

IR: infrared LSCM: laser-scanning confocal microscope







Kagawa-Nagamura et al Figure S4

Strain	Genotype Reference or Source	
N2	wild type	Brenner, Genetics, 1974
RB993	tdc 1 (ak014) II	The C. elegans Deletion Mutant
		Consortium, G3, 2012. OMRF ^a
RB1161	tbb-1 (ok1196) X	The C. elegans Deletion Mutant
		Consortium, G3, 2012. OMRF ^a
RB1690	ser-2 (ok2103) X	The C. elegans Deletion Mutant
		Consortium, G3, 2012. OMRF°
VC125	tyra-3 (ok325) X	The C. elegans Deletion Mutant
<u></u>		Consortium, G3, 2012. OMRE
GR1321	tpn-1 (mg280) cam-1(vs166) II	Sze Jy et al., Nature, 2000
MT14680	Igc-55 (n4331) V	Ringstad N et al., Science, 2009
QJ6	cat-2 (jq6)	Usami A, Ph.D. Thesis, 2012
QJ3063	jqls6	This study
QJ3064	jqls71	This study
QJ3065	jqls72	This study
QJ3067	jqls75	This study
QJ1412	jqEx439	This study
QJ4110	tph-1 (mg280) II; jqIs6	This study
QJ4111	cat-2 (jq6) II ; jqls6	This study
QJ1359	ser-2 (ok2103) X ; jqEx386	This study
QJ1501	ser-2 (ok2103) X ; jqEx528	This study
QJ1502	ser-2 (ok2103) X ; jqEx529	This study
QJ1526	ser-2 (ok2103) X ; jqEx553	This study
QJ1543	ser-2 (ok2103) X ; jqEx570	This study
QJ1544	ser-2 (ok2103) X ; jqEx571	This study
QJ4113	jąls73 ; jąls75	This study
QJ4114	jqls6 ; jqEx318	This study
QJ1489	jqls75 ; jqEx516	This study
QJ4116	tdc-1 (ok914) II ; jqls6 ; jqEx318	This study
QJ4117	tbh-1 (ok1196) X ; jqls6 ; jqEx318	This study
QJ4118	ser-2 (ok2103) X ; jqls6 ; jqEx318	This study
QJ4119	lgc-55 (n4331) V ; jqls6 ; jqEx318	This study
QJ4120	tyra-3 (ok325) X ; jqls6 ; jqEx318	This study
QJ4122	ser-2 (ok2103) X ; jqls6 ; jqEx386	This study
QJ4123	ser-2 (ok2103) X ; jqls6 ; jqEx422	This study
QJ4112	ser-2 (ok2103) X ; jqls6 ; jqEx528	This study

 Table S1
 C. elegans
 strains used in this study

^aOMRF: Oklahoma Medical Research Foundation, The *C. elegans* Deletion Mutant Consortium

Transgenic array	DNA injected		
jqls6	Рипс-25::G-CaMP7 (100 ng/µl) Punc-25::DsRedx (50 ng/µl)		
jqls71	Punc-25::G-CaMP7 (ce) (50 ng/μl) Punc-25::DsRedx (50 ng/μl) pBluescript II SK+ (100 ng/μl)		
jqls72	Punc-25::G-CaMP7(ce) (50 ng/μl) Punc-25::DsRedx (50 ng/μl) pBluescript II SK+ (100 ng/μl)		
jqls73	Punc-25::G-CaMP7(ce) (50 ng/μl) Punc-25::DsRedx (50 ng/μl) pBluescript II SK+ (100 ng/μl)		
jqls75	Ptdc-1::G-CaMP7(ce) (50 ng/μl) Ptdc-1::DsRedx (50 ng/μl) Ptbh-1::EBFP2 (50 ng/μl) pBluescript II SK+ (50 ng/μl)		
jqEx318	Psra-6::ChR2::mCherry (100 ng/μl) Punc-122::mCherry (50 ng/μl) pBluescript II SK+ (50 ng/μl)		
jqEx386	Punc-25::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		
jqEx422	Punc-17::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		
jqEx439	Punc-25::R-CaMP2 (50 ng/μl) Punc-47::EGFP (50 ng/μl) pBluescript (50 ng/μl)		
jqEx516	Punc-47::EGFP (50 ng/μl) pBluescript II SK+ (150 ng/μl)		
jqEx528	Pser-2prom2::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		
jqEx529	Pser-2prom2::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		
jqEx553	Punc-25::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (110 ng/μl)		
jqEx570	Punc-30::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		
jqEx571	Punc-30::ser-2 (+) (20 ng/μl) Punc-122::EYFP (50 ng/μl) pBluescript II SK+ (130 ng/μl)		

