

Supplementary information

A Peptide-based Synthetic Transcription Factor Selectively Activates Transcription in a Mammalian Cell

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Experimental Section

Materials

All amino acids and Rink amide PEGA resin were either purchased from Novabiochem (Switzerland) or Anaspec (China). TFA, phenol and thioanisole were purchased from Merck (Germany). Alexa-488 succinimidyl ester and lipofectamineTM 2000 were purchased from Invitrogen (NY, USA). Luciferase reagent was purchased from Promega (Madison, Wisconsin, USA). C6-aminolink oligonucleotides were purchased from Trilink (USA) and O_R3-O_R3 oligonucleotide was purchased from IDT (USA). The sequences of the DNA used in this study are shown in Table S1.

Peptide Synthesis

Peptide synthesis were followed as described in Dhar et al.¹ Briefly, the peptides were synthesized in solid phase on Rink Amide PEGA Resin in an automated peptide synthesizer (Protein Technology International, AZ, USA). The N terminal amino group of amino acids was Fmoc protected and other reactive side chains were protected with appropriate protecting groups. Standard Fmoc chemistry was used to synthesize the peptides. DMF and piperidine were kept overnight in molecular sieve to produce peptide synthesis grade anhydrous reagent before use. Rink Amide PEGA Resin were swelled in DMF for 45 min. Amino acids were dissolved in DMF followed by addition of 1.7x molar excess of N,N-Diisopropylethylamine (DIPEA) and then transferred to the reaction vessel containing the resin. The coupling reactions were performed under nitrogen gas. Coupling of each amino acid was performed twice with 3-10 fold molar excess of the amino acid in each reaction and coupling times were 20-45 min. The amount of amino acid and the time of coupling reaction depended on the length of the peptide. The resin was washed 3 times with DMF to ensure removal of excess uncoupled amino acids. Fmoc group was removed by mixing with 20% piperidine for 3-10

min and Fmoc removal were repeated twice. The Fmoc deprotection reaction was performed under nitrogen gas pressure. The resin was washed 3 times with DMF to ensure removal of the excess piperidine. Fmoc-Lys(Fmoc)-OH was used as the branching point for the dimer. The dimeric chains were grown simultaneously through α and ϵ amine of Lys. The solid phase resin and side chain protections were removed from peptide by a cocktail of 90% trifluoro acetic acid (TFA), 5% water, 5% phenol and 5% thioanisole for 3 h followed by precipitation with cold diethyl ether. The peptides were purified by RP-HPLC using acetonitrile-water mixture (10-60 % gradient of water to acetonitrile with 0.1% TFA).

Labeling oligonucleotides

The oligonucleotide bearing 5'-C6 amino link was labeled with 10 times molar excess of alexa-488 succinimidyl ester as described previously².

Measurement of Fluorescence Anisotropy

The fluorescence anisotropy measurements were performed with fluorescein labeled 1 nM oligodeoxynucleotide duplexes in 50 mM Tris buffer pH 8.0 in a PTI Quantimaster T-geometry fluorimeter at 25°C. The experiments were carried out in a 0.5cm path length quartz cuvette. The excitation wavelength was at 480 nm and emission was monitored at 530 nm, the excitation and emission band passes were 5 nm each. The anisotropy measurement data were fitted to an equation considering 1:1 binding stoichiometry. The anisotropy values were fitted to the following single site binding equation using Kyplot as described previously³.

$$A_{obs} = A_0 + \frac{[(A_\infty - A_0) * \{(K_d + X + [DNA]) - \sqrt{(K_d + X + [DNA])^2 - 4 \cdot X \cdot [DNA]}\}]}{2 * [DNA]}$$

Where, A_{obs} is the observed anisotropy, A_0 is the initial anisotropy, A_∞ is the final limiting anisotropy, $[DNA]$ is the fluorescein end-labeled DNA concentration, X is the total peptide concentration and K_d is the dissociation constant. All concentrations were expressed in Molar units.

Isothermal Titration Calorimetry

All ITC experiments were done in a VPITC instrument from Microcal.Inc. The titrations were done in 50 mM Tris-HCl buffer (pH 8.0). The peptides (8 μ M) was taken in the cell (volume 1.47 mL), and either O_R3 or C12-O_R3 DNA (80 μ M) was added from a syringe with continuous stirring in 50 mM Tris-HCl buffer, pH 8.0. The data were fitted to a single site binding isotherm using MS Origin 7.0 software as described previously⁴

Cell viability assay

10^4 B16F10 cells/ well were plated in a 96 well plate in 200 μ l volume. Non adherent cells were removed by washing after 16 h. Adherent cells were treated with 5 μ M, 10 μ M, 20 μ M, 30 μ M, 40 μ M and 50 μ M peptides for 36 h. 20 μ l of MTT (stock 5 mg/ml) was added per well and kept for 4 h. Cells were centrifuged and the resulting precipitate was dissolved in 100 μ l of 0.1 (N) HCl in isopropanol. The absorbance was measured at 595 nm.

Confocal microscopy

10^4 B16F10 cell/ well were plated in a 4 well chamber slide (Nunc, UK). 2 μ M FITC labelled peptide was added and incubated for 2 h. Then cells were washed thoroughly and fixed with Ultracruz mounting media containing DAPI (Santacruz Biotechnology, sc-24941). The confocal images were captured in Nikon (A1R). The images were captured at DIC channel (cell morphology), green channel (peptide uptake) and blue channel (nuclear marker).

Cloning of O_R3-O_R3 variants in pGL3

The plasmid pGL3 containing the mouse IL-10 promoter (pGL3-IL-10) was a kind gift from Dr. Syamal Roy. The sequence of the O_R3-O_R3 used in this study is as follows, TATATCCCTTGCGGTGATAGATATTATCCCTTGCGGTGATAT, with KpnI overhang on both the 5' and 3' side. This O_R3-O_R3 sequence was inserted in the KpnI restriction site in the pGL3-IL-10 plasmid (Figure S7A). The plasmid containing multiple O_R3-O_R3 sites is represented as pGL3-nX O_R3 where “n” represents 2, 4 and 6 (Figure S7B).

Transfection and luciferase assay

The plasmids used for transfection were pGL3 (blank), pGL3-IL-10 (control), pGL3-2X O_R3, pGL3-4X O_R3, pGL3-6X O_R3. B16F10 cells were grown in a 6 well plate with 4×10^5 cells/ well. The cells were allowed to adhere for 16 h and non-adherent cells were removed by washing. The transfections were performed with 1 μ g of plasmid using lipofectamine according to manufacturer protocol. Transfected cells were then kept in resting condition for 12-14 h in DMEM with 10% FCS followed by treated with ATF and incubated for 36 h. Floating cells were removed by washing with PBS. Attached cells were removed from the plate using 200 μ L trypsin-EDTA followed by quench with 20 μ L FCS with addition of 500 μ L media. The cells were centrifuged at 400 rcf for 5 min at room temperature. The cell

pellet was resuspended in 1 mL PBS. 20 μ L were kept for cell counting and the rest of the cells were centrifuged for 400 rcf for 5 min at room temperature. The cells were counted by staining with trypan blue. Trypan blue staining showed that > 95% of the cells were viable. The cell count was very similar among the different conditions tested. The total cell count was approximately 2.2×10^6 cells/ well. Thus, neither the cell viability nor the cell count changes significantly due to transfection with different plasmids. The cells were lysed with 200 μ l of passive lysis buffer and 20 μ l of lysate were used to assay luciferase activity using luciferase kit/reagent as manufacturer protocol (Promega). The luciferase activity was normalized with respect to the total cellular protein concentration⁵. To further confirm and reduce the effect of experimental variability, luciferase activity was normalized with respect to viable cell number as well. The luciferase activity obtained by cell number normalization and protein normalization produces similar results.

Whole genome microarray

To study the effect of the wild type peptide on the global transcription of B16F10 DNA, microarray experiments were performed. The treatments were done as described previously. After 36 h cells were resuspended in RNA later solution. The microarray experiments were conducted by Genotypic Inc. (Bangalore, India) on payment for service basis. c-DNA was prepared from extracted RNA and then labeled c-RNA was synthesised by T7-promoter based amplification. Labeled RNA were hybridized *in situ* with 60 mer oligonucleotides. The hybridisation was carried at 65°C for 17 h. The samples were run on slide, Mouse 8x60K Array AMADID, and the data were collected and analysed.

Promoter Analysis

The sequence of the promoter (region near the transcription starts site) of the upregulated (3 fold or more) gene following treatment with Kix-Br-Ala β -Cro were obtained using UCSC

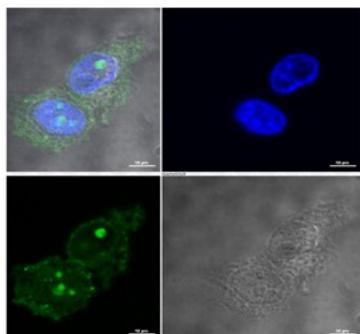
genome browser. The database provides the information regarding the localization of the gene in the genome such as chromosome number and position within the chromosome. Though promoter sequence of the all gene is yet not characterized. We looked for the O_R3 sequence within the promoter with maximum 2 bp mismatch using NCBI sequence alignment tool.

References

1. A. Dhar, S. Mallick, P. Ghosh, A. Maiti, I. Ahmed, S. Bhattacharya, T. Mandal, A. Manna, K. Roy and S. Singh, *Pep Sci*, 2014, **102**, 344-358.
2. S. Chatterjee, Y. N. Zhou, S. Roy and S. Adhya, *Proc Natl Acad Sci U S A*, 1997, **94**, 2957-2962.
3. S. Polley, S. Guha, N. S. Roy, S. Kar, K. Sakaguchi, Y. Chuman, V. Swaminathan, T. Kundu and S. Roy, *J Mol Biol*, 2008, **376**, 8-12.
4. A. Mazumder, A. Maiti, K. Roy and S. Roy, *ACS Chem Biol*, 2012, **7**, 1084-1094.
5. S. Sen, K. Roy, S. Mukherjee, R. Mukhopadhyay and S. Roy, *PLoS pathogens*, 2011, **7**, e1002229.

Figures

A



B

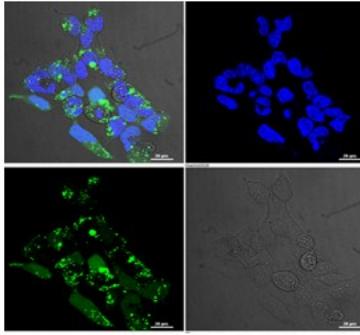


Figure S1. Nuclear localisation of the fluorescein label (A) Kix-Br-Ala β -Cro and (B) No-NLS- Kix-Br-Ala β -Cro. Cells were treated with 2 μ M of FITC labelled peptide (green) for 2 h and nucleus was stained with DAPI (blue). Right lower corner represent DIC image, left lower corner represent green fluorescence of fluorescein, right upper corner represent DAPI fluorescence and left upper corner represent overlay of all the three channel.

Figure S2. Helical wheel diagram for a single peptide chain of the branched peptide used in the study. The output presents the hydrophilic residues as circles, hydrophobic residues as diamonds, and potentially positively charged as pentagons. The red starred residues correspond to residues which make contacts with the Operator DNA (O_{R3}) in the crystal structure (PDB id: 6cro). The blue starred residues were replaced with Aib. The residues here are numbered starting from residue 26 in the Cro (residue 1 here and so on). Thus, residue 30 is I5, residue 34 is I9 and residue 36 is A11 here.

