Continual inactivation of stem cell functional identity genes stabilizes progenitor commitment

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Summary Statement

Expanding the pool of stem cells that indirectly generate differentiated cell types through intermediate progenitors drives vertebrate brain evolution. Due to lack of lineage information, mechanistic investigation of their competency to generate intermediate progenitors remain impossible. Fly larval brain neuroblasts provide excellent *in vivo* models for investigating the regulation of stem cell functionality during neurogenesis. Type II neuroblasts undergo indirect neurogenesis by repeatedly dividing asymmetrically to generate a neuroblast and a progeny that commits to an intermediate neural progenitor (INP) identity. We identified Tailless as a unique regulator that maintains type II neuroblast functional identity including the competency to generate INPs. Successive inactivation during INP commitment renders *tll* refractory to activation by Notch signaling, preventing INPs from re-acquiring neuroblast functionality. We propose that continual inactivation of neural stem cell functional identity genes by histone deacetylation allows intermediate progenitors to stably commit to generate diverse differentiated cell types during indirect neurogenesis.

Keywords: *Drosophila*, intermediate progenitors, neural stem cells, Notch signaling, neurogenesis

Introduction

The outer subventricular zone (OSVZ) is unique to gyrencephalic mammals, and directly contributes to the immense disparity in appearance and functionality of their brains as compared to brains in lissencephalic mammals (Cárdenas and Borrell, 2019; Delaunay et al., 2017; Di Lullo and Kriegstein, 2017). A key functional feature of OSVZ neural stem cells is their dependence on intermediate progenitors to indirectly generate neurons (Cárdenas and Borrell, 2019; Delaunay et al., 2017; Di Lullo and Kriegstein, 2017). Intermediate progenitors function as transit amplifying cells to increase cell number and cell diversity, allowing for the formation of deep folding that is characteristic of gyrencephalic brains. Despite recent advance in our understanding of the generation of OSVZ neural stem cells (Fujita et al., 2019; Namba et al., 2019), key open questions regarding the mechanisms that regulate their unique functionality such as the competency to generate intermediate progenitors remain.

Two distinct neuroblast lineages function together to generate the requisite number of neurons for the development of adult fly brains (Farnsworth and Doe, 2017; Homem et al., 2015; Janssens and Lee, 2014). Similar to ventricular zone neural stem cells, type I neuroblasts undergo direct neurogenesis. A type I neuroblast undergoes repeated asymmetric division to generate one daughter cell that remains as a neuroblast and a sibling cell (ganglion mother cell or GMC) that divides once to generate two neurons. By contrast, type II neuroblasts undergo indirect neurogenesis, resembling OSVZ neural stem cells. A type II neuroblast continually undergoes asymmetric division to self-renew and to generate a sibling cell that commits to an intermediate neural progenitor (INP) identity (Bello et al., 2008; Boone and Doe, 2008; Bowman et al., 2008). The Sp8 family transcription factor Buttonhead (Btd) and the ETS-1 transcription factor PointedP1 (PntP1) are specifically expressed in type II neuroblasts (Komori et al., 2014a; Xie et al., 2014; Zhu et al., 2011). Btd and PntP1 expression diminishes as a newly born immature INP transitions into an non-Asense expressing (Ase^-) immature INP. Three to four hours after this transition, an Ase^- immature INP up-regulates Ase expression as it progresses through INP commitment. Once INP commitment is complete, an Ase^+ immature INP transitions into an INP that undergoes 6-8 rounds of asymmetric division. These molecularly defined intermediate stages during INP commitment provide critical landmarks to search for genes that control type II neuroblast functional identity.

Notch signaling maintains type II neuroblasts in an undifferentiated state partly by poising transcription of the master regulator of differentiation earmuff (*erm*) (Janssens et al., 2017). During asymmetric neuroblast division, the TRIM-NHL protein Brain tumor (Brat) segregates into the newly born immature INP, and targets transcripts encoded by downstream effector genes of Notch for RNA decay (Bello et al., 2006; Betschinger et al., 2006; Komori et al., 2018; Lee et al., 2006; Xiao et al., 2012). *brat-null* brains accumulate thousands of supernumerary type II neuroblasts that originate from the reversion of newly born immature INPs due to defects in the down-regulation of Notch signaling (Komori et al., 2014b). In parallel to Brat, the nuclear protein Insensible (Insb) inhibits the activity of Notch downstream effector proteins during asymmetric neuroblast division, and Insb over-expression efficiently triggers premature differentiation in type II neuroblasts (Komori et al., 2018). This multi-layered gene control allows for the onset of Erm expression coinciding with termination of PntP1 expression. Erm expression is maintained in immature INPs, but rapidly diminishes in INPs (Janssens et al., 2014). Erm belongs to a family of transcription factors that are highly expressed in neural progenitors, and can bind histone deacetylase 3 (Hdac3) (Hirata et al., 2006; Koe et al., 2014; Levkowitz et al., 2003; Weng et al., 2010). Erm prevents INP reversion by repressing gene transcription. In *erm* null brains, INPs spontaneously revert into type II neuroblasts; this phenotype can be suppressed by knocking down *Notch* function (Weng et al., 2010). Thus, Erm likely inactivates type II neuroblast functionality genes by promoting histone deacetylation.

By comparing mRNAs enriched in type II neuroblasts or immature INPs, we identified the *tailless* (*tll*) as the master regulator of type II neuroblast functional identity. Tll is expressed in type II neuroblasts but not in INPs, and is necessary and sufficient for type II neuroblast functionality. Tll over-expression is sufficient to transform type I neuroblasts into type II neuroblasts as indicated by changes in gene expression and acquisition of the competence to generate intermediat progenitors. We identified and demonstrated *hamlet* (*ham)* as a novel negative regulator of type II neuroblast maintenance. *ham* is expressed after *erm*, and Erm and Ham function through Hdac3 to continually inactivate *tll*. Sequential inactivation during INP commitment renders *tll* refractory for activation by Notch signaling in INPs. We propose that continual inactivation of the master regulator of stem cell functional identity by histone

deacetylation allows intermediate progenitors to stably commit to generate differentiated cell types but not re-acquire stem cell functionality instead.

Results

A novel transient over-expression strategy to identify regulators of type II neuroblast functional identity

Genes that regulate neuroblast functional identity should be expressed in type II neuroblasts and become rapidly down-regulated in Ase^- and Ase^+ immature INPs in wild-type brains (Figure 1A). To enrich for transcripts that fulfills this criterium, we tested if supernumerary type II neuroblasts transiently over-expressing a *UAS-insb* transgene synchronously transition into INPs in *brat-null* brains as in wild-type brains. The combination of *Wor-Gal4,AseGal80*, which is only active in type II neuroblasts, and *Tub-Gal80^{ts}*, which loses its inhibitory effect on Gal4 activity under non-permissive temperatures, allows for spatial and temporal control of Insb over-expression in all type II neuroblasts. We allowed larvae to grow at 25°C, and transiently over-expressed Insb for 6, 12, 18, or 24 hours by shifting them to a non-permissive temperature of 33°C. Larvae that remained at 25°C for the entire duration of this experiment served as the control (time 0). We assessed the effect of this transient over-expression strategy on cell identity by quantifying total type II neuroblasts and INPs per brain lobe based on Deadpan (Dpn) and Ase expression (Figure 1A). Over-expressing Insb for 6 hours did not significantly affect the supernumerary neuroblast phenotype in *brat-null* brains (Figure S1). By contrast, 12 or more hours of Insb over-expression led to a time-dependent decrease in supernumerary type II neuroblasts and a corresponding increase in INPs. These results demonstrate that transient Insb over-expression can induce supernumerary type II neuroblasts to synchronously transition into INPs in *brat-null* brains.

We next assessed if supernumerary type II neuroblasts transiently over-expressing Insb indeed transition through identical intermediate stages to INPs in *brat-null* brains as in wild-type brains (Figure 1A). We found that the level of *pntP1* mRNA increased within the first 6 hours of Insb over-expression, and then continually declined (Figure 1B). By contrast, *erm* transcription rapidly increased in the first 12 hours of Insb over-expression, and plateaued between 12 and 18 hours. *ase* transcription showed little

change in the first 6 hours of Insb over-expression, and then steadily increased between 12 and 24 hours. The combined temporal patterns of *pntP1*, *erm* and *ase* transcription strongly suggest that type II neuroblasts transiently over-expressing Insb for 6 hours in *brat-null* brains are at the equivalent stage as a newly born immature INP transitioning to an Ase^- immature INP in wild-type brains. *brat-null* type II neuroblasts are at the stage equivalent to Ase^- immature INPs following 6-12 hours of Insb over-expression, and at the stage equivalent to Ase^+ immature INPs following 12-24 hours of Insb over-expression. Indeed, most type II neuroblasts in *brat-null* brains expressed cell identity markers indicative of Ase^- immature INPs (Erm::$V5^+Ase^-$) following 9 hours of Insb over-expression, and then Ase^+ immature INPs (Erm::$V5^+Ase^+$) following 18 hours of Insb over-expression (Figures 1C-F). These data indicate that supernumerary type II neuroblasts transiently over-expressing Insb indeed transition through identical intermediate stages during INP commitment in *brat-null* brains as in wild-type brains.

