**miR-219 Regulates Neural Precursor Differentiation by Direct Inhibition of Apical Par Polarity Proteins**

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**SUMMARY**

Asymmetric self-renewing division of neural precursors is essential for brain development. Partitioning-defective (Par) proteins promote self-renewal, and their asymmetric distribution provides a mechanism for asymmetric division. Near the end of neural development, most asymmetric division ends and precursors differentiate. This correlates with Par protein disappearance, but mechanisms that cause downregulation are unknown. MicroRNAs can promote precursor differentiation but have not been linked to Par protein regulation. We tested a hypothesis that microRNA miR-219 promotes precursor differentiation by inhibiting Par proteins. Neural precursors in zebrafish larvae lacking miR-219 function retained apical proteins, remained in the cell cycle, and failed to differentiate. *miR-219* inhibited expression via target sites within the 3' untranslated sequence of *pard3* and *prkci* mRNAs, which encode Par proteins, and blocking *miR-219* access to these sites phenocopied loss of *miR-219* function. We propose that negative regulation of Par protein expression by *miR-219* promotes cell-cycle exit and differentiation.

**INTRODUCTION**

Following neural induction in vertebrate embryos, neural precursor cells initially divide symmetrically to expand the precursor population. After a period of tissue growth, many precursors switch to asymmetric division, producing both new precursors to maintain the population and cells that differentiate as neurons or glial cells, either directly or after a limited number of transit-amplifying divisions. Near the end of embryogenesis, most precursors exit the cell cycle and differentiate, although some may become quiescent or be maintained into adulthood as stem cells. A balance of symmetric proliferative divisions, asymmetric self-renewing divisions, and terminal differentiation assures production of appropriate numbers of neural cells to build a functional brain (reviewed in Huttner and Kosodo, 2005; Gönçzy, 2008). Mechanisms that maintain that balance remain incompletely understood.

One mechanism that clearly contributes to neural precursor maintenance and differentiation involves neuroepithelial polarity (reviewed in Shitamukai and Matsuzaki, 2012; Fietz and Huttner, 2011; Götz and Huttner, 2005). After closure of the neural tube, apical membranes of neural precursors line a lumen, which subsequently forms the ventricles and central canal of the central nervous system. Associated with the apical membrane are various protein complexes, including one consisting of the partitioning-defective (Par) proteins Pard3, Pard6, and Prcki, also known as atypical protein kinase C (aPKC). Experiments performed in *Drosophila* provide strong evidence that unequal distribution of apical Par proteins during neural precursor division determines the fate of the progeny cells. In particular, cells that inherit apical Par proteins remain as precursors, whereas those that do not enter a differentiation pathway (Prehoda, 2009).

Data from vertebrate models are generally compatible with the idea that, in asymmetric divisions, high levels of apical Par proteins are associated with neural precursor self-renewal, whereas low levels lead to differentiation (Bultje et al., 2009; Costa et al., 2008; Kosodo et al., 2004), although studies performed in zebrafish indicated that Pard3 can inhibit self-renewal and promote differentiation (Alexandre et al., 2010; Dong et al., 2012). The basis for these apparently conflicting conclusions is not known.

At the end of embryogenesis, the uniform disappearance of Par proteins from apical membranes of neural precursors correlates with their terminal exit from the cell cycle and differentiation (Costa et al., 2008). Although the mechanistic basis of unequal Par protein distribution to progeny cells undergoing asymmetric self-renewing division has been thoroughly investigated, no mechanism that promotes disappearance of Par proteins from precursors fated to undergo terminal differentiation has been described.

