The Ciliary Marginal Zone of the Zebrafish Retina: Clonal and Time-Lapse Analysis of a Continuously Growing Tissue

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Summary Statement: A quantitative study of cell proliferation and fate choice in a continuously growing neural tissue, including clonal and time-lapse analyses.
ABSTRACT

Clonal analysis is helping us understand the dynamics of cell replacement in homeostatic adult tissues (Simons and Clevers, 2011). Such an analysis, however, has not yet been achieved for continuously growing adult tissues, but is essential if we wish to understand the architecture of adult organs. The retinas of lower vertebrates grow throughout life, from retinal stem cells (RSCs) and retinal progenitor cells (RPCs) at the rim of the retina, called the ciliary marginal zone (CMZ). Here, we show that RSCs reside in a niche at the extreme periphery of the CMZ and divide asymmetrically along a radial (peripheral to central) axis, leaving one daughter in the peripheral niche and the other more central where it becomes an RPC. We also show that RPCs of the CMZ have clonal sizes and compositions that are statistically similar to progenitor cells of the embryonic retina and fit the same stochastic model of proliferation. These results link embryonic and postembryonic cell behavior, and help to explain the constancy of tissue architecture that has been generated over a lifetime.

KEY WORDS: stem cells, progenitor cells, retina, ciliary marginal zone, live imaging, clonal analysis, zebrafish.
INTRODUCTION

The retinas of fish and amphibians grow continuously throughout much of life by adding rings of new cells at the periphery, the so-called ciliary marginal zone (CMZ) (Johns, 1977; Straznicky and Gaze, 1971). The CMZ provides an excellent model for studying the proliferation in continuously growing tissues. In the past decades, the CMZ of lower vertebrates has been under investigation with respect to the genes, signaling pathways and transcriptional profiles that drive stem cells at the peripheral edge of the CMZ through proliferation and differentiation into the adult neurons that emerge at the central edge of the CMZ (Borday et al., 2012; Cerveny et al., 2012; Moshiri et al., 2004).

Early clonal analysis of the CMZ (Wetts et al., 1989) showed it to consist of two types of proliferative cells: those that give rise to large growing clones that stay in contact with the CMZ, and those that give rise to small differentiated clones that lose contact with the CMZ. The former, Wetts and colleagues assumed to be RSCs, and the latter RPCs. More recent work in medaka fish using indelible genetic markers shows that there are indeed dedicated RSCs in the CMZ, which give rise to arched continuous stripes (ArCoSs) containing all retinal cell types (Centanin et al., 2011). The fact that these ArCoSs are of constant width suggests that RSCs undergo invariant asymmetric divisions (Centanin et al., 2014).

Molecular studies reveal that cells from peripheral to central CMZ have different expression profiles of transcription factors, signaling molecules and cell cycle genes (Agathocleous and Harris, 2009; Cerveny et al., 2012; Harris and Perron, 1998; Johns, 1977; Raymond et al., 2006). Genes that are expressed at earlier developmental stages of retinogenesis are expressed in more peripheral regions of the CMZ while genes that are expressed later in development are expressed more centrally, suggesting that what happens in time through embryonic development is recapitulated spatially in the CMZ (Harris and Perron, 1998).
Analysis of markers combined with BrdU uptake and label retention show that the cells located at the extreme peripheral edge of the CMZ and presumed to be RSCs divide slowly, while cells in the central part of the CMZ divide more rapidly (Ohnuma et al., 1999; Perron et al., 1998; Xue and Harris, 2012).

To appreciate the relationship between embryonic and postembryonic development in quantitative detail, it makes sense to investigate the proliferative characteristics of CMZ cells in the light of what has recently been learned about retinal proliferation during embryogenesis. In zebrafish, the cells that give rise to the optic vesicle divide slowly between 14 and 22hpf. Then, a wave of rapid proliferation starting from the central region of the optic cup starts to spread outwards (He et al., 2012; Hu and Easter, 1999). By 72hpf this wave is at the very the edge of the retina. It leaves in its wake cells that have exited the cell cycle and differentiated into functional retinal neurons and glia. In front is the rim of undifferentiated and dividing cells at the periphery of the retina, which defines the CMZ.

Live imaging studies show that when embryonic retinal progenitor cells (RPCs) first arise in this wave, they initially divide a few times symmetrically and proliferatively (PP), they then enter a short stochastic period in which they undergo PP, PD (a mode in which one daughter differentiates (D) while other remains proliferative), or DD (a symmetric differentiative) division. This is followed by a final stochastic period in which the probability of a DD division becomes high and most cells quickly exit the cell cycle (He et al., 2012). These early RPCs thus produce clones of variable sizes with a mean of about 12 differentiated cells per initial RPC. This provides a simple quantitative framework for asking whether postembryonic RPCs of the CMZ behave in a statistically similar manner as embryonic RPCs.