We predicted that a candidate regulator of type II neuroblast functional identity should become down-regulated in *brat-null* brains following 9 and 18 hours of Insb over-expression. By sequencing mRNA in triplicate following the transient over-expression strategy (Figure 1C), we identified 76 genes that were reproducibly down-regulated by 1.5-fold or more in *brat-null* brains over-expressing Insb for 9 hours (Figure 1G). *tll* is the most down-regulated gene among these initial candidates as well as in *brat-null* brains over-expressing Insb for 18 hours. We validated *tll* expression pattern in the type II neuroblast lineage using a bacterial artificial chromosome (BAC) transgene (*tll::GFP(BAC))* where green fluorescent protein (*GFP*) is fused in frame with the *tll* reading frame. Consistent with a previous study (Bayraktar and Doe, 2013), we detected Tll::GFP in type II neuroblasts, but not in immature or mature INPs (Figure 1H). Thus, *tll* is uniquely expressed in type II neuroblasts, and is an excellent candidate for regulating type II neuroblast functionality.

tll is the master regulator of type II neuroblast functional identity

We defined the type II neuroblast functional identity as the maintenance of an undifferentiated state and the competency to generate INPs. We first tested if *tll* is required for maintaining type II neuroblasts in an undifferentiated state by over-expressing a *UAS-tll*^{RNAi} transgene to knock down *tll*

function in larval brains. While control brains always contained 8 type II neuroblasts per lobe, brains with *tll* function knocked down contained 4.3 ± 1.2 type II neuroblasts per lobe (Figure 2B; n = 10 brains per genotype). Closer examination revealed that the remaining *tll* mutant neuroblasts showed reduced cell diameter and ectopically expressed Ase, two characteristics that are typically associated with INPs in the type II neuroblast lineage (Figures 2A, 2C, 2D). This result indicates that *tll* is required for maintaining type II neuroblast in a undifferentiated state. We next tested if *tll* is sufficient to re-establish a type II neuroblast-like undifferentiated state in INPs by mis-expressing a *UAS-tll* transgene under the control of the *Erm-Gal4(III)* driver (Figure 1A). Brains with *tll* mis-expressed in INPs contained 150 ± 50 type II neuroblasts per lobe, whereas control brains always contained 8 type II neuroblasts per lobe (Figures 2E and 2F; Figure S2A; n = 10 brains per genotype). Thus, Tll mis-expression is sufficient to revert INPs into type II neuroblasts. These data together led us to conclude that *tll* is necessary and sufficient for maintaining type II neuroblasts in a undifferentiated state.

The requirement of *tll* in maintaining type II neuroblasts in a undifferentiated state hindered direct assessment of its role in regulating the competency to generate INPs. As an alternative, we tested if Tll over-expression is sufficient to transform type I neuroblasts into type II neuroblasts. To unambiguously distinguish the two types of neuroblasts, we searched for robust protein markers for type II neuroblasts and their progeny. *Sp-1* mRNAs are uniquely detected in type II neuroblasts (Yang et al., 2016). We proceeded to determine Sp1 protein expression by using a *Sp-1::GFP(BAC)* transgene. We found that Sp1::GFP is expressed in all type II neuroblasts and their progeny, which are exclusively located on the dorsal surface of the brain (Figure 2G). Importantly, Sp1::GFP was undetectable in all type I neuroblasts and their progeny in the brain as well as the ventral nerve cord (Figures 2G, 2H, 2J). Thus, Sp1::GFP is a new marker for type II neuroblast lineages (Figure 2A). Many type I neuroblasts ectopically expressing *tll* lost Ase expression but gained Sp1::GFP expression in the brain as well as the ventral nerve cord (Figures 2I and 2K). Thus, ectopic *tll* expression is sufficient to molecularly transform type I neuroblasts into type II neuroblasts (Figure 2A). Next, we tested if type I neuroblasts ectopically expressing *tll* can generate immature INPs by examining the expression of Erm::V5, which is expressed in all but newly born immature INPs (Figure 2A). Indeed, we detected Ase^- immature INPs ($Erm::V5^+Ase^-$) and Ase^+

immature INPs (Erm::V5^{+}Ase^{+}) in many type I neuroblast lineages that ectopically express *tll* (Figure 2K; data not presented). These data led us to conclude that ectopic *tll* expression is sufficient to molecularly and functionally transform type I neuroblasts into type II neuroblasts. We conclude that *tll* is a master regulator of type II neuroblast functional identity.

Ham is a new regulator of INP commitment

tll is a putative Notch target in neuroblasts (Zacharioudaki et al., 2016). Because Notch signaling becomes re-activated in INPs, *tll* must become inactivated during INP commitment to prevent supernumerary type II neuroblast formation at the expense of generation of differentiated cell types (Figure 2A). We predicted that genes required for inactivating *tll* should become be up-regulated in immature INPs in wild-type brains as well as in *brat-null* brains following 9 and 18 hours of Insb over-expression. From our RNA sequencing dataset, we identified a list of genes that encode transcription factors and are up-regulated in *brat-null* brains over-expressing Insb for 9 or 18 hours. We performed reverse transcriptase PCR and confirmed that the transcript levels of these genes indeed become up-regulated in *brat-null* brains over-expressing Insb (Figure 1B; data not presented). Thus, these genes are candidates for inactivating *tll* during INP commitment.

erm^{hypo} brains are highly sensitive to changes in gene activity required to prevent INPs from reverting into type II neuroblasts (Janssens et al., 2017). Because ectopic *tll* expression in INPs leads to supernumerary type II neuroblast formation, reducing the function of genes that inactivate *tll* should enhance the supernumerary neuroblast phenotype in *erm^{hypo}* brains. *erm^{hypo}* brains alone contained 31.5 ± 6.8 type II neuroblasts per lobe (Figure 3A; n = 10 brains). While reducing the function of most candidate genes did not enhance the supernumerary neuroblast phenotype in *erm^{hypo}* brains, knocking down *ham* function with two different *UAS-RNAi* transgenes reproducibly enhanced the phenotype (Figure 3A; n = 10 brains per transgene). Consistent with the effect of reducing *ham* function by RNAi, the heterozygosity of a deficiency that deletes the entire *ham* locus also enhanced the supernumerary neuroblast phenotype in *erm^{hypo}* brains (Figure S3A). Thus, *ham* likely plays a role in preventing INP reversion into supernumerary type II neuroblasts.

Our findings contradicted a previous study that concludes *ham* playing no roles in suppressing supernumerary type II neuroblast formation (Eroglu et al., 2014). This discrepancy might be due to the ham^{1} allele used in the study, which encodes a nearly full-length protein and exhibits similar protein stability with wild-type Ham (Figure 3B) (Moore et al., 2002). We took two approaches to test if *ham* is required for preventing supernumerary type II neuroblast formation. First, we examined if *ham* deficiency heterozygous brains might display a supernumerary type II neuroblast phenotype. We reproducibly observed a mild supernumerary type II neuroblast phenotype in *ham* deficiency heterozygous brains (9 ± 0.9 per lobe, n = 12 brains) (Figure S3A). Second, we knocked down *ham* function by over-expressing a *UAS-RNAi* transgene. We found that knocking down *ham* function also led to a mild but statistically significant increase in type II neuroblasts per lobe (9.8 ± 1.8; n = 20 brains) as compared to control brains (8 ± 0; n = 10 brains) (Figure S3B). To confirm that *ham* is indeed required for suppressing supernumerary type II neuroblast formation, we generated two new *ham* alleles, ham^{SK1} and ham^{SK4}, by CRISPR-CAS9 (Figure 3B). We confirmed that ham^{SK1} or ham^{SK4} homozygous brains show undetectable Ham protein expression by using an antibody specific for Ham (Figures 3C and 3D; data not presented). We then tested if Ham protein is required for suppressing supernumerary type II neuroblast formation. Indeed, ham^{SK1} homozygous brains contained 40 ± 10 type II neuroblasts per lobe (Figures 3E and 3F; n = 15 brains). Because Ham becomes detectable in Ase^{+} immature INPs and remains expressed in all INPs (Eroglu et al., 2014), *ham* is a novel regulator of INP commitment.

ham knock-down drastically enhanced the supernumerary neuroblast phenotype in erm^{hypo} brains, leading us to hypothesize that *ham* functions together with *erm* to suppress INP reversion into supernumerary type II neuroblasts. We tested this hypothesis by examining if *erm* and *ham* genetically interact during INP commitment. Although the heterozygosity of *erm* did not increase total type II neuroblasts (8.1 ± 0.3 per lobe; n = 11 brains), it did enhance the supernumerary neuroblast phenotype in *ham* deficiency heterozygous brains (14 ± 2.2 per lobe; n = 14 brains) as well as in ham^{SK1} homozygous brains (76.2 ± 18.3 per lobe; n = 14 brains) (Figures 3G-I). We conclude that Ham functions synergistically with Erm to suppress supernumerary type II neuroblast formation.

