The Drosophila Perlecan gene trol regulates multiple signaling pathways in different developmental contexts

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Abstract

Background: Heparan sulfate proteoglycans modulate signaling by a variety of growth factors. The mammalian proteoglycan Perlecan binds and regulates signaling by Sonic Hedgehog, Fibroblast Growth Factors (FGFs), Vascular Endothelial Growth Factor (VEGF) and Platelet Derived Growth Factor (PDGF), among others, in contexts ranging from angiogenesis and cardiovascular development to cancer progression. The Drosophila Perlecan homolog trol has been shown to regulate the activity of Hedgehog and Branchless (an FGF homolog) to control the onset of stem cell proliferation in the developing brain during first instar. Here we extend analysis of trol mutant phenotypes to show that trol is required for a variety of developmental events and modulates signaling by multiple growth factors in different situations.

Results: Different mutations in trol allow developmental progression to varying extents, suggesting that trol is involved in multiple cell-fate and patterning decisions. Analysis of the initiation of neuroblast proliferation at second instar demonstrated that trol regulates this event by modulating signaling by Hedgehog and Branchless, as it does during first instar. Trol protein is distributed over the surface of the larval brain, near the regulated neuroblasts that reside on the cortical surface. Mutations in trol also decrease the number of circulating plasmatocytes. This is likely to be due to decreased expression of pointed, the response gene for VEGF/PDGF signaling that is required for plasmatocyte proliferation. Trol is found on plasmatocytes, where it could regulate VEGF/PDGF signaling. Finally, we show that in second instar brains but not third instar brain lobes and eye discs, mutations in trol affect signaling by Decapentaplegic (a Transforming Growth Factor family member), Wingless (a Wnt growth factor) and Hedgehog.

Conclusion: These studies extend the known functions of the Drosophila Perlecan homolog trol in both developmental and signaling contexts. These studies also highlight the fact that Trol function is not dedicated to a single molecular mechanism, but is capable of regulating different growth factor pathways depending on the cell-type and event underway.
Background

Heparan sulfate proteoglycans (HSPGs) are a family of cell-surface and extracellular proteins modified by the attachment of glycosaminoglycan chains. The general structure of the protein core determines the family the HSPG belongs to; Syndecans contain a transmembrane domain, Glypicans are tethered to the cell surface via a GPI linkage and Perlecans are secreted components of the extracellular matrix. Both the protein core and glycan chains play important roles in HSPG function through protein-protein and sugar-protein interactions. Genetic studies, first in Drosophila and later in mouse and zebrafish, demonstrated the importance of the heparan sulfate chains on all three types of HSPGs for signaling by multiple growth factors such as the Fibroblast Growth Factors (FGFs), Hedgehogs, Wnts and Transforming Growth Factors (TGFβs) (reviewed in [1]).

Perlecan is the largest member of the HSPG family with a core protein of approximately 450kD in size. Perlecan has been linked to signaling by the heparan-dependent growth factors FGF2, Vascular Endothelial Growth Factor (VEGF) and Sonic Hedgehog (SHH) in mammalian systems (reviewed in [2,3]). Studies of Perlecan knock-out mice have demonstrated roles for Perlecan in vascular development and chondrogenesis as well as maintenance of basement membrane integrity [4-7]. Additional mammalian studies have revealed Perlecan’s functions in angiogenesis and carcinogenesis ([8-11], reviewed in [2,12]). Mutation of Perlecan in humans leads to the muscle tone syndrome, presumably through effects on chondrogenesis ([13-15].

Studies of Perlecan in invertebrate model systems have led to additional insights into Perlecan function. The single Perlecan gene in C. elegans is encoded by the unc-52 locus [16]. Mutations in unc-52 result in embryonic or adult paralysis due to defects in body wall muscle cells ([16,17], reviewed in [18]). Mutations in unc-52 also enhance cell migration defects caused by decreased netrin, FGF, TGFβ or Wnt signaling. In Drosophila, Perlecan is encoded by the trol gene on the X chromosome [19,20], which was initially implicated in the control of stem cell division in the developing larval brain [21,22]. In the larval brain, trol promotes the cell cycle progression of mitotically arrested neuroblasts [23,24] through modulation of FGF and Hedgehog signaling [19]. These Drosophila studies were the first to link Perlecan to Hedgehog signaling. More recently, studies of oogenesis in Drosophila have uncovered a role for Perlecan in the maintenance of epithelial cell polarity through interactions with the extracellular matrix receptor Dystroglycan [25].

The many signaling pathways associated with HSPGs in general and Perlecan in particular led us to ask what other biological processes may require Perlecan function. We used a series of trol mutants to investigate several phenotypes ranging from overall developmental progress to specific alterations of stem cell division and hemocyte production. Furthermore, analysis of signaling pathway response genes revealed that while mutations in Perlecan decrease signaling in multiple pathways, at least some of these effects are tissue specific.

