Mitotic spindle orientation can direct cell fate and bias Notch activity in chick neural tube

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Inheritance of apical membrane is proposed to maintain vertebrate neural stem cell proliferation. However, evidence for this is contradictory. Using direct clonal analysis and live imaging in chick neural tube, we show that divisions that separate apical and basal components generate an apical daughter, which becomes a neuron, and a basal daughter, which rapidly re-establishes apico-basal polarity and divides again. Using a recently described real-time reporter of Notch activity, we confirm progenitor status and demonstrate that division orientation can influence Notch signalling. In addition, we reveal loss of apical complex proteins on neuronal differentiation onset, suggesting that removal of this inherited complex is part of the neuronal differentiation mechanism. These findings reconcile contradictory data, link asymmetric division to Notch signalling dynamics and identify apical complex loss as a new step towards neuronal differentiation.

Keywords: vertebrate neurogenesis; asymmetric cell fate; mitotic spindle; Notch signalling; chick

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INTRODUCTION

The role of mitotic spindle orientation with respect to the apical/luminal surface of the neuroepithelium and the decision to adopt neuronal or progenitor cell fate remains controversial. In the cortex, data suggest that apical membrane promotes self-renewing neural stem cell divisions [1–6], but also indicate that inheritance of both apical and basal cell compartments is required to maintain neural progenitor status [7,8]. The spinal cord is a simpler tissue and is amenable to real-time imaging of individual cells [9,10]. In early chick neural tube, divisions that give rise to a neuron and a progenitor have cleavage planes non-perpendicular to the luminal surface, whereas divisions that generate progenitors divide mainly with a perpendicular cleavage plane [9]. One explanation for the more varied orientation of the latter is a lack of proneural gene expression at early stages that confers potential to adopt a neuronal fate [9,11]. These observations suggest that as proneural gene expression commences mitotic spindle orientation comes to influence cell fate choice. However, randomization of mitotic spindle position in chick neural tube analysed at a cell population level is reported to disrupt this tissue, but not to alter cell fates [12].

RESULTS

Apico-basal divisions increase neuron production

To investigate the role of mitotic spindle orientation in cell fate choice, a green fluorescent protein (GFP)-inscuteable (GFP-Insc)
expression vector was transfected into the stage HH10-12 chick neural tube by in ovo electroporation (which expresses low-level *Insc*; supplementary Fig S1 online). A minimal plasmid concentration that systematically generated apico-basally orientated divisions was determined (Fig 1A–C) and the consequences of such divisions were assessed after 48 h. Many GFP-expressing cells were found in the mantle zone with neuronal morphology (Fig 1D). In contrast, in control pCAGGS-GFP-transfected embryos, most GFP-expressing cells retained contact with the apical surface (Fig 1D, bottom panels). Increase in neuron numbers was then confirmed with the neuronal marker HuC/D (Fig 1D′) and post-mitotic marker p27 (Fig 1D′′; supplementary Fig S2A online), and decreased cell proliferation was indicated by reduced numbers of BrdU-incorporating cells (Fig 1E,F). These data demonstrate increased neuronal differentiation and depletion of the neural progenitor pool on induction of A/B divisions.

**Apico-basal divisions generate ectopic neural progenitors**

*Insc* mis-expression also led to abnormal distribution of BrdU-positive cells in the mantle zone (12/14 embryos; Fig 1E), which also expressed the neural progenitor marker Sox2 (5/5 embryos;
supplementary Fig S2B online). Previous work found that such ectopic proliferative cells lack A/B polarity [12] and such change is associated with oncogenic transformation of tissue [18]. However, in embryos transfects with GFP-Insc, apical Par-complex proteins Par3, aPKC and the sub-apical adhesion protein N-cadherin were additionally ectopically localized in the mantle zone (Par3, 10/12 embryos; aPKC, 9/10 embryos; N-cadherin 5/5 embryos; Fig 1F; supplementary Fig S2C,D online far right panels). The presence of ectopic apical protein in this progenitor cell population was then confirmed by localization of Par3 in BrdU-incorporating cells (Fig 1G); co-expression of N-cadherin and the G2/M phase marker pH3 (Fig 1H); and non-overlapping expression of Par3 and the post-mitotic marker p27 (26/26 cells; Fig 1I). Closer analysis revealed that Par3 was localized to one end of each ectopic progenitor cell (Fig 1I, white arrows) further supporting the retention of A/B polarity. In addition, experiments using a truncated version of the G-protein regulator LGN (Ct-cLGN) that interferes dominantly with endogenous LGN was used to compare the effects of Insc-induced A/B divisions and the randomization of the mitotic spindle (after [12]). These experiments showed that although less effective, randomization also altered cell fates and lead to ectopic Par-complex protein expression in the mantle layer (supplementary Fig S3A,A online). Overall, induction of A/B divisions in the spinal cord increases neuron numbers and creates an ectopic population of neural progenitor cells that exhibit A/B polarity.

