Functional analysis of the Drosophila twist promoter reveals a dorsal-binding ventral activator region

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Twist is one of the earliest expressed zygotically active genes required for dorsal–ventral pattern formation in the Drosophila embryo. Genetic studies suggest that this gene is activated in the ventral part of the blastoderm by maternally expressed dorsal gene product. Using P-element-mediated germ-line transformation, we have mapped a small (260 bp) dorsal-dependent ventral activator region (VAR) in the 5'-flanking region of the twist promoter that can direct the early ventral expression of a heterologous promoter. The VAR contains binding sites for a number of proteins present in extracts of Drosophila embryos. One of these sites bears homology to known binding sites for the dorsal transcription factor and is specifically bound by bacterially expressed dorsal protein. Furthermore, a 37-bp deletion that removes the dorsal-binding sequences abolishes the ventral-specific activity of the twist promoter constructs. Our data also show that additional sequences within the VAR are required to render the dorsal-binding sites functional. Finally, reverse genetic and biochemical data suggest that the transcription factor, encoded by the zeste gene, may help control the overall level, if not the pattern, of twist expression.

[Key Words: twist, dorsal, transcription, Drosophila embryogenesis, dorsal–ventral pattern formation]

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A large number of genes that influence the process of Drosophila embryogenesis have been identified, and many of them have been found to encode DNA-binding proteins that serve to control the transcription of downstream genes [Ingham 1988; Govind and Steward 1991]. Intensive genetic and molecular analysis has succeeded in organizing these genes into intricate regulatory hierarchies. Most developmentally important genes can be grouped into one of two classes: [1] those that determine cell fate as a function of position along the anterior–posterior axis and [2] those that determine cell fate as a function of position along the dorsal–ventral axis of the early embryo. The former class specifies the segmental structure of the embryo, whereas the latter set determines the germ layers of the developing organism. The initial designation of position along the dorsal–ventral axis involves the action of a large set of maternally active genes, most of which belong to the "dorsal group" of maternal effect genes. The role of these genes is to generate an activity gradient of the product of the dorsal (dl) gene (itself a member of the dorsal group) emanating from the ventral midline of the blastoderm embryo. It is this activity gradient of the dl protein that specifies the dorsal–ventral coordinate axis of the embryo.

The dl gene exhibits extensive sequence similarity to the nuclear proto-oncogene c-rel [Steward 1987] and the gene encoding the human transcription factor NF-kB [Ghosh et al. 1990; Kieran et al. 1990], and thus is believed to encode a transcription factor. In support of this idea, it has been shown that the dl protein is a DNA-binding protein that specifically recognizes sequences similar to those bound by NF-kB [Ip et al. 1991]. Furthermore, in accord with genetic data suggesting that dl is a repressor of zerknäfft (zen), the zen ventral repressor element was found to contain binding sites for the dl protein [Ip et al. 1991]. The site of action of the dl protein is the nucleus [Steward 1987], and a number of researchers have demonstrated that the activity gradient of the dl protein is caused by a gradient of nuclear localization in the syncytial blastoderm embryo [Roth et al. 1989; Rushlow et al. 1989; Steward 1989]. Specifically, the dl protein is transported efficiently into ventral nuclei, less efficiently into lateral nuclei, and not at all into dorsal nuclei.

Partially in response to the dl activity gradient, a number of early zygotically active genes are expressed in discrete domains along the dorsal–ventral axis of the embryo [Govind and Steward 1991]. These include the dorsally active genes zen and decapentaplegic (dpp), which are initially expressed in the anlagen of the dorsal ectoderm and the amnioserosa, and the ventrally active genes snail (sna) and twist (twi), which are initially expressed in the mesodermal anlagen. Genetic evidence
strongly suggests that *dl* is a repressor of *zen* and *dpp* and an activator of *twi* and *sna*. Thus, studies of *dl* should reveal information about the mechanisms of both transcriptional activation and repression.

As indicated above, *twi* is first activated in the ventral 20–25% of the nuclei of the blastoderm embryo in the mesodermal anlagen (Thiss et al. 1987, 1988). This gene is also expressed in the anterior and posterior poles of the early embryo. Genetic analysis demonstrates that the early ventral activation of *twi* is completely dependent on *dl* (Thiss et al. 1987). *twi* expression is absolutely required for gastrulation and for the differentiation of mesoderm. *twi* appears to encode a helix–loop–helix transcription factor (Murre et al. 1989); therefore, it presumably carries out its functions in gastrulation and cell differentiation by activating or repressing downstream genes at the transcriptional level.

