Drosophila Hox genes induce melanised pseudotumours when misexpressed in hemocytes

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Abstract

**Background:** Hox genes are key early determinants of cell identity along the anterior-posterior body axis across bilaterians. Recently, several late non-homeotic functions of Hox genes have emerged in a variety of processes involved in organogenesis in several organisms, including mammals. Being crucial factors in determining cell identity and organogenesis, the misregulation of Hox genes is likely to be associated with defects in these processes. Several studies have reported the misexpression of Hox genes in a variety of malignancies including acute myeloid leukaemia.

**Methods:** The Hox genes *Dfd, Ubx, abd-A* and *Abd-B* were overexpressed via the UAS-Gal4 system using *Cg-Gal4, Lsp2-Gal4, He-Gal4* and *HmlD3-Gal4* as specific drivers. Genetic interaction was tested by bringing overexpression lines in heterozygous mutant backgrounds of Polycomb and trithorax group factors. Larvae were visually scored for melanised bodies. Hemocytes were quantified by dissecting larvae for lymph in 4mm wells and staining nuclei with DAPI and tested for differentiation by staining them with anti-myospheroid and for proliferation with anti-PH3. Pupal lethality was carried out by letting pupae eclose and scoring those that failed after the time point.

**Results:** Expression of *Dfd, Ubx* and *abd-A*, but not *Abd-B* in the hematopoietic compartment of *Drosophila* led to the appearance of circulating melanised bodies, and increase in cell numbers, cell-autonomous proliferation and differentiation of hemocytes. Pupal lethality and the melanised pseudo-tumor phenotype were suppressed by the mutations in *Psc* and *esc* background while polycomb group member mutations *Pc* and *Su(z)12* and trithorax group member mutation *TrlR* increased the phenotype.

**Conclusions:** *Dfd, Ubx* and *abd-A* are leukemogenic. Mutations in Polycomb and trithorax group members, which are responsible for maintaining the expression state of the Hox genes, lead to repression in the leukemogenic phenotype. *Drosophila*, widely used as a model for myeloid leukemias, can serve as a testbed for Hox expression induced leukemias.

Background

Life comes in a variety of body forms. Despite this variety, there is similarity at the genetic and molecular level in the developmental mechanisms that lead to this variety across species. For example, in spite of the evolutionary distance between vertebrates and *Drosophila*, many organ and tissue types show a degree of homology with each other and many key developmental pathways governing their development and function are conserved.

The hematopoietic system is no exception. Hemocytes of *Drosophila* resemble the myeloid lineage of blood cells. The most abundant cells, plasmatocytes, are the equivalent of macrophages and are involved in a variety of processes such as responses to pathogens, removal of apoptotic cells, deposition of the extracellular matrix during embryonic development, etc. The next most abundant cells are Crystal
cells, specialised to induce melanisation reactions in the presence of pathogens and wound healing, resemble the granulocytes, and contribute about four per cent of the blood cells. Lamellocytes are the least abundant population of blood cells, usually only appearing in circulation upon the larva being challenged by any object too large to be cleared off by the macrophages, such as the eggs of a parasitoid wasp.

This conservation includes the molecular pathways involved in the development of these cell types. For example, *Drosophila Serpent* is related to GATA 1, 2 and 3 of vertebrates. GATA-2 is responsible for blood progenitor proliferation and survival, while GATA-1 is required for progenitor differentiation into erythrocytes, megakaryocytes and eosinophils. Similar to GATA-2, Srp is required for progenitor maintenance and proliferation. Loss of function in Srp leads to a reduced number of progenitors and a loss of all hemocytes. It is also required in plasmatocyte differentiation, similar to GATA-1. Additionally, *Drosophila ushaped* is related to the Friend of GATA (FOG) family. FOG-1 and GATA-1 together are required for erythrocyte and megakaryocyte differentiation. FOG-1 interacts with GATA-1 to repress eosinophil differentiation and must be downregulated for eosinophil differentiation. Similarly, *ush* is expressed in hemocyte precursors and plasmatocytes, and must be downregulated for crystal cell development.