The clonal analysis and imaging studies in this paper show that RSCs reside at the extreme peripheral edge of the CMZ, and that as the retina develops, they divide increasingly
asymmetrically in the radial orientation. This axis of cell division leaves one daughter in the stem cell niche where it remains an RSC and one that is pushed centrally, where it becomes an RPC. We also show that the fate behavior of RPCs of the CMZ is statistically similar to embryonic RPCs. These results provide an explanation for why the retinas of fish and amphibians can grow throughout a lifetime to produce a neural tissue with a similar cellular architecture, from the embryonically generated centre to the adult periphery.
MATERIALS AND METHODS

Animals and Transgenic Lines. Zebrafish were maintained and bred at 28.5 °C. Embryos were staged in hours post fertilization (hpf) before 72 hpf and in days post fertilization (dpf) afterwards. Embryos were treated with 0.003% phenylthiourea (PTU, Sigma) at 8 hpf to delay pigmentation, and anaesthetized with 0.035%-0.04% MS-222 (Sigma) prior to live imaging. The Tg(UAS:Kaede) (Scott and Baier, 2009), the MAZe (Collins et al., 2010) and the Tg(Rx2:GFPcaax) (Heermann et al., 2015) transgenic lines have been described previously. The double transgenic line Tg(MAZe,Rx2:GFPcaax) was generated by crossing the MAZe with the Tg(Rx2:GFPcaax) line. For photoconversion experiments, the Tg(MAZe,Rx2:GFPcaax) line was crossed with Tg(UAS:Kaede) line. The nlsRFP of the original MAZe line caused retinal cells to be sick. Therefore, as in Jie et al. (2012), we photoconverted Kaede expressing cells that showed no nlsRFP expression.

Heat shock and photoconversion. For clonal analysis, Tg(MAZe,Rx2:GFPcaax,UAS:Kaede) embryos were heat-shocked either at 10hpf or 54hpf for between 5-20 min at 37°C, and allowed to recover at 28.5°C. At 72hpf, the embryos were screened on a Nikon upright fluorescent microscope with a 40x water immersion objective for Rx2 positive retinas with Kaede-expressing cells. The selected embryos were then embedded in 3% methylcellulose and transferred to a spinning-disk microscope (Perkin Elmer) equipped for photoconversion coupled with an Olympus 40x water immersion objective (NA 1.3), where photoconversion was performed on green Kaede expressing cells. For short-term analysis, at 3dpf or 5dpf, single cells in the CMZ region (defined by Rx2 membrane marker) were randomly targeted by applying a 5s train of 405 nm laser pulses until the cell turns red. For long-term clonal analysis, the same photoconverting procedure was applied initially at 5dpf but on all the cells in a connected Kaede cell patch in the CMZ region. The photoconversion was then repeated.
on the converted cells that still remain in the CMZ at subsequent time points, such as 6, 8, 12, and 16dpf or until the death of the fish.

**Confocal image acquisition and analysis.** For short-term clonal analysis, 5dpf or 7dpf embryos were collected, anaesthetized and embedded in 3% methylcellulose with the proper orientation. For transverse sections, 5dpf fish retinas were fixed in 4%PFA, cryosectioned, stained with DAPI and mounted using Fluorsave (Calbiochem). Retina clones and sections were imaged using an Olympus 60x silicon oil objective (NA=1.35) on an inverted laser-scanning confocal microscope (Olympus FV1000), as the resolution of the image acquired on LSCM is higher than one from the photo-conversion equipped spinning disc. All the images were acquired using comparable settings (1024 x 1024 resolution, 10 µs/pixel scanning speed, 1-1.2 µm optical section). For long-term clonal analysis, the images were acquired using the spinning disk confocal microscope right after photoconversion (1000 x 1000 resolution, 250 ms/frame exposure time, 1 µm optical section). Image analysis was performed using Volocity software (Improvision).

