Lef1-dependent hypothalamic neurogenesis inhibits anxiety

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Abstract

While innate behaviors are conserved throughout the animal kingdom, it is unknown whether common signaling pathways regulate the development of neuronal populations mediating these behaviors in diverse organisms. Here, we demonstrate that the Wnt/β-catenin effector Lef1 is required for the differentiation of anxiolytic hypothalamic neurons in zebrafish and mice, although the identity of Lef1-dependent genes and neurons differ between these 2 species. We further show that zebrafish and Drosophila have common Lef1-dependent gene expression in their respective neuroendocrine organs, consistent with a conserved pathway that has diverged in the mouse. Finally, orthologs of Lef1-dependent genes from both zebrafish and mouse show highly correlated hypothalamic expression in marmosets and humans, suggesting co-regulation of 2 parallel anxiolytic pathways in primates. These findings demonstrate that during evolution, a transcription factor can act through multiple mechanisms to generate a common behavioral output, and that Lef1 regulates circuit development that is fundamentally important for mediating anxiety in a wide variety of animal species.

Author summary

Humans, mice, fish, and even flies exhibit anxiety-like behavior despite the fact that their brain anatomy varies widely. This study reveals another common thread that runs through these diverse animals: the molecular origins of their shared behavior. Gene knockout experiments in mouse and zebrafish show that the molecular signal Wnt acts through the transcription factor Lef1 to inhibit anxiety in both species. The pathway is required for formation
of anxiolytic neurons in a highly conserved brain region, the hypothalamus. From there, however, the process diverges. In the fish, the pathway triggers genes including corticotropin-releasing hormone binding protein (crhbp), but in mice the same pathway calls into action a different gene, Pro-melanin concentrating hormone (Pmch). By comparison, the fruit fly Dro sophila activates crhbp, similar to zebrafish. Furthermore, CRHBP and PMCH show extraordinarily coordinated expression in the primates hypothalamus, indicating that they may act together downstream of Wnt and Lef1 to regulate human behavior. This work reveals the surprising finding that conserved signaling pathways can regulate common behavioral outputs through diverse brain circuits during evolution.

Introduction

Recent work has demonstrated that innate behaviors can be highly conserved across diverse animal models [1]. Individual neuronal populations that mediate these behaviors are specified during embryogenesis by transcription factors that can also be conserved across species [2]. However, molecular signaling pathways that regulate the development of common behavioral circuits have not been identified. As brain anatomy and connectivity change through evolution, it is possible that a single pathway could act through diverse molecular and cellular targets to establish a single behavioral output, which is the ultimate constraint on gene function.

Wnt/β-catenin signaling plays important evolutionarily conserved roles in brain development, and thus represents an ideal candidate pathway to link gene regulation with the evolution of behavioral circuits. The Wnt pathway acts through Tcf/Lef transcription factors [3], and both Wnt signaling and Lef1 are required for neurogenesis in the zebrafish hypothalamus [4], an evolutionarily ancient brain structure that regulates innate behaviors [5]. However, the identity and behavioral function of Lef1-dependent hypothalamic neurons, and their degree of evolutionary conservation, are unknown. Here, we show that Lef1 is required for the differentiation of hypothalamic neurons that inhibit anxiety in both zebrafish and mice, but through divergent molecular and cellular mechanisms in the 2 species. Generation of neurons expressing corticotropin-releasing hormone binding protein (crhbp) requires Lef1 in zebrafish but not in mice, whereas neurons expressing Pro-melanin concentrating hormone (Pmch) are Lef1-dependent in mice but not in zebrafish. Furthermore, zebrafish and Dro sophila have common Lef1-dependent crhbp expression in their respective neuroendocrine organs, consistent with an ancient conserved pathway that has diverged in mammals. Finally, the Genotype-Tissue Expression (GTEx) project [6] reveals a top-ranked positive correlation between CRHBP and PMCH in the human hypothalamus, suggesting co-expression and/or co-regulation. Both genes are also correlated with LEF1 expression in humans, and are expressed in the same region of the marmoset hypothalamus, consistent with a conserved regulatory pathway in primates. These findings suggest that the gene expression network regulated by a transcription factor can change during evolution while still generating a common behavioral output. Our data also suggest an anxiolytic role for Wnt signaling in the human hypothalamus, with potential implications for the etiology and treatment of anxiety disorders.

Results

Lef1 is required for the differentiation of hypothalamic neurons in zebrafish

We sought to first characterize the earliest cellular defect in lef1 null zebrafish mutants [4], so that we could perform a transcriptome analysis at that stage to identify Lef1-dependent genes.
Despite grossly normal morphology, mass, and brain size, lef1 mutants have a smaller caudal hypothalamus (Hc) at 15 days post-fertilization (dpf) [4], and we found that the size reduction occurred at as early as 3–4 dpf (Fig 1A and S1A and S1B Fig). At 3 dpf the tissue already contained fewer Wnt-responsive cells [7] (Fig 1B), as well as fewer serotonergic cells and ventricular GABAergic HuC/D+ neurons (Fig 1C and S1C Fig). However, th2;GFP+ dopaminergic neurons [8] were unaffected (S1D Fig), indicating that not all neuronal subtypes are Lef1-dependent. In addition, the number of BLBP+ cells was increased (S1E Fig), confirming an inhibitory role of Wnt signaling in the formation of hypothalamic radial glia [4,9].

To determine the cellular mechanism underlying the decreased populations in lef1 mutants, we measured apoptosis and proliferation. We observed an increase in p53-dependent apoptosis within the Hc at 3 dpf (Fig 1D), but no change in proliferation at 3 dpf and beyond (Fig 1E and S1F–S1H Fig). Rescue of apoptosis by loss of p53 (Fig 1D) did not restore HuC/D
expression in \textit{lef1} mutants (Fig 1F), consistent with a primary defect in progenitor differentiation. To confirm a failure in neurogenesis, we performed BrdU pulse-chase experiments, and observed fewer newly born serotonergic and ventricular HuC/D+ cells in \textit{lef1} mutants (S1I Fig). To test whether Lef1 functions cell-autonomously, we transplanted cells from \textit{lef1+/-} donors into the hypothalamic anlage of \textit{lef1} mutant hosts during gastrulation, and observed rescue of ventricular HuC/D expression only in donor cells (Fig 1G). Together these data suggest that Lef1 functions cell-autonomously to promote hypothalamic neurogenesis; in \textit{lef1} mutants, neural progenitors fail to differentiate and subsequently undergo cell death, leading to a smaller Hc. Our data also justified 3 dpf as the optimal time point to perform a transcriptome analysis.

