Disruption of the Abdominal-B Promoter Tethering Element Results in a Loss of Long-Range Enhancer-Directed Hox Gene Expression in Drosophila

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Abstract

There are many examples within gene complexes of transcriptional enhancers interacting with only a subset of target promoters. A number of molecular mechanisms including promoter competition, insulators and chromatin looping are thought to play a role in regulating these interactions. At the Drosophila bithorax complex (BX-C), the IAB5 enhancer specifically drives gene expression only from the Abdominal-B (Abd-B) promoter, even though the enhancer and promoter are 55 kb apart and are separated by at least three insulators. In previous studies, we discovered that a 255 bp cis-regulatory module, the promoter tethering element (PTE), located 5′ of the Abd-B transcriptional start site is able to tether IAB5 to the Abd-B promoter in transgenic embryo assays. In this study we examine the functional role of the PTE at the endogenous BX-C using transposon-mediated mutagenesis. Disruption of the PTE by P element insertion results in a loss of enhancer-directed Abd-B expression during embryonic development and a homeotic transformation of abdominal segments. A partial deletion of the PTE and neighboring upstream genomic sequences by imprecise excision of the P element also results in a similar loss of Abd-B expression in embryos. These results demonstrate that the PTE is an essential component of the regulatory network at the BX-C and is required in vivo to mediate specific long-range enhancer-promoter interactions.

Introduction

To ensure a high fidelity of gene expression patterns in embryos, a very strict functional requirement exists for the interaction of cis-regulatory modules (CRMs) in the genome of animals during development [1,2,3,4]. Embryonic transcriptional enhancers, which direct specific spatio-temporal patterns of gene expression in the embryo, are not permitted to promiscuously activate transcription from non-target promoters. Chromatin structural organization, insulators and promoter competition are thought to play a role in regulating enhancer-promoter interactions [5,6,7,8,9], for recent reviews see [10,11,12]. At the Drosophila bithorax complex (BX-C), an extensive network of CRMs located in over 300 kb of infraabdominal (iab) non-genic sequence is responsible for directing embryonic expression of just three homeotic genes: Ultrabithorax (Ubx), abdominal-A (abd-A) and Abdominal-B (Abd-B) (Fig. 1) [13,14]. These three homeotic genes are critical for establishing cellular identities in the presumptive thoracic and abdominal segments during development [2,15].

We recently identified a 255 bp promoter tethering element (PTE) located 5′ of the Abd-B transcriptional start site that is responsible for specifically recruiting the IAB5 enhancer from the BX-C to the Abd-B promoter in competition assays on transgenes [16]. Furthermore, the PTE also demonstrates anti-insulator activity by enabling the IAB5 enhancer to bypass an insulator from the BX-C to activate a target promoter on transgenes [17]. The ability of the PTE to facilitate specific enhancer-promoter interactions may explain how certain IAB enhancers, such as IAB5, IAB6, IAB7a and IAB7b [8,18,19], are able to bypass the Frontabdominal (Fab) insulator Fab-6, Fab-7 and Fab-8 [19,20,21,22] and direct transcription from the Abd-B morphogenetic (m) promoter at the endogenous BX-C (Fig. 1). As such, the PTE may be part of the complex regulatory network of CRMs, including the insulators, Promoter Targeting Sequences (PTSs) [23,24] and additional tethering sequences located 5′ of the Abd-B gene [25,26], responsible for mediating enhancer-promoter interactions in the BX-C. In contrast to the PTSs, which are located adjacent to the Fab-7 and Fab-8 insulators, the PTE is adjacent to the Abd-B promoter (Fig. 1). The PTSs have been implicated in mediating insulator bypass for the IAB enhancers on transgenic constructs [23,24]. However, deletion of either of the characterized PTSs from the endogenous BX-C did not result in a significant phenotype [8,18,19], suggesting that there might be other sequences capable of mediating the long-range enhancer-promoter specificity at the BX-C.

