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## Sgt1 acts via an LKB1/AMPK pathway to establish cortical polarity in larval neuroblasts

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### Abstract

*Drosophila* neuroblasts are a model system for studying stem cell self-renewal and the establishment of cortical polarity. Larval neuroblasts generate a large apical self-renewing neuroblast, and a small basal cell that differentiates. We performed a genetic screen to identify regulators of neuroblast self-renewal, and identified a mutation in *sgt1* (*suppressor-of-G2-allele-of-skp1*) that had fewer neuroblasts. We found that *sgt1* neuroblasts have two polarity phenotypes: failure to establish apical cortical polarity at prophase, and lack of cortical Scribble localization throughout the cell cycle. Apical cortical polarity was partially restored at metaphase by a microtubule-induced cortical polarity pathway. Double mutants lacking Sgt1 and Pins (a microtubule-induced polarity pathway component) resulted in neuroblasts without detectable cortical polarity and formation of “neuroblast tumors.” Mutants in *hsp83* (encoding the predicted Sgt1-binding protein Hsp90), *LKB1*, or *AMPK $\alpha$*  all show similar prophase apical cortical polarity defects (but no Scribble phenotype), and activated AMPK $\alpha$  rescued the *sgt1* mutant phenotype. We propose that an Sgt1/Hsp90-LKB1-AMPK pathway acts redundantly with a microtubule-induced polarity pathway to generate neuroblast cortical polarity, and the absence of neuroblast cortical polarity can produce neuroblast tumors.

### INTRODUCTION

The precise regulation of stem cell self-renewal versus differentiation is essential for normal development, required for tissue homeostasis, and may suppress tumorigenesis. Despite its importance, the molecular mechanisms regulating stem self-renewal are only beginning to be uncovered. Recently, *Drosophila* larval neuroblasts have proven to be an effective model for characterizing the mechanisms regulating stem cell self-renewal (Doe, 2008; Januschke and Gonzalez, 2008).

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*Drosophila* larval neuroblasts undergo repeated asymmetric cell divisions that involve formation of molecularly distinct apical and basal cortical domains, and alignment of the mitotic spindle along the apical/basal polarity axis. The apical cortex contains two protein complexes: the Par complex (Bazooka, Baz; atypical protein kinase C, aPKC; and Partitioning defective-6, Par-6) and the Pins complex (Partner of Inscuteable, Pins; Gai/o, and Discs large,Dlg). These two complexes are thought to be linked by the protein Inscuteable (Insc) (Doe, 2008; Knoblich, 2008). The differentiation factors Numb, Brain tumor (Brat), and Prospero (Pros) accumulate on the basal surface of neuroblasts; the adaptor protein Miranda (Mira) is required for both Brat and Pros localization (Doe, 2008; Knoblich, 2008). In larval neuroblasts, apical polarity is first established at late G2/early prophase, whereas basal cortical polarity is first detectable at the prophase/metaphase transition (Siller et al., 2006). The apical domain is partitioned into the larger daughter cell which self-renews as a neuroblast, while the basal domain segregates into the smaller daughter cell which has a more restricted developmental potential (Doe, 2008; Knoblich, 2008).

Despite the importance of the apical polarity proteins in regulating neuroblast self-renewal, it remains unknown exactly how the Par and Pins complexes are initially localized to the apical cortex. The conserved polarity protein Cdc42 binds Par-6 and is required for apical localization of Par-6 and aPKC, but *cdc42* mutants still localize Baz to the apical cortex, albeit weakly (Atwood et al., 2007). The only mutant reported to abolish Baz apical localization is *lkb1* (Bonaccorsi et al., 2007), which encodes a serine/threonine kinase in the Par-4 kinase family; however, the authors only observed weak Baz localization in wild type neuroblasts, so the significance of the mutant phenotype is difficult to interpret.

Here we describe a forward genetic screen that identified a mutant with a reduced number of brain neuroblasts and defects in the earliest steps of generating neuroblast cortical polarity. We mapped the mutant to a region of the third chromosome and used a novel method to identify the mutant lesion, revealing a small deletion in the *sgt1* gene. Sgt1 is an evolutionarily conserved protein that directly interacts with Hsp90 using its “CHORD/Sgt1” domain (Lee et al., 2004); it regulates kinetochore assembly in yeast and mammalian cells; cell cycle progression in yeast, mammals, and *Drosophila*; and pathogen sensing in plant and animal cells (da Silva Correia et al., 2007; Kitagawa et al., 1999; Martins et al., 2009; Mayor et al., 2007; Shirasu, 2009; Steensgaard et al., 2004).

## RESULTS

### A genetic screen for neuroblast self-renewal mutants identifies *sgt1*

To find potential regulators of neuroblast self-renewal, we screened homozygous P-element induced mutations for altered larval neuroblast numbers, using Miranda (Mira) and Deadpan (Dpn) to identify neuroblasts and Scribble (Scrib) to outline the neuroblast cell cortex and highlight general neuroanatomical landmarks (Lee et al., 2006c) (Figure 1A). One mutant that showed a modest decrease in neuroblast number, and larval/pupal lethality, was *l(3)s2383* which contains a P element at 66E1-66E2 on the left arm of the third chromosome (Figure 1A,B). To confirm that the P element insertion caused the phenotype, we assayed the *l(3)s2383* chromosome *in trans* to a deficiency that removed chromosomal region

66E1-66E2. Surprisingly, these larvae were completely normal for neuroblast number and viability, showing that the mutation mapped to elsewhere on the third chromosome. We used deficiency mapping to localize the mutation to the 84F6-84F13 region of the right arm of the third chromosome (Figure 1B). We found that *l(3)s2383 / Df(3R)6147* larvae showed the same neuroblast number and larval/pupal lethality phenotypes as *l(3)s2383* homozygotes. Phenotypic analysis of overlapping deficiencies in the region allowed us to map the mutation to a 60kb interval, and we confirmed the mutation was in this interval by genomic DNA rescue of the phenotype (data not shown; and see Materials and Methods).

To identify the DNA lesion responsible for the phenotype, we used a novel “sequence capture” strategy combined with Illumina deep sequencing (Figure 1B) to identify an in frame 15 nucleotide deletion in the first exon of the *sgt1* gene, also known as *CG9617* (Figure 1C; and see Materials and Methods). We henceforth call this allele *sgt1<sup>s2383</sup>*. To confirm that the *sgt1<sup>s2383</sup>* lesion is responsible for the observed neuroblast and viability phenotype, we crossed *sgt1<sup>s2383</sup>* with the known *sgt1<sup>c01428</sup>* mutation (Martins et al., 2009). We found that *sgt1<sup>s2383</sup>* fails to complement *sgt1<sup>c01428</sup>*, and that *sgt1<sup>s2383</sup> / sgt1<sup>c01428</sup>* larvae have the same phenotype as *sgt1<sup>s2383</sup>* homozygotes. Furthermore, the fact that the phenotype of *sgt1<sup>s2383</sup> / sgt1<sup>c01428</sup>* is identical to that of *sgt1<sup>s2383</sup> / Df(3R)6147* indicated that *sgt1<sup>s2383</sup>* is a strong or null allele for CNS function. The lesion in *sgt1<sup>s2383</sup>* resulted in a five amino acid deletion in the Chord and Sgt1 (CS) domain that mediates Sgt1-Hsp90 protein interactions in other organisms (Zhang et al., 2010). We conclude that *Drosophila* Sgt1 is required for maintaining normal numbers of larval neuroblasts and for viability.

