



Asymmetric cell division: recent developments and their implications for tumour biology

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Abstract | The ability of cells to divide asymmetrically is essential for generating diverse cell types during development. The past 10 years have seen tremendous progress in our understanding of this important biological process. We have learned that localized phosphorylation events are responsible for the asymmetric segregation of cell fate determinants in mitosis and that centrosomes and microtubules play important parts in this process. The relevance of asymmetric cell division for stem cell biology has added a new dimension to the field, and exciting connections between asymmetric cell division and tumorigenesis have begun to emerge.

Centrosome

(Also called the microtubule-organizing centre or spindle pole). A structure that nucleates microtubules and is important for signalling processes.

The development of multicellular organisms involves the specification of diverse cell types from a single fertilized egg. To generate this diversity, some cells can undergo an asymmetric cell division, during which they differentially segregate protein or RNA determinants into the two daughter cells, thereby determining distinct cell fates.

The process of asymmetric cell division was originally described almost 100 years ago by Conklin¹, who found that during division of early ascidian embryos an area of yellow cytoplasm always co-segregates with cells that will become muscle cells. It was not until 1994, however, that an asymmetrically segregating cell fate determinant from *Drosophila melanogaster*, called Numb, was functionally and molecularly characterized². During mitosis, Numb was found to localize to one edge of the cell, forming a crescent-shaped pattern, and to segregate into only one of the two daughter cells^{2,3} ([Supplementary information S1](#) (movie)); in the absence of Numb, normally different cells assume the same fate in *D. melanogaster* external sensory organs⁴. These observations suggested that high levels of Numb in one of the two daughter cells cause the division to become asymmetric.

In *Caenorhabditis elegans*, a similar asymmetric localization was found for partitioning defective (Par) proteins, which are also involved in other processes that require polarization⁵⁻⁷. During the first division of the *C. elegans* zygote, PAR-3 (REF. 8), PAR-6 (REF. 9) and protein kinase C-like 3 (PKC-3)¹⁰ accumulate at the anterior cell cortex, and PAR-1 (REF. 11) and PAR-2 (REF. 12) accumulate posteriorly. Thus, these proteins differentially segregate into one of the two daughter cells. In contrast to *D. melanogaster*

Numb, however, Par proteins are also required for other aspects of asymmetric cell division, including the establishment of different daughter cell sizes and the orientation and position of the mitotic spindle in *C. elegans*^{6,7}. In fact, it is the *D. melanogaster* homologues of the anterior Par proteins that direct the asymmetric localization of Numb into one of the two daughter cells¹³⁻¹⁷.

A simple model of asymmetric cell division postulates that it is a three-step process: in interphase, Par proteins set up a polarity axis¹⁸; in mitosis, this axis is used for spindle orientation and for the asymmetric localization of cell fate determinants; and in telophase, the tight coordination of spindle orientation and asymmetric localization ensures that cell fate determinants are inherited by only one of the two daughter cells.

Since this model was first proposed almost 10 years ago¹⁸, new findings have emerged. In this Review I highlight the recent discoveries that have changed our view of how determinants are asymmetrically localized. I also summarize recent findings revealing a surprising role for centrosomes in maintaining the polarity axis over many divisions. Finally, I describe how the connections between asymmetric cell division and tumorigenesis have opened unexpected and challenging avenues for this dynamic and rapidly moving field.

Asymmetric cell division: the basics

The mechanisms of asymmetric cell division have been derived from studies of invertebrates — specifically, *D. melanogaster* and *C. elegans*. Below, I describe the basic principles of this process in these organisms.

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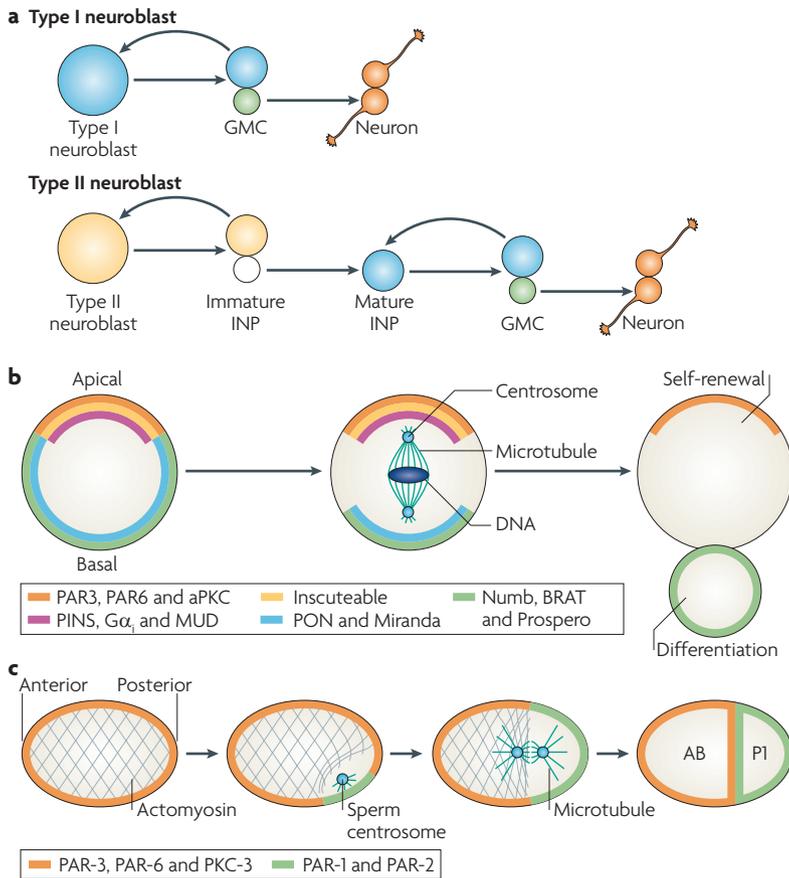


Figure 1 | Models for asymmetric cell division. a | *Drosophila melanogaster* type I neuroblasts divide asymmetrically into one neuroblast and one ganglion mother cell (GMC). The neuroblast self-renews, and the GMC divides terminally into two neurons. Type II neuroblasts divide into one self-renewing type II neuroblast and one immature intermediate neural precursor (INP). The INP starts expressing the neuroblast markers *Asense* and *Deadpan* to become a mature INP, which divides asymmetrically into one GMC and one mature INP. Differential expression of the markers *Deadpan*, *Asense*, *Prospero* and *Embryonic lethal abnormal vision (ELAV)* allows the unique identification of individual cell types in type I, type II and optic lobe (not shown) neuroblast lineages (see [Supplementary information S2](#) (figure)). **b** | In *D. melanogaster* neuroblasts, the apically localized Partitioning defective 3 (PAR3)–PAR6–atypical protein kinase C (aPKC) complex is connected to partner of Inscuteable (PINS; also known as RAPS)–G protein α_1 -subunit ($G\alpha_1$)–MUD by the adaptor protein Inscuteable. During mitosis, this apical complex directs the orientation of the mitotic spindle and the asymmetric localization of the adaptor proteins Partner of Numb (PON) and Miranda and, consequently, of the cell fate determinants Numb, Brain tumour (BRAT) and Prospero to the basal cell cortex. After mitosis, Numb, BRAT and Prospero act together to prevent self-renewal and induce cell cycle exit and differentiation. **c** | In the *Caenorhabditis elegans* zygote, the anterior Par proteins PAR-3, PAR-6 and PKC-3 segregate into the anterior AB cell, and the posterior Par proteins PAR-1 and PAR-2 segregate into the posterior P1 cell. Polarization starts after fertilization, when interactions between the sperm centrosome and cortex allow PAR-2 to accumulate at the posterior cortex. This initiates an anterior contraction of the cortical actin cytoskeleton, which allows anterior movement of PAR-3, PAR-6 and PKC-3.

Neuroblast
A. D. melanogaster neural progenitor cell that generates all of the neurons and glial cells in the brain.

Asymmetric cell division in *D. melanogaster*. During the past 10 years, most of the progress in understanding asymmetric cell division in *D. melanogaster* has been made in neuroblasts, which are cells that delaminate from the ventral neuroectoderm during embryogenesis. In embryos, neuroblasts undergo up to 20 rounds of asymmetric cell division to generate the neurons of the larval

nervous system, and they become quiescent at the end of embryogenesis. During the larval stages of development, neuroblasts re-enter the cell cycle and continue to divide asymmetrically to generate the neurons of the adult fly brain¹⁹. Several types of larval neuroblasts can be distinguished on the basis of lineage and location (FIG. 1a), and unique markers exist to allow their identification ([Supplementary information S2](#) (figure)). Most prevalent are the type I neuroblasts, which divide into a large cell that remains a neuroblast and a smaller ganglion mother cell (GMC); the GMC subsequently divides into two terminally differentiated neurons. Type II neuroblasts are located in the dorsoposterior region of each central brain hemisphere and divide to give rise to a different cell lineage to type I neuroblasts^{20–22}. The smaller daughter cell of type II neuroblasts becomes an intermediate neural precursor (INP), which continues to undergo self-renewing asymmetric divisions, each division generating one INP and one GMC. Furthermore, specialized kinds of type I neuroblasts exist in the mushroom bodies^{19,23} and the optic lobes²⁴.

The basic mechanism of asymmetric cell division is common to all *D. melanogaster* neuroblasts^{25–28} (FIG. 1b). The endocytic protein Numb²⁹ (which inhibits Notch–Delta signalling) and the translation inhibitor Brain tumour (BRAT)³⁰ transiently accumulate at the basal plasma membrane in late prometaphase^{3,31–33}. Their asymmetric localization is facilitated by two adaptor proteins that localize asymmetrically at the same time as Numb and BRAT. BRAT localizes by binding Miranda^{31,33}, and Numb localization is facilitated by (but does not depend on) the adaptor protein Partner of Numb (PON)^{34,35}. In type I neuroblasts and INPs, Miranda also transports the transcription factor Prospero into the GMC^{36–40}. Slightly after the basal determinants localize, the mitotic spindle is set up in an apical–basal orientation so that these determinants are inherited by the basal daughter cell.

The asymmetric localization of basal determinants also requires another set of proteins that accumulate at the apical cell cortex before mitosis. These include the PDZ domain-containing proteins PAR3 and PAR6 and the protein kinase atypical PKC (aPKC^{13–17}; the *D. melanogaster* homologue of *C. elegans* PKC-3). The group of proteins also includes the adaptor protein Inscuteable^{41,42}, which links PAR3–PAR6–aPKC to a second protein complex containing the heterotrimeric G protein α_1 -subunit ($G\alpha_1$)⁴³ and the adaptor protein Partner of Inscuteable (PINS; also known as RAPS)^{43–45}. PINS binds to the microtubule-associated dynein-binding protein MUD^{46–48} and thereby provides a cortical attachment site for astral microtubules to ensure the apical–basal orientation of the mitotic spindle.

The initial apical localization of PAR3, PAR6 and aPKC is inherited from epithelial cells of the ventral neuroectoderm when the neuroblasts delaminate^{13,14,16,17}. In these epithelial cells, Par proteins localize apically and are required for establishing and maintaining apicobasal polarity. In fact, PAR3, PAR6 and aPKC — and their homologues in other organisms — play a key part in almost all known cell polarity events, including

epithelial polarity, axon outgrowth, synapse formation and specification of the anteroposterior body axis^{6,7}. How Par proteins direct the asymmetric localization of cell fate determinants during asymmetric cell division and how the apical localization of Par proteins is maintained during subsequent neuroblast cell cycles have become clear only recently and are discussed below.