microRNAs (miRNAs) are small noncoding RNAs that in most circumstances inhibit gene expression by mediating degradation or translational inhibition of target mRNAs (reviewed in Bartel, 2009). miRNAs regulate numerous developmental processes, but identification of authentic miRNA targets and the mechanistic functions that targeting fulfills in vivo remain daunting because most miRNAs potentially bind dozens, if not hundreds, of target mRNAs, and many miRNAs have multiple sequences that could be bound by miRNAs (reviewed in Bartel, 2009; Asuelime and Shi, 2012). Nevertheless, recent data have begun to identify some roles for miRNAs in regulating neural precursor division and differentiation. In particular, several studies using mouse, zebrafish, and frog have shown that *miR-9* can drive precursors from the cell cycle and promote neuronal...
determination by regulating expression of multiple transcription factors, including Foxg1, Meis2, Gsh2, Hes1, Her6, Zic5, and Hairy1 (Shibata et al., 2011; Tan et al., 2012; Coolen et al., 2012; Bonev et al., 2011). Neuronal differentiation also is driven by miR-9 and let-7b through inhibition of the nuclear receptor TLX (Zhao et al., 2009) and by miR-124, which represses expression of the RNA-binding protein PTBP (Makeyev et al., 2007) and the transcription factor Sox9 (Cheng et al., 2009). Given the large number of miRNAs expressed by neural cells and their multitudes of potential target mRNAs, these advances likely provide only a partial understanding of the mechanisms by which miRNAs regulate neural precursor maintenance and differentiation.

Here, we report evidence of neural precursor regulation by miR-219, an evolutionarily conserved miRNA expressed in zebrafish brain (Kapsimali et al., 2007). We previously noted that knockdown of miR-219 in zebrafish caused a deficit of oligodenodcyte progenitor cells (OPCs) (Zhao et al., 2010), raising the possibility that miR-219 promotes formation of glial cells from neural precursors. Target prediction software identified potential miR-219 target sites within the 3′ UTR sequences of Pard3 and Prkci genes in multiple species. Therefore, we hypothesized that miR-219 promotes transition of neural precursors from self-renewal to differentiation by negatively regulating Pard3 and Prkci. Our work now shows that pardi3 and prkci mRNAs are functionally relevant targets of miR-219 in zebrafish. Loss of miR-219 function results in prolonged maintenance of apically localized proteins and retention of neural precursors in the cell cycle at the expense of late-born neurons and glial cells, indicating that miR-219 contributes to a microRNA-based mechanism that promotes neural cell differentiation.

RESULTS

miR-219 Promotes Neural Precursor Exit from Proliferative Division

To test the hypothesis that miR-219 promotes neural precursor differentiation, we first set out to better characterize changes in precursor characteristics during zebrafish spinal cord development. In cat (Böhme, 1988) and rat (Sevc et al., 2009), a primitive lumen, extending across nearly the entire extent of the dorsoventral axis of the neural tube, transforms into a much smaller, ventrally positioned central canal. Initiation of this transformation coincides with exit of neural precursors from the cell cycle and completion of neurogenesis in late embryonic stage. The zebrafish spinal cord undergoes a similar transformation. At 1 and 2 days postfertilization (dpf), the tight junction-associated protein ZO-1 was enriched at the apical membranes of cells lining a primitive lumen spanning the spinal cord (Figures 1A and 1B). At 3 and 5 dpf, apical ZO-1 enrichment was no longer evident dorsally but remained associated with a central canal in ventral spinal cord (Figures 1C and 1D). F-actin was similarly localized, lining the primitive lumen at 1 and 2 dpf (Figures 1E and 1F) but limited to the central canal at 3 and 5 dpf (Figures 1G and 1H). This loss of apical protein enrichment coincided with loss of neural precursors. Whereas numerous cells lining the primitive lumen expressed Sox2, a marker of neural precursors (Ellis et al., 2004), at 1 and 2 dpf (Figures 1E and 1F), fewer Sox2+ cells remained at 3 and 5 dpf, and these were mostly restricted to the area bordering the central canal (Figures 1G and 1H). Similarly, many cells lining the primitive lumen incorporated the thymidine analog bromodeoxyuridine (BrDU), a marker of cells in S phase, at 1 and 2 dpf (Figures 1I and 1J), but few spinal cord cells incorporated BrDU at 3 and 5 dpf (Figures 1K and 1L). Therefore, by early larval stage most spinal cord precursors differentiate or become mitotically quiescent, concomitant with transformation of a primitive lumen to a central canal and loss of apically enriched proteins.

