen #15140) for 5 days before the cells were trypsinized and collected. Total RNA was isolated from these cells using the RNeasy Plus Mini Kit (QIAGEN) according to the manufacturer's instructions. RNA was reverse transcribed to cDNA using the VILO cDNA synthesis kit (Life Technologies #11754) and 1.5 micrograms of RNA for 2 hrs at 42° C. according to the manufacturer's instructions. The target loci were amplified by 35 cycles of PCR with the AccuPrime High Fidelity PCR kit (Invitrogen) using primers annealing to exons 44 and 52 to detect exon 51 deletion by CR1/5 or CR2/5 or primers annealing to exons 44 and 60 to detect exon 45-55 deletion by CR6/36 (Table 4). PCR products were run on TAE-agarose gels and stained with ethidium bromide for analysis. The resolved PCR bands were cloned and analyzed by Sanger sequencing to verify the expected exon junctions. Table 5 lists the sequences of primers used in Example 4.

TABLE 5

SEQ ID NO.	Primer name	Primer sequence	Notes
205	CelI-CR1/2-F	GAGAGGTTATGTGGCTTTACCA	Forward Surveyor primer for CR1/2
206	CelI-CR1-R	AAAAATGCTTCCCACTTTGC	Reverse Surveyor primer for CR1
207	CelI-CR2-R	CTCATTCTCATGCCTGGACA	Reverse Surveyor primer for CR2
208	CelI-CR3-F	GAGTTTGGCTCAAATTGTTACTCTT	Forward Surveyor primer for CR3

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
209	CelI-CR3-R	GGGAAATGGTCTAGGAGAGTAAAGT	Reverse Surveyor primer for CR3
210	CelI-CR4/31-F	GTTTGGCTCAAATTGTTACTCTTCA	Forward Surveyor primer for CR4 or CR31
211	CelI-CR4/31-R	GTGAGAGTAATGTGTTTGCTGAGAG	Reverse Surveyor primer for CR4 or CR31
212	CelI-CR5-F	CGGGCTTGGACAGAACTTAC	Forward Surveyor primer for CR5
213	CelI-CR5-R	CTGCGTAGTGCCAAAACAAA	Reverse Surveyor primer for CR5
214	CelI-CR6-F	TAATTTCATTGAAGAGTGGCTGAA	Forward Surveyor primer for CR6
215	CelI-CR6-R	AAGCCCTGTGTGGTAGTAGTCAGT	Reverse Surveyor primer for CR6
216	CelI-CR7-F	TGAGTCATGTTGGATAACCAGTCT	Forward Surveyor primer for CR7
217	CelI-CR7-R	GAAGGTCAGGAACATACAATTCAA	Reverse Surveyor primer for CR7
218	CelI-CR10/11-F	GATATGGGCATGTCAGTTTCATAG	Forward Surveyor primer for CR10 or CR11
219	CelI-CR10/11-R	TGCTGTTGATTAATGGTTGATAGG	Reverse Surveyor primer for CR10 or CR11
220	CelI-CR12/13-F	TTTTAAATTGCCATGTTTGTGTC	Forward Surveyor primer for CR12 or CR13
221	Cell-CR12/13-R	ATGAATAACCTAATGGGCAGAAAA	Reverse Surveyor primer for CR12 or CR13
222	CelI-CR14/15-F	TCAAGTCGCTTCATTTTGATAGAC	Forward Surveyor primer for CR14 or CR15
223	CelI-CR14/15-R	CACAACAAAACATATAGCCAAAGC	Reverse Surveyor primer for CR14 or CR15
224	CelI-CR16/17-F	TGCTGCTAAAATAACACAAATCAGT	Forward Surveyor primer for CR16 or CR17
225	CelI-CR16/17-R	CTGTGCCTATTGTGGTTATCCTG	Reverse Surveyor primer for CR16 or CR17
226	CelI-CR18/19-F	ATTGATCTGCAATACATGTGGAGT	Forward Surveyor primer for CR18 or CR19
227	CelI-CR18/19-R	TTTGCCTCTGCTATTACAGTATGG	Reverse Surveyor primer for CR18 or CR19
228	CelI-CR20/21-F	TGTAGGGTGGTTGGCTAAAATAAT	Forward Surveyor primer for CR20 or CR21
229	CelI-CR20/21-R	TTTTTGCACAGTCAATAACACAAA	Reverse Surveyor primer for CR20 or CR21
230	CelI-CR22/23-F	GGCTGGTCTCACAATTGTACTTTA	Forward Surveyor primer for CR22 or CR23
231	CelI-CR22/23-R	CATTATGGACTGAAAATCTCAGCA	Reverse Surveyor primer for CR22 or CR23
232	CelI-CR24/25-F	ATCATCCTAGCCATAACACAATGA	Forward Surveyor primer for CR24 or CR25
233	CelI-CR24/25-R	TTCAGCTTTAACGTGATTTTCTGT	Reverse Surveyor primer for CR24 or CR25
234	CelI-CR26/27-F	GGATTCAGAAGCTGTTTACGAAGT	Forward Surveyor primer for CR26 or CR27

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
235	CelI-CR26/27-R	TTTAGCTGGATTGGAAAAACAAAT	Reverse Surveyor primer for CR26 or CR27
236	CelI-CR28/29-F	AACTCACCCCATTGTTGGTATATT	Forward Surveyor primer for CR28 or CR29
237	CelI-CR28/29-R	CCTTGTCCAAATACCGAAATACAT	Reverse Surveyor primer for CR28 or CR29
238	CelI-CR33-F	CACATAATTCATGAACTTGGCTTC	Forward Surveyor primer for CR33
239	CelI-CR33-R	TAGTAGCTGGGGAGGAAGATACAG	Reverse Surveyor primer for CR33
240	CelI-CR34-F	TTTTTGTTTTAATTGCGACTGTGT	Forward Surveyor primer for CR34
241	CelI-CR34-R	AGAAAAGGGGTTTTCTTTTGACTT	Reverse Surveyor primer for CR34
242	CelI-CR35-F	CATTGTGACTGGATGAGAAGAAAC	Forward Surveyor primer for CR35
243	CelI-CR35-R	AACGGCTGTTATTAAAGTCCTCAG	Reverse Surveyor primer for CR35
244	CelI-CR36-F	CAAGTCAGAAGTCACTTGCTTTGT	Forward Surveyor primer for CR36
245	CelI-CR36-R	TTTTATGTGCAGGAATCAGTCTGT	Reverse Surveyor primer for CR36
246	Dys-E44-F	TGGCGGCGTTTTCATTAT	Forward RT-PCR primer binding in exon 44
247	Dys-E52-R	TTCGATCCGTAATGATTGTTCTAGCC	Reverse RT-PCR primer binding in exon 52
248	Dys-E60-R	GGTCTTCCAGAGTGCTGAGG	Reverse RT-PCR primer binding in exon 60
249	CR3-CelI-OT1-F	TGTGTGCTTCTGTACACATCATCT	Forward Surveyor primer for CR3 off-target 1
250	CR3-CelI-OT1-R	AGATTTCAACCCTCAAAAACTGAG	Reverse Surveyor primer for CR3 off-target 1
251	CR3-CelI-OT2-F	TAAACTCTTTCTTTTCCGCAATTC	Forward Surveyor primer for CR3 off-target 2
252	CR3-CelI-OT2-R	CAAGGTGACCTGCTACCTAAAAAT	Reverse Surveyor primer for CR3 off-target 2
253	CR3-CelI-OT3-F	TATGACCAAGGCTATGTGTTCACT	Forward Surveyor primer for CR3 off-target 3
254	CR3-CelI-OT3-R	ACAGCCTCTCTCCAGTAACATTCT	Reverse Surveyor primer for CR3 off-target 3
255	CR3-CelI-OT4-F	TATTCTTGCAGTGGTTTCACATTT	Forward Surveyor primer for CR3 off-target 4
256	CR3-CelI-OT4-R	ATATTTTAAGCCAAGACCCAACAA	Reverse Surveyor primer for CR3 off-target 4
257	CR3-CelI-OT5-F	CTTTCAACTGTCTGTCTGATTGCT	Forward Surveyor primer for CR3 off-target 5
258	CR3-CelI-OT5-R	AACAGCCTCTCTTCATTGTTCTCT	Reverse Surveyor primer for CR3 off-target 5
259	CR3-CelI-OT6-F	CTCTGGAACTTGTCTCTGTCTTGA	Forward Surveyor primer for CR3 off-target 6
260	CR3-CelI-OT6-R	CTTTCCTGCGTTCTCATGTTACTA	Reverse Surveyor primer for CR3 off-target 6
261	CR3-CelI-OT7-F	CCTTATATCCGTATCGCTCACTCT	Forward Surveyor primer for CR3 off-target 7

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
262	CR3-CelI-OT7-R	CATATCTGTCTAACTTCCGCACAC	Reverse Surveyor primer for CR3 off-target 7
263	CR3-CelI-OT8-F	ACAGGTGTTATGTTGTCTGCATCT	Forward Surveyor primer for CR3 off-target 8
264	CR3-CelI-OT8-R	ACTCCATTCCCAGATTAGTTATGC	Reverse Surveyor primer for CR3 off-target 8
265	CR3-CelI-OT9-F	CTGTTTTCTTTGTGAGAGTGGAGA	Forward Surveyor primer for CR3 off-target 9
266	CR3-CelI-OT9-R	TGTAAGGTGGTCAAACTTGCTCTA	Reverse Surveyor primer for CR3 off-target 9
267	CR3-CelI-OT10-F	TTTTTCCTAGTACCCACAGATTTTT	Forward Surveyor primer for CR3 off-target 10
268	CR3-CelI-OT10-R	TCCCTGATTCTCTCATTTGTGTTA	Reverse Surveyor primer for CR3 off-target 10
269	CR1-CelI-OT1-F	TTGGGAACATCAGAGAAAGTATGA	Forward Surveyor primer for CR1 off-target 1
270	CR1-CelI-OT1-R	ACAAATTACAGTCTCCTGGGAAAG	Reverse Surveyor primer for CR1 off-target 1
271	CR1-CelI-OT2-F	AGTAGCTTACCTTGGCAGAGAAAA	Forward Surveyor primer for CR1 off-target 2
272	CR1-CelI-OT2-R	TGACATACTGTTACCCTTTGCAGT	Reverse Surveyor primer for CR1 off-target 2
273	CR1-CelI-OT3-F	GAAAGGCTCAGTGAATGTTTGTT	Forward Surveyor primer for CR1 off-target 3
274	CR1-CelI-OT3-R	CACTGCATCATCTCATTAAATCAA	Reverse Surveyor primer for CR1 off-target 3
275	CR1-CelI-OT4-F	CCCATATATTCATGATTACCCACA	Forward Surveyor primer for CR1 off-target 4
276	CR1-CelI-OT4-R	TATCAGAACGAGCACTAAAAGCAC	Reverse Surveyor primer for CR1 off-target 4
277	CR1-CelI-OT5-F	TTGGGAGGCTGAGGTACAAG	Forward Surveyor primer for CR1 off-target 5
278	CR1-CelI-OT5-R	GAATGAAAAACAAACAGAAGGTGA	Reverse Surveyor primer for CR1 off-target 5
279	CR1-CelI-OT6-F	CTCCTCATCTGTACCCTTCAATCT	Forward Surveyor primer for CR1 off-target 6
280	CR1-CelI-OT6-R	AGAGTGGCATCTAGTGTCAGTGAG	Reverse Surveyor primer for CR1 off-target 6
281	CR1-CelI-OT7-F	TACCAAAAGCTTCTCCTGTTTACC	Forward Surveyor primer for CR1 off-target 7
282	CR1-CelI-OT7-R	GTAAGTTGGATGGCCTATTCTTTG	Reverse Surveyor primer for CR1 off-target 7
283	CR1-CelI-OT8-F	GAAGGAAATGCAAGGATACAAGAT	Forward Surveyor primer for CR1 off-target 8
284	CR1-CelI-OT8-R	TGATTGAAAGAATCATTCCAGAAA	Reverse Surveyor primer for CR1 off-target 8
285	CR1-CelI-OT9-F	TCAGAAGGAAAATTGAAATTGGTT	Forward Surveyor primer for CR1 off-target 9

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
286	CR1-CelI-OT9-R	CAGATGTGTTCTTCATCATTCCTC	Reverse Surveyor primer for CR1 off-target 9
287	CR1-CelI-OT10-F	TTCTCTTTAGGGAAAGCTCTCAAA	Forward Surveyor primer for CR1 off-target 10
288	CR1-CelI-OT10-R	GGGTATAGATCATATGGAGGGAAG	Reverse Surveyor primer for CR1 off-target 10
289	CR5-CelI-OT1-F	AGATGATCTGCCCACCTCAG	Forward Surveyor primer for CR5 off-target 1
290	CR5-CelI-OT1-R	CTTTCTTCCTCATTTAGTGGCAAT	Reverse Surveyor primer for CR5 off-target 1
291	CR5-CelI-OT2-F	ATGAATTGCAGATTGATGGTACTG	Forward Surveyor primer for CR5 off-target 2
292	CR5-CelI-OT2-R	TCTCACCAAGAACCAAATTGTCTA	Reverse Surveyor primer for CR5 off-target 2
293	CR5-CelI-OT3-F	GTAGGATACCTTGGCAACAGTCTT	Forward Surveyor primer for CR5 off-target 3
294	CR5-CelI-OT3-R	TTAACGAATTGTGAGATTTGCTGT	Reverse Surveyor primer for CR5 off-target 3
295	CR5-CelI-OT4-F	TCAGAAAGTCAAGTAGCACACACA	Forward Surveyor primer for CR5 off-target 4
296	CR5-CelI-OT4-R	AGAAGCACACACTCAGGTAAAGC	Reverse Surveyor primer for CR5 off-target 4
297	CR5-CelI-OT5-F	TCTTTGGGGGAATAATGACTAAAA	Forward Surveyor primer for CR5 off-target 5
298	CR5-CelI-OT5-R	TTTGGCATTTATGGGAATAAAACT	Reverse Surveyor primer for CR5 off-target 5
299	CR5-CelI-OT6-F	ACTAATTCTGGTCAAGCCCATCA	Forward Surveyor primer for CR5 off-target 6
300	CR5-CelI-OT6-R	TTAAGACATCGGATGAACAGAAAG	Reverse Surveyor primer for CR5 off-target 6
301	CR5-CelI-OT7-F	AGAAGCTTTCTGACATGATCTGC	Forward Surveyor primer for CR5 off-target 7
302	CR5-CelI-OT7-R	TCAATTGCATTAGGACTTAGACCA	Reverse Surveyor primer for CR5 off-target 7
303	CR5-CelI-OT8-F	GTTAAATTACCTGTGAAGCCCTTG	Forward Surveyor primer for CR5 off-target 8
304	CR5-CelI-OT8-R	CGGAAAACAGATCCACTTTATGAT	Reverse Surveyor primer for CR5 off-target 8
305	CR5-CelI-OT9-F	AAATCCACTGGAAACATCTTGAGT	Forward Surveyor primer for CR5 off-target 9
306	CR5-CelI-OT9-R	AGTCTCTTCAGAATCATGCCCTAT	Reverse Surveyor primer for CR5 off-target 9
307	CR5-CelI-OT10-F	GCTTGGTGGCACATACCTGTAG	Forward Surveyor primer for CR5 off-target 10
308	CR5-CelI-OT10-R	GGTAGGTAGATTTGCTTGCTTGTT	Reverse Surveyor primer for CR5 off-target 10
309	CR6-CelI-OT1-F	AGCTCTCAGCAGAGTAGGGATTTA	Forward Surveyor primer for CR6 off-target 1

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
310	CR6-CelI-OT1-R	GTGAGTCTACTGCACCCCATC	Reverse Surveyor primer for CR6 off-target 1
311	CR6-CelI-OT2-F	TGACACTGTGAAGTCAATTCTGTC	Forward Surveyor primer for CR6 off-target 2
312	CR6-CelI-OT2-R	TCAAGAACTTGACAATGAGCAAAT	Reverse Surveyor primer for CR6 off-target 2
313	CR6-CelI-OT3-F	TATCCGATCCACTGTTGTGTGT	Forward Surveyor primer for CR6 off-target 3
314	CR6-CelI-OT3-R	CAGGAGACCCAAAACCACTCTAC	Reverse Surveyor primer for CR6 off-target 3
315	CR6-CelI-OT4-F	TTGTTCTACAAATAGGGCTTCCTT	Forward Surveyor primer for CR6 off-target 4
316	CR6-CelI-OT4-R	TGTTAAGTTTGGGCTTATGTTCCT	Reverse Surveyor primer for CR6 off-target 4
317	CR6-CelI-OT5-F	CACAAGTCTCACTGCACAAACAT	Forward Surveyor primer for CR6 off-target 5
318	CR6-CelI-OT5-R	TGACCCATGATTATCTCTCTTTGA	Reverse Surveyor primer for CR6 off-target 5
319	CR6-CelI-OT6-F	TTCAGCTTCTGATTGGTTTTAATG	Forward Surveyor primer for CR6 off-target 6
320	CR6-CelI-OT6-R	CCAATTCCTTAATTTTCCCTACAG	Reverse Surveyor primer for CR6 off-target 6
321	CR6-CelI-OT7-F	ATCTCAGACCAGGAGGGAGAC	Forward Surveyor primer for CR6 off-target 7
322	CR6-CelI-OT7-R	CCTCAGGGTCAGTACATTTTTCAG	Reverse Surveyor primer for CR6 off-target 7
323	CR6-CelI-OT8-F	TTCTTAGGACATTGCTCCACATAC	Forward Surveyor primer for CR6 off-target 8
324	CR6-CelI-OT8-R	GCAAACATAATGCAACTCGTAATC	Reverse Surveyor primer for CR6 off-target 8
325	CR6-CelI-OT9-F	GCAAGGGAGTCTGTGTCTTTG	Forward Surveyor primer for CR6 off-target 9
326	CR6-CelI-OT9-R	TCATTTAAGTGGCTGTTCTGTGTT	Reverse Surveyor primer for CR6 off-target 9
327	CR6-CelI-OT10-F	ACAAAACAGAGAGAAAAGGCAGAG	Forward Surveyor primer for CR6 off-target 10
328	CR6-CelI-OT10-R	GTTTTGATTTCTGGTGCCTACAG	Reverse Surveyor primer for CR6 off-target 10
329	CR36-CelI-OT1-F	ACTGAAGCTGAAGCCCAGTC	Forward Surveyor primer for CR36 off-target 1
330	CR36-CelI-OT1-R	ACATGAGCTCTCAGGTTTCTGAC	Reverse Surveyor primer for CR36 off-target 1
331	CR36-CelI-OT2-F	TCAAACTTAGATGGTTCCCTATGTT	Forward Surveyor primer for CR36 off-target 2
332	CR36-CelI-OT2-R	GTACCCTGAAAATGTAGGGTGACT	Reverse Surveyor primer for CR36 off-target 2
333	CR36-CelI-OT3-F	CACTTCCCAAGTGAGGCAAT	Forward Surveyor primer for CR36 off-target 3

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
334	CR36-CelI-OT3-R	CTATACTTGGGGCTGACTTGCTAC	Reverse Surveyor primer for CR36 off-target 3
335	CR36-CelI-OT4-F	TCGTATAGGTTACTTTGGCTCACA	Forward Surveyor primer for CR36 off-target 4
336	CR36-CelI-OT4-R	AGGGATCTTTACTCCTCAGTGTGT	Reverse Surveyor primer for CR36 off-target 4
337	CR36-CelI-OT5-F	TGTAGAAGTTGGAATATCCTGCTG	Forward Surveyor primer for CR36 off-target 5
338	CR36-CelI-OT5-R	GTCAACAATTTGATCTCAGGCTTC	Reverse Surveyor primer for CR36 off-target 5
339	CR36-CelI-OT6-F	CTCAGTACTAAAGATGGACGCTTG	Forward Surveyor primer for CR36 off-target 6
340	CR36-CelI-OT6-R	AATCATTTCAGTCTTCCCAACAAT	Reverse Surveyor primer for CR36 off-target 6
341	CR36-CelI-OT7-F	GGGAATCACAGTAGATGTTTGTCA	Forward Surveyor primer for CR36 off-target 7
342	CR36-CelI-OT7-R	AGACCAGGAGGTAAGAACATTTTG	Reverse Surveyor primer for CR36 off-target 7
343	CR36-CelI-OT8-F	CCACATAGAAAGAGACTTGCAGAA	Forward Surveyor primer for CR36 off-target 8
344	CR36-CelI-OT8-R	AGAGATGCCAAAAGAACAGTCAAT	Reverse Surveyor primer for CR36 off-target 8
345	CR36-CelI-OT9-F	TGTGCCTTAGGCTATGTAAACTGT	Forward Surveyor primer for CR36 off-target 9
346	CR36-CelI-OT9-R	AAACCCTTGTAACCAAAATTACCA	Reverse Surveyor primer for CR36 off-target 9
347	CR36-CelI-OT10-F	TAACTGCATCAGAAGTCCTTGCTA	Forward Surveyor primer for CR36 off-target 10
348	CR36-Cell-OT10-R	GGAGACCAAGCTGCTAAAGTCA	Reverse Surveyor primer for CR36 off-target 10
349	CelI-CR3-F-nested	GTGGTGccgcggGAGTTTGGCTCAAAT TGTTACTCTT	Nested PCR first round primers
350	CelI-CR3-R-nested	GTGGTGccgcggGGGAAATGGTCTAGG AGAGTAAAGT	Nested PCR first round primers
351	CelI-CR1-F-nested	GTGGTGccgcggGAGAGGTTATGTGGC TTTACCA	Nested PCR first round primers
352	CelI-CR1-R-nested	GTGGTGccgcggCTCATTCTCATGCCTG GACA	Nested PCR first round primers
353	CelI-CR5-F-nested	GTGGTGccgcggCGGGCTTGGACAGAA CTTAC	Nested PCR first round primers
354	CelI-CR5-R-nested	GTGGTGccgcggCTGCGTAGTGCCAAA ACAAA	Nested PCR first round primers
355	CelI-CR6-F-nested	GTGGTGccgcggTAATTTCATTGAAGA GTGGCTGAA	Nested PCR first round primers
356	CelI-CR6-R-nested	GTGGTGccgcggAAGCCCTGTGTGGTA GTAGTCAGT	Nested PCR first round primers
357	CelI-CR36-F-nested	GTGGTGccgcggCAAGTCAGAAGTCAC TTGCTTTGT	Nested PCR first round primers

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
358	CelI-CR36-R-nested	GTGGTGccgcggTTTTATGTGCAGGAA TCAGTCTGT	Nested PCR first round primers
359	CR3-CelI-OT1-F-nested	GTGGTGccgcggTGTGTGCTTCTGTACA CATCATCT	Nested PCR first round primers
360	CR3-CelI-OT1-R-nested	GTGGTGccgcggAGATTTCAACCCTCA AAAACTGAG	Nested PCR first round primers
361	CR1-CelI-OT1-F-nested	GTGGTGccgcggTTGGGAACATCAGAG AAAGTATGA	Nested PCR first round primers
362	CR1-CelI-OT1-R-nested	GTGGTGccgcggACAAATTACAGTCTC CTGGGAAAG	Nested PCR first round primers
363	CR36-CelI-OT3-F-nested	GTGGTGccgcggCACTTCCCAAGTGAG GCAAT	Nested PCR first round primers
364	CR36-CelI-OT3-R-nested	GTGGTGccgcggCTATACTTGGGGCTG ACTTGCTAC	Nested PCR first round primers
365	CR3-P1/P3-F	GTGGTGccgcggTTGGCTCTTTAGCTTG TGTTTC	Nested PCR second round primers
366	CR3-P1/P3-R	GTGGTGccgcggTGAGACTCCCAAAGG CAATC	Nested PCR second round primers
367	CR3-P1/P4-F	GTGGTGccgcggTTGGCTCTTTAGCTTG TGTTTC	Nested PCR second round primers
368	CR3-P1/P4-R	GTGGTGccgcggACTGAGGGGTGATCT TGGTG	Nested PCR second round primers
369	CR3-P2/P3-F	GTGGTGccgcggGCAGAGAAAGCCAGT CGGTA	Nested PCR second round primers
370	CR3-P2/P3-R	GTGGTGccgcggTGAGACTCCCAAAGG CAATC	Nested PCR second round primers
371	CR3-P2/P4-F	GTGGTGccgcggGCAGAGAAAGCCAGT CGGTA	Nested PCR second round primers
372	CR3-P2/P4-R	GTGGTGccgcggACTGAGGGGTGATCT TGGTG	Nested PCR second round primers
373	CR1-P1/P5-F	GTGGTGccgcggCCAGAGTTCCTAGGG CAGAG	Nested PCR second round primers
374	CR1-P1/P5-R	GTGGTGccgcggAGCTAGTCCCCACAT TCCAC	Nested PCR second round primers
375	CR1-P1/P6-F	GTGGTGccgcggCCAGAGTTCCTAGGG CAGAG	Nested PCR second round primers
376	CR1-P1/P6-R	GTGGTGccgcggGGTGGAGGGAAACTT TAGGC	Nested PCR second round primers
377	CR1-P2/P5-F	GTGGTGccgcggCTCATTCTCATGCCTG GACA	Nested PCR second round primers
378	CR1-P2/P5-R	GTGGTGccgcggAGCTAGTCCCCACAT TCCAC	Nested PCR second round primers
379	CR1-P2/P6-F	GTGGTGccgcggTCTCATGCCTGGACA AGTAACT	Nested PCR second round primers
380	CR1-P2/P6-R	GTGGTGccgcggGGTGGAGGGAAACTT TAGGC	Nested PCR second round primers
381	CR5-P3/P5-F	GTGGTGccgcggGGCTTGGACAGAACT TACCG	Nested PCR second round primers

TABLE 5-continued

SEQ ID NO.	Primer name	Primer sequence	Notes
382	CR5-P3/P5-R	GTGGTGccgcggCACCACTGTCTGCCTAAGGA	Nested PCR second round primers
383	CR5-P4/P6-F	GTGGTGccgcggGGCTTGGACAGAACTTACCG	Nested PCR second round primers
384	CR5-P4/P6-R	GTGGTGccgcggGGTGGAGGGAAACTTTAGGC	Nested PCR second round primers
385	CR5-P3/P5-F	GTGGTGccgcggCGTAGTGCCAAAACAAACAGT	Nested PCR second round primers
386	CR5-P3/P5-R	GTGGTGccgcggCACCACTGTCTGCCTAAGGA	Nested PCR second round primers
387	CR5-P4/P6-F	GTGGTGccgcggCGTAGTGCCAAAACAAACAGT	Nested PCR second round primers
388	CR5-P4/P6-R	GTGGTGccgcggGGTGGAGGGAAACTTTAGGC	Nested PCR second round primers
389	CR6-P1/P5-F	GTGGTGccgcggGCGAGGGCCTACTTGATATG	Nested PCR second round primers
390	CR6-P1/P5-R	GTGGTGccgcggCTTCCCAAGTGAGGCAATGC	Nested PCR second round primers
391	CR6-P1/P6-F	GTGGTGccgcggACGTTTTGTGCTGCTGTAACA	Nested PCR second round primers
392	CR6-P1/P6-R	GTGGTGccgcggCTGCAGGCACATTCTCTTCC	Nested PCR second round primers
393	CR6-P2/P5-F	GTGGTGccgcggGCCCTGTGTGGTAGTAGTCA	Nested PCR second round primers
394	CR6-P2/P5-R	GTGGTGccgcggCTTCCCAAGTGAGGCAATGC	Nested PCR second round primers
395	CR6-P2/P6-F	GTGGTGccgcggCAGTATTAAGGGGTGGGAGCT	Nested PCR second round primers
396	CR6-P2/P6-R	GTGGTGccgcggTCTCTTCCTCACACAGCTGA	Nested PCR second round primers
397	CR36-P3/P5-F	GTGGTGccgcggGGAGCTTGGAGGGAAGAGAA	Nested PCR second round primers
398	CR36-P3/P5-R	GTGGTGccgcggCTTCCCAAGTGAGGCAATGC	Nested PCR second round primers
399	CR36-P4/P6-F	GTGGTGccgcggATGGATGGGGAAGACACTGG	Nested PCR second round primers
400	CR36-P4/P6-R	GTGGTGccgcggCTGCAGGCACATTCTCTTCC	Nested PCR second round primers
401	CR36-P3/P5-F	GTGGTGccgcggGGATGAAACAGGGCAGGAAC	Nested PCR second round primers
402	CR36-P3/P5-R	GTGGTGccgcggTTCCCAAGTGAGGCAATGC	Nested PCR second round primers
403	CR36-P4/P6-F	GTGGTGccgcggTTTGCAGAGCCATGATGAGG	Nested PCR second round primers
404	CR36-P4/P6-R	GTGGTGccgcggCGACAGCCAAAACAGCCG	Nested PCR second round primers

[0333] Western blot analysis. To assess dystrophin protein expression, immortalized myoblasts were differentiated into myofibers by replacing the growth medium with DMEM supplemented with 1% insulin-transferrin-selenium (Invitrogen) and 1% antibiotic/antimycotic (Invitrogen) for 4-7 days, such as 6 or 7 days. Fibroblasts were transdifferentiated into myoblasts by inducing MyoD overexpression and incubating the cells in DMEM supplemented with 1% insulin-transferrin-selenium (Invitrogen), 1% antibiotic/antimycotic (Invitrogen) and 3 μg/mL doxycycline for 15 days. Dystrophin expression was assessed at 3 days after transfecting HEK293T cells. Cells were trypsinized, collected and lysed in RIPA buffer (Sigma) supplemented with a protease inhibitor cocktail (Sigma) and the total protein amount was quantified using the bicinchoninic acid assay according to the manufacturer's instructions (Pierce). Samples were then mixed with NuPAGE loading buffer (Invitrogen) and 5% β-mercaptoethanol and heated to 85° C. for 10 minutes. Twenty-five micrograms of protein were separated on 4-12% NuPAGE Bis-Tris gels (Invitrogen) with MES buffer (Invitrogen). Proteins were transferred to nitrocellulose membranes for 1-2 hrs in transfer buffer containing 10-20% methanol, such as 10% methanol, and 0.01% SDS. The blot was then blocked for 1 hr with 5% milk-TBST at room temperature. Blots were probed with the following primary antibodies: MANDYS8 to detect dystrophin (1:100, Sigma D8168) and rabbit anti-GAPDH (1:5000, Cell Signaling 2118S). Blots were then incubated with mouse or rabbit horseradish peroxidase-conjugated secondary antibodies (Santa Cruz) and visualized using the ChemiDoc chemilumescent system (BioRad) and Western-C ECL substrate (BioRad).

[0334] Transplantation into immunodeficient mice. All animal experiments were conducted under protocols approved by the Duke Institutional Animal Care & Use Committee. Cells were trypsinized, collected and washed in 1× Hank's Balanced Salt Solution (HBSS, Sigma). Two million cells were pelleted and resuspended in five μL 1×HBSS (Sigma) supplemented with cardiotoxin (Sigma #C9759) immediately prior to injection. These cells were transplanted into the hind limb tibialis anterior (TA) muscle of NOD.SCID.gamma (NSG) mice (Duke CCIF Breeding Core) by intramuscular injection. Four weeks after injection, mice were euthanized and the TA muscles were harvested.

[0335] Immunofluorescence staining. Harvested TA muscles were incubated in 30% glycerol overnight at 4° C. before mounting and freezing in Optimal Cutting Temperature compound. Serial 10 micron sections were obtained by cryosectioning of the embedded muscle tissue at −20° C. Cryosections were then washed in PBS to remove the OCT compound and subsequently blocked for 30-60 minutes at room temperature in PBS containing 10% heat-inactivated fetal bovine serum for spectrin detection or 5% heat-inactivated fetal bovine serum for dystrophin detection. Cryosections were incubated overnight at 4° C. with the following primary antibodies that are specific to human epitopes only: anti-spectrin (1:20, Leica NCL-SPEC1) or anti-dystrophin (1:2, Leica NCL-DYS3). After primary staining, spectrin or dystrophin expression was detected using a tyramide-based immunofluorescence signal amplification detection kit (Life Technologies, TSA Kit #22, catalog #T-20932,). Briefly, cryosections were incubated with 1:200 goat anti-mouse biotin-XX secondary (Life Technologies #B2763) in blocking buffer for 1 hr at room temperature. The signal was then amplified using streptavidin-HRP conjugates (1:100, from TSA Kit) in blocking buffer for 1 hr at room temperature. Finally, cryosections were incubated with tyramide-AlexaFluor488 conjugates (1:100, TSA kit) in manufacturer-provided amplification buffer for 10 minutes at room temperature. Stained cryosections were then mounted in ProLong AntiFade (Life Technologies #P36934) and visualized with conventional fluorescence microscopy.

[0336] Cytotoxicity assay. To quantitatively assess potential sgRNA or SpCas9 nuclease-associated cytotoxicity, HEK293T cells were transfected with 10 ng of a GFP reporter and 100 ng SpCas9 expression vector and 100 ng sgRNA expression vector using Lipofectamine 2000 according to the manufacturer's instructions (Invitrogen). The percentage of GFP positive cells was assessed at 2 and 5 days by flow cytometry. The survival rate was calculated as the decrease in GFP positive cells from days 2 to 5 and normalized to cells transfected with an empty nuclease expression vector as described (Cornu et al., Meth Mol Biol 649:237-245 (2010)).

Example 4

CRISPRs Targeting the Dystrophin Gene—Results

[0337] The CRISPR/Cas9-based system was designed to target the dystrophin gene. Various gRNAs were chosen to target different regions of the human and mouse dystrophin gene based on NNNNN NNNNN NNNNN NNNNN NGG (SEQ ID NO: 677) and GNNNN NNNNN NNNNN NNNNN NGG (SEQ ID NO: 678) (see Tables 6, 7 and 8).

