torso-like encodes the localized determinant of Drosophila terminal pattern formation

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Differentiation of the anterior and posterior poles of the Drosophila embryo requires seven maternally expressed genes including torso-like (tsl) and torso (tor). The tor gene encodes a receptor tyrosine kinase that is expressed throughout the embryo but is activated specifically at the poles. Genetic mosaic analysis has shown that tsl is required during oogenesis in follicle cells at each end of the oocyte. We cloned the tsl locus and showed that it was expressed specifically in follicle cells at the anterior and posterior ends of the oocyte. tsl encodes a novel protein with a putative amino-terminal signal sequence. Ectopic expression of tsl produced embryos with a phenotype similar to that resulting from constitutively active Tor alleles. These results suggest that localized TSL controls the localized activation of TOR.

[Key Words: torso-like gene; Drosophila embryo; pattern formation; oogenesis; ectopic expression]

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Anterior-posterior pattern formation in Drosophila depends on spatially localized gene products deposited in the egg during oogenesis (for review, see St. Johnston and Nüsslein-Volhard 1992). Three distinct determinant systems are required for specification of the anterior-posterior axis, and they are referred to as the anterior, posterior, and terminal systems. The anterior and posterior systems determine the segmented regions of the head, the thorax and the abdomen and rely on localization of specific mRNAs to the anterior and posterior ends of the embryo, respectively. Proper development of the nonsegmented head and tail (terminal) regions of the Drosophila embryo depends on the action of seven maternal effect genes: torso [tor (Schupbach and Wieschaus 1986)]; torso-like [tsl (Stevens et al. 1990); fs(1) Nasrat [fs(1)N] (Degelmann et al. 1986)]; fs(1)polehole [fs(1)ph] (Perri- mon et al. 1986)]; trunk [trk (Schupbach and Wieschaus 1986)]; corkscrew [csw (Perkins et al. 1992)]; and l(1)polehole [l(1)ph] (Perrimon et al. 1985)]. Recessive mutations in any of these genes give rise to the same phenotype, loss of the anterior portion of the head, and of all structures posterior to and including the eighth abdominal segment.

A key step in terminal patterning is the localized activation of the receptor tyrosine kinase (RTK) encoded by the tor locus (Sprenger et al. 1989). This receptor is found uniformly distributed throughout the plasma membrane of the oocyte and early embryo (Casanova and Struhl 1989); however, activation of the receptor is normally restricted to the extreme anterior and posterior ends. Tor gain-of-function mutations cause the production of constitutively active receptor leading to a disruption of segmentation in the trunk region of the embryo and ectopic production of terminal structures (Klingler et al. 1988; St. Johnston and Nüsslein-Volhard 1993). Epistasis analysis with Tor gain-of-function mutations has revealed that l(1)ph (also known as D-raf) and csw act downstream of the receptor, whereas the other maternal effect loci act before activation of the receptor (Ambrosio et al. 1989; Perkins et al. 1992).

An important question that remains is how localized activation of the TOR receptor is achieved. The tsl gene is a particularly good candidate for encoding a localized signal leading to activation of the TOR receptor. The requirement for tsl+ function has been shown by mosaic analysis to be restricted to a few follicle cells at each end of the egg chamber during oogenesis (Stevens et al. 1990), whereas the other maternal effect genes are required in the germ line (Perrimon and Gans 1983; Schupbach and Wieschaus 1986; Perkins et al. 1992). tsl could encode an extracellular protein, either the ligand for the TOR receptor or some product involved in the production of an active TOR ligand. Alternatively, tsl could encode a regulatory protein controlling the expression of such a signal.