Asymmetric cell division in *C. elegans*. In *C. elegans*, the first cell division during development generates an anterior AB cell and a posterior P1 cell (for excellent reviews, see REFS 49,50). The size and fate of these two daughter cells are different, and the mechanisms that generate this asymmetry are similar to those that act in neuroblasts (FIG. 1c). Polarization of the zygote starts when the entire cortical actin cytoskeleton moves towards the anterior pole⁵¹. This movement is initiated by the sperm centrosome^{52,53} and by the Rho guanine nucleotide exchange factor (RhoGEF) cytokinesis defect 4 (CYK-4)⁵⁴, which is contributed during fertilization and remains localized close to the posterior male pronucleus. As a result of anterior cortical movement, surface contractions that initially occur throughout the cell are progressively confined to the anterior half of the zygote, whereas the posterior side becomes smooth⁵⁰. PAR-3, PAR-6 and PKC-3 are initially uniformly cortical but concentrate at the anterior side after fertilization⁵¹, although a second, actomyosin-independent mechanism has been described⁵⁵. PAR-1 and PAR-2 become enriched in the posterior, non-contracting cell cortex, and inhibitory interactions between the anterior and posterior Par proteins ensure that the groups maintain their localization to opposite cortical domains. PAR-2, for example, prevents the cortical localization of PKC-3 (REF. 56) and PKC-3 phosphorylates PAR-2; this removes it from the plasma membrane. Thus, in contrast to those in *D. melanogaster*, Par proteins in *C. elegans* are involved in regulating both asymmetric cell division and the symmetry-breaking events that establish the anteroposterior axis in the zygote.

The distinction between segregating determinants and proteins establishing polarity is not as clear in *C. elegans* as in *D. melanogaster*⁵⁷. In addition to the effects of the Par proteins, the asymmetric division of the zygote is influenced by the CCCH-Zn finger proteins muscle excess 1 (MEX-1), MEX-5, MEX-6, posterior segregation protein 1 (POS-1) and pharynx and intestine in excess protein 1 (PIE-1), the RNA-binding proteins MEX-3 and spindle orientation defective protein 4 (SPN-4; also known as PIP-1) and the homeodomain protein posterior alae in males protein 1 (PAL-1)⁵⁷. PIE-1 is inherited by the posterior P1 cell⁵⁸, where it blocks transcriptional elongation⁵⁹ and prevents the expression of genes that would promote somatic differentiation in the germline blastomeres⁶⁰. MEX-5 and MEX-3 segregate into the anterior AB daughter cell and inhibit the specification of muscle cell fate in its progenitors^{61,62}. Par proteins are essential for asymmetric segregation of PIE-1, MEX-5 and MEX-3. However, the accumulation of Par proteins themselves, as well as actomyosin flow, is regulated redundantly by MEX-5 and

the highly related MEX-6 (REF. 63). In fact, most of these proteins are also involved in the asymmetric segregation of other factors, with the notable exception of PIE-1 and PAL-1, and they are therefore considered to be polarity mediators rather than segregating determinants⁵⁷.

Asymmetric localization of determinants

The mechanisms that lead to the asymmetric localization of Numb, BRAT and Prospero in *D. melanogaster* neuroblasts had remained a mystery for many years. Similarly, it was unclear how cytoplasmic determinants are segregated into the AB or P1 cell in *C. elegans*. Initial experiments using chemical inhibitors in *D. melanogaster* showed that the process does not require microtubules but depends on actin and myosin^{36,64-66}. This led to the formulation of a model in which an actomyosin-dependent process moves asymmetrically, segregating cell fate determinants along the cell cortex to concentrate them on the basal side^{18,67}. Support for this model came from the demonstration that myosin VI is important for asymmetric cell division⁶⁸ and from the finding that the cytoskeletal protein Lethal (2) giant larvae (L(2)GL) is important for the basal localization of Numb but not for the apical localization of Par proteins^{69,70}. L(2)GL binds and inhibits cytoplasmic non-muscle myosin^{71,72}, and this interaction is inhibited through phosphorylation by the apical protein aPKC^{73,74}. As a result, L(2)GL is inhibited in the apical half but active in the basal half, where it could potentially inhibit myosin. Consistent with this hypothesis, myosin II is concentrated apically in neuroblasts⁷⁵ and, when it is inhibited by mutation or chemical inhibitors of Rho-associated protein kinase (ROCK), Numb and its interacting protein Miranda (see below) no longer concentrate on the basal side⁷⁵.

Although the cortical transport model is attractive, it has been challenged by several recent observations. The asymmetry in myosin localization is not observed in external sensory organs⁷⁶ and could not be confirmed in more recent reports, which actually describe myosin localization to the basal side of the neuroblast⁷⁷. Furthermore, the ROCK inhibitor that was used to demonstrate the requirement of myosin for Numb and Miranda localization can also inhibit aPKC⁷⁸. Finally, fluorescence recovery after photobleaching (FRAP) experiments did not reveal unidirectional cortical transport of the Numb adaptor PON⁷⁹. Instead, FRAP recovery rates showed that PON and Numb rapidly exchange between cortex and cytoplasm and that local differences in cortical 'on' and 'off' rates, rather than in cortical transport, are responsible for the asymmetric localization of these proteins⁸⁰. Therefore, the cortical transport model has been replaced by more dynamic models, in which the differential mobility or cortical attachment of protein determinants to the apical and basal plasma membranes regulates their asymmetric localization. Below, I discuss how those models explain asymmetric segregation of determinants in *D. melanogaster* and describe similar models that explain the asymmetric localization of cytoplasmic proteins in *C. elegans*.

Mushroom body

A mushroom-shaped paired-neuropil structure that is found in the *D. melanogaster* brain and functions in learning and memory.

Optic lobe

A morphologically distinct part of the developing *D. melanogaster* brain that forms the visual processing centres.

Blastomere

A cell that is generated during embryonic cleavage divisions.

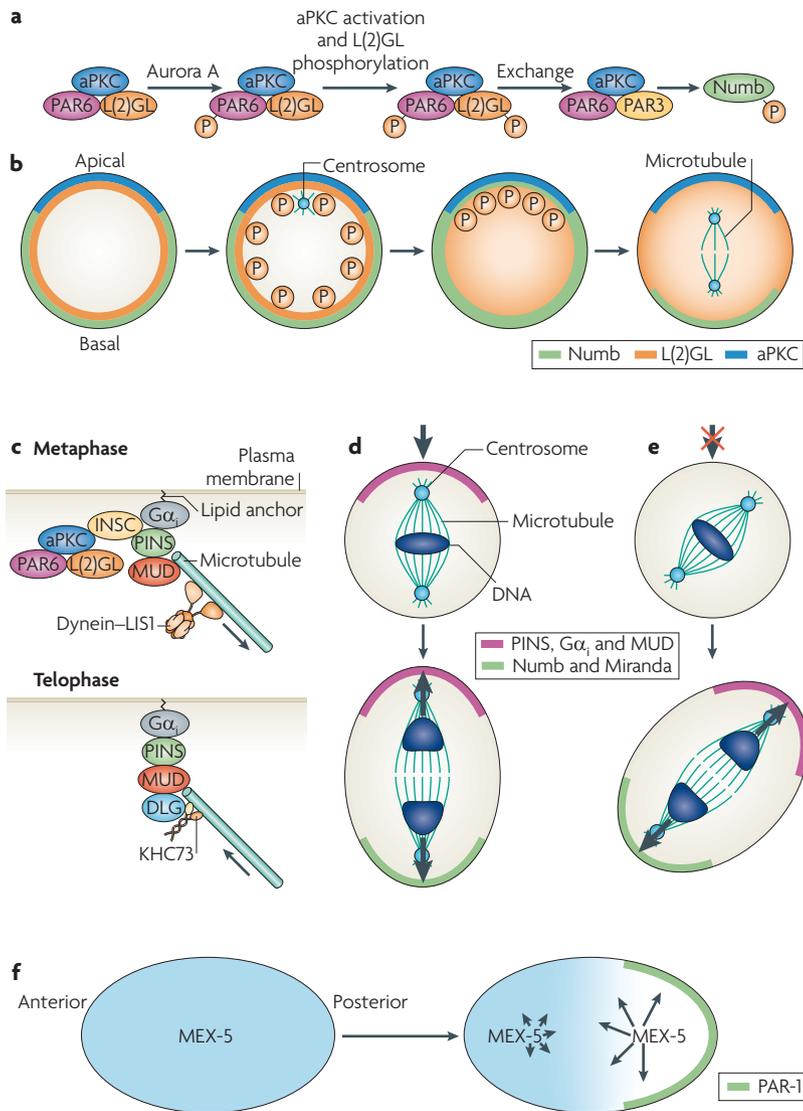


Figure 2 | Asymmetric segregation of protein determinants. a In *Drosophila melanogaster* neuroblasts, activation of Aurora A results in the phosphorylation of Partitioning defective 6 (PAR6), which in turn activates atypical protein kinase C (aPKC), leading to Lethal (2) giant larvae (L(2)GL) phosphorylation and exit from the complex. L(2)GL is exchanged for PAR3, which acts as an adaptor that allows aPKC to phosphorylate Numb. Phosphorylated Numb is then released into the cytoplasm. As aPKC is restricted to the apical cortex, Numb is retained on the basal side and segregates into the basal daughter cell. **b** Localization of Numb, L(2)GL and aPKC in the cell during mitosis. **c** In metaphase, G protein α_1 -subunit ($G\alpha_1$), Partner of Inscuteable (PINS; also known as RAPS) and MUD establish a cortical attachment site for astral microtubules to orient the mitotic spindle. In telophase, however, it is the mitotic spindle that influences cortical polarity of neuroblasts through a microtubule-dependent pathway. In this case, kinesin KHC73, which is transported on astral microtubules, binds Discs large (DLG). This, in turn, recruits $G\alpha_1$ and PINS, which then recruits MUD. This results in the accumulation of determinants over one spindle pole. **d, e** | Normally, the telophase pathway (**d**) is not essential. When components of the apical complex are missing, however, it rescues the formation of opposing cortical domains in anaphase and telophase. The new polarity axis aligns with the mitotic spindle and not necessarily with the apicobasal axis (**e**). **f** | In *Caenorhabditis elegans*, muscle excess 5 (MEX-5) and pharynx and intestine in excess protein 1 (PIE-1; not shown) exist as fast- and slow-diffusing forms. The fast-diffusing form of MEX-5 is more abundant posteriorly, and the fast-diffusing form of PIE-1 is concentrated anteriorly, resulting in the asymmetric distribution of these cytoplasmic proteins. For MEX-5, phosphorylation by posteriorly localized PAR-1 may be responsible for the faster diffusion rate. INSC, Inscuteable; LIS1, Lissencephaly 1.

The prophase pathway: asymmetric phosphorylation. In *D. melanogaster* neuroblasts, Numb is recruited to the plasma membrane through the phospholipid interactions of positively charged amino acids in its amino terminus⁶⁵. Next to those residues are several phosphorylation sites for aPKC, mutation of which to Ala abolishes the asymmetric localization of Numb in mitosis⁸¹. These observations suggest that aPKC-mediated phosphorylation neutralizes positive charges and thereby inhibits the membrane association of Numb⁷⁶.