**2-photon imaging acquisition and analysis.** 5dpf Tg(Rx2:GFPcaax) embryos were selected and embedded in 1% low melting agarose in a standard 90mm petri dish. Following the addition of the imaging medium (0.3x Danieau’s solution, 1x PTU, 0.8x MS222), the embedded embryos were placed in a 28.5°C heated chamber and imaged with a LaVision BioTec TriM ScopeII 2-photon microscope, using a 25x (NA 1.05) water immersion objective. The movies were acquired using comparable settings (0.75-1µm optical section, 2.5-3min interval, for 3-5h). The 3d+t data was analyzed using Fiji and Matlab software. The data analysis consists of the following three stages: detecting the position of the CMZ edge, identifying the divisions and the orientation of the division planes and finally, computing the relevant statistics for each division (angle of division and approximate distance from the CMZ edge). Detailed methods are described in Supplemental Methods.
**Numerical simulation of the RPC clone size.** The RPC clone size distribution predicted by the stochastic models in the text was based on the previously described simulation of clone size in the central retina (He et al., 2012). Two key points were inherited from the original model: 1) the cell cycles of the central retina progenitor cells follow a shifted gamma distribution, with a refractory period of 4hr, mean of 6hr, and width of 1hr; 2) the transition from symmetrical proliferation (PP) followed by asymmetrical (PD) and then terminal (DD) divisions were simplified to an internal clock starting from the first mitosis, with the probability initiated by pure PP division in the period 0-8 hr, to 20% PP, 40% PD and 40% DD during the period 8-15 hr, and then finally to 20% PP and 80% DD after 15 hr.

The model was adapted to simulate the RPC proliferation based on the assumptions discussed in the text. The mathematical simulation was implemented using Monte Carlo methods in a custom program written in MATLAB (MathWorks Inc).
RESULTS

CMZ consists of RSCs and RPCs with distinct potency. To resolve how individual cells in the CMZ behave in the postembryonic retina, we labeled single cells in 72hpf fish, which is when the central retina development is complete, and proliferation is restricted to the CMZ. We used the MAZe-Kaede system (He et al., 2012) in transgenic fish that carry the membrane-localized progenitor (and Müller Glial) marker Rx2:GFPcaax, which was functionally characterized as labeling the CMZ cell population including neural stem cells (Chuang and Raymond, 2001; Heermann et al., 2015; Reinhardt et al., 2015) (Figure 1A,B). At 54 hpf, a defined heat shock was applied to trigger Cre-mediated recombination in a random subset of cells, driving the expression of the transcriptional activator Gal4:VP16, which then activates the expression of the cytoplasmic photoconvertible protein Kaede (Ando et al., 2002) from an upstream activating sequence (UAS) promoter (Collins et al., 2010). At this stage, single GFP labeled cells in the CMZ were randomly selected for photoconversion to red fluorescence. We found that red fluorescence after photoconversion provided a durable signal for lineage tracing for up to 5dpf (Figure 1E).

This assay revealed two populations of CMZ cells with surprisingly distinct proliferation/differentiation behaviors. 88% (104/118) of the CMZ cells that were photoconverted at 72hpf produced fully differentiated clones consisting of retinal neurons or glia cells by 120hpf (Figure 1D, E top panel). By contrast, 9% (11/118) of labeled cells generated clones in which all the progenitor cells remained in the CMZ (Figure 1D, E bottom panel), arranged in a stripe-like structure on the eye. The remaining 3% (3/118) of clones had a mixture of cells that remained in the CMZ and cells that had differentiated and left the CMZ. This bimodality of proliferative characteristics in the CMZ is similar, although not numerically comparable, to what was previously shown in the CMZ for a different species using a different methodology (Wetts et al., 1989), and suggests that we are looking at RSCs and RPCs.
RSCs reside in the most peripheral edge of the CMZ. The low percentage of RSC derived clones together with the lack of knowledge of the exact location within the CMZ of the photoconverted cells in the above experiment led us to adapt the MAZe-Kaede system further so that we could study the RSCs with more precision (Figure 2A). In order to label more RSCs, we heat shocked the fish at 54 hpf, but instead of photoconverting at 3 dpf, we left the retina growing for 2 more days and applied the initial photoconversion at 5 dpf. As the vast majority of RPC clones induced at 3dpf terminate within 2 days, we reasoned that the clones that remained in the CMZ at 5 dpf were likely to contain at least one stem cell. We then photoconverted all the Kaede-positive cells in a connected cell patch in the CMZ at 5 dpf, in effect labeling a polyclone that consists of multiple Rx2 positive cells in the CMZ. We then imaged the photoconverted polyclones over an extended period of time (up to 16dpf, see Methods).

