

Plasticity within the niche ensures the maintenance of a Sox2⁺ stem cell population in the mouse incisor

Maria Sanz-Navarro^{1,2}, Kerstin Seidel³, Zhao Sun⁴, Ludivine Bertonnier-Brouty^{1,5}, Brad A. Amendt^{4,6}, Ophir D. Klein^{3,7} and Frederic Michon^{1,8,*}

ABSTRACT

In mice, the incisors grow throughout the animal's life, and this continuous renewal is driven by dental epithelial and mesenchymal stem cells. *Sox2* is a principal marker of the epithelial stem cells that reside in the mouse incisor stem cell niche, called the labial cervical loop, but relatively little is known about the role of the Sox2⁺ stem cell population. In this study, we show that conditional deletion of *Sox2* in the embryonic incisor epithelium leads to growth defects and impairment of ameloblast lineage commitment. Deletion of *Sox2* specifically in Sox2⁺ cells during incisor renewal revealed cellular plasticity that leads to the relatively rapid restoration of a Sox2-expressing cell population. Furthermore, we show that *Lgr5*-expressing cells are a subpopulation of dental Sox2⁺ cells that also arise from Sox2⁺ cells during tooth formation. Finally, we show that the embryonic and adult Sox2⁺ populations are regulated by distinct signalling pathways, which is reflected in their distinct transcriptomic signatures. Together, our findings demonstrate that a Sox2⁺ stem cell population can be regenerated from Sox2⁻ cells, reinforcing its importance for incisor homeostasis.

KEY WORDS: Incisor, Stem cells, *Sox2*, *Lgr5*, Hierarchy, Morphogenesis, Renewal

INTRODUCTION

Renewing organs, such as hair, intestine and certain types of teeth, rely on the ability of stem cells (SCs) to self-renew and differentiate. To ensure tissue homeostasis, the number of SCs in a niche must be kept stable, and conditions such as tissue damage can trigger an SC population increase (Fuchs and Chen, 2012). When the damage is too great or when it affects SCs themselves, the early SC progeny or the niche cells can exhibit plasticity and de-differentiate in order to replenish the SC compartment (Rompolas et al., 2013; Tian et al., 2011). These capacities reflect the potential of the SC niche to control cell fate (Lane et al., 2014).

To compensate for its constant wear, the mouse incisor grows continuously. This life-long growth is fuelled by dental epithelial

SCs located at the proximal end of the incisor, in a structure called the labial cervical loop (laCL). The laCL arises from the dental epithelium around embryonic day (E) 14, and its various cell types are well defined prior to birth (E19) (Fig. 1A). The stellate reticulum (SR) is a pool of epithelial cells located at the core of the laCL. It is surrounded posteriorly and labially by the columnar outer enamel epithelium (OEE), and anteriorly and lingually by the columnar inner enamel epithelium (IEE). The IEE houses the early SC progeny, namely the transient-amplifying (TA) cells and the stratum intermedium (SI) cells (Harada et al., 2006). The TA cells generate pre-ameloblasts, which then differentiate into enamel-secreting ameloblasts (Fig. 1A) (Thesleff and Tummars, 2008). Initial reports suggesting that dental epithelial SCs are present in the SR (Harada et al., 1999) were followed by *in vivo* genetic fate mapping experiments demonstrating that *Gli1* (Seidel et al., 2010), *Sox2* (Juuri et al., 2012), *Bmi1* (Biehs et al., 2013), *Lrig1* and *Igfbp5* (Seidel et al., 2017) mark SCs in the laCL; a number of potential dental SC markers that have not yet been tested through lineage tracing were recently identified using gene co-expression analysis (Seidel et al., 2017). Moreover, the expression of some genes that mark SCs in other organs, such as *Lgr5* (Suomalainen and Thesleff, 2009), *Abcg2*, *Oct3/4* (*Pou5f1*), *Tbx1*, *Pitx2* and *Yap* (*Yap1*) (Cao et al., 2013; Gao et al., 2015; Hu et al., 2017; Li et al., 2011), has also been detected in the incisor SC niche.