Signalling pathways involved in regulating hematopoiesis, as would be anticipated, are conserved between vertebrates and *Drosophila*. For example, Jagged-1, the vertebrate homolog of *Serrate* and a ligand of Notch, is produced by the stromal cells of the bone marrow, to regulate Hematopoietic Stem Cell (HSC) proliferation and survival. *Ser* performs a similar role, being secreted by cells of the Posterior Signalling Centre (PSC), a set of regulatory cells at the posterior end of the Lymph Gland (LG). Vertebrate JAK2 is required for erythropoiesis, while STAT5 is required for proper progenitor and myeloid cell function. The *Drosophila* JAK/STAT pathway is required within the LG for the maintenance of prohemocytes, among other things. Transformations in JAK2 can lead to leukemogenesis in vertebrates, similar to how gain of function *hop* mutants behave. The Toll pathway is also conserved, playing a major role in innate immunity in both vertebrates and flies.

One aspect of vertebrate hematopoiesis that has not been mirrored in *Drosophila* is the role of Hox genes. Hox genes are well known for their conserved role in body axis formation across all bilaterians, but also play roles in vertebrate hematopoiesis, autophagy, as well as cell proliferation, differentiation, migration and apoptosis. Hox genes are transcribed in HSCs as well as lineage progenitors, and are suppressed in differentiated blood cells. Overexpression models show blockages in certain stages of development, expansion of HSCs, the circulation of blast cells, etc. For example, *Hoxa7* and *Hoxa9* have been shown to have a role in the development of hematopoietic progenitors of different lineages in mice. On the other hand, in *Drosophila*, other than *Antp*, which is implicated in setting up the location of the LG, as well as later marking the PSC, *Hox* genes have not been reported to play any role.
The expression of genes of the *Hox* cluster during, and after development is regulated by two chromatin remodelers, Polycomb and trithorax group (PcG and trxG) of proteins, which were discovered as transcriptional repressors (PcG) and activators (trxG) of *Hox* genes in *Drosophila*[^42]. Later, these proteins were shown to regulate many biological processes such as cell fate and lineage, cellular memory, stem cell function, and tissue homeostasis in cell lines and mouse models[^43–46]. The deregulation of *Hox* genes via Polycomb or trithorax proteins can lead to leukemogenesis by misregulation of hematopoiesis. Furthermore, PcG members EZH2, a human homolog of *Drosophila* E(z) protein, EED (Esc in *Drosophila*), SUZ(12) (*Drosophila* Su(z)12) and BMI-1 (homolog of *Drosophila* Psc) have been shown to have a role in different cancers in knock out studies carried out in cell lines as well as mouse model[^47–50]. Mixed Lineage Leukemia (MLL), a human homolog of *Drosophila* Trithorax (Trx) protein, regulates *Hoxa* expression in HSCs. MLL is a frequent fusion protein partner in acute leukemia[^51]. Evidence for the role of PcG and trxG genes in regulating HSC development in *Drosophila* remains largely to be explored[^52,53].

In this study, we show that overexpression of the *Hox* genes, *Dfd*, *Ubx* and *abd-A* in blood cells not only leads to melanised pseudo-tumors, but also to a significant increase in blood cell number and the induction of lamellocyte differentiation. Further, we present genetic evidence to show the role of PcG members, *Psc* and *Esc*, in the melanised pseudo-tumor formation induced by *Hox* genes. These findings will be helpful in understating the biological events associated with leukaemia in humans, which may open new possibilities of markers and therapy.

**Methods**

**Fly strains and culture:** Flies were cultured in standard commeal and sucrose agar. The wild-type flies used in this study were Canton-S. Flies were maintained at 25°C. For all experiments, flies were allowed to lay eggs for 6 hours before being transferred to a fresh vial. Larvae were screened and used for immunohistochemistry at 96-102 hours post egg laying, before the onset of metamorphosis. Supplementary Table S1 and S2 list the fly stocks used in this study.