In this study, we demonstrate the functional importance of the PTE at the endogenous BX-C. Disruption of the PTE by a P element insertion in vivo results in a loss of IAB enhancer-directed Abd-B expression during development and a homeotic phenotype

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when placed in hemizygosity with an Abd-B null allele. Further molecular analysis of a mutant generated by imprecise excision of the P element reveals that a partial deletion of the PTE and neighboring upstream genomic sequences results in a similar loss of enhancer-directed Abd-B expression. These results demonstrate the critical functional role that the PTE plays in regulating promoter-enhancer interactions at the endogenous *Drosophila* BX-C.

### Results

**Insertions at the PTE disrupt Abd-B expression and result in homeotic transformation**

Genetic studies were carried out to determine whether the activity of the PTE is important for *in vivo* IAB enhancer-Abd-B interactions in the context of the BX-C. The previously identified *Abd-B* T2N mutation [27] is derived from a P element insertion in the endogenous PTE sequence (~241 bp relative the *Abd-B* transcription start site). The original element insertion line (*Abd-B*iso) contains a 5.6 kb Abd-B promoter region which includes 4.3 kb of 5’ sequence harboring the PTE sequence, fused to a lacZ reporter gene. This P element also contains the Fab-B insulator element, an IAB8 enhancer element and the raf gene. In contrast, the derived *Abd-B* T2N line contains only a truncated Abd-B promoter-lacZ reporter gene, generated by imprecise excision of the 5’ region of the P element (Fig. 2a). In the *Abd-B* T2N line the P element is inserted within the endogenous 255 bp PTE sequence, while leaving the entire Abd-B morphogenic (m) transcription unit intact [27]. The *Abd-B* T2N mutant is described as a Class I Abd-B mutant, capable of complementing mutations in the Abd-B regulatory (r) transcript, but affecting Abd-B m transcript expression [27]. In hemizygosity with the *Abd-B* M1 null allele, the *Abd-B* T2N insertion line shows a phenotypic transformation of abdominal segments 5-8 towards a more anterior abdominal segment identity. This is seen most clearly in the complete transformation of the seventh abdominal tergite to a more anterior identity in the *Abd-B* T2N insertion line (Fig. 2c) when compared to wild-type (WT) (Fig. 2b). The homeotic transformation is easiest to see in the cuticles of adult females due to their lighter abdominal pigmentation compared to males (Fig. 2b and c), although it is apparent in both sexes. This phenotype is characteristic of Abd-B m mutations [27,28], suggesting that the P element insertion into the PTE is disrupting expression from the adjacent Abd-B m promoter and is consistent with a reduction in IAB5-7 enhancer-Abd-B interactions.

In *D. melanogaster* embryos, the expression pattern of the *Abd-B* m transcript extends from parasegment (PS) 10–13 during the germ-band elongation stage of development, while the r transcript is predominantly restricted to PS14 at this stage [29,30].

*In situ* hybridization with a RNA probe that detects both the m and r transcripts in germ-band elongation stage embryos generated from crosses of heterozygous balanced *Abd-B* T2N adults (Fig. 2f) [27], demonstrates a loss of *Abd-B* expression in PS10–12 (presumptive abdominal segments 5–7) when compared with expression in WT embryos (Fig. 2d). These results indicate that the transposon insertion in the PTE impairs the ability of the IAB5–7 enhancers in the BX-C to drive expression from the endogenous Abd-B m promoter. The detectable expression of Abd-B in PS13 in *Abd-B* T2N embryos raises the question of why there is a homeotic transformation of the eighth abdominal segment in hemizygous *Abd-B* T2N/*Abd-B* M1 adults. While this phenotype is significant, it is only a partial transformation of A8 towards a more anterior identity (Fig. 2c), suggesting that there is not a complete loss of Abd-B patterning function. It is possible that there may be subtle modulation of the level of Abd-B gene expression in PS13 which, while responsible for the phenotypic transformation, is not readily detectable using RNA *in situ* hybridization. The expression of Abd-B persists in PS13 and 14 at least through stage 13 of development (Fig. 2h and i). However, it is also possible that transcription of Abd-B in late embryonic or larval stages may be modulated from the *Abd-B* T2N allele.