To investigate if miR-219 expression correlates with the morphological and neural precursor changes described above, we first performed quantitative PCR. This showed that miR-219 levels were relatively low prior to 1 dpf and then peaked at 2 dpf (Figure S1 available online). We then carried out in situ hybridization using a locked nucleic acid (LNA) probe designed to detect the mature miR-219 sequence. At 1 dpf, spinal cord cells expressed miR-219 very weakly (Figure 2A), but, by 2 dpf, miR-219 expression was robust (Figure 2B). Notably, miR-219 expression appeared to be highest in the cells just outside the proliferative ventricular zone and somewhat lower in the neural precursors immediately adjacent to the primitive lumen. At 3 dpf, after transformation of the primitive lumen to the central canal, cells throughout the medial spinal cord expressed miR-219 at uniform levels (Figure 2C). At 5 dpf, following completion of embryonic neurogenesis and gliogenesis, miR-219 expression in spinal cord was no longer evident (data not shown). Therefore, the pattern of miR-219 expression is consistent with the possibility that it regulates cell division and differentiation.

To test miR-219 function, we investigated markers of neural precursors in control larvae and larvae injected with antisense morpholino oligonucleotides (MOs) (Zhao et al., 2010) designed to bind the mature form of miR-219, thereby blocking function. In 3 dpf control larvae, few spinal cord cells incorporated BrDU, indicating a low level of neural precursor division (Figure 2D). By contrast, numerous BrDU+ cells occupied the medial spinal cord of miR-219 MO-injected larvae (Figure 2E). Quantification revealed elevated number of BrDU+ cells in both dorsal and ventral spinal cord, with an approximately 2-fold greater increase in dorsal spinal cord than in ventral spinal cord (Figure 2F). Similarly, immunohistochemistry to detect phosphohistone H3 (PH3), which reveals cells in M phase, showed that miR-219 MO-injected larvae had substantially more PH3+ spinal cord cells than control larvae (Figures 2G–2I). These data indicate that miR-219 drives neural precursors from the cell cycle. To determine if neural cells retain other precursor characteristics in the absence of miR-219 function, we investigated expression of Sox2 and localization of apically associated proteins. In 3 dpf control larvae, Sox2+ cells were primarily located around the central canal, revealed by F-actin labeling, in the ventral spinal cord (Figure 2J). In miR-219 MO-injected larvae, Sox2+ cells lined the entire dorsoventral axis of the medial spinal cord (Figure 2K). Quantification revealed that miR-219 MO-injected larvae had approximately 1.75-fold more Sox2+ cells than control larvae, but that the difference was limited to dorsal spinal cord (Figure 2L). Additionally, F-actin labeling revealed an enlarged lumen extending into dorsal spinal cord (Figures 2K and 2L’). Similarly, whereas ZO-1 was limited to membranes surrounding the central canal of 3 dpf control larvae (Figures 2M and 2M’), ZO-1 labeling was evident along the entire dorsoventral extent.
of an enlarged lumen of miR-219 MO-injected larvae (Figures 2N and 2N’). These phenotypic features are characteristic of early neural development. Therefore, loss of miR-219 function results in retention of embryonic spinal cord characteristics into post-embryonic, larval stage.

Our data show that larvae lacking miR-219 function maintain excess spinal cord precursors, raising the possibility that the OPC deficit we observed previously (Zhao et al., 2010) resulted from a failure of precursor differentiation. To test this, we examined radial glial and neuronal markers in combination with EGFP expression driven by an olig2:EGFP transgene, which marks pMN precursors, motor neurons, and oligodendrocyte lineage cells (Shin et al., 2003). In wild-type zebrafish, radial glia appear by 1 dpf and persist throughout development and into adulthood (Park et al., 2007). Accordingly, numerous GFAP+ radial glia were evident in the spinal cords of 3 dpf control larvae (Figure 3A). By contrast, GFAP+ radial glia were completely absent from miR-219 MO-injected larvae, except in the most dorsal and ventral portions of the spinal cord (Figure 3B). Consistent with our previous results, miR-219 MO-injected larvae also had few OPCs. Expression of the radial glia marker BLBP showed a similar pattern, marking numerous cells in control larvae (Figure 3C), but only a few cells in the most dorsal and ventral portions of the spinal cord of miR-219 MO-injected larvae (Figure 3D). To investigate neuronal differentiation, we examined expression of Elavl3, which marks newly born neurons (Marusich et al., 1994). This revealed no obvious differences in neurons in wild-type and miR-219 MO-injected larvae (Figures 3E and 3F). To quantify a specific population of neurons, we counted motor neurons, which are produced by ventral spinal cord precursors and marked by Islet (Isl) expression. Indeed, motor neuron number in miR-219 MO-injected larvae was not different from control at 3 dpf (Figure 3G). To investigate whether late stages of neurogenesis might be specifically affected by miR-219 loss of function, we performed a fate-mapping experiment. In particular, we labeled dividing cells at 24 hr postfertilization (hpf) with a pulse of the thymidine analog EdU, waited 24 hr to permit cell differentiation, and examined EdU distribution. In control embryos,
Figure 2. miR-219 Knockdown Causes Retention of Embryonic Spinal Cord Precursors into Postembryonic Stage

