

# Mutations in *Serac1* or *Synj2* cause proximal *t* haplotype-mediated male mouse sterility but not transmission ratio distortion

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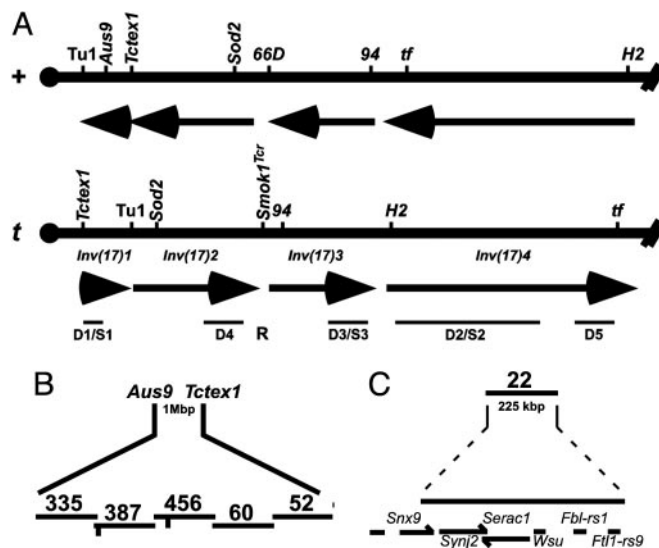
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Transmission ratio distortion (TRD) and sterility are male-specific quantitative trait phenomena associated with the mouse *t* haplotype. TRD occurs in *t* haplotype-heterozygous males and is caused by the deleterious action of distorter products on sperm bearing a wild-type responder locus. It has been proposed that *t*-mediated male sterility is a severe manifestation of TRD caused by homozygosity for distorter loci; thus, the distorter and sterility loci would be identical. In this study a transgenic approach was used to identify the proximal sterility locus, *tcs1* (S1), and test its role in TRD. Mice transgenic for a wild-type bacterial artificial chromosome (BAC) derived from the S1-critical region were bred onto *t* haplotype backgrounds. Mating results conclusively showed that the BAC is sufficient to restore fertility in otherwise sterile males. Multiple mutations were identified in the *t* alleles of *Synj2* and *Serac1*, two genes in the BAC; thus, they are candidates for S1. In addition, whereas the BAC transgene rescued sterility, it had no effect on TRD. These results uncouple the proximal *t* haplotype sterility locus, S1, from TRD, demonstrating that S1 and the proximal distorter locus, D1, are not the same gene.

synaptojanin 2 | *t* complex

The *t* complex spans  $\approx 20$  cM of mouse chromosome 17 and exists in two forms: a variant known as the *t* haplotype (*t*) present in 20% of the feral house mouse population, and the wild-type form (+) (Fig. 1). The *t* haplotype differs from + by the presence of four nonoverlapping inversions (1), *Inv(17)1-Inv(17)4*, that effectively suppress recombination between *t* and +; thus, the *t* haplotype is inherited *en bloc*. Two male-specific phenomena occur in *t* haplotype mice: transmission ratio distortion (TRD) and sterility (reviewed in ref. 2). TRD, or non-Mendelian segregation, occurs in *t/+* males and is characterized by the transmission of the *t* haplotype to up to 99% of offspring (reviewed in ref. 3). Several recessive lethal and semilethal *t* alleles of multiple loci affecting embryonic development have been discovered. These alleles are subdivided into complementation groups such that mice homozygous for two complementing *t* haplotypes are viable, yet the males are invariably sterile because of homozygosity of sterility loci. Females are equally affected by the lethal loci but do not exhibit TRD or sterility.

