Fig. 1

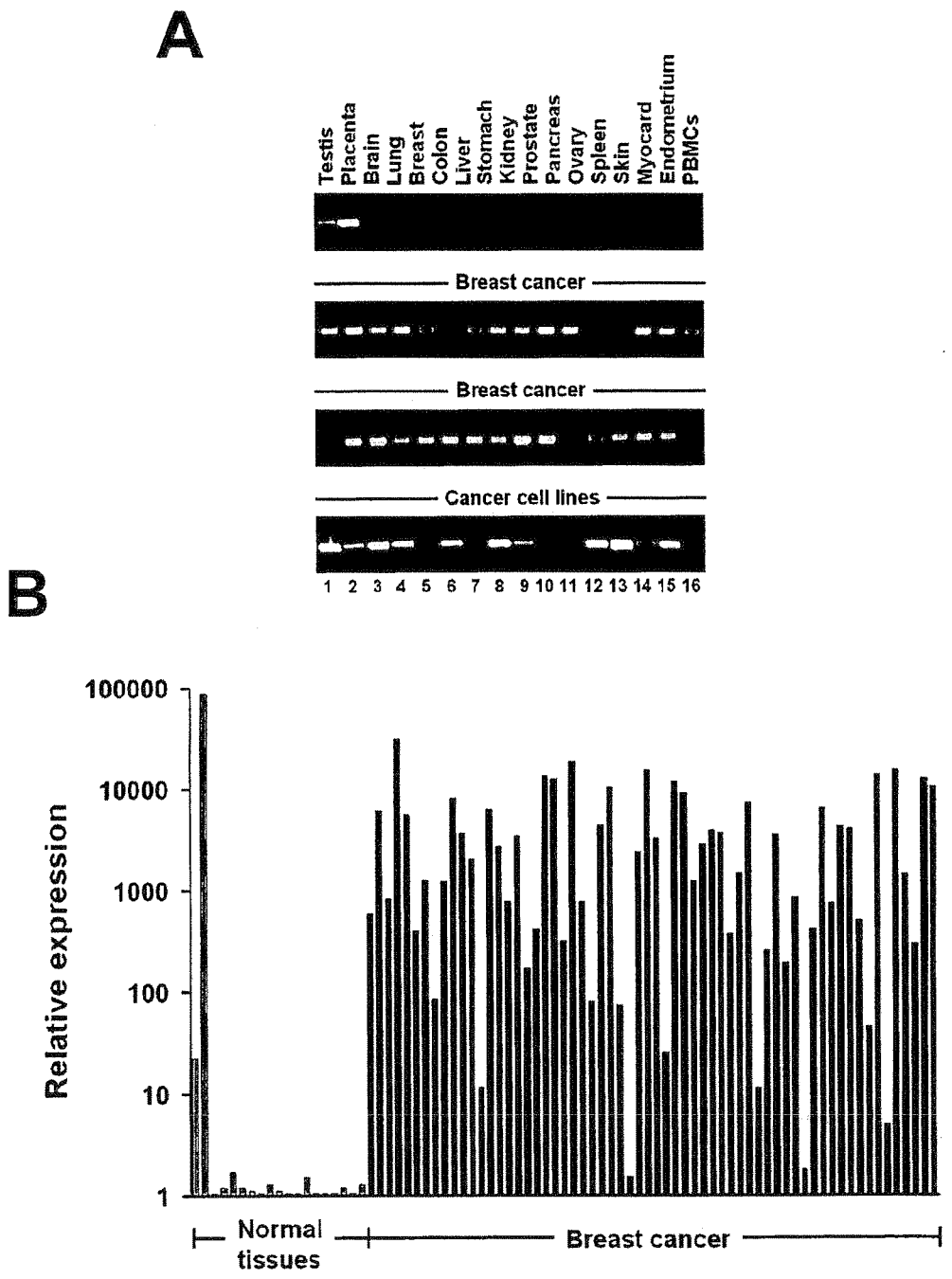

A
Testis
Placenta
Brain
Lung
Breast
Colon
Liver
Stomach
Kidney
Prostate
Pancreas
Ovary
Spleen
Skin
Myocard
Endometrium
PBMCs
Breast cancer
Breast cancer
Cancer cell lines
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16
B
Relative expression
100000
10000
1000
100
10
1
Normal tissues
Breast cancer

Fig. 1

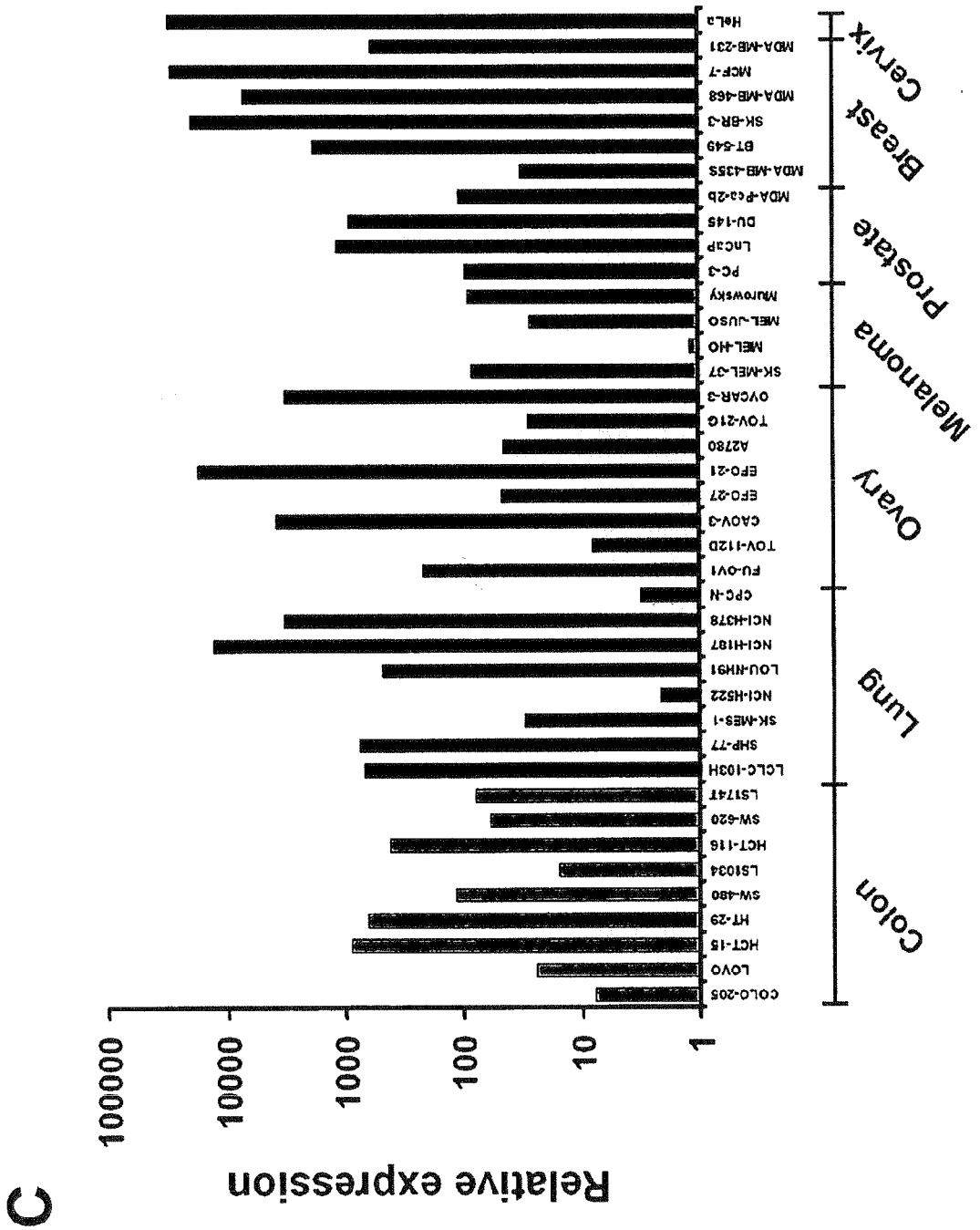

C
Relative expression
100000
10000
1000
100
10
1
Colon
Lung
Ovary
Melanoma
Prostate
Breast
Cervix
COLO-205
LOVO
HCT-15
HT-29
SW-480
LS1034
HCT-116
SW-620
LS174T
LCLC-103H
SHP-77
SK-MES-1
NCI-H522
LOU-NH91
NCI-H187
NCI-H378
CPC-N
FU-OV1
TOV-112D
CAOV-3
EFO-27
EFO-21
A2780
TOV-21G
OVCAR-3
SK-MEL-37
MEL-HO
MEL-UUSO
Murowsky
PC-3
LnCaP
DU-145
MDA-Pca-2b
MDA-MB-435S
BT-549
SK-BR-3
MDA-MB-468
MCF-7
MDA-MB-231
HeLa

Fig. 1

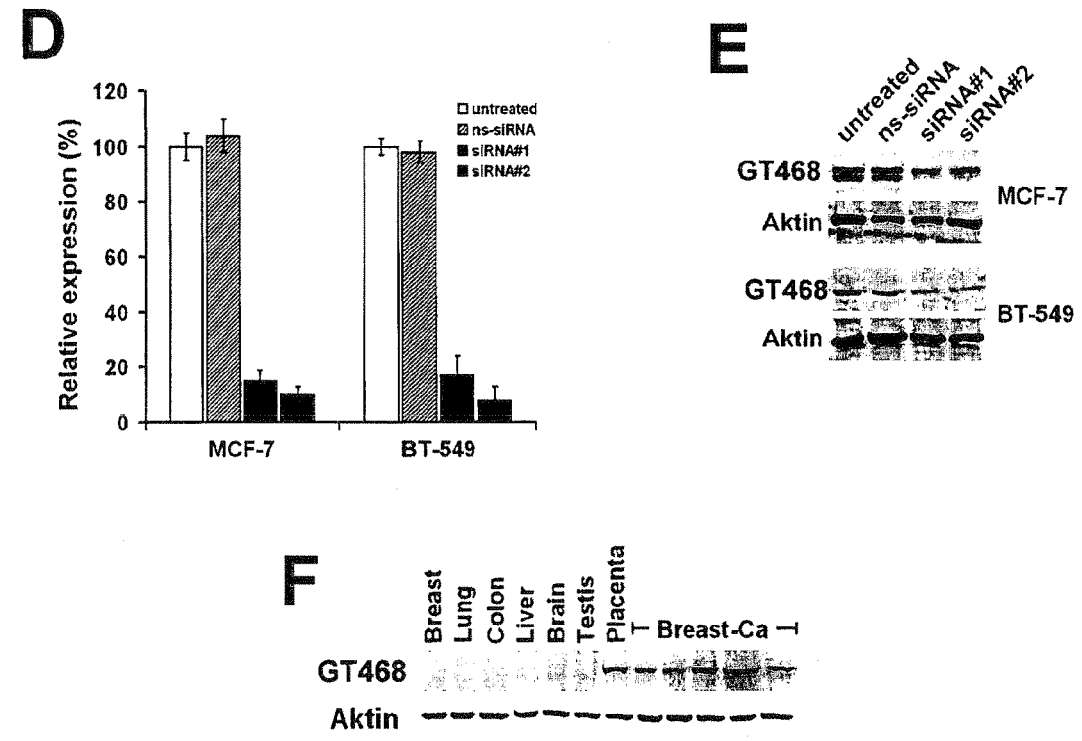

D
Relative expression (%)
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
siRNA#2
MCF-7
BT-549
E
untreated
ns-siRNA
siRNA#1
siRNA#2
GT468
Aktin
MCF-7
GT468
Aktin
BT-549
F
Breast
Lung
Colon
Liver
Brain
Testis
Placenta
Breast-Ca
GT468
Aktin
G