All images show representative transverse sections through trunk spinal cord with dorsal up.

(A–C) In situ hybridization to detect the mature form of miR-219 reveals low expression at 1 dpf but prominent expression at 2 dpf. Staining appears most intense in cells (asterisks) lateral to ventricular zone cells (vz, bracket) lining the primitive lumen. At 3 dpf, miR-219 expression appears uniform in medial spinal cord. Boxed areas are shown in higher magnification in (A0–C0).

(D and E) miR-219 (3 dpf) MO-injected larva has more cells that incorporate BrdU than a control larva.

(F) Graph showing the number of BrdU+ cells in the ventral, dorsal, and entire spinal cord at 3 dpf. Data represent the mean ± SEM (n = 10 larvae, five to ten sections each). ****p < 0.0001, unpaired t test.

(G and H) miR-219 (3 dpf) MO-injected larva has more PH3+ cells than a control larva.

(I) Graph showing the number of spinal cord cells labeled with anti-PH3 antibody in control and miR-219 MO-injected larvae. Data represent the mean ± SEM (n = 10 sections obtained from 15 larvae per group, with two replicates). *p = 0.0328, unpaired t test.

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many of the EdU+ cells were also Elavl3+, representing precursors that had left the cell cycle and differentiated as neurons (Figure 3H). By contrast, miR-219 MO-injected embryos retained most of the EdU label within the proliferative zone and had correspondingly few EdU+ Elav3+ neurons (Figure 3I). Quantification confirmed this, showing that miR-219 MO-injected embryos had approximately half as many EdU+ Elav3+ neurons as control embryos (Figure 3J). Together with the data presented above, these results indicate that miR-219 promotes exit of neural precursors from the cell cycle and their differentiation as neurons and glia at late stages of development.

miR-219 Regulates pard3 and prkci via 3’ UTR Target Sites

Because zebrafish larvae retain neuroepithelial precursor characteristics in the absence of miR-219 function, we hypothesized that miR-219 promotes downregulation of factors necessary for maintaining precursors in an undifferentiated, dividing state. To search for such factors, we used target prediction software to identify miR-219 binding sites. Strikingly, this revealed single putative target sites within the 3’ UTR sequences of the Par genes pard3 and prkci that are conserved among numerous vertebrate species (Figures 4A and 4B). Target prediction analysis also

Figure 3. miR-219 Is Required for Differentiation of Glia and Late-Born Neurons
All images show representative transverse sections through trunk spinal cord with dorsal up.
(A and B) Whereas numerous GFAP+ radial glia occupy the spinal cord of a 3 dpf control larva, a miR-219 MO-injected larva has few radial glia except for the most dorsal and ventral regions of spinal cord (brackets). Asterisks mark oligodendrocyte lineage cells.
(C and D) Images showing a deficit of BLBP+ radial glia in a miR-219 MO-injected larva compared to control.
(E and F) miR-219 MO-injected larvae appear to have a normal number and distribution of neurons, marked by Elavl3 expression, but fewer oligodendrocyte lineage cells (asterisks) than control larvae.
(G) Graph showing number of Isl+ motor neurons in control and miR-219 MO-injected larvae. Data are presented as mean ± SEM (n = 10 sections obtained from 15 larvae per group, with two replicates). p > 0.05, unpaired t test.
(H and I) Confocal images of embryos pulsed with EdU at 1 dpf and fixed at 2 dpf. Numerous EdU+ cells are also Elavl3+ (arrowheads) in control larvae (H), whereas most EdU label persists within cells lining the spinal cord lumen, and fewer neurons are labeled by EdU in miR-219 MO-injected larva (I).
(J) Graph showing number of Elavl3+ EdU+ neurons in control and miR-219 MO-injected larvae. Data represent mean ± SEM (n = 10 sections obtained from five larvae per group). ***p < 0.0001, unpaired t test. Scale bar equals 10 μm.