TABLE 6

Name (SEQ ID NO)	Species	Gene	Target	Strand	19 bp for chimeric (add G on 5' end)	PAM
DCR1 (628)	Human	DMD	Intron 50	+	attggctttgatttcccta (SEQ ID NO: 65)	GGG
DCR2 (66)	Human	DMD	Intron 50	-	tgtagagtaagtcagccta (SEQ ID NO: 679)	TGG
DCR3 (629)	Human	DMD	Exon 51-55'	+	cctactcagactgttactc (SEQ ID NO: 67)	TGG
DCR4 (68)	Human	DMD	Exon 51-53'	+	ttggacagaacttaccgac (SEQ ID NO: 680)	TGG

TABLE 6-continued

Name (SEQ ID NO)	Species	Gene	Target	Strand	19 bp for chimeric (add G on 5' end)	PAM
DCR5 (630)	Human	DMD	Intron 51	-	cagttgcctaagaactggt (SEQ ID NO: 69)	GGG
DCR6 (631)	Human	DMD	Intron 44	-	GGGCTCCACCCTCACGAG T (SEQ ID NO: 70)	GGG
DCR7 (71)	Human	DMD	Intron 55	+	TTTGCTTCGCTATAAAACG (SEQ ID NO: 681)	AGG
DCR8 (72)	Human	DMD	Exon 41	+	TCTGAGGATGGGGCCGCA A (SEQ ID NO: 682)	TGG
DCR9 (73)	Human	DMD	Exon 44	-	GATCTGTCAAATCGCCTGC (SEQ ID NO: 683)	AGG
DCR10 (74)	Human	DMD	Exon 45	+	CCAGGATGGCATTGGGCA G (SEQ ID NO: 684)	CGG
DCR11 (75)	Human	DMD	Exon 45	+	CTGAATCTGCGGTGGCAG G (SEQ ID NO: 685)	AGG
DCR12 (76)	Human	DMD	Exon 46	-	TTCTTTTGTTCTTCTAGCc (SEQ ID NO: 686)	TGG
DCR13 (77)	Human	DMD	Exon 46	+	GAAAAGCTTGAGCAAGTC A (SEQ ID NO: 687)	AGG
DCR14 (78)	Human	DMD	Exon 47	+	GAAGAGTTGCCCCTGCGC C (SEQ ID NO: 688)	AGG
DCR15 (79)	Human	DMD	Exon 47	+	ACAAATCTCCAGTGGATA A (SEQ ID NO: 689)	AGG
DCR16 (80)	Human	DMD	Exon 48	-	TGTTTCTCAGGTAAAGCTC (SEQ ID NO: 690)	TGG
DCR17 (81)	Human	DMD	Exon 48	+	GAAGGACCATTTGACGTTa (SEQ ID NO: 691)	AGG
DCR18 (82)	Human	DMD	Exon 49	-	AACTGCTATTTCAGTTTCc (SEQ ID NO: 692)	TGG
DCR19 (83)	Human	DMD	Exon 49	+	CCAGCCACTCAGCCAGTG A (SEQ ID NO: 693)	AGG
DCR20 (84)	Human	DMD	Exon 50	+	gtatgcttttctgttaaag (SEQ ID NO: 694)	AGG
DCR21 (85)	Human	DMD	Exon 50	+	CTCCTGGACTGACCACTAT (SEQ ID NO: 695)	TGG
DCR22 (86)	Human	DMD	Exon 52	+	GAACAGAGGCGTCCCCAG T (SEQ ID NO: 696)	TGG
DCR23 (87)	Human	DMD	Exon 52	+	GAGGCTAGAACAATCATT A (SEQ ID NO: 697)	CGG
DCR24 (88)	Human	DMD	Exon 53	+	ACAAGAACACCTTCAGAA C (SEQ ID NO: 698)	CGG
DCR25 (89)	Human	DMD	Exon 53	-	GGTTTCTGTGATTTTCTTT (SEQ ID NO: 699)	TGG
DCR26 (90)	Human	DMD	Exon 54	+	GGCCAAAGACCTCCGCCA G (SEQ ID NO: 700)	TGG
DCR27 (91)	Human	DMD	Exon 54	+	TTGGAGAAGCATTCATAA A (SEQ ID NO: 701)	AGG
DCR28 (92)	Human	DMD	Exon 55	-	TCGCTCACTCACCctgcaa (SEQ ID NO: 702)	AGG

TABLE 6-continued

Name (SEQ ID NO)	Species	Gene	Target	Strand	19 bp for chimeric (add G on 5' end)	PAM
DCR29 (93)	Human	DMD	Exon 55	+	AAAAGAGCTGATGAAACA A (SEQ ID NO: 703)	TGG
DCR30 (94)	Human	DMD	5'UTR/Exon 1	+	TAcACTTTTCaAAATGCTT (SEQ ID NO: 704)	TGG
DCR31 (95)	Human	DMD	Exon 51	+	gagatgatcatcaagcaga (SEQ ID NO: 705)	AGG
DCR32 (96)	Mouse	DMD	mdx mutation	+	ctttgaaagagcaaTaaaa (SEQ ID NO: 706)	TGG
DCR33 (97)	Human	DMD	Intron 44	-	CACAAAAGTCAAATCGGA A (SEQ ID NO: 707)	TGG
DCR34 (98)	Human	DMD	Intron 44	-	ATTTCAATATAAGATTCGG (SEQ ID NO: 708)	AGG
DCR35 (99)	Human	DMD	Intron 55	-	CTTAAGCAATCCCGAACTC (SEQ ID NO: 709)	TGG
DCR36 (632)	Human	DMD	Intron 55	-	CCTTCTTTATCCCCTATCG (SEQ ID NO: 100)	AGG
DCR40 (104)	Mouse	DMD	Exon 23	-	aggccaaacctcggcttac (SEQ ID NO: 710)	NNGRR
DCR41 (105)	Mouse	DMD	Exon 23	+	TTCGAAAATTTCAGgtaag (SEQ ID NO: 711)	NNGRR
DCR42 (106)	Mouse	DMD	Exon 23	+	gcagaacaggagataacag (SEQ ID NO: 712)	NNGRRT
DCR43 (107)	Mouse	ACVR2B	Exon 1	+	gcggccctcgcccttctct (SEQ ID NO: 713)	ggggat
DCR48 (108)	Human	DMD	Intron 45	-	TAGTGATCGTGGATACGA G (SEQ ID NO: 714)	AGG
DCR49 (109)	Human	DMD	Intron 45	-	TACAGCCCTCGGTGTATAT (SEQ ID NO: 715)	TGG
DCR50 (110)	Human	DMD	Intron 52	-	GGAAGGAATTAAGCCCGA A (SEQ ID NO: 716)	TGG
DCR51 (111)	Human	DMD	Intron 53	-	GGAACAGCTTTCGTAGTTG (SEQ ID NO: 718)	AGG
DCR52 (112)	Human	DMD	Intron 54	+	ATAAAGTCCAGTGTCGAT C (SEQ ID NO: 719)	AGG
DCR53 (113)			Intron 54	+	AAAACCAGAGCTTCGGTC A (SEQ ID NO: 720)	AGG
DCR54 (114)	Mouse	Rosa26	ZFN region	+	GAGTCTTCTGGGCAGGCTT AAAGGCTAACC (SEQ ID NO: 720)	TGG
DCR55 (115)	Mouse	Rosa26	mRNA	-	TCGGGTGAGCATGTCTTTA ATCTACCTCGA (SEQ ID NO: 721)	TGG
DCR49 (116)	Human	DMD	Ex 51	-	gtgtcaccagagtaacagt (SEQ ID NO: 722)	ctgagt
DCR50 (117)	Human	DMD	Ex 51	+	tgatcatcaagcagaaggt (SEQ ID NO: 723)	atgag
DCR60 (118)	Mouse	DMD	Exon 23	+	AACTTCGAAAATTTCAGgta (SEQ ID NO: 724)	agccgagg
DCR61 (119)	Mouse	DMD	Intron 22	+	gaaactcatcaaatatgcgt (SEQ ID NO: 725)	gttagtgt

TABLE 6-continued

Name (SEQ ID NO)	Species	Gene	Target	Strand	19 bp for chimeric (add G on 5' end)	PAM
DCR62 (120)	Mouse	DMD	Intron 22	-	tcatttacactaacacgcat (SEQ ID NO: 726)	atttgatg
DCR63 (121)	Mouse	DMD	Intron 22	+	gaatgaaactcatcaaatat (SEQ ID NO: 727)	gcgtgtta
DCR64 (122)	Mouse	DMD	Intron 23	-	tcatcaatatctttgaagga (SEQ ID NO: 728)	ctctgggt
DCR65 (123)	Mouse	DMD	Intron 23	-	tgttttcataggaaaaatag (SEQ ID NO: 729)	gcaagttg
DCR66 (124)	Mouse	DMD	Intron 23	+	aattggaaaatgtgatggga (SEQ ID NO: 730)	aacagata
DCR67 (125)	Human	DMD	Exon 51	+	atgatcatcaagcagaaggt (SEQ ID NO: 731)	atgagaaa
DCR68 (126)	Human	DMD	Exon 51	+	agatgatcatcaagcagaag (SEQ ID NO: 732)	gtatgaga
DCR69 (127)	Human	DMD	Exon 51	-	catttttctcataccttct (SEQ ID NO: 733)	gcttgatg
DCR70 (128)	Human	DMD	Exon 51	+	tcctactcagactgttactc (SEQ ID NO: 734)	tggtgaca
DCR71 (129)	Human	DMD	Exon 51	-	acaggttgtgtcaccagagt (SEQ ID NO: 735)	aacagtct
DCR72 (130)	Human	DMD	Exon 51	-	ttatcatttttctcatacc (SEQ ID NO: 736)	ttctgctt
DCR73 (131)	Human	DMD	Intron 51	-	ttgcctaagaactggtggga (SEQ ID NO: 737)	aatggtct
DCR74 (132)	Human	DMD	Intron 51	-	aaacagttgcctaagaactg (SEQ ID NO: 738)	gtgggaaa
DCR75 (133)	Human	DMD	Intron 51	+	tttcccaccagttcttaggc (SEQ ID NO: 739)	aactgttt
DCR76 (134)	Human	DMD	Intron 50	+	tggctttgatttccctaggg (SEQ ID NO: 740)	tccagctt
DCR77 (135)	Human	DMD	Intron 50	-	tagggaaatcaaagccaatg (SEQ ID NO: 741)	aaacgttc
DCR78 (136)	Human	DMD	Intron 50	-	gaccctagggaaatcaaagc (SEQ ID NO: 742)	caatgaaa
DCR79 (137)	Human	DMD	Intron 44	-	TGAGGGCTCCACCCTCAC GA (SEQ ID NO: 743)	GTGGGTTT
DCR80 (138)	Human	DMD	Intron 44	-	AAGGATTGAGGGCTCCAC CC (SEQ ID NO: 744)	TCACGAGT
DCR81 (139)	Human	DMD	Intron 44	-	GCTCCACCCTCACGAGTG GG (SEQ ID NO: 745)	TTTGGTTC
DCR82 (140)	Human	DMD	Intron 55	-	TATCCCCTATCGAGGAAA CC (SEQ ID NO: 746)	ACGAGTTT
DCR83 (141)	Human	DMD	Intron 55	+	GATAAAGAAGGCCTATTT CA (SEQ ID NO: 747)	TAGAGTTG
DCR84 (142)	Human	DMD	Intron 55	-	AGGCCTTCTTTATCCCCTA T (SEQ ID NO: 748)	CGAGGAAA
DCR85 (143)	Human	DMD	Intron 44	-	TGAGGGCTCCACCCTCAC GA (SEQ ID NO: 749)	GTGGGT

TABLE 6-continued

Name (SEQ ID NO)	Species	Gene	Target	Strand	19 bp for chimeric (add G on 5' end)	PAM
DCR86 (144)	Human	DMD	Intron 55	+	GATAAAGAAGGCCTATTT CA (SEQ ID NO: 750)	TAGAGT

TABLE 7

Name	Notes	% Mod
DCR1	Delete exon 51	6.6
DCR2	Delete exon 51	10.3
DCR3	Frameshift	13
DCR4	Delete exon 51	11.9
DCR5	Delete exon 51	12.4
DCR6	As close to exon 44 as possible in intron 44 (in case of patient deletions)	16.1
DCR7	As close to exon 56 as possible in intron 55 (in case of patient deletions)	6.8
DCR8	Can correct exon 42-43 deletion (−1/+2) only, (−2/+1) is not correctable by this	17.3
DCR9	Skip exon 44 (5')	14.4
DCR10	Frameshift	14.9
DCR11	Correct downstream of exon 45	<1
DCR12	5' splice acceptor/frameshift	<1
DCR13	Correct downstream of exon 46	16.9
DCR14	Frameshift	17.2
DCR15	Correct downstream of exon 47	15.4
DCR16	Frameshift	11.5
DCR17	Correct downstream of exon 48	<1
DCR18	5' splice acceptor/frameshift	1.8
DCR19	Correct downstream of exon 49	33.7
DCR20	5' splice acceptor	14.9
DCR21	Correct downstream of exon 50	24.1

TABLE 7-continued

Name	Notes	% Mod
DCR22	Frameshift	25.9
DCR23	Correct downstream of exon 52	25.2
DCR24	Frameshift (can only correct +1 frame)	24.8
DCR25	Correct downstream of exon 53	2.6
DCR26	Frameshift	24.5
DCR27	Correct downstream of exon 54	13.4
DCR28	5' splice acceptor	21.6
DCR29	Correct downstream of exon 55	19.2
DCR30	Integrate minidys in exon 1	not tested
DCR31	Correct downstream of exon 51	18.9
DCR32	Delete stop codon	not tested
DCR33	Alternative to CR6	1.3
DCR34	Alternative to CR6	13.2
DCR35	Alternative to CR7	22.5
DCR36	Alternative to CR7	26.4
DCR40	Disrupt exon 23 5' splice donor (correct mdx mutation)	
DCR41	Disrupt exon 23 5' splice donor (correct mdx mutation)	
DCR42	Delete exon 53 mdx4cv mutation	
DCR43	Disrupt myostatin receptor	

TABLE 8

Name	Cas9		Notes	Cas9 used
DCR49	*S. Aureus*		Frameshift in exon 51	SaCas9 (from Zhang pX441) (NNGRRT PAM)
DCR50	*S. Aureus*		Disrupt 5' end of exon 51	SaCas9 (from Zhang pX441) (NNGRR PAM)
DCR60	*N. Meningitidis*	NNNNGANN	Target 3' splice donor of exon 23 to bypass mdx mutation	NmCas9 (NNNNGANN PAM)
DCR61	*N. Meningitidis*	NNNNGNNT	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGNNT PAM)
DCR62	*N. Meningitidis*	NNNNGANN	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGANN PAM)
DCR63	*N. Meningitidis*	NNNNGTTN	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGTTN PAM)
DCR64	*N. Meningitidis*	NNNNGNNT	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGNNT PAM)
DCR65	*N. Meningitidis*	NNNNGTTN	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGTTN PAM)
DCR66	*N. Meningitidis*	NNNNGANN	Delete exon 23 and bypass mdx mutation	NmCas9 (NNNNGANN PAM)
DCR67	*N. Meningitidis*	NNNNGANN	Target 3' splice donor of exon 51 to skip exon	NmCas9 (NNNNGANN PAM)
DCR68	*N. Meningitidis*	NNNNGANN	Target 3' splice donor of exon 51 to skip exon	NmCas9 (NNNNGANN PAM)

TABLE 8-continued

Name	Cas9		Notes	Cas9 used
DCR69	*N. Meningitidis*	NNNNGANN	Target 3' splice donor of exon 51 to skip exon	NmCas9 (NNNNGANN PAM)
DCR70	*N. Meningitidis*	NNNNGANN	Frameshift in exon 51	NmCas9 (NNNNGANN PAM)
DCR71	*N. Meningitidis*	NNNNGNNT	Frameshift in exon 51	NmCas9 (NNNNGNNT PAM)
DCR72	*N. Meningitidis*	NNNNGNNT	Target 3' splice donor of exon 51 to skip exon	NmCas9 (NNNNGNNT PAM)
DCR73	*N. Meningitidis*	NNNNGNNT	Delete exon 51 (bind as close to DCR5 as possible)	NmCas9 (NNNNGNNT PAM)
DCR74	*N. Meningitidis*	NNNNGANN	Delete exon 51 (bind as close to DCR5 as possible)	NmCas9 (NNNNGANN PAM)
DCR75	*N. Meningitidis*	NNNNGTTN	Delete exon 51 (bind as close to DCR5 as possible)	NmCas9 (NNNNGTTN PAM)
DCR76	*N. Meningitidis*	NNNNGNNT	Delete exon 51 (bind as close to DCR1/2 as possible)	NmCas9 (NNNNGNNT PAM)
DCR77	*N. Meningitidis*	NNNNGTTN	Delete exon 51 (bind as close to DCR1/2 as possible)	NmCas9 (NNNNGTTN PAM)
DCR78	*N. Meningitidis*	NNNNGANN	Delete exon 51 (bind as close to DCR1/2 as possible)	NmCas9 (NNNNGANN PAM)
DCR79	*N. Meningitidis*	NNNNGNNT	Delete exons 45-55 - overlaps NNNNGTTN PAM, bind as close to DCR6 as possible	NmCas9 (NNNNGNNT PAM)
DCR80	*N. Meningitidis*	NNNNGANN	Delete exons 45-55, bind as close to DCR6 as possible	NmCas9 (NNNNGANN PAM)
DCR81	*N. Meningitidis*	NNNNGTTN	Delete exons 45-55, bind as close to DCR6 as possible	NmCas9 (NNNNGTTN PAM)
DCR82	*N. Meningitidis*	NNNNGNNT	Delete exons 45-55, bind as close to DCR36 as possible - overlaps NNNNGTTN PAM	NmCas9 (NNNNGNNT PAM)
DCR83	*N. Meningitidis*	NNNNGTTN	Delete exons 45-55, bind as close to DCR36 as possible	NmCas9 (NNNNGTTN PAM)
DCR84	*N. Meningitidis*	NNNNGANN	Delete exons 45-55, bind as close to DCR36 as possible	NmCas9 (NNNNGANN PAM)
DCR85	*S. Aureus*	NNGRRT	Delete exons 45-55, bind as close to DCR6 as possible	SaCas9 (from Zhang pX441) (NNGRRT PAM)
DCR86	*S. Aureus*	NNGRRT	Delete exons 45-55, bind as close to DCR36 as possible	SaCas9 (from Zhang pX441) (NNGRRT PAM)

[0338] In particular, 400 ng of Cas9 was co-transfected into HEK 293T cells with either 400 ng of empty vector or gRNA that targets the region encompassing Exon 51, i.e., CR1, CR2, CR3, CR4, and CR5 (see FIG. **11**B). Genomic DNA was harvested at 2 days post-transfection and analyzed using the Surveyor assay (see FIG. **11**A and FIG. **11**C).

[0339] The CRISPR/Cas9-based system was used in DMD 8036 (del148-50) cells to determine if the system could repair a mutant dystrophin gene. 5 μg of Cas9 was co-transfected into DMD 8036 (del48-50) cells with either 7.5 μg of empty vector or gRNA. In particular, 7.5 μg of CR1 ("DCR1"), 7.5 μg of CR5 ("DCR5"), 15 μg of CR3 ("DCR3") or 7.5 μg of a combination of CR1 and CR5 (DCR1+DCR5) were used. Genomic DNA was harvested at 3 days post-transfection and analyzed using the Surveyor assay (FIG. **12**) or PCR analysis across the entire locus (FIG. **13**). This locus was amplified by PCR using primers flanking the region containing the genomic targets for CR1 and CR5 (the forward primer: 5'-gagaggttatgtggctttacca (SEQ ID NO:457), the reverse primer: 5'-ctgcgtagtgccaaaacaaa (SEQ ID NO:458)), resulting in a 1447 bp band for the wild-type locus or an expected size of approximately 630 bp for the deleted locus. After 7 days of differentiation, western blot of the treated cells shows expression of dystrophin protein (see FIG. **14**).

Example 5

Targeting CRISPR/Cas9 to Hotspot Mutations in the Human Dystrophin Gene

[0340] To utilize the CRISPR/Cas9 gene editing platform for correcting a wide range of dystrophin mutations, dozens of sgRNAs targeted to the hotspot mutation region between exons 45-55 were created (FIGS. **16** A-**16**D). The *S. pyogenes* system that utilizes a human-codon optimized SpCas9 nuclease and a chimeric single-guide RNA (sgRNA) expression vector to guide efficient site-specific gene editing was used. Similar to Example 4 targeting exon 51 with TALENs, protospacers were selected to target the 5' and 3' ends of exons 45 through 55 which meet the 5'-NRG-3' PAM requirement of SpCas9. Small insertions or deletions created by NHEJ-based DNA repair within these exons can generate targeted frameshift mutations that address various dystrophin mutations surrounding each exon (FIGS. **16**A-**16**B). For example, CR3 was designed to correct dystrophin mutations or deletions surrounding exon 51 by introducing small insertions or deletions in the 5' end of exon 51 to restore the downstream dystrophin reading frame (FIG. **16**B). Additionally, sgRNAs were designed to employ the multiplex capability of the CRISPR/Cas9 system and specifically delete individual exons or a series of exons to restore the dystrophin reading frame, similar to the methods of oligonucleotide-based exon skipping. For this purpose, sgRNAs were targeted to the intronic regions surrounding exon 51 (FIG. **16**C) or exons 45-55 (FIG. **16**D). These sgRNAs were intentionally targeted to sites nearest to the downstream or upstream exon intended to be included in the resulting transcript to minimize the likelihood that the background patient deletion would include the intronic sgRNA target sites.

Example 6

Screening of sgRNAs Targeted to the Dystrophin Gene in Human Cells

[0341] Gene editing frequency in the human HEK293T cell line was assessed to rapidly determine different sgRNA targeting efficiencies. HEK293 Ts were transfected with constructs encoding human codon-optimized SpCas9 and the indicated sgRNA. Each sgRNA was designed to modify the dystrophin gene as indicated. The frequency of gene modification at day 3 or day 10 post-transfection was determined by the Surveyor assay. The ratio of measured Surveyor signal at day 3 and day 10 was calculated to quantify the stability of gene editing frequencies for each sgRNA in human cells. As quantified by the Surveyor assay 3 days post-transfection, 29/32 (~90%) of the sgRNAs tested were able to mediate highly efficient gene modification at the intended locus (Table 9, FIG. **17**). The gene editing frequencies were stable for almost all of the sgRNAs (<25% signal change from day 3 to day 10, Table 9, FIG. **18**), indicating that gene editing mediated by each individual sgRNA was well-tolerated. A notable exception is CR33, which had no detectable activity at day 10, although activity may be below the sensitivity of the Surveyor assay (est. ~1%).

TABLE 9

Measured activity of sgRNAs in human cells

Target	sgRNA #	% modified alleles at day 3	% modified alleles at day 10	% change day 10/day 3
		Multiplex deletion of exon 51		
Int 50	CR1	6.6	9.3	41.8
Int 50	CR2	10.3	14.0	36.2
Ex 51	CR4	11.9	14.4	21.3
Int 51	CR5	12.4	13.3	7.8
		Multiplex deletion of exons 45-55		
Int 44	CR6	16.1	16.9	4.3
Int 44	CR33	1.3	<1	n.d.
Int 44	CR34	13.2	11.0	−16.6
Int 55	CR7	6.8	7.1	5.3
Int 55	CR35	22.5	20.9	−7.1
Int 55	CR36	26.4	24.7	−6.4
		Targeted frameshifts		
Ex 45	CR10	14.9	16.3	9.3
Ex 45	CR11	<1	<1	n.d.
Ex 46	CR12	<1	<1	n.d.
Ex 46	CR13	16.9	18.4	9.2
Ex 47	CR14	17.2	17.6	2.9
Ex 47	CR15	15.4	15.3	−0.9
Ex 48	CR16	11.5	10.9	−5.0
Ex 48	CR17	<1	<1	n.d.
Ex 49	CR18	1.8	2.2	20.1
Ex 49	CR19	33.7	38.4	13.9
Ex 50	CR20	14.9	13.7	−7.6
Ex 50	CR21	24.1	20.8	−13.5
Ex 51	CR3	13.0	16.7	28.0
Ex 51	CR31	18.9	16.9	−10.2
Ex 52	CR22	25.9	20.3	−21.6
Ex 52	CR23	25.2	24.0	−4.8
Ex 53	CR24	24.8	23.6	−4.6
Ex 53	CR25	2.6	2.9	9.5
Ex 54	CR26	24.5	22.0	−10.1
Ex 54	CR27	13.4	12.6	−5.9
Ex 55	CR28	21.6	19.8	−8.4
Ex 55	CR29	19.2	19.6	2.2

Example 7

Enrichment of Gene-Edited Cells Using a Fluorescence-Based Reporter System

[0342] sgRNAs were selected to correct specific mutations in DMD patient myoblast cell lines. After transfection into DMD myoblasts, unexpectedly low or undetectable gene modification activity was observed as measured by the Surveyor assay (FIG. **19**C, bulk population). Flow cytometry was used to select for transfected cells co-expressing GFP through a 2A ribosomal skipping peptide linked to the SpCas9 protein (FIG. **19**A). The addition of this fluorescent reporter to the SpCas9 expression vector did not seem to significantly impact gene editing activity in HEK293T cells (FIG. **19**B). A low percentage of transfected myoblasts (~0.5-2%) expressed the fluorescent reporter at 3 days after electroporation, despite high transfection efficiencies of control GFP expression plasmids (typically >70%, FIG. **19**D, pmaxGFP). Given the high levels of CRISPR/Cas9 activity in the easily transfected HEK293T line, inefficient transgene expression after electroporation of SpCas9-T2A-GFP and sgRNA constructs into the DMD cells may explain the low observed gene editing efficiencies in unsorted cells. After sorting the GFP-positive DMD myoblasts, a substantial increase was observed in detectable activity at most sgRNA target loci (FIG. **19**C). Therefore, all subsequent experiments used cells sorted for SpCas9 expression by expression of this fluorescent reporter.

Example 8

Restoration of Dystrophin Expression by Targeted Frameshifts

[0343] Small insertions and deletions created by NHEJ DNA repair may be used to create targeted frameshifts to correct aberrant reading frames. A sgRNA, CR3, was designed to restore the dystrophin reading frame by introducing small insertions and deletions within exon 51 (FIG. **16**B and FIG. **20**A). The types of insertions and deletions generated by CRISPR/Cas9 at this locus were assessed by Sanger sequencing of alleles from the genomic DNA of HEK293T cells co-transfected with expression plasmids for SpCas9 and the CR3 sgRNA (FIG. **20**B). Notably, the insertions and deletions resulted in conversion to all three reading frames (FIG. **20**B, FIG. **20**C). To demonstrate genetic correction in a relevant patient cell line, expression plasmids for SpCas9 and the CR3 sgRNA were electroporated into a DMD myoblast line with a deletion of exons 48-50 that is correctable by creating frameshifts in exon 51. The treated cells were sorted, verified to have gene modification activity by the Surveyor assay (CR3, FIG. **19**C sorted population), and differentiated into myotubes to test for restored dystrophin expression. Expression of dystrophin protein was observed concomitant with the detectable nuclease activity (FIG. **20**D). The *S. pyogenes* CRISPR/Cas9 system presents a powerful method to quickly generate targeted frameshifts to address a variety of patient mutations and restore expression of the human dystrophin gene.

Example 9

Multiplex CRISPR/Cas9 Gene Editing Mediates Genomic Deletion of Exon 51 and Rescues Dystrophin Protein Expression

[0344] The multiplexing capability of the CRISPR/Cas9 system presents a novel method to efficiently generate genomic deletions of specific exons for targeted gene correction. DMD patient myoblasts with background deletions correctable by exon 51 skipping were treated with two combinations of sgRNAs flanking exon 51 (CR1/CR5 or CR2/CR5) and sorted to enrich for gene-edited cells as in FIGS. **19**A-**19**D. As detected by end-point PCR of the genomic DNA from these treated cells, the expected genomic deletions were only present when both sgRNAs were electroporated into the cells with SpCas9 (FIG. **21**A). Sanger sequencing confirmed the expected junction of the distal chromosomal segments (FIG. **21**B) for both deletions. After differentiating the sorted myoblasts, a deletion of exon 51 from the mRNA transcript was detected only in the cells treated with both sgRNAs (FIG. **21**C). Finally, restored dystrophin protein expression was detected in the treated cells concomitant with observed genome- and mRNA-level deletions of exon 51 (FIG. **21**D).

Example 10

Dystrophin Rescue by a Multi-Exon Large Genomic Deletion

[0345] Although addressing patient-specific mutations is a powerful use of the CRISPR/Cas9 system, it would be advantageous to develop a single method that can address a myriad of common patient deletions. For example, a promising strategy is to exclude the entire exon 45-55 region as a method to correct up to 62% of known patient deletions. Multiplex CRISPR/Cas9-based gene editing was tested to determine if it may be able to generate efficient deletion of the exon 45-55 locus in human cells. After transfection into HEK293T cells, the expected deletion of ~336,000 bp was detected by PCR of the genomic DNA (FIG. **22**A). Similarly, this deletion was detected by PCR of the genomic DNA from SpCas9/sgRNA-treated DMD patient cells harboring a background deletion of exons 48-50 of unknown length (FIG. **22**A). Sanger sequencing of this deletion band from the genomic DNA of treated DMD cells revealed the expected junctions of intron 44 and intron 55 immediately adjacent to the sgRNA target sites (FIG. **22**B). After differentiation of treated DMD cells, the expected deletion of exons 45-55 was detected in the dystrophin mRNA transcript and verified to be a fusion of exons 44 and 56 by Sanger sequencing (FIG. **22**C). Restored protein expression was observed by western blot in the sorted cell populations containing the CRISPR/Cas9-induced deletion of exons 45-55 from the genome and resulting mRNA transcripts (FIG. **22**D). These data demonstrate that multiplex CRISPR/Cas9 editing presents a single universal method to restore the dystrophin reading frame in more than 60% of DMD patient mutations.

Example 11

Transplantation of Corrected Myoblasts into Immunodeficient Mice

[0346] A promising method for DMD therapy is to correct a population of autologous patient muscle progenitor cells that can be engrafted into the patient's skeletal muscle tissue to rescue dystrophin expression. To demonstrate the ability of the corrected cells to express human dystrophin in vivo, a population of DMD myoblasts that were treated with sgRNAs CR1 and CR5, which flank exon 51, was transplanted and sorted for expression of GFP as before (FIGS. **19**A-**19**D, FIG. **23**). After 4 weeks, muscle fibers positive for human spectrin, which is expressed by both corrected and uncorrected cells, were detected in cryosections of injected muscle tissue (FIG. **24**). A number of these fibers were also positive for human dystrophin with expression localized to the sarcolemma, demonstrating functional gene correction in these cells (FIG. **24**, FIGS. **25**A-**25**F). No fibers positive for human dystrophin were observed in sections from mice injected with the untreated DMD myoblasts (FIG. **24**, FIGS. **25**A-**25**F), indicating that the CRISPR/Cas9-modified cells were the source of human dystrophin expression.

Example 12

Off-Target and Cytotoxicity Analysis

[0347] The relative cytotoxicity of the CRISPR/Cas9 system was assessed in human cells for select sgRNAs by adapting a flow cytometry-based GFP retention assay as previously described (Ousterout et al., Mol Ther 21:1718-1726 (2013)). Minimal cytotoxicity was observed for SpCas9 co-expressed with or without sgRNAs after transfection into human cells (FIG. **26**A). Publicly available tools are available to assess and prioritize potential CRISPR/Cas9 activity at off-target loci based on predicted positional bias of a given mismatch in the sgRNA protospacer sequence and the total number of mismatches to the intended target site

(Hsu et al., Nat Biotechnol 31:827-826 (2013)). This public webserver was used to predict the most likely off-target sites for the sgRNAs used to correct the dystrophin gene in this study (Table 4). The top ten potential off-target sites were assessed by the Surveyor assay in HEK293T cells treated with SpCas9 and individual sgRNA expression cassettes for CR1, CR3, CR5, CR6, or CR36. CR1, CR3 and CR36 each had one of these ten predicted off-target loci demonstrate significant levels of gene modification (Table 4 and FIG. **27**). Interestingly, the CR3 off-target sequence had substantial homology and similar modification frequencies to the intended on-target (9.3% at OT-1 vs. 13.3% at intended site (Table 4 and FIG. **27**). Notably, CR3-OT1 was the only one of these three off-target sites to show significant levels of activity in the sorted hDMD cells by the Surveyor assay (FIG. **26**B).

[0348] Nuclease activity at off-target sites may cause unintended chromosomal rearrangements by distal re-ligation between cleaved target and off-target loci on distinct chromosomes. This presents a significant concern for deletion-based gene correction strategies due to the increased potential for off-target activity by using two or more nucleases, such as in multiplex CRISPR/Cas9 gene editing. Potential translocations were probed for using a highly sensitive nested genomic PCR assay to detect translocations at the validated off-target loci (Table 4) during both single and multiplex CRISPR/Cas9 editing strategies. Using this assay, translocations were readily detected between on-target and off-target sites in the model HEK293T cell line that also shows high levels of off-target activity (FIG. **26**C and FIG. **28**A, **28**B). Sanger sequencing of the PCR amplicons confirmed the identity of the predicted translocation event for each primer pair (FIGS. **29**-**30**). A subset of the translocations detected in the HEK293T cells were also detectable by nested PCR in the sorted hDMD myoblasts, although the signal was considerably weaker and the sequence identity was not confirmed due to low yield of product (FIG. **26**D, FIG. **28**A, and FIG. **28**C). Translocations were not detected using this assay in HEK293T or sorted hDMD cells treated with CR6 or CR6/CR36, respectively, (FIGS. **28**A-**28**C) that had low levels of off-target activity at CR6-OT3 only in HEK293T cells (Table 4). These results underscore the importance of selecting highly specific sgRNAs, particularly for multiplex editing applications, and show that this approach can benefit from ongoing efforts to improve the specificity of the CRISPR/Cas9 system. These data suggest that the selected sgRNAs are able to correct the dystrophin gene without significant toxicity and with only a single strongly predicted off-target site with detectable levels of activity.

Example 13

Discussion

[0349] Genome editing is a powerful tool for correcting genetic disease and the recent development of the CRISPR/Cas9 system is dramatically accelerating progress in this area. The correction of DMD, the most common genetic disease that also currently has no approved therapeutic options, was demonstrated. Many gene- and cell-based therapies for DMD are in preclinical development and clinical trials, and genome editing methods are compatible with many of these approaches. For example, genome editing may be combined with patient-specific cell-based therapies for DMD. The CRISPR/Cas9 system may function in human pluripotent stem cells and other human cell lines, as well as human skeletal myoblasts, as shown. Importantly, gene editing with CRISPR/Cas9 did not abolish the myogenic capacity of these cells, as demonstrated by efficient dystrophin expression in vitro and in vivo after transplantation into immunodeficient mice. Thus, this strategy should be compatible with cell-based therapies for DMD.