### ***sgt1* is required for apical protein localization in prophase neuroblasts**

To investigate the cellular origin of the neuroblast depletion phenotype in *sgt1* mutants, we assayed neuroblast cell cycle progression and neuroblast cell polarity. Prophase neuroblasts were identified as PH3+ without a bipolar spindle; metaphase neuroblasts were identified as PH3+ with a bipolar spindle. We confirmed a previous report (Martins et al., 2009) that *sgt1* mutants have slower cell cycle progression, cytokinesis failure, polyploidy, multiple centrosomes, and malformed mitotic spindles (Supplemental Figure 1 and data not shown). In this paper, we focus on the previously unreported cell polarity aspects of the *sgt1* mutant phenotype. First, we assayed neuroblast apical cortical polarity (“apical cortex” is defined as the cortical domain that is partitioned into the sibling neuroblast, whereas “basal cortex” is defined as the domain segregated into the GMC). The first sign of neuroblast asymmetry occurs during late G2 and early prophase when Par complex proteins (e.g. Baz, aPKC), Pins complex proteins (e.g. Pins), and the Par/Pins linker protein Insc are localized to an apical cortical crescent (Figure 2A; quantified in A’). In contrast, *sgt1* mutant prophase neuroblasts typically showed cytoplasmic or undetectable localization of these proteins (Figure 2B; quantified in B’). Interestingly, during metaphase we found a substantial rescue of apical cortical polarity (Figure 2C,D; quantified in C’,D’); the basal proteins Miranda and Numb were also fairly normal at metaphase (Supplemental Figure 2). We conclude that Sgt1 is required for establishing apical cortical polarity during prophase, and during metaphase an Sgt1-independent pathway can generate apical/basal cortical protein localization.

We next analyzed Dlg and Scrib cortical localization in wild type and *sgt1* mutant neuroblasts, for two reasons. First, we had observed a complete lack of Scrib cortical localization in our original screen (Figure 1A), and second, Dlg is part of the Pins complex (Siegrist and Doe, 2005) so we wanted to know if it was delocalized in prophase neuroblasts similar to Pins. In wild type interphase and mitotic neuroblasts, Scrib was always uniform cortical (Figure 3A; 100% uniform cortical, n=50), while Dlg was invariably uniform cortical throughout the cell cycle plus enriched with Pins at the apical cortex during mitosis (Figure 3B; 100% uniform cortical n=53, 94% apical enriched, n=34). In contrast, *sgt1* mutant neuroblasts showed a strong loss of cortical Scrib localization during metaphase (Figure 3C; 3% cortical, n=75) and interphase (Figure 1A, and data not shown); they also showed a loss of the uniform cortical pool of Dlg protein at interphase and mitosis, with retention of the pool of Dlg asymmetrically co-localized with Pins at the apical cortex at metaphase (Figure 3D; 2% uniform cortical n=45, 83% apical enriched, n=29). We conclude that Sg1 is essential for targeting Dlg/Scrib to the neuroblast cortex, except where Dlg is co-localized with Pins.

Martins et al. (2009) showed that overexpression of wild type Polo kinase can rescue aspects of the *sgt1* neuroblast cell cycle defects; we found that Polo overexpression increases the number of mitotic neuroblasts in both wt and *sgt1* mutant brains (Supplemental Figure 1B), but does not rescue neuroblast prophase cortical polarity (7% normal Pins, n=45; 4% normal aPKC, n=27) or Scrib cortical localization (0% normal Scrib, n=75).

### Microtubules induce apical cortical polarity in *sgt1* metaphase neuroblasts

What is the Sg1-independent pathway that generates Pins and Par complex apical cortical localization in metaphase neuroblasts? We previously defined a microtubule-induced cortical polarity pathway that could induce Pins/Gai/Dlg cortical polarity (Siegrist and Doe, 2005), so here we test whether this pathway induces Pins and Par complex apical cortical polarity at metaphase in *sgt1* mutant neuroblasts. We used the microtubule inhibitor colcemid to abolish spindle microtubules in *sgt1* mutant and control neuroblasts. Wild type neuroblasts treated with the colcemid retain apical cortical crescents of Dlg, Pins and Insc (Figure 4B, top row; quantified in figure legend). Strikingly, *sgt1* mutant neuroblasts treated with colcemid lack all detectable apical cortical polarity (Figure 4B, bottom row; quantified in figure legend). We conclude that microtubules are required to establish apical cortical polarity in the absence of Sg1.

### *pins sgt1* double mutants neuroblasts lack all detectable cortical polarity and form neuroblast brain “tumors”

The developmental consequence of a complete loss of neuroblast polarity has never been assayed, and the loss of both Sg1-dependent and microtubule-dependent cortical polarity pathways is predicted to generate a completely apolar neuroblast (see above). We can not assay sibling cell fate in *sgt1*-colcemid-treated neuroblasts because they are arrested in mitosis, so we examined a double mutant lacking Sg1 and the microtubule-induced polarity component Pins. We tested (a) whether *sgt1 pins* double mutants abolish both Par and Pins complex polarity, and if so (b) what is the fate of such apolar neuroblasts. We found that *sgt1 pins* double mutant neuroblasts establish little or no apical or basal cortical polarity

(Figure 5B and data not shown). The developmental consequences of this loss of polarity are the formation of ectopic neuroblasts throughout the brain (Figure 5F). This is in striking contrast to the *sgt1* and *pins* single mutant brains, which are each smaller with reduced neuroblast numbers (Figure 5D,E). We propose that loss of all cortical polarity in these double mutants results in symmetric neuroblast division to form two neuroblasts and expand the pool of brain neuroblasts (see Discussion).

### Reducing Hsp90 levels mimics the *sgt1* mutant phenotype

To determine the mechanism by which Sgt1 promotes apical cortical polarity and Dlg/Scrib cortical localization, we started by determining if Sgt1 worked together with its evolutionarily-conserved binding partner, Hsp90. Sgt1 directly interacts with Hsp90 via its CS domain in all organisms tested (Catlett and Kaplan, 2006; Lee et al., 2004; Lingelbach and Kaplan, 2004; Nony et al., 2003), and our *sgt1*<sup>s2383</sup> allele generates an Sgt1 protein lacking 5 amino acids within CS domain (Figure 1C). We obtained mutants in *hsp83* (which encodes the Hsp90 protein) and examined the larval brain neuroblast phenotype in *hsp83*<sup>13F3</sup>/*hsp83*<sup>582</sup> transheterozygotes (Lange et al., 2000). We find that *hsp83* mutants have neuroblast phenotypes that are similar to that of *sgt1* mutants: loss of apical cortical polarity at prophase (Figure 6B,C; quantified in Figure 6G). However, there is no loss of cortical Scrib in *hsp83* mutants (Figure 6C), suggesting that Sgt1 promotes Dlg/Scrib localization by an Hsp90-independent mechanism (see Discussion). The similarity of the *sgt1* and *hsp83* apical cortical polarity phenotypes, and the fact that the Sgt1<sup>s2383</sup> protein has a deletion within the Hsp90-binding CS domain, leads us to hypothesize that Sgt1/Hsp90 act together to promote apical cortical polarity in prophase neuroblasts.

### Sgt1 acts via an LKB1/AMPK pathway to generate apical cortical polarity, but not for Scrib cortical localization

Sgt1 has been shown to act through Polo kinase to regulate neuroblast cell cycle progression and cytokinesis (Martins et al., 2009). We confirmed that misexpression of Polo rescues the neuroblast cell cycle phenotypes, but found no rescue of the neuroblast cortical polarity phenotypes (data not shown). Thus, we investigated other downstream effectors of Sgt1 for a role in establishing apical protein localization.

*lkb1* mutants are reported to have defects in neuroblast apical polarity that are similar to the *sgt1* mutant phenotype (Bonaccorsi et al., 2007). We confirm that *lkb1* mutants had a nearly identical neuroblast polarity phenotype as *sgt1* mutants, with the striking exception of showing normal Scrib cortical localization (Figure 6D; quantified in Figure 6G). In addition, *lkb1* mutants – similar to *sgt1* mutants – have slower cell cycle progression, cytokinesis failure, polyploidy, multiple centrosomes, and malformed mitotic spindles (Bonaccorsi et al., 2007)(data not shown). We conclude that Sgt1 and LKB1 are both required to establish apical cortical polarity in neuroblasts.