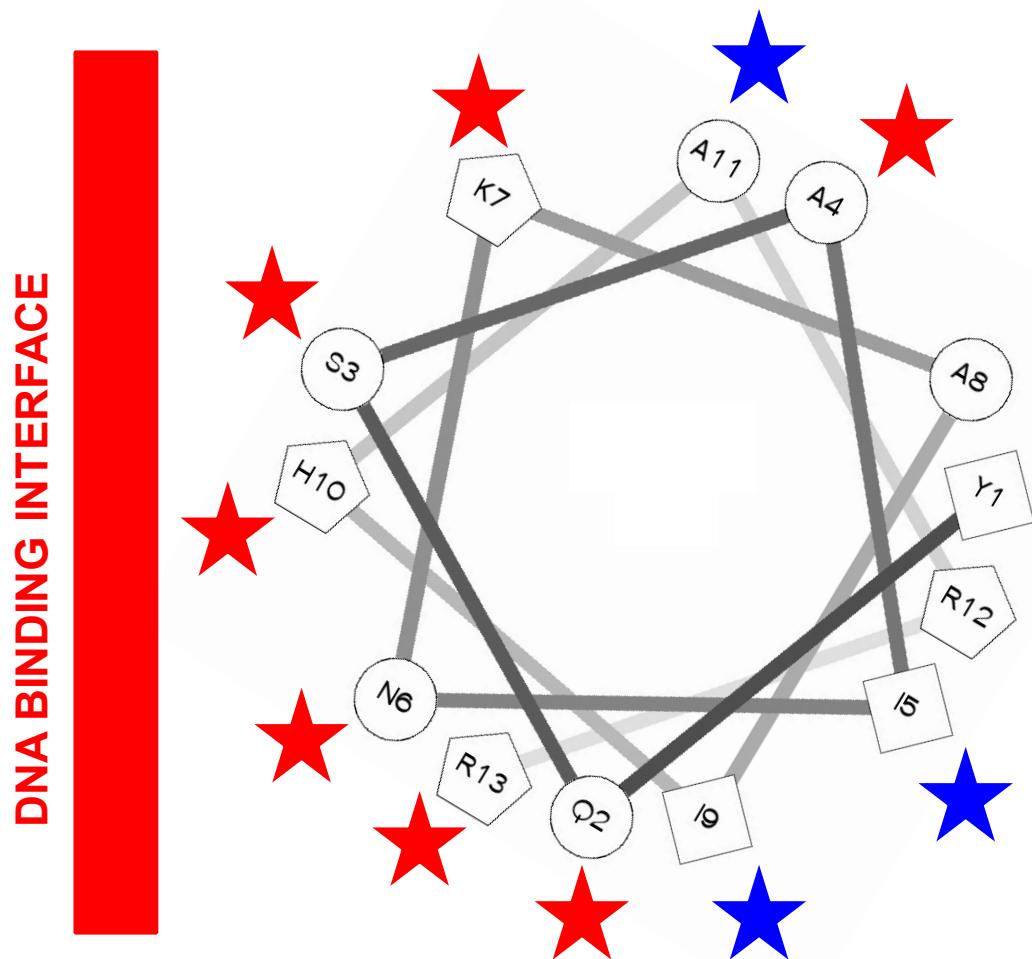
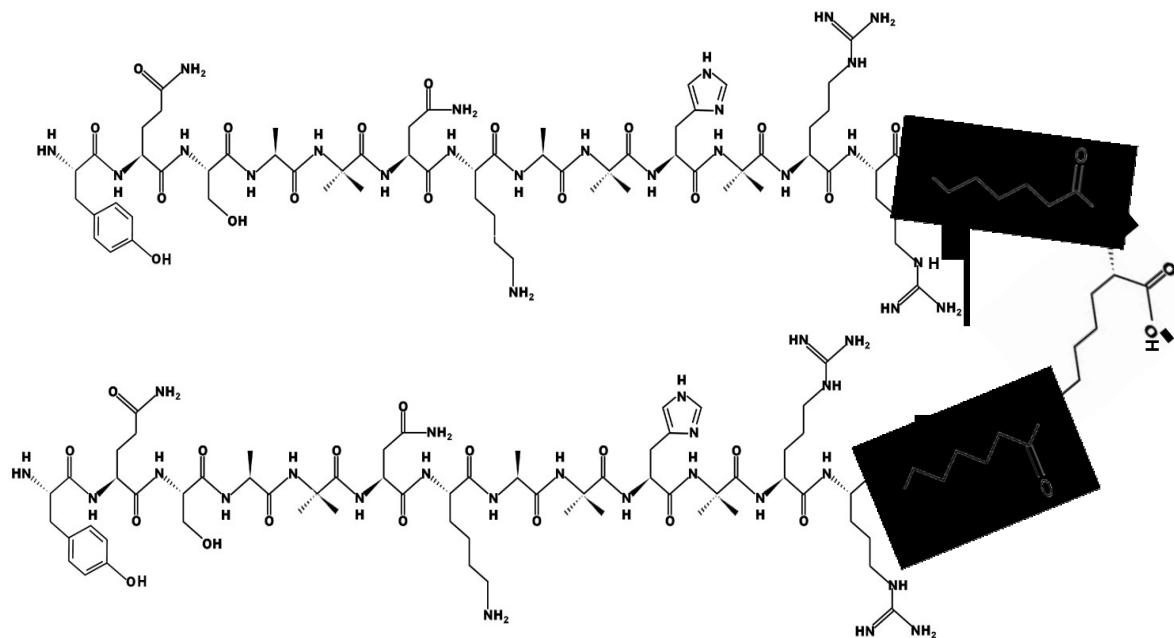


Figure S3. Design and sequence of the peptides used as DBD. The red color amino acids and terminal ‘K’ was used as linker. Using amino hexanoic acid (abbreviated as A_h in the diagram) as linker the distance is ~24 Å and the peptide is designated as Br-Ahx-Cro. Using β-alanine (abbreviated as A_b in the diagram) as linker the distance is ~19 Å and the peptide is designated as Br-Ala_β-Cro. Using glycine (G) as linker the distance is ~16 Å and the peptide is designated as Br-G-Cro. The scramble Br-Ala_β-Cro peptide contains scramble sequence of the Br-Ala_β-Cro and the peptide is designated as S-Br-Ala_β-Cro. α-amino isobutyric acid is represented as B.

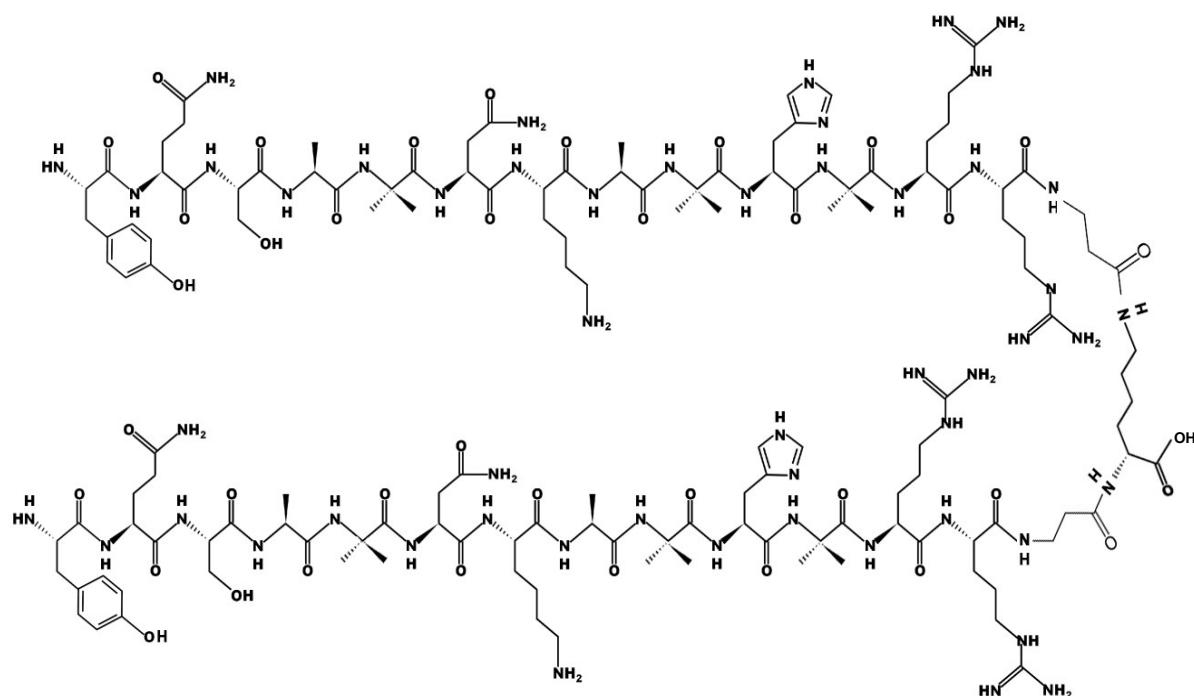
<u>Peptide</u>	<u>Sequence</u>	<u>Linker length</u>
Br-Ahx-Cro	YQSABNKABHBRRA_h > K	~ 24 Å
	YQSABNKABHBRRA_h	
Br-Ala_β-Cro	YQSABNKABHBRRA_b > K	~ 19 Å
	YQSABNKABHBRRA_b	
Br-G-Cro	YQSABNKABHBRRG > K	~ 16 Å
	YQSABNKABHBRRG	
S- Br- Ala_β-Cro	RBSRNKHQAABBYA_b > K	~ 19 Å
	RBSRNKHQAABBYA_b	

Figure S4. Chemical structures of (A) Br-Ahx-Cro (B) Br-Ala_β-Cro (C) Br-G-Cro (D) S-Br-Ala_β-Cro (E) Kix-Br-Ala_β-Cro (F) S-Kix-Br-Ala_β-Cro (G) No-NLS-Kix-Br-Ala_β-Cro (H) VP1-Br-Ala_β-Cro and (I) VP2-Br-Ala_β-Cro.

