Zebrafish telencephalon acquires an everted morphology by a two-step process that occurs from 1 to 5 days post-fertilization (dpf). Little is known about how this process affects the positioning of discrete telencephalic cell populations, hindering our understanding of how eversion impacts telencephalic structural organization. In this study, we characterize the neurochemistry, cycle state and morphology of an EGFP positive (+) cell population in the telencephalon of Et(gata2:EGFP)bi105 transgenic fish during eversion and up to 20dpf. We map the transgene insertion to the early-growth-response-gene-3 (egr3) locus and show that EGFP expression recapitulates endogenous egr3 expression throughout much of the pallial telencephalon. Using the gata2:EGFPbi105 transgene, in combination with other well-characterized transgenes and structural markers, we track the development of various cell populations in the zebrafish telencephalon as it undergoes the morphological changes underlying eversion. These datasets were registered to reference brains to form an atlas of telencephalic development at key stages of the eversion process (1dpf, 2dpf, and 5dpf) and compared to expression in adulthood. Finally, we registered gata2:EGFPbi105 expression to the Zebrafish Brain Browser 6dpf reference brain (ZBB, see Marquart et al., 2015, 2017; Tabor et al., 2019), to allow comparison of this expression pattern with anatomical data already in ZBB.

Keywords: telencephalon, eversion, telencephalon development, zebrafish, atlas, egr3

INTRODUCTION

The telencephalon, consisting of the olfactory bulbs and the telencephalic lobes, lies in the alar portion of the secondary prosencephalon (see Puelles and Rubenstein, 2003, 2015; Affaticati et al., 2015). It is responsible for higher brain functions, such as memory, emotions, cognition and higher-level multisensory integration and motor control. In zebrafish, as in all other ray-finned fish, the telencephalon is everted, a markedly different morphology to that of other vertebrates that have an evaginated telencephalon (Wullimann and Mueller, 2004b; Folgueira et al., 2012).
The morphological differences between everted and evaginated telencephali complicate comparisons and establishing homologies. In ray-finned fish, the telencephalon consists of two solid lobes separated by a midline ventricle that also extends over the dorsal surface under the tela choroidea (Butler, 2000; Yamamoto et al., 2007; Braford, 2009; Mueller and Wullimann, 2009; Nieuwenhuys, 2009a; Folgueira et al., 2012; Porter and Mueller, 2020). In other vertebrates, the telencephalon is evaginated and consists of two hollow hemispheres that surround an inflated, lateral ventricle. There is consensus that the dorsal and ventral portions of the everted telencephalon are homologous to the pallium and subpallium, respectively, but establishing further homologies has proven to be challenging, especially for the pallium (Wullimann and Mueller, 2004b; Northcutt, 2008; Nieuwenhuys, 2009a; Mueller et al., 2011; Ganz et al., 2012; Porter and Mueller, 2020). This is because eversion is more complex than just a simple lateral out-folding of the neural tube (Yamamoto et al., 2007; Mueller et al., 2011; Folgueira et al., 2012; Porter and Mueller, 2020), as first proposed by Gage (1893) and Studnička (1894, 1896). For instance, based on the analysis of early forebrain development, we showed that there are two early morphogenetic events critical to formation of an everted telencephalon in zebrafish (Folgueira et al., 2012). The first event is the formation of the ventricle, including the formation of the anterior intracerebral sulcus (AIS). The second event is the rostrocaudal expansion of the pallium. During this expansion the roof of the AIS gets stretched to form the tela choroidea and the ventricular zone reaches the dorsal surface of the telencephalon (see Folgueira et al., 2012 for more details). Although our model showed a more complex process of eversion than classically proposed, its implications are yet to be determined by detailed analysis of the developmental changes in locations of discrete cell populations during eversion (Briscoe and Ragsdale, 2019).

After the early developmental events that generate an everted telencephalon, other processes, such as cell differentiation and migration, will certainly contribute to generate the organization of pallial subdivisions observed in the adult (Yamamoto et al., 2007; Mueller et al., 2011; Porter and Mueller, 2020).

**MATERIALS AND METHODS**

**Fish Stocks and Maintenance**

Adult zebrafish (*Danio rerio*, Cyprinidae) were maintained under standard conditions that meet FELASA guidelines (Aleström et al., 2020): 28°C and 14 h light/10 h dark periods (Westerfield, 2000) at University College London (UCL) Fish Facility. Embryos were raised in fish water at 28°C and staged according to Kimmel et al. (1995). Phenylthiocarbamide (PTU, Sigma) at a concentration of 0.003% w/v was added to the fish water at 24 h post fertilization (hpf) to prevent pigment formation in larvae. All experimental procedures were conducted under license from the United Kingdom Home Office, following United Kingdom Home Office regulations and/or European Community Guidelines on animal care and experimentation, and were approved by animal care and use committees.

The following zebrafish strains were used in this study: AB and TL wild types, Et(gata2:EGFP)1105 (Folgueira et al., 2012), Tg(1.4dlx5a-dlx6a:GFP)1127 (Zerucha et al., 2000), Tg(isl1:GFP)1110 (Higashijima et al., 2000) and Tg(-10lx2a:EGFP)1176 (Miyasaka et al., 2009, 2014).

**BrdU Staining**

BrdU (5-bromo-2’-deoxyuridine) incorporation assay was performed on 4dpf larvae by first embedding fish in 1% low-melting-point agarose and then injecting approximately 1 nl of 10 mM BrdU (Sigma) into the heart. Larvae were then removed from the agarose and allowed to freely swim for 2 h at 28.5°C. After anesthesia and fixation, larvae were processed for immunohistochemistry (see below).

**Anesthesia and Fixation**

Specimens were deeply anesthetized in 0.2% tricaine methanesulphonate (MS222, Sigma) in fresh water and fixed in 4% paraformaldehyde (PFA) in phosphate buffered saline (PBS) with 4% sucrose by immersion. The tissue was then post-fixed in the same fixative for 24 h at room temperature. After this, cranial skin, jaw and other tissues were removed from embryos and larvae by fine dissection using two pairs of sharp forceps on embryos pinned in Sylgard (Turner et al., 2014). After
washing in PBS and dehydration in methanol, they were kept at -20°C for at least 24 h. Brains from 20dpf fish and adults were dissected and kept in PBS at 4°C until use.

**Immunostaining**

Embryos and larvae were stained as whole mounts following standard procedures (Shanmugalingam et al., 2000; Turner et al., 2014). In brief, specimens were rehydrated to phosphate buffered saline with 0.5% Triton X-100 (PBS-T). After a brief proteinase digestion to improve permeability of the tissue, specimens were incubated with primary antibody overnight at 4°C. Specimens were then washed in PBS-T and incubated with fluorescent secondary antibodies again overnight at 4°C. After washing in PBS-T, embryos and larvae were mounted in 2% low-melting point agarose and imaged on a Leica confocal laser scanning microscope. For a detailed protocol on larval dissection, antibody labeling and mounting for imaging see Turner et al. (2014).

Brains from 20dpf fish were cryoprotected, frozen with liquid nitrogen-cooled methylbutane and cut in sections (12 μm thick) on a cryostat. After mounting the sections on gelatinized slides, they were rinsed in PBS-T and preincubated with normal goat serum (Sigma, 1:10) for 1 h. Next, they were incubated with a primary antibody overnight at room temperature and then with secondary fluorescent antibodies for 1 h, followed by PBS washes. Sections then stained for 5 min in a solution of Sytox Orange to detect nuclei and distribution of different nuclei. Following antibody labeling, larval brains were washed in PBS and then stained for 30 min in a solution of Sytox Orange diluted in PBS-T (dilution 1:20,000). Subsequently, brains were washed in PBS and mounted for imaging.

**Primary Antibodies**

Rabbit anti-green fluorescent protein (GFP; Torrey Pines Biolabs, Cat# TP401, dilution 1:1,000), rat anti-GFP (Nacalai Tesque; Cat# GF090R, dilution 1:1,000), mouse anti-bromodeoxyuridine (BrdU; Roche; Cat# B8434, dilution 1:300), mouse anti zona occludens 1 (ZO1, Invitrogen; Cat# 33-9100, dilution 1:300), mouse anti-acetylated tubulin antibody (IgG2b; α-tubulin; Sigma; Cat# T7451, dilution 1:250), rabbit anti-γ-aminobutyric acid (GABA; Sigma; Cat# A2052, dilution 1:1,000), mouse anti-synaptic vesicle protein 2 (IgG1; SV2; DSHB; Cat# AB 2315387, dilution 1:250), rabbit anti-DSRed antibody (Living Colors; Clontech; Cat# 632496, dilution 1:300), rabbit anti-red fluorescent protein (RFP, MBL; Cat#PM005; dilution 1:2,000).