LKB1 kinase can activate multiple members of the AMP-activated kinase (AMPK) family, such as Par-1, AMPK, Sik, NUA, and others (Lizcano et al., 2004). Both Par-1 and AMPK activity are required for generating cell polarity in multiple cell types and organisms (Goldstein and Macara, 2007; Williams and Brenman, 2008), so we tested whether Par-1 or

AMPK functions downstream of Sgt1/LKB1 in generating apical localization of Par complex proteins in neuroblasts. First we examined the localization and function of Par-1; we observed high levels of the protein in cortex glia that ensheath the neuroblast, but no sign of polarized localization within the neuroblast and no change in neuroblast polarity in *par-1* mutant MARCM clones (data not shown). We conclude that loss of Par-1 from larval neuroblasts has no detectable effect on cell polarity.

We next assayed *AMPK $\alpha$*  mutants, and found that they – like *sgt1* and *lkb1* mutants – showed a loss of apical protein localization at prophase (Figure 6E; quantified in Figure 6G), slower cell cycle progression, cytokinesis failure, polyploidy, multiple centrosomes, and malformed mitotic spindles (Lee et al., 2007)(data not shown). As with *lkb1* mutants, we saw normal cortical localization of Scrib, confirming that Sgt1 acts in an LKB1/AMPK-independent pathway to promote Scrib cortical localization. Importantly, we found that expression of constitutively activated AMPK $\alpha$  (Lee et al., 2007) (*tub-gal4 UAS-ampk $\alpha$ <sup>TD</sup>*) substantially rescued the *sgt1* mutant phenotype (Figure 6F; quantified in Figure 6G). We conclude that an Sgt1 – Hsp90 – LKB1 – AMPK pathway is used to establish apical cortical polarity in prophase neuroblasts via an unknown link between AMPK $\alpha$  and the apical Par complex, whereas an Sgt1-specific pathway is used to establish corticalDlg/Scrib localization (Figure 6H).

The AMPK pathway promotes myosin contractility in epithelia, and an activated myosin regulatory light chain, called Sqh in Drosophila (Sqh<sup>20E21E</sup>), can rescue the *ampka* embryonic epithelial polarity phenotype (Lee et al., 2007; Mirouse et al., 2007). Thus, we tested if active Sqh<sup>21E</sup> (Jordan and Karess, 1997) could rescue the *sgt1* neuroblast cortical polarity phenotype, or whether loss of *sgt1*, *lkb1* or *ampka* altered cortical myosin activity in larval neuroblasts. First, we found that expression of activated Sqh<sup>21E</sup> or the non-activatable Sqh<sup>20A21A</sup> (Jordan and Karess, 1997; Lee et al., 2007) both failed to rescue the *sgt1* neuroblast cortical polarity phenotype (Supplementary Figure 3A). Second, we found that all three pathway mutants (*sgt1*, *lkb1*, and *ampka*) showed an increase in uniform or patchy cortical activated Sqh<sup>20E21E</sup> (Supplemental Figure 3B-E). In addition, time-lapse imaging of *sgt1* mutant larval neuroblasts showed highly abnormal cortical protrusions, misshapen cells, and failure of cytokinesis (Supplemental Figure 3F, Movie 1); these phenotypes are never seen in wild type larval neuroblasts (Cabernard and Doe, 2009; Cabernard et al., 2010). The failure to execute cytokinesis is consistent with a loss of myosin activity, whereas the elevated cortical phospho-Sqh is consistent with an increase in myosin activity; thus all three pathway mutants (*sgt1*, *lkb1*, and *ampka*) have defects in the regulation of cortical myosin activity. However, the lack of cortical polarity phenotype in neuroblasts homozygous for *zipper*, which encodes myosin heavy chain (Barros et al., 2003; Peng et al., 2000), and the inability of Sqh<sup>21E</sup> to rescue the cortical polarity phenotype (this work) suggests that simple loss of myosin activity can not explain the *sgt1*, *lkb1*, or *ampka* mutant neuroblast polarity defects (Figure 6H).

## DISCUSSION

Here we present evidence that the evolutionary-conserved protein Sgt1 acts with Hsp90, LKB1 and AMPK to promote apical localization of the Par and Pins complexes in prophase

neuroblasts. We propose that Sgt1/Hsp90 proteins function together based on multiple lines of evidence: (1) they show conserved binding from plants to humans (Catlett and Kaplan, 2006; Lee et al., 2004; Lingelbach and Kaplan, 2004; Nony et al., 2003); (2) our *sgt1*<sup>s2383</sup> mutant results in a five amino acid deletion within the CS domain, which is the Hsp90 binding domain (Catlett and Kaplan, 2006; Zhang et al., 2008); (3) *sgt1* and *hsp83* have similar cell cycle phenotypes (Supplemental Figure 1)(Lange et al., 2000); and (4) *sgt1* and *hsp83* have similar neuroblast polarity phenotypes (Figure 6B,C). The Sgt1/Hsp90 complex either stabilizes or activates client proteins (Zuehlke and Johnson, 2010); we suggest that Sgt1 activates LKB1, rather than stabilizing it, because we were unable to rescue the *sgt1* mutant phenotype by simply overexpressing wild type LKB1 protein (data not shown). We have not tested for direct interactions between Sgt1 and LKB1 proteins, and thus the mechanism by which Sgt1 activates LKB1 remains unknown.

LKB1 is a “master kinase” that activates at least 13 kinases in the AMPK family (Lizcano et al., 2004). We suggest that LKB1 activates AMPK to promote neuroblast polarity because overexpression of phosphomimetic, activated AMPK $\alpha$  can rescue the *lkb1* and *sgt1* mutant phenotype (Figure 6F and data not shown). It remains unclear how AMPK activity promotes apical protein localization. An antibody to activated AMPK $\alpha$  (anti-phosphoT385-AMPK $\alpha$  , Mirouse et al., 2007) shows spindle and cytoplasmic staining that is absent in *ampka* mutants, and centrosomal staining that persists in AMPK $\alpha$  null mutants, but no sign of asymmetric localization in neuroblasts (data not shown). AMPK activity is thought to directly (Lee et al., 2007) or indirectly (Bultot et al., 2009) activate myosin regulatory light chain to promote epithelial polarity (Lee et al., 2007; Miranda et al., 2010). AMPK is activated by a rise in AMP/ATP levels that occur under energy stress or high metabolism; AMP binds to the  $\gamma$  regulatory subunit of the heterotrimeric complex and results in allosteric activation of the  $\alpha$  subunit (Hardie et al., 2007). *ampka* mutants grown under energy stress have defects in apical/basal epithelial cell polarity in follicle cells within the ovary (Mirouse et al., 2007). In contrast, AMPK $\alpha$  mutants grown on nutrient rich food still show defects in embryonic epithelial polarity (Lee et al., 2007), neuroblast apical polarity (this work), and visceral muscle contraction (Bland et al., 2010). Larval neuroblasts, embryonic ectoderm, and visceral muscle may have a high metabolic rate, require low basal AMPK activity, or use a different mechanism to activate AMPK than epithelial cells. What are the targets of AMPK signaling for establishing apical cortical polarity in larval neuroblasts? AMPK could directly phosphorylate Baz to destabilize the entire pool of apical proteins, but currently we have no evidence supporting such a direct model. AMPK may act via regulating cortical myosin activity: we have seen clear defects in cortical motility, ectopic patchy activated myosin at the cortex, and failure of cytokinesis in *sgt1*, *lkb1*, and *ampka* mutants (Movie 1; Supplemental Figure 3). This strongly suggests defects in the regulation of myosin activity, but how or if gain/loss/mispositioning of myosin activity leads to failure to establish apical cortical polarity remains unknown. Lastly, the defects in apical cell polarity seen at prophase could be due to the prometaphase cell cycle delays.