In interphase, aPKC forms a complex with PAR6 and L(2)GL (FIG. 2a,b); this complex cannot phosphorylate Numb, presumably because the substrate-binding site is blocked. On entry into mitosis, the kinase Aurora A phosphorylates PAR6 (REF. 76), leading to the activation of aPKC and consequent phosphorylation of L(2)GL. This reduces the affinity of L(2)GL for PAR6 and aPKC, thereby releasing it from the complex and allowing PAR3 to enter⁸². PAR3 can bind to both Numb and aPKC and might act as an adaptor between kinase and substrate. This subunit exchange initiates the phosphorylation of Numb because aPKC phosphorylates Numb only when it is bound to PAR3 and not when bound to L(2)GL. Therefore, the function of L(2)GL is not to recruit determinants to the cortex, as previously thought, but to regulate the substrate specificity and maybe also the activity of aPKC. In *l(2)gl* mutants, for example, it is premature aPKC phosphorylation, rather than myosin defects, that prevents Numb localization; moreover, the effects of overexpression of an L(2)GL that cannot be phosphorylated are due to inhibition of aPKC⁷⁸ rather than active recruitment of asymmetric determinants to the cortex. These new findings have converted Numb localization from a complete mystery to one of the best-understood mitotic events.

In fact, aPKC-dependent phosphorylation is a general mechanism for asymmetric protein localization during mitosis, at least in *D. melanogaster*. aPKC can also phosphorylate Miranda⁷⁶ and regulate its cortical localization, similarly to how it controls Numb localization⁷⁸. Furthermore, the E3 ubiquitin ligase Neuralized, which segregates asymmetrically in sensory-organ precursor cells, contains aPKC consensus sites in its N-terminal phosphoinositide-binding domain⁸³, suggesting that it might also be regulated by aPKC. This new model of phosphorylation-dependent asymmetric cell division does not implicate actomyosin as a major player in asymmetric protein localization. Consistent with this, the weak actin inhibitor cytochalasin D does not inhibit the process, although it can prevent cytokinesis³⁶. This model might also explain why asymmetric segregation of aPKC alone is sufficient to generate different fates, even when Numb and Miranda are inherited by both daughter cells in mutants with altered spindle orientation⁸⁴. As both proteins need to be membrane bound to carry out their functions (Numb acts on endocytic vesicles and Miranda recruits other proteins to the cortex), they can be inhibited by aPKC phosphorylation in one of the two daughter cells. Therefore, it is the ratio between aPKC and basal determinants that ultimately determines the fate of each daughter cell⁸⁴.

The telophase pathway: microtubule–cortex interactions. Numb and Miranda still segregate asymmetrically in mutants in which asymmetric localization in prophase and metaphase is completely abolished^{15,69,70}. This is due to a second pathway for asymmetric localization of determinants that acts in anaphase and telophase of the cell cycle (reviewed in REF. 85). In contrast to the prophase pathway, the telophase pathway is sensitive to microtubule-depolymerizing drugs or to mutations affecting astral microtubules^{85,86} (FIG. 2c). In wild-type *D. melanogaster*, the pathway is not required for asymmetric protein localization in metaphase, as disruption of microtubules has no effect³⁶. However, in *inscuteable* mutants — in which PAR3–PAR6–aPKC is delocalized in interphase^{43,44} and mitotic spindles are no longer oriented along the apicobasal axis — the microtubule-dependent pathway is responsible for PINS and Gα_i accumulation over one of the two spindle poles in mitosis and for asymmetric segregation of determinants (FIG. 2d,e) so that cell fate specification occurs normally in a large subset of neuroblasts⁸⁶. The microtubule-dependent pathway for neuroblast polarization depends on the PINS-binding partner Discs large (DLG), which is a membrane-associated guanylate kinase that also plays a part in Numb and Miranda localization in wild-type embryos. The pathway also requires the kinesin KHC73, which localizes to microtubule plus ends and can bind DLG. These observations have suggested a model in which KHC73, transported on astral microtubules, is responsible for the accumulation of DLG and PINS over one spindle pole. PINS then recruits the microtubule-binding protein MUD, and this mutual microtubule–cortex interaction stabilizes spindle orientation (FIG. 2c).

Although this model is attractive and consistent with all of the available data, several key questions remain. For example, it is unknown how DLG and PINS establish the localization of basal determinants when the Par proteins are not asymmetrically localized. Furthermore, the phenotype of *dlg* mutants is not entirely consistent with the model: basal determinants do not localize correctly in metaphase in these mutants, but the rescue of asymmetric cell division in telophase still occurs^{69,70}. Therefore, DLG is required for the telophase pathway when other regulators are missing, but its function can be replaced when the rest of the machinery is intact. The precise molecular function of the telophase pathway still needs to be defined.

Asymmetric protein segregation in *C. elegans*. The mechanisms regulating asymmetric cell division in *C. elegans* are remarkably similar to those in *D. melanogaster*, even though the segregating determinants PIE-1 and MEX-5 localize asymmetrically in the cytoplasm and not at the cortex. Their asymmetric localization is mediated by regulated protein degradation and a reaction–diffusion mechanism in which asymmetry is established through different ratios of slowly and rapidly diffusing isoforms in the anterior and posterior halves.

Protein degradation contributes to PIE-1 asymmetry in late-stage embryos but not in the zygote^{87–89}. PIE-1 degradation during these late cycles is mediated by the

suppressor of cytokine signalling (SOCS) box protein Zn finger-interacting factor 1 (ZIF-1)⁸⁷, which interacts with the CCCH-Zn fingers of PIE-1 and also binds to a ubiquitin ligase complex containing elongin C, cullin 2 (CUL-2) and E2 ubiquitin-conjugating enzyme 5 (UBC-5). Together, these proteins degrade PIE-1 in somatic cells and thereby restrict its expression to the germ line. Interestingly, MEX-5 activates ZIF-1 and is also required for restricting PIE-1 to the germ line. This degradation mechanism explains the antagonistic expression of MEX-5 and PIE-1 in later embryos.

In the zygote, the asymmetric localization of PIE-1 and MEX-5 is thought to be established through a reaction–diffusion mechanism^{88,90} (FIG. 2f). This mechanism is used to describe chemical reactions and involves two substances that can be converted into each other by a chemical reaction and that move in space with different kinetics. The mechanism was initially applied to biology by Turing⁹¹, and it is now well established that reaction–diffusion mechanisms are responsible for pattern generation in many biological systems⁹². In this case, the two substances can be differentially modified forms of a protein or a free and a complex-associated form. FRAP and fluorescence correlation spectroscopy (FCS) experiments have shown that PIE-1 and MEX-5 exist as rapidly and slowly diffusing isoforms^{88,90}. The ratio between these isoforms is different in the anterior and posterior parts of the zygote, with more slowly diffusing PIE-1 localized posteriorly and more slowly diffusing MEX-5 localized anteriorly (FIG. 2f). In both cases, mathematical modelling of the protein distributions that would result from the measured diffusion coefficients predicts the observed asymmetric protein distributions.

So how are the apparent differences in cytoplasmic mobility established? MEX-5 needs to be phosphorylated by PAR-1 to localize asymmetrically⁸⁹. PAR-1 is concentrated posteriorly and can locally change the mobility of MEX-5 by modifying its association with the actin cytoskeleton. This explains the actin dependence of MEX-5 asymmetry, although the asymmetric movement of the actin meshwork itself adds an additional complication. For PIE-1, differential association with posterior P granules was proposed⁸⁸. P granules segregate asymmetrically in a Par protein-dependent manner, so this would explain PIE-1 asymmetry, although biochemical evidence for this is still lacking.

Thus, differential association with membranes or other cellular components, rather than directional transport, establishes the asymmetric localization of cell fate determinants in both *D. melanogaster* and *C. elegans*.

A new role for the centrosome

Ten years ago, microtubules were thought not to have a role during asymmetric cell division in *D. melanogaster*^{18,67}. Now, it is clear that microtubules play an important part in the telophase pathway. In addition, microtubule-dependent cortical interactions are integral to maintain polarity over many divisions.

D. melanogaster neuroblasts repeatedly divide along the apicobasal axis. Real-time analysis of spindle orientation has revealed that the mitotic spindle is established

P granule

A type of ribonucleoprotein particle that segregates with and marks all cells of the *C. elegans* germ line.

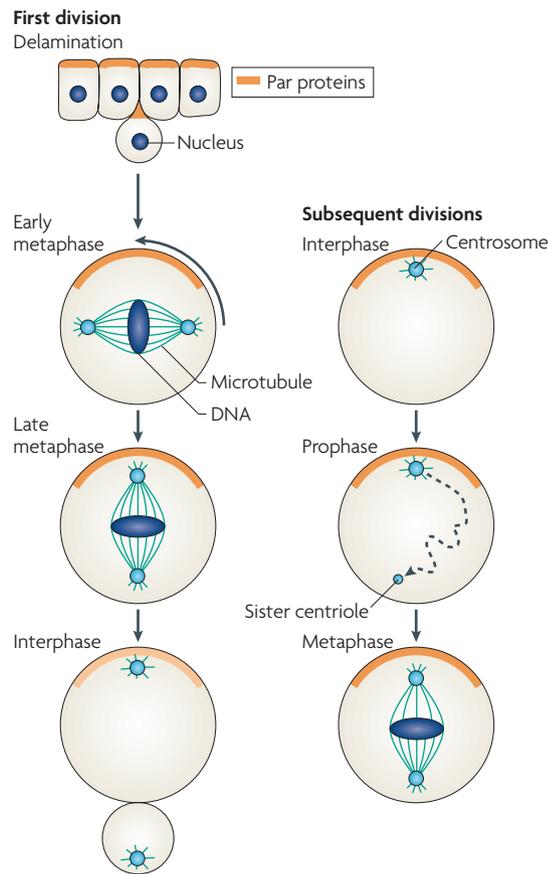


Figure 3 | Par proteins and centrosomes establish cortical polarity in *Drosophila melanogaster* neuroblasts. During the first neuroblast cycle, Partitioning defective (Par) proteins are inherited from the apical cortex of the overlying epithelium. Through a series of adaptor proteins, they recruit MUD, which forms cortical attachment sites for astral microtubules and thereby induces spindle rotation into an apicobasal orientation. During subsequent cell cycles, centrosomes are responsible for maintaining cortical polarity in interphase. Cortical Partner of Inscuteable (PINS; also known as RAPS), in turn, is required for maintaining the apical centrosome while the sister centriole migrates through the cytoplasm. On the basal side, this centriole recruits pericentriolar material to form a bipolar spindle in the proper apicobasal orientation.

parallel to the embryonic surface but then rotates by 90° into its final vertical position⁹³ (FIG. 3). It was thought that both centrosomes organize microtubule asters simultaneously at the onset of mitosis and set up a bipolar mitotic spindle in prophase. More recently, it became clear that this mechanism applies only to the first division of embryonic neuroblasts. During subsequent divisions, the apical position of the neuroblast centrosome that results from the previous cytokinesis is maintained throughout interphase^{94,95}. After centriole duplication, the daughter centriole is devoid of pericentriolar material when it migrates to the basal side of the neuroblast. A second microtubule aster appears in prophase, shortly before breakdown of the nuclear envelope. As a result, the mitotic spindle is already set up in its final, vertical orientation and does not rotate

Centriole

A small organelle (consisting of two short, barrel-like arrays of microtubules) that organizes the centrosome and contributes to cytokinesis and cell-cycle progression.

substantially in metaphase. Thus, contrary to what was previously thought, the orientation of most neuroblast divisions is established early in the cell cycle.