In this paper we describe the isolation and characterization of the tsl gene. We report that expression of tsl is normally restricted to a subset of follicle cells at each end of the egg chamber. Ectopic expression of tsl caused patterning defects similar to those of dominant tor alleles, indicating that the localized expression of tsl is critical for proper terminal patterning. Sequence analysis revealed an amino-terminal signal sequence, suggesting that the TSL protein is a secreted product.
Results

Identification of a P-element insertion allele of tsl

In studying the migration and function of the specialized follicle cells known as the border cells (King 1970; Montell et al. 1992), we screened 7800 single P-element enhancer trap insertions on the second and third chromosomes (Karpen and Spradling 1992) for their β-galactosidase staining patterns in the ovary and for lethal and female sterile phenotypes (Montell et al. 1992). One line displayed a weakly fertile phenotype and specific β-galactosidase expression in the border cells and a group of posterior follicle cells (Fig. 1). Approximately 90% of eggs laid by homozygous PZ0617 females failed to hatch into larvae.

The single P-element insertion in this line was mapped to 93F on the polytene chromosomes by in situ hybridization (data not shown), the same location as the tsl locus (Stevens et al. 1990). Because of this chromosomal location and the female sterile phenotype, we tested females heterozygous for the P-element insertion and a point mutation in the tsl locus for complementation and found that they were also weakly fertile. Based on this complementation data and the excision data described below, we refer to the P-element insertion as a new allele of tsl, ts\textsuperscript{0617} (Table 1).

To generate stronger alleles and to test whether the apparent tsl mutation was attributable to insertion of the P-element, we remobilized the P-element using a standard excision protocol (Cohen et al. 1992). The weakly fertile phenotype completely reverted to wild-type in 6/16 independent ros\textsuperscript{y} lines, indicating that the mutation was attributable to the P-element insertion and not to a second mutation elsewhere on the chromosome. In addition, one excision line, ts\textsuperscript{0617e8}, was completely sterile both when homozygous and when heterozygous with a ts\textsuperscript{l} point mutation (Table 1). Furthermore, the cuticularized embryos all displayed a typical "terminal" phenotype (Fig. 2), including head and tail deletions. Taken together, these data indicated that ts\textsuperscript{0617} was a hypomorphic allele of the tsl locus.

Identification of the tsl gene

DNA flanking the ts\textsuperscript{0617} element was cloned, and the site of P-element insertion determined [see Materials and methods]. A transcript map for the region was generated by using subfragments of the λ phage clone to probe Northern blots containing poly(A)\textsuperscript{+} RNA from ovaries and embryos (Fig. 3B). Probes from one side of the P-element hybridized to a 7-kb RNA present in embryos but not in ovaries, whereas probes from the other side of the P-element hybridized to a 1.9-kb RNA present in both ovaries and embryos (Fig. 3C). Furthermore, the 1.9-kb mRNA appeared to be reduced in concentration in ovaries from ts\textsuperscript{0617} mutant females (Fig. 3B). Because the

Table 1. Complementation analysis of tsl alleles

| Allele   | ts\textsuperscript{1691} | ts\textsuperscript{0617} | ts\textsuperscript{0617e8} | ts\textsuperscript{0617e1} |
|----------|--------------------------|--------------------------|---------------------------|---------------------------|
| Female   | sterile                  | weakly fertile           | weakly fertile            | fertile                   |
| sterile  | female                   | female                   | female                    | N.T.                      |
| tsl\textsuperscript{0617} |                                  |                          |                           |                           |
| tsl\textsuperscript{0617e8} |                                  |                          |                           |                           |
| tsl\textsuperscript{0617e1} |                                  |                          |                           |                           |

[N.T.] not tested. ts\textsuperscript{0617e1} was representative of six lines that reverted to wild type.

Figure 1. β-Galactosidase expression from the enhancer trap.
Stage 9 egg chamber from a PZ0617 female, stained for β-galactosidase activity. Migrating border cells (bc, arrowhead) and posterior follicle cells (arrowhead) stain specifically. Nurse cells (nc), oocyte (o), and columnar follicle cells (fc) are indicated. Anterior is to the left.

