

**Supplementary file 1A: Plasmid used in this study**

Short name	Plasmid backbone	Insert	Cloning technique	Primers <sup>a</sup> or gBlocks <sup>b</sup>	
pIK155	published in (Katic et al., 2015)				
pIK198	published in (Katic et al., 2015)				
pCFJ90	published in (Frøkjær-Jensen et al., 2008)				
pCFJ104	published in (Frøkjær-Jensen et al., 2008)				
pFA27	published in (Aeschmann et al., 2019)				
pFA224	pENTR L4-R1	<i>col-10</i> promoter	gift from Matyas Ecsedi. The <i>col-10</i> promoter starts is 1320 bp long and starts with the following sequence: 5'-CCTCGGGTGGTCATCATCTCAAATCCAATCTTCTTTTCATTTTCAATCTCTAACAAATTTCAAATTACTAGAACCCACAAACCAATGTTTCGATA ... - 3'		
pFA198	pENTR L1-L2	3xFLAG tag followed by BamHI restriction site (can be used for Gibson assembly instead of BP reaction)	BP reaction <sup>c</sup>	fwd primer	ggggacaagttgtacaaaaaagcaggctATGGATTATAAAGACGATGACGATAAGCGTGA
				rev primer	ggggaccactttgtacaagaaagctgggtGGATCCTCTCTTGTTCATCGTCATCCTTGTAAATC
pFA199	pENTR L1-L2	3xFLAG:: <i>lin-29a</i> (ORF and 3'UTR; including introns except for intron 4)	Gibson assembly <sup>d</sup>	fwd(1)	acaaggatgacgatgacaagagaATGGATCAAAGTCTTAGATTCGGCA
				rev(1)	gctttgttcgaaCTGTGGATATCTTCAAATTTGTGA
				fwd(2)	tggaagatccacagTTCGAACAAAAGCCGGACGT
				rev(2)	tgcccactttgtacaagaaagctgggtCCACTGTGCATGAAAAAGCTGGT
pFA206	pENTR L1-L2	3xFLAG:: <i>lin-29b</i> (ORF and 3'UTR, including introns)	Gibson assembly <sup>e</sup>	fwd primer	acaaggatgacgatgacaagagaATGCAGATGCGGGAAGCAAAAC
				rev primer	tgcccactttgtacaagaaagctgggtCCACTGTGCATGAAAAAGCTGGT
pFA207	pENTR L1-L2	3xFLAG:: <i>mab-10</i> (ORF and 3'UTR, including introns)	Gibson assembly <sup>e</sup>	fwd primer	acaaggatgacgatgacaagagaATGCATCATCGTCGTCGTC
				rev primer	tgcccactttgtacaagaaagctgggtACTATTGTTACGGGAATCATGTCT
pFA218	published in (Aeschmann et al., 2019)				
pFA219	pENTR L1-L2	FLAG-HA-degron:: <i>lin-29a</i> (ORF and 3'UTR; including introns except for intron 4)	Gibson assembly <sup>f</sup>	fwd primer	aggcggcggcgttcgtgaagATGGATCAAAGTCTTAGATTCGGCA
				rev primer	tgcccactttgtacaagaaagctgggtCCACTGTGCATGAAAAAGCTGGT
pFA220	pENTR L1-L2	FLAG-HA-degron:: <i>lin-29b</i> (ORF and 3'UTR, including introns)	Gibson assembly <sup>f</sup>	fwd primer	aggcggcggcgttcgtgaagATGCAGATGCGGGAAGCAAAAC
				rev primer	tgcccactttgtacaagaaagctgggtCCACTGTGCATGAAAAAGCTGGT
pFA221	pENTR L1-L2	FLAG-HA-degron:: <i>mab-10</i> (ORF and 3'UTR, including introns)	Gibson assembly <sup>f</sup>	fwd primer	aggcggcggcgttcgtgaagATGCATCATCGTCGTCGTC
				rev primer	tgcccactttgtacaagaaagctgggtACTATTGTTACGGGAATCATGTCT
pENTR_R2-L3 operon-GFP-H2b	published in (Ecsedi et al., 2015) based on (Merritt et al., 2008)				
pFA238	pCFJ150	pFA224, pFA219, pENTR_R2-L3 operon-GFP-H2b	LR reaction		
pFA239	pCFJ150	pFA224, pFA220, pENTR_R2-L3 operon-GFP-H2b	LR reaction		
pFA240	pCFJ150	pFA224, pFA221, pENTR_R2-L3 operon-GFP-H2b	LR reaction		

<sup>a</sup>Overhangs are in lowercase, the part annealing to the template in uppercase. If not stated otherwise, the PCR products were amplified from *C. elegans* genomic DNA.