SOX2, the focus of this study, is an important transcription factor in the maintenance of pluripotency (Takahashi et al., 2006), formation of endodermal organs (Que et al., 2007; Xie et al., 2014) and development of ectodermal tissues (Arnold et al., 2011; Clavel et al., 2012). We previously reported that SOX2 is a marker for dental epithelial SCs in the mouse incisor and that it is not expressed in the mesenchyme (Juuri et al., 2012). Recently, we showed that deletion of *Sox2* in the dental epithelium at E10.5 (*Pitx2^{Cre/+};Sox2^{fl/fl}*) drastically impairs incisor formation and leads to disappearance of the organ by E18. We also showed that *Sox2* deletion using a ubiquitous promoter during incisor renewal (*Rosa26^{CreER/+};Sox2^{fl/fl}*) slowed down incisor growth (Sun et al., 2016).

Here, we deleted *Sox2* in the epithelium at E11 using *Shh^{GFP-Cre/+}* (Dassule and McMahon, 1998) to analyse the effects on cell differentiation. We found that SOX2 is necessary for ameloblast lineage commitment. Also, we specifically deleted *Sox2* expression from Sox2⁺ cells (*Sox2^{CreER/fl}*) and assessed the consequences of short- and long-term deletion on the laCL. We analysed the effect on laCL shape and on the expression pattern of *Sox2* as well as *Lgr5*, a marker that has been suggested to be expressed by SCs in the laCL (Chang et al., 2013; Suomalainen and Thesleff, 2009). We found that loss of *Sox2* led to a change in laCL morphology and to the disappearance of *Lgr5* expression. Moreover, our data suggest that SR cells are capable of re-establishing a cell population expressing *Sox2* and *Lgr5*. Together, these data indicated the importance of maintaining a Sox2⁺ SC population within the adult laCL. Moreover,

¹Helsinki Institute of Life Sciences, Institute of Biotechnology, University of Helsinki, 00014 Helsinki, Finland. ²Orthodontics, Department of Oral and Maxillofacial Diseases, University of Helsinki, 00290 Helsinki, Finland. ³Department of Orofacial Sciences and Program in Craniofacial Biology, UCSF, San Francisco, CA 94143, USA. ⁴Department of Anatomy and Cell Biology, and the Craniofacial Anomalies Research Center, The University of Iowa, Iowa City, IA 52242, USA. ⁵Département de Biologie, École Normale Supérieure de Lyon, Université de Lyon, 69007 Lyon, France. ⁶College of Dentistry, The University of Iowa, Iowa City, IA 52242, USA. ⁷Department of Pediatrics and Institute for Human Genetics, University of California San Francisco, San Francisco, CA 94143, USA. ⁸Keele Medical School and Institute for Science and Technology in Medicine, Keele University, Keele ST5 5BG, UK.

*Author for correspondence (frederic.michon@helsinki.fi)

 F.M., 0000-0003-4305-305X

we have observed that the transcriptomic signature of the *Sox2*⁺ cells varies between embryonic and adult stages, reflecting their distinct potential. Our data reveal a complex hierarchy in the laCL, and a degree of cellular plasticity not previously identified in the incisor SC niche.

RESULTS

Sox2 expression pattern changes during the transition from embryonic to adult incisor

Our previous use of a reporter mouse strain (*Sox2*^{GFP}), immunohistochemistry, and RNA *in situ* hybridisation pointed to distinct *Sox2*-expressing populations (Juuri et al., 2012), and thus the *Sox2*⁺ cell population in the mouse incisor had not yet been definitively identified. Therefore, we used the highly sensitive RNAscope single mRNA *in situ* hybridisation method (Wang et al., 2012) to investigate the expression pattern of *Sox2* during tooth morphogenesis (Fig. 1). Consistent with previous reports (Juuri et al., 2012; Sun et al., 2016; Zhang et al., 2012), *Sox2* was expressed throughout the dental epithelium at E13.5 (Fig. 1B) and became gradually restricted to the laCL perinatally (Fig. 1C). At postnatal day (P) 60, the *Sox2* transcripts appeared more scattered than at P3, when most cells within the laCL are *Sox2*⁺ (Juuri et al., 2012). The use of this more sensitive method allowed the detection of *Sox2* transcripts in several epithelial lineages of the P60 incisor (Fig. 1D). Although most of the *Sox2*⁺ cells were found in the SR and enamel epithelium (EE) of the laCL, we detected transcripts in the TA cells, pre-ameloblasts, ameloblasts, and the SI (Fig. 1D'). We have previously shown that *Sox2* and its upstream regulator *Fgf8*

are regulated by miRNAs in the laCL (Juuri et al., 2012; Michon et al., 2010), and this miRNA regulation could be the cause of the more restricted SOX2 protein domain.