Knowing that expression from the endogenous Abd-B m promoter was perturbed in *Abd-B* T2N mutants, we examined whether the enhancers from the BX-C were re-directed to the intact ectopic *Abd-B* m promoter (containing the PTE sequence), which drives lacZ on the P element insertion. In *Abd-B* T2N embryos, lacZ expression was detected in a pattern extending from PS10 to 14 (Fig. 2g). The lacZ expression appears strongest in PS13 (presumptive abdominal segment 8) and weaker in PS10 (Fig. 2g), presumably due to the proximity of the endogenous IAB8 regulatory sequences to the target promoter, as previously observed [27]. This pattern of lacZ expression is consistent with the Abd-B expression pattern detected in wild-type embryos (Fig. 2d) and indicates that the IAB enhancers from the endogenous BX-C are now being re-directed to the intact ectopic Abd-B promoter (containing the PTE) on the P element to drive expression of lacZ in *Abd-B* T2N embryos.

**Deletion of PTE and neighboring upstream sequences results in a loss of enhancer-directed Abd-B transcription**

In the *Abd-B* T2N mutant endogenous PTE function is disrupted by P element insertion. In order to address the loss-of-functional activity of the PTE more directly we generated a mutant line in which the 5’ portion of the PTE and neighboring upstream genomic sequences are deleted from the endogenous BX-C locus.
Figure 2. P element insertion into the endogenous PTE results in homeotic transformation and loss of Abd-B expression. (A) Schematic diagram of the Abd-B\textsuperscript{T2N} P element line, showing the insertion site –241 bp 5’ of the Abd-B\textsuperscript{m} transcription start site in the endogenous PTE sequence. The same symbols and color scheme as in Fig. 1 indicate the neighboring cis-regulatory modules. (B) Dorsal cuticles were prepared from adult females. A normal abdominal pigmentation pattern is observed in wild-type (WT) adults. (C) In Abd-B\textsuperscript{T2N}/Abd-B\textsuperscript{M1} hemizygotes the seventh abdominal tergite (T7) is fully developed, indicating a transformation toward a more anterior abdominal segment identity. The eighth abdominal tergite (T8) also shows a partial transformation toward a more anterior identity. Abd-B\textsuperscript{M1} is a Class III null allele for both the Abd-B\textsuperscript{m} and r transcripts\cite{15,27}. Abd-B gene expression pattern in wild-type (WT) (D) and Abd-B\textsuperscript{T2N} (F) germ-band elongation stage 9 embryos. The staining patterns for the Abd-B\textsuperscript{T2N} line show reduced Abd-B expression in parasegments (PS) 10, 11 and 12 at stage 9 (F), stage 11 (H) and stage 13 (I) of development. The lacZ reporter gene is expressed in developing posterior regions of the Abd-B\textsuperscript{T2N} embryos (G). Expression is strongest in PS13, although the pattern extends from PS 10–14 and is very similar to the endogenous Abd-B transcription pattern. No lacZ expression is detectable in WT embryos (E). doi:10.1371/journal.pone.0016283.g002
This was accomplished by performing an imprecise excision of the P element insertion from the Abd-B^{ΔDN} mutant (Fig. S1). The Abd-B^{ΔDN} mutant was generated by a P element replacement strategy and carries an insertion containing the GAL4 and white genes in the identical location within the PTE (~241 bp relative to the Abd-B transcription start site) as the Abd-B^{ΔDN} line, but does not contain an ectopic copy of the PTE (Fig. 3a) [31]. In the Abd-B^{ΔDN} line, both the white and GAL4 genes are only expressed clearly in PS13 and very weakly in PS14 in embryos (data not shown), indicating that the distantly located IAB5–7 enhancers in the BX-C are not activating transcription of either reporter gene. The Abd-B^{ΔDN} line was utilized to generate an imprecise excision mutant as it carries the readily detectable white reporter gene (Fig. 3a). As a result, adult flies in which P element excision had occurred were easily identified by loss of red eye color (Fig. S1). PCR-based screening with primers from the proximal Abd-B m promoter region and from 0.5 kb, 1 kb and 1.5 kb 5′ of the Abd-B transcription start site was used to identify a 1.2 kb deletion in the Abd-B^{PTE-UP} allele (Fig. 3a–c). The molecular nature of the deletion was characterized by sequencing and found to have removed 53 bp of the PTE sequence 5′ of the original P element insertion site, as well as 1134 bp of endogenous genomic sequence 5′ of the defined PTE. In this Abd-B^{PTE-UP} allele, a portion of the GAL4 gene from the original P element remains, along with 202 bp of the 3′ PTE sequence (Fig. 3b).