**Lef1-dependent genes in the zebrafish hypothalamus are associated with anxiety**

To identify Lef1-dependent genes, we next performed RNA sequencing (RNA-seq) analysis of whole hypothalami dissected from 3 dpf control and \textit{lef1} mutant zebrafish embryos, and found 144 genes with an adjusted \textit{P} value (AdjP) <0.1, among which 53 genes had a fold change >2 (Fig 2A, S2 Table). Most of these genes had reduced expression in \textit{lef1} mutants (Fig 2A), consistent with Lef1 functioning as a Wnt transcriptional activator [10]. Surprisingly, Ingenuity Pathway Analysis (IPA) identified Lef1-dependent genes as being most highly associated with anxiety and depressive disorder (Fig 2B and S3 and S4 Tables). In contrast, genes associated with other hypothalamus-mediated behaviors, such as feeding (\textit{neuropeptide Y} [\textit{npy}], \textit{agouti-related protein} [\textit{agrp}], and \textit{proopiomelanocortin} [\textit{pomc}]) or sleep (\textit{hypocretin} [\textit{hcrt}]), were unaffected (S2 Table). We performed in situ hybridization on 3 dpf offspring of \textit{lef1+/-} incrosses and confirmed that all Lef1-dependent genes with specific detectable hypothalamic expression showed predicted changes in approximately 25% of embryos, consistent with Mendelian segregation (Fig 2C and 2D and S2A–S2C Fig). These included several known Wnt targets such as \textit{sp5a} and \textit{sp5l} [11] (Fig 2C), and anxiety-related genes identified from IPA (Fig 2B and 2D). Expression of neuronal markers such as \textit{crhbp} and 5-hydroxytryptamine receptor 1A b (\textit{htr1ab}), was lost specifically in the Hc of \textit{lef1} mutants while remaining intact in the rostral hypothalamus (Fig 2D), resulting in their relatively small fold change in whole hypothalamus RNA-seq analysis (S2 Table). In contrast, expression of other genes, such as 2 phosphodiesterase 9a (\textit{pde9a}) paralogs, was lost in the rostral hypothalamus and Hc of \textit{lef1} mutants (Fig 2D and S2A Fig), consistent with \textit{lef1} expression in both regions (Fig 2C). We also observed expression of Lef1-dependent genes in the Hc of wild-type (wt) adult zebrafish (S2D Fig), suggesting the presence of Wnt activity and Lef1-dependent neuronal populations throughout life. Together these results suggested that \textit{lef1} mutants might have an anxiety-related behavioral phenotype.

**Zebrafish \textit{lef1} mutants exhibit increased anxiety**

\textit{lef1} mutants raised with siblings had decreased survival and size (S3A and S3C Fig). When separated at 15 dpf, mutants survived normally (S3B and S3C Fig), but were still smaller than control siblings at culture densities that maximized their growth (Fig 3A and S3D Fig), a phenotype potentially due to enhanced anxiety [12]. We then performed a novel tank diving test to measure anxiety-related behavior [13]. We found that \textit{lef1} mutant larvae had a longer latency to enter the upper half of a novel tank and spent less overall time in this zone during the initial exploration phase (Fig 3B and 3C and S1 Video), consistent with elevated anxiety. Notably, \textit{lef1} mutants travelled less distance during this phase, partially due to more frequent freezing behavior as indicated by increased time in immobility (Fig 3D and 3E and S1 Video), and again consistent with elevated anxiety. Importantly, \textit{lef1} mutants no longer displayed
Fig 2. Lef1 activates expression of zebrafish hypothalamic genes associated with anxiety. (A) Volcano plot of zebrafish RNA sequencing (RNA-seq) shows differentially expressed genes in the 3 days post-fertilization (dpf) hypothalamus of left1 mutants compared to control. Only genes with adjusted P value (AdjP) < 0.1 (green line) are shown. Genes with an absolute value of log2 ratio > 1 (blue lines) are shown in red; others are shown in black. Node size represents the averaged fragments per kilobase of transcript per million mapped reads of a gene in the control. (B) Ingenuity Pathway Analysis (IPA) for zebrafish hypothalamic Lef1-dependent genes revealed 20 genes associated with anxiety and depressive disorder, listed in the table. (C and D) Representative images of whole mount in situ hybridization on 3 dpf control and left1 mutant embryos for known Wnt targets (C) and genes associated with anxiety and depressive disorder (D). Red and yellow arrows indicate expression in
anxiety-related behavior after the exploration phase (Fig 3F). The body growth and anxiety phenotypes in \textit{lef1} mutants could be explained by reduced expression of multiple hypothalamic genes including \textit{crhbpp} (Fig 2D), which encodes a corticotropin-releasing hormone (CRH) inhibitor [14]. However, pleiotropic phenotypes in zebrafish \textit{lef1} mutants [4,15] could also contribute to defects in growth or motor behavior. Therefore, we sought to create a tissue-specific mouse knockout model to examine the hypothalamic function of Lef1, and to determine whether it is evolutionarily conserved.

### Hypothalamic Lef1 inhibits anxiety in mice

\textit{Lef1} is expressed in the mouse Hc from embryonic day (E) 10.5 to adulthood [16,17], and while previously characterized \textit{Lef1} null mutants exhibit postnatal lethality and a smaller body size, no hypothalamic phenotypes were reported [18,19]. We created a mouse hypothalamic knockout model using \textit{Nkx2-1}^-\text{Cre} and \textit{Lef1}^{\text{flx}} alleles [20,21]. We also introduced the Cre reporter \textit{Rosa}^-tdTomato [22] to create the conditional knockout allele \textit{Nkx2-1}^-\text{Cre/;}\textit{Lef1}^{\text{flx}}/;\textit{Rosa}^-tdTomato/ (herein referred to as \textit{Lef1}^-\text{CKO}) and control littermates \textit{Nkx2-1}^-\text{Cre/;}\textit{Lef1}^{\text{flx}}/;\textit{Rosa}^-tdTomato/ (herein referred to as \textit{Lef1}^-\text{CON}), which were used for all experiments. We confirmed successful recombination by tdTomato expression (S5A Fig), and loss of hypothalamic \textit{Lef1} and Wnt reporter [23] expression in \textit{Lef1}^-\text{CKO} mice (S5B and S5C Fig), which were viable, caudal and rostral hypothalamus, respectively. Lateral (\textit{axin2, dkk1b, lef1, notum1a, crhbpp, and grin2cb}) or ventral (other genes) views were selected for optimal expression visualization. Scale bar: 100 μm.

![Fig 2.](https://doi.org/10.1371/journal.pbio.2002257.g002)

Fig 3. Lef1 regulates growth and anxiety in zebrafish. (A) Size of 30 days post-fertilization (dpf) fish when raised at 5 fish per tank separated by genotype. \(n=25, 30\) for control and mutant, respectively. (B-F) Novel tank diving test. Sixteen dpf larvae were analyzed between 1–3 minutes (C–E) or 4–6 minutes (F) after entering a novel tank. \(n=9\) for both controls and mutants. Data are mean ± SEM. *\(P<0.05\), **\(P<0.01\), ***\(P<0.001\), ns, \(P>0.05\) by unpaired Student \textit{t} tests. Raw data can be found in S1 Data.
fertile, and morphologically indistinguishable from Lef1<sup>CON</sup> littermates. However, both male and female Lef1<sup>CKO</sup> mice gained weight more slowly after weaning (Fig 4A), similar to the phenotype we observed in zebrafish lef1 mutants (Fig 3A), and again consistent with elevated anxiety [12].
To directly measure anxiety-related behavior, we used an elevated plus maze (EPM) test and found that male \textit{Lef1}^{CKO} mice spent significantly less time in the open arms and more time in the closed arms (Fig 4B) despite normal mobility (S4A Fig). In an open field test (OFT), male \textit{Lef1}^{CKO} mice spent significantly less time in the center zone (Fig 4C) despite normal mobility (S4B Fig). These results are consistent with elevated anxiety in male \textit{Lef1}^{CKO} mice. We also observed enhanced anxiety specifically in OFT with estrous female \textit{Lef1}^{CKO} mice, but not with diestrous or all females, or with EPM testing of any females (Fig 4B and 4C and S4A and S4B Fig), likely due to reported variations in anxiety-related behavior between different sexes [24] and different behavioral assays [25]. Together, these results suggest a conserved role of hypothalamic Lef1 in inhibiting anxiety.