A



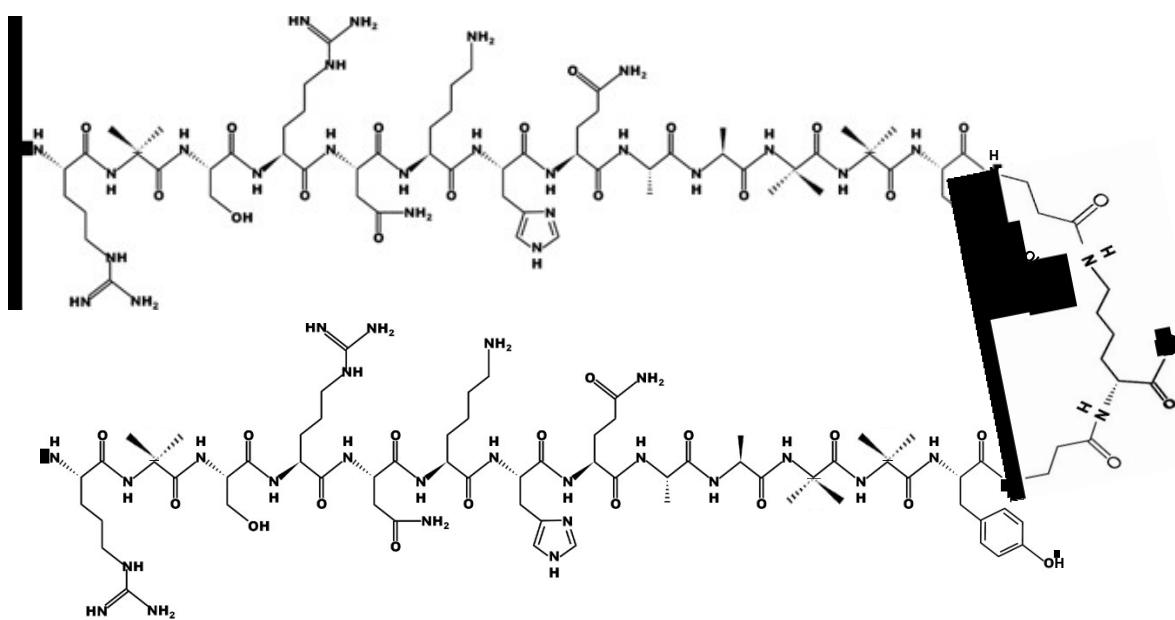
B



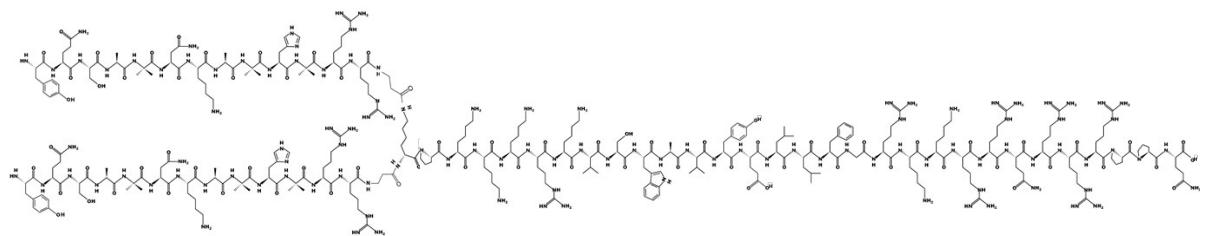
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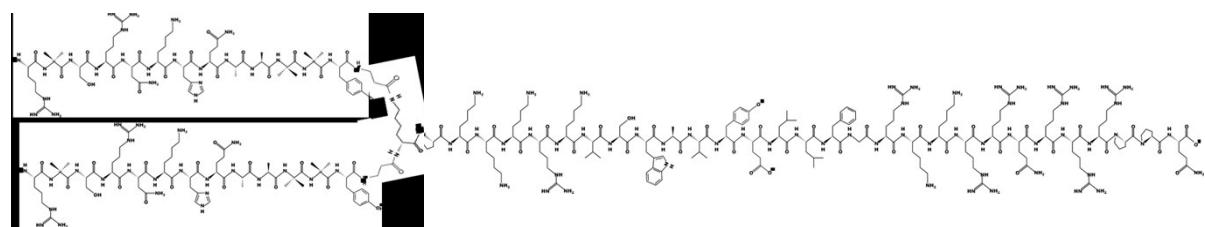
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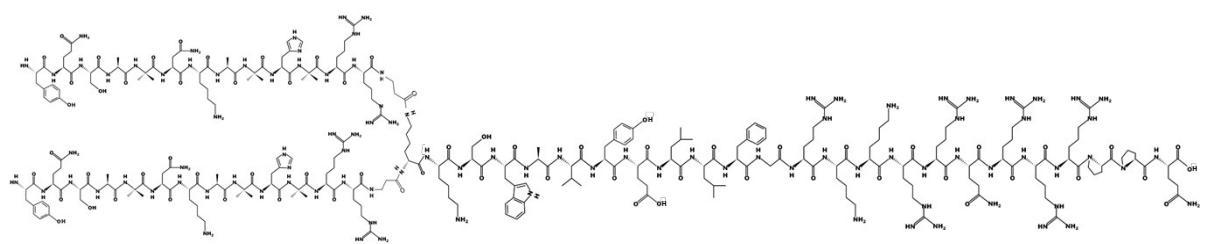
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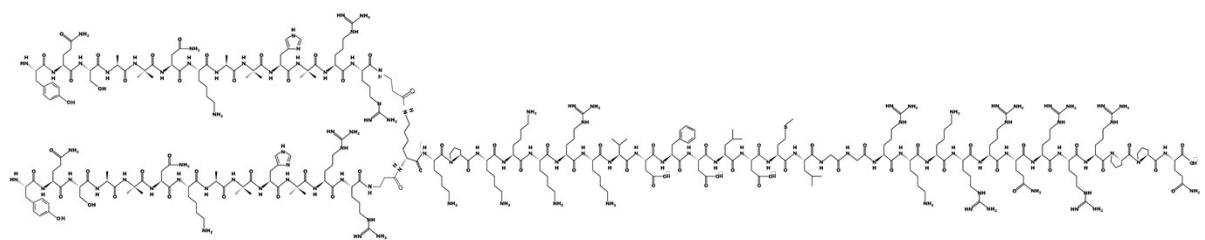
F



G



H



I

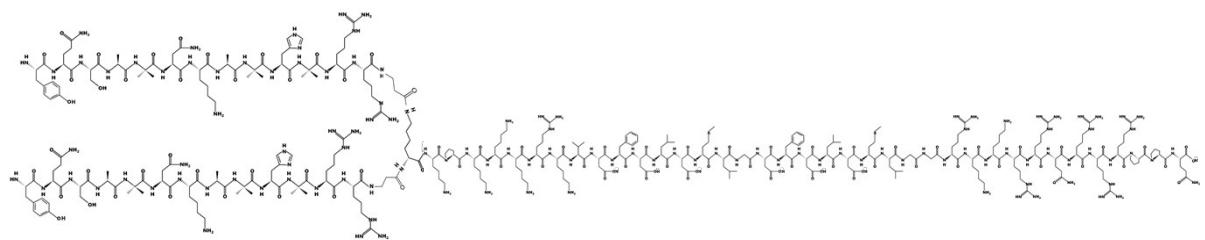
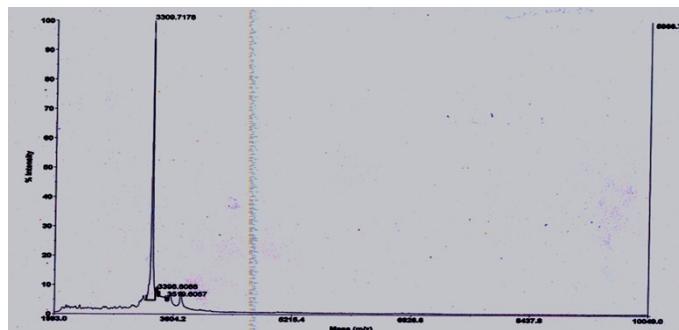
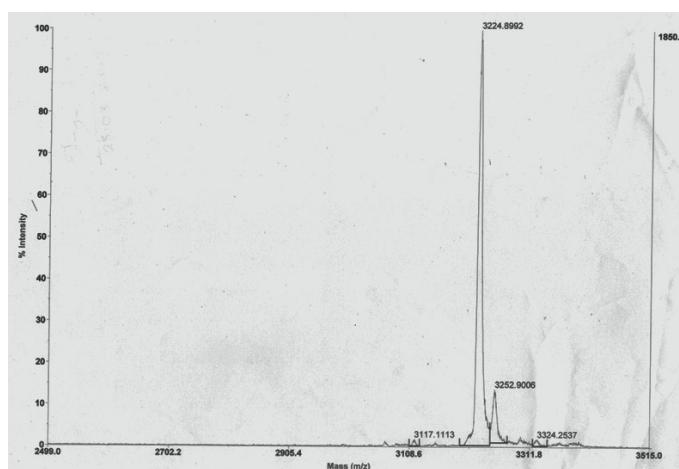


Figure S5. Mass spectrra of the A) Br-Ahx-Cro. (B) Br-Ala β -Cro. (C) Br-G-Cro (D) S-Br-Ala β -Cro (E) No-NLS-Kix-Br-Ala β -Cro (F) VP1-Br-Ala β -Cro (G) VP2-Br-Ala β -Cro (H) Kix-Br-Ala β -Cro.

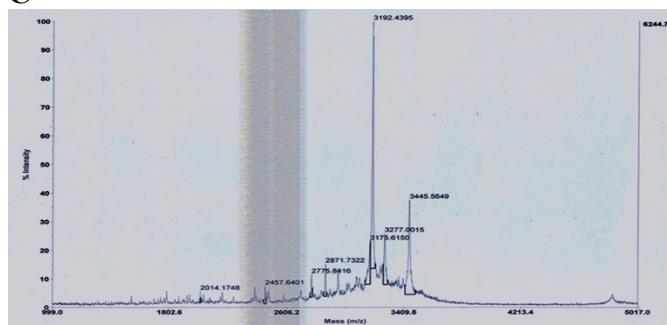
A



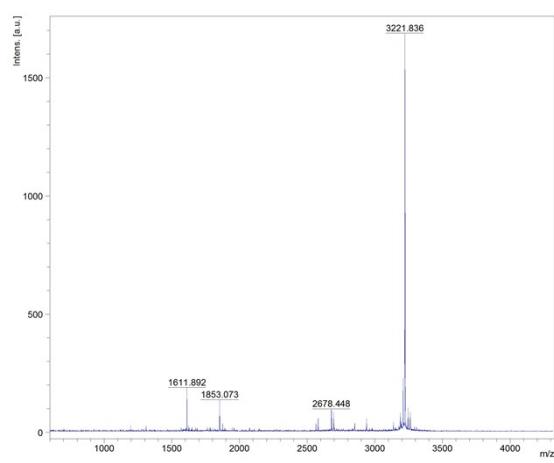
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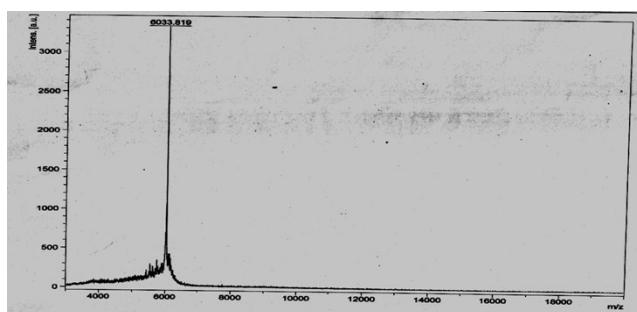
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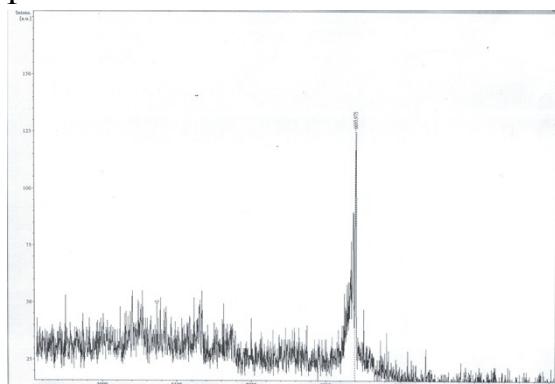
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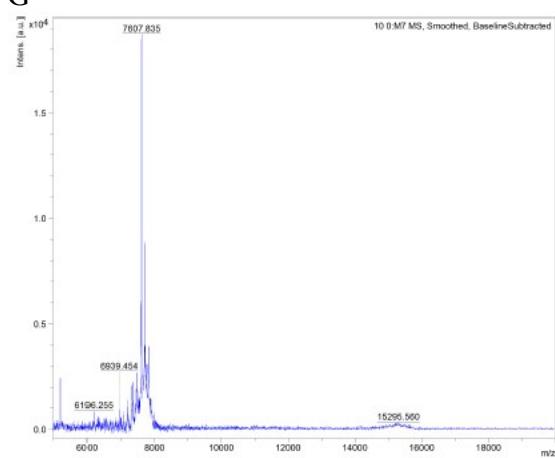
E