What activates the Sgt1-LKB1-AMPK pathway to promote cell polarity during prophase? In budding yeast, Sgt1 requires phosphorylation on Serine 361 (which is conserved in

Drosophila Sgt1) for dimerization and function (Bansal et al., 2009); this residue is conserved in Drosophila Sgt1 but its functional significance is unknown.

Sgt1/Hsp90/LKB1/AMPK are all required for apical Par/Pins complex localization, but Sgt1 must act via a different pathway to promote Dlg/Scrib cortical localization, because only the *sgt1* mutant affects Dlg/Scrib localization, and overexpression of activated AMPK $\alpha$  is unable to restore cortical Scrib in *sgt1* mutants (Figure 6). The mechanism by which Sgt1 promotes Dlg/Scrib cortical localization is unknown.

We have shown that *sgt1* mutants lack Par/Pins apical polarity in prophase neuroblasts, but these proteins are fairly well polarized in metaphase neuroblasts. The rescue of cortical polarity is microtubule dependent, probably occurring via the previously described microtubule-dependent cortical polarity pathway containing Pins, Dlg and Khc-73 (Siegrist and Doe, 2005). The weak polarity defects still observed in *sgt1* metaphase neuroblasts may be due to the poor spindle morphology (Supp. Figure 1). The lack of microtubule-induced polarity at prophase, despite a robust microtubule array in prophase neuroblasts, suggests that the microtubule-induced cortical polarity pathway is activated at metaphase. Activation of the pathway could be via expression of the microtubule-binding protein Khc-73; via phosphorylation of Pins, Dlg or Khc-73 by a mitotic kinase like Aurora A (Johnston et al., 2009); or via a yet unknown pathway.

It was somewhat surprising that the *sgt1 pins* double mutants had increased numbers of brain neuroblasts, because each single mutant had reduced neuroblast numbers (this work) (Lee et al., 2006b). The double mutant phenotype may be due to loss of both Pins and cortical Dlg/Scrib, as the *sgt1 pins* double mutant phenotype is similar to the *dlg pins* double mutant phenotype (data not shown). It could also be due to a change in an unknown downstream effector of both Sgt1 and Pins. A not mutually exclusive possibility is that the *sgt1 pins* double mutant phenotype is due to loss of all Par/Pins cortical polarity. This model is consistent with our observation that *sgt1* or *pins* single mutants retain some neuroblast cortical polarity, whereas the *sgt1 pins* double mutants lack all known neuroblast cortical polarity. We propose that the apolar double mutant neuroblasts partition cell fate determinants equally to both siblings, and that both siblings frequently assume a neuroblast identity. This is supported by our recent finding that when the neuroblast spindle is aligned orthogonal to a normal apical/basal polarity axis, such that both siblings inherit equal amounts of apical cortical proteins, the siblings always acquire a neuroblast identity (Cabernard and Doe, 2009). Thus, equal partitioning of apical/basal cell fate determinants (in spindle orientation mutants) or failure to establish any cortical polarity (*sgt1 pins* mutants) may result in neuroblast/neuroblast siblings and an expansion of the neuroblast population.

## METHODS

### Identification and sequencing of the *sgt1*<sup>S2383</sup> mutant

The *sgt1*<sup>S2383</sup> mutant allele was originally found in a P-element screen of the 3rd chromosome. The P-element mapped to 66E1-2, but deficiency mapping revealed that the lethality and CNS phenotype mapped to *Df(3R)6147* at 84F6-F13. To further define the



region, rescue constructs were made using gap-repair and  $\Phi$ C31 mediated transgenesis (Venken et al., 2009). Using the tilling BAC 33N15, we made two overlapping 60kb and 28kb rescue constructs; only the 60kb construct rescued the *l(3)s2383* phenotype, narrowing the relevant region to approximately 13 genes. To identify the lesion in this interval, we used a biotin mediated “sequence capture” technique followed by deep sequencing. Genomic DNA from both wild type and mutant larva were prepared for deep sequencing. Using BAC33N15 as a template, biotin incorporated “probes” were created. The biotin labeled probes were individually hybridized with wild type or mutant DNA and purified over streptavidin beads. The genomic DNA was then eluted from the beads, and sequenced using Illumina deep sequencing, revealing a 15 nucleotide in frame deletion within the first exon of *sgt1<sup>s2383</sup>*; this allele will be called *sgt1* for simplicity. In addition, a P element allele of *sgt1* (C01428; from the Harvard Exelixis collection) failed to complement and showed the same phenotype as *sgt1<sup>s2383</sup>*.

### Fly Stocks

*Df(3R)6147* was obtained from the Bloomington stock center. The *lkb1<sup>4A4-2</sup>* and *lkb1<sup>4A4-11</sup>* mutant stocks (Mirouse et al., 2007) were kindly provided by Daniel St. Johnston (Cambridge, UK); the strong or null *ampk $\alpha$*  mutants *ampk<sup>1</sup>*, *ampk<sup>2</sup>* and the *UAS-GFP-lkb1* and *UAS-ampk $\alpha$  (T-D)* stocks were gifts from Jay Brennan (Chapel Hill, NC); *UAS-apkcRNAi* was obtained from the Vienna Stock Center. *pins<sup>p62</sup>* was obtained from Bill Chia (Singapore); *hsp83<sup>13F3</sup>* and *hsp83<sup>582</sup>* were obtained from Howard Lipshitz (Toronto); *sqh<sup>21E</sup>*, *sqh<sup>20A21A</sup>*, *sqh<sup>21A</sup>* were obtained from the Bloomington stock center; *UAS-polo* was obtained from Claudio Sunkel (Porto, Portugal), and *worniu-gal4 UAS-miranda:GFP* (Cabernard and Doe, 2009) was used for live imaging. The *sgt1<sup>s2383</sup> pins<sup>p62</sup>* double mutants were generated by recombination. *insc-gal4* (aka *1407-gal4*), *worniu-gal4*, and *tubulin-gal4* were all used for rescue experiments. Mutant larva were identified by lack of balancer chromosome markers *Tubby* or GFP.

### Antibody staining, drug treatment, and imaging

Third instar larval brains were dissected and fixed as previously described (Lee et al., 2006a). Neuroblast counting and BrdU labeling were done as previously described (Lee et al., 2006a). Dissected brains were washed several times in PBS-T (PBS with 0.1% TritonX-100) and blocked for 1hr in PBS-BT (PBS with 0.1% Triton and 1% BSA). Brain preparations were incubated overnight at 4C with primary antibodies diluted in PBS-BT. All analyses were performed with the mitotic markers rat anti- $\alpha$ -tubulin (1:3000; Serotec, Kidlington, Oxford, UK) or rabbit anti-phospho-histone H3 (1:1000, Upstate, West Grove, PA) and one or more of the following antibodies: guinea pig anti-Bazooka (1:1000, Doe lab), rabbit anti-aPKC (1:1000; Sigma, St. Louis, MO), rabbit anti-Par-6 (1:100; Doe lab), rat anti-Pins (1:500; Doe lab), rabbit anti-Inscuteable (1:500; Bill Chia), rabbit anti-Gai (1:500; Doe lab), mouse anti-Discs Large (1:500; Developmental Hybridoma Studies Bank [DHSB], Iowa), anti-rabbit Scribble (1:5000; Doe lab), guinea pig anti-Miranda (1:400; Doe lab), rabbit anti-Miranda (1:1000; Doe lab), rabbit anti-phosphoT385AMPK $\alpha$  (1:100; Cell Signaling, Beverly, MA), anti-phospho-Ser19 human MRLC (Cell Signaling, no. 3671 1:100 to detect phospho-Ser22 Sqh (Myosin regulatory light chain), and guinea pig anti-Numb (1:100; Jim Skeath). After 3 washes with PBS-BT the samples were incubated for 2

hours at room temperature with the appropriate secondary antibody (FITC-conjugated IgG, Rhodamine Red-X-conjugated IgG, and Cy5-conjugated IgG; Invitrogen, Eugene, OR). All secondary antibodies were diluted from a 0.5 mg/ml stock solution and used at 1:400. Brains washed in PBT, mounted in VectaShield medium, and examined with a Bio-Rad Radiance 2000 confocal microscope using a 63X 1.4NA oil immersion objective or a Zeiss LSM 700 confocal microscope using a 40X 1.3NA oil immersion objective. Image processing was performed using ImageJ (NIH) and Photoshop (Adobe), and figures were assembled in Illustrator (Adobe).