Based on the Abd-B expression observed in the Abd-B^{ΔDN} insertion line, we hypothesized that the expression pattern of the endogenous Abd-B gene from the Abd-B^{PTE-UP} allele may also be perturbed. In situ hybridization with an RNA probe that can detect both the m and r transcripts shows that Abd-B gene expression is lost specifically in PS10-12, but not in PS13-14 in germ-band elongation stage embryos generated from crosses of heterozygous balanced Abd-B^{ΔDN} adults (Fig. 3d). Statistical analysis demonstrates that the number of embryos demonstrating this restricted Abd-B expression pattern from both the Abd-B^{PTE-UP} and Abd-B^{ΔDN} alleles is highly significant (p<0.01) when compared to the number of embryos with the WT Abd-B expression pattern. This restricted pattern of Abd-B expression persists at least through stage 13 of development (data not shown). One possible explanation for the loss of Abd-B expression in PS10-12 is that the genomic region deleted in the Abd-B^{PTE-UP} allele harbors a CRM capable of driving transcription in these specific parasegments. However, when tested in a transgenic reporter gene assay the 1.2 kb region does not exhibit embryonic enhancer activity (data not shown).

To confirm that mutations in the PTE only affect expression from the Abd-B m promoter, but not the Abd-B r promoter, in situ hybridization with a RNA probe (BPP, [32]) that specifically detects only the Abd-B r transcript was performed on embryos collected from WT, Abd-B^{ΔDN} and Abd-B^{PTE-UP} balanced lines. The expression pattern of the Abd-B r transcript was confirmed to be identical in the WT, Abd-B^{ΔDN} and Abd-B^{PTE-UP} embryos, appearing only in PS14 of germ-band elongation stage embryos (Fig. 3d). These observations are consistent with the known pattern of expression from the Abd-B r promoter [29] and confirm that the disruption to the PTE in these mutants only affects enhancer-mediated transcription from the Abd-B m promoter (see discussion for more detail).

**Discussion**

Parasegment-specific interactions between the IAB enhancers and Abd-B m promoter in the BX-C

The absence of Abd-B expression in PS10–12 of Abd-B^{PTE-UP} germ-band elongation stage embryos is consistent with a loss of IAB-enhancer directed expression from the Abd-B m promoter [29]. This suggests that deletion of the Abd-B m promoter tethering sequence (PTE) and the neighboring 1.1 kb 5′ sequence in the Abd-B^{PTE-UP} line leads to a disruption of the long-range interactions between the Abd-B m promoter and enhancers from the iab-5, iab-6 and iab-7 regions in PS10, 11 and 12, respectively. For example, in PS12 of WT embryos the tethering sequences upstream of the Abd-B m promoter enable the IAB7α and IAB7β embryonic enhancers to bypass the Fab-α chromatin insulator and drive expression from the Abd-B m promoter (Fig. 4a). In Abd-B^{PTE-UP} mutant embryos, removal of the tethering sequences appears to disrupt the ability of the IAB7α and IAB7β enhancers to activate the Abd-B m promoter, resulting in an absence of Abd-B expression in PS12 (Fig. 4a and 3d). Similarly, the IAB6 and IAB5 enhancers are unable to bypass intervening insulators to activate Abd-B m expression in PS11 and PS10 in Abd-B^{PTE-UP} embryos. In contrast to PS10–12, the specific enhancer-promoter interactions at the BX-C in PS13 appear to be intact in Abd-B^{PTE-UP} embryos. In WT embryos the IAB8 embryonic enhancer, located 3′ of the Abd-B gene, is solely responsible for directing expression from the Abd-B m promoter in PS13 [33]. In Abd-B^{PTE-UP} mutant embryos the Abd-B m transcript remains strongly expressed in PS13 (Fig. 3d). This result indicates that the interaction between the IAB8 enhancer and the Abd-B m promoter may not require tethering activity, likely due to the physical proximity of the enhancer to the promoter and lack of an intervening chromatin insulator (Fig. 4a). However, given the partial transformation phenotype observed for PS13 in flies carrying a disruption of the PTE sequences (as in the case of the Abd-B^{ΔDN} allele, Fig. 2c) it remains possible that loss of PTE activity may be responsible for subtle effects on transcription of Abd-B in PS13. In PS14, Abd-B expression in germ-band elongation stage embryos is not driven by the 3′ IAB enhancers, as it is initiated from the r transcriptional start site located 5′ of the PTE sequence (Fig. 1) [29,34]. Consequently, expression of the r transcript in PS14 is not perturbed by the loss of tethering activity in Abd-B^{PTE-UP} mutant embryos (Fig. 4a and 3d).