Fig. 2

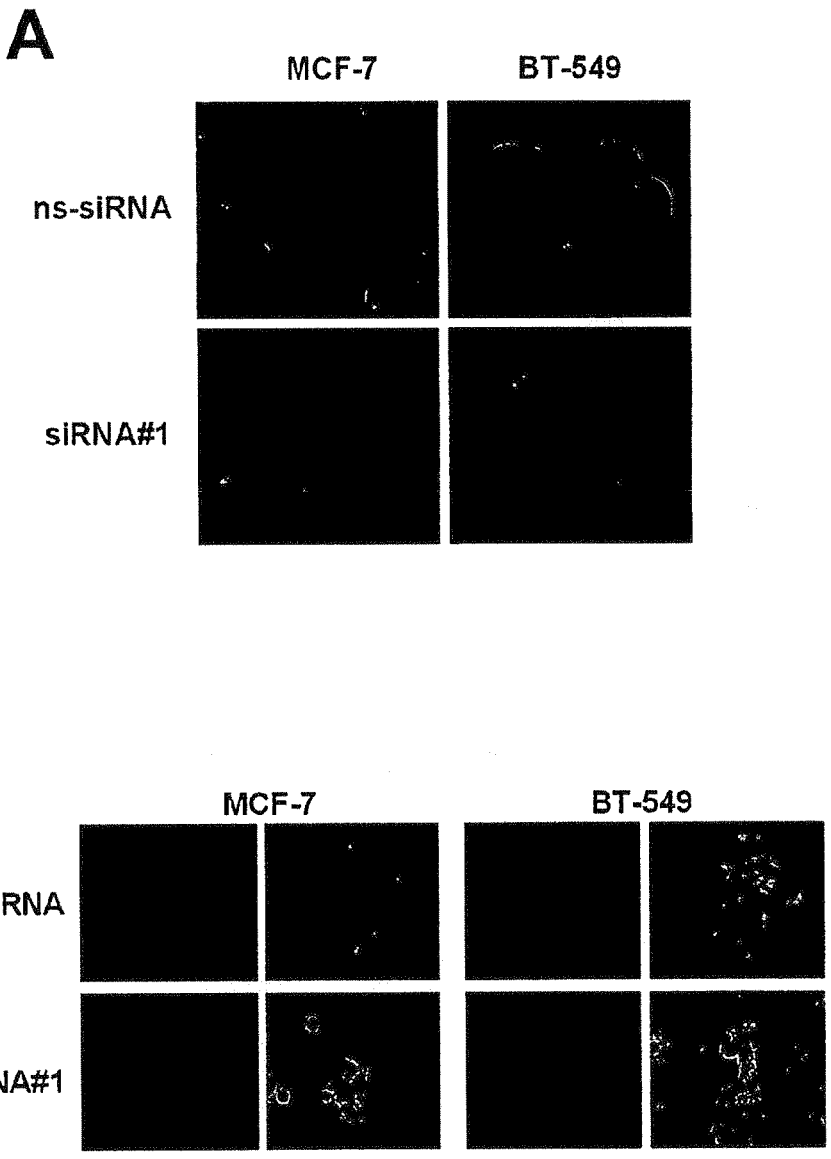


Fig. 3

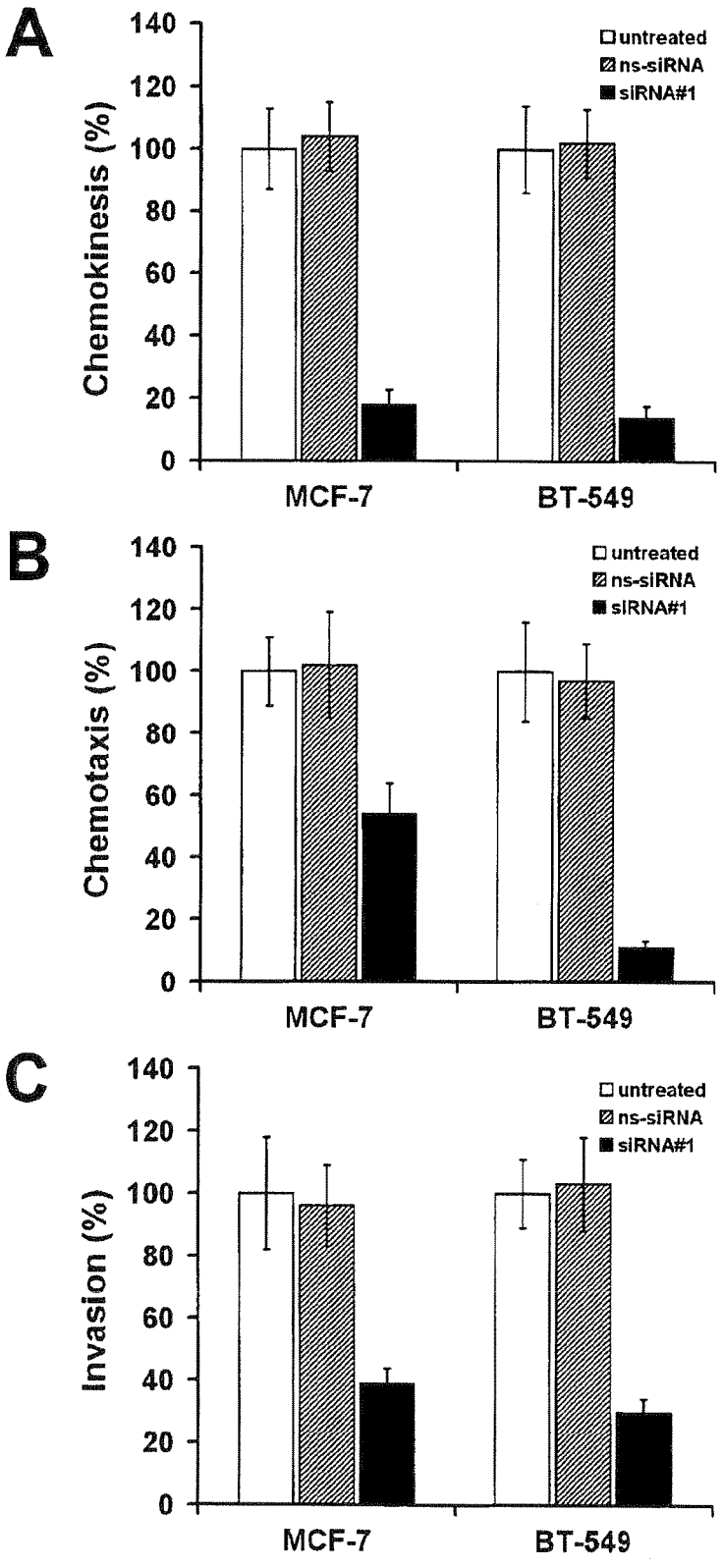

A
Chemokinesis (%)
140
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
MCF-7
BT-549
B
Chemotaxis (%)
140
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
MCF-7
BT-549
C
Invasion (%)
140
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
MCF-7
BT-549

Fig. 4

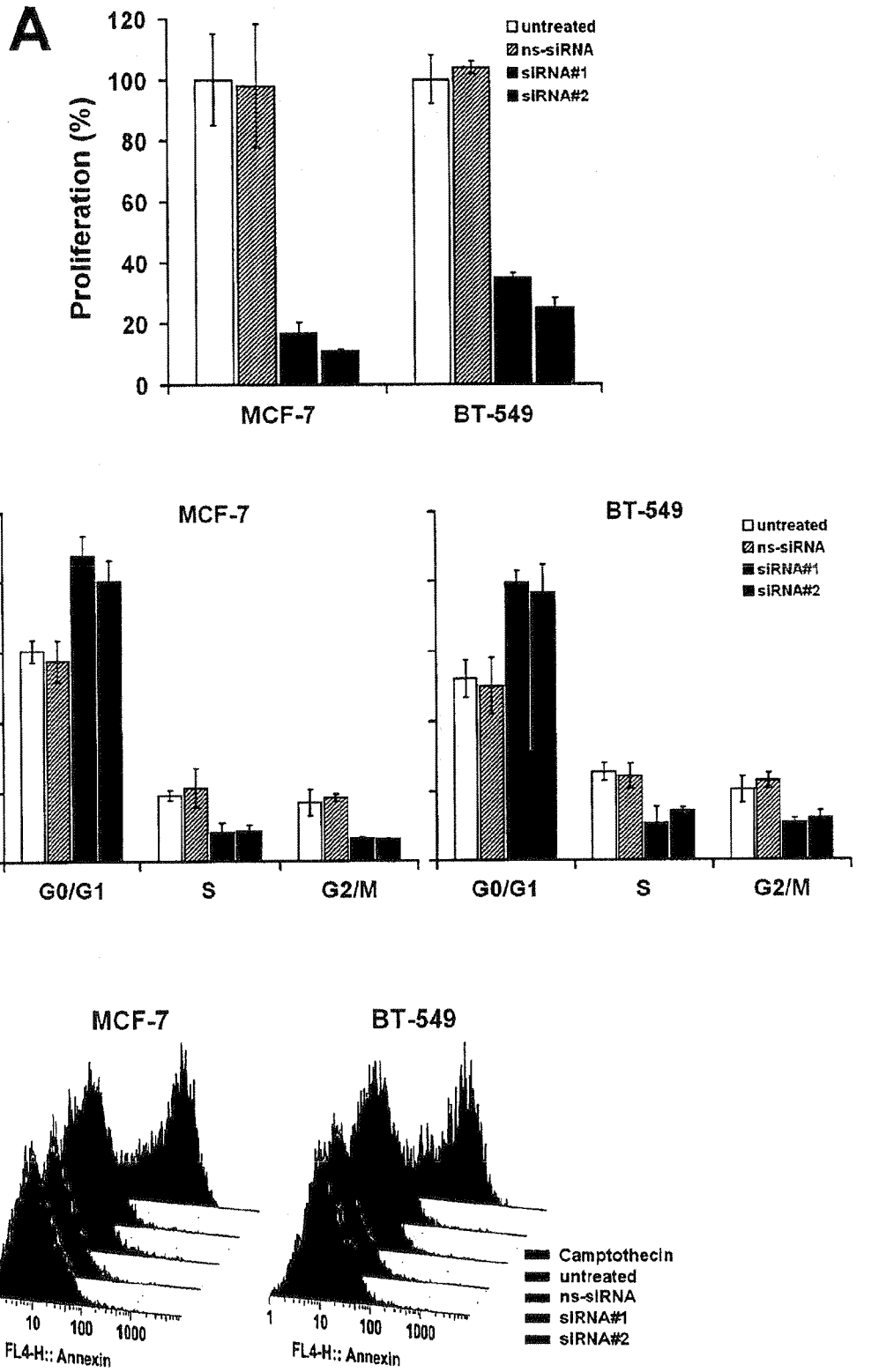

A
Proliferation (%)
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
siRNA#2
MCF-7
BT-549
B
Cells (%)
100
80
60
40
20
0
MCF-7
BT-549
G0/G1
S
G2/M
G0/G1
S
G2/M
untreated
ns-siRNA
siRNA#1
siRNA#2
C
MCF-7
BT-549
1
10
100
1000
FL4-H:: Annexin
1
10
100
1000
FL4-H:: Annexin
Camptothecin
untreated
ns-siRNA
siRNA#1
siRNA#2