To inhibit microtubule formation, dissected wild type and *sgt1* mutant brains were placed in 10 $\mu$ g/ml Colcemid (Sigma, St. Louis, MO) diluted in Schneider's medium (Sigma, St. Louis, MO) for 1hr. Brains were fixed and antibody stained as described above.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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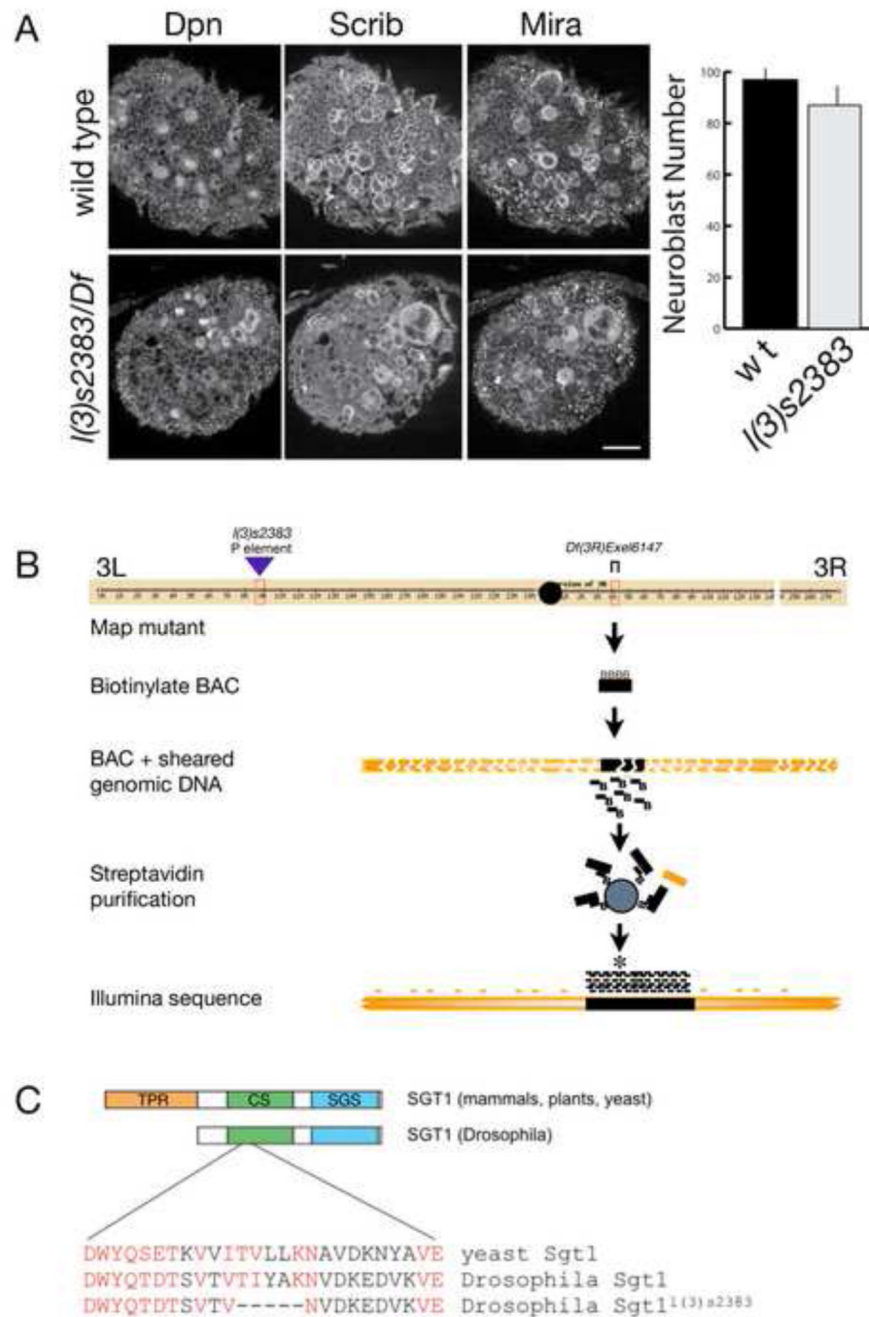
### Highlights

We used a novel sequence-capture genomic method to identify *sgt1*.

*sgt1* is required for Bazooka localization in neuroblasts.

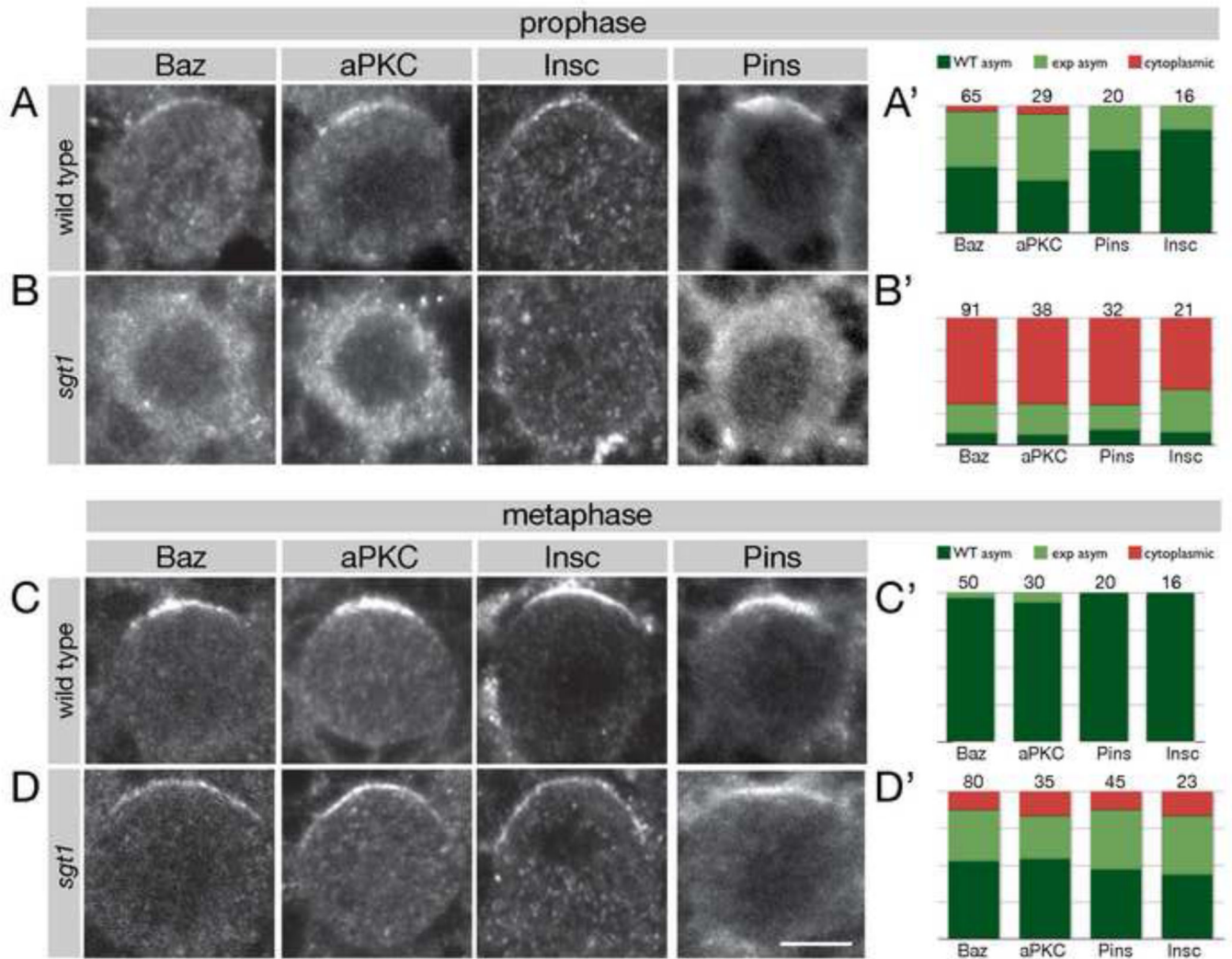
*sgt1* is required for Scribble localization in neuroblasts.