Deletion of *Sox2* leads to incisor defects during morphogenesis

To decipher the function of SOX2 during incisor morphogenesis, we conditionally deleted the gene in the dental epithelium. We have previously demonstrated that the timing of Cre-driven recombination can dramatically impact the dental phenotype (Cao et al., 2010; Michon et al., 2010; Seidel et al., 2010). As the *Pitx2*-driven *Sox2*^{cKO} led to the absence of incisors at late stages of morphogenesis (Sun et al., 2016), we decided to use *Shh*^{Cre-GFP/+} to delete *Sox2*. *Shh* is expressed later than *Pitx2*, and almost all dental epithelial cells derive from early *Shh*⁺ cells (Juuri et al., 2013b). The *Shh*-driven *Sox2*^{cKO} mice have a hyperplastic dental epithelium in the second and third molars (Juuri et al., 2013a), but no incisor phenotype has been described. As the incisors of the *Shh-Cre; Sox2*^{fl/fl} mice had a different phenotype to that previously reported in *Pitx2-Cre; Sox2*^{fl/fl} mice (Sun et al., 2016), and the incisor was present until the end of embryogenesis, this gave us the opportunity to analyse the dental phenotype at later developmental stages.

We used RNAscope to determine the efficiency of *Sox2* ablation in *Shh-Cre; Sox2*^{fl/fl} mice. By E13.5, essentially no *Sox2* transcripts were detected in the incisor epithelium (Fig. S1A,B). Moreover, the incisor shape was drastically affected in the mutants. At this stage, the control incisor had invaginated into the dental mesenchyme, and the forming laCL contained a large *Sox2*⁺ cell population (Fig. S1A).

