Localization of the *fushi tarazu* Protein during Drosophila Embryogenesis

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Summary

The *fushi tarazu* (*ftz*) gene of *Drosophila* acts early in embryogenesis to regulate body segmentation. The localization of the *ftz* protein product in embryos was examined using indirect immunofluorescence microscopy. Antibodies were prepared against a β-galactosidase-*ftz* hybrid protein made in *E. coli*. The *ftz* protein was first detectable in blastoderm-stage embryos as seven stripes of nuclei encircling the embryos transversely. The stripes persist through the early events of gastrulation, but disappear before overt segmentation is visible. The *ftz* protein is expressed a second time in some nuclei of the developing nervous system. In contrast to the early pattern, at the later stage, *ftz* is expressed in each of fifteen metameric subunits of the embryo.

Introduction

The establishment of the segmental body pattern in *Drosophila* requires the coordinated function of three classes of zygotically active genes early in development (Nüsslein-Volhard and Wieschaus, 1980; Nüsslein-Volhard et al., 1984; Jürgens et al., 1984; Wieschaus et al., 1984). These three classes have been called gap, polarity, and pair-rule loci based on morphological defects that occur in the absence of wild-type gene activity. It is thought that information incorporated into the oocyte by maternally encoded protein product, the *ftz* protein was produced as a hybrid protein in *E. coli*. Antibodies specific to the *ftz* portion of this fusion protein were purified. Immunofluorescence analysis of fixed whole-mount embryos was used to study the cellular localization of *ftz* gene product, and the time and site of its expression. We observe that *ftz* protein is located in nuclei in a striped pattern, that not all of the *ftz*-expressing blastoderm stripes are of equal width, that the stripes become narrower during gastrulation, and that *ftz* protein is also detected in the developing ventral nervous system in a pattern quite different from its earlier distribution in the embryo.

Results

Expression of the *ftz* Protein in *E. coli*

For the generation of antibodies directed against the *ftz* protein, a gene fusion was constructed in *λgt11* to produce the protein in *E. coli* (Young and Davis, 1983). To express *ftz* as a hybrid protein, a 1228 bp Ava II fragment from the pDmG20 *ftz* cDNA (Laughon and Scott, 1984), with appropriate Eco RI linkers ligated to it, was introduced into the Eco RI site of *λgt11* (Figure 1). The phage (*λftzA10*) produces a hybrid protein containing 399 amino acids of *ftz* protein and lacking only the five N-terminal and nine C-terminal amino acids of the deduced *ftz* protein (Laughon and Scott, 1984).

Lysogens containing the *λftzA10* phage were induced to express the hybrid β-galactosidase-*ftz* hybrid protein. Upon thermal induction a fusion protein migrating with an apparent molecular weight of 175 kd was produced (Figure 2A). The protein migrates more slowly than expected from the known sequence (160 kd is predicted). The high proline content (10%) of *ftz* may be responsible for the anomalous migration of the hybrid protein. The hybrid protein was indeed a hybrid; it reacted more strongly with a slightly faster migrating band, presumably a degradation product of the full length fusion protein, perhaps caused by protein conformation effects.

Antibodies against *ftz* Protein

Antiserum for use in immunofluorescence studies was elicited by immunization with *ftz*-βgal fusion protein purified using amino-phenyl thiogalactopyranoside affinity chromatography (see Experimental Procedures). The immune serum was affinity-purified using hybrid protein cross-linked to anti-β-galactosidase antibody attached to Sepharose 4B. The specificity of the final antibody prepa-
clear division, ftz protein was detected within nuclei in a pattern of seven stripes, each about four nuclei wide in the cellular blastoderm stage, some 30 min after the last anti-ffz antibodies (Figures 3c and 3d). However, by the localized immunofluorescent labeling was detected with was observed at any developmental stage examined.  

Expression of ftz during Embryogenesis  

After fertilization, the Drosophila embryo undergoes a series of nearly synchronous nuclear divisions (Foe and Alberts, 1983). At about the eighth nuclear division the nuclei begin to migrate to the periphery of the embryo, a number of anti-ftz labeled bands were presumably because of the presence of breakdown or prematurely terminated products of the hybrid protein. To localize ftz antigen during embryogenesis, whole fixed embryos were stained with anti-ftz antibodies according to the method of Mitchison and Sedat (1983).

