Development and postnatal neurogenesis in the retina: a comparison between altricial and precocial bird species

Introducción

Los pájaros se clasifican en un arco prerreconocimiento-prerral a traves de la descripción de comportamiento y morfológico del desarrollo de las camadas. En el momento de la camada, los libreros de camada están cubiertos con plumas que se reemplazan por plumas de adulto. También tienen ojos abiertos y un sistema muscular y esquelético desarrollado (Figura 1A). En contraste, los camados prerreconocimiento tienen poco o ningún plumaje en su piel, son ciegos, tienen menos sistema locomotor y están altamente dependientes de la atención parental (Nice, 1962; Figura 1B).

Estudios embrionarios también han confirmado una gama de diferencias de desarrollo entre los prerreconocimiento y los camados prerreconocimiento (Murray et al., 2013; de Almeida et al., 2015; Álvarez-Hernán et al., 2013). Además, varios estudios maturation features occur at later stages in the altricial birds than in the precocial birds (Murray et al., 2013; de Almeida et al., 2015; Álvarez-Hernán et al., 2018; 2020), and, while different tissues mature at later stages of development in altricial embryos, this period is primarily characterized by growth in precocial birds (Hamburger and Hamilton, 1951; Murray et al., 2013).

Precocial birds also differ from altricial birds in the timing of brain structures differentiation (Charvet and Striedter, 2011). Thus, several brain regions in altricial birds are relatively immature at hatching (Striedter and Charvet, 2008). Therefore, altricial birds exhibit a great post-hatching brain growth associated with intense neurogenesis in certain brain regions such as the telencephalon (DeWulf and Bottjer, 2005; Striedter and Charvet, 2008; Charvet and Striedter, 2009, 2011). Neurogenesis is intense in T. guttata chicks during the first week of life, but this process persists even into adulthood (DeWulf and Bottjer, 2005; Charvet and Striedter, 2009).

Abstract

The visual system is affected by neurodegenerative diseases caused by the degeneration of specific retinal neurons, the leading cause of irreversible blindness in humans. Throughout vertebrate phylogeny, the retina has two kinds of specialized niches of constitutive neurogenesis: the retinal progenitors located in the circumferential marginal zone and Müller glia. The proliferative activity in the retinal progenitors located in the circumferential marginal zone in precocial birds such as the chicken, the commonest bird model used in developmental and regenerative studies, is very low. This region adds only a few retinal cells to the peripheral edge of the retina during several months after hatching, but does not seem to be involved in retinal regeneration. Müller cells in the chicken retina are not proliferative under physiological conditions, but after acute damage some of them undergo a reprogramming event, dedifferentiating into retinal stem cells and generating new retinal neurons. Therefore, regenerative response after injury occurs with low efficiency in the precocial avian retina. In contrast, it has recently been shown that neurogenesis is intense in the retina of altricial birds at hatching. In particular, abundant proliferative activity is detected both in the circumferential marginal zone and in the outer half of the inner nuclear layer. Therefore, stem cell niches are very active in the retina of altricial birds. Although more extensive research is needed to assess the potential of proliferating cells in the adult retina of altricial birds, it emerges as an attractive model for studying different aspects of neurogenesis and neural regeneration in vertebrates.

Key Words: altricial; birds; circumferential marginal zone; Müller glia; postnatal neurogenesis; precocial regeneration; retinogenesis

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Those authors conclude that postnatal maturation of brain structures facilitates learning in altricial birds. Therefore, the brain of altricial birds emerged as an influential model for the study of adult neurogenesis.

