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Nuclear Progesterone Receptors Are Up-Regulated by Estrogens in Neurons and Radial Glial Progenitors in the Brain of Zebrafish

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Abstract

In rodents, there is increasing evidence that nuclear progesterone receptors are transiently expressed in many regions of the developing brain, notably outside the hypothalamus. This suggests that progesterone and/or its metabolites could be involved in functions not related to reproduction, particularly in neurodevelopment. In this context, the adult fish brain is of particular interest, as it exhibits constant growth and high neurogenic activity that is supported by radial glia progenitors. However, although synthesis of neuroprogestagens has been documented recently in the brain of zebrafish, information on the presence of progesterone receptors is very limited. In zebrafish, a single nuclear progesterone receptor (pgr) has been cloned and characterized. Here, we demonstrate that this pgr is widely distributed in all regions of the zebrafish brain. Interestingly, we show that Pgr is strongly expressed in radial glial cells and more weakly in neurons. Finally, we present evidence, based on quantitative PCR and immunohistochemistry, that nuclear progesterone receptor mRNA and proteins are upregulated by estrogens in the brain of adult zebrafish. These data document for the first time the finding that radial glial cells are preferential targets for peripheral progestagens and/or neuroprogestagens. Given the crucial roles of radial glial cells in adult neurogenesis, the potential effects of progestagens on their activity and the fate of daughter cells require thorough investigation.

Introduction

In adult vertebrates, production of sex steroids has long been considered to be restricted to the gonads and, on a lower level, the adrenals. However, it is now acknowledged that de novo synthesis of steroids from cholesterol also occurs within the brain. The evidence is based on the fact that all steroidogenic enzymes are expressed in the brain and exhibit biological activity. The synthesis of such “neurosteroids” was initially demonstrated in mammals and subsequently in other vertebrates [1,2,3,4,5,6,7].

While many studies have focused on the feedback effects of sex steroids on the organization and regulation of neuroendocrine and behavioral circuits, there is increasing interest for non-reproductive functions of steroids in the brain, notably on neurogenesis and brain repair. This is particularly the case of estrogens and progesterone, whose effects on brain development and brain repair are more and more studied [8,9]. In the case of progesterone and its metabolites, accumulating data indicate possible involvement in neurodevelopment and neuroprotection [10,11,12]. Neuroperotective effects of progesterone were notably demonstrated in experimental models of traumatic brain injury or ischemia [10,11,13,14]. In addition, in vitro studies using brain slices of neonate rats showed that progesterone promotes dendritic growth and dendritic spine formation in Purkinje cells and that these effects are blocked by the progesterone receptor antagonist mifepristone [15,16]. Recent studies in adult mice showed that progesterone enhances the survival and the migration of newborn neurons in the dentate gyrus [17] or in the subventricular zone (SVZ) of the hippocampus after a focal cerebral ischemia [18]. There is indication that neuroprotective effects of progesterone could be mediated by various mechanisms such as reduction of neuronal vulnerability to neurotoxic molecules [19], reduction of cell loss, inhibition of lipid peroxidation and expression of pro-inflammatory genes [11,20].

In both the adult and developing rodent brain, multiple hypothalamic and extrahypothalamic sites are targeted by progesterone through activation of either nuclear or membrane receptors. In mammals, two nuclear progesterone receptor isoforms encoded by a single gene, PRA and PRB, mediate the genomic effects of progesterone [21]. More recently, it was shown that PRA and PRB can interact as dimers not only with DNA progesterone responsive element but also with signaling proteins of the Src/Ras/Erk pathway outside the nucleus [22,23,24]. The recent identification of membrane progesterone receptors adds further complexity to the picture [25,26]. Interestingly, in rodents, studies have shown that various forebrain regions express nuclear and
receptors for progesterone (PRA and PRB) during specific periods of brain development. These include the preoptic-hypothalamic areas involved in reproductive neuroendocrine functions [27], but also regions not classically associated with reproduction, such as the thalamus, the hippocampus [27] and the neocortex [28,29]. These data suggest that progesterone could also be involved in neural development.