**Secondary Antibodies**

Alexa Fluor 488 (Invitrogen, Goat anti-Rabbit, Cat# A-11034, Goat anti-Rat Cat# A-11006, Goat anti-Mouse, Cat# A-11029, dilution all 1:200), Alexa Fluor 568 (Invitrogen, Goat anti-Mouse IgG, Cat# A-11031, Goat anti-Mouse IgG2b, Cat# A-21144, dilution all 1:200) and Alexa Fluor 633 (Invitrogen, Goat anti-Mouse IgG, Cat# A-21052; Goat anti-Mouse IgG1, Cat# A-21126, dilution all 1:200). To detect anti-acetylated tubulin and anti-SV2 in the same sample, isotype-specific secondary antibodies were used: Alexa Fluor 568 (IgG2b, Cat# A-21144, dilution 1:200) and Alexa Fluor 633 (IgG1, Cat# A-21126, dilution 1:200).

**Whole Mount Fluorescent in situ Hybridisation**

The FISH protocol was adapted from Jülich et al. (2005) and performed as described in Turner et al. (2016). Digoxigenin probes were made by standard protocols and were detected using the anti-DIG POD antibody (Roche, 1:1,000) and stained using Cy3-tetramethyl substrate (Perkin Elmer, 1:50 in amplification buffer). After staining, antibody labeling for GFP was performed as above using rabbit anti-green fluorescent protein (GFP, Torrey Pines Biolabs, dilution 1:1,000) and goat anti-rabbit Alexa Fluor 488 (Invitrogen, dilution 1:200) antibodies without further PK digestion. Cell nuclei were labeled with TOPRO-3-iodide (Invitrogen, dilution 1:5,000). Embryos were mounted in 1% agarose in 80% glycerol/PBS solution and imaged on a Leica SP8 confocal microscope.

**Fluorescent Nucleic Acid Staining**

Larval brains were whole mount stained with Sytox Orange (S11368; ThermoFisher) and used for studying cell architecture and distribution of different nuclei. Following antibody labeling, larval brains were washed in PBS and then stained for 30 min in a solution of Sytox Orange diluted in PBS-T (dilution 1:20,000). Subsequently, brains were washed in PBS and mounted for imaging.

**Genome Walker Mapping of Enhancer Trap Insertion**

The genomic location of the insertion in the enhancer-trap line Et(gata2;EGFP)bi105 was mapped using a linker-mediated approach with the Universal Genome Walker TM 2.0 Kit (Clontech; see online for user manual). The Et(gata2;EGFP)bi105 line was made using the Tol2-transposase system for genomic insertion (Kawakami et al., 2000; Kawakami, 2004). Using this information, nested primers specific to the 5’ and 3’ ends of the tol2 sequence were designed: one for primary PCR (GSP1) and one for secondary nested PCR (GSP2) (sequences below). These primers have minimal overlap (5’ GSP1 had 16bp overlap with 5’ GSP2, but 3’ primers had no overlap), annealing to sequences as close to the end of the known sequence as possible. A combination of these primers (sequences below) and the ligated linkers/primer from the kit were used to generate four Genome Walker PCR amplicons from two restriction-generated genomic libraries. These amplicons were purified (Qiagen QIAquick PCR purification kit) and checked for purity and concentration using a NanoDrop 2000C (ThermoScientific). Sanger-sequencing (Source BioScience, United Kingdom) using either stock primers such as M13F (within linker annealed kit constructs) or gene specific primers was checked for novel genomic sequence adjacent to the tol2 sequence in each fragment, these sequences were aligned by BLAST/BLAT against the zebrafish genome. All four amplicons aligned to the same region of chromosome 8 (Figure 1A). The insertion site was confirmed
using PCR with primers specific to the specific EGFP employed in the transgenesis construct and the newly sequenced adjacent genomic sequence.

Nested sets of primers used with the adapter linked primers. GTCAGGTGGTGGATTGGTAGATAA, tol2_3′_GSP1 GGACCAATGAAATGCTGACCAATT, tol2_3′_GSP2 TACAACCTTGTGCATGCTGACTGGATT, tol2_5′_GSP1 GCGTGTACTGGCATTAGATTGTCTGCTT, tol2_5′_GSP2

### Genetic Linkage Analysis

To confirm linkage between transgene and genome hits derived from Genome Walker use we employed traditional linkage mapping using simple sequence length polymorphisms (SSLPs) (Knapik et al., 1998). Briefly, nearby SSLPs were identified from zebrafish information network (ZFlin; zfin.org) webpages and Ensembl. A small panel of 48 transgene-positive, and 48 transgene-negative embryos were arrayed on a 96 well plate and genominc DNA was extracted by proteinase K digestion. PCRs for SSLPs were carried out on this arrayed DNA followed by gel electrophoresis. Where nearby SSLP markers were polymorphic between positive and negative groups, linkage was tested by measuring differential meiotic recombination between transgene positive and negative groups. The SSLP markers and primer sequences employed are listed below.

| Marker name | F primer | R primer | Genbank ref |
|-------------|----------|----------|-------------|
| z2979       | TGGCTAAATTGCA | TCTGGGCTTAAAATGGC | G40762       |
| z21390     | CAAAAAACAGAACGAGG | GGGAGATGTGCAAGA | G40246       |
| z23039     | CCCTACTGTTGGCAGG | GAAGCTGAGGAATAGCAGGA | G39541       |

### Preparation of sgRNAs, Cas9 mRNA

Template DNA for short guide RNA (sgRNAs) synthesis was digested with DraI, and sgRNAs were transcribed using the MAXIscript T7 kit (Life Technologies). The pT3TS-nCas9 plasmid (Addgene) (Jao et al., 2013) was linearized with XbaI (Promega) and mRNA synthesized with the mMessage mMachine T3 Transcription Kit (Ambion). Transcription reactions were incubated in a water bath at 37°C for 2 h or longer. To digest template DNA, 1 μL of TURBO DNase was added and reaction incubated for a further 15 min at 37°C. The Cas9 mRNA was polyadenylated using the polyA tailing kit (Ambion). Cas9 mRNA and sgRNAs were purified using either the RNeasy Minikit (Qiagen) or Zymo columns.

### Microinjections

Microinjection of DNA and RNA was performed using a borosilicate glass capillary needle attached to a Picospritzer III injector. Adult zebrafish were paired with dividers the night before injections. Dividers were removed the following morning and embryos were collected soon after to ensure they were at early 1 cell stage at time of injections. Embryos were aligned against a glass slide inside an upturned petri dish lid. The needle was calibrated to inject 1 nl per embryo. Injections were performed into the cell for DNA and RNA.

### Single Cell Labeling Using EGFP to Gal4 Switching With CRISPR/Cas9

To achieve somatic switching of transgenes and mosaically label single or small groups of neurons within the Et(gata2:EGFP)bii05 transgenic line we adapted a technique developed by Auer et al. (2014) using the modified donor plasmid hs:Gbait (Kimura et al., 2014). This CRISPR/Cas9-mediated knock-in of DNA cassettes into the zebrafish genome uses homology-independent double-strand break repair pathways. Co-injection of a donor plasmid with a sgRNA and Cas9 nuclease mRNA results in concurrent cleavage of the donor plasmid and the selected chromosomal integration site resulting in the targeted integration of donor DNA. Using an EGFP target sequence in the donor plasmid and a sgRNA against EGFP Auer et al. (2014) effectively converted EGFP transgenic lines into Gal4 versions. Kimura et al. (2014) modified the donor plasmid (hs:Gbait) to include a heat-shock promoter to increase the efficiency of the switching of EGFP to Gal4, as the Gal4 cassette would be expressed regardless of which orientation the hs:Gbait integrates into the genome.

To achieve sparse labeling of single EGFP expressing neurons in Et(gata2:EGFP)bii05 injected embryos we added 5xUASTdTom DNA to the injection mixture. This led to very mosaic labeling with TdTom of neurons within the original EGFP expression pattern. All plasmids, including hs:Gbait (Kimura et al., 2014), were a kind gift from Thomas Auer. For the acute substitution, each embryo was injected with a solution containing 60 pg of sgRNA EGFP1, 150 pg of Cas9 mRNA, 7 pg of Gbait-hsp70:Gal4 and 5 pg UAS:TdTom (Zhang et al., 2012). Embryos were screened for RFP expression from 24hpf onward. Specimens with desired expression were fixed and processed for immunohistochemistry. Technical limitations of this labeling technique, due to genome editing occurring early in development, made it challenging to label later born neurons.

### Imaging

Fish were mounted in 2% low melting point agarose, either in lateral or dorsal view, under a dissecting scope equipped with bright field and fluorescent filters. Imaging was performed using either Leica SP2 or SP8 Confocal Microscopes equipped with filter-free AOBS systems tuned to the appropriate GFP/RFP spectra, with either water 25x (NA: 0.95) or 40x (NA: 0.8) immersion lenses and 3x line averaging. Imaging parameters varied for different data sets and scale references are included on figures to account for these differences. Due to the opacity of the tissue, imaging beyond a total depth of 200–300 μm from the surface of the fish brain was not possible.