The visual system is a powerful model for advancing in knowledge about ontogenetic differences among vertebrates. Visual system morphogenesis and retinogenesis are well conserved throughout the vertebrate phylogeny (see below). However, while most of these developmental features occur during the embryonic period in precocial birds, some of these species complete retinal development during postnatal life. In precocial Chondrichthyes (Bejarno-Escobar et al., 2013; Sánchez-Farias and Candral, 2015, 2016), osteichthyes (Candal et al., 2005, 2008; Álvarez-Hernán et al., 2019), reptiles (Francisco-Morcillo et al., 2006), and mammals (Loeliger and Rees, 2005; Gudurich-Fuchs et al., 2009), retinogenesis is completed during embryological stages. In contrast, in altricial fish (Doldán et al., 1999; Pavón-Muñoz et al., 2016; Álvarez-Hernán et al., 2019) and mammals (Rapaport et al., 2004; Bejarno-Escobar et al., 2011), retinal development occurs postnatally.

Regarding avian retinal development, it is well known that events such as cell differentiation (Prada et al., 1991; Martín-Partido and Francisco-Morcillo, 2015; de Mera Rodríguez et al., 2019) and cell death (Cook et al., 1998; Marin-Teva et al., 1999; Francisco-Morcillo et al., 2014) within the precocial retina occur embryonically. At birth, the layers of the retina of precocial birds are clearly distinguished and show differentiated cells (Prada et al., 1991; Rojas et al., 2007; de Mera-Rodríguez et al., 2019). Neurogenesis is almost absent in the chicken retina, and is restricted to a few cells located in the circumferential marginal zone (CMZ) (Fischer and Reh, 2000). In contrast, many features of immaturity have been described in the retina of newly hatched altricial birds (Bagnoli et al., 1985; Rojas et al., 2007; Álvarez-Hernán et al., 2018, 2020). Recent studies conducted in our laboratory have shown that neurogenesis is intense in the retina of newly hatched altricial birds (zebra finch, Taeniopygia guttata Vieilliot 1817) (Álvarez-Hernán et al., 2018, 2020).

In this review, we shall describe the variation in the developmental timetable of visual system ontogeny from the altricial/precocial perspective. We shall also highlight the remarkable variation in proliferative activity of avian cell progenitors in the retina of altricial and precocial birds at hatching and during the first postnatal days.

Search Strategy and Selection Criteria

Studies cited in this review were found on the PubMed database, between December 2019 and February 2020, using the search terms: adult neurogenesis, retinogenesis, visual system development, altricial birds, precocial birds, retinal progenitors, retinal regeneration, altricial development, precocial development, and various combinations of the above terms.

Morphogenesis, Histogenesis, and Cell Differentiation in the Vertebrate Retina

The vertebrate retina is formed embryonically from the brain and plays a key role in vision. This sensory tissue consists of several layers of neurons with two neuropils where synaptic contacts occur. There are three layers of cell somata which contain the cell nuclei: outer nuclear layer (ONL), inner nuclear layer, and ganglion cell layer; there are also two layers that contain synaptic connections between axons and dendrites: outer plexiform layer and inner plexiform
one of seven classes of retinal cells. This process of cell determination and differentiation is termed retinogenesis (Xiang, 2013; Amini et al., 2018) (Figure 1C).

Timing Differences during Visual System Development in Altricial and Precocial Birds

Many of the developmental events that occur during visual system embodiment in birds begin earlier in the retina of precocial species than in altricial species. Thus, the optic vesicle and optic cup formation and the emergence of the plexiform layers occur at later stages in the *T. guttata* retina (Álvarez-Hernán et al., 2018) than in the chicken (Hamburger and Hamilton, 1951). Furthermore, the first differentiating ganglion cells are detected in the *G. gallus* retina at HH13 (48–52 hours of incubation) (Prada et al., 1991; Snow and Robson, 1994), while in the *T. guttata* retina, the first ganglion cells become post-mitotic at St24 (108 hours) (Álvarez-Hernán et al., 2020). The onset of visinin expression, an early marker of photoreceptor differentiation, in the chicken OLN occurs at HH27 (120 hours) (Bruhn and Cepko, 1996; Bradford et al., 2005), while in the *T. guttata* retina it is first detected at St28 (132 hours) (Álvarez-Hernán et al., 2020). Finally, the onset of expression of Prox1, a transcription factor that is expressed during early horizontal cell differentiation, is detected in the *T. guttata* retina at St34 (172 hours) (Álvarez-Hernán et al., 2020), while the first Prox1-immunoreactive progenitors in the chicken retina are detected at HH30 (156 hours) (Edqvist and Hallböök, 2004). Thus, morphogenesis, histogenesis, and cell differentiation in the developing *T. guttata* retina are delayed with respect to *G. gallus*, a precocial bird species.