F



G



H

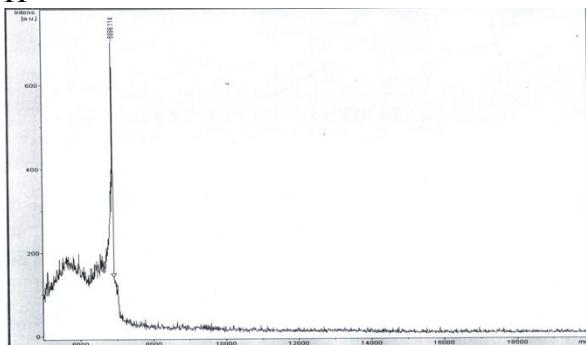
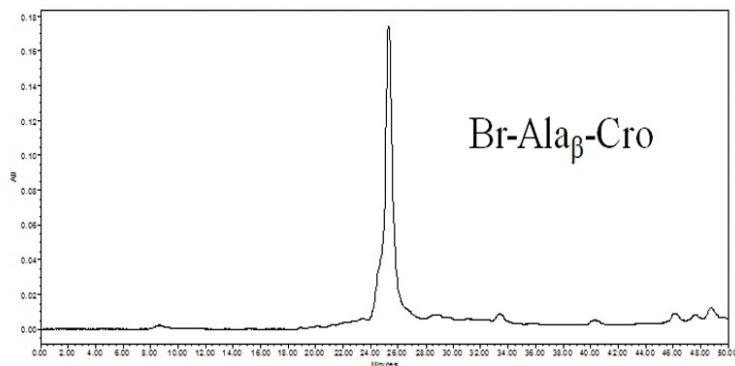
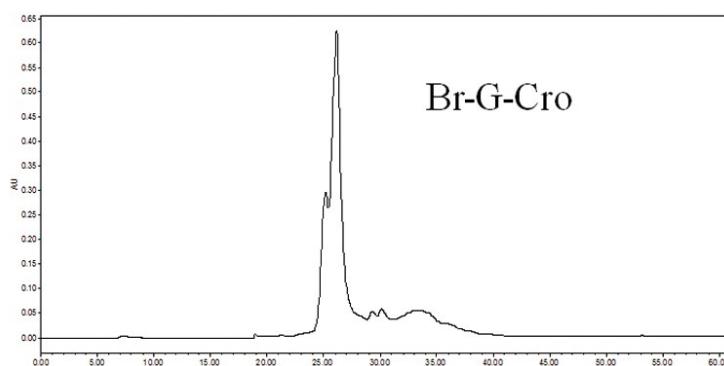


Figure S6. HPLC profiles of some representative peptides to show quality of peptides used.
(A) Crude Br-Ala β -Cro (B) Crude Br-G-Cro (C) Crude Br-Ahx-Cro (D) Crude S-Br-Ala β -Cro
(E) Purified S-Br-Ala β -Cro (F) Purified VP1-Br-Ala β -Cro (G) Purified Kix- Br-Ala β -Cro. (H)
Purified S-Kix-Br-Ala β -Cro.

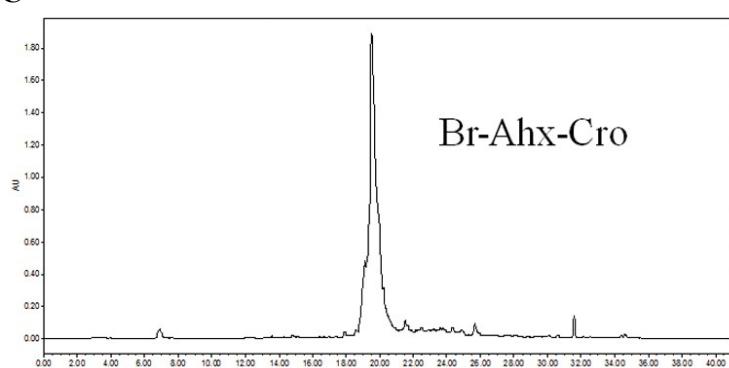
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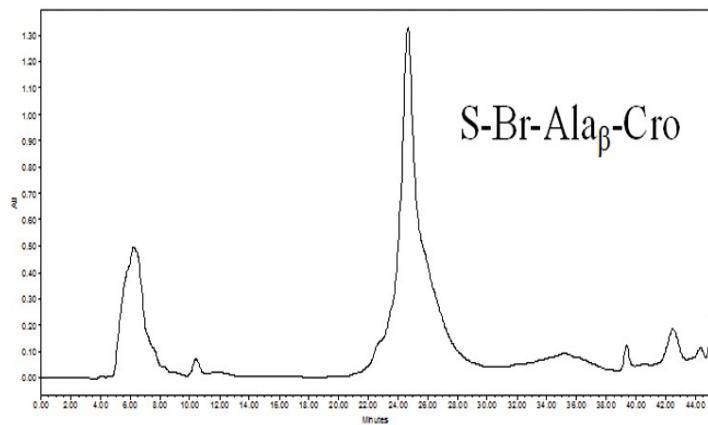
B



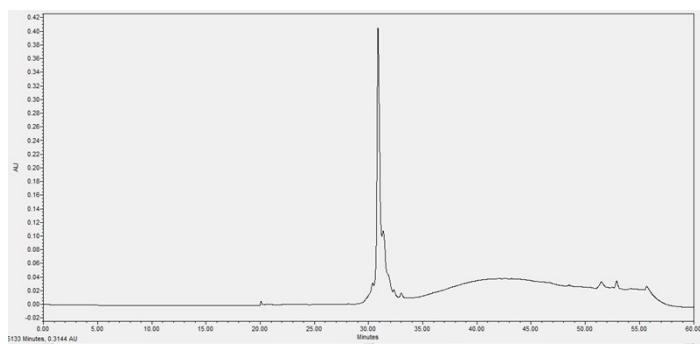
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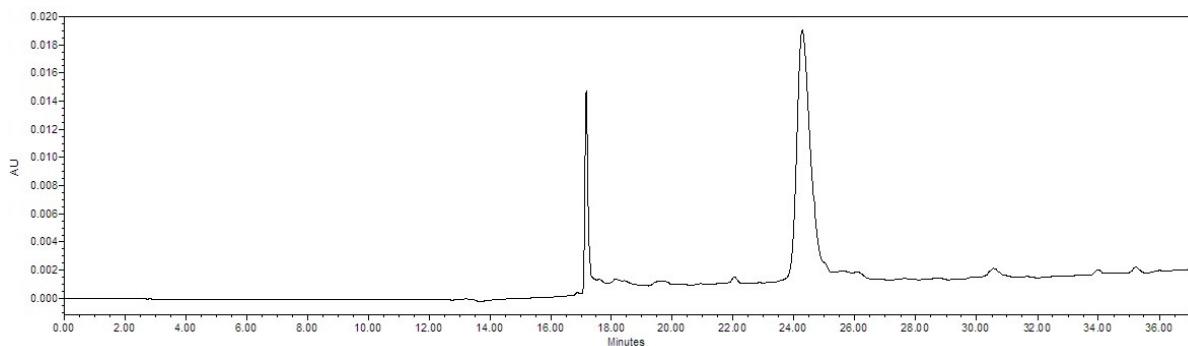
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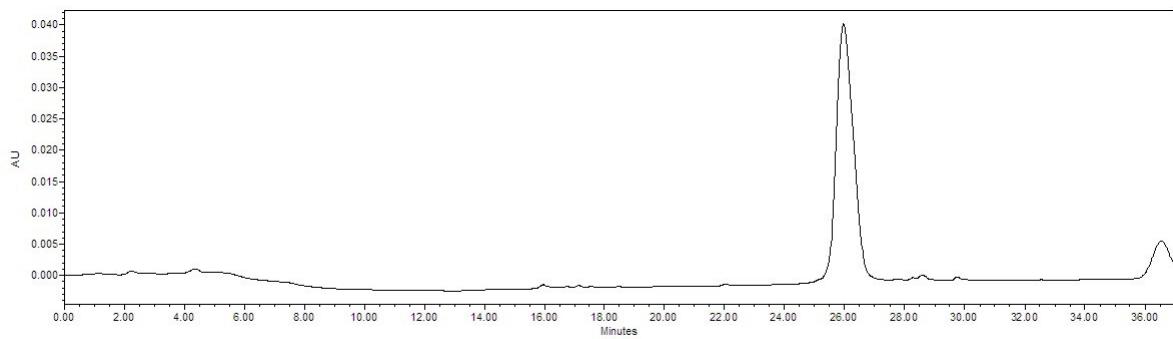
(E)



F



G



H

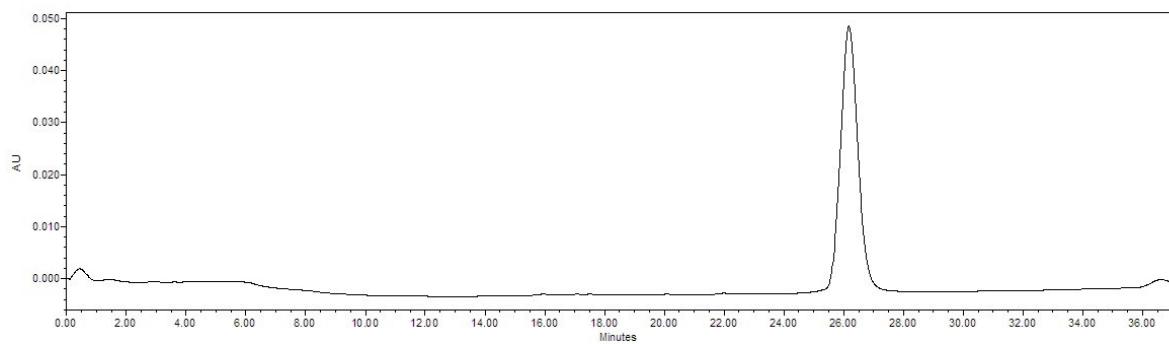
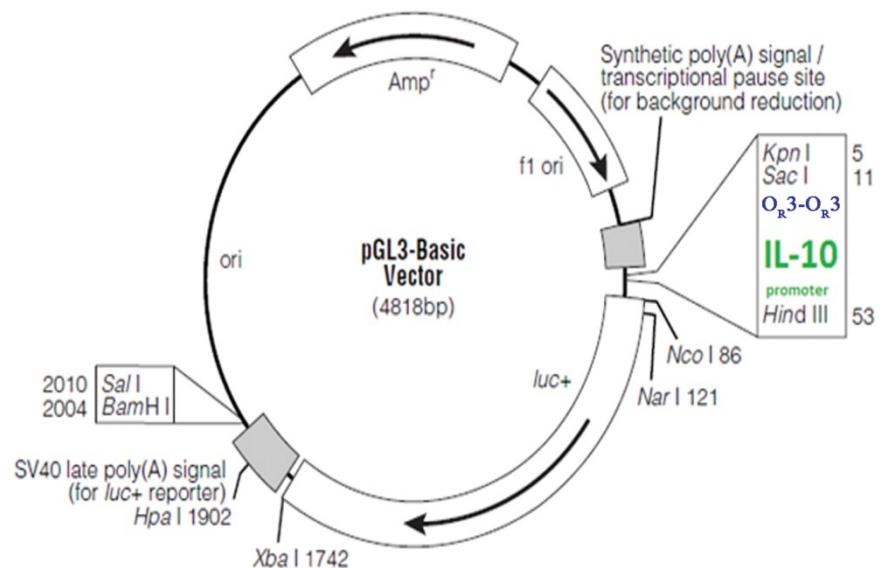


Figure S7. (A) Vector map of the pGL3 containing (O_R3-O_R3)_n and mouse IL-10 promoter. (B) Schematic representation of the reporter plasmid. n represents number of O_R3 in the upstream of IL-10 promoter, n=1 represent 2X O_R3 ; n=2 represent 4X O_R3 and n=3 represent 6X O_R3 .