Fig. 5

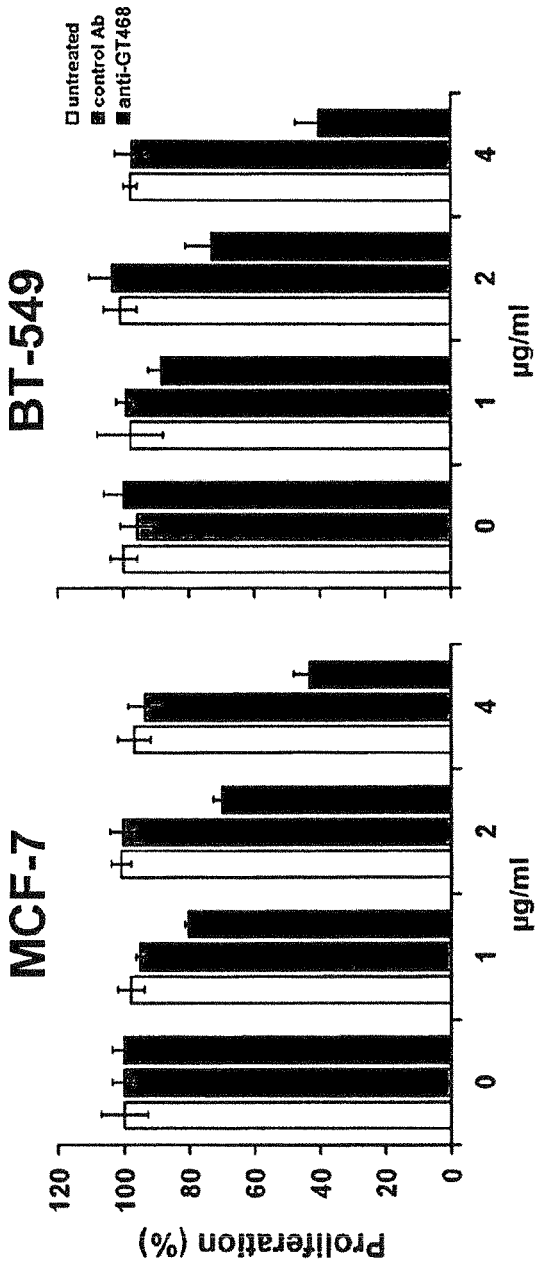

untreated
control Ab
anti-GT468
BT-549
MCF-7
µg/ml
0
1
2
4
Proliferation (%)
120
100
80
60
40
20
0

Fig. 6

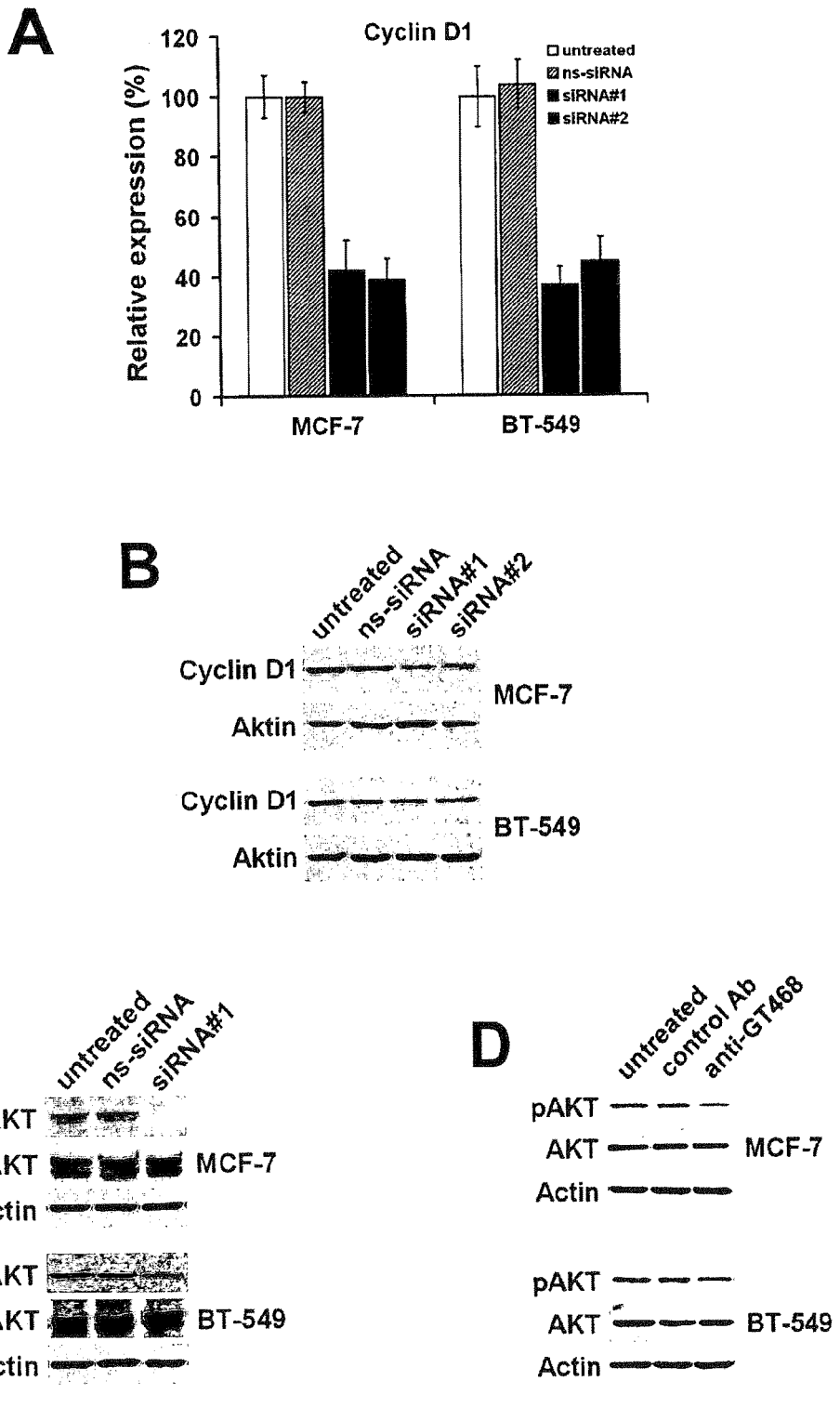

A
Cyclin D1
Relative expression (%)
120
100
80
60
40
20
0
untreated
ns-siRNA
siRNA#1
siRNA#2
MCF-7
BT-549
B
untreated
ns-siRNA
siRNA#1
siRNA#2
Cyclin D1
Aktin
MCF-7
Cyclin D1
Aktin
BT-549
C
untreated
ns-siRNA
siRNA#1
pAKT
AKT
Actin
MCF-7
pAKT
AKT
Actin
BT-549
D
untreated
control Ab
anti-GT468
pAKT
AKT
Actin
MCF-7
pAKT
AKT
Actin
BT-549

Fig. 7

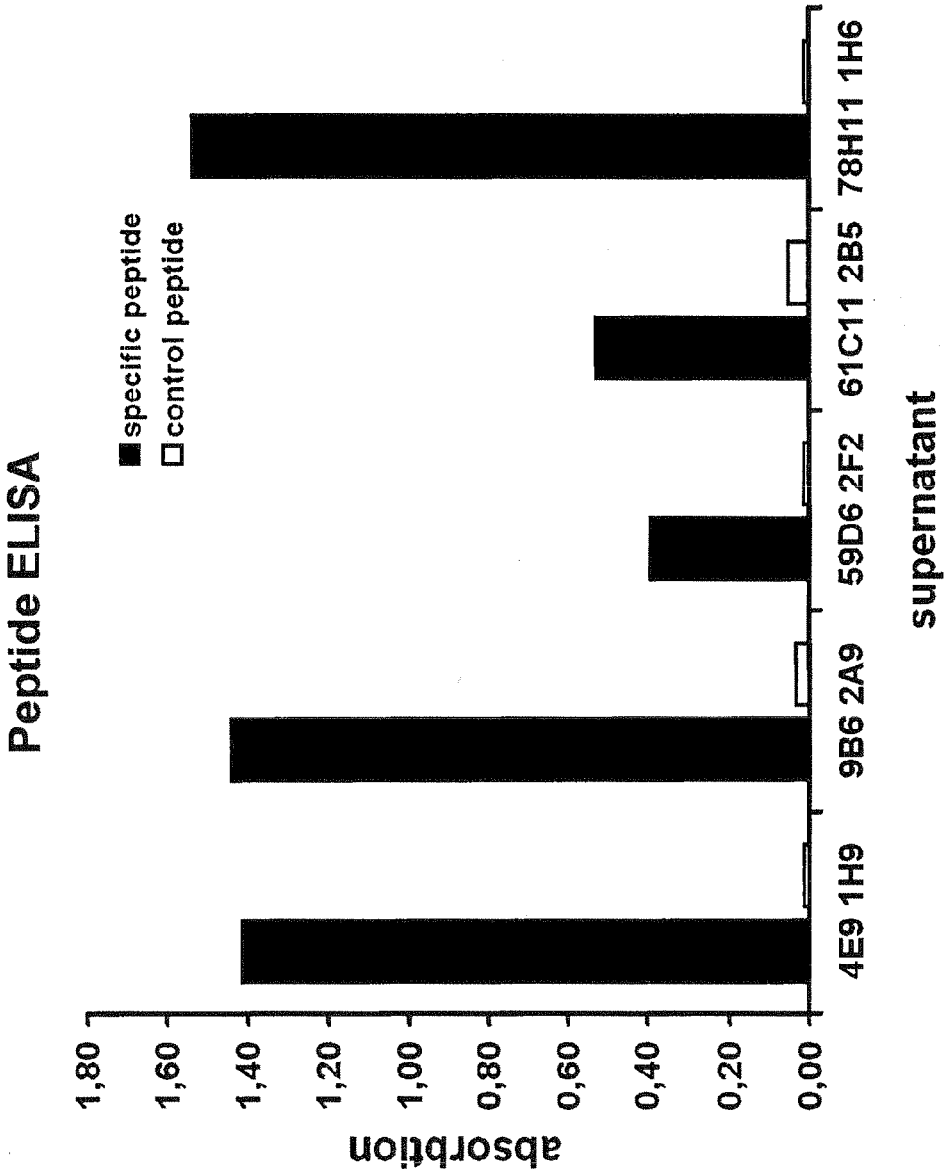

Peptide ELISA
specific peptide
control peptide
absorbtion
1,80
1,60
1,40
1,20
1,00
0,80
0,60
0,40
0,20
0,00
4E9 1H9
9B6 2A9
59D6 2F2
61C11 2B5
78H11 1H6
supernatant