One possible outcome of the disruption of IAB enhancer interactions with the Abd-B m promoter at the Abd-B^{PTE-UP} allele may be the re-direction of those enhancers to the neighboring abd-A promoter in the BX-C. However, no change in the expression pattern of abd-A (extending from PS7 to PS13 in germ-band elongation stage embryos [35]) could be detected from the Abd-B^{PTE-UP} or Abd-B^{ΔDN} alleles (data not shown).

**Functional dissection of critical sequences at the PTE**

The functional activity of the PTE at the endogenous BX-C prompts the question of exactly what sequences associated with the PTE are necessary for confer promoter-enhancer tethering. Our earlier studies demonstrated that a 255 bp region located between −40 and −294 bp 5′ of the Abd-B m transcription start site is sufficient to tether the IAB5 enhancer to an ectopic promoter in a transgene competition assay (Fig. 4b) [16]. In contrast, an approximately 200 bp region extending from −100 to +100 relative to the Abd-B m transcriptional start site is not sufficient to mediate promoter-enhancer tethering when tested in similar transgenic assays [36]. In agreement with this observation, the 202 bp of sequence from the 3′ end of the PTE remaining at the endogenous BX-C in the Abd-B^{PTE-UP} allele is also not sufficient to mediate tethering of the IAB enhancers to the Abd-B m promoter (Fig. 4b). A previous study by Sipos and colleagues, utilizing an Abd-B mutant allele harboring a small deletion at the Abd-B m promoter region and a deletion at the iab-7 region carried on opposite chromosomes resulted in Abd-B transcriptional activity
Figure 3. Deletion of the PTE and upstream sequences at the endogenous BX-C disrupts Abd-B expression. (A) The Abd-B^LDN mutant features a P element inserted within the PTE (yellow box) (−241 bp relative to the Abd-B transcription start site (black arrow)). The P element contains GAL4 (purple box) and white (white box) reporter genes. PCR primers (black triangles) designed against the Abd-B promoter region and 0.5 kb, 1 kb and 1.5 kb upstream of the Abd-B m transcription start site were used to characterize the approximately 1.2 kb deletion (dashed box) in the Abd-B^PTE-UP mutant generated by imprecise excision of the P element. (B) The resulting Abd-B^ΔPTE-UP deletion removes 53 bp of the PTE sequence 5’ of the original insertion site and an additional 1134 bp of endogenous sequence 5’ of the PTE. A 476 bp portion of the GAL4 coding sequence (purple polygon) from the original P element remains, as well as 202 bp of the 3’ PTE sequence (yellow polygon). (C) Primers designed against the Abd-B promoter region and 1.5 kb upstream of the Abd-B transcription start site were used to genotype adult flies carrying the deletion allele by amplifying the 1.5 kb wild-type (WT) band and the 0.8 kb Abd-B^ΔPTE-UP band in balanced Abd-B^ΔPTE-UP (ΔPTE-UP/TM3,Ser) individuals. (D) In situ hybridization.
with an RNA probe that can detect both the Abd-B m and r transcripts (Abd-B) shows WT expression in parasegments (PS) 10–14 in yw67 embryos, while expression is found only in PS13–14 in a portion of embryos collected from crosses of balanced Abd-B\(^{\text{PTE-UP}}\) and Abd-B\(^{\text{LDN}}\) lines. In situ hybridization with a probe specifically designed against the Abd-B r transcript (Abd-B r) detects identical patterns of expression in PS14 of germ-band elongation stage embryos collected from WT, mutant Abd-B\(^{\text{LDN}}\) and Abd-B\(^{\text{PTE-UP}}\) balanced lines.