Hypothalamic \textit{Lef1} is required for generation of \textit{Pmch}+ neurons in mice

Consistent with the neurogenesis defect we observed in zebrafish, we found fewer HuC/D+ cells in the mouse hypothalamic ventricular zone in \textit{Lef1}^{CKO} embryos at E14.5 (Fig 5A). Importantly, this effect was restricted to coronal sections in which endogenous Lef1 is expressed (S5B Fig). To identify Lef1-dependent genes in the mouse hypothalamus, we performed RNA-seq analysis of hypothalami dissected from E14.5 \textit{Lef1}^{CON} and \textit{Lef1}^{CKO} embryos, and surprisingly identified only 1 protein-coding gene that mapped to a unique locus with an AdjP < 0.1 and a fold change > 2, \textit{Pmch} (Fig 5B and S5 Table). \textit{Pmch} expression normally overlaps with \textit{Lef1} in the premammillary hypothalamus, and extends into the lateral hypothalamus (Fig 5C) [17,26]. We confirmed loss of \textit{Pmch} expression in E14.5 \textit{Lef1}^{CKO} embryos by quantitative real-time PCR (qPCR) and immunostaining (Fig 5D and S5D and S5E Fig). The only other significantly affected protein-coding gene identified by RNA-seq, \textit{Ribosomal Protein L34} (\textit{Rpl34}) (Fig 5B, S5 and S6 Tables), is a repetitive processed pseudogene that could not be conclusively mapped to a single genomic locus, although one copy is located adjacent to \textit{Lef1}.

Reduced \textit{Pmch} expression in \textit{Lef1}^{CKO} embryos was unexpected because its orthologs were not significantly affected in RNA-seq analysis of zebrafish \textit{lef1} mutants (S2 Table). To determine if any Lef1-dependent genes were conserved with zebrafish later in development, we performed another RNA-seq analysis at postnatal day (P) 22, when \textit{Lef1}^{CKO} mice begin to exhibit a growth defect (Fig 4A). In this experiment, we identified only 2 affected protein-coding genes mapped to unique loci with an AdjP < 0.1: \textit{Pmch} and \textit{Tachykinin receptor 3} (\textit{Tacr3}) (Fig 5B, S6 Table). \textit{Tacr3} is known to be co-expressed in \textit{Pmch}+ neurons, along with \textit{CART prepropeptide} (\textit{Cartpt}) [27]. We confirmed their reduced expression in the lateral hypothalamus of P22 \textit{Lef1}^{CKO} mice by qPCR and in situ hybridization (Fig 5D and S5E and S5F Fig), consistent with loss of \textit{Pmch}+ neurons. Decreased body weight observed after ablating \textit{Pmch}+ neurons [28,29] may therefore be related to an anxiolytic role for these cells [12], which is further supported by characterization of their inputs and activity [30].

Orthologs of multiple Lef1-dependent anxiety-related genes in zebrafish are expressed near \textit{Lef1} in the mouse hypothalamus, such as \textit{Pde9a} and \textit{Nitric oxide synthase 1} (\textit{Nos1}) at E14.5 [26], and \textit{Crhbp} and \textit{Histidine decarboxylase} (\textit{Hdc}) in adults [16]. However, RNA-seq analysis indicated that expression of these genes was Lef1-independent in mice (S5 and S6 Tables), and we confirmed this result for \textit{Crhbp} by qPCR and in situ hybridization (Fig 5D and S5E and S5F Fig). In addition, we confirmed that expression of zebrafish \textit{pmch} orthologs [31] does not depend on Lef1 at either 3 dpf or 15 dpf (S6A–S6C Fig). While we cannot rule out the possibility that our RNA-seq analysis of the mouse hypothalamus lacked the sensitivity to identify other conserved Lef1-dependent genes, it is clear that the identity of Lef1-dependent neurons relevant for anxiety differs between zebrafish and mice.
Fig 5. Hypothalamic Lef1 is required for Pmch+ neuron formation in mice. (A) Immunostaining of HuC/D+ cells in the hypothalamic ventricular zone of E14.5 CON-M and CKO-M, with quantification shown on the right (n = 4). Images are z-projections of 16 μm confocal optical slices, shown with dorsal side on top, and higher magnification views of yellow squares in the insets. (B) Volcano plot of mouse RNA sequencing (RNA-seq) shows differentially expressed genes in the hypothalamus of CKO-M compared to CON-M at E14.5 (left) and P22 (right), using the same format as in Fig 2A. (C) E14.5 sagittal in situ hybridization images (www.genepaint.org) show expression of Lef1 (red arrows) and Pmch in the wild-type (wt) hypothalamus [26]. (D) Quantitative real-time PCR (qPCR) analysis for male shows hypothalamic gene expression in E14.5 and P22 CKO-M relative to CON-M. (E) P22 coronal in situ hybridization images show expression of Pro-melanin concentrating hormone (Pmch), CART prepropeptide (Cartpt), and Tachykinin receptor 3 (Tacr3) in the lateral hypothalamus. 3V, third ventricle. Data are mean ± SEM. ***P<0.001, ns. P > 0.05 by unpaired Student t tests. Scale bars: 400 μm in (C); 30 μm in (E). Raw data can be found in S1 Data.

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Lef1 dependence of \textit{crhbp} expression is conserved between zebrafish and \textit{Drosophila}

Interestingly, many Lef1-dependent genes in zebrafish encoding components of anxiety-mediating transmitter pathways, such as GABA, 5-HT, and CRH (Fig 2B), have a conserved function in \textit{Drosophila} anxiety-like behavior [1]. Therefore, we hypothesized that hypothalamic Lef1-dependent neurons in zebrafish may represent an evolutionarily ancient pathway. The \textit{Drosophila} pars intercerebralis (PI) and pars lateralis (PL) represent neuroendocrine organs equivalent to the vertebrate rostral hypothalamus and Hc, respectively [32]. In \textit{Drosophila}, a single Lef/Tcf family member, \textit{pangolin (pan)}, functions as a Wnt activator [33,34]. Consistent with our hypothesis, we detected specific \textit{pan} expression at stage 14 and the \textit{crhbp} ortholog \textit{CG15537} expression at stage 16 in the \textit{Drosophila} PL primordium [32] (Fig 6A–6C). Furthermore, we observed a loss of \textit{crhbp} expression in the PL of \textit{pan} mutants [34] at stage 16, despite intact expression in the PI and normal PL morphology (Fig 6C–6E). \textit{Drosophila crhbp} in the PL may also be anxiolytic by inhibiting CRH/CRH-like diuretic hormone in the PI [1,32,35], thus these results support a relationship between neuroendocrine Lef1 function and the development of anxiolytic \textit{Crhbp}+ neurons dating back to a common bilaterian ancestor. By contrast, \textit{Pmch} is a vertebrate specific gene, and Lef1-dependent \textit{Pmch}+ neuronal circuitry in mice may reflect a more recent mammalian divergence that co-evolved with new brain structures [36].

Coordinated expression of \textit{PMCH} and \textit{CRHBP} in the human hypothalamus

Our animal models suggest that in humans Lef1 may also regulate the formation of Pmch+ and/or Crhbp+ hypothalamic neurons. To test this hypothesis, we compared the hypothalamic RNA-seq transcriptomes of 96 human individuals from the GTEx project [37] (S7 Table). Despite the fact that these data did not include prenatal samples, we found that expression of

![Fig 6. Loss of Drosophila corticotropin-releasing hormone binding protein (crhbp) expression in pangolin (pan) mutants.](https://doi.org/10.1371/journal.pbio.2002257.g006)
PMCH and CRHBP are both moderately correlated with LEF1, which is expressed at a relatively low level in the adult human hypothalamus (Fig 7A and 7B). Notably, PMCH and

![Image](https://doi.org/10.1371/journal.pbio.2002257.g007)

Fig 7. Correlation analysis in the human hypothalamus. (A-G) Pearson correlations for hypothalamic gene expression among 96 postmortem human samples obtained from the Genotype-Tissue Expression (GTEx) project [39]. All the Pearson’s $r$ and $P$ values were calculated between 2 genes, and displayed in the graphs or tables after sorting by $r$ values. Correlation expression profiles are shown for gene pairs Pro-melanin concentrating hormone (PMCH) versus LEF1 (A), Corticotropin-releasing hormone binding protein (CRHBP) versus LEF1 (B), CRHBP versus PMCH (F), and Agouti-related protein (AGRP) versus Neuropeptide Y (NPY) (G), with reads per kilobase of transcript per million mapped reads (RPKM) at log$_{10}$ scale used on both axes. Note that 1 data point (NPY: 0.1395; AGRP: 0) was not included in (G) due to the inability of plotting a 0 value on the logarithmic axis. Three tables of correlated genes for LEF1 (C), CRHBP (D), and PMCH (E) list the top 9 positively correlated genes plus selected genes, including those involved in canonical Wnt signaling labeled in red. See the full list in S8 Table.

