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Lateral thalamic eminence – a novel origin for mGluR1/lot cells

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Running head: lot cells are born in the thalamic eminence.
A unique population of cells, called lot cells, circumscribes the path of the lateral olfactory tract (LOT) in the rodent brain and acts to restrict its position at the lateral margin of the telencephalon. Lot cells were believed to originate in the dorsal pallium (DP). We show that Lhx2 null mice that lack a DP show a significant increase in the number of mGluR1/lot cells in the piriform cortex, indicating a non-DP origin of these cells. Since lot cells present common developmental features with Cajal-Retzius cells, we analyzed Wnt3a- and Dbx1-reporter mouse lines and found that mGluR1/lot cells are not generated in the cortical hem, ventral pallium or septum, the best characterized sources of Cajal-Retzius cells. Finally, we identified a novel origin for the lot cells by combining in utero electroporation assays and histochemical characterization. We show that mGluR1/lot cells are specifically generated in the lateral thalamic eminence and that they express mitral cell markers, although a minority of them expresses ΔNp73 instead. We conclude that most mGluR1/lot cells are prospective mitral cells migrating to the accessory olfactory bulb, while mGluR1+, ΔNp73+ cells are Cajal-Retzius cells that migrate through the LOT to the piriform cortex and the olfactory bulb.

**Keywords:** accessory olfactory bulb, Cajal-Retzius cells, lateral thalamic eminence, lot cells, Tbr2.
Introduction

The olfactory circuit is one of the oldest sensory modalities in the phylogenetic history of mammals. Besides the detection of odorants, the olfactory system plays a significant role in the regulation of behavioral responses such as sexual behavior, the recognition of predators and the care of offspring (Halpern and Martinez-Marcos, 2003; Ashwell, 2012). The olfactory system is the only sensory system lacking a relay station in the thalamus (Kay and Sherman, 2007), and the information integrated by the principal neurons in the olfactory bulb (OB) is conveyed directly to the higher olfactory centers in the brain by axons that course along the lateral olfactory tract (LOT).

During development, mitral cells, the principal projection neurons of the OB, elongate their axons within a prospective LOT territory, which is populated by an early-generated population of neurons, the lot cells. The lot cells are considered guidepost cells for mitral cell axons, since chemically induced ablation of those cells prevents LOT formation in organotypic cultures (Sato et al. 1998; Hirata et al., 2012).

Lot cells supposedly derive from progenitor cells in the DP and migrate tangentially within the pallial marginal zone to enter into the LOT (Sato et al., 1998; Tomioka et al., 2000; Kawasaki et al., 2006; for reviews, see Marin et al., 2010; Bielle and Garel, 2013; Squarzoni et al., 2015). Lot cells are visualized using the monoclonal antibody (mAb) lot1 that recognizes the glutamate metabotropic receptor mGluR1 (Sato et al. 1998; Hirata et al., 2012). According to these authors, mGluR1 expression does not occur within the proposed migration territory of lot cells within the pallial marginal zone but only once the lot cells arrive at the LOT territory. However, the absence of specific cell markers precludes the identification of lot cells during their migration within the pallial marginal zone. Thus, it is unclear whether mGluR1/lot cells have a pallial origin.
as previously suggested or, on the contrary, if they are akin to the AP-2α/Lhx5 presumptive mitral cell destined for the posterior accessory olfactory bulb (pAOB) that arise in the thalamic eminence (Huilgol et al., 2013).

The pAOB is a structure that resides in the dorsocaudal region of the OB and forms a parallel yet independent pathway (Mohedano-Moriano et al., 2007) directly related to the neural control of aggressive and defensive behaviors, just as the anterior tier of the AOB (aAOB) serves sexual behaviors (Kumar et al., 1999). Unlike mitral cells of the main OB and the aAOB that are generated in the OB neuroepithelium, Lhx5/AP-2α presumptive pAOB mitral cells have their origin in the thalamic eminence (TE), a transient anatomical domain of the developing brain located at the rostral end of the diencephalon and topologically related to the telencephalic roof including the choroid plexus neuroepithelium and the cortical hem (Roy et al 2014). In addition to being a source of Lhx5/AP-2α presumptive pAOB mitral cells, the TE has been proposed as a neurogenic source of Trp73⁺ Cajal-Retzius (CR) cells that will go to the entorhinal and piriform cortex (Meyer et al., 2002, 2004; Tissir et al. 2009, Meyer, 2010).

Here we use a combination of genetic fate mapping in mice, in utero electroporation experiments and histochemical characterization to clarify the origins and neurochemical features of mGluR1/lot cells. Our data indicate that mGluR1/lot cells originate in the lateral tier of the TE (LTE) and migrate rostrally along the LOT. These cells are a heterogeneous population that includes a large group of prospective mitral cells that migrate to the pAOB and a small group of CR cells going to the piriform cortex and the olfactory bulb.
Materials and Methods

Animals

$\Delta Np73^{\text{Cre-IRESEGFP}}$ and B6.129P2-Grm1tm1Dgen/J mice and wild type C57BL/6J and ICR mice were maintained in the animal facilities of Universidad Miguel Hernández, San Juan de Alicante, Spain. Fixed $Lhx2^{-/-}$ mouse embryos and wild type littermates were from the TATA Institute for Fundamental Biology. Fixed $Wnt3a^{\text{IRCRE}}; A19^{\text{td-Tomato}}$ embryos were provided by Sonia Garel, École Normale Supérieure Paris, France, and $Dbx1^{\text{Cre}}; ROSA26^{\text{YFP}}$ embryos were provided by Alessandra Pierani, Institut Jacques Monod, CNRS, University of Paris Diderot, Paris, France. The care and handling of the animals prior or during the experimental procedures followed European Union (2010/63/UE) and NIH guidelines and were approved by the Animal Care and Use Committees of the participating institutions. The day a vaginal plug was detected was considered as embryonic day 0.5 (E0.5).

Immunohistochemistry and in situ hybridization

Embryos were fixed in 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer saline (PBS), pH 7.4, at room temperature (RT) for 2 h for immunohistochemistry or at 4°C overnight for in situ hybridizations, and rinsed in PBS for 2 h. For thick sections (100 μm), embryonic brains were embedded in 4% agarose diluted in PBS and cut in a Vibratome (Leica VT1000S). For thinner sections, brains were cryoprotected in 30% sucrose in 0.1M PB under agitation for 4 hours and frozen in liquid N$_2$ chilled isopentane. Immunohistochemical staining was performed as described (Espinosa et al., 2009) in 100 μm Vibratome sections. Primary antibodies used were: chicken anti-GFP (Aves Labs, 1:2000); rabbit anti-GFP (Abcam, 1:1000); rabbit polyclonal anti-DsRed (Clontech, 1:500); mouse monoclonal anti-reelin G10 (provided by A. Goffinet, 1:2000); mouse
monoclonal 3B5 anti-AP-2α (Santa Cruz 1:20); mouse monoclonal 4F2 anti-Lim1+2, i.e., Lhx1 + Lhx5 (Developmental Studies Hybridoma Bank, 1:200); rabbit anti-calretinin (Swant, 1:2000); rabbit anti-Tbr1 (Millipore, 1:500); chicken anti-Tbr2 (Millipore, 1:500); rabbit anti-COUP-TFI (provided by M. Studer, 1:500); rabbit polyclonal anti-β-galactosidase (Cappel, MP Biomedicals, 1:1000); rat monoclonal antibody lot1 (provided by T. Hirata, 1:200); rabbit polyclonal anti-Vglut2 (Synaptic Systems, 1:500); and mouse monoclonal antibody Tuj1 (Covance, 1:500). Different combinations of AlexaFluor secondary antibodies (Invitrogen, 1:500) were used.

*In situ* hybridization was performed as described (Martinez-Lopez et al., 2015). The following probes were used: *BMP4* (Jones et al., 1991), *Dbx1* (Lu et al., 1994), *Emx2* (Simeone et al., 1992), *Fgf15* (McWhirter et al., 1997), *Gdf10* (Zhao, Lawler et al., 1999), *Grml* (Zhu et al., 1999), *Gsh2* (Hsieh-Li et al., 1995), *Lhx5* (Zhao, Sheng et al., 1999), *Mash1* (Guillenot and Joyner, 1993), *Neurog2* (Gradwohl et al., 1996), *Neuropilin 2* (NP2; Tamamaki et al., 2003) *Pax6* (Walther and Gruss, 1991), *Sfrp2* (Kim et al., 2001), *Tbet* (Faedo at al., 2002), *Trp73* (Tissir et al., 2009), *Wnt3a* (Roelink and Nusse, 1991), and *Wnt8b* (Richardson et al., 1999).