To help elucidate the mechanisms that generate the spatial and temporal pattern of *twi* transcription, we have used P-element-mediated germ-line transformation to map cis-regulatory transcriptional control elements in this gene. In addition, we have initiated a biochemical analysis of the protein factors that interact with *twi* control regions. These experiments have allowed us to identify a number of elements in the 5′-flanking region of the gene that contribute to the normal *twi* pattern of expression. In particular, we have mapped a small region that directs the ventral activation of *twi*. This region is bound by several nuclear proteins in the *Drosophila* embryo, including the *dl* protein. Our results suggest that *dl* bound to this region interacts either directly or indirectly with other DNA-bound regulatory factors to activate *twi* in the presumptive mesoderm.

**Results**

5′-Deletion analysis of the *twi* promoter defines a small region sufficient for ventral-specific expression

In Figure 1 (A,B), we show whole-mount in situ hybridizations in which wild-type blastoderm stage *Drosophila* embryos have been probed with a digoxigenin-labeled probe containing the *twi*-coding region to reveal the spatial distribution of the *twi* mRNA. In agreement with data published previously (Thiss et al. 1987, 1988), we find that *twi* is first expressed in the ventral 20–25% of the blastoderm (in the mesodermal anlagen) and that this early domain of expression also includes the anterior and posterior poles of the embryo.

To locate regions of the *twi* gene sufficient to direct this ventral-specific RNA synthesis, we created a series of transgenic fly lines bearing various portions of the *twi* 5′-flanking region fused to the *lacZ*-coding region. In our first series of experiments, we examined a set of 5′-deletion mutants extending from a variable position upstream of the *twi* transcriptional start site to a fixed position just downstream of the *twi* initiation codon (+236). Embryos from the various transgenic lines were fixed and subjected to in situ hybridization with a digoxigenin-labeled probe specific for the *lacZ* mRNA. For each construct, the embryos shown are representative of multiple independently derived transgenic fly lines. A construct containing 5 kb of 5′-flanking sequences fused to the *lacZ*-coding region gives strong and specific ventral expression in the cellular blastoderm that is essentially indistinguishable from endogenous *twi* expression (−5000 *twi/lacZ*, Fig. 1C,D). Likewise, deletion mutants containing only 1829 bp (−1829 *twi/lacZ*, Fig. 1E,F), 920 bp (−920 *twi/lacZ*, Fig. 1G,H), or 648 bp (−648 *twi/lacZ*, Fig. 1I,J) of 5′-flanking sequences exhibit specific ventral expression [although, as discussed below, the exact expression patterns of −920 *twi/lacZ* and −648 *twi/lacZ* differ somewhat from those of the longer constructs]. In contrast, a construct containing only 269 bp of 5′-flanking sequences does not exhibit ventral expression in the cellular blastoderm (−269 *twi/lacZ*, Fig. 1K). Thus, these experiments (summarized in Fig. 1L) suggest that the region extending from −648 to −269 relative to the transcriptional start site contains an element that is required for ventral activation of *twi* in the *Drosophila* embryo.

A detailed examination of the embryos bearing the −920 *twi/lacZ* or −648 *twi/lacZ* constructs (Fig. 1G–J) shows that they exhibit expression patterns that differ in a number of ways from those observed with the longer constructs (Fig. 1C–F). First, the level of activity is somewhat reduced. Second, the domain of expression does not include the anterior or posterior poles of the embryo (see arrows, Fig. 1E,G). Third, the width of the ventral *lacZ*-expressing domain is narrowed (see arrows, Fig. 1F,H). Finally, these promoter constructs exhibit aberrant expression in the head region, which appears slightly later than the ventral expression [late stage 5; see arrow, Fig. 2A]. Thus, the region upstream of −920 bp seems to contain regulatory elements required to direct expression in the termini, to strengthen and broaden the ventral domain and to repress aberrant expression in the head region.