*sgt1* acts at the top of an Hsp90-LKB1-AMPK polarity pathway.



**Figure 1. Identification of a mutant in *sgt1* that is required for larval brain development**  
**(A)** Wild type and *l(3)s2383 / Df(3R)Exel6147* (*l(3)s2383 / Df*) third instar larval brains stained for the neuroblast markers Miranda (Mira) and Deadpan (Dpn) plus Scribble (Scrib) which decorates the cortex of all cells in the brain. Right: histogram of the average neuroblast number per brain lobe (bar, standard deviation) in each genotype. Note that the mutant brains have fewer neuroblasts and Scrib protein is cytoplasmic. **(B)** Schematic of the “sequence capture” and deep sequencing strategy used to identify the lesion in the *l(3)s2383* mutant chromosome. Although the chromosome was generated in a P element mutagenesis,

we found a 15 nucleotide deletion unrelated to the P element insertion that showed the mutant phenotype when transheterozygous to *Df(3R)Exel6147*. (C) The *l(3)s2383* mutation resulted in an in-frame five amino acid deletion in the Drosophila Sgt1 protein, within the N-terminal CS domain (green); the C-terminal Sgt1-specific SGS domain is shown (white).



**Figure 2. *sgt1* mutant neuroblasts have defects in establishing apical cortical polarity**

Wild type and *sgt1*<sup>s2383</sup> / *sgt1*<sup>s2383</sup> (*sgt1*) mutant larval brain neuroblasts stained for the indicated cortical proteins plus  $\alpha$ -tubulin (not shown) and phospho-histone H3 (PH3; not shown). Prophase neuroblasts were identified as PH3+ without a bipolar spindle; metaphase neuroblasts were identified as PH3+ with a bipolar spindle. Scale bar, 5  $\mu$ m.

(A) Wild type prophase neuroblasts show apical enrichment of the Par complex (Baz, aPKC) and Pins complex (Pins) and Insc; quantification in (A').

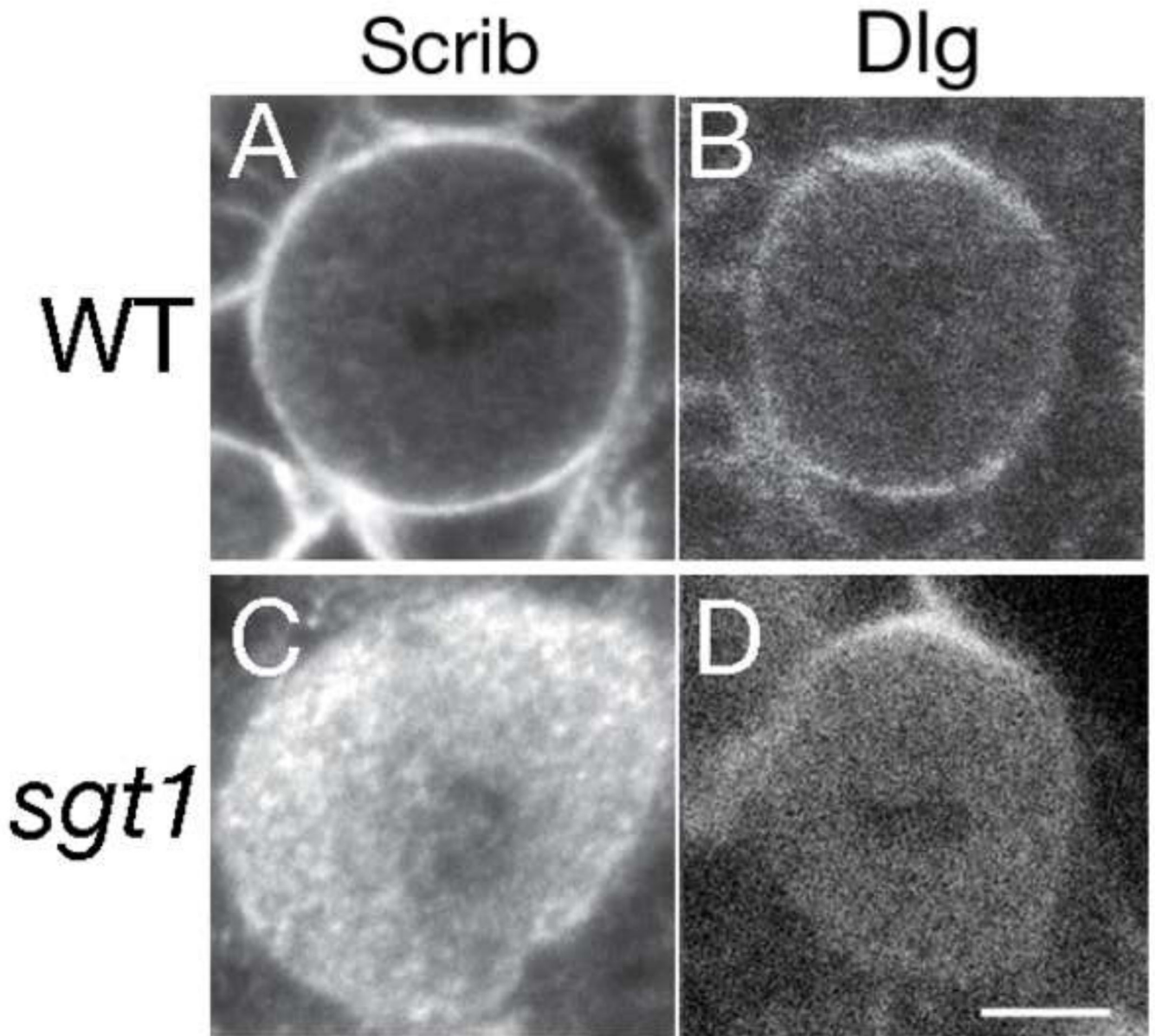
(B) *sgt1* prophase neuroblasts show a nearly complete loss of apical localized Par complex (Baz, aPKC), Pins complex (Pins) and Insc; quantification in (B').

(C) Wild type metaphase neuroblasts show apical enrichment of the Par complex (Baz, aPKC) and Pins complex (Pins) and Insc; quantification in (C').

(D) *sgt1* metaphase neuroblasts have recovered substantial apical protein localization compared to their prophase phenotype, although it is not fully back to wild type levels; quantification in (D').