Figure 2. Terminal class phenotype of ts\textsuperscript{0617e8} excision allele. Dark-field micrographs of cuticle preparations of embryos produced by wild-type [a] and heterozygous ts\textsuperscript{0617e8}/ts\textsuperscript{0617} [b] showing deletions of terminal structures. Eight ventral denticle belts, a well-formed head skeleton, and posterior spiracles can be seen in a. In b the head skeleton is reduced, only 6 ventral denticle belts are clearly distinguished, and posterior spiracles are missing.
tsl gene is required in ovaries, we focused on the 1.9-kb transcript as the best candidate for tsl.

To investigate the spatial pattern of expression of the 1.9-kb mRNA, several cDNA clones were isolated. Single-stranded, digoxygenin-labeled probes were made and used for in situ hybridization to ovaries. The result is shown in Figure 4. Specific hybridization to border cells and posterior follicle cells was seen with one strand only, in a pattern closely resembling the β-galactosidase expression from tslP0617 (Fig. 1) and closely resembling that predicted for the tsl gene based on mosaic analysis (Stevens et al. 1990). Expression was first detected quite early in oogenesis, at stage 3 (for staging, see King 1970), in just two follicle cells at each end of the egg chamber (not shown). And at later stages, a larger number of anterior follicle cells, namely the centripetal follicle cells, were also labeled (not shown).

To test for tsl function, the 10-kb EcoRI genomic DNA fragment, which truncates the neighboring embryonic transcript but includes all transcribed sequences from the tsl gene, was cloned into a P-element transformation vector. This DNA was injected into embryos, and six independent germ-line transformants were obtained. The presence of a single copy of the transgene was sufficient to rescue the strongest tsl allele to fertility, providing strong evidence that the 1.9-kb transcript corresponded to tsl.

The nucleotide and deduced amino acid sequences of a putative full-length tsl cDNA are shown in Figure 5. Conceptual translation revealed a single long open reading frame of 353 amino acids with good Drosophila codon usage (not shown). Amino acids 1–22 were found to have a high probability of forming a signal sequence as predicted by the algorithm of von Heijne (1986). All of the features of a signal sequence were present, including a positively charged amino acid following the methionine, a hydrophobic sequence long enough to span the bilayer, followed by a more polar sequence and a poten-
have been reported previously {Stevens et al. 1990}. They revealed two introns, which are indicated on the transcript map.

Five ethylmethane sulfonate (EMS)-induced tsl alleles have been reported previously [Stevens et al. 1990]. They have been ordered in an allelic series of decreasing severity such that 174 = 691 > 146 > 035 > 135. To determine the lesions in these mutant alleles, we amplified genomic DNA corresponding to the tsl-coding region from each mutant, using the PCR, and sequenced the products directly. A single nucleotide change causing an amino acid substitution was found in each case as indicated in Figure 5. The sequence from nucleotides -230 to 1 is genomic sequence, and the site of P-element insertion is indicated by the open inverted triangle. The sequence from nucleotide 1 (arrow) to 1857 is cDNA sequence. The positions of the two introns are indicated by solid inverted triangles. The putative signal sequence is underlined (solid line), as are the two leucine-rich regions [broken lines]. Nucleotide and amino acid (boldface) numbers are indicated at right.

Figure 6. Schematic diagram of the deduced TSL protein. Amino acid substitutions found in the five EMS-induced mutations are indicated (for details, see text).
an asparagine, and the mutated tryosines occurred within similar sequences [YSYVLVRVV and YEYYVKVY]. No other nucleotide differences were observed except in ts169, where a silent T-C was observed in codon 248, which encodes a proline. The molecular lesion in the tsP17 excision line, which had a stronger phenotype than the original tsP17 insertion [see Table 1], was determined by probing genomic Southern blots with probes from each end of the P-element [data not shown]. We found that only internal P-element sequences and no flanking DNA had been deleted.