The central effects of progesterone and/or its metabolites are poorly documented in teleost fishes. In fact, it is only known that 4-pregnen-17,20β-diol-3-one (17,20β-DHP) secreted by the ovary at the time of spawning acts as a pheromone stimulating sexual behavior [30,31]. However, recent reports have established that the zebrafish brain exhibits 3β-hydroxysteroid dehydrogenase/Δ5-4 isomerase (3β-HSD) activity and expression [7,32]. In addition, biochemical studies indicated that the progesterone synthetic pathway is very active, resulting in brain production of progesterone and progesterone derivatives, such as 17-hydroxyprogesterone and dihydro-progesterone. This suggests that, similar to the situation in mammals, progestagens actions in the fish brain result from both peripheral production and local synthesis.

Until now, virtually nothing is known regarding the expression of either nuclear or membrane-bound progesterone receptors in fish. The recent characterization of a unique nuclear progesterone receptor (pgr) in zebrafish and the development of antibodies to zebrafish Pgr now opens the possibility of investigating in detail the distribution of Pgr in the brain, and the phenotype of the cells expressing this Pgr [33,34]. Nuclear PR receptor binding assays and dual transactivation/transcription assays clearly established that progestins, including progesterone, 17-hydroxyprogesterone, dihydro-progesterone and 17,20β-DHP specifically bind and activate zebrafish Pgr [33,34]. Preliminary data indicated that Pgr is widely distributed in the brain of zebrafish [33], notably in the ventricular proliferative regions. In adult fish, the brain ventricles are lined by radial glial cells acting as neuronal progenitors and supporting the migration of newborn neurons [35,36,37,38,39,40]. In contrast to mammals, in which radial glial cells become astrocytes at the end of migration of newborn neurons [35,36,37,38,39,40], these cells in zebrafish persist in the whole brain during adulthood, supporting its constant growth. In the brain of adult zebrafish, radial glial cells express classical markers such as GFAP (Glia Fibrillary Acid Protein), BLBP (Brain Lipid Binding Protein), protein S100β, CXCR4 (a chemokine receptor) or nestin [35,37,38,47,48,49]. In contrast to mammals, in which radial glial cells become astrocytes at the end of embryogenesis [41,42,43,44,45,46], radial glial cells persist in the whole brain during adulthood, supporting its constant growth. In the brain of adult zebrafish, radial glial cells express classical markers such as GFAP (Glia Fibrillary Acid Protein), BLBP (Brain Lipid Binding Protein), protein S100β, CXCR4 (a chemokine receptor) or nestin [35,37,38,47,48,49]. In contrast to mammals, in which radial glial cells become astrocytes at the end of embryogenesis [41,42,43,44,45,46], radial glial cells persist in the whole brain during adulthood, supporting its constant growth.

In situ hybridization

Brains (n = 6) were partially dissected through skull opening and fish were immersed overnight at 4°C in 4% paraformaldehyde (PFA) in saline phosphate buffer (PBS, pH 7.4). On the next day, brains were taken out and fixed at 4°C in 4% PFA, in order to be processed for paraffin embedding and cut at 6 μm. Sections were mounted on poly-lysine slides for subsequent processing using the in situ hybridization protocol described previously by Diotel and colleagues [49]. Riboprobe synthesis was previously described by Hanna and colleagues [33]. In order to ascertain the in situ hybridization specificity, sense and antisense probes were always hybridized on parallel sets of slides.

Immunohistochemistry

All staining was performed on frozen larval sections or frozen brain sections (12 μm) from adult zebrafish (n = 6) prepared with a cryostat. Samples were fixed in 4% PFA overnight at 4°C. The samples were next immersed in PBS containing 30% of sucrose overnight, embedded in Tissue-tek before freezing at −80°C. Immunohistochemistry experiments were processed as previously described [33]. Briefly, tissue sections were first rinsed in PBS and non-specific binding was blocked for 1 hour at room temperature in PBS containing 0.2% triton and 0.5% milk powder. The antibody used was raised against peptides based on the N-terminal part (aa100–112) of zebrafish Pgr [33]. Sections were then incubated with rabbit anti-Pgr antibody (1:2500–1:5000) alone or with mouse anti-acetylated tubulin (1:100; Acetyl-Tub; clone 6-11B-1; Reference T 6793, Sigma) overnight at room temperature. After several washes in PBS 0.2% triton, tissue sections were incubated with the appropriate secondary antibodies (Goat anti-rabbit Alexa 594 or goat anti-mouse Alexa 488, Invitrogen) and slides were then rinsed in PBS-Triton. After immunohistochemical processing, cell nuclei were visualized with a DAPI counterstaining (Vectorshield with DAPI, Vector Laboratories). The Pgr staining specificity has already been demonstrated [33]. Furthermore, omission of the primary or secondary antibodies caused no reaction.