A

Vector map of pGL3-IL-10



B



Figure S8. The cell viability was measured by MTT assay. The cells were treated with increasing concentrations of (5, 10, 20, 30, 40 and 50 μ M) Kix-Br-Ala $_{\beta}$ -Cro. The cell viability was measured at 36 h. The data are plotted as % viable cell against concentration of the peptide. Untreated cells were considered as 100% viable.

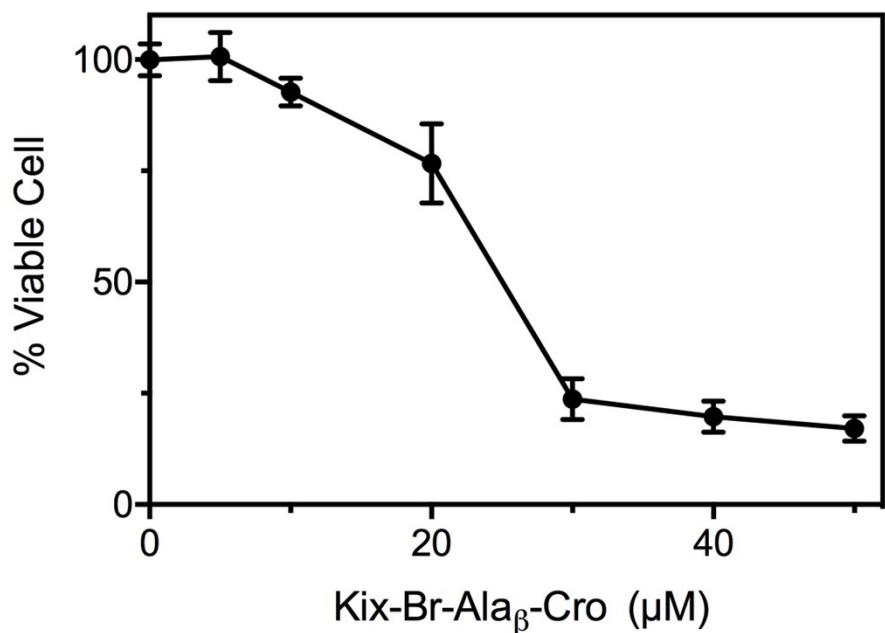


Table S1: Name and sequence of the oligonucleotide used in the study

Oligonucleotide	sequence
O_R3	5'-TATCACCGCAAGGGATA-3' 3'-ATAGTGGCGTTCCCT AT-5'
C12-O_R3	5'-TATCACCGCA <u>AGGGG</u> TA-3' 3'-ATAGTGGCGTTCCC <u>C</u> AT-5'

Table S2: Sequences of activation domain and artificial transcription factor (ATF) used in this study

KBP	SWAVYELLF
VP1	DFDLDMLG
VP2	DFDLDMLGDFDLDMLG
Kix-Br-Ala _β -Cro	(YQSABNKABHBRRA _b) ₂ K PKKKRKV SWAVYELLF GRKKRRQRRPPQ
S-Kix-Br-Ala _β -Cro	(RBSRNKHQAABBYA _b) ₂ K KKKRKV SWAVYELLF GRKKRRQRRPPQ
VP1-Br-Ala _β -Cro	(YQSABNKABHBRRA _b) ₂ K PKKKRKV DFDLDMLG GRKKRRQRRPPQ
VP2-Br-Ala _β -Cro	(YQSABNKABHBRRA _b) ₂ K PKKKRKV DFDLDMLGDFDLDMLG GRKKRRQRRPPQ

NLS and CPP sequence are marked in red.

Table S3: Gene upregulated more than 3 fold upon treatment with Kix-Br-Ala_β-Cro. Gene name with fold upregulation. The location of the promoter in the genome with availability of O_{R3} like sequence within the promoter region.

Gene Name	Fold Change	Location of promoter in the genome	O _{R3} site in promoter
Endou	16	Chromosome 15: 97,717,800-97,718,201	absent
Golph3l	15	Chromosome 3:95587401-95588001	absent
Nlrp4a	12	Promoter not characterized yet	-
Pi4k2b	11	Chromosome 5:52741200-52743600	absent
Cwc22	10	Chromosome 2:77945200-77947000	absent
Dym	6	Chromosome 18:75018201-75020001	absent
Pdkp1	7	Chromosome 17:24140200-24142600	absent
LOC100040563	7	Chromosome 4:102986001-102986401	absent
Slc5a12	6	Chromosome 2:110597200-110603400	absent
LOC665635	5	Chromosome 2:30677000-30677801	absent
Tmx3	4	Chromosome 18:90508400-90511601	absent
Man2a2	4	Chromosome 7:80369000-80369601	absent
1700013B16Rik	4	Promoter not characterized yet	-
Col9a2	4	Promoter not characterized yet	-
LOC100047593	4	Promoter not characterized yet	-
Rps18	4	Chromosome 17:33954400-33957801	absent
LOC668861	4	Promoter not characterized yet	-
XM_001480358	4	Promoter not characterized yet	-
Zfp397	4	Chromosome 18:23953801-23956201	absent
ENSMUST00000067307	4	Promoter not characterized yet	-
Ints9	4	Chromosome 14:64948001-64951201	absent
0610009D07Rik	3	Chromosome 12:4816600-4818800	absent

Table S4: The location of the promoter in the mouse genome with chromosome number and position in the genome. Promoter sequences

Endou promoter:

Location: chromosome 15:97717800-97718201

Sequence:

TAGCCCAGTGGTTCTAACCTTACCTTGGCAAACATCATCTCTAAAAACATGTTAC
ATTATGATTCTACAAACAGTAGTAAAATTACTGTTATGAAGTAGCAAGGAAAATAATTTAT
GGTTGGGGTTGACACAACATAAGGAACGTATTAAGGGCAGCAGCGTCCGAAGGTTGA
GAAACACCGGCATGCCTGTCTTGTGTTACTGCAAAGCAGGCACCAGGGGCATCC
CAGGATTGCGTGTATCTAAAGAATCCCTCCGCGTTCACCTCTAACACAGTTAGTAA
CTGTTGTTGCTAAGGTTGCAGACAGGTTGCATGCAGCCCCAGGGCCAGCCACG
GAGACCCATACTGATCCCGTCTGAGACAGAATCCAGCTTGC

Golphl3 promoter

Location: chromosome 3:95587401-95588001

Sequence:

TAGTCAAGAGTCAGTGCTGAAGAAAAGGAGAGTTAGTGAACCAAAGGAACATACCTAAAA
ATTATTTCTTATGCAAATTGCACTGAGTTCTAACATTAAACTAATATGCTGTATTATT
GAAGGTGTGCCTCTGTACCTCTGAAAATTATTTGTAGTTCATAGTTATTAAACTAATA
AAAGTTTAGCTATTATATTGTGTTACCTGAGATAAAATCCAAGGGAACTTTGACTAAA
CAAAAATACAATGTCATTGATTCTGCTCATATCCACTGAATACAACCTGCCTTCTATT
TAAAGAAAGCTTTTTTTAAATGTGTTAAGTTAAGGCCCTCTAAACCACTTTGAA
GTGTTTATTCTGACTTTAAATCTGTAAGGATAACACTTTACTTATGTTAAACTGTA
CCATTTACTGGTGCTATTAAACAATGTTGCTATTGATATAGTGTGTTATTACAA
CTTTTTTTTTAAATATTATTATTATGTAAGTACACTGTAGCTGTCTTCAGA
CACACCAGAAGAGGGCATCAGATTTCATTACAGATGGTATGCCACCATGTGGTGCT
G

Pi4k2b promoter

Location: chromosome 5:52741200-52743600

Sequence:

TTAGTAGCAAATAATGCATATAATGCACAGATTATGATTACAAACTATATTTAAC
TATGGACTATAGTAATAGAAGTAGCTCCAGCCTCATGAACCTTAAAGGAAAGTTAGT
GAAATAATGTATATTATGTATATAGTAATAAGAGAGGGACTATTAGGTTGGAAGGGATG
CACCTCCCGCGAAATGACGCTGGCTGGCTTCTGCTGGATCCACTGCAATGGCTGGC
AACGCTCTGGAGTTGAGACACAGGTGGCCATCAATGCCCTACCCGCCCGCCCCGCC
CACCTCGCTGCCCAACCCCTGCTCTGCGCAGGCGTAAGGAAGGCGTCCGGGG
TCGAGAGGCGGGCGTGGCCGGCTGGCGTGGCTGGCCGGCGGGCTGGTGGACTGGTGA
GGTGGCGTCGGTGGAGCAGTCCCCCGGGCTGGCGTGTGCTGGGTCCAGTCCGGA
GCCCGCGGCTCTGAGCGCCGGAGCCGCCCGGGACATCATGGCGGAGGCCTGCGAGCCC
ACCCGCCCTCGGAGGACGAGGACGAGGAGCAGGGAGCCGCTGCTGCCCTCGCTGG
GCCCGCCGGAGGGTCGCGCCGGAGCGCCGTGAGAAATGCAAGGCGACGAGGGCGCG
GATGTCCTCCCGAGCCCGCTACCGACGAGCCGCCGGTGTCCGGGGAAAGGGTCGATC
TCCCGAGCTTGTCCACCGAGCTGGATCGGACCCGACTACGAGCTCAGGTGCGACCCAT
GCGCGCGCAACCCGCCCTGCCCTGCTCAGTTCCCTCACCTTCCCTGTGGCAG
GAGAACGGGACCACAAATCTACTCTGAGGCTTCCCAAGAGTCAGTCCAAGTGG
GAACCCAACACTGAGGGAAACTTCATCCAGGACTGTGACGTAAGTTGTTCAAGTT
GGACTTAAGTTCTGAATTAAGAACGACAACGATCCAGAAAGTTCTCGTGGAGTCATGA
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ACAAGTTCAAACCTTCTATTAAATGTTGCTATTGAGCTTAGCCAGAGGGGGCGGAG
GCGGGCGAACTCCTGCCGCCAGCTCTGCAAAGTATATTGCGCGCTAAAAGAAAAA
CACCTTTCTTATCTCCCTGGGCTGATTTAAAGTGGACCTTGAAGAGAGGGCTAACTAA
CAATGTTAAACTAAAAATTAAAGGCATGCTATTGCAAGGCCCTGGGTGTGTGGTAGT
TTGGACCGGTGGAGTGGAGTGTGGCTGAAATGGAGTAGATTGATGAGATCCATCTGGA
GTCTAGACTGGCTTTTTTTTTTTCCCTTTCTCCTACTTGATGATGATA
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TTGCCATGCAACACTGAATAGAGCAAAGGAATGTGAAGTCCACTTCACAAATCTCT
GTTCAGCCAGCTGCAGTGGACACTTGGCCCTATCTGAGGTAGAATCTAGTCTGTGAGGT
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GAGCCTGAAATAGAGTCTGAAAAACTTTACTGTTGAGTGGGTTGGATTGGTTGTG
TGTTTCTTGTGCAACACTTATATTAAAGGCTAAAGTCAAAAGGACGAGAATGTGGTA
GCTAAAATGCAAATGGATTATAAGAACTCTTAGCTCCTCGGTGGTGTAGCTAGGTAGA
GTTCACAGTTAAGATGAGTTGACATTGGTCTCACAGCAGCACAATGGGTGAGTTGT
CGCTGACCCCTTTGTGAAACTGAAACTTAGTAAAACGGTAATAATGGGGCTCCATT
CCAAGATGTTGCACAGTCACAGATCCAGGAAGTGGTCAAGTTATGACTGAGCCACTTCT
TTCTCTTCAGGTCTGAACTTGTGTTACTCTAGGAGTCTGTGTTACAAAACAA
CTACTGCTTGGGGTATTATGTAGGTTCTTTAAAAGGATTTTTCTGACTCTTA
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Cwc22 promoter