**Fig 2 | Insc mis-expression induces stem cell mode divisions.** (A) Clonal analysis reveals induction of a neuron and a progenitor by GFP-Insc mis-expression (see text for details). (B) Following an A/B division (1 h 52 min–2 h 55 min), the apical daughter cell extends a new basal process (4 h 19 min, white arrows) and the basal daughter extends a new apical process (5 h 43 min, yellow arrows), which re-establishes apical surface (white dotted line) contact (11 h 47 min, yellow asterisk). (C) Cell indicated with white dot undergoes A/B division, generating a basal daughter (red dot) that remains as a progenitor and divides again (20 h 39 min) and an apical daughter (yellow dot) that extends a new basal process (6 h 32 min) and undergoes neuronal differentiation (27 h 53 min). (D) Following an A/B division (starting at 21 min), Par3 GFP is restricted to an apical crescent that is inherited by the apical daughter (41–54 min). Scale bars, 10 μm. GFP, green fluorescent protein; Insc, Insuteable.

**Apico-basal divisions generate a neuron and a progenitor** To determine the fates of daughter cells generated by A/B divisions, single cells mis-expressing GFP-Insc or control plasmid pCAGGS-GFP and KiKGR, a photo-convertible fluorescent protein (see Methods) were marked by UV-laser-mediated photo-conversion and slices incubated for 24 h (at least one cell cycle) and then imaged (Fig 2A). The majority of cells (7/9) co-expressing GFP-Insc and KiKGR gave rise to one cell that spanned the neural tube width (in the manner of a progenitor), whereas the other adopted a neuronal morphology (lacking apical process attachment and projecting an axon along the neural tube perimeter). In contrast, all cells expressing control pCAGGS-GFP and KiKGR (8/8) produced progeny with cell shape profiles that extended across the neural tube width and lacked neuronal morphology. This direct clonal analysis demonstrates that A/B divisions produce cells with asymmetric fates, a neuron and a progenitor.
Apical cell differentiation and basal cell re-polarization

Long-term time lapse was used to determine cell behaviour following Insc-induced division. Cells transfected with pCAGGS-inscuteable-IRE5-nucGFP (pCIG-Insc) and mApple-Farn (membrane marker) divided apico-basally, with the basal daughter retaining the original basal process while its apical sibling produced a new one (n = 7/7; Fig 2B; supplementary Movie S1 online). In all cases, the basal cell also extended a new apical process; even when these cells became located laterally within the neural tube this new process strikingly re-attached the cell to the luminal surface (Fig 2B; supplementary Movie S1 online). To extend these observations, time-lapses of cells expressing pCIG-Insc and GFP-GPI (monitored through a single filter set, giving increased cell viability) were performed. In divisions where the fate of the apical daughter was clear, the majority of cells became neurons (13/15). Conversely, most basal daughter cells divided again (17/19; Fig 2C; supplementary Movie S2 online) with a small subset (4/19) dividing ectopically in the mantle zone (supplementary Fig S4A online; supplementary Movie S3 online). This determines the identity of the ectopic BrDU-positive progenitor cell population (see Fig 1E) as mis-placed basal daughter cells and as such mis-localized cells express apical markers this further supports re-establishment of A/B polarity in basal daughters.

To address directly whether basal daughter cells re-acquire apical polarity or inherit a small portion of apical membrane, Par3 localization was analysed in cells co-transfected with pCAGGS--inscuteable-IRE5-H2B RFP and low-level (25 ng/μl) pCAGGS-Par3-GFP (Par3-GFP). In the resulting A/B divisions, all detectable Par3-GFP was inherited by the apical daughter (7/7 divisions; Fig 2D; supplementary Movie S4 online). To confirm this, inheritance of endogenous Par3 protein was also analysed using immunocytochemistry and the apical cell was again found to inherit most of this protein (36/36 A/B divisions; supplementary Fig S4B online). These data suggest that the rapid re-polarization of the basal daughter involves de-novo synthesis of apical complex proteins.