Zebrafish treatment and RNA extraction

Adult male zebrafish were exposed to either to 10−7 M of 17β-estradiol (17β-estradiol, Sigma-Aldrich, St. Louis, MO, USA) for 100 hours, or to 10−8 M of 1,4,6-androstatriene-3,17-dione (ATD) for 7 days, in glass tanks maintained at 28.5°C. Control fish were exposed to ethanol. Larvae were treated with 17β-estradiol (10−8 M in EtOH) or with EtOH alone, from 0 to 8 days post-fertilization. In both cases, water and hormones were replaced every day. After exposure, individual brains, pools of 5 brains of adult zebrafish, and pools of 20 larvae were sonicated.
(10 sec, three times) in 1 mL Trizol Reagent (Gibco, Carlsbad, CA, USA), and total RNA was extracted according to the manufacturer’s protocol. Each experiment was performed at least twice.

Quantitative real-time PCR
Reverse transcription was carried out by incubating 2 μg total RNA with 1 μg of random primer oligonucleotides, 2.5 mM dNTPs and 50 U MMLV-RT (Promega) in the appropriate buffer 10 minutes at 65°C and 60 minutes at 37°C. Polymerase Chain Reactions (PCRs) were performed with an iCycler thermocycler coupled to the MyiQ detector (Bio-Rad, Hercules, CA, USA) using iQ SYBR-Green Supermix (Bio-Rad) according to the manufacturer’s protocol. The following primers were used: EF-1 (fw) 5’-AGCAGCAGCAGTGGAGATGAT-3’; EF-1 (rev) 5’-CGCGGATTCTAGATGGTAT-3’; pgr (fw) 5’-GAGCAAT-GATGCGTGAGAAGG-3’; pgr (rev) 5’-TCCAGAAAGGAA-CAGTGTGAGG-3’. Expression levels of EF-1 mRNA were used to normalize the expression levels between each sample. Triplicates of each sample were loaded in the 96 well-plate for each analyzed gene. Melting curve and PCR efficiency analyses were performed to confirm correct amplification. The Delta-Delta CT method was used for calculating the relative expression. The were performed to confirm correct amplification. The Delta-Delta

Microscopy
Observations were carried out on an Olympus Provis photomicroscope equipped with a DP50 digital camera or with an epifluorescence Zeiss (Imager Z1, equipped with the Apotome module). Micrographs were taken in the TIFF format using the Analysis software, allowing image superposition. Images were then prepared with Adobe Photoshop CS4 for light or contrast adjustment before preparation of the plates.

Pgr fluorescence intensity quantification
In order to quantify Pgr staining intensity in radial glial and parenchymal cells, pictures were converted into 8 bit. Using the ImageJ software, Pgr-positive cell nuclei located in the regions of interest were given a value according to a gray scale. In order to determine the effect of 17β-estradiol on Pgr fluorescence intensity, pictures of control and treated larvae were taken at the same exposure time. Numerous cells were then selected and the same method was used to analyze Pgr intensity. Significant differences were determined using a Student’s t test. Three independent samples were analyzed.

Nomenclature
The nomenclature for brain nuclei is according to that established in the zebrafish atlas by Wullimann and colleagues (1996) [52].

Results
Pgr expressing cells are largely distributed in the adult brain
The distribution of nuclear progesterone receptor mRNA and protein in the forebrain of adult zebrafish was assessed by in situ hybridization and immunohistochemistry. Overall, these two techniques yielded identical results and thus reciprocally validate each other in addition to the other controls.

Figure 1 shows transverse sections hybridized with the pgr antisense probe in the tel-, di- and mesencephalon (Figures 1A to 1D). In agreement with the RT-PCR data showing high pgr gene expression in the brain [33], in situ hybridization also revealed strong pgr mRNA expression, widely distributed in all brain regions. Figure 1 also presents parallel sections hybridized with the anti-sense and sense probes (Figures 1E-1H).