To determine whether the observed ventral expression is dependent on *dl*, homozygous males containing *twi/lacZ* fusion constructs were crossed to females homozygous for a null mutation in *dl*. The resulting embryos were analyzed as described above, and one example (with the −920 *twi/lacZ* construct) is shown in Figure 2. Similar results were observed with other constructs [data not shown]. In the absence of *dl* gene product (Fig. 2B), the embryos lack the ventral domain of *lacZ* expression observed in the wild-type background (Fig. 2A). Thus, the 5′-deletion series appears to define a small *dl*-dependent ventral activator element. Interestingly, unlike the ventral-specific activity, the aberrant head-specific expression characteristic of some of the deletion constructs seems to be independent of *dl* and is retained in the *dl* background (see arrows, Fig. 2B,C). The *dl*-independent head expression was observed with multiple transformants of the −920 *twi/lacZ* and −648 *twi/lacZ* constructs but not with the longer constructs [which use the same vector]. This suggests that this expression is an inherent property of certain *twi* promoter deletion mutants and not a position or vector effect.
The ventral activator can direct the expression of a heterologous promoter

To determine whether the ventral activator can work in conjunction with a heterologous promoter, we fused various portions of the 660-bp proximal promoter region to a basal level hsp70 promoter (Fig. 3). Embryos from transgenic fly lines bearing these constructs were collected, and the expression patterns of the lacZ transgenes were assessed as described above. Sequences extending from −660 to −55 (twi/hsp-1) were found to direct ventral-specific expression of the hsp70 basal level promoter element (Fig. 3A), demonstrating that the ventral activator can mediate the ventral-specific expression of a heterologous promoter. This expression is approximately five- to eightfold weaker than that observed for constructs containing the normal twi promoter and 648 or 920 bp of 5′-flanking sequences. Thus, the constructs shown in Figure 3 were stained longer than those in Figure 1.

When the 605-bp element extending from −660 to −55 (twi/hsp-1) was deleted from the 3′ end, it was found that deletion to −205 (twi/hsp-2; Fig. 3D,E) or −269 (twi/hsp-3; Fig. 3F,G) resulted in a sharp drop in overall promoter strength, but weak ventral-specific activity was retained. Further deletion to −306 (twi/hsp-4) resulted in the complete loss of early ventral activity. Thus, these 3′ deletions define a 37-bp region (from −306 to −269) that contains sequences essential for ventral activation. When the 605-bp element was deleted from the 5′ end, a deletion to −537 (twi/hsp-7) was found to have no effect (Fig. 3B,C), whereas a deletion to −323 (twi/hsp-8) eliminated ventral expression. Taken together, these results (which are summarized in Fig. 3H) suggest that the 605-bp region contains at least two functional units (Fig. 4B). Part or all of the first unit (which we have named element z) lies between −55 and −205. This element is not required for ventral-specific activity.

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**Figure 1.** Expression of twi/lacZ transgenes in Drosophila embryos. Patterns of expression were revealed by in situ hybridization for twi [A and B] or lacZ [C–K] mRNA. Stage 5 (cellular blastoderm) embryos are shown in both sagittal [A,C,E,G,I,K] and ventral [B,D,F,H,J] views. The various constructs used are shown in L. All frames show anterior to the left; in the sagittal views, dorsal is up. [A,B] Wild-type (Oregon R) embryos probed to reveal normal twi pattern of expression. (C,D) −5000 twi/lacZ. (E,F) −1829 twi/lacZ. (G,H) −920 twi/lacZ. (I,J) −648 twi/lacZ. In contrast to the −5000 twi/lacZ and −1829 twi/lacZ constructs, the domains of lacZ expression in the −920 twi/lacZ and the −648 twi/lacZ constructs do not include the termini (arrows in E,G). Also, these latter two constructs exhibit ventral domains of lacZ expression that are narrower (cf. arrows in F,H) and of lower intensity. (K) −269 twi/lacZ (no staining).
but influences overall promoter strength. The second functional element lies somewhere between −537 and −269 and constitutes a ventral activator region (VAR), as it is able to mediate the ventral activation of the heterologous hsp70 basal level promoter.

To initiate an analysis of the trans-acting factors responsible for the ventral-specific expression of twi, we scanned the 600-bp proximal promoter region using a footprinting assay. Probes covering the entire region were incubated with crude Drosophila embryo extracts before being subjected to DNase I digestion. Representative footprinting data is shown in Figure 4A and summarized in Figure 4C. The footprint analysis reveals three prominent clusters of footprints (Fig. 4B). One cluster consists of four strong protections between −10 and −120, three of which lie within the boundaries of element z. Three protections are found between −240 and −312, a region that overlaps the 3′ boundary of the VAR. Finally, the third cluster includes three regions of protection that are located between −604 and −484, a region that overlaps the 5′ boundary of the VAR.