**Larval screening for percent penetrance and severity of the phenotype:** For the over-expression of different *Hox* genes, the UAS-Gal4 binary system was used. To assess the effect of PcG and trxG members had in modifying the phenotype, heterozygous mutant lines were recombined with the *Cg-Gal4* driver (Supplementary Table S2 for all recombined stocks made in the lab). Confirmation of recombination was based on expression of *w*+ linked with the *Cg-Gal4* transgene and lethality when backcrossed with the mutant line. Recombined mutants with *Cg-Gal4* were maintained over the *CyO-GFP* balancer for GFP screening. Third chromosome mutants were crossed with homozygous *Cg-Gal4* lines and maintained over TM6B for screening via the *Tubby* phenotype. Experimental crosses were set between recombined strains (*Cg-Gal4* with mutant) and *UAS-abd-A* at a density of 12 females and 6 males for each cross. Egg lay was allowed for 6 hours and progeny were collected after 96 hours post egg lay, at the L3F stage. Screening was done using a stereomicroscope. Penetrance was calculated by calculating the percentage
of melanotic pseudo-tumor manifesting larvae. Severity of the phenotype was assessed visually. One-way ANOVA (Dunnett’s multiple comparisons) was performed to test the significance.

Pupal lethality count: To assess the pupal lethality, larvae were allowed to develop into pupae and were observed beyond 10 days post egg-lay. Eclosed progenies were considered as survivors. Dead pupae were counted manually.

For heterozygous mutant experimental pupae, larvae were first screened to confirm the presence of the Cg-Gal4 driven expression of UAS-abd-A and the presence of the mutation before being transferred to fresh vials. Second chromosome mutants were confirmed by selecting non-GFP larvae while third chromosome mutations were Tb+.

**Immunostaining and cell quantification:** For staining proliferative cells we made use the M-phase marker, Anti-PhosphoHistone 3 at serine 10, from Upstate (cat# 07-212, 1ng/μL). For confirming the presence of lamellocytes, we used anti-myospheroid (DSHB #CF.6G11, 27pg/μL). Blood cells were prepared using an established protocol. Blood cells numbers were quantified using a modified version of the protocol by Petraki, Alexander, & Bruckner, 2015. Larvae were dissected in 4mm wells, their hemolymph allowed to settle down, before being fixed with 1% formaldehyde and stained with DAPI. Each well was scanned using an Olympus IX83 at 20X, with 32 images stitched. Cells were quantified using CellProfiler by counting individual nuclei. Significance was tested using an unpaired t-test with Welch's correction between control and overexpression genotypes.

**Results**

**Tumor phenotype correlates with the tissue specificity and strength of the driver:**

In *Drosophila*, the collagen-Gal4 (Cg-Gal4) driver induces the strong expression UAS tagged genes in the fatbody as well as in the hematopoietic system. The different UAS Hox genes lines, *Dfd, Ubx, abd-A* and *Abd-B*, when brought under the Cg-Gal4 driver, induced melanised pseudo-tumors in larvae. This phenotype manifested in 26% of Cg-Gal4>UAS Dfd larvae, 60% of Cg>Ubx larvae, 82% of Cg-Gal4>UAS abd-A larvae and 4% of Cg-Gal4>UAS Abd-B larvae (Figure 1, 2A, 2B and Supplementary Table S3).