[0350] Additionally, an enriched pool of gene-corrected cells demonstrated expression of human dystrophin in vivo following engraftment into immunodeficient mice. CRISPR/Cas9 gene editing did not have significant toxic effects in human myoblasts as observed by stable gene editing frequencies and minimal cytotoxicity of several sgRNAs. However, gene editing activity was confirmed at three out of 50 predicted off-target sites across five sgRNAs and CRISPR/Cas9-induced chromosomal translocations between on-target and off-target sites were detectable. The CRISPR/Cas9 technology is an efficient and versatile method for correcting a significant fraction of dystrophin mutations and can serve as a general platform for treating genetic disease.

[0351] Additionally, direct transfection of the sgRNA and Cas9 mRNA, in contrast to the plasmid-based delivery method used here, may be used to increase specificity and safety by reducing the duration of Cas9 expression and eliminating the possibility of random plasmid integration. Alternatively, delivery of the CRISPR/Cas9 system directly to skeletal and/or cardiac muscle by viral, plasmid, or RNA delivery vectors may be used for in vivo genome editing and translation of this approach. The large size of *S. pyogenes* Cas9 gene (~4.2 kilobases) presents a challenge to its use in size-restricted adeno-associated viral vectors. However, Cas9 genes from other species, such as *N. meningitidis* and *S. thermophilus*, are short enough to efficiently package both Cas9 and sgRNA expression cassettes into single AAV vectors for in vivo gene editing applications.

[0352] The CRISPR/Cas9 system enabled efficient modification of nearly 90% of tested targets, consistent with other reports of robust activity of this system at diverse loci. The robustness and versatility of this technology is a significant advancement towards at-will creation of patient-specific gene editing. Low levels of dystrophin, including as little as 4% of wild-type expression, may be sufficient to improve survival, motor function, and cardiac function in a mouse model. The levels of CRISPR/Cas9 activity may be sufficient for therapeutic benefit.

[0353] The use of multiplexing with CRISPR/Cas9 to delete exons also presents a unique set of opportunities and challenges. The deletion of complete exons from the genome to restore dystrophin expression was performed, in contrast to restoring the reading frame of the dystrophin gene with

small indels generated by NHEJ-based DNA repair following the action of a single nuclease. The protein product of the edited gene is predictable and already characterized in Becker muscular dystrophy patients with the naturally occurring deletion, in contrast to the random indels created by intraexonic action of a single nuclease that will lead to the creation of novel epitopes from each DNA repair event. Furthermore, the product resulting from the exon deletions will lead to restored dystrophin for every successful gene editing event, whereas modifying the gene with random indels within exons will only restore the reading frame in the one-third of editing events that leads to the correct reading frame.

[0354] All of the sgRNAs tested were not associated with significant cytotoxic effects in human cells. Three potential off-target sites out of 50 total tested sites for the five sgRNAs used were identified to restore dystrophin expression. Furthermore, chromosomal translocations between the intended on-target sites and these off-target sites were detectable by highly sensitive nested PCR assays in HEK293T cells expressing high levels of Cas9 and sgRNAs. Notably, the off-target activity and translocations identified in HEK293T cells, which is an immortalized and aneuploid cell line that expresses very high levels of Cas9 and sgRNA, did not occur at as high a level and in some cases were undetectable in the hDMD myoblasts. Importantly, this level of specificity may be tolerable given the severity of DMD, the lack of an apparent cytotoxic effect in human cells

Example 14

[0355] An engineered AAV capsid, termed SASTG (FIG. **34**; SEQ ID NOs: 436 and 437), was developed for enhanced cardiac and skeletal muscle tissue tropism (Piacentino et al. (2012) Human Gene Therapy 23:635-646). A ZFN targeting the Rosa26 locus ("Rosa26 ZFN"; FIG. **33**; SEQ ID NO: 434 and 435) was shown to be highly active in mouse cells (Perez-Pinera et al. Nucleic Acids Research (2012) 40:3741-3752). AAV-SASTG vectors encoding the Rosa26 ZFN protein were designed and subsequently generated and purified by the UNC Viral Vector Core. The Surveyor assay (Guschin et al., Methods Mol Biol 649, 247-256 (2010)) was used to demonstrate NHEJ mutagenesis at the Rosa26 locus following delivery of AAV-SASTG Rosa26 ZFNs in cultured C2C12 myoblasts that were actively cycling or forced into differentiation by serum removal (not shown).

[0356] To verify that adult post-mitotic skeletal muscle were efficiently targeted by the Rosa26 ZFN following AAV transfer, AAV-SASTG vectors encoding the Rosa26 ZFN were injected directly into the tibialis anterior (TA) muscle of 6 week old C57BL6/J mice at titers of 1e10 vector genomes (vg) or 2.5e10 vg per muscle. Mice were sacrificed 4 weeks after injection and the TA muscles were harvested and partitioned into several fragments for genomic DNA extraction and analysis (FIG. **31**). Genomic DNA was PCR amplified and subjected to the Surveyor assay to detect NHEJ mutations characteristic of ZFN mutagenesis at the Rosa26 target site (FIGS. **32**A-**32**C). FIGS. **32**A-**32**C shows Surveyor analysis of Rosa26 ZFN activities in skeletal muscle in vitro and in vivo following delivery of AAV-SASTG-ROSA. Proliferating C2C12s were transduced with the indicated amount of virus and harvested at 4 days post-infection (FIG. **32**A). C2C12s were incubated in differentiation medium for 5 days and then transduced with the indicated amount of AAV-SASTG-ROSA virus in 24 well plates (FIG. **32**B). Samples were collected at 10 days post-transduction. The indicated amount of AAV-SASTG-ROSA was injected directly into the tibialis anterior of C57BL/6J mice and muscles were harvested 4 weeks post-infection. The harvested TA muscles were partitioned into 8 separate pieces for genomic DNA analysis, each shown in a separate lane (FIG. **32**C). Notably, high levels of gene modification were detected in all fragments at the highest dose (2.5e10 vg).

Example 15

AAV-CRISPR Constructs for Targeting Mutant Dystrophin Genes

[0357] AAV constructs are designed to therapeutically correct mutations of the dystrophin gene that cause Duchenne muscular dystrophy and skeletal and cardiac muscle degeneration. CRISPR/Cas9 systems can be delivered using the AAV to restore the dystrophin reading frame by deleting exon 51, deleting exons 45-55, disrupting splice donor or acceptor sites, or creating frameshifts within exon 51 (Ousterout et al., Molecular Therapy 2013) to restore the dystrophin reading frame and protein expression. The CRISPR/Cas9 system will include a Cas9 having a sequence of SEQ ID NO: 64 or 114 (See FIGS. **40** and **41**). gRNAs that could be combined with these Cas9s, targeting their respective PAM sequences, are provided (see FIGS. **40** and **41**; see also Tables 2 and 3).

Example 16

Generation of Induced Neurons (iNs)

[0358] The generation of induced neurons (iNs) from other cell lineages has potential applications in regenerative medicine and the study of neurological diseases. The direct conversion of mouse embryonic fibroblasts (MEFs) to functional neuronal cells may occur through the delivery of a cocktail of three neuronal transcription factors—BRN2, ASCL1, and MYT1L (BAM factors, FIG. **48**). Other methods may include additional factors to induce various neuronal subtypes. These experiments require transcription factors to be delivered ectopically, and the activation of the corresponding endogenous loci to sustain the neuronal phenotype. The CRISPR/Cas9 system was engineered as a versatile transcription factor to activate endogenous genes in mammalian cells with the capacity to target any promoter in the genome through an RNA-guided mechanism (FIGS. **49**A-**49**B).

[0359] Materials & Methods. The CRISPR/Cas9-transcription factor was used to activate the endogenous genes encoding ASCL1 and BRN2 to directly reprogram MEFs to functional induced neurons.

[0360] Cell Culture: MEFs were seeded in either 24-well TCPS plates or on poly-D-lysine/laminin-coated coverslips. Following transduction of dCas9-VP64 and transfection of the gRNAs (see Tables 10 and 11 for sequences of gRNAs), the cells were cultured in MEF medium (Adler et al. Mol Ther Nucleic Acids 1:e32 (2012)) for 24 hrs and then transferred to N3 neural induction medium (Vierbuchen et al. *Nature* 463:1035-1041 (2010)) for the duration of the experiment (FIG. **49**B).

TABLE 10

gRNAs for mouse ASCL1 (CR13):

Oligo (5' to 3')	5' overhang	ASCL1 Target Sequence	SEQ ID NO
CR13-1_S:	cacc	G CAGCCGCTCGCTGCAGCAG (SEQ ID NO: 468)	492
CR13-1_AS:	AAAC	CTGCTGCAGCGAGCGGCTG (SEQ ID NO: 469) C	493
CR13-2_S:	cacc	G GCTGGGTGTCCCATTGAAA (SEQ ID NO: 470)	494
CR13-2_AS:	AAAC	TTTCAATGGGACACCCAGC (SEQ ID NO: 471) C	495
CR13-3_S:	cacc	G GTTTATTCAGCCGGGAGTC (SEQ ID NO: 472)	496
CR13-3_AS:	AAAC	GACTCCCGGCTGAATAAAC (SEQ ID NO: 473) C	497
CR13-4_S:	cacc	G TGGAGAGTTTGCAAGGAGC (SEQ ID NO: 474)	498
CR13-4_AS:	AAAC	GCTCCTTGCAAACTCTCCA (SEQ ID NO: 475) C	499
CR13-5_S:	cacc	G CCCTCCAGACTTTCCACCT (SEQ ID NO: 476)	500
CR13-5_AS:	AAAC	AGGTGGAAAGTCTGGAGGG (SEQ ID NO: 477) C	501
CR13-6_S:	cacc	G AATTTTCTTCCAAGTTCTC (SEQ ID NO: 478)	502
CR13-6_AS:	AAAC	GAGAACTTGGAAGAAAATT (SEQ ID NO: 479) C	503
CR13-7_S:	cacc	G CTGCGGAGAGAAGAAAGGG (SEQ ID NO: 480)	504
CR13-7_AS:	AAAC	CCCTTTCTTCTCTCCGCAG (SEQ ID NO: 481) C	505
CR13-8_S:	cacc	G AGAGCCACCCCCTGGCTCC (SEQ ID NO: 482)	506
CR13-8_AS:	AAAC	GGAGCCAGGGGGTGGCTCT (SEQ ID NO: 483) C	507
CR13-9_S:	cacc	G cgaagccaaccgcggcggg (SEQ ID NO: 484)	508
CR13-9_AS:	AAAC	cccgccgcggttggcttcg (SEQ ID NO: 485) C	509
CR13-10_S:	cacc	G agagggaagacgatcgccc (SEQ ID NO: 486)	510
CR13-10_AS:	AAAC	gggcgatcgtcttccctct (SEQ ID NO: 487) C	511
CR13-11_S:	cacc	G ccccttaactttcctccg (SEQ ID NO: 488)	512
CR13-11_AS:	AAAC	cggaggaaagttaaagggg (SEQ ID NO: 489) C	513
CR13-12_S:	cacc	G gcagccccgcttccttcaa (SEQ ID NO: 490)	514
CR13-12_AS:	AAAC	ttgaaggaagcggggctgc (SEQ ID NO: 491) C	515

TABLE 11

gRNAs for mouse BRN2 (CR16):

Oligo (5' to 3')	5' overhang	BRN2 Target Sequence	SEQ ID NO
CR16-1_S:	cacc	G CGAGAGCGAGAGGAGGGAG (SEQ ID NO: 516)	540
CR16-1_AS:	AAAC	CTCCCTCCTCTCGCTCTCG (SEQ ID NO: 517) C	541
CR16-2_S:	cacc	G GAGAGAGCTTGAGAGCGCG (SEQ ID NO: 518)	542
CR16-2_AS:	AAAC	CGCGCTCTCAAGCTCTCTC (SEQ ID NO: 519) C	543
CR16-3_S:	cacc	G GGTGGAGGGGGCGGGGCCC (SEQ ID NO: 520)	544
CR16-3_AS:	AAAC	GGGCCCCGCCCCCTCCACC (SEQ ID NO: 521) C	545

TABLE 11-continued

qRNAs for mouse BRN2 (CR16):

Oligo (5' to 3')	5' overhang		BRN2 Target Sequence		SEQ ID NO
CR16-4_S:	cacc	G	GGTATCCACGTAAATCAAA	(SEQ ID NO: 522)	546
CR16-4_AS:	AAAC		TTTGATTTACGTGGATACC	(SEQ ID NO: 523) C	547
CR16-5_S:	cacc	G	CCAATCACTGGCTCCGGTC	(SEQ ID NO: 524)	548
CR16-5_AS:	AAAC		GACCGGAGCCAGTGATTGG	(SEQ ID NO: 525) C	549
CR16-6_S:	cacc	G	GGCGCCCGAGGGAAGAAGA	(SEQ ID NO: 526)	550
CR16-6_AS:	AAAC		TCTTCTTCCCTCGGGCGCC	(SEQ ID NO: 527) C	551
CR16-7_S:	cacc	G	GGGTGGGGGTACCAGAGGA	(SEQ ID NO: 528)	552
CR16-7_AS:	AAAC		TCCTCTGGTACCCCCACCC	(SEQ ID NO: 529) C	553
CR16-8_S:	cacc	G	CCGGGGACAGAAGAGAGGG	(SEQ ID NO: 530)	554
CR16-8_AS:	AAAC		CCCTCTCTTCTGTCCCCGG	(SEQ ID NO: 531) C	555
CR16-9_S:	cacc	G	gagagagagtgggagaagc	(SEQ ID NO: 532)	556
CR16-9_AS:	AAAC		gcttctcccactctctctc	(SEQ ID NO: 533) C	557
CR16-10_S:	cacc	G	aaagtaactgtcaaatgcg	(SEQ ID NO: 534)	558
CR16-10_AS:	AAAC		cgcatttgacagttacttt	(SEQ ID NO: 535) C	559
CR16-11_S:	cacc	G	ttaaccagagcgcccagtc	(SEQ ID NO: 536)	560
CR16-11_AS:	AAAC		gactgggcgctctggttaa	(SEQ ID NO: 537) C	561
CR16-12_S:	cacc	G	cgtcggagctgcccgctag	(SEQ ID NO: 538)	562
CR16-12_AS:	AAAC		ctagcgggcagctccgacg	(SEQ ID NO: 539) C	563

[0361] qRT-PCR & IF: Activation of endogenous ASCL1 was assessed by qRT-PCR and immunofluorescence in MEFs on day 3 following delivery of either dCas9-VP64 and gRNAs, ASCL1 cDNA, or a negative control vector encoding luciferase. The generation of iNs was evaluated by TUJ1 and MAP2 co-staining and identification of cells with neuronal morphology and extended processes.

[0362] Live Cell Reporters: After 7-8 days in N3 medium, MEFs cultured on poly-D-lysine/laminin-coated coverslips were transduced with viruses carrying hSyn-RFP and MAP2-GCamP5 reporters to identify the most mature iNs for functional characterization via calcium imaging and electrophysiology (FIG. **49**B).

[0363] Results. dCas9-VP64 and gRNAs targeted to the ASCL1 promoter activated the endogenous gene in MEFs. Co-delivery of 8 gRNAs activated the endogenous gene 400-fold, a significant increase ($p<0.05$) over the 100-fold activation induced by the co-delivery of 4 gRNAs (FIG. **50**A). Nuclear-localized Ascl1 protein was detected by immunofluorescence in MEFs. Ectopic Ascl1 expression produced more Ascl1 protein than dCas9-VP64 with either gRNA cocktail, but did not activate the endogenous locus by day 3 (FIGS. **50**A-**50**B). TUJ1 and MAP2 co-positive cells with extended processes were identified by immunofluorescence after 13 days in neurogenic medium following delivery of dCas9-VP64 and gRNAs targeting the ASCL1 and BRN2 promoters (FIG. **51**A first row). A similar number of TUJ1 and MAP2 co-positive cells were identified with ectopic expression of the BAM factors (FIG. **51**A second row). Cells with a neuronal morphology expressing the hSyn-RFP reporter were visible in culture as early as day 11 in neurogenic medium (FIG. **51**B). A cell expressing the MAP2-GCaMP5 calcium indicator exhibited KCl-induced depolarization detected with a fluorescent microscope (FIGS. **52**A-**52**B).

[0364] The direct conversion of mouse embryonic fibroblasts to TUJ1 and MAP2 co-positive cells with a neuronal morphology was accomplished through activation of endogenous BRN2 and ASCL1 by CRISPR/Cas9-based transcription factors. Though dCas9-VP64 produces less protein than ectopic expression of ASCL1 (FIG. **50**B), the generation of neuronal-like cells is similar. The activation of the endogenous loci may induce a reprogramming cascade of events that is not mechanistically identical to that generated with ectopic expression.

[0365] dCas9-VP64 was able to penetrate heterochromatin and activated stably silenced endogenous genes, a characteristic of only a subset of "pioneer" transcription factors. As a result, converting cell lineage with CRISPR/Cas9-transcription factors may better overcome epigenetic barriers to reprogramming than ectopic expression of transcription factors, particularly in hard-to-reprogram cell-types, such as adult human cells. This may have clinical importance in the

field of regenerative medicine, as it is often desired to use autologous sources in cell replacement therapies.

Example 17

Multiplex CRISPR/Cas9-Based Genome Engineering—Materials and Methods

[0366] Plasmid constructs. The expression cassettes for the *S. pyogenes* sgRNA and human codon optimized Cas9 (hCas9) nuclease were used, as described above. Additional promoters for mU6 (Ohshima et al., *Nucleic Acids Res* 9:5145-5158 (1981)), H1 (Myslinski et al., *Nucleic Acids Res* 29:2502-2509 (2001)), and 7SK (Murphy et al., Cell 51:81-87 (1987)) pol-III promoters were synthesized using GeneBlocks (IDT) and cloned in place of the hU6 sgRNA expression cassette. A GeneBlock (IDT) was cloned onto the 3' end of the Cas9 coding sequence to fuse a T2A skipping peptide and eGFP gene immediately after Cas9 to monitor vector expression. The coding region for hCas9-T2A-GFP (SEQ ID NO: 145) was then transferred into a lentiviral expression vector containing the human ubiquitin C (hUbC) promoter to drive expression of hCas9-T2A-GFP, as well as restriction sites to facilitate Golden Gate cloning of sgRNA expression cassettes immediately upstream of the hUbC promoter (FIG. **42**A).

[0367] Protocol for assembly of custom lentiviral vectors. Assembly of custom lentiviral vectors expressing up to four sgRNAs of choice and active Cas9, dCas9, or dCas9-VP64 was accomplished in less than five days. The cloning method used the Golden Gate cloning and type IIS restriction enzymes that cleave outside their recognition sequence to create unique overhangs. Golden Gate assembly expedited cloning as all four expression cassettes were ligated into the final lentiviral vector in one step. The lentiviral vector expressed active Cas9, cCas9, or dCas9-VP64 in addition to one, two, three, or four sgRNAs expressed from independent promoters.

[0368] Step 1: Single stranded oligos containing each 20 bp protospacer were annealed in such a fashion to create sticky ends and were ligated into the desired pZDonor-promoter vector. Order two single stranded oligos for each desired genomic target. To anneal the complimentary oligos, mix 8 μL sense oligo+8 μL antisense oligo (both at 10 mM)+2 μL 10× ligase buffer. The oligos are melted and reannealed in a PCR machine with the program: 96° C. for 300 seconds, followed by 85° C. for 20 seconds, 75° C. for 20 seconds, 65° C. for 20 seconds, 55° C. for 20 seconds, 45° C. for 20 seconds, 35° C. for 20 seconds, and 25° C. for 20 seconds with a −0.3° C./second rate between steps. To phosphorylate the sticky ends, add 1 μL 25 mM ATP+1 microliter T4 Polynucleotide Kinase (NEB) and incubate at 37° C. for 60 minutes followed by 65° C. for 20 minutes to heat inactivate the enzyme. Each protospacer was ligated into the desired expression vector using T4 DNA ligase (NEB) incubated at 16° C. for 60 minutes using 50 ng of vector and 1 μL of annealed oligonucleotides in a 10 μL reaction volume according to manufacturer's instructions. Five microliters of each ligation was transformed into XL1 blue chemically competent bacteria (Agilent) following the manufacturer's instructions. Plate transformation onto LB agar plates containing 50 μg/mL kanamycin (Sigma) and incubate overnight at 37° C. In our experience, >90% of the colonies will contain the desired ligation product. Sequencing using the M13 reverse standard sequencing primer was performed to validate each final sgRNA construct prior to moving onto step 2.

[0369] Step 2: Construct the four promoter-gRNA cassettes into a lentiviral destination vector using Golden Gate assembly. After completion of step 1, there are four independent plasmids each expressing a different sgRNA from a different promoter. To assemble the four different promoter-sgRNA constructs into the desired destination vector, combine 200 ng of each sgRNA expression plasmid and desired lentiviral destination vector with 1 μL of T4 DNA ligase (NEB), 1 μL BsmBI FastDigest (Fisher Scientific), and 2 μL 10× T4 ligase buffer (NEB) in a 20 μL reaction volume. Incubate the reaction as follows: 37° C. for 10 minutes, 16° C. for 15 minutes, 37° C. for 30 minutes, 80° C. for 5 minutes. Transform 5 μL of ligation reaction into SURE 2 chemically competent cells (Agilent) following the manufacturer's instructions. Plate transformations onto LB agar plates containing 100 μg/mL ampicillin and incubate overnight at 37° C. Optionally, colonies can be screened by lacZ-based blue/white screening using IPTG and X-gal; however, in our experience, >90% of the transformants contain the proper ligation product. Due to the inverted repeats formed by the opposing sgRNA expression cassettes, the final constructs may be unstable and thus we recommend maintaining these plasmids in the SURE 2 cell line and screening the final plasmid with a test PCR using the sense primer 5'-TCGGGTTTATTACAGGGACAGCAG-3' (SEQ ID NO:464) and antisense primer 5'-TCTAAGGCCGAGTCTTATGAGCAG-3' (SEQ ID NO:465). These primers amplify across the four promoter-gRNA region. Due to the repetitive nature, a distinct banding pattern should be observed with the largest product approximately 1800 bp in size.

[0370] Cell culture and transfection. HEK293T cells were obtained from the American Tissue Collection Center (ATCC, Manassas, Va.) through the Duke University Cancer Center Facilities and were maintained in DMEM supplemented with 10% FBS and 1% penicillin streptomycin. Primary human dermal fibroblasts (Catalog ID: GM03348) were obtained from Coriell Institute (Camden, N.J.) and were maintained in DMEM supplemented with 10% FBS and 1% penicillin streptomycin. All cells were cultured at 37° C. with 5% CO_2. HEK293T cells were transfected with Lipofectamine 2000 (Life Technologies) with 200 ng of each sgRNA expression vector (800 ng total pDNA) according to the manufacturer's protocol in 24 well plates.

[0371] Viral production and transduction. All lentiviral vectors used is this study are second generation and were produced using standard viral production methods. Briefly, 3.5 million HEK293 Ts were plated per 10 cm dish. The following day, cells were transfected by the calcium phosphate transfection method with 20 μg of transfer vector, 6 μg of pMD2G, and 10 μg psPAX2. The media was changed 12-14 hrs post transfection. The viral supernatant was collected 24 and 48 hrs after this media change, passed through a 0.45 micron filter and pooled. For transduction, the cell medium was replaced with viral supernatant supplemented with 4 μg/mL polybrene. The viral supernatant was exchanged for fresh medium 12-24 hrs later.

[0372] Reverse Transcription PCR. RNA was isolated using the miRNeasy Mini RNA isolation kit (Qiagen). DNAse digestion was performed using the DNA-free Kit (Applied Biosystems). cDNA synthesis was performed

using the SuperScript VILO cDNA Synthesis Kit (Invitrogen). cDNA was PCR amplified using Taq DNA polymerase (NEB) and the resulting product was run on TAE agarose gels. Images were captured using a ChemiDoc XRS+ System and processed using ImageLab software (Bio-Rad).

[0373] Quantitative Real Time PCR. RNA was isolated using the RNeasy Plus RNA isolation kit (Qiagen). cDNA synthesis was performed using the SuperScript VILO cDNA Synthesis Kit (Invitrogen). Real-time PCR using PerfeCTa SYBR Green FastMix (Quanta Biosciences) was performed with the CFX96 Real-Time PCR Detection System (Bio-Rad). Primer specificity was confirmed by agarose gel electrophoresis and melting curve analysis. Reaction efficiencies over the appropriate dynamic range were calculated to ensure linearity of the standard curve. The results are expressed as fold-increase mRNA expression of the gene of interest normalized to β-Actin expression using the ΔΔCt method. Reported values are the mean and S.E.M. from two independent experiments (n=2) where technical replicates were averaged for each experiment.

[0374] Western Blot. Cells were lysed in RIPA Buffer (Sigma) supplemented with protease inhibitor cocktail (Sigma). Protein concentration was measured using BCA protein assay reagent (Thermo Scientific) and BioTek Synergy 2 Multi-Mode Microplate Reader. Lysates were mixed with loading buffer and boiled for 5 min; 25 g of protein were run in NuPage 10% Bis-Tris Gel polyacrylamide gels (Bio-Rad) and transferred to nitrocellulose membranes. Nonspecific antibody binding was blocked with TBST (50 mM Tris, 150 mM NaCl and 0.1% Tween-20) with 5% nonfat milk for 1 hr at room temperature. The membranes were incubated with primary antibodies: (anti-MyoD (1:250, Santa Cruz Sc-32758) in 5% BSA in TBST overnight at 4° C.; anti-Myogenin (1:250, Santa Cruz Sc-12732) in 5% BSA overnight at 4° C.; anti-FLAG-HRP (1:1000, Cell Signaling 2044) in 5% milk in TBST for 60 min at room temperature; anti-GAPDH (1:5000, Cell Signaling, clone 14C10) in 5% milk in TBST for 30 min at room temperature. Membranes were then washed three times with TBST for 15 minutes total. Membranes were incubated with anti-rabbit HRP-conjugated antibody (Sigma, A 6154) or anti-mouse HRP-conjugated antibody (Santa Cruz, SC-2005) diluted 1:5,000 for 30 min and washed with TBST three times for 15 min each. Membranes were visualized using the ImmunStar WesternC Chemiluminescence Kit (Bio-Rad) and images were captured using a ChemiDoc XRS+ System and processed using ImageLab software (Bio-Rad).

[0375] Cel-I quantification of endogenous gene modification. CRISPR/Cas9 nuclease lesions at the endogenous target site were quantified using the Surveyor nuclease assay, which can detect mutations characteristic of nuclease-mediated NHEJ. After transfection or transduction, cells were incubated for 3-10 days at 37° C. and genomic DNA was extracted using the DNeasy Blood and Tissue kit (Qiagen). The target locus was amplified by 35 cycles of PCR with the AccuPrime High Fidelity PCR kit (Invitrogen). The resulting PCR products were randomly melted and reannealed in a PCR machine with the program: 95° C. for 240 seconds, followed by 85° C. for 60 seconds, 75° C. for 60 seconds, 65° C. for 60 seconds, 55° C. for 60 seconds, 45° C. for 60 seconds, 35° C. for 60 seconds, and 25° C. for 60 seconds with a −0.3° C./second rate between steps. Following re-annealing, 8 μL of PCR product was mixed with 1 μL of Surveyor Nuclease S and 1 μL of Enhancer S (Transgenomic, Omaha, Nebr.) and incubated at 42° C. for 1 hr. After incubation, 6 μL of digestion product was loaded onto a 10% TBE polyacrylamide gel and run at 200 V for 30 minutes. The gels were stained with ethidium bromide and quantified using ImageLab (Bio-Rad) by densitometry (Perez-Pinera et al., *Nucleic Acids Res* 40:3741-3751 (2012)).

[0376] Statistical Analysis. At least two independent experiments were compiled as means and standard errors of the mean. Effects were evaluated with multivariate ANOVA and Dunnett's post hoc test using JMP 10 Pro.

Example 18

Development of a Single Lentiviral Vector for Multiplex CRISPR/Cas9 Applications

[0377] A limitation of current CRISPR/Cas9 gene editing systems, particularly transactivator systems, is the simultaneous and efficient delivery of multiple sgRNAs and Cas9 protein for multiplex gene editing and synergistic gene activation applications, especially in difficult to transfect cell types. To overcome this limitation, we developed a single lentiviral vector that efficiently expresses Cas9 and up to four sgRNAs. In order to maximize the expression efficiency of each sgRNA, this vector expresses four sgRNAs from four independent pol III promoters (human U6 promoter, mouse U6 promoter, 7SK, and H1). We validated sgRNA expression from each promoter using end-point RT-PCR to detect a sgRNA targeting the AAVS1 locus (FIG. **42**A). To test the activity of each sgRNA expression construct, we co-transfected each promoter construct expressing an sgRNA targeting AAVS1 independently with an active Cas9 expression construct into human HEK293T cells. Notably, we detected consistent and high levels of gene modification at the target locus for each sgRNA that are comparable to a well-characterized zinc-finger nuclease with high activity at the AAVS1 locus (FIG. **42**B). Furthermore, lentiviral delivery of different Cas9-based constructs, including an active Cas9 nuclease, dead Cas9, and dead Cas9 fused to the VP64 transactivator domain, resulted in expression of full-length Cas9 protein in HEK293T cells as determined by western blot (FIG. **42**C).

[0378] Using these components, we developed a Golden Gate cloning method to facilitate rapid and efficient cloning of multiple sgRNA expression cassettes into a single lentiviral vector expressing the desired Cas9 effector (FIG. **43**). In the first step, oligonucleotides encoding sgRNA protospacer sequences are cloned independently into different expression vectors, each with a distinct promoter driving sgRNA expression. In the second step, each sgRNA expression construct is subcloned into a lentiviral Cas9 expression vector of choice by Golden Gate assembly. This strategy allows for robust and rapid cloning of up to four sgRNAs into a single lentiviral vector for gene editing or activation applications. To express less than four sgRNAs, a PolyT terminator sequence is cloned down-stream of unused promoters to prevent transcription from the unused promoters. Each vector co-expresses the choice Cas9 with eGFP via a 2A skipping peptide to enable fluorescence-activated flow sorting and enrichment of cells with a high multiplicity of infection. Finally, the entire region containing the sgRNA and Cas9 expression cassettes is flanked by loxP sites to mediate removal by Cre-lox excision.

Example 19

Validation of a Single Lentiviral sgRNA/Cas9 Expression Vector for Multiplex Genome Engineering

[0379] To validate the independent activity of each sgRNA, we assembled a single lentiviral vector expressing active Cas9 and four sgRNAs, each targeting an independent loci (FIG. **44**A). As control vectors, we assembled constructs expressing only one sgRNA along with polyT protospacers in the other three positions. We transduced HEK293 Ts and primary fibroblasts with lentiviral vectors expressing the indicated sgRNAs and monitored gene modification frequencies at 7 or 10 days post-transduction, respectively (FIG. **44**B). In both cell types, the single lentiviral vector mediated highly efficient multiplex editing at all four loci (FIG. **44**B). Interestingly, expression of all four sgRNAs together resulted in higher modification frequencies than a single sgRNA alone at 3 out of 4 loci in fibroblasts (FIG. **44**B). We observed efficient multiplex gene editing in fibroblasts, which are conventionally a difficult to transfect cell type. These data demonstrate that a single lentivirus can express four active sgRNAs efficiently and that this lentiviral platform can be used to target four distinct loci for multiplex CRISPR/Cas9 gene editing.

Example 20

Transient RNA-Guided Gene Activation in Cell Lines Stably Expressing a Lentiviral Cas9-Based Transactivator

[0380] Next, we were interested in developing a system that enables transient gene activation by transfecting sgRNAs into model cell lines stably expressing Cas9. HEK293 Ts were transduced with different Cas9-T2A-GFP and GFP expression was monitored using flow cytometry. Following normal passaging every 2-3 days, each cell line exhibited stable GFP expression for up to 35 days post transduction. Transduced HEK293 Ts were then transfected with one to four separate sgRNA expression constructs targeting either the IL1RN or HBG1 promoter. Transient transfection of these sgRNA constructs in stable dCas9-VP64 expressing cells lines resulted in tunable endogenous gene activation (FIGS. **45**A-**45**B). Gene activation following transient transfection of sgRNA constructs in cells expressing dCas9-VP64 reached a maximum level of activation approximately 3-6 post-transfection and fell to undetectable levels by 20 days post-transfection (FIGS. **45**C-**45**D). Furthermore, we were able to re-activate each gene by a second transfection of all four sgRNA constructs targeting each promoter, although activation levels were significantly lower than observed from the first transfection (FIGS. **45**C-**45**D). This reduction in activity after the second transfection may be due to reduced vector expression or competitive growth of untransduced cells. Despite this, these data demonstrate that lentiviral Cas9 combined with transient sgRNA delivery can be used as a versatile system to tunably and transiently activate and re-activate target genes in a Cas9 stably tranduced cell line.

Example 21

Stable Gene Activation in HEK293T Cells Using a Single Lentiviral sgRNA/Cas9 Transactivator Expression Vector

[0381] Lentiviral delivery may enable stable, long-term gene activation by CRISPR/Cas9 transactivation. To test this, HEK293 Ts were transduced using a single lentiviral vector encoding dCas9-VP64 and one to four sgRNA expression cassettes. Similar to our transient transfection results (FIGS. **45**A-**45**D), we were able to tunably and robustly activate expression of endogenous IL1RN and HBG1 genes (FIGS. **46**A-**46**B). Gene activation induced by co-transfection of HEK293T cells with dCas9-VP64 and four sgRNAs targeted to the IL1RN and HBG1 promoters peaked three-five days post-transfection and gene expression returned to background levels 15-20 days post-transfection (FIGS. **46**A-**46**B). In contrast, lentiviral delivery of dCas9-VP64 and the same four IL1RN or HBG1-targeted sgRNAs induced sustained gene activation for more than 20 days post-transduction (FIGS. **46**C-**46**D). Thus, single lentiviral delivery of multiplex dCas9-VP64 transactivators is a useful platform to efficiently and stably upregulate target endogenous genes.