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Promoter tethering as a general regulatory mechanism

Additional examples of genetic complexes in which long-range interactions between CRMs have been characterized include the human beta-globin locus [37], the Drosophila Antennapedia complex [38,39] and mouse HoxD complex [3]. In the large genomes of vertebrates, where extensive global control regions have been identified, promoter-enhancer tethering is emerging as a critical general mechanism for regulation of gene expression [3,37]. More recently, in Drosophila, other PTE sequences have been located in the even-skipped [40] and engrailed [41] loci that mediate enhancer-promoter communication. The long-range CRM communication in these critical developmental genes has been shown to be dynamic, changing through the course of development in some cases [40] and perhaps even capable of mediating the evolution of novel patterns of gene expression in different insect species [42].

Our current model for the molecular function of the PTE in the Drosophila BX-C is that regulatory interactions that enable the PTE to tether the IAB enhancers to the Abd-B m promoter in specific parasegments during embryonic development may be mediated by chromatin looping [17]. A number of studies have indicated that chromatin looping may facilitate promoter-enhancer tethering through the action of different transcription factors. For example, the sea urchin GCF1 protein is able to form higher order multimeric loop structures when added to target site oligonucleotides in vitro [43]. The abundant mammalian transcription factor Sp1 has also been shown to form multimers and to strongly facilitate in vivo activation of a promoter by distantly located enhancer CRMs [44,45]. More recently, molecular studies have used high-magnification confocal imaging and 2D RNA fluorescence in situ hybridization (FISH) to visualize specific physical associations between distantly located cis-regulatory sequences in the nucleus at the human beta-globin locus [37].

Direct evidence for chromatin looping at the Drosophila BX-C comes from a study examining physical chromosomal interactions using probes designed against the IAB5 and IAB8 enhancers and a promoter-proximal region upstream of the Abd-B m transcription unit, containing the PTE sequence [46]. In a portion of nuclei taken from the eighth abdominal segment region of a germ-band elongation stage embryo, FISH signals from the Abd-B m promoter region co-localize with signals from the distal IAB5 enhancer, while the more proximal IAB8 enhancer remains distantly located and disassociated [46]. This result suggests that physical associations may indeed occur between the IAB5 enhancer and the 5′ upstream region of the Abd-B m promoter to facilitate expression of the Abd-B m transcript. These data support our model that tethering mediates the interaction of long-range enhancers (such as IAB5) to the Abd-B m promoter, but that it is not required for the functional interaction of the 3′ proximal IAB8 enhancer to the Abd-B m promoter.

In our current model the PTE may bind sequence-specific protein factors which interact with complementary factors bound near the IAB enhancers, allowing a molecular bridge or loop to form between the Abd-B m promoter and the IAB enhancers [17]. This model does not exclude the possibility that additional molecular interactions between known regulatory regions in the BX-C, including insulator and PTS sequences, may also mediate enhancer-promoter communication. In future studies other techniques, such as three-dimensional chromatin conformation capture (3C) [47] and Dam methylase identification [48], will be well suited to more fully elucidate the nature of such interactions and the molecular mechanisms of PTE function. In addition, molecular dissection of the functional sequences within the PTE and 5′ associated sequences in transgenic tethering assays and the biochemical identification of putative sequence-specific DNA-binding trans factors that may mediate the PTE activity will be essential.

Materials and Methods

Genetic insertion lines

The Abd-B\(^{\text{PTE}}\) and Abd-B\(^{\text{LDN}}\) P element fly lines inserted ~241 bp 5′ of the Abd-B m transcription start site were provided by Ernesto Sanchez-Herrero [27,31].