Results and discussion

Development and lethal phase

We had previously shown that the viable trol22 and the lethal trolb, trol1, and trol4 alleles form an allelic series of increasing severity based on their onset of neuroblast proliferation phenotype in first instar larval brain lobes [24]. Identification [19] and phenotypic analysis of a fifth trol allele, trol7, revealed that trol7 is the strongest allele with respect to the first instar proliferation phenotype (Fig. 1A). Unexpectedly, trol7 mutant larvae appeared healthier overall than other trol mutant larvae, suggesting that the order of allelic severity determined by analysis of first instar brain lobes would be different from one based on developmental progression. To test this hypothesis, we examined the lethal stage and developmental progression of larvae mutant for trol22, trolb, trol1, trol4 and trol7. In all the experiments, crosses were designed to use sibling controls in order to minimize the effects of genetic background, which can be significant in fly stocks kept in reproductive isolation from each other for years in our laboratory. For the lethal trol alleles, y trol /Binsn stocks were used as the source of mutant and control larvae. At this stage of first instar, mutations in y produce one of the few reliable phenotypic markers. Thus trol mutant animals were identified as y mutant larvae that are y trolb hemizygous males and sibling controls were a mixed population of y’ animals: y trolb/Binsn heterozygotes, Binsn homozygous females and Binsn hemizygous males. Note that while Binsn homozygous females and hemizygous males can become viable adults, not all Binsn/Binsn or Binsn/Y larvae reach adulthood. Thus our comparison provides a measure of developmental progression and lethal phase that will err on the side of minimizing the trol mutant phenotype. For analysis of the viable trol22 allele, additional crosses were required to produce wild-type sibling controls from the homozygous y trol22 stock. y trol22 animals were crossed to the wild-type strain Canton Special (CS) to produce trol22/CS heterozygous females. These females were mated to CS males to generate hemizygous y trol22 male larvae and y’ sibling control larvae (a mixture of heterozygous y trol22/CS female, homozygous CS female and hemizygous CS male) for the developmental studies. One hundred mutant and sibling control animals for each allele were collected at early first
instar and monitored at 24 hour intervals for developmental progression and viability. Of these, only 1 mutant trol4 and no trol4 animals pupariated. However, when the same numbers of trolB22, trol8 and trol7 mutant larvae were analyzed and compared to sibling controls, 102%, 38% and 23% of the animals were able to pupariate, respectively (Fig. 1B). The pupariation assay resulted in shifts of perceived functional severity for both trol4 and trol7, with trol4 appearing stronger and trol7 appearing weaker.

Why would animals mutant for trol7 (that has a strong effect on neuroblast proliferation) be able to progress further in development than animals mutant for trol4 which causes a weaker neuroblast proliferation phenotype? One possibility is that trol modulates the activity of different signaling pathways in different tissues. For example, a mutation that affects the ability of Trol to function in the Hh pathway would have a severe effect on developmental signaling pathways in different tissues. For example, setting controls to a value of 4 to control for genetic background effects between experiments. Setting controls to a value of 4 on our 5 point scale was chosen to allow evaluation of over-proliferation (>4) as well as under-proliferation (<4) mutant phenotypes. To obtain the TNb phenotype score for each mutant allele we normalized the score for each sample to the respective sibling control and calculated the average and standard error of the mean (Table 1). Surprisingly, trolB22 mutants had a significantly higher than normal level of TNb proliferation (TNb score >4) while the remaining trol mutants showed decreased TNb cell division compared to controls. The differences between mutant and control BrdU incorporation were statistically significant (p < 0.05) for each mutant allele. Comparison between mutants showed a phenotypic trend from trolB22 having hyperactive TNb proliferation to trolB4 as the mutant with the fewest labeled TNbs. In this assay trol7 mutants appear to have a weaker phenotype than trolB4.

**Effects of trol mutations on TNb proliferation**

**trol** was initially identified as a mutation on the X chromosome that affected the proliferation pattern of neuroblasts in the brain lobes and ventral ganglion [21,22]. Since neuroblasts in the thoracic region of the ventral ganglion begin proliferation in early second instar [21,26,27], we evaluated the ability of thoracic neuroblasts (TNbs) to enter S phase in trol mutant animals. We adapted the idea of phenotypic classes to produce a scale for the extent of TNb proliferation at four hours post molt (Fig. 1C–G). Five TNb classes were defined as follows: Class 1, no neuroblasts labeled; Class 2, a small number of labeled neuroblasts with no distinct segmental pattern; Class 3, labeled neuroblasts in a segmentally repeated lines with very few labeled neuroblasts in between the lines or in the medial region of the ventral ganglion; Class 4, labeled neuroblasts in a segmentally repeated line with some labeled neuroblasts in between the lines or in the medial region of the ventral ganglion; and Class 5, labeled neuroblasts in heavily populated segmental pattern with many labeled neuroblasts in the medial portion of the ventral ganglion. When both sides of a ventral ganglion did not conform to a single class, the sample was scored as the higher class. This will have a conservative effect of scoring a partial loss-of-proliferation TNb phenotype as more wild-type. Thus we can have greater confidence in the significance of TNb proliferation phenotypes observed compared to controls.