The activity of element z may be mediated by the zeste transcription factor

As mentioned above, the proximal 120 bp of the twi promoter was found to contain four strongly protected regions of ~10–15 bp each, three of which lie within the bounds of element z. Inspection of the protected sequences (Fig. 4C) reveals that they each contain either a five of six or a six of six match for the binding site of the transcription factor encoded by the zeste gene (Biggin et al. 1988). These sites are protected by a homogeneous preparation of Drosophila zeste protein purified on the basis of its affinity for sites found in the Ultrabithorax (Ubx) promoter (data not shown). Thus, the activity of element z may be mediated, at least in part, by the zeste transcription factor.

Distant regulatory elements are not normally active in conventional in vitro transcription systems. However, as the zeste-binding sites lie within the 120-bp region proximal to the transcriptional start site, we were encouraged to examine the activity of these sites in an in vitro transcription system. Thus, we analyzed a series of 5′-deletion mutants for transcriptional activity in an extract derived from Drosophila embryos (Fig. 5). Deletion of the most distal zeste site has little or no effect on promoter activity (Fig. 5A, cf. lanes 1 and 2 with lanes 3 and 4). In contrast, a deletion that leaves only the two proximal zeste sites intact (lanes 5,6) causes a sharp reduction in promoter activity. The further deletion of one of the two remaining zeste sites does not cause a further reduction in promoter activity (lanes 7,8). Thus, the region containing three of four zeste-binding sites seems to be required for efficient in vitro transcription.

The experiments described in the preceding paragraph were carried out with transcription extracts prepared by the method of Heiermann and Pongs (1985; HP extracts). A different result is obtained if we use extracts prepared by the method of Soeller et al. (1988; SK extracts). In this case, high levels of transcription are observed for all the deletion templates and there does not seem to be any dependence on sequences upstream of −51 (Fig. 5B, cf. lanes 1 and 2). It has recently been demonstrated that a major difference between these two types of extracts is
the presence of histone H1 in the HP extracts and the absence of this protein in SK extracts (Croston et al. 1991; Kerrigan et al. 1991). We have found that the addition of H1 to the SK extracts has two effects on the in vitro activity of the twi promoter [lanes 3–7]. First, this protein has an overall negative effect on promoter activity. (This is also true of the Adh proximal promoter, which was used as an internal control in these in vitro transcription experiments.) Second, the addition of this protein restores the activity of sequences upstream of −51. Thus, in the HP extracts, the zeste-binding sites seem to act to overcome a generalized repression by histone H1. This conclusion is strengthened by the finding that in the presence of excess DNA [lanes 10–12], the HP extracts behave similarly to the SK extracts [i.e., high levels of expression are observed both in the absence and presence of multiple zeste-binding sites]. This is presumably because free H1, which is a nonspecific DNA-binding protein, is titrated out of solution by excess DNA.

The essential 37-bp segment of the VAR contains binding sites for the dl transcription factor

As noted above, the VAR is sufficient to drive the dl-dependent ventral expression of a heterologous promoter. Ip et al. (1991) have shown that dl is a sequence-specific DNA-binding protein with a binding specificity very similar to that of the human transcription factor NF-κB. Presumably, therefore, dl controls transcription by binding directly to cis-regulatory elements and influencing the rate of transcriptional initiation. Thus, we expect to find one or more binding sites for the dl protein in the VAR. Inspection of the 600-bp proximal promoter region reveals no perfect matches for the consensus sequence defined by Ip et al. (1991). However, we do find a single 9 of 10 match [nucleotides −299 to −290]. This best match falls near the 3′ border of the VAR and is contained within one of the footprints in that region [Fig. 4C]. Furthermore, this sequence falls within the 37-bp region that the 3′-deletion analysis in Figure 3 demonstrated to be necessary [but not sufficient] for ventral activation.