The current model proposes that in *t/+* males, *t* distorters expressed during spermatogenesis interact deleteriously with the + allele of the cis-acting responder (*Ter*; R); thus, R<sup>+</sup> sperm are incapacitated. However, R<sup>t</sup> is refractory to distorter effects, and R<sup>t</sup> sperm are normal and capable of fertilization (4). Distorter and sterility factors are genetically inseparable, leading to the proposal that distortion and sterility are caused by the same mutations, and the end effect is distorter-dosage-dependent. Accordingly, a double dose of distorters overcomes the refractory nature of R<sup>t</sup> and causes sterility in viable *t/t* males (5). Evidence exists for three distorter/sterility loci (6–8): *Distorter 1/Sterility 1* (D1/S1), D2/S2, and D3/S3; and two distortion-only loci: D4 and D5 (Fig. 1). Only R<sup>t</sup> has been cloned and shown to encode a mutant kinase (SMOK1<sup>Ter</sup>), consisting of the C terminus of SMOK1, a sperm motility kinase, fused to the N terminus of RSK6, a ribosomal kinase (9).



**Fig. 1.** The *t* complex and the S1-critical region. (A) Structure and organization of the *t* complex. The complex exists as two variants, + and *t* haplotype, distinguished by four nonoverlapping inversions (arrows). Markers defining the inversions are shown. TRD and sterility loci are shown below the *t* haplotype (S1/D1, etc.) (B) BAC tile of the S1-critical region. The RPCI-23 BACs are 335F22 (335), 387F2 (387), 456P8 (456), 60J11 (60), 52A7 (52), and 22M9 (22). The structure and organization of 335, 387, 456, 60, and 52 are published elsewhere (13). BAC 22M9 is 225 kb and overlaps 387 and 456 (tick marks). (C) The following are in 22M9. Genes: *Snx9* (3' half only), *Synj2*, *Serac1*, and *D17Wsu155e*. Pseudogenes: *Fibrillarlin*-like and *Ft1-rs9*. The transcriptional orientations are shown.

Distortion and sterility effects attributable to D1/S1 are caused by amorphic mutations because a deletion of the wild-type genomic region harboring D1/S1 recapitulates TRD and sterility when in trans to distal partial *t* haplotypes or complete *t* haplotypes, respectively (10). Various combinations and dosages of S1, D2/S2, and D3/S3 result in sterility or reduced fertility. Whereas homozygosity of S1 alone does not cause infertility, mice homozygous for S1 and heterozygous for D2/S2 (abbreviated as S1 S2/S1 +) are sterile (5). A similar dosage effect has been observed for TRD (8). For example, males heterozygous for partial *t* haplotypes typically transmit the *t* haplotype chromosome at a lower frequency than do

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Abbreviations: TRD, transmission ratio distortion; BAC, bacterial artificial chromosome; Tg, transgenic; OFM, offspring per female per month; SERAC1, serine active site containing 1; IP<sub>3</sub>, inositol-3,4,5-trisphosphate; ADAM2,  $\beta$ -fertilin; INPP5, inositol polyphosphate 5-phosphatase.

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**Table 1. Analysis of S1 candidate genes within BAC RPCI-23-22M9**

Gene symbol	Function	Human homolog	Mutation		Coordinates*	RT-PCR
<b>RPCI-23-22M9</b>			DNA	Protein	3110767–3368420	
<i>Synj2</i>	Phosphoinositide 5-phosphatase	<i>SYNJ2</i>	A <sup>562</sup> →G	N94D	3216322–3318403	Y
			C <sup>1436</sup> →T	S385F		
			A <sup>1603</sup> →G	NC		
			C <sup>3884</sup> →T	NC		
			A <sup>3905</sup> →C	Q1208P		
			A <sup>4016</sup> →G	3' UTR		
<i>Serac1</i>	Unknown	<i>SERAC1</i>	C <sup>74</sup> →A, C <sup>98</sup> →A	5' UTR	3180874–3216545	Y
			T <sup>112</sup> →C	V4A		
			T <sup>118</sup> →A, T <sup>119</sup> →C	F6Y		
			T <sup>121</sup> →G	F7C		
			T <sup>141</sup> →C	NC		
			C <sup>185</sup> →G	T25M		
			A <sup>301</sup> →C, C <sup>302</sup> →A	T64Q		
			T <sup>1215</sup> →C	NC		
			A <sup>1574</sup> →G	N488S		
			T <sup>1866</sup> →C	NC		
			T <sup>1976</sup> →G	NC		
			G <sup>1994</sup> →A	3' UTR		
<i>D17Wsu155e</i>	Unknown	BC060317			3175122–3180761	Y
<i>Fibrillarin</i>	Pseudogene				3138435–3139391	N
<i>Ftl2</i>	Pseudogene				3127986–3128906	N