Example 22

dCas9-KRAB—Targeting the HS2 Enhancer

[0382] The HS2 enhancer is a well-characterized distal regulatory element necessary for activation of the globin gene locus. dCas9-KRAB with gRNAs targeted to the HS2 enhancer were delivered to determine if this system would repress γ-, ε-, and β-globin expression in the K562 human erythroid leukemia cell line (FIG. **54**). A panel of gRNAs was created targeting different sites along the core region of the HS2 enhancer SEQ ID NO: 467). See Table 12.

TABLE 12

HS2 gRNA Target Sequences

Cr#	Protospacer	Complete gRNA Sequence
1	gagacacacagaaatgtaac (SEQ ID NO: 564)	gagacacacagaaatgtaacgtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 585)
2	ggtggggcactgaccccgac (SEQ ID NO: 565)	ggtggggcactgaccccgacgtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 586)
3	ctagagtgatgactcctatc (SEQ ID NO: 566)	ctagagtgatgactcctatcgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 587)

TABLE 12-continued

HS2 gRNA Target Sequences

Cr#	Protospacer	Complete gRNA Sequence
4	gactaaaactccacctcaaa (SEQ ID NO: 567)	gactaaaactccacctcaaagtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 588)
5	aatatgtcacattctgtctc (SEQ ID NO: 568)	aatatgtcacattctgtctcgtttTAGAGCTAGAAATAGCAAGTTAAAATAAG GCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGTG CTTTTTTTCC (SEQ ID NO: 589)
6	ggactatgggaggtcactaa (SEQ ID NO: 569)	ggactatgggaggtcactaagttttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 590)
7	gctcatgcttggactatggg (SEQ ID NO: 570)	gctcatgcttggactatggggtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 591)
8	gttctggccaggcccctgtc (SEQ ID NO: 571)	gttctggccaggcccctgtcgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 592)
9	agtgccccacccccgccttc (SEQ ID NO: 572)	agtgccccacccccgccttcgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 593)
10	gtggggcactgaccccgaca (SEQ ID NO: 573)	gtggggcactgaccccgacagtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 594)
11	aaccttctaagcaaaccttc (SEQ ID NO: 574)	aaccttctaagcaaaccttcgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 595)
12	gttacacagaaccagaaggc (SEQ ID NO: 575)	gttacacagaaccagaaggcgtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 596)
13	gaaggttacacagaaccaga (SEQ ID NO: 576)	gaaggttacacagaaccagagtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 597)
14	agtcatgatgagtcatgctg (SEQ ID NO: 577)	agtcatgatgagtcatgctggtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 598)
15	gatgagtcatgctgaggctt (SEQ ID NO: 578)	gatgagtcatgctgaggcttgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 599)
16	actctaggctgagaacatct (SEQ ID NO: 579)	actctaggctgagaacatctgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 600)
17	gtccccagcaggatgcttac (SEQ ID NO: 580)	gtccccagcaggatgcttacgtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 601)
18	gccctgtaagcatcctgctg (SEQ ID NO: 581)	gccctgtaagcatcctgctggtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 602)
19	cagggcagatggcaaaaaaa (SEQ ID NO: 582)	cagggcagatggcaaaaaaagtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 603)
20	gaggtggagttttagtcagg (SEQ ID NO: 583)	gaggtggagttttagtcagggtttTAGAGCTAGAAATAGCAAGTTAAAATAA GGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGGT GCTTTTTTTCC (SEQ ID NO: 604)
21	aaacggcatcataaagaaaa (SEQ ID NO: 584)	aaacggcatcataaagaaaagtttTAGAGCTAGAAATAGCAAGTTAAAATA AGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCGG TGCTTTTTTTCC (SEQ ID NO: 605)

[0383] Screening single gRNAs at the globin locus by CRISPR/dCas9. dCas9 and dCas9-KRAB effectors were delivered lentivirally 5-8 days prior to electroporation with 5 μg of a plasmid encoding the U6-sgRNA expression (FIG. **55**A). Cells that were not electroporated with gRNA (no gRNA) and cells treated with a gRNA targeting a different locus (IL1RN) were included as controls. Multiple gRNAs effected potent repression of ε-, γ-, and β-globin genes when assayed 3 days post-transfection, with up to 80% knockdown achieved (FIGS. **55**B, **55**C, **55**D). Expressing a gRNA with either dCas9 or dCas9-KRAB inhibited gene expression at the globin locus. Generally, treatment with dCas9-KRAB resulted in stronger repression for a given gRNA compared to dCas9 alone, suggesting an important role for the KRAB domain in recruiting heterochromatin factors that enhance repression. The repression levels achieved are dependent on the amount of gRNA plasmid delivered by transfection only in the dCas9-KRAB treated cells (FIGS. **56**A, **56**B, **56**C). Increasing the dose of Cr4 gRNA plasmid up to 10 μg increases silencing levels of the globin genes in dCas9-KRAB-treated cells.

[0384] Stable silencing of globin genes by dCas9-KRAB. dCas9/dCas9-KRAB were co-expressed with single gRNAs lentivirally in K562s (FIG. **57**A). Cells that were not treated with lentivirus (NT), treated with dCas9/dCas9-KRAB without a gRNA (no gRNA), and with dCas9/dCas9-KRAB and gRNA targeting a different locus (IL1RN) were included as controls. Cells treated with lentivirus were selected from days 4 to 7. Multiple gRNAs effected potent transcriptional repression of ε-, γ-, and β-globin genes when assayed 7 days after transduction, with up to 95% knockdown achieved (FIGS. **57**B, **57**C, **57**D). Expression of β-globin was silenced the most in response gRNAs targeted to the HS2 enhancer. Treatment with dCas9-KRAB with a gRNA resulted in dramatically more repression than treatment with dCas9 and gRNA.

[0385] These findings demonstrate that dCas9-KRAB targeted to the HS2 enhancer by gRNAs effects potent repression of the distal globin genes. This is the first example of targeted epigenetic control of distal regulatory elements in mammalian cells by the CRISPR/Cas9 system. Enhancers regulate development and disease, and this disclosure provides a method to probe and control enhancer function and may be used to determine the effects of dCas9-KRAB on local chromatin accessibility and genome-wide expression.

Example 23

dCas9-p300

[0386] A dCas9-p300 fusion protein was designed and compared to dCas9-VP64 fusion protein (see FIG. **59**). The amino acid constructs of dCas9 constructions are shown in FIGS. **61**A-**61**C. Cells from the Human Embryonic Kidney tissue culture line HEK293T (ATCC; CRL-11268) were seeded at a density of 1.5e5 cells per well in 24-well tissue culture dishes one day prior to transfection with Lipofectamine 2000 transfection reagent (Life Technologies). 24 hrs later cells were transfected with 1 μL Lipofectamine 2000, 375 ng dCas9 expression construct (dCas9, dCas9VP64, or dCas9p300 respectively), and 125 ng of pooled gRNA expression plasmids (4 each at equimolar ratios). Table 13 shows the gRNA information.

TABLE 13

gRNA information

Target Location	Protospacer Sequence (5'-3')	Genomic Location (GRCh38 Primary Assembly)	Reference
IL1RN Promoter	TGTACTCTCTGAGGTGCTC (SEQ ID NO: 606)	Chr2: 113117865-113117883	Perez-Pinera et al., Nat Methods, 2013
IL1RN Promoter	ACGCAGATAAGAACCAGTT (SEQ ID NO: 607)	Chr2: 113117714-113117732	Perez-Pinera et al., Nat Methods, 2013
IL1RN Promoter	CATCAAGTCAGCCATCAGC (SEQ ID NO: 608)	Chr2: 113117781-113117799	Perez-Pinera et al., Nat Methods, 2013
IL1RN Promoter	GAGTCACCCTCCTGGAAAC (SEQ ID NO: 609)	Chr2: 113117749-113117767	Perez-Pinera et al., Nat Methods, 2013
MyoD Promoter	CCTGGGCTCCGGGGCGTTT (SEQ ID NO: 610)	Chr11: 17719509-17719527	Perez-Pinera et al., Nat Methods, 2013
MyoD Promoter	GGCCCCTGCGGCCACCCCG (SEQ ID NO: 611)	Chr11: 17719422-17719440	Perez-Pinera et al., Nat Methods, 2013
MyoD Promoter	CTCCCTCCCTGCCCGGTAG (SEQ ID NO: 612)	Chr11: 17719350-17719368	Perez-Pinera et al., Nat Methods, 2013

TABLE 13-continued

gRNA information			
Target Location	Protospacer Sequence (5'-3')	Genomic Location (GRCh38 Primary Assembly)	Reference
MyoD Promoter	AGGTTTGGAAAGGGCGTGC (SEQ ID NO: 613)	Chr11: 17719290-17719308	Perez-Pinera et al., Nat Methods, 2013
Oct4 Promoter	ACTCCACTGCACTCCAGTCT (SEQ ID NO: 614)	Chr6: 31170953-31170934	Hu et al., Nucleic Acids Res, 2014
Oct4 Promoter	TCTGTGGGGGACCTGCACTG (SEQ ID NO: 615)	Chr6: 31170885-31170866	Hu et al., Nucleic Acids Res, 2014
Oct4 Promoter	GGGGCGCCAGTTGTGTCTCC (SEQ ID NO: 616)	Chr6: 31170855-31170836	Hu et al., Nucleic Acids Res, 2014
Oct4 Promoter	ACACCATTGCCACCACCATT (SEQ ID NO: 617)	Chr6: 31170816-31170797	Hu et al., Nucleic Acids Res, 2014

[0387] The 3 days post-transfection cells were harvested and assayed by RT-QPCR for mRNA expression. The RT-QPCR primer sequences are listed in Table 14.

TABLE 14

RT-QPCR Primers		
Primer Target	Primer Sequence (5'-3')	Reference
GAPDH Forward	CAATGACCCCTTCATTGACC (SEQ ID NO: 618)	Perez-Pinera et al., Nat Methods, 2013
GAPDH Reverse	TTGATTTTGGAGGGATCTCG (SEQ ID NO: 619)	Perez-Pinera et al., Nat Methods, 2013
IL1RN Forward	GGAATCCATGGAGGGAAGAT (SEQ ID NO: 620)	Perez-Pinera et al., Nat Methods, 2013
IL1RN Reverse	TGTTCTCGCTCAGGTCAGTG (SEQ ID NO: 621)	Perez-Pinera et al., Nat Methods, 2013
MyoD Forward	CTCTCTGCTCCTTTGCCACA (SEQ ID NO: 622)	Perez-Pinera et al., Nat Methods, 2013
MyoD Reverse	GTGCTCTTCGGGTTTCAGGA (SEQ ID NO: 623)	Perez-Pinera et al., Nat Methods, 2013
Oct4 Forward	CGAAAGAGAAAGCGAACCAGTAT CGAGAAC (SEQ ID NO: 624)	Hu et al., Nucleic Acids Res, 2014
Oct4 Reverse	CGTTGTGCATAGTCGCTGCTTGA TCGC (SEQ ID NO: 625)	Hu et al., Nucleic Acids Res, 2014

[0388] RT-QPCR was normalized to GAPDH expression using the $\Delta\Delta C_t$ method. Results are expressed as fold-increase expression of the gene of interest relative to cells treated with Lipofectamine only without DNA transfected ("No DNA") (See FIGS. **60**A-**60**C).

[0389] FIG. **62** shows that mutating residues in the p300 HAT domain causes a loss of its ability to activate gene expression. FIGS. **63**A-**63**B shows that multiple gRNAs work synergistically with dCas-9-p300, as showed with dCas-9-VP64.

Example 24

[0390] FIGS. **64**A-**64**C shows TALEN mediated integration of minidystrophin at the 5'UTR of the Dp427m skeletal muscle isoform of dystrophin in skeletal myoblast cell lines derived from human DMD patients carrying different deletions in the dystrophin gene. DMD patient cells were electroporated with constructs encoding a TALEN pair active at the 5'UTR locus and a donor template carrying the minidystrophin gene. FIG. **64**A shows a schematic of how minidystrophin was integrated into the 5'UTR. FIG. **64**B shows that hygromycin-resistant clonal cell lines were isolated and screened by PCR for successful site-specific integrations at the 5'UTR using the primers shown in FIG. **64**A. Asterisks indicate clones selected for further analysis in FIG. **64**C. FIG. **64**C shows that clonally isolated DMD myoblasts with detected integration events were differentiated for 6 days and assessed for expression of an HA tag fused to the C terminus of minidystrophin.

[0391] It is understood that the foregoing detailed description and accompanying examples are merely illustrative and are not to be taken as limitations upon the scope of the invention, which is defined solely by the appended claims and their equivalents.

[0392] Various changes and modifications to the disclosed embodiments will be apparent to those skilled in the art. Such changes and modifications, including without limitation those relating to the chemical structures, substituents, derivatives, intermediates, syntheses, compositions, formulations, or methods of use of the invention, may be made without departing from the spirit and scope thereof.

APPENDIX

hCas9-T2A-GFP DNA sequence: SEQ ID NO: 145 (SpCas9 human optimized sequence, [illegible], T2A peptide, eGFP sequence)

gccaccATGGACAAGAAGTACTCCATTGGGCTCGATATCGGCACAAACAGCGTCGGCTGGGCCGTCATTACGGACGAGTA
CAAGGTGCCGAGCAAAAAATTCAAAGTTCTGGGCAATACCGATCGCCACAGCATAAAGAAGAACCTCATTGGCGCCCTCC
TGTTCGACTCCGGGGAGACGGCCGAAGCCACGCGGCTCAAAAGAACAGCACGGCGCAGATATACCCGCAGAAAGAATCGG
ATCTGCTACCTGCAGGAGATCTTTAGTAATGAGATGGCTAAGGTGGATGACTCTTTCTTCCATAGGCTGGAGGAGTCCTT
TTTGGTGGAGGAGGATAAAAAGCACGAGCGCCACCCAATCTTTGGCAATATCGTGGACGAGGTGGCGTACCATGAAAAGT
ACCCAACCATATATCATCTGAGGAAGAAGCTTGTAGACAGTACTGATAAGGCTGACTTGCGGTTGATCTATCTCGCGCTG
GCGCATATGATCAAATTTCGGGGACACTTCCTCATCGAGGGGGACCTGAACCCAGACAACAGCGATGTCGACAAACTCTT
TATCCAACTGGTTCAGACTTACAATCAGCTTTTCGAAGAGAACCCGATCAACGCATCCGGAGTTGACGCCAAAGCAATCC
TGAGCGCTAGGCTGTCCAAATCCCGGCGGCTCGAAAACCTCATCGCACAGCTCCCTGGGGAGAAGAAGAACGGCCTGTTT
GGTAATCTTATCGCCCTGTCACTCGGGCTGACCCCCAACTTTAAATCTAACTTCGACCTGGCCGAAGATGCCAAGCTTCA
ACTGAGCAAAGACACCTACGATGATGATCTCGACAATCTGCTGGCCCAGATCGGCGACCAGTACGCAGACCTTTTTTTGG
CGGCAAAGAACCTGTCAGACGCCATTCTGCTGAGTGATATTCTGCGAGTGAACACGGAGATCACCAAAGCTCCGCTGAGC
GCTAGTATGATCAAGCGCTATGATGAGCACCACCAAGACTTGACTTTGCTGAAGGCCCTTGTCAGACAGCAACTGCCTGA
GAAGTACAAGGAAATTTTCTTCGATCAGTCTAAAAATGGCTACGCCGGATACATTGACGGCGGAGCAAGCCAGGAGGAAT
TTTACAAATTTATTAAGCCCATCTTGGAAAAAATGGACGGCACCGAGGAGCTGCTGGTAAAGCTTAACAGAGAAGATCTG
TTGCGCAAACAGCGCACTTTCGACAATGGAAGCATCCCCCACCAGATTCACCTGGGCGAACTGCACGCTATCCTCAGGCG
GCAAGAGGATTTCTACCCCTTTTTGAAAGATAACAGGGAAAAGATTGAGAAAATCCTCACATTTCGGATACCCTACTATG
TAGGCCCCCTCGCCCGGGGAAATTCCAGATTCGCGTGGATGACTCGCAAATCAGAAGAGACCATCACTCCCTGGAACTTC
GAGGAAGTCGTGGATAAGGGGGCCTCTGCCCAGTCCTTCATCGAAAGGATGACTAACTTTGATAAAAATCTGCCTAACGA
AAAGGTGCTTCCTAAACACTCTCTGCTGTACGAGTACTTCACAGTTTATAACGAGCTCACCAAGGTCAAATACGTCACAG
AAGGGATGAGAAAGCCAGCATTCCTGTCTGGAGAGCAGAAGAAAGCTATCGTGGACCTCCTCTTCAAGACGAACCGGAAA
GTTACCGTGAAACAGCTCAAAGAAGACTATTTCAAAAAGATTGAATGTTTCGACTCTGTTGAAATCAGCGGAGTGGAGGA
TCGCTTCAACGCATCCCTGGGAACGTATCACGATCTCCTGAAAATCATTAAAGACAAGGACTTCCTGGACAATGAGGAGA
ACGAGGACATTCTTGAGGACATTGTCCTCACCCTTACGTTGTTTGAAGATAGGGAGATGATTGAAGAACGCTTGAAAACT
TACGCTCATCTCTTCGACGACAAAGTCATGAAACAGCTCAAGAGGCGCCGATATACAGGATGGGGGCGGCTGTCAAGAAA
ACTGATCAATGGGATCCGAGACAAGCAGAGTGGAAAGACAATCCTGGATTTTCTTAAGTCCGATGGATTTGCCAACCGGA
ACTTCATGCAGTTGATCCATGATGACTCTCTCACCTTTAAGGAGGACATCCAGAAAGCACAAGTTTCTGGCCAGGGGGAC
AGTCTTCACGAGCACATCGCTAATCTTGCAGGTAGCCCAGCTATCAAAAAGGGAATACTGCAGACCGTTAAGGTCGTGGA
TGAACTCGTCAAAGTAATGGGAAGGCATAAGCCCGAGAATATCGTTATCGAGATGGCCCGAGAAGAACCAAACTACCCAGA
AGGGACAGAAGAACAGTAGGGAAAGGATGAAGAGGATTGAAGAGGGTATAAAAGAACTGGGGTCCCAAATCCTTAAGGAA
CACCCAGTTGAAAACACCCAGCTTCAGAATGAGAAGCTCTACCTGTACTACCTGCAGAACGGCAGGGACATGTACGTGGA
TCAGGAACTGGACATCAATCGGCTCTCCGACTACGACGTGGATCATATCGTGCCCCAGTCTTTTCTCAAAGATGATTCTA
TTGATAATAAAGTGTTGACAAGATCCGATAAAAATAGAGGGAAGAGTGATAACGTCCCCTCAGAAGAAGTTGTCAAGAAA
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        515                 520                 525

Arg Met Thr Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro
    530                 535                 540

Lys His Ser Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr
545                 550                 555                 560

Lys Val Lys Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser
                565                 570                 575

Gly Glu Gln Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg
            580                 585                 590

Lys Val Thr Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu
        595                 600                 605

Cys Phe Asp Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala
    610                 615                 620

Ser Leu Gly Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp
625                 630                 635                 640

Phe Leu Asp Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu
                645                 650                 655

Thr Leu Thr Leu Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys
            660                 665                 670

Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg
        675                 680                 685

Arg Arg Tyr Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly
    690                 695                 700

Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser
705                 710                 715                 720

Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser
                725                 730                 735

Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly
            740                 745                 750

Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile
        755                 760                 765

Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys
    770                 775                 780
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Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg
785                 790                 795                 800

Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met
                805                 810                 815

Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys
            820                 825                 830

Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu
        835                 840                 845

Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp
    850                 855                 860

Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp Ala Ile Val Pro Gln Ser
865                 870                 875                 880

Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp
                885                 890                 895

Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys
            900                 905                 910

Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr
        915                 920                 925

Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser
    930                 935                 940

Glu Leu Asp Lys Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg
945                 950                 955                 960

Gln Ile Thr Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr
                965                 970                 975

Lys Tyr Asp Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr
            980                 985                 990

Leu Lys Ser Lys Leu Val Ser Asp  Phe Arg Lys Asp Phe  Gln Phe Tyr
        995                 1000                1005

Lys Val  Arg Glu Ile Asn Asn  Tyr His His Ala His  Asp Ala Tyr
    1010                1015                1020

Leu Asn  Ala Val Val Gly Thr  Ala Leu Ile Lys Lys  Tyr Pro Lys
    1025                1030                1035

Leu Glu  Ser Glu Phe Val Tyr  Gly Asp Tyr Lys Val  Tyr Asp Val
    1040                1045                1050

Arg Lys  Met Ile Ala Lys Ser  Glu Gln Glu Ile Gly  Lys Ala Thr
    1055                1060                1065

Ala Lys  Tyr Phe Phe Tyr Ser  Asn Ile Met Asn Phe  Phe Lys Thr
    1070                1075                1080

Glu Ile  Thr Leu Ala Asn Gly  Glu Ile Arg Lys Arg  Pro Leu Ile
    1085                1090                1095

Glu Thr  Asn Gly Glu Thr Gly  Glu Ile Val Trp Asp  Lys Gly Arg
    1100                1105                1110

Asp Phe  Ala Thr Val Arg Lys  Val Leu Ser Met Pro  Gln Val Asn
    1115                1120                1125

Ile Val  Lys Lys Thr Glu Val  Gln Thr Gly Gly Phe  Ser Lys Glu
    1130                1135                1140

Ser Ile  Leu Pro Lys Arg Asn  Ser Asp Lys Leu Ile  Ala Arg Lys
    1145                1150                1155

Lys Asp  Trp Asp Pro Lys Lys  Tyr Gly Gly Phe Asp  Ser Pro Thr
    1160                1165                1170

Val Ala  Tyr Ser Val Leu Val  Val Ala Lys Val Glu  Lys Gly Lys
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    1175                1180                1185

Ser Lys  Lys Leu Lys Ser Val  Lys Glu Leu Leu Gly  Ile Thr Ile
    1190                1195                1200

Met Glu  Arg Ser Ser Phe Glu  Lys Asn Pro Ile Asp  Phe Leu Glu
    1205                1210                1215

Ala Lys  Gly Tyr Lys Glu Val  Lys Lys Asp Leu Ile  Ile Lys Leu
    1220                1225                1230

Pro Lys  Tyr Ser Leu Phe Glu  Leu Glu Asn Gly Arg  Lys Arg Met
    1235                1240                1245

Leu Ala  Ser Ala Gly Glu Leu  Gln Lys Gly Asn Glu  Leu Ala Leu
    1250                1255                1260

Pro Ser  Lys Tyr Val Asn Phe  Leu Tyr Leu Ala Ser  His Tyr Glu
    1265                1270                1275

Lys Leu  Lys Gly Ser Pro Glu  Asp Asn Glu Gln Lys  Gln Leu Phe
    1280                1285                1290

Val Glu  Gln His Lys His Tyr  Leu Asp Glu Ile Ile  Glu Gln Ile
    1295                1300                1305

Ser Glu  Phe Ser Lys Arg Val  Ile Leu Ala Asp Ala  Asn Leu Asp
    1310                1315                1320

Lys Val  Leu Ser Ala Tyr Asn  Lys His Arg Asp Lys  Pro Ile Arg
    1325                1330                1335

Glu Gln  Ala Glu Asn Ile Ile  His Leu Phe Thr Leu  Thr Asn Leu
    1340                1345                1350

Gly Ala  Pro Ala Ala Phe Lys  Tyr Phe Asp Thr Thr  Ile Asp Arg
    1355                1360                1365

Lys Arg  Tyr Thr Ser Thr Lys  Glu Val Leu Asp Ala  Thr Leu Ile
    1370                1375                1380

His Gln  Ser Ile Thr Gly Leu  Tyr Glu Thr Arg Ile  Asp Leu Ser
    1385                1390                1395

Gln Leu  Gly Gly Asp Pro Ile  Ala Gly Ser Lys Ala  Ser Pro Lys
    1400                1405                1410

Lys Lys  Arg Lys Val Gly Arg  Ala Asp Ala Leu Asp  Asp Phe Asp
    1415                1420                1425

Leu Asp  Met Leu Gly Ser Asp  Ala Leu Asp Asp Phe  Asp Leu Asp
    1430                1435                1440

Met Leu  Gly Ser Asp Ala Leu  Asp Asp Phe Asp Leu  Asp Met Leu
    1445                1450                1455

Gly Ser  Asp Ala Leu Asp Asp  Phe Asp Leu Asp Met  Leu Ile Asn
    1460                1465                1470

Tyr Pro  Tyr Asp Val Pro Asp  Tyr Ala Ser
    1475                1480


<210> SEQ ID NO 2
<211> LENGTH: 350
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polynucleotide

<400> SEQUENCE: 2

gagggcctat ttcccatgat tccttcatat ttgcatatac gatacaaggc tgttagagag     60

ataattggaa ttaatttgac tgtaaacaca aagatattag tacaaaatac gtgacgtaga    120
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aagtaataat ttcttgggta gtttgcagtt ttaaaattat gttttaaaat ggactatcat 180

atgcttatcg taacttgaaa gtatttcgat ttcttggctt tatatatctt gtggaaagga 240

cgaaacaccg ggtcttcgag aagacctgtt ttagagctag aaatagcaag ttaaaataag 300

gctagtccgt tatcaacttg aaaaagtggc accgagtcgg tgctttttt 350

<210> SEQ ID NO 3
<211> LENGTH: 338
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 3

acccccggca gcagcccgcc ccgagcgcgc cgcctgttta ttcagccggg agtccggcac 60

gcgccaggcg cacgcactgc aacaacaaac ccagctgaat ggagagtttg caaggagcgg 120

gagaaaggaa cgggaggggg ggagaggaga ggaggagggg gagtttaggg agtgggtggg 180

aggaagaggt aagaggaggg gggggagtgg gggctgcagc cgctcgctgc agcagcgggg 240

agtgggggggc gaggcggggc cagggctgcg cgtggggctg ggtgtcccat tgaaaaggcg 300

gacgcactcc ggcagcccag cactctctca cttctggc 338

<210> SEQ ID NO 4
<211> LENGTH: 305
<212> TYPE: DNA
<213> ORGANISM: Mus musculus

<400> SEQUENCE: 4

ccgccccgag cgcgcggcct gtttattcag ccgggagtcc ggcacgcgcc gggcgcacgc 60

actgcaacaa caaaccaggc tgaatggaga gtttgcaagg agcgggcgcg ggcaactgga 120

gggggggggg gcgagaggga gggagctgag gaggtggggg aagaggaggg gtagtggggg 180

ctgcagccgc tcgctgcagc agcggggagt ggggggcgag gcggggccag ggctgcgcgt 240

ggggctgggt gtcccattga aaaggcggcc gcaccgcagc cgcccagcag tctctcactt 300

ctggc 305

<210> SEQ ID NO 5
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 5

gctgggtgtc ccattgaaa 19

<210> SEQ ID NO 6
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 6

cagccgctcg ctgcagcag 19

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<210> SEQ ID NO 7
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 7

tggagagttt gcaaggagc 19

<210> SEQ ID NO 8
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 8

gtttattcag ccgggagtc 19

<210> SEQ ID NO 9
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 9

cgccaggagg ggtgggtcta 20

<210> SEQ ID NO 10
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 10

ccttggtgag actggtaga 19

<210> SEQ ID NO 11
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 11

gtcttcaggt tctgttgct 19

<210> SEQ ID NO 12
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 12

atattcctga tttaaaagt 19

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<210> SEQ ID NO 13
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 13

ttaaaagtcg gctggtagc 19

<210> SEQ ID NO 14
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 14

cgggccgggg gcggggtcc 19

<210> SEQ ID NO 15
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 15

gcccgagccg cgtgtggaa 19

<210> SEQ ID NO 16
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 16

ccttcattgc ggcgggctg 19

<210> SEQ ID NO 17
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 17

ccgacccctc ccgggtccc 19

<210> SEQ ID NO 18
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 18

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caggaccgcg cttcccacg 19

<210> SEQ ID NO 19
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 19

tgcaccctgg gagcgcgag 19

<210> SEQ ID NO 20
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 20

ccgcacgcac ctgttccca 19

<210> SEQ ID NO 21
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 21

aaaacagcga gggagaaac 19

<210> SEQ ID NO 22
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 22

ttaacttgat tgtgaaatc 19

<210> SEQ ID NO 23
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 23

aaaacaatgc atatttgca 19

<210> SEQ ID NO 24
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 24

aaaatccagt attttaatg 19

<210> SEQ ID NO 25
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 25

acccagcact gcagcctgg 19

<210> SEQ ID NO 26
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 26

aacttatgcg gcgtttcct 19

<210> SEQ ID NO 27
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 27

tcactttaaa accacctct 19

<210> SEQ ID NO 28
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 28

gcatcttttt ctctttaat 19

<210> SEQ ID NO 29
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 29

tgtactctct gaggtgctc 19

<210> SEQ ID NO 30
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 30

acgcagataa gaaccagtt 19

<210> SEQ ID NO 31
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 31

catcaagtca gccatcagc 19

<210> SEQ ID NO 32
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 32

gagtcaccct cctggaaac 19

<210> SEQ ID NO 33
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 33

gctagggatg aagaataaa 19

<210> SEQ ID NO 34
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 34

ttgaccaata gccttgaca 19

<210> SEQ ID NO 35
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 35

tgcaaatatc tgtctgaaa 19

<210> SEQ ID NO 36
<211> LENGTH: 19
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 36

aaattagcag tatcctctt 19

<210> SEQ ID NO 37
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 37

cctgggctcc ggggcgttt 19

<210> SEQ ID NO 38
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 38

ggcccctgcg gccaccccg 19

<210> SEQ ID NO 39
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 39

ctccctccct gcccggtag 19

<210> SEQ ID NO 40
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 40

aggtttggaa agggcgtgc 19

<210> SEQ ID NO 41
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 41

ggagcttctc gacttcacca 20

<210> SEQ ID NO 42

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<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 42

gatttgtggg cctgaagaaa 20

<210> SEQ ID NO 43
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 43

aaggaggagg gcagaatcat 20

<210> SEQ ID NO 44
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 44

aaaccttcct cagctatgcc c 21

<210> SEQ ID NO 45
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 45

agctgatggc cctaaacaga 20

<210> SEQ ID NO 46
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 46

caggaggact ctggcaccta 20

<210> SEQ ID NO 47
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 47

ggaatccatg gagggaagat 20

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<210> SEQ ID NO 48
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 48

gctgagtgaa ctgcactgtg a 21

<210> SEQ ID NO 49
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 49

ctctctgctc ctttgccaca 20

<210> SEQ ID NO 50
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 50

caatgacccc ttcattgacc 20

<210> SEQ ID NO 51
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 51

ggaacaagag ctgctggact 20

<210> SEQ ID NO 52
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 52

aactttggca ttgtggaagg 20

<210> SEQ ID NO 53
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 53

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aacgccactg acaagaaagc 20

<210> SEQ ID NO 54
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 54

cagatccatg gaggaaggaa 20

<210> SEQ ID NO 55
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 55

gggtactcct ggaagatgtc c 21

<210> SEQ ID NO 56
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 56

gtttgcgacg catgttcctc 20

<210> SEQ ID NO 57
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 57

aagcccttgc tgtagtggtg 20

<210> SEQ ID NO 58
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 58

cggcaggaaa gcatctgtat 20

<210> SEQ ID NO 59
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 59

tgttctcgct caggtcagtg 20

<210> SEQ ID NO 60
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 60

gaattctttg ccgaaatgga 20

<210> SEQ ID NO 61
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 61

gtgctcttcg ggtttcagga 20

<210> SEQ ID NO 62
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 62

ttgattttgg agggatctcg 20

<210> SEQ ID NO 63
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 63

gtttttctgc ctccccattt 20

<210> SEQ ID NO 64
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 64

ggatgcaggg atgatgttct 20

<210> SEQ ID NO 65
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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oligonucleotide

<400> SEQUENCE: 65

gattggcttt gatttcccta 20

<210> SEQ ID NO 66
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 66

gtgtagagta agtcagccta tgg 23

<210> SEQ ID NO 67
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 67

gcctactcag actgttactc 20

<210> SEQ ID NO 68
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 68

gttggacaga acttaccgac tgg 23

<210> SEQ ID NO 69
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 69

gcagttgcct aagaactggt 20

<210> SEQ ID NO 70
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 70

ggggctccac cctcacgagt 20

<210> SEQ ID NO 71
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 71

gtttgcttcg ctataaaacg agg 23

<210> SEQ ID NO 72
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 72

gtctgaggat ggggccgcaa tgg 23

<210> SEQ ID NO 73
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 73

ggatctgtca aatcgcctgc agg 23

<210> SEQ ID NO 74
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 74

gccaggatgg cattgggcag cgg 23

<210> SEQ ID NO 75
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 75

gctgaatctg cggtggcagg agg 23

<210> SEQ ID NO 76
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 76

gttcttttgt tcttctagcc tgg 23

<210> SEQ ID NO 77
<211> LENGTH: 23

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 77

ggaaaagctt gagcaagtca agg 23

<210> SEQ ID NO 78
<211> LENGTH: 23
<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 78

ggaagagttg cccctgcgcc agg 23

<210> SEQ ID NO 79
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 79

gacaaatctc cagtggataa agg 23

<210> SEQ ID NO 80
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 80

gtgtttctca ggtaaagctc tgg 23

<210> SEQ ID NO 81
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 81

ggaaggacca tttgacgtta agg 23

<210> SEQ ID NO 82
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 82

gaactgctat ttcagtttcc tgg 23

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<210> SEQ ID NO 83
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 83

gccagccact cagccagtga agg 23

<210> SEQ ID NO 84
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 84

ggtatgcttt tctgttaaag agg 23

<210> SEQ ID NO 85
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 85

gctcctggac tgaccactat tgg 23

<210> SEQ ID NO 86
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 86

ggaacagagg cgtccccagt tgg 23

<210> SEQ ID NO 87
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 87

ggaggctaga acaatcatta cgg 23

<210> SEQ ID NO 88
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 88

gacaagaaca ccttcagaac cgg 23

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<210> SEQ ID NO 89
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 89

gggtttctgt gattttcttt tgg 23

<210> SEQ ID NO 90
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 90

gggccaaaga cctccgccag tgg 23

<210> SEQ ID NO 91
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 91

gttggagaag cattcataaa agg 23

<210> SEQ ID NO 92
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 92

gtcgctcact caccctgcaa agg 23

<210> SEQ ID NO 93
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 93

gaaaagagct gatgaaacaa tgg 23

<210> SEQ ID NO 94
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 94

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gtacactttt caaaatgctt tgg 23