We first examined the onset of TNb proliferation in wild-type sibling controls to determine the time point at which to assay the trol mutants (data not shown). In our hands, high levels of 5-Bromodeoxyuridine (BrdU)-labeled TNbs were first observed in control samples between 2–5 hours post molt depending on genetic background. This timing is slightly earlier than the previous observation that TNb mitosis begins between 28–34 hours post hatching, or 4–10 hours post molt (pm) to second instar [26]. To evaluate the TNb proliferation phenotype produced by the different trol alleles, at least twenty samples for each mutant and sibling control (generated as described above) were allowed to incorporate BrdU from 4–5 hours pm and scored for TNb class. The average score and standard error of the mean were calculated for each group of sibling controls. The control value for each study was set to a value of 102%, 38%, and 23% of the animals able to pupariate, respectively (Fig. 1B). The pupariation assay resulted in shifts of perceived functional severity for both trol4 and trol7, with trol4 appearing stronger and trol7 appearing weaker.

To determine if trol affects TNb proliferation through modulation of Bnl and Hh signaling, we used genetic interaction studies with the weak trol allele trolB22. As we have shown, trolB22 animals have over proliferation of TNbs compared to sibling controls (Table 1). For the genetic interaction assay, y trolB22 females were crossed to bmB6916/TM3y+ males to generate y larvae that were y trolB22 ; bmB6916/+ and y+ siblings that were a combination of y trolB22 ; +/TM3y+ males, y trolB22/+ ; +/TM3y+ females and trolB22/+ ; bmB6916/+ females. None of the sibling controls had TNb proliferation scores outside of the normal (CS) range at this timepoint. y trolB22 males carrying a single copy of the bmB6916 allele had fewer BrdU labeled TNbs at 2–3 hours post molt to second instar (pm) compared to siblings that were hemizygous or heterozygous for trolB22 alone or heterozygous for both trolB22 and bmB6916 (Fig. 2). The decreased TNb proliferation in samples heterozygous for bmB6916 in a trolB22 background compared to controls versus the increased proliferation in trolB22 animals wild-type for bnl compared to controls sug-
Figure 1
Phenotypic series of trol alleles. A) First instar neuroblast proliferation phenotype presented as % of samples with numbers of BrdU labeled neuroblasts falling within the control range, some data originally published in Park et al, 2003a. B) Lethal phase phenotype presented as the percentage of trol mutant animals capable of survival and development to third instar (grey bars) or to pupal formation (black bars) compared to sibling controls. Error bars indicate s.e.m. C) Cartoon of second instar larval brain with dividing TNBs in ventral ganglion. Boxed area indicates portion of brain shown in panels D-G below. A = anterior, P = posterior. D-G) Examples of the five classes of BrdU incorporation into TNBs are shown. In all panels anterior is to the left, posterior is to the right. Scale bar in panel D indicates 25 um. D) None (class 1). E) Few (class 2). F) Segmentally repeated lines with few extra neuroblasts (class 3). G) Segmentally repeated lines with several scattered neuroblasts (class 4). H) Heavily populated segmental pattern (class 5).
gests that the \textit{trol}^{22} mutation affects signaling by Bnl in the ventral ganglion at second instar.

We also used genetic interactions to evaluate the possibility that \textit{trol} might affect Hedgehog signaling in the ventral ganglion. For this study \textit{y \textit{trol}^{22}} females were crossed to \textit{hhAC/TM3\textit{y}} males to generate \textit{y \textit{trol}^{22} ; hhAC/+} and \textit{y \textit{trol}^{22} /+ ; hhAC/+} sibling controls that were a combination of \textit{y \textit{trol}^{22} ; +/TM3\textit{y}} males, \textit{y \textit{trol}^{22} /+ ; +/TM3\textit{y}} females and \textit{trol^{22} /+ ; hhAC/+} females. \textit{trol^{22}} animals carried a single copy of the \textit{hhAC} allele, also had fewer dividing TNbs at 2–3 hours pm compared to sibling controls (Fig. 2). The decrease in the number of BrdU labeled TNbs in \textit{trol} hemizygotes upon heterozygosity for \textit{hhAC} suggest that mutations in \textit{trol} also weaken the signaling action of Hh in the ventral ganglion. To further test our hypotheses, we examined the signaling activity of Bnl and Hh in \textit{trol} mutants directly by quantitative RealTime PCR (qRT-PCR) in the central nervous system (CNS). To avoid interfering signals from the lobes of the second instar brain that might overwhelm differences in signal in the ventral ganglion, we isolated ventral ganglia from second instar \textit{trol} mutant and sibling control brains at one hour post molt. First instar brains were dissected at 20 hours post hatching which correlates with the end of the BrdU labeling period used to assess neuroblast proliferation in first instar ([24] and this manuscript). RNA was isolated, cDNA synthesized and amplified and the level of expression of the Hh response gene \textit{ptc} (Fig. 3) and the Bnl response gene \textit{pnt} (Fig. 3) assayed. Our qRT-PCR data demonstrate that mutations in \textit{trol} affect the strength of signaling by both Hh and Bnl in the larval ventral ganglion and in first instar larval brains (data not shown).

