BDNF Expression in Larval and Adult Zebrafish Brain: Distribution and Cell Identification

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Abstract

Brain-derived neurotrophic factor (BDNF), a member of the neurotrophin family, has emerged as an active mediator in many essential functions in the central nervous system of mammals. BDNF plays significant roles in neurogenesis, neuronal maturation and/or synaptic plasticity and is involved in cognitive functions such as learning and memory. Despite the vast literature present in mammals, studies devoted to BDNF in the brain of other animal models are scarce. Zebrafish is a teleost fish widely known for developmental genetic studies and is emerging as model for translational neuroscience research. In addition, its brain shows many sites of adult neurogenesis allowing higher regenerative properties after traumatic injuries. To add further knowledge on neurotrophic factors in vertebrate brain models, we decided to determine the distribution of bdnf mRNAs in the larval and adult zebrafish brain and to characterize the phenotype of cells expressing bdnf mRNAs by means of double staining studies. Our results showed that bdnf mRNAs were widely expressed in the brain of 7 days old larvae and throughout the whole brain of mature female and male zebrafish. In adults, bdnf mRNAs were mainly observed in the dorsal telencephalon, preoptic area, dorsal thalamus, posterior tuberculum, hypothalamus, synencephalon, optic tectum and medulla oblongata. By combining immunohistochemistry with in situ hybridization, we showed that bdnf mRNAs were never expressed by radial glial cells or proliferating cells. By contrast, bdnf transcripts were expressed in cells with neuronal phenotype in all brain regions investigated. Our results provide the first demonstration that the brain of zebrafish expresses bdnf mRNAs in neurons and open new fields of research on the role of the BDNF factor in brain mechanisms in normal and brain repairs situations.

Introduction

Brain derived neurotrophic factor (BDNF) is a protein that belongs to the neurotrophin family, including nerve growth factor (NGF) and neurotrophin (NT) 3, NT 4/5 and NT 6/7 [1]. All
neurotrophins interact with two types of receptors, tropomyosin-related receptor kinase (Trk) and p75 neurotrophin receptor (p75NTR). In the brain of mammals, BDNF promotes through TrkB receptor, neuronal survival, growth, differentiation and synaptic plasticity [2]. In addition, BDNF has shown to modulate, through p75NTR receptor, neuronal migration [3], myelination [4] and neuronal apoptosis [5, 6]. Furthermore, BDNF seems to be involved in modulating memory, learning processes [7], age-dependent alterations of hippocampus [8, 9] and neurodegenerative diseases [10–12]. The bdnf gene is composed of multiple alternative exons: 10 in human, 8 in mouse, and 6 in non-mammalian vertebrates [13]. Multiple promoters can modulate the tissue specific transcription of the bdnf gene [14–17]. Multiple promoters can modulate the tissue specific transcription of the bdnf gene [14–17]. This gene is well conserved across vertebrate evolution [14, 18, 19] and some regulatory sequences in the 5′ UTR of the bdnf gene appear highly conserved between zebrafish and mammals [18], suggesting conserved functions.

Bdnf mRNAs and protein distributions have been described in the brain of different vertebrates species during development and in adults. In rat, bdnf transcripts dramatically increased between embryonic days 11 and 12, a timing coinciding with neurogenesis [20]. Studies performed in rat, mouse and pig showed that bdnf transcripts and proteins were detected at different postnatal stages (newborn, adult) in regions such as hippocampus, hypothalamus, cerebral cortex, amygdala and brain stem adrenergic nuclei [20–27]. In the human brain, bdnf mRNAs were also reported in the hippocampus, the amygdala and the septum [28, 29]. There are few studies in birds and amphibians, where BDNF has been described. In songbirds, bdnf is expressed in brain nuclei involved in sensorimotor integration of song learning [30–32]. In amphibians, bdnf mRNAs and immunoreactivity were described in optic tectum and hypothalamus [33, 34]. In adult teleost fishes, bdnf mRNAs are reported in the brain of eel [35], perciform (Cichlasoma dimerus) [36] and Turquoise killifish (Nothobranchius furzeri) [37]. In zebrafish, bdnf mRNAs are present early during embryonic development and whole-mount in situ hybridization experiments at early developmental stages demonstrated the presence of bdnf transcripts in the forebrain, midbrain and hindbrain [38].