<210> SEQ ID NO 95
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 95

ggagatgatc atcaagcaga agg 23

<210> SEQ ID NO 96
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 96

gctttgaaag agcaataaaa tgg 23

<210> SEQ ID NO 97
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 97

gcacaaaagt caaatcggaa tgg 23

<210> SEQ ID NO 98
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 98

gatttcaata taagattcgg agg 23

<210> SEQ ID NO 99
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 99

gcttaagcaa tcccgaactc tgg 23

<210> SEQ ID NO 100
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 100

gccttcttta tcccctatcg 20

<210> SEQ ID NO 101

<400> SEQUENCE: 101

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<210> SEQ ID NO 102

<400> SEQUENCE: 102

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<210> SEQ ID NO 103

<400> SEQUENCE: 103

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<210> SEQ ID NO 104
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (21)..(22)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 104

gaggccaaac ctcggcttac nngrr 25

<210> SEQ ID NO 105
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (21)..(22)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 105

gttcgaaaat ttcaggtaag nngrr 25

<210> SEQ ID NO 106
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (21)..(22)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 106

ggcagaacag gagataacag nngrrt 26

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<210> SEQ ID NO 107
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 107

ggcggccctc gcccttctct ggggat 26

<210> SEQ ID NO 108
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 108

gtagtgatcg tggatacgag agg 23

<210> SEQ ID NO 109
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 109

gtacagccct cggtgtatat tgg 23

<210> SEQ ID NO 110
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 110

gggaaggaat taagcccgaa tgg 23

<210> SEQ ID NO 111
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 111

gggaacagct ttcgtagttg agg 23

<210> SEQ ID NO 112
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 112

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gataaagtcc agtgtcgatc agg 23

<210> SEQ ID NO 113
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 113

gaaaaccaga gcttcggtca agg 23

<210> SEQ ID NO 114
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 114

ggagtcttct gggcaggctt aaaggctaac ctgg 34

<210> SEQ ID NO 115
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 115

gtcgggtgag catgtcttta atctacctcg atgg 34

<210> SEQ ID NO 116
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 116

ggtgtcacca gagtaacagt ctgagt 26

<210> SEQ ID NO 117
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 117

gtgatcatca agcagaaggt atgag 25

<210> SEQ ID NO 118
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 118

gaacttcgaa aatttcaggt aagccgagg 29

<210> SEQ ID NO 119
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 119

ggaaactcat caaatatgcg tgttagtgt 29

<210> SEQ ID NO 120
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 120

gtcatttaca ctaacacgca tatttgatg 29

<210> SEQ ID NO 121
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 121

ggaatgaaac tcatcaaata tgcgtgtta 29

<210> SEQ ID NO 122
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 122

gtcatcaata tctttgaagg actctgggt 29

<210> SEQ ID NO 123
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 123

gtgttttcat aggaaaaata ggcaagttg 29

<210> SEQ ID NO 124
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 124

gaattggaaa atgtgatggg aaacagata 29

<210> SEQ ID NO 125
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 125

gatgatcatc aagcagaagg tatgagaaa 29

<210> SEQ ID NO 126
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 126

gagatgatca tcaagcagaa ggtatgaga 29

<210> SEQ ID NO 127
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 127

gcatttttc tcataccttc tgcttgatg 29

<210> SEQ ID NO 128
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 128

gtcctactca gactgttact ctggtgaca 29

<210> SEQ ID NO 129
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 129

gacaggttgt gtcaccagag taacagtct 29

<210> SEQ ID NO 130
<211> LENGTH: 29
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 130

gttatcattt tttctcatac cttctgctt 29

<210> SEQ ID NO 131
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 131

gttgcctaag aactggtggg aaatggtct 29

<210> SEQ ID NO 132
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 132

gaaacagttg cctaagaact ggtgggaaa 29

<210> SEQ ID NO 133
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 133

gtttcccacc agttcttagg caactgttt 29

<210> SEQ ID NO 134
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 134

gtggctttga tttccctagg gtccagctt 29

<210> SEQ ID NO 135
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 135

gtagggaaat caaagccaat gaaacgttc 29

<210> SEQ ID NO 136

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<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 136

ggaccctagg gaaatcaaag ccaatgaaa 29

<210> SEQ ID NO 137
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 137

gtgagggctc caccctcacg agtgggttt 29

<210> SEQ ID NO 138
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 138

gaaggattga gggctccacc ctcacgagt 29

<210> SEQ ID NO 139
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 139

ggctccaccc tcacgagtgg gtttggttc 29

<210> SEQ ID NO 140
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 140

gtatccccta tcgaggaaac cacgagttt 29

<210> SEQ ID NO 141
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 141

ggataaagaa ggcctatttc atagagttg 29

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<210> SEQ ID NO 142
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 142

gaggccttct ttatccccta tcgaggaaa 29

<210> SEQ ID NO 143
<211> LENGTH: 27
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 143

gtgagggctc caccctcacg agtgggt 27

<210> SEQ ID NO 144
<211> LENGTH: 27
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 144

ggataaagaa ggcctatttc atagagt 27

<210> SEQ ID NO 145
<211> LENGTH: 4932
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 145

gccaccatgg acaagaagta ctccattggg ctcgatatcg gcacaaacag cgtcggctgg 60

gccgtcatta cggacgagta caaggtgccg agcaaaaaat tcaaagttct gggcaatacc 120

gatcgccaca gcataaagaa gaacctcatt ggcgccctcc tgttcgactc cggggagacg 180

gccgaagcca cgcggctcaa aagaacagca cggcgcagat atacccgcag aaagaatcgg 240

atctgctacc tgcaggagat ctttagtaat gagatggcta aggtggatga ctctttcttc 300

cataggctgg aggagtcctt tttggtggag gaggataaaa agcacgagcg ccacccaatc 360

tttggcaata tcgtggacga ggtggcgtac catgaaaagt acccaaccat atatcatctg 420

aggaagaagc ttgtagacag tactgataag gctgacttgc ggttgatcta tctcgcgctg 480

gcgcatatga tcaaatttcg gggacacttc ctcatcgagg gggacctgaa cccagacaac 540

agcgatgtcg acaaactctt tatccaactg gttcagactt acaatcagct tttcgaagag 600

aacccgatca acgcatccgg agttgacgcc aaagcaatcc tgagcgctag gctgtccaaa 660

tcccggcggc tcgaaaacct catcgcacag ctccctgggg agaagaagaa cggcctgttt 720

ggtaatctta tcgccctgtc actcgggctg acccccaact ttaaatctaa cttcgacctg 780

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tcctactcag actgttactc tgg 23

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tcctactcac actgttactc agg 23

<210> SEQ ID NO 152
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<210> SEQ ID NO 153
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gcattctcaa actgttactc agg 23

<210> SEQ ID NO 154
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acatacttat actgttactc tag 23

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tattcctaag actgttactc aag 23

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gagctctcat actgttactc tag 23

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<400> SEQUENCE: 159

gcaaaatgag actgttactc cag 23

<210> SEQ ID NO 160
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 160

cctcattcag actgttactc aag 23

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<400> SEQUENCE: 161

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aattggcatt gatttcccta gag 23

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gataggctgt gatttcccta gag 23

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gaatagcctt gatttcccta aag 23

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<213> ORGANISM: Artificial Sequence
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<400> SEQUENCE: 166

aatttgcttt gatttccctg agg 23

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<400> SEQUENCE: 167

gatgtgcttt gatttccctt ggg 23

<210> SEQ ID NO 168
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<400> SEQUENCE: 168

aattggtttt aatttcccta aag 23

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gatgggtttt tatttcccta gag 23

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gaatggtttt gatttccctg gag 23

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<400> SEQUENCE: 172

acagttgcct aagaactggt ggg 23

<210> SEQ ID NO 173
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<400> SEQUENCE: 173

ccagttgtct aagaactggg gag 23

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<400> SEQUENCE: 174

gcagttgcct gtgaactggt agg 23

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gcagatgcag aagaactggt gag 23

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<400> SEQUENCE: 176

gcagttccag aagaactggt gag 23

<210> SEQ ID NO 177
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<400> SEQUENCE: 177

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<400> SEQUENCE: 178

acacctgcct aagaactgga ggg 23

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<400> SEQUENCE: 179

tcaggtggct aagaactggg tgg 23

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<400> SEQUENCE: 180

gaagttggcc aagaactgga gag 23

<210> SEQ ID NO 181
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<400> SEQUENCE: 181

gctgctgccc aagaactggc agg 23

<210> SEQ ID NO 182
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<400> SEQUENCE: 182

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<400> SEQUENCE: 184

gcagctcagc cctcacgagt cag 23

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 185

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<210> SEQ ID NO 186
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<400> SEQUENCE: 186

ggggctctcc cctcactagt gag 23

<210> SEQ ID NO 187
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<400> SEQUENCE: 187

ggggatccac cttcaccagt cag 23

<210> SEQ ID NO 188
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<212> TYPE: DNA
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 188

agggctggac cctcacaagt aag 23

<210> SEQ ID NO 189
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<212> TYPE: DNA
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<220> FEATURE:
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<400> SEQUENCE: 189

tggtctcctc ccccacgagt ggg 23

<210> SEQ ID NO 190
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<400> SEQUENCE: 190

agggctccca ccccacgagt gag 23

<210> SEQ ID NO 191
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 191

gaggctccat actcaccagt gag 23

<210> SEQ ID NO 192
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<212> TYPE: DNA
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 192

ggagctgccc cttcacgagt ggg 23

<210> SEQ ID NO 193
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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 193

atgactccac cctcaagagt aag 23

<210> SEQ ID NO 194
<211> LENGTH: 23
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 194

gccttcttta tcccctatcg agg 23

<210> SEQ ID NO 195
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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oligonucleotide

<400> SEQUENCE: 195

gtctgctgtg tcccctatcg ggg 23

<210> SEQ ID NO 196
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 196

cccttctcta tcccctgtcg tgg 23

<210> SEQ ID NO 197
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 197

gccttcttta tcccctctct tgg 23

<210> SEQ ID NO 198
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 198

gcgctctttt tcccctatct tag 23

<210> SEQ ID NO 199
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 199

gccctctgtc tcccctgtcg cag 23

<210> SEQ ID NO 200
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 200

tccatctttg tcccctattg agg 23

<210> SEQ ID NO 201
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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 201

accttctctc tcccctatag agg 23

<210> SEQ ID NO 202
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 202

gttttctttt tcccctatgg gag 23

<210> SEQ ID NO 203
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 203

tgcttcttaa tcccctatca aag 23

<210> SEQ ID NO 204
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 204

accttcttac tcccctatcc ggg 23

<210> SEQ ID NO 205
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 205

gagaggttat gtggctttac ca 22

<210> SEQ ID NO 206
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 206

aaaaatgctt cccactttgc 20

<210> SEQ ID NO 207
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 207

ctcattctca tgcctggaca 20

<210> SEQ ID NO 208
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 208

gagtttggct caaattgtta ctctt 25

<210> SEQ ID NO 209
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 209

gggaaatggt ctaggagagt aaagt 25

<210> SEQ ID NO 210
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 210

gtttggctca aattgttact cttca 25

<210> SEQ ID NO 211
<211> LENGTH: 25
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 211

gtgagagtaa tgtgtttgct gagag 25

<210> SEQ ID NO 212
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<220> FEATURE:
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cgggcttgga cagaacttac 20

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<400> SEQUENCE: 213

ctgcgtagtg ccaaaacaaa 20

<210> SEQ ID NO 214
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<400> SEQUENCE: 214

taatttcatt gaagagtggc tgaa 24

<210> SEQ ID NO 215
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aagccctgtg tggtagtagt cagt 24

<210> SEQ ID NO 216
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tgagtcatgt tggataacca gtct 24

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gaaggtcagg aacatacaat tcaa 24

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gatatgggca tgtcagtttc atag 24

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tgctgttgat taatggttga tagg 24

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ttttaaattg ccatgtttgt gtc 23

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atgaataacc taatgggcag aaaa 24

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tcaagtcgct tcattttgat agac 24

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cacaacaaaa catatagcca aagc 24

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tgctgctaaa ataacacaaa tcagt 25

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ctgtgcctat tgtggttatc ctg 23

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attgatctgc aatacatgtg gagt 24

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tttgcctctg ctattacagt atgg 24

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<400> SEQUENCE: 228

tgtagggtgg ttggctaaaa taat 24

<210> SEQ ID NO 229
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tttttgcaca gtcaataaca caaa 24

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<400> SEQUENCE: 230

ggctggtctc acaattgtac ttta 24

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<400> SEQUENCE: 231

cattatggac tgaaaatctc agca 24

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<400> SEQUENCE: 232

atcatcctag ccataacaca atga 24

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<400> SEQUENCE: 233

ttcagcttta acgtgatttt ctgt 24

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 234

ggattcagaa gctgtttacg aagt 24

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<212> TYPE: DNA
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 235

tttagctgga ttggaaaaac aaat 24

<210> SEQ ID NO 236
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 236

aactcacccc attgttggta tatt 24

<210> SEQ ID NO 237
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 237

ccttgtccaa ataccgaaat acat 24

<210> SEQ ID NO 238
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 238

cacataattc atgaacttgg cttc 24

<210> SEQ ID NO 239
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<400> SEQUENCE: 239

tagtagctgg ggaggaagat acag 24

<210> SEQ ID NO 240
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 240

tttttgtttt aattgcgact gtgt 24

<210> SEQ ID NO 241
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 241

agaaaagggg ttttcttttg actt 24

<210> SEQ ID NO 242
<211> LENGTH: 24
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 242

cattgtgact ggatgagaag aaac 24

<210> SEQ ID NO 243
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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 243

aacggctgtt attaaagtcc tcag 24

<210> SEQ ID NO 244
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 244

caagtcagaa gtcacttgct ttgt 24

<210> SEQ ID NO 245
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 245

ttttatgtgc aggaatcagt ctgt 24

<210> SEQ ID NO 246
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 246

tggcggcgtt ttcattat 18

<210> SEQ ID NO 247
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 247

ttcgatccgt aatgattgtt ctagcc 26

<210> SEQ ID NO 248

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<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 248

ggtcttccag agtgctgagg 20

<210> SEQ ID NO 249
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 249

tgtgtgcttc tgtacacatc atct 24

<210> SEQ ID NO 250
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 250

agatttcaac cctcaaaaac tgag 24

<210> SEQ ID NO 251
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 251

taaactcttt cttttccgca attc 24

<210> SEQ ID NO 252
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 252

caaggtgacc tgctacctaa aaat 24

<210> SEQ ID NO 253
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 253

tatgaccaag gctatgtgtt cact 24

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<210> SEQ ID NO 254
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 254

acagcctctc tccagtaaca ttct 24

<210> SEQ ID NO 255
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 255

tattcttgca gtggtttcac attt 24

<210> SEQ ID NO 256
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 256

atattttaag ccaagaccca acaa 24

<210> SEQ ID NO 257
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 257

ctttcaactg tctgtctgat tgct 24

<210> SEQ ID NO 258
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 258

aacagcctct cttcattgtt ctct 24

<210> SEQ ID NO 259
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 259

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ctctggaact tgtctctgtc ttga 24

<210> SEQ ID NO 260
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 260

ctttcctgcg ttctcatgtt acta 24

<210> SEQ ID NO 261
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 261

ccttatatcc gtatcgctca ctct 24

<210> SEQ ID NO 262
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 262

catatctgtc taacttccgc acac 24

<210> SEQ ID NO 263
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 263

acaggtgtta tgttgtctgc atct 24

<210> SEQ ID NO 264
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 264

actccattcc cagattagtt atgc 24

<210> SEQ ID NO 265
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 265

ctgttttctt tgtgagagtg gaga 24

<210> SEQ ID NO 266
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 266

tgtaaggtgg tcaaacttgc tcta 24

<210> SEQ ID NO 267
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 267

tttttcctag tacccacaga ttttt 25

<210> SEQ ID NO 268
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 268

tccctgattc tctcatttgt gtta 24

<210> SEQ ID NO 269
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 269

ttgggaacat cagagaaagt atga 24

<210> SEQ ID NO 270
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 270

acaaattaca gtctcctggg aaag 24

<210> SEQ ID NO 271
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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<400> SEQUENCE: 271

agtagcttac cttggcagag aaaa 24

<210> SEQ ID NO 272
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 272

tgacatactg ttacccttg cagt 24

<210> SEQ ID NO 273
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 273

gaaaggctca gtgaatgttt gtt 23

<210> SEQ ID NO 274
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 274

cactgcatca tctcattaaa tcaa 24

<210> SEQ ID NO 275
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 275

cccatatatt catgattacc caca 24

<210> SEQ ID NO 276
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 276

tatcagaacg agcactaaaa gcac 24

<210> SEQ ID NO 277
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 277

ttgggaggct gaggtacaag 20

<210> SEQ ID NO 278
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 278

gaatgaaaaa caaacagaag gtga 24

<210> SEQ ID NO 279
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 279

ctcctcatct gtacccttca atct 24

<210> SEQ ID NO 280
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 280

agagtggcat ctagtgtcag tgag 24

<210> SEQ ID NO 281
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 281

taccaaaagc ttctcctgtt tacc 24

<210> SEQ ID NO 282
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 282

gtaagttgga tggcctattc tttg 24

<210> SEQ ID NO 283
<211> LENGTH: 24

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 283

gaaggaaatg caaggataca agat 24

<210> SEQ ID NO 284
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 284

tgattgaaag aatcattcca gaaa 24

<210> SEQ ID NO 285
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 285

tcagaaggaa aattgaaatt ggtt 24

<210> SEQ ID NO 286
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 286

cagatgtgtt cttcatcatt cctc 24

<210> SEQ ID NO 287
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 287

ttctctttag ggaaagctct caaa 24

<210> SEQ ID NO 288
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 288

gggtatagat catatggagg gaag 24

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<210> SEQ ID NO 289
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 289

agatgatctg cccacctcag 20

<210> SEQ ID NO 290
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 290

ctttcttcct catttagtgg caat 24

<210> SEQ ID NO 291
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 291

atgaattgca gattgatggt actg 24

<210> SEQ ID NO 292
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 292

tctcaccaag aaccaaattg tcta 24

<210> SEQ ID NO 293
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 293

gtaggatacc ttggcaacag tctt 24

<210> SEQ ID NO 294
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 294

ttaacgaatt gtgagatttg ctgt 24

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<210> SEQ ID NO 295
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 295

tcagaaagtc aagtagcaca caca 24

<210> SEQ ID NO 296
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 296

agaagcacac actcaggtaa agc 23

<210> SEQ ID NO 297
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 297

tctttggggg aataatgact aaaa 24

<210> SEQ ID NO 298
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 298

tttggcattt atgggaataa aact 24

<210> SEQ ID NO 299
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 299

actaattctg gtcaagccca tca 23

<210> SEQ ID NO 300
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 300

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ttaagacatc ggatgaacag aaag 24

<210> SEQ ID NO 301
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 301

agaagctttc tgacatgatc tgc 23

<210> SEQ ID NO 302
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 302

tcaattgcat taggacttag acca 24

<210> SEQ ID NO 303
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 303

gttaaattac ctgtgaagcc cttg 24

<210> SEQ ID NO 304
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 304

cggaaaacag atccacttta tgat 24

<210> SEQ ID NO 305
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 305

aaatccactg gaaacatctt gagt 24

<210> SEQ ID NO 306
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 306

agtctcttca gaatcatgcc ctat 24

<210> SEQ ID NO 307
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 307

gcttggtggc acatacctgt ag 22

<210> SEQ ID NO 308
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 308

ggtaggtaga tttgcttgct tgtt 24

<210> SEQ ID NO 309
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 309

agctctcagc agagtaggga ttta 24

<210> SEQ ID NO 310
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 310

gtgagtctac tgcaccccat c 21

<210> SEQ ID NO 311
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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 311

tgacactgtg aagtcaattc tgtc 24

<210> SEQ ID NO 312
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 312

tcaagaactt gacaatgagc aaat 24

<210> SEQ ID NO 313
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 313

tatccgatcc actgttgtgt gt 22

<210> SEQ ID NO 314
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 314

caggagaccc aaaaccactc tac 23

<210> SEQ ID NO 315
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 315

ttgttctaca aatagggctt cctt 24

<210> SEQ ID NO 316
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 316

tgttaagttt gggcttatgt tcct 24

<210> SEQ ID NO 317
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 317

cacaagtctc actgcacaaa cat 23

<210> SEQ ID NO 318
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<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 318

tgacccatga ttatctctct ttga 24

<210> SEQ ID NO 319
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 319

ttcagcttct gattggtttt aatg 24

<210> SEQ ID NO 320
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 320

ccaattcctt aattttccct acag 24

<210> SEQ ID NO 321
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 321

atctcagacc aggagggaga c 21

<210> SEQ ID NO 322
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 322

cctcagggtc agtacatttt tcag 24

<210> SEQ ID NO 323
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 323

ttcttaggac attgctccac atac 24

<210> SEQ ID NO 324

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<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 324

gcaaacataa tgcaactcgt aatc 24

<210> SEQ ID NO 325
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 325

gcaagggagt ctgtgtcttt g 21

<210> SEQ ID NO 326
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 326

tcatttaagt ggctgttctg tgtt 24

<210> SEQ ID NO 327
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 327

acaaaacaga gagaaaaggc agag 24

<210> SEQ ID NO 328
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 328

gttttgattt ctggtgccta cag 23

<210> SEQ ID NO 329
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 329

actgaagctg aagcccagtc 20

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<210> SEQ ID NO 330
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 330

acatgagctc tcaggtttct gac 23

<210> SEQ ID NO 331
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 331

tcaaacttag atggttccct atgtt 25

<210> SEQ ID NO 332
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 332

gtaccctgaa aatgtagggt gact 24

<210> SEQ ID NO 333
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 333

cacttcccaa gtgaggcaat 20

<210> SEQ ID NO 334
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 334

ctatacttgg ggctgacttg ctac 24

<210> SEQ ID NO 335
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 335

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tcgtataggt tactttggct caca 24

<210> SEQ ID NO 336
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 336

agggatcttt actcctcagt gtgt 24

<210> SEQ ID NO 337
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 337

tgtagaagtt ggaatatcct gctg 24

<210> SEQ ID NO 338
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 338

gtcaacaatt tgatctcagg cttc 24

<210> SEQ ID NO 339
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 339

ctcagtacta aagatggacg cttg 24

<210> SEQ ID NO 340
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 340

aatcatttca gtcttcccaa caat 24

<210> SEQ ID NO 341
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 341

gggaatcaca gtagatgttt gtca 24

<210> SEQ ID NO 342
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 342

agaccaggag gtaagaacat tttg 24

<210> SEQ ID NO 343
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 343

ccacatagaa agagacttgc agaa 24

<210> SEQ ID NO 344
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 344

agagatgcca aaagaacagt caat 24

<210> SEQ ID NO 345
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 345

tgtgccttag gctatgtaaa ctgt 24

<210> SEQ ID NO 346
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 346

aaaccttgt aaccaaaatt acca 24

<210> SEQ ID NO 347
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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<400> SEQUENCE: 347

taactgcatc agaagtcctt gcta 24

<210> SEQ ID NO 348
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 348

ggagaccaag ctgctaaagt ca 22

<210> SEQ ID NO 349
<211> LENGTH: 37
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 349

gtggtgccgc gggagtttgg ctcaaattgt tactctt 37

<210> SEQ ID NO 350
<211> LENGTH: 37
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 350

gtggtgccgc gggggaaatg gtctaggaga gtaaagt 37

<210> SEQ ID NO 351
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 351

gtggtgccgc gggagaggtt atgtggcttt acca 34

<210> SEQ ID NO 352
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 352

gtggtgccgc ggctcattct catgcctgga ca 32

<210> SEQ ID NO 353
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 353

gtggtgccgc ggcgggcttg gacagaactt ac 32

<210> SEQ ID NO 354
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 354

gtggtgccgc ggctgcgtag tgccaaaaca aa 32

<210> SEQ ID NO 355
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 355

gtggtgccgc ggtaatttca ttgaagagtg gctgaa 36

<210> SEQ ID NO 356
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 356

gtggtgccgc ggaagccctg tgtggtagta gtcagt 36

<210> SEQ ID NO 357
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 357

gtggtgccgc ggcaagtcag aagtcacttg ctttgt 36

<210> SEQ ID NO 358
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 358

gtggtgccgc ggttttatgt gcaggaatca gtctgt 36

<210> SEQ ID NO 359
<211> LENGTH: 36

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 359

gtggtgccgc ggtgtgtgct tctgtacaca tcatct 36

<210> SEQ ID NO 360
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 360

gtggtgccgc ggagatttca accctcaaaa actgag 36

<210> SEQ ID NO 361
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 361

gtggtgccgc ggttgggaac atcagagaaa gtatga 36

<210> SEQ ID NO 362
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 362

gtggtgccgc ggacaaatta cagtctcctg ggaaag 36

<210> SEQ ID NO 363
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 363

gtggtgccgc ggcacttccc aagtgaggca at 32

<210> SEQ ID NO 364
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 364

gtggtgccgc ggctatactt ggggctgact tgctac 36

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<210> SEQ ID NO 365
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 365

gtggtgccgc ggttggctct ttagcttgtg tttc 34

<210> SEQ ID NO 366
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 366

gtggtgccgc ggtgagactc ccaaaggcaa tc 32

<210> SEQ ID NO 367
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 367

gtggtgccgc ggttggctct ttagcttgtg tttc 34

<210> SEQ ID NO 368
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 368

gtggtgccgc ggactgaggg gtgatcttgg tg 32

<210> SEQ ID NO 369
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 369

gtggtgccgc gggcagagaa agccagtcgg ta 32

<210> SEQ ID NO 370
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 370

gtggtgccgc ggtgagactc ccaaaggcaa tc 32

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<210> SEQ ID NO 371
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 371

gtggtgccgc gggcagagaa agccagtcgg ta 32

<210> SEQ ID NO 372
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 372

gtggtgccgc ggactgaggg gtgatcttgg tg 32

<210> SEQ ID NO 373
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 373

gtggtgccgc ggccagagtt cctagggcag ag 32

<210> SEQ ID NO 374
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 374

gtggtgccgc ggagctagtc cccacattcc ac 32

<210> SEQ ID NO 375
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 375

gtggtgccgc ggccagagtt cctagggcag ag 32

<210> SEQ ID NO 376
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 376

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gtggtgccgc ggggtggagg gaaactttag gc 32

<210> SEQ ID NO 377
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 377

gtggtgccgc ggctcattct catgcctgga ca 32

<210> SEQ ID NO 378
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 378

gtggtgccgc ggagctagtc cccacattcc ac 32

<210> SEQ ID NO 379
<211> LENGTH: 34
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 379

gtggtgccgc ggtctcatgc ctggacaagt aact 34

<210> SEQ ID NO 380
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 380

gtggtgccgc ggggtggagg gaaactttag gc 32

<210> SEQ ID NO 381
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 381

gtggtgccgc ggggcttgga cagaacttac cg 32

<210> SEQ ID NO 382
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 382

gtggtgccgc ggcaccactg tctgcctaag ga 32

<210> SEQ ID NO 383
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 383

gtggtgccgc ggggcttgga cagaacttac cg 32

<210> SEQ ID NO 384
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 384

gtggtgccgc ggggtggagg gaaactttag gc 32

<210> SEQ ID NO 385
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 385

gtggtgccgc ggcgtagtgc caaaacaaac agt 33

<210> SEQ ID NO 386
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 386

gtggtgccgc ggcaccactg tctgcctaag ga 32

<210> SEQ ID NO 387
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 387

gtggtgccgc ggcgtagtgc caaaacaaac agt 33

<210> SEQ ID NO 388
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 388

gtggtgccgc ggggtggagg gaaactttag gc 32

<210> SEQ ID NO 389
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 389

gtggtgccgc gggcgagggc ctacttgata tg 32

<210> SEQ ID NO 390
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 390

gtggtgccgc ggcttcccaa gtgaggcaat gc 32

<210> SEQ ID NO 391
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 391

gtggtgccgc ggacgttttg tgctgctgta aca 33

<210> SEQ ID NO 392
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 392

gtggtgccgc ggctgcaggc acattctctt cc 32

<210> SEQ ID NO 393
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 393

gtggtgccgc gggccctgtg tggtagtagt ca 32

<210> SEQ ID NO 394
<211> LENGTH: 32
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 394

gtggtgccgc ggcttcccaa gtgaggcaat gc 32

<210> SEQ ID NO 395
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 395

gtggtgccgc ggcagtatta aggggtggga gct 33

<210> SEQ ID NO 396
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 396

gtggtgccgc ggtctcttcc tcacacagct ga 32

<210> SEQ ID NO 397
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 397

gtggtgccgc ggggagcttg gagggaagag aa 32

<210> SEQ ID NO 398
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 398

gtggtgccgc ggcttcccaa gtgaggcaat gc 32

<210> SEQ ID NO 399
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 399

gtggtgccgc ggatggatgg ggaagacact gg 32

<210> SEQ ID NO 400

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<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 400

gtggtgccgc ggctgcaggc acattctctt cc 32

<210> SEQ ID NO 401
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 401

gtggtgccgc ggggatgaaa cagggcagga ac 32

<210> SEQ ID NO 402
<211> LENGTH: 31
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 402

gtggtgccgc ggttcccaag tgaggcaatg c 31

<210> SEQ ID NO 403
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 403

gtggtgccgc ggtttgcaga gccatgatga gg 32

<210> SEQ ID NO 404
<211> LENGTH: 30
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 404

gtggtgccgc ggcgacagcc aaaacagccg 30

<210> SEQ ID NO 405
<211> LENGTH: 44
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 405

aaaatatttt agctcctact cagactgtta ctctggtgac acaa 44

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<210> SEQ ID NO 406
<211> LENGTH: 44
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 406

ttgtgtcacc agagtaacag tctgagtagg agctaaaata tttt 44

<210> SEQ ID NO 407
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 407

tagctcctac tcagactgtt actctggtga cacaac 36

<210> SEQ ID NO 408
<211> LENGTH: 28
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 408

tagctcctac tcagactggt gacccaac 28

<210> SEQ ID NO 409
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 409

tagctcctac tctggtgaca caac 24

<210> SEQ ID NO 410
<211> LENGTH: 28
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 410

tagctcctac tcagactggt gacacaac 28

<210> SEQ ID NO 411
<211> LENGTH: 16
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 411

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tagctcctac tcagac 16

<210> SEQ ID NO 412
<211> LENGTH: 27
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 412

tagctcctac tcagactgtt acacaac 27

<210> SEQ ID NO 413
<211> LENGTH: 30
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 413

tagctcctac tcagactgtg gtgaggtgac 30

<210> SEQ ID NO 414
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 414

tagctcctac tcagactctc tggtgacaca ac 32

<210> SEQ ID NO 415
<211> LENGTH: 31
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 415

tagctcctac tcagacctct ggtgacacaa c 31

<210> SEQ ID NO 416
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 416

tagctcctac tcaggctgtc tggtgacaca ac 32

<210> SEQ ID NO 417
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 417

tagctcctac tcagactact ctggtgacac aac 33

<210> SEQ ID NO 418
<211> LENGTH: 28
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 418

tagctcctac tcagactgtt gacacaac 28

<210> SEQ ID NO 419
<211> LENGTH: 13
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 419

ctggtgacac aac 13

<210> SEQ ID NO 420
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 420

tagctcctac tcagactgtt agacacaac 29

<210> SEQ ID NO 421
<211> LENGTH: 33
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 421

tagctcctac tcagactgct ctggtgacac aac 33

<210> SEQ ID NO 422
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 422

cagactgtta ctctggtgac 20

<210> SEQ ID NO 423
<211> LENGTH: 29
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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oligonucleotide

<400> SEQUENCE: 423

cagaccacct gtggtctcct actggtgac 29

<210> SEQ ID NO 424
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 424

gctttgattt ccctaggg 18

<210> SEQ ID NO 425
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 425

gctttgattt ccagttctta ggcaa 25

<210> SEQ ID NO 426
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 426

gctttgattc ttaggcaa 18

<210> SEQ ID NO 427
<211> LENGTH: 62
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (13)..(53)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 427

gctttgattt ccnnnnnnnn nnnnnnnnnn nnnnnnnnnn nnnnnnnnnn nnncttaggc 60

aa 62

<210> SEQ ID NO 428
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 428

tcttaaccat taccatag 18

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<210> SEQ ID NO 429
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 429

tcttaaccat taccatagag ttcttaggca ac 32

<210> SEQ ID NO 430
<211> LENGTH: 30
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 430

tcttaaccat taccaaagtt cttaggcaac 30

<210> SEQ ID NO 431
<211> LENGTH: 31
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 431

tcttaaccat taccataggt tcttaggcaa c 31

<210> SEQ ID NO 432
<211> LENGTH: 14
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 432

gaaccaaacc cact 14

<210> SEQ ID NO 433
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 433

gaaccaaacc cacttagggg ataa 24

<210> SEQ ID NO 434
<211> LENGTH: 2328
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 434