**Slice cultures and CMFDA injections**

Timed pregnant dams were killed by cervical dislocation. Uterine horns were removed and isolated in cold Krebs solution. E12.5 brain embryos were embedded in 4% low melting point agarose (Sea Plaque Agarose, Cambrex) in PBS. 300 μm Vibratome coronal sections were transferred to Millicell CM culture plate inserts (Millipore) previously placed in 6-well culture plastic dishes (Nunc, Thermo Scientific) containing 1 ml of DMEM-F12 medium supplemented with N2 supplement (5 μl/ml), L-glutamine (0.1 mM), glucose (2.4 mg/ml), Penicillin–Streptomycin (500 U/ml) and 10% fetal bovine
serum (all these reagents provided by Invitrogen). Slices were maintained at 37°C in 5% CO₂ in a standard sterile incubator for 1 h. Next, resin beads (Bio-Rad), previously soaked in CellTracker CMFDA (Molecular Probes – Invitrogen) were placed on selected microanatomical areas of the slices. Then, DMEM-F12 medium was replaced by Neurobasal medium supplemented with B27 (1x), glucose (2.4 mg/ml), Penicillin – Streptomycin (500 U/ml) and L-glutamine (0.1 mM) and the slices were kept in the incubator at 37°C and 5% CO₂ for 2 days (Stoppini et al., 1991). Slices were then fixed in 4% PFA in PBS for 2 h and mounted on glass slides.

**Primary cultures**

TE and DP explants from E11.5 ICR wild type embryos were carefully dissected in chilled L15 medium (Gibco) supplemented with glucose (6g/L, Sigma), HEPES (5mM; Gibco), glutamine (2mM, Gibco) and Penicillin – Streptomycin (1x, Gibco). The TE and DP explants were incubated in 500 μl of differentiation medium DMEM/F12 (Gibco), glucose (6g/L Sigma), HEPES (5mM, Gibco), glutamine (2mM, Gibco), Penicillin – Streptomycin (1x, Gibco), N2 (1:100, Gibco), B27 (1:50, Gibco), FBS (5%, Gibco) and mechanically dissociated by repeated pipetting to isolate individual cells. 250,000 cells/well were incubated on laminin (5μg/ml, Sigma) and poly-lysine (100μg/ml, Sigma) coated coverslips and maintained in a sterile incubator at 37°C and 5% CO₂. Medium was daily replaced by 500 μl of fresh differentiation medium at 37°C. After 4 D.I.V., cells were fixed in 4% PFA in PBS at 4°C for 20 min.

**In utero electroporation**

The procedures were as previously described (Borrell et al., 2005; Garcia-Frigola et al., 2007) with some modifications. Plasmid pCAG–GFP (Matsuda and Cepko, 04; Addgene,
Teddington, Middlesex, UK) was purified with a midiprep endofree kit (Macherey-Nagel, Düren, Germany). The DNA solution (2µg/µl in PBS, with 0.05% Fast-green added) was injected in the third ventricle or in the lateral ventricle using pulled glass pipettes. Embryos were electroporated using tweezers-type electrodes. Five square electric pulses were passed at 1 s intervals (50 ms; 30V for E10.5 and 35 V for E11.5 embryos).

Quantification and statistical analysis

Images were captured with a digital camera coupled to a Leica MZ APO stereomicroscope or a Leica MD5000 fluorescence microscope. Confocal microscope analyses were carried out in a Leica TCS SP2 AOBS or an Olympus FluoView FV1200 Laser Scanning confocal microscope. Figures were prepared using Adobe Photoshop CS5 and Adobe Illustrator CS5, and 2D mosaic reconstructions were produced when needed using the Photomerge tool of Photoshop CS5 software package. A minimum of 3 animals and 3 slices of each animal were used for all the analyses and quantifications. InStat (GraphPad, San Diego, CA) software was used for statistical analysis.

Results

Comparative expression of lot cell markers

To assess the origin of lot cells, we first compared the expression patterns of several markers with known expression in the TE and/or the LOT areas: (i), the LIM-homeobox transcription factor Lhx5, expressed in future mitral cells (Huilgol et al., 2013) and also in CR cells (Abellan et al., 2010; Miquelajáuregui et al., 2010), (ii), the transcription factor Trp73, expressed by CR cells (Meyer et al., 2002, 2004; Yoshida et al., 2006; Tissir et al., 2009; Meyer, 2010) and, (iii), the metabotropic glutamate receptor mGluR1, expressed in lot cells when already located in the LOT territory (Hirata et al., 2012).
In situ hybridization in coronal sections of E12.5 wild type embryos revealed that Lhx5 was expressed in the TE, in the diencephalic-telencephalic boundary (DTB) and in the LOT territory (Fig. 1A1 – A3). Trp73 was expressed in the lateral part of the TE (LTE), a territory otherwise called strionuclear neuroepithelium (Altman and Bayer, 1995; Meyer, 2010) that is continuous with the caudal end of the cortical hem (hereafter, caudal hem), and showed a continuous distribution pattern from the LTE to the TE mantle, the DTB, the posterior part of the LOT (pLOT) and along the LOT territory (Fig. 1B1 – B3). Trp73 was expressed in postmitotic cells and also in progenitor cells of the ventricular zone of both the LTE and the caudal hem.

The expression of Grm1, coding for mGluR1, showed a pattern similar to Lhx5 and Trp73 in the TE and LOT (Fig. 1C1 – C3) with the notable exception that Grm1 was not expressed in the TE ventricular zone. At E11.5, Grm1 was already expressed in the TE, DTB and LOT territory (Suppl. Fig. 1). Unlike Grm1, at E12.5 Lhx5 and Trp73 were also expressed in the marginal zone of the cortex, likely in CR cells as previously reported (Yamazaki et al., 2004; Tissir et al., 2009; Abellan et al., 2010; Miquelajáuregui et al., 2010). Most notably, Grm1 and Lhx5 in situ hybridization signals overlapped in the LOT territory at different rostrocaudal levels (Suppl. Fig. 2). Monoclonal antibody (mAb) lot1 recognizes mGluR1 (Sato et al., 1998; Hirata et al., 2012) and we observed that Grm1 (mGluR1) in situ hybridization (Fig. 1D1 – D3) provided equivalent results to mAb lot1 immunostaining (Fig. 2A, B). Even further, reporter mGluR1 mice B6.129P2-Grm1<sup>tm1Dgen</sup>/J, where LacZ was inserted in frame with the mGluR1 coding sequence (Sachs et al., 2007), confirmed that mAb lot1 specifically recognizes the mGluR1 protein (Suppl. Fig. 3).

The vast majority of the mGluR1/lot cells do not originate in the dorsal pallium
It has been previously proposed that cells located at the LOT territory originate in the DP neuroepithelium (Tomioka et al., 2000; Kawasaki et al., 2006). We decided to re-examine the pallial origin of mGluR1/lot cells by analyzing mice lacking the transcription factor Lhx2 ($Lhx2^{-/-}$ mice). In this animal, whereas the DP is absent, the cortical hem and the ventral pallium (VP) are considerably enlarged (Bulchand et al., 2001; Monuki et al., 2001; Vyas et al., 2003; Mangale et al., 2008; Subramanian and Tole, 2009; Subramanian et al., 2009, Roy et al., 2013, 2014), and the TE is aberrantly expanded (Roy et al., 2014).

In $Lhx2^{-/-}$ mice, the OB mitral cells are specified but they reside in an ectopic olfactory bulb-like structure (OBLS) in which mitral cells are not able to pioneer their axonal projections towards the LOT (Saha et al., 2007). Additionally to these previously reported observations, we found that AOB marker neuropilin2 (NP2), the aAOB marker Tbet and the pAOB marker $Lhx5$ were all expressed in the OBLS region (Suppl. Fig. 4). In contrast, the $Lhx2^{-/-}$ brain displayed a considerably enlarged lot1 (mGluR1) immunoreactive territory. Compared to E12.5 wild type mice (Fig. 2A – B), $Lhx2$ mutants showed an intensely lot1-immunoreactive TE field that expanded along the DTB, the pLOT and the LOT territories at mid-rostrocaudal levels (Fig. 2C – D). The expansion of the mGluR1 expression domains remained conspicuous in E14.5 $Lhx2^{-/-}$ mice (Fig. 2E, F). We based our identification of the enlarged TE in $Lhx2^{-/-}$ mutant mice first on the intense calretinin immunostaining of the TE mantle (Fig. 2C'; see Abbott and Jacobowitz, 1999) and, second, on the presence of a Tbr2-immunostained TE subventricular zone encircling the lot1+ TE mantle (Fig. 2E'). Tbr2 also decorated the subventricular zone of the VP and highlighted the enlarged cortical hem of these animals (Fig. 2E’').