The orientation of the spindle across several neuroblast divisions is maintained by crosstalk between the centrosome and apical proteins. In *pins* mutants, the apical aster loses its microtubule-nucleating activity and starts to migrate basally, resulting in two identical centrosomes and random spindle orientation⁹⁵. This suggests that apical proteins maintain the apical position of the centrosome in interphase. However, the positioning of apical proteins can also be instructed by the centrosome itself. When microtubules are transiently inactivated, the apical centrosome assumes a random position and induces the localized accumulation of Par proteins at its new position⁹⁶. This symmetry-breaking property of the neuroblast centrosome is strikingly similar to what occurs in the *C. elegans* zygote, in which the sperm centrosome breaks symmetry and establishes the localization of Par proteins to the anterior and posterior domains. In contrast to *D. melanogaster*, however, in the *C. elegans* zygote the centrosome removes, rather than attracts, the PAR-3–PAR-6–PKC-3 complex. One important implication of these new findings is that the sister centrosomes are not identical in neuroblasts and could therefore be involved in maintaining asymmetric cell division. In yeast, it has been shown that the newly born centriole (known as the spindle pole body in this case) is always inherited by the bud cell and never by the mother cell. During the asymmetric divisions of *D. melanogaster* testes, the mother centriole remains anchored at the stem cell niche and is always inherited by the daughter cell, which retains the self-renewal capacity. These observations have raised speculations about centrosomes having fate-determining properties⁹⁷. For example, during brain development in vertebrates, the mother centriole is preferentially inherited by the progenitor cell⁹⁸. In this case, removal of ninein, a protein that ensures this inheritance pattern, causes randomization of centriole inheritance and a defect in progenitor cell maintenance. Although this is just a correlation, this finding indicates that centrosome asymmetry might contribute to asymmetric cell division in vertebrate cells.

Specifying daughter cell sizes

Besides having different cell fates, the daughter cells of both *D. melanogaster* neuroblasts and the *C. elegans* zygote are different in size. Identification of the mechanisms through which this asymmetry is established has revealed an exciting role for heterotrimeric G proteins in mediating microtubule–cortex interactions (FIG. 4). Although the involvement of G proteins was clear 10 years ago, how they interact with microtubules and establish cell asymmetry was discovered only recently.

In *C. elegans*, size asymmetry during the first division is due to an asymmetric displacement of the mitotic spindle towards the posterior end of the cell (FIG. 4a). This is thought to be due to increased pulling forces exerted on the spindle at the posterior end that are mediated by heterotrimeric G proteins and their binding partners, the *C. elegans* PINS homologues G protein regulator 1

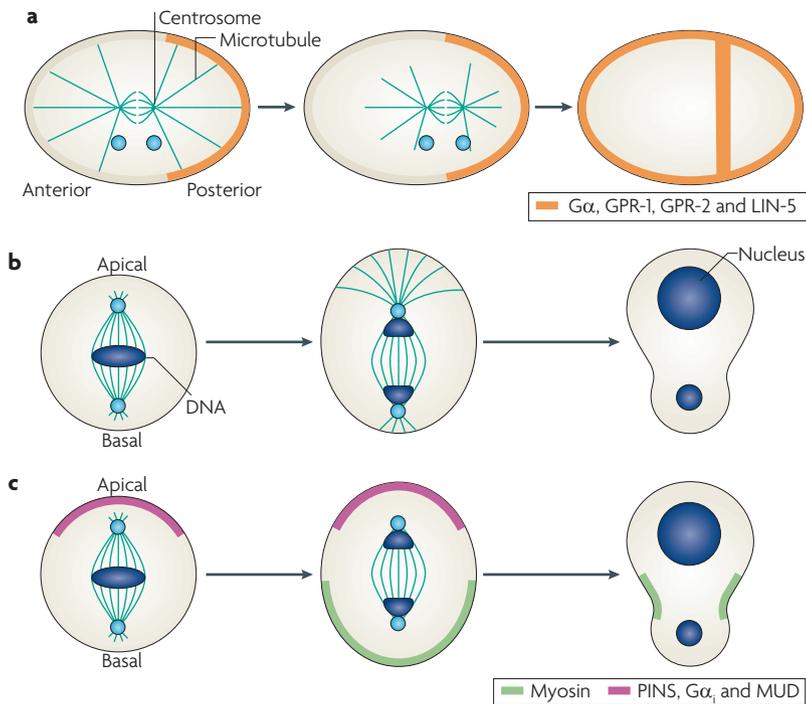


Figure 4 | Three ways to generate different daughter cell sizes. **a** | In the *Caenorhabditis elegans* zygote, heterotrimeric G protein α -subunit ($G\alpha$) and the GoLoco proteins G protein regulator 1 (GPR-1) and GPR-2 recruit the dynein-binding protein LIN-5 to the cortex to facilitate microtubule–cortex interactions. Higher concentrations of GPR-1 and GPR-2 on the posterior side result in a net pulling force and posterior spindle displacement. **b** | In *Drosophila melanogaster* neuroblasts, the apical microtubule aster is larger, resulting in spindle displacement towards the basal side and asymmetric cleavage. **c** | In neuroblasts, the apical Partner of Inscuteable (PINS; also known as RAPS)– $G\alpha$ –MUD complex induces a basal shift of cortical myosin, resulting in basal displacement of the cleavage furrow. This pathway can generate asymmetric daughter cell sizes even in the absence of a mitotic spindle.

(GPR-1) and GPR-2 (REFS 99,100 and reviewed in REF. 49). GPR-1 and GPR-2 carry a GoLoco domain that binds to the GDP-bound form of $G\alpha$. $G\alpha$ –GPR-1 and $G\alpha$ –GPR-2 recruit LIN-5 (REF. 101) (the *C. elegans* homologue of the human microtubule-binding protein nuclear mitotic apparatus protein 1 (NuMA)), the microtubule minus-end-directed motor dynein and the dynein-binding protein lissencephaly 1 (LIS-1)¹⁰². LIN-5, dynein and LIS-1 form a complex in the cytoplasm and are recruited to the plasma membrane by binding to $G\alpha$. After it is recruited to the plasma membrane, the complex can form an attachment site for the plus ends of astral microtubules, thereby exerting pulling force on the mitotic spindle. As the concentration of GPR-1 and GPR-2 is higher at the posterior cortex, the mitotic spindle is pulled towards this end.

It is likely that the mechanism identified in *C. elegans* applies to *D. melanogaster* and vertebrates, as all components of the system are conserved and their subcellular localization and biochemical interactions are similar to those seen in *C. elegans*. In *D. melanogaster* neuroblasts, however, the mechanisms regulating daughter cell sizes are different from those in the *C. elegans* zygote. First, the mitotic spindle itself is asymmetric in shape, with a large apical and a much smaller basal microtubule

aster (FIG. 4b). Second, recent experiments suggest that, in addition to the spindle-induced pathway, the site of cytokinesis is determined by a second, cortical pathway⁷⁷ (FIG. 4c). Evidence for this pathway comes from live-imaging experiments showing that the cleavage furrow proteins Anillin, Pavarotti (a *D. melanogaster* guanine nucleotide exchange factor for Rho) and myosin accumulate in the basal side of the cell before the mitotic spindle becomes asymmetric, and this is mediated by the apical PINS– $G\alpha$ –MUD complex. Surprisingly, this cortical asymmetry and the resulting asymmetric cleavage furrow can even be established when spindle formation is blocked by microtubule-depolymerizing drugs, and the resulting checkpoint arrest is overcome by a mutation in a kinetochore protein. In mutants with abnormal spindle orientation but normal cortical polarity, the cortical polarity pathway and the classical spindle-induced pathway are both active, resulting in the formation of anucleate lobes of cytoplasm that are cleaved from the mother cell. These surprising recent findings that challenge the dogma for how cytokinesis is established will certainly spark new insights into this important process. As myosin asymmetry has recently also been described in *C. elegans*, the new mechanism seems to be conserved and might also exist in higher organisms¹⁰³.

Asymmetric division in tumour formation

The connection between asymmetric cell division and tumorigenesis has been one of the most surprising and important findings in the field in the past 10 years. Furthermore, studies in mammals have identified a link between tumorigenesis and dysregulated asymmetric cell division of stem cells.

Tumorigenesis in *D. melanogaster*. Genetic screens carried out in the 1970s for brain tumour formation in *D. melanogaster* revealed an involvement for the genes *l(2)gl*, *dlg*, *lethal (2) giant discs (l(2)gd)*, *brat* and *lethal (3) malignant brain tumour (l(3)mbr)*^{104,105}. Neuroblasts fail to differentiate in *D. melanogaster* embryos that are mutated for any of these genes, leading to tumour-like overproliferation. After they have been transplanted into the abdomen of another fly, the tumours continue to grow, undergo metastasis and become aneuploid.

The identification of L(2)GL and DLG as key regulators of asymmetric cell division^{69,70} and of BRAT as a segregating determinant^{31,33} suggested that these tumours actually arise from defects in asymmetric cell division. Indeed, transplantable tumours also form in mitotic neuroblast clones that are mutated for *numb* or *prospero*^{20,32} or on overexpression of activated aPKC¹⁰⁶. Subsequent analysis showed that tumours can also occur in mutants for the mitotic Ser/Thr protein kinases Aurora A^{107,108} and Polo³⁵, following overactivation of Notch¹⁰⁸, or in mutants with aberrant spindle orientation^{46–48,84,109}. Tumours can even occur when neuroblasts divide with an excess of centrosomes^{110,111}. In all these cases, defects in asymmetric cell division are the root cause of tumour formation. Notably, however, mutations in apical proteins such as aPKC or PINS have the opposite phenotype, resulting in fewer neuroblasts.

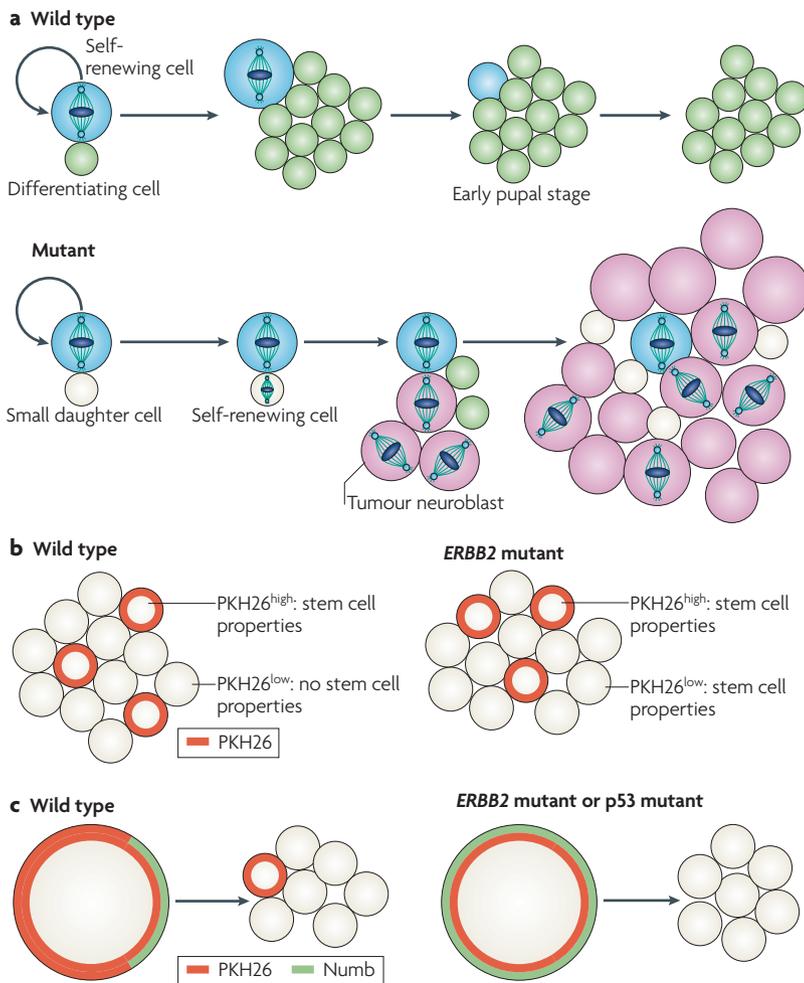


Figure 5 | Asymmetric cell division and tumour formation. **a** | Wild-type *Drosophila melanogaster* neuroblasts generate one large self-renewing daughter cell and one small differentiating daughter cell. The differentiating daughter cell exits the cell cycle after a terminal division (not shown). The neuroblast shrinks during pupal stages and undergoes apoptosis. In mutants that are defective in asymmetric cell division, the smaller daughter cell cannot differentiate. After some time, it undergoes mitosis and reverts to a tumour neuroblast. These tumour neuroblasts are abnormal because they do not exit the cell cycle during pupal stages. Whether the original neuroblast (blue) disappears or also continues to proliferate is unclear. **b** | Mammospheres that are grown from wild-type mammary gland tissue or erythroblastosis oncogene B2 (*ERBB2*)-mutant tumour tissue contain the same number of slowly proliferating (PKH26 dye-retaining) cells (PKH26^{high}). In wild-type tissue, only the cells retaining the dye can form secondary mammospheres, but in tumour tissue mammospheres can be grown from any cell. **c** | In wild-type tissue, PKH26^{high} cells localize Numb asymmetrically. When cultured, one dye-retaining cell remains, indicating that the initial division was asymmetric. When isolated from an *ERBB2* tumour model or from p53-mutant mice, PKH26^{high} cells do not divide asymmetrically and all daughter cells lose the dye, indicating that the initial division was symmetric.