The pgr transcripts are found in numerous cells from the olfactory bulbs to the mesencephalon. Similarly, immunohistochemistry showed an identical widespread distribution of the Pgr protein (Figures 1, 2 and 3). Progesterone receptor expressing cells are distributed in all brain regions in many cells as shown by the DAPI counterstaining (Figures 2A-2F). However, DAPI counterstaining clearly indicated that a significant proportion of cells in every brain region do not express Pgr (Figures 2D-2F). Similar to the hybridization signal, a marked heterogeneity in the intensity of the immunohistochemical staining was also observed (Figures 2A-2F).

The most anterior cells expressing Pgr were located in the granular cell layer of the olfactory bulbs (data not shown). Numerous labeled cells were detected in the telencephalic hemispheres (Figures 1A, 1E, 3A, 3D and 3G). Indeed, both in situ hybridization and immunohistochemistry generated intense labeling in both pallial and subpallial regions (Figures 1A, 1E, 3A, 3D and 3G). An obvious characteristic of the hybridization and immunohistochemical signals in the telencephalon was that the periventricular regions consistently exhibited stronger staining than cells in the parenchyma, which was also true at other brain levels (see below). For example, cells bordering the ventricles in the subpallial and pallial regions always expressed stronger Pgr staining compared to cells located in the parenchyma (Figures 3A, 3D and 3G). Quantification of Pgr staining by image analysis clearly showed that in the subpallium and the anterior hypothalamus, cells lining the ventricle exhibit significantly stronger staining than cells in the parenchyma (Figure 4).

Both in situ hybridization and immunohistochemistry produced very consistent staining in the anterior (PPa) and posterior part of the preoptic region and also in the post-commissural nucleus of the ventral telencephalic area (Vp) (Figure 1B, 2A and 2D). In the anterior preoptic region, Pgr protein and mRNA expressing cells were abundant in the parvo cellular preoptic nucleus (PPa) (Figures 1B and 2A). Again, immunohistochemistry clearly showed that the ependymal layer was more heavily labeled than the parenchyma (Figures 2A and 4). We also observed expression of nuclear progesterone receptors in the diencephalon notably in the ventromedial and ventrolateral thalamic nuclei (not shown), the ventral habenular nucleus, the periventricular nucleus of posterior tuberculum, and the diffuse nucleus of the inferior lobe (Figures 1D, 2J and 3J). Very high expression of pgr messengers and protein was also detected at all levels of the mediobasal hypothalamus (Figures 1C, 1G, 2G and 2J), in the periventricular regions of the lateral (Figure 1D and 1G) and posterior recesses (Figure 3M). Furthermore, we observed that cells surrounding the lateral recess (LR) strongly express pgr (Figure 1D and 1G). A remarkable finding was the very high expression of mRNA and protein in numerous cells of the posterior tuberal nucleus (PTN) and in cells lining the ventral zone of periventricular hypothalamus (Figures 1C, 1G, 2G and 2J). On the contrary, more laterally, in the lateral hypothalamic nucleus (LH) and in the anterior tuberal nucleus (ATN), the intensity of the Pgr labeling decreases while the number of stained cells remains constant (Figure 2J).

We also observed Pgr expressing cells in the torus longitudinalis (Figure 1C) and in the optic tectum, notably in the densely packed small cells of the periventricular gray zone (PGZ) (Figure 1C, 1D and 2M). The valvula cerebelli and the torus semicircularis also clearly expressed pgr (Figure 1D).
Pgr is differentially expressed in neurons and radial glial cells

A major characteristic of the expression pattern of nuclear progesterone receptors, at both the mRNA and protein levels, was the consistently stronger signal identified along the brain ventricles, suggesting higher Pgr expression in radial glial cells than in surrounding neurons. In order to further characterize the Pgr expressing cells, we performed double staining experiment using a monoclonal antibody to acetylated tubulin, an established marker of neurons. We found that many cells located in the parenchyma exhibited a cytoplasm labeled by acetylated tubulin, but also a Pgr positive nucleus. As shown in Figure 2G–2O, this was observed in many regions. As expected, the cells bordering the ventricles did not contain acetylated-tubulin labeling (Figure 2G–2L). By performing Pgr immunohistochemistry on tg(cyp19a1b-GFP) transgenic fish, which express GFP in radial glial cells under control of the cyp19a1b promoter, we demonstrated that cells strongly expressing Pgr along the ventricles correspond to cyp19a1b-GFP positive radial glial cells (Figure 3A to 3O). Consequently, it appears that Pgr expression is higher in the estrogen-synthesizing radial glial cells.