Mutations are numbered according to the following: *Serac1*, NM\_177311 (DNA) and NP\_796285 (protein); *Synj2*, NM\_011523 (DNA) and NP\_035653 (protein). NC, no change. Mutations in bold are correlated (see text).

\*Coordinates are derived from Build 30 (Ensembl).

important in sperm function, as demonstrated by the sterility observed in males homozygous for a *Inpp5b* null allele (24); thus, we proceeded to generate BAC Tgs to test the hypothesis that S1 is encoded by the *t* allele of *Synj2*.

Shortly afterward, the sequence from RPCI-23 BACs 387F2, 456P8, and 60J11 became available, closing the S1 critical region (Fig. 1). GENSCAN (20) analysis of the BAC sequences predicted a gene distal to *Synj2* which has been named *Serac1*. Further analysis confirmed that *Serac1* is contained within 22M9.

The function of serine active site containing 1 (SERAC1) is unknown. INTERPRO analysis of the ORF predicts an esterase/lipase/thioesterase domain (accession no. IPR000379) and an ARM repeat fold (accession no. IPR008938). RT-PCR analysis confirmed that the gene is expressed in the testis (Fig. 2). A comparison of wild-type and *t* alleles revealed a cluster of *t*-specific mutations (Table 1), located within the first 25 codons of the *Serac1* ORF, and several other mutations dispersed throughout the transcript. Several mutations are nonsynonymous and introduce non-conservative amino acid replacements (Phe7Cys and Thr25Met), whereas others are less drastic (Val4Ala, Phe6Tyr, and Asn488Ser). One mutation in particular, Thr64Gln, represents one of two alleles found in wild-type strains, because this site is Gln in *t* and BALB/cByJ, but is Thr in (NMRI/GSF × CD1) F1 mice (GenBank accession no. NM\_011523) (30). In addition, several mutations were found within UTRs.

*Synj2* and *Serac1* overlap tail to tail such that the same genomic segment encodes a fragment of the ORFs of both genes (Fig. 3). At the mRNA level, nucleotides 1847–2034 of *Serac1* are the complements of nucleotides 3848–4035 of *Synj2*, and a subset of the mutations in *Serac1* correlates with mutations in *Synj2*: mutations 1866, 1976, and 1994 in *Serac1* correspond to mutations 4016, 3905, and 3884, respectively, in *Synj2* (Table 1).

Mapping of BAC ends onto the mouse assembly (Ensembl Build 31) using global survey sequences (National Center for Biotechnology Information accession IDs AZ083735 and AZ083736) identified three more genes: *Fibrillarin*-like, *Ftl-rs9*, and *D17Wsu155e*. *Fibrillarin*-like and *Ftl-rs9* are inactive pseudogenes, whereas *D17Wsu155e* is a small ORF encoding a hypothetical 71-aa protein of unknown function that is 38% identical to the 78-aa

protein Rex1-S of *Chlamydomonas reinhardtii*, required in DNA repair (31). Testis expression of *D17Wsu155e* was confirmed by RT-PCR (data not shown), and a comparison of the genomic sequence of the *t* and wild-type alleles yielded no differences that would lead to amino acid changes, thus *D17Wsu155e* was excluded as an S1 candidate.

**BAC Effect on Fertility Rescue and TRD Experiments.** To test the hypotheses that S1 is encoded in 22M9, and that D1 and S1 are distinct genetic elements, we generated male mice with various combinations of complete and partial *t* haplotypes with or without the BAC. Fecundity and transmission ratios were tested with the following results.