(J–L) miR-219 (3 dpf) MO-injected larvae have more Sox2+ cells than control larvae. Graph showing the number of Sox2+ cells in ventral, dorsal, and entire spinal cord. Data in graph represent mean ± SEM (n = 10 sections obtained from 15 larvae per group, with three replicates). *p = 0.0369, ****p < 0.0001, unpaired t test. (J, K, M, and N) miR-219 (3 dpf) MO-injected larvae maintain a primitive lumen marked by apically localized F-actin and ZO-1 (brackets). Boxed areas are shown at higher magnification in (J’), (K’), (M’), and (N’). Scale bar equals 10 μm for low-magnification images and 5 μm for high-magnification images. See also Figure S1.
revealed single sites within the 3' UTR sequences of *pard6b* and *pard6ga* but not those of *pard6a* and *pard6gb*. Because apical localization of Par proteins is a hallmark feature of neuroepithelial precursors and Par protein functions promote neural precursor self-renewal in flies and mice (Bultje et al., 2009; Rolls et al., 2003; Kuchinke et al., 1998; Lee et al., 2006; Costa et al., 2008), we chose *pard3* and *prkci* for further investigation.

To investigate whether Par protein localization coincides with neural precursor state, we performed immunohistochemistry using an anti-Prkci antibody (Horne-Badovinac et al., 2001; Roberts and Appel, 2009). Similar to ZO-1 and F-actin, Prkci was enriched at apical membranes along the entire spinal cord dorsoventral axis at 1 and 2 dpf, but then limited to membranes lining the central canal by 3 dpf (Figures 4C–4E). By contrast, apical Prkci persisted throughout the spinal cord of *miR-219* MO-injected larvae at 3 and 5 dpf (Figures 4F and 4G). These data are consistent with the possibility that *miR-219* promotes the disappearance of Prkci from differentiating neuroepithelial precursors with a concomitant loss of apical membrane characteristics.

We lack an antibody that reliably detects Pard3 protein in zebrafish tissue. Therefore, to test whether *miR-219* can regulate *pard3* expression, we cloned the *pard3* 3' UTR sequence containing the predicted target site into a dual luciferase reporter plasmid (Figure 4H), cotransfected the construct into HEK293T cells with either a negative control mimic or *miR-219* mimic, and measured luciferase activity. The *miR-219* mimic, but not the control mimic, reduced luciferase expression by 70% (Figure 4I). Introduction of one and two base mutations in the target site abrogated *miR-219*-mediated inhibition of luciferase expression (Figure 4I), indicating that *miR-219* regulates *pard3* expression by binding the predicted target site.

To test the ability of *mir-219* to regulate target expression in vivo by fusing the *pard3* and *prkci* 3' UTRs to cDNA encoding EGFP and using the constructs as templates for in vitro synthesis of mRNA, which we injected into zebrafish embryos at early cleavage stage. We coinjected some embryos with *miR-219* or, as a control, *miR-216a*, which has no predicted target sites within the *pard3* and *prkci* 3' UTRs. To quantify EGFP expression, we measured fluorescence intensity within a defined region of the spinal cord. Whereas EGFP fluorescence appeared to be similar in embryos injected only with RNA including the *prkci* 3' UTR and those coinjected with *miR-216a*, fluorescence intensity

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**Figure 4. miR-219 Has Single, Conserved Target Sites within prkci and pard3 3' UTRs**

(A and B) Schematic representations of prkci and pard3 transcripts with predicted miR-219 target sites conserved among various species.

(F and G) Prkci labeling persists along a primitive lumen extending across the spinal cord dorsoventral axis in 3 dpf and 5 dpf *miR-219* MO-injected larvae.

(H) Sequences (220 bp) from the pard3 3' UTR containing wild-type and mutated miR-219 target sites were cloned into dual luciferase vectors.

