Data in Brief 7 (2016) 1670-1677



Contents lists available at ScienceDirect

Data in Brief

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Data Article

Data for amino acid alignment of Japanese stingray melanocortin receptors with other gnathostome melanocortin receptor sequences, and the ligand selectivity of Japanese stingray melanocortin receptors



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ARTICLE INFO

Article history: Received 28 March 2016 Received in revised form 14 April 2016 Accepted 19 April 2016 Available online 26 April 2016

Keywords: Adrenocorticotropic hormone (ACTH) Dasyatis akajei Melanocortin receptor (MCR) Melanocyte-stimulating hormone (MSH) Stingray

ABSTRACT

This article contains structure and pharmacological characteristics of melanocortin receptors (MCRs) related to research published in "Characterization of melanocortin receptors from stingray *Dasyatis akajei*, a cartilaginous fish" (Takahashi et al., 2016) [1]. The amino acid sequences of the stingray, *D. akajei*, MC1R, MC2R, MC3R, MC4R, and MC5R were aligned with the corresponding melanocortin receptor sequences from the elephant shark, *Callorhinchus milii*, the dogfish, *Squalus acanthias*, the goldfish, *Carassius auratus*, and the mouse, *Mus musculus*. These alignments provide the basis for phylogenetic analysis of these gnathostome melanocortin receptor sequences. In addition, the Japanese stingray melanocortin receptors were separately expressed in Chinese Hamster Ovary cells, and stimulated with stingray ACTH, α -MSH, β -MSH, γ -MSH, δ -MSH, and β -endorphin. The dose response curves reveal the order of ligand selectivity for each stingray MCR.

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DOI of original article: http://dx.doi.org/10.1016/j.ygcen.2016.03.030

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http://dx.doi.org/10.1016/j.dib.2016.04.050

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Subject area	Biology
More specific sub- ject area	Endocrinology
Type of data	Text files, graphs
How data was acquired	Amino acid sequences were aligned using MEGA 6.0. Ligand selectivity assays were done using the CRE/Luciferase reporter assay [2]. Luminescence was measured using a Bio-Tek Synergy HT plate reader (Bio Tek, Winooski, VT, USA), and the data were analyzed and graphed using Kaleidagraph software (Synergy Software, Reading, PA, USA)
Data format	Raw
Experimental	Melanocortin DNAs were cloned from stingray genomic DNA or brain mRNA.
factors	Cloned DNA were expressed in Chinese Hamster Ovary cells
Experimental	Sequence alignment was done using MEGA 6.0. The ligand selectivity assays
features	were done as described in reference [3].
Data source	Kitasato University, Sagamihara, Kanagawa, Japan. University of Denver,
location	Denver, Colorado, USA
Data accessibility	Data is within this article

Specifications Table

Value of the data

- These data are valuable for researchers participated in endocrinology of primitive fish and evolution of melanocortin systems.
- These could be used as probes to explore orthologs in other cartilaginous fish such as skates, sharks and chimaeras.
- The data on ligand selectivity could be useful tools for structure–function relationship studies in endocrinology and pharmacology.

1. Data

Data provided in this article show amino acid sequence comparison of melanocortin receptors (MCRs) in vertebrates and ligand selectivity of stingray MC peptides on these receptors. The amino acids sequences of MC1R (Fig. 1), MC2R (Fig. 2), MC3R (Fig. 3), MC4R (Fig. 4), and MC5R (Fig. 5) of stingray (*Squalus acanthias*) which determined by us [1] were compared to corresponding sequences from two species of other cartilaginous fishes (i.e., *Callorhinchus milii*, elephant shark and *S. acanthias*, dogfish), a teleost (*Carassius auratus*, goldfish), and a mammal (*Mus musculus*, mouse). Data are also provided for ligand selectivity include effects of stingray Des-acetyl- α -MSH, β -MSH, γ -MSH, δ -MSH, ACTH(1-24) and β -endorphin on MC1R, MC3R, MC4R, and MC5R (Figs. 6, 8–10) and those of stingray Des-acetyl- α -MSH, ACTH(1-24), human ACTH(1-24) and NDP-MSH on stingray MC2R (Fig. 7).