The simplest explanation for tumour formation is that defects in segregating determinants result in symmetric divisions, giving rise to two neuroblasts (FIG. 5a). The resulting exponential increase in neuroblast number would explain certain aspects of tumour formation. However, it does not explain why tumour neuroblasts do not exit the cell cycle but continue to proliferate even in adult brains³² or after transplantation¹¹². In addition, a detailed analysis of *brat*-mutant clones showed that tumour formation does not simply involve a series of symmetric divisions²⁰.

After an initial delay phase in which the BRAT-inheriting cell fails to initiate correct marker expression and enters a prolonged cell cycle block, it divides and enters a second phase in which it proliferates rapidly and indefinitely. Thus, defects in asymmetric cell division cause the formation of tumour neuroblasts that lack the mechanisms responsible for cell cycle exit during pupal development.

The formation of tumour neuroblasts in mutants with aberrant asymmetric cell division can be explained by genetic or epigenetic defects or by the intrinsic properties of the growth control mechanism. In the genetic model, DNA mutations are responsible for immortalizing neuroblasts; however, although aneuploidy does occur in transplanted neuroblast tumours¹⁰⁴ and may be responsible for their metastatic behaviour¹¹², it has not been described in primary tumours. As mutations causing genome instability do not result in brain tumours¹¹¹, it is more likely that transcriptional and/or epigenetic changes alter the behaviour of the mutant neuroblasts. The transcriptional network governing self-renewal in neuroblasts needs to be reprogrammed towards a stable and irreversible differentiation state after asymmetric cell division. Defects in this process could create a new stable state, in which the self-renewal programme is active but the modules controlling exit from proliferation are missing. For example, neuroblasts serially express different transcription factors at different developmental stages¹¹³, and a reset of this developmental timer during each defective asymmetric cell division could explain immortalization. Finally, it is possible that the growth control mechanism acting in pupae can deal with only a limited number of neuroblasts — for example, because a growth inhibitor is limiting or because neuroblasts secrete an autocrine growth-promoting factor that competes with a systemic extrinsic factor.

Several redundant mechanisms have been proposed to stop neuroblast proliferation in wild-type flies. In the abdomen of the ventral nerve chord, transient expression of the homeotic gene *abdominal A* eliminates neuroblasts by inducing apoptotic cell death¹¹⁴. In the central brain, a decrease in insulin and phosphoinositide 3-kinase signalling causes a reduction in neuroblast size followed by caspase-mediated cell death during the pupal stages of development¹¹⁵. When caspase activation is prevented, neuroblast size is still reduced, and the cells are eliminated by a caspase-independent autophagic pathway that is regulated by the transcription factor Forkhead box O (FOXO). When both FOXO and caspases are inhibited, neuroblasts continue to proliferate and generate functional neurons, even in adult flies. Surprisingly, however, this does not result in a tumour, indicating that both an increase in neuroblast number and inhibition of the elimination pathways contribute to tumour formation.

Clearly, identifying the molecular events that connect asymmetric cell division to cell immortalization is one of the greatest new challenges in the field. This is particularly important because defects in asymmetric cell division are relevant for human tumorigenesis¹¹⁶ (see below)^{117,118} and may be part of the mechanisms that convert a normal mammalian stem cell into what is known as a cancer stem cell (BOX 1).

Box 1 | Tumour stem cells

The tumour stem cell hypothesis¹¹⁶ states that tumours contain a rare population of cells that have stem cell properties and are the only tumour cells that can generate all other cell types in the tumour. The hypothesis is based on xenotransplantation experiments in which transplantation of human tumours into immunocompromised mice recapitulates the human tumour histology. It had long been known that only a few cells in a tumour could initiate tumour formation in those transplantation experiments. In the 1990s, it was shown that these few cells in leukaemia express stem cell markers and that tumour formation involves a cellular hierarchy that is similar to the one in normal haematopoiesis^{132,133}. These findings formed the basis of the tumour stem cell hypothesis and, soon after, similar experiments identified tumour stem cells in brain and mammary tumours^{134,135} and in almost all other types of human cancer¹³⁶.

Whether or not the formation of cancer stem cells is an intrinsic property of tumorigenesis is intensely debated¹³⁷. Opponents of the theory argue that cancer stem cells are simply a subtype of human tumour cells that adapt more easily to the environment of the mouse host. Whether human tumours actually arise from stem cells is debated. In mouse models, intestinal cancer can be induced by mutating the adenomatous polyposis coli (APC) tumour suppressor in stem cells but not in non-stem-cell types¹³⁸. In a mouse glioblastoma model, tumour formation coincides with the appearance of abnormal stem cell populations¹³⁹. These results suggest that DNA mutations in stem cells might be the initial event in those tumours. Experiments in *Drosophila melanogaster*, in which this hypothesis can be stringently tested, might shed light on the mechanisms that cause stem cells to become malignant.

Tumorigenesis and mammalian stem cells. The ability to generate both self-renewing and differentiating daughter cells is a defining feature of any stem cell, and asymmetric cell division is one of the mechanisms used to establish this. Evidence for asymmetric cell division exists for stem cells in muscle¹¹⁹, skin¹²⁰, the gut¹²¹, mammary glands¹¹⁷, the haematopoietic system¹¹⁸ and the developing brain^{98,122}. Nevertheless, the mechanisms that guide these asymmetric cell divisions are generally not well understood (BOX 2). In fact, transferring our detailed understanding of the process from *D. melanogaster* and *C. elegans* to vertebrates has been much more challenging than expected. Almost all of the molecular players are

conserved in vertebrates, but they often act in distinct ways. Numb, for example, is polarized in vertebrate neural progenitors, but this is because it regulates trafficking of E-cadherin at adherens junctions¹²³. Par proteins are apical but, unlike in neuroblasts, only a few progenitor divisions are aligned along the axis of Par protein polarity¹²⁴. Nevertheless, building on the results from flies and worms, some exciting connections between asymmetric cell division and tumorigenesis have recently been identified.

Stem cells from mouse mammary glands can grow into spherical cultures known as mammospheres that recapitulate the mammary morphogenic programme¹²⁵. The stem cells can be isolated because they retain a lipophilic vital dye following labelling, whereas dividing cells do not¹¹⁷ (FIG. 5b). When purified from wild-type mammary glands, these stem cells divide asymmetrically and segregate Numb into one of their two daughter cells (FIG. 5c). In a mouse mammary tumour model, the number of stem cells is increased (FIG. 5b) and they divide symmetrically — Numb is no longer asymmetrically localized, and both daughter cells behave identically in terms of dye dilution (FIG. 5c). Similar observations have been made in p53-mutant mice. As p53 degradation is regulated by Numb¹²⁶, it is possible that the asymmetric inheritance of Numb regulates p53 levels and restricts stem cell fate to only one of the two daughter cells. Consistent with this, Numb is a major tumour suppressor in breast cancer¹²⁷. Numb also acts in the haematopoietic system, where it can inhibit the progression of chronic myeloid leukaemia (CML). Cultured haematopoietic progenitors normally divide and segregate Numb asymmetrically, but their divisions become symmetric following the expression of the fusion protein NUP98–HOXA9 (REF. 118) (encoded by a fusion of two genes that occurs during tumour formation and is characteristic for a specific form of

Box 2 | Asymmetric cell division in vertebrates

Almost all of the molecules regulating asymmetric cell division in *Drosophila melanogaster* and *Caenorhabditis elegans* are conserved in vertebrates. Similarly to those in invertebrate model organisms, Partitioning defective 3 (PAR3), PAR6 and atypical protein kinase C (aPKC) homologues act together to establish cell polarity in vertebrates⁷. The vertebrate Partner of Inscuteable (PINS; also known as RAPS) homologue, G protein α -subunit ($G\alpha$), and nuclear mitotic apparatus protein 1 (NuMA; the LIN-5 and Mushroom body defect (MUD) homologue) regulate spindle orientation and microtubule–cortex association¹⁴⁰, and Numb controls endocytosis¹⁴¹. Nevertheless, the function of these proteins during asymmetric cell division in vertebrates is not clear.

Asymmetric cell division in neural progenitor cells is the best understood asymmetric cell division process in mammals¹⁴². After an initial expansion phase by symmetric division, progenitors undergo asymmetric divisions, giving rise to two daughter cells: one progenitor cell and one cell that either differentiates into a neuron or becomes an intermediate progenitor, which forms two neurons after a terminal symmetric division. Progenitors are located in the apical side of the neuroepithelium, where Par proteins accumulate in the apical cortex. Unlike in *D. melanogaster*, however, the apical membrane domain in dividing progenitors is very narrow, and even slight twists of the cleavage plane lead to asymmetric inheritance¹²⁴. Numb is expressed by the progenitors and concentrates on apical adherens junctions and on the basolateral plasma membrane. This has led to a model in which the asymmetric inheritance of PAR3 during oblique divisions (divisions occurring at $\sim 45^\circ$ angle) inactivates Numb in one of the two daughter cells so that it no longer inhibits Notch, and two daughters with unequal Notch signalling levels are formed¹⁴³. As PAR3 is a key factor promoting Numb phosphorylation by aPKC in *D. melanogaster*⁷⁶, it is possible that, in vertebrates, differential phosphorylation of Numb might be responsible for the different activity in the two daughter cells. In addition to this Par protein-mediated asymmetry, the asymmetric inheritance of apical and basal processes¹⁴⁴, the polarized localization of the vertebrate Brain tumour (BRAT) homologue E3 ubiquitin–protein ligase tripartite motif-containing protein 3 (TRIM3)¹⁴⁵ and extracellular signals might have a role in establishing asymmetry.

leukaemia). NUP98–HOXA9 induces the expression of Musashi 2, which in turn inhibits Numb, potentially triggering the enormous expansion of undifferentiated progenitors in advanced-stage CML¹²⁸. Thus, the conserved connection with tumorigenesis establishes an unprecedented clinical relevance for research on asymmetric cell division.