As recently described by Roy et al. (2013), $Lhx2^{-/-}$ mice display an abnormal optic-like vesicle at the DTB (Fig. 2C, C', D, G, arrows). This vesicle was calretinin positive (Fig 2C'), and combined lot1/calretinin immunostaining (Fig. 2C') or Grm1 in
situ hybridizations (Fig. 2G) showed that the putative migration path of mGluR1 TE-derived cells surrounded the optic-like vesicle, which in the mutants is anatomically very close to the pLOT territory.

In conclusion, because the Lhx2−/− mice lack a DP, it is very unlikely that the mGluR1/lot cells in the LOT territory originate in the neocortical primordium.

**mGluR1/lot cells do not originate from Wnt3a progenitors in the hem, or from Dbx1 progenitors in the ventral pallium or the septum**

We then wondered whether mGluR1/lot cells originate from the hem. To test this hypothesis, we used a Wnt3aCre mouse line (Yoshida et al 2006) crossed to Ai9TdT-Tomato mice to drive the expression of the Td-Tomato reporter in Wnt3a-lineage cells. We did not find Wnt3a-lineage, td-Tomato cells in the TE mantle or in the LOT territory (Fig 3A, A’). At E12.5, we observed strong expression of td-Tomato in the cortical hem and in the marginal zone of the DP that corresponds to cortical hem-derived CR cells (Fig. 3A, B). At rostral levels, there were scattered Wnt3a/td-Tomato cells in the LOT territory, but they were not mGluR1 positive (Fig 3B, B’). This result demonstrates that Wnt3a hem progenitors are not the precursors of mGluR1/lot cells.

Another possible source of lot cells could be the VP or the septum. To test this possibility, we used a Dbx1Cre mouse line (Bielle et al., 2005) crossed to ROSA26RYFP mice to detect Dbx1-lineage cells. Dbx1 is a homeodomain transcription factor expressed in progenitor cells in the VP and pallial septum that generate glutamatergic neurons destined to the cortex and amygdala (Bielle et al., 2005; Hirata et al., 2009; Teissier et al., 2010, Griveau et al., 2010; Zimmer et al., 2010). In addition, the ventricular zone of the medial tier of the TE displayed strong Dbx1 expression (see Suppl. Fig. 5C). Dbx1-lineage cells distributed widely in the hypothalamus and the basal telencephalon, but very
few of them invaded the LOT territory (Fig. 3C). In the posterior LOT, we found that only $3.3 \pm 0.85 \%$ (mean $\pm$ s.e.m.) of mGluR1/lot cells displayed Dbx1-YFP immunoreactivity (Fig. 3C, C’, white arrowheads). At rostral levels, the percentage of mGluR1/lot cells displaying Dbx1-YFP immunoreactivity was also very low ($4.16 \pm 0.76 \%$) (Fig 3D, D’, white arrowheads). These results indicate that Dbx1 progenitors from the VP or the pallial septum are not the source of mGluR1/lot cells.

**The TE has the potential to produce lot cells**

Given that the mGluR1/lot cells do not originate from the DP, hem, VP, or septum we considered the possibility that they are generated in TE. To explore the potential of the TE to produce lot cells, we dissociated cells from TE or DP from E11.5 embryos ($n = 17$) and put them in culture ($n = 3$). After 4 days-in-vitro (D.I.V.), cells were immunostained for Tuj1 and mGluR1 using mAb lot1 (Fig. 4A, B) and a minimum of 500 Tuj1$^+$ newly differentiated neurons were counted. In the case of TE-derived primary cultures, $19.5 \pm 2.4 \%$ of the Tuj1$^+$ cells also expressed mGluR1, while in the case of pallial cultures only $0.5\% \pm 0.2 \%$ of the Tuj1$^+$ cells were also mGluR1$^+$ (Fig 4A – C). This experiment demonstrates that the E11.5 TE has the potential to generate mGluR1 cells and confirms that the dorsal pallium has a very limited capacity of generating lot cells at this embryonic stage (Tomioka et al., 2000), consistent with the results obtained with the $Lhx2^{-/-}$ mice.

Organotypic slice cultures (Stoppini et al., 1991) provided a further line of evidence in favor of a TE origin for mGluR1 cells in the LOT territory. We placed a minute crystal of *CellTracker* CMFDA over the cross-sectioned TE in coronal slices from E12.5 wild type embryos and cultured them for 24h. CMFDA-labeled cells were then detected in the DTB and in the LOT (Fig. 4D), and mAb lot1 immunostaining revealed that the CMFDA labeled cells were mGluR1$^+$ (Fig. 4D – D”).
The thalamic eminence expresses genes associated with glutamatergic neurogenesis

The prethalamus is the neurogenic site of GABAergic neurons that will populate the thalamus in the adult brain (Inamura et al., 2010) and thus expresses genes encoding transcription factors involved in GABAergic neurogenesis such as *Mash1* and *Gsh2* (Fig. 5A, B). However, we observed that, unlike the prethalamus, the TE is negative for genes related to GABAergic neurogenesis (Fig. 5A – B).

The TE was, by contrast, positive for genes that participate in glutamatergic neurogenesis such as *Pax6*, *Neurog2* and *Tbr2* (Schuurmans and Guillemot, 2002; Mattar et al., 2004;Englund et al., 2005; Hevner et al., 2006). The paired domain transcription factor *Pax6* was expressed in the ventricular zones of prethalamus, TE (with a medial\textsuperscript{high}–lateral\textsuperscript{low} gradient), and pallium (Fig. 5C). The proneural gene *Neurog2* was expressed in the TE ventricular zone (Fig. 5D) and the T-box transcription factor *Tbr2* was expressed in the TE subventricular zone (Fig 5E). We also observed that cell populations in the TE mantle expressed Vglut2, calretinin and Tbr1 (Fig 5F – H), three markers of glutamatergic neurons (Bulfone et al., 1995; Abbott and Jacobowitz, 1999; Ina et al., 2007). All these traits of molecular identity are also compatible with the idea that the TE is a potential source of lot cells.

*mGluR1* cells are generated in the lateral TE

While analyzing the expression of glutamatergic markers in the TE we realized that the TE encompasses two well-differentiated neuroepithelial compartments, i.e., the medial TE (MTE) and the lateral TE (LTE), which surround a centrally located mantle zone. In order to properly characterize the TE, we decided to carefully define the expression patterns of genes that have been previously identified in this brain region. We found that certain genes were expressed in the entire TE, while some others highlighted specific TE
subdomains, sometimes also encompassing hem subregions (Suppl. Figs. 5 and 6). In particular, \textit{Wnt3a} expression delimited the complete cortical hem and its expression domain ceased at the point where the caudal hem continues with the TE (Suppl. Fig. 5B, D1 – D3). Similarly, \textit{BMP4} expression highlighted the cortical hem (Suppl. Fig. 6A). By contrast, \textit{Wnt8b} (Suppl. Fig. 5E1 – E3) and \textit{Emx2} (Suppl. Fig. 6B) expressions revealed the existence of a genoarchitectonic continuum between the caudal hem and the LTE, although the \textit{Emx2} expression domain extended far more dorsally than \textit{Wnt8b}. Other genes, such as \textit{Sfrp2}, a secreted modulator of Wnt signaling (Suppl. Fig. 5C, F1 – F3), and the transcription factor \textit{Dbx1} (Suppl. Fig. 6C), delimited the neuroepithelium of the MTE. Interestingly, \textit{Sfrp2} and \textit{Dbx1} also labeled the VP at the pallium-subpallium boundary. \textit{Gdf10 (BMP3b)} (Suppl. Fig. 5G1 – G3; Shimogori et al., 2010) and \textit{Fgf15} (Suppl. Fig. 6D) were detected in both the LTE and the MTE ventricular zones but not so in the cortical hem.

Therefore, the differential anatomical distributions of these markers indicate that the MTE and the LTE have different molecular identities and support the hypothesis that they might be programmed to generate distinct sets of neuronal populations.