Ham promotes stable INP commitment by repressing gene transcription

Re-activation of Notch signaling in INPs drives supernumerary type II neuroblast formation in *erm* null brains (Weng et al., 2010). We tested if supernumerary type II neuroblasts in *ham-null* brains also originate from INPs. Whereas *erm,ham* heterozygous brains alone contained 12.7 ± 1.6 type II neuroblasts per lobe (n = 9 brains), over-expressing *ham* in Ase^+ immature INPs and INPs driven by *Erm-Gal4(III)* rescued the supernumerary neuroblast phenotype in *erm,ham* double heterozygous brains (8.2 ± 0.8 neuroblasts per lobe; n = 24 brains) (Figure 3J). This result strongly suggests that supernumerary type II neuroblasts originate from INPs in *ham-null* brains. We generated GFP-marked mosaic clones derived from single type II neuroblasts to confirm the origin of supernumerary type II neuroblasts in *ham-null* brains. In wild-type clones (n = 9 clones), the parental type II neuroblast was always surrounded by Ase^- and Ase^+ immature INPs (Figure 4A). By contrast, supernumerary neuroblasts in ham^{SK1} homozygous clones (2.5 ± 1.7 type II neuroblasts per clone; n = 11 clones) were always located far away from parental neuroblasts and surrounded by Ase^+ cells that were most likely Ase^+ immature INPs and ganglion mother cells (Figures 4B and 4C). It was highly unlikely that supernumerary neuroblasts in ham^{SK1} homozygous clones originated from symmetric neuroblast division based on their relative locations to parental neuroblasts and the cell types that surrounded them. Thus, we conclude that Ham suppresses INP reversion into supernumerary type II neuroblasts.

Because Erm suppresses supernumerary type II neuroblast formation by repressing gene transcription, Ham likely prevents INP reversion by functioning as a transcriptional repressor. We reasoned that Ham functions through its N-terminal zinc finger to prevent INP reversion based on our finding that ham^{SK1} but not ham^{1} homozygous clones displayed a supernumerary neuroblast phenotype (Figures 3D and 4C). Consistent with this hypothesis, over-expressing $Ham^{\Delta C\text{-}ZF}$ rescued the supernumerary neuroblast phenotype in ham^{SK1} homozygous brains to a similar extend as over-expressing full-length Ham (11.1 ± 1.4 neuroblasts per lobe vs. 11.9 ± 1.3 neuroblasts per lobe; n = 11 or 9 brains, respectively) (Figure 4E). Furthermore, mis-expressing full-length Ham or $Ham^{\Delta C\text{-}ZF}$ triggered premature differentiation in type II neuroblasts (n = 10 brains per genotype) (Figure 4F). These data indicate that Ham functions through the N-terminal zinc-finger motif to prevent INP reversion. Under

identical conditions, over-expressing a constitutive transcriptional repressor form of Ham containing only the N-terminal zinc fingers (ERD::Ham$^{N\text{-}ZF}$) also rescued the supernumerary neuroblast phenotype in *ham*^{SK1} homozygous brains (14 ± 2.3 neuroblasts; n = 9 brains) (Figures 4D and 4E). These results indicate that Ham functions through the N-terminal zinc finger to repress the transcription of genes that can trigger INP reversion into supernumerary type II neuroblasts.

Finally, we tested if the N-terminal zinc finger of Ham mediate target gene recognition. We mis-expressed a constitutive transcriptional activator form of Ham containing only the N-terminal zinc-finger motif (VP16::Ham$^{N\text{-}ZF}$) (Figure 4D). We found that VP16::Ham$^{N\text{-}ZF}$ mis-expression in type II neuroblasts was sufficient to induce supernumerary neuroblast formation (17.1 ± 4.7; n = 10 brains; Figure 4G). Because VP16::Ham$^{N\text{-}ZF}$ can exert dominant negative effect, we conclude that Ham suppresses INP reversion by recognizing target genes through its N-terminal zinc-finger motif and repressing their transcription.

Inactivation during INP commitment relinquishes the competence to activate *tll* by Notch in INPs

Our findings strongly suggest that Erm and Ham likely inactivate *tll* during INP commitment rendering re-activation of Notch signaling unable to trigger *tll* expression in INPs (Figure 5A). We tested if Erm and Ham indeed inactivate *tll* by reducing *tll* function in *erm,ham* double heterozygous brains. *tll* heterozygous brains contained 8.2 ± 0.4 type II neuroblasts per lobe (n = 10 brains; Figure 5B). *erm,ham* double heterozygous brains contained 11 ± 1.2 type II neuroblasts per lobe (n = 25 brains; Figure 5B). The heterozygosity of *tll* consistently suppressed the supernumerary neuroblast phenotype in *erm,ham* double heterozygous brains (8.4 ± 0.6 type II neuroblasts per lobe; n = 22 brains; Figure 5B). This result strongly supports our hypothesis that Erm and Ham inactivate *tll* during INP commitment. Consistent with this interpretation, we found that Tll::GFP becomes ectopically expressed in INPs in *erm,ham* double heterozygous brains (Figure 5C). In these brains, we reproducibly observed small cells that expressed Tll::GFP and Deadpan (Dpn) but not Ase, and were most likely supernumerary type II neuroblasts newly derived from INP reversion (Figure 5D). Furthermore, we found that 1 copy of the *tll::GFP(BAC)* transgene mildly enhanced the supernumerary neuroblast phenotype in *erm,ham* double heterozygous

brains (15.4 ± 1.8 type II neuroblasts per lobe vs. 11.1 ± 1 type II neuroblasts per lobe; n = 12 or 11 brains) (Figure 5E). These data indicate that sequential inactivation by Erm and Ham during INP commitment renders *tll* refractory for activation in INPs.

Because *tll* is a putative Notch target in neuroblasts (Zacharioudaki et al., 2016), we tested if inactivation by Erm and Ham relinquishes the competence of Notch signaling to activate *tll* in INPs. We knocked down *Notch* function in INPs in wild-type or ham^{SK1} homozygous brains. Knock-down of *Notch* function in INPs had no effect on type II neuroblasts in wild-type brains (7.9 ± 0.3; n = 17 brains; Figure 5F). Whereas ham^{SK1} homozygous brains alone contained 128 ± 39.6 type II neuroblasts per lobe (n = 9 brains), ham^{SK1} homozygous brains with knocking down *Notch* function in INPs contained 36.4 ± 11.6 type II neuroblasts per lobe (n = 9 brains) (Figure 5F). This result indicates that re-activation of Notch signaling triggers INP reversion into supernumerary type II neuroblasts in *ham-null* brains. We extended our analyses to test if inactivation by Erm and Ham indeed reduces the competency to respond to Notch signaling. Consistently, over-expressing constitutively activated Notch ($Notch^{intra}$) induced a drastically more severe supernumerary neuroblast phenotype in *erm*,*ham* double heterozygous brains (3,437.6 ± 586.8 type II neuroblasts per lobe; n = 9 brains) than in *erm* or *ham* single heterozygous brains (687 ± 134.3 type II neuroblasts per lobe vs. 588.6 ± 77.6 type II neuroblasts per lobe, respectively; n = 9 or 11 brains) (Figure 5G). Together, these data strongly suggest that inactivation by Erm and Ham during INP commitment renders *tll* refractory to respond to Notch signaling in INPs (Figure 5A).