### Image Analysis and Editing

Confocal stacks were processed using Fiji (ImageJ) and/or Volocity and/or Imaris. In some z-stacks, a FIRE Temporal-Color
FIGURE 1 | Et\textit{(gata2:EGFP)}

\textsuperscript{205} insertion is located near \textit{egr3} and expression of \textit{egr3} matches transgene expression. (A,B) Mapping of \textit{Et(gata2:EGFP)}
\textsuperscript{205} insertion showing the position of Genome Walker BLAST hits on assembly GRCz11, relative position of neighboring genes (black lines) and linkage markers with meiotic recombination data (A), the lower blue line in panel (A) indicates the region around \textit{egr3} shown in panel (B), with BLAST hits (insertion) locations. BLAST/BLAT coordinates: 8:50720195-50720677 (C–F′) Immunostaining and \textit{in situ} hybridization showing EGFP and \textit{egr3} mRNA co-expression in the telencephalon and optic tectum of \textit{Et(gata2:EGFP)}
\textsuperscript{205} fish at 48 hpf (C–D′) and 6 dpf (E–F′) in frontal view. Panels (C–C′, E–E′) show z-projections and panels (D–D′, F–F′) show single z-slices. Note the co-expression of EGFP and \textit{egr3} in pallial (arrow) and subpallial (arrowheads) cells. Scale bars: 50 µm.
RESULTS

The Et(gata2:EGFP)\textsuperscript{bi105} Transgene Neighbors Early Growth Response 3 and EGFP Expression Closely Matches Endogenous egr3 Expression

EGFP is expressed in the telencephalon of Et(gata2:EGFP)\textsuperscript{bi105} larvae and is a useful marker to understand telencephalic morphogenesis (Folgueira et al., 2012; Turner et al., 2016). The genomic location of this transgene was, however, not known.

To identify this locus, we mapped the position of the Et(gata2:EGFP)\textsuperscript{bi105} insertion using linker-mediated PCR, confirmed by genetic linkage mapping. Our results showed that the genomic insertion of the transgene in Et(gata2:EGFP)\textsuperscript{bi105} fish lies 6,543 bp upstream of the first exon of Early Growth Response 3 (egr3) (see methods and Figures 1A,B) which encodes a transcription factor that functions as an immediate-early growth response gene in some systems (Yamagata et al., 1994).

With the insertion point established, we then assessed whether Et(gata2:EGFP)\textsuperscript{bi105} transgene expression recapitulates egr3 expression. We examined egr3 mRNA expression using fluorescent in-situ hybridisation in combination with immunohistochemistry for EGFP in Et(gata2:EGFP)\textsuperscript{bi105} embryos. There was extensive co-localisation of EGFP protein and egr3 mRNA in the telencephalon from 48hpf (Figures 1C–D'). At this stage, there is also strong egr3 mRNA expression in the midline of the optic tectum, but little EGFP protein observed in this location (Figures 1C–D'). By 6dpf, egr3 mRNA expression overlaps with EGFP protein in the pallium, subpallium and optic tectum of Et(gata2:EGFP)\textsuperscript{bi105} embryos (Figures 1E–F').

In summary, this demonstrates that the Et(gata2:EGFP)\textsuperscript{bi105} transgene insertion maps near egr3, and EGFP expression is nested within the endogenous egr3 expression.

gata2:EGFP Expression in 3dpf and 20dpf Larval Stages

We first characterize the expression of Et(gata2:EGFP)\textsuperscript{bi105} in the 4 and 20 dpf larval zebrafish brain, which can be used as a useful reference tool for developmental studies of various brain areas in this species. Given the scarcity of transgenic lines with well-characterized expression in the telencephalon and the difficulty in characterizing the development of distinct telencephalic domains during the process of eversion, we next focus our study on this region (see below).

3dpf Zebrafish

To characterize gata2:EGFP\textsuperscript{bi105} expression, we used anti-GFP immunocytochemistry combined with anti-SV2 counterstaining to highlight anatomical landmarks (Turner et al., 2014). Rostrally, EGFP expression was observed in cell bodies in the pallium (Figures 2A–C'; see also Folgueira et al. (2012)) and a small cluster in the subpallium, just ventral to the olfactory bulb (Figures 2A–B'). In the secondary prosencephalon, outside the telencephalon, we found EGFP expression in the lateral hypothalamus (Figures 2A–B'). In the diencephalon, we
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FIGURE 2 | Etgata2:EGFP<sup>Pr105</sup> line EGFP expression at 3 dpf. Lateral (A,A'), ventral (B,B') and dorsal (C,C') views of a 4 dpf Etgata2:EGFP<sup>Pr105</sup> larvae labeled with anti-EGFP (green) and anti-SV2 antibodies (magenta). Major areas of EGFP expression are annotated in the schematics on the right (A'',B'',C''), these include: pallium, subpallial nucleus, pretectum, hypothalamic areas, optic tectum and hindbrain areas, among others. Scale bars: 100 µm.

| Lateral | Ventral | Dorsal |
|---------|---------|--------|
| ![A, A'] 3dpf | ![B, B'] 3dpf | ![C, C'] 3dpf |

observed a few neurons in the left-sided parapineal organ (Turner et al., 2016 and Figures 2C–C')). In the midbrain, scattered EGFP+ cell bodies in the periventricular gray zone of the optic tectum extended dendrites into the superficial tectal neuropil (Figures 2A–C'). Based on location and morphology, these EGFP+ neurons could represent periventricular interneurons [Robles et al., 2011; type XIV cells of Meek and Schellart (1978)/small periventricular cells of Vanegas et al. (1974)]. In the hindbrain, several EGFP+ cell bodies were present in the superior raphe, areas of the reticular formation (Figures 2B–B') and caudally in areas of the medulla oblongata (Figures 2B–B'). In all regions, expression appeared to be predominantly in neurons rather than ventricular cells/glia.

Despite the EGFP being a cytoplasmic variant, transgene expression (detected by immunohistochemistry) labeled several tracts and neuropil domains (Figure 2B). The superficial tectal neuropil was EGFP+ (Figures 2A–A',C–C'); most of these fibers appeared to originate from cell bodies located in the periventricular gray zone, however, some could have other origins. The horizontal commissure in the hypothalamus also contained EGFP+ fibers (Figures 2B–B'). Finally, at caudal rhombencephalic levels, EGFP+ fibers were present in the medial (mlf) and lateral (llf) longitudinal fascicles.

20dpf Zebrafish

As the anatomical divisions between regions are clearer by 20dpf, we could more precisely determine the identity of the different EGFP+ populations at this stage. EGFP expression was more scattered at 20dpf (Figure 3) than 3dpf (Figure 2), but expression areas were consistent with those at 3dpf. At 20dpf, there was more mosaicism in expression within...
and between individuals, suggesting transgene silencing over time. This mosaicism was even more prevalent in adult transgenic fish (not shown). The only regions with de novo expression observed were in the cerebellum and vagal lobe (Figures 3K,L).

Rostrally, many EGFP+ fibers (but no cell bodies) were present in the internal cell layer of the olfactory bulbs (Figures 3A,B). In the telencephalic lobes, we observed that EGFP expression was not consistently strong throughout the dorsal telencephalon/pallium (Figures 3B–E). Following
nomenclature of Castro et al. (2006; see also Yáñez et al. (2021)), the principal areas of EGFP expression in the pallium were the medial zone (Dma and Dm of Castro et al., 2006), lateral zone (Dl) (Figures 3B–E), central zone (Dc) (Figures 3C,D) and scattered cells in the dorsal zone (Dd) (Figures 3C–E). In contrast to 3dpf larval expression (apparently in the whole pallium, excluding pallial derived olfactory bulb cells), we found very few EGFP+ cells in the anterior region of the lateral zone (Dla) and in the posterior zone (Dp) (Figures 3B,E). EGFP expression was also excluded from the pallial ventricular zone (Figures 3B–E). This suggests that EGFP expression is excluded from radial glia and proliferative cells which occupy these areas. A few EGFP+ cells were found in the lateral subpallium and close to the lateral forebrain bundle (lfb) (Figure 3C, inset i). Based on this location, these subpallial cells could be part of the lateral nucleus of the ventral telencephalon (Vl; see Mueller et al., 2008; Mueller and Guo, 2009; Ganz et al., 2012).