**Sterility Rescue Test.** Three assays were performed. The first assay tested *t*<sup>H49</sup>/*t*<sup>w2</sup> males, the second assay tested 5J males, and the third assay tested *t*<sup>w82</sup>/*t*<sup>w2</sup> males. In each case, sibling males with or without the Tg were tested, and in all three assays, 22M9 provided strong evidence of sterility rescue.



**Fig. 3.** *Serac1* and *Synj2* overlap. Mutated nucleotides (small boxes; see Table 1), and coding-coding overlap (large boxes) are shown. Translation stop sites (all caps) are shown. The transcriptional orientations are indicated below the alignment.

**Table 2. Results of fertility assay by using Tg and non-Tg males**

Genotype	Mating test results					
	Fertile	Semisterile	Sterile	OFM range	OFM average	SD
$t^{w2}/Aus9^{df5};Tg18/+$	5	2	5	0–8.66	2.52	2.88
$t^{w2}/Aus9^{df5};Tg26/+$	1	1	2	0–12.40	3.62	5.94
$t^{w2}/Aus9^{df5};+/+$	0	1	14	0–1.09	0.08	0.28
$t^{w2}/t^{w82};Tg18/+$	9	0	0	4.71–11.54	7.81	2.38
$t^{w2}/t^{w82};Tg26/+$	2	0	0	—	10.0	0.86
$t^{w2}/t^{w82};+/+$	2	1	3	0–6.05	2.12	2.65
$t^{w2}/t^{h49};Tg18/+$	9	1	1	0–11.33	7.72	3.78
$t^{w2}/t^{h49};Tg26/+$	3	0	0	3.88–8.09	6.64	2.40
$t^{w2}/t^{h49};+/+$	1	0	11	0–0.18	0.02	0.06

**$t^{h49}/t^{w2}$ .** The first test used a partial *t* haplotype,  $t^{h49}$ , in trans to  $t^{w2}$ , a complete *t* haplotype.  $t^{h49}$  harbors S1 but not the stronger sterility locus, S2 (4). Thus, the genotype of non-Tg males is (S1+ +/S1, S3, S2) and they are expected to be sterile. Control and experimental males were mated with two females each for several months. The number of offspring sired by each male was normalized (5) and compared with the fecundity of non-Tg control male siblings. The fertility of Tg males was remarkably similar to wild-type mice (*Tg18*: 7.72 OFM and *Tg26*: 6.64 OFM), whereas the non-Tg sibling controls were infertile (0.02 OFM), with the exception of one male (Table 2 and Fig. 4). The fecundity of the fertile non-Tg  $t^{h49}/t^{w2}$  male was indistinguishable from wild-type males and is likely due to complex genetic background effects or an undetected change in  $t^{h49}$  or  $t^{w2}$ . These results unequivocally demonstrate that 22M9 has a significant effect on fertility.

**5J/ $t^{w2}$ .** The second test used 5J, a deletion that spans the S1/D1 critical region, in trans to  $t^{w2}$ ; therefore, non-Tg males are (null+ +/S1, S3, S2) and considered homozygous for the S1 amorphic mutation, whereas BAC Tgs would be (+ + +/S1, S3, S2) if the wild-type allele of S1 were in 22M9. Twelve 5J/ $t^{w2};Tg18/+$  and 4 5J/ $t^{w2};Tg26/+$  males were tested, and their fecundity was compared with that of 15 non-Tg siblings. Non-Tg male fecundity was ex-

tremely low (0.08 OFM), whereas Tg male fecundity was significantly increased (*Tg18*: 2.52 OFM and *Tg26*: 3.62 OFM). These results are consistent with the first assay and confirm that 22M9 is capable of rescuing S1 infertile males. However, a minority of Tg males remained sterile, a phenomenon that may be due to the health effects of harboring a large deletion and not to factors affecting fertility *per se* (11).