Inactivation by Hdac3 relinquishes the competence to activate *tll* by Notch in INPs

To render Notch signaling unable to activate *tll* in INPs, Erm and Ham must function through some chromatin modifying proteins to inactivate *tll* during INP commitment. We first tested if Erm and Ham inactivate *tll* by promoting sequential chromatin changes during INP commitment. We over-expressed full-length Ham or $Ham^{\Delta C\text{-}ZF}$ driven by *Erm-Gal4(III)* in *erm* null brains. *erm* null brains alone contained 1,824.6 ± 520.7 type II neuroblasts per lobe (n = 13 brains) (Figures 6A and S4A). Over-expressing full-length Ham or $Ham^{\Delta C\text{-}ZF}$ strongly suppressed the supernumerary neuroblast phenotype in *erm* null brains (34.5 ± 8.3 vs. 64.6 ± 18.3 neuroblasts per lobe; n = 13 vs. 7 brains) (Figures 6A and

S4B). This result indicates that Ham can replace endogenous Erm function to suppress INP reversion, suggesting that these two transcriptional repressors function through an identical chromatin modifying protein to inactivate *tll* during INP commitment.

We predicted that decreasing the activity of chromatin modifying protein required for Ham-mediated gene inactivation during INP commitment should enhance the supernumerary neuroblast phenotype in *ham* heterozygous brains. We proceeded to knock down the function of genes known to contribute to the inactivation of gene transcription in *ham* heterozygous brains. Neither reducing the activity of Polycomb Repressive Complex 2 nor Heterochromatin Protein 1 or decreasing the recruitment of Heterochromatin Protein 1 had any effect on the supernumerary neuroblast phenotype in *ham* heterozygous brains (Figure 6B). While reducing the activity of nucleosome remodelers alone led to a mild supernumerary type II neuroblast phenotype, it did not further enhance the supernumerary neuroblast phenotype in *ham* heterozygous brains (Figure 6C). By contrast, reducing the activity of histone deacetylase 3 (Hdac3) alone resulted in a mild supernumerary type II neuroblast phenotype, and significantly enhanced the supernumerary neuroblast phenotype in *ham* heterozygous brains (Figure 6C). These results led us to conclude that Ham likely functions through Hdac3 to prevent INP reversion. We next tested if Erm also functions through Hdac3 to prevent INP reversion. *erm* heterozygous brains alone did not display a supernumerary type II neuroblast phenotype (Figure 6D). Knocking down *hdac3* function in *erm* heterozygous brains led to a more than 3-fold increase in supernumerary type II neuroblasts as compared to wild-type brains (Figure 6D). Thus, Erm and Ham likely function through Hdac3 to prevent INP reversion into supernumerary type II neuroblasts by continually inactivating *tll*.

Discussion

The expansion of OSVZ neural stem cells, which indirectly produce neurons by first generating intermediate progenitors, drives the evolution of lissencephalic brains to gyrencephalic brains (Cárdenas and Borrell, 2019; Delaunay et al., 2017; Di Lullo and Kriegstein, 2017). Recent studies have revealed important insights into genes as well as cell biological changes that lead to the formation of OSVZ neural stem cells (Fujita et al., 2019; Namba et al., 2019). However, the mechanisms that control the functional

identity of OSVZ neural stem cells including the competency to generate intermediate progenitors remain completely unknown. In this study, we provided compelling evidence to demonstrate that Tll is necessary and sufficient for the maintenance of a undifferentiated state and the competency to generate intermediate progenitors in type II neuroblasts (Figure 7). We also showed that two sequentially activated transcriptional repressors, Erm and Ham, inactivate *tll* during INP commitment to ensure normal indirect neurogenesis in larval brains. We identified Hdac3 as the key chromatin modifying protein that functions together with Erm and Ham to prevent aberrant INP reversion into supernumerary type II neuroblasts. We propose that continual inactivation of stem cell functional identity genes by histone deacetylation allow intermediate progenitors to stably commit to generate the number and the diversity of differentiated cells during neurogenesis.

Regulation of the maintenance of a undifferentiated state and the competency to generate intermediate progenitors in neural stem cells

Stem cell functional identity encompasses the maintenance of a undifferentiated state as well as other unique functional features such as the competency to generate intermediate progenitors. Because genetic manipulation of Notch signaling perturbs the regulation of differentiation during asymmetric stem cell division (Imayoshi et al., 2010; Kageyama et al., 2008), the role of Notch in regulating other stem cell functions remain poorly understood. In the fly type II neuroblast lineage, over-expressing Notch^{intra} or downstream transcriptional repressors Dpn, E(spl)mγ, and Klumpfuss (Klu) induces the formation of supernumerary type II neuroblasts at the expense of generating immature INPs by inhibiting *erm* activation (Janssens et al., 2017). Thus, Notch signaling prevents type II neuroblast from initiating differentiation by maintaining *erm* in a poised but inactivated state. By contrast, over-expressing Notch^{intra} but not Dpn, E(spl)mγ and Klu in combination is sufficient to re-establish a type II neuroblast-like undifferentiated state in INPs driving supernumerary type II neuroblast formation. Thus, the Notch-Dpn/E(spl)mγ/Klu axis maintains type II neuroblasts in a undifferentiated state, and Notch alone is a competent of a regulatory circuit that endows type II neuroblasts with the competency to generate INPs.

We propose that *tll* is the master regulator of type II neuroblast functional identity (Figure 7). A

previous study found that Suppressor of Hairless, the DNA-binding partner of Notch, binds the *tll* locus in larval brain neuroblasts (Zacharioudaki et al., 2016). Identical to *Notch*, *tll* is necessary and sufficient for inducing a undifferentiated state that is associated with type II neuroblasts. However, over-expressing *tll* but not Notch^{intra} is sufficient to molecularly and functionally transform a type I neuroblast into a type II neuroblast. Thus, Notch likely contributes to *tll* activation, but additional activator that is specifically expressed in type II neuroblasts must also exist. Transcription factor Zelda (Zld) is expressed in type II neuroblasts but not in INPs, and binds the *cis*-regulatory region of *tll* in embryos (Harrison et al., 2011; Nien et al., 2011; Reichardt et al., 2018). However, all type I neuroblasts show a high level of activated Notch as well as Zld expression, but only a subset of type I neuroblasts show detectable Tll (Kurusu et al., 2009; Reichardt et al., 2018). Thus, it is unlikely that Zld play a key role in activating *tll* expression in type II neuroblasts. The vertebrate homolog of Tll, Tlx, has been shown to function as a transcriptional activator during neurogenesis (Sun et al., 2017). Thus, Tll might function in a positive feedback loop to amplify its own expression in type II neuroblasts. Elucidating the mechanisms that function cooperatively with Notch signaling to activate *tll* expression in type II neuroblasts might provide novel insights into the specification of OSVZ neural stem cells.

Successive transcriptional repressor activity inactivates stem cell functional identity genes during progenitor commitment

Identification of *ham* as a putative regulator of INP commitment was unexpected given a previously published study concluded that Ham functions to limit INP proliferation (Eroglu et al., 2014). Ham is the fly homolog of Prdm16 in vertebrates, and has been shown to play a key role in regulating cell fate decisions in multiple stem cell lineages (Baizabal et al., 2018; Harms et al., 2015; Moore et al., 2002; Shimada et al., 2017). Prdm16 contains two separately defined zinc finger motifs, each of which likely recognizes unique target genes. Prdm16 can also function through a variety of co-factors to activate or repress target gene expression independent of its DNA-binding capacity. Thus, Ham could potentially inactivate stemness genes via one of several mechanisms. By using a combination of previously isolated alleles and new protein-null alleles, we were able to show that the N-terminal zinc-

finger motif is required for Ham function in immature INPs. By over-expressing a series of chimeric proteins containing the N-terminal zinc-finger motif, our data indicate that Ham prevents INP reversion into supernumerary type II neuroblasts by recognizing target genes through the N-terminal zinc-finger motif and repressing their transcription. Our results indicate that Ham prevents INPs from reverting into supernumerary type II neuroblasts by repressing target gene transcription.