A wealth of data obtained from rodent studies points to a role for BDNF in neurogenesis. BDNF contributes to the regulation of neurogenesis during development by promoting neuronal survival and differentiation [39, 40], but also plays roles in adult neurogenesis. However, BDNF effects in adult neurogenesis are not completely established and studies have reported conflicting results. In some studies, BDNF infusion into the ventricle of adult rat substantially increased the number of newly formed cells in many regions, the majority of which differentiate into neurons [41, 42]. BDNF administration in the hippocampus was associated with an increased neurogenesis of granule cells in the dentate gyrus [43]. However, more recently, Galvao and colleagues obtained contrasting results suggesting that BDNF delivered intraventricularly in mice and rats failed to enhance subventricular zone neurogenesis and even reduced it [44]. Transgenic mouse models with bdnf gene expression or signalling disruption have been produced to decipher the role of BDNF in regulating neural stem cells but again a clear role of BDNF cannot be stated. Mutant mice lacking BDNF did not survive and had severe neuronal deficits [45, 46]. In heterozygous BDNF knockout mice, proliferation of neural stem cells was decreased in the dentate gyrus of the hippocampus [47]. In contrast, conditional knockout mice with depletion of BDNF in mature neurons exhibited an increase in hippocampal proliferation [48]. The role of BDNF in brain regenerative processes was also investigated in rodents. After experimentally induced brain injury, the expression of bdnf was up-regulated close to the lesioned area, probably as part of the repair mechanism following trauma [49–51]. In the mouse, BDNF treatment improved long-term survival and maturation of newly formed cells derived from the subcallosal zone after cortex injury [52]. Conversely, in transgenic mice in which bdnf gene expression was abolished in astrocytes, a low number of new generated
oligodendrocytes and consequently a larger damaged area were observed after hypoperfusion of the carotid artery [53]. All together, those reports on the role of BDNF in neurogenesis generated conflicting results, which need deeper investigations.

Over the last ten years zebrafish has become an interesting model organism to study the molecular and cell biology of the vertebrate brain [54, 55] and has emerged as a model species for translational research in various neuroscience areas, such as depressive disorders [56], neurodegeneration [57], motor neuron disease [58], autism spectrum disorders [59] and Alzheimer’s disease [60]. In addition, compared to mammals, the brain of adult zebrafish exhibits a high number of proliferative areas distributed along the ventricles of the telencephalon, diencephalon and mesencephalon [61–65], a feature tightly linked to the persistence of radial glial progenitors [61]. Moreover, as shown by several studies, teleost fishes, including zebrafish, exhibit an outstanding capacity to regenerate after brain injury [66–69]. The factors supporting this intense neurogenic activity in normal physiological and reparative conditions are not identified to date.

Virtually nothing is known on BDNF functions in the central nervous system of fish. Given the potential role of BDNF in neurogenesis mentioned above and the strong proliferative activity in the brain of adult zebrafish, we have investigated the potential link between this neurotrophin and neurogenesis in zebrafish. First, we studied in detail the distribution of bdnf mRNAs in the brain of zebrafish and determined, by means of double staining studies, the phenotype (proliferating cells, glial or neuronal cells) of bdnf-expressing cells.

Materials and Methods

Animals and tissue processing

Animals were handled and sacrificed in agreement with the guidelines for the use and care of laboratory animals and in compliance with French and European regulations on animal welfare. Animals used in this study were housed in our zebrafish facilities (INRA LPGP, BIOSIT, Rennes, France, agreement number: B 35-238-6) under standard conditions of photoperiod (14 hours light and 10 hours dark) and temperature (28°C). This project was approved by the local animal care and ethics committee (Comité Rennais d’Ethique en matière d’Expérimentation Animale, Rennes, France), under the number EEA B-35-040. Zebrafish did not receive medical treatment prior or during the experience. No deaths occurred in the facilities before the sacrifice of animals used for in situ hybridization and immunohistochemistry experiments. For studies on larvae, adults were spawned, embryos were collected (n = 20) and kept for 7 days in an incubator in 100 ml glass bottles before being sacrificed in ice water and fixed overnight 4°C in phosphate-buffered saline (PBS; pH 7.4) containing 4% paraformaldehyde (PAF). For studies in adults, four months old females (n = 6) and males (n = 6) were sacrificed by overdose of tricaine methanesulfonate (MS-222, 300mg/l). The sex was then determined by direct examination of the gonads under a binocular stereoscopic microscope. After skull opening, the brain was removed and fixed overnight in PBS-PAF. Both larvae and adult brains were processed for paraffin embedding and microtome sections were mounted on poly-lysine slides.