Ectopic expression of tsl

If the normal pattern of tsl expression were crucial for providing spatially restricted activation of the TOR receptor, uniform tsl expression would be expected to cause uniform TOR activation and a phenotype similar to that observed in mutants with constitutive activation of the TOR receptor. Because the heat shock promoter drives uniformly high levels of expression in all follicle cells [Xu and Rubin 1993], one of the putative full-length cDNAs was cloned into a P-element transformation vector behind the hsp70 heat-inducible promoter. This DNA was injected into embryos, and eight independent germ-line transformants were obtained. We found that when a single 1-hr heat shock was administered to females bearing the hs-tsl construct, 50–60% of the cuticularized embryos that developed displayed a phenotype similar to that of the dominant Tor allele TorRL3 [Fig. 7]. The characteristics of the TorRL3 phenotype are that abdominal segmentation is suppressed and/or disrupted, ectopic filzkörper material and/or mouth parts are occasionally produced, and the embryos are about half the length of wild-type embryos. Embryos from females carrying the hs-tsl construct were short and had suppressed and disrupted abdominal segmentation; however, we did not observe ectopic terminal structures. These phenotypes were dependent on the hs-tsl construct and were not artifacts of the heat shock treatment because we did not observe such defects in embryos derived from w1118 females that had been treated similarly.

Discussion

An early event in terminal patterning is the localized activation of the TOR RTK at the two poles of the blastoderm. The genes acting downstream of tor include Ras1 [Lu et al. 1993], the Raf serine–threonine kinase [Nishida et al. 1988; Ambrosio et al. 1989; Sprenger et al. 1993], and a tyrosine phosphatase encoded by the csw locus [Perkins et al. 1992]. This signal transduction cascade ultimately results in the spatially restricted expression of transcription factors encoded by the tailless and huckebein loci. A key question that remains in understanding terminal pattern formation is how the TOR receptor becomes activated only at the two poles. Immunocytochemistry, as well as functional studies, has shown that the receptor is present throughout the plasma membrane surrounding the embryo [Casanova and Struhl 1989] yet activated only at the two ends [Casanova and Struhl 1993; Sprenger and Nüsslein-Volhard 1993]. Our results indicate that tsl expression is restricted to follicle cells at each end of the oocyte during oogenesis, that the product is likely to be a secreted protein, and that ectopic tsl expression is sufficient to cause ubiquitous TOR activation.

Evidence that the 1.9-kb transcript corresponds to tsl

Several lines of evidence indicate that the 1.9-kb transcript that we identified was the product of the tsl locus. First, we rescued the mutant phenotype by germ-line transformation. Second, the restricted spatial expression pattern of this transcript correlated well with that predicted from mosaic analysis [see below]. Third, we have detected changes in the coding region in each of five EMS-induced tsl alleles and a decrease in the level of mRNA in the weaker, P-element-induced allele. Finally, we obtained the predicted gain-of-function phenotype by expressing the cDNA ectopically.

Spatially restricted tsl expression is critical for terminal patterning

Tsl is the only maternal effect, terminal class gene identified to date with a function that is required in the somatic follicle cells and not in the germ line [St. Johnston and Nüsslein-Volhard 1992]. Furthermore, mosaic analysis demonstrated that tsl function is required only in follicle cells at the anterior and posterior poles of the oocyte, rather than in all of the follicle cells [Stevens et al. 1990]. On the basis of the expression of the lacZ reporter gene in the tsl enhancer trap allele and in situ

Figure 7. Ectopic expression of tsl. Dark-field micrographs of cuticle preparations from TorRL3 gain-of-function allele [a], hs-tsl [b], and w1118 heat-shocked control [c]. Note the similarity between a and b: Embryos are shorter than wild type, and abdominal segmentation is severely disrupted, however, the termini remain relatively normal.
hybridization of the cDNA to ovaries, we found tsl expression to be restricted to follicle cells at the anterior and posterior poles of the oocyte. Thus, the expression pattern and genetic requirement corresponded closely.