Fig. 8

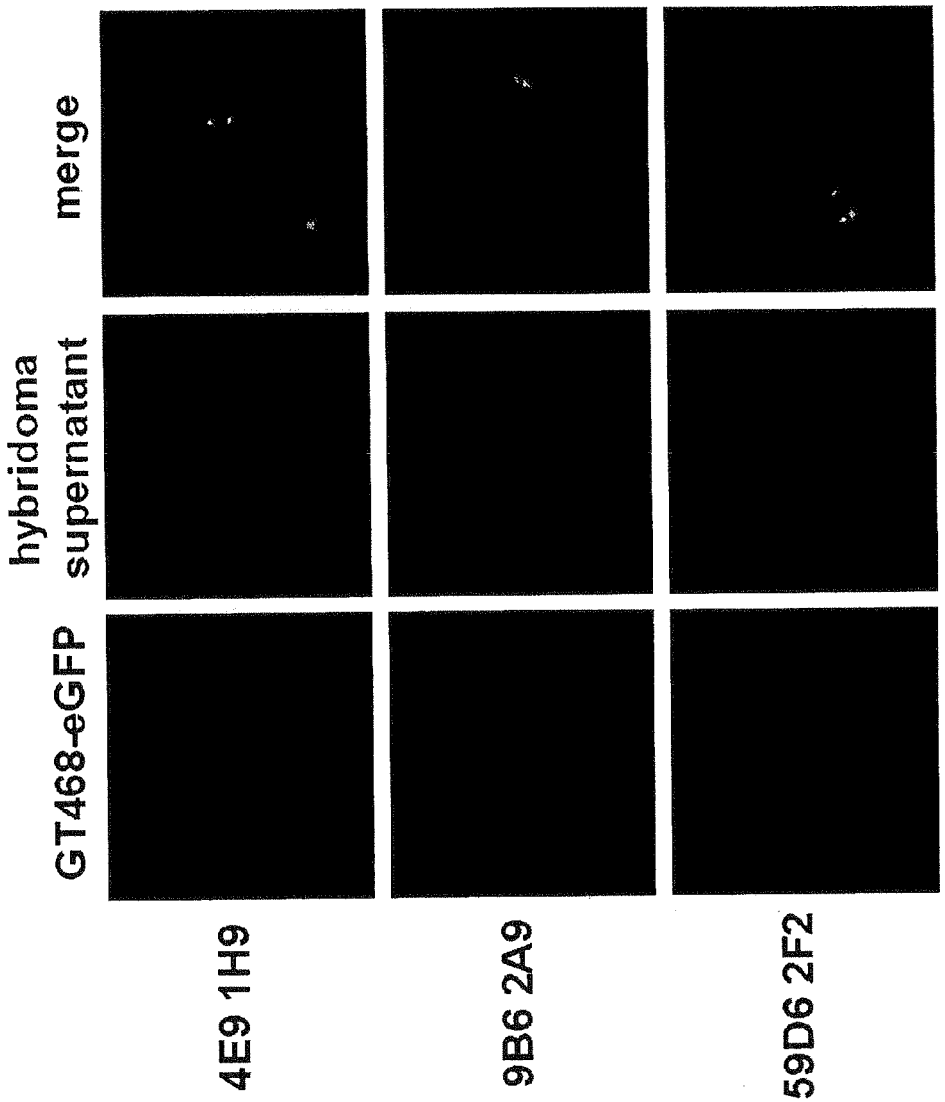

merge
hybridoma supernatant
GT468-eGFP
4E9 1H9
9B6 2A9
59D6 2F2

Fig. 9

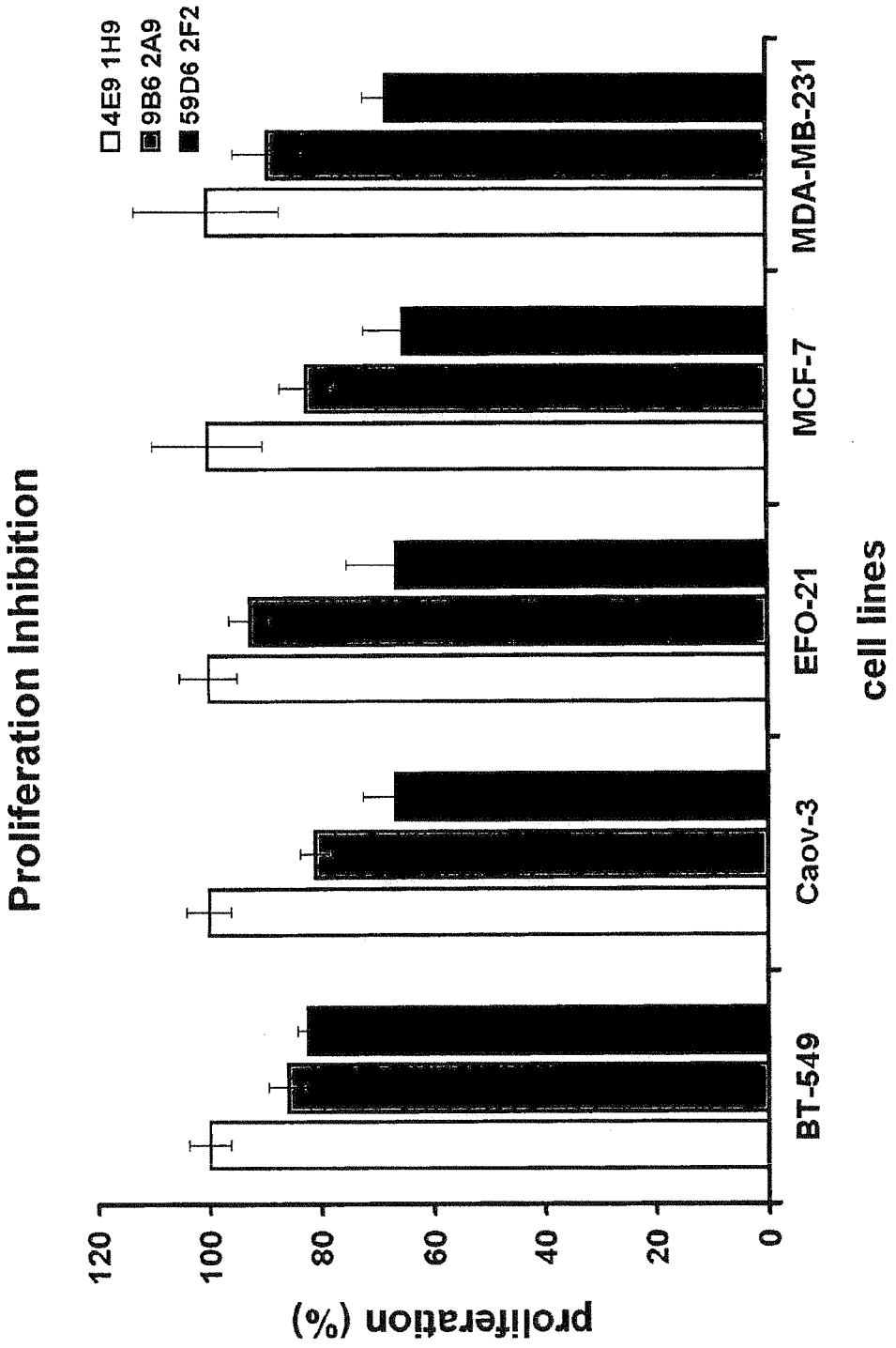

Proliferation Inhibition
4E9 1H9
9B6 2A9
59D6 2F2
proliferation (%)
120
100
80
60
40
20
0
BT-549
Caov-3
EFO-21
MCF-7
MDA-MB-231
cell lines