We then used the Hemese-Gal4 (He-Gal4) driver, which expresses throughout the lymph gland as well as in circulating hemocytes, and the HemolectinD3-Gal4 (HmlD3-Gal4) driver, which expresses in the cortical region of the lymph gland as well as in mature circulating hemocytes. While melanised pseudo tumours were observed in these genotypes, they appeared smaller and the penetrance of the phenotype was very low, manifesting in 3% of He-Gal4>UAS Dfd, 6% in HmlD3-Gal4>UAS Dfd, 9% in He-Gal4>UAS Ubx, 2% in HmlD3-Gal4>UAS Ubx, 8% in He-Gal4>UAS abd-A, 4% in HmlD3-Gal4>UAS abd-A, 3% in He-Gal4>UAS Abd-B and 2% in HmlD3-Gal4>UAS Abd-B (Figure 1, 2A, 2B and Supplementary Table S3). He-Gal4 induces expression throughout the lymph gland and in sessile cell pockets which reside underneath the larval cuticle. Thus, it expresses in all areas involved in hematopoiesis. Over-expression of Hox genes with the He-Gal4 driver always showed a higher penetrance of the phenotype when compared to HmlD3-Gal4.
Lamellocytes are responsible for the encapsulation mechanism in combating an immune challenge, and they do not express *Hemolectin*. The low penetrance of the phenotype in *HmlD3-Gal4* could be due to a lack of expression in lamellocytes \(^{57}\). Also, *Hemolectin* does not express in the medullary zone of the lymph gland, where cell proliferation and differentiation takes place \(^{58}\). It shows the phenotype is associated with active proliferation and differentiation of hemocytes of developing larvae. To test that the phenotype was not due to expression of the Hox genes in the fatbody (as *Cg-Gal4* expresses in both blood cells as well as the fatbody) we over-expressed these genes using the fatbody specific driver *Lsp2-Gal4*. No melanised spots were observed in such larvae, indicating that the pseudo-tumor phenotype is not induced by the misexpression of of Hox gene in the fatbody. We also observed the degree of penetrance is in proportion to the strength of the respective drivers in hemocytes (Figure 1, 2A, 2B and Supplementary Table S3). Taken together, this implied that the melanised pseudo-tumour phenotype we observe is of hemocyte origin.

**Tumor phenotype leads to lethality at pupal stage:**

We also noticed a significant level of pupal lethality when Hox genes were misexpressed in these conditions. Pupal lethality with the *Cg-Gal4* driver was highest when it drives *UAS-abd-A* (99 %). *Cg-Gal4>*UAS Dfd* (53%) and *Cg>Ubx* (24%) also show an increased lethality at pupal stage. It was negligible in *Cg-Gal4>*UAS-Abd-B* (2%). We observed lethality when the same genes were over expressed in the fatbody with *Lsp2-Gal4*. However, *Lsp2-Gal4* driven Hox expression induced lethality was lower compared to *Cg-Gal4* driven Hox expression induced lethality. But it must be noted that it was greater than that induced by the blood specific drivers used by us. Pupal lethality with *Lsp2-Gal4* driver was observed 9% in *Lsp2-Gal4>*UAS Dfd*, 26% in *Lsp2-Gal4>*UAS Ubx* and 31% in *Lsp2-Gal4>*UAS abd-A*. It has previously been shown that aberrant blood cells can induce pupal lethality \(^{59}\). However, while we did observe some pupal lethality when the Hox genes were expressed under He and Hml, the lethality was most prominent in when the *Cg-Gal4* or *Lsp2-Gal4* driver was used (Figure 2B, Supplementary Table S4) which supports the earlier report suggesting that Hox genes are repressors of autophagy in the fatbody \(^{27}\). Thus, while we do observe insignificant lethality with blood specific drivers since the expression of Hox genes in the fatbody does indeed induce lethality, the greater lethality when *Cg-Gal4* is used may be due to the concomitant expression induced in the fatbody as well as blood cells.

**Hox genes over-expression induces hemocyte proliferation and differentiation:**

Change in number cells and types of cells become important considering the phenotype observed upon misexpression of Hox genes. We quantified the number of blood cells in our overexpression lines using a modified version of established methods \(^{54,55}\). When expressed by blood specific driver, *Dfd, Ubx* and *abd-A* led to a significant increase in the number of circulating hemocytes (Figure 3A and 3B, Supplementary Table S5-8). Interestingly, while the penetrance of melanised spots was lower, blood specific drivers showed a larger number of blood cells (Figure 3B). Under the control of, *Lsp2*, the fatbody exclusive driver, however, *Ubx* and *abd-A* gave a significant increase in hemocyte number, despite them not manifesting melanised spots. Our results show that melanised spots (or pseudo-tumors), which have been reported as
the hallmarks of a “leukemia-like” phenotype in *Drosophila*, may not reflect an actual increase in hemocytes. Additionally, many studies have used the strong driver *Cg-Gal4*, which drives expression in fatbody as well as the blood cells. As our results show that perturbations in the fatbody may indeed lead to an increase in circulating hemocytes.