\textit{In utero} electroporation allows gene targeting to specific locations of the developing brain (Borrell et al., 2005; García-Frigola et al., 2007). Therefore, we used this approach to express eGFP in progenitors residing in the developing TE of E11.5 embryos. By manipulating the position and polarity of the electrodes, different subdomains of the TE were targeted (Fig. 6 and Suppl. Fig. 7). When the MTE region was targeted by electroporation, no eGFP$^+$ cells were detected in the LOT territory or the ventral telencephalon at E14.5 (Suppl. Fig. 7). However, when progenitors in the LTE were targeted, groups of eGFP$^+$ cells in the TE mantle were systematically found at E14.5 together with a larger population of labeled cells scattered along the DTB migratory path,
the pLOT (Fig. 6B2, C2) and the LOT (Fig. 6B4, C3, C4). eGFP+ cells derived from the TE localized around the LOT and those located in the DTB and in the LOT were mGluR1+ (Fig. 6B2', C4'). In sum, these experiments showed that mGluR1/lot cells generated in the LTE at E11.5 migrate to the TE mantle and then ventrally to the posterior LOT to subsequently turn rostrally along the LOT. In agreement with this interpretation, whole mount in situ hybridizations for Grm1 (Fig. 6D – G) revealed the progressive invasion of the pLOT by mGluR1+ cells between E11.5 and E13.5.

mGluR1/lot cells exhibit neurochemical features typical of pAOB mitral cells

Our observations suggested that a majority of mGluR1/lot cells may be prospective pAOB mitral cells, similar to the cells described by Huilgol et al. (2013) and therefore, we explored whether mGluR1/lot cells have neurochemical properties in common with mitral cells. To this end, we performed double immunostainings for mGluR1 and reelin or Tbr2, which are general markers of mitral cells, and for mGluR1 and COUP-TFI, Lhx5, or AP-2α, which are selective markers of pAOB mitral cells (Suppl. Fig. 8; see also, Huilgol et al., 2013).

Coronal sections at different rostrocaudal levels of E12.5 embryos showed that reelin was strongly expressed in the LOT territory, but not along the migratory route between the TE and the pLOT (Fig. 7A). In the pLOT, 70.7 ± 1.5% of the mGluR1/lot cells co-expressed reelin (Fig. 7A’, F) while in the aLOT the percentage was 89.74 ± 3.04% (data not shown). mGluR1/lot cells along the DTB co-expressed Tbr2 (Fig. 7B). In the pLOT territory, 69.8 ± 4.25% of the mGluR1/lot cells co-expressed Tbr2 (Fig 7B', F), similar to the aLOT (76.10 ± 4.04%; data not shown). Interestingly, Tbr2 expression discriminated the Tbr2+ mGluR1/lot cells from the presumed Tbr2− CR cells also present in the LOT (Figs. 8A' and 9A', B', C – C').
COUP-TFI is a transcription factor expressed in the pAOB at E15.5 (Suppl. Fig 8B, Fig. 8F, F’) and E18.5 (Fig. 8G, G’). At E12.5, 74.25 ± 4.37 % of mGluR1/lot cells in the pLOT territory co-expressed COUP-TFI (Fig. 7C, C’, F) vs. 59.56 ± 3.17% in the aLOT (data not shown). We also observed that the majority of the lot cells were positive for the mAb 4F2 that recognizes Lhx1/5 in both the pLOT (95.41 ± 0.91 %; Fig 7D, D’, F) and the aLOT (92.39 ± 1.47 %; data not shown).

At E12.5, we found mGluR1/lot cells positive for AP-2α in the aLOT (Suppl. Fig. 9B’, C’, white arrowheads), but not in the pLOT (Fig 8E, E’ and Suppl. Fig. 9A, A’). Within the AP-2α population in the aLOT, we also found cells co-expressing AP-2α and COUP-TFI (Suppl. Fig 9B’, C’, yellow arrowheads).

Altogether, these results indicate, first, that the Tbr2+ mGluR1/lot cells co-express reelin and pAOB mitral cell markers such as COUP-TFI, Lhx5, or AP-2α and, second, that the AP-2α cells are a subpopulation of the mGluR1/lot cells.

**Most mGluR1/lot cells follow a rostral migratory path to finally populate the pAOB**

The number of Tbr2+/mGluR1+ cells in the LOT decreased from E13.5 to E18.5 (Fig. 8A’ – C’, D). By E13.5, when the axons of the OB mitral cells have already formed the LOT, there was an average of 37.2 ± 2.25 Tbr2+ mGluR1/lot cells per section around this axonal tract at mid-rostrocaudal levels. This number significantly dropped to 8.9 ± 1.03 (q = 20.15, P<0.001) by E15.5 (Fig 8A’, D), a result compatible with the hypothesis that these cells migrate rostrally.

AP-2α and COUP-TFI immunolabeling distinguished two groups of mGluR1+ mitral cells in the AOB. At E13.5, when the OB evagination was already apparent, we detected abundant mGluR1+/AP-2α+ cells in the pAOB (Fig 8E, E’, arrows, and Fig.
At this early stage, there were also some (8.83 ± 1.80 cells per section) mGluR1+/COUP-TFI+ cells negative for AP-2α in the pAOB (Fig 8E’, white arrowheads). At subsequent stages, while the amount of AP-2α+/mGluR1+ cells was constant in the pAOB (Fig. 8H), the number of AP-2α-negative/mGluR1+/COUP-TFI+ cells (marked in Figs. 8F’, G’ with white arrowheads) increased significantly from E13.5 to E15.5 (Fig. 8F’, G’, I). These results strongly suggest the existence of an early arriving wave of mGluR1/AP-2α expressing neurons that complete their migration to the pAOB by E13.5 – E14.5 (in agreement with the data by Huilgol et al., 2013), and a second wave of mGluR1/COUP-TFI expressing cells that reach the pAOB at later stages, between E15.5 and E18.5.

**A small percentage of mGluR1/lot cells express p73 and do not integrate as mitral cells in the AOB**

A recent article by Dixit el al. (2014) suggests that mGluR1/lot cells are p73 immunoreactive, and thus propose that lot cells are a subset of CR cells associated with the LOT. Given that our results show that most mGluR1/lot cells express Tbr2 (Fig. 9C – C’’), which is not a marker of CR cells, we decided to assess the degree of co-expression of p73 in mGluR1/lot cells to clarify this issue. To this end we used a ΔNp73Cre−IRESeGFP reporter mouse line (Tissir et al., 2009) in which eGFP expression recapitulates Trp73. It has been shown that ΔNp73, the N-truncated isoform of Trp73 (Meyer et al., 2004; Tissir et al., 2009), is expressed in cortical hem, septum (Meyer et al., 2002, 2004, Hernandez-Acosta et al., 2010) and TE-derived CR cells (Tissir et al., 2009), while it is not so in VP-derived CR cells (Hanashima et al., 2007). So, in the TE, ΔNp73-eGFP was expressed in the ventricular zone and the subpial side of the lateral TE (LTE), the TE mantle and the putative migration path along the DTB to the pLOT (Fig. 9A), reflecting the distribution
of the *in situ* hybridization signals in the same territories (Fig. 1B3, B3'). ΔNp73-eGFP immunoreactivity and *Trp73* transcripts were also observed in the caudal cortical hem (Figs. 1B, B', 10A). We found that in ΔNp73Cre−IRESeGFP mice only a minority of E12.5 mGluR1/lot cells co-expressed ΔNp73 indicating that most mGluR1/lot cells are not CR cells. In the pLOT territory, only 15.36 ± 2.76 % of the mGluR1/lot cells were ΔNp73-eGFP immunoreactive, quite similar to the 17.97 ± 3.13 % detected in the LOT territory at mid-rostrocaudal levels (Fig. 9A’, B’, white arrowheads). Moreover, ΔNp73-eGFP cells were in all instances Tbr2-negative unlike mGluR1/lot cells destined to the pAOB (Figs. 9C – C’). While mGluR1/lot cells tended to locate deeper within the LOT territory, most ΔNp73-eGFP+ CR cells accumulated near the subpial side of the LOT at E12.5 (Fig. 9A, B). At subsequent developmental stages (E15.5 and E18.5), mGluR1/ΔNp73 cells extended into the piriform cortex but they never invaded the neocortex primordium (Fig. 9B – B”, C – C’). At E15.5, some of the mGluR1/ΔNp73 cells (Suppl. Fig. 10A’, B’, white arrowheads) reached the olfactory bulb and located around (but not within) the mitral cell layers (Suppl. Fig. 10A – C).

We also noticed that nearly half of the ΔNp73-eGFP positive cells did not co-express mGluR1 (52.98 ± 1.98 %) (Fig. 9A’, B’, hollow arrowheads) and, therefore, by definition they should not be considered lot cells. However, p73 cells of any of these two groups, mGluR1-positive or mGluR1-negative, likely are CR cells. These data reveal that in contrast to what has been previously proposed, only a small percentage of the mGluR1/lot cells arising in the LTE are CR cells.