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atgaggtctg actacaaaga ccatgacggt gattataaag atcatgacat cgattacaag 60

gatgacgatg acaagatggc ccccaagaag aagaggaagg tgggcctcga gcccggagaa 120

aaaccgtaca agtgccctga gtgcgggaaa tcattctccg accctggggc gctcgtccgg 180

caccaaagga cgcatacagg ggaaaagccg tataagtgcc ccgagtgtgg aaagagcttc 240

tcgcagagag cccaccttga acgacaccaa agaacacaca ctggtgagaa accctataag 300

tgtccagagt gcggcaaatc gtttagcaga tccgatgact tggtgcgcca ccagcggaca 360

cacacgggtg aaaagcccta caaatgcccg gagtgtggga agtcgttttc aaggtcggat 420

catctgacta cccatcagcg cacccatacg ggagcggccg cccgcgccct ggtgaagagc 480

gagctggagg agaagaagtc cgagctgcgg cacaagctga agtacgtgcc ccacgagtac 540

atcgagctga tcgagatcgc caggaacccc acccaggacc gcatcctgga gatgaaggtg 600

atggagttct tcatgaaggt gtacggctac aggggagagc acctgggcgg aagcagaaag 660

cctgacggcg ccatctatac agtgggcagc cccatcgatt acggcgtgat cgtggacaca 720

aaggcctaca gcggcggcta caatctgcct atcggccagg ccgacgagat gcagagatac 780

gtgaaggaga accagacccg gaataagcac atcaacccca acgagtggtg gaaggtgtac 840

cctagcagcg tgaccgagtt caagttcctg ttcgtgagcg gccacttcaa gggcaactac 900

aaggcccagc tgaccaggct gaaccgcaaa accaactgca atggcgccgt gctgagcgtg 960

gaggagctgc tgatcggcgg cgagatgatc aaagccggca ccctgacact ggaggaggtg 1020

cggcgcaagt tcaacaacgg cgagatcaac ttcgagggca gaggaagtct tctaacatgc 1080

ggtgacgtgg aggagaatcc cggccctaga tctgactaca aagaccatga cggtgattat 1140

aaagatcatg acatcgatta caaggatgac gatgacaaga tggcccccaa gaagaagagg 1200

aaggtgggcc tcgagccggg agagaagccg tacaagtgtc ccgaatgtgg aaagagcttc 1260

tcacagtcgg gggaccttcg gcgccaccag cgcacacata ctggtgaaaa gccgtataag 1320

tgtccagaat gtggcaaatc attctccaca tcagggagcc tggtcaggca ccagcgaacc 1380

cacacgggtg agaagcccta taagtgcccc gaatgcggga agtccttttc gcagagagcc 1440

cacttggaga ggcaccagag gacccatacg ggggagaaac cttacaagtg ccctgaatgc 1500

ggaaagtcgt tctcgaccca tctggatctc atcagacatc agagaacgca cactggagag 1560

aaaccctaca aatgtcccga gtgtgggaag tcgtttagcc gaaaggacaa tctcaaaaac 1620

catcaacgga cacacacggg tgaaaaacca tacaaatgcc cggagtgcgg caaatcgttt 1680

tcccaacttg cgcacttgcg ggcacaccaa cgcacgcata ctggagcggc cgcccgcgcc 1740

ctggtgaaga gcgagctgga ggagaagaag tccgagctgc ggcacaagct gaagtacgtg 1800

ccccacgagt acatcgagct gatcgagatc gccaggaacc ccacccagga ccgcatcctg 1860

gagatgaagg tgatggagtt cttcatgaag gtgtacggct acaggggaga gcacctgggc 1920

ggaagcagaa agcctgacgg cgccatctat acagtgggca gccccatcga ttacggcgtg 1980

atcgtggaca caaaggccta cagcggcggc tacaatctgc ctatcggcca ggccgacgag 2040

atggagagat acgtggagga gaaccagaca cgggataagc acctcaaccc caacgagtgg 2100

tggaaggtgt accctagcag cgtgaccgag ttcaagttcc tgttcgtgag cggccacttc 2160

aagggcaact acaaggccca gctgaccagg ctgaaccaca tcaccaactg caatggcgcc 2220

gtgctgagcg tggaggagct gctgatcggc ggcgagatga tcaaagccgg caccctgaca 2280

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ctggaggagg tgcggcgcaa gttcaacaac ggcgagatca acttctga 2328

<210> SEQ ID NO 435
<211> LENGTH: 775
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polypeptide

<400> SEQUENCE: 435

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

Ile Asp Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg
20 25 30

Lys Val Gly Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys
35 40 45

Gly Lys Ser Phe Ser Asp Pro Gly Ala Leu Val Arg His Gln Arg Thr
50 55 60

His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe
65 70 75 80

Ser Gln Arg Ala His Leu Glu Arg His Gln Arg Thr His Thr Gly Glu
85 90 95

Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Arg Ser Asp
100 105 110

Asp Leu Val Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys
115 120 125

Cys Pro Glu Cys Gly Lys Ser Phe Ser Arg Ser Asp His Leu Thr Thr
130 135 140

His Gln Arg Thr His Thr Gly Ala Ala Ala Arg Ala Leu Val Lys Ser
145 150 155 160

Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu Lys Tyr Val
165 170 175

Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr Gln
180 185 190

Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met Lys Val Tyr
195 200 205

Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro Asp Gly Ala
210 215 220

Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile Val Asp Thr
225 230 235 240

Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp Glu
245 250 255

Met Gln Arg Tyr Val Lys Glu Asn Gln Thr Arg Asn Lys His Ile Asn
260 265 270

Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr Glu Phe Lys
275 280 285

Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys Ala Gln Leu
290 295 300

Thr Arg Leu Asn Arg Lys Thr Asn Cys Asn Gly Ala Val Leu Ser Val
305 310 315 320

Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly Thr Leu Thr
325 330 335

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Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe Glu
            340                 345                 350

Gly Arg Gly Ser Leu Leu Thr Cys Gly Asp Val Glu Glu Asn Pro Gly
        355                 360                 365

Pro Arg Ser Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp
    370                 375                 380

Ile Asp Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg
385                 390                 395                 400

Lys Val Gly Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys
                405                 410                 415

Gly Lys Ser Phe Ser Gln Ser Gly Asp Leu Arg Arg His Gln Arg Thr
            420                 425                 430

His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe
        435                 440                 445

Ser Thr Ser Gly Ser Leu Val Arg His Gln Arg Thr His Thr Gly Glu
    450                 455                 460

Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Ala
465                 470                 475                 480

His Leu Glu Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys
                485                 490                 495

Cys Pro Glu Cys Gly Lys Ser Phe Ser Thr His Leu Asp Leu Ile Arg
            500                 505                 510

His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys
        515                 520                 525

Gly Lys Ser Phe Ser Arg Lys Asp Asn Leu Lys Asn His Gln Arg Thr
    530                 535                 540

His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe
545                 550                 555                 560

Ser Gln Leu Ala His Leu Arg Ala His Gln Arg Thr His Thr Gly Ala
                565                 570                 575

Ala Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu
            580                 585                 590

Leu Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile
        595                 600                 605

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

Met Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly
625                 630                 635                 640

Gly Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile
                645                 650                 655

Asp Tyr Gly Val Ile Val Asp Thr Lys Ala Tyr Ser Gly Gly Tyr Asn
            660                 665                 670

Leu Pro Ile Gly Gln Ala Asp Glu Met Glu Arg Tyr Val Glu Glu Asn
        675                 680                 685

Gln Thr Arg Asp Lys His Leu Asn Pro Asn Glu Trp Trp Lys Val Tyr
    690                 695                 700

Pro Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe
705                 710                 715                 720

Lys Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn His Ile Thr Asn
                725                 730                 735
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Cys Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu
740 745 750

Met Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe
755 760 765

Asn Asn Gly Glu Ile Asn Phe
770 775

<210> SEQ ID NO 436
<211> LENGTH: 4725
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 436

tggccactcc ctctatgcgc actcgctcgc tcggtggggc ctggcgacca aaggtcgcca 60

gacggacgtg ctttgcacgt ccggccccac cgagcgagcg agtgcgcata gagggagtgg 120

ccaactccat cactagaggt atggcagtga cgtaacgcga agcgcgcgaa gcgagaccac 180

gcctaccagc tgcgtcagca gtcaggtgac ccttttgcga cagtttgcga caccacgtgg 240

ccgctgaggg tatatattct cgagtgagcg aaccaggagc tccattttga ccgcgaaatt 300

tgaacgagca gcagccatgc cggggttcta cgagattgtc ctgaaggtcc cgagtgacct 360

ggacgagcac ctgccgggca tttctaactc gtttgttaac tgggtggccg agaaggaatg 420

ggagctgccg ccggattctg acatggatcc gaatctgatt gagcaggcac ccctgaccgt 480

ggccgaaaag cttcagcgcg agttcctggt ggagtggcgc cgcgtgagta aggccccgga 540

ggccctcttt tttgtccagt tcgaaaaggg ggagacctac ttccacctgc acgtgctgat 600

tgagaccatc ggggtcaaat ccatggtggt cggccgctac gtgagccaga ttaaagagaa 660

gctggtgacc cgcatctacc gcggggtcga gccgcagctt ccgaactggt tcgcggtgac 720

caaaacgcga aatggcgccg ggggcgggaa caaggtggtg gacgactgct acatccccaa 780

ctacctgctc cccaagaccc agcccgagct ccagtgggcg tggactaaca tggaccagta 840

tttaagcgcc tgtttgaatc tcgcggagcg taaacggctg gtggcgcagc atctgacgca 900

cgtgtcgcag acgcaggagc agaacaaaga gaatcagaac cccaattctg acgcgccggt 960

catcaggtca aaaacctcag ccaggtacat ggagctggtc gggtggctgg tggaccgcgg 1020

gatcacgtca gaaaagcaat ggattcagga ggaccaggcc tcgtacatct ccttcaacgc 1080

cgcctccaac tcgcggtccc agatcaaggc cgcgctggac aatgcctcca agatcatgag 1140

cctgacaaag acggctccgg actacctggt gggcagcaac ccgccggagg acattaccaa 1200

aaatcggatc taccaaatcc tggagctgaa cgggtacgat ccgcagtacg cggcctccgt 1260

cttcctgggc tgggcgcaaa agaagttcgg gaagaggaac accatctggc tctttgggcc 1320

ggccacgacg ggtaaaacca acatcgcgga agccatcgcc cacgccgtgc ccttctacgg 1380

ctgcgtaaac tggaccaatg agaactttcc cttcaacgat tgcgtcgaca agatggtgat 1440

ctggtgggag gagggcaaga tgacggccaa ggtcgtggag agcgccaagg ccattctggg 1500

cggaagcaag gtgcgcgtgg accaaaaagtg caagtcatcg gcccagatcg aacccactcc 1560

cgtgatcgtc acctccaaca ccaacatgtg cgccgtgatt gacgggaaca gcaccacctt 1620

cgagcatcag cagccgctgc aggaccggat gtttaaattt gaacttaccc gccgtttgga 1680

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ccatgacttt gggaaggtca ccaaacagga agtaaaggac tttttccggt gggcttccga 1740

tcacgtgact gacgtggctc atgagttcta cgtcagaaag ggtggagcta agaaacgccc 1800

cgcctccaat gacgcggatg taagcgagcc aaaacggcag tgcacgtcac ttgcgcagcc 1860

gacaacgtca gacgcggaag caccggcgga ctacgcggac aggtaccaaa acaaatgttc 1920

tcgtcacgtg ggcatgaatc tgatgctttt tccctgtaaa acatgcgaga gaatgaatca 1980

aatttccaat gtctgtttta cgcatggtca aagagactgt ggggaatgct tccctggaat 2040

gtcagaatct caacccgttt ctgtcgtcaa aaagaagact tatcagaaac tgtgtccaat 2100

tcatcatatc ctgggaaggg cacccgagat tgcctgttcg gcctgcgatt tggccaatgt 2160

ggacttggat gactgtgttt ctgagcaata aatgacttaa accaggtatg gctgctgacg 2220

gttatcttcc agattggctc gaggacaacc tttctgaagg cattcgtgag tggtgggctc 2280

tgaaacctgg agtccctcaa cccaaagcga accaacaaca ccaggacaac cgtcggggtc 2340

ttgtgcttcc gggttacaaa tacctcggac ccggtaacgg actcgacaaa ggagagccgg 2400

tcaacgaggc ggacgcggca gccctcgaac acgacaaagc ttacgaccag cagctcaagg 2460

ccggtgacaa cccgtacctc aagtacaacc acgccgacgc cgagtttcag gagcgtcttc 2520

aagaagatac gtcttttggg ggcaaccttg gcagagcagt cttccaggcc aaaaagagga 2580

tccttgagcc tcttggtctg gttgaggaag cagctaaaac ggctcctgga aagaagaggc 2640

ctgtagatca gtctcctcag gaaccggact catcatctgg tgttggcaaa tcgggcaaac 2700

agcctgccag aaaaagacta aatttcggtc agactggcga ctcagagtca gtcccagacc 2760

ctcaacctct cggagaacca ccagcagccc ccacaagttt gggatctaat acaatggctt 2820

caggcggtgg cgcaccaatg gcagacaata acgagggtgc cgatggagtg ggtaattcct 2880

caggaaattg gcattgcgat tcccaatggc tgggcgacag agtcatcacc accagcacca 2940

gaacctgggc cctgcccact tacaacaacc atctctacaa gcaaatctcc agcgcttcaa 3000

cgggagcttc aaacgacaac cactactttg gctacagcac cccttggggg tattttgact 3060

ttaacagatt ccactgccac ttctcaccac gtgactggca gcgactcatt aacaacaact 3120

ggggattccg gcccaagaaa ctcagcttca agctcttcaa catccaagtt aaagaggtca 3180

cgcagaacga tggcacgacg actattgcca ataaccttac cagcacggtt caagtgttta 3240

cggactcgga gtatcagctc ccgtacgtgc tcgggtcggc gcaccaaggc tgtctcccgc 3300

cgtttccagc ggacgtcttc atggtccctc agtatggata cctcaccctg aacaacggaa 3360

gtcaagcggt gggacgctca tccttttact gcctggagta cttcccttcg cagatgctaa 3420

ggactggaaa taacttccaa ttcagctata ccttcgagga tgtacctttt cacagcagct 3480

acgctcacag ccagagtttg gatcgcttga tgaatcctct tattgatcag tatctgtact 3540

acctgaacag aacgcaagga acaacctctg gaacaaccaa ccaatcacgg ctgcttttta 3600

gccaggctgg gcctcagtct atgtctttgc aggccagaaa ttggctacct gggccctgct 3660

accggcaaca gagactttca aagactgcta acgacaacaa caacagtaac tttccttgga 3720

cagcggccag caaatatcat ctcaatggcc gcgactcgct ggtgaatcca ggaccagcta 3780

tggccagtca caaggacgat gaagaaaaat ttttccctat gcacggcaat ctaatatttg 3840

gcaaagaagg gacaacggca agtaacgcag aattagataa tgtaatgatt acggatgaag 3900

aagagattcg taccaccaat cctgtggcaa cagagcagta tggaactgtg gcaaataact 3960

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tgcagagctc aaatacagct cccacgacta gaactgtcaa tgatcagggg gccttacctg 4020

gcatggtgtg gcaagatcgt gacgtgtacc ttcaaggacc tatctgggca aagattcctc 4080

acacggatgg acactttcat ccttctcctc tgatgggagg ctttggactg aaacatccgc 4140

ctcctcaaat catgatcaaa aatactccgg taccggcaaa tcctccgacg actttcagcc 4200

cggccaagtt tgcttcattt atcactcagt actccactgg acaggtcagc gtggaaattg 4260

agtgggagct acagaaagaa aacagcaaac gttggaatcc agagattcag tacacttcca 4320

actacaacaa gtctgttaat gtggacttta ctgtagacac taatggtgtt tatagtgaac 4380

ctcgccctat tggaacccgg tatctcacac gaaacttgta atcctggtta atcaataaac 4440

cgtttaattc gtttcagttg aactttggct cttgtgcact tcttatctta tcttgtttcc 4500

atggctactg cgtagataag cagcggcctg cggcgcttgc gcttcgcggt ttacaactgc 4560

tggttaatat ttaactctcg ccatacctct agtgatggag ttggccactc cctctatgcg 4620

cactcgctcg ctcggtgggg ccggacgtgc aaagcacgtc cgtctggcga cctttggtcg 4680

ccaggcccca ccgagcgagc gagtgcgcat agagggagtg gccaa 4725

<210> SEQ ID NO 437
<211> LENGTH: 737
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polypeptide

<400> SEQUENCE: 437

Met Ala Ala Asp Gly Tyr Leu Pro Asp Trp Leu Glu Asp Asn Leu Ser
1 5 10 15

Glu Gly Ile Arg Glu Trp Trp Ala Leu Lys Pro Gly Val Pro Gln Pro
20 25 30

Lys Ala Asn Gln Gln His Gln Asp Asn Arg Arg Gly Leu Val Leu Pro
35 40 45

Gly Tyr Lys Tyr Leu Gly Pro Gly Asn Gly Leu Asp Lys Gly Glu Pro
50 55 60

Val Asn Glu Ala Asp Ala Ala Ala Leu Glu His Asp Lys Ala Tyr Asp
65 70 75 80

Gln Gln Leu Lys Ala Gly Asp Asn Pro Tyr Leu Lys Tyr Asn His Ala
85 90 95

Asp Ala Glu Phe Gln Glu Arg Leu Gln Glu Asp Thr Ser Phe Gly Gly
100 105 110

Asn Leu Gly Arg Ala Val Phe Gln Ala Lys Lys Arg Ile Leu Glu Pro
115 120 125

Leu Gly Leu Val Glu Glu Ala Ala Lys Thr Ala Pro Gly Lys Lys Arg
130 135 140

Pro Val Asp Gln Ser Pro Gln Glu Pro Asp Ser Ser Ser Gly Val Gly
145 150 155 160

Lys Ser Gly Lys Gln Pro Ala Arg Lys Arg Leu Asn Phe Gly Gln Thr
165 170 175

Gly Asp Ser Glu Ser Val Pro Asp Pro Gln Pro Leu Gly Glu Pro Pro
180 185 190

Ala Ala Pro Thr Ser Leu Gly Ser Asn Thr Met Ala Ser Gly Gly Gly
195 200 205

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Ala Pro Met Ala Asp Asn Asn Glu Gly Ala Asp Gly Val Gly Asn Ser
    210                 215                 220

Ser Gly Asn Trp His Cys Asp Ser Gln Trp Leu Gly Asp Arg Val Ile
225                 230                 235                 240

Thr Thr Ser Thr Arg Thr Trp Ala Leu Pro Thr Tyr Asn Asn His Leu
                245                 250                 255

Tyr Lys Gln Ile Ser Ser Ala Ser Thr Gly Ala Ser Asn Asp Asn His
            260                 265                 270

Tyr Phe Gly Tyr Ser Thr Pro Trp Gly Tyr Phe Asp Phe Asn Arg Phe
        275                 280                 285

His Cys His Phe Ser Pro Arg Asp Trp Gln Arg Leu Ile Asn Asn Asn
    290                 295                 300

Trp Gly Phe Arg Pro Lys Lys Leu Ser Phe Lys Leu Phe Asn Ile Gln
305                 310                 315                 320

Val Lys Glu Val Thr Gln Asn Asp Gly Thr Thr Thr Ile Ala Asn Asn
                325                 330                 335

Leu Thr Ser Thr Val Gln Val Phe Thr Asp Ser Glu Tyr Gln Leu Pro
            340                 345                 350

Tyr Val Leu Gly Ser Ala His Gln Gly Cys Leu Pro Pro Phe Pro Ala
        355                 360                 365

Asp Val Phe Met Val Pro Gln Tyr Gly Tyr Leu Thr Leu Asn Asn Gly
    370                 375                 380

Ser Gln Ala Val Gly Arg Ser Ser Phe Tyr Cys Leu Glu Tyr Phe Pro
385                 390                 395                 400

Ser Gln Met Leu Arg Thr Gly Asn Asn Phe Gln Phe Ser Tyr Thr Phe
                405                 410                 415

Glu Asp Val Pro Phe His Ser Ser Tyr Ala His Ser Gln Ser Leu Asp
            420                 425                 430

Arg Leu Met Asn Pro Leu Ile Asp Gln Tyr Leu Tyr Tyr Leu Asn Arg
        435                 440                 445

Thr Gln Gly Thr Thr Ser Gly Thr Thr Asn Gln Ser Arg Leu Leu Phe
    450                 455                 460

Ser Gln Ala Gly Pro Gln Ser Met Ser Leu Gln Ala Arg Asn Trp Leu
465                 470                 475                 480

Pro Gly Pro Cys Tyr Arg Gln Gln Arg Leu Ser Lys Thr Ala Asn Asp
                485                 490                 495

Asn Asn Asn Ser Asn Phe Pro Trp Thr Ala Ala Ser Lys Tyr His Leu
            500                 505                 510

Asn Gly Arg Asp Ser Leu Val Asn Pro Gly Pro Ala Met Ala Ser His
        515                 520                 525

Lys Asp Asp Glu Glu Lys Phe Phe Pro Met His Gly Asn Leu Ile Phe
    530                 535                 540

Gly Lys Glu Gly Thr Thr Ala Ser Asn Ala Glu Leu Asp Asn Val Met
545                 550                 555                 560

Ile Thr Asp Glu Glu Glu Ile Arg Thr Thr Asn Pro Val Ala Thr Glu
                565                 570                 575

Gln Tyr Gly Thr Val Ala Asn Asn Leu Gln Ser Ser Asn Thr Ala Pro
            580                 585                 590

Thr Thr Arg Thr Val Asn Asp Gln Gly Ala Leu Pro Gly Met Val Trp
        595                 600                 605

Gln Asp Arg Asp Val Tyr Leu Gln Gly Pro Ile Trp Ala Lys Ile Pro
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    610                 615                 620

His Thr Asp Gly His Phe His Pro Ser Pro Leu Met Gly Gly Phe Gly
625                 630                 635                 640

Leu Lys His Pro Pro Pro Gln Ile Met Ile Lys Asn Thr Pro Val Pro
                645                 650                 655

Ala Asn Pro Pro Thr Thr Phe Ser Pro Ala Lys Phe Ala Ser Phe Ile
            660                 665                 670

Thr Gln Tyr Ser Thr Gly Gln Val Ser Val Glu Ile Glu Trp Glu Leu
        675                 680                 685

Gln Lys Glu Asn Ser Lys Arg Trp Asn Pro Glu Ile Gln Tyr Thr Ser
    690                 695                 700

Asn Tyr Asn Lys Ser Val Asn Val Asp Phe Thr Val Asp Thr Asn Gly
705                 710                 715                 720

Val Tyr Ser Glu Pro Arg Pro Ile Gly Thr Arg Tyr Leu Thr Arg Asn
                725                 730                 735

Leu


<210> SEQ ID NO 438
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 438

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Arg Lys Asp Ala Leu Arg Gly His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

His Arg Thr Thr Leu Thr Asn His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Asn Ala
            100                 105                 110

Leu Ala Gly His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser His Lys Asn Ala Leu Gln Asn His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Asp Pro Gly His Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Thr Ser Gly Asn Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
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    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Glu Arg Tyr Val Glu Glu Asn Gln
305                 310                 315                 320

Thr Arg Asp Lys His Leu Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn His Ile Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400

Asn Gly Glu Ile Asn Phe
                405


<210> SEQ ID NO 439
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 439

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Gln Gln Arg Ser Leu Val Gly His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

Asp Lys Lys Asp Leu Thr Arg His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Thr Ser Gly His
            100                 105                 110

Leu Val Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Ala His Leu Glu Arg His
    130                 135                 140
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Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Thr Ser Gly Ser Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Thr Ser Gly Asn Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Gln Arg Tyr Val Lys Glu Asn Gln
305                 310                 315                 320

Thr Arg Asn Lys His Ile Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400

Asn Gly Glu Ile Asn Phe
                405


<210> SEQ ID NO 440
<211> LENGTH: 320
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 440

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
        35                  40                  45

Arg Asn Phe Ser Ser Lys Gln Ala Leu Ala Val His Thr Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65                  70                  75                  80
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Gln Ser Thr Thr Leu Lys Arg His Leu Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser Arg Ser Asp His
            100                 105                 110

Leu Ser Leu His Leu Lys Thr His Leu Arg Gly Ser Gln Leu Val Lys
        115                 120                 125

Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu Lys Tyr
    130                 135                 140

Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr
145                 150                 155                 160

Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met Lys Val
                165                 170                 175

Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro Asp Gly
            180                 185                 190

Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile Val Asp
        195                 200                 205

Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp
    210                 215                 220

Glu Met Gln Arg Tyr Val Lys Glu Asn Gln Thr Arg Asn Lys His Ile
225                 230                 235                 240

Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr Glu Phe
                245                 250                 255

Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys Ala Gln
            260                 265                 270

Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys Asn Gly Ala Val Leu Ser
        275                 280                 285

Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly Thr Leu
    290                 295                 300

Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
305                 310                 315                 320


<210> SEQ ID NO 441
<211> LENGTH: 320
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 441

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
        35                  40                  45

Arg Asn Phe Ser Arg Arg Ala His Leu Gln Asn His Thr Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65                  70                  75                  80

Gln Ser Thr Thr Leu Lys Arg His Leu Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser Asp Gly Gly His
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            100                 105                 110

Leu Thr Arg His Leu Lys Thr His Leu Arg Gly Ser Gln Leu Val Lys
        115                 120                 125

Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu Lys Tyr
    130                 135                 140

Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr
145                 150                 155                 160

Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met Lys Val
                165                 170                 175

Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro Asp Gly
            180                 185                 190

Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile Val Asp
        195                 200                 205

Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp
    210                 215                 220

Glu Met Glu Arg Tyr Val Glu Glu Asn Gln Thr Arg Asp Lys His Leu
225                 230                 235                 240

Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr Glu Phe
                245                 250                 255

Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys Ala Gln
            260                 265                 270

Leu Thr Arg Leu Asn His Ile Thr Asn Cys Asn Gly Ala Val Leu Ser
        275                 280                 285

Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly Thr Leu
    290                 295                 300

Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
305                 310                 315                 320


<210> SEQ ID NO 442
<211> LENGTH: 42
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      oligonucleotide

<400> SEQUENCE: 442

caaactagaa atgccatctt ccttgatgtt ggaggtacct gc                        42


<210> SEQ ID NO 443
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 443

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Arg Lys Asp Ala Leu Arg Gly His Gln Arg Thr His
    50                  55                  60
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```
Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

His Arg Thr Thr Leu Thr Asn His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Asn Ala
            100                 105                 110

Leu Ala Gly His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser His Lys Asn Ala Leu Gln Asn His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Asp Pro Gly His Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Thr Ser Gly Asn Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Glu Arg Tyr Val Glu Glu Asn Gln
305                 310                 315                 320

Thr Arg Asp Lys His Leu Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn His Ile Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400