\textbf{Trol localization in the larval brain}

Previously we had isolated complexes containing either Trol and FGF2 or Trol and Hh by co-immunoprecipitation [19]. In combination with our genetic studies, these complexes suggested that the Trol protein regulates neuroblast division by binding growth factors that stimulate neuroblast proliferation in a manner similar to Perlecan-mediated promotion of ligand-receptor binding described in mammalian systems [28]. This model predicts that Trol protein should be localized near the regulated neuroblasts, i.e. the optic lobe and central brain neuroblasts of

\begin{table}
\centering
\begin{tabular}{|l|c|c|}
\hline
\textbf{\textit{trol} allele} & \textbf{TNb score*} & \textbf{S.E.M.} \\
\hline
Control & 4.00 & 0.27 \\
\textit{trol}^{22} & 4.55 & 0.16 \\
\textit{trol}^{R} & 3.37 & 0.21 \\
\textit{trol}^{H} & 3.00 & 0.27 \\
\textit{trol}^{F} & 2.95 & 0.2 \\
\textit{trol}^{D} & 2.45 & 0.21 \\
\hline
\end{tabular}
\caption{TNb BrdU incorporation phenotype of \textit{trol} mutants at 4–5 hours pm.}
\end{table}
the first instar brain [19-24,29] and the thoracic region of
the second instar brain ([22] and Fig. 4A). In contrast,
in situ hybridization studies in the third instar larval brain by
Voigt et al [20] had revealed that only a few isolated cells
at a distance from the optic lobe proliferation centers
express trol. This led the authors to suggest that Trol is
unlikely to regulate neuroblast proliferation by promot-
ing binding of FGF-type ligands to their receptors since
this would require Trol protein localization near the
responding cells. However, since Trol is a secreted protein
with a long half-life, mRNA expression patterns may not
accurately portray protein localization. In addition, Voigt
et al conducted their in situ analysis at late third instar, 2–
3 days after the activation of neuroblast division at late
first or early second instar. Thus the expression pattern
observed for trol message at late third instar may not
reflect expression of trol at earlier larval stages. Furth-
more, a study of trol mRNA localization by in situ hybrid-
ization in embryos showed either no obvious staining in
the CNS [30] or expression in a small subset of glial cells
in the CNS [20]. However, analysis of Trol protein locali-
sation with an anti-Trol antibody in embryos revealed
localization to the basement membrane of the CNS [30].
This evidence further suggests that trol message patterns
may not reflect Trol protein localization. To address the
conflicting models, we took advantage of a Trol protein
trap in which the GFP gene is inserted within the endog-
enous trol locus [31]. Analysis of GFP localization in larval
brains demonstrates that Trol-GFP is found in a layer, pre-
sumably the basal lamina, encompassing the entire outer
surface of the larval brain with little to no signal detecta-
bale at internal sites within the brain (Fig. 4B–E). Trol-GFP
was also observed in the basal lamina surrounding nerves
emanating from the larval brain. The distribution of Trol
over the entire brain was further verified by immunohis-
tochemistry using an anti-Trol antibody (Fig. 4F). This
localization of the Trol protein is consistent with the
model that Trol binds Bnl and Hh and facilitates their sig-
naling to promote neuroblast proliferation, as the regu-
lated neuroblasts are found at the surface of the cellular
cortex in both the brain lobes and the ventral ganglion
[21,26,32]. To determine if the localization of Trol-GFP to
the basal lamina was unique to the larval brain, we exam-
nined Trol-GFP in the salivary glands. As in our larval brain
studies, Trol-GFP is found on the surface of the gland, pre-
sumably as a component of the basal lamina (Fig. 4G).