2. Experimental design, materials and methods

In order to align the amino acid sequences of the melanocortin receptors for the Japanese stingray, *D. akajei*, the dogfish, *S. acanthias*, the elephant shark, *C. milii*, the goldfish, *C. auratus*, and the mouse, *M. musculus*, it was essential to identify putative transmembrane domains in each receptor sequence. To this end, the program "MEMSAT3" (http://bioinf.cs.ucl.ac.uk/psipred/) was used. The amino acid sequences where then aligned using the program MEGA 6.0.

To functionally express and determine the ligand selectivity of the stingray (sr) MC1R, srMC2R, srMC3R, srMC4R, and srMC5R paralogs, the nucleotide sequences for the *srmcrs* were separately

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Dasyatis akajei Callorhinchus milii Carassius auratus Mus musculus	MMNITTLAPR MNDSSRH MS.QE	GNEQKDISFH E.LRL. YFSM.HMDYI SLLGS	WLPRNVN HGA. YNID.NITL. LNSNATSHLG	NSY-NASSMQ S.SM.AVA TTLGEMNATG LATQ.EPW	CKHINIPEEV .Q.VSV IAQ.M.Q.L .LYVSDGL	FLTLGILSFV SV.L. MLI.L. SLV.L.	56 42 57 52
Dasyatis akajei Callorhinchus milii Carassius auratus Mus musculus	ENILVIIAII MT VA VVT	KNQNLHSPMY RH R	YLICCLAMAD .FA. .FVS. .FLS.	TLVSMSNTIE MVMV. MVVV. LMV.IVL.	TIVLILMEKE RG .LFML.K.HG .TIIL.L.AG	VLTVQNHILK .MVYL L.L.TAKM.Q I.VARVALVQ	116 102 117 112
Dasyatis akajei	QIDNIIDLMI	CTSMVSSLSF	LAAIAADRYI	TIFYALRYHV	IMTTRKAVII	IVGIWIVSCT	176
Callorhinchus milii	LM	.S	.G	T	RGV	MVI.	162
Carassius auratus	HLVI	.S.V	.CT	S	QRA.	.AVV.LT.I.	177
Mus musculus	.LLVL.	.GC.	.GII	SS	.V.LPR.RRA	VMIV	172
Dasyatis akajei	SSIMFIVYSE	SSAVIICLIS	FFFMMLVIMG	GLYFHMFMLA	QMHTKKIMAQ	RKKRP-THQA	235
Callorhinchus milii	AI	NV	V.VIF	ALT	RI.A.R	HT-L	221
Carassius auratus	SLHT	DNAVT	GLTFTA	VLI	HV.SRRL	H.SR	234
Mus musculus	TLT.YK	HTLLVT	LAAL.A	IATR.	CQ.AQG.AQL	H.R.RSIR.G	232
Dasyatis akajei	ANMKGAITLT	ILLGLFLICW	SPFFLHLLLI	ISCPKNPYCL	CFNSHFNMFL	ILIICNSVFD	295
Callorhinchus milii	TS	V	I	.LTQ	T	I.	281
Carassius auratus	TS	V.V	GI	LITK	.YFL.	LI.	294
Mus musculus	FCLA	I.FL.	G	VLQH.T.S	.IFKNL.	LVLS.TV.	292
Dasyatis akajei Callorhinchus milii Carassius auratus Mus musculus	PIIYAFRSQE .LY .L	LRKTLKEFIP	CSW 318 284 LFAM 322 315	3 1 1 5			

Fig. 1. Amino acid sequence comparison of MC1R used for phylogenetic analysis. Species names are *Dasyatis akajei* for stingray, *Callorhinchus milii* for elephant shark, *Carassius auratus* for goldfish, and *Mus musculus* for mouse. Dot shows identical amino acid to stingray sequence. Hyphen shows gap. Accession numbers: LC108746 (*Dasyatis akajei*), BR000855 (*Callorhinchus milii*), AB618067 (*Carassius auratus*), and BC119296 (*Mus musculus*). The percent identify for the MC1R orthologs was 33%.