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CRHBP were both within the top 100 LEF1-correlated genes, along with known Wnt targets such as Sal-like protein 4 (SALL4) [38] and SP5 [11] (Fig 7C and S8 Table).

In the course of this analysis, we noticed similar correlation profiles for CRHBP and PMCH (Fig 7A and 7B), suggesting a possible expression correlation between these 2 genes. Surprisingly, we found CRHBP and PMCH to be the most highly correlated genes with each other (Fig 7D–7F and S8 Table), a relationship that has never been reported previously. Among the top 200 PMCH- or CRHBP-correlated genes, we also found 2 Wnt ligands and 1 Wnt co-activator: R-Spondin 1 (RSPO1) [40] (Fig 7D and 7E). As a comparison, AGRP is the most highly correlated gene with Neuropeptide Y (NPY) (Fig 7G and S8 Table), consistent with their co-expression in the same hypothalamic neurons [41]. Interestingly, while Pmch and Crhbp are expressed in different regions of the mouse hypothalamus [16], they are expressed in the same hypothalamic nuclei in another primate, the marmoset according to the Marmoset Gene Atlas (https://gene-atlas.bminds.brain.riken.jp). Importantly, the results of all our correlation analyses are recapitulated on GeneNetwork (www.genenetwork.org) [42], which imported an older version of GTEx’s datasets and calculated Pearson correlation across a population (See Materials and methods). Together these data suggest co-expression of PMCH and CRHBP in the primate hypothalamus and potential regulation by LEF1-mediated Wnt signaling in humans.

Discussion

In this study, we demonstrate that Lef1-mediated hypothalamic Wnt signaling plays an evolutionarily conserved role in regulating the formation of anxiolytic neurons (See Fig 8 for summary). In zebrafish lef1 mutants, neural progenitors fail to differentiate and undergo apoptosis, resulting in a smaller Hc (alternatively named the hypothalamic posterior recess, the posterior part of the paraventricular organ, or the caudal zone of the periventricular hypothalamus [4,43,44]). Any or all of the 20 anxiety-related genes that are misregulated in the zebrafish mutant (Fig 2B) may contribute to the behavioral phenotypes that we observe. Likewise, our data do not conclusively prove that crhbp+ neurons, or indeed any individual Lef1-dependent...
neuronal populations, mediate the effect of Lef1 on anxiety. Such a conclusion would require either rescue of the lef1 mutant phenotype by restoration of missing neurons, or phenocopy by specific ablation of the cells. However, the specific loss of Pmch+ neurons in our mouse conditional knockout (Fig 3B), combined with the unexpected expression correlation between PMCH and CRHBP in the human hypothalamus (Fig 7F), is consistent with a common role for these 2 genes in behavior. While we also cannot rule out the possibility that Lef1 mutants may have other behavioral defects, genes that are known to regulate other hypothalamus-driven behaviors, such as Npy, Agrp, Pomc, and Hcrt, are unaffected in our mutants (S2, S5 and S6 Tables). In addition, pure assessment of other behaviors cannot distinguish a direct phenotype from an anxiety-related secondary phenotype.

While the major product of Pmch, melanin-concentrating hormone (MCH), is an anxiolytic factor in teleosts [45], studies in mammals have reported it to be either anxiolytic, anxiogenic or having no effect [46,47]. In addition, the Pmch propeptide makes at least 2 more neuropeptides, neuropeptide-glutamic acid-isoleucine (NEI), and neuropeptide-glycine-glutamic acid (NGE), which are also involved in stress response and anxiety [48]. Germline Pmch mouse knockouts gain weight more slowly than controls, a phenotype originally attributed to decreased food intake [49]. However, on a different background strain, the same group reported that the knockout mice were not hypophagic, while retaining a growth phenotype [50]. Interestingly, all rodent models ablating Pmch [49–53] or Pmch+ neurons [28,29] exhibit a reduced growth rate. One possible underlying mechanism could be enhanced anxiety [12], which was not directly tested in any of these studies. Therefore, we hypothesize that in Lef1CKO mice, loss of hypothalamic Pmch+ neurons is responsible for elevated anxiety, leading to a secondary growth phenotype.

Our data suggest that the gene expression and neuronal subtypes dependent on Lef1 can change during evolution while maintaining a common behavioral output. While transcriptional networks can undergo rapid rewiring at the level of enhancer binding sites during yeast, insect and mammalian evolution [54,55], the direct transcriptional targets of Lef1 mediating hypothalamic neurogenesis are still unknown. We have identified Tcf/Lef consensus binding sites in zebrafish and mouse Crhbp and Pmch loci, but it remains important to determine whether these 2 genes are direct targets of Lef1, or are instead lost as a secondary result of neurogenesis defects in mutants. In either case, it will also be useful to understand the circuitry of Lef1-dependent neurons. While the targets of Crhbp+ neurons in Drosophila and zebrafish are unknown, the projections of Pmch+ neurons in the hypothalamus of mice and other mammals are well characterized, and the regulation of these circuits by Lef1 in these species may be linked to anatomical and functional expansion of target brain regions such as the cortex [36]. Importantly, the coordinated expression of CRHBP and PMCH in the human hypothalamus suggests that they may be co-expressed in a single neuronal cell type.

Loss of other genes important for hypothalamic neurogenesis has been shown to affect behavior [2]. Interestingly, mice lacking hypothalamic Dbx1 also exhibit a loss of Pmch+ neurons along with other populations [56]. In that study, Lef1-expressing hypothalamic nuclei were hypothesized to regulate innate behaviors outside the hypothalamic-pituitary-adrenal (HPA) axis, partly due to the observation of expanded Wnt activity in Dbx1 knockout animals. However, because our work demonstrates that Lef1 is in fact required for the genesis of Pmch+ neurons and for HPA-related behaviors, an alternative explanation is that Dbx1 functions in a parallel pathway to Lef1.

Together these results identify Wnt signaling as a link between brain development and function that allows essential behaviors to be maintained even as anatomical structures change through evolution. In addition, given the function for hypothalamic Wnt signaling in regulating postembryonic zebrafish neurogenesis [4], and the continuous expression of Lef1 in the
hypothalamus of fish (S2D Fig) and mammals [16] throughout life, it would be interesting to test a possible contribution to adult behavior using temporal conditional knockout models. While Wnt signaling in the mammalian hippocampus and nucleus accumbens has been associated previously with anxiety and depression [57,58], our data demonstrate a novel requirement for pathway activity in a brain region that is highly conserved throughout the vertebrate lineage, and may prove useful for the diagnosis and treatment of hypothalamus-related anxiety disorders.

Materials and methods

Ethics statement

All experimental protocols were approved by the University of Utah Institutional Animal Care and Use Committee and were in accordance with the guidelines from the National Institutes of Health. Approval number: 16–09011. Zebrafish were euthanized by ice water immersion. Mice were euthanized by CO2 or ketamine/xylazine.