To test the idea that this site represents a binding site for the dl transcription factor, a truncated dl cDNA containing the sequences encoding the DNA-binding domain was expressed in Escherichia coli. The resulting protein was tested for binding to the twi promoter by a DNase I footprinting assay. As shown in Figure 6A, the bacterially expressed protein binds specifically to a 40-bp region in the twi promoter, including the 9 of 10 match for the dl consensus-binding site. This protection, which is the only such footprint observed in the VAR, extends in both directions to include additional weaker matches...
Figure 4. Footprint analysis of the 600-bp twi proximal promoter region. [A] Three uniquely end-labeled footprinting probes were generated by using exonuclease III/mung bean nuclease deletion derivatives of the twi promoter region. These probes were 5'-end-labeled at -155 (left), -434 (middle), or -648 (right). Probes were incubated with no extract (lanes 1, 4, 5, 7, 8, 10), 4 μl of SK embryo extract (lanes 2, 6, 9), or 8 μl of SK embryo extract (lane 3). Reaction products were then displayed on a 6% sequencing gel. The approximate positions of the footprints discussed in the text are shown by the boxes at right. Lanes M contain the appropriate G + A chemical sequencing ladders. Additional data (not shown) indicates that there are no significant protections between -200 and -130. [B] The positions of the various protected regions are indicated by boxes. The lines above the sequence represent the two transcriptional elements inferred from our deletion analysis (in particular, Fig. 3). [C] Sequence of the twi proximal promoter region. The shading indicates the sequences protected from DNase I digestion by the crude Drosophila embryo extracts. The bracket under the sequence shows the extent of the footprint generated by bacterially expressed dl protein (see Fig. 6). Also indicated are the dl and zeste recognition sequences discussed in the text. The zeste consensus-binding site is (C/T)GAG(C/T)G (Biggin and Tjian 1989). The dl consensus binding site is GGGA(A/G)A(A/T)(A/C)C(A/C) (Ip et al. 1991). The central dl binding site is labeled in larger type because it represents the best match for the dl consensus-binding sequence.

for the consensus sequence on either side of the best match (Fig. 6B). Thus, this protection seems to represent the binding of dl protein to three consecutive sites within the VAR. Interestingly, all three sites contain one half-site that matches the zen consensus perfectly and a second half-site of lesser quality. In conclusion, the activity of the VAR appears to be the result, at least in part, of its ability to serve as a binding site for the dl transcription factor. However, additional sequences (between -537 and -323) upstream of the dl-binding sites are also required for ventral activation.

Discussion

Our data show that a region of twi extending from -1829 to +236 can direct a pattern of transcription that is very similar to the wild-type twi pattern of transcription. Furthermore, these experiments allow us to identify a small region of the twi promoter that can mediate the dl-dependent ventral activation of a heterologous promoter. This region, which we have termed the VAR, lies between -537 and -269 relative to the twi transcriptional start site. The VAR includes a 40-bp region (from -277 to -317) that is specifically protected from DNase I digestion by bacterially expressed dl protein. A 37-bp deletion (from -269 to -306) that removes most of this region inactivates the VAR. Thus, the dl-binding region is essential for ventral-specific transcriptional activation.

DNase I footprinting experiments with Drosophila embryo extracts reveal that the VAR overlaps two clusters of protected regions centered at about -275 and -540. The proteins responsible for these footprints remain largely uncharacterized. However, the most 5' footprint in the cluster centered around -275 is con-
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Figure 5. In vitro transcription of the twi promoter. (A) Reactions (25 &x12;1) were programmed with 150 ng of one of the twi promoter deletion mutants shown at the bottom. In addition, reactions contained 6 &x12; of HP transcription extract. The arrow indicates the position of the correctly initiated twi transcripts. (B) Reactions (25 &x12;1) were programmed with 50 ng of the indicated twi promoter 5'-deletion mutants and 100 ng (lanes 1–9) or 400 ng (lanes 10–12) of a construct containing a 55 5' deletion of the Adh proximal promoter. In addition, the reactions contained 2 &x12; of SK extract (lanes 1–6) or 6 &x12; of HP extract (lanes 7–12), as well as the indicated amounts of histone H1. The positions of the correctly initiated twi and Adh transcripts are indicated by arrows. For all experiments in A and B, transcription products were visualized by S1 nuclease analysis as described in Materials and methods.