Figure 1. Construction of lacZ-ftz Gene Fusion in λgt11  

An Ava II subclone of the ftz pDmG20 cDNA (Laughon and Scott, 1984) was inserted into pUC8 at the unique Eco RI site using 10 bp (CGGAAATTCCG) Eco RI linkers ligated to the 1228 bp coding fragment of ftz. H-BOX, the 160 bp homeobox sequence. The subcloned, linked fragment was then transferred, in the proper orientation, into the unique Eco RI site of the λgt11 expression vector (see Experimental Procedures).

Figure 2. Expression of a β-galactosidase-ftz Fusion Protein in E. coli and Production of Specific Antibodies to the ftz-Encoded Domain  

Lysogens were grown for 2 hr at either 30°C (uninduced) or 42°C (induced), and cell lysates were prepared for electrophoresis on a 7.5% SDS-polyacrylamide gel. Part of the gel was stained with Coomassie blue (A), and replicate lanes were transferred to nitrocellulose and were probed with affinity-purified antibodies against: β-galactosidase (B); an 11 amino acid peptide of the homeodomain (C); or the ftz portion of the hybrid protein (D). The molecular weight markers indicated in A are myosin (200 kd), β-galactosidase (116 kd), phosphorylase b (97 kd), bovine serum albumin (66 kd), and ovalbumin (45 kd). The arrows indicate the position of the full-length hybrid protein (migrating as 175 kd). Note that a small amount of hybrid protein is present in uninduced cultures because of the absence of repression in strains without lac I. Many antigenic degradation products are present in induced cells. U, uninduced; I, induced; M, markers.

Expression for the ftz-encoded portion of the fusion protein was demonstrated using protein blot (Figure 2). The affinity-purified antibody reacted with ftz fusion protein but not with β-galactosidase or other E. coli proteins. The larger number of anti-ftz labeled bands were presumably because of the presence of breakdown or prematurely terminated products of the hybrid protein. To localize ftz antigen during embryogenesis, whole fixed embryos were stained with anti-ftz antibodies according to the method of Mitchison and Sedat (1983).
ies. The most anterior stripe is located just posterior to the cephalic furrow, although the stripe and the furrow do not precisely align at all times (Figure 5a). As gastrulation proceeds, the stripes become wedge-shaped, about two to three nuclei wide on the dorsal side, and about four nuclei wide ventrally (Figures 5a and 5d). The posterior stripe remains the widest and is also somewhat wedge-shaped. These changes appear to be due to rearrangements of cells, although we cannot rigorously eliminate the possibility that a different subset of the cells begins to produce ftz antigen. After formation of the furrows, the germ band, a ventral strip of ectodermal cells in which segments will first become visible, begins to form and to extend around the posterior end of the embryo. The lengthening of the germ band is thought to be due largely to cell movement rather than to cell division (Sander, 1976; Hartenstein and Campos-Ortega, 1985).

The indentation that will become the posterior midgut invagination begins to form posteriorly to the posterior-most stripe (the flattened posterior end in Figure 5b), while the first signs of the posterior transverse fold (pf in Figure 5b) appear dorsally at the location of the stripe fifth from the anterior. The stripes can still be seen dorsally and ventrally as the pole cells enter the posterior midgut invagination (Figures 5c and 5e). Parts of the fourth and fifth stripes disappear into the posterior transverse fold. Eventually part of the sixth stripe is drawn into the fold as well, at a time after the pole cells have completely disappeared into the posterior midgut invagination (Figure 5a). The posterior end of the embryo moves dorsally and then anteriorly during the germ-band elongation phase of development (Poulson, 1950; Turner and Mahowald, 1977). During elongation, the ftz stripes can be followed until nearly the time of full germ-band extension (not shown). As the germ band extends, ftz expression decreases in the dorso-lateral cells while remaining present in the germ band. The stripes disappear before the germ band is fully extended.

While the germ band is at its maximum extension, a dramatic change in the pattern of staining occurs. Concomi-

Figure 3. Control and Early Staining Patterns
Antibodies against β-galactosidase were affinity-purified from serum from the same rabbit from which the anti-ftz antibodies were prepared. (a) The immunofluorescent staining of an embryo in the early stages of gastrulation, stained (as in Experimental Procedures) with anti-β-galactosidase. (b) The same embryo as in a, stained with the DNA-marking dye, DAPI.
(c) Embryos later than the thirteenth nuclear division stained with anti-ftz antibodies. (d) Same embryo as in c, stained with DAPI.
Figure 4. Blastoderm and Early Gastrulation Stage Embryos

In each micrograph, the anterior end of the embryo is at the top. (a, b, and c) Whole-mount embryos incubated with anti-ftz antibodies and secondarily stained with fluorescein-conjugated goat anti-rabbit antibody. (d, e, and f) The same three embryos as in a, b, and c (respectively) stained with DAPI.