A key question raised by our study is why two transcriptional repressors that seemingly function in a redundant manner are required to prevent INPs from reverting into supernumerary type II neuroblasts. INP commitment lasts approximately 10-12 hours following the birth of an immature INP, after which time an immature INP transitions into an INP. *erm* is poised for activation in type II neuroblasts, and becomes rapidly activated in the newly born immature INP less than 90 minutes after birth (Janssens et al., 2017). As such, Erm-mediated transcriptional repression allows for rapid inactivation of type II neuroblast functional identity genes. Because Erm is no longer expressed in INPs where Notch signaling becomes re-activated, a second transcriptional repressor that becomes activated after Erm and whose expression is maintained throughout the life of an INP is required to continually inactivate type II neuroblast functional identity genes. Ham is an excellent candidate because it becomes expressed in immature INPs 4-6 hours after the onset of Erm expression, and is detected in all INPs. Similar to Erm, Ham recognizes target genes and repress their transcription. Furthermore, *ham* functions synergistically with *erm* to prevent INP reversion into supernumerary type II neuroblasts, and Ham over-expression can partially substitute for endogenous Erm. Thus, Erm- and Ham-mediate transcriptional repression renders type II neuroblast functional identity genes refractory for activation by Notch signaling throughout the lifespan of an INP, ensuring the generation of differentiated cell types but not supernumerary type II neuroblasts instead.

Sustained inactivation of stem cell functional identity genes distinguishes intermediate progenitors from stem cells

Genes that specify stem cell functional identity become refractory for activation during differentiation, but the mechanisms that restrict their expression are poorly understood due to the lack of

lineage information. Several epigenetic regulator complexes have been proposed to restrict neural stem cell-specific gene expression in neurons (Hirabayashi and Gotoh, 2009; Ronan et al., 2013). We knocked down the function of genes that were implicated in restricting neural stem cell gene expression during differentiation in search for chromatin regulators that are required for inactivating type II neuroblast functional identity genes during INP commitment. Surprisingly, we found that only Hdac3 is required for both Erm- and Ham-mediated suppression of INP reversion into type II neuroblasts. Our finding is consistent with a recent study showing that blocking apoptosis in lineage clones derived from type II neuroblasts mutant for Polycomb Repressive Complex 2 did not lead to supernumerary neuroblast formation (Abdusselamoglu et al., 2019). Our data strongly suggest that genes specifying type II neuroblast functional identity such as *tll* likely are inactivated rather than repressed in INPs. This result is supported by the result that over-expressing Notch^{intra} but not Notch downstream transcriptional repressors in INPs can re-establish a type II neuroblast-like undifferentiated state. We speculate that the chromatin in the *tll* locus remains open and accessible in INPs, and continual histone deacetylation is sufficient to counter the transcriptional activator activity of endogenous Notch and maintain *tll* in an inactive state (Figure 7). By contrast, the chromatin in the *tll* locus might be close and inaccessible by the Notch transcriptional activator complex, and over-expressing Notch^{intra} cannot transform type I neuroblasts into type II neuroblasts. If this were to be true, a key question to answer is what transcription factor is required to keep the chromatin in the *tll* loci in an open state. Insights into regulation of the competency of the *tll* locus to respond to activated Notch signaling might improve our understanding of the molecular determinants of OSVZ neural stem cells.

Author Contributions

N.Q-R., H.K., and D.H. conducted the experiments. N.Q-R., H.K., D.H., and C.L. designed the experiments. S.K, Q.D., and A.M contributed key reagents. N.Q-R., H.K., and C.L. wrote the manuscript.

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Declaration of Interests

The authors declare no competing interests.

Figure Legends

Figure 1. Identification of candidate regulators of type II neuroblast functional identity.

(A) A diagram of the type II neuroblast lineage showing the expression patterns of genes and *Gal4* drivers. The color scheme of arrows and arrowheads used to identify various cell types in the type II neuroblast lineage in all figures is shown. (B) A time course analysis of gene transcription patterns in *brat*-null brains following transient Insb over-expression. Transient Insb over-expression triggers supernumerary type II neuroblasts in *brat*-null brains to activate gene transcription patterns that mimic immature INPs undergoing INP commitment in wild-type brains. (C) A strategy to synchronize the identities of supernumerary type II neuroblasts in *brat*-null brains to undergo INP commitment induced by transient Insb over-expression. Larvae were collected and aged at 25°C. Transient Insb over-expression was induced by shifting one third of larvae to 33°C at 126 or 135 hours after hatching. The last one third remained at 25°C and served as the source enriched for type II neuroblast-specific transcripts (time 0). (D-F) Confocal images of *brat*-null brains with transient Insb over-expression for 0, 9, or 18 hours. Transient Insb over-expression triggers supernumerary type II neuroblasts in *brat*-null brains to sequentially activate Erm and then Ase expression, recapitulating changes in marker expression during INP commitment in wild-type brains. Larvae used in this experiment carry the *erm::V5* allele generated by CRISPR-CAS9, allowing for the detection of endogenous Erm expression. (G) Volcano plots showing fold-change of gene expression in *brat*-null brains with transient Insb over-expression for 9 or 18 hours.

(H) Tll expression pattern in the type II neuroblast lineage. Endogenous Tll is detected in the type II neuroblast but not in immature INPs and INPs in larval brains carrying one copy of the *tll::GFP(Bac)* transgene. Scale bar, 10 μm.

Figure 2. *tll* is necessary and sufficient for type II neuroblast functional identity.

(A) A diagram of the type I and II neuroblast lineages showing the expression patterns of genes. (B) Quantification of total type II neuroblasts per lobe in larval brains with knock-down of *tll* function. Knocking down *tll* function led to a reduced number of type II neuroblasts. (C-D) Knocking down *tll* function led to premature differentiation in type II neuroblasts as indicated by reduced cell diameter and ectopic Ase expression (white arrow). Type I neuroblasts (orange arrow) served as the control. (E-F) Confocal images of larval brains with *tll* ectopically expressed in Ase^+ immature INPs. *tll* mis-expression in Ase^+ immature INPs led to supernumerary type II neuroblast formation. (G-H) Sp-1::GFP marks the type II neuroblast lineages. Type I neuroblast lineages (orange arrow) can be found on the dorsal and ventral surface of the brain (Br), but type II neuroblast lineages (white arrow) are only found on the dorsal surface. (I) Tll over-expression molecularly transforms type I neuroblasts into type II neuroblasts as indicated by loss of Ase expression and gain of Sp-1::GFP expression. (J) Only type I neuroblasts are found on the wild-type ventral nerve cord. The thoracic segments (Th) are shown. (K) Tll over-expression molecularly transforms type I neuroblasts on the ventral nerve cord into type II neuroblasts. The thoracic segments (Th) are shown. (L) Tll over-expression functionally transforms type I neuroblasts into type II neuroblasts as indicated by the generation of Erm::$V5^+Ase^-$ immature INPs (white arrowhead) and Erm::$V5^+Ase^+$ immature INPs (yellow arrow). Scale bar, 10 μm. P-values: *** <0.005.

Figure 3. Ham is a novel regulator of INP commitment

(A) Quantification of total type II neuroblasts per lobe in erm^{hypo} brains with knock-down of candidate regulators of type II neuroblast functional identity. Knocking down *ham* function consistently enhanced the supernumerary type II neuroblast phenotype in erm^{hypo} brains. (B) A diagram summarizing the lesions in *ham* alleles. (C-D) ham^{SK1} is a protein-null allele. Ham was detected in immature INPs and INPs in

wild-type brains, but was undetectable in *ham*^{SK1} homozygous brains. (E) Quantification of total type II neuroblasts per lobe in *ham* mutant brains. *ham*^{SK1} homozygous brains display a supernumerary type II neuroblast phenotype but not *ham*^{1} homozygous brains. (F-H) The heterozygosity of *erm* alone had no effect on type II neuroblasts, but enhanced the supernumerary type II neuroblast phenotype in *ham*^{SK1} homozygous brains. (I) The heterozygosity of *erm* enhanced the supernumerary type II neuroblast phenotype in *ham* deficiency heterozygous brains and *ham*^{SK1} homozygous brains. (J) Over-expressing Ham in Ase^{+} immature INPs rescued the supernumerary type II neuroblast phenotype in *erm,ham* double heterozygous brains. Scale bar, 10 μm. P-values: *** $<$0.005. ns: not significant.