Subjects: Zebrafish

Zebrafish (Danio rerio) were bred and maintained in a 14:10 hour light/dark cycle as previously described [59]. Zebrafish per tank were fed with similar amount of food and treated by the staff who were blinded to the experiments. Wt strains were AB. The following mutant and transgenic strains were used: lef1^td11 [4], Tg(top:GFP)^b25 [7], Tg(dlx6a-1.4dlx5a-dlx6aGFP)^p61 [60], Tg(h2afv-GFP)^ko6 [61], Tg(th2:GFP-Aequorin)^z201 [8], p53^e7 [62]. lef1^− homozygous mutants were identified between 3 dpf and 10 dpf by DASPEI staining as described previously [15] and at or after 15 dpf by loss of caudal fin [4]; wt and heterozygous siblings were used as controls. All the zebrafish were from at least 1 single-pair breeding. Genotyping was done as described before for lef1^td11 [4] and p53^e7 [63], except primers used for lef1^td11 (forward primer: 5'-CACTCTCTCCAGC CCAACATT-3', reverse primer: 5'-TGTTACTGTTGG GACTGATTCTG-3').

Subjects: Mice

Male and female C57BL/6J mice (Mus musculus) were group-housed with 2–5 mice per cage in a reverse 12 hour light/dark cycle with ad libitum access to food and water. Mice were 19–20 and 15–20 weeks old at the time of behavioral tests for male and female animals, respectively. Ai9 reporter Rosa^tdTomato (line 007905) [22], Nkx2-1^Cre (line 008661) [21], and TCF/Lef: H2B-GFP mice (line 013752) [23] were purchased from Jackson Laboratories. Lef1^flox/flox mice were provided by HHX [20]. All strains were maintained on a C57BL/6J background except TCF/Lef:H2B-GFP mice, which were originally on a C57BL/6 × 129 background. Male Nkx2-1^Cre/Cre;Lef1^flox/flox;Rosa^tdTomato/tdTomato mice were used to generate conditional knockout (Lef1^CKO: Nkx2-1^Cre/Cre;Lef1^flox/flox;Rosa^tdTomato/+ ) and control (Lef1^CON: Nkx2-1^Cre/Cre;Lef1^flox/+ ;Rosa^tdTomato/+ ) offspring. Females breeders were maintained by inbreeding. Male breeders were maintained by interbreeding Nkx2-1^Cre/Cre;Lef1^+/+ and Nkx2-1^Cre/Cre; Lef1^flox/+ for no more than 5 generations to avoid potential artifacts caused by Cre homozygous inbreeding [64]. In occasional litters, Ai9 reporter expression was observed throughout the body of approximately 10% of experimental animals, consistent with published literature [21]; such animals were not used for experiments. All the mice were from at least 3 litters unless otherwise noted. Sex at E14.5 was determined by genotyping by Jarid 1c [65]. When generating experimental mice for body weight measurement and behavioral tests, each litter was culled to 8 pups at P0. Genotyping for Rosa^tdTomato and TCF/Lef:H2B-GFP animals was done according
to available Jackson Laboratory protocols for these strains. Genotyping for Nkx2-1<sup>Cre</sup> mice was done using primers for Cre recombinase detection (forward primer: 5'-ATGCTTCTGTCCGT TTGCCG-3', reverse primer: 5'-CCTGTTTTGACGTCACCG-3'). Genotyping for Lef1<sup>flax</sup> mice was done using primers contributed by HHX (forward primer: 5'-GCAGATATAGACTAGCACC-3', reverse primer: 5'-TCCACACACTAAGGCTAC-3').

Subjects: *Drosophila*

Canton-S wild-type and *pan<sup>2</sup>* mutant (BL4759) *Drosophila melanogaster* strains were obtained from Bloomington Stock Center.

**Zebrafish transplantation experiments**

At the sphere stage, 10–50 blastula cells from donor embryos were transplanted using a glass micropipette into the dorsal side of shield stage host embryos, 20–40 degrees from the animal pole, representing the hypothalamus anlage [66]. Embryos were then raised to 5 dpf for immunohistochemistry. Donor and host embryos were retained for genotyping to identify *lef1* mutants.

**BrdU labeling**

Four dpf zebrafish embryos were incubated in E3 media containing 10 mM BrdU (Sigma-Aldrich, St. Louis, MO) at 28.5˚C for indicated time before being washed in E3 media for at least 3 times.

**Immunohistochemistry: Zebrafish**

Embryos and larvae were fixed in 4% paraformaldehyde (PFA) for 3 hours at room temperature (RT) or overnight (O/N) at 4˚C followed by brain dissection. Brains were either dehydrated in methanol and stored at −20˚C, or immediately processed for immunohistochemistry. For 3 dpf embryos, 5% sucrose was included in the fixative to ease dissection. Brains were treated with 0.5 U dispase (Gibco #17105–041) in 2% PBST (PBS/2% Triton X-100) for 60 minutes at RT. For BrdU, PCNA, pH3 or Caspase-3 staining, brains were washed in water for 5 minutes twice, followed by incubation in 2 N HCl for 60 minutes at RT, followed by 2 more water washes. Brains were then blocked in 5% to 10% goat serum in 0.5% PBST for 60 minutes at RT. Embryos were incubated in primary antibodies in block O/N at 4˚C and secondary antibodies and Hoechst 33342 (Life Technologies, H3570) in block O/N at 4˚C before mounting in Fluormount-G (SouthernBiotech, Birmingham, AL) with the ventral hypothalamus facing the coverslip. Primary antibodies were all used at 1:500 dilution except as noted: chicken anti-GFP (Aves Labs, GFP-1020), rabbit anti-GFP (Molecular Probes, A11122), mouse anti-HuC/D (Molecular Probes, A21271), rabbit anti-5-HT (Immunostar, 541016), rabbit anti-pH3 (1:400, Cell Signaling, 9713), rabbit anti-active Caspase-3 (BD Pharmingen, 55965), rabbit anti-BLBP (Abcam, ab32432), mouse anti-PCNA (Sigma, P8825), and chicken anti-BrdU (ICL, CBDU-65A-Z). Secondary antibodies were all used at 1:500 dilution: goat anti-mouse Alexa Fluor 488 (Invitrogen, A11001), goat anti-rabbit Alexa Fluor 488 (Invitrogen, A11008), donkey anti-chicken Alexa Fluor 488 (Jackson ImmunoResearch, 703-545-155), goat anti-rabbit cy3 (Jackson ImmunoResearch, 116-165-003), goat anti-mouse cy3 (Jackson ImmunoResearch, 115-165-003), goat anti-mouse Alexa Fluor 647 (Invitrogen, A21235), goat anti-rabbit Alexa Fluor 647 (Invitrogen, A21244), and goat anti-chicken Alexa Fluor 647 (Invitrogen, A21449). Hoechst 33342 (1:10,000) was used to stain nuclei. All the primary antibodies were validated previously [4,67].
Immunohistochemistry: Mice
E14.5 embryo heads were dissected in PBS and fixed in 4% PFA at RT for 1.5 hours or O/N at 4˚C. Brains were dissected and cryoprotected in 15% and then 30% sucrose, embedded in OCT, and stored at −80˚C. Brains were cryosectioned at a thickness of 16 μm, air dried and stored at −80˚C. Air-dried sections were then washed in PTW (PBS+0.1% Tween 20) 3 times, followed by permeabilization in 0.25% PBST for 5 minutes and blocking in 10% goat serum in PTW for 60 minutes. Sections were incubated in primary antibodies in blocking solution O/N at 4˚C and secondary antibodies in blocking solution for 2 hours at RT, followed by Hoechst 33342 stain for 10 minutes at RT before mounting in Fluoromount-G. Antibodies used were as described above except rabbit anti-LEF1 (1:200, Cell Signaling, 2230), goat anti-PMCH (1:500, Santa Cruz, sc14509) and donkey anti-goat Alexa Fluor 647 (1:400, Invitrogen, A21447). All primary antibodies were validated by absence of staining in Lef1 CKO animals. For HuC/D staining, incubation for 30 minutes in 0.5 U dispase was performed in 0.25% PBST.