In situ hybridization

Bdnd expression (ie bdnf mRNAs presence in cells) was investigated using in situ hybridization. For bdnf riboprobes synthesis, we used a pCMV-Sport 6.1 plasmid containing the full-length BDNF cDNA (Unigene DR.132862; Entrez Gene 58118). Antisense and sense riboprobes were generated with DIG RNA Labelling Mix (Roche Diagnostic, Indianapolis, US) by in vitro transcription, using T7 polymerase (Roche-Diagnostic) and SP6 polymerase (Roche-Diagnostic) on plasmid linearized by EcoRI and NotI. To check the specificity of the staining, the sense and antisense riboprobes were always hybridized on adjacent sections. No staining was
observed on sections hybridized with the sense riboprobe either in larvae and adults (S1 and S2 Figs) as demonstrated in a previous study in zebrafish [38]. Paraffin sections (7 μm) were deparaffinized with OTTIX and rehydrated through a series of graded ethanol (100–30%). Sections were washed in PBS-NaCl (0.85%) and post-fixed for 20 minutes in PBS-PFA 4%. Tissues were rinsed in PBS and treated for 7 minutes with protease K (2 mg/ml) diluted in PBS at 37°C. The reaction was stopped in PBS before post-fixation for 20 minutes in 4% PBS-PAF and the slides were rinsed 10 minutes in PBS and 10 minutes in standard saline citrate (SSC 2x).

The sections were incubated overnight at 62.5°C in a moist chamber with the probes (1.5 μg/ml) diluted in hybridization buffer (formamide 50%; SSC 2X, Denhart 5X, yeast tRNA 50 μg/ml, EDTA 4 mM, dextran sulfate 2.5%). On the following day, slides were rinsed with SCC 2x, SCC 2x/formamide 50%, SCC 0.2x and SSC 0.1x. Next, they were dipped in Tris-HCl/NaCl buffer (100 mM, Tris-HCl pH 7.5, 150 mM NaCl) and washed in the same buffer containing 0.1% Triton and 0.5% of milk powder. Sections were incubated overnight at room temperature with anti-digoxigenin alkaline phosphatase Fab fragments (1:2000, Roche Diagnostic). On the next day, slides were rinsed in Tris-HCl/NaCl buffer and washed three times with Tris-HCl 100 mM (pH 8) containing NaCl (100 mM) and MgCl2 (10mM). The hybridization signal was revealed with the HNPP/Fast-Red detection kit (Roche Diagnostic) for 6 or 12 hours according to the manufacturer’s instructions. Sections were washed several times in PBS, mounted in Vectashield medium containing DAPI and coverslipped for microscopic analysis.

**Immunohistochemistry**

To get further insights in the identification of bdnf-expressing cells (ie cells that express bdnf mRNAs), sections were processed for immunohistochemistry just after in situ hybridization. To stop in situ hybridization reaction, sections were dipped 5 minutes in PBS-PAF 4%, washed in PBS and PBS/Triton (0.2%) to perform immunohistochemistry as follows: sections were incubated overnight at room temperature with different primary antibodies diluted in PBS containing 0.5% milk powder. Sections were washed three times in PBS-Triton 0.2% and incubated with goat anti-rabbit or anti-mouse Alexa Fluor 488 (1:200, Invitrogen, ThermoFisher Scientific, France) for 2 hours. Tissue sections were washed in PBS-Triton 0,2%, and slides were mounted with the Vectashield medium containing DAPI for nuclei counterstaining (Vector Laboratories, Burlingame, CA).

Radial glial cells were identified with a rabbit anti-zebrafish aromatase B (this antibody was raised in our laboratory, 1:200) and with a rabbit anti-mouse BLBP (Brain Lipid Binding Protein, 1:100, Chemicon, Temecula, CA, Cat. No. AB9558). The specificity of these antibodies was previously demonstrated in zebrafish [70, 71].

Three neuronal markers were used to identify post-mitotic neurons. The labeling was performed with a mouse monoclonal anti-acetylated-tubulin (1:100, Sigma-Aldrich, France, clone 6-11B-1, Cat. no. T 6793), with a mouse monoclonal anti-HuC/D (1:20, Invitrogen, ThermoFisher Scientific, France, clone 16A11, Cat. no. A21271) or with a mouse monoclonal anti-MAP2 (Microtube-Associated Protein2, 1:100, Abcam, France, clone AP-20, Cat. no. 11268). The specificity of these antibodies was previously confirmed in zebrafish [61, 72].

Proliferative cells were visualized with a monoclonal antibody raised against PCNA (Proliferating Cell Nuclear Antigen, 1:100, Dako, France, clone PC10, Cat. no. M0879). The specificity of this antibody for PCNA has been validated in many vertebrate species including zebrafish [61, 73].

**Microscopy**

Sections were observed with an epifluorescence microscope (Olympus Provis, equipped with a DP71 digital camera), an epifluorescence Zeiss (Imager Z1, equipped with the Apotome...
module) or a confocal microscope Leica SP2. Images were processed with either the Olympus (CellB) Zeiss (AxioVision4) or Leica (LCS Lite) software. Micrographs were acquired in TIFF format and adjusted for light and contrast before being assembled on plates using Photoshop CS4. The nomenclature for brain nuclei in adults and developing zebrafish were taken from Wullimann et al, and Mueller and Wullimann, respectively [74, 75].