Figure 3
Hh and Bnl signaling activity in the ventral ganglion of trol mutant animals. Quantitative RT-PCR analysis of the
expression of the Hh response gene ptc (black bars) and the Bnl response gene pnt (grey bars) in the ventral ganglia of trol
mutant normalized to controls at one hour post molt to second instar. β-actin was used to as an internal control to normalize
message levels. All analyses were done in triplicate and three different concentrations to ensure samples were within linear
range of amplification. Error bars indicate standard deviation.
Effects of trol mutations on larval hemocyte number

A second system where we thought trol might have an effect on development is the production of hemocytes during larval life. A number of studies have elegantly shown that the larval lymph gland is the source of larval hemocytes [33]. In the primary lobe of the third instar lymph gland prohemocytes arise in the medullary zone while maturing hemocytes are found in the adjacent cortical zone. Hemocytes are then released into the hemolymph and are present as three types of circulating cells: plasmatocytes (95%), lamellocytes (1–5%) and crystal cells (rare). Each cell-type has characteristic morphology and can be easily identified under a compound microscope. Mature circulating larval hemocytes are still undergoing cell division, albeit at a low rate, as shown by staining of hemocytes with phosphohistone H3, an M phase marker [34,35]. Expression of an activated Ras (Rasv12) in circulating hemocytes increases the percentage of circulating hemocytes that stain for phosphohistone H3 and results in a 40-fold increase in the number of hemocytes through activation of the Ras-MAPK pathway [34]. The Ras-MAPK pathway is activated by Vascular Endothelial Growth Factor (VEGF) and Platelet Derived Growth Factor (PDGF) among others. Signaling by mammalian homologs of both growth factors has been linked to mammalian Perlecan [3]. Furthermore, studies of PDGF/VEGF receptor (PVR) in Drosophila revealed that PVR is expressed in hemocytes and that decreased PVR function leads to increased hemocyte cell death [33]. Thus it seemed likely that mutations in trol could decrease PDGF/VEGF signaling in circulating plasmatocytes, resulting in decreased numbers of circulating plasmatocytes in trol mutants. To address this hypothesis, we determined the relative number of circulating plasmatocytes in third instar trolb22 or trol7 and sibling control larvae (Fig. 5A). Our analysis demonstrates a significant (p < 0.05) drop in the number of plasmatocytes in trol mutant versus sibling control larvae.

Trol localization and function in hemocytes

The decrease in the number of circulating plasmatocytes in trol mutants versus controls suggested that trol might indeed function to promote Ras-MAPK signaling by PDGF/VEGF in circulating plasmatocytes. This predicts that Trol protein would be localized on these plasmatocytes. We used Trol-GFP protein trap to examine the plasmatocytes for the presence or absence of Trol protein. Fluorescence microscopy revealed that Trol-GFP is indeed found on circulating plasmatocytes in third instar larvae (Fig. 5B,C), but not in the lymph gland (data not shown).

This result is consistent with the requirement for Ras-MAPK activation in plasmatocytes for plasmatocyte proliferation and for PVR in plasmatocytes to avert apoptosis, and supports the hypothesis that Trol modulates PVR-Ras-MAPK signaling in plasmatocytes. The ETS-transcription factor pnt is a MAPK-response gene and will drive plasmatocyte proliferation [36]. Therefore we asked if trol mutant plasmatocytes show decreased levels of pnt compared to controls. Plasmatocytes were collected by bleeding third instar trolb22 and trol7 mutant larvae and sibling controls, RNA was extracted and amplified, and subjected to qRT-
Mutations in *trol* decrease circulating plasmatocyte number and *pnt* expression. A) Quantification of circulating plasmatocytes in *tro*\[^b22\] and *tro*\[^7\] mutants compared to controls. Each sample consisted of hemolymph pooled from three third instar larvae. Five squares were counted for each sample. Each genotype was analyzed in triplicate. B) Brightfield image of plasmatocytes from Trol-GFP stock. C) Fluorescence image of plasmatocytes from Trol-GFP stock demonstrating presence of Trol on plasmatocytes. D) Expression of the VEGF/PDGF response gene *pnt* in *tro*\[^b22\] and *tro*\[^7\] mutant hemocytes compared to sibling controls by qRT-PCR. Samples of hemolymph from three third instar larvae of each genotype were pooled, RNA extracted, amplified and analyzed. All reactions were carried out in triplicate at three different template concentrations to ensure amplification was in the linear range. *β*-actin was used as an internal normalization control.
Dpp, Wg and Hh signaling are affected in trol mutants in second instar brains but not in third instar brain lobes/eye discs. qRT-PCR analysis of A) the expression levels of dpp, its response gene salm, wg and its response gene slp in the ventral ganglia of second instar larvae. Data for the Hh response gene ptc is shown in Fig. 3. B) Expression levels of dpp, its response gene salm, wg and its response gene slp, hh and its response gene ptc in trolsd mutant third instar brain lobes/eye discs. C) Expression of dpp and salm in hsGAL4/+; +/(EP)dad second instar ventral ganglia and hsGAL4 controls. D) Expression of wg and slp in hsGAL4/+; +/(EP)sgg second instar ventral ganglia and hsGAL4 controls. In all panels, error bars indicate standard error. All reactions were carried out in triplicate at three different template concentrations to ensure amplification was in the linear range. β-actin was used as an internal normalization control.
PCR analysis. qRT-PCR studies demonstrated that plasmatocytes isolated from either trol^{B22} or trol^{7} mutants show decreased expression of pnt compared to controls, further evidence that trol modulates Ras-MAPK signaling in plasmatocytes (Fig. 5D).