(I) Quantification of light units revealed a miR-219-mediated reduction of reporter gene expression that was abrogated by 1 and 2 bp mutations within the target site. Data represent ± SEM (three independent experiments). Brackets indicate pairwise comparisons. ****p < 0.0001, unpaired t test. Scale bar equals 10 μm.
in embryos coinjected with \( \text{miR-219} \) appeared to be less than control (Figures 5A–5C). Quantification confirmed this, revealing an approximately 30% reduction in fluorescence intensity in \( \text{miR-219} \)-injected embryos relative to the controls (Figure 5D). The \( \text{pard3} \) 3′ UTR mediated a similar effect (Figures 5E–5H), although the degree to which fluorescence intensity was reduced was less than that mediated by the \( \text{prkci} \) 3′ UTR. EGFP fluorescence produced by RNA having neither the \( \text{pard3} \) nor \( \text{prkci} \) 3′ UTRs was not affected by coinjection of \( \text{miR-219} \) (Figures 5I–5K).

We next performed a series of experiments to investigate whether \( \text{pard3} \) and \( \text{prkci} \) are functionally relevant in vivo targets of \( \text{miR-219} \). First, we reasoned that if the oligodendrocyte deficit of \( \text{miR-219} \) MO-injected larvae results from elevated levels of Par proteins, then reduction of Par protein function should restore oligodendrocyte number. To test this prediction, we assessed dorsally migrated oligodendrocytes, marked by Tg(olig2:EGFP) reporter expression, formed in larvae injected with only \( \text{miR-219} \) MO or coinjected with \( \text{miR-219} \) MO and MO designed to block translation of \( \text{pard3} \) mRNA (Alexandre et al., 2010). Consistent with our previous observations, \( \text{miR-219} \) MO-injected larvae had few dorsally migrated OPCs compared to control (Figures 6A and 6B). By contrast, coinjection of \( \text{miR-219} \) MO and \( \text{pard3} \) MO partially restored OPC number (Figure 6C). To quantify this effect, we classified phenotypes as normal, mild, or severe based on the number of dorsally migrated OPCs. Whereas most larvae lacking only \( \text{miR-219} \) function were classified as severe, significantly more larvae lacking both \( \text{miR-219} \) and...
**DISCUSSION**

The progression of neural precursors from symmetric proliferative to asymmetric self-renewing division and, subsequently, to symmetric cell-cycle exit and differentiation represent key transitions in neural development. Data drawn from investigations of invertebrate and vertebrate neural precursors provide the basis for a model in which symmetric or asymmetric distribution
of apical Par proteins to sibling cells at mitosis influences whether each remains as a precursor or enters a differentiation pathway (reviewed in Homem and Knoblich, 2012; Shitamukai and Matsuzaki, 2012; Huttner and Kosodo, 2005). Still missing from this model is a mechanism that would cause both progeny of a dividing precursor to leave behind the self-renewing activity of apical Par proteins and undergo differentiation. Here, we propose that miRNA-mediated downregulation of Par proteins contributes to this transition.

In Drosophila, neuroblasts delaminate from the neuroectoderm, retaining an apico-basal polarity evident in asymmetric localization of Par proteins (reviewed in Homem and Knoblich, 2012). Apical Par proteins, through interaction with Inscutable, Gαi, Pins, and Mud, position the mitotic spindle so that neuroblast division occurs perpendicularly to the plane of the neuroectoderm (reviewed in Siller and Doe, 2009). Consequently, apical Par proteins are retained within one progeny cell, which continues as a neuroblast, and absent from the other, which undergoes differentiation, indicating that apical proteins are associated with self-renewal. Consistent with this, reduction of aPKC function resulted in failure of neuroblasts to undergo self-renewing divisions, whereas aPKC overexpression caused formation of excess neuroblasts (Lee et al., 2006; Haenfler et al., 2012). Neuroblasts exit the cell cycle prior to adulthood, under control of a temporally regulated sequence of transcription factors (Maurange et al., 2006), but whether...
apical Par proteins are downregulated to help promote this transition and the mechanisms by which this might occur remain unknown.