Immunohistochemistry: Drosophila
Drosophila immunohistochemistry was performed as previously described [68] except that a fluorescent secondary antibody was used. Antibodies used were as described above except mouse anti-FasII (1:5, DSHB, 1D4), which was validated previously [32].

Probes for in situ hybridization
In situ hybridization probes were made by a clone-free method as described previously [69,70], with DNA templates purified using Zymo Research DNA Clean & Concentrator-5 kit. Primers were designed by Primer-BLAST [71] except for mouse genes with primer sequences available from the Allen Brain Atlas (ABA) [16] or GenePaint Atlas [26]. A full list of primers used to make probes is in S9 Table. cDNA made from 3 dpf zebrafish embryos, P2, and P60 mouse hypothalamus, and adult Drosophila (gift from C. Thummel) was used as the initial template for PCR to generate T7 promoter-containing DNA. RNA probes for zebrafish lef1 [72] and axin2 [73] were previously described. The RNA probe for Drosophila pan was generated from the Drosophila Gene Collection T7 promoter-containing cDNA GM04312 [74].

Whole mount in situ hybridization: Zebrafish
Zebrafish whole mount in situ hybridization was performed as described previously [75] except that 15 dpf and adult zebrafish were fixed in 4% PFA O/N at 4˚C followed by washing in PBS and brain dissection. All tissues were treated for 30 minutes with 10 μg/ml Proteinase K. Pigmented embryos were bleached in 1% H2O2/5% formamide/0.5× SSC O/N at RT after in situ hybridization. 3 dpf embryos and postembryonic brains were imaged in 100% glycerol and PBS, respectively. For automated whole mount in situ hybridization, all steps following probe hybridization and before color reaction were performed using a BioLane HTI (Intavis, Chicago, IL).

Section in situ hybridization: Mice
Twenty-five μm brain cryosections were collected and post-fixed as previously described [76] (http://developingmouse.brain-map.org/docs/Overview.pdf). In situ hybridization was then performed as described [77].

Whole mount in situ hybridization: Drosophila
Drosophila whole mount in situ hybridization was performed as described previously [68].
Body length: Zebrafish

Zebrafish from a single home tank were anesthetized using tricaine (Sigma-Aldrich, St. Louis, MO) in shallow water. Images were acquired of immobilized, non-overlapping fish with a ruler for scale. Body length was calculated by measuring the distance between the mouth and the anterior edge of the tail fin, using ImageJ.

Novel tank diving test

Five fish from lef1+/- incrosses were raised per tank starting at 5 dpf. lef1 mutants and controls were separated at 15 dpf. Novel tank diving tests [13] were performed on 16 dpf larvae during the early afternoon of the same days, before lef1 mutants start to display surface behavior at 20 dpf. Novel rectangular tanks (16.6 cm × 9.5 cm × 12.3 cm) were illuminated by a centered white light, and videos were acquired with a mounted Nokia Lumia 640 phone 1080p camera. For each experiment, single mutant and control larvae were netted and then removed simultaneously from their home cages and transferred to novel tanks with identical water volume. The order of netting mutant and control fish was rotated between trials. Videos were viewed in MPlayerX to manually analyze the latency of larvae to enter the upper half of the tank after initial sinking. Videos were then imported and analyzed using Ethovision XT version 11.5 (Noldus, Leesburg, VA) during the initial exploration phase, with a tracking period of 2 minutes beginning 1 minute after release into the novel tank to decrease water agitation resulting from netting. Videos were also analyzed after the initial exploration phase with a tracking period during the 4 to 6 minute interval. Tracks were analyzed for distance travelled, time in upper half of the tank and time of immobility.

Body weight: Mice

All pups were weaned at P21 immediately following the first weighing. Pups weighing less than 6.5 g were excluded from analysis. All mice were weighed during the morning of the same days of the following weeks.

Behavior tests: Mice

Group-housed mice were allowed to acclimate to the animal facility for behavioral tests 9 days after an on-campus transfer. Each mouse was handled daily for 2 minutes, during midmorning for 7 days before commencement of behavioral testing using the cupped hand method [78]. To avoid behavioral variation caused by female estrous cycle [79], a vaginal lavage procedure was done after daily handling for estrous phase evaluation for 7 days, as previously reported [80]. Female mice in their proestrus or estrus phases were collectively grouped as “Estrus” and females in their metestrus and diestrus phases were collectively grouped as “Diestrus.” All mice were acclimated to the behavior room for 1 hour under red light (69 lux) before commencement of tests. Open field and EPM behavioral tests were performed in order, once daily for 2 days, from 9 AM to 5 PM. The experimenter was blinded to genotype.

OFT

Each mouse was placed in a circular plexiglass chamber (4.5” diameter × 3” height) located inside an illuminated (330 lux) circular open field arena (110 cm diameter) and allowed to acclimate for 1 minute to decrease movement bias resulting from experimenter handling. After 1 minute, the plexiglass chamber was removed from the arena, and the mouse was allowed to freely explore the arena for 10 minutes. Movement was video recorded and analyzed using Ethovision version 9 (Noldus, Leesburg, VA).
EPM

The EPM apparatus was elevated 60 cm from the floor, having 2 open arms (35 cm × 5 cm) and 2 closed arms (35 cm × 16 cm) connected by a central platform (5 cm × 5 cm). The EPM was illuminated by a white light (205 lux) at the center platform. Each mouse was placed in a rectangular opaque white plexiglass chamber (2” × 3” × 5”) located on the center platform, and allowed to acclimate for 1 minute before commencement of the test. The white chamber was removed and the mouse was allowed to freely explore the EPM for 5 minutes. Behavior was video recorded and analyzed using Ethovision version 9 (Noldus, Leesburg, VA).

RNA-seq: Zebrafish

Embryos were fixed for 1.5 hours in 4% PFA/5% sucrose in PBS at RT, followed by whole hypothalamus dissection with super-fine forceps (FST, 11252–00). For each biological replicate, 28 to 38 dissected hypothalami were pooled for lef1 mutant and control samples from at least 1 single-pair breeding. RNA was extracted using a RecoverAll Total Nucleic Acid Isolation Kit for FFPE (Ambion, AM1975) according to the manufacturer’s instructions. Three biological replicates were obtained on different days from offspring of different breedings. A total of 300 ng RNA per sample was submitted to the High Throughput Genomic Core at the University of Utah for RNA quality control by High Sensitivity R6K ScreenTape. RNA concentration by vacuum drying, cDNA library prep by Illumina TruSeq Stranded RNA Kit with Ribo-Zero Gold and sequencing by HiSeq 50 Cycle Single-Read Sequencing version 3. RNA-seq was analyzed by the Bioinformatics Core at the University of Utah. A transcriptome reference was created by combining GRCz10 chromosome sequences with Ensembl build 84 splice junction sequences generated with USeq (version 8.8.8) MakeTranscriptome. RNA-seq reads were mapped to the GRCz10 zebrafish transcriptome reference using Novoalign (version 2.08.03). Splice junction alignments were converted back to genomic space using USeq SamTranscriptomeParser. USeq DefinedRegionDifferentialSeq was used to generate per gene read counts, which were used in DESeq2 to determine differential expression. RNA-seq graph in Fig 2A was made by IPython Notebook with package NetworkX.