Asn Gly Glu Ile Asn Phe
                405


<210> SEQ ID NO 444
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide
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<400> SEQUENCE: 444

```
Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                  10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Gln Gln Arg Ser Leu Val Gly His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

Asp Lys Lys Asp Leu Thr Arg His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Thr Ser Gly His
            100                 105                 110

Leu Val Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Ala His Leu Glu Arg His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Thr Ser Gly Ser Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Thr Ser Gly Asn Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Gln Arg Tyr Val Lys Glu Asn Gln
305                 310                 315                 320

Thr Arg Asn Lys His Ile Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400
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Asn Gly Glu Ile Asn Phe
405

<210> SEQ ID NO 445
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (1)..(1)
<223> OTHER INFORMATION: May or may not be present
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (25)..(25)
<223> OTHER INFORMATION: May or may not be present

<400> SEQUENCE: 445

tatctgccca tgactggcgc aggga 25

<210> SEQ ID NO 446
<211> LENGTH: 320
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polypeptide

<400> SEQUENCE: 446

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1 5 10 15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
20 25 30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
35 40 45

Arg Asn Phe Ser Ser Lys Gln Ala Leu Ala Val His Thr Arg Thr His
50 55 60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65 70 75 80

Gln Ser Thr Thr Leu Lys Arg His Leu Arg Thr His Thr Gly Glu Lys
85 90 95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser Arg Ser Asp His
100 105 110

Leu Ser Leu His Leu Lys Thr His Leu Arg Gly Ser Gln Leu Val Lys
115 120 125

Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu Lys Tyr
130 135 140

Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr
145 150 155 160

Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met Lys Val
165 170 175

Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro Asp Gly
180 185 190

Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile Val Asp
195 200 205

Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp

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    210                 215                 220

Glu Met Gln Arg Tyr Val Lys Glu Asn Gln Thr Arg Asn Lys His Ile
225                 230                 235                 240

Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr Glu Phe
                245                 250                 255

Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys Ala Gln
            260                 265                 270

Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys Asn Gly Ala Val Leu Ser
        275                 280                 285

Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly Thr Leu
    290                 295                 300

Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
305                 310                 315                 320


<210> SEQ ID NO 447
<211> LENGTH: 320
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 447

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
        35                  40                  45

Arg Asn Phe Ser Arg Arg Ala His Leu Gln Asn His Thr Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65                  70                  75                  80

Gln Ser Thr Thr Leu Lys Arg His Leu Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser Asp Gly Gly His
            100                 105                 110

Leu Thr Arg His Leu Lys Thr His Leu Arg Gly Ser Gln Leu Val Lys
        115                 120                 125

Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu Lys Tyr
    130                 135                 140

Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr
145                 150                 155                 160

Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met Lys Val
                165                 170                 175

Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro Asp Gly
            180                 185                 190

Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile Val Asp
        195                 200                 205

Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp
    210                 215                 220

Glu Met Glu Arg Tyr Val Glu Glu Asn Gln Thr Arg Asp Lys His Leu
225                 230                 235                 240
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Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr Glu Phe
                245                 250                 255

Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys Ala Gln
            260                 265                 270

Leu Thr Arg Leu Asn His Ile Thr Asn Cys Asn Gly Ala Val Leu Ser
        275                 280                 285

Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly Thr Leu
    290                 295                 300

Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
305                 310                 315                 320


<210> SEQ ID NO 448
<211> LENGTH: 42
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      oligonucleotide

<400> SEQUENCE: 448

actagaaatg ccatcttcct tgatgttgga ggtacctgct ct                      42


<210> SEQ ID NO 449
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 449

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser His Arg Thr Thr Leu Thr Asn His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

Gln Arg Asn Ala Leu Ala Gly His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser His Lys Asn Ala
            100                 105                 110

Leu Gln Asn His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser Asp Pro Gly His Leu Val Arg His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Thr Ser Gly Asn Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Gln Ser Ser Asn Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
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        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Glu Arg Tyr Val Glu Glu Asn Gln
305                 310                 315                 320

Thr Arg Asp Lys His Leu Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn His Ile Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400

Asn Gly Glu Ile Asn Phe
                405


<210> SEQ ID NO 450
<211> LENGTH: 406
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 450

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Gln Arg Asn Ala Leu Ala Gly His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

Gln Gln Arg Ser Leu Val Gly His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Asp Lys Lys Asp
            100                 105                 110

Leu Thr Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125
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Pro Glu Cys Gly Lys Ser Phe Ser Thr Ser Gly His Leu Val Arg His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Gln Arg Ala His Leu Glu Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
            180                 185                 190

Thr Ser Gly Ser Leu Val Arg His Gln Arg Thr His Thr Gly Ala Ala
        195                 200                 205

Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu
    210                 215                 220

Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu
225                 230                 235                 240

Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met
                245                 250                 255

Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly
            260                 265                 270

Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp
        275                 280                 285

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

Pro Ile Gly Gln Ala Asp Glu Met Gln Arg Tyr Val Lys Glu Asn Gln
305                 310                 315                 320

Thr Arg Asn Lys His Ile Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro
                325                 330                 335

Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys
            340                 345                 350

Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys
        355                 360                 365

Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met
    370                 375                 380

Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn
385                 390                 395                 400

Asn Gly Glu Ile Asn Phe
                405


<210> SEQ ID NO 451
<211> LENGTH: 378
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 451

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Gln Arg Asn Ala Leu Ala Gly His Gln Arg Thr His
    50                  55                  60
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Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

His Lys Asn Ala Leu Gln Asn His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Asp Pro Gly His
            100                 105                 110

Leu Val Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser Thr Ser Gly Asn Leu Val Arg His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Gln Ser Ser Asn Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Ala Ala Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys
            180                 185                 190

Lys Ser Glu Leu Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile
        195                 200                 205

Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu
    210                 215                 220

Met Lys Val Met Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu
225                 230                 235                 240

His Leu Gly Gly Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly
                245                 250                 255

Ser Pro Ile Asp Tyr Gly Val Ile Val Asp Thr Lys Ala Tyr Ser Gly
            260                 265                 270

Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp Glu Met Glu Arg Tyr Val
        275                 280                 285

Glu Glu Asn Gln Thr Arg Asp Lys His Leu Asn Pro Asn Glu Trp Trp
    290                 295                 300

Lys Val Tyr Pro Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser
305                 310                 315                 320

Gly His Phe Lys Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn His
                325                 330                 335

Ile Thr Asn Cys Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile
            340                 345                 350

Gly Gly Glu Met Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg
        355                 360                 365

Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
    370                 375


<210> SEQ ID NO 452
<211> LENGTH: 378
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 452

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
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            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
        35                  40                  45

Lys Ser Phe Ser Gln Gln Arg Ser Leu Val Gly His Gln Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser
65                  70                  75                  80

Asp Lys Lys Asp Leu Thr Arg His Gln Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Tyr Lys Cys Pro Glu Cys Gly Lys Ser Phe Ser Thr Ser Gly His
            100                 105                 110

Leu Val Arg His Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys
        115                 120                 125

Pro Glu Cys Gly Lys Ser Phe Ser Gln Arg Ala His Leu Glu Arg His
    130                 135                 140

Gln Arg Thr His Thr Gly Glu Lys Pro Tyr Lys Cys Pro Glu Cys Gly
145                 150                 155                 160

Lys Ser Phe Ser Thr Ser Gly Ser Leu Val Arg His Gln Arg Thr His
                165                 170                 175

Thr Gly Ala Ala Ala Arg Ala Leu Val Lys Ser Glu Leu Glu Glu Lys
            180                 185                 190

Lys Ser Glu Leu Arg His Lys Leu Lys Tyr Val Pro His Glu Tyr Ile
        195                 200                 205

Glu Leu Ile Glu Ile Ala Arg Asn Pro Thr Gln Asp Arg Ile Leu Glu
    210                 215                 220

Met Lys Val Met Glu Phe Phe Met Lys Val Tyr Gly Tyr Arg Gly Glu
225                 230                 235                 240

His Leu Gly Gly Ser Arg Lys Pro Asp Gly Ala Ile Tyr Thr Val Gly
                245                 250                 255

Ser Pro Ile Asp Tyr Gly Val Ile Val Asp Thr Lys Ala Tyr Ser Gly
            260                 265                 270

Gly Tyr Asn Leu Pro Ile Gly Gln Ala Asp Glu Met Gln Arg Tyr Val
        275                 280                 285

Lys Glu Asn Gln Thr Arg Asn Lys His Ile Asn Pro Asn Glu Trp Trp
    290                 295                 300

Lys Val Tyr Pro Ser Ser Val Thr Glu Phe Lys Phe Leu Phe Val Ser
305                 310                 315                 320

Gly His Phe Lys Gly Asn Tyr Lys Ala Gln Leu Thr Arg Leu Asn Arg
                325                 330                 335

Lys Thr Asn Cys Asn Gly Ala Val Leu Ser Val Glu Glu Leu Leu Ile
            340                 345                 350

Gly Gly Glu Met Ile Lys Ala Gly Thr Leu Thr Leu Glu Glu Val Arg
        355                 360                 365

Arg Lys Phe Asn Asn Gly Glu Ile Asn Phe
    370                 375


<210> SEQ ID NO 453
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      oligonucleotide
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<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (1)..(1)
<223> OTHER INFORMATION: May or may not be present
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (26)..(26)
<223> OTHER INFORMATION: May or may not be present

<400> SEQUENCE: 453

tgccatcttc cttgatgttg gaggta 26

<210> SEQ ID NO 454
<211> LENGTH: 322
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polypeptide

<400> SEQUENCE: 454

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1 5 10 15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
20 25 30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
35 40 45

Arg Asn Phe Ser Ser Pro Ser Lys Leu Ala Arg His Thr Arg Thr His
50 55 60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65 70 75 80

Val Arg His Asn Leu Thr Arg His Leu Arg Thr His Thr Gly Glu Lys
85 90 95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser Gln Arg Asn Asn
100 105 110

Leu Gly Arg His Leu Lys Thr His Thr Gly Ala Ala Ala Arg Ala Leu
115 120 125

Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu
130 135 140

Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn
145 150 155 160

Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met
165 170 175

Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro
180 185 190

Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile
195 200 205

Val Asp Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln
210 215 220

Ala Asp Glu Met Gln Arg Tyr Val Lys Glu Asn Gln Thr Arg Asn Lys
225 230 235 240

His Ile Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr
245 250 255

Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys
260 265 270

Ala Gln Leu Thr Arg Leu Asn Arg Lys Thr Asn Cys Asn Gly Ala Val

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        275                 280                 285

Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly
    290                 295                 300

Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile
305                 310                 315                 320

Asn Phe


<210> SEQ ID NO 455
<211> LENGTH: 322
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 455

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Leu Glu Pro Gly Glu Arg Pro Phe Gln Cys Arg Ile Cys Met
        35                  40                  45

Arg Asn Phe Ser Ile Pro Asn His Leu Ala Arg His Thr Arg Thr His
    50                  55                  60

Thr Gly Glu Lys Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser
65                  70                  75                  80

Gln Ser Ala His Leu Lys Arg His Leu Arg Thr His Thr Gly Glu Lys
                85                  90                  95

Pro Phe Gln Cys Arg Ile Cys Met Arg Asn Phe Ser His His Asn Ser
            100                 105                 110

Leu Thr Arg His Leu Lys Thr His Thr Gly Ala Ala Ala Arg Ala Leu
        115                 120                 125

Val Lys Ser Glu Leu Glu Glu Lys Lys Ser Glu Leu Arg His Lys Leu
    130                 135                 140

Lys Tyr Val Pro His Glu Tyr Ile Glu Leu Ile Glu Ile Ala Arg Asn
145                 150                 155                 160

Pro Thr Gln Asp Arg Ile Leu Glu Met Lys Val Met Glu Phe Phe Met
                165                 170                 175

Lys Val Tyr Gly Tyr Arg Gly Glu His Leu Gly Gly Ser Arg Lys Pro
            180                 185                 190

Asp Gly Ala Ile Tyr Thr Val Gly Ser Pro Ile Asp Tyr Gly Val Ile
        195                 200                 205

Val Asp Thr Lys Ala Tyr Ser Gly Gly Tyr Asn Leu Pro Ile Gly Gln
    210                 215                 220

Ala Asp Glu Met Glu Arg Tyr Val Glu Glu Asn Gln Thr Arg Asp Lys
225                 230                 235                 240

His Leu Asn Pro Asn Glu Trp Trp Lys Val Tyr Pro Ser Ser Val Thr
                245                 250                 255

Glu Phe Lys Phe Leu Phe Val Ser Gly His Phe Lys Gly Asn Tyr Lys
            260                 265                 270

Ala Gln Leu Thr Arg Leu Asn His Ile Thr Asn Cys Asn Gly Ala Val
        275                 280                 285

Leu Ser Val Glu Glu Leu Leu Ile Gly Gly Glu Met Ile Lys Ala Gly
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    290                 295                 300

Thr Leu Thr Leu Glu Glu Val Arg Arg Lys Phe Asn Asn Gly Glu Ile
305                 310                 315                 320

Asn Phe


<210> SEQ ID NO 456
<211> LENGTH: 6626
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polynucleotide

<400> SEQUENCE: 456

gggggggggg gggggggttt ggccactccc tctctgcgcg ctcgctcgct cactgaggcc     60

gggcgaccaa aggtcgcccg acgcccgggc tttgcccggg cggcctcagt gagcgagcga    120

gcgcgcagag agggagtggc caactccatc actaggggtt cctagatctg aattcggtac    180

ccgttacata acttacggta aatggcccgc ctggctgacc gcccaacgac ccccgcccat    240

tgacgtcaat aatgacgtat gttcccatag taacgccaat agggactttc cattgacgtc    300

aatgggtgga gtatttacgg taaactgccc acttggcagt acatcaagtg tatcatatgc    360

caagtacgcc ccctattgac gtcaatgacg gtaaatggcc cgcctggcat tatgcccagt    420

acatgacctt atgggacttt cctacttggc agtacatcta cgtattagtc atcgctatta    480

ccatggtgat gcggttttgg cagtacatca atgggcgtgg atagcggttt gactcacggg    540

gatttccaag tctccacccc attgacgtca atgggagttt gttttggcac caaaatcaac    600

gggactttcc aaaatgtcgt aacaactccg ccccattgac gcaaatgggc ggtaggcgtg    660

tacggtggga ggtctatata agcagagctc gtttagtgaa ccgtcagatc gcctggagac    720

gccatccacg ctgttttgac ctccatagaa gacaccggga ccgatccagc ctccggactc    780

tagaggatcc ggtactcgag gaactgaaaa accagaaagt taactggtaa gtttagtctt    840

tttgtctttt atttcaggtc ccggatccgg tggtggtgca aatcaaagaa ctgctcctca    900

gtggatgttg cctttacttc taggcctgta cggaagtgtt acttctgctc taaaagctgc    960

ggaattgtac ccgcggcccg ggatccaccg gtggctagcg tctataggcc cacccccttg   1020

gtggaattcg ccatgaggtc tgactacaaa gaccatgacg gtgattataa agatcatgac   1080

atcgattaca aggatgacga tgacaagatg gcccccaaga agaagaggaa ggtgggcctc   1140

gagcccggag aaaaaccgta caagtgccct gagtgcggga aatcattctc cgaccctggg   1200

gcgctcgtcc ggcaccaaag gacgcataca ggggaaaagc cgtataagtg ccccgagtgt   1260

ggaaagagct tctcgcagag agcccacctt gaacgacacc aaagaacaca cactggtgag   1320

aaaccctata agtgtccaga gtgcggcaaa tcgtttagca gatccgatga cttggtgcgc   1380

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<210> SEQ ID NO 457
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 457

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<210> SEQ ID NO 458
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 458

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<210> SEQ ID NO 459
<211> LENGTH: 3252
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 459

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attgactatg aaacaaggga cgtgatcgac gcaggcgtca gactgttcaa ggaggccaac 120

gtggaaaaca atgagggacg gagaagcaag aggggagcca ggcgcctgaa acgacggaga 180

aggcacagaa tccagagggt gaagaaactg ctgttcgatt acaacctgct gaccgaccat 240

tctgagctga gtggaattaa tccttatgaa gccagggtga aaggcctgag tcagaagctg 300

tcagaggaag agttttccgc agctctgctg cacctggcta agcgccgagg agtgcataac 360

gtcaatgagg tggaagagga caccggcaac gagctgtcta caaaggaaca gatctcacgc 420

aatagcaaag ctctggaaga gaagtatgtc gcagagctgc agctggaacg gctgaagaaa 480

gatggcgagg tgagagggtc aattaatagg ttcaagacaa gcgactacgt caaagaagcc 540

aagcagctgc tgaaagtgca gaaggcttac caccagctgg atcagagctt catcgatact 600

tatatcgacc tgctggagac tcggagaacc tactatgagg gaccaggaga agggagcccc 660

ttcggatgga aagacatcaa ggaatggtac gagatgctga tgggacattg cacctatttt 720

ccagaagagc tgagaagcgt caagtacgct tataacgcag atctgtacaa cgccctgaat 780

gacctgaaca acctggtcat caccagggat gaaaacgaga aactggaata ctatgagaag 840

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gctactaaga aagctggtca agctaagaaa aagaaaggat cctacccata cgatgttcca 3240

gattacgctt aa 3252

<210> SEQ ID NO 460
<211> LENGTH: 83
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 460

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caacttgttg gcgagatttt ttt 83

<210> SEQ ID NO 461
<211> LENGTH: 3333
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 461

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gaaaacccca tccgcctgat tgatttgggc gtgcgcgtat ttgagcgtgc cgaagtaccg 180

aaaacaggcg actcccttgc catggcaagg cgtttggcgc gcagtgttcg ccgcctgacc 240

cgccgtcgcg cccaccgcct gcttcggacc cgccgcctat tgaaacgcga aggcgtatta 300

caagccgcca attttgacga aaacggcttg attaaatcct taccgaatac accatggcaa 360

cttcgcgcag ccgcattaga ccgcaaactg acgcctttag agtggtcggc agtcttgttg 420

catttaatca aacatcgcgg ctatttatcg caacggaaaa acgagggcga aactgccgat 480

aaggagcttg gcgctttgct taaaggcgta gccggcaatg cccatgcctt acagacaggc 540

gatttccgca caccggccga attggcttta aataaatttg agaaagaaag cggccatatc 600

cgcaatcagc gcagcgatta ttcgcatacg ttcagccgca aagatttaca ggcggagctg 660

attttgctgt ttgaaaaaca aaaagaattt ggcaatccgc atgtttcagg cggccttaaa 720

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gctccagcag cgaagaaaaa gaagctggat taa 3333

<210> SEQ ID NO 462
<211> LENGTH: 120
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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polynucleotide

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<210> SEQ ID NO 463
<211> LENGTH: 144
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 463

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gaaccgttgc tacaataagg ccgtctgaaa agatgtgccg caacgctctg ccccttaaag 120

cttctgcttt aaggggcttt tttt 144

<210> SEQ ID NO 464
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 464

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<210> SEQ ID NO 465
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 465

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<210> SEQ ID NO 466
<211> LENGTH: 4564
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 466

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tccattgggc tcgccatcgg cacaaacagc gtcggctggg ccgtcattac ggacgagtac 180

aaggtgccga gcaaaaaatt caaagttctg ggcaataccg atcgccacag cataaagaag 240

aacctcattg gcgccctcct gttcgactcc ggggaaaccg ccgaagccac gcggctcaaa 300

agaacagcac ggcgcagata tacccgcaga aagaatcgga tctgctacct gcaggagatc 360

tttagtaatg agatggctaa ggtggatgac tctttcttcc ataggctgga ggagtccttt 420

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gtggcgtacc atgaaaagta cccaaccata tatcatctga ggaagaagct tgtagacagt 540

actgataagg ctgacttgcg gttgatctat ctcgcgctgg cgcatatgat caaatttcgg 600

ggacacttcc tcatcgaggg ggacctgaac ccagacaaca gcgatgtcga caaactcttt 660

atccaactgg ttcagactta caatcagctt ttcgaagaga acccgatcaa cgcatccgga 720

gttgacgcca aagcaatcct gagcgctagg ctgtccaaat cccggcggct cgaaaacctc 780

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gaagactatt tcaaaaagat tgaatgtttc gactctgttg aaatcagcgg agtggaggat 1860

cgcttcaacg catccctggg aacgtatcac gatctcctga aaatcattaa agacaaggac 1920

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tttgaagata gggagatgat tgaagaacgc ttgaaaactt acgctcatct cttcgacgac 2040

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ggtggcctgt ctgagttgga taaagccggc ttcatcaaaa ggcagcttgt tgagacacgc 2880

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ggtggagaca gcagggctga ccccaagaag aagaggaagg tggctagcga tgctaagtca 4260

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ttggagaagg gagaagagcc ctggctggtg gagagagaaa ttcaccaaga gacccatcct 4500

gattcagaga ctgcatttga aatcaaatca tcagttccga aaaagaaacg caaagttgct 4560

agcg 4564

<210> SEQ ID NO 467
<211> LENGTH: 388
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 467

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ccttctggtt ctgtgtaacc ttctaagcaa accttctggc tcaagcacag caatgctgag 180

tcatgatgag tcatgctgag gcttagggtg tgtgcccaga tgttctcagc ctagagtgat 240

gactcctatc tgggtcccca gcaggatgct tacagggcag atggcaaaaa aaaggagaag 300

ctgaccacct gactaaaact ccacctcaaa cggcatcata aagaaaatgg atgcctgaga 360

cagaatgtga catattctag aatatatt 388

<210> SEQ ID NO 468
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 468

cagccgctcg ctgcagcag 19

<210> SEQ ID NO 469
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 469

ctgctgcagc gagcggctg 19

<210> SEQ ID NO 470
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 470

gctgggtgtc ccattgaaa 19

<210> SEQ ID NO 471
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 471

tttcaatggg acacccagc 19

<210> SEQ ID NO 472
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 472

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gtttattcag ccgggagtc 19

<210> SEQ ID NO 473
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 473

gactcccggc tgaataaac 19

<210> SEQ ID NO 474
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 474

tggagagttt gcaaggagc 19

<210> SEQ ID NO 475
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 475

gctccttgca aactctcca 19

<210> SEQ ID NO 476
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 476

ccctccagac tttccacct 19

<210> SEQ ID NO 477
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 477

aggtggaaag tctggaggg 19

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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 478

aattttcttc caagttctc 19

<210> SEQ ID NO 479
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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 479

gagaacttgg aagaaaatt 19

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 480

ctgcggagag aagaaaggg 19

<210> SEQ ID NO 481
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 481

ccctttcttc tctccgcag 19

<210> SEQ ID NO 482
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 482

agagccaccc cctggctcc 19

<210> SEQ ID NO 483
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 483

ggagccaggg ggtggctct 19

<210> SEQ ID NO 484
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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<400> SEQUENCE: 484

cgaagccaac cgcggcggg 19

<210> SEQ ID NO 485
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 485

cccgccgcgg ttggcttcg 19

<210> SEQ ID NO 486
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 486

agagggaaga cgatcgccc 19

<210> SEQ ID NO 487
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 487

gggcgatcgt cttccctct 19

<210> SEQ ID NO 488
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 488

ccccttaac tttcctccg 19

<210> SEQ ID NO 489
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 489

cggaggaaag ttaaagggg 19

<210> SEQ ID NO 490
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 490

gcagccccgc ttccttcaa 19

<210> SEQ ID NO 491
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 491

ttgaaggaag cggggctgc 19

<210> SEQ ID NO 492
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 492

caccgcagcc gctcgctgca gcag 24

<210> SEQ ID NO 493
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 493

aaacctgctg cagcgagcgg ctgc 24

<210> SEQ ID NO 494
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 494

caccggctgg gtgtcccatt gaaa 24

<210> SEQ ID NO 495
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 495

aaactttcaa tgggacaccc agcc 24

<210> SEQ ID NO 496
<211> LENGTH: 24

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 496

caccggttta ttcagccggg agtc 24

<210> SEQ ID NO 497
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 497

aaacgactcc cggctgaata aacc 24

<210> SEQ ID NO 498
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 498

caccgtggag agtttgcaag gagc 24

<210> SEQ ID NO 499
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 499

aaacgctcct tgcaaactct ccac 24

<210> SEQ ID NO 500
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 500

caccgccctc cagactttcc acct 24

<210> SEQ ID NO 501
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 501

aaacaggtgg aaagtctgga gggc 24

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<210> SEQ ID NO 502
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 502

caccgaattt tcttccaagt tctc 24

<210> SEQ ID NO 503
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 503

aaacgagaac ttggaagaaa attc 24

<210> SEQ ID NO 504
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 504

caccgctgcg gagagaagaa aggg 24

<210> SEQ ID NO 505
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 505

aaaccccttt cttctctccg cagc 24

<210> SEQ ID NO 506
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 506

caccgagagc caccccctgg ctcc 24

<210> SEQ ID NO 507
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 507

aaacggagcc agggggtggc tctc 24

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<210> SEQ ID NO 508
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 508

caccgcgaag ccaaccgcgg cggg 24

<210> SEQ ID NO 509
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 509

aaaccccgcc gcggttggct tcgc 24

<210> SEQ ID NO 510
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 510

caccgagagg gaagacgatc gccc 24

<210> SEQ ID NO 511
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 511

aaacgggcga tcgtcttccc tctc 24

<210> SEQ ID NO 512
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 512

caccgcccct ttaactttcc tccg 24

<210> SEQ ID NO 513
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 513

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aaaccggagg aaagttaaag gggc 24

<210> SEQ ID NO 514
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 514

caccggcagc cccgcttcct tcaa 24

<210> SEQ ID NO 515
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 515

aaacttgaag gaagcggggc tgcc 24

<210> SEQ ID NO 516
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 516

cgagagcgag aggagggag 19

<210> SEQ ID NO 517
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 517

ctccctcctc tcgctctcg 19

<210> SEQ ID NO 518
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 518

gagagagctt gagagcgcg 19

<210> SEQ ID NO 519
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 519

cgcgctctca agctctctc 19

<210> SEQ ID NO 520
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 520

ggtggagggg gcggggccc 19

<210> SEQ ID NO 521
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 521

gggccccgcc ccctccacc 19

<210> SEQ ID NO 522
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 522

ggtatccacg taaatcaaa 19

<210> SEQ ID NO 523
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 523

tttgatttac gtggatacc 19

<210> SEQ ID NO 524
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 524

ccaatcactg gctccggtc 19

<210> SEQ ID NO 525
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 525

gaccggagcc agtgattgg 19

<210> SEQ ID NO 526
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 526

ggcgcccgag ggaagaaga 19

<210> SEQ ID NO 527
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 527

tcttcttccc tcgggcgcc 19

<210> SEQ ID NO 528
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 528

gggtgggggt accagagga 19

<210> SEQ ID NO 529
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 529

tcctctggta cccccaccc 19

<210> SEQ ID NO 530
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 530

ccggggacag aagagaggg 19

<210> SEQ ID NO 531
<211> LENGTH: 19
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 531

ccctctcttc tgtccccgg 19

<210> SEQ ID NO 532
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 532

gagagagagt gggagaagc 19

<210> SEQ ID NO 533
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 533

gcttctccca ctctctctc 19

<210> SEQ ID NO 534
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 534

aaagtaactg tcaaatgcg 19

<210> SEQ ID NO 535
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 535

cgcatttgac agttacttt 19

<210> SEQ ID NO 536
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 536

ttaaccagag cgcccagtc 19

<210> SEQ ID NO 537

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<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 537

gactgggcgc tctggttaa 19

<210> SEQ ID NO 538
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 538

cgtcggagct gcccgctag 19

<210> SEQ ID NO 539
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 539

ctagcgggca gctccgacg 19

<210> SEQ ID NO 540
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 540

caccgcgaga gcgagaggag ggag 24

<210> SEQ ID NO 541
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 541

aaacctccct cctctcgctc tcgc 24

<210> SEQ ID NO 542
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 542

caccggagag agcttgagag cgcg 24

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<210> SEQ ID NO 543
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 543

aaaccgcgct ctcaagctct ctcc 24

<210> SEQ ID NO 544
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 544

caccgggtgg agggggcggg gccc 24

<210> SEQ ID NO 545
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 545

aaacgggccc cgccccctcc accc 24

<210> SEQ ID NO 546
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 546

caccgggtat ccacgtaaat caaa 24

<210> SEQ ID NO 547
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 547

aaactttgat ttacgtggat accc 24

<210> SEQ ID NO 548
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 548

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caccgccaat cactggctcc ggtc 24

<210> SEQ ID NO 549
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 549

aaacgaccgg agccagtgat tggc 24

<210> SEQ ID NO 550
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 550

caccgggcgc ccgagggaag aaga 24

<210> SEQ ID NO 551
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 551

aaactcttct tccctcgggc gccc 24

<210> SEQ ID NO 552
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 552

caccggggtg ggggtaccag agga 24

<210> SEQ ID NO 553
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 553

aaactcctct ggtaccccca cccc 24

<210> SEQ ID NO 554
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 554

caccgccggg gacagaagag aggg 24

<210> SEQ ID NO 555
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 555

aaacccctct cttctgtccc cggc 24

<210> SEQ ID NO 556
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 556

caccggagag agagtgggag aagc 24

<210> SEQ ID NO 557
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 557

aaacgcttct cccactctct ctcc 24

<210> SEQ ID NO 558
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 558

caccgaaagt aactgtcaaa tgcg 24

<210> SEQ ID NO 559
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 559

aaaccgcatt tgacagttac tttc 24

<210> SEQ ID NO 560
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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oligonucleotide

<400> SEQUENCE: 560

caccgttaac cagagcgccc agtc 24

<210> SEQ ID NO 561
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 561

aaacgactgg gcgctctggt taac 24

<210> SEQ ID NO 562
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 562

caccgcgtcg gagctgcccg ctag 24

<210> SEQ ID NO 563
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 563

aaacctagcg ggcagctccg acgc 24

<210> SEQ ID NO 564
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 564

gagacacaca gaaatgtaac 20

<210> SEQ ID NO 565
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 565

ggtggggcac tgaccccgac 20

<210> SEQ ID NO 566
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 566

ctagagtgat gactcctatc 20

<210> SEQ ID NO 567
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 567

gactaaaact ccacctcaaa 20

<210> SEQ ID NO 568
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 568

aatatgtcac attctgtctc 20

<210> SEQ ID NO 569
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 569

ggactatggg aggtcactaa 20

<210> SEQ ID NO 570
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 570

gctcatgctt ggactatggg 20

<210> SEQ ID NO 571
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 571

gttctggcca ggcccctgtc 20

<210> SEQ ID NO 572
<211> LENGTH: 20

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 572

agtgccccac ccccgccttc 20

<210> SEQ ID NO 573
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 573

gtggggcact gaccccgaca 20

<210> SEQ ID NO 574
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 574

aaccttctaa gcaaaccttc 20

<210> SEQ ID NO 575
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 575

gttacacaga accagaaggc 20

<210> SEQ ID NO 576
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 576

gaaggttaca cagaaccaga 20

<210> SEQ ID NO 577
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 577

agtcatgatg agtcatgctg 20

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<210> SEQ ID NO 578
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 578

gatgagtcat gctgaggctt 20

<210> SEQ ID NO 579
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 579

actctaggct gagaacatct 20

<210> SEQ ID NO 580
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 580

gtccccagca ggatgcttac 20

<210> SEQ ID NO 581
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 581

gccctgtaag catcctgctg 20

<210> SEQ ID NO 582
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 582

cagggcagat ggcaaaaaaa 20

<210> SEQ ID NO 583
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 583

gaggtggagt tttagtcagg 20

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<210> SEQ ID NO 584
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 584

aaacggcatc ataaagaaaa 20

<210> SEQ ID NO 585
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 585

gagacacaca gaaatgtaac gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 586
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 586

ggtggggcac tgaccccgac gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 587
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 587

ctagagtgat gactcctatc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 588
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 588

gactaaaact ccacctcaaa gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 589

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<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 589

aatatgtcac attctgtctc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 590
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 590

ggactatggg aggtcactaa gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 591
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 591

gctcatgctt ggactatggg gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 592
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 592

gttctggcca ggcccctgtc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 593
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 593

agtgccccac ccccgccttc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 594
<211> LENGTH: 105

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 594

gtggggcact gaccccgaca gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 595
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 595

aaccttctaa gcaaaccttc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 596
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 596

gttacacaga accagaaggc gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 597
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 597

gaaggttaca cagaaccaga gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 598
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 598

agtcatgatg agtcatgctg gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 599
<211> LENGTH: 105
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 599

gatgagtcat gctgaggctt gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 600
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 600

actctaggct gagaacatct gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 601
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 601

gtccccagca ggatgcttac gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 602
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 602

gccctgtaag catcctgctg gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 603
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 603

cagggcagat ggcaaaaaaa gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 604
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 604

gaggtggagt tttagtcagg gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 605
<211> LENGTH: 105
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polynucleotide

<400> SEQUENCE: 605

aaacggcatc ataaagaaaa gttttagagc tagaaatagc aagttaaaat aaggctagtc 60

cgttatcaac ttgaaaaagt ggcaccgagt cggtgctttt tttcc 105

<210> SEQ ID NO 606
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 606

tgtactctct gaggtgctc 19

<210> SEQ ID NO 607
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 607

acgcagataa gaaccagtt 19

<210> SEQ ID NO 608
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 608

catcaagtca gccatcagc 19

<210> SEQ ID NO 609
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 609

gagtcaccct cctggaaac 19

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<210> SEQ ID NO 610
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 610

cctgggctcc ggggcgttt 19

<210> SEQ ID NO 611
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 611

ggcccctgcg gccaccccg 19

<210> SEQ ID NO 612
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 612

ctccctccct gcccggtag 19

<210> SEQ ID NO 613
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 613

aggtttggaa agggcgtgc 19

<210> SEQ ID NO 614
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 614

actccactgc actccagtct 20

<210> SEQ ID NO 615
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 615

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tctgtggggg acctgcactg 20

<210> SEQ ID NO 616
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 616

ggggcgccag ttgtgtctcc 20

<210> SEQ ID NO 617
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 617

acaccattgc caccaccatt 20

<210> SEQ ID NO 618
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 618

caatgacccc ttcattgacc 20

<210> SEQ ID NO 619
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 619

ttgattttgg agggatctcg 20

<210> SEQ ID NO 620
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 620

ggaatccatg gagggaagat 20

<210> SEQ ID NO 621
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

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<400> SEQUENCE: 621

tgttctcgct caggtcagtg 20

<210> SEQ ID NO 622
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 622

ctctctgctc ctttgccaca 20

<210> SEQ ID NO 623
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 623

gtgctcttcg ggtttcagga 20

<210> SEQ ID NO 624
<211> LENGTH: 30
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 624

cgaaagagaa agcgaaccag tatcgagaac 30

<210> SEQ ID NO 625
<211> LENGTH: 27
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 625

cgttgtgcat agtcgctgct tgatcgc 27

<210> SEQ ID NO 626
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 626

gagtttggct caaattgtta ctctt 25

<210> SEQ ID NO 627
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 627

gggaaatggt ctaggagagt aaagt 25

<210> SEQ ID NO 628
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 628

gattggcttt gatttcccta ggg 23

<210> SEQ ID NO 629
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 629

gcctactcag actgttactc tgg 23

<210> SEQ ID NO 630
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 630

gcagttgcct aagaactggt ggg 23

<210> SEQ ID NO 631
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 631

ggggctccac cctcacgagt ggg 23

<210> SEQ ID NO 632
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 632

gccttcttta tcccctatcg agg 23

<210> SEQ ID NO 633
<211> LENGTH: 48
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: CDS
<222> LOCATION: (1)..(45)

<400> SEQUENCE: 633

gaa gga cca ttt gac gtt cag ctc cta ctc aga ctg tta ctc tgg tga 48
Glu Gly Pro Phe Asp Val Gln Leu Leu Leu Arg Leu Leu Leu Trp
1 5 10 15

<210> SEQ ID NO 634
<211> LENGTH: 15
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic peptide

<400> SEQUENCE: 634

Glu Gly Pro Phe Asp Val Gln Leu Leu Leu Arg Leu Leu Leu Trp
1 5 10 15

<210> SEQ ID NO 635
<211> LENGTH: 46
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: CDS
<222> LOCATION: (1)..(42)

<400> SEQUENCE: 635

gaa gga cca ttt gac gtt cag ctc cta ctc aga ctg tct ctg gtga 46
Glu Gly Pro Phe Asp Val Gln Leu Leu Leu Arg Leu Ser Leu
1 5 10

<210> SEQ ID NO 636
<211> LENGTH: 14
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic peptide

<400> SEQUENCE: 636

Glu Gly Pro Phe Asp Val Gln Leu Leu Leu Arg Leu Ser Leu
1 5 10

<210> SEQ ID NO 637
<211> LENGTH: 20
<212> TYPE: RNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 637

gccuacucag acuguuacuc 20

<210> SEQ ID NO 638
<211> LENGTH: 19

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 638

cccaccagtt cttaggcaa 19

<210> SEQ ID NO 639
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 639

cccaccagtt cttaggcaac 20

<210> SEQ ID NO 640
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 640

tttgatttcc agttcttagg 20

<210> SEQ ID NO 641
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 641

attaccatag agttcttag 19

<210> SEQ ID NO 642
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 642

gtggataaag gcaacaatg 19

<210> SEQ ID NO 643
<211> LENGTH: 16
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 643

cctcgatagg ggataa 16

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<210> SEQ ID NO 644
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 644

accaaaccca cttaggggat aaa 23

<210> SEQ ID NO 645
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 645

ggtatcttaa ggacctccaa g 21

<210> SEQ ID NO 646
<211> LENGTH: 47
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 646

taggaagtaa attaatttga agcttgccaa ttaatccaaa tcttacc 47

<210> SEQ ID NO 647
<211> LENGTH: 83
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 647

tttccttttt gcaaaaccca aaatatttta gctcctactc agactgttag agtaggagga 60

gaaggagggc tgtggctctt gtg 83

<210> SEQ ID NO 648
<211> LENGTH: 83
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (20)..(20)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (31)..(31)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (69)..(69)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 648

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aaaaccccca aaccctcccn ctttttcctc ntactctaac agtgtgagta ggaggccaca 60

cttagctgng gctctagtaa gtc 83

<210> SEQ ID NO 649
<211> LENGTH: 94
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (50)..(50)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 649

cttctttttt ccttttttgca aaaacccaaa atattttagc tcctactcan accgttttcg 60

gggggtattt tgtatttgga tgaaacactt gatg 94

<210> SEQ ID NO 650
<211> LENGTH: 93
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 650

tcttcttttt tccttcccgc aaaaacccaa aatattttcg ctcctactca ggacctggtc 60

gtggggtatt ttgtatttgg atgaaacact tga 93

<210> SEQ ID NO 651
<211> LENGTH: 104
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (21)..(22)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 651

tcaaggaaga tggcatttct nntttggaga tggcagtttc cttagtaacc acaggttgtg 60

tcaccatggc cagtaagaag gatgccacac ttgtgtgtgt cgct 104

<210> SEQ ID NO 652
<211> LENGTH: 86
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (35)..(36)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (42)..(44)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 652

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ttggagatag cagtttcctt ggtaagtaca ggttnntcac cnnnacagtg tgagtaggag 60

gccacactta gctgtggctc tagtaa 86

<210> SEQ ID NO 653
<211> LENGTH: 71
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (54)..(54)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (61)..(63)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (69)..(69)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 653

tttctagttt ggagatggca gtttccttag taaccacagg ttgtgtcacc aggnctcgag 60

nnntggttnt g 71

<210> SEQ ID NO 654
<211> LENGTH: 87
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 654

cagttgccat agtcttcact aaccgcgtca ccagagctca ggacctggtc gtggggtatt 60

ttgtatttgg atgaaacact tgatggt 87

<210> SEQ ID NO 655
<211> LENGTH: 92
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (52)..(52)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (59)..(59)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (78)..(78)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 655

tattgtatct tggtttttct tcactgctgg ccagtttact aacaatctga antaagccna 60

ttaatccaaa tcttaccnga cgaaaaatca aa 92

<210> SEQ ID NO 656
<211> LENGTH: 89