P element imprecise excision from the Abd-B\(^{\text{LDN}}\) line

Abd-B\(^{\text{LDN}}\) flies were crossed with a transposase expressing line (Δ2–3) carrying a Stubble (Sb) dominant phenotypic marker (BL Stock 1798) in Cross 1 (Fig. S1). Cross 1 male progeny were screened for Sb (marking the presence of Δ2–3 transposase) and variegated red eye color (rather than white eye color). The selected Cross 1 males were crossed with female with D, a dominant marker, and a TM3 balancer chromosome carrying Serrate (Ser), a dominant phenotypic marker (BL Stock 7198) in Cross 2 (Fig. S1). The male Cross 2 progeny were screened for Ser (marking the presence of the TM3 balancer), white eyes (indicating excision of the Abd-B-GAL4\(^{\text{LDN}}\) P element construct), and the absence of Sb (indicating loss of the Δ2–3 transposase) as well as against D. The selected Cross 2 male flies were crossed with BL Stock 7198 flies again in Cross 3 (Fig. S1). After a few days, the male parents were recovered from Cross 3 vials. PCR amplification of the Abd-B region with primers located 0.5 kb, 1 kb, and 1.5 kb upstream of the Abd-B transcription start site on the genomic DNA prepared from approximately 100 selected Cross 2 male flies was used to detect deletions in the PTE sequence.
Figure 4. PTE sequence is required for specific promoter-enhancer tethering at the Abd-B gene. (A) Abd-B transcription is disrupted in PTE mutants. In wild-type (WT) germ-band elongation stage embryos expression from the Abd-B m promoter is observed in parasegments (PS) 10–13 and from the Abd-B r promoter in PS14 [29]. The loss of Abd-B expression in PS10–12 in Abd-B^{PTE-UP} (ΔPTE-UP) embryos is consistent with a disruption of expression from the Abd-B m promoter. In PS12 of WT embryos, the tethering sequences upstream of the Abd-B m promoter enable the IAB7a and IAB7b enhancers (7a and 7b, orange circles) to act over the long-range of intervening sequence and drive expression from the Abd-B m promoter. In PS12 of Abd-B^{PTE-UP} mutants the removal of the tethering sequences disrupts the ability of the IAB7 enhancers to activate the Abd-B m promoter across the intervening Fab-8 insulator. A similar disruption of promoter-enhancer tethering activity at Abd-B^{PTE-UP} is also seen in PS11 (IAB6 enhancer) and PS10 (IAB5 enhancer). In PS13 of both WT and Abd-B^{PTE-UP} embryos, IAB8 (8, orange circle) is able, due to its physical proximity and lack of intervening chromatin insulator, to drive expression from the Abd-B m promoter. In PS14 of both WT and Abd-B^{PTE-UP} embryos, Abd-B r is transcribed independently of IAB embryonic enhancer activity and represses transcription from the Abd-B m promoter [28]. (B) Functional dissection of critical Abd-B tethering sequences. Schematic diagram of the regulatory region of the Abd-B gene shows enhancers (orange boxes) in the infraabdominal regions iab-5 to iab-8 (green) that are responsible for activating expression of the Abd-B gene in abdominal PS 10–13 during embryogenesis. Color scheme for other types of cis-regulatory modules is same as in Figure 1. Transgenic studies (highlighted in yellow box) show that a 1.4 kb of sequence from +1228 to −294, (including the 255 bp PTE sequence located 40 bp 5’ of the Abd-B m transcription start site), is sufficient for promoter-enhancer tethering [16]. Additional transgenic studies have shown that an approximately 200 bp sequence spanning the Abd-B m transcription start site (+100 to −100) is not sufficient for tethering activity [36]. P element insertion at the endogenous BX-C (as in the Abd-B^{LDN} or Abd-B^{T2N} mutations) demonstrates that the 242 bp sequence 5’ of the Abd-B m transcription
Primers used:

**Abd-B** promoter out: 5'-CGA CAA CAT ATC CAC ATC GCT-3'

**Abd-B** promoter 0.5 kb in: 5'-AAG TGC GAT ACC ATC TTT-3'

**Abd-B** promoter 1 kb in: 5'-TGG CTT TGG AAG TGA GAC AA-3'

**Abd-B** promoter 1.5 kb in: 5'-GGA AAT AGA TGG CCG CAG TTA A-3'

The PCR products were ligated into pGEM-T Easy vector (Promega) and sequenced using T3 and SP6 sequencing primers. Progeny from Cross 3 vials seeded with a male parent exhibiting a disruption of the PTE were screened against D and for Ser and self-crossed to generate a balanced mutant line.