**Discussion**

As demonstrated by Sato et al. (1998), an early-generated population of neurons, the lot cells, act as guidepost cells for mitral cell axons along the LOT. Here we demonstrate
that, in contrast to the current belief, the lot cells predominantly originate from the TE, a transient structure localized in the boundary between the telencephalon and diencephalon, and specifically from its lateral region, the lateral thalamic eminence (LTE). The present findings highlight the TE as a neurogenic hub of diverse neuronal sets destined to different telencephalic domains that include CR cells (Meyer et al., 2002; Takiguchi-Hayashi et al., 2004; Tissir et al., 2009; Roy et al., 2014), Lhx5/AP-2α prospective pAOB mitral cells (Huilgol et al., 2013), and COUP-TFI+ mGluR1/lot cells that also invade the pAOB (present results).

**The thalamic eminence as a primary source of mGluR1/lot cells**

Tomioka et al. (2000) and Kawasaki et al. (2006) cultured pallial and subpallial progenitor cells at E10.5 and found that the pallial progenitors, unlike the subpallial ones, could differentiate into the mGluR1-positive lot cells described previously by Sato et al. (1998). We replicated these experiments at E11.5 including cells derived from the TE. Similar to Tomioka et al. (2000), we detected a small proportion of cells derived from the E11.5 pallium that express mGluR1 in culture. However, the percentage of mGluR1 cells derived from the TE was almost 20-fold higher than that of mGluR1 cells derived from the pallium. Thus, our results show that the E11.5 TE progenitor cells have the potential to produce mGluR1 cells.

Migration experiments performed in whole embryo cultures (Tomioka et al., 2000) and in organotypic cultures (Kawasaki et al., 2006; Ito et al., 2008) supported the view that mGluR1/lot cells are generated in the DP and then migrate tangentially within the pallial marginal zone to reach the LOT territory in the piriform cortex, prior to the arrival of mitral cell axons. Lot cells share this purported pallial migration path with several early-differentiated neurons such as preplate cells from the rostromedial
telencephalic wall (Pedraza et al., 2014), CR cells from the cortical hem and pallial septum (Yoshida et al., 2006; Zhao et al., 2006, Garcia-Moreno et al., 2007; Zimmer et al., 2010) and olfactory cortex neurons from dorsal pallium (Nomura et al. 2006; Garcia-Moreno et al., 2008). All these neurons migrate ventrally within the pallial marginal zone (MZ) to reach the LOT territory, i.e. the external pallial-subpallial boundary. However, although Tomioka et al. (2000), Kawasaki et al. (2006) and Ito et al. (2008) clearly demonstrated a ventrally directed tangential migration within the pallium marginal zone at E10.5 – E11.5, the hypothesis that these migrating cells were lot cells could not be ascertained in their experiments because these cells were non-immunoreactive to mAb lot-1. Moreover, the present study confirms the absence of mGluR1 expression in the pallial marginal zone at the stages considered.

In contrast, we report here that mGluR1 expression occurs in a continuous domain that comprises the TE, the DTB, and the LOT territory as early as E11.5 (Suppl. Fig. 1), which strongly suggests a TE origin for these cells.

Lhx2 null mice, which lack a DP and display an enlarged TE and ventral pallium (Vyas et al., 2003; Roy et al., 2014), provided further evidence that mGluR1/lot cells originate from the TE similar to the prospective pAOB mitral cells described by Huilgol et al. (2013). In Lhx2 /− mice, the spread of mGluR1/lot cells in the piriform cortex was considerably larger than normal and, therefore, it is unlikely that these cells come from the DP. In a similar vein, the OB is specified as an olfactory bulb like structure (OBLS) in Lhx2 /− mice, in which mitral cell axons failed to pioneer the LOT projection. This anomaly is compounded by defects in axon guidance associated to the described mislocalization of the lot guidepost cells (Saha et al., 2007). However, the present findings show that lot cells are exuberant in Lhx2 /− mice. Intriguingly, Saha et al. (2007) detected in these mutants a notable overexpression of Sema6A in a rostrolateral region of
the telencephalon that topographically corresponds to the expanded mGluR1 immunoreactive domain described in the present report, and we found that lot cells express Sema6A in wild type mice (not shown). It remains to be seen in a future study how lot cell exuberance, the overexpression of Sema6A or other anomalies affect axon guidance along the LOT in Lhx2\(^{-/-}\) mice.

An argument that has also been used to suggest a DP origin for the lot cells was the phenotype exhibited by the Gli3 null mice Gli3\(^{ Xu/Xu}\). At E13.5, these animals show subpial clusters of mGluR1/lot cells limited to the DP (Tomioka et al., 2000) together with a convincing reduction of tangential cell migrations in the pallial surface attributed to the absence of Gli3 function (Kawasaki et al., 2006). The authors proposed that the lack of Gli3 function arrested the migration of lot cells originated in the pallium, causing these cells to stall abnormally within the pallium itself instead of reaching the LOT territory. However, contrasting with this interpretation, it has been amply demonstrated that the choroid plexus, the cortical hem and the medial pallium are missing in Gli3 null mice juxtapositioning the DP and the TE (Grove et al., 1998, Theil et al., 1999, Tole et al., 2000). Moreover, the poorly developed DP contains a mixture of dorsal pallial, ventral pallial and TE cells (Fotaki et al., 2006; Friedrichs et al., 2008). Furthermore, lot cells are correctly positioned in the piriform cortex in a conditional Gli3 mouse (Emx1\(^{Cre};\) Gli3\(^{fl/fl}\)) where the pallium lacks Gli3 expression (Amaniti et al., 2015). This finding supports the interpretation that lot cells originate from outside the Emx1\(^{+}\) dorsal pallium. Even further, our own unpublished data indicate that the Grm1\(^{+}\) clusters in the pallium of Gli3 null mice correspond to TE ectopias resulting from a faulty specification of the TE (NR, TT, AF, ms. in preparation). Thus, these new findings clearly discard the cortical hem and the medial pallium, but also the DP, as possible origin of mGluR1/lot cells.
Additionally, we explored the hem and the VP as potential sources of lot cells using two reporter mouse lines for Wnt3a and Dbx1 progenitors in the hem, VP and septum, and demonstrated that mGluR1 cells do not arise from these progenitors. Finally, \textit{in utero} electroporation provided decisive evidence that mGluR1/lot cells derive from neural progenitors in the lateral domain of the TE, the lateral thalamic eminence (LTE).

\textbf{The mGluR/lot cells are a heterogeneous cell population}

Our results indicate that most mGluR1/lot cells are akin to the cells described by Huilgol et al. (2013) because they both express mitral cell markers and incorporate into the pAOB. Moreover, the \textit{in utero} electroporation experiments together with our immunohistochemical results suggest that there are at least two waves of lot cells generation: an early wave of cells that reach the pAOB at E13.5, which seems to be the AP-2α/Lhx5 cell population previously described by Huilgol et al. (2013), and a late wave of mGluR1/COUP-TFI cells that reach the pAOB at later stages.

On the other hand, we also found a population of mGluR1/lot cells in the LOT territory that express p73 and, therefore, may be considered as CR cells. These LOT-associated CR cells migrate rostrally to reach the AOB but they never integrate into this structure as mitral cells. Instead, most of these cells spread over the paleocortical primordium by E18.5 without invading the neocortical primordium. This result differs from that of Dixit et al. (2014), who proposed that almost all the mGluR1/lot cells are p73 positive and, therefore, that lot cells are a subset of CR cells. The evidence shown by these authors was based on p73 immunohistochemical experiments performed with a p73 antibody whose specificity was not adequately tested. In contrast, we have performed cell lineage experiments using the ΔNp73\textsuperscript{eGFP} mice and shown that eGFP labeling in these mice closely matched the \textit{in situ} hybridization images for \textit{Trp73}. Previous studies
defining the expression and functional roles of p73 in CR cell biology (Meyer et al., 2002, 2004) have noted that the expression of p73 is a characteristic of most subtypes of CR cells, and only CR cells derived from the VP lack p73 expression (Hanashima et al., 2007).

In summary, we conclude that the mGluR1/lot cells are heterogeneous and form at least two different cell populations. First, a large group of Tbr2+ cells that are prospective pAOB mitral cells and that is divided into at least two subpopulations that migrate into the pAOB in two partially overlapping waves: the early arriving AP-2α+ cells and the late arriving COUP-TFI+ cells (Fig. 10). Second, ΔNp73+/mGluR1+ cells positioned around the LOT territory and in the marginal zone of the piriform cortex that, together with the ΔNp73+/mGluR1− cells likely are CR cells.