Figure 4. Ham suppresses INP reversion by repressing gene transcription

(A-B) Phenotypic characterization of *ham*^{SK1} homozygous type II neuroblast clones. Supernumerary neuroblasts (-15 μm) were always located far from the parental neuroblast (0 μm), and were surrounded by Ase^{+} cells in *ham*^{SK1} homozygous clones. (C) *ham*^{SK1} clones contained supernumerary type II neuroblasts but *ham*^{1} clones did not. (D) A diagram summarizing *ham* transgenes. (E) Quantification of total type II neuroblasts per lobe in *ham*^{SK1} homozygous brains that over-expressed various Ham transgenic proteins in Ase^{+} immature INPs. Over-expressing full-length Ham, truncated Ham lacking the C-terminal zinc-finger motif or ERD::Ham$^{N\text{-}ZF}$ rescued the supernumerary neuroblast phenotype in *ham*^{SK1} homozygous brains. (F) Brains over-expressing full-length Ham or Ham lacking the C-terminal zinc-finger motif contained fewer type II neuroblasts than control brains. (G) Over-expressing VP16::Ham$^{N\text{-}ZF}$ led to supernumerary type II neuroblast formation. Scale bar, 10 μm. P-values: *** $<$0.005. ns: not significant.

Figure 5. Erm- and Ham-mediated repression renders *tll* refractory for activation by Notch.

(A) A diagram depicting our model that ectopic activation of *tll* triggers INP reversion into supernumerary type II neuroblasts in *erm*,*ham* double heterozygous brains. (B) Heterozygosity of *tll* suppressed the supernumerary neuroblast phenotype in in *erm,ham* double heterozygous brains. (C-D) The *tll::GFP(BAC)* transgene becomes ectopically expressed in INPs and

supernumerary type II neuroblasts (*) in *erm,ham* double heterozygous brains. (E) 1 copy of the *tll::GFP(BAC)* transgene enhanced the supernumerary type II neuroblast phenotype in *erm,ham* double heterozygous brains. (F) Knocking down *Notch* function in INPs suppressed the supernumerary type II neuroblast phenotype in ham^{SK1} homozygous brains. (G) Over-expressing N^{intra} induced INP reversion into supernumerary neuroblasts much more efficiently in *erm,ham* double heterozygous brains than *erm* or *ham* heterozygous brains. Scale bar, 10 μm. P-values: ** <0.05, *** <0.005. ns: not significant.

Figure 6. Erm and Ham function through Hdac3 to suppress INP reversion into type II neuroblasts.

(A) Over-expressing full-length Ham or truncated Ham lacking the C-terminal zinc-finger motif suppressed the supernumerary type II neuroblast phenotype in *erm*-null brains. (B) Reducing activity of the Polycomb Repressive Complex 2 or the chromatin complex that promotes heterochromatin formation did not enhance the supernumerary neuroblast phenotype in *ham* heterozygous brains. (C) Reducing the activity of Hdac3 but not the Mi-2 or Brm complex enhanced the supernumerary neuroblast phenotype in *ham* heterozygous brains. (D) Reducing the activity of Hdac3 enhanced the supernumerary neuroblast phenotype in *erm* heterozygous brains. P-values: * <0.5, ** <0.05, *** <0.005. ns: not significant.

Figure 7. A proposed model for the regulation of type II neuroblast functionality.

Figure S1. Time course analysis of transient Insb over-expression in *brat*-null brains.

Quantification of total type II neuroblasts or INPs per lobe in *brat*-null brains following transient Insb over-expression.

Figure S2. Tll mis-expression in Ase^{+} immature INPs led to supernumerary type II neuroblast

formation.

Figure S3. Ham functions synergistically with Erm to suppress supernumerary type II neuroblast formation.

(A) *erm,ham* double heterozygous brains displayed the supernumerary type II neuroblast phenotype. (B) Knock-down of *ham* function led to a mild supernumerary type II neuroblast phenotype.

Figure S4. Ham over-expression in Ase$^+$ immature INPs suppressed INP reversion in *erm-null* brains.

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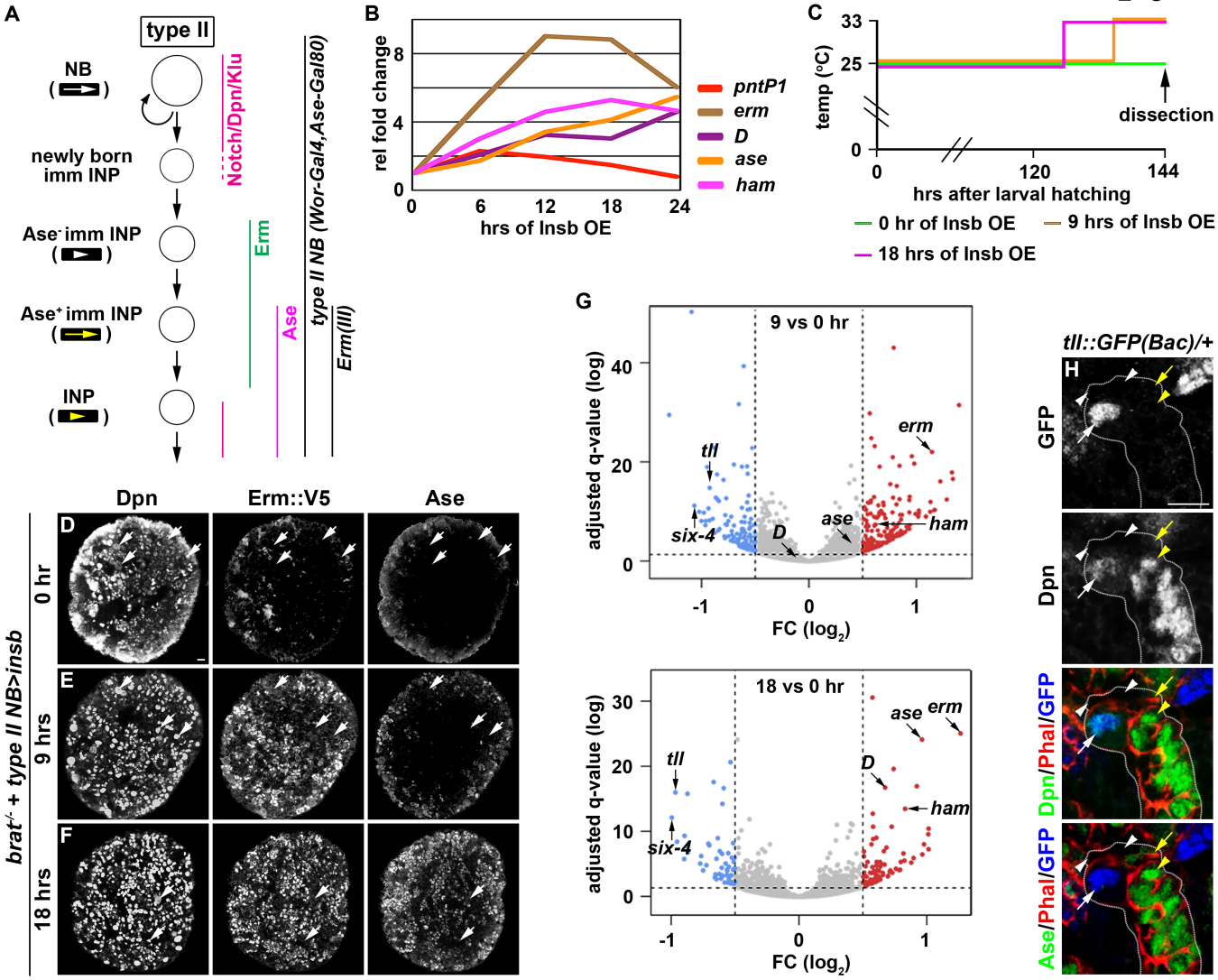