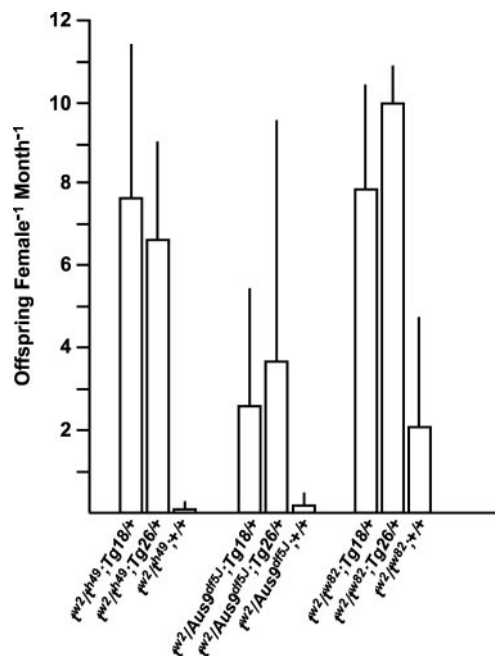
**$t^{w82}/t^{w2}$ .** The last assay used male mice heterozygous for a partial proximal *t* haplotype ( $t^{w82}$ ) harboring S1 but not the stronger sterility locus, S2, and a complete *t* haplotype ( $t^{w2}$ ). Accordingly, non-Tg males are (S1+ +/S1, S3, S2) and are expected to be sterile. Interestingly, non-Tg males were semisterile (2.12 OFM); however, the Tg counterparts were considerably more fecund (*Tg18*: 7.72 OFM and *Tg26*: 6.64), which is consistent with the previous assays.  $t^{w82}$  is derived from  $t^{w32}$ , a complete *t* haplotype known to exhibit variability in fertility assays, sometimes resulting in unexpected fertility in carrier males (10). Although it is possible that a hypomorphic  $t^{w82}$  allele of S1 with residual wild-type function caused the semisterility of non-Tg males, it is more likely that background genetic modifiers affected the results because none of the mice were inbred.

In conclusion, the results of all three assays demonstrate that 22M9 rescues S1-induced male sterility. Of a total of 41 Tg males tested, 33 were fertile (average OFM = 6.4) whereas only 5 of 33 non-Tg control siblings exhibited any fertility (average OFM = 0.74; Table 2).

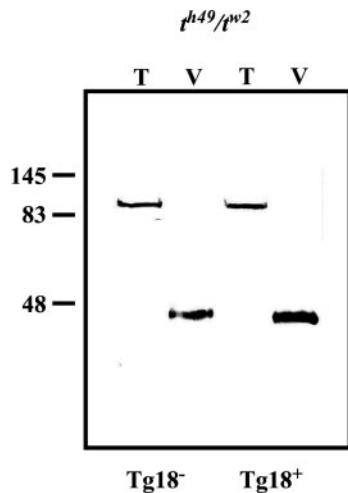
**TRD Test.** ES cell-derived deletions in chromosome 17 segregate into two groups when introduced in male mice: (i) deletions that cause sterility when in trans to complete *t* haplotypes, or TRD when in trans to partial *t* haplotypes; and (ii) deletions that cause sterility when in trans to complete *t* haplotypes but not TRD when in trans to partial *t* haplotypes (11). If D1 and S1 are the same, then  $t^{w2}$  in the presence of the Tg is functionally equivalent to a long distal partial *t* haplotype (+ R D3, D2) such as  $t^6$  (4), because 22M9 would supply the D1/S1 wild-type allele.

$t^6/+$  males transmit  $t^6$  to  $\approx 65\%$  of their offspring on average (4, 6, 10, 11), which is significantly lower than the 95% observed for  $t^{w2}$  (16, 32). Therefore, we tested the transmission of  $t^{w2}$  from  $t^{w2}/+$  males with or without *Tg18*. If the wild-type allele of D1 is in 22M9, we would observe a significant decrease in the transmission of  $t^{w2}$  from  $t^{w2}/+;Tg18/+$  males. The results clearly showed that *Tg18* had no effect on  $t^{w2}$  because  $t^{w2}/+;Tg18/+$  males transmitted  $t^{w2}$  to 94% of offspring compared with 97% transmission by  $t^{w2}/+$  males ( $\chi^2 = 0.10$ ,  $P = 0.75$ ).