Among the 50 polyclones that were followed, 31 continued to retain cells in the CMZ by 16dpf (n=14) or up to the time of death of the fish if earlier (n=17, 10 of which died on 12dpf, and 7 died on 16 dpf), while 19 clones became fully differentiated during postembryonic retinal development (Figure S1). Although it is possible that a maintained polyclone is not equivalent to a polyclone that contains an RSC, to find which cells in our polyclones were most likely to be the RSCs, we looked at the spatial distribution of the photoconverted cells. In a 3D image of the CMZ clone, the stripe-like structure was rotated to an orientation where the lens was facing the viewer, and the surface of the eye at the position of the clone stripe was parallel to the viewing plane (Figure 2C). If a cell in the clone was within the annulus or ring of CMZ cells that were closest to the lens, it was defined to be in “position 1” or “ring 1” (see also Figure 1B). The cells that were one cell diameter further away from the lens were deemed to be in position or ring 2, etc. Thus the lower the position or ring number of a cell, the more peripheral that cell is in the CMZ. This provided a metric of the spatial distribution
of cells within the clones with respect to a central-peripheral (radial) axis in the CMZ at any observed time point. This is shown for a single maintained and a single terminated clone in Figure 2B and C respectively. (See Figure S1 for all polyclones.) Taking into account the spatial arrangement of cells in a clone, it became immediately evident that the maintained clones have their most peripheral cells at position 1 or 2, while the terminated clones detach from these most peripheral rings before termination (Figure 2D). This positional distribution of maintained and terminated clones is not only observed in the long-term polyclones, but also in single clones traced from 5-7 dpf in older larvae (Figure S2). Thus, we conclude that rings 1 and 2 of the CMZ contain most or all or the RSCs, while higher rings contain the RPCs.

**RSC divisions become increasingly asymmetric.** It has been proposed that each RSC in the CMZ of larval medaka fish divides asymmetrically to produce 1 RSC and 1 RPC to generate an ArCoS that stays in touch with the CMZ (Centanin et al., 2011). Such asymmetric divisions would also assure that the CMZ maintains a constant population of RSCs and provides a natural explanation for the relatively constant width of ArCoSs (Centanin et al., 2014). However, because the space at the extreme edge of the CMZ is strictly limited, it is possible that at early stages of development, there is competition for this niche and that in some cases both daughters of an RSC may end up within the niche while, in others, neither daughter will. As our protocol is designed to label RSCs in polyclones, the fact that a third of the polyclones become detached from the putative niche region suggests that, though there may be a strong bias toward asymmetry, not all RSC divisions are asymmetric at 5 dpf. However, by following polyclones further in time, we find that the rate of clonal detachment drops significantly and that by 8 dpf, no further detachment of clones from rings 1-2 was observed (Figure 3A). The higher detachment rate at early time points could be due to the fact that the CMZ has not been fully stabilized and/or that the polyclones that do not contain
a RSC detach by 8dpf. The fact that the detachment rate drops to near zero indicated that the
RSCs in the stem cell niche at the extreme periphery of the CMZ divide in a radial orientation
that leads to asymmetric fate outcomes, with one cell always remaining in the ring 1-2. A
prediction of this asymmetry, and of the radial geometry seen here and in ArCoS clones, is
that the number of RSCs per clone should remain constant (Centanin et al., 2014). Indeed, as
all the RSCs seem to be in ring 1 and 2, when we looked at the number of cells in rings 1-2 of
maintained clones, we found that this was indeed the case (Figure 3B). Figure 3C shows a
simple model of RSC and RPC divisions that are compatible with these data.

**Orientations of RSC and RPC cell divisions in the CMZ.** If RSCs of the zebrafish CMZ
divide asymmetrically with one daughter remaining as an RSC in the stem cell niche and the
other daughter leaving the niche, one could expect that cell divisions of RSCs should be
oriented along the radial axis. In this case, we should see evidence of oriented cell divisions
in the most peripheral part of the CMZ. To investigate the distribution of cell division
orientations within the CMZ, we used 2-photon confocal microscopy over several hours
using the Tg(Rx2:caaxGFP) line (Figure 4A). We captured more than 150 cell divisions in
two 4-dimensional time-lapse movies of 5dpf larvae (Figure 4B,C). We noted that, as in the
embryonic retina (Das et al., 2003), the vast majority of these divisions take place at the
apical surface, and are parallel to this surface (see also Figure 1B and S3), i.e. both daughter
cells remain in contact with the apical surface. Although there may be small deviations along
the apicobasal axis, we focused on the orientation of cell division as a two-dimensional issue;
i.e., within the plane of the apical surface of the CMZ. Here, the orientation of cell division
can vary from radial, which is defined as orthogonal to the local tangent at the circumference
of the CMZ, to circumferential, which is parallel to this tangent (Figure 4D). For each
observed cell division, we then calculated the distance to the edge of the CMZ and the
orientation of the division (Figure 4E,F and Supplemental Methods). Significantly, we found
that the most peripheral cells (row 1) divide predominantly radially, while divisions of more central CMZ cells are unconstrained in their orientation (Figure 4E). This significant bias of RSCs to divide radially would tend to keep one daughter close to the peripheral edge, next to the ring of blood vessels that is forming at this time (Kitambi et al., 2009) (Figure 4B), where it could remain in a RSC niche and push the other daughter out of the niche and towards a more central RPC fate.