Results

Bdnf mRNAs in the brain of 7 days old larvae

We first studied the presence of bdnf mRNAs at early developmental stages. To analyze in detail the distribution of bdnf transcripts in the brain of 7 days old larva, we performed in situ hybridization on series of transverse sections of the whole zebrafish brain. The bdnf sense riboprobe did not generate any signal (S1 Fig,[38]), confirming the specificity of the labeling obtained with the antisense riboprobe. The antisense riboprobe generated a positive labeling in several brain regions, demonstrating the presence of bdnf mRNAs. The Table 1 and Fig 1 summarize the results. From anterior to posterior, bdnf mRNAs were detected in the olfactory rosettes and in the anterior telencephalon (Fig 1A–1D). At this level, positive cells were mostly located in the dorsal and medial parts of the dorsal telencephalon. More caudally, labeled cells were still observed in the dorsal telencephalon, preferentially in the central telencephalon and also in the preoptic area (Fig 1E–1H). Bdnf mRNAs were particularly abundant in the thalamic area (Fig 1I and 1J) and were also consistently observed in the optic tectum, particularly in the periventricular layer (Fig 1G and 1H–1K and 1L). Bdnf mRNAs were also detected at different levels of the midbrain tegmentum, in particular in the reticular formation (Fig 2H–2J).

Attempts to identify the phenotype of these bdnf-expressing cells were made by using double staining with either PCNA, aromatase B or HuC/D as markers of cell proliferation, radial glial cells or neurons respectively [61, 76]. Data showed that, in each studied region, cells expressing bdnf mRNAs never corresponded to PCNA-positive cells indicating that bdnf-positive cells are post-mitotic (Fig 2A–2D). Bdnf-expressing cells did not co-express aromatase B attesting that they were not radial glia progenitors (data not shown). Regarding the neuronal marker HuC/D, although the coupling of immunohistochemistry with in situ hybridization resulted in some loss of bdnf transcripts, it appeared that in many regions such as the telencephalon, the optic tectum and the midbrain tegmentum, the signals generated overlapped,

Table 1. Bdnf mRNAs in juvenile zebrafish. + = few cells; ++ = moderate number of cells; +++ = numerous cells.

| Region                        | Bdnf mRNAs |
|-------------------------------|------------|
| Olfactofy bulb                | +++        |
| Pallium                       | ++         |
| Subpallium                    | +          |
| Preoptic region               | ++         |
| Habenula                      | +          |
| Dorsal thalamus               | +++        |
| Dorsal part of posterior tuberculum | +++    |
| Ventral part of posterior tuberculum | ++    |
| Intermediate hypothalamus     | ++         |
| Rostral hypothalamus          | ++         |
| Caudal hypothalamus           | +++        |
| Optic tectum                  | +++        |
| Tegmentum                     | +++        |

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suggesting that \textit{bdnf}-expressing cells may rapidly gained at least a neuronal phenotype (Fig 2E–2J).

\textbf{Bdnf mRNAs in the brain of adult zebrafish}

The distribution of \textit{bdnf} mRNAs was studied on transverse brain sections of sexually mature males and females from the olfactory bulbs to the medulla oblongata. No labeling was observed with the sense probe (S2 Fig, [38]). The distribution of \textit{bdnf} mRNAs, revealed by the in situ reaction with the antisense riboprobe, is shown in Table 2 and Figs 3 and 4.

The overall pattern of \textit{bdnf} expression was identical in all animals. In particular, \textit{bdnf} mRNAs were expressed in the same areas/regions in males and in females. There was no obvious difference between sexes, either in staining intensity and labeled-cells number. For this
reason, micrographs presented in Figs 3 and 4 include both male and female brains. The olfactory bulbs displayed few bdnf-positive cells in the glomerular cell layer and in the external and internal cell layers (data not shown). A strong hybridization staining was observed in a large number of small round cells localized in the dorsal telencephalon particularly in its medial, lateral and posterior divisions (Fig 3A–3C and 3E). The ventral part of the telencephalon exhibited fewer and weakly labeled cells in the posterior zone and intensely stained cells in the entopeduncular nucleus (Fig 3A and 3B and 3E). In the diencephalon, we observed an intense positive signal in the parvocellular (Fig 3D and 3E), and magnocellular nuclei of the preoptic area and in the entopeduncular and suprachiasmatic nuclei (Fig 3G). Bdnf mRNAs are strongly expressed in cells of the habenula, specifically in its dorsal component (Fig 3H–3I) and in the ventrolateral and ventromedial nuclei of the ventral thalamus (Figs 3F and 3F’ and 4F), but the intensity of labeling was weaker than in the dorsal thalamic nuclei (see below). Transcripts