Figure 6. dl protein binds to the ventral activator region. (A) A probe that had been 5'-end-labeled at -434 was incubated with a bacterial extract containing the dl-378 protein or a control extract of nonexpressing bacteria. The probes were then digested with DNase I before electrophoresis on a 6% sequencing gel. (Lane 1) no protein; (lanes 2,3) 25 &x12; of the control extract; (lane 4) 10 &x12; of the control extract; (lanes 5,6) 25 &x12; of the dl-378 extract; (lane 7) 10 &x12; of dl-378 extract; (lane 8) no protein. Lanes 2 and 3 and 5 and 6 differ in the extent of DNase I digestion, with lanes 3 and 6 being under-digested relative to lanes 2 and 5. Lane M contains the A + G chemical sequencing ladder. (B) Alignment of the dl recognition sequences in the twi VAR, with the consensus-binding site derived from the sites in the zen ventral repressor (Ip et al. 1991). Stars indicate matches with the consensus.
hsp-7, which contains additional sequences upstream of the essential dl-binding sites, displays ventral activity. These additional sequences do not bind dl with any detectable affinity. This finding may reflect the ability of dl to serve as both a transcriptional activator (e.g., for twi and snai) and a transcriptional repressor (e.g., for zeste and dpp). It is likely that sequence elements outside the dl recognition sequence are responsible for modifying the activity of bound dl protein and converting it into either an activator or a repressor. One intriguing possibility is that the DNase I-protected region at the 3′ end of the VAR serves the function of turning dl protein bound to the 3′ end of the VAR into an activator, and we are currently testing this possibility.

The differential activity of the zen ventral repressor and the twi ventral activator could also be due to the existence of multiple dl-containing heteromeric complexes in the Drosophila embryo. As mentioned above, the dl protein bears sequence similarity to the human transcription factor NF-κB, and the two proteins have similar binding site specificities. The predominant DNA-binding form of NF-κB may be a heterodimer between two polypeptides termed p50 and p65 [Baeuerle and Baltimore 1989; Nolan et al. 1991]. In all of the experiments with bacterially expressed dl, the observed DNA-binding affinity is extremely low [Ip et al. 1991; J. Shirokawa and A.J. Courey, unpubl.). Although this could be the result of partially denatured bacterially expressed protein, it may also suggest that dl protein normally binds to DNA and regulates transcription as a part of heteromeric complexes with other polypeptides. To test these possibilities, it will be necessary to use biochemical methods to examine the protein–protein interactions between the dl protein and other proteins present in extracts of Drosophila embryos. These studies will be facilitated greatly by our ability to detect dl-binding activity in crude Drosophila embryo extracts.

Wild-type levels of twi expression appear to require sequences upstream and downstream of the VAR. In particular, sequences between −205 and −55 (element z) were found to boost the activity of the VAR severalfold. Sequences between −1829 and −920 also stimulate the activity of the twi promoter. In addition, these distal sequences are required to broaden the ventral domain, for early terminal expression, and to repress aberrant head expression of twi. We are currently carrying out experiments to determine whether this region can serve as an independent ventral activator region.

Element z contains multiple tandem binding sites for the zeste transcription factor. Our in vitro transcription experiments show that under certain conditions, promoter activity decreases significantly with the deletion of two of the zeste-binding sites. Specifically, we found that the activity of the region containing the zeste-binding sites was critically dependent on the presence of histone H1 in the transcription system. Thus, as with the Drosophila GAGA factor [Croston et al. 1991; Kerrigan et al. 1991], zeste protein may stimulate transcription by excluding H1 from the promoter region.

zeste is required for the process of transvection, whereby an enhancer element on one chromosome activates a promoter element on the homologous chromosome [Wu and Goldberg 1989]. Transvection requires normal chromosome pairing, and the zeste protein may act as a link to hold the promoter and remote enhancer together [Benson and Pirrotta 1988; Biggin et al. 1988]. Thus, the zeste-binding sites in element z may mediate the effects of the more distal regulatory elements in the twi promoter by anchoring a chromosomal loop. In this regard, it is interesting to note the presence of an additional possible zeste-binding site just 3′ of the dl-binding sites in the VAR (Fig. 4C).

In conclusion, the experiments described in this paper should lead to an understanding of the biochemical basis of transcriptional activation and repression. In addition, they will illuminate the mechanisms used to generate discrete germ layers during development. Ultimately, we hope to learn how the positional information present in morphogenetic gradients can result in the formation of complex multicellular organisms.