We also showed that ectopic expression of tsl in all follicle cells was sufficient to cause the disruption of abdominal segmentation that is characteristic of Tor gain-of-function mutations. We concluded that ectopic tsl expression was sufficient to cause ubiquitous TOR activation, suggesting that all of the other components required for TOR activation must be present all around the embryo. Therefore TSL is the only localized signal in terminal pattern formation prior to TOR activation.

The TSL product

The most striking feature of the TSL-deduced amino acid sequence was the presence of a putative amino-terminal signal sequence. Because tsl function is required in the follicle cells, it has been proposed previously that tsl might encode a secreted protein deposited into the perivitelline space, or a transcription factor or some other type of regulatory protein involved in the production of a secreted signal (Stevens et al. 1990; St. Johnston and Nüsslein-Volhard 1992). Our results indicate that tsl does not encode a transcriptional or post-transcriptional regulatory protein. Rather, the presence of a putative signal sequence strongly supports the model that the TSL product is secreted into the vitelline membrane or perivitelline space and, therefore, that TSL participates directly in the signaling pathway.

Although sequence analysis did not reveal significant homologies with proteins of known biochemical function, two regions of leucine-rich sequence with some similarity to the leucine-rich repeats found in Toll, chaptin, and connectin (Nose et al. 1992) were observed in the TSL sequence. The precise function of such repeats has not been characterized; however, they have been implicated in mediating protein–protein and protein–lipid interactions (Krantz et al. 1991).

All of the EMS-induced alleles were characterized by single amino acid substitutions; thus, it is possible that even the strongest alleles are not null mutations. Because all of the tsl alleles were recovered in screens for female sterile mutants, it is possible that the null phenotype is actually lethality. Consistent with this idea, we have recovered lethal excision lines from the tsl^P0617 P-element insertion (C. Andrews and D. Montell, unpubl.), however, it is not yet clear that the lethality is attributable to loss of tsl function. The observations that two of the stronger mutations resulted in replacement of tyrosine residues by asparagine residues and that the mutated tyrosines occurred within somewhat similar sequences suggest that these two motifs serve similar functions. It was surprising to find that the relatively conservative replacement of alanine by valine in tsl^P091 results in a strong female sterile phenotype. However an alanine-to-valine substitution in the superoxide dismutase gene has recently been characterized, which also leads to a strong mutant phenotype (Deng et al. 1993).

The role of tsl in terminal patterning

Whereas tsl appears to encode the localized signal for TOR activation, a remaining question is whether TSL is the TOR ligand or whether TSL somehow converts a uniformly distributed, inactive ligand precursor into an active form. The active TOR ligand has been demonstrated to have three important characteristics: It is present in limiting amount; it is spatially localized; and it is freely diffusible (Casanova and Struhl 1993; Sprenger and Nüsslein-Volhard 1993). Although TSL is very likely to be spatially localized based on the expression pattern of the mRNA, TSL is unlikely to be freely difusible. This is because TSL is made during oogenesis and must be kept from diffusing for hours (or even days if females are holding their eggs), until after fertilization when the ligand acts (Sprenger and Nüsslein-Volhard 1993).

Three additional genes, namely trk, fs(1)N, and fs(1)ph, are required for normal activation of TORSO. fs(1)ph and fs(1)N are candidates for genes with products that might be involved in anchoring TSL to the vitelline membrane. Females homozygous for most alleles of these loci produce eggs that collapse, indicating a role for these gene products in maintaining eggshell integrity (Degelmann et al. 1990). trk mutants, on the other hand, produce embryos with only terminal patterning defects. It has been suggested that the TRK protein may be a secreted, inactive ligand for TOR (Casanova and Struhl 1993). If so, this would suggest that TSL is more likely to be involved in converting inactive TRK into active TRK. This model could explain why ectopic terminal structures were not observed in hs–tsl embryos. Ectopic expression of TSL, while producing ectopic active TRK and disrupted abdominal segmentation, might not produce as high a level of TOR activity as mutations in TOR itself, if TRK were the product that was limiting in amount.