Location: chromosome 2:77945200-77947000

Sequence:

AAATAAACAAAAGGCAGTTAAATACAGAGCACAATTAACAGAAAAGGTAGCTCATTTTCA
ACATGTATGGATCTTTTGCCCTCATATACATATAGGGACAGTTCAAGTTGCTGGTGCA
AGTGGCTATCAGCAAGCAGGTATCCAGTCCCCTGAATCTGATATTATAATTGTTGTGAT
CCATCTTGTGGGTGCTTGGATCGAACCTAGATCCCTCTGAAAGAGTGCTTTAACCACTG
GGCAAACCTACCAACGGCTTGGGGGCTCAGTTACTGTTCTATTAAATACCTTCCCCA
AATTATTCTCCGGGGGGCGGGGGGGCGAGTTAAGTGGCTACAATGTATCAAAAAGTCT
TGCATTAGAGCAGGTCAAACAGCTGGCACAGTGTGTTGACTGTAGGGCCTGCAGCT
CGCTAGGGGCAGGAACGTCAAACACCAGTTGGAAAGTTGCAAATAGTAGAGGCTTTAGA
GTTAAGGTTAAGTACCGTATCGAACCTAAACACAGAAAGATCGAAACATGGATCACTAGGG
CTAAACATGACAGGGGTTCAAGTCAAAGTCAAACAGATCCATAAATGAAATGAAATA
GAATTGGCTACTGGGATAAGCTGGAAATGAGGCAAACCGCGTTACCGAAATTATTAAATGCC
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CAACCCAGACCAAGGCCAGACACGCACGCCGCCGCCGCGCTAGGCCAGCCGGCTC
ACACGGCACCTCGCGGGCGCCGGTGGAGACGGCGCAGCCACATGAGGAAAGCAAGGCC
ACCGGGCAAGCGCCACTCGCAAGTCCCTGCCCTCCACCGCACCTAGCAACCCGCC
CCCCACTCTGGCTCTAGTCTATCCGAGTCTCAGAGTCCGTAATTACCTAAAGCTCC
ACCGCGCCGCATCACCGGAAGAGCGTCAGTCCCTTCCGGTGAAAGTAGCGCGA
GAAATTAGTAAAAACTGGTAGTGAATTGGCAGGTATGTGATACGATTAATGATAATGCG
GTATAGTAAACACCTTGAGTCAAAGCAATCCAGCTAGTAACACTGTCAATTCTACAAAGA
ACACGAGAAAAGAACATGTGAGTCGAGTGAACATAAAAGTCCAGCATGCTCTGGCAGCG
CGGACGGAAATGAGTTGAGTGGACCGCAGTGCATGACGGGAATTGTAGTCGTTGG
TAGGAGGTACACTAGAGACACTTTGAGAGCTAGGTTGAGGCATGCTTATTCT
AACACTACTCATCTCTAAAGCCATAAAAGTAGCTTCTAGTAGACACCGTAGCTTTCT
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GCAGAGGCTTACTATATAACCTTGCTGAAAATAATAAAAAAGTAGCGATCCCTTCT
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ATAAAATGATATGTTGTAAGGACAGTGAGAAGTAATGGAGGCCAACATGGGGTGTCT
GCCTTACTCCAAATATTTGGAGGCTGAGACTGGCACTGTGTTCCAAGACATCCAGA
GCTACATAATGAGCCCTGTCTGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG
A

Dym promoter

Location: chromosome 18:75018201-75020001

Sequence:

TACGAAGCAGAACGGTCTTGCCATTGAGTACTTAGATTATGGGCTAAATTCCATCTT
TTGTTGTTGTTGTTGTTGAGACAGGGCTTGTATGCGACTGTGGGGCTTCT
GATTCAAATATGCAGCCCTACGTGACTTGAACACTGTACAATCCCTTCTAGCTCTCA
GGTAATACCATCTTGACGCTACACAGTATATGGCTTCTATTGGTGCACAAATCC
TGATTATATTGCTGAGATCTTCTGTTAAAATATTATTCTTAAATTGACAGGCCC
ACAAAAGACTGTATGACATTGGTGTGATTGTCAGCAGGAATTAACAGGCACCTAGC
TCAATTAAAGTTACCCATGCGCAAGAGAGCTCTCAGTCCCAGGAGCCTAGAGGTTGG
TT

TACACTTGCCACGTGGCTATGGTGCAGGGAGACGCTAGGAGGATGGAACGCCGGA
AGGACTACATTCCCAGAAGCTCGCTGGCGTCCGACCGGGACCGCCCGCTGGC
GCGCGCTTAGCCGTCGGTCCCCAGACCGCTCCGCTCTTCCCCCTCTCACCCCAA
CTCCTCCTGCGTCTCCGACGGACAGCCGCTGGGCTGGAACCGCGTGGGAAACCGGAGCC
CCTGACCTACGGAACCATGGAGGCCTAAGCCGCCGTGCACACCCGGGGCCCTGTGCGCCC
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TGGCCGGCCCGCCGGAACCAAGGGTTGCCGGAACGGTTGGAGCTTCCGACACTCCGGA
CAGCCGAGGTTGGCTGGACAAATGGCGTGTGATGGCAGGGCGAGCCTAGTGTGTTGC
GTGTCTGACACACAGTTTATTCGATATCATCCTATACTGTAAAAGCACAAGTACC
CTCTGAAGTACAGCTCACTAGCTTTAGTCTCGTACAGCTCGTACCTCTACGGCAAT
CCCTCAACATTATCTCGAAAGAGGCTGTACCCATTGCAACCCCTCTACTCC
CTCCTCAGCCCTGCAACCATTGGTCTATTGGCTGGGATTTGCTTGAGTCAT
TGGGTGTTAGCTGAAGGATGGAAAGGGCACATTGGGTGTGAGGGAAAGTAGAGCGATAAA
GCAGGTGGAAGAAAGTCCTGGACAACAGCGAAGGGATGTAACGGGCTAGCAAAGCCCTA
CCCTGAGACTGGGGATAGGAGGAAGTGCACATCCTGAAGGCTGAGGAGCAGTGTAGCCTCCCAGAACCA
GGTGCCTGCTGAGTTGAGGGCTGGGCTGGAGGTAGTTGTAAGTGAATGGTTGC
TTAATAGGATGGACTTCATGGTTCTAGAACGACTATCCTGCTCTGGAGTTTAGATA
ATCTTTTGCTCAGAGAAACTGGACAAGGGTAACTATCCCCATCTAGAAAACGAAGAAC
CGGCCTTTAGAGGTTGGGTACTGGCTAAGCTCCCAAAGTTAGTAAGTAGCTGTGAGT
T

Pdpk1 promoter

Location: chromosome 17:24140200-24142600

Sequence:

GGTAAAGACACAAGTGTGCAACACCCCTTTGCCCTTACCAATTTCAGCTGGAT
AGGTCCCTCCTCTTAAATTGATTAATTACCATCCATCTACAGTAGAGGGTTATA
ACTTCAAGTCCATGGATGAGATGCAAGGGCTTGTGAGCCTCTGGTAGCAACAAATTCC
GTGCAACTGTGTATGTTCTCAAAGGAGCCATACTTGTATTCTAAAGGGGCTCATGA
CCCTCCAAAAACCACTACCACAGACACATCCTAGACCATACTGAACCAAGTGAGTTCTT
CAGTCCCCTCGAATAAAAGAGACCAACAACCCCTTCCCATCCGCCCACTCCCTGAAAG
CTGGAGTGAGGATGAAACCAAGAGCCCTGAACATGTGAAGCAAGCAGATTACCACTGAA
CCACAGCCCTAGCCCAAGGAGGTTGCCCTTGAACCAAATCTGCTGGCGTGTGACAT
CTTCTCATCAGCCACTTCATATCTCATACAAGGCAGACACGCTGATCTAAAGAACTCTAA
ACTTACCAAGAAAATACAACATGGAACTTATTAAACAACACAGAGAAAACAAACATTTC
ACCCAACAAGTGGTGGTTGTGTCCTCGTCCTCCGCCCTCAAGCCGGCTCCAGGGCTTCC
TCCCTGTCAAACCTAGGCTGCCCTCGCTGGGCCCTAAGCCGGCTCCAGGGCTTCC
CACCTCACTGAGCGTAGAACTGCAGGAAGAGGAGGACGAAATGCCCGCTGACCAAGA
ATCTGGAAAGGAGCCTCCAGCATCAGGTGCTACAAGCAGGAACCTACATCTCTGCTTCC
CAGGAACCTCCAGGTAAAGGATGGAGAATGCTCTCTCGGGCAGGACCTAAGGTA
AGCCCAGAGGCTCGGCTCCAGGCTGCACAGAGCCAAAACAAACTTCCAGGCACCTCCA
CCCCGCCCGACCGGGTACACCTCCGCCCTCCCCGCCCTCGTGCCTGTAGCGCC
CCGGCTGCCGCCAGCCCCGAGCCGGGCCATTCCCTGCCAGCCGAGAGGAAC
CTCGGCCGGTGCCAACAGTCTTGGCCGCCACCCAGCCGAGGTGAGGGGCCGCCGG
GCGACACGGGCCGCCCTCGCCAGCAGAACCCAGGGGACGCCACGCTCCGG
AACGGACGGGTGCGCCCTACAGCAGCCGCGCTCAGCTGGCTGGTGGTCTGGCCA
TGAGCCCCAAGTCGGCGGGCGCTCCCCCGAGCGTGTGCGCTGCTGGCGCTCTCA
GCGTCCTCTCCCCCGGGAGCCCCAGCAATGGCGCCCTCATCCTGCGCCCGCC
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TCACGAGTAATCTGGAGAGAAATTGGCTATGATTGAAGCAGAAAACAAGCCGTC
TCACTCCTGATGAACCTGTTTGCCTAGCAACTATGAGGACGCTTTGTGAGCTGCTT
TTTTTTTTTCCCTCCAGCAGGATCTCACTATGAGAACAGACTGGCTCCTACTT
CGGCAATCCTGTCAGCCTCCAGCTGAAATGACAGGTATGCCACATCCTCCTT