Nuclear progesterone receptor mRNA and protein are positively regulated by estrogens

Given that Pgr expression was higher in aromatase B expressing radial glial cells, we decided to investigate the potential involvement
of the estrogenic environment on Pgr expression by treating fish either with an aromatase inhibitor (ATD) or with 17β-estradiol. The data generated by real time-qPCR clearly demonstrated that treatment with ATD (10^{-6} M) leads to a significant 50% decrease of pgr expression after 7 days of treatment (Figure 5A). In contrast, treatment of adult fish with 17β-estradiol (10^{-7} M) for 100 hours significantly increases pgr expression levels (Figure 5B). Treatment of eggs and larvae with 17β-estradiol (10^{-8} M) for 8 days also leads to a two fold increase in pgr mRNA expression as assessed by real time-qPCR (Figure 5C). This result was confirmed at the protein level using immunohistochemistry. Figure 6 shows that, as expected, E2 treatment increases cyp19a1b-GFP levels in radial glial cells and also increases the number of cyp19a1b-GFP positive radial glial cells [50]. In addition, 17β-estradiol causes increased expression of Pgr protein in radial glial cells along the brain ventricles, but also within neurons in the parenchyma (Figure 6). Although we observed that estradiol leads to an increase in the number of both parenchymal and cyp19a1b-GFP radial glial cells expressing Pgr, this increase was not significantly different (Figures 7 A and B). Furthermore, we noted that there were more cyp19a1b-GFP radial glial cells expressing Pgr in the E2-treated larvae compared to control larvae (Figure 7C). Finally, it appeared that the Pgr staining intensity was significantly stronger in the E2-treated larvae (Figures 7D and E). Such data suggest that estrogen treatment leads to an increase in Pgr expression, but also to a higher number of Pgr expressing cells corresponding to both neurons and radial glial cells.

**Discussion**

The present study further documents the sites of Pgr expression in the brain of the zebrafish, a teleost fish. We clearly establish by
both in situ hybridization and immunohistochemistry that the zebrafish brain is widely targeted by progesterone and/or progesterone derivatives, in particular in the preoptic area and the hypothalamus. However, we also show that Pgr is expressed in many cells of non-neuroendocrine regions. Additionally, we demonstrate that Pgr staining is stronger in radial glial cells than in neurons, implicating crucial progenitor cells as preferential targets of progesterone. Finally, we provide evidence that estrogens up-regulate Pgr expression in the brains of fish, similar to what is known in mammals.

A wide expression of Pgr in the zebrafish brain

The localization and functions of Pgr in the fish brain are very poorly documented [33,53]. The present study shows a very strong and widely distributed expression of pgr mRNAs in the whole brain, confirming the data obtained by immunohistochemistry.

This further reinforces the specificity of the zebrafish Pgr antibody previously validated by western blotting [33]. In all cases, the staining is nuclear as expected for nuclear receptors. Apart from zebrafish, expression of Pgr in teleosts has only been documented in the African cichlid, *Astatotilapia burtoni*, in which similar results were obtained in terms of abundance and localization [53]. In mammals, Pgr is also widely expressed in the brain, notably in the preoptic area, the hypothalamus or also in the hippocampus [27,54]. Interestingly, Pgr expression levels in rodents seem to be developmentally regulated [27]. In the developing forebrain of rat, numerous brain nuclei indeed appear to express Pgr, while, at postnatal day 28, expression is much lower in most brain regions [27]. In mammals, the phenotype of Pgr-expressing cells is partly documented. Nuclear progesterone receptors were notably found in dopaminergic neurons of the preoptic area and hypothalamus, but not in the ventral tegmental area [55]. In addition, Pgr in

![Figure 3. Radial glial cells strongly express Pgr.](https://example.com/figure3.jpg)
mammals were also observed in glial cells, including oligodendrocytes and astrocytes [8,56,57].

Despite this wide expression of Pgr in the brain of fish, there is virtually no data on the functions that could be potentially affected by progestagens.