**Have the lot cells a diencephalic or a telencephalic origin?**

All our results unambiguously demonstrate that most lot cells originate in the TE. However, it is not clear whether the TE should be considered as a diencephalic or a telencephalic structure. Our chemoarchitectonic observations summarized in Figs. 5 and Suppl. Fig. 6 and 9 highlight several remarkable properties that the TE has in common with the telencephalic pallium. On the one hand, the TE displays a Tbr2-positive subventricular zone that is anatomically continuous with the pallial SVZ (Fig. 5E), and the TE mantle expresses Tbr1 as the cohorts of pallium-generated earliest neurons (Englund et al., 2005). Second, the LTE and the telencephalic caudal cortical hem share anatomically continuous gene expression domains such as Wnt8b (Suppl. Fig. 5E1 – E3) and Emx2 (Suppl. Fig. 6B). Third, the MTE also displays gene expression features in common with the pallium such as Pax6, Neurog2, Dbx1, and Sfrp2. In fact, the caudal hem and the LTE are one long piece of Wnt2b- and Neurog2-expressing tissue that
generates, and is divided by, the lateral ventricle choroid plexus (see Figs. 1 and 2 in Grove et al., 1998). Such an anatomical continuum caudal hem – LTE was also suggested by Gorski et al. (2001) in their Suppl. Fig. 1. Moreover, Yoshida et al. (2006) found that crossing a \textit{Wnt3a}-floxed \textit{dt} mouse with an \textit{Emx1}-Cre mouse produced progeny lacking the hem – except in a very caudal region (see Yoshida et al., 2006, their Fig. 3B) that likely corresponds to the LTE – caudal hem continuum. A very caudal tip of choroid plexus in the lateral ventricle is also present in the mutant, most likely generated from this region that could deserve to be called telencephalic. However, given that these arguments only suggest, but do not prove, a telencephalic identity for the TE, we conclude that the attribution of a telencephalic origin to the LTE-derived cells present in the LOT cannot be considered definitive and needs further research.

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Figure legends

Abbreviations used in Figures: aAOB, anterior accessory olfactory bulb; AOB, accessory olfactory bulb; C-hem, caudal cortical hem; CGE, caudal ganglionic eminence; cp, choroid plexus; D-hem, dorsal cortical hem; DB, diagonal band; DTB, diencephalic-telencephalic boundary; IC, internal capsule; LGE, lateral ganglionic eminence; LOT, lateral olfactory tract; LTE, lateral thalamic eminence; MGE, medial ganglionic eminence; MOB, main olfactory bulb; MTE medial thalamic eminence; OB, olfactory bulb; OBLS, olfactory bulb-like structure; pAOB, posterior accessory olfactory bulb; pCx, piriform cortex; pLOT posterior end of the LOT; pTh, prethalamus; SE, septum; SVZ, subventricular zone; TCA, thalamocortical axons; TE, thalamic eminence; Th, thalamus; VP, ventral pallium; ZLI, zona limitans intrathalamica.

Figure 1. Comparing expression patterns of three genes expressed in the thalamic eminence and the LOT territory: Lhx5, Trp73 and Grm1 (coding for mGluR1). Coronal sections of E12.5 wild type embryos at three rostrocaudal levels.

A1 – A3'. Lhx5 is expressed in the septum, the nuclei of the diagonal band in the basal forebrain, the whole thalamic eminence, and the supraventricular domain of the hypothalamus (asterisk in A3). A3' is a close-up image of the boxed area in A3 that displays Lhx5 expression in the lateral TE (LTE), including its ventricular zone. Arrowheads point to the subpial side of the LTE.

B1 – B3'. Trp73 is expressed in the ventricular zone of the pallial septum (B1, arrow), the dorsal cortical hem (B2), the choroid plexus (B2, B3), the caudal cortical hem, and the lateral TE (B3, B3'). In B3, the ventricular zones of the dorsal hem, the caudal hem and the LTE display Trp73 positive progenitors, whereas the MTE ventricular zone lacks
Trp73 progenitors. Note subpial accumulations of transcripts in the dorsal and caudal hem and in the lateral TE (arrowheads in B3'). The distribution of discrete transcript clumps suggests the invasion of the TE mantle by p73 cells derived from the lateral TE (in B3).

C1 – C3'. Compared to Lhx5 and Trp73, Grm1 expression is spatially much more limited, but is outstanding in the TE. Note in C3' the absence of Grm1 expression in the ventricular zone of the LTE, and the modest accumulation of transcripts in the subpial side of the LTE (arrowheads in C3').

Note, for all these three genes, the strong hybridization signal in the LOT at all rostrocaudal levels, and the presence of the respective transcripts within the space between the lateral hypothalamus and the caudal ganglionic eminence (the diencephalic-telencephalic boundary, DTB), suggesting a putative migrating path between the thalamic eminence and the posterior end of the LOT (pLOT).

Scale bars: A1 – C3, 100 μm; A3' – D3', 50 μm.

**Figure 2. Absence of dorsal pallium in Lhx2± mice does not prevent an increased invasion of the piriform cortex by mGluR1/lot cells**

A – D. mGluR1 immunohistochemistry with mAb lot1 in coronal sections of E12.5 wild type (A, B) and Lhx2 knockout littermates (C, D) at two different rostrocaudal levels. The immunohistochemical signal was more extended and more intense in the TE and LOT of the Lhx2± mice than in the wild type embryo. Arrows in C, C', D, G indicate the abnormal optic-like vesicle located at the DTB, typical of these mutants. Asterisks signal the presence of artefactual staining over the meninges.
C'. From the boxed area in C. Calretinin immunostaining in the putative migration path from the TE to the pLOT in \( Lhx2^{-/-} \) mice. Note the co-expression of mGluR1 and calretinin except in the optic-like vesicle, which is calretinin⁺ but mGluR1⁻.

G. *In situ* hybridization shows *Grm1* transcripts surrounding the optic-like vesicle in a section similar to that in C'.

E, F. mGluR1 immunostaining in E14.5 \( Lhx2^{-/-} \) mouse embryos shows the notable enlargement of the putative migration pathway of mGluR1 cells along the DTB to the posterior end of the LOT (pLOT, in E) and in the LOT at mid-rostrocaudal levels (in F).

E'. From the boxed area E' in E. Tbr2-immunoreactive subventricular zones surround the mGluR1⁺ mantle of the thalamic eminence and serve to corroborate its TE identity.

E''. From the boxed area E'' in E. Tbr2 immunoreactive secondary progenitor cells in the expanded VP and the cortical hem, both known features of \( Lhx2^{-/-} \) mice. A white line signals the limit between these two structures.

All calibration bars, 100 µm. Bar in A applies to A – F; in E’, the bar applies to E’ – E”.

**Figure 3.** mGluR1/lot cells do not derive from Wnt3a progenitors in the cortical hem or Dbx1 progenitors in the ventral pallium.

A, B. Coronal sections of an E12.5 \( Wnt3a^{Cre};Ai9^{td-Tom} \) mouse embryo. td-tom⁺ cells are Wnt3a-lineage neurons that concentrate in the dorsal cortical hem. Td-Tom⁺ cells were conspicuous in the pallial preplate, but virtually no td-Tom⁺ cells were found in the LOT territory.
A', B'. Close-ups of the boxed areas in A, B show the absence of td-Tom$^+$ cells in the pLOT and occasional td-tom$^+$ (but mGluR1$^-$) cells at more rostral levels of the LOT, thus discarding that Wnt3a progenitors in the cortical hem could generate mGluR1/lot cells.

C, D. Coronal sections of an E12.5 $Dbx1^{Cre,R26R^{YFP}}$ mouse embryo. Dbx1-lineage, YPF$^+$ cells were abundant in the TE, in the basal telencephalon mantle and in the septum, but very scarce in the LOT territory.

C', D'. High magnification images of the boxed areas in C, D show that occasional Dbx1-lineage cells may co-express mGluR1 (white arrowheads) both in the pLOT and at rostral levels of the LOT (see text for details).

Scale bars: A – D, 100 μm; A’ – D’, 50 μm.

Figure 4. The E11.5 TE has the potential to generate mGluR1 cells.

A, B. Primary cultures of E11.5 cells from DP (A, A') and TE (B, B') after 4 D.I.V. In both cases, we counted the percentages of cells co-expressing mGluR1$^+$ and Tuj1$^+$ cells over the total of Tuj1$^+$ young neurons.

C. The percentages of mGluR1-expressing neurons were much higher in TE than in DP cultures.

D. Organotypic culture of a 300 μm thick coronal section at the level of the TE from one wild type embryo. A tiny crystal of CellTracker CMFDA (green) was placed on the TE.

D' – D''. High magnification of the boxed area in D shows the migratory pathway of lot cells along the diencephalic-telencephalic boundary after 2 D.I.V. The cells that had incorporated CMFDA while in the TE are mGluR1$^+$. 
Figure 5. The TE is a source of glutamatergic cells

A, B. *In situ* hybridizations for *Mash1* and *Gsh2* in coronal sections of an E12.5 wild type embryo. Both transcription factors, implicated in GABAergic neurogenesis, are expressed in the ventricular zone of the caudal ganglionic eminence and prethalamus but not in the TE.