Fig. 10

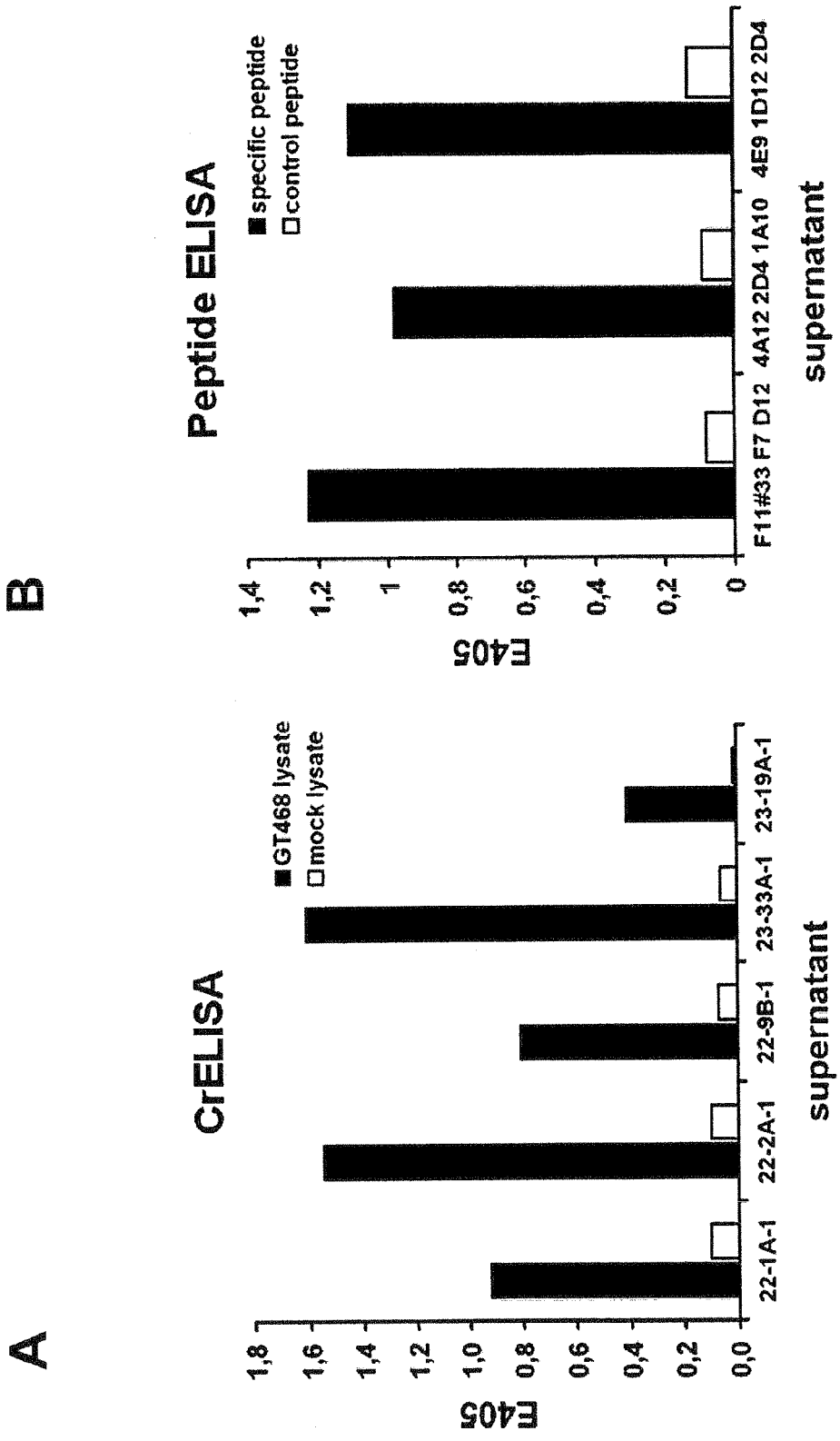

A
CrELISA
GT468 lysate
mock lysate
E405
1,8
1,6
1,4
1,2
1,0
0,8
0,6
0,4
0,2
0,0
22-1A-1
22-2A-1
22-9B-1
23-33A-1
23-19A-1
supernatant
B
Peptide ELISA
specific peptide
control peptide
E405
1,4
1,2
1
0,8
0,6
0,4
0,2
0
F11#33 F7 D12
4A12 2D4 1A10
4E9 1D12 2D4
supernatant

Fig. 11

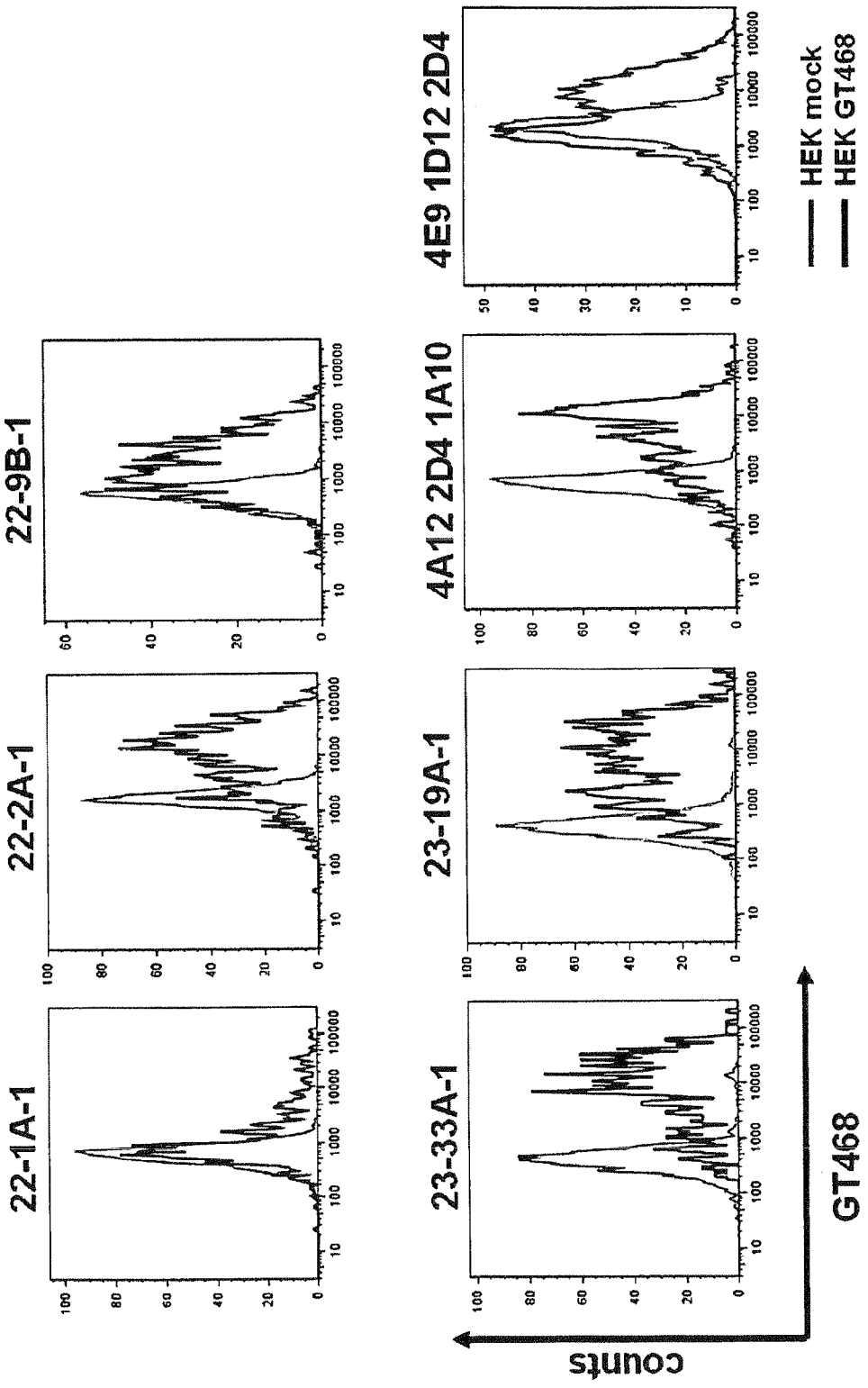

22-1A-1
22-2A-1
22-9B-1
23-33A-1
23-19A-1
4A12 2D4 1A10
4E9 1D12 2D4
HEK mock
HEK GT468
counts
GT468