Pgr is strongly expressed in radial glial cells in adult zebrafish

A major finding of this study is the discovery that Pgr is not only detected in neurons, but that is also expressed in radial glial cells. Such cells were first described in the developing brain of mammals by Magini at the end of the 1800s [58]. By definition, they exhibit a small soma localized along the ventricles and extend two cytoplasmic processes, one to the ventricular surface and the other to the pial surface [58,59]. In the developing brain of mammals, radial glia were first documented as a support system for the migration of newborn neurons [41,44,58,60,61], but it is now admitted that radial glial cells have more crucial functions. Indeed, they are acknowledged as neural stem cells that can give birth, directly or indirectly, to all cell types of the brain, including neurons, oligodendrocytes, ependymal cells and astrocytes [45,46,62]. In contrast to mammals in which radial glial cells transform into astrocytes at birth [42,60,62,63], they persist in the brain of fish during adulthood.

Using the transgenic tg(cyp19a1b-GFP) zebrafish line that expresses GFP in radial glial cells [51], we unambiguously demonstrate Pgr expression in GFP expressing cells lining the ventricles. This result is particularly interesting in view of the fact that radial glial cells act as neural stem cells and sustain constant brain growth in adult fish [36,37,38,64]. Recent data in zebrafish indicate that radial glial cells actively divide symmetrically or asymmetrically to generate both neurons and radial cells [36,37,40,48]. These data have led some authors to suggest that the adult fish brain is a developing organ that conserves some

figure 4. expression of pgr is stronger in radial glial cells than in neurons. Image analysis of the pgr staining intensity in the ventral subpallium and in the anterior hypothalamus showing that radial glial cells exhibit a stronger pgr immunoreactivity than parenchymal cells (p<0.001 Student’s t test). The graphs present the mean value +/- the standard deviation. doi:10.1371/journal.pone.0028375.g004

figure 5. Nuclear progestin receptor is up-regulated by estrogen in both the developing and adult brain. Fold induction of pgr expression after treatment of adult zebrafish with an aromatase inhibitor (A, 10^-6 M of ATD) or estradiol (B, 10^-7 M of 17β-estradiol), and larvae with E2 (C, 10^-8 M of 17β-estradiol). In A, the aromatase inhibitor (ATD) leads to a significant decrease of pgr expression in individual brains of adult zebrafish (n = 4). In B, the estrogenic treatment leads to a significant increase of pgr expression in pools of 5 brains of adult zebrafish (n = 3). In C, the estrogenic treatment leads to a significant increase of pgr expression in pools of 20 larvae (n = 2). Asterisk (*) indicates significant differences (p<0.05, Student’s t test). The graphs present the mean value +/- the standard deviation. doi:10.1371/journal.pone.0028375.g005
features of the embryonic mammalian brain \[36,38,39,40,65\]. This could explain the abundance of Pgr in the brain of fish. Indeed, it was recently proposed that the transient expression of Pgr in many brain regions during development in rodents reflects a role for progesterone and Pgr in fundamental mechanisms of neural development \[27\]. During brain development in rats, the nuclear progesterone receptor is transiently expressed in regions not typically associated with neuroendocrine functions, such as the neocortex, the dentate gyrus, the caudato-putamen, the periventricular striatum \[27\]. This suggests a developmental function for progesterone in these regions and reinforces the view that steroid hormones could play a significant role in neurogenesis \[8,9,27,38,64\]. Studies in rats also suggest that Pgr expressed in the subventricular zone participate in neural stem cell proliferation \[27\]. Whether progesterone is involved in radial glial cell proliferation in developing or adult fish will have to be investigated. In any case, this raises the question of the source, peripheral or central, of progesterone available for Pgr binding during embryonic brain development in mammals or adult neurogenesis in fish.

**Radial glial cells: Source and target for neuroprogesterone**

Our own results in zebrafish suggest that, the brain, in addition to the gonads, is a source of neurosteroids \[7,32\]. Similar to other vertebrates \[1,2,3,4,5,6\], the brain of fish is able to produce a number of steroids, including progesterone and progesterone derivatives, suggesting a role for progestagens in brain functions \[7,32\]. In particular, the steroidogenic enzyme 3β-HSD which catalyzes the oxidation and isomerization of pregnenolone into progesterone is expressed in the brain of zebrafish. In zebrafish, 3β-HSD activity was first reported in the adult brain by Sakamoto.
and colleagues [32]. Our recent experiments confirmed the presence of 3\beta-HSD activity and expression in the brain of adult zebrafish, which also expresses other steroidogenic enzymes (Cyp11a1, Cyp17; 3\beta-Hsd, 3\alpha-Hsd, 5\alpha-reductase and aromatase) [7]. Interestingly, it also appears that radial glial cells could express 3\beta-hsd, cyp11a1 and cyp17 mRNAs in addition to the estrogen synthesizing enzyme aromatase [36,50]. Thus, the brain of adult zebrafish appears to be a steroidogenic organ, able to synthesize its own (neuro)steroids which act locally to regulate some brain functions, potentially neurogenesis and plasticity [7].