Open questions and future challenges

The progress in our understanding of asymmetric cell division during the past 10 years has been enormous. We have learned that the phosphorylation of cell fate determinants by asymmetrically distributed kinases is the driving force for the asymmetric localization of these determinants, whereas polarized transport seems to have a minor role. It has become clear that microtubules, which were originally thought not to be involved in asymmetric cell division, mediate essential interactions between centrosomes and the cell cortex. These interactions maintain the polarity axis over multiple divisions and guide asymmetric protein localization during late mitosis. An exciting connection between asymmetric cell division and tumorigenesis has emerged in flies, mice and humans and has given rise to major challenges, in part because of our still incomplete understanding of asymmetric cell division in vertebrates. In addition, newly emerging technologies lay the groundwork for a systems-level understanding of the process.

Although we have learned the basic principles of asymmetric determinant segregation, our understanding of the cell fate choices that are influenced by those determinants is limited. We know that Numb acts on Notch, that Prospero is a transcription factor and that BRAT regulates post-transcriptional events, but how these factors cooperate to prevent self-renewal is unclear. In fact, we do not understand the transcriptional network that governs and maintains self-renewal in *D. melanogaster* neuroblasts. We also do not know how the initial bias in this network is stabilized over time and results in a daughter cell that terminally exits proliferation. And in particular, we do not know how defects in fate specification result in the formation of misguided tumour-initiating cells.

The solutions to these problems may come from the spectacular technological advances in the field. The establishment of genome-wide transgenic RNA interference libraries in flies¹²⁹ allows us to test gene functions at an unprecedented speed and on a near genome-wide level¹³⁰. In addition, the development of new sequencing technologies has opened new dimensions for genome-wide profiling of transcription, RNA splicing and chromatin association¹³¹. It is likely that these technologies will establish *D. melanogaster* neuroblasts as one of the best model systems for the establishment and stabilization of cell fate choices and will shed light on the mechanisms of stem cell-derived tumour formation. The potential clinical relevance of those findings will be a strong motivation to embark on these difficult tasks.

1. Conklin, E. G. The organization and cell-lineage of the ascidian egg. *J. Acad. Nat. Sci. Philadelphia* **13**, 1–119 (1905).
2. Rhyu, M. S., Jan, L. Y. & Jan, Y. N. Asymmetric distribution of numb protein during division of the sensory organ precursor cell confers distinct fates to daughter cells. *Cell* **76**, 477–491 (1994). **This study shows for the first time that the asymmetric segregation of a cytoplasmic determinant is important for asymmetric cell division in somatic cells.**
3. Spana, E. P., Kocczynski, C., Goodman, C. S. & Doe, C. Q. Asymmetric localization of numb autonomously determines sibling neuron identity in the *Drosophila* CNS. *Development* **121**, 3489–3494 (1995).
4. Uemura, T., Shepherd, S., Ackerman, L., Jan, L. Y. & Jan, Y. N. *numb*, a gene required in determination of cell fate during sensory organ formation in *Drosophila* embryos. *Cell* **58**, 349–360 (1989).
5. Kempthues, K. J., Priess, J. R., Morton, D. G. & Cheng, N. S. Identification of genes required for cytoplasmic localization in early *C. elegans* embryos. *Cell* **52**, 311–320 (1988). **This study describes the identification of Par proteins, which are key regulators of asymmetric cell division and cell polarity.**
6. Ohno, S. Intercellular junctions and cellular polarity: the PAR–aPKC complex, a conserved core cassette playing fundamental roles in cell polarity. *Curr. Opin. Cell Biol.* **13**, 641–68 (2001).
7. Suzuki, A. & Ohno, S. The PAR–aPKC system: lessons in polarity. *J. Cell Sci.* **119**, 979–987 (2006).
8. Etemad-Moghadam, B., Guo, S. & Kempthues, K. J. Asymmetrically distributed PAR-3 protein contributes to cell polarity and spindle alignment in early *C. elegans* embryos. *Cell* **83**, 743–752 (1995).
9. Hung, T. J. & Kempthues, K. J. PAR-6 is a conserved PDZ domain-containing protein that colocalizes with PAR-3 in *Caenorhabditis elegans* embryos. *Development* **126**, 127–135 (1999).
10. Tabuse, Y. *et al.* Atypical protein kinase C cooperates with PAR-3 to establish embryonic polarity in *Caenorhabditis elegans*. *Development* **125**, 3607–3614 (1998).
11. Guo, S. & Kempthues, K. J. *par-1*, a gene required for establishing polarity in *C. elegans* embryos, encodes a putative Ser/Thr kinase that is asymmetrically distributed. *Cell* **81**, 611–620 (1995).
12. Boyd, L., Guo, S., Levitan, D., Stinchcomb, D. T. & Kempthues, K. J. PAR-2 is asymmetrically distributed and promotes association of P granules and PAR-1 with the cortex in *C. elegans* embryos. *Development* **122**, 3075–3084 (1996).
13. Wodarz, A., Ramrath, A., Kuchinke, U. & Knust, E. Bazoooka provides an apical cue for Inscuteable localization in *Drosophila* neuroblasts. *Nature* **402**, 544–547 (1999).
14. Schober, M., Schaefer, M. & Knoblich, J. A. Bazoooka recruits Inscuteable to orient asymmetric cell divisions in *Drosophila* neuroblasts. *Nature* **402**, 548–551 (1999).
15. Petronczki, M. & Knoblich, J. A. DmPAR-6 directs epithelial polarity and asymmetric cell division of neuroblasts in *Drosophila*. *Nature Cell Biol.* **3**, 43–49 (2001).
16. Wodarz, A., Ramrath, A., Grimm, A. & Knust, E. *Drosophila* atypical protein kinase C associates with Bazoooka and controls polarity of epithelia and neuroblasts. *J. Cell Biol.* **150**, 1361–1374 (2000).
17. Rolls, M. M., Albertson, R., Shih, H. P., Lee, C. Y. & Doe, C. Q. *Drosophila* aPKC regulates cell polarity and cell proliferation in neuroblasts and epithelia. *J. Cell Biol.* **163**, 1089–1098 (2003).
18. Knoblich, J. A. Asymmetric cell division during animal development. *Nature Rev. Mol. Cell Biol.* **2**, 11–20 (2001).
19. Ito, K. & Hotta, Y. Proliferation pattern of postembryonic neuroblasts in the brain of *Drosophila melanogaster*. *Dev. Biol.* **149**, 134–148 (1992).
20. Bowman, S. K. *et al.* The tumor suppressors Brat and Numb regulate transit-amplifying neuroblast lineages in *Drosophila*. *Dev. Cell.* **14**, 535–546 (2008).
21. Boone, J. Q. & Doe, C. Q. Identification of *Drosophila* type II neuroblast lineages containing transit amplifying ganglion mother cells. *Dev. Neurobiol.* **68**, 1185–1195 (2008).
22. Bello, B. C., Izerghina, N., Caussinus, E. & Reichert, H. Amplification of neural stem cell proliferation by intermediate progenitor cells in *Drosophila* brain development. *Neural Develop.* **3**, 5 (2008). **Together with references 20 and 21, this article describes type II neuroblasts that generate a transit-amplifying population of intermediate neural progenitors and have become a valuable model system for stem cell biology.**
23. Ito, K., Awano, W., Suzuki, K., Hiroimi, Y. & Yamamoto, D. The *Drosophila* mushroom body is a quadruple structure of clonal units each of which contains a virtually identical set of neurons and glial cells. *Development* **124**, 761–771 (1997).
24. Egger, B., Boone, J. Q., Stevens, N. R., Brand, A. H. & Doe, C. Q. Regulation of spindle orientation and neural stem cell fate in the *Drosophila* optic lobe. *Neural Develop.* **2**, 1 (2007).
25. Neumüller, R. A. & Knoblich, J. A. Dividing cellular asymmetry: asymmetric cell division and its implications for stem cells and cancer. *Genes Dev.* **23**, 2675–2699 (2009).
26. Knoblich, J. A. Mechanisms of asymmetric stem cell division. *Cell* **132**, 583–597 (2008).
27. Wu, P. S., Egger, B. & Brand, A. H. Asymmetric stem cell division: lessons from *Drosophila*. *Semin. Cell Dev. Biol.* **19**, 283–293 (2008).
28. Doe, C. Q. Neural stem cells: balancing self-renewal with differentiation. *Development* **135**, 1575–1587 (2008).
29. Berdnik, D., Török, T., González-Gaitán, M. & Knoblich, J. A. The endocytic protein α -Adaptin is required for Numb-mediated asymmetric cell division in *Drosophila*. *Dev. Cell* **3**, 221–231 (2002).
30. Sonoda, J. & Wharton, R. P. *Drosophila* Brain tumor is a translational repressor. *Genes Dev.* **15**, 762–773 (2001).
31. Lee, C. Y., Wilkinson, B. D., Siegrist, S. E., Wharton, R. P. & Doe, C. Q. Brat is a Miranda cargo protein that promotes neuronal differentiation and inhibits neuroblast self-renewal. *Dev. Cell* **10**, 441–449 (2006).