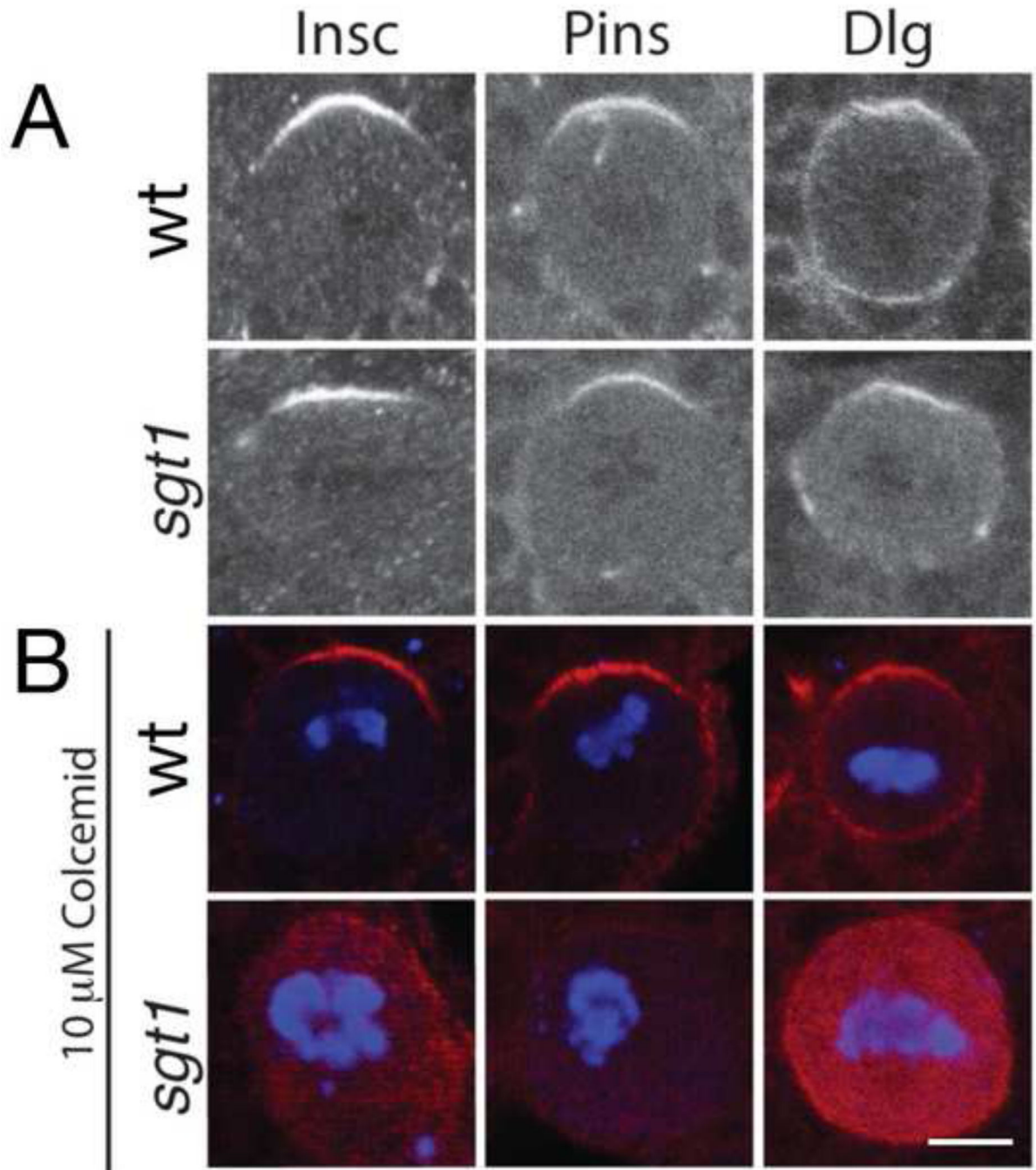
(A'-D') Quantification of the phenotypes shown in A-D. Dark green, normal asymmetric; light green, expanded asymmetric; red, cytoplasmic. Number of neuroblasts scored shown in each bar.



**Figure 3. *sgt1* mutant neuroblasts have defects in Dlg/Scrib cortical localization**

(A, B) Wild type prophase or metaphase larval neuroblasts; Scrib is uniform cortical, and Dlg is uniform cortical with apical enrichment at metaphase.

(C, D) *sgt1<sup>s2383</sup> / sgt1<sup>s2383</sup> (sgt1)* metaphase larval neuroblasts; Scrib is cytoplasmic, and Dlg is mostly cytoplasmic except for persistent apical enrichment at metaphase. Scale bar, 5  $\mu$ m.

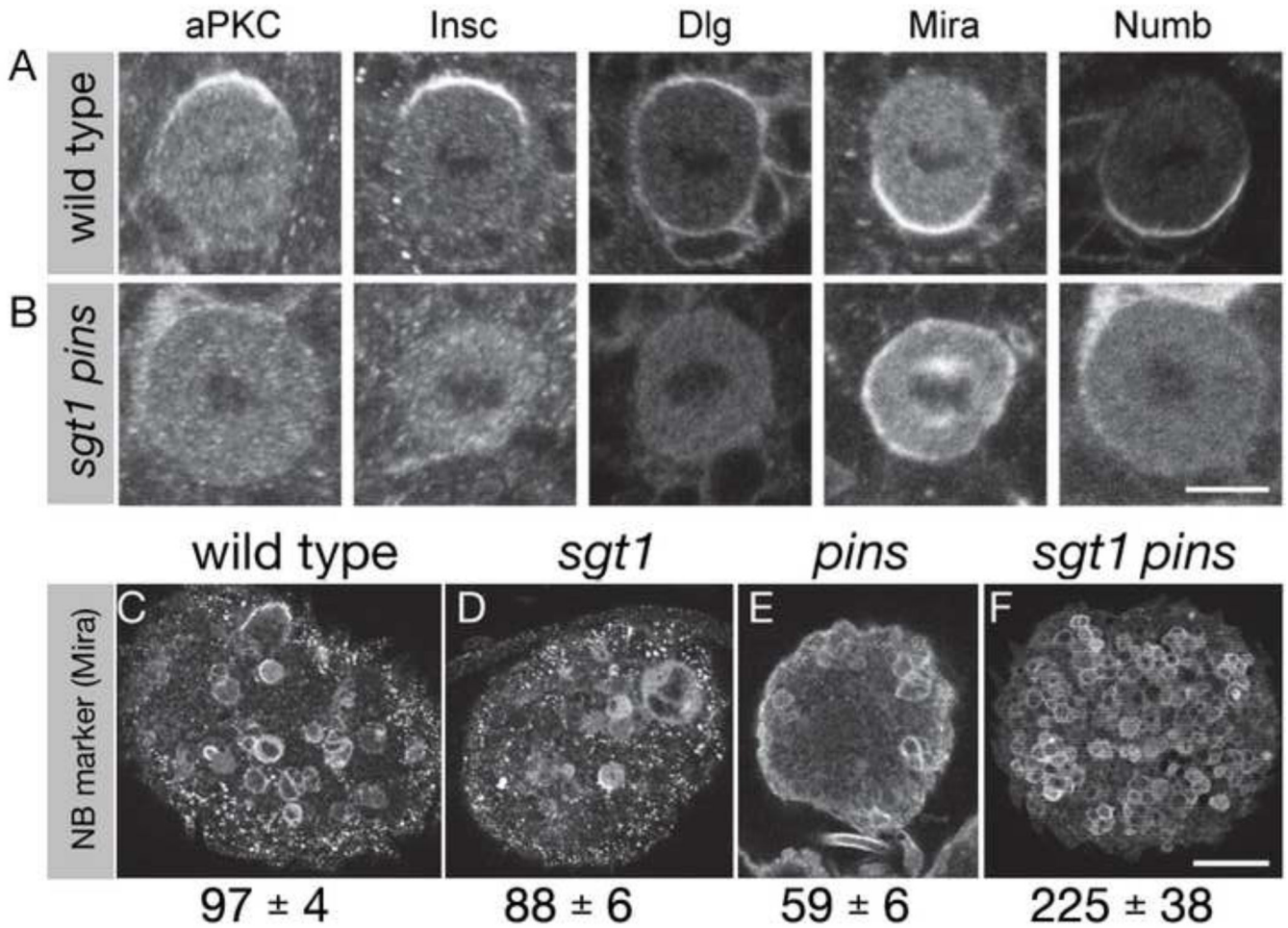


**Figure 4. Sgt1 and microtubules act in redundant pathways to generate apical protein localization in metaphase neuroblasts**

Wild type and *sgt1<sup>s2383</sup> / sgt1<sup>s2383</sup>* (*sgt1*) metaphase larval neuroblasts stained for the indicated Pins complex proteins and the mitotic marker mitotic marker phospho-histone H3 (not shown in A; blue in B). Scale bar, 5  $\mu$ m.