Previous studies have shown that cells of the LG do not enter into circulation until the onset of metamorphosis. However, *Hml* and *Cg* express in the cortical region of the LG, and *He* expresses throughout. Thus, the question arose as to whether the increase in cell number was due to an increase in cell proliferation at the LG or were circulating cells proliferating in a cell-autonomous manner. Hence, we checked for the presence of the mitotic cell marker PH3. We observed cells positive for PH3, when Hox genes were expressed in the blood cells, and not when expressed exclusively in the fatbody (Figure 3A, Supplementary Figure 1A-D). Unlike previous reports, we did not find proliferative cells in our control experiments. This may be due to a loss of cells in our preparations or more robust immunostaining on our part. Thus, while we cannot rule out the possibility that LG cells contribute to this increase, at least a fraction of the increase takes place due to the cell autonomous division of Hox overexpressing cells.

While imaging the blood cells, we noticed that there were larger, flattened cells in circulation, reminiscent of lamellocytes. To test whether they were bonafide lamellocytes, we stained the hemocytes for the lamellocyte marker myospheroid (Figure 4A, Supplementary Figure 1A-D). Control larvae infrequently showed the presence of lamellocytes. In our overexpression lines, however, we noticed that a significant number of cells were lamellocytes mys+. Some plasmatocytes also stained positive for mys. None of the plasmatocytes in the control flies or those overexpressing Abd-B were positive for mys. However, *Lsp2-Gal4>UASUbx* also had a significant number of lamellocytes. This is in keeping with reports that signals from the fat body can drive lamellocyte differentiation. Thus, we speculate that these cells, upon Hox overexpression, are pushed toward the lamellocyte fate (Figure 4A, 4B).

**Effect of PcG and trxG genes**

PcG members are known to function primarily through two distinct complexes, PRC1 (consisting of Pc, Psc, Su(z)2 and Sce) and PRC2 (consisting of E(z), Su(z)12, Esc and Caf 1-55). Members of the PcG and trxG have been shown to have a role in hematological malignancies in different clinicopathological data in leukemic patients and mice models. To determine their role in melanized pseudo-tumor formation in flies, we over-expressed abd-A using *Cg-Gal4* in the background of different PcG and trxG mutants. We selected *Psc*¹, *Pc*¹, *Su(z)2*, *Su(z)12*, *E(z)* and *esc2* from the PcG and *brm*², *Trl* from the trxG. Melanotic pseudo-tumor phenotype was used in our study to assay the effect of the mutants as it is convenient and robust. All experiments were performed in biological triplicates. The PcG mutants *Pc*¹, *Su(z)12*³, and trxG member *brm*² showed an increase in melanotic body formation (Figure 5A, 5B and Supplementary Table S9), and enhanced the phenotype upto 100 per cent. *Pc*¹ and *Su(z)12*² not only enhanced the penetrance (percentage phenotype showing larvae) but showed an increase in severity (scored as number and size of the black spots) compared to abd-A over-expressed in absence of mutants (Figure 5A). Pc and Su(z)12, both are the core proteins of PRC1 complex and play a role in negative
regulation of their target genes. Our results indicate these proteins might regulate melanotic body formation. Surprisingly, E(z) does not show any significant effect on penetrance. On the other hand, esc² (PRC2 member) and Psc¹ (PRC1 member) showed a significant decrease in penetrance 15% and 17% respectively (Figure 5B, Supplementary Table S9). The severity of the phenotype is also reduced in both the mutant background. These results indicate that genes involved in melanotic pseudo-tumor causing phenotype might be the target of the Esc and Psc proteins. Although it has been shown that Esc-E(z) complex is a thousand times effective to E(z) alone, our results suggest that Esc regulates its targets independent of E(z) activity or, for that matter, any other member of the PRC2 complex in the observed phenotype. Similarly, Psc mutation rescued the phenotype.