**RPCs of the CMZ recapitulate the developmental history of embryonic RPCs.** Our recent work in the zebrafish suggests that embryonic RPCs are equipotent in terms of their proliferative and fate potential and that they behave stochastically according to highly constrained intrinsic probabilities (He et al., 2012). The distributions of clone size and fate compositions of embryonic RPCs are characteristics of these probabilities. Obvious questions therefore arise about the RPCs that are generated in the CMZ. Do these postembryonic RPCs follow the same statistical rules as the embryonic RPCs? To answer these questions, we looked in detail at clones generated by RPCs. By the end of the short-term clonal analysis at 5 dpf, we identified the 4 basic neuronal types – retinal ganglion cells (RGCs), amacrine cell (ACs), bipolar cell (BPs), and photoreceptors (PRs) – in all of the photoconverted clones, and compared them to the lineages reconstructed from live imaging in the central retina during embryonic development (He et al., 2012) (Figure 5A,B and Figure S4). There is indeed a striking similarity to the cell-type distributions in the clones generated by embryonic and CMZ generated RPCs. In the CMZ, 2-cell clones from a terminal division have strong preference to specific fate combinations: the clones composed of PR, BC and AC pairs are overrepresented, which is consistent with the pattern observed in the embryonic retina (Figure 5A). Similarly, the cell fate combinations in 3-cell clones is also in agreement, with RGC+2PRs being the prominent type. In the central retina, 93% of the 3-cell clones with 2 PRs contain either and AC or an RGC, while the ratio is 94% in the CMZ. Furthermore, the
percentage of ACs, BC, HCs and PRs in the clones larger than 5 cells is similar to those originating from 24hpf progenitors in the central retina (Figure 5B). We also photoconverted single RPCs that gave rise to terminated clones at 5dpf, and analyzed these clones at 7 dpf. Although this data set was more difficult to obtain for technical reasons, these 5dpf RPC clones also show similar compositions to the embryonic clones assessed by He et al., (2012) and of course the 3dpf RPC clones (Table S1).

**Proportions and Clone Sizes.** We next focused on the clone size of the RPC lineages. A prototypical clone of 12 cells would be generated by 4 rounds of mitoses (Figure 5D): two rounds of PP divisions that turn the initial RPC (I₀) into 2 primary progenitors (I₁) and then 4 secondary progenitors (I₂), followed by one round of a mixture of PP, PD and DD divisions that gives 4 tertiary progenitors (I₃) and 4 differentiated cells, and finally the round of DD divisions that terminate the clone (He et al., 2012). In reality, because of the stochastic nature of some of these division modes, few if any clones follow this exact scheme. Yet such a scheme, representing average proliferative behavior, is useful for approximating the distribution of different staged RPCs in the CMZ. As we excluded the single clones in the analysis and only focused on the behavior of RPCs with proliferation potential, we additionally assumed that RPCs can reside in the CMZ for a limited time (17h in the case of this model based on the 3 dpf data) after their first division (Figure 5C) before being pushed into the central retina and forced to differentiate. If we initialize the RPC pool in the CMZ by a mixture of many cell lineages originated from equipotent I₀s (like the typical lineage in Figure 5D), randomly select one of the descendent RPCs in the lineage, and allow it to fully differentiate following the probabilistic model in Figure 5C, the experimental clone size distribution of the RPC cells stays well within 95% confidence interval of the predicted distribution from the model (Figure 5E).
In the 24hpf embryonic retina, a prototypical I₀ RPC generates a clone of 12 neurons, the outcome of 11 divisions. In the CMZ, we have shown that an RSC typically makes an asymmetrical division to give rise one copy of itself and one early (I₀) RPC. Thus, if these early RPCs of the CMZ are equivalent to embryonic RPCs in terms of their proliferative potential, we might expect a ratio of 1 RSC division to 11 RPC divisions. How does this prediction match the assumption that RSCs reside in ring 1 and RPCs reside in the more central rings? We found that 12 cells in ring 1 went through cytokinesis in the same time interval that 153 cells in ring 2 and higher did, yielding a ratio of 1:13. This is broadly consistent with the prediction and might be even closer if, as seems likely from our clonal imaging analyses, a small fraction of cells in ring 2 were RSCs. It is also clear from these time-lapse recordings that the proportion of cells that can be seen to go through cytokinesis during the defined period of these movies depends on the distance from the periphery. The proliferation rate is low in the most peripheral annulus corresponding to ring 1, and it then increases in the middle of the CMZ and decreases again toward the central CMZ (Figure 4F). This agrees qualitatively with our hypothesis that, from peripheral to central, the CMZ consists of RSCs that divide slowly, young RPCs that divide quickly, and finally old RPCs that are leaving the cell cycle and beginning differentiate. This too, is in accordance with what has been found within the embryonic retina, where RSCs divide slowly (Hu and Easter, 1999), early RPCs divide symmetrically in a PP mode, and late RPCs slow down their cell cycles and differentiate (He et al., 2012).
DISCUSSION