**Trol and other growth factor signaling pathways**

Two other growth factor signaling pathways that have been linked to HSPGs are the wingless (wg/Wnt) and decapentaplegic (dpp/TGFβ) signaling pathways. Both of these pathways are active in the developing Drosophila eye disc and/or third instar brain along with Hh and Ras-MAPK signaling [37,38]. To ask if Trol might modulate the Dpp and Wg pathways we evaluated the expression of dpp and wg and their target genes splt major (salm, [39]) and sloppy paired (slp, [40]), respectively, in second instar ventral ganglia and third instar brains and eye discs from trol mutant larvae by qRT-PCR (Fig. 6A–B). We also assayed expres-
sion of hh and its response gene ptc in third instar brains and eye discs. In the trol^{B22} second instar ventral ganglion we observed a significant drop in the level of both dpp and wg compared to controls. The trol^{B22} mutation also resulted in diminished signaling efficiency by both growth factors as indicated by a larger drop in the level of their response genes salm and slp compared to the ligands themselves. In contrast, the trol^{4d} mutation decreased only dpp expression, but the efficiency of both dpp and wg signal-
ing was impaired. Thus in the second instar ventral gan-
glion, wild-type function of trol appears to be required for normal signaling by hh, bnl, dpp and wg (Figs. 3 and 6A). The decreased expression of dpp and wg in trol^{B22} mutants and of dpp in trol^{4d} expression may be due to secondary effects on dpp and wg expression caused by the changes in Hh and Bnl signaling in trol mutants. Alternatively, decreased expression of dpp and wg could be due to positive feedback between Dpp signaling and dpp expression and Wg signaling and wg expression, respectively. To test the latter possibility, we blocked Dpp signaling by over-expression of daughters against dpp (dad) [41], and assayed for dpp message levels (Fig. 6C). (EP)dad females were crossed to hsGAL4 males to drive expression of dad. Embryogenesis and first instar larval development were carried out at 18°C to limit expression of dad and inhibition of Dpp signaling at early stages. Upon molt to second instar, larvae were moved to 25°C for one hour to induce expression of dad. Larval brains were dissected and the ventral ganglia harvested for RNA isolations. Inhibition of Dpp signaling was confirmed by analysis of salm mRNA levels. Similarly, we inhibited Wg signaling by over-expression of shaggy (sgg), and assayed for wg message levels (Fig. 6D). Decreased Wg signaling was verified by analysis of slp expression levels. As shown by our qRT-PCR analysis, inhibition of Dpp signaling by over-expression of dad resulted in a drop in expression of the dpp ligand itself. Inhibition of Wg signaling by over-expression of sgg also produced a drop in the expression of wg. As these studies were conducted in flies wild-type for trol, they eliminate the possibility that the decreased expression of dpp and wg in trol mutants was due solely to reduced Trol-mediated signaling by Hh and/or Bnl. These data indicate the presence of a positive feedback loop for Dpp and Wg in the ventral ganglion.

To determine if Trol is necessary for growth factor signaling in other tissues at other stages we assayed for dpp, wg and hh expression and activity in trol^{B22} and trol^{4d} third instar brain lobes and eye discs. No significant changes in either growth factor expression or signaling were observed in trol^{B22} samples (data not shown). In trol^{4d} samples, expression of all three growth factors decreased by 65–85%, as did the expression of their response genes (Fig. 6B). The sole exception is wg/slp, where wg expression decreased about 65% and slp expression decreased only about 50%. These data indicate that mutations in trol do not dramatically decrease the signaling efficiency of Dpp, Wg or Hh in third instar brain lobes and eye discs, unlike the effect of those same trol mutations in second instar.