32. Bello, B., Reichert, H. & Hirth, F. The brain tumor gene negatively regulates neural progenitor cell proliferation in the larval central brain of *Drosophila*. *Development* **133**, 2639–2648 (2006).
33. Betschinger, J., Mechtler, K. & Knoblich, J. A. Asymmetric segregation of the tumor suppressor *Brat* regulates self-renewal in *Drosophila* neural stem cells. *Cell* **124**, 1241–1253 (2006).
Together with references 31 and 32, this study shows that asymmetrically segregating determinants can act as tumour suppressors in *D. melanogaster*.
34. Lu, B., Rothenberg, M., Jan, L. Y. & Jan, Y. N. Partner of Numb colocalizes with Numb during mitosis and directs Numb asymmetric localization in *Drosophila* neural and muscle progenitors. *Cell* **95**, 225–235 (1998).
35. Wang, H., Ouyang, Y., Somers, W. G., Chia, W. & Lu, B. Polo inhibits progenitor self-renewal and regulates Numb asymmetry by phosphorylating Pon. *Nature* **449**, 96–100 (2007).
36. Knoblich, J. A., Jan, L. Y. & Jan, Y. N. Asymmetric segregation of Numb and Prospero during cell division. *Nature* **377**, 624–627 (1995).
37. Spana, E. P. & Doe, C. Q. The prospero transcription factor is asymmetrically localized to the cell cortex during neuroblast mitosis in *Drosophila*. *Development* **121**, 3187–3195 (1995).
38. Shen, C. P., Jan, L. Y. & Jan, Y. N. Miranda is required for the asymmetric localization of Prospero during mitosis in *Drosophila*. *Cell* **90**, 449–458 (1997).
39. Ikeshima-Kataoka, H., Skeath, J. B., Nabeshima, Y., Doe, C. Q. & Matsuzaki, F. Miranda directs Prospero to a daughter cell during *Drosophila* asymmetric divisions. *Nature* **390**, 625–629 (1997).
40. Matsuzaki, F., Ohshiro, T., Ikeshima-Kataoka, H. & Izumi, H. Miranda localizes staufer and prospero asymmetrically in mitotic neuroblasts and epithelial cells in early *Drosophila* embryogenesis. *Development* **125**, 4089–4098 (1998).
41. Kraut, R. & Campos-Ortega, J. A. *inscuteable*, a neural precursor gene of *Drosophila*, encodes a candidate for a cytoskeleton adaptor protein. *Dev. Biol.* **174**, 65–81 (1996).
42. Kraut, R., Chia, W., Jan, L. Y., Jan, Y. N. & Knoblich, J. A. Role of *inscuteable* in orienting asymmetric cell divisions in *Drosophila*. *Nature* **383**, 50–55 (1996).
43. Schaefer, M., Petronczki, M., Dorner, D., Forte, M. & Knoblich, J. A. Heterotrimeric G proteins direct two modes of asymmetric cell division in the *Drosophila* nervous system. *Cell* **107**, 183–194 (2001).
44. Schaefer, M., Shevchenko, A., Shevchenko, A. & Knoblich, J. A. A protein complex containing Inscuteable and the α -binding protein Pins orients asymmetric cell divisions in *Drosophila*. *Curr. Biol.* **10**, 353–362 (2000).
45. Yu, F., Morin, X., Cai, Y., Yang, X. & Chia, W. Analysis of *partner of inscuteable*, a novel player of *Drosophila* asymmetric divisions, reveals two distinct steps in Inscuteable apical localization. *Cell* **100**, 399–409 (2000).
46. Siller, K. H., Cabernard, C. & Doe, C. Q. The NuMA-related Mud protein binds Pins and regulates spindle orientation in *Drosophila* neuroblasts. *Nature Cell Biol.* **8**, 594–600 (2006).
47. Izumi, Y., Ohta, N., Hisata, K., Raabe, T. & Matsuzaki, F. *Drosophila* Pins-binding protein Mud regulates spindle-polarity coupling and centrosome organization. *Nature Cell Biol.* **8**, 586–593 (2006).
48. Bowman, S. K., Neumüller, R. A., Novatckova, M., Du, Q. & Knoblich, J. A. The *Drosophila* NuMA homolog Mud regulates spindle orientation in asymmetric cell division. *Dev. Cell* **10**, 731–742 (2006).
Together with references 46 and 47, this study shows that a cortical microtubule binding protein interacts with the asymmetric cell division machinery to orient the mitotic spindle in asymmetric cell division.
49. Gönczy, P. Mechanisms of asymmetric cell division: flies and worms pave the way. *Nature Rev. Mol. Cell Biol.* **9**, 355–366 (2008).
50. Cowan, C. R. & Hyman, A. A. Asymmetric cell division in *C. elegans*: cortical polarity and spindle positioning. *Annu. Rev. Cell Dev. Biol.* **20**, 427–453 (2004).
51. Munro, E., Nance, J. & Priess, J. R. Cortical flows powered by asymmetrical contraction transport PAR proteins to establish and maintain anterior-posterior polarity in the early *C. elegans* embryo. *Dev. Cell* **7**, 413–424 (2004).
This study shows that there is anterior cortical myosin flow in *C. elegans* and reveals that it is crucial for Par protein localization.
52. Cowan, C. R. & Hyman, A. A. Centrosomes direct cell polarity independently of microtubule assembly in *C. elegans* embryos. *Nature* **431**, 92–96 (2004).
53. Tsai, M. C. & Ahiring, J. Microtubules are involved in anterior-posterior axis formation in *C. elegans* embryos. *J. Cell Biol.* **179**, 397–402 (2007).
54. Jenkins, N., Saam, J. R. & Mango, S. E. CYK-4/GAP provides a localized cue to initiate anteroposterior polarity upon fertilization. *Science* **313**, 1298–1301 (2006).
55. Zonies, S., Motegi, F., Hao, Y. & Seydoux, G. Symmetry breaking and polarization of the *C. elegans* zygote by the polarity protein PAR-2. *Development* **137**, 1669–1677 (2010).
56. Hao, Y., Boyd, L. & Seydoux, G. Stabilization of cell polarity by the *C. elegans* RING protein PAR-2. *Dev. Cell* **10**, 199–208 (2006).
57. Gönczy, P. & Rose, L. S. Asymmetric cell division and axis formation in the embryo. *WormBook* 15 Oct 2005 (doi:10.1895/wormbook.1.30.1).
58. Mello, C. C. *et al.* The PIE-1 protein and germline specification in *C. elegans* embryos. *Nature* **382**, 710–712 (1996).
59. Zhang, F., Barboric, M., Blackwell, T. K. & Peterlin, B. M. A model of repression: CTD analogs and PIE-1 inhibit transcriptional elongation by P-TEFb. *Genes Dev.* **17**, 748–758 (2003).
60. Mello, C. C., Draper, B. W., Krause, M., Weintraub, H. & Priess, J. R. The *pie-1* and *mex-1* genes and maternal control of blastomere identity in early *C. elegans* embryos. *Cell* **70**, 163–176 (1992).
61. Schubert, C. M., Lin, R., de Vries, C. J., Plasterk, R. H. & Priess, J. R. MEX-5 and MEX-6 function to establish soma/germline asymmetry in early *C. elegans* embryos. *Mol. Cell* **5**, 671–682 (2000).
62. Draper, B. W., Mello, C. C., Bowerman, B., Hardin, J. & Priess, J. R. MEX-3 is a KH domain protein that regulates blastomere identity in early *C. elegans* embryos. *Cell* **87**, 205–216 (1996).
63. Cuenca, A. A., Schetter, A., Aceto, D., Kempthues, K. & Seydoux, G. Polarization of the *C. elegans* zygote proceeds via distinct establishment and maintenance phases. *Development* **130**, 1255–1265 (2003).
64. Broads, J. & Doe, C. Q. Extrinsic cues, intrinsic cues and microfilaments regulate asymmetric protein localization in *Drosophila* neuroblasts. *Curr. Biol.* **7**, 827–835 (1997).
65. Knoblich, J. A., Jan, L. Y. & Jan, Y. N. The N terminus of the *Drosophila* Numb protein directs membrane association and actin-dependent asymmetric localization. *Proc. Natl Acad. Sci. USA* **94**, 13005–13010 (1997).
66. Shen, C. P. *et al.* Miranda as a multidomain adapter linking apically localized Inscuteable and basally localized Staufer and Prospero during asymmetric cell division in *Drosophila*. *Genes Dev.* **12**, 1837–1846 (1998).
67. Jan, Y. N. & Jan, L. Y. Asymmetric cell division in the *Drosophila* nervous system. *Nature Rev. Neurosci.* **2**, 772–779 (2001).
68. Petritsch, C., Tavosanis, G., Turck, C. W., Jan, L. Y. & Jan, Y. N. The *Drosophila* myosin VI jaguar is required for basal protein targeting and correct spindle orientation in mitotic neuroblasts. *Dev. Cell* **4**, 273–281 (2003).
69. Ohshiro, T., Yagami, T., Zhang, C. & Matsuzaki, F. Role of cortical tumour-suppressor proteins in asymmetric division of *Drosophila* neuroblast. *Nature* **408**, 593–596 (2000).
70. Peng, C. Y., Manning, L., Albertson, R. & Doe, C. Q. The tumour-suppressor genes *lg1* and *dlg* regulate basal protein targeting in *Drosophila* neuroblasts. *Nature* **408**, 596–600 (2000).
71. Strand, D. *et al.* The *Drosophila* lethal(2)giant larvae tumor suppressor protein forms homo-oligomers and is associated with nonmuscle myosin II heavy chain. *J. Cell Biol.* **127**, 1361–1373 (1994).
72. Strand, D., Raska, I. & Mechler, B. M. The *Drosophila* lethal(2)giant larvae tumor suppressor protein is a component of the cytoskeleton. *J. Cell Biol.* **127**, 1345–1360 (1994).
73. Betschinger, J., Mechtler, K. & Knoblich, J. A. The Par complex directs asymmetric cell division by phosphorylating the cytoskeletal protein Lgl. *Nature* **422**, 326–330 (2003).
74. Betschinger, J., Eisenhaber, F. & Knoblich, J. A. Phosphorylation-induced autoinhibition regulates the cytoskeletal protein Lethal(2) giant larvae. *Curr. Biol.* **15**, 276–282 (2005).
75. Barros, C. S., Phelps, C. B. & Brand, A. H. *Drosophila* nonmuscle myosin II promotes the asymmetric segregation of cell fate determinants by cortical exclusion rather than active transport. *Dev. Cell* **5**, 829–840 (2003).
76. Wirtz-Peitz, F., Nishimura, T. & Knoblich, J. A. Linking cell cycle to asymmetric division: Aurora-A phosphorylates the Par complex to regulate Numb localization. *Cell* **135**, 161–173 (2008).
This study connects the phosphorylation of segregating determinants to the cell cycle machinery and provides a model for how determinants localize asymmetrically during mitosis.
77. Cabernard, C., Prehoda, K. E. & Doe, C. Q. A spindle-independent cleavage furrow positioning pathway. *Nature* **467**, 91–94 (2010).
This paper identifies an unprecedented pathway for cytokinesis that involves asymmetric cortical localization of myosin.
78. Atwood, S. X. & Prehoda, K. E. aPKC phosphorylates Miranda to polarize fate determinants during neuroblast asymmetric cell division. *Curr. Biol.* **19**, 723–729 (2009).
79. Lu, B., Ackerman, L., Jan, L. Y. & Jan, Y. N. Modes of protein movement that lead to the asymmetric localization of partner of Numb during *Drosophila* neuroblast division. *Mol. Cell* **4**, 883–891 (1999).
80. Mayer, B., Emery, G., Berdnik, D., Wirtz-Peitz, F. & Knoblich, J. A. Quantitative analysis of protein dynamics during asymmetric cell division. *Curr. Biol.* **15**, 1847–1854 (2005).
81. Smith, C. A. *et al.* aPKC-mediated phosphorylation regulates asymmetric membrane localization of the cell fate determinant Numb. *EMBO J.* **26**, 468–480 (2007).
This investigation shows that phosphorylation of the segregating determinant Numb by the asymmetrically localized kinase aPKC regulates membrane association and provides a model for how Numb localizes asymmetrically in mitosis.
82. Yamanaka, T. *et al.* Mammalian Lgl forms a protein complex with PAR-6 and aPKC independently of PAR-3 to regulate epithelial cell polarity. *Curr. Biol.* **13**, 734–743 (2003).
83. Skwarek, L. C., Garroni, M. K., Commisso, C. & Boulianne, G. L. Neuralized contains a phosphoinositide-binding motif required downstream of ubiquitination for Delta endocytosis and Notch signaling. *Dev. Cell* **13**, 783–795 (2007).
84. Cabernard, C. & Doe, C. Q. Apical/basal spindle orientation is required for neuroblast homeostasis and neuronal differentiation in *Drosophila*. *Dev. Cell* **17**, 134–141 (2009).
85. Siegrist, S. E. & Doe, C. Q. Microtubule-induced cortical cell polarity. *Genes Dev.* **21**, 483–496 (2007).
86. Siegrist, S. E. & Doe, C. Q. Microtubule-induced Pins/Gai cortical polarity in *Drosophila* neuroblasts. *Cell* **123**, 1323–1335 (2005).
This article provides a molecular explanation for the telophase rescue pathway that acts during anaphase and telophase.
87. DeRenzo, C., Reese, K. J. & Seydoux, G. Exclusion of germ plasm proteins from somatic lineages by cullin-dependent degradation. *Nature* **424**, 685–689 (2003).
This work shows that protein degradation is important for asymmetric localization of cytoplasmic determinants in *C. elegans*.
88. Daniels, B. R., Perkins, E. M., Dobrowsky, T. M., Sun, S. X. & Wirtz, D. Asymmetric enrichment of PIE-1 in the *Caenorhabditis elegans* zygote mediated by binary counterdiffusion. *J. Cell Biol.* **184**, 473–479 (2009).
89. Tenlen, J. R., Molk, J. N., London, N., Page, B. D. & Priess, J. R. MEX-5 asymmetry in one-cell *C. elegans* embryos requires PAR-4- and PAR-1-dependent phosphorylation. *Development* **135**, 3665–3675 (2008).
90. Daniels, B. R., Dobrowsky, T. M., Perkins, E. M., Sun, S. X. & Wirtz, D. MEX-5 enrichment in the *C. elegans* early embryo mediated by differential diffusion. *Development* **137**, 2579–2585 (2010).
91. Turing, A. M. The chemical basis of morphogenesis. *Phil. Trans. R. Soc. Lond. B* **237**, 37–72 (1952).
92. Kondo, S. The reaction-diffusion system: a mechanism for autonomous pattern formation in the animal skin. *Genes Cells* **7**, 535–541 (2002).

