

Table S1 Molecular identity of Sacy mutations in *tn* alleles

Gene	Allele	Nucleotide change	Amino acid change ^a
<i>pde-6</i>	<i>tn1237</i>	CAG -> <u>T</u> AG	Q668*
	<i>tn1242</i>	TGG -> T <u>G</u> A	W444*
	<i>tn1336</i>	TGG -> T <u>G</u> A	W664*
<i>tom-1</i>	<i>tn1454</i>	Tc1 insertion	
	<i>tn1463</i>	GGT -> <u>A</u> GT	G498S
<i>sacy-1</i>	<i>tn1385</i>	GGA -> <u>A</u> GA	G533R
	<i>tn1391</i>	GGA -> <u>A</u> GA	G473R
	<i>tn1440</i>	GGA -> <u>A</u> GA	G331R
<i>twk-1</i>	<i>tn1397</i>	TGG -> T <u>A</u> G	W177*
	<i>tn1398</i>	TGG -> T <u>G</u> A	W330*
	<i>tn1403</i>	GAA -> T <u>A</u> A	E272*
<i>spr-4</i>	<i>tn1383</i>	CGA -> T <u>G</u> A	R1230*
	<i>tn1402</i>	CGA -> T <u>G</u> A	R1230*
	<i>tn1404</i>	CAT -> T <u>A</u> T	H845Y
	<i>tn1438</i>	CGA -> T <u>G</u> A	R486*
	<i>tn1444</i>	CAG -> T <u>A</u> G	Q1128*
	<i>tn1467</i>	CAG -> T <u>A</u> G	Q45*
<i>spr-5</i>	<i>tn1378</i>	TGG -> T <u>A</u> G	W666*
	<i>tn1379</i>	GAG -> <u>A</u> AG	E164K
	<i>tn1394</i>	GGT -> <u>G</u> AT	G619D
<i>uev-1</i>	<i>tn1381</i>	GGA-> <u>G</u> AA	G17E
	<i>tn1382</i>	GGT -> <u>G</u> AT	G47D
<i>spr-2</i>	<i>tn1380</i>	GAT-> <u>G</u> CT	D42A
	<i>tn1436</i>	Tc1 insertion	

^aPremature stop codons are indicated by asterisks. The numbering of amino acids refers to the isoforms PDE-6A, TOM-1A, SPR-4A, and SPR-2B.