AGTTGTCATTATGACAACCGCTGTAGATGTTTTAAATTATGGTAAACGCATAAAT
TTTGTAACCTTTGCGCTGTGTACAGTGTATGTGTGTCACCTTGCTGGCTGGCTGGC
CAGATGTCTCCCTATACCACTCGCTACCTGGAACTTGCTGATCCCTCTGGCTGGCTGGC
TGGCCAGTGGAGATGCAGTGGCCCCCTGCTCTGTTTCCCAGGGGTGAAGGTACAGACA
CACCCCCACCATCTCAGCTTCTTACAGGAGTGCTGAGGACTTGCTGCATAGCAGG
CATCATACTGAGCCATCTACAGACCTCCCCATTTTTCAGTGTGGAAATTAAAC
TCAAGACCTCACACATGCTAGATCATCTCAAATTAAATGTTCTAAATTACACCCATAG
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LOC100040563 (predicted Tctex1d1) promoter

Location: chromosome 4:102986001-102986401

Sequence:

GAGTGAGCAGTTATTATTGCATTATGTGAATTCTTAAGTAATTAAAGTCTTTTGCTCTA
AAATAAATTCTGCTTACAAGTCATTATAGTAAGTCCGGAGTGACTTGTAAAGATGAAT
TTACAACCTAGGAAAAGTGTATTCAAGTGTGGTACAAAGCAAGCTTATTTAGCT
TTCTTATTGAAAAAGTCCATTGCTATTCCAAAAGCAAAGGAAGGGAGAAGAAGGA
AGGAAGAACAAAGGAGGAGAGTAAGGGTTCTGCCTGCACTGGAAATATGGCAGGGCTGGC
ACACCCCTCGTTAATGACGTACAGTCATGTGACATAGCCAGGCCTCAAACACATCCAGC
ATCACCTCTCTGATTGCACTGTGGCTTGAACCAGTCCT

Slc5a12 promoter

Location: chromosome 2:110597200-110603400

Sequence:

CTGAAGTAGTCAAATTATCACCTCCATGGTCCTAATAATGTTAACCTTCAATTAA
AAAAATCTAGGAGATAATAAAAGCATGCCATGGGCCAGCCTACTCCCTCTACCCCCCAG
AAACTTCAGGAGTCTCTGTCAGATGTTGAGAAGAAAACGCAGACAGCTAGTGGCAGAA
TCTGATCCCAATTGTCATTTCCTGGTGTAGTTCACAGCCAGCTTAGAGA
ACTTGAGGACTGTTGAGTATCTGAAAACCTAGACATGGATGGATGATCTCAACTCTC
AGAGAACCAAATTCTCCTCCATCTGAAAAAAAGTTGGCTGTTGGAAATATCTCAGTG
AGCCCTGCTTGAAGAGACTCTTGTGTTCACTTCCAAGCATGAGGGTAAGAACCTT
GAAGCTTGGATTATGTTGTTGAGGCTCTTCGTCATTTCCTGTTGAAATTGGCGTG
TTCTTGCCATAAAGGAGAGAAAAAGACACATCTCGGGAAATTCTTAGTAGGAGGAAGG
CAAATGAGCTTGGCCCTGTAGCTTGCTCTGACAGCCAGCTCATGTCAGCTGCACT
GTCCTGGGACTCCTGCTGAAGTCTACCGCTTGGGCATCCTCTTCTCCTCATT
TCATATGTTGTTGCTTTACATCTGAGCTTTCTGTTGTTGTTGTTACAGGTCT
GGCATCACAAAGCACCTATGAGGAAGAGACCACCTCCCTGCCCATCACAGCAAGGAGA
TGGCACTGCTTCCACACAACAGAGCTTCTCTTCTGCTTGTGTTCTATGAC
ATCTTGACATTAGTAGTAGATGACTATTTCAGGGCAATTCCCACTGTTAAAAAAAGA
CAATATGTTGTTGTTGCTTGTGTTGTTGCTTGTGTTGTTGTTGTTGTTGTTGTT
TTGTCACTCTGTTGAGGTAACTAGTAATGTCACTCCCGCTGGCTAGTTCCATGATTAA
TAAGGATCCATGATAAAAAAAATGGATGCACTATTAAATTGATGAAAGATAAGTAAGAT
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CCTGAAAGTCAAGCAGAGACTCATAAAAAAACTACTTAAAGGAAATTCTGAGATAAA
TGATCACAAAATGTTCTAGTCTTCAAGCTGTTGCTGAGATACATATTCTCCAGT
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CATCCTCTGCTACATACGCAGCTGGAGCCATGGGCTCCATGGGTACACTTGTTGG
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GCAGCATTAAAGAAACAAACCAAGTGTGTTCTGTTGAAAGTGTGGTACCGGCTT
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CTATTCAGTGTGAGTAACCATGGAAAGTAGATTAAATGTGCTCGAGTTAG
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GGAGAAAATAGTCTAGTCAAGAGAGAACAATCCTCCAGGGCGATGGAGC
GGAAAGGACTGGCAACTAAGAGTCCAGAGAAGGTCAAAGGGGAAACTAG
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CTCAGAAATGAAAGGTTATAAGTTCTTAAAGCCACCATATT
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LOC665635 (Gm14486) promoter

Location: chromosome 2:30677000-30677801

Sequence:

CGCCCCTGTGAGTGAGTGAGTGCCCCGGCGCGCTGCAGGCCGGCAGGGATTGAAG
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GCTGGCAGGCAGCAAGAAGCTGTCGGCCAGCGCTATCAGCCCACGGAGGGAGGG
CGGCAGGAGGGCGGGAGGGAGGGAAAGTGTTCAGCCCTTAATACCGGAATTGAA
ATCCCAGCGAGTCGGTGCCTTACATCTCCGGCTCCACGGCTGGATCCGGTGG
AAGAAAGGAGGCTGCGCATTTACATCTCCGGCTCCACGGCTAGGGCTGGGTGGCA
GGAGAGGCCAGCGGTGGCTCTGGACTTCCACGGCTAGGGCTGGGTGGCA
GATGGGCCTGCTGAGTGAGGTGCTGGCGGGCATCACATTATCCCATAAGAAC
CCTGATCTCCCCACCCAAAATTGCCCCGGTCCGGGGACTCCGGAGACTCTGCCAAC
CTCTTACCCCCCTCAAGCTAACGGGGCTGAGGCACCGCAGGACAAGCCCCTCTGT
GCAGATATGGACACTGAGCCCTGAGACTGAGTTCTGGAGATCTTGTAGGCCAAGGTACA
CAATGGCAGACCGAGACCAGGAGGGAGGGCTGGGATGAGGGTACACAGGAATGGAGAGG
GAGCTGTCAAGGCTGAAAGCTCTTGTCACTGACTTACCCACTGTCCCTCTGACC
TTGATGTGGAAAGTTAGGGACC

Tmx3 promoter

Location: chromosome 18:90508400-90511601

Sequence: CCAGCTGGACGGAAATCTGTGCCAGGCAGTAAACAAGCCCATAATTGGTGGATGGCTCGC
CCTTAATCCCTGATTGGCTGAGCAAGCCTGCCCTGGTGAGGTGATGTCACCTGTGGTGATT
GGTTGAGGCGTGGTTGTGGCTTGAGTGGATAAAAAAAACGCTTGTACCCCGAGGGCGCGGG
GGATTGAAAGAGAGAACGCTGCCAGTAAGCTTAAGAGGAACCGGTTGGCGTT
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CATTGCCAATTGGTCTGGAGTTGTTAACATGCCTCCATTACATGCAAAATGGAGG
AGTGCAGAGAGTCCCTAAAGTCTAAATGAAAACCAGCAATGTTCCATAT
TGTTCTTCTTTGGTTAAT

Man2a2 promoter

Location: chromosome 7:80369000-80369601

Sequence:

AAAAAAAAACAAGAACAGAAAGGCATCATCACTCAGACCACGGAAGGCAAACATTCCTC
TCACATGCCGTGCTCAGCGACCTACATGTTCTCTGTCTCAAGCACGGTGCTGAGGAG
GACTGCGTTCAAGCTTAATGGAAAAGCAATACGGTGGTCAATCAGCAGTGTCTGCCAGAAA
CTTAAGTCAGGCTATTACACAAGGCAAATTCTGTTCCCTCTCCCTTCAGCATTGCTGTTTC
CCTCCTCTCCCTCTCCCTCCCTCCCTCCCTCCCTTCAGCATTGCTGTTTC
ACTTCCTGTTACAGCCAGGAGCATAGCTCGTGGTAGAGCATGTGTTAGCATGCCAG
TCCTGGGCATCCATCTCCACAAGAAAGACAATCTGCCTAGGAATGTCCCTGGCTACA
GCTACATAACTCAGTTAGATCGTTAGTGTATATGCACGCATGCCATGAGGAGGCTC
TAGGTCAAATCTAAAGACAATGGGGTGGGGGGTGTCTTGTGCTGACAAGAC
TCTCCTGAGCCCAGGTGACTTGCATTGAAGAAGACAAGGTGTGACTCACCTGGGG
AA

1700013B16Rik (Spata31d1a) promoter

Sequence: no sequence found

Col9a2 promoter

Sequence: no sequence found

LOC100047593 promoter

Sequence: no sequence found

Rps18 promoter

Location: chromosome 17:33954400-33957801

Sequence:

CCAGGCACCTCTGCTGTCTACATTGTTGCCAAAGTGAGAATTAAAAAAAGTCAC
ATGGCAAGGACTAAAGTGAGACTTCACAAGGTTAAAAAAATTACTCCAACCTTATG
TCCACCTGACATTACCTCGGCTCCCCAGGCCTCTCAAACCTCTCAGCAGCCACCCAGT
TCCATCTATACCCTAGCATCCCACAAGGTCCCACCACTGACAGGTCTCATGGCTAACAC