Retinal Maturity at Hatching

Precocial birds hatch with a fully differentiated and functional retina (Fischer and Reh, 2000; Kubota et al., 2002). Mitotic figures, detected with antibodies against phospho-histone H3, were absent in the central retina of newly hatched individuals in the chicken (Figure 2A) and in the quail (Figure 2C). However, sparse phospho-histone H3-immunoreactive mitotic figures are found in the CMZ in both precocial species (Figure 2B and D). This proliferative activity in the peripheral-most region of the retina in hatchlings and adults of these birds suggests ongoing growth of the retina during postnatal life (Fischer et al., 2014). These proliferative cells also express retinal progenitor markers. Intra-ocular injections of different growth factors can greatly enhance the proliferative activity in the CMZ (Fischer and Reh, 2003). Furthermore, a sub-population of cells located in a specific region of the ciliary body also have neurogenic potential and can be stimulated by several growth factors to proliferate and differentiate into retinal neurons (Fischer and Reh, 2003).

 Müller glia plays a essential role in the maintenance of the retinal structure and participates in essential homeostatic processes in the developing and mature retinal tissue (Bejarano-Escobar et al., 2017). These cells have a significant proliferative and neurogenic capacity under physiological (Julian et al., 1998; Pavón-Muñoz et al., 2016) and experimental conditions (Bernardos et al., 2007) in the fish retina. However, Müller glia are not proliferative in the healthy bird retina at hatching. Avian Müller cells can exhibit neurogenic properties under the effect of acute damage or by the addition of specific growth factors in the uninjured chicken retina (Fischer and Reh, 2000). These stimuli induce proliferation of Müller cells and re-initiation of the expression of retinal progenitor markers (Fischer and Reh, 2000). Therefore, Müller glia need to be stimulated to acquire the proliferative phenotype in precocial birds.

Figure 2 | Immunodetection of pH3H3 in *G. gallus* (A, B) and *C. coturnix* (C, D) in the retina of newly hatched animals.

pH3H3 immunoreactivity is absent in the central retina in both species (A, C). Sparse immunoreactive mitotic figures are observed in the most peripheral retina, arrows in (B, D). CMZ: circumferential marginal zone; GCL: ganglion cell layer; INL: inner nuclear layer; IPL: inner plexiform layer; OLN: outer nuclear layer. Scale bar: 50 μm. Figure 2 is sourced from our laboratory (unpublished).

At hatching, the central retina of altricial birds such as the zebra finch (*T. guttata*) shows the typical layering of the mature tissue. However, signs of immaturity are found such as a increased thickness of the ganglion cell layer, very thin plexiform layers, and photoreceptors displaying immature morphology (Rojas et al., 2007; Álvarez-Hernán et al., 2018, 2020). Nevertheless, the main feature of immaturity is that neurogenesis is intense in the altricial retina in newly hatched animals (Figure 3). Proliferating cell nuclear antigen (PCNA) is expressed by retinal progenitors in the avian retina (Fischer and Reh, 2000; Álvarez-Hernán et al., 2018, 2020). Thus, on hatching day (post-hatching day 0, P0), abundant PCNA-immunoreactive nuclei are detected in the outer half of the inner nuclear layer, the region where the bipolar/Müller cell bodies are located (Figure 3A and B). Bipolar and Müller cells are the last cell types that exit the cell cycle in the developing retina (Amini et al., 2018). It has been described that strong expression of PCNA is detected in early degenerating cells (Borges et al., 2004). During perinatal stages, sparse pyknotic nuclei are detected in the inner nuclear layer of altricial birds (unpublished data) and therefore, some of the PCNA-positive nuclei could correspond to apoptotic cells.