**Western Blot Analysis of ADAM2.** Male mice homozygous for the *Inpp5b* knock-out are sterile and exhibit defects in the proteolytic maturation of ADAM2 (24). ADAM2 and ADAM1 form a heterodimer necessary for sperm–egg binding (33) and maturation of ADAM2, which occurs in the epididymis, is affected by inositol



**Fig. 4.** Results of Tg rescue analysis. The results are expressed as a mean of all experiments for each cluster of genotypes (described in *Materials and Methods*) plus the SE of the mean, for each experiment.



**Fig. 5.** Western Blot analysis of ADAM2. The  $t^{h49}/t^{w2}$  males (described in *Materials and Methods*) are sterile in the absence of the transgene ( $Tg18^-$ ), and fertile in its presence ( $Tg18^+$ ). ADAM2 was derived from testis sperm (T) or vas deferens sperm (V). Molecular mass markers (kDa) are shown.

metabolism (24). Although the maturation protease is unknown, it is thought to be a serine protease (34).

The prediction that SERAC1 contains a serine active site characteristic of serine proteases, prompted us to test ADAM2 maturation in fertile Tg animals and their non-Tg siblings. The results show that ADAM2 is processed normally in rescued and sterile males (Fig. 5). Consequently, the  $t$  alleles of *Serac1* and *Synj2* do not affect ADAM2 maturation.

## Discussion

Genetic studies have uncovered five distorter (D1-D5) and three sterility (S1-S3) loci, although the genes remain unknown (2, 3). D1/S1, D2/S2, and D3/S3 comprise distorter/sterility loci; it is hypothesized that each is encoded by a single-mutant allele. However, the results of our Tg rescue analysis, and deletion (12) and high-resolution mapping studies (11), do not support this hypothesis for D1/S1; instead, we propose that D1 and S1 are different mutations.

TRD is caused by a deleterious interaction between distorters and SMOK1, resulting in the inactivation of + gametes. SMOK1<sup>Tr</sup>, however, is resistant to distorter action (3). The trans-acting inactivation effect of distorters may be facilitated by diffusion across cytoplasmic bridges connecting synchronously developing sperm (35), whereby they gain access to + gametes. Alternatively, distorter gene expression before meiosis may distribute distorters throughout meiotic partners where they could affect SMOK1. SMOK1 however, is postmeiotically expressed and restricted to the gamete of origin, despite apparently being soluble (9). The resistance of SMOK1<sup>Tr</sup> to harmful distorter effects may be due to multiple mutations and reduced kinase activity (9), although the chemistry of distorter/SMOK1 interactions is unknown.

To determine whether BAC 22M9 ameliorates TRD by providing a wild-type copy of D1, we bred it onto the  $t^{w2}$  background. If 22M9 harbors wild-type D1, Tg  $t^{w2}$  males should have the same levels of TRD as  $t^6$  males because  $t^6$  is a long distal characterized by (+ D3 and D2) alleles. However, the expected decrease of  $t^{w2}$  transmission was not observed. Instead Tg males transmitted  $t^{w2}$  at a rate equal to non-Tg  $t^{w2}$  siblings ( $\approx 95\%$ ). The persistence of a mutant D1 locus in Tg  $t^{w2}$  males used for TRD analysis was deemed unimportant, because D1 is amorphic and its effects are elicited by deleting the genomic interval that harbors it (10). Thus, D1 encodes a functionless product, or the product is not expressed. We conclude that the wild-type allele of D1 is not encoded in 22M9.

$t$  haplotype sterility is thought to be the consequence of a double dose of distorters, referred to as sterility factors in this context (5). In  $t/+$  males, SMOK1<sup>Tr</sup> counteracts the effect of the distorters (4) but this resistance is absent in  $t/t$  males; thus, the type of responder makes no difference on the outcome (36), and  $R^+/R^+$ ,  $R'/R^+$ , or  $R'/R'$  males are sterile provided that at a minimum, the male is homozygous for S2. If the male is heterozygous for S2, then it must be homozygous for S1 to be infertile (5).