Dasyatis akajei Callorhinchus milii Carassius auratus Mus musculus	MPDMMIPGYG	TLLDSNGILP	MPPDATISPH MSG	SHPTISPWLP ADTSAA MK	YGTEVVIDTI NV.TA.MN.S MNSS.E HIINSYEH.N	NQTNMNATEE GFM.G-SGGI ALSTHPTD DTARNNSD	60 32 14 20
Dasyatis akajei	CSQIEIPTEV	YLILGLVSLL	ENLLVVIAVL	KNKKLHFPMY	FFICSLAVSD	ILLCLSKAWE	120
Callorhinchus milii	.R.LL	G.GM.	IV	N.RNS	LMA.	M.VSVGS.	92
Carassius auratus	.AEVQV.SQ.	FMAIAVAS	I.ILI	RNS	CNFN	TISS.C.SL.	74
Mus musculus	.PDVVL.E.I	FFTISVIGI.	I.LL.I	N.QS	I.	M.GS.Y.IL.	80
Dasyatis akajei	AFTISLVNNH	EDLFIQTFLL	SLDNVFDTLI	CISFLASIFN	LAAITTDRYI	SIFHALRYHN	180
Callorhinchus milii	.VI.F.DQ	SH.LTE.LID	HYLS	LILS	.GAL	TQ	151
Carassius auratus	TILLLFKEAG	HLNGR.E.	NI.DIM.S.L	.MCGS	ILT.AV	T	132
Mus musculus	NIL.MFR.MG	YLKPRGS.ES	TA.DII.CMF	IL.L.GS	.SV.AA	TQS	140
Dasyatis akajei	IMTGKRVAFA	IAGIWVFCTA	TGILMINFHN	SQGIISFYII	FFLLSVVLIV	SLYIYMFLLA	240
Callorhinchus milii	VA.LI	.SAL.TF	S.SFI.K.NR	KNAFPGSL.T	MYFTTLFV	V	211
Carassius auratus	LMRVVT	LSTGT	S.VG.S.	AATVKISSLC	SSSTALLL	LVH	192
Mus musculus	.V.MR.TIIT	LTIMG	ST.VI.SH	HIPTVLTFTS	L.P.ML.F.L	CH	200
Dasyatis akajei Callorhinchus milii Carassius auratus Mus musculus	QMHARKIRIL RRQCS. RHNR.ASM RSST.	PG-HTAHQGI QRVT L.ARQRQS RT	NFKGAFTVTV SLI.L.I GLRL.L.I .MM.L.I	LLGVFIFCWA II .IVA	PLSLHFILFL .FFLV. .FLLISM .FVVL.MT	LCPSDPYCAC ANT. IENE. FNNV.	299 270 252 253
Dasyatis akajei	FMSLFQIDLI	FIMCHSIIDP	LIYAFRDPEL	SNTFKKMMFC	HKKQWYFHAS	PSFLNI 355	
Callorhinchus milii	YV	LN	FS	RCI.	FNL.	316	
Carassius auratus	YRLHVL	LLVS.AV.E.	AST	RYVFL.	SASRIFKECV	302	
Mus musculus	YVNGM	LNAV	FS	RDAR.L	NRY	296	

Fig. 2. Amino acid sequence comparison of MC2R used for phylogenetic analysis. Species names are *Dasyatis akajei* for stingray, *Callorhinchus milii* for elephant shark, *Carassius auratus* for goldfish, and *Mus musculus* for mouse. Dot shows identical amino acid to stingray sequence. Hyphen shows gap. Accession numbers: LC108747 (*Dasyatis akajei*), BR000856 (*Callorhinchus milii*), AB618068 (*Carassius auratus*), and NM_008560 (*Mus musculus*). The percent identity for the MC2R orthologs was 24%.

synthesized with a V-5 epitope tag at the N-terminal of the receptor, and inserted into a pcDNA3.1 expression vector (GenScript; Picataway, NJ, USA). Each *srmcr* cDNA was separately transiently transfected into Chinese Hamster Ovary (CHO) cells. The CHO cells were grown at 37 °C in a humidified 5%

Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	MNSTPSISF H MHI. DSYLQFL SCCL.S	FHPAMRNGTE LFDLQLSG LQLPTM.S KGQKPA.S.S VS.MLP.LS.	DLNESSILNN K NV LPPNG.TVD- HPAAPPAS	RNGTGFCEQV .SNP .SSA PPAGAL .S.S	PIKAELFFCL S.V.LT. V.LI. Q.Q.V.LT. FP.V.LA.	GIISFLENVL LI. L.LI. V.LI. V.LM.I.	59 55 60 58 58
Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	VILAVAKNKN IL V VR.G.	LHSPMYLFLC F F F F	SLAVADMLVS	VSNALETIVM SI LSMI	AFLKNGFLIA .L.NY.V. NY.V. .V.NSRL.V. .VINSDS.TL	NDQLIQQMDN F.I S.HFVRL EF.H	119 115 120 118 118
Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	VFDSMICISL .IL 	VASICNLLVI	AIDRYITIFY 	ALRYHSIMTV	KRAIILIVVI LI L.DRSY RLVA.AG. RK.LTG.	WIFCIFCGII A LDCLY .LV.VVV .VC.GIVM	179 175 180 178 178
Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	FIIYSESQTV D.K.A VK. KM.	IICLITMFFV T .VA .VA	MLFLMTTLYV V .SS .V.A .VL.GI	HMFMLARLHI 	KRIATLPVPG .KAD. N. QAPAA QVPA.	VVHQ I.RP M.R AAAGNPAPR. APQ.	233 229 234 238 234
Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	RTCMKGAITI 	TILLGIFIIC V. SI.V.VC. V.F.	WAPFFLHLIM L L VL	IISCPKNPYC A LVHH.L. TT	ICYTSHFNTY 	LILIMCNSVI 	293 289 294 298 294
Dasyatis akajei Callorhinchus milii Squalus acanthias Carassius auratus Mus musculus	DPIIYAFRSQ M LCL LL	EMRKTFKEIL A I 	CCYCMNFNFR G.L.S. L.L. FGCQPPL .GCNSMNLG	CK 325 Y 322 326 327 323			

Fig. 3. Amino acid sequence comparison of MC3R used for phylogenetic analysis. Species names are *Dasyatis akajei* for stingray, *Callorhinchus milii* for elephant shark, *Squalus acanthias* for dogfish, *Carassius auratus* for goldfish, and *Mus musculus* for mouse. Dot shows identical amino acid to stingray sequence. Hyphen shows gap. Accession numbers: LC108748 (*Dasyatis akajei*), BR000857 (*Callorhinchus milii*), AY560605 (*Squalus acanthias*), AB618069 (*Carassius auratus*), and NM_008561 (*Mus musculus*). The percent identity for the MC3R orthologs was 52%.

Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	MDLSYTRGPA .NS.FHHRLP .NT.HHHH .NSTHHH.MY	DTTQNRNQSV E.P.L.H HSY.H.Q TSLHLW.R.S	SGFTG-ANIL AR.ASGS GALPV-GKPD Y.LH.N.SES	HSN-GSSSGC R.DF QGERT LGKGHPDG	NEQLWISTEV Y YL YFV.P.	FVMLGIVSLL .LTF .LTL TVI	58 58 57 60
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	ANILVVAAIV I EII EIVA	KNKNLHSPMY	FFICSLAVAD	MLISVSNAWE V L.VS. VGS.	TITIAMLKSR F .VVM.LITGG V.TL.N.T	HLLAQDKLIK T.PEN N.TYRESI DTDS-FTV	118 118 117 119
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	SIDNVFDSVI NMM. NMIM. NI.	CSSLLASICS	LLAVAVDRYI I.I I SIF	TIFYALRYHN	IMTVRRALTV .VMI QG.I VGII	IAGIWAVCIG AA.T. .TCTL.TV .SCA.TV	178 178 177 179
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	SGILFIIYSE V VV VD	STTAVICLIA AVIT VLS .SAVIS	MFFAMLAIMA L TL TVL	SLYVHMFMLA L. LM.	RLHLKRIAAL V M V.	PSSGAICQAA .GNVR .GN.P.W .GT.T.R.GT	38 238 237 239
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	NMKGAITLTI	LLGVFVVCWA	PFFLHLILMI M LFY.	SCPRNPYCIC QV. QV.	FMSHFNMYLI	LILCNSIIDP MV MV MAV	298 298 297 299
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	LIYAFRSQEM	RKTFKEIICC	YS-LRGSCDL P.L WYG.ASL.V .PG.I.E.	LSNINTH 33 T.EY 33 32 S.RY 33	4 1 6 2		