Although the correlation of spindle orientation, cleavage plane, and cell fate is not as obvious in the vertebrate CNS as in flies, several studies provided evidence that vertical cleavage planes that bisect apical membrane of neural precursors equally are generally associated with symmetric proliferative divisions, whereas oblique or horizontal cleavage planes that cause unequal segregation of apical membrane to progeny cells are asymmetric with respect to cell fate (Bultje et al., 2009; Kosodo et al., 2004; Haydar et al., 2003; Chenn and McConnell, 1995). Consistent with these observations, Pard3 can be asymmetrically localized to the progeny of dividing radial glial cells in mice, although the mechanism of unequal distribution was not clear (Bultje et al., 2009). Notably, whereas overexpression of Pard3 promoted symmetric divisions producing progenitor fate, Pard3 knockdown caused symmetric divisions producing neurons (Bultje et al., 2009). Similarly, Pard3 knockdown reduced proliferation of mouse cortical progenitors and favored formation of neurons, whereas Pard3 or Pard6x overexpression drove progenitor proliferation (Costa et al., 2008). These observations support the idea that, like invertebrates, apically localized Par complex proteins promote a self-renewing precursor state in the vertebrate nervous system.

Birthdating studies showed that the majority of dividing cells within the mouse cerebral cortex ventricular zone exit the cell cycle and differentiate by the end of embryogenesis (Takahashi et al., 1996). This cessation of proliferation correlates temporally with depletion of apical Par complex proteins from the ventricular surface (Costa et al., 2008). We showed here that the zebrafish spinal cord undergoes a similar transition. During early stages of neural development, a primitive lumen occupies nearly the entire dorsoventral extent of the neural tube. Apically associated proteins, including the Par complex protein Prkci, are localized to cell membranes lining the ventricular surface of the lumen. Cells that border the lumen, and thereby have apically concentrated Par proteins, have precursor characteristics in that they divide and express the precursor marker Sox2. At the end of the embryonic period, the lumen transforms to a small central canal in the ventral spinal cord. Concomitantly, the apparent concentration of apically associated proteins is reduced from all spinal cord cells except those bordering the central canal, and the number of dividing neural precursors is greatly diminished.

Our functional studies provide evidence that termination of neural precursor division and subsequent neuronal and glial differentiation at late embryonic stage is driven, in part, by negative regulation of Pard3 and Prkci by mir-219. We noted, though, that our functional manipulations affected primarily dorsal spinal cord. In fact, transitions in lumen morphogenesis, apical protein distribution, and neural precursor characteristics during development occur predominantly in dorsal spinal cord. In ventral spinal cord, apical proteins and Sox2+, putative neural precursors that also express olig2 (Park et al., 2007), remain localized to the central canal into postembryonic stage, raising the possibility that ventral neural precursors are not subject to mir-219 regulation. However, most spinal cord OPCs arise from ventral cells that express olig2. Nevertheless, prkci mutant zebrafish larvae had excess OPCs (Roberts and Appel, 2009) and mir-219 knock-down larvae had a deficit of OPCs (Zhao et al., 2010). One possible explanation is that OPCs originate dorsal to the central canal from precursors that are regulated by mir-219. More careful fate-mapping of OPCs origins should resolve this question.

Our data now extend current models in which, during early stages of neural development, mechanisms that influence whether the progeny cells of dividing neural precursors receive equal or unequal amounts of apical Par proteins influence whether divisions are symmetric and proliferative or asymmetric with respect to cell fate. We now propose that, near the end of embryogenesis, inhibition of new Par protein translation by mir-219 depletes Par proteins from dividing precursors below a limit necessary for their self-renewing functions, causing precursors to exit the cell cycle and differentiate as late born neurons and glia.

Previous work identified mir-219 as an important regulator of oligodendrocyte differentiation and myelination (Zhao et al., 2010; Dugas et al., 2010). Our study identifies an additional earlier and broader role for mir-219 in promoting the transition of neural precursors to specified cell types, including oligodendrocyte lineage cells. Additionally, because neural precursors continue to divide in the absence of mir-219 function, our data raise the possibility that alterations of mir-219 expression contribute to tumor formation. Consistent with this, mir-219 expression was significantly lower in childhood medulloblastoma relative to neural stem cells (Ferretti et al., 2009; Genovesi et al., 2011).

EXPERIMENTAL PROCEDURES

The animal work in this study was approved by the Institutional Animal Care and Use Committee the University of Colorado School of Medicine.

Zebrafish Lines

The experiments conducted in this study used the following strains of zebrafish: AB, Tg(olfi:EGFP)mi172 (Shin et al., 2003) and Tg(hsp70;pard3-EGFP)mi14.