Using quantitative clonal strategies, we confirmed that RSCs are located at the peripheral extremity of the CMZ, which functions as a stem cell niche (Raymond et al., 2006; Xue and Harris, 2012), where they divide mainly in the radial orientation promoting asymmetric fate outcome ensuring a constant production of RPCs. We also show that the same simple model (using the same parameters) that was shown to describe the clonal evolution in embryonic RPCs (He et al., 2012) fits the proliferative statistics of RPCs in the CMZ suggesting the functional equivalency of the embryonic and CMZ progenitor cells.

The location of RSCs at the extreme periphery of the CMZ is consistent with previous studies based on molecular markers and label retention (Ahmad et al., 2004; Johns, 1977; Ohnuma et al., 2002). The position of RSCs in the CMZ, with access to the apical epithelial surface and close association with the ring blood vessel, is a feature that is common to other neurogenic niches in the vertebrate brain (Raymond et al., 2006), and supports the idea that there is a growth factor dependent stem cell niche at the peripheral edge of CMZ (Xue and Harris, 2012). This study also demonstrated that RSCs divide in a persistently asymmetrical manner to produce one RSC and one early stage RPC, as was strongly suggested by the constant thickness of ArCoSs (Centanin et al., 2014; Centanin et al., 2011).

The multipotency of clones derived from RPCs in the *Xenopus* CMZ (Wetts et al., 1989) suggested that adult RPCs and embryonic RPCs share some fundamental properties. This notion was reinforced by later studies, using a variety of differentiation and cell cycle markers, showing that the CMZ spatially recapitulates, from peripheral to central, the temporal progression of embryonic retinal development (Johns, 1977; Ohnuma et al., 2002; Raymond et al., 2006). Here, we show that CMZ derived RPCs are not significantly different in terms of their statistical proliferation patterns to embryonic RPCs, suggesting that they are functionally equivalent cell types, which helps to explain the constancy of retinal tissue
architecture in zebrafish from centre to periphery. We did not see any Müller Glia in our 3-5dpf terminated clones. This is not unexpected due to the low percentage of Müller Glia in the retina and our small sample size, yet it raises the question of whether the central Müller Glia contribute to the cellular architecture of the peripheral retina or whether it all arises from the CMZ. Although our work here does not address this question, Centanin et al. (2011) showed that the ArCoS clones contain all retinal neurons and Müller Glia, and thickly label all cells within their width, suggesting that the cellular architecture of the retina arises from clones that originate in the CMZ. Our paper builds on their work by showing that RPCs share the same proliferative potential and fate behavior of embryonic RPCs, which offers a quantitative explanation for the homogeneity of retinal architecture.

The key difference between the embryonic generation of the central retina and the postembryonic generation of the peripheral retina, which continues throughout much of life in frogs and fish, is that the latter is fueled by a population of self-renewing RSCs in the CMZ. During the early formation of the optic vesicle in zebrafish, the cell cycle is very slow and then, at about 24hpf, a wave of proliferation spreads from the centre of the retina reaching the periphery by 72hpf (He et al., 2012). The peripheral rim that remains proliferative is the initial CMZ and at its extreme periphery is the stem cell niche.