Male sterility can be induced by deleting, in + chromosomes, the genetic interval harboring the S1 locus (10–12), indicating that S1 is an amorphic mutation whose lack of function has deleterious consequences on fertility provided one S2 allele is present. In fact, the fecundity of males carrying S1 only is indistinguishable from wild type (10), suggesting that the + allele of S1 is redundant, except when its absence is compounded by heterozygosity for S2.

We bred 22M9 onto various backgrounds that otherwise produce sterile or semisterile males, expecting that if 22M9 harbors the + allele of S1, Tg mice would be heterozygous for the S1 mutation and exhibit normal fertility. With one exception, this is what we observed (Fig. 4). The exception was when 22M9 was bred onto the  $5J/t^{w2}$  background; these males are sterile in the absence of the BAC, whereas their Tg counterparts were mildly fecund, averaging three OFM (Table 2). The diminished rescue effect is likely due to adverse health effects brought on by the large size of the  $5J$  deletion, which spans several cM. Taken together, our results demonstrate that 22M9 rescues S1-mediated male sterility, which is in stark contrast to its lack of effect on TRD. Therefore, different genes encode S1 and D1.

Analysis of 22M9 uncovered *Synj2*, *Serac1*, and *D17Wsu155e*, and several pseudogenes. *D17Wsu155e* was ruled out because the  $t$  allele sequence and expression profile is indistinguishable from +. The *Synj2* and *Serac1*  $t$  alleles, however, harbor multiple mutations, making them strong candidates for S1 (Table 1). Interestingly, the  $t$  alleles do not qualitatively differ in expression relative to the + alleles, insofar as both genes are expressed in sterile males at levels similar to wild-type and rescued Tg males.

SYNJ2 functions in the inositol-signaling pathway. It has two catalytic motifs: an N-terminal *SacI*-like domain that dephosphorylates positions 3 and 4 of phosphatidylinositol, and a central domain that dephosphorylates position 5 of lipidic and soluble inositol, including inositol-3,4,5-trisphosphate (IP<sub>3</sub>) (27, 28). SYNJ2 functions in clathrin-mediated endocytosis (37) and has been localized to various cellular compartments, including the outer mitochondrial membrane (38), the plasma membrane, where it functions as a Rac1 effector (39), and the spermatid manchette (29); its function in spermatogenesis or sperm function is unknown. However, regulation of the inositol signaling pathway is important in spermatogenesis and sperm function as seen by the effect of knocking out *Inpp5b* (24, 40), and by the expression of *Inpp5d* (SHIP) in round spermatids (41). Both genes encode proteins with substrate specificity similar to SYNJ2.

SYNJ2 mutations fall in three categories. N94D and S385F are in the *SacI* domain. Interestingly, residue 94 is D in rat and human SYNJ2, suggesting that the mutation originated in wild-type mice after the divergence of  $t$  and + lineages. S385F is in a highly conserved site in mouse, rat, and human SYNJ2, and is drastic. S385 is in the vicinity of R381, a critical Arg residue that when mutated to His, abolishes *SacI*-phosphatase activity (29). The same effect is observed in the yeast homolog of SAC1 (42). Thus, we suspect that S385F may adversely affect enzyme activity by disrupting the three-dimensional structure in the vicinity of R381.

The C-terminal mutation Q1208P, is in a unique region of SYNJ2 arising from an alternative splicing event not observed in human or rat; therefore, this residue does not have an equivalent in human or rat SYNJ2. The last category is characterized by mutations within the 3' UTR, none of which appear to affect the splicing or expression of the gene.