Effect of PcG mutants on the melanized pseudo-tumor related pupal lethality:

To test the effect of mutations on pupal lethality, L3F larvae from each combination, which manifested melanised pseudo-tumours, were transferred to fresh vials and allowed to pupate and eclose. Larvae from overexpressed abd-A (driven by Cg-Gal4) with melanotic body showed up to 99% lethality at the pupal stage. Further, we checked pupal lethality in mutant background. Since all mutants are maintained over balancers (Table S2), we selected overexpressed progenies without balancer to confirm mutant in the same progeny and transferred them in new food vials. Pupal lethality in Su(z)12³, Pc¹ and Su(z)2¹.a¹ was always 100% while we could get a few survivors from Cg-Gal4>UAS abd-A (Figure 3C, Table S10). A decline in lethality was seen in Psc¹, esc² brm² and TrlR85. The survivors from Psc¹ and esc² were quite healthy as compared to the survivors of Cg-Gal4>UAS abd-A. This reduction in lethality indicates that Esc and Psc proteins are strongly suppressing the melanotic pseudo-tumour phenotype and its consequences on development. Although brm² showed an increase in penetrance it decreases pupal lethality 89% compare to abd-A alone. TrlR85 showed a decrease in pupal lethality (79%).

Conclusion

Homeotic genes or Hox genes determine the cell identity across the anterior-posterior body axis early during development, a function that is conserved across all bilaterians. The later functions played by these genes, however, are less well studied. A number of reports indicate that they play a variety of non-homeotic functions later in development. Our lab has shown that abd-A, one of the three Hox genes of the bithorax complex of Drosophila, acts as a growth promoter in Histoblast Nest Cells. We show this to be a normal additional function of abd-A which involved adult cuticle formation during pupation. In the present study, we tested four Hox genes of Drosophila by ectopically expressing them in the blood cells and show that they are capable of inducing melanised bodies in circulation. These melanotic spots appear only when expressed in the lymph gland and circulating blood cells. The ectopic expression of the Hox genes also triggers cell proliferation. The cells appear to divide in a cell-autonomous manner, which is reflected in the detection of PH3+ cells in circulation. The presence of myospheroid positive elongated cells, seen in circulation, also suggests that Hox overexpression leads to the differentiation of the circulating blood cells into lamellocytes. Overexpression of abd-A, shows a relatively stronger phenotype
while Abd-B overexpression does not. It supports an earlier finding in which we observed a non-homeotic growth promoter role of abd-A but not of Abd-B during the formation of the adult cuticle during pupation.

Our results indicate that Hox genes are causal in leukaemia, reinforcing previous studies in vertebrate model systems, and extending these findings to Drosophila. This also opens the possibility that Hox gene induced leukaemias, especially those of the myeloid lineage, can be studied and modelled in Drosophila. Till date, the only known Hox gene to participate in Drosophila hematopoiesis is Antp, which marks in the PSC, and provides spatial signals for the development of the LG. In vertebrates, Hox genes have been shown to express within progenitor cells and are rapidly switched off during cell maturation. As our overexpression lines perturb both cell number and differentiation, it is possible that multiple Drosophila Hox genes are involved in finetuning the precise programme of Drosophila blood cell development as well.

The fact that the cells appear to be phenotypically confined to plasmatocytes and lamellocytes implies that expression of these genes works in tandem with, and above the specific programme of the cell types. It would be interesting to know which genes are being modulated in our overexpression lines, by profiling their transcription states as well as the binding sites of the individual Hox proteins. In the absence of this information, we speculate that Hox gene overexpression leads to the aberrant transcription of genes. It is known from previous studies that Hox dysregulation in leukaemia is usually concomitant with gain or loss of function mutations in upstream regulators, most commonly in Mixed Lineage Leukemia-1 (MLL-1) fusion proteins, or loss of function Enhancer of Zeste Homolog 2 (EZH2) mutations. It has been reported that EZH2 mutations have the lowest number of co-operating mutations to induce leukemogenesis.

