Supplementary Information for

Genetic and Chemical Controls of Sperm Fate and Spermatocyte Dedifferentiation in Caenorhabditis elegans

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Table S1. C. elegans strains used in this study

Strain	Genotype	Citation
N2	wild-type (Bristol strain)	[1]
JK3231	puf-8(q725)	[34]
AH102	lip-1(zh15)	[68]
JK3961	puf-8(q725)/mln[mls14 dpy-10(e128)]; lip-1(zh15)	[23]
SD939	mpk-1(ga111) unc-79(e1068)	[31, 43]
JK3997	puf-8(q725)/mln[mls14 dpy-10(e128)]; lip-1(zh15); mpk-	[23]
	1(ga111) unc-79(e1068)	
JK816	fem-3(q20) gain-of-function	[35]
MHL58	puf-8(q725)/mln[mls14 dpy-10(e128)]; fem-3(q20gf)	This study
DG3913	tn1541[GFP::tev::s::lin-41]	[37]
MHL106	puf-8(q725)/mln[mls14 dpy-10(e128)]; fem-3(q20gf);	This study
	tn1541[GFP::tev::s::lin-41]	

Table S2. Antibodies used in this study

Antibody against	Resource
MSP (a marker for sperm fate)	DSHB (Cat#: 4A5)
HIM-3 (a marker for meiotic cells)	NOVUS (Cat#: 53470002)
GFP (Green Fluorescent Protein)	Abcam (Cat#: Ab290)
pERK1/2	Cell Signaling (Cat#: 9101)
ERK1/2	Cell Signaling (Cat#: 9102)
Tubulin	Sigma Aldrich (Cat#: T8203)

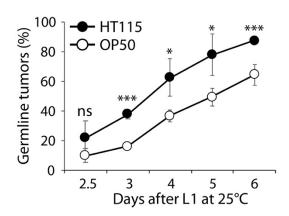


Figure S1. The effect of bacterial foods on spermatocyte dedifferentiation-mediated tumorigenesis in *puf-8(q725); fem-3(q20gf)* mutants at **25°C.** The *puf-8(q725); fem-3(q20gf)* mutants fed HT115 *E. coli* had significantly more germline tumors than those fed OP50 *E. coli.* ***, p<0.001; **, p<0.01; **, p<0.05; ns, not statistically significant.