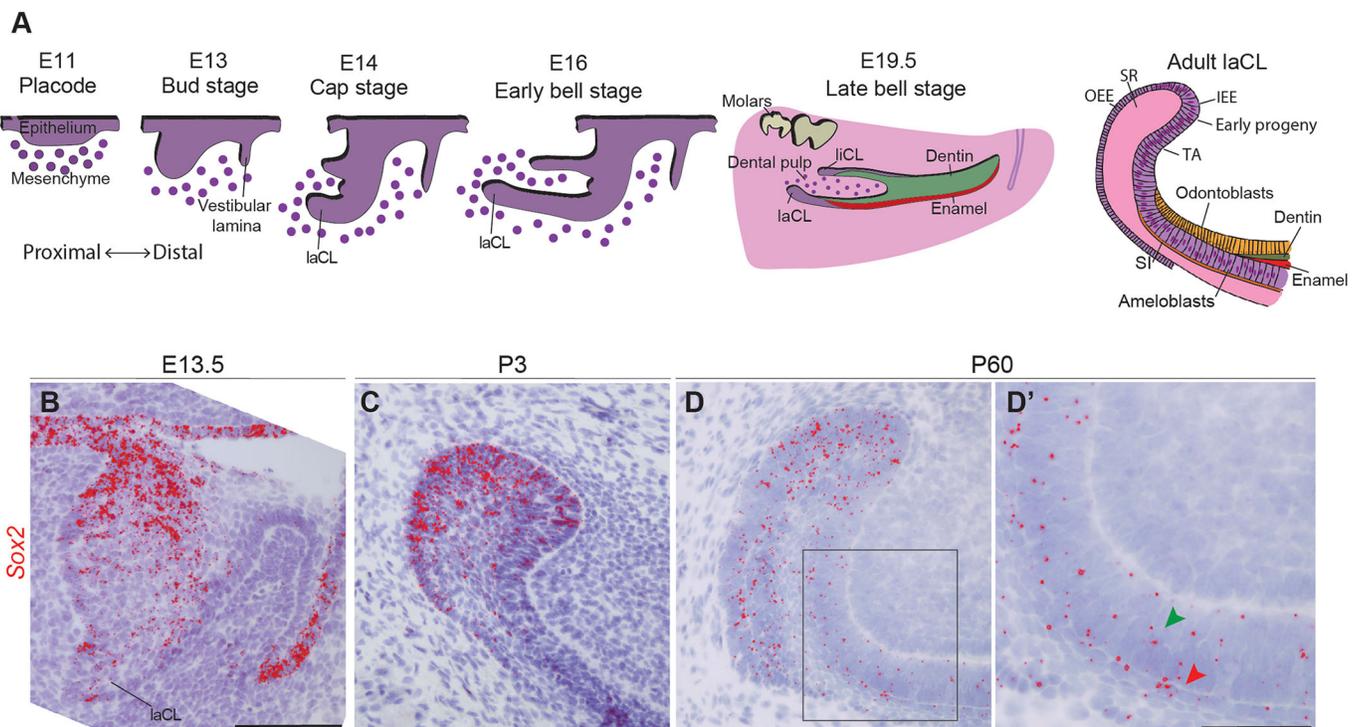


Fig. 1. *Sox2* expression during incisor morphogenesis. (A) Illustration of mouse incisor development, representing the morphological steps from placode stage to adult. At E14, the dental lingual epithelium gives rise to the lingual cervical loop (liCL), while the labial side originates a larger structure: the labial cervical loop (laCL). The adult laCL is composed of the stellate reticulum (SR), the outer enamel epithelium (OEE) and inner enamel epithelium (IEE). The latter gives rise to the stem cell early progeny, namely the transient-amplifying cells (TA), the stratum intermedium (SI) and the ameloblasts. (B) *Sox2* expression (red) is present throughout the entire dental epithelium at E13.5, at highest levels in the lingual side. (C) At P3, *Sox2* expression is more sparse, and restricted to the laCL. (D, D') In adult mice (P60), *Sox2* is expressed in the laCL (SR, IEE, OEE and TA cells), as well as in the pre-ameloblasts, ameloblasts (green arrowhead) and SI (red arrowhead). The boxed region in D is magnified in D'. Scale bars: B-D, 100 µm; D', 50 µm.

The incisor of the *Sox2^{CKO}* littermates displayed a shallow laCL and a wider dental lamina (Fig. S1B). As previously reported (Sun et al., 2016), this phenotype was accompanied by an enlargement of the *Shh*⁺ population (Fig. S1C,D). At E13.5, both the basal (high P-cadherin) and suprabasal (low P-cadherin) (Jussila et al., 2015) cell compartments were present in the mutant incisors (Fig. S1E,F).

To determine the consequences of this early phenotype for subsequent dental morphogenesis, we reconstructed the dental epithelium from micro-computed tomography (micro-CT) scans of *Sox2^{CKO}* embryos and their control littermates from E13.5 to E18.5. The three-dimensional (3D) renderings showed that the morphology of the *Sox2^{CKO}* incisor differed from that of control littermates at all developmental stages (Fig. 2Aa-h), and the shape and length of the incisors varied both among and within individual embryos (Fig. 2Ae-h; data not shown). A recurrent trait in the *Sox2^{CKO}* incisors was the presence of clefts in the labial epithelium (Fig. 2Af-h). These defects were visualised in histological sections as discontinuities in the epithelial tissue (Fig. 2B, red arrowhead). Histological sections also evidenced defective cell differentiation at E18.5, with ameloblast-like cells present in the lingual region (Fig. 2B, green arrowhead).

We observed no significant differences in incisor length at E15.5 (Fig. 2C). However, *Sox2^{CKO}* lower incisors were significantly shorter than those of control littermates at E17.5, and they did not grow further after this stage. The tooth size defect was not attributable to decreased cell proliferation, as we did not detect any significant difference in the density of phospho-histone H3 (pH3)⁺ cells in the dental epithelium at E13.5, nor in the laCL at E18.5 (Fig. 3A). These results are in line with the report on the molars of *Shh^{Cre-GFP/+}; Sox2^{fl/fl}* embryos (Juuri et al., 2013a). Moreover, we did not observe any obvious increase in apoptotic cells by TUNEL assay in the *Sox2^{CKO}* (Fig. S2), indicating that *Sox2* loss does not affect cell death rate, in agreement with other *Sox2