(a and d) An embryo at the cellular blastoderm stage, just beginning gastrulation movements. The large internal bright spots in d are the yolk nuclei. One yolk nucleus is indicated (yn).

(b and e) An embryo (ventral view) that has begun gastrulation. The ventral furrow (vf) through which the mesodermal cells move into the interior and the cephalic furrow (cf) are visible. By this stage, the stripes of stained nuclei have narrowed to an average three-cell width, except for the most posterior stripe, rather than the initial four-cell width seen in a.

(c and f) A dorsal view of a gastrulating embryo. The immunofluorescent stripes (in c) can be seen to encircle the embryo. As cells move from the dorsal and lateral regions toward the ventral side, the stripes take on a wedge-shaped appearance.

The appearance of the ectodermal invaginations that form tracheal pits, detectable antigen appears in fifteen clusters of germ-band nuclei within the dividing cells of the developing nervous system (Pouloon, 1960, Lohmann et al., 1983). From a lateral view, it can be seen that the initial staining is in one cell layer (not shown). From a ventral or dorsal perspective, the earliest staining is of nuclei arranged in a chain of linked hexagons (Figure 6b). The laterally extended vertices of the hexagons are just posterior to the tracheal pits and are also in the approximate positions where the clusters of nuclei staining with anti-ftz subsequently coalesce. The clusters (Figure 6c) remain visible during germ band shortening (Figure 6d). The staining is repeated bilaterally in each of fifteen segmental units, including the regions where the gnathocephalic segments are forming (Figure 6e and Figure 7a).

To assess whether this unexpected spatial and temporal pattern of anti-ftz staining is caused by the bona fide gene...
Figure 5. Embryos during Gastrulation

(a) A dorsolateral view of an embryo showing the anterior stripe of stained nuclei just posterior to the cephalic furrow (cf). The distorted shape of the nuclei along the ventral (v) side is associated with the movement of the cells in that region into the ventral furrow.

(b) A lateral view of an embryo in which the posterior transverse fold (pf) is beginning to form. The indentation at the posterior (bottom) end is the beginning of the posterior midgut invagination. The cephalic furrow (cf) is still visible. The ventral furrow is visible by both immunofluorescent staining and by the DAPI staining (in c), along the ventral (v) side.

(c) A dorsolateral view of an embryo that is beginning germ-band elongation. Most of the posterior endoderm has moved into the posterior midgut invagination, leaving the double-width stripe of ftz protein near the posterior end. The cephalic furrow (cf) and posterior transverse fold (pf) are visible. The pole cells (pc) are in the process of entering the posterior midgut invagination in this embryo. Some of the cells around that invagination are distorted in shape as they bend into the opening. Most of the cells that contain detectable ftz protein are not distorted in this way (compare c and e).

We examined homozygous mutant ftz\textsuperscript{woo} embryos for staining. The ftz\textsuperscript{woo} allele has a 5 kb transposon inserted into the protein-coding sequence of the gene (Weiner et al., 1984). Staining of the neural cells was not detectable at any stage of mutant embryo development (example in Figures 7c and 7d). Therefore, the neural staining is indicative of either authentic ftz protein product or a cross-reactive protein that requires ftz\textsuperscript{+} function for its expression. The fifteen-unit staining pattern disappears shortly after germ-band shortening is completed (Figure 7b), and no staining with the anti-ftz serum is seen at later embryonic stages (data not shown).
Figure 6. Protein Patterns Later in Embryogenesis
(a) A dorsal view of the posterior end of an embryo. The pole cells have moved into the interior. The fourth, fifth, and sixth stripes have merged in the posterior transverse fold. (b) A ventral view of an embryo with an elongated germ band, stained with anti-\(ftz\). The tracheal pits (tp) are visible. The first detectable antigen is arranged in linked hexagons. The antigen is in the nuclei. (c) Lateral view of an embryo with an elongated germ band, stained with anti-\(ftz\). The clypeolabrum (cl) marks the anterior end of the embryo. Fifteen units of staining are visible. (d) Same as c, but during germ-band shortening. (e) Ventral view of an embryo near the end of germ-band retraction. The clypeolabrum (cl) is at the anterior end of the embryo. Antigen is visible in all of the subdivisions of the germ band and is located in patches flanking the ventral midline. (f) DAPI-stained view of the embryo shown in e.