**Conclusion**

**trol and Drosophila development**

We have previously demonstrated that mutations in trol prevent the onset of neuroblast division in the first instar brain and that most trol mutations are lethal. Mutations in a second gene, anachronism, also affect the onset of neuroblast proliferation but in the opposite manner: in anachronism mutants, mitotically regulated neuroblasts begin cell division too early [42]. However, when a lethal trol mutation was combined with a viable allele of anachronism, the lack of neuroblast division was rescued (double mutants exhibited the anachronism phenotype of premature neuroblast division) but lethality was not [21]. This outcome suggested that trol function is required for other developmental events necessary for survival. Further analyses revealed that trol modulates Hh and Bnl signaling in the first instar brain [19]. Here we have demonstrated that trol function is required for developmental progression to third instar and for pupariation. Analogous to its function in the first instar brain, trol is required to initiate the division of a second, independent and spatially distinct population of neuroblasts in the second instar brain (Table 1, Fig. 1). This initiation of division is also dependent on Bnl and Hh signaling (Fig. 2). We have also demonstrated that the Trol protein is localized to the surface of the brain at all larval stages, which places it in close proximity to the regulated neuroblasts. This localization is consistent with our model where Trol regulates Bnl and Hh signaling to cells adjacent to the regulated neuroblasts by binding the growth factors directly [19]. Trol protein localization to the basal lamina is not limited to the larval brain, as Trol-GFP studies also showed Trol protein in the basal lamina.
surrounding the salivary glands (Fig. 4G). trol function is not limited to the nervous system, as mutations in trol also diminish the number of circulating plasmatocytes by decreasing expression of pnt, a PVR response gene in plasmatocytes (Fig. 5). We speculate that trol may be necessary for signaling by the Drosophila PDGF and/or VEGF growth factor, just as mammalian Perlecan has been shown to function during angiogenesis [3]. Our studies of Dpp and Wg indicate a positive feedback between dpp expression and Dpp signaling and wg expression and Wg signaling in the second instar ventral ganglion. Signaling by Dpp and Wg is also dependent on trol in the second instar brain, but not (or very little) in the third instar brain lobes and eye discs (Fig. 6), despite the fact that Dpp and Wg signaling are taking place in those tissues. In fact, even Hh signaling appears to be independent of trol in this context. These results highlight an important concept in trol, and indeed, in proteoglycan function: that the Trol protein will be used at different times and places to regulate the signaling of different growth factors. Deciphering the role of trol in different developmental decisions will require that we examine each event individually, as trol will not necessarily mediate the same molecular mechanism each time.

Involvement of HSPGs in growth factor signaling

The requirement for heparan sulfate proteoglycans in signaling by different families of growth factors is well established [43], but what is not yet clear is why different organs and tissue types use different HSPGs to modulate these signaling pathways. One possibility is that the specific mechanism(s) through which these molecules modulate signaling activity allows for site-specific variations in the regulation of signaling activity. HSPGs with varied amino acid sequence can act in the same signaling pathway, such as Syndecan-4 and Perlecan for FGF2 [28,44] or Glypicans, Syndecan-3 and Perlecan for Hh [19,43,45]. Mutations that affect heparan sulfate synthesis or modification strongly affect FGF2 and Hh signaling [43]. Furthermore, Perlecan isolated from various endothelial cell sources has different binding affinities for FGF2 [46]. These data initially suggested that the protein core of the HSPG might have little to do with signaling specificity and that the main functional domain of HSPGs is concentrated in the sequence of the heparan sulfate chains.

The carbohydrate-centric view is being challenged by studies that indicate a role for the protein-protein interactions of HSPGs with growth factors and other signaling molecules. For example, expression of chimeric molecules has shown that the cytoplasmic tail of Syndecan is specifically required for FGF2 signaling in addition to its heparan sulfate chains [47]. Perlecan protein-protein interactions include the ability of Perlecan to bind growth factors and extracellular matrix molecules at various sites on its protein core. Further mechanisms that allow for differential regulation include processing of HSPGs. These studies suggest a reason for the use of a particular HSPG during an individual developmental decision – the flexibility of combining both carbohydrate-based regulation and protein-based regulation of cell-cell signaling may make a specific HSPG uniquely suited for a given situation.

In the context of combined carbohydrate and protein inputs into HSPG function, it becomes clear that a given HSPG may be expressed and function in very specific contexts that take advantage of its unique regulatory abilities. It is interesting to note that we have connected Perlecan with FGF and Hh signaling in the developing fly brain while mouse studies have shown that Perlecan knock-out mice have cerebral cortex abnormalities [6,19,21]. trol mutant larvae have decreased numbers of circulating hemocytes that are likely due to decreased Ras-MAPK signaling by VEGF/PDGF. Perlecan knockout mice also have defects in chondrogenesis and cardiovascular development and mammalian studies have demonstrated a role for Perlecan in angiogenesis driven by FGFs, VEGF and PDGF [3]. Finally, we have shown that Perlecan is required for SHH signaling during human prostate cancer growth [8], which reveals a new system for the investigation of the mechanism of Perlecan action. Further analysis of the ability of HSPGs to substitute for each other in cell fate decisions and the means by which they individually regulate cell-cell communication will lead to a clearer understanding of the inputs necessary for cells to carry out a developmental or disease progression.