 Table S2
 Transgenic lines generated in this study

Genotype	Mutation	Primer name	Sequence
cat-2 (jq6) II	320bp deletion + 9bp insertion	cat-2_jq6_forward	CTTCACACGACGTGCTCCTA
		cat-2_jq6_reverse1	GTTGTCTGGAAGACCCGAAA
		cat-2_jq6_reverse2	CTTGAACTCTAACGCCTGATC
lgc-55 (n4331) V	1986bp deletion	lgc-55_n4331_forward	CGGTGGTCAGTGGATAATTTTGTGC
		lgc-55_n4331_reverse1	GCGATTTGTGAATTGCAGAGCC
		lgc-55_n4331_reverse2	CGTCATCTATGAATTCGACCGGTTG
	2043bp deletion	ser-2_ok2103_forward1	GTGATGGTACAAGCTTCACC
ser-2 (ok2103) X		ser-2_ok2103_reverse1	CTTCGTCATTTGTGGTTTCG
		ser-2_ok2103_forward2	CTCCATCATATTCTCCTGCC
	981bp deletion	tbh-1_ok1196_forward	GGAAAGGACAATGGCACGTTGA
tbh-1 (ok1196) X		tbh-1_ok1196_reverse1	AGGTGCTGAATTTTGCCTGCC
		tbh-1_ok1196_reverse2	CGAAGGTCCAACTGATCAGCCAA
	629bp deletion	tdc-1_ok914_forward	GCTATGGCTTCATGTTGATGCTGC
tdc-1 (ok914) II		tdc-1_ok914_reverse1	GCCGTTATCAAAGGCGGAAACT
		tdc-1_ok914_reverse2	GCACATACACAGAAGCGGATCAC
	1305bp deletion	tph-1_mg280_forward	GCAAAGACCCCTCTCAACCTCATT
tph-1 (mg280) II		tph-1_mg280_reverse1	GGAAACCATTCAGAACCGGTTGTC
		tph-1_mg280_reverse2	GCATGATGGCGAACGTATTGAGTG
	X 781bp deletion	tyra-3_ok325_forward1	GTTCTGAAACAGTTCACGGG
tyra-3 (ok325) X		tyra-3_ok325_forward2	GATTGCCTCATGAGAGCTTC
		tyra-3_ok325_reverse1	GGAGAAGAGTCCGTTTATCG

Table S3 Mutations and primers for genotyping

Table S4 Primers for molecular biology

No.	primer name	Sequence	Comments	
1	G-CaMP7(CE)_Nhel#F1	CTTACTAGTGCTAGCATGAGAGGAAGTCATCA TCA	1 and 2 were used to amplify the 1.4 kb codon-optimized G-CaMP7 cDNA for <i>C</i> .	
2	G-CaMP7(CE)_NheI#R1	TTTTTTTGCTAGCTTATTTGGCAGTCATCATC T	elegans	
3	R-CaMP2_Bgl#R1	TTTTAGATCTCTATGGGTTGGACTCCACGT	3 and 4 were used to amplify the 1.5 kb R- CaMP2 cDNA	
4	R-CaMP2_Not#F1	CTTACTAGTGCGGCCGCATGGGTTCTCATCAT CATCATC		
5	DsRedx_NheI#F1	CTTACTAGTGCTAGCATGGCCTCCTCCGAGGA CGT	5 and 6 were used to amplify the 0.68 kb DsRed-Express-1 cDNA	
6	DsRedx_NheI#R1	TTTTTTTGCTAGCCTACAGGAACAGGTGGTG GC		
7	ser-2 cDNA Bgl#R1	TTTTAGATCTCTAAGGTTGCGCACTCATTC	7 and 8 were used to amplify the 1.3 kb full-length <i>ser-2</i> cDNA	
8	ser-2 cDNA Not#F1	CTTACTAGTGCGGCCGCATGTTCCGAAATTAC ACTGA		
9	ser-2prom2#F1	ATTGGCTGCAGATTTTATGACTTTCACTAGAA ATG	9 and 10 were used to amplify the 4.7 kb ser-2prom2 promoter	
10	ser-2prom2#R1	AGCTGGATCCTTTTGCAAATTACTTGAGGCTG		
11	tbh-1 promoter#F1_Bam	AGCTGGATCCCCTTATGTATCTCATTGCTATTT	11 and 12 were used to amplify the 2.8 kb <i>tbh-1</i> promoter	
12	tbh-1 promoter#R1_Not	CTTACTAGTGCGGCCGCTTTCTGAAATCGTAT AGTAATGG		
13	tdc-1p_BamHI#R1	AGCTGGATCCTTGGGCGGTCCTGAAAAATG	13 and 14 were used to amplify the 4.4 kb <i>tdc-1</i> promoter	
14	tdc-1p_HindIII#F1	TTTTTTTTAAGCTTAAGGGAGAGAGAGTATTGCA GTGG		
15	unc-25p_BamHI#R1	AGCTGGATCCTCTTTTTTGGCGGTGAACTG	15 and 16 were used to amplify the 1.9 kb <i>unc-25</i> promoter	
16	unc-25p_HindIII#F1	TTTTTTTTAAGCTTTGTGCATGCAAAAAAACACC C		
17	unc-30_SphI#F1	TTTTTTTTGCATGCTCGATGACACTTTTTGATG TGT	17 and 18 were used to amplify the 2.4 kb <i>unc-30</i> promoter	
18	unc-30_BamHI#R1	AGCTGGATCCGTGTTGATCTTTTGCCGGAG		
19	unc-54 3'#R1	GACTTAGAAGTCAGAGGCAC	19-25 were used to verify the sequences of DNA constructs	
20	unc-86_UTR#R1	CAAAGCGAATTGCTAAACCC		
21	pFX#F1	GCAAGGCGATTAAGTTGGGT		
22	pPD49.26#F1	CGTTGGCCGATTCATTAATG		
23	ser-2#R2	GTAGACAACCCGCAACTATC		
24	ser-2 cDNA #F2	CATTGGGATTATAGTCACGC		
25	unc-25p#F1	CGGAATAGGGAAAAGCTCAG		