Discussion

The \(ftz\) Protein Accumulates in Embryonic Nuclei

The molecular functions of proteins encoded by segmentation genes and homeotic genes are unknown. Based on structural homology between part of the homeodomain and bacterial DNA-binding proteins, it has been hypothesized that one function of the family of homeodomain-containing proteins is to bind to DNA (Laughon and Scott, 1984). The nuclear location of the \(ftz\) protein is consistent with this hypothesis. Two other Drosophila homeodomain-containing proteins have been localized in both embryonic and imaginal disc nuclei, the Ultrabithorax protein (White and Wilcox, 1984; Beachy et al., 1985), and the engrailed protein (Di Nardo et al., 1985). Other genes containing homeoboxes include the homeotic genes Antennapedia, Sex combs reduced, Deformed, Infraabdominal-2, and Infraabdominal-7 (Scott and Weiner, 1984;
The arrangement of segmental precursor cells

The analysis of whole-mount embryos with ftz protein-specific antibody allows the detection of ftz protein up to the 10 hr stage of embryogenesis; ftz mRNA has been detected only up to about the four and a half hr stage (Hafen et al., 1984). When the ftz protein is first detected, the stripes of immunofluorescent labeling are about four nuclei wide, in the anterior-posterior axis. A precise number cannot be determined because the nuclei are arranged (approximately) in a hexagonally close-packed array. Also, at the edges of each stripe, some nuclei appear to stain weakly. At the edges of the embryo where the signal is strongest because of vertical stacking of stained nuclei (Figures 4a and 4b), four stained nuclei appear to alternate with four unstained nuclei.

As gastrulation begins, the ftz stripes become narrower, averaging three nuclei across instead of four. The rapid disappearance of ftz protein from some of the blastoderm nuclei suggests that a specific mechanism for elimination of the protein may exist. Later still, the stripes become wedge-shaped, averaging two nuclei wide dorsally and four nuclei wide ventrally. The most posterior stripe appears slightly wider than the other stripes at the blastoderm stage (Figure 4a) and does not become distinctly narrower during early gastrulation movements (Figure 5). As a result, during gastrulation the posteriormost stripe remains about twice as wide as the others.

The identity and number of segments at the posterior end of Drosophila is a matter of some uncertainty. Most primitive insects have eleven abdominal segments plus a terminal segment, while Drosophila have only eight obvious segments. The double-width posterior ftz stripe may be due to the evolutionary disappearance of one or more segments that in more primitive insects would divide the

Figure 7. Protein Patterns after Germ-Band Shortening and in ftz- Embryos

(a) Ventrolateral view of embryo stained with anti-ftz at the end of germ-band shortening. The light patch seen at the right (dorsally) is in the region of the remaining yolk and is not due to immunofluorescence. (b) Ventrolateral view of an embryo at a slightly later stage than the embryo in a. The ventral staining with anti-ftz has disappeared. (c and d) Immunofluorescent (c) and DAPI (d) staining of an embryo after germ-band shortening. The embryo has about half the usual number of body divisions because it is homozygous for a ftz null mutation (ftz^{null}). No immunofluorescent staining of embryos of this genotype (recognized by their diminished segment number) is observed. The bars in d indicate the remaining body divisions.
double-width ftz stripe. Perhaps the "lost" segments are the eighth, tenth, and eleventh, and the seventh and ninth grouped. One repeating unit would be segmental, units of may not correspond to segmental primordia. If the segments remain, fused together.