ACCTGACCAGCAACTCCTCAAATCACCAACTTCAAGTCCTGCCCTGTCTACAAGACTTT
TAGCTCTGCAGAGAAACATCCCAGCTTCCCTACCGACCTCCCTTTGCCATAATGGGG
AAAAGTGCAGCTTCTGTGTCACCTTCACTGTGGCATCTGCCAATAGCTCTGCAGACC
ACCCCCAACCATCACCGCTAACGCCTGGACACCTCTAGGATGTTAGTTACAGGAATTCT
ACTCAAGGTTGCTGACAGCAGAGACTGCTCTCGGAAGCTGAAGATCTCGGGAAACCTCT
GCAAGGTGGGACATGACTACACTCACACCCCACTACCCCTGGAGCAATGTTGGTTAA
CCCTGGGTACATCATCAGGTTTATTGAAACGGGTTGGGTGAGTTAGTGTCTCTCTT
TGCCCCTAACACTTTAACCTTAATGGCAGTGATGGCAAGGCTATTTCCGCCGCCATC
GATGTTGGTGGTGGAGTACTCGAAATGTGCTGGAACTTCTCAGGGATCACTAGAGACTG
AAAGAACACGGGCAAGTCAAGACAGGTTACCCCCAACCGGGAAAGGTGGCGAGGCAG
GGTTGGATTGTCTGCTAACGTTACACAACCTCAGTCACCGGGCTCACCCCTAACG
CGCTCAGGGCGGGAAATCGAACCTCTGACAAGAGGGTGGACGTCGCCGCC
ATTCCCTGGACAGGGGAAGCGGGAGGAGCCCACACAGCCAAGCTCTGGCAGCTTGC
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GTCCCCCTCTCGGGTACACTCCATCGCTCACAGAGGGCTGCCCCCGCAGCATGGGC
ACTCTCTGAGTCTGACGAAATCTCTCCCTGCCCCGGCTCCGATGTCGAGGATCCGG
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TCCCCGGAGGTCCACTCCGGAAATTGCTCTCCCCGTCTCCCCCTAAGGGAGGAGTT
GTATTGAGACAGTTTGGAAAGGGGACGGAGCTATAACTCCGGTTCGGCCTCTGTG
TCGTACCGTTCTGTGTTGGCGGGAGCGAAGGTTCCGGAACAGAGTTGTGAGACG
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TGTAGCTTGCTCGGGAAACGGACTTCTGAGACTGACAGCTGCCGAAGTCGGAAAGTG
AAGCTCGGGGGAAATGGCAGCCGAGCGAACCATGGCGGCCCTGCCGGAGCTGGT
GGGGCTGGGGCTCAGATGTAGAGGAGGAGGAGGGCCCTGGTGAAGACACGGGGCT
TAGGTCTCGGAGTATCGTCAGGTTGCTCCCTCTCCTCAGGCGTAGGCTCCATTAGACA
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GTCCCACCTCCAGCCCCACCGCTGAAAGCTATGCTGTTCTGGTCCAAACTTCCC
ACACCTTACAGTGCAAACGCTTCTGAGACATACCTTATAGGTGGCTTAGGATTAGCC
GAATGAGCATGTTAATGGTACCGAGACACAGGGTAAACACTATGTAAGCTGGCGATT
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CAAAGGCAAGTTGACTAGCTCTAGTTTCTCATTTGAACAACCTCCAGCACTCCATG
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GATACCCGATTATGTTTGTGTTCAACATCTTCTGTATCCCACATTCAACCTGT
CCCTCTCAGGGGGGGTCTGGACTCCAAGAACCCCTACAACCTGGAGAGTTGGACAT
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AGTCACAGCAATAACACTATAAAATCGCAGCATCACCTTCCCTTGGACTATTG
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AGCGAGAGGGTCTCTAGCTCTCAGAACATGCCACACATTAGATCTTACAAGCCACACT
TTGGAAATCCCTTTAATATTTCCATCTGTGGTCAGTAGACCATGAGGAGCAGAAC
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LOC668861 promoter

Sequence: no sequence found

XM_001480358 promoter

Sequence: no sequence found

Zfp397 promoter

Location: chromosome 18:23953801-23956201

Sequence:

TTAGCTAACAAAAGAGAATTAGTTGTTCTATGTGATCGTCTTATTACCCCTTTACTAA
CTTTTGAAATAATTACAACAAATGTAAGACTATTAAATAAGAAATAACTTGGTGCCTT
AACAAATAAAATCGCTCACAAAGTAAAGTACCTAGCTTTAAAGGGGAAAGCTGGGG
CTGGAGAAAGGGCTACCAATTAAACATTGCTGTTCTCCAGAGAAACACTCACATATAT
CAGTAACATACAGTCCAAGGGACCGATCTCCATAGGCACACTGCACACTCACGCCATATAA
CGAAGGCAGTTACACATAAACAAAAATAATAAAAGTAAATATTAAATGGAGACAA
CAGGGCCTAAAAGATGGTCAGAGCATTGATGCTTTGAGAGGACCAAGGTTGGTT
CTCTCAGTACGTATAAGGTGGTCACAACCATTGTCACTGGATTCAACGCCCTCTTAG
CCTCTACCAACAAAGAGTGGTGTGGCAAAGTATTATACACCGAAAGAGTCATATT
AAAGGCGTGTGGAAATCCGATCTGAGAGCAGCTTGCTTGCTAGGGCTGCAGAGGA
CGCTGCGGCTGCCGCCAGAGCGTGAAGAGCGCGGTTGGACCGTTACTCCGTCCATTAC
CCACAGGTTACTCTCATCAGAGCTCACGAAAAACAGCCGTAAGAGGAAATCCGATT
ATGGGACTGGGAAACTGAGCGAGAAGGGAGAGACCGGACAATGGGACACAGCGGGCTGCA
AAGCCGCGGGCGGGGTGGGCAAGAATTGTGCGCGCGAGCGGAAGGGCGGGACA
GACGGGAGGTCGGACGCAGGCAGCGGAAGGGCGGGACGCGCCTCAGGCCTTGTGGC
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CTGCGGAGATCCGAAGTCGCCGCTGTCAGCTGGAGAGCCCCGTGGAGGCCCTCGGGC
ACGCGCTACGCTGTGCTTAACCCGAAGCTAAAATAGGGTGCCTGCAGCCGAGCCGCG
AGGAAGGCAGGTGTGGAGCTGGTGTCTCGTAGCGGGCGCACGGAGCTGTGGTGC
CTCGGGGTCCTCCCCATGGGACTGAGGGAGAGGGAGAGCGGGACCCGGGACTGAGAA
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CAGTACGGGGCCAGGAGGTCCGGCTGGCAGGGACGCAGGCTTAGTTAATGTGACCTAG
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GTCTAGCATGAACCTGAGTTCACTGCAAGGAGTGGCAAAGATGTAGCTACTAGCAAAG
TAAAAGTGTACCTATCATTCTAACAGACCTGAGTCGTTCTCCAGCACTGAGAAAAGTTC
CCGTTACAGAGTTAGGAATTAAATGAGGACTGAAAGATAACTGAAAAGGAGATAATCTC
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G

ENSMUST00000067307 promoter

Sequence: no sequence found

Ints9 promoter

Location: chromosome 14:64948001-64951201

Sequence:

GATAATGACTTGTGGCATAAATGCAAAATAACAACAAATTGGGCTTTCACTTAACG
ATGCATTTATAGTGAACACCACAAAAGCACAGCCCATATTAGCACAAACTAAGCAAAA
AGTACAGGAGGTATTATGTCATCGGGAAAGAGAAATGTTCTGTGCCAAGGAAGGAAGGAA
GGAAGGAAGGAAGGAAGGAAGGGGGTGGGGAGAGAGAGGAAGCTAAGGAAGATAGG
AAAGAGAAGGCAAAGATTCTATTATCTGAGGTCCCCAGAAAAAGGGGCACGTGACTAT
ACCTATATAACAGACGAGGGTTTTAAGCTTCACTGACACAAAAATAAGCAATCAC
TTTATCATTCAACTTACGAAGCGTGTACTTACTGCTAAGTGCATTGCTTCAC

TCACCCAATGCTCCTCACTCTCCATACCAGTTCAAGCATTGTGACACAGAACAGG
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ACACAGCGCCTTCAGCGCACAGTCAACCCCTTCTCGAAAAAAATTCCAACAGTT
TTCAACTACGAAAATAAGTCGCCTTATCTACCTCAGGATACAGAGCAGACGTTCTGTCT
CATTAGAAGACAGCGATGCGGATGGCATTAGCACTCGAGATCCCAGCCCCACGTTCCC
GAATTAAATAACTGCAAGCTACTGAGCATGAAGATGACAAGATACTCTACTTACTG
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CACGGTCCCGAATAAAACTAAAGAGACACTTCTTCCACCTTAGCTGCCGACT
TGAGGCAAAGATGCATCTCACATCACATCATCTATAGTCCTGCCGGAGTTGGGT
GGAGACTGCCAGAGTAGGGCACCGAGTGTGAGCTGGGCCAGACACTAACGAGTTA
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CGTCTTCCAAGGCCAGGCCCTATCTGACTCCAGCCATCTAACCTACATCACA
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ACACTCCCCAACAAACAGCACACCTCGAAACACCTTACAGTTCTCAAACCTCCAAG
CAGGAAGACCCCCAACTCCAGGATTCAAACCTCCGCCGGCTGAAGACCAGAGAGAA
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CCGCGCACGGCCGCCCTCGGAAGTACCCCTACTTGTCTCCCTATGCTCAACCTGGCTA
CTCCAGAGACCCATCTGTCGCCCTACTCATTCTCTCCCCCAAACATCTGACATT
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TGCAGTTGAATTACAAGAAAGAGATGCTAAAGGGTGGTGCCTTGAAGGATTCCCGTCT
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GCACACCGTTCTAACAGTTTATTCAAGCTTAATGGAATAAGTAATAGGTAATAA
GGAAATTGAAACGCAGGTGTGAAATGCTGTCGCCCTCATTCCCTACATTCTCACT
GCCAACTTACTGTAGAATCTCTTAATTGCTACGTGGAGAGAAATACCTCAATAGAGTG
AAGTGTGTTGGTATGTTCTG

0610009D07Rik (0610009D07Rik) promoter

Location: chromosome 12:4816600-4818800

Sequence:

TGTTTCAGGGTAGACCTCTCACTATACCAAAAAAGCCTATGTAAGCTGGACCTTC
GACAGGTTATGTCCTAAATCCTCTGTAGCATAGAAGGAGCCAGCTTGGGAGTA
TATCCCACAGTTCGCAGCGATAAGGCTTCAGCATTCTTGTGTTAAGCTATGTAAG
TCTGTCACTGCACTAGTATTTACTGCGCCCTCAGTCTACCTGGTACAGAATGAGGG
TGACATTTGAAAGCATGCTCTCTGATGGGAAGTGAAGGGTGTGCTTTAGATA
CTCAAAACATAGGGTAGGCAGGATGGAGAAACAGGGTAAGGCCTTCTCAGGCT
GATAATCTTCTCTAGTACAAACGCCGGAAAACCTCCACGGAGCAGATTCTGCC
CTCTGCAAACAGCCCCAACGCTGCTGCTGGTTCTGATAGAGTCTACCCGGCAACTTA
TATTCAAGTGGCTAATATGTTGGGCTAACAGGAGCCGGGAAAATCGTCTGGAGCTG
CCCTAGTGAAGGAGGACCTCCGAGCAAACCTGGAGTGGCCTCTTCCCATGTC
AAGCTGGCTAAGGAAATAAAAGATGCATGTGACGGGTTGTGCTCTGTTGCGTGA
CAGGGGCCGGGACCGGGTGAGGGAGTTGACGGTCAGGTGAAGCGCGACCTGAGGCC
CAGGGCCCCGGGATCGGCCTCCGAGCAAAGTCAGATTGGACCTCTGTTGGAAGCGCT

GGCGAGCCGGGTTCTCTCCGAGACCTGGCGCGCTAAATCGCAGGCTCTGGGTCTGATC
TCTCTCATGGACCCCGCCCCCTCGCTCCTTAAGCCACACCCCCCAAGCGGAACACCTCCC
TTCCCTTGGCATTGCCCGCCCTCCGGTCCGCAATCGGAAGCGTCTGGAGCTTGC
TAGTCGCTTCCGGCTTGAGAATGCGTTCTGTAGTGTGACTTGGAGCATCTTTCTA
GGAAGTAAACGCGTCTAAATTGAGACGCGGACTTCCGGCATTGCTGAAAAGCTTCCG
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