(A) Wild type and *sgt1* mutant neuroblasts show apical localization of Insc and Pins proteins, and apical enrichment of Dlg protein. Quantification of apical asymmetry: WT (Pins, 100%, n=50; Insc, 100%, n=40; Dlg, 94%, n=34), *sgt1* (Pins, 83%, n=35; Insc, 82%, n=23; Dlg, 83%, n=29)

**(B)** Wild type and *sgt1* mutants treated with 10  $\mu$ M Colcemid to depolymerize microtubules. Wild type neuroblasts show normal apical localization of the Pins complex (Pins, 100%, n=20; Insc, 100%, n=26; Dlg, 100%, n=20), whereas *sgt1* mutants lack detectable cortical localization of Pins complex proteins (Pins, 6%, n=31; Insc, 14%, n=22; Dlg, 7%, n=15).



**Figure 5. *sgt1 pins* double mutants larval brains have ectopic, apolar neuroblasts**

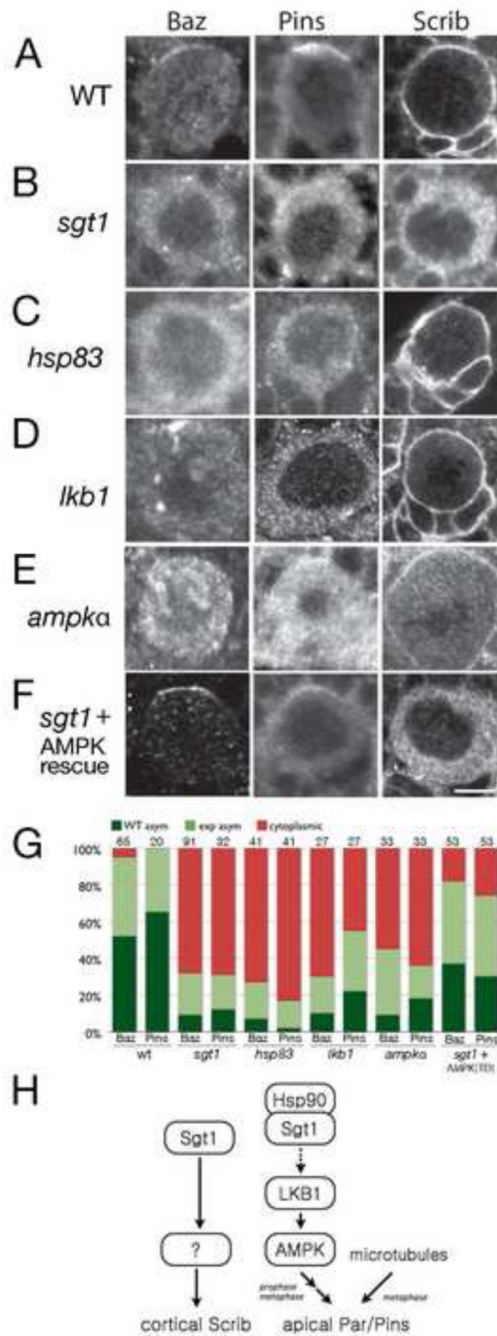
Wild type, *sgt1*<sup>S2383</sup> / *sgt1*<sup>S2383</sup> (*sgt1*) single mutants, *pins*<sup>b62</sup> / *pins*<sup>b62</sup> (*pins*) single mutants, or *sgt1 pins* double mutant larval brains stained for the indicated cortical polarity proteins and the mitotic marker phospho-histone H3 (not shown). Scale bar, 5  $\mu$ m (A-J) and 25  $\mu$ m (K-N).

(A) Wild type metaphase neuroblasts showing normal cortical polarity; apical up.

Quantification of normal asymmetric localization for each protein: aPKC, 100%, n=30; Insc, 100%, n=16; Dlg, 83%, n=29; Mira, 100%, n=20; Numb, 100%, n=20.

(B) *sgt1 pins* double mutant metaphase neuroblasts lack all cortical polarity. Quantification of asymmetric localization for each protein: aPKC, 6%, n=64; Insc, 5%, n=39; Dlg, 0%, n=45; Mira, 10%, n=79; Numb, 7%, n=41.

(C-F) Third instar larval brains stained for the neuroblast marker Miranda (Mira); the number of central brain neuroblasts in each genotype is given below each panel (average  $\pm$  standard deviation). (C) Wild type showing normal neuroblast numbers. (D,E) *sgt1* and *pins* single mutant brains showing reduced neuroblast numbers. (F) *sgt1 pins* double mutant brain showing a large increase in neuroblast numbers. (Ahmed et al., 2003)



**Figure 6. *sgt1*, *hsp83*, *LKB1*, and *AMPK* have similar neuroblast polarity phenotypes**  
 Prophase larval brain neuroblasts stained for the Par complex protein Bazooka (Baz), the Pins complex protein Pins, and the cortical protein Scribble (Scrib). Quantification in (G). Scale bar, 5  $\mu$ m.

- (A) Wild type. Baz and Pins form apical cortical crescents; Scrib is uniform cortical.  
 (B) *sgt1*<sup>s2383</sup> / *Df(3R)6147* (*sgt1*). Baz and Pins are delocalized; Scrib is cytoplasmic.  
 (C) *hsp83*<sup>13F3</sup> / *hsp83*<sup>582</sup> (*hsp83*). Baz and Pins are delocalized; Scrib is uniform cortical.  
 (D) *lkb1*<sup>4A4-2</sup> / *lkb1*<sup>4A4-2</sup> (*lkb1*). Baz and Pins are delocalized; Scrib is uniform cortical.

- (E)** *ampk<sup>1</sup> / ampk<sup>1</sup> (ampk $\alpha$ )*. Baz and Pins are delocalized; Scrib is uniform cortical.
- (F)** *sgt1<sup>s2383</sup> / Df(3R)6147; tub-gal4 UAS-ampk<sup>TD</sup>(sgt1 + AMPK rescue)*. Baz and Pins form apical cortical crescents; Scrib is cytoplasmic.
- (G)** Quantification of the Baz and Pins phenotypes for the genotypes listed in A-F. Dark green, normal asymmetric; light green, expanded asymmetric; red, cytoplasmic. Number of neuroblasts scored shown in each bar.
- (H)** Model for Sgt1 regulation of neuroblast cortical polarity. Sgt1/Hsp90 activate LKB1 which activates AMPK to promote apical Par/Pins complex localization by an unknown mechanism; spindle microtubules provide a redundant pathway generating Par/Pins apical localization at metaphase. Sgt1 acts by an Hsp90/LKB1/AMPK-independent mechanism to promote Scribble (Scrib) uniform cortical localization.