Materials and methods

P-element-mediated transformation and analysis of expression patterns

Germ-line transformation was carried out as described previously [Rubin and Spradling 1982]. Constructs were injected into w^1118 flies with p25.7wc helper DNA [Karess and Rubin 1984]. Multiple independent lines were established for each P-element construct and examined for β-galactosidase activity. In situ hybridization to whole-mount embryos was carried out according to standard procedures [Tautz and Pfeifle 1989] with digoxigenin-ATP-labeled probes (Genius kit, Boehringer Mannheim). The stained embryos were examined and photographed with a Zeiss Axioshot microscope and Nomarski optics. Embryo stages are according to Campos-Ortega and Hartenstein [1985].

To examine the dl dependence of the various P-element constructs we used a strain containing a balanced null allele of dl (dl^b, en sc/Cyo, DTS-100). w^1118 male flies homozygous for a P insertion were crossed to females homozygous for dl^b. The resulting embryos were analyzed as described above.

Plasmid constructions

We have constructed two series of plasmids for use in P-element-mediated germ-line transformation. The first contains various lengths of twi 5′-flanking region fused to the E. coli lacZ gene. These plasmids were constructed in two steps. First, a BamHI–PstI fragment (from −11 to +236) was cloned into the BamHI site of cosWhiteGal [Pirodda 1988] by adding a BglII linker (CAGATCTG, New England Biolabs) to the PstI end. This results in an in-frame fusion of the first 26 amino acids of twi to the eighth amino acid of β-galactosidase, with the additional sequence CCA GAT CCC (Pro Asp Pro) at the junction. To examine the dl dependence of the various P-element constructs we used a strain containing a balanced null allele of dl (dl^b, en sc/Cyo, DTS-100). w^1118 male flies homozygous for a P insertion were crossed to females homozygous for dl^b. The resulting embryos were analyzed as described above.

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The second series of P-element plasmids contains deletions of the 660-bp proximal promoter region fused to an hsp70/lacZ gene in a P-element vector that was constructed by G. Liaw and J. Lengyel (unpubl.). This vector contains the hsp70 promoter from −50 to the cap site fused to lacZ [Hiromi and Gehring 1987] and a mini-white gene as a marker. The deletions of the 660-bp proximal promoter region shown in Figure 3H were generated by polymerase chain reaction (PCR). For each pair of primers, one has an overhang bearing an overhang. This allows the PCR products to be cloned directionally into Xbal and NotI sites in the polylinker region of the vector.

The −648, −434, −269, −155, −99, −79, and −51 5′-deletion mutations of the twi promoter used in the footprinting and in vitro transcription analysis and in the subsequent construction of some of the P-element constructs were generated by deletion with exonuclease III and mung bean nuclease. A fragment extending from −1829 to +236 of the twi gene was subjected to unidirectional exonuclease digestion and the various size products were cloned in the Bluescript SK + vector.

The −55 deletion of the Adh proximal promoter and the S1 probe used for analysis of the twi transcripts was generated by cleavage of the −99 5′ deletion of the twi promoter with HindIII and Xbal, labeling with polynucleotide kinase, and isolating the nonspecific strand on a strand-separating gel. The correctly initiated transcript is expected to give a 263-bp protected species.

Expression of the truncated dl protein in E. coli

The truncated dl protein [dl-378] was expressed by using a T7 expression construct that is essentially identical to the one described by Ip et al. (1991). Expression was carried out with a strain of E. coli [BL21] lysogenic for a λ phage containing the gene for T7 RNA polymerase under the control of the lac promoter. Cells were grown to mid-log (OD_{560} = 0.5), and expression was induced with 0.4 mM IPTG. After an additional hour at 37°C, inclusion body extracts were prepared as described previously [Bohmann and Tjian 1989].

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Note added in proof

Since the submission of this paper, Thisse et al. [Cell 65: 1191–1201] have published an analysis of the twi promoter that reaches some of the same conclusions reported here. These investigators present evidence suggesting that dl protein binds specifically to the twi promoter, and in tissue culture cell cotransfection assays, they show that dl can activate twi transcription. The findings of Jiang et al. (this issue) overlap and complement the studies reported in this paper. In addition to the proximal VAR that is the focus of our study, these researchers have also identified a more distal dl-binding element [from 1.2 to 0.8 kb upstream of the twi transcriptional start site] that can independently activate ventral specific transcription from the twi promoter.

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