If *Synj2* is S1, we assume that its phosphatase activity is lost in S1/S1 homozygotes because deletion of the genomic interval harboring it mimics the S1 effect, but only in the presence of the D2/S2 mutation (10–12). The lost activity could be compensated by other testis-expressed phosphatases, including inositol polyphosphate 5-phosphatase (INPP5B) and INPP5D. An interaction between S1 and S2 is suggested because at least one S2 *t* allele is necessary to elicit sterility in S1/S1 homozygotes (10). Perhaps S2 encodes a general regulatory protein that enhances phosphatase activity in SYNJ2, INPP5B, and INPP5D. In the absence of SYNJ2, up-regulation of other phosphatases by the wild-type product of S2 could be sufficient to restore normal function. However, a single S2 *t* allele would create a conditional haploinsufficiency, characterized by up-regulation of phosphatase activity below a critical threshold, thus severely compromising all sperm. Regulation of the inositol pathway by modulating IP<sub>3</sub> levels is critical to sperm function. IP<sub>3</sub> is required for mobilizing intracellular Ca<sup>2+</sup>, an action that affects sperm hypermotility (43) and the acrosomal reaction (44), required at critical steps during fertilization. S2 mutations might affect this regulatory network, allowing IP<sub>3</sub> accumulation and misregulation of calcium release. Although levels of IP<sub>3</sub> or phosphatase activity in *t/t* males have not been measured relative to *+/+* males, this hypothesis is consistent with the fact that S2 homozygotes are sterile in the absence of other mutant *t* sterility loci (5). The region harboring S2 is several cM, making candidate gene analysis impractical. Interestingly, there are reports of abnormal hypermotility and flagellar morphology in *t/t* males as a consequence of improper calcium regulation (45, 46). In addition, the differential expression in developing sperm of calmodulin and other cytoplasmic Ca<sup>2+</sup>-binding proteins (47), and the localization of IP<sub>3</sub> receptors to various sperm organelles, including the acrosome (48), demonstrates that spermatogenesis is affected by Ca<sup>2+</sup> regulation. Our hypothesis predicts the presence of abnormally high levels of IP<sub>3</sub> in *t/t* spermatozoa.

*Serac1* is ubiquitously expressed and its expression in *t* haplotypes is not qualitatively different from wild type. Its function remains unknown, yet it appears to belong to the family of serine active-site-containing proteins characteristic of many hydrolases, including serine proteases. Interestingly, in the *Inpp5b* knock-out males, defects in ADAM2 maturation were observed (24). ADAM2 is essential for normal sperm function (49), and its maturation

depends on an as-of-yet identified protease, although it has been suggested to be a serine protease (34). Consequently, we tested the maturation of ADAM2 in sterile *t* haplotypes and their rescued counterparts and observed normal processing in both, thus ruling out SERAC1 as the ADAM2 maturation protease (Fig. 5).

The tail-to-tail overlap in *Serac1* and *Synj2* (Fig. 3) suggests that they might be coevolving to a limited extent, because silent mutations in one could be harmful to the other. Interestingly, this overlap is not observed in the human, chimp, or rat orthologs (Ensembl). A recent analysis of overlapping genes within the mouse genome found that 31 of 578 overlapping genes include so-called coding-coding overlap such as that observed with *Serac1* and *Synj2* (50). A similar analysis of the human genome did not find an instance in which coding-coding overlap in one organism is conserved in the other (50), thus substantiating our finding that the *Serac1-Synj2* overlap is unique to mouse. What effect, if any, this overlap has on the expression of these genes is unknown. However, if the genes have the same spatial-temporal expression profiles, the possibility exists that the transcripts associate by means of complementarity and this in turn could affect their expression profiles and/or stability.

We conclude that S1 is encoded either by *Synj2* or *Serac1*. Further studies requiring a knock-out approach could unravel which of the two genes is S1. If the *t* allele of *Synj2* encodes S1, it raises the interesting question of why are multiple inositol-metabolizing enzymes, with overlapping substrate specificity, required for proper sperm function? The answer may lie in nonoverlapping subcellular localization or in temporally regulated differences in enzyme activity. Alternatively, if *Serac1* encodes S1, further work will be required to dissect the role of SERAC1 in spermatogenesis, beginning with identifying its function.

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