

<b>Supplementary Table 2. <i>lite-1</i> mutations</b>			
Allele	Genomic sequence in N2 (wild type)	Genomic sequence in mutant	Effect on <i>lite-1</i> gene or protein
<i>ce302</i>	TTG <b>GT</b> AGT	TTT <b>AT</b> AGT	Splice donor site after end of 9 <sup>th</sup> exon
<i>ce303</i>	CTG <b>GAT</b> TC	CTG <b>AAT</b> TC	D214N
<i>ce304</i>	CTC <b>TGG</b> GA	CTC <b>TGA</b> GA	W419 Stop
<i>ce314</i>	GCG <b>GT</b> ATT	GCG <b>AT</b> ATT	Splice donor site after end of 6 <sup>th</sup> exon
<i>ce318</i>	CTT <b>CAG</b> GA	CTT <b>TAG</b> GA	Q106 Stop
<i>ce319</i>	TCT <b>CAA</b> AT	TCT <b>TAA</b> AT	Q376 Stop
<i>ce320</i>	TCC <b>AG</b> GGA	TCC <b>AA</b> GGA	Splice acceptor site before start of 9 <sup>th</sup> exon
<i>ce327</i>	TCC <b>AG</b> TGT	TCC <b>AA</b> TGT	Splice acceptor site before start of 6 <sup>th</sup> exon
<i>ce329</i>	TTA <b>CCA</b> GC	TTA <b>TCA</b> GC	P342S
<i>ce338</i>	CTT <b>TTG</b> GC	CTT <b>TTI</b> GGC	Insertion of T in 178 <sup>th</sup> codon causing frameshift
<i>ce340</i>	TAT <b>CAA</b> AG	TAT <b>TAA</b> AG	Q276 Stop
<i>ce341</i>	TGC <b>CCG</b> CT	TGC <b>TCG</b> CT	P259S
<i>ce342</i>	AAT <b>CAA</b> TC	AAT <b>TAA</b> TC	Q62 Stop
<i>ce343</i>	TCC <b>CGT</b> GC	TCC <b>TGT</b> GC	R222C
<i>ce345</i>	TTG <b>AG</b> GTG	TTG <b>AA</b> GTG	S126N (mutation is in last base of exon; cDNA change is <b>AGC</b> to <b>AAC</b> )
<i>ce312</i> , <i>ce316</i> , and <i>ce331</i> , isolated from separate mutageneses, have the same mutation as <i>ce302</i> .			