![Fig 2. Immunohistochemical characterization of bdnf-expressing cells in the brain of 7 days old zebrafish larvae.](Image)

Double staining for bcdnf mRNA (red) and PCNA protein (green) on cross-sections through the telencephalon (A–B) and the thalamus (C–D). Double staining for bcdnf mRNA (red) and the neuronal marker Hu (green) in the thalamus (E,F and G), the optic tectum (H, I and J) and at the level of the superior reticular formation (H, I and J). In F and I cells nuclei are counterstained with DAPI. The dotted lines indicate the ventricles. OT: optic tectum; SRF: superior reticular formation; Tel: telencephalon; Thal: thalamus. Scale bar: 150 μm in H, I and J. 120 μm in A, B, C. 60 μm in D, E, F and G.

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Table 2. Expression of *bdnf* in the brain of adult zebrafish. + = few cells; ++ = moderate number of cells; +++ = numerous cells.

| Brain Region                        | Expression |
|------------------------------------|------------|
| **Olfactory bulbs**                |            |
| Glomerular layer of olfactory bulb | ++         |
| External cellular layer of olfactory bulb | +       |
| Internal cellular layer of olfactory bulb | +       |
| Lateral olfactory tract            | +         |
| Medial olfactory tract             | +         |
| **Dorsal telencephalic area**      |            |
| Lateral zone of dorsal telencephalic area | ++       |
| Medial zone of dorsal telencephalic area | +++      |
| Central zone of dorsal telencephalic area | +++     |
| Dorsal zone of dorsal telencephalic area | ++      |
| Posterior zone of dorsal telencephalic area | +++   |
| **Ventral telencephalic area**     |            |
| Entopeduncular nucleus, dorsal part | +       |
| Postcommissural nucleus of ventral telencephalic area | +       |
| **Preoptic area**                  |            |
| Parvocellular preoptic nucleus, anterior part | ++  |
| Magnocellular preoptic nucleus     | +         |
| Parvocellular preoptic nucleus, posterior part | +     |
| Subglomerular nucleus              | +         |
| **Epithalamus**                    |            |
| Dorsal habenular nucleus           | ++        |
| Ventral habenular nucleus          | ++        |
| **Dorsal thalamus**                |            |
| Anterior thalamic nucleus          | +         |
| Posterior zone of dorsal telencephalic area | +    |
| Central posterior thalamic nucleus | +         |
| **Ventral thalamus**               |            |
| Ventromedial thalamic nucleus      | +         |
| Ventrolateral thalamic nucleus     | ++        |
| **Posterior Tuberculum**           |            |
| Posterior tuberal nucleus          | +++       |
| Anterior preglomerular nucleus     | ++        |
| Lateral preglomerular nucleus      | ++        |
| Medial preglomerular nucleus       | +         |
| **Hypothalamus**                   |            |
| Diffuse nucleus of the inferior lobe | +       |
| Ventral zone of periventricular hypothalamus | ++ |
| Anterior tuberal nucleus           | +         |
| Lateral hypothalamic nucleus       | ++        |
| Central nucleus of the inferior lobe | ++     |
| Dorsal zone of periventricular hypothalamus | +   |
| Mamillary body                     | +         |
| **Synencephalon**                  |            |
| Nucleus of the medial longitudinal fascicle | +   |
| Periventricular pretectal nucleus, ventral part | ++ |
| Periventricular pretectal nucleus, dorsal part | +++ |
were also abundantly reported in the anterior, dorsal, posterior and central posterior nuclei of the dorsal thalamus (Fig 4B and 4B’ and 4G). More ventrally, bdnf mRNAs were expressed in the posterior tuberal nucleus (Fig 4A and 4A’, 4D, 4D’, 4F and 4G), in the diffuse and central nuclei of the inferior lobe and in the mammillary body. Between diencephalon and mesencephalon, in the so-called synencephalon, the nucleus of the medial longitudinal fascicle displayed few bdnf-positive cells, whereas bdnf mRNAs were highly expressed in the dorsal and ventral periventricular pretectal nuclei. In the mesencephalon, bdnf mRNAs were also observed in the torus longitudinalis. The periventricular gray zone of the optic tectum exhibited numerous cells expressing bdnf transcripts (Fig 4C and 4C’ and 4G). In the tegmentum, bdnf mRNA were observed in the central nucleus of the torus semicircularis and in the interpeduncular nucleus. In the medulla oblongata, bdnf mRNAs were highly expressed in the secondary gustatory nucleus and in few cells of the vagal lobe. Finally, large bdnf-positive perikarya were seen in the superior and inferior reticular formation (Fig 4E and 4E’ and 4H).