Fig. 4. Amino acid sequence comparison of MC4R used for phylogenetic analysis. Species names are *Dasyatis akajei* for stingray, *Squalus acanthias* for dogfish, *Carassius auratus* for goldfish, and *Mus musculus* for mouse. Dot shows identical amino acid to stingray sequence. Hyphen shows gap. Accession numbers: LC108749 (*Dasyatis akajei*), AY169401 (*Squalus acanthias*), AJ534337 (*Carassius auratus*), and BC116959 (*Mus musculus*). The percent identity for the MC4R orthologs was 55%.

Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	MQDQSPVNRR	FNSQKPPGTR	MNLTKL G. MTSEA EESCLP.RGA	S QS TLS EQNGKSDAKK	LMSV REPWPKNL LWAISANS WGHSLPA.NS	TLAEAMVNGT .P.NDIT.R. SPVLDLL.T. SSTLTVL.L.	21 26 28 60
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	RSST K.TS ETPSHA LNASEDGILG	ALC G KPKA. SNVKNKSLA.	EQVSVAVEVF I .LNI.T .EMGI	LILGIISLLE .TM .TLV	NILVIAAIIK C.V. G.V.	NRNLHSPMYF .K .K .K	68 73 79 120
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	FVCSLAVADM Y.G	LVSVSNAWET	IVITLLHSRH AN YTN.Q VT.YNNK.	LVVKDSFVKH .I.EQ E.H.IRQ IA.TR.	VDNVFDSMIC M I	TSVVASMCSL I I I	128 133 139 180
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	LAIAVDRYIT	IFYALRYHHI N.	MSMKRAAFII .TVT .TVR .TAR.SGV	AGIWALCIGC TF GTF.TS. .CTFS.	GIIFIIYSES DN VDN VY	PTVIICLVTM IA. TSVS. KYIS.	188 193 199 240
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	FFIMLLIMAS VL AL TFF.V.	LYSHMFLLAR M I	SHAKRIAAMS L. VLP N.VSP	SSNSIHQQAS .YR GYR RYVR.RT.	MKGAITLTIL 	LGIFIVCWAP	248 253 259 300
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	LFLHLILMIS F F F	CPRNLYCTCF GV M Q.VS	MSHFNLYLIL M YM	IMCNSIIDPL	IYSFRSQEMR AW A AL	KTLKEIICCH FY Y R.FV	308 313 319 360
Dasyatis akajei Squalus acanthias Carassius auratus Mus musculus	SLRAVCRLTV A.G.SG N.FGMSR GF.RPLG	K 319 . 324 329 GY 372					

Fig. 5. Amino acid sequence comparison of MC5R used for phylogenetic analysis. Species names are *Dasyatis akajei* for stingray, *Squalus acanthias* for dogfish, *Carassius auratus* for goldfish, and *Mus musculus* for mouse. Dot shows identical amino acid to stingray sequence. Hyphen shows gap. Accession numbers: LC108750 (*Dasyatis akajei*), AY562212 (*Squalus acanthias*), AJ576322 (*Carassius auratus*), and BC100720 (*Mus musculus*). The percent identity for the MC5R orthologs was 61%.



Fig. 6. Ligand selectivity of stingray MC1R. (A) Functional activation of the stingray MC1R after stimulation with the following stingray melanocortins: Des-acetyl- α -MSH (Des-Ac- α -MSH), ACTH(1-24), or β -MSH. (B) Functional activation of stingray MC1R after stimulation with the following stingray melanocortins: Des-Ac- α -MSH, γ -MSH, δ -MSH or β -endorphin(1-20). As described in methods, CHO cells were transiently transfected with a stingray *m*c1r cDNA construct and a *cre/luc* cDNA construct. Two days post-transfection, wells containing 1 × 10⁵ cells were stimulated with the stingray melanocortin ligands at concentrations ranging from 10⁻⁶ M to 10⁻¹² M. Results are expressed as mean \pm S.E.M.; *n*=3.