Immunohistochemistry

Immunohistochemistry was performed on frozen sections as described previously (Park et al., 2003). F-actin was labeled using Rhodamine Phalloidin (Invitrogen). To detect EdU incorporation, we used the EdU Detection Reaction mix (Invitrogen). Images were collected on a Zeiss Axio Observer microscope equipped with a PerkinElmer UltraVIEW VoX spinning disc confocal system and Volocity imaging software (PerkinElmer).

In Situ RNA Hybridization

Antisense probes were transcribed using the Roche digoxigenin-labeling reagents and T3, T7, or SP6 RNA polymerases (New England Biolabs). To detect mir-219 expression, we used a dre-mir-219 miRCURY LNA probes (Exiqon 35172-01) (Kloosterman et al., 2006).

 Luciferase Assay

A 220 bp sequence containing the predicted target site was placed in the 3’ UTR of a renilla luciferase reporter gene using the commercially available plasmid, psiCHECK-2 (Promega). This construct and microRNA mimics were cotransfected into HEK293T cells. Luciferase activity was detected using the SpectraMax L Luminescence microplate reader (Molecular Devices).

Morpholino Injections

Antisense morpholino oligonucleotides were purchased from Gene Tools, LLC. We injected 1–2 nl of 0.25 mM morpholino in injection buffer into the yolk just below the single cell of fertilized embryos. All morpholino-oligonucleotide-injected embryos were raised in embryo medium at 28.5 °C.
BrdU and EdU Labeling
Dechorionated embryos were labeled with 20 μM 5-bromo-2′-deoxyuridine (BrdU) (Roche). For labeling with 5-ethyl-2′-deoxyuridine (EdU), embryos were incubated in 2 mM EdU (Click-IT EdU Alexa Fluor 555 detection kit, Invitrogen #c10338).

GFP Injections and Quantification
pard3 (1,200 bp) and prkci (800 bp) UTRs were cloned in an EGFP containing vector. These mRNAs were injected with or without miR-219 at one cell stage and raised to 30 hpf. Images of 20 embryos per group were collected using a Leica M165 FC microscope equipped with a SPOT RT3 camera and SPOT imaging software (Diagnostic Instruments). The experiment was performed three times and readings from 60 total embryos per group are reported.

Tg(hsp70:pard3-EGFP) Construction and Heat Shock Procedure
Tg(hsp70:pard3-EGFP) (53 hpf) and nontransgenic control embryos were placed in a 15 ml conical tube in approximately 10 ml of embryo medium and raised to 30 hpf. Images of 20 embryos per group were collected using a Leica M165 FC microscope equipped with a SPOT RT3 camera and SPOT imaging software (Diagnostic Instruments). The experiment was performed three times and readings from 60 total embryos per group are reported.

Quantitative PCR
RNA was isolated from 15–20 pooled larvae for each control or experimental condition. Samples for each condition were collected in triplicate. Real-time qPCR was performed in triplicate for each cDNA sample using an Applied Biosystems StepOne Plus machine and software version 2.1.

Quantification and Statistical Analysis
Cell counts were obtained by direct observation of sections using the microscopes described above. For Sox10, Sox2, PH-3, and Isl quantification, ten sections per embryo from 15 embryos per group with two or three replicates were counted to produce the average number per section. p values were generated using an unpaired t test using GraphPad Prism software. Dorsally migrated OPCs were assessed based on lateral views of Tg(olig2:EGFP) embryos at 3 dpf. olig2:EGFP positive cells were counted over the entire spinal cord. Larvae classified as normal had the number of dorsally migrated OPCs typical of wild-type. Larvae were classified as severe if fewer than five OPCs had migrated and mild in all other circumstances. p values were generated using an unpaired t test comparing the number of normal embryos when injected with miR-219 MO alone or miR-219 together with pard3 MO.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures and two figures and can be found with this article online at http://dx.doi.org/10.1016/j.devcel.2013.10.015.

AUTHOR CONTRIBUTIONS
L.I.H. performed all the experiments and acquired the data. A.J.B provided the transgenic line used to produce the data in Figure S2. L.I.H. and B.A. conceived the project, analyzed data, and wrote the paper.

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