Another phase would give parasegments (Martinez-Arias an anterior compartment plus a posterior compartment. derm stage (Lohs-Schardin et al., 1979). The ftz results One argument supporting a nonsegmental unit as the cor-

there are at least four "phases" in which the cells can be blastoderm-stage repeating units are four cells wide, then the difference between a segment and a parasegment is only a frameshift of one cell wide (Kornberg et al., 1985; DiNardo (Laughon and Scott, 1984) which causes a homeotic transformation of the posterior compartment of metathorax (T3) into mesothorax (T2). Only if ftz+ is expressed in this phase of ftz expression comes from the ftzR@ allele

It has been estimated that the precursor cells for each segment lie in a band three to five cells wide at the blasto-
derm stage. In ftz- embryos, the segment boundaries be-
tween MX and LB, T1 and T2, T3 and A1, A2 and A3, and so on, fail to form. Therefore, we have indicated two frames that straddle the seg-
ment boundary and are consistent with the pattern elements deleted when ftz+ is absent.

Figure 6. Diagram of a Model for the Blastoderm-Stage Segmental Primordial Cells

The blastoderm cells are represented in a lateral view as repeating segmental primordia, each four cells wide in the anterior-posterior axis. Only part of the embryo is represented. The cells that will form posterior compartment structures are shown as filled circles; cells that will form anterior compartment structures are shown as open circles. The segments (maxillary, labial, first, second, and third thoracic, and first, second, and third abdominal) and some of the parasegments (lower brackets; PS 2, 4, 6, and 8) are indicated. The two rows of brackets indicate two possible frames in which the ftz+ protein might be expressed when it first appears in four cell wide stripes at the cellular blastoderm stage. In ftz- embryos, the segment boundaries be-

Between MX and LB, T1 and T2, T3 and A1, A2 and A3, and so on, fail to form. Therefore, we have indicated two frames that straddle the seg-
ment boundary and are consistent with the pattern elements deleted when ftz+ is absent.

ftz and the Ventral Nervous System

One of the unexpected results of the in situ hybridization of homeotic gene DNA probes to the RNA of sectioned embryos was the high concentration of transcripts in the nervous system. Both Ultrabithorax (Akam, 1983) and Antennapedia (Levine et al., 1983) transcripts are located in specific parts of the ventral ganglia. In contrast to these homeotic genes, segmentation genes such as ftz and engrailed have not been reported to have major accumulations of transcripts in neural cells (Hafen et al., 1984; Kornberg et al., 1985). It was therefore a surprise to find a nuclear protein in parts of the nervous system that is rec-
ognized by the anti-ftz serum (Figure 6).

There are several possible explanations for the neural staining. First, the staining may be due to authentic ftz protein. The ftz RNA may be at a low level in the nervous system, and may have been missed. Some recent in situ hybridization analyses of sectioned embryos with ftz probes have detected transcripts in neural tissue in a pat-
tern similar to that reported here for ftz protein (M. Levine, personal communication). Northern blot analysis of staged embryonic RNA indicated that ftz mRNA is detect-
able up to 10–12 hr of development, although at a lower level than its expression at cellular blastoderm (Weiner et al., 1984). The absence of the neuroblast staining in ftz· embryos is consistent with the signal being due to ftz-encoded protein, but another possibility is that the antiserum cross-reacts with one or more proteins the expression of which requires ftz· function. A third possibility is that only part of the observed staining is due to authentic ftz protein, for example, that found in alternate segments.

One likely source of cross-reactivity would be the 60 amino acid ftz homeodomain structure that is shared with several other proteins. However, the antiserum against ftz contains only a very low level of reactivity with other homeodomain-containing proteins, as assayed by protein blotting (Carroll and Scott, unpublished observations). Furthermore, the transient post-specific signal observed with anti-ftz antibody is not consistent with the previously described patterns of homeotic gene transcription in the nervous system (Akam, 1983; Levine et al., 1983; McGinnis et al., 1984). Also, an antiserum made against a synthetic peptide representative of part of the homeodomain reacts with nuclei from most or all parts of the embryo as well as all nuclei in the nervous system (Carroll et al., 1985). The faint staining of all nuclei seen with the anti-ftz serum might be accounted for by homeodomain cross-reactivity. Experiments to identify on blots the proteins detected by the ftz antibodies have not yet succeeded convincingly, and we do not know the number of forms of the native ftz protein or of any cross-reactive polypeptides.