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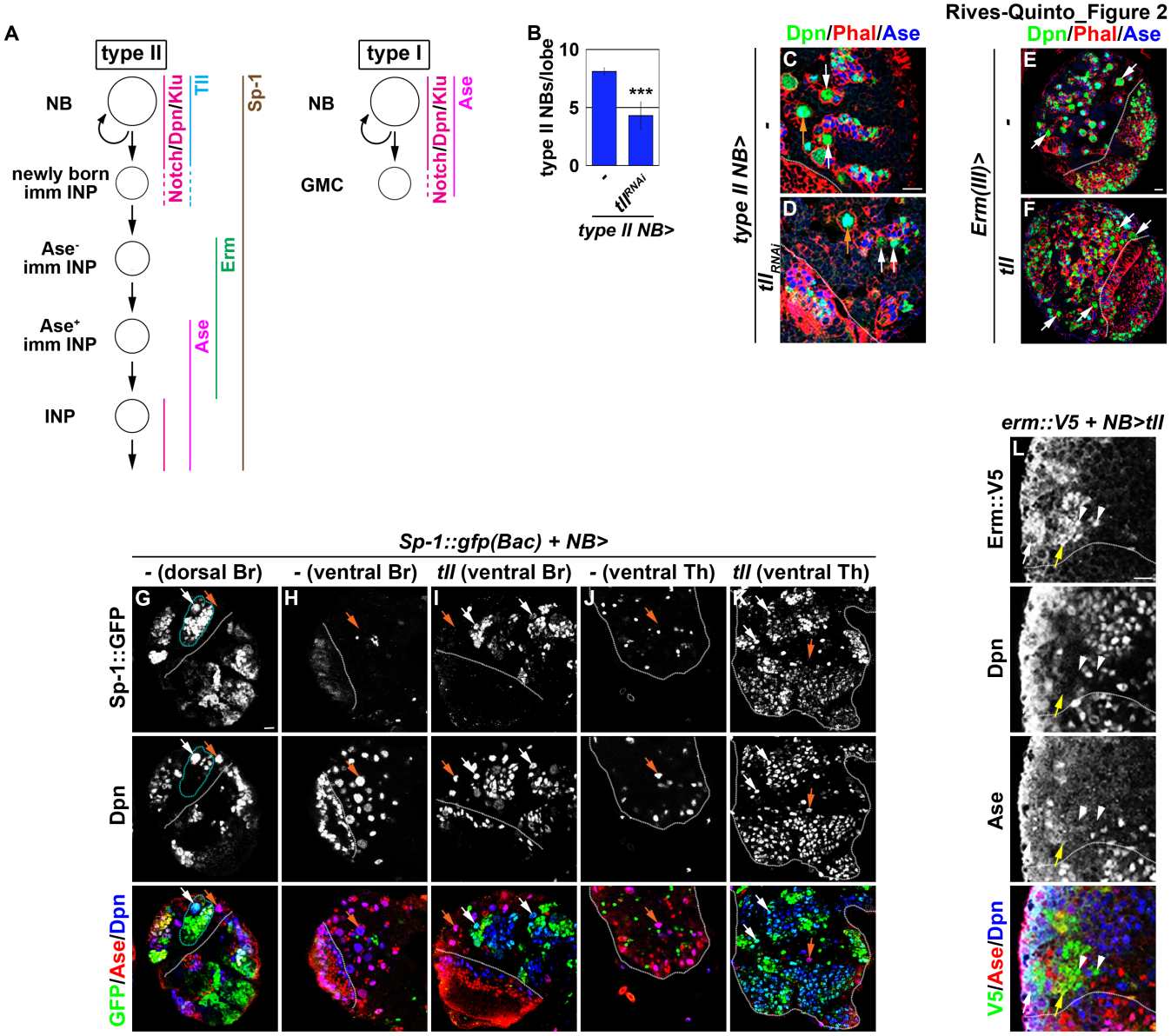

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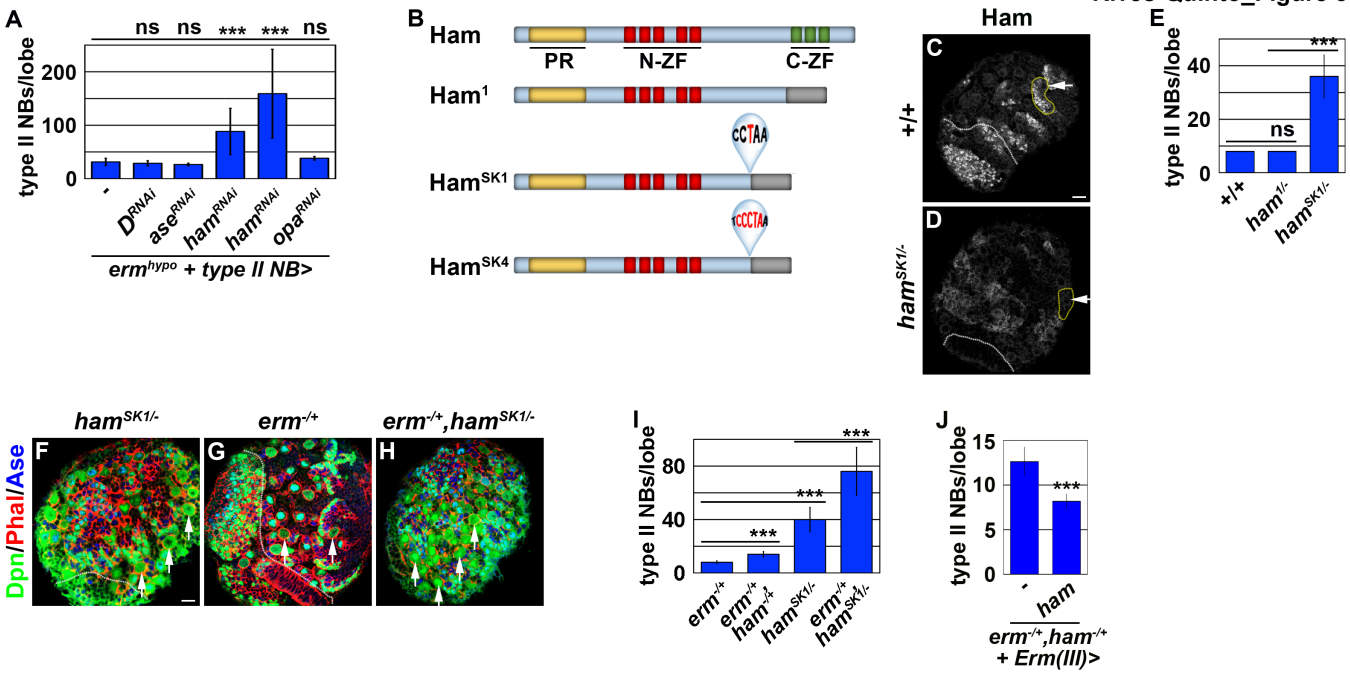

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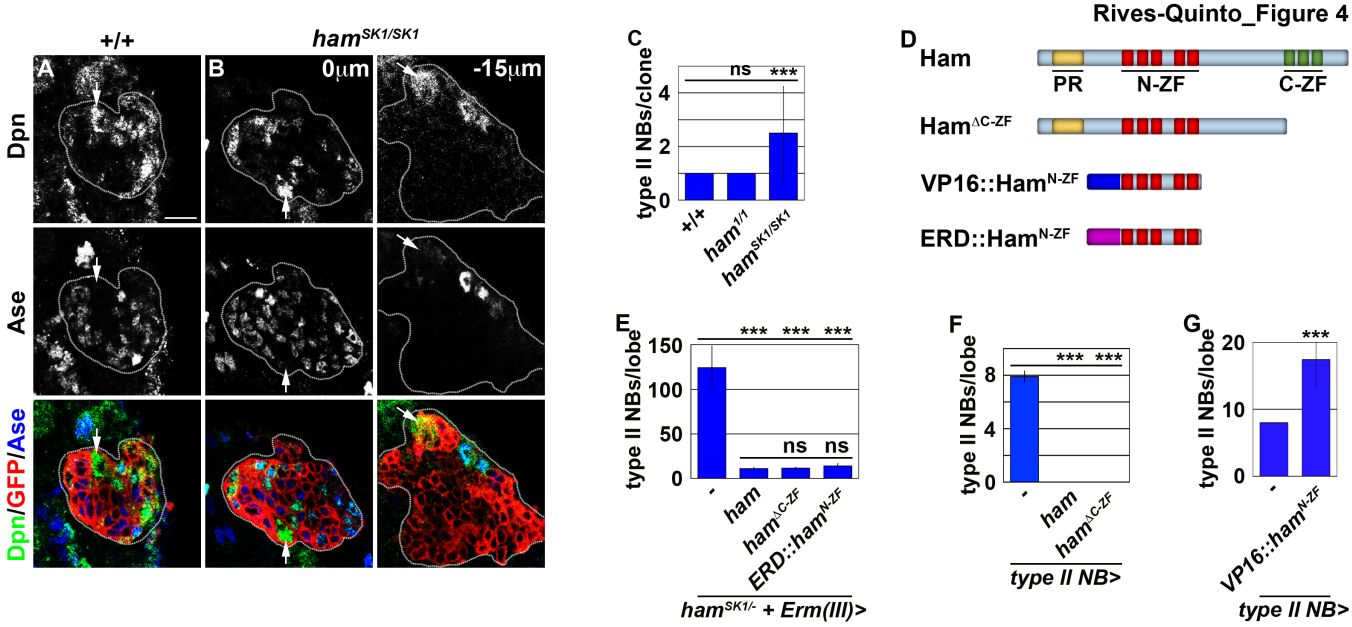