In the basal secondary prosencephalon, we observed EGFP+ cells in the inferior hypothalamic lobes (Figures 3G–J). In the alar diencephalon, we observed that the parapineal (ppn) was labeled, as previously described in Turner et al. (2016) at 1mpf (Figure 3F and inset ii). EGFP+ fibers from the left-sided parapineal innervate the left dorsal habenula. At the same level as the habenulae, a small group of EGFP+ pretectal cells were present just above the forebrain bundle (Figure 3F). In the basal diencephalon, EGFP+ cells were found in the pregglomerular complex, torus lateralis, scattered cells in anterior and posterior tuberal regions (Figures 3G–J).

In the midbrain and hindbrain there were only a few areas of expression. Transverse sections at the level of the optic tectum showed EGFP+ cell bodies in the PGZ and in the stratum album central of the optic tectum (Figures 3G–I) that extended processes, most probably dendrites, dorsally into the stratum fibrosum griseum superficially and stratum opticum, consistent with 4dpf (see above). Also consistent with 4dpf, in the hindbrain, we observed EGFP+ neurons in the superior and intermediate reticular formation (Figure 3K) and superior raphe (Figure 3J and inset iii). In addition, we observed faintly EGFP+ cells in the valvula and corpus cerebelli (Figure 3K and inset iv), likely to represent eurydendroid and/or Purkinje cells, and EGFP+ cells in the vgal lobe (Figure 3L). These last two populations are additional to those observed in the 4dpf larva.

In summary, our analyses show EGFP expression in Et(gata2:EGFP)bi105 transgenic fish is restricted to specific areas in the brain that are broadly consistent between 3dpf and 20dpf.

Changes in Telencephalic Topography Between 18hpf and 20dpf

We used whole mount immunohistochemistry against EGFP in Et(gata2:EGFP)bi105 fish from 18hpf to 20dpf to follow the development of pallial and subpallial telencephalic populations, combined with anti-acetylated tubulin to highlight the progressive development of major axon pathways (Figures 4A–F; Wilson et al., 1990; Chitnis and Kuwada, 1990). Following EGFP expression over time does not allow us to definitively identify the same cells at each stage, but our observations strongly support there being stable transgene expression within the same discrete telencephalic cell populations over time.

By 20hpf, the first few EGFP+ cells were evident ventrally in the telencephalon (not shown), dorsal to the anterior commissure and forebrain bundle (telencephalic tract of Chitnis and Kuwada, 1990; supraoptic tract of Wilson et al., 1990). By 24 hpf, EGFP+ cells are divided in two populations (Figure 4A,A′): “population A” is located just rostral to the anterior commissure (subpallial nucleus, see below) and “population B” is located more caudally and dorsally (pallial population, see below). At this stage, EGFP+ axons and growth comes from these neurons were observed within the forebrain bundle (Figure 4A).

By 36hpf, an increase in the number of EGFP+ cells was evident in both populations (Figures 4A–B′), especially in “population B” (pallial population). The cells of “population A” (subpallial population) extended processes to the dorsal part of the anterior commissure (Figure 4B inset), where they intermingled with processes from the EGFP+ cells of “population B”. Some fibers extended along the forebrain bundle toward caudal forebrain areas (Figure 4B inset).

From 36hpf to 3dpf (Figures 4B–C′) the number of EGFP+ cell bodies again increased in population B (pallial + population). By 3dpf, a few EGFP+ fibers (but no cell bodies) were observed in the inner layer of the olfactory bulb. From 3dpf to 5dpf, the number of EGFP+ pallial cell bodies in the telencephalon increased substantially, as did the overall size of the pallial territory (Figures 4C–D′).

From 5dpf to 20dpf, we observed a dramatic increase in the size of the telencephalic lobes relative to other neighboring areas (Figures 4D–F′), such as the olfactory bulb and habenula. Although the EGFP+ subpallial cells were initially positioned rostral to the EGFP+ pallial cells, by 20dpf they sat ventral to this cell population (Figures 4E,F). Note that this change affects topography of the telencephalic populations, not topology (subpallial cells are always located anteriorly and pallial cells posteriorly). Large numbers of EGFP+ fibers were observed coursing toward caudal areas via the forebrain bundle (Figure 4F) and crossing the midline in the anterior commissure (Figure 4F). At this stage, a number of fibers originating from the subpallial population can be observed in the anterior commissure (Figures 4E,F).

The changes in the relative positions and sizes of the subpallial and pallial EGFP+ populations over time (from 24hpf to 20dpf) reflect the morphological changes that occur as a result of eversion (24hpf to 5dpf) and telencephalic growth (5dpf to 20dpf) (Folgueira et al., 2012; Dirian et al., 2014; Furlan et al., 2017).

To confirm the regional identity of the EGFP+ telencephalic cells at early stages, Et(gata2:EGFP)bi105 embryos were labeled with immunohistochemistry against EGFP and fluorescent in situ hybridisation against dixl1a and tbr1 at 36 hpf (MacDonald et al., 2010; Figure 5). tbr1 encodes a transcription factor that marks developing glutamatergic cells (Puelles et al., 2000; Hevner et al., 2001; Englund et al., 2005; Hevner, 2006). In zebrafish, tbr1 is mainly expressed in the pallium, with two smaller additional expression sites in the septal areas and...
Figure 4 | Time series showing $E\text{tgata}2:EGFP^{bi105}$ expression. Top box: sagittal view of a 3dpf zebrafish larval brain illustrating the body axis coordinates ("rostral," "caudal," "dorsal," and "ventral") predominantly used in this paper (adapted from Herget et al., 2014). For reference, the bent neuraxis of the neural tube is also shown (see Herget et al., 2014) with the alar/basal subdivision of the brain defined according to the neuromeric model (Puelles and Rubenstein, 2003, 2015). Lateral (A–E) and ventral (F) views of zebrafish brains stained against EGFP only inset in (B,F) or EGFP in combination with acetylated tubulin (A–E) from 24 hpf to 20 dpf. Arrow points to the ventral EGFP+ population identified as a subpallial nucleus. Arrowheads in panels (A,B) inset point to processes exiting the telencephalon along the forebrain bundle toward caudal areas. Arrow in panel (B) inset points to processes extending to the dorsal part of the anterior commissure. (A′–F′) Schematics showing the changes in location of the EGFP+ populations A and B relative to telencephalic regions and tracts over time. Rostral to the left. Scale bars: (A–D) 50 µm; (E,F). Axis coordinates abbreviations: A, anterior; C, caudal; D, dorsal; P, posterior; R, rostral; V, ventral. 100 µm.
in telencephalic populations that derive from the prethalamic eminence (Mione et al., 2001; Wullimann and Mueller, 2004a,b; Mueller et al., 2008; Turner et al., 2016). *dlx1a* is expressed in proliferative and immature neurons of the subpallium and preoptic area (Wullimann and Mueller, 2004a,b; MacDonald et al., 2010). Using these markers we found that for “population B”, EGFP+ cells were nested within the *tbr1*+ domain (Figures 5A–A′′,E–E′′) and mostly co-express EGFP+ and *tbr1*, with the exception of cells located ventral in the domain. “Population B” cells were predominantly *dlx1a* negative (-) (Figures 5B,B′,F,F′). These observations confirmed the pallial character of this population. The transgene does not label the full set of pallial cells, as there are cells that are *tbr1*+ but EGFP-, including the ventricular zone (Figures 5A–A′′,E–E′′). Single confocal slices through 36hpf larvae showed the EGFP+ cells of “population A” located at the border of the *tbr1*+ pallial domain (Figures 5C–C′′,G–G′′), nested within the subpallial marker *dlx1a* (Figure 5D–D′,H–H′′). The location of “population A” in relationship to *tbr1* and *dlx1a* domains confirms its subpallial identity.

In addition to our analysis of *tbr1* at 36hpf, we also analyzed *tbr2a* by 24 and 48hpf. *Tbr2a* is a broader marker for the pallium and septal cells in the subpallium (Mueller et al., 2008; Figure 6). At 24hpf the EGFP+ pallial group of cells, “population B”, showed overlap with *tbr2a* similar to *tbr1*. Again, there is a ventral intermediate group of cells that did not overlap with the *tbr2a* expression. Subpallial cells “population A” lie directly adjacent to the *tbr2a* expression domain but do not express *tbr2a*. By 48hpf the pallial and subpallial groups have moved further apart. The ventral intermediate *tbr2a*+ population observed by 24hpf within “population B” (pallial domain) has now resolved into a distinct pallial (*tbr2a*+) cell group.

**Et(gata2:EGFPb105)** Drives EGFP Expression in Post-mitotic Glutamatergic Telencephalic Neurons

The position of EGFP+ cells distant from the ventricular zone (Figures 7A–C), suggested that they are post-mitotic neurons. To ascertain if this is correct, we treated Et(gata2:EGFP)b105 larvae with short pulses of BrdU at 4dpf and then performed immunostaining against EGFP and BrdU. The gata2:EGFP+ cells did not incorporate BrdU and, consequently, they are most probably non-proliferative, post-mitotic neurons (Figures 7D,D′).

