

Table S1. Reagents for CRISPR-mediated gene editing.		
Reagent	Sequence (5'-3')	Concentration injected
<i>Y51F10.10</i> crRNA	UUCAUCAAUGGAGCUUUUCCGUUUUAGAGCUAUG CUGUUUUUG	8 µg/mL
<i>Y51F10.10</i> repair DNA	ATTTTTACAGATTGAATACTACATGAAAGCATGCC GAACAGTCGAAAAAA <u>T</u> CGGGAA <u>G</u> AGCTCCATTGAT GAAACGTTGCCGTTCTTGATTCAATGCAATT ¹	0.6 µg/mL
<i>dpy-10</i> crRNA	GCUACCAUAGGCACCACGAGGUUUUAGAGCUAUG CUGUUUUUG	8 µg/mL
<i>dpy-10</i> repair DNA	CACTTGAACCTCAATACGGCAAGATGAGAATGACT GGAAACCGTACCGCATGCGGTGCCTATGGTAGCGGA GCTT CACATGGCTTCAGACCAACAGCCTAT	0.6 µg/mL
tracrRNA	AACAGCAUAGCAAGUAAAAUAAGGCUAGUCCGU UAUCAACUUGAAAAAGUGGCACCGAGUCGGUGCU UUUUUU	4 µg/mL
¹ Edits introduced into <i>Y51F10.10</i> by CRISPR are underlined and indicated in red; 'T' replaces 'C' and introduces the <i>hc85</i> mutation (Thr272Ile); 'G' replaces 'A' as a synonymous mutation that introduces a <i>SacI</i> restriction site for screening.		