In many homeostatic adult epithelial tissues, stem cells can frequently commit to terminal differentiation while the loss of these stem cells is compensated by the multiplication of neighboring stem cells (Simons and Clevers, 2011). In such homeostatic self-renewing tissues, where stem cell duplication happens with the same probability as termination, the tissue is eventually taken over by clones that dominate through neutral competition (Vogel et al., 1969). In contrast to such scenarios, indelible genetic markers used for the long-term tracking of clones originating in the CMZ of medaka fish (Centanin et al., 2011) show that
retinal clones derived from stem cells do not take over, but rather form long thin ArCoSs, comprised of all types of retinal cells that stretch from the central retina to the still growing CMZ. The fact that such ArCoSs rarely terminate and rarely gain width strongly suggests the absence of such neutral competition and suggests that the RSCs generating these clones divide strictly asymmetrically (Centanin et al., 2014). Our polyclonal analysis at a cellular level of resolution strongly supports these observations that RSC division is asymmetric in terms of fate. We also find that these asymmetric divisions tend to be radially oriented. One unifying explanation for these two observations is that RSC competence is ensured by factors located at the extreme edge of the CMZ, near the ring blood vessel that lies between the lens and the retina (Kitambi et al., 2009). Clone terminations were observed in our young but not older fish, suggesting that the CMZ is stabilized during the first few days of postembryonic development.

Asymmetric divisions along particular axes have been seen to be important during the development of both plants and animals. For example, in the developing nervous system, asymmetric divisions along the apicobasal axis, are thought to lead to the unequal inheritance of intrinsic determinants such as Numb (Roegiers and Jan, 2004). In many systems, however, the oriented division of stem cells can lead to one cell remaining with a niche that provides short-range signals, such as growth factors, which maintains the “stemness” of the cells that remain within the niche (Januschke and Nathke, 2014). This seems to be the case in the CMZ of the zebrafish, where the critical orientation is not apicobasal but radial, which helps ensure that one daughter cell remains at the extreme periphery of the retina, while the other is forced out. This raises many interesting questions. For example, do these radial divisions lead to the asymmetric segregation of intrinsic factors that lead to differential fates, or is it just that one daughter remains in the stem cell niche while the other moves out? If the latter is the case, one would also like to know how this orientation is achieved, and whether the blood
vessel is an important component of the niche and whether the same signals that regulate proliferation in adult mammalian stem cell niches (Li and Clevers, 2010) are used in this system. These questions must await further investigation of this fascinating proliferative neural tissue.

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COMPETING INTERESTS

The authors declare no competing or financial interests.

AUTHOR CONTRIBUTIONS

Y.Wan did some of the experimental work, wrote the code for theoretical modeling and simulation and helped write the manuscript. A.D.A., N.C., Y.Wu, and J.H. all did experimental work. SR did the statistics, theory and helped with the modeling, LM did the image analysis of the 4D movie sets and the graphical work associated with that. BDS provided theoretical framework and W.A.H. supervised the work and helped to write the manuscript.