**In situ analysis of abd-A and Abd-B expression**

In situ hybridization probes to detect transcription of **Abd-B** and **abd-A** were PCR-amplified using D. melanogaster yw**+/** adult genomic DNA as a template. The previously described DNA sequences for the Bexon region (exon 8 of the D. melanogaster Abd-B gene), BPP (specific to the r transcript) and Aexon region [32] were PCR amplified and cloned into pGEMT-Easy (Promega). PCR primer sequences were as follows:

Bexon s: 5'-GAAACAAGAAGACCTCAACGAC-3';

Bexon as: 5'-TGGAGCATGGTGATGGTAGG-3';

BPP s: 5'-TAATTATCCGTTCCAGTCG-3';

BPP as: 5'-CCCGATTGATGGGGTTTG-3';

Aexon s: 5'-CAACCAAGCCAGCAACACAGCAGC-3' (173566);

Aexon as: 5'-CATGTTATCAACCCTTGCG-3' (174736);

Antisense RNA probes (relative to the direction of **Abd-B** and **abd-A** transcription) were prepared using a digoxigenin (DIG) RNA-labeling kit (Roche, Gipf-Oberfrick, Switzerland). Embryos from each of the wild-type D. melanogaster, and mutant **Abd-B**DNA, **Abd-B**D**DN** and **Abd-B**PTE-UP lines were collected, fixed and hybridized with the appropriate probes as previously described [32]. Anti-sense Bexon RNA probes enable the detection of both the m and r transcript and anti-sense BPP RNA probes enable the specific detection of only the **Abd-B** r transcript [32].

**Supporting Information**

**Figure S1** Crosses to generate **Abd-B**PTE-UP mutant. (A) Schematic diagram of the **Abd-B**D**DN** P element insertion line, showing the insertion site −241 bp 5’ of the **Abd-B** m transcription start site in the endogenous PTE sequence. The P element insertion in the **Abd-B**D**DN** line contains GAL4 (purple box) and white (white box) reporter genes. The same symbols and color scheme as shown in Fig. 1 are used to show the cis-regulatory modules. (B) **Abd-B**D**DN** flies were crossed with a transposase expressing line (Δ2–3) carrying Stubble (Sb) (BL Stock 1798) (Cross 1). Cross 1 male progeny were screened for Sb and variegated red eye color (rather than white eye color). (C) The selected Cross 1 males were crossed with a female with D, a dominant marker, and a TM3 balancer chromosome carrying Serrate (Ser), a dominant phenotypic marker (BL Stock 7198) (Cross 2). Cross 2 male progeny were screened for Ser, white eyes (indicating excision of the GAL4 D**DN** P element construct), and the absence of Sb and D. (D) The selected Cross 2 male flies were crossed with BL Stock 7198 flies again (Cross 3). After a few days, the male parents were recovered from the Cross 3 vials. PCR amplification of the **Abd-B** promoter region with primers (black triangles) located 0.5 kb, 1 kb, and 1.5 kb upstream of the **Abd-B** transcription start site on the genomic DNA prepared from these selected Cross 2 male flies was used to detect deletions in the PTE sequence. Progeny originating from Cross 3 vials seeded with a male parent exhibiting a disruption of the PTE (ΔPTE) were recovered and screened against D and for Ser. (E) These selected Cross 3 progeny were then self-crossed to generate the **Abd-B**PTE-UP/TM3,Ser balanced line. (TIF)

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**Author Contributions**

Conceived and designed the experiments: MCWH BJS RAD. Performed the experiments: MCWH BJS OSA EB RAD. Analyzed the data: MCWH BJS OAS EB RAD. Contributed reagents/materials/analysis tools: MCWH BJS RAD. Wrote the paper: MCWH BJS RAD.

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