C, D. *In situ* hybridization for *Pax6* and *Neurog2* in coronal sections of an E12.5 wild type embryo. Both transcription factors, implicated in glutamatergic neurogenesis, are expressed in the ventricular zone of the TE. Note that *Pax6* is also expressed in the prethalamus, and that it is expressed with low intensity in the LTE and more intensely in the MTE.

E – H. Immunolabeling for different markers of glutamatergic neurons in coronal sections of E12.5 wild type embryos. Tbr2 (in E) is expressed in the subventricular zone of both the TE (asterisks) and the pallium (arrows). Vglut2 (F), calretinin (G) and Tbr1 (H), three glutamatergic neuron markers, are expressed in the TE mantle, indicating that the TE produces glutamatergic neurons. Note that VGlut2, calretinin and Tbr1 are expressed in the migration path between the TE and the pLOT along the DTB.

Scale bar, 100 μm.

Figure 6. mGluR1 cells are specifically generated in the LTE
A. Schemata of the *in utero* electroporation experiments. Plasmid pCAG-GFP was injected in the lateral ventricle (green) of E11.5 embryos, and tweezers-type electrodes were oriented to electroporate progenitor cells in the ventricular zone of LTE, facing the lateral ventricle. Three days after electroporation, at E14.5, the embryos were removed, fixed and processed for immunohistochemistry.

B1 – B4, C1 – C4. Coronal sections of E14.5 embryos in which the LTE was targeted at E11.5. The sections show that the cells electroporated in the LTE express eGFP (green) and have migrated to the TE mantle, the DTB and the LOT in the piriform cortex at successive caudorostral levels. Note that the contralateral thalamus was also labeled (arrows) due to the direction of the current applied.

B2’, C4’. Close-ups of the boxed areas in B2 and C4, showing eGFP+ cells co-expressing mGluR1 (white arrowheads) while they are migrating along the DTB (B2’) and also when they locate in the LOT (C4’).

D – G. Whole mount *in situ* hybridizations for *Grml* at different embryonic ages, from E11.5 to E13.5. Note the progressive invasion of the rostral levels of the LOT and the pAOB (arrowhead in F) by mGluR1/lot cells as revealed by the distribution of *Grml* transcripts.

Scale bars: B1 – B4, C1 – C4, 100 μm; B2’, C4’, 50 μm; D – G, 200 μm.

**Figure 7. Immunohistochemical characterization of the mGluR1+ cells in the pLOT.**

A – E. Immunohistochemistry of mGluR1 and additional markers of lot cells in coronal sections of E12.5 embryo. The dashed lines highlight the pathway between the TE and the pLOT through the DTB. Note that reelin started its co-expression in mGluR1+ cells
when in the pLOT territory, that Tbr2 was co-expressed in mGluR1+ cells along the DTB and in the LOT, and mGluR1+ cells started to co-express COUP-TFI and Lim1/2 within the TE. AP-2α was not detected in the pLOT territory at E12.5.

A’ – E’. High magnifications of the pLOT territory (boxed areas in A - E) showing in detail the co-expression of different markers with mGluR1. Note the co-expression of mGluR1 and COUP-TFI in C', and of mGluR1 and Lhx5 in D'.

F. Percentages of mGluR1 cells that express additional markers in the pLOT territory.

Scale bars: A – E, 100 μm in A; A’ – E’, 50 μm in A'.

**Figure 8. mGluR1/lot cells follow a rostral migratory path along the LOT.**

A – C. Immunostaining for eGFF, mGluR1 and Tbr2 in ΔNp73eGFP embryos in rostral coronal sections at E13.5, E15.5 and E18.5.

A’ – C’. High magnifications of the white-boxed areas in A – C that contain the LOT. Asterisks indicate the space occupied by the (unstained) mitral cell axons. Most of the mGluR1+ cells at E13.5 were Tbr2+, while only a small proportion of them were ΔNp73+ but Tbr2−. Note the apparent descent in the number of mGluR1+/Tbr2+ cells with age (see graph in D).

A’’ – C’’. High magnifications of the yellow-boxed areas in A – C showing details of the mGluR1+/ΔNp73+ cells in the pallial marginal zone. The arrowheads point to mGluR1/ΔNp73 cells that have reached the boundary between piriform cortex and neocortex.
D. Graph representing the average numbers per section of mGluR1\(^+\)/Tbr2\(^+\) cells in the LOT territory at E13.5, and in the LOT at E15.5 and E18.5. The number of cells descended significantly between E13.5 and E15.5 (p<0.001; Tukey-Kramer multiple comparisons test).

E – G. Sagittal sections of E13.5, E15.5 and E18.5 embryos showing mitral cells in the pAOB immunostained for COUP-TFI, mGluR1, and AP-2\(\alpha\). Note in E the presence of AP-2\(\alpha^+\) cells in the pAOB already at E13.5, and the drastic increase in number of COUP-TFI\(^+\) cells between E13.5 and E15.5, in E – F'.

E’ – G’. High magnification views of the pAOB in sagittal sections to highlight the presence of co-expressions of mGluR1, COUP-TFI and AP-2\(\alpha\). Arrows point to mGluR1\(^+\)/AP-2\(\alpha^+\) cells, and white arrowheads point to COUP-TFI\(^+\)/AP-2\(\alpha^-\) cells.

H. Graph indicating the number of mGluR1\(^+\)/AP-2\(\alpha^+\) cells in the pAOB between E13.5 and E18.5. Note the number of these cells reaches a plateau already at E13.5.

I. Graph indicating the statistically significant increase in the number of COUP-TFI\(^+\)/AP-2\(\alpha^-\) mitral cells in the pAOB from E13.5 and E15.5 (p<0.001; Tukey-Kramer multiple comparisons test).

Scale bars, A – G, E, 100 \(\mu\)m; A' – C' (in C'), A'' – B'' (in A''), and E' – G (in G), 50 \(\mu\)m.

**Figure 9. A small percentage of mGluR1/lot cells express p73.**

A, B. Coronal sections of an E12.5 \(\Delta Np73^{\text{eGFP}}\) embryo. \(\Delta Np73\)-eGFP\(^+\) expressing cells were observed in the TE, the diencephalic-telencephalic boundary and the LOT territory.
A', B'. Close-up images of the boxed areas in A, B. A small number of mGluR1+ cells in
the LOT territory were ΔNp73-eGFP+ (white arrowheads). Hollow arrowheads point to
ΔNp73-expressing, mGluR1-negative cells. See text for details.

C – C'. Close-up image of an area similar to that boxed in A. mGluR1/lot cells (arrows)
were Tbr2+. Note that both the mGluR1+/ΔNp73-eGFP+ cells and the mGluR1−/ΔNp73-
eGFP+ cells were Tbr2-negative.

Scale bars: A, B, 100 μm; A’, B’, C – C”, 50 μm

Figure 10. Temporal evolution of early-generated cells in the LOT: mGluR1/lot cells
and CR cells.

Schemata summarizing the temporospatial changes in the distribution of early-generated
cells associated with the LOT. At E12.5 mGluR1/lot cells positive for AP-2α (blue dots)
localize at rostral levels of the LOT, close to the prospective OB. The mGluR1/lot cells
positive for COUP-TF1 but negative for AP-2α (red dots) concentrate in the TE and
extend rostrally along the LOT territory. Less abundant are the LTE-derived, mGluR1-
positive and ΔNp73-positive CR cells (green dots).

At E18.5, the mGluR1+/AP-2α+ cells and the mGluR1+/COUP-TFI+ cells have advanced
rostrally to be conspicuous in the pAOB, while the mGluR1+/ΔNp73+ CR cells have
extended dorsally within the paleocortex (PCx) and rostrally within the AOB, where they
do not differentiate as mitral cells. Mitral cell axons were depicted in blue.
Figure 1
Figure 2
Figure 3
Figure 4

E11.5 + 4 DIV  Tuj1  ltl1  DAPI

Dorsal pallium

A  A'

B  B'

E12.5 + 2 DIV  CMFDA  ltl1  DAPI

C

% ltl1 cells

TE  pallium

D  D'  D''

Th  CGE

pLOT

TE
Figure 5

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- **Mash1**: pTh, CGE, TE, LTE, MTE
- **Gsh2**: pTh, CGE, TE, LTE, MTE
- **Pax6**: pTh, CGE, TE, LTE, VP
- **Neurog2**: TE, LTE, VP

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- **Tbr2 DAPI**: CGE, TE
- **Vglut2 DAPI**: CGE, TE
- **Calretinin DAPI**: CGE, TE, pLOT
- **Tbr1 DAPI**: TE
Figure 6

**eGFP loph**

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Caudal → Rostral

**Gmi1**

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Figure 7
Figure 8
Figure 9
Lateral thalamic eminence – a novel origin for mGluR1/lot cells

Supplementary Figures

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Legends to Supplemental Figures

Supplemental Figure 1. mGluR1 expression in the forebrain at E11.5

A1 – A3. Grm1 in situ hybridization in an E11.5 wild type embryo; coronal sections at three rostrocaudal levels. Expression is limited to the lateral olfactory tract at rostral levels (A1, B1). In A3, note the intense Grm1 expression in a domain that includes the TE, the posterior end of the LOT territory and the supraoptic–paraventricular region of the hypothalamus.