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (6)..(6)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (10)..(10)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (19)..(19)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (30)..(30)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (57)..(57)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 656

atggtntttn gtcattgcng gccagttttn taacaatgtg atcaatgcca attaatncaa 60

atcttaccag aggaaaaatc aagccacag 89

<210> SEQ ID NO 657
<211> LENGTH: 94
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (93)..(94)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 657

aataaaaaga aaaaaatatg ataaactgct cccagtataa aatacagagc taagacaaga 60

acgtttcatt ggctgttaaa agtcggaaat cgnn 94

<210> SEQ ID NO 658
<211> LENGTH: 94
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 658

ttctgattta aaattccctt gaataggaag taaattaatt tgaagcttgc caattaatcc 60

aaatcttacc agaggaaaaa tcaagccaca gata 94

<210> SEQ ID NO 659
<211> LENGTH: 90
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 659

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caaaacaaga tgggataaaa tcttctttct gatttaaaat tcccttgaat aggaagtaaa 60

ttactagagc tgttcaaggc ctgaactcta 90

<210> SEQ ID NO 660
<211> LENGTH: 96
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (49)..(49)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (51)..(52)
<223> OTHER INFORMATION: a, c, t, g, unknown or other
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (54)..(54)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 660

ctgatttaaa attcccttga ataggaagta aattactaga gctgttcang nncngaactc 60

taggctagca tttccagccc aaaacaaaca cagttc 96

<210> SEQ ID NO 661
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (8)..(26)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 661

ttgtttgnnn nnnnnnnnnn nnnnnn 26

<210> SEQ ID NO 662
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (6)..(24)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 662

caccgnnnnn nnnnnnnnnn nnnn 24

<210> SEQ ID NO 663
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base

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<222> LOCATION: (6)..(24)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 663

tcccannnnn nnnnnnnnnn nnnn 24

<210> SEQ ID NO 664
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (6)..(24)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 664

cctcgnnnnn nnnnnnnnnn nnnn 24

<210> SEQ ID NO 665
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (5)..(23)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 665

aaacnnnnnn nnnnnnnnnn nnncaa 26

<210> SEQ ID NO 666
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (5)..(23)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 666

aaacnnnnnn nnnnnnnnnn nnnc 24

<210> SEQ ID NO 667
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (5)..(23)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 667

aaacnnnnnn nnnnnnnnnn nnnt 24

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<210> SEQ ID NO 668
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (5)..(23)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 668

aaacnnnnnn nnnnnnnnnn nnnc 24

<210> SEQ ID NO 669
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 669

ttgtttgtct tcgagaagac ctgttt 26

<210> SEQ ID NO 670
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 670

caccgggtct tcgagaagac ctgttt 26

<210> SEQ ID NO 671
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 671

tcccaggtct tcgagaagac ctgttt 26

<210> SEQ ID NO 672
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 672

cctcgggtct tcgagaagac ctgttt 26

<210> SEQ ID NO 673
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 673

ggcctaccgc taaggctagc ctagt 25

<210> SEQ ID NO 674
<211> LENGTH: 1411
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic polypeptide

<400> SEQUENCE: 674

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1 5 10 15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
20 25 30

Gly Arg Gly Met Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr
35 40 45

Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser
50 55 60

Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys
65 70 75 80

Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala
85 90 95

Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn
100 105 110

Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val
115 120 125

Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu
130 135 140

Asp Lys Lys His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu
145 150 155 160

Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys
165 170 175

Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala
180 185 190

Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp
195 200 205

Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val
210 215 220

Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly
225 230 235 240

Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg
245 250 255

Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu
260 265 270

Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys
275 280 285

Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp
290 295 300

Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln
305 310 315 320

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Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu
                325                 330                 335

Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu
            340                 345                 350

Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr
        355                 360                 365

Leu Leu Lys Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu
    370                 375                 380

Ile Phe Phe Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly
385                 390                 395                 400

Gly Ala Ser Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu
                405                 410                 415

Lys Met Asp Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp
            420                 425                 430

Leu Leu Arg Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln
        435                 440                 445

Ile His Leu Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe
    450                 455                 460

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

Phe Arg Ile Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg
                485                 490                 495

Phe Ala Trp Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn
            500                 505                 510

Phe Glu Glu Val Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu
        515                 520                 525

Arg Met Thr Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro
    530                 535                 540

Lys His Ser Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr
545                 550                 555                 560

Lys Val Lys Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser
                565                 570                 575

Gly Glu Gln Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg
            580                 585                 590

Lys Val Thr Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu
        595                 600                 605

Cys Phe Asp Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala
    610                 615                 620

Ser Leu Gly Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp
625                 630                 635                 640

Phe Leu Asp Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu
                645                 650                 655

Thr Leu Thr Leu Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys
            660                 665                 670

Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg
        675                 680                 685

Arg Arg Tyr Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly
    690                 695                 700

Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser
705                 710                 715                 720

Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser
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               725                 730                 735

Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly
           740                 745                 750

Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile
        755                 760                 765

Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys
    770                 775                 780

Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg
785                 790                 795                 800

Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met
               805                 810                 815

Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys
           820                 825                 830

Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu
        835                 840                 845

Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp
    850                 855                 860

Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser
865                 870                 875                 880

Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp
               885                 890                 895

Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys
           900                 905                 910

Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr
        915                 920                 925

Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser
    930                 935                 940

Glu Leu Asp Lys Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg
945                 950                 955                 960

Gln Ile Thr Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr
               965                 970                 975

Lys Tyr Asp Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr
           980                 985                 990

Leu Lys Ser Lys Leu Val Ser Asp  Phe Arg Lys Asp Phe  Gln Phe Tyr
        995                 1000                1005

Lys Val  Arg Glu Ile Asn Asn  Tyr His His Ala His  Asp Ala Tyr
    1010                1015                1020

Leu Asn  Ala Val Val Gly Thr  Ala Leu Ile Lys Lys  Tyr Pro Lys
    1025                1030                1035

Leu Glu  Ser Glu Phe Val Tyr  Gly Asp Tyr Lys Val  Tyr Asp Val
    1040                1045                1050

Arg Lys  Met Ile Ala Lys Ser  Glu Gln Glu Ile Gly  Lys Ala Thr
    1055                1060                1065

Ala Lys  Tyr Phe Phe Tyr Ser  Asn Ile Met Asn Phe  Phe Lys Thr
    1070                1075                1080

Glu Ile  Thr Leu Ala Asn Gly  Glu Ile Arg Lys Arg  Pro Leu Ile
    1085                1090                1095

Glu Thr  Asn Gly Glu Thr Gly  Glu Ile Val Trp Asp  Lys Gly Arg
    1100                1105                1110

Asp Phe  Ala Thr Val Arg Lys  Val Leu Ser Met Pro  Gln Val Asn
    1115                1120                1125
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Ile Val  Lys Lys Thr Glu Val  Gln Thr Gly Gly Phe  Ser Lys Glu
    1130                1135                1140

Ser Ile  Leu Pro Lys Arg Asn  Ser Asp Lys Leu Ile  Ala Arg Lys
    1145                1150                1155

Lys Asp  Trp Asp Pro Lys Lys  Tyr Gly Gly Phe Asp  Ser Pro Thr
    1160                1165                1170

Val Ala  Tyr Ser Val Leu Val  Val Ala Lys Val Glu  Lys Gly Lys
    1175                1180                1185

Ser Lys  Lys Leu Lys Ser Val  Lys Glu Leu Leu Gly  Ile Thr Ile
    1190                1195                1200

Met Glu  Arg Ser Ser Phe Glu  Lys Asn Pro Ile Asp  Phe Leu Glu
    1205                1210                1215

Ala Lys  Gly Tyr Lys Glu Val  Lys Lys Asp Leu Ile  Ile Lys Leu
    1220                1225                1230

Pro Lys  Tyr Ser Leu Phe Glu  Leu Glu Asn Gly Arg  Lys Arg Met
    1235                1240                1245

Leu Ala  Ser Ala Gly Glu Leu  Gln Lys Gly Asn Glu  Leu Ala Leu
    1250                1255                1260

Pro Ser  Lys Tyr Val Asn Phe  Leu Tyr Leu Ala Ser  His Tyr Glu
    1265                1270                1275

Lys Leu  Lys Gly Ser Pro Glu  Asp Asn Glu Gln Lys  Gln Leu Phe
    1280                1285                1290

Val Glu  Gln His Lys His Tyr  Leu Asp Glu Ile Ile  Glu Gln Ile
    1295                1300                1305

Ser Glu  Phe Ser Lys Arg Val  Ile Leu Ala Asp Ala  Asn Leu Asp
    1310                1315                1320

Lys Val  Leu Ser Ala Tyr Asn  Lys His Arg Asp Lys  Pro Ile Arg
    1325                1330                1335

Glu Gln  Ala Glu Asn Ile Ile  His Leu Phe Thr Leu  Thr Asn Leu
    1340                1345                1350

Gly Ala  Pro Ala Ala Phe Lys  Tyr Phe Asp Thr Thr  Ile Asp Arg
    1355                1360                1365

Lys Arg  Tyr Thr Ser Thr Lys  Glu Val Leu Asp Ala  Thr Leu Ile
    1370                1375                1380

His Gln  Ser Ile Thr Gly Leu  Tyr Glu Thr Arg Ile  Asp Leu Ser
    1385                1390                1395

Gln Leu  Gly Gly Asp Pro Lys  Lys Lys Arg Lys Val  Gly
    1400                1405                1410


<210> SEQ ID NO 675
<211> LENGTH: 1483
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 675

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                  10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Gly Met Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr
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        35                  40                  45

Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser
    50                  55                  60

Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys
65                  70                  75                  80

Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala
                85                  90                  95

Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn
            100                 105                 110

Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val
        115                 120                 125

Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu
    130                 135                 140

Asp Lys Lys His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu
145                 150                 155                 160

Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys
                165                 170                 175

Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala
            180                 185                 190

Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp
        195                 200                 205

Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val
    210                 215                 220

Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly
225                 230                 235                 240

Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg
                245                 250                 255

Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu
            260                 265                 270

Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys
        275                 280                 285

Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp
    290                 295                 300

Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln
305                 310                 315                 320

Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu
                325                 330                 335

Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu
            340                 345                 350

Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr
        355                 360                 365

Leu Leu Lys Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu
    370                 375                 380

Ile Phe Phe Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly
385                 390                 395                 400

Gly Ala Ser Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu
                405                 410                 415

Lys Met Asp Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp
            420                 425                 430

Leu Leu Arg Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln
        435                 440                 445
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```
Ile His Leu Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe
    450                 455                 460

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

Phe Arg Ile Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg
                485                 490                 495

Phe Ala Trp Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn
            500                 505                 510

Phe Glu Glu Val Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu
        515                 520                 525

Arg Met Thr Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro
    530                 535                 540

Lys His Ser Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr
545                 550                 555                 560

Lys Val Lys Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser
                565                 570                 575

Gly Glu Gln Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg
            580                 585                 590

Lys Val Thr Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu
        595                 600                 605

Cys Phe Asp Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala
    610                 615                 620

Ser Leu Gly Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp
625                 630                 635                 640

Phe Leu Asp Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu
                645                 650                 655

Thr Leu Thr Leu Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys
            660                 665                 670

Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg
        675                 680                 685

Arg Arg Tyr Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly
    690                 695                 700

Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser
705                 710                 715                 720

Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser
                725                 730                 735

Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly
            740                 745                 750

Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile
        755                 760                 765

Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys
    770                 775                 780

Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg
785                 790                 795                 800

Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met
                805                 810                 815

Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys
            820                 825                 830

Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu
        835                 840                 845
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Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp
    850                 855                 860

Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp Ala Ile Val Pro Gln Ser
865                 870                 875                 880

Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp
                885                 890                 895

Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys
            900                 905                 910

Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr
        915                 920                 925

Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser
    930                 935                 940

Glu Leu Asp Lys Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg
945                 950                 955                 960

Gln Ile Thr Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr
                965                 970                 975

Lys Tyr Asp Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr
            980                 985                 990

Leu Lys Ser Lys Leu Val Ser Asp  Phe Arg Lys Asp Phe  Gln Phe Tyr
        995                 1000                1005

Lys Val  Arg Glu Ile Asn Asn  Tyr His His Ala His  Asp Ala Tyr
    1010                1015                1020

Leu Asn  Ala Val Val Gly Thr  Ala Leu Ile Lys Lys  Tyr Pro Lys
    1025                1030                1035

Leu Glu  Ser Glu Phe Val Tyr  Gly Asp Tyr Lys Val  Tyr Asp Val
    1040                1045                1050

Arg Lys  Met Ile Ala Lys Ser  Glu Gln Glu Ile Gly  Lys Ala Thr
    1055                1060                1065

Ala Lys  Tyr Phe Phe Tyr Ser  Asn Ile Met Asn Phe  Phe Lys Thr
    1070                1075                1080

Glu Ile  Thr Leu Ala Asn Gly  Glu Ile Arg Lys Arg  Pro Leu Ile
    1085                1090                1095

Glu Thr  Asn Gly Glu Thr Gly  Glu Ile Val Trp Asp  Lys Gly Arg
    1100                1105                1110

Asp Phe  Ala Thr Val Arg Lys  Val Leu Ser Met Pro  Gln Val Asn
    1115                1120                1125

Ile Val  Lys Lys Thr Glu Val  Gln Thr Gly Gly Phe  Ser Lys Glu
    1130                1135                1140

Ser Ile  Leu Pro Lys Arg Asn  Ser Asp Lys Leu Ile  Ala Arg Lys
    1145                1150                1155

Lys Asp  Trp Asp Pro Lys Lys  Tyr Gly Gly Phe Asp  Ser Pro Thr
    1160                1165                1170

Val Ala  Tyr Ser Val Leu Val  Val Ala Lys Val Glu  Lys Gly Lys
    1175                1180                1185

Ser Lys  Lys Leu Lys Ser Val  Lys Glu Leu Leu Gly  Ile Thr Ile
    1190                1195                1200

Met Glu  Arg Ser Ser Phe Glu  Lys Asn Pro Ile Asp  Phe Leu Glu
    1205                1210                1215

Ala Lys  Gly Tyr Lys Glu Val  Lys Lys Asp Leu Ile  Ile Lys Leu
    1220                1225                1230

Pro Lys  Tyr Ser Leu Phe Glu  Leu Glu Asn Gly Arg  Lys Arg Met
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    1235                1240                1245

Leu Ala  Ser Ala Gly Glu Leu  Gln Lys Gly Asn Glu  Leu Ala Leu
    1250                1255                1260

Pro Ser  Lys Tyr Val Asn Phe  Leu Tyr Leu Ala Ser  His Tyr Glu
    1265                1270                1275

Lys Leu  Lys Gly Ser Pro Glu  Asp Asn Glu Gln Lys  Gln Leu Phe
    1280                1285                1290

Val Glu  Gln His Lys His Tyr  Leu Asp Glu Ile Ile  Glu Gln Ile
    1295                1300                1305

Ser Glu  Phe Ser Lys Arg Val  Ile Leu Ala Asp Ala  Asn Leu Asp
    1310                1315                1320

Lys Val  Leu Ser Ala Tyr Asn  Lys His Arg Asp Lys  Pro Ile Arg
    1325                1330                1335

Glu Gln  Ala Glu Asn Ile Ile  His Leu Phe Thr Leu  Thr Asn Leu
    1340                1345                1350

Gly Ala  Pro Ala Ala Phe Lys  Tyr Phe Asp Thr Thr  Ile Asp Arg
    1355                1360                1365

Lys Arg  Tyr Thr Ser Thr Lys  Glu Val Leu Asp Ala  Thr Leu Ile
    1370                1375                1380

His Gln  Ser Ile Thr Gly Leu  Tyr Glu Thr Arg Ile  Asp Leu Ser
    1385                1390                1395

Gln Leu  Gly Gly Asp Pro Ile  Ala Gly Ser Lys Ala  Ser Pro Lys
    1400                1405                1410

Lys Lys  Arg Lys Val Gly Arg  Ala Asp Ala Leu Asp  Asp Phe Asp
    1415                1420                1425

Leu Asp  Met Leu Gly Ser Asp  Ala Leu Asp Asp Phe  Asp Leu Asp
    1430                1435                1440

Met Leu  Gly Ser Asp Ala Leu  Asp Asp Phe Asp Leu  Asp Met Leu
    1445                1450                1455

Gly Ser  Asp Ala Leu Asp Asp  Phe Asp Leu Asp Met  Leu Ile Asn
    1460                1465                1470

Tyr Pro  Tyr Asp Val Pro Asp  Tyr Ala Ser
    1475                1480


<210> SEQ ID NO 676
<211> LENGTH: 2048
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      polypeptide

<400> SEQUENCE: 676

Met Asp Tyr Lys Asp His Asp Gly Asp Tyr Lys Asp His Asp Ile Asp
1               5                   10                  15

Tyr Lys Asp Asp Asp Asp Lys Met Ala Pro Lys Lys Lys Arg Lys Val
            20                  25                  30

Gly Arg Gly Met Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr
        35                  40                  45

Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser
    50                  55                  60

Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys
65                  70                  75                  80
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Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala
                85                  90                  95

Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn
            100                 105                 110

Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val
        115                 120                 125

Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu
    130                 135                 140

Asp Lys Lys His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu
145                 150                 155                 160

Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys
                165                 170                 175

Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala
            180                 185                 190

Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp
        195                 200                 205

Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val
    210                 215                 220

Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly
225                 230                 235                 240

Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg
                245                 250                 255

Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu
            260                 265                 270

Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys
        275                 280                 285

Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp
    290                 295                 300

Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln
305                 310                 315                 320

Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu
                325                 330                 335

Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu
            340                 345                 350

Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr
        355                 360                 365

Leu Leu Lys Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu
    370                 375                 380

Ile Phe Phe Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly
385                 390                 395                 400

Gly Ala Ser Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu
                405                 410                 415

Lys Met Asp Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp
            420                 425                 430

Leu Leu Arg Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln
        435                 440                 445

Ile His Leu Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe
    450                 455                 460

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

Phe Arg Ile Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg
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               485                 490                 495

Phe Ala Trp Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn
            500                 505                 510

Phe Glu Glu Val Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu
        515                 520                 525

Arg Met Thr Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro
    530                 535                 540

Lys His Ser Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr
545                 550                 555                 560

Lys Val Lys Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser
                565                 570                 575

Gly Glu Gln Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg
            580                 585                 590

Lys Val Thr Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu
        595                 600                 605

Cys Phe Asp Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala
    610                 615                 620

Ser Leu Gly Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp
625                 630                 635                 640

Phe Leu Asp Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu
                645                 650                 655

Thr Leu Thr Leu Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys
            660                 665                 670

Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg
        675                 680                 685

Arg Arg Tyr Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly
    690                 695                 700

Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser
705                 710                 715                 720

Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser
                725                 730                 735

Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly
            740                 745                 750

Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile
        755                 760                 765

Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys
    770                 775                 780

Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg
785                 790                 795                 800

Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met
                805                 810                 815

Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys
            820                 825                 830

Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu
        835                 840                 845

Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp
    850                 855                 860

Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp Ala Ile Val Pro Gln Ser
865                 870                 875                 880

Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp
                885                 890                 895
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Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys
            900                 905                 910

Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr
        915                 920                 925

Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser
    930                 935                 940

Glu Leu Asp Lys Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg
945                 950                 955                 960

Gln Ile Thr Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr
                965                 970                 975

Lys Tyr Asp Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr
            980                 985                 990

Leu Lys Ser Lys Leu Val Ser Asp  Phe Arg Lys Asp Phe  Gln Phe Tyr
        995                 1000                1005

Lys Val  Arg Glu Ile Asn Asn  Tyr His His Ala His  Asp Ala Tyr
    1010                1015                1020

Leu Asn  Ala Val Val Gly Thr  Ala Leu Ile Lys Lys  Tyr Pro Lys
    1025                1030                1035

Leu Glu  Ser Glu Phe Val Tyr  Gly Asp Tyr Lys Val  Tyr Asp Val
    1040                1045                1050

Arg Lys  Met Ile Ala Lys Ser  Glu Gln Glu Ile Gly  Lys Ala Thr
    1055                1060                1065

Ala Lys  Tyr Phe Phe Tyr Ser  Asn Ile Met Asn Phe  Phe Lys Thr
    1070                1075                1080

Glu Ile  Thr Leu Ala Asn Gly  Glu Ile Arg Lys Arg  Pro Leu Ile
    1085                1090                1095

Glu Thr  Asn Gly Glu Thr Gly  Glu Ile Val Trp Asp  Lys Gly Arg
    1100                1105                1110

Asp Phe  Ala Thr Val Arg Lys  Val Leu Ser Met Pro  Gln Val Asn
    1115                1120                1125

Ile Val  Lys Lys Thr Glu Val  Gln Thr Gly Gly Phe  Ser Lys Glu
    1130                1135                1140

Ser Ile  Leu Pro Lys Arg Asn  Ser Asp Lys Leu Ile  Ala Arg Lys
    1145                1150                1155

Lys Asp  Trp Asp Pro Lys Lys  Tyr Gly Gly Phe Asp  Ser Pro Thr
    1160                1165                1170

Val Ala  Tyr Ser Val Leu Val  Val Ala Lys Val Glu  Lys Gly Lys
    1175                1180                1185

Ser Lys  Lys Leu Lys Ser Val  Lys Glu Leu Leu Gly  Ile Thr Ile
    1190                1195                1200

Met Glu  Arg Ser Ser Phe Glu  Lys Asn Pro Ile Asp  Phe Leu Glu
    1205                1210                1215

Ala Lys  Gly Tyr Lys Glu Val  Lys Lys Asp Leu Ile  Ile Lys Leu
    1220                1225                1230

Pro Lys  Tyr Ser Leu Phe Glu  Leu Glu Asn Gly Arg  Lys Arg Met
    1235                1240                1245

Leu Ala  Ser Ala Gly Glu Leu  Gln Lys Gly Asn Glu  Leu Ala Leu
    1250                1255                1260

Pro Ser  Lys Tyr Val Asn Phe  Leu Tyr Leu Ala Ser  His Tyr Glu
    1265                1270                1275
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Lys Leu  Lys Gly Ser Pro Glu  Asp Asn Glu Gln Lys  Gln Leu Phe
    1280                1285                1290

Val Glu  Gln His Lys His Tyr  Leu Asp Glu Ile Ile  Glu Gln Ile
    1295                1300                1305

Ser Glu  Phe Ser Lys Arg Val  Ile Leu Ala Asp Ala  Asn Leu Asp
    1310                1315                1320

Lys Val  Leu Ser Ala Tyr Asn  Lys His Arg Asp Lys  Pro Ile Arg
    1325                1330                1335

Glu Gln  Ala Glu Asn Ile Ile  His Leu Phe Thr Leu  Thr Asn Leu
    1340                1345                1350

Gly Ala  Pro Ala Ala Phe Lys  Tyr Phe Asp Thr Thr  Ile Asp Arg
    1355                1360                1365

Lys Arg  Tyr Thr Ser Thr Lys  Glu Val Leu Asp Ala  Thr Leu Ile
    1370                1375                1380

His Gln  Ser Ile Thr Gly Leu  Tyr Glu Thr Arg Ile  Asp Leu Ser
    1385                1390                1395

Gln Leu  Gly Gly Asp Pro Ile  Ala Gly Ser Lys Ala  Ser Pro Lys
    1400                1405                1410

Lys Lys  Arg Lys Val Gly Arg  Ala Ile Phe Lys Pro  Glu Glu Leu
    1415                1420                1425

Arg Gln  Ala Leu Met Pro Thr  Leu Glu Ala Leu Tyr  Arg Gln Asp
    1430                1435                1440

Pro Glu  Ser Leu Pro Phe Arg  Gln Pro Val Asp Pro  Gln Leu Leu
    1445                1450                1455

Gly Ile  Pro Asp Tyr Phe Asp  Ile Val Lys Ser Pro  Met Asp Leu
    1460                1465                1470

Ser Thr  Ile Lys Arg Lys Leu  Asp Thr Gly Gln Tyr  Gln Glu Pro
    1475                1480                1485

Trp Gln  Tyr Val Asp Asp Ile  Trp Leu Met Phe Asn  Asn Ala Trp
    1490                1495                1500

Leu Tyr  Asn Arg Lys Thr Ser  Arg Val Tyr Lys Tyr  Cys Ser Lys
    1505                1510                1515

Leu Ser  Glu Val Phe Glu Gln  Glu Ile Asp Pro Val  Met Gln Ser
    1520                1525                1530

Leu Gly  Tyr Cys Cys Gly Arg  Lys Leu Glu Phe Ser  Pro Gln Thr
    1535                1540                1545

Leu Cys  Cys Tyr Gly Lys Gln  Leu Cys Thr Ile Pro  Arg Asp Ala
    1550                1555                1560

Thr Tyr  Tyr Ser Tyr Gln Asn  Arg Tyr His Phe Cys  Glu Lys Cys
    1565                1570                1575

Phe Asn  Glu Ile Gln Gly Glu  Ser Val Ser Leu Gly  Asp Asp Pro
    1580                1585                1590

Ser Gln  Pro Gln Thr Thr Ile  Asn Lys Glu Gln Phe  Ser Lys Arg
    1595                1600                1605

Lys Asn  Asp Thr Leu Asp Pro  Glu Leu Phe Val Glu  Cys Thr Glu
    1610                1615                1620

Cys Gly  Arg Lys Met His Gln  Ile Cys Val Leu His  His Glu Ile
    1625                1630                1635

Ile Trp  Pro Ala Gly Phe Val  Cys Asp Gly Cys Leu  Lys Lys Ser
    1640                1645                1650

Ala Arg  Thr Arg Lys Glu Asn  Lys Phe Ser Ala Lys  Arg Leu Pro
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    1655                1660                1665

Ser Thr  Arg Leu Gly Thr Phe  Leu Glu Asn Arg Val  Asn Asp Phe
    1670                1675                1680

Leu Arg  Arg Gln Asn His Pro  Glu Ser Gly Glu Val  Thr Val Arg
    1685                1690                1695

Val Val  His Ala Ser Asp Lys  Thr Val Glu Val Lys  Pro Gly Met
    1700                1705                1710

Lys Ala  Arg Phe Val Asp Ser  Gly Glu Met Ala Glu  Ser Phe Pro
    1715                1720                1725

Tyr Arg  Thr Lys Ala Leu Phe  Ala Phe Glu Glu Ile  Asp Gly Val
    1730                1735                1740

Asp Leu  Cys Phe Phe Gly Met  His Val Gln Glu Tyr  Gly Ser Asp
    1745                1750                1755

Cys Pro  Pro Pro Asn Gln Arg  Arg Val Tyr Ile Ser  Tyr Leu Asp
    1760                1765                1770

Ser Val  His Phe Phe Arg Pro  Lys Cys Leu Arg Thr  Ala Val Tyr
    1775                1780                1785

His Glu  Ile Leu Ile Gly Tyr  Leu Glu Tyr Val Lys  Lys Leu Gly
    1790                1795                1800

Tyr Thr  Thr Gly His Ile Trp  Ala Cys Pro Pro Ser  Glu Gly Asp
    1805                1810                1815

Asp Tyr  Ile Phe His Cys His  Pro Pro Asp Gln Lys  Ile Pro Lys
    1820                1825                1830

Pro Lys  Arg Leu Gln Glu Trp  Tyr Lys Lys Met Leu  Asp Lys Ala
    1835                1840                1845

Val Ser  Glu Arg Ile Val His  Asp Tyr Lys Asp Ile  Phe Lys Gln
    1850                1855                1860

Ala Thr  Glu Asp Arg Leu Thr  Ser Ala Lys Glu Leu  Pro Tyr Phe
    1865                1870                1875

Glu Gly  Asp Phe Trp Pro Asn  Val Leu Glu Glu Ser  Ile Lys Glu
    1880                1885                1890

Leu Glu  Gln Glu Glu Glu Glu  Arg Lys Arg Glu Glu  Asn Thr Ser
    1895                1900                1905

Asn Glu  Ser Thr Asp Val Thr  Lys Gly Asp Ser Lys  Asn Ala Lys
    1910                1915                1920

Lys Lys  Asn Asn Lys Lys Thr  Ser Lys Asn Lys Ser  Ser Leu Ser
    1925                1930                1935

Arg Gly  Asn Lys Lys Lys Pro  Gly Met Pro Asn Val  Ser Asn Asp
    1940                1945                1950

Leu Ser  Gln Lys Leu Tyr Ala  Thr Met Glu Lys His  Lys Glu Val
    1955                1960                1965

Phe Phe  Val Ile Arg Leu Ile  Ala Gly Pro Ala Ala  Asn Ser Leu
    1970                1975                1980

Pro Pro  Ile Val Asp Pro Asp  Pro Leu Ile Pro Cys  Asp Leu Met
    1985                1990                1995

Asp Gly  Arg Asp Ala Phe Leu  Thr Leu Ala Arg Asp  Lys His Leu
    2000                2005                2010

Glu Phe  Ser Ser Leu Arg Arg  Ala Gln Trp Ser Thr  Met Cys Met
    2015                2020                2025

Leu Val  Glu Leu His Thr Gln  Ser Gln Asp Tyr Pro  Tyr Asp Val
    2030                2035                2040
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Pro Asp Tyr Ala Ser
2045

<210> SEQ ID NO 677
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Dystrophin gene reference sequence
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (1)..(21)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 677

nnnnnnnnnn nnnnnnnnnn ngg 23

<210> SEQ ID NO 678
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Dystrophin gene reference sequence
<220> FEATURE:
<221> NAME/KEY: modified_base
<222> LOCATION: (2)..(21)
<223> OTHER INFORMATION: a, c, t, g, unknown or other

<400> SEQUENCE: 678

gnnnnnnnnn nnnnnnnnnn ngg 23

<210> SEQ ID NO 679
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 679

gtgtagagta agtcagccta 20

<210> SEQ ID NO 680
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 680

gttggacaga acttaccgac 20

<210> SEQ ID NO 681
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 681

gtttgcttcg ctataaaacg 20

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<210> SEQ ID NO 682
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 682

gtctgaggat ggggccgcaa 20

<210> SEQ ID NO 683
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 683

ggatctgtca aatcgcctgc 20

<210> SEQ ID NO 684
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 684

gccaggatgg cattgggcag 20

<210> SEQ ID NO 685
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 685

gctgaatctg cggtggcagg 20

<210> SEQ ID NO 686
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 686

gttctttttgt tcttctagcc 20

<210> SEQ ID NO 687
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 687

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ggaaaagctt gagcaagtca 20

<210> SEQ ID NO 688
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 688

ggaagagttg cccctgcgcc 20

<210> SEQ ID NO 689
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 689

gacaaatctc cagtggataa 20

<210> SEQ ID NO 690
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 690

gtgtttctca ggtaaagctc 20

<210> SEQ ID NO 691
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 691

ggaaggacca tttgacgtta 20

<210> SEQ ID NO 692
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 692

gaactgctat ttcagtttcc 20

<210> SEQ ID NO 693
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 693

gccagccact cagccagtga 20

<210> SEQ ID NO 694
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 694

ggtatgcttt tctgttaaag 20

<210> SEQ ID NO 695
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 695

gctcctggac tgaccactat 20

<210> SEQ ID NO 696
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 696

ggaacagagg cgtccccagt tgg 23

<210> SEQ ID NO 697
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 697

ggaggctaga acaatcatta 20

<210> SEQ ID NO 698
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 698

gacaagaaca ccttcagaac 20

<210> SEQ ID NO 699
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 699

gggtttctgt gattttcttt 20

<210> SEQ ID NO 700
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 700

gggccaaaga cctccgccag 20

<210> SEQ ID NO 701
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 701

gttggagaag cattcataaa 20

<210> SEQ ID NO 702
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 702

gtcgctcact caccctgcaa 20

<210> SEQ ID NO 703
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 703

gaaaagagct gatgaaacaa 20

<210> SEQ ID NO 704
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 704

gtacactttt caaaatgctt 20

<210> SEQ ID NO 705
<211> LENGTH: 20
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 705

ggagatgatc atcaagcaga 20

<210> SEQ ID NO 706
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 706

gctttgaaag agcaataaaa 20

<210> SEQ ID NO 707
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 707

gcacaaaagt caaatcggaa 20

<210> SEQ ID NO 708
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 708

gatttcaata taagattcgg 20

<210> SEQ ID NO 709
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 709

gcttaagcaa tcccgaactc 20

<210> SEQ ID NO 710
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 710

gaggccaaac ctcggcttac 20

<210> SEQ ID NO 711

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<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 711

gttcgaaaat ttcaggtaag 20

<210> SEQ ID NO 712
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 712

ggcagaacag gagataacag 20

<210> SEQ ID NO 713
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 713

ggcggccctc gcccttctct 20

<210> SEQ ID NO 714
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 714

gtagtgatcg tggatacgag 20

<210> SEQ ID NO 715
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 715

gtacagccct cggtgtatat 20

<210> SEQ ID NO 716
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 716

gggaaggaat taagcccgaa 20

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<210> SEQ ID NO 717
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 717

gggaacagct ttcgtagttg 20

<210> SEQ ID NO 718
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 718

gataaagtcc agtgtcgatc 20

<210> SEQ ID NO 719
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 719

gaaaaccaga gcttcggtca 20

<210> SEQ ID NO 720
<211> LENGTH: 31
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 720

ggagtcttct gggcaggctt aaaggctaac c 31

<210> SEQ ID NO 721
<211> LENGTH: 31
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 721

gtcgggtgag catgtcttta atctacctcg a 31

<210> SEQ ID NO 722
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 722

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ggtgtcacca gagtaacagt 20

<210> SEQ ID NO 723
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 723

gtgatcatca agcagaaggt 20

<210> SEQ ID NO 724
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 724

gaacttcgaa aatttcaggt a 21

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<400> SEQUENCE: 725

ggaaactcat caaatatgcg t 21

<210> SEQ ID NO 726
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<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 726

gtcatttaca ctaacacgca t 21

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<211> LENGTH: 21
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<213> ORGANISM: Artificial Sequence
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<400> SEQUENCE: 727

ggaatgaaac tcatcaaata t 21

<210> SEQ ID NO 728
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

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<400> SEQUENCE: 728

gtcatcaata tctttgaagg a 21

<210> SEQ ID NO 729
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 729

gtgttttcat aggaaaaata g 21

<210> SEQ ID NO 730
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 730

gaattggaaa atgtgatggg a 21

<210> SEQ ID NO 731
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 731

gatgatcatc aagcagaagg t 21

<210> SEQ ID NO 732
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 732

gagatgatca tcaagcagaa g 21

<210> SEQ ID NO 733
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 733

gcatttttc tcataccttc t 21

<210> SEQ ID NO 734
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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oligonucleotide

<400> SEQUENCE: 734

gtcctactca gactgttact c 21

<210> SEQ ID NO 735
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 735

gacaggttgt gtcaccagag t 21

<210> SEQ ID NO 736
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<400> SEQUENCE: 736

gttatcattt tttctcatac c 21

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 737

gttgcctaag aactggtggg a 21

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 738

gaaacagttg cctaagaact g 21

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<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 739

gtttcccacc agttcttagg c 21

<210> SEQ ID NO 740
<211> LENGTH: 21
<212> TYPE: DNA
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 740

gtggctttga tttccctagg g 21

<210> SEQ ID NO 741
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 741

gtagggaaat caaagccaat g 21

<210> SEQ ID NO 742
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 742

ggaccctagg gaaatcaaag c 21

<210> SEQ ID NO 743
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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 743

gtgagggctc caccctcacg a 21

<210> SEQ ID NO 744
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 744

gaaggattga gggctccacc c 21

<210> SEQ ID NO 745
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 745

ggctccaccc tcacgagtgg g 21

<210> SEQ ID NO 746
<211> LENGTH: 21

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 746

gtatccccta tcgaggaaac c 21

<210> SEQ ID NO 747
<211> LENGTH: 21
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 747

ggataaagaa ggcctatttc a 21

<210> SEQ ID NO 748
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 748

gaggccttct ttatccccta t 21

<210> SEQ ID NO 749
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 749

gtgagggctc caccctcacg a 21

<210> SEQ ID NO 750
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic oligonucleotide

<400> SEQUENCE: 750

ggataaagaa ggcctatttc a 21

What is claimed is:

1. A DNA targeting system for deleting exon 51 of a dystrophin gene, the system comprising Cas9 and at least one guide RNA (gRNA), wherein the at least one gRNA binds and targets a polynucleotide sequence comprising SEQ ID NO: 67, SEQ ID NO: 70, SEQ ID NO: 100, or SEQ ID NO: 680.

2. An isolated polynucleotide encoding the DNA targeting system of claim **1**.

3. A vector comprising the isolated polynucleotide of claim **2**.

4. A cell comprising the isolated polynucleotide of claim **2**.

5. A method of treating a subject in need thereof having a mutant dystrophin gene, the method comprising administering to the subject the DNA targeting system of claim **1**.

6. The method of claim **5**, wherein the subject is suffering from Duchenne muscular dystrophy.

7. A method of correcting a mutant dystrophin gene in a cell, the method comprising administering to a cell containing a mutant dystrophin gene the DNA targeting system of claim **1**.

8. A composition for genome editing in a muscle of a subject, the composition comprising a modified adeno-associated virus (AAV) vector and a nucleotide sequence encoding the DNA targeting system of claim **1**, wherein the muscle is skeletal muscle or cardiac muscle.

9. The composition of claim **8**, wherein the modified AAV vector has enhanced cardiac and skeletal muscle tissue tropism.

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