Interestingly, we observe polycomb members Psc and Esc have a role in suppressing melanised pseu
don tumour formation. Both Psc and Esc mutants rescued the phenotype significantly which suggests that tumour suppressor genes may be their targets for repression. BMI1, a mammalian counterpart of Psc, was discovered as a proto-oncogene. Overexpression of BMI1 in mouse models has been shown to induce both types of leukaemia (lymphoid and myeloid origin). BMI1 cooperates with c-myc in the generation of lymphomas. MYC is a transcription factor that regulates many cellular processes, including proliferation and apoptosis. c-myc is an attractive target for chemotherapy as it has a role in multiple converging signalling cascades. In many cancers, MYC protein is overexpressed due to various processes like translocation, duplication or epigenetic misregulation. BMI1 regulates MYC protein expression by inhibiting its downstream target ink4a-ARF, a tumour suppressor gene. The role of EED, a mammalian homolog of Drosophila Esc, in leukaemia is not very well understood. Mutations in EED have been indicated to impair polycomb complex (PRC2) functionality and it is associated with myelodysplastic neoplasm.

Although Drosophila does not have ink4a homolog, it will be interesting to see the mechanism of regulation of melanotic tumor formation and over-proliferation of blood cells by Psc. Taken together, we
speculate that Hox gene activation in hemocytes causes cell-autonomous proliferation and differentiation and induces leukaemia via aberrant transcription.

In summary, *Drosophila* the Hox genes *Dfd, Ubx and particularly abd-A*, when expressed in blood cells, are leukemogenic. This causal link of Hox genes to the pseudo-tumor phenotype supports the non-homeotic role of *abd-A* as a growth promoter later during development. The disease phenotype is modified by select PcG/trxG members. This reinforces previous studies in vertebrates that report the misregulation of Hox genes in several cancers and implicate the role of epigenetic factors in them. Hox induced leukaemia in *Drosophila* offers advantages of the fly model to explore the biology of the process and develope novel potential markers and therapeutic options.

**Declarations**

-Ethics approval and consent to participate- Not applicable

-Consent for publication

All the authors have consented for publication of this work.

-Availability of data and material

Data and material are available on request.

-Competing interests

Authors declare no competing interests.

-Funding

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-Authors’ contributions

TP carried out experiments, contributed to the design, interpretation of the experiments and writing the manuscript, RS carried out experiments on epigenetic effects, and RKM conceived the project, wrote the manuscript and supervises the work.

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Abbreviations

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<th>Description</th>
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<tr>
<td>PH3</td>
<td>Phospho Histone 3 at Serine 10</td>
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<tr>
<td>LG</td>
<td>Lymph Gland</td>
</tr>
<tr>
<td>PcG</td>
<td>Polycomb Group of Proteins</td>
</tr>
<tr>
<td>TrxG</td>
<td>Trithorax Group of Proteins</td>
</tr>
<tr>
<td>PRC1</td>
<td>Polycomb Repressive Complex 1</td>
</tr>
<tr>
<td>PRC2</td>
<td>Polycomb Repressive Complex 2</td>
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57. Goto, A., Kadowaki, T. & Kitagawa, Y. Drosophila hemolectin gene is expressed in embryonic and larval hemocytes and its knock down causes bleeding defects11Two hml-GAL4 lines (w1118; P{w + mc = GAL4-Hml} 5-6 and w1118; P{w + mc = GAL4-Hml}6-4) and one homozygote of d-hml-GAL4, UAS-gfpnlacZ, UAS-gfp[S65T]/d-hml-GAL4, UAS-gfpnlacZ, UAS-gfp[S65T] (w1118; P{w + mc = GAL4-Hml} 6-4 P{w + mc = UAS-GFP::lacZ.nls} 15.1) transgenic lines are available from the Bloomington Stock Center with the stock numbers of 6395, 6396, and 6397, respectively. *Dev. Biol.* **264**, 582–591 (2003).


**Figures**
Larvae with subcutaneous tumors. Dfd, Ubx, abd-A and Abd-B, when expressed under the drivers Cg, He and Hml D3 lead to melanised bodies in the viscera.
Tumor phenotype in larvae and pupal lethality. A) The size and penetrance of melanised bodies was maximum when expressed under Cg, while tumors do manifest when He and Hml are used, they are much rarer and smaller. Expression under Lsp2-Gal4 does not lead to the formation of such bodies. B) Percentage of pupal lethality, indicated by larvae that fail to eclose. When Cg-Gal4 drives the genes Dfd, Ubx, abd-A and Abd-B do cause lethality, so does expressing them in the fatbody under Lsp2-Gal4. Driving
these genes in the blood cells (He-Gal4 and HmlD3-Gal4) leads to a much lower penetrance of this phenotype.