Materials and methods

Fly stocks and isolation of P-element tsl allele

EMS-induced tsl mutant alleles were gifts from Dr. L. Stevens (Albert Einstein University, New York). The tsl^P0617 allele was obtained in an enhancer trap screen (Karpen and Spradling 1992) using the PlacZ element described in Mlodzik et al. (1990).

Cloning

DNA flanking the tsl^P0617 P-element was cloned by making a library in AZAP (Stratagene), following complete digestion of tsl^P0617 genomic DNA with EcoRI, and screening with a P-element-specific probe. Flanking sequence (3 kb) was recovered and used as a probe to isolate a λ phage clone from a wild-type genomic library. The site of P-element insertion was determined by comparing the restriction map of the 3-kb flanking DNA with that of the wild-type phage clone and by genomic Southern blotting using P-element and flanking DNA probes on blots containing DNA from tsl^P0617 and wild-type flies. The site was subsequently confirmed by DNA sequencing. Embryonic and ovarian cDNA libraries (gifts of S. Hawley (University of California, Davis) and A. Spradling (Carnegie Institute of Wash-
from the ovarian library; however, one clone was obtained from the embryonic library. This clone was later determined by sequencing to be a genomic DNA fragment spanning 1.6 kb of transcribed sequence, including the 330-bp intron within the coding sequence. This clone was used to isolate several cDNAs from a 9- to 12-hr embryonic library (a generous gift from K. Zinn, California Institute of Technology, Pasadena), including several that were 1.9 kb in length.

**β-Galactosidase staining and in situ hybridization in ovaries**

Staining for β-galactosidase activity was carried out as described (Montell et al. 1992). For in situ hybridizations, egg chambers were dissected in Ringer’s solution and fixed in 4% paraformaldehyde (Polysciences, E.M. grade) in 1 × PBS (PP) for 20 min. Following three washes in 1 × PBS with 0.1% Tween 20 (Sigma, PBT) for 5 min each, egg chambers were digested with 100 μg/ml of proteinase K (Boehringer Mannheim) in PBT for 1 hr at room temperature, rinsed once with 0.2% glycine in PBT, and refixed for 20 min in PP; they were then washed three times for 20 min each in PBT alone. Egg chambers were then treated with 90% methanol/10% 0.2 M EGTA for 1 hr at −20°C, washed with PBT three times for 20 min each, and equilibrated in hybridization buffer (50% formamide, 5 × SSC, 100 μg/ml of salmon sperm DNA, 50 μg/ml of heparin, 50 μg/ml of cRNA, 0.1% Tween 20). Egg chambers were prehybridized in the same solution at 42°C before the probe was added and incubated overnight. Single-stranded, digoxigenin-labeled probes were made by asymmetric PCR. The PCR reaction contained 0.1 μl of plasmid DNA, 0.05 μl of KCl, 0.01 μl of Tris-HCl (pH 8.3), 1.5 mm MgCl₂, 0.001% gelatin, 0.2 mM each dATP, dCTP, and dGTP, and 0.13 mM dTTP, 0.075 mM digoxigenin-11-dUTP (Boehringer Mannheim), 150 ng KS or SK primer, and 2 units of AmpliTaq polymerase (Cetus Corporation). Forty microliters of mineral oil was added and the reaction was subjected to 35 cycles of the following temperature paradigm: 95°C for 45 min, 55°C for 30 min, 72°C for 1 min and 30 sec. The product was ethanol precipitated and resuspended in hybridization buffer, boiled for 1 hr to reduce the size of the probe, chilled, and added to egg chambers. Subsequent washing and developing of the reaction was carried out as described (Tautz and Pfeifle 1989).