Bdnf-expressing cells do not proliferate and are not radial glial cells but neurons

In order to identify the nature of bdnf-expressing cells in the brain of adult zebrafish, we performed bdnf in situ hybridization in combination with immunohistochemistry using antibodies directed against different types of markers (proliferation, neuronal or glial markers), as described above. Bdnf-expressing cells were visualized close to the ventricular surface in many regions such as the telencephalon, the preoptic area, the hypothalamus, the thalamus, and the optic tectum. Previous studies showed that in the brain of adult zebrafish, many proliferating cells were located along the ventricles in those regions [61, 63]. To investigate if cells undergoing proliferation could be bdnf-expressing cells, we performed PCNA labeling in combination with bdnf in situ hybridization. As expected, PCNA-labeled cells were positioned along the ventricular cavities in the telencephalon (Fig 5A–5A’), in the thalamus (Fig 5B and 5B’) and in

Table 2. (Continued)

| Optic Tectum         |   |
|----------------------|---|
| Periventricular grey zone of optic tectum | ++ |
| Deep white zone      | + |
| Central zone of optic tectum | + |
| Superficial white zone | + |
| Longitudinal torus   | + |

| Torus semicircularis |   |
|----------------------|---|
| Central nucleus of semicircular torus | + |

| Tegmentum              |   |
|------------------------|---|
| Interpeduncular nucleus | + |
| Superior reticular formation | + |

| Cerebellum             |   |
|------------------------|---|
| Granular cell layer    | + |

| Medulla oblongata      |   |
|------------------------|---|
| Secondary gustatory nucleus | ++ |
| Inferior reticular formation | ++ |
| Vagal lobe             | + |
the hypothalamus (Fig 5B and 5B’). In those regions, bdnf-expressing cells were observed very close to PCNA-labeled nuclei, but bdnf mRNAs were never expressed by the proliferating cells (PCNA-positive cells).

**Fig 3.** *Bdnf* mRNA distribution in cross-sections through the adult zebrafish forebrain. Telencephalon (A, B, C, C’ and E), preoptic area (B, D, D’ and E, G), thalamus (F, F’ and I) and dorsal habenula (H, H’). In C’, D’, F’, H’ cell nuclei are labeled in blue with DAPI. E, G and I are representative sections taken from the zebrafish atlas (Wullimann et al., 1996). *Bdnf*-expressing cells are represented by red dots. Dc: central zone of the dorsal telencephalon; DiV: diencephalic ventricle; Di: lateral zone of the dorsal telencephalon; Dm: medial zone of the dorsal telencephalon; Dp: posterior zone of the dorsal telencephalon; EN: endopeduncular nucleus; Had: dorsal habenular nucleus; PM: magno cellular preoptic nucleus; OC: optic chiasma; POA: preoptic area; TelV: telencephalic ventricle; VL: ventrolateral thalamic nuclei. Scale bar: 120 μm except in C and C’: 60 μm.

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To test whether some of the bdnf-expressing cells could be radial glial cells, we combined bdnf hybridization with immunohistochemistry against aromatase B or BLBP, two well-
established markers of radial glial cells in fish and other vertebrates [61, 76–78]. As shown in Fig 6, aromatase B-positive cells never co-expressed bdnf mRNAs in the ventral telencephalon (A to A’'), the thalamus (B and B’) and the ventral hypothalamus (B and B’’). A and B are representative sections taken from the zebrafish atlas (Wullimann et al., 1996). Bdnf-expressing cells are represented by red dots and PCNA-labeled cells are green dots. DI: lateral zone of the dorsal telencephalon; Dm: medial zone of the dorsal telencephalon; Hyp: hypothalamus; TelV: telencephalic ventricle; Thal: thalamus; Vv: ventral zone of the ventral telencephalon. Scale bar = 200 μm in A’; 100 μm in A’; 50 μm in A’’.