Figure 3

Cell proliferation and quantification of hemocytes. A) Anti-PH3 staining for comparative study of cell proliferation with over-expression of abd- A gene driven by Cg-Gal4, He-Gal4, HmlD3-Gal4 and Lsp2-Gal4. Phalloidin iFluor-488 was used to stain actin filaments. B) The number of circulating hemocytes
increases significantly when Dfd is over expressed using Cg-Gal4 (p=0.0003), He-Gal4 (p<0.0001) and HmlD3-Gal4 (p<0.0001), Ubx when over expressed with Lsp2-Gal4 (p<0.0001) and HmlD3-Gal4 (p=0.0011), abd-A when over expressed with Cg-Gal4 (p<0.0001), Lsp2-Gal4 (p=0.0028), He-Gal4 (p=0.0018) and HmlD3-Gal4 (p=0.0002), Abd-B when over expressed with Cg-Gal4 (p<0.0001). B) PhosphoHistone3+ nuclei appear when these genes are expressed in the blood cells, indicating that some of the increase in cell number maybe due to cell autonomous proliferation.

**Figure 4A**

Control | UAS-abd-A | UAS-Abd-B
---|---|---
Myospheroid | Merged (+DIC) | Myospheroid | Merged (+DIC) | Myospheroid | Merged (+DIC)
Cg-Gal4
He-Gal4
HmlD3-Gal4
Lsp2-Gal4

**Figure 4B**

![Graph showing percentage lamellocytes](image)

**Figure 4**
Myospheroid staining and Quantification of lamellocytes. A) When Dfd, Ubx, abd-A, but not Abd-B are driven in blood cells (under Cg-Gal4, He-Gal4 or HmlD3-Gal4), but not in the fatbody (Lsp2-Gal4), large, dorsoventrally flattened cells begin to appear in circulation. These stain positive for mys. Some circulating plasmatocytes also appear to mys+. This indicates that they might be in the process of differentiating. B) Percentage of lamellocyte is plotted on Y-axis. Ubx over-expression with Cg-Gal4 (p<0.0001) and He-Gal4 (p=0.0455) has a significant increase while Lsp2-Gal4 and HmlD3-Gal4 do not show any significant increase. abd-A with Cg-Gal4 and He-Gal4 has comparatively high percent of increase to control as well as Ubx over-expressed.
Effect of PcG and trxG mutations on the abd-A induced tumor phenotype. A) Effect of PcG and trxG on subcutaneous melanotic tumor formation. Over-expression of abd-A gene is driven by Cg-Gal4. Most of PcG and trxG mutants have effect on the phenotype. Polycomb mutants, esc2 (PRC2 member) and Psc1 (PRC1 member) show a decrease in severity of phenotype (size and numbers of melanotic body). Su(z)123 and Pc1 have increased severity. Brm2, TrlR85, Su(z)21.a1 and E(z)731 do not show any
change in severity compare to Cg-Gal4 driven abd-A over-expressed individual. B) Comparative quantification of melanotic tumor formation phenotype in abd-A over-expressed (driven by Cg-Gal4) individual in different PcG and trxG background. Percentage of tumor showing individuals is plotted on Y-axis (no. of animal screened is >80 in each case, error bars represent the standard error). Phenotype is rescued in esc2 (p<0.0001), Psc1(p<0.0001), and TrlR85 (p= 0.0223) mutants while Pc1 (p=0.0109), Su(z)123 (p= 0.0028) and Brm2 (p=0.0088) have a significant increase in melanotic tumor formation. C) Percent pupal lethality in PcG and trxG mutant background. Psc1 (p< 0.0001), esc2 (p<0.0001), brm2 (p=0.0042) and TrlR85 (p<0.0001) show decrease in pupal lethality.

**Supplementary Files**

This is a list of supplementary files associated with this preprint. Click to download.

- SupplementaryFigure1AD.pdf
- SupplementaryTablesS110.pdf