<sup>b</sup>Overhangs for Gibson assembly reactions are in lowercase.

<sup>c</sup>PCR product was amplified from pFA27.

<sup>d</sup>Assembly with BamHI-digested plasmid pFA198 and two PCR products.

<sup>e</sup>Assembly with BamHI-digested plasmid pFA198 and one PCR product.

<sup>f</sup>Assembly with BamHI-digested plasmid pFA218 and one PCR product.

**Supplementary file 1B: oligonucleotides and sgRNAs used in this study**

Identifier	Sequence	Source	Description
CA1-2	ctaagcctaagtctatgcctaagcccaagcttgatctcaca atthaagctttcaaactagtgctaattaaag ggaatacggtttcagaattaaggagacacctagatcaatt gagctctaaagatccactatattcaatgtac ttacagttcgaacaaaagccggacgtgggggtgcttcag caacagatgcagatgcgg	This study	HR oligo for generation <i>lin-29(xe61 xe114)</i> and <i>lin-29(xe116)</i>
FA419	gaacggaatgatccgaaacctagatcaattgagctctaa agatccactatattcaatgtacttacagttc gaacaaaagccagatgtcgggggtgcttcagcaacagat gcagatgcgggaagcaaaccttaca gtgcacgcagtggtcaaggtatgtgt	This study	HR oligo for generation <i>lin-29(xe61 xe121)</i> and <i>lin-29(xe120)</i>
FA434	cctttcagcttcgagaggatcaagccaacttctcaacgca atgttttcacaattggaagataccacag gtttgataggggaacacattcaaacgaggggta	This study	HR oligo for generation <i>lin-29(xe61 xe133)</i> and <i>lin-29(xe200)</i>
3-AC15-16	acagaattaaggagacaagg	This study	sgRNA for generation <i>lin-29(xe61 xe114)</i> and <i>lin-29(xe116)</i>
2-AC2-3	agctcaattgatctaggttt	This study	sgRNA for generation <i>lin-29(xe61 xe114)</i> and <i>lin-29(xe116)</i>
L29_ATG1	gttcgaacaaaagccggacg	This study	sgRNA for generation <i>lin-29(xe61 xe121)</i> and <i>lin-29(xe120)</i>
L29_ATG3	agctgaagcaccaccacgctc	This study	sgRNA for generation <i>lin-29(xe61 xe121)</i> and <i>lin-29(xe120)</i>
L29_031	gctggaaccaccactggctc	This study	sgRNA for generation <i>lin-29(xe61 xe133)</i> and <i>lin-29(xe200)</i>
L29_041	gtggcaggagagaattctga	This study	sgRNA for generation <i>lin-29(xe61 xe133)</i> and <i>lin-29(xe200)</i>
L29_if1	agccaacttctcaacgcaa	This study	sgRNA for generation <i>lin-29(xe61 xe133)</i> and <i>lin-29(xe200)</i>
L29_if3	gtgaaaacatatgatgtggc	This study	sgRNA for generation <i>lin-29(xe61 xe133)</i> and <i>lin-29(xe200)</i>
lin-29_F4	ccagcacatcattcgatcact	(Aeschimann et al., 2017)	qPCR primer for <i>lin-29a</i> specific detection (exon3-exon4 junction)
lin-29_R4	gaagttcagtagatccgcttga	(Aeschimann et al., 2017)	qPCR primer for <i>lin-29a</i> specific detection (exon3-exon4 junction)

lin-29_F5	acccaagtttgagttcgaaca	(Aeschimann et al., 2017)	qPCR primer for <i>lin-29b</i> specific detection (SL1-exon5 junction)
lin-29_R5	gatgagttggcaaatgccttga	(Aeschimann et al., 2017)	qPCR primer for <i>lin-29b</i> specific detection (SL1-exon5 junction)
JKq17	gtcggagaccacgcatcaa	This study	qPCR primer for <i>act-1</i> specific detection (exon1)
JKq18	agggtaaggatacctctcttga	This study	qPCR primer for <i>act-1</i> specific detection (exon1-exon3 junction)