**Plasmid construction and germ-line transformation**

The 1.9-kb *tsl* cDNA insert was amplified from phage DNA using PCR [Hgt11 primers: 5’-AGCCAGCGCGCTACGGTGCATAATC-3’ and 5’-GGAGGCGCGCTACGATCGCCGAAAT-TC-3’] and cloned into the EcoRI site of pBluescript (SK+, Stragtagene). The hs–tsl construct was made by subcloning the same 1.9-kb EcoRI fragment into pCaSpeR–hs [a gift from C. Thummel, University of Utah, Salt Lake City). The rescue construct was made by subcloning the same 1.9-kb EcoRI genomic fragment containing the *tsl* gene into pCaSpeR–1 [a gift from C. Thummel] to create pCaSpeR–*Gtsl*. Plasmids pCaSpeR–*Gtsl* (0.4 mg/ml) and pCaSpeR–*Gtsl* (0.4 mg/ml) were microinjected into w¹¹⁸ embryos, along with the helper plasmid pB25.2wc (0.1 mg/ml), following a standard P-element mediated germ-line transformation protocol (Rubin and Spradling 1982; Grigliatti et al. 1986). In subsequent generations progeny expressing the w¹ gene were selected as transformants. Insertions were mapped to a chromosome by segregation analysis, balanced, and underwent homozygosis when possible. To test for rescue, a pCaSpeR–*Gtsl* insertion on the X chromosome was crossed into a ts¹⁶⁹¹ mutant background. Females homozygous for ts¹⁶⁹¹ and heterozygous for pCaSpeR–*Gtsl* were tested for fertility.

**Heat shock treatment and analysis of hs–tsl**

Healthy 4- to 5-day-old adults were heat-shocked for 1 hr at 36°C. The flies were allowed to recover for 10 hr at 25°C. Embryos were collected from 10 to 22 hr after heat shock and aged for 30 hr at 25°C. For cuticle preparations embryos were dechorionated, and transferred to a drop of 50% lactic acid:50% Hoyer’s solution (Grigliatti et al. 1986) on a microscope slide, coverslipped, and baked at 65°C overnight.

**RNA preparations and Northern blot analysis**

Total RNA from embryos was prepared by homogenization in 6 M guanidine hydrochloride in 0.1 M sodium acetate (pH 5.5), followed by centrifugation through a 5.7 M cesium chloride cushion in 0.1 M sodium acetate (pH 5.5), in a swinging bucket ultracentrifuge rotor at 25,000 rpm for 18 hr at 20°C. Total RNA from ovaries was isolated by dissecting out ovaries from fat females and homogenizing them in extraction buffer [50 mM Tris (pH 7.5), 10 mM EDTA, 100 mM NaCl, 0.5% SDS] containing 400 μg/ml of proteinase K. The RNA was extracted with phenol/chloroform, followed by chloroform, and precipitated with 0.15 M NaCl. Poly(A)* RNA was isolated from the total embryo and ovary RNA, electrophoresed in a denaturing 1% agarose–formaldehyde gel, and transferred to GeneScreen Plus (NEN-DuPont) membrane. Membranes were hybridized in 50% formamide, 6× SSPE, 5× Denhardt’s solution, and 0.1% SDS at 42°C overnight. The blots were washed in 0.1% SDS, 0.1% SSC, at 65°C for 2 hr.

**DNA sequencing**

DNA sequence was obtained using the Sequenase II kit (U.S. Biochemical). The 1.9-kb cDNA was sequenced on both strands using both dGTP and dITP to resolve compressions. Genomic DNA was sequenced using oligonucleotide primers derived from the cDNA sequence. The second intron was sequenced completely, but only the intron/exon boundaries were sequenced from the larger first intron. Most but not all genomic sequence was obtained from both strands. One nucleotide difference was observed between the cDNA and genomic sequences. Nucleotide 790 was found to be a C rather than a T in the genomic sequence, causing codon 84 to encode a proline rather than a leucine. The genomic sequence was confirmed in each of the five *tsl* strains sequenced. Therefore, it is likely that the cDNA sequence resulted from an error in reverse transcription.

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Note added in proof

The EMBL accession number for the sequence in Figure 5 is X75614.

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