RNA-seq: Mice

E14.5 and P22 nonweaned male Lef1\textsuperscript{CON} and Lef1\textsuperscript{CKO} hypothalami were dissected using a fluorescent microscope in ice-cold PBS, while tail tissue was retained for genotyping. E14.5 tissues were immediately immersed in RNAlater (Thermo Fisher, Waltham, MA) and stored at 4°C for up to 7 days until RNA extraction. P22 tissues dissected from at least 2 litters were immediately homogenized in TRIzol (Thermo Fisher, Waltham, MA) and stored at −80°C. Three biological replicates were prepared from either 5 pooled hypothalami (E14.5) or a single hypothalamus (P22) from Lef1\textsuperscript{CON} and Lef1\textsuperscript{CKO} mice, and RNA was extracted on the same day using TRIzol followed by purification with an RNeasy Mini Kit (Qiagen, Hilden, Germany) and on-column DNase digestion (Sigma-Aldrich, St. Louis, MO). One μg of RNA per sample was submitted to the High Throughput Genomic Core at the University of Utah for RNA quality control with Agilent RNA ScreenTape, cDNA library prep with Illumina TruSeq Stranded RNA Kit with Ribo-Zero Gold, and sequencing using HiSeq 50 Cycle Single-Read Sequencing version 4. RNA-seq reads were mapped to GRCm38. Differential gene expression analysis and graph plotting were carried out using the same methods as for zebrafish RNA-seq.
qPCR

Three biological replicates of RNA from male and female mice were prepared as described above for RNA-seq. Two and a half μg RNA was used for cDNA synthesis with a SuperScript III Reverse Transcriptase kit (Invitrogen, Carlsbad, CA). qPCR was performed in triplicate using Platinum SYBR Green master mix (Invitrogen, Carlsbad, CA) on 96-well CFX Connect (Bio-Rad, Hercules, CA) plates or 384-well QuantStudio 12K Flex (Life Technologies, Durham, NC) plates at the Genomics Core at the University of Utah, according to manufacturer’s instructions. Gapdh was used to normalize quantification, and reverse transcriptase was omitted for controls. qPCR analysis was performed with the ΔΔCt method to determine relative expression change [81]. Dissociation curve analysis was performed to confirm the specificity of amplicons. qPCR primers were designed from PrimerBank [82] as follows (forward primer first, reverse primer second, in 5’ to 3’ orientation with PrimerBank ID in the parentheses), Pmch (12861395a1): GTCTGGCTGTAAAACCTTA CCTC, CCTGAGCATGTCAAAAT CTCTCC; Tacr3 (10946720a1): CTGGGCTTGCCAGTGA CAT, CGCTTGTGGGCCAAGA TGAT; Crhbp (162287189c2): CTTACCCTCGGACACT TGCAT, GGTCTGCTAAGGGCA TCATCT.

Image analysis and cell counting

Fluorescent images of dissected zebrafish and mouse brains were obtained with an Olympus FV1000 confocal microscope at the Cell Imaging Core at the University of Utah. Z-stack images were all maximum intensity z-projections of 3 μm slices; single- or double-labeled cells were manually counted in FV1000 ASW 4.2 Viewer. All the zebrafish and mouse in situ hybridization images were obtained with an Olympus SZX16 dissecting microscope except those in Fig 5E, S2C Fig and S6B Fig, which were obtained with an Olympus BX51WI compound microscope. Two months post-fertilization (mpf) zebrafish images (S3A and S3B Fig) were acquired using a Leica MZ16 microscope. Drosophila in situ hybridization images were obtained with a Zeiss Axioskop.

IPA

IPA (QIAGEN, Redwood City, CA) was performed with 129 mouse orthologs of the 138 zebrafish protein-coding genes identified from RNA-seq with AdjP <0.1 (S4 Table). Analysis was performed by the Bioinformatics Core at the University of Utah according to QIAGEN’s instructions and “diseases and functions” were extracted from the software (S3 Table).

Human correlation analysis

Publically available GTEx raw datasets were downloaded from www.gtexportal.org in April 2017 as a single file: GTEx_Analysis_v6p_RNA-seq_RNA-SeQCv1.1.8_gene_rpkm.gct.gz. Ninety-six hypothalamic samples were identified according to their specific strong PMCH expression, and extracted into S7 Table by IPython Notebook with packages gzip and xlwt. Pearson correlation was calculated by gene reads per kilobase of transcript per million mapped reads (RPKM) using IPython Notebook with function scipy.stats.stats.pearsonr, followed by result writing into S8 Table by IPython Notebook with package xlwt. The same Pearson correlation r values were confirmed using Excel’s CORREL function. A similar correlation result was obtained when searching for the top 200 correlated genes by Pearson on GeneNetwork (www.genenetwork.org) in April 2017. Several differences are noted between our analyses and GeneNetwork’s analyses. First, GeneNetwork imported an older version of GTEx’s datasets (GTExv5 Human Brain Hypothalamus RefSeq [Sep15] RPKM log2). Second, GeneNetwork
calculated Pearson correlation using RPKM log₂ rather than RPKM in our case. Third, GeneNetwork calculated Pearson’s sample correlation across a population, with an adjustment across the genome, and also based on the number of the top correlated genes requested by the users; in our case, we calculated Pearson correlation between 2 genes, and simply ranked all the genes by their Pearson’s r values calculated for the gene of interest. Lastly, GeneNetwork’s imported older GTEx datasets had 102 hypothalamic samples, 6 among which were left out in current GTEx’s server. The complete overlapping of the 96 samples further confirmed our successful extraction of hypothalamic datasets from the GTEx project.

Statistical analysis
No statistical methods were used to predetermine sample size. For behavioral assays, sample size was determined based on accepted practice. The experiments were not randomized. Due to visible phenotypes, the investigators were not blinded to outcome assessment except for whole mount in situ hybridization of zebrafish lef1+/− crosses, Drosophila pan+/− incrosses, and mouse body weight and behavioral assays. Two-tailed unpaired Student t tests were performed for all statistical analysis, except mouse body weight (2-way ANOVA with repeated measures), using GraphPad Prism software version 6. Outliers were identified by Grubbs’ test for behavioral assays with significance assigned at \( P < 0.05 \) (alpha = 0.01). All the criteria for excluding data points were established prior to data collection.

Supporting information
S1 Fig. Lef1 promotes neurogenesis in the zebrafish caudal hypothalamus (Hc). (A and B) Hc size in control and lef1 mutants (A) estimated by the area of confocal ventricular slice (B). Hc was defined as an oval indicated by red outline in (B). The lengths of \( a_1, a_2, b_1, b_2 \) in the representative image (B) were measured by ImageJ, and the area of the oval was calculated by the following equations: Estimated area = \( \pi \cdot \frac{a \cdot b}{4} \); \( a = a_1 + a_2; b = \frac{(b_1 + b_2)}{2} \). (C) Co-immunostaining of HuC/D and GABAergic lineage marker dlx5/6:GFP [83] in the 3 dpf Hc. Three confocal channel-split magnified images of the region depicted by the yellow rectangle are shown on the right. A representative image is shown for at least 3 embryos tested. (D and E) Immunostaining of th2:GFP+ (D) and BLBP+ cells (E) in the Hc of 3 dpf control and lef1 mutant. Representative images are shown on the left, and quantifications are shown on the right. Higher magnification views of yellow rectangles in single channel are shown in the insets in (E). (F-H) Measurement of proliferation in the Hc of 5 dpf control and lef1 mutant as shown by pH3+ (F) and PCNA+ cells (G; representative image on the left and quantification on the right; cells adjacent to the horizontal ventricle were counted), and 1 day BrdU labeling (H; schematic on the left). (I), BrdU pulse-chase (schematic on the left) to measure birth of 5-HT+ and ventricular HuC/D+ cells after 4 dpf. Data are mean ± SEM, except mean ± SD in (A) and (I). * * * \( P < 0.001 \), * * \( P < 0.01 \), * \( P < 0.05 \), ns. \( P > 0.05 \) by unpaired Student t tests. All images are confocal ventricular slices. All scale bars are 25 \( \mu \)m except 12.5 \( \mu \)m in the magnified image in (C). See S1 Table for description of quantification and experimental n. Raw data can be found in S1 Data.