Fig. 12

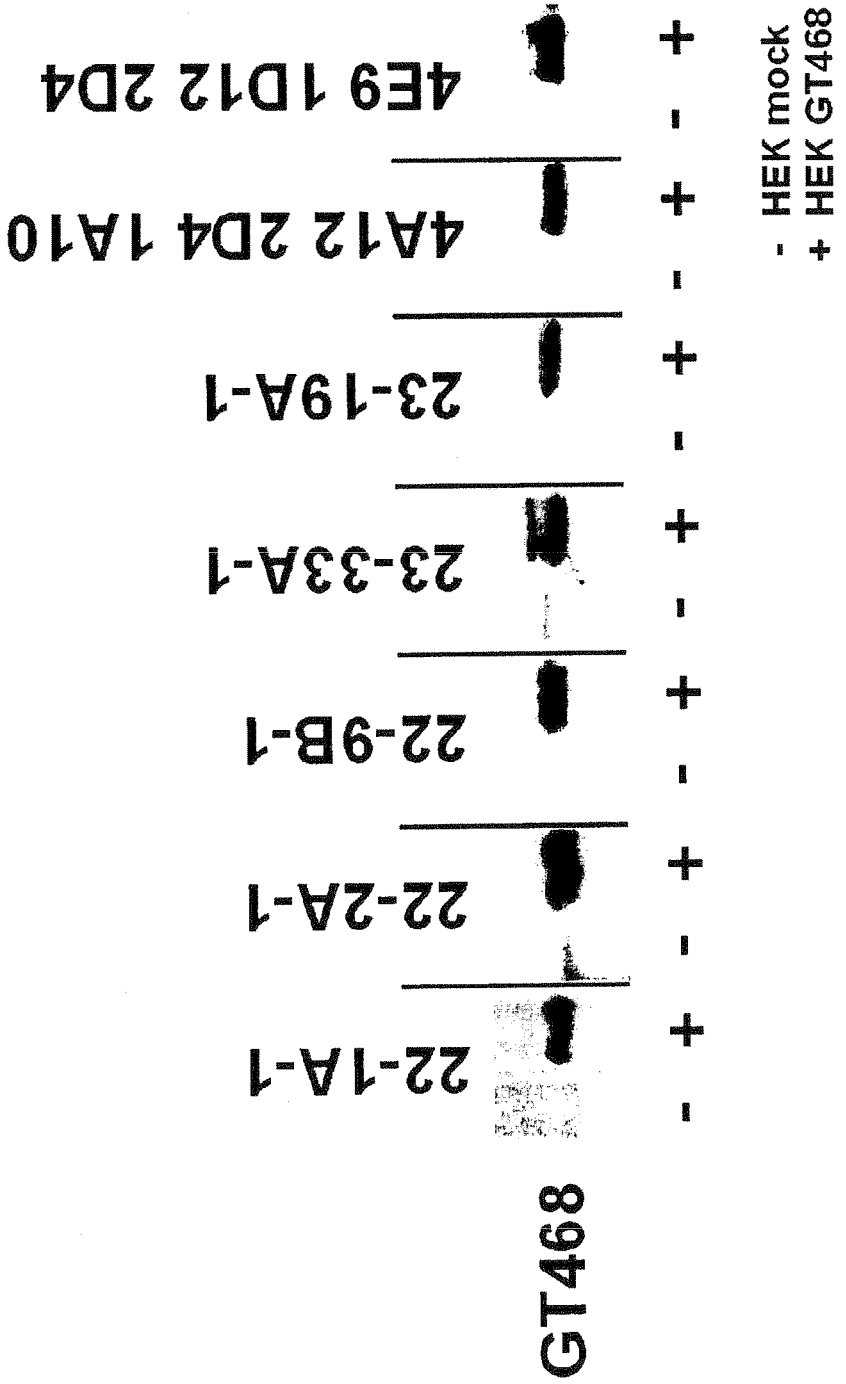

- HEK mock
+ HEK GT468
GT468
22-1A-1
22-2A-1
22-9B-1
23-33A-1
23-19A-1
4A12 2D4 1A10
4E9 1D12 2D4

Fig. 13

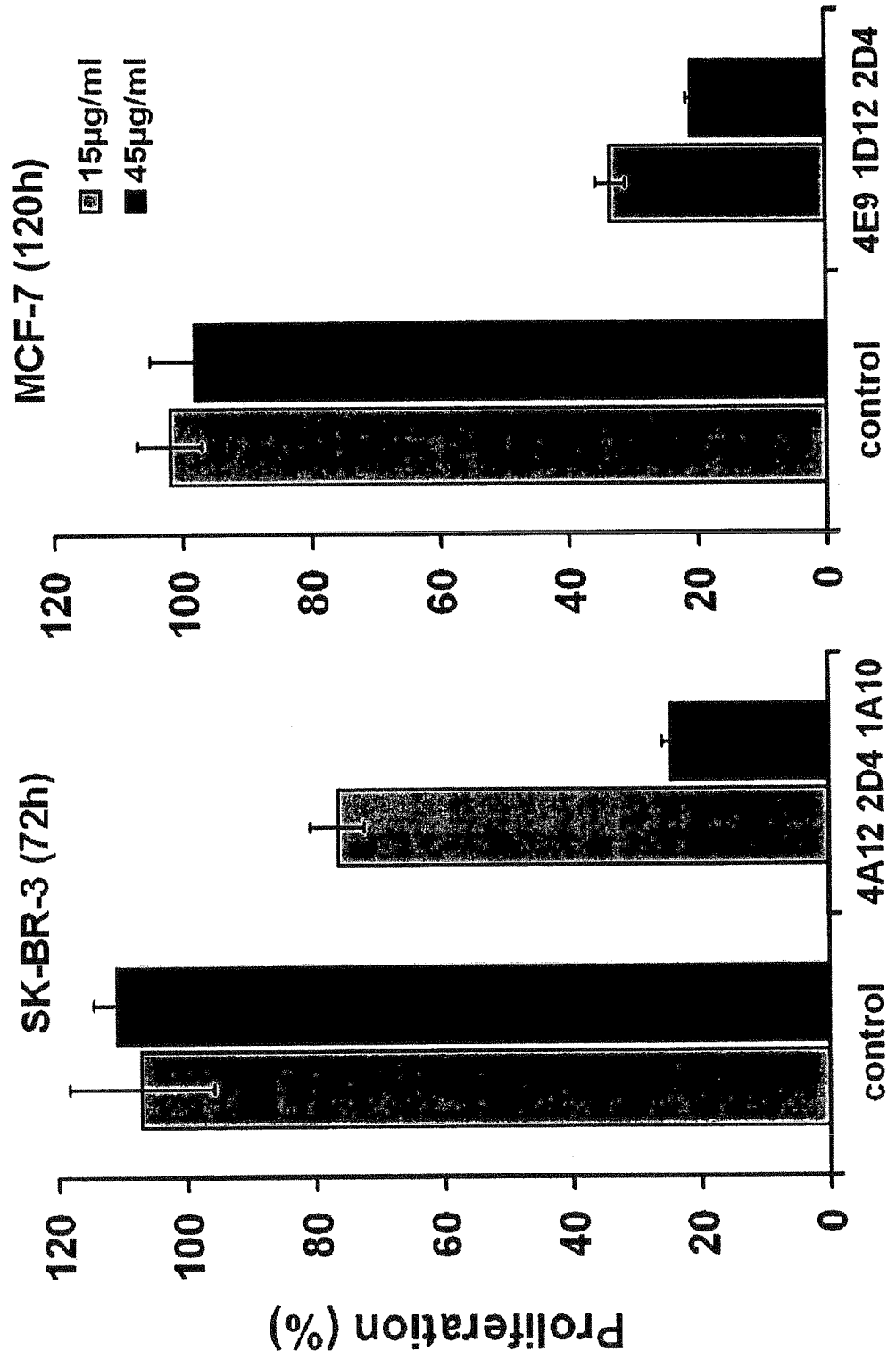
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1	MKVFKFIGLMILLTS
2	KFIGLMILLTSAFSA
3	LMILLTSAFSAGSGQ
4	LTSAFSAGSGQSPMT
5	FSAGSGQSPMTVLCS
6	SGQSPMTVLCSIDWF
7	PMTVLCSIDWFMVTV
8	LCSIDWFMVTVHPFM
9	DWFMVTVHPFMLNND
10	VTVHPFMLNNDVCVH
11	PFMLNNDVCVHFHEL
12	NNDVCVHFHELHLGL
13	CVHFHELHLGLGCPP
14	HELHLGLGCPPNHVQ
15	LGLGCPPNHVQPHAY
16	CPPNHVQPHAYQFTY
17	HVQPHAYQFTYRVTE
18	HAYQFTYRVTECGIR
19	FTYRVTECGIRAKAV
20	VTECGIRAKAVSQDM
21	GIRAKAVSQDMVIYS
22	KAVSQDMVIYSTEIH
23	QDMVIYSTEIHYSSK
24	IYSTEIHYSSKGTPS
25	EIHYSSKGTPSKFVI

Peptide	Sequence
26	SSKGTPSKFVIPVSC
27	TPSKFVIPVSCAAPQ
28	FVIPVSCAAPQKSPW
29	VSCAAPQKSPWLTKP
30	APQKSPWLTKPCSMR
31	SPWLTKPCSMRVASK
32	TKPCSMRVASKSRAT
33	SMRVASKSRATAQKD
34	ASKSRATAQKDEKCY
35	RATAQKDEKCYEVFS
36	QKDEKCYEVFSLSQS
37	KCYEVFSLSQSSQRP
38	VFSLSQSSQRPNCDC
39	SQSSQRPNCDCPPCV
40	QRPNCDCPPCVFSEE
41	CDCPPCVFSEEEHTQ
42	PCVFSEEEHTQVPCH
43	SEEEHTQVPCHQAGA
44	HTQVPCHQAGAQEAQ
45	PCHQAGAQEAQPLQP
46	AGAQEAQPLQPSHFL
47	EAQPLQPSHFLDISE
48	LQPSHFLDISEDWSL
49	HFLDISEDWSLHTDD
50	ISEDWSLHTDDMIGS
51	SEDWSLHTDDMIGSM

Hybridoma	Reactivity (Peptide)
22-1A-1	50+51
22-2A-1	18, 26-31, 42
22-9B-1	27, 29-31
23-33A-1	47+48
23-19A-1	48+49

Hybridoma	Reactivity (Sequence)
F11#33 F7 D12	VFSLSQSSQRPNC
4A12 2D4 1A10	APQKSPWLTKPC
4A9 1D12 2D4	APQKSPWLTKPC

Fig. 14


Proliferation (%)
SK-BR-3 (72h)
MCF-7 (120h)
15µg/ml
45µg/ml
0
20
40
60
80
100
120
control
4A12 2D4 1A10
4E9 1D12 2D4

SEQUENCE LISTING

<110> Ganymed Pharmaceuticals AG et al.

<120> MONOCLONAL ANTIBODIES FOR TREATMENT OF CANCER

<130> 342-35PCT

<150> EP07005258.4
<151> 2007-03-14

<150> US60/894,860
<151> 2007-03-14

<160> 79

<170> PatentIn version 3.3

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<211> 1126
<212> DNA
<213> Homo sapiens

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acccctcttc agttccagtg actgagatac atttttccaa tcctgggggc aaatacagac 120

acagcaagtt ccttcttccc tttggaaatt tggcagctgc cttcaccagt gagcacaaag 180

ccacatttca aaggaaactg acaaattatc cccagctgcc agaagaagaa atcctcactg 240

gacggcttcc tgtttcctgt ggttcattat ctgattggct gcagggatga aagtttttaa 300

gttcatagga ctgatgatcc tcctcacctc tgcgttttca gccggttcag gacaaagtcc 360

aatgactgtg ctgtgctcca tagactggtt catggtcaca gtgcaccccc tcatgctaaa 420

caacgatgtg tgtgtacact ttcatgaact acacttgggc ctgggttgcc ccccaaacca 480

tgttcagcca cacgcctacc agttcaccta ccgtgttact gaatgtggca tcagggccaa 540

agctgtctct caggacatgg ttatctacag cactgagata cactactctt ctaagggcac 600

gccatctaag tttgtgatcc cagtgtcatg tgctgccccc caaaagtccc catggctcac 660

caagccctgc tccatgagag tagccagcaa gagcagggcc acagcccaga aggatgagaa 720

atgctacgag gtgttcagct tgtcacagtc cagtcaaagg cccaactgcg attgtccacc 780

ttgtgtcttc agtgaagaag agcataccca ggtcccttgt caccaagcag gggctcagga 840

ggctcaacct ctgcagccat ctcactttct tgatatttct gaggattggt ctcttcacac 900

agatgatatg attgggtcca tgtgatcctc aggtttgggg tctcctgaag atgctatttc 960

tagaattagt atatagtgta caaatgtctg acaaataagt gctcttgtga ccctcatgtg 1020

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ttatattcat gtgtaaacaa aaaataaaat aaaattctga tcgcat 1126

<210> 2

<211> 212
<212> PRT
<213> Homo sapiens

<400> 2

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

Phe Ser Ala Gly Ser Gly Gln Ser Pro Met Thr Val Leu Cys Ser Ile
20 25 30

Asp Trp Phe Met Val Thr Val His Pro Phe Met Leu Asn Asn Asp Val
35 40 45

Cys Val His Phe His Glu Leu His Leu Gly Leu Gly Cys Pro Pro Asn
50 55 60

His Val Gln Pro His Ala Tyr Gln Phe Thr Tyr Arg Val Thr Glu Cys
65 70 75 80

Gly Ile Arg Ala Lys Ala Val Ser Gln Asp Met Val Ile Tyr Ser Thr
85 90 95

Glu Ile His Tyr Ser Ser Lys Gly Thr Pro Ser Lys Phe Val Ile Pro
100 105 110

Val Ser Cys Ala Ala Pro Gln Lys Ser Pro Trp Leu Thr Lys Pro Cys
115 120 125

Ser Met Arg Val Ala Ser Lys Ser Arg Ala Thr Ala Gln Lys Asp Glu
130 135 140

Lys Cys Tyr Glu Val Phe Ser Leu Ser Gln Ser Ser Gln Arg Pro Asn
145 150 155 160

Cys Asp Cys Pro Pro Cys Val Phe Ser Glu Glu Glu His Thr Gln Val
165 170 175

Pro Cys His Gln Ala Gly Ala Gln Glu Ala Gln Pro Leu Gln Pro Ser
180 185 190

His Phe Leu Asp Ile Ser Glu Asp Trp Ser Leu His Thr Asp Asp Met
195 200 205

Ile Gly Ser Met
210

<210> 3
<211> 12

<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 3

Ala Pro Gln Lys Ser Pro Trp Leu Thr Lys Pro Cys
1 5 10

<210> 4
<211> 13
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 4

Pro Leu Gln Pro Ser His Phe Leu Asp Ile Ser Glu Asp
1 5 10

<210> 5
<211> 11
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 5

Ile Tyr Ser Thr Glu Ile His Tyr Ser Ser Lys
1 5 10

<210> 6
<211> 12
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 6

Trp Ser Leu His Thr Asp Asp Met Ile Gly Ser Met
1 5 10

<210> 7
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 7

Cys Ser Ile Asp Trp Phe Met Val Thr Val His Pro Phe Met Leu
1 5 10 15

<210> 8
<211> 12
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 8

Cys Val His Phe His Glu Leu His Leu Gly Leu Gly
1 5 10

<210> 9
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 9

Ser Ser Lys Gly Thr Pro Ser Lys Phe Val Ile Pro Val Ser Cys
1 5 10 15

<210> 10
<211> 13
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Immunization