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Discussion

In the present study, we reported, for the first time, that bdnf mRNAs are consistently expressed in the brain of larval and adult zebrafish. Our results also demonstrated that cells expressing bdnf mRNAs do not correspond to radial glial cells or progenitor cells but rather to neurons.
Fig 6. Immunohistochemical characterization of **bdnf**-expressing cells in adult zebrafish brain. A, B, C, D and E are representative sections taken from the zebrafish atlas (Wullimann et al., 1996). **Bdnf**-expressing cells are represented by red dots and Aromatase B (A-D) or BLBP-labeled cells (E) are represented by black dots with thin lines indicating radial glia cytoplasmic processes. Double staining for **bdnf** mRNA (red) and Aromatase B protein (green) on cross-sections through the telencephalon (A-A'), the preoptic area (B-B'), the entopedoncular nucleus (B-B''), the thalamus (C-C') and the ventral hypothalamus (D-D''). Double staining for **bdnf** mRNA (red) and BLBP protein (green) on cross-sections through the telencephalon (E-E''). DiV: diencephalic ventricle; Di: lateral zone of the dorsal telencephalon; Dm: medial zone of the dorsal telencephalon; ENv: endopedoncular...
The present study shows that *bdnf* mRNAs are widely expressed in the brain of 7 days old larvae, which is in agreement with the data previously obtained by RT-PCR and showing that *bdnf* expression is maternally inherited, drops down at 1 day post-fertilization (dpf) and then progressively increases from 1 to 8 dpf [38]. Our results bring new information by showing that the strongest signal is observed in the dorsal telencephalon (pallium), the preoptic region, the thalamus and the optic tectum. In general, our expression is in good agreement with what has been shown by previous authors [38]. Indeed, the distribution of *bdnf* mRNAs was addressed by in toto hybridization and established that *bdnf* was strongly expressed in the pallium, hypothalamus, posterior tuberculum and optic tectum at 1, 2 and 3 dpf. However, there were no detailed information on the precise structures expressing these messengers. In addition, at 7 dpf, *bdnf* mRNAs level seems to be lower, possibly due to a problem of access of the probe to the brain [38]. Our results show that in all cases, cells expressing *bdnf* mRNAs were located in the parenchyma and are identified as neurons. Furthermore, *bdnf*-expressing cells never exhibited PCNA, aromatase B or BLBP staining, indicating that *bdnf* mRNAs are not...
expressed in radial glia and/or progenitors. Although the function of BDNF in the developing brain of zebrafish was not addressed in the present study, these data suggest that this factor could be involved in the differentiation and maintenance of newborn neurons, mainly in the forebrain.

In adult zebrafish, *bdnf* expression has a similar distribution as in the larvae with the most prominent staining in dorsal telencephalon, preoptic area, dorsal thalamus, posterior tectum, hypothalamus, synencephalon and optic tectum. A more diffuse and weaker labelling is detected in other brain regions. The pattern of distribution of *bdnf* mRNAs is relatively similar to that reported in the European eel and the Turquoise killifish [35, 37] with some slight differences regarding the ventral telencephalon, diencephalon, tegmentum and rhombencephalon. Compared to the European eel and the turquoise killifish, *bdnf* expression in zebrafish is less expressed in the ventral telencephalon. By contrast, the staining in the diencephalon appears more widespread in the zebrafish than in the eel. Some slight differences were seen in other diencephalic nuclei, such as nucleus corticais and nucleus glomerulosus, which were intensely positive in the killifish and are lacking in the brain of zebrafish. Finally, the tegmentum and rhombencephalon of zebrafish showed fewer *bdnf*-expressing cells than eel and turquoise killifish. Thus, we can conclude that, in zebrafish, *bdnf* is expressed mostly in the forebrain and just in few areas of midbrain and hindbrain.

The localization of *bdnf* transcripts in adult zebrafish is similar in both sexes, despite the fact that sex hormones have been recognized to influence the expression of *bdnf* through a combination of genomic and epigenetic mechanisms [79, 80]. The general idea about the regulation of BDNF (protein and gene activation) by estradiol is that estradiol increases *bdnf* mRNAs levels. However, if we examine at the literature into more detail, the impacts of estrogens on *bdnf* mRNAs are complex and fluctuate according to the species, age, brain region and treatment [79–81]. Preliminary experiments performed in the laboratory and based on RT-PCR in whole brains failed to show any significant effect of estradiol exposure (10^-7M to 10^-10M) on *bdnf* expression in 7dpf zebrafish larvae.