The expression of ftz in the nervous system raises the question of a late function for the product. In an analysis of the temperature-sensitive period of a ftz ts allele (Wakimoto et al., 1984), the major period of sensitivity was from 2 to 4 hr of development, at the blastoderm stage. However, in addition, some lethality was observed if mutant embryos were shifted to the restrictive temperature as late as 6 or 8 hr into development. These data provide a hint that ftz may function later than blastoderm stage. It is also possible that lack of ftz function in the nervous system does not cause much lethality, or that prior function of ftz in neuroblasts at the early stages of gastrulation partially protects embryos from a subsequent loss of ftz function. Ciona analysis has shown that no ftz function is required for differentiation of adult cuticular structures later than 12 hr of development (Wakimoto et al., 1994). ftz mRNA is not detectable after 12 hr of development (Weiner et al., 1984). The distinct temporal and spatial patterns of ftz expression in the ectoderm and neural tissues suggests that the regulation of the ftz gene expression is more complex than was previously thought. The identification of those genes influencing ftz expression at each stage of development and the elucidation of ftz function at the cellular and molecular level remain the most pressing issues for eventually understanding the role of this pair-rule gene in Drosophila embryonic pattern formation.

Experimental Procedures

Gene Fusion

The ftz cDNA clone pDmG20 has been described previously (Weiner et al., 1984; Laughon and Scott, 1984). A 1228 bp Ava II fragment containing nearly all of the coding region was purified, its recessed 3' termini filled in with the Klenow fragment of E. coli DNA polymerase, and Eco RI 10-mer linkers (New England Biolabs) that had been phosphorylated with T4 polynucleotide kinase were ligated to the now blunt-ended Ava II fragment. After ligation, the DNA was digested with Eco RI, and the fragment was purified. To amplify the fragment and remove tandem linkers, it was cloned into pUC8, the purified plasmid DNA was digested with Eco RI, and the 1.2 kb Eco RI fragment was purified on a preparative gel. The fragment was then ligated into Eco RI-digested λgt11 (Young and Davis, 1983) at the unique site near the 3' end of the lacZ gene. The phage were packaged, plated on RY1000, and scored for colorless plaques. The DNA from plaque-purified phage was prepared and screened for insertion of the ftz fragment in the proper orientation.

Expression of Hybrid Proteins

Lysogens of λgt11-ftz Av10 were established in RY1089. The pMC9 plasmid carrying the lac Z gene was cured from the strain to permit optimal induction. (Even at 10 mM IPTG induction of hybrid protein is not as efficient as when no lac Z repressor gene is present.) A culture of the lysogen was grown at 30°C overnight in NZCYM to saturation and then was diluted 1/100 and grown to a density of 4 × 10^4 cells/ml at 30°C. To induce the production of hybrid protein, the culture was
shifted to a 45°C waterbath for 15 min. After 2 hr at 37°C, cells were harvested by centrifugation and solubilized in 1% SDS-polyacrylamide gel sample buffer in 1/10 of the original cell culture volume and then heated for 5 min at 100°C. Lysates were clarified by centrifugation for 5 min in an Eppendorf centrifuge, and 25 μl of the supernatant was loaded per lane onto an analytical 7.5% SDS-polyacrylamide gel (Weber and Orns, 1975).

### Purification of Hybrid Proteins

The β-galactosidase-ftz Av10 fusion protein was purified by several methods. First, to obtain pure protein in soluble form, it was extracted and purified by amphenothyinolgalactopyranoside-agarose (Sigma) affinity chromatography (Germann, 1983). About 1 mg of intact protein per liter of induced culture was obtained in this fashion. About 90% of the fusion protein was not extracted by the above method, but could be solubilized in a buffer containing 1% (vol/vol) of Triton X-100 (PBT) for 3 hr at 4°C. The embryos were incubated overnight at 4°C with agitation in a solution of purified antibodies (1-2 μg/ml) diluted in the PBT buffer. After the primary incubation, the embryos were washed 3-4 hr with 5-6 changes of PBT. Goat anti-rabbit IgG-FITC, which had been previously incubated with fixed embryos, was diluted 1:500 in PBT. The embryos were incubated in the secondary antibody for 3 hr. The embryos were washed as above, stained with 50 ng/ml diamidino-phenylindole (DAPI) for 3 min, and washed for 30 min more. The embryos were mounted in Tris buffer containing 4 mM sodium acid under a coverslip and were viewed by epifluorescence microscopy. Photography was with Kodak Tri-X film used at ASA 1600 and developed in Diafine (Acufine, Inc., Chicago, IL).