Scale bar, 100 μm.

Supplemental Figure 2. The expression domains of Grm1 and Lhx5 overlap at E12.5.

A – D. In situ hybridizations for Grm1 and Lhx5 in consecutive coronal sections of the same brain. Note that Grm1 and Lhx5 transcripts occupied the same territories in the TE mantle, DTB, pLOT and aLOT, suggesting co-expression.

Supplemental Figure 3. mAb lot1 recognizes mGluR1 specifically

In this photomicrograph, the TE mantle shows mGluR1 positive cells immunostained with mAb lot1 in an E12.5 Grm1tm1Dgen/J mouse embryo. In these animals, LacZ was inserted in frame with the mGluR1 coding sequence resulting in the expression of a β-Gal fusion protein. Note the co-expression of mGluR1 and β-Gal, indicating the specificity of mAb lot1.

Scale bar, 50 μm.

Supplemental Figure 4. The altered OB (OBSL) in Lhx2+/– mice retain a normal expression of markers of Lhx5 prospective mitral cells

The expression of the AOB marker NP2 in the AOB of Lhx2+/+ or Lhx2+/– mice (a) did not differ from that found in the OBSL of Lhx2–/– mice (b) at E16.5. Likewise, Tbet
expression in the aAOB of $Lhx2^{+/+}$ or $Lhx2^{+/r}$ mice (c) remained unchanged in the OBLS of $Lhx2^{-/-}$ mice (d) at E14.5. A similar situation was encountered for the pAOB marker $Lhx5$ at E14.5 (e, f).

Scale bar, 500 μm.

Supplemental Figure 5. Definition of the medial and lateral subdomains of the TE based on gene expression patterns

A. Schematic displaying the cortical hem (green) as it appears in a hemi-dissected forebrain at E12.5, viewed from the medial side. Note the characteristically curved J-shape of the hem and the distinction we made of a dorsal hem, D-hem, and a caudal hem (hitherto named ventral hem), C-hem. The choroid plexus is colored in light gray, and the projection of the medial TE over the caudal hem is colored in red. The genes selected display expression domains that encompass different territories within the TE and the cortical hem, some of which do not respect the accepted limits between these two structures. Note that their names are color-coded according to the schematics, to remark their main expression territories.

B, C. Medial views of in toto in situ hybridizations for $Wnt3a$ and $Sfrp2$ in hemi-dissected forebrains at E12.5. $Wnt3a$ is expressed in the cortical hem with a rostral$^{\text{high}}$ - caudal$^{\text{low}}$ gradient while $Sfrp2$ labels the MTE intensely.

1 – 3. Schematized coronal sections of E12.5 wild type embryos at three successive rostrocaudal levels. The colors represent the different parts of the thalamic eminence and some adjacent structures. The ventricular zone of the LTE is labeled in blue; the ventricular zone of the MTE in red; the TE mantle in pink and the cortical hem (D-hem and C-hem) in green. Colors also highlight the transitions in gene expression observed between the caudal hem and the LTE, in 2. Other labeled structures are the zona limitans intrathalamica in yellow, and the prethalamus in orange.

D – G. In situ hybridization in coronal sections of E12.5 embryos at the three rostrocaudal levels schematized in 1 – 3. $Wnt3a$ labels the cortical hem including its dorsal and caudal tiers; note that the expression domain of $Wnt3a$ does not invade the LTE. On its part, the expression domain of $Wnt8b$ in the caudal hem extends into the LTE (E2). Note the anatomical continuation between the dorsal hem and the caudal hem in the coronal sections taken at caudal levels (D3, E3). The ventricular zone of the MTE
corresponds to an Sfrp2 expression domain, although Sfrp2 is also expressed in the VP. Expression of Gdf10 occurs in the ventricular zone of the whole thalamic eminence (Shimogori et al., 2010).

Scale bars: B, C, 200 μm in B; D – G, 100 μm in D1.

**Supplemental Figure 6. Definition of the medial and lateral subdomains of the TE based on gene expression patterns**

*In situ* hybridizations in coronal sections of E12.5 embryos at the rostrocaudal level schematized in the diagram.

A – D. Of the four genes selected here, BMP4 highlights the cortical hem and the choroid plexus, Emx2 the LTE and the caudal hem, Dbx1 the MTE, and Fgf15 the ventricular zone of the TE. Note that the Emx2 expression domain in the LTE extends into the caudal hem, and that Dbx1 and Sfrp2 label also the VP.

Scale bar: 100 μm in A.

**Supplemental Figure 7. In utero electroporations in the MTE.**

A. Schemata showing *in utero* electroporations at E11.5 that labeled precursor cells in the MTE. Plasmid pCAG-GFP was injected in the third ventricle (green) and a current was applied that permitted electroporation of progenitor cells in the MTE. Note that cells in the prethalamus and thalamus were labeled to a minor degree. Three days after electroporation, at E14.5, the embryos were removed, fixed and processed for immunohistochemistry.

B1 – B4, C1 – C4. Coronal section series of two E14.5 embryos electroporated in the MTE at E11.5. The cells expressing eGFP remained in the TE mantle while a few of them migrated towards the hypothalamus (yellow arrowheads) but never to the telencephalon. Thalamocortical axons were eGFP labeled as a consequence of the electroporation in the thalamus. Note the TE subventricular zone highlighted by Tbr2 immunostaining, and the reelin immunostaining of the LOT.

Scale bar, 100 μm in B1.
Supplemental Figure 8. mGluR1$^+$ cells in the pAOB exhibit molecular features typical of mitral cells.

A – C. Mitral cell markers in sagittal sections of E15.5 embryos. mGluR1, Tbr2 and reelin were expressed in all mitral cells while COUP-TFI and AP-2α were expressed in the pAOB exclusively.

Scale bar: 100 μm in A.

Supplemental Figure 9. AP-2α cells in the LOT are a subpopulation of the mGluR1/lot cells.

A – C. Immunohistochemistry for COUP-TFI, lot1 and AP-2α in coronal sections of an E12.5 embryo.

A’ – C’. Close-ups of the boxed areas in A – C containing the pLOT and aLOT territories. AP-2α positive cells were not detected in the pLOT at this embryonic age (A’). At mid-rostrocaudal levels (B’), the majority of the mGluR1/lot cells were positive for COUP-TFI and negative for AP-2α. At more rostral levels (C’), most mGluR1/lot cells were AP-2α positive (white arrowheads). Note that some mGluR1$^+$/AP-2α$^+$ cells expressed COUP-TFI (yellow arrowheads).

Calibration bars, A – C, μm in A; A’ – C’, μm in A’.

Supplemental Figure 10. p73 cells in the olfactory bulb

A, B. Mitral cell markers in the olfactory bulb of an E15.5 ΔNp73$^{eGFP}$ embryo.

A’ and B’. Close-ups of the boxed areas in A, B. pAOB mitral cells expressed Tbr2 and mGluR1. ΔNp73 was expressed in some neurons around the OB mitral cell layer, but not in mitral cells. All the ΔNp73 cells were Tbr2 negative and some of them were also mGluR1 positive (white arrowheads).

C. Trp73 in situ hybridization in a sagittal section of an E18.5 embryo showing the distribution of Trp73 transcripts in the OB around the mitral cell layer.

Scale bars: A, B, 50 μm in A; A’, B’ μm; H, 100 μm.
Supplemental Figure 1

E11.5

A1 A2 A3

Rostral → Caudal

LGE LOT LGE LOT TE CGE pLOT MGE
Supplemental Figure 2

E12.5

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- LTE
- pLOT
- aLOT

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Supplemental Figure 3
Supplemental Figure 4
Supplemental Figure 5
Supplemental Figure 6
Supplemental Figure 7
Supplemental Figure 8

Supplemental Figure 9
Supplemental Figure 10