(TIF)

S2 Fig. Whole mount in situ hybridization for zebrafish Lef1-dependent genes identified from RNA-seq. (A) Representative images of whole mount in situ hybridization on 3 dpf control and lef1 mutant embryos. Red and yellow arrows indicate gene expression in caudal and rostral hypothalamus, respectively. Lateral (adarb2, ccdc129, foxb2, klf17, mmp17b, and slc18a2) or ventral (other genes) views were selected for optimal expression visualization. (B)
Quantification of expression following whole mount in situ hybridization on 3 dpf offspring from *lef1*+/− incrosses. Fifty to eighty-five embryos were analyzed per gene. (C) Images of 3 dpf control brains centered on Hc from ventral view. (D) Gene expression in the hypothalamus of 4 months post-fertilization (mpf) female wild-type zebrafish from ventral view. Representative images are shown in (C) and (D) for at least 2 samples tested. Images of ventral view have anterior on top; images of lateral view have dorsal on top and anterior on the left. Red dashed outlines in (C) and (D) depict the caudal hypothalamus. Scale bars: 0.1 mm in (A); 5 μm in (C); 0.2 mm in (D). Raw data can be found in S1 Data.

S3 Fig. Physiological and behavioral analysis of zebrafish *lef1* mutants. (A–C) Body size and survival rate of *lef1* mutants under different culture conditions. Offspring of *lef1*+/− incrosses were either unsorted or sorted by genotype at 15 dpf, and raised at 25 fish per tank. Body length and number of surviving fish at 2 mpf are shown in (A). (B) Elevated plus maze. (C) Open field test. In (A) and (B), n = 12, 9 for male CON, CKO. In (A), n = 11, 11 for female CON, CKO in estrus; n = 12, 11 for female CON, CKO in diestrus. In (B), n = 12, 6 for female CON, CKO in estrus; n = 11, 16 for female CON, CKO in diestrus. Data are mean ± SEM. **P < 0.01, ns. P > 0.05 by unpaired Student t tests. Outliers depicted in black (B) were excluded from statistical analysis using the Grubbs’ test (P < 0.05). Raw data can be found in S1 Data.

S4 Fig. Mouse anxiety tests. (A) Elevated plus maze. (B) Open field test. In (A) and (B), n = 12, 9 for male CON, CKO. In (A), n = 11, 11 for female CON, CKO in estrus; n = 12, 11 for female CON, CKO in diestrus. In (B), n = 12, 6 for female CON, CKO in estrus; n = 11, 16 for female CON, CKO in diestrus. Data are mean ± SEM. **P < 0.01, ns. P > 0.05 by unpaired Student t tests. Outliers depicted in black (B) were excluded from statistical analysis using the Grubbs’ test (P < 0.05). Raw data can be found in S1 Data.

S5 Fig. Cellular and molecular phenotypes in mouse *Left<sup>CKO</sup>* hypothalamus. (A) P50 female *Nkx2-1<sup>Cre<sup>+</sup></sup>;*Lef1<sup>flk</sup>‡<sup>+</sup>;*Rosa<sup>tdTomato</sup>‡<sup>+</sup> (CON-F) expresses tdTomato in the hypothalamus. Bright field (left) and red fluorescence (right) ventral view images of the same brain with anterior on top are shown. Representative images are shown for at least 3 adult brains dissected. (B) Immunostaining for Lef1 in the hypothalamus of E14.5 *Lef1<sup>CON</sup>* (CON) and *Lef1<sup>CKO</sup>* (CKO). Coronal images are z-projections of 16 μm confocal optical sections, shown with dorsal side on top. Representative images are shown for at least 2 replicates tested. (C) Immunostaining for Wnt reporter *TCF/LefH2B-GFP*. Hypothalamic green fluorescence (below) views of yellow rectangles in bright field (above) view images of the same brain are shown, respectively. Images are whole mount ventral views with anterior side on top, acquired with the same setting for CON and CKO. Representative images are shown for at least 3 replicates tested. (D) Immunostaining for *Pmch* in the E14.5 hypothalamus. Higher magnification views of yellow rectangles are shown in the insets. Coronal images are z-projections of 16 μm confocal optical slices, shown with dorsal side on top. (E) qPCR analysis for female shows hypothalamic gene expression in E14.5 and P22 CKO-F relative to CON-F. Data are mean ± SEM. ***P < 0.001, ns. P > 0.05 by unpaired Student t tests. (F) Twenty-five μm coronal section in situ hybridization for *Crhbp* in the male P22 ventral premammillary and posterior hypothalamus, shown with dorsal side on top. Representative images are shown for at least 2 replicates tested. 3V: third ventricle. All scale bars are 100 μm except 500 μm in (F). Raw data can be found in S1 Data.

S6 Fig. Normal expression of *pmch* and *pmchl* in zebrafish *lef1* mutants. (A–C) Whole mount in situ hybridization images for *pmch* and *pmchl* (*pmchl*, like) in the hypothalamus of 3 dpf (A and B) and 15 dpf (C) zebrafish control and *lef1* mutant embryos. Images of dorsal views (anterior on top) and lateral views (dorsal on top and anterior on the left) of the same
individual \textit{lef1}+/− or \textit{lef1}−/− fish were shown in (A). Representative ventral view images of 3 dpf \textit{lef1}+/− (B), 15 dpf control and \textit{lef1} mutant (C) brains centered on the caudal hypothalamus (dashed red outlines) with anterior on top. Number of fish with representative gene expression among total number of fish is labeled on the right upper corner of each image in (C). Scale bar: 100 μm in (A and C); 5 μm in (B).

\textbf{S1 Table.} Details of confocal images, quantification and number of samples.

\textbf{S2 Table.} Zebrafish RNA-seq at 3 dpf.

\textbf{S3 Table.} Ingenuity Pathway Analysis (IPA) for diseases & functions.

\textbf{S4 Table.} IPA input gene list. Same zebrafish genes in Tab “AdjP<0.1” of \textbf{S2 Table} are listed with orthologous mouse gene information used for IPA.

\textbf{S5 Table.} Mouse RNA-seq at E14.5.

\textbf{S6 Table.} Mouse RNA-seq at P22.

\textbf{S7 Table.} Extracted GTEx RNA-seq data from 96 human hypothalamic samples.

\textbf{S8 Table.} Pearson correlation with hypothalamic GTEx RNA-seq data for \textit{PMCH}, \textit{CRHPR}, \textit{LEF1} and \textit{NPY}. Gene name followed with “\_r” indicates Pearson’s \(r\) value and gene name followed with “\_p” indicates \(P\) value.

\textbf{S9 Table.} Primer sequences for synthesizing in situ hybridization probes. Reverse primers also included a T7 promoter-containing sequence “CCAAGCTTCTAATACGACTCACTATAGGGAGA” that was added 5’ to the sequences listed in the table [70]. All primers were designed by Primer-BLAST except mouse genes \textit{Cartpt} (ABA experiment 72077479), \textit{Crhbp} (ABA experiment 77455017), \textit{Pmch} (GenePaint set MH227) and \textit{Tacr3} (ABA experiment 80342167).

\textbf{S1 Data.} Excel spreadsheet containing, in separate sheets, the underlying numerical data for all the figure panels.

\textbf{S1 Video.} One representative video of novel tank diving test. The resolution of the video was reduced from original 1080p to 540p, and its dimension was cropped to remove unnecessary space using software HandBrake.

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