93. Kaltschmidt, J. A., Davidson, C. M., Brown, N. H. & Brand, A. H. Rotation and asymmetry of the mitotic spindle direct asymmetric cell division in the developing central nervous system. *Nature Cell Biol.* **2**, 7–12 (2000).
94. Rebollo, E., Roldán, M. & Gonzalez, C. Spindle alignment is achieved without rotation after the first cell cycle in *Drosophila* embryonic neuroblasts. *Development* **136**, 3393–3397 (2009).
95. Rebollo, E. *et al.* Functionally unequal centrosomes drive spindle orientation in asymmetrically dividing *Drosophila* neural stem cells. *Dev. Cell* **12**, 467–474 (2007).
This study reveals how mitotic spindle orientation is established in *D. melanogaster* larval neuroblasts.
96. Januschke, J. & Gonzalez, C. The interphase microtubule aster is a determinant of asymmetric division orientation in *Drosophila* neuroblasts. *J. Cell Biol.* **188**, 693–706 (2010).
This paper shows how spindle orientation is kept constant over many divisions in *D. melanogaster* neuroblasts.
97. Spradling, A. C. & Zheng, Y. Developmental biology. The mother of all stem cells? *Science* **315**, 469–470 (2007).
98. Wang, X. *et al.* Asymmetric centrosome inheritance maintains neural progenitors in the neocortex. *Nature* **461**, 947–955 (2009).
99. Gotta, M., Dong, Y., Peterson, Y. K., Lanier, S. M. & Ahninger, J. Asymmetrically distributed *C. elegans* homologs of AGS3/PINS control spindle position in the early embryo. *Curr. Biol.* **13**, 1029–1037 (2003).
100. Colombo, K. *et al.* Translation of polarity cues into asymmetric spindle positioning in *Caenorhabditis elegans* embryos. *Science* **300**, 1957–1961 (2003).
101. Srinivasan, D. G., Fisk, R. M., Xu, H. & Van Den Heuvel, S. A complex of LIN-5 and GPR proteins regulates G protein signaling and spindle function in *C. elegans*. *Genes Dev.* (2003).
102. Nguyen-Ngoc, T., Afshar, K. & Gönczy, P. Coupling of cortical dynein and Gα proteins mediates spindle positioning in *Caenorhabditis elegans*. *Nature Cell Biol.* **9**, 1294–1302 (2007).
This study reveals how cortical polarity proteins connect to microtubules to position the mitotic spindle in *C. elegans*.
103. Ou, G., Stuurman, N., D'Ambrosio, M. & Vale, R. D. Polarized myosin produces unequal-size daughters during asymmetric cell division. *Science* **330**, 667–680 (2010).
104. Gateff, E. Malignant neoplasms of genetic origin in *Drosophila melanogaster*. *Science* **200**, 1448–1459 (1978).
105. Gateff, E. Tumor suppressor and overgrowth suppressor genes of *Drosophila melanogaster*: developmental aspects. *Int. J. Dev. Biol.* **38**, 565–590 (1994).
106. Lee, C. Y., Robinson, K. J. & Doe, C. Q. Lgl, Pins and aPKC regulate neuroblast self-renewal versus differentiation. *Nature* **439**, 594–598 (2006).
This study shows that mutations in genes regulating asymmetric cell division can cause overproliferation in the *D. melanogaster* brain.
107. Lee, C. Y. *et al.* *Drosophila* Aurora-A kinase inhibits neuroblast self-renewal by regulating aPKC/Numb cortical polarity and spindle orientation. *Genes Dev.* **20**, 3464–3474 (2006).
108. Wang, H. *et al.* Aurora-A acts as a tumor suppressor and regulates self-renewal of *Drosophila* neuroblasts. *Genes Dev.* **20**, 3453–3463 (2006).
109. Kitajima, A., Fuse, N., Isshiki, T. & Matsuzaki, F. Progenitor properties of symmetrically dividing *Drosophila* neuroblasts during embryonic and larval development. *Dev. Biol.* **347**, 9–23 (2010).
110. Basto, R. *et al.* Centrosome amplification can initiate tumorigenesis in flies. *Cell* **133**, 1032–1042 (2008).
111. Castellanos, E., Dominguez, P. & Gonzalez, C. Centrosome dysfunction in *Drosophila* neural stem cells causes tumors that are not due to genome instability. *Curr. Biol.* **18**, 1209–1214 (2008).
112. Caussinus, E. & Gonzalez, C. Induction of tumor growth by altered stem-cell asymmetric division in *Drosophila melanogaster*. *Nature Genet.* **37**, 1125–1129 (2005).
This article shows that defects in asymmetric cell division cause the formation of tumours on transplantation of defective stem cells in *D. melanogaster*.
113. Maurange, C., Cheng, L. & Gould, A. P. Temporal transcription factors and their targets schedule the end of neural proliferation in *Drosophila*. *Cell* **133**, 891–902 (2008).
114. Bello, B. C., Hirth, F. & Gould, A. P. A pulse of the *Drosophila* Hox protein Abdominal-A schedules the end of neural proliferation via neuroblast apoptosis. *Neuron* **37**, 209–219 (2003).
115. Siegrist, S. E., Haque, N. S., Chen, C. H., Hay, B. A. & Hariharan, I. K. Inactivation of both *foxo* and *reaper* promotes long-term adult neurogenesis in *Drosophila*. *Curr. Biol.* **20**, 643–648 (2010).
116. Reya, T., Morrison, S. J., Clarke, M. F. & Weissman, I. L. Stem cells, cancer, and cancer stem cells. *Nature* **414**, 105–111 (2001).
117. Cicalese, A. *et al.* The tumor suppressor p53 regulates polarity of self-renewing divisions in mammary stem cells. *Cell* **138**, 1083–1095 (2009).
This paper connects tumour formation in breast cancer with defects in asymmetric cell division.
118. Wu, M. *et al.* Imaging hematopoietic precursor division in real time. *Cell Stem Cell* **1**, 541–554 (2007).
119. Shinin, V., Gayraud-Morel, B., Gomes, D. & Tajbakhsh, S. Asymmetric division and cosegregation of template DNA strands in adult muscle satellite cells. *Nature Cell Biol.* **8**, 677–682 (2006).
120. Lechler, T. & Fuchs, E. Asymmetric cell divisions promote stratification and differentiation of mammalian skin. *Nature* **437**, 275–280 (2005).
121. Quyn, A. J. *et al.* Spindle orientation bias in gut epithelial stem cell compartments is lost in precancerous tissue. *Cell Stem Cell* **6**, 175–181 (2010).
122. Costa, M. R., Wen, G., Lepier, A., Schroeder, T. & Götz, M. Par-complex proteins promote proliferative progenitor divisions in the developing mouse cerebral cortex. *Development* **135**, 11–22 (2008).
123. Rasin, M. R. *et al.* Numb and Numb1 are required for maintenance of cadherin-based adhesion and polarity of neural progenitors. *Nature Neurosci.* **10**, 819–827 (2007).
This study changes our view of how mammalian homologues act during asymmetric cell division in the mouse brain.
124. Kosodo, Y. *et al.* Asymmetric distribution of the apical plasma membrane during neurogenic divisions of mammalian neuroepithelial cells. *EMBO J.* **23**, 2314–2324 (2004).
This study provides important insight into the mechanism of asymmetric cell division in the mouse brain.
125. Dontu, G. *et al.* *In vitro* propagation and transcriptional profiling of human mammary stem/progenitor cells. *Genes Dev.* **17**, 1253–1270 (2003).
126. Colaluca, I. N. *et al.* Numb controls p53 tumour suppressor activity. *Nature* **451**, 76–80 (2008).
127. Pece, S. *et al.* Loss of negative regulation by Numb over Notch is relevant to human breast carcinogenesis. *J. Cell Biol.* **167**, 215–221 (2004).
128. Ito, T. *et al.* Regulation of myeloid leukaemia by the cell-fate determinant Musashi. *Nature* **466**, 765–768 (2010).
129. Dietzl, G. *et al.* A genome-wide transgenic RNAi library for conditional gene inactivation in *Drosophila*. *Nature* **448**, 151–156 (2007).
130. Mummery-Widmer, J. L. *et al.* Genome-wide analysis of Notch signalling in *Drosophila* by transgenic RNAi. *Nature* **458**, 987–992 (2009).
131. Shendure, J. & Ji, H. Next-generation DNA sequencing. *Nature Biotech.* **26**, 1135–1145 (2008).
132. Lapidot, T. *et al.* A cell initiating human acute myeloid leukaemia after transplantation into SCID mice. *Nature* **367**, 645–648 (1994).
133. Bonnet, D. & Dick, J. E. Human acute myeloid leukemia is organized as a hierarchy that originates from a primitive hematopoietic cell. *Nature Med.* **3**, 730–737 (1997).
134. Singh, S. K. *et al.* Identification of a cancer stem cell in human brain tumors. *Cancer Res.* **63**, 5821–5828 (2003).
135. Dick, J. E. Breast cancer stem cells revealed. *Proc. Natl Acad. Sci. USA* **100**, 3547–3549 (2003).
136. Lobo, N. A., Shimono, Y., Qian, D. & Clarke, M. F. The biology of cancer stem cells. *Annu. Rev. Cell Dev. Biol.* **23**, 675–699 (2007).
137. Rosen, J. M. & Jordan, C. T. The increasing complexity of the cancer stem cell paradigm. *Science* **324**, 1670–1673 (2009).
138. Barker, N. *et al.* Crypt stem cells as the cells-of-origin of intestinal cancer. *Nature* **457**, 608–611 (2009).
139. Kwon, C. H. *et al.* *Pten* haploinsufficiency accelerates formation of high-grade astrocytomas. *Cancer Res.* **68**, 3286–3294 (2008).
140. Siller, K. H. & Doe, C. Q. Spindle orientation during asymmetric cell division. *Nature Cell Biol.* **11**, 365–374 (2009).
141. Santolini, E. *et al.* Numb is an endocytic protein. *J. Cell Biol.* **151**, 1345–1352 (2000).
142. Götz, M. & Huttner, W. B. The cell biology of neurogenesis. *Nature Rev. Mol. Cell Biol.* **6**, 777–788 (2005).
143. Bultje, R. S. *et al.* Mammalian Par3 regulates progenitor cell asymmetric division via notch signaling in the developing neocortex. *Neuron* **63**, 189–202 (2009).
144. Miyata, T., Kawaguchi, A., Okano, H. & Ogawa, M. Asymmetric inheritance of radial glial fibers by cortical neurons. *Neuron* **31**, 727–741 (2001).
145. Schwamborn, J. C., Berezikov, E. & Knoblich, J. A. The TRIM-NHL protein TRIM32 activates microRNAs and prevents self-renewal in mouse neural progenitors. *Cell* **136**, 913–925 (2009).

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Competing interests statement

The author declares no competing financial interests.

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