Asymmetric inheritance during normal neurogenesis

It has been suggested that small changes in spindle orientation are sufficient to partition A/B components in such divisions [1,2,19]. To investigate this in the chick neural tube, we monitored inheritance of Par3 at mitosis in cells transfected with either H2B-RFP or mApple-Farn together with low levels of Par3-GFP to visualize normal Par3 localization. This revealed divisions with moderately non-perpendicular cleavage planes in which Par3-GFP was inherited largely by only one daughter cell (n = 7/7; Fig 3A; supplementary Movie S5 online). Furthermore, in divisions in which cellular processes were visible and subsequent cell fates of both daughter cells could be established, neuronal cell fate correlated with retention of an apical process, whereas the basal sibling grew a new apical process and divided again (n = 6; Fig 3B; supplementary Movie S6 online). These observations suggest that small changes in mitotic spindle orientation during normal neurogenesis can lead to asymmetric segregation of apical membrane and that re-growth of an apical process by basal cells is characteristic of normal neural stem cell mode divisions in the chick spinal cord.

Notch signalling is elevated in basal daughter cells

Using a novel live reporter for Notch signalling pHeS5-VNP, based on the chicken Hes5-1 promoter-driving expression of a destabilized nuclear Venus fluorescent protein coupled with the Hes5-1 3′ UTR to confer instability [20] (supplementary Methods online), we and our collaborators have shown that most Notch-active cells undergoing mitosis in the chick neural tube at HH10-12 generate two Notch-active daughters, consistent with most cells remaining progenitors at this stage [20]. However, a subset of cells generated only one Notch-active daughter; this could reflect the influence of non-cell autonomous signals and/or asymmetric cell division. Here we investigated Notch signalling following induction of A/B divisions by co-transfection of pCAGGS-Insc-IRE5-H2B-RFP and the pHeS5-VNP reporter. This revealed that all basal daughters increased Notch signalling, whereas the apical daughter had low or no activity (a threefold increase above that in the apical daughter was detected between 7 and 10 h, n = 6; the precise time of activation is influenced by level of transfected reporter plasmid; Fig 3C; supplementary Figs S4 and S5 online; supplementary Movie S7 online). This differential activation indicates that Notch signalling can be influenced by mitotic spindle orientation.

Loss of apical protein complex by apical daughter cells

Work in the mouse cortex suggests that Par3 promotes progenitor status [21] via Notch signalling [22]. Par3 mis-expression in the chick neural tube also generates a phenotype consistent with this [23]. These observations appear to contradict the asymmetric inheritance of the apical complex by the apical daughter as it then differentiates into a neuron. However, we show above that A/B polarity is rapidly re-established in basal cells that behave as progenitors and so the Par-complex is also characteristic of progenitor cells. One possibility is that apical daughters, although inheriting the majority of the Par-complex, quickly lose this as they differentiate. Tuj-1 marks cells commencing neuronal differentiation [24]. We found that such Tuj-1 cells lack both Par3 (n = 76) and aPKC (n = 54) at the tip of the apical process, which is withdrawn as differentiation commences [9] (Fig 3D). This suggests that rapid loss of the apical protein complex is part of the neuronal differentiation mechanism.

DISCUSSION

We show here that induction of A/B divisions generates an increase in neuron numbers, and by carrying out direct clonal analysis and real-time monitoring of individual cells we demonstrate that A/B divisions generate a neuron and a progenitor. We conclude that mitotic spindle orientation can influence cell fate choice in the chick neural tube (summarized in Fig 4). This contrasts with a previous study in this tissue, in which mitotic spindle position was randomised and no change in cell fates was detected [12]. We reconcile this by demonstrating that randomization of mitotic spindle position can elicit a slight increase in neurons, consistent with the differing incidence of A/B divisions generated by these two approaches. These different findings might also be explained by use of a different technical approach, retrospective clonal analysis dependent on a co-transfection strategy [12].

Randomizing mitotic spindle orientation or inducing A/B divisions also generates ectopic neural progenitor cells ([7,12] and this study) and we find that these cells possess A/B polarity. By monitoring the behaviour of individual Insc-expressing cells in real-time, we further identify the origin of these ectopic
progenitors as displaced basal daughters that fail to re-contact the apical surface (supplementary Fig S4A online). The finding that on division Par3 largely localizes in the apical daughter further suggests that re-polarization of the basal cell may involve synthesis of new apical complex proteins as well as re-localization of any inherited proteins. Our observations of normal neurogenesis reveal that moderately non-perpendicular cleavage planes can generate daughter cells with asymmetric inheritance of the Par-complex and that basal daughter cells re-grow an apical process. This strongly suggests that re-establishment of apical polarity is a normal neurogenesis step (Fig 4), in keeping with observations in the zebrafish [10].