In mammals, astrocytes are also known to be both a source of neurosteroids, including progesterone [66,67,68,69,70], and a target for steroids and/or neurosteroids [3,6,56,57,71]. Until now, there is no solid evidence that the forebrain of fish contains bona fide star shaped astrocytes. This is understandable in view of the fact that radial glial cells persist throughout life in fish while they transform into astrocytes at birth in mammals [72,73]. Furthermore, radial glial cells in adult fish express many of the canonical markers of the astrocyte lineage in mammals [35,36,37,48,49,74]. Thus, all these data suggest that radial glial cells in zebrafish are both a source and a target for neuroprogesterone.

Estrogens up-regulate nuclear progesterone receptor expression

While it is well known that Pgr is up-regulated by estrogens in the brain of mammals, our data are the first to demonstrate that nuclear progesterone receptors are also up-regulated by estrogens in the brain of zebrafish at both the mRNA and the protein levels. Evidence based on qPCR experiments shows that fish treated with ATD, an aromatase inhibitor, exhibit lower brain pgr expression, whereas fish treated with estrogens exhibit a significant increase in pgr mRNA levels. Experiments in larvae and adult brains further reinforced this result by showing that E2 increased pgr mRNAs (qPCR) and Pgr immunoreactivity as measured by image analyses. Such data demonstrate that estradiol treatment significantly increased progesterone receptor expression [8,67,68,69,73,76]. According to our immunohistochemical results, the pgr up-regulation upon E2 treatment would concern both neurons and radial glial cells. In the brain of teleosts, there is evidence for the presence of nuclear estrogen receptor esr1, esr2a and esr2b in many brain regions, in particular in the periventricular regions of the forebrain, in neurons and probably in radial glial cells [77,78,79] (N. Diotel et. al, unpublished).

Conclusions

Complementing our previous studies on the expression of steroidogenic enzymes in the brain of zebrafish [7,38], the present report provides for the first time detailed information on the distribution and nature of Pgr positive cells. The data indicate a widespread localization of Pgr in many brain regions, suggesting actions outside the reproductive sphere. This receptor is present in many neurons throughout the brain, but it is obvious that radial glial cells exhibit stronger Pgr staining. Such data are particularly significant given the fact that radial glial cells are now recognized as progenitors sustaining brain growth in adult fishes. Additionally, such cells are also unique in strongly expressing brain aromatase and, to a lesser extent, other steroidogenic enzymes. Radial progenitors thus appear to be both sources and targets of neurosteroids, notably progestagens. In addition, we also show that, similar to mammals, pgr expression is up-regulated by E2.

Figure 7. Effects of estradiol on Pgr expression in the preoptic area. The total number of Pgr, cyp19a1b-GFP and DAPI expressing cells were counted in the same region of the preoptic area in controls and E2-treated larvae, respectively (n = 3). A: Effect of estrogens on the proportion of DAPI positive cells expressing Pgr. B: Number of cyp19a1b-GFP expressing radial glial cells per section of control and E2-treated larva. C: Effect of estrogens on the proportion of cyp19a1b-GFP positive radial glial cells showing Pgr expression. D and E: Analysis of the fluorescence intensity of Pgr staining in control and estradiol-treated larvae shown in Figure 6 (D corresponds to Figures 6 B and F; E corresponds to Figures J and N). n = number of cells counted. The asterisk (*) means that the differences observed are significant (p<0.001, Student’s t test). The graphs present the mean value +/- the standard deviation.

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Altogether, these results suggest that the potential functions of progestagens, of peripheral and central origins, have been overlooked until now and should be investigated in particular in relation to the neurogenic activity of radial progenitors under physiological conditions, but also under situations of brain repair.

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Author Contributions

Conceived and designed the experiments: ND EP CV OK IA. Performed the experiments: ND M-MG SM AS IA. Analyzed the data: ND IA YZ OK. Contributed reagents/materials/analysis tools: YZ. Wrote the paper: ND IA OK.

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