### Production, Purification, and Assay of Antibodies

Rabbits were immunized with 500 μg of antigen in complete Freund’s adjuvant on day 0 and again with 200 μg on days 14 and 21 (and monthly thereafter) until the final exsanguination. Rabbits were bled on day 26 and about monthly thereafter until the final exsanguination.

### Immunofluorescence

Sera were absorbed to remove anti-β-galactosidase antibody by repeated passages on β-galactosidase-Sepharose 4B. The resin was made by coupling 1 mg of enzyme per ml of CNBr-activated resin. The columns were loaded with 5-20 ml of serum, washed with phosphate-buffered saline (PBS), PBS/Tween (0.1 M boric acid 25 mM sodium borate, 1 M NaCl, and 0.1% Tween 20) and PBS again. The bound antibody was eluted with 4 M Guanidine-HCl, 10 mM Tris (pH 8.0), and the eluate was dialyzed against PBS overnight.

The flow-through fraction of the β-galactosidase column was fractionated on an anti-β-galactosidase-DNP-avt 10 affinity column as follows: the column was prepared by attaching 1 mg of affinity-purified anti-β-galactosidase antibody per ml of CNBr-activated Sepharose 4B. A dialyzed urea extract containing 1 mg/ml of intact fusion protein was then passed over the column. The column was washed, until the effluent was free of protein, with PBS, PBS/Tween, and 0.2 M triethanolamine (pH 8.2). To attach the fusion protein covalently, the column beads were suspended in 10 ml of 0.2 M triethanolamine, 50 mM dithymethylmelamine-2 HCl and adjusted to pH 8.2. The antibody-antigen complexes were cross-linked by rocking the beads gently in this solution at room temperature for 45 min (Schneider et al., 1982). The beads were poured into a column and washed with PBS, noncovalently bound protein was removed with 4 M Guanidine-HCl washes, and the column was equlibrated in PBS. The serum from which anti-β-galactosidase antibodies had been removed was then applied, the column was washed as above, and the eluted antibody was dialyzed against PBS. The purified antibody was passed once more over the β-galactosidase column, to remove residual contaminating anti-β-galactosidase antibodies in order to obtain the final anti-ftz antibody.

The specificity of purified antibody was tested on proteins resolved by PAGE and transferred to nitrocellulose (Towbin et al., 1979) from lysogen extracts and from purified proteins by incubating a blot with primary antibody (1 μg/ml) for 2 hr, washing with PBS, PBS/Tween, and PBS, and then incubating the blot with affinity-purified alkaline phosphatase-conjugated goat anti-rabbit IgG (Cappell). After washing, the color was developed with bromochloro indolyl phosphate and nitroblue tetrazolium at pH 9.5 in 50 mM NaCO3.

### Immunofluorescence

Whole Drosophila embryos were dechorionated, permeabilized in heptane, fixed in formaldehyde, and devitellinized in heptane/methanol-acetic acid to 4°C with agitation in a solution of purified antibodies (1-2 μg/ml) diluted in the PBT buffer. After the primary incubation, the embryos were washed 3-4 hr with 5-6 changes of PBT. Goat anti-rabbit IgG-FITC, which had been previously incubated with fixed embryos, was diluted 1:500 in PBT. The embryos were incubated in the secondary antibody for 3 hr. The embryos were washed as above, stained with 50 ng/ml diamidino-phenylindole (DAPI) for 3 min, and washed for 30 min more. The embryos were mounted in Tris buffer containing 4 mM sodium acid under a coverslip and were viewed by epifluorescence microscopy. Photography was with Kodak Tri-X film used at ASA 1600 and developed in Diafine (Acufine, Inc., Chicago, IL).

### Acknowledgments

We thank Dr. Allen Laughon for his continued interest in, and enthusiasm for, all phases of this project, Robert Laymon for superb technical assistance, and Dr. Bionon DiNardo and Patrick O’Farrell for valuable discussions and for communicating unpublished data. We are grateful to Dr. Michael Klymkowsky for advice on immunofluorescence procedures and for the use of his microscope. Ur. 11 mm for advice on staining embryos, Drs. Susan Dutcher and Michael Klymkowsky for criticizing the manuscript, and Cathy S. Inouye and Karen Brown for secretarial help.

This work was supported by NIH grant #1 RO1 HD18163 to M. P. S. and a Damon Runyon-Walter Winchell Postdoctoral Fellowship #DRG-659 to S. B. C.

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

Received July 12, 1985

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