To further characterize the EGFP+ cells in the telencephalon, we assessed if they are glutamatergic and/or GABAergic. Co-immunostaining against GFP and GABA at 4dpf showed that...
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**FIGURE 6 | Etgata2:EGFP<sup>bi105</sup> line expression in combination with tbr2a.** Lateral view of Etgata2:EGFP<sup>bi105</sup> fish labeled with anti-GFP (green) and FISH for tbr2a (magenta) at 24hpf (A–C') and 48hpf (D–D''). Rostral to the left. Images are projections for z-stacks (A–A'', D–F') or single-confocal z-slices (B, B', C, C', E, E', F, F').

(A', D'') Blue whisker box contains EGFP+ pallial cells; orange whisker box contains subpallial cells. (B, B', E, E') Sagittal section through pallial cells. (C, C', F, F') Sagittal section through subpallial cells. Arrows indicate pallial EGFP+ cells that express tbr2a. Arrowheads indicate subpallial EGFP+ cells that do not express tbr2a. Yellow arrows indicate EGFP+ cells in the ventral pallial domain (population B) that do not express tbr2a at 24hpf. By 48hpf the intermediate cells (yellow arrow) have resolved into the two distinct pallial/tbr2a+ and subpallial/tbr2a- populations. Dotted white line indicates the telencephalic ventricle. Scale bars: 50 µm.

| 24hpf  | 48hpf  |
|--------|--------|
| z-projection | z-projection |
| single z-slice | single z-slice |

**EGFP/Tbr2a**

| A | A' | A'' |
|---|----|-----|
| A | A' | A'' |

| B | B' |
|---|----|
| C | C' |

| D | D' | D'' |
|---|----|-----|
| E | E' | F | F' |

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FIGURE 7 (Continued)
most EGFP+ cells in the subpallium were GABAergic (6 out of 8 cells analyzed; arrowheads in Figure 7F), but that there is no co-expression in the pallium (Figure 7E). Pallial GABAergic cell bodies were intermingled with the EGFP+ cell bodies in a salt and pepper manner (Figures 7Eiii,iv). The number of GABAergic cells in the pallium is less than that of EGFP+ cells. GABAergic neurons in the pallium are likely to be non-proliferative and postmitotic (E–F). Transverse sections through the telencephalon of a 4dpf Et(gata2:EGFP)bi105 brain stained with anti-GABA (magenta) to label GABAergic cells and anti-EGFP (green). EGFP+ cells in the pallium (E) are for the most part GABA-. Gaps where GABAergic pallial interneurons intermingle with EGFP+ pallial neurons are visible (arrowheads in panel (Ei)). In subpallial telencephalic domains (F) some cells do co-express EGFP and GABA (arrowheads in panels (Fi–iv)). Different z-sections through the same Et(gata2:EGFP)bi105 brain stained with anti-GABA (magenta) to label GABAergic cells and anti-EGFP (green). EGFP+ cells in the pallium (E) are for the most part GABA-. Gaps where GABAergic pallial interneurons intermingle with EGFP+ pallial neurons are visible (arrowheads in panel (Ei)). In subpallial telencephalic domains (F) some cells do co-express EGFP and GABA (arrowheads in panels (Fi–iv)).

**Individual Morphologies of EGFP+ Neurons in Et(gata2:EGFP)bi105 Larvae**

The density of pallial EGFP expression in Et(gata2:EGFP)bi105 larvae prevents any analysis of the morphology and projections of individual neurons. To visualize sparsely labeled neurons, we employed the CRISPR/Cas9 approach developed by Auer et al. (2014) and further modified by Kimura et al. (2014) to excise the EGFP transgene and mosaically insert Gal4. We modified this method slightly by adding a UAS:TDTomato construct to the injection mix. The mosaic inheritance of this DNA construct, layered upon the already mosaic conversion to Gal4, together with fine process staining properties of TDTomato, permitted both increased mosaicism and better labeling of neuronal processes (Schematic Figure 8).

Injection of the CRISPR/Cas9/EGFP guideRNA/Gal4/UAS cocktail into one cell stage Et(gata2:EGFP)bi105 embryos lead to mosaic RFP labeling of small numbers of pallial neurons (Figures 8A–I). All observations of morphology were made at 6 dpf. Labeled pallial neurons fell into two obviously distinct morphologies:

1. A population of neurons surrounding a central core of telencephalic neuropil (demarcated by SV2 expression, Figure 8B). These cells had monopolar morphology with their single process extending toward the neuropil core before ramifying into multiple dendrites that formed a dense tangle with dendrites of adjacent neurons. Some labeled processes were also present in the contralateral telencephalic neuropil core (Figure 8B). We were unable to ascertain whether these contralateral processes originated from these monopolar pallial neurons, perhaps crossing the anterior commissure, or other subpallial cells labeled in this specimen. Some processes also projected to caudal regions through the forebrain bundle. A survey of the single neuron morphologies registered to the mapzebrain atlas (Kunst et al., 2019) with soma located in the pallium (n = 28) revealed some pallial neurons projecting through the forebrain bundle to innervate caudal regions, such as the hypothalamus. None of the pallial neurons in this mapzebrain dataset had processes that crossed in the anterior commissure to elaborate processes in the contralateral pallium. So, although this cannot be definitively ruled out, it seems more likely such processes originate from subpallial gata2:EGFP+ neurons.

2. A population of multipolar neurons with stellate morphology (Figures 8G–I). These neurons were positioned either at the dorsal surface or within the central neuropil core. These cells extended multiple processes directly from the cell body that appeared to encase small areas of neuropil or groups of cell bodies (Figures 8G–I).

These experiments also allowed us to analyze the morphology of subpallial cells (Figures 8J–Q). Figures 8K, I shows one such neuron whose cell body lay just rostral to the anterior commissure. This neuron was monopolar, with an elaborate dendritic arbor, some with spines, in the vicinity of the anterior commissure (Figures 8K, I). The cell extended a bifurcated axon bilaterally down the left and right forebrain bundles to reach downstream targets that may include the ventral entopeduncular nucleus (vENT, arrow Figure 8K), pregeleral, tuberal and hypothalamic areas (Figure 8K). While the dendrites of this cell ramified only ipsilaterally, axon collaterals were bilateral and overtly symmetrical. Another example of a subpallial projection neuron with predominantly ipsilateral projections (Figures 8M, N) innervated similar caudal regions.

[FIGURE 7 | Et(gata2:EGFP)bi105 is expressed in post-mitotic glutamatergic pallial neurons. Sagittal (A,B) and horizontal (C) sections through a 4dpf Et(gata2:EGFP)bi105 brain stained with sytox (magenta) to label cell nuclei and anti-EGFP (green). EGFP expression is largely excluded from the dorsal and medial ventricular zones of the telencephalon. (D,D’) Dorsal views of a 4dpf Et(gata2:EGFP)bi105 brain labeled with anti-BrdU (red) and anti-EGFP (green) antibodies. (D) z-projection and (D’) shows a single coronal z-slice through the same brain. BrdU is excluded from the EGFP+ neurons in the dorsal telencephalon indicating that these cells are likely to be non-proliferative and postmitotic. (E–F) Transverse sections through the telencephalon of a 4dpf Et(gata2:EGFP)bi105 brain stained with anti-GABA (magenta) to label GABAergic cells and anti-EGFP (green). EGFP+ cells in the pallium (E) are for the most part GABA-. Gaps where GABAergic pallial interneurons intermingle with EGFP+ pallial neurons are visible (arrowheads in panel (EI)). In subpallial telencephalic domains (F) some cells do co-express EGFP and GABA (arrowheads in panels (Fi–iv)). Different z-sections through the same Et(gata2:EGFP)bi105 brain showing subpallial EGFP+ neurons on both sides of the brain. Dorsal (G) and lateral (H) views of Et(gata2:EGFP)bi105 brains at 5dpf labeled with FISH using a “cocktail” probe for vglut2a/slc17a6b and vglut2b/slc17a6a (magenta) and antibody labeling with anti-GFP (green). EGFP+ cells in the pallium (gata2:EGFP) bi105 embryos. This probe cocktail into one cell stage Et(gata2:EGFP)bi105 embryos lead to mosaic RFP labeling of small numbers of pallial neurons.
FIGURE 8 | Morphologies of pallial and subpallial neurons. Schematic: Schematic parts (i,ii) adapted from Kimura et al., 2014 explaining the Crisp/Cas9 mediated method to switch EGFP to Gal4. (iii) Shows our adaptation of this method to sparsely label single neurons and their processes by adding 5XUASTdTom to the injection mix. (A–Q) Etgata2:EGFP;bi105 larvae injected with CRISPR cocktail and labeled with anti-RFP and other markers. (A–I) Pallial Neurons. (A–I) Dorsal views of 6dpf Etgata2:EGFP;bi105 larvae injected with CRISPR cocktail. Labeled with anti-RFP (FIRE depth color code) (A,D) and FIRE LUT (B,C,E,F), anti-SV2 (gray) (B), anti-tERK (gray) (E). (C) Detail of “monopolar” dorsal pallial neurons from larva shown in panel (A,B) [area indicated by whisker box in panel (B)]. (F) Detail of single monopolar pallial neuron from larva shown in panels (D,E) [area indicated by whisker box in panel (E)]. (G–I) Three examples of basket shaped cells all found in the dorsal pallium of larva shown in panels (D,E). (J–Q): Subpallial neurons. (J–J”) A cluster of midline subpallial cells from larva shown in panels (D,E) exhibiting (Continued)
from this neuron also crossed the midline in the anterior commissure to terminate in the vicinity of the contralateral ventral entopeduncular nucleus. Axonal processes also crossed the midline in the hypothalamus. The subpallial neurons shown in Figures 8J–J, with cell bodies located at the midline, showed similar innervation of commissural regions and long descending axons. We observed one cell within the anterior commissure (Figures 8O–Q). This cell exhibited cocoon-like dendrites, tightly coiling within the ac itself. It sent one process rostrally into the contralateral pallium at the border with the olfactory bulb and another caudally through the lateral forebrain bundle to the posterior tuberculum.