Because in some regions, *bdnf*-expressing cells in adults were located close to the ventricular surface, we examined whether such cells could correspond to radial glia cells. Indeed, we and others have shown that in adult fish radial glial cells line entirely the ventricles [61, 70, 82]. Furthermore, it is now recognized that radial glial cells are progenitors and sustain the constant growth of the brain throughout life [61, 83]. To investigate if radial glial cells express or not *bdnf* transcripts, we used aromatase B and BLBP immunohistochemistry in combination with *bdnf* in situ hybridization [70, 71, 84]. Interestingly, we could not detect *bdnf* mRNAs/aromatase B or *bdnf* mRNAs/BLBP co-staining, suggesting that radial glial cells do not express *bdnf* under physiological conditions. However, in mammals, *bdnf* expression has been documented in glial cells under pathological conditions, in particular after brain lesions or around amyloid plaques [28, 85, 86]. We previously developed a model of telencephalic lesion in adult zebrafish and showed that proliferation increased rapidly after injury in the parenchyma of the injured telencephalon [68]. Thus, it would be interesting to investigate, in this model, if *bdnf* expression could be induced in glial cells after brain injuries (as in mammals). Double staining with the cell proliferation marker PCNA failed to show any co-expression of *bdnf* mRNAs and PCNA in the same cells. However, this does not rule out the possibility that they may express *bdnf* mRNAs later. A study performed in zebra finch showed that, 15 days after bromodeoxyuridine injection (BrdU, a marker for new born cells), BDNF protein is present in new generated cells labeled with BrdU [87]. In our study, double staining with neuronal markers, notably HuC/D, MAP2 and acetylated-tubulin clearly showed that *bdnf*-expressing cells have a neuronal phenotype in the brain of adult zebrafish. This finding is consistent with the fact that BDNF is a factor of neuronal origin in mammals [22, 24, 29]. The neuronal marker HuC/D specifically
characterizes early-differentiated neurons [88], whereas the markers MAP2 and acetylated-tubulin characterize both early and late differentiated neurons [89, 90]. Therefore, we can assume that bdnf mRNAs could be expressed in the early stages of differentiation processes.

It is interesting to note that the distribution of bdnf mRNAs in the brain of zebrafish is quite similar to that reported in mammals, notably in rat, mouse and pig [21–26]. Indeed, strong bdnf expression was reported in the cortex and hippocampus, two pallial structures. Because the telencephalon of fish develops by eversion, it is usually considered that the hippocampus equivalent in zebrafish is the dorsolateral region of the dorsal telencephalon [91–93], in which abundant bdnf mRNAs are detected. It is also noteworthy that the central part of the telencephalon, regarded as the presumptive equivalent of the isocortex of mammals, also strongly expresses bdnf [22]. Similarly, the preoptic area and in particular the magnocellular neurons express bdnf transcripts, similar to their mammalian counterparts, the paraventricular and supraoptic nuclei. Other structures exhibiting bdnf mRNAs in both fishes and mammals include the habenula, the thalamic region, the mediobasal hypothalamus and the optic tectum (inferior colliculus). These similarities suggest that BDNF functions are conserved between fishes and mammals and have emerged early in the vertebrate lineage. Studies in Drosophila have recently reported the existence of a neurotrophin-related gene (DNT1) [94]. According to this study, this gene would represent the ancestor of the neurotrophin family that diversified through whole genome duplications. Interestingly, in Drosophila, DNT1 exerts trophic and maintenance functions on neurons.

In conclusion, this study presents a detailed pattern of bdnf mRNAs distribution in the brain of larva and adult zebrafish under normal conditions. We show that bdnf transcripts are most abundant in forebrain regions (that are homolog to bdnf-expressing areas in the mammalian brain). We also bring evidences that bdnf mRNAs are expressed in neurons and not in radial glial cells or progenitor cells. As BDNF contributes to the organization and plasticity of neural network in the intact brain in mammals, further studies in fish should aim at investigating its potential roles in the developing and adult brain in physiological conditions, but also in the case of brain recovery after injuries.

Supporting Information

S1 Fig. The bdnf sense riboprobe did not generate any signal in the brain of 7 days old zebrafish. Telencephalon (A), Preoptic area (B), Optic Tectum (B and C). Hyp: hypothalamus; POA: preoptic area; OR: olfactory rosettes; OT: optic tectum. Teg: tegmentum; Tel: telencephalon. Scale bar: 120 μm in A and C. Scale bar: 60 μm in B. (TIF)

S2 Fig. The bdnf sense riboprobe did not generate any signal in the brain of adult zebrafish. Telencephalon (A and B), Preoptic area (C), Thalamus (D), Hypothalamus (E) and Optic Tectum (F). Dc: central zone of the dorsal telencephalon; DiV: diencephalic ventricle; Di: lateral zone of the dorsal telencephalon; Dm: medial zone of the dorsal telencephalon; Dp: posterior zone of the dorsal telencephalon; DP: dorsal thalamic nucleus; Hd: dorsal zone of the periventricular hypothalamus; Hv: ventral zone of the periventricular hypothalamus; OC: optic chiasma; OT: optic tectum; PGZ: periventricular gray zone of the optic tectum; POA: preoptic area. Scale bar: 120 μm in A, C. Scale bar: 60 μm in B, D, E, F. (TIF)

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**Author Contributions**

Conceived and designed the experiments: EP OK LDA CL. Performed the experiments: P. Cacialli MMG P. Coumailleau. Analyzed the data: P. Cacialli OK EP CL. Wrote the paper: EP P. Coumailleau OK CL.

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