In the vertebrate cortex, the prevailing hypothesis identifies apical properties as critical determinants of the self-renewing progenitor cell state, reviewed in Gotz and Huttner [19]. However, there is now evidence that possession of both apical and basal components correlates with progenitor status in the cortex [7,8] and in the neural tube ([10], this study). Furthermore, cells that inherit only apical components tend to exit the cell cycle in the cortex [7] and become neurons in fish [10] and here in the chick neural tube. In the cortex, the incidence of differential apical process inheritance is reported to be insufficient to account for cortical neurogenesis [7]. However, given the rapid regulation of A/B polarity in basal daughter cells observed by live imaging, here and in the zebrafish [10], the incidence of this event may have been under-estimated.

Our data and that of Alexandre et al [10] and Konno et al [7] point to inheritance of the original basal process as the identifier of the progenitor daughter. Although there is no clear correlation between bisection of the basal process and adoption of symmetrical cell fates in cortex or spinal cord [14,25,26], the basal process of radial glial cells, which appear at later stages in the cortex, is clearly inherited by the daughter cell that continues as a progenitor [27,28]. This is consistent with observations in the neural tube and supports a requirement for basal process inheritance for maintenance of this established neural stem cell.

It is clear that mitotic spindle orientation is just one influence on cell fate choice and that signalling via Notch and other pathways also directs this process. This study shows, for the first time, that
Notch signalling is differentially activated following induction of an A/B division in the live neuroepithelium, suggesting that mitotic spindle orientation influences Notch activity during normal neurogenesis. The systematic detection of elevated Notch activity in the basal daughter also confirmed the progenitor status of this cell and strongly suggests that asymmetric division results in a cell intrinsic difference in the ability of sibling cells to respond to Notch signalling; although it is also formally possible that asymmetric partition of cellular components leads to changes that affect the ability of neighbouring cells to deliver Notch signalling.

These data are consistent with the finding that the ability of Par3 to promote the neural progenitor cell state depends on Notch signalling [22]. These authors and others [21,23] conclude that inheritance of Par3 promotes progenitor cell fate, but it is likely that the cellular context in which Par3 is expressed determines its action. We show that while the basal daughter expresses Par3 and exhibits Notch activity, the apical daughter cell inherits Par3 but lacks Notch signalling. However, we also find that the apical daughter rapidly downregulates Par3 as neuronal differentiation commences. Retraction of the apical end-foot by differentiating neurons occurs in spinal cord and cortex [9,29] and must involve detachment from abutting apical end-feet of neighbouring cells. Indeed, this loss of the apical protein complex may facilitate release from the luminal surface; a critical physical requirement for normal neuronal differentiation. In the future, it will be important to investigate the mechanisms underlying re-establishment of A/B polarity in neural progenitors and the loss of Par-complex proteins as cells embark on neuronal differentiation.

**METHODS**

**In ovo electroporation and plasmids.** Spinal cords were transfected using standard conditions. GFP-Insc (pCAGGS-GFP-Insc) was a kind gift of Domingos Henrique. The Ct-cLGN-IRES-GFP-GPI construct was kindly provided by Xavier Morin. See supplementary Methods online for further detail.

**Immunofluorescence.** Immunocytochemistry was performed on a 20-μm thick cryosections following standard procedures (supplementary Methods online).

**Embryo slice culture and time-lapse imaging.** Slices were prepared, cultured and imaged as previously described [9,30] (supplementary Methods online).

**Clonal analysis.** Single cells were labelled and their lineages traced using the photo-convertible fluorescent protein KiKGR (MBL International; supplementary Methods online).

**Angle measurements.** Cleavage plane orientations were measured in anaphase and telophase cells (after [9]; supplementary Methods online).

**Statistical analysis.** All cell counts involving percentages of GFP-positive cells expressing markers were compared using unpaired t-tests; electroporated versus unelectroporated sides were assessed using paired t-tests.

**Supplementary information** is available at EMBO reports online (http://www.emboreports.org).

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**Author contributions.** K.G.S. conceived the project. K.G.S. and R.M.D. designed experiments. R.M.D. carried out all the experiments. K.G.S. and R.M.D. analysed the data and wrote the manuscript.
CONFLICT OF INTEREST
The authors declare that they have no conflict of interest.

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