A survey of single neuron morphologies registered to mapzebrain atlas (Kunst et al., 2019) with somata located in the subpallium (n = 139) showed long-projecting neurons comparable to those shown in 8K-M to be very common with around 70% projecting either bilaterally or ipsilaterally to caudal regions, predominantly the hypothalamus, interpeduncular nucleus, posterior tuberculum and raphe nuclei. Unlike pallial cells in mapzebrain, the subpallial neurons frequently innervate the contralateral subpallium and pallium via the anterior commissure.

Mosaic analysis of neuron morphologies in Tg(gata2:EGFP)bi105 larvae demonstrated that, even at larval stages, telencephalic cells already have elaborated intricate processes and dendritic arbors.

Registration of Transgene Expression Reveals the Morphogenetic Rearrangements of Telencephalic Regions During Development

As we have described, the Tg(gata2:EGFP)bi105 transgene labels post-mitotic pallial cells, as well as a small group of subpallial cells. The restricted pallial expression of EGFP makes it a useful tool to track telencephalic cell populations through key stages of telencephalic development, particularly relating to eversion.

To build on our previous analyses of pallial eversion in zebrafish (Folgueira et al., 2012), we have taken advantage of non-linear volumetric image registration to register brains with labeling of different cell populations/structures to reference brains (Figures 9, 10). We used acetylated tubulin labeling, common between all datasets, as a reference channel (Figures 9E,K,Q; not shown in 10). These registered datasets describe how major telencephalic cell populations/structures (olfactory bulb, pallium, subpallium, and tela choroidea) change position relative to each other during development.

Olfactory Bulb

To view changes in olfactory bulb position and cell populations during early telencephalon development, the following transgenes were used in combination with the Tg(gata2:EGFP)bi105 transgene in registered data sets:

1. Tg(-10lx2a:EGFP)zf176 (Figures 9D,J,P, 10D,J,P) labels a subset of mitral cells, the principle output neurons of the olfactory bulb (Miyasaka et al., 2009, 2014).

2. Tg(1.4dlx5a-dlx6a:GFP)et1 (Figures 9B,H,N,T, 10B,H,N,T) labels a subset of olfactory bulb GABAergic interneurons (Li J. et al., 2005). Figure 11 shows Tg(1.4dlx5a-dlx6a:GFP)et1 in comparison with Tg(gata2:EGFP)bi105 in frontal view.

3. Anti-acetylated tubulin immunohistochemistry labels olfactory bulb glomeruli, amongst other structures (Figures 9E,K,Q, 11D,F); see Folgueira et al., 2012.

The first mitral cells to differentiate by 1dpf (Figures 9D,F, 10D,E) were located within the dorsal telencephalon, close to the anterior intrencephalic sulcus (AIS). By this stage, mitral cells had already extended processes (Figure 9D). Between 2dpf (Figures 9J, 10J) and 5dpf, the telencephalic domain caudal to the olfactory bulb expanded greatly, increasing the distance between the OB and the AIS (Figures 9BR, 10PQ). Mitral cell axons reached the pallium and right habenula via the olfactory tracts and stria medullaris, respectively (Figures 9J,P, 10PQ; see also Miyasaka et al. (2009, 2014). Olfactory bulb interneurons, visualized with the 1.4dlx5a-dlx6a:GFP transgene, could be differentiated based on cell morphology. By 2dpf, these interneurons already extended processes toward the olfactory glomeruli (not shown), which increase in morphological complexity from 2dpf to 5dpf (Figures 9N, 10N, 11Fi).

Pallium

The development of the pallium, as visualized through the changes in expression of gata2:EGFP in Tg(gata2:EGFP)bi105 fish (Figures 9A,G,M,S, 10A,G,M,S), has already been well covered in earlier parts of this paper. Here we compared EGFP expression with that in Tg(1.4dlx5a-dlx6a:GFP)et1 fish, in which pallial GABAergic interneurons of subpallial origin are labeled (Mione et al., 2008; Yu et al., 2011), and in Tg(-10lx2a:EGFP)zf176 fish. We observed that 1.4dlx5a-dlx6a:GFP+ interneurons had a salt and pepper distribution in the pallium (Figures 9N, 10N), being more sparse in number than gata2:EGFP+ cells (Figures 9M, 10M,Q; compare
Figures 11Ci–vi with Figures 11Fi–vi). Note that registration of Tg(1.4dlx5a-dlx6a:GFP)ø1 data in lateral view (4dpf) in combination with the other markers was not satisfactory (hence not shown), but this was not the case for the dorsal view (Figure 10Q). The morphological complexity of 1.4dlx5a-dlx6a:GFP+ pallial interneurons increased over time, such that by 4/5dpf they ramified dense processes throughout the pallial neuropil (Figures 9N, 10N,Q, 11Fiii,iv). The -10lhx2a:EGFPzf176 transgene labels a neuropil within the pallium, within which axons from olfactory mitral cells terminate. This neuropil, which is nested within the gata2:EGFP+ pallial domain, may give rise to the posterior zone of the dorsal telencephalic area (labeled Dp in Supplementary Figures 1D,E; see Miyasaka et al. (2009, 2014)).

Subpallium
To observe how eversion affects subpallial populations, two transgenic lines were imaged in combination with the Et(gata2:EGFP)ø105 line.

1. Tg(1.4dlx5a-dlx6a:GFP)ø1 (Figures 9B,H,N,T, 10B,H,N,T, 11D–F) labels neuronal precursors and GABAergic neurons in regions of the subpallium (MacDonald et al., 2010; Yu et al., 2011). Based on the expression pattern of dlx5a mRNA in the adult, this transgene is likely to label the dorsal (Vd), ventral (Vv), supracommissural (Vs), central (Vc), lateral (Vl) and posterior divisions of subpallium (Vp) (see Ganz et al., 2012). These divisions are not well differentiated by 5dpf, so we were unable to confirm this expression pattern.

2. Tg(isl1:GFP)ø0 (Figures 9C,I,O,U, 10C,I,O,U; see Higashijima et al., 2000) shows a more restricted expression in the subpallium than the 1.4dlx5a-dlx6a:GFPø1 transgene. Recently, GFP expression in the adult brain has been extensively described in this transgenic line (see Baeuml et al., 2019) where the transgene labels neurons in the ventral subpallium (Vv), ventral part of the dorsal subpallium (Vdv) and ventral domain of the supracommissural subpallium (Vs) (see Baeuml et al., 2019; Porter and Mueller, 2020).
At 1dpf, the ventral gata2:EGFP+ cluster overlapped with the 1.4dlx5a-dlx6a:GFP+ domain (Figures 9A,B,F, 10A,B,E, 11A,B,Cii,D,E,Fii), but not with the isl1:GFP+ one (Figures 9C,F, 10C,E,F; see also Supplementary Figure 1). Pallial gata2:EGFP+ cells were located dorso-laterally to the 1.4dlx5a-dlx6a:GFP+ domain at this stage (Figures 10A,B,E). At 2dpf, 1.4dlx5a-dlx6a:GFP expression was largely complementary to gata2:GFP expression (Figures 9G,H,L, 10G,H,K,L), marking the pallial- subpallial boundary (Figure 10K). At this stage, viewing from a dorsal aspect (Figure 10K) showed that the gata2:EGFP+ (pallium) and 1.4dlx5a-dlx6a:GFP+ regions are distributed radially, with gata2:EGFP+ pallial cells located dorsolateral to the 1.4dlx5a-dlx6a:GFP+ domain. This arrangement changes with elongation of the telencephalon along its rostro-caudal axis from 2dpf to 5dpf (Figures 10M–W).