<400> 10

Val Phe Ser Leu Ser Gln Ser Ser Gln Arg Pro Asn Cys
1 5 10

<210> 11
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: siRNA

<400> 11
ccaugagagu agccagcaat t 21

<210> 12
<211> 21
<212> DNA
<213> Artificial

<220>

<223> Description of artificial sequence: siRNA

<400> 12
uugcuggcua cucucaugga g 21

<210> 13
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: siRNA

<400> 13
gguucaggac aaaguccaat t 21

<210> 14
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: siRNA

<400> 14
uuggacuuug uccugaaccg g 21

<210> 15
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: Oligonucleotide

<400> 15
aaatttggca gctgccttca c 21

<210> 16
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: Oligonucleotide

<400> 16
tgatgccaca ttcagtaaca c 21

<210> 17
<211> 326
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Translation of PCR product

<400> 17

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1               5                  10                  15

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

Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
        35                  40                  45

Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
    50                  55                  60

Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
65                  70                  75                  80

Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Lys Val Glu Pro
                85                  90                  95

Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
            100                 105                 110

Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
        115                 120                 125

Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
    130                 135                 140

Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
145                 150                 155                 160

Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
                165                 170                 175

Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
            180                 185                 190

Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro
        195                 200                 205

Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
    210                 215                 220

Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu Leu Thr Lys Asn
225                 230                 235                 240

Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
                245                 250                 255
```

Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
260 265 270

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

Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
290 295 300

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

Ser Leu Ser Pro Gly Lys
325

<210> 18
<211> 107
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Translation of PCR product

<400> 18

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1 5 10 15

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
20 25 30

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
35 40 45

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
50 55 60

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
65 70 75 80

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
85 90 95

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
100 105

<210> 19
<211> 324
<212> DNA
<213> Artificial

<220>

<223> Description of artificial sequence: PCR product

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tggaaggtgg ataacgccct ccaatcgggt aactcccagg agagtgtcac agagcaggac 180

agcaaggaca gcacctacag cctcagcagc accctgacgc tgagcaaagc agactacgag 240

aaacacaaag tctacgcctg cgaagtcacc catcagggcc tgagctcgcc cgtcacaaag 300

agcttcaaca ggggagagtg ttag 324

<210> 20
<211> 981
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: PCR product

<400> 20
ggcccatcgg tcttcccccc ggcaccctcc tccaagagca cctctggggg cacagcggcc 60

ctgggctgcc tggtcaagga ctacttcccc gaaccggtga cggtgtcgtg gaactcaggc 120

gccctgacca gcggcgtgca caccttcccg gctgtcctac agtcctcagg actctactcc 180

ctcagcagcg tggtgaccgt gccctccagc agcttgggca cccagaccta catctgcaac 240

gtgaatcaca agcccagcaa caccaaggtg gacaagaaag ttgagcccaa atcttgtgac 300

aaaactcaca catgcccacc gtgcccagca cctgaactcc tgggggggacc gtcagtcttc 360

ctcttccccc caaaacccaa ggacaccctc atgatctccc ggaccccctga ggtcacatgc 420

gtggtggtgg acgtgagcca cgaagaccct gaggtcaagt tcaactggta cgtggacggc 480

gtggaggtgc ataatgccaa gacaaagccg cgggaggagc agtacaacag cacgtaccgt 540

gtggtcagcg tcctcaccgt cctgcaccag gactggctga atggcaagga gtacaagtgc 600

aaggtctcca acaaagccct cccagccccc atcgagaaaa ccatctccaa agccaaaggg 660

cagccccgag aaccacaggt gtacaccctg cccccatccc gggatgagct gaccaagaac 720

caggtcagcc tgacctgcct ggtcaaaggc ttctatccca gcgacatcgc cgtggagtgg 780

gagagcaatg ggcagccgga gaacaactac aagaccacgc ctcccgtgct ggactccgac 840

ggctccttct tcctctatag caagctcacc gtggacaaga gcaggtggca gcaggggaac 900

gtcttctcat gctccgtgat gcatgaggct ctgcacaacc actacacgca gaagagcctc 960

tccctgtctc cgggtaaatg a 981

<210> 21
<211> 324
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: codon optimized nucleic acid

<400> 21
cgtacggtgg ccgctcccag cgtgttcatc ttcccccca gcgacgagca gctgaagtcc 60

ggcaccgcca gcgtggtgtg cctgctgaac aacttctacc cccgggaggc caaggtgcag 120

tggaaggtgg acaacgccct gcagagcggc aacagccagg agagcgtcac cgagcaggac 180

agcaaggact ccacctacag cctgagcagc accctgaccc tgagcaaggc cgactacgag 240

aagcacaagg tgtacgcctg cgaggtgacc caccagggcc tgtccagccc cgtgaccaag 300

agcttcaaca ggggcgagtg ctag 324

<210> 22
<211> 107
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: codon optimized protein

<400> 22

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
1 5 10 15

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
20 25 30

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
35 40 45

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
50 55 60

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
65 70 75 80

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
85 90 95

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
100 105

<210> 23
<211> 981
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: codon optimized nucleic acid

<400> 23
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ctgggctgcc tggtgaagga ctacttcccc gagcccgtga ccgtgagctg gaacagcgga 120

gccctgacct ccggcgtgca caccttcccc gccgtgctgc agagcagcgg cctgtacagc 180

ctgagcagcg tggtgaccgt gcccagcagc agcctgggca cccagaccta catctgcaac 240

gtgaaccaca agcccagcaa caccaaggtg gacaagagag tggagcccaa gagctgcgac 300

aagacccaca cctgcccccc ctgcccagcc ccagagctgc tgggcggacc cagcgtgttc 360

ctgttccccc ccaagcccaa ggacaccctg atgatcagca ggacccccga ggtgacctgc 420

gtggtggtgg acgtgagcca cgaggaccca gaggtgaagt tcaactggta cgtggacggc 480

gtggaggtgc acaacgccaa gaccaagccc agagaggagc agtacaacag cacctacagg 540

gtggtgtccg tgctgaccgt gctgcaccag gactggctga acggcaagga atacaagtgc 600

aaggtctcca acaaggccct gccagccccc atcgaaaaga ccatcagcaa ggccaagggc 660

cagccacggg agccccaggt gtacaccctg ccccccagcc gggaggagat gaccaagaac 720

caggtgtccc tgacctgtct ggtgaagggc ttctacccca gcgacatcgc cgtggagtgg 780

gagagcaacg gccagcccga gaacaactac aagaccaccc ccccagtgct ggacagcgac 840

ggcagcttct tcctgtacag caagctgacc gtggacaagt ccaggtggca gcagggcaac 900

gtgttcagct gcagcgtgat gcacgaggcc ctgcacaacc actacaccca gaagtccctg 960

agcctgagcc ccggcaagta g 981

<210> 24
<211> 326
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: codon optimized protein

<400> 24

Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly
1 5 10 15

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

Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
35 40 45

Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
50 55 60

Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
65 70 75 80

Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro
85 90 95

Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
100 105 110

Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
115 120 125

Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
130 135 140

Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
145 150 155 160

Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
165 170 175

Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
180 185 190

Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro
195 200 205

Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
210 215 220

Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn
225 230 235 240

Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
245 250 255

Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
260 265 270

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

Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
290 295 300

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

Ser Leu Ser Pro Gly Lys
325

<210> 25
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: Oligonucleotide

<400> 25
tgacactggc aaaacaatgc a 21

<210> 26
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: Oligonucleotide

<400> 26
ggtccttttc accagcaagc t 21

<210> 27
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: siRNA

<400> 27
uaacuguaua aucgacuagt t 21

<210> 28
<211> 21
<212> DNA
<213> Artificial

<220>
<223> Description of artificial sequence: siRNA

<400> 28
cuagucgauu auacaguuag a 21

<210> 29
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 29

Met Lys Val Phe Lys Phe Ile Gly Leu Met Ile Leu Leu Thr Ser
1 5 10 15

<210> 30
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 30

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

<210> 31
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 31

Leu Met Ile Leu Leu Thr Ser Ala Phe Ser Ala Gly Ser Gly Gln
1 5 10 15

<210> 32
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 32

Leu Thr Ser Ala Phe Ser Ala Gly Ser Gly Gln Ser Pro Met Thr
1 5 10 15

<210> 33
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 33

Phe Ser Ala Gly Ser Gly Gln Ser Pro Met Thr Val Leu Cys Ser
1 5 10 15

<210> 34
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 34

Ser Gly Gln Ser Pro Met Thr Val Leu Cys Ser Ile Asp Trp Phe
1 5 10 15

<210> 35
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 35

Pro Met Thr Val Leu Cys Ser Ile Asp Trp Phe Met Val Thr Val
1 5 10 15

<210> 36
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 36

Leu Cys Ser Ile Asp Trp Phe Met Val Thr Val His Pro Phe Met
1 5 10 15

<210> 37
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 37

Asp Trp Phe Met Val Thr Val His Pro Phe Met Leu Asn Asn Asp
1 5 10 15

<210> 38
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 38

Val Thr Val His Pro Phe Met Leu Asn Asn Asp Val Cys Val His
1 5 10 15

<210> 39
<211> 15

<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 39

Pro Phe Met Leu Asn Asn Asp Val Cys Val His Phe His Glu Leu
1 5 10 15

<210> 40
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 40

Asn Asn Asp Val Cys Val His Phe His Glu Leu His Leu Gly Leu
1 5 10 15

<210> 41
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 41

Cys Val His Phe His Glu Leu His Leu Gly Leu Gly Cys Pro Pro
1 5 10 15

<210> 42
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 42

His Glu Leu His Leu Gly Leu Gly Cys Pro Pro Asn His Val Gln
1 5 10 15

<210> 43
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 43

Leu Gly Leu Gly Cys Pro Pro Asn His Val Gln Pro His Ala Tyr
1 5 10 15

<210> 44
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 44

Cys Pro Pro Asn His Val Gln Pro His Ala Tyr Gln Phe Thr Tyr
1 5 10 15

<210> 45
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 45

His Val Gln Pro His Ala Tyr Gln Phe Thr Tyr Arg Val Thr Glu
1 5 10 15

<210> 46
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 46

His Ala Tyr Gln Phe Thr Tyr Arg Val Thr Glu Cys Gly Ile Arg
1 5 10 15

<210> 47
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 47

Phe Thr Tyr Arg Val Thr Glu Cys Gly Ile Arg Ala Lys Ala Val
1 5 10 15

<210> 48
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 48

Val Thr Glu Cys Gly Ile Arg Ala Lys Ala Val Ser Gln Asp Met
1 5 10 15

<210> 49
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 49

Gly Ile Arg Ala Lys Ala Val Ser Gln Asp Met Val Ile Tyr Ser
1 5 10 15

<210> 50
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 50

Lys Ala Val Ser Gln Asp Met Val Ile Tyr Ser Thr Glu Ile His
1 5 10 15

<210> 51
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 51

Gln Asp Met Val Ile Tyr Ser Thr Glu Ile His Tyr Ser Ser Lys
1 5 10 15

<210> 52
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 52

Ile Tyr Ser Thr Glu Ile His Tyr Ser Ser Lys Gly Thr Pro Ser
1 5 10 15

<210> 53
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 53

Glu Ile His Tyr Ser Ser Lys Gly Thr Pro Ser Lys Phe Val Ile
1 5 10 15

<210> 54
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 54

Ser Ser Lys Gly Thr Pro Ser Lys Phe Val Ile Pro Val Ser Cys
1 5 10 15

<210> 55
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 55

Thr Pro Ser Lys Phe Val Ile Pro Val Ser Cys Ala Ala Pro Gln
1 5 10 15

<210> 56
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 56

Phe Val Ile Pro Val Ser Cys Ala Ala Pro Gln Lys Ser Pro Trp
1 5 10 15

<210> 57
<211> 15
<212> PRT
<213> Artificial

<220>

<223> Description of artificial sequence: Peptide for Screening

<400> 57

Val Ser Cys Ala Ala Pro Gln Lys Ser Pro Trp Leu Thr Lys Pro
1 5 10 15

<210> 58
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 58

Ala Pro Gln Lys Ser Pro Trp Leu Thr Lys Pro Cys Ser Met Arg
1 5 10 15

<210> 59
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 59

Ser Pro Trp Leu Thr Lys Pro Cys Ser Met Arg Val Ala Ser Lys
1 5 10 15

<210> 60
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 60

Thr Lys Pro Cys Ser Met Arg Val Ala Ser Lys Ser Arg Ala Thr
1 5 10 15

<210> 61
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 61

Ser Met Arg Val Ala Ser Lys Ser Arg Ala Thr Ala Gln Lys Asp
1 5 10 15

<210> 62
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 62

Ala Ser Lys Ser Arg Ala Thr Ala Gln Lys Asp Glu Lys Cys Tyr
1 5 10 15

<210> 63
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 63

Arg Ala Thr Ala Gln Lys Asp Glu Lys Cys Tyr Glu Val Phe Ser
1 5 10 15

<210> 64
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 64

Gln Lys Asp Glu Lys Cys Tyr Glu Val Phe Ser Leu Ser Gln Ser
1 5 10 15

<210> 65
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 65

Lys Cys Tyr Glu Val Phe Ser Leu Ser Gln Ser Ser Gln Arg Pro
1 5 10 15

<210> 66
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 66

Val Phe Ser Leu Ser Gln Ser Ser Gln Arg Pro Asn Cys Asp Cys
1 5 10 15

<210> 67
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 67

Ser Gln Ser Ser Gln Arg Pro Asn Cys Asp Cys Pro Pro Cys Val
1 5 10 15

<210> 68
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 68

Gln Arg Pro Asn Cys Asp Cys Pro Pro Cys Val Phe Ser Glu Glu
1 5 10 15

<210> 69
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 69

Cys Asp Cys Pro Pro Cys Val Phe Ser Glu Glu Glu His Thr Gln
1 5 10 15

<210> 70
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 70

Pro Cys Val Phe Ser Glu Glu Glu His Thr Gln Val Pro Cys His
1 5 10 15

<210> 71
<211> 15

<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 71

Ser Glu Glu Glu His Thr Gln Val Pro Cys His Gln Ala Gly Ala
1 5 10 15

<210> 72
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 72

His Thr Gln Val Pro Cys His Gln Ala Gly Ala Gln Glu Ala Gln
1 5 10 15

<210> 73
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 73

Pro Cys His Gln Ala Gly Ala Gln Glu Ala Gln Pro Leu Gln Pro
1 5 10 15

<210> 74
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 74

Ala Gly Ala Gln Glu Ala Gln Pro Leu Gln Pro Ser His Phe Leu
1 5 10 15

<210> 75
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 75

Glu Ala Gln Pro Leu Gln Pro Ser His Phe Leu Asp Ile Ser Glu
1 5 10 15

<210> 76
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 76

Leu Gln Pro Ser His Phe Leu Asp Ile Ser Glu Asp Trp Ser Leu
1 5 10 15

<210> 77
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 77

His Phe Leu Asp Ile Ser Glu Asp Trp Ser Leu His Thr Asp Asp
1 5 10 15

<210> 78
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 78

Ile Ser Glu Asp Trp Ser Leu His Thr Asp Asp Met Ile Gly Ser
1 5 10 15

<210> 79
<211> 15
<212> PRT
<213> Artificial

<220>
<223> Description of artificial sequence: Peptide for Screening

<400> 79

Ser Glu Asp Trp Ser Leu His Thr Asp Asp Met Ile Gly Ser Met
1 5 10 15

BUDAPEST TREATY ON THE INTERNATIONAL
RECOGNITION OF THE DEPOSIT OF MICROORGANISMS
FOR THE PURPOSES OF PATENT PROCEDURE

DSMZ
Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH

INTERNATIONAL FORM

Ganymed Pharmaceuticals AG
Freiligrathstr. 12
55131 Mainz
Deutschland

RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
issued pursuant to Rule 7.1 by the
INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 22-1A-1	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2895

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit) and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

Form DSMZ-BP/4 (sole page) 08/2006

BUDAPEST TREATY ON THE INTERNATIONAL
RECOGNITION OF THE DEPOSIT OF MICROORGANISMS
FOR THE PURPOSES OF PATENT PROCEDURE

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Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH

INTERNATIONAL FORM

Ganymed Pharmaceuticals AG
Freiligrathstr. 12
55131 Mainz
Deutschland

RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
issued pursuant to Rule 7.1 by the
INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 22-2A-1	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2893

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

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and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

Form DSMZ-BP/4 (sole page) 08/2006

BUDAPEST TREATY ON THE INTERNATIONAL
RECOGNITION OF THE DEPOSIT OF MICROORGANISMS
FOR THE PURPOSES OF PATENT PROCEDURE

DSMZ
Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH

INTERNATIONAL FORM

Ganymed Pharmaceuticals AG
Freiligrathstr. 12
55131 Mainz
Deutschland

RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
issued pursuant to Rule 7.1 by the
INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 22-9B-1	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2896

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

BUDAPEST TREATY ON THE INTERNATIONAL
RECOGNITION OF THE DEPOSIT OF MICROORGANISMS
FOR THE PURPOSES OF PATENT PROCEDURE

DSMZ
Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH

INTERNATIONAL FORM

Ganymed Pharmaceuticals AG
Freiligrathstr. 12
55131 Mainz
Deutschland

RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
issued pursuant to Rule 7.1 by the
INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 23-33A-1	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2897

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

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INTERNATIONAL DEPOSITARY AUTHORITY
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I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 23-19A-1	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2891

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Wenz Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

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RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
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INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: F11#33F7D12	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2894

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

BUDAPEST TREATY ON THE INTERNATIONAL
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RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
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INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 4A12 2D4 1A10	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2892

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

BUDAPEST TREATY ON THE INTERNATIONAL
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RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT
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INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. IDENTIFICATION OF THE MICROORGANISM	
Identification reference given by the DEPOSITOR: 4E9 1D12 2D4	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2898

II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION

The microorganism identified under I. above was accompanied by:

( ) a scientific description
( ) a proposed taxonomic designation

(Mark with a cross where applicable).

III. RECEIPT AND ACCEPTANCE

This International Depositary Authority accepts the microorganism identified under I. above, which was received by it on 2008-03-11 (Date of the original deposit)[1].

IV. RECEIPT OF REQUEST FOR CONVERSION

The microorganism identified under I above was received by this International Depositary Authority on (date of original deposit)
and a request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion).

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

[1] Where Rule 6.4 (d) applies, such date is the date on which the status of international depositary authority was acquired.

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Deutschland

VIABILITY STATEMENT
issued pursuant to Rule 10.2 by the
INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Freiligrathstr. 12 Address: 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2891 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

1 Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
2 In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
3 Mark with a cross the applicable box.
4 Fill in if the information has been requested and if the results of the test were negative.

BUDAPEST TREATY ON THE INTERNATIONAL
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Deutschland

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I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2892 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

1 Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
2 In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
3 Mark with a cross the applicable box.
4 Fill in if the information has been requested and if the results of the test were negative.

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I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2893 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

[1] Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
[2] In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
[3] Mark with a cross the applicable box.
[4] Fill in if the information has been requested and if the results of the test were negative.

BUDAPEST TREATY ON THE INTERNATIONAL
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Freiligrathstr. 12
55131 Mainz
Deutschland

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INTERNATIONAL DEPOSITARY AUTHORITY
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I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2894 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Werz Date: 2008-03-25

1 Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
2 In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
3 Mark with a cross the applicable box.
4 Fill in if the information has been requested and if the results of the test were negative.

BUDAPEST TREATY ON THE INTERNATIONAL
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Freiligrathstr. 12
55131 Mainz
Deutschland

VIABILITY STATEMENT
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INTERNATIONAL DEPOSITARY AUTHORITY
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I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2895 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

[1] Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
[2] In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
[3] Mark with a cross the applicable box.
[4] Fill in if the information has been requested and if the results of the test were negative.

BUDAPEST TREATY ON THE INTERNATIONAL
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55131 Mainz
Deutschland

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INTERNATIONAL DEPOSITARY AUTHORITY
identified at the bottom of this page

I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2896 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. Weiss Date: 2008-03-25

1 Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
2 In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
3 Mark with a cross the applicable box.
4 Fill in if the information has been requested and if the results of the test were negative.

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Deutschland

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INTERNATIONAL DEPOSITARY AUTHORITY
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I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2897 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): U. We-Ls Date: 2008-03-25

[1] Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
[2] In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
[3] Mark with a cross the applicable box.
[4] Fill in if the information has been requested and if the results of the test were negative.

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identified at the bottom of this page

I. DEPOSITOR	II. IDENTIFICATION OF THE MICROORGANISM
Name: Ganymed Pharmaceuticals AG Address: Freiligrathstr. 12 55131 Mainz Deutschland	Accession number given by the INTERNATIONAL DEPOSITARY AUTHORITY: DSM ACC2898 Date of the deposit or the transfer[1]: 2008-03-11

III. VIABILITY STATEMENT

The viability of the microorganism identified under II above was tested on 2008-03-11 [2].
On that date, the said microorganism was

(X)[3] viable

( )[3] no longer viable

IV. CONDITIONS UNDER WHICH THE VIABILITY TEST HAS BEEN PERFORMED[4]

V. INTERNATIONAL DEPOSITARY AUTHORITY

Name: DSMZ-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH Address: Inhoffenstr. 7 B D-38124 Braunschweig	Signature(s) of person(s) having the power to represent the International Depositary Authority or of authorized official(s): V. Weiss Date: 2008-03-25

1 Indicate the date of original deposit or, where a new deposit or a transfer has been made, the most recent relevant date (date of the new deposit or date of the transfer).
2 In the cases referred to in Rule 10.2(a) (ii) and (iii), refer to the most recent viability test.
3 Mark with a cross the applicable box.
4 Fill in if the information has been requested and if the results of the test were negative.