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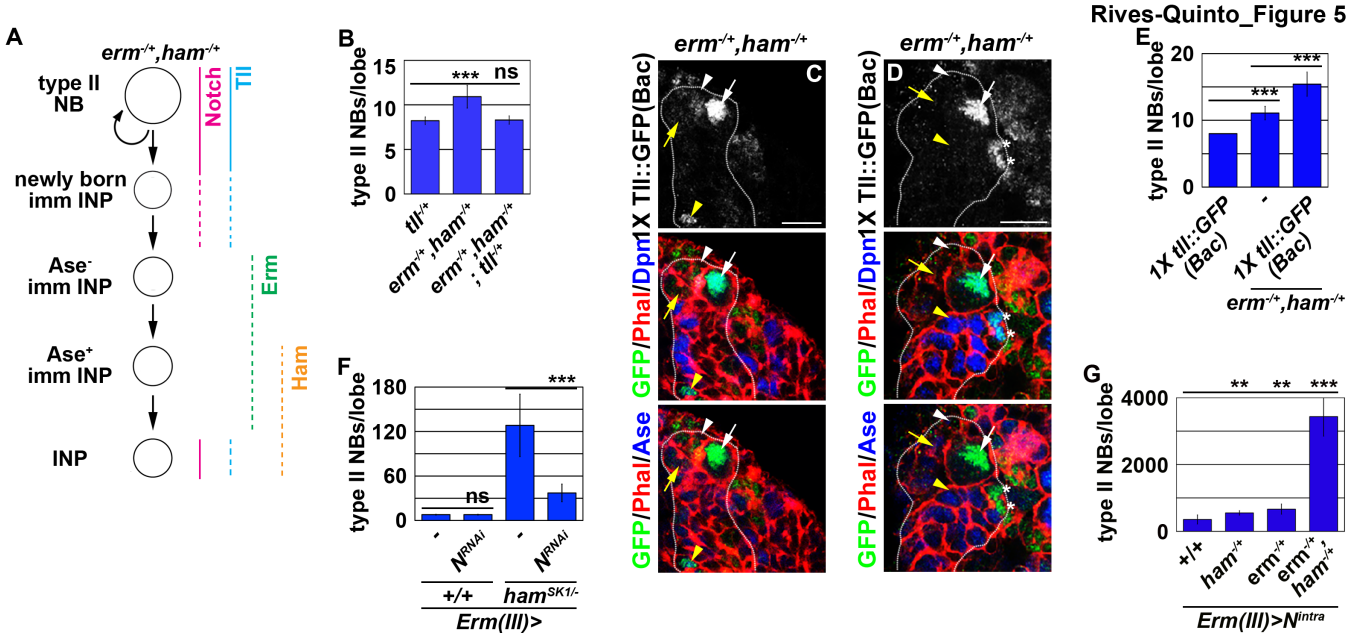

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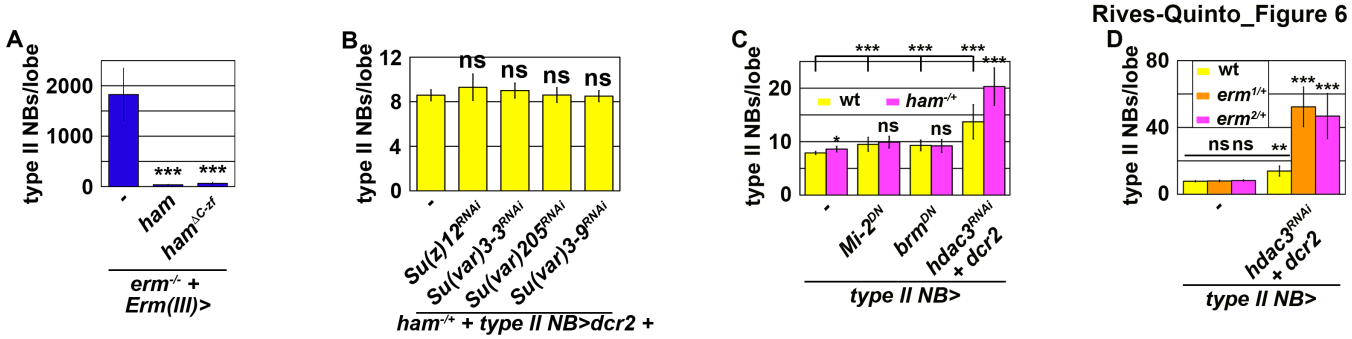
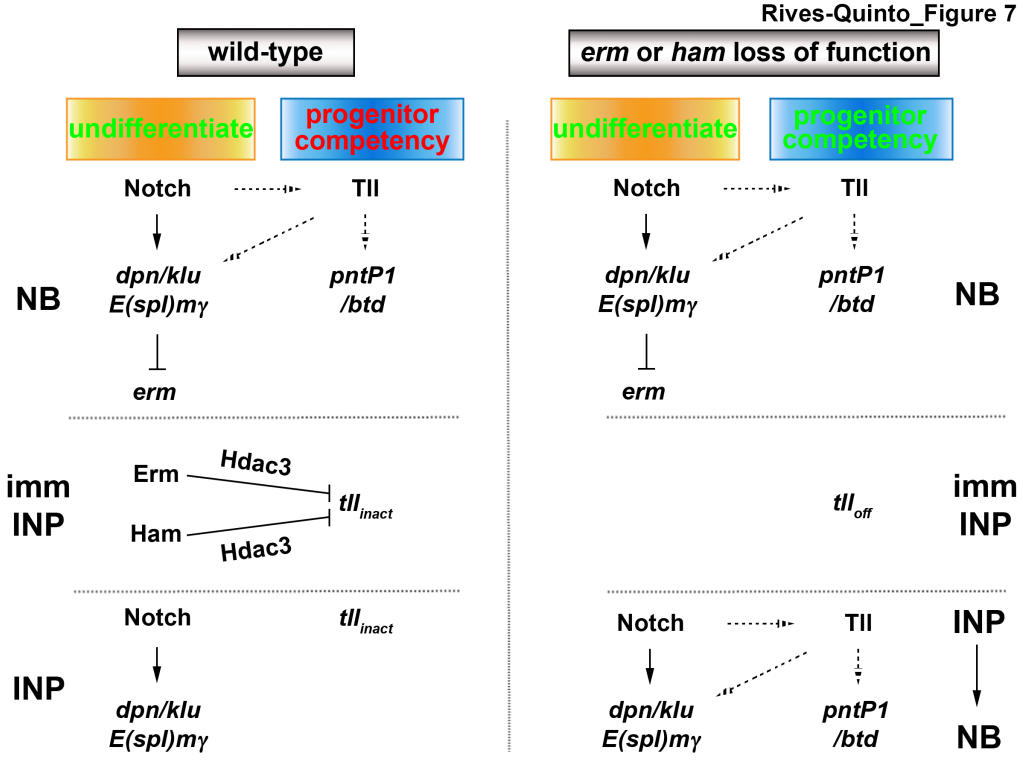


Rives-Quinto_Figure 7



Rives-Quinto_Figure S1

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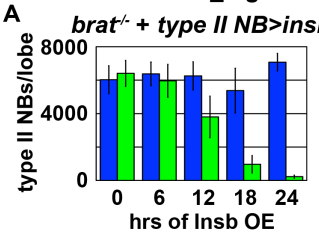

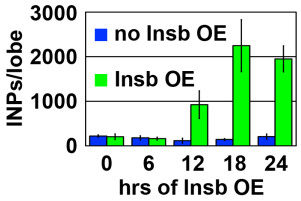

Rives-Quinto_Figure S2

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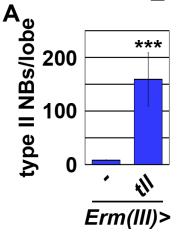
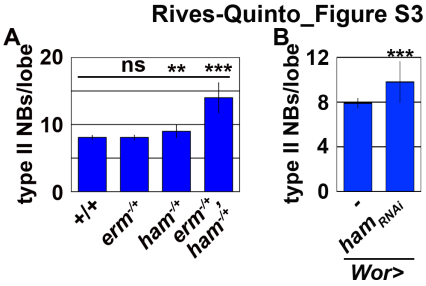
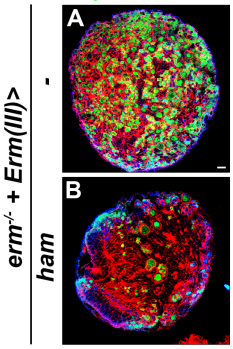


Rives-Quinto_Figure S3



Rives-Quinto_Figure S4

Dpn/Phal/Ase