Methods

Fly stocks

Stocks of the viable trolR22 allele and the lethal trol4, trol7, trol8 and trol24 alleles have been described previously [19,21,22,29]. All trol mutant stocks with the exception of trolR22 are y trolw/Binsn where the chromosome carrying the trol mutation is marked with y to facilitate identification of y trol mutant versus y control larvae. The trol-GFP protein trap was obtained from Dr. Stephane Noselli. The bnlB6916 and hhAC stocks were obtained from the Bloomington stock center and used to construct y; bnlB6916/TM3y and y; hhAC/TM3y stocks for genetic studies.

Lethal phase

Early first instar larvae were collected and placed on apple juice plates with yeast. Each plate initially had 50 mutant or control animals per plate, segregated to prevent competition between mutant and wildtype siblings. Two plates of each genotype were examined. The number and stage of larvae still present on each plate were assayed every 24 hours and the survivors transferred to a fresh plate. Since none of the trol mutants with the exception of trolR22 pro-
duce viable adults, individual animals were followed only until pupariation.

**Developmental staging**

Developmental synchronization was carried out as previously described [19,21,23,48]. Flies were allowed to lay eggs on apple juice agar plates with fresh yeast overnight or for about 24 hours. For staging of synchronized first instar larvae, the plate was first cleared of any larvae and newly hatched larvae collected in one hour windows and placed on new apple juice plates with yeast at 25°C for aging. For staging of second instar larvae, late first instar larvae were placed on fresh apple juice plates with yeast. Newly molted second instars were collected in one hour windows and placed on apple juice plates with yeast at 25°C for aging or dissected immediately.

**Proliferation assay**

BrdU assays were carried out as previously described [19,21,23,48]. Briefly, animals were fed BrdU-containing artificial medium for one hour, dissected in PBST and fixed with Histochoice (Amresco) for 10 minutes. Brain samples were denatured in PBST-HCl for 30 minutes, washed and blocked in PBNT for one hour. Primary anti-BrdU antibody (Becton-Dickinson) was added at 1:200 overnight at 4°C. Samples were washed and incubated with HRP-conjugated secondary antibody at 1:400 for 2–4 hours at room temperature. Signal was developed using a DAB substrate (Sigma).

**Larval hemocyte assay**

Hemocytes from three third instar larvae were harvested using a Pasteur pipette pulled to generate a capillary end, pooled and counted on a standard hemacytometer slide. Five 16-square regions were counted for each pooled sample. Three replicates were assayed for each genotype.

**Quantitative RealTime PCR**

Whole first instar brains or ventral ganglia dissected from the brains of second instar larvae were used for RNA isolation. For first instar brain samples, total RNA was isolated using Trizol (Invitrogen) following manufacturer's directions. Samples were DNAsed and reverse transcribed using oligo dT primers. The resulting cDNA was used to perform quantitative Real Time PCR with SYBR Green dye. For ventral ganglia isolated during second instar RNA was extracted and the sequences amplified as described in [49,50]. Hemocyte studies were carried out on pooled hemolymph from three third instar larvae per sample. RNA was extracted and amplified as for ventral ganglia. All qRT-PCR reactions were carried out in triplicate at three different template concentrations to ensure that we were within linear template range. Primer sequences are available upon request. β-actin expression was used as an internal control. Data were analyzed using the delta-delta calculation method to yield fold change compared to controls.

**Statistics**

Determination of significance was accomplished by use of Student’s t test or ANOVA, depending on the design of the study.

**List of Abbreviations**

- **Bnl**: Branchless
- **BrdU**: 5-Bromodeoxyuridine
- **dpp**: decapentaplegic
- **FGF**: Fibroblast Growth Factor
- **GFP**: Green Fluorescent Protein
- **Hh**: Hedgehog
- **HSPG**: Heparan Sulfate Proteoglycan
- **PDGF**: Platelet Derived Growth Factor
- **PVR**: PDGF- and VEGF-Receptor Related
- **qRT-PCR**: quantitative Real Time PCR
- **SHH**: Sonic Hedgehog
- **TGFβ**: Transforming Growth Factors
- **TNb**: Thoracic neuroblast
- **VEGF**: Vascular Endothelial Growth Factor
- **wg**: wingless

**Authors’ contributions**

JL carried out all of the TNb genetic and molecular analyses, localization of Trol-GFP in the larval brain, hemocyte counting and qRT-PCR in hemocytes. JL also oversaw the trol developmental/lethality phenotypic analyses.

PH carried out the trol developmental/lethality analyses and participated in the trol TNb studies.

AB oversaw the trol hemocyte counting studies.

MC participated in the trol hemocyte counting studies.

TP carried out the qRT-PCR analyses of first instar trol larval brains.
YP carried out the BrdU study of the trol first instar neuroblast phenotype.

SD conceived of the studies, contributed to the experimental design and interpretation of all studies and wrote the manuscript.

All authors have read and approved the final manuscript.

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