Expansion of the Tela Choroidea

The tela choroidea is a thin sheet of neuroepithelial origin that encloses the dorsal part of telencephalic ventricle of the everted telencephalon. At 1dpf, the telencephalic ventricle, delineated by anti-ZO1 labeling (this antibody labels apical cell junction proteins in both the tela choroidea and the cells lining the ventricle, see Supplementary Figure 2), is restricted medially, becoming larger caudally at the level of the AIS (Figures 10A–D,F). At this stage, both 10lxh2a:EGFP+ mitral cells (Figures 10D,F) and gata2:EGFP+ pallial cells (Figure 10A,F) were not covered by the ventricular surface and the tela choroidea dorsally, thus the telencephalon was not yet everted. By 2dpf, the number of gata2:EGFP+ and 1.4dlx5a-dlx6a:GFP positive cells increased, with the caudalmost telencephalic parenchyma expanding into the AIS (Figures 10G,H,K,L). Thus, a portion of the pallium becomes
covered by the ventricular surface and associated tela choroidea, so from this stage the telencephalon (pallium) can be described as everted (Figures 10G,L). By 5dpf most pallial cells are covered by the tela choroidea with the exception of very few gata2:EGFP+ pallial cells located rostrolaterally (Figures 10M,R). This shows that the pallium is not completely everted by 5dpf (Figures 10M,R), but it will be at later stages. Thus, it seems that eversion process does continue after 5dpf, at least at extreme
rostral locations. The olfactory bulb was clearly not everted, as the rostral attachment of the tela choroidea or taenia (for full details on the taeniae and eversion see Nieuwenhuys, 2009b) was just caudal to the olfactory bulb (Figures 10P,V).

In addition to registering these transgenes at key stages during telencephalic eversion, we also imaged the Et(gata2:EGFP)bi105 transgenic line at 6dpf, the stage of development used in the Zebrafish Brain Browser atlas (ZBB). Supplementary Figure 3 shows a snapshot of the registered Et(gata2:EGFP)bi105 expression pattern as viewed in the online ZBB viewer. A.PNG file is available for download (see Supplementary Material) so that users may upload this expression pattern and compare/combine it with any other labels registered to ZBB.

**DISCUSSION**

In this study, we have performed a detailed analysis of spatially restricted transgene expression in the telencephalon, in combination with other markers, to follow eversion and regionalisation of the zebrafish telencephalon during larval development. The registration of data from transgenic lines and other structural markers onto reference telencephali at key timepoints during development provides a framework atlas of major regions, structures and cell populations within the developing telencephalon. These datasets are available for further refinement through addition of further datasets and incorporation within other atlases, such as the Zebrafish Brain Browser (Marquart et al., 2015, 2017) within which we have registered a 6dpf Et(gata2:EGFP)bi105 dataset (Supplementary Figure 3).

**An Atlas of Telencephalon Development**

To understand how neural circuits control behavior it is necessary to integrate well-annotated structural data with functional data (Arrenberg and Driever, 2013). This is challenging with respect to the teleost telencephalon that exhibits an everted morphology, rendering direct comparison with other vertebrate telencephali difficult. The repertoire of tools available for dissecting the anatomy and function of neural circuits has increased dramatically in the last few years (Scott, 2009; Leung et al., 2013; Feierstein et al., 2015; Förster et al., 2017, 2018; Robles, 2017), and includes the creation and characterization of many transgenic lines expressing fluorescent proteins in discrete brain areas (Kwan et al., 2007; Scott et al., 2007; Kawakami et al., 2010; Suster et al., 2013; Randlett et al., 2015; Takeuchi et al., 2015; Marquart et al., 2017). Despite this wealth of tools and resources, there is still need of well-annotated baseline structural data that can be used to annotate reference brains (Ronneberger et al., 2012; Marquart et al., 2015, 2017; Randlett et al., 2015).

Existing neuroanatomical resources for zebrafish include 3D atlases based on confocal data during development and in the adult (Bryson-Richardson et al., 2007; Ullmann et al., 2010; Ronneberger et al., 2012; Marquart et al., 2015, 2017; Randlett et al., 2015; Kunst et al., 2019; Kenney et al., 2021; zebrafishbrain.org). However, these atlases are usually based on a single developmental timepoint, so cannot communicate the gross morphological changes happening early in development. Here we present an atlas of the zebrafish telencephalon during key stages of development (1dpf, 2dpf and 5dpf). Our 5dpf annotations are easily extended to 6dpf, the stages presented in the Zebrafish Brain Browser (Marquart et al., 2015, 2017), Zebrafish Brain Atlas (Ronneberger et al., 2015) and mapzebrain (Kunst et al., 2019).

We have used anti-acetylated tubulin as an anatomical reference instead of cytoplasmic/nuclear staining often used in other atlases (Ronneberger et al., 2012; Marquart et al., 2015; Randlett et al., 2015). We observed that, as previously shown in VibeZ (Ronneberger et al., 2012), this labeling works well as a reference channel and has the added bonus that one can view the development of the axonal scaffold through development. We also include the marker ZO1, which delineates the extension of the ventricular zone and allows the degree of eversion to be tracked. Given there are portions of the ventricle that, as a result of eversion, are difficult to visualize without markers (see Supplementary Figure 2; see also Nieuwenhuys, 2009b), ZO1 or other ventricular zone markers are crucial when building a reference telencephalon resource.

**Eversion and “Outside-In” Construction Mode of the Teleost Telencephalon**

Eversion models have failed so far to explain how morphogenetic events during development lead to the organization of the pallial regions in the adult (Yamamoto et al., 2007; Mueller et al., 2011; Porter and Mueller, 2020). After the primary events of eversion from 1dpf to 5dpf (Folgueira et al., 2012), there is a subsequent major expansion of the telencephalon that affects its organization (present results; Dirian et al., 2014; Furlan et al., 2017). With the main events of eversion occurring so early (mostly by 1-5dpf) and growth so late, this complicates the eversion narrative of a simple lateral out-folding of the neural tube during early development (Butler and Hodos, 2005; Yamamoto et al., 2007; Mueller et al., 2011; Nieuwenhuys, 2011; Porter and Mueller, 2020). In addition, eversion models currently do not adequately account for differential timing in the development of different telencephalic areas.

The relative size of different telencephalic domains changes dramatically over the first weeks of development. For instance, early on, the nascent olfactory bulbs constitute much of the telencephalon whereas there is disproportionate, massive expansion of the pallial telencephalic lobes from 5dpf to 20dpf. Within the expanding pallium, Dirian et al. (2014) found two distinct populations of neural stem cells that are segregated in space: a dorso-medial domain and a lateral domain. Consistent with late expansion of pallial territories, the lateral domain that contributes neurons to the lateral domain of the pallium, DI, only becomes neurogenic from 5dpf onward.

Following morphogenesis of the ventricle, tangential growth of the ventricular zone, driven by telencephalic neuronal differentiation, may be one of the driving forces to expand the pallial ventricular surface (Abotiz and Montiel, 2019; present results). By 5dpf, the zebrafish telencephalon shows everted...
morphology (Folgueira et al., 2012; present results), but there is a subsequent expansion of the telencephalic ventricular zone, which grows in lockstep with general telencephalic lobe growth. Cells are added in an outside-in manner following a “sequential stacking” mode, with neurons arranged in age-related layers surrounding a central “core” of earliest-born neurons (Furlan et al., 2017). This addition of newborn cells toward the pallial ventricular surface may force tangential expansion of the ventricular zone as the telencephalon grows. We observed that the first cells to express EGFP in Et(gata2:EGFP)bi105 are also post-mitotic and located centrally in the telencephalon, so these first EFGP+ cells could be part of this central “core”.

**gata2:EGFPbi105 + Pallial Neuron Morphologies**

Using a Crispr-based labeling technique, we revealed morphologies of cell types in the pallium and subpallium of 6dpf larvae. We observed two different cell types in the pallium by 6dpf that, despite the early age of the animals, show quite elaborate dendritic arbors, similar to those reported in adult fish. Studies in various adult teleosts have shown different neuronal cell types occupying different areas of the pallium (Demski and Beaver, 2001; Demski, 2013). Deep pallial territories host large efferent projection neurons, while periventricular neurons are predominantly small stellate shaped cells (Demski and Beaver, 2001; Demski, 2013). These findings match well with the cell types described in this study. Stellate cells reported here, for instance, resemble the morphology of interneurons present in lateral and medial regions (Dm) of the pallium in adult fish (Yamane et al., 1996; Demski and Beaver, 2001; Giassi et al., 2012; see Demski (2013) for a review). Thus, even at early stages, differentiated pallial neurons already show complex arborization, indicative of being integrated in active neural circuits.