Fig. 7. Ligand selectivity of stingray MC2R. (A) Functional activation of the stingray MC2R after stimulation with stingray Desacetyl- α -MSH (Des-Ac- α -MSH) or stingray ACTH(1-24). (B) Functional activation of stingray MC2R after stimulation with human ACTH(1-24) (hACTH(1-24)) or NDP-MSH. The activation assays were performed as described in the figure legend for Fig. 6. Results are expressed as mean \pm S.E.M.; n=3.



Fig. 8. Ligand selectivity of stingray MC3R. (A) Functional activation of the stingray MC3R after stimulation with the following stingray melanocortins: Des-acetyl- α -MSH (Des-Ac- α -MSH), ACTH(1-24), or β -MSH. (B) Functional activation of the stingray MC3R after stimulation with the following stingray melanocortins: Des-acetyl- α -MSH (Des-Ac- α -MSH), γ -MSH, δ -MSH or β -endorphin(1-20). The activation assays were performed as described in the figure legend for Fig. 6. Results are expressed as mean \pm S.E.M.; n=3.

CO₂ incubator in DMEM/F12 with 5% fetal calf serum. Each sr cDNA was co-expressed with a CRE/ Luciferase reporter plasmid [2] using the Solution T Cell Line Nucleofector Kit (Amaxa Inc., Gaithersburg, MD, USA) and program U-23 [4]. The transiently transfected cells were seeded on a 96-well plate at a density of 1×10^{-5} cells/well. After 48 h in culture, the transfected cells were stimulated with either synthetic srACTH(1-24), srDes-acetyl- α -MSH, sr β -MSH, sr γ -MSH, sr δ -MSH, sr β -endorphin or hACTH (1-24), or NDP-MSH at concentrations ranging from 10^{-6} M to 10^{-12} M, in serum-free CHO media for four hours at 37 °C. At the end of the incubation period, 100 µl of Bright-Glo luciferase assay reagent



Fig. 9. Ligand selectivity of stingray MC4R. (A) Functional activation of the stingray MC4R after stimulation with the following stingray melanocortins: Des-acetyl-α-MSH (Des-Ac-α-MSH), ACTH(1-24), or β-MSH. (B) Functional activation of stingray MC4R after stimulation with the following stingray melanocortins: Des-acetyl-α-MSH (Des-Ac-α-MSH), γ-MSH, δ-MSH or β-endorphin(1-20). The activation assays were performed as described in the figure legend for Fig. 6. Results are expressed as mean ± S. E.M.; n=3.



Fig. 10. Ligand selectivity of stingray MC5R. (A) Functional activation of the stingray MC5R after stimulation with the following stingray melanocortins: Des-acetyl- α -MSH (Des-Ac- α -MSH), ACTH(1-24), or β -MSH. (B) Functional activation of stingray MC5R after stimulation with the following stingray melanocortins: Des-acetyl- α -MSH (Des-Ac- α -MSH), β -MSH or β -endorphin(1-20). The activation assays were performed as described in the figure legend for Fig. 6. Results are expressed as mean \pm S. E.M.; n=3.

(Promega Inc., Madison, WI, USA) was added to each well, and incubated for 5 min at room temperature. Luminescence was measured with a Bio-Tek Synergy HT plate reader (Bio Tek, Winooski, VT, USA), and the dose response curves were analyzed by using Kaleidagraph software (Synergy Software, Reading, PA, USA). All experimental treatments were performed in triplicate.

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Acknowledgments

We thank Mr. Kosuke Arai, Kitasato University, and Dr. Yasuhisa Kobayashi, Dr. Naoaki Tsutsui and Mr. Kazuhiro Saito, Okayama University, for technical assistance. Partial support for this study was provided by the Long Research Fund (RMD) and National Science Foundation, U.S.A. Grant IOB 0516958 Supplement (RMD).

Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at http://dx.doi. org/10.1016/j.dib.2016.04.050.

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