Furthermore, the peripheral-most retina shows abundant PCNA immunoreactive nuclei (Figure 3A and C), suggesting that the altricial retina is still undergoing intense growth via the addition and integration of concentric rings of newly generated cells. The PCNA immunostaining observed in the P0 *T. guttata* retina is similar to that described in the E12 *G. gallus* retina, about nine days before hatching (Ghai et al., 2008).

Also, abundant mitotic activity is found in the ventricular retinal surface in both the central (Figure 3D) and peripheral (Figure 3E) retina of newborn *T. guttata*, but also in hatching chicks (PS) (Figure 3F and G), showing that neurogenesis is intense in the altricial retina after hatching (Álvarez-Hernán et al., 2018, 2020).

Conclusion

The neural retina has been the tissue of choice to study the main aspects of embryonic and postnatal neurogenesis. Current research in teleost fish and amphibians demonstrates that adult neural stem cells exist in the retina of these species (Araki, 2014; Pavón-Muñoz et al., 2016; Madelaine and...
Mourrain, 2017). These populations of cells are crucial to the function of this region of the visual system, since they can both permit physiological cellular turnover and replace cells lost from injury or disease. Proliferating cells within the mature retina are also present in other vertebrates, including reptiles (Todd et al., 2015; Eymann et al., 2019) and birds (Fischer and Reh, 2000). These non-mammalian species provide excellent comparative models for examining the molecular and cellular mechanisms of adult retinal stem cell maintenance and fate. Precocial birds such as the chicken or quail generate most of their retinal tissue at the peripheral retinal margin (Fischer and Reh, 2000). Müller cells are considered to be astrocyte-related cells that acquire retinal stem cells characteristics after injury (Hoang et al., 2020). In our current work (Álvarez-Hernán et al., 2018, 2020), we have shown that the patterns of proliferation reveal that neurogenesis is intense in the retinas of altricial birds at hatching and even at early stages post-hatching. Strong proliferative activity is detected in both the CMZ and the bipolar/Müller cell layer. Therefore, the altricial retina continues to grow during post-embryonic stages, even until day 5 after hatching. Furthermore, the differentiation of bipolar and Müller cells extends into the post-hatching period. It is necessary to determine whether neurogenesis is still detected in the retina of juveniles or adult altricial birds, but long-life neurogenesis in the retina of altricial birds opens up an additional field beyond developmental neurogenesis. It could constitute a powerful tool to determining how post-embryonic neurogenesis is regulated under physiological and regenerative conditions. The understanding of the molecular mechanisms involved in neurogenesis could have crucial implications for regenerative therapies. In conclusion, the retina of altricial birds such as canaries or zebra finches, models used for the study of brain plasticity, may constitute a useful model system for eye research, both in development and, given its postnatal neurogenic activity, regeneration research.

Figure 3- PCNA (A–C) and pHisH3 (D–G) immunoreactivity in the P0 (A–E) and P5 (F, G) T. guttata retina. Abundant PCNA-positive nuclei are detected in the bipolar/Müller cell layer (A–C) and in the CMZ (A, C). Abundant pHisH3-positive cells are found in the vitreal surface of the central (D, F) and peripheral (E, G) retina. CMZ: Circumferential marginal zone; GCL: ganglion cell layer; INL: inner nuclear layer; IPL: inner plexiform layer; L: lens; ONL: outer nuclear layer; OPL: outer plexiform layer. Scale bars: 500 µm in A; 100 µm in B and C; 50 µm in D–G. Figure 3 is sourced from our laboratory (unpublished).

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