**Establishment of Neural Circuits and the Emergence of Control of Behaviors in the Telencephalon**

The adult telencephalon is involved in many functions, such as motor control, sensory processing, memory, learning, emotions and social interaction. What is the sequence of maturation of the telencephalic circuits involved in these functions? Our results indicate that areas of the pallium and subpallium show considerable expansion after 5dpf suggesting late maturation of these regions. This is in agreement with behavioral studies, which have shown that complex behaviors appear from 2 to 6 weeks post-fertilization (Valente et al., 2012; Aoki et al., 2013; Cheng et al., 2014; Dreosti et al., 2015; Lal et al., 2018; Stednitz et al., 2018; Tunbak et al., 2020; Bartoszek et al., 2021).

As a consequence, most functional studies likely involving the role of the telencephalon, such as fear conditioning, social interaction, learning and memory, were performed on older zebrafish.

Despite late maturation of some telencephalic circuits, the animal needs to process information from the environment and coordinate movement from an early age. By 4dpf, the larva has already hatched and is transitioning into independent feeding. In this sense, we observed quite mature cell morphologies already by 2dpf in the olfactory bulb, indicative of early olfactory processing. We also observed that telencephalic gata2:EGFP+ cells show complex morphologies and connections by 5-6dpf. In the case of the subpallial gata2:EGFP+ cells, by 6dpf they already show a complex connection pattern with caudal regions, which resembles the connections described in the adult for the subpallium (Rink and Wullimann, 2004). Thus, from an early age, this circuit might allow the young animal to exert telencephalic control over certain larval behaviors.

**gata2:EGFP+ Subpallial Cells May Be Part of the Zebrafish Septum**

Various cell populations and nuclei in the zebrafish subpallium have been described based on topographical location and neurochemistry. Among these, the ventral area (Vv) and the lateral area (Vl) of the subpallium are proposed to be homologous to septal areas in tetrapods (Rink and Wullimann, 2004; Wullimann and Mueller, 2004b; Ganz et al., 2012; Baeuml et al., 2019). Here we were able to follow the development of a gata2:EGFP+ subpallial cluster located at pre-commissural and commissural levels of the zebrafish telencephalon and analyze its connections. We identify gata2:EGFP+ subpallial cells with elaborate dendritic arbors that span around the anterior commissure and axons that bifurcate and project unilaterally or bilaterally to tuberal and hypothalamic areas. These cell populations could correspond, at least partially, with early-migrated telencephalic area M4 (see Mueller and Wullimann, 2005; Mueller et al., 2008). Previously we showed the prethalamic origin of another group of cells in the zebrafish ventral telencephalon, the ventral entopeduncular nucleus (see Turner et al., 2016). However, we did not analyze cell migrations into the telencephalon in the present study, so we cannot comment on the exact origin of the subpallial gata2:EGFP+ cells, or, for that matter, any other population in the telencephalon.

Based on location and projection pattern, we believe gata2:EGFP+ subpallial cells could correspond to septal cholinergic populations described in the lateral subpallium of adult zebrafish (Mueller et al., 2004) and other teleosts (trout: Pérez et al., 2000). Both subpallial gata2:EGFP+ cells and cells immunoreactive for choline acetyl transferase (ChAT) share similar location in Vl and similar projection pattern to the hypothalamus (Mueller et al., 2004; Rink and Wullimann, 2004). Cholinergic cells also show varicose fibers in the subpallium, especially dense around the anterior commissure (Mueller et al., 2004), similar to the gata2:EGFP+ subpallial cells described in this study. We do not think that the location of the subpallial gata2:EGFP+ cells correspond with neuropeptide Y+ cells previously described in Vl (Castro et al., 2006; Turner et al., 2016), which project to the dorsomedial pallium (not shown), or with the proper entopeduncular nucleus (former ventral entopeduncular nucleus; Turner et al., 2016). In any case, we have not looked at colocalization of either neuropeptide Y or ChAT in Et(gata2:EGFP)bi105 fish, so the putative identity of these gata2:EGFP+ subpallial neurons remains speculative.
The heterogeneity of cell types in the lateral subpallium indicates that further detailed analysis is needed to dissect nucleus identity, anatomy, connectivity and development of this area.

**egr3 Expression and Function, Future Directions for Looking at Telencephalic Activity**

The insertion point in Et(gata2:EGFP)bi105 fish is 7 kb upstream of the first exon of *Early Growth Response 3* gene (*egr3*). Although so far we have used this transgenic line for neuroanatomical studies only (Folgueira et al., 2012; Turner et al., 2016; present study), we believe it will be a useful tool for other analyses. Although the line expresses a stable form of EGFP, the CRISPR–Cas9 switching method (Auer et al., 2014; Kimura et al., 2014) could be used to convert this EGFP transgene to Gal4 to generate a more versatile transgenic line for such studies. Our results, using this method to interrogate single cell morphology, already show that the eGFP sgRNAs can effectively lead to transgene replacement.

*egr3* is a member of the immediate early gene (IEG) family that encodes transcription factors with almost identical zinc finger DNA binding domains (O’Donovan et al., 1999). Among other functions, IEGs such as *egr1*, *egr2/krox20*, *egr3* and *egr4* have been implicated in neural plasticity in response to neuronal activation (Li L. et al., 2005, Li et al., 2007; Kim et al., 2012). In mice, *egr3* expression is induced by synaptic activity and is required for hippocampal long-term potentiation and long-term depression (Galâtano-Mendel et al., 2007), as well as for hippocampal and amygdala dependant learning and memory (Li et al., 2007). The persistent expression of *egr3* in spatially restricted populations of CNS cells from early stages (this study, Deguchi et al., 2009) suggests that in addition to roles in synaptic plasticity, the gene may also have a role in early neuronal differentiation or function. Indeed, a closely related gene *egr2/krox20*, is required for normal rhombomere development (Mechta-Grigoriou et al., 2000).

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Animal Welfare and Ethical Review Body (AWERB)-University College London; Animal research performed under Home Office License.

**AUTHOR CONTRIBUTIONS**

MF, KT, and TH conceived and designed the work, with inputs from SW and acquired and analyzed the data. PH registered the transgenic lines. KT, TH, and LV developed the single cell labeling with Crispr/Cas9 technique. IB provided comments on the final manuscript. All authors contributed to interpretation of data and to writing the article.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnana.2022.840924/full#supplementary-material

**Supplementary Figure 1** | Registered brain (lateral view) showing Elgata2:EGFPbi105 (green), Tg10lhx2a:EGFPzf176 (FIRE) and Tg Isl1:GFPwbi (yellow) transgene expression at 5dpf. (A-F) Single z-slices showing transverse sections (A′-F′) made using orthogonal view tool (YZ) in ImageJ. Panel (E) is more lateral than panel (F). The rostro-caudal level of transversal sections (A′-F′) are indicated by a pink line on accompanying lateral z-slice in panels (A-F), (D,E) same rostro-caudal level, (D) closer z-slice to the midline than panel (E), (D′-E′) show the location of Tg10lhx2a:EGFPzf176 mitral cell processes in putative Dp. Scale bars: 50 μm.

**Supplementary Figure 2** | Z1 as a marker of the ventricular surface and tela choroidea. Lateral (A,B) and dorsal views (C,D) of the telencephalon of 4dpf (A,B) and 6dpf (C,D) fish stained against Z1 (magenta) and counterstained with sytox orange (cyan). (A,B) White arrowheads point to the portions of the ventricle that are clearly visible with a nuclear stain only (AIS dorsally, optic recess ventrally). (B) Yellow arrowheads point to the extension of the ventricle visible by ZO1 staining. Double arrow marks the extension of the tela choroidea. This could be easily missed with a nuclear staining only (compare (A,B)). (C,D) Full extension of the ventricle and tela choroidea is not clearly visible with a nuclear stain only in dorsal view (C), but clear when Z01 staining is used (dotted line marks the full extension of the ventricle and tela choroidea dorsally). (E) Transverse section of the telencephalon of a 4dpf larva labeled against Z01 (magenta) and tubulin (cyan). Note the T-shape of the ventricle. The tela choroidea extends dorsally (places of attachment or taeniae are marked with arrows). (F) Schematic representation of E, showing the ventricular surface (VS) and tela choroidea (tc) labeled by Z01. Brain parenchyma is represented in pale blue. Taeniae marked with arrows. Scale bars: 100 μm.

**Supplementary Figure 3** | Elgata2:EGFPbi105 expression at 6dpf registered to Zebrafish Brain Browser (ZBB). Screenshot of the online ZBB viewer showing a 6dpf Elgata2:EGFPbi105 larvae labeled with anti-EGFP (green) registered to the ZBB standard brain. HuC channel (gray) shows full brain structure.

**Supplementary Figure 4** | A 6dpf Elgata2:EGFPbi105 stack in PNG format. This dataset can be uploaded to Zebrafish Brain Browser atlas (see "Materials and Methods").
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