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SUPPLEMENTARY MATERIAL

Supplementary material available online at http://dev.biologists.org/lookup/suppl/
Figures
Figure 1 | Short-term clonal analysis. Rx2 functions as a marker for the CMZ of the zebrafish retina. (A) A maximum intensity projection of a transverse 12µm section of a 5dpf zebrafish retina, showing the expression of the transgenic reporter Tg(Rx2:GFPcaax). Rx2 positive cells (green) are found in the CMZ as well as in two cell types of the neural retina, photoreceptors (PR) and Muller glia (MG). (B) High magnification (2x) 1µm section from a region of the CMZ (dotted line in A), revealing that both RSCs (row 1-2) and RPCs (rows 3-6) are Rx2 positive. Numbers indicate rings. Nuclei are labeled with DAPI (magenta). ONL, outer nuclear layer; INL, inner nuclear layer; RGC, retinal ganglion cell layer. (C) Experimental flow of labeling CMZ cells and capturing the short-term clonal composition. (D) Composition of short-term CMZ clones reveals the distinct proliferative fates in CMZ population. (E) In these sections all the Rx2 positive cells have green membranes while the MAZe:Kaede cells have green cytoplasms that become red upon photoconversion. (Top panel) Single CMZ cell is photoconverted to red at 3dpf (left, indicated by arrow) and the resultant terminated clone at 5dpf (right, red) is shown with identified cell fates (RGC, retinal ganglion cell; PR, photoreceptors). (Bottom panel) A single CMZ cell photoconverted at 3dpf (left, indicated by arrow) and the resultant maintained clone observed at 5dpf (right, red). Note the dilution of the label in the cells further from the lens suggestive of more rapid division. Scale bars for A, B and E (10µm).
Figure 2 | Long-term polyclonal analysis. (A) Experimental flow of labeling 5 dpf CMZ clones and following the maintaining CMZ polyclones at subsequent time points. (B) Colors are the same as in Fig. 1E. Photoconverted CMZ cells (red) at 5, 6, 8, 12 and 16dpf, with the upper rows showing a maintained polyclone and lower rows a terminated polyclone at 16dpf. Scale bar: 20 µm. The final panels in each series show the clonal geometry of each of these two clones. Black circles indicate ring 1, grey = ring 2, dark green to light green = ring 3 to ring 8. (C) Frontal view of an Rx2 membrane-marked retina, with the cell ring position coded in colors (as above). (D) The distribution of the position of the most peripheral cells in clones at the last observed day of all the maintained clones (n=31) and the corresponding value for all terminated clones before termination (n=19). Error bar represents one sigma confidence interval of Poisson distribution.
Figure 3 | Stabilization of CMZ clones is due to asymmetrical SC division. (A) The rate of clone detachment events is plotted as a function of age. Detachment rate is defined as the number of detachment cases (clones move away from ring 1-2 in a specific period of time) divided by the total cases survived up to that time point, and then divided by the number of days in the time period. (B) Plot showing the average number of Ring 1 plus Ring 2 cells in the maintained clones as a function of age. Note that this number remains relatively constant in the maintained clones, indicating potential asymmetrical division of RSCs. (C) Model of SC and RPC division, with $\alpha$ and $\beta$ indicating the division rate of SC and RPC respectively.
Figure 4 | Live imaging of cell divisions in the CMZ. (A) Schematic illustration of the 2-photon imaging set up. (B) 3D projection of one movie frame (t=69, z=16.5µm), showing the CMZ region imaged. Arrows indicate areas of retinal pigment epithelium (RPE). Yellow rectangle represents the area shown in the following panels. The ring blood vessel is shaded light green (C) Selected frames of a 4D time-lapse movie showing several cell divisions in a variety of orientation planes. Dividing cells are labeled with a colored dot. A presumptive stem cell, dividing parallel to the edge of the retina close to the blood vessel, is labeled in red. Each frame represents a maximum intensity projection (6 frames, z=4.5µm). Elapsed time shown in min:sec. Dotted line indicates the edge of the blood vessel. Scale bar:10µm. (D) Computation of division angle. The 3D image stack is segmented into retina (blue) and blood vessel (green). Inner panel: The pixels belonging to the divisions are detected by analysing the difference between two consecutive or temporally close frames (depending on the time resolution of the imaging) and finding large contiguous volumes of statistically significant
pixels in the difference image, followed by a manual validation step (see Supplemental Methods). Once the divisions are detected, the normal of the division plane (yellow) is estimated and projected in the CMZ plane and the angle between the projected normal and the tangent to the CMZ circle at the closest point is computed. (E) Box-plot of division angles detected in each ring of cells in the CMZ. Ring 1 cells divide more strictly axially compared to the random distribution in higher rings. Boxes span from the first to the third quartile. Whiskers extend from the lowest value still within 1.5 interquartile range (IQR) of the lower quartile to the highest value still within 1.5 IQR of the upper quartile. (F) Estimated division rate in each ring of cells in the CMZ. Division rate is defined as the number of divisions observed divided by the number of cells imaged in the corresponding ring. Error bar represents one sigma confidence interval of Poisson distribution.
**Figure 5** | RPCs recapitulate the developmental history of central retina progenitors. (A) The frequency of fate combinations of sister cells in 2-cell clones is comparable in the CMZ and the central retina. Color in the matrix is normalized to the ratio of dominant fate combinations. Brightest red color indicates the most common composition, and black means the fate combination does not exist. (B) Clonal composition of large CMZ clones (≥5 cells) resembles the composition of central retina clones induced at 24hpf. None of the cell types except for RGCs show significant difference in the percentage of the clone (α=5%). (C) Schematic showing the theoretical model used to simulate the RPC clone size. Time-dependent mitosis and division mode probability in central retina development are plotted (adapted from He et al, 2012), with the CMZ progenitors randomly selected from the pool of progenitors in the first 17 hours within the lineage, indicated in the purple box. (D) An example of the RPC clone in the CMZ recapitulating the age distribution within a prototypical cell lineage in a central retina progenitor. I₀: initial progenitor, I₁: primary progenitor, I₂: secondary progenitor, I₃: tertiary progenitor, residing in the CMZ region (green).
and D: differentiated cells residing in the central retina (blue). (E) Fit between model predictions (red line with shaded pink region showing 95% plausible intervals due to finite sampling) and size distribution (blue crosses) of RPC clones. In this figure the right panel of A, the red bars in B and the model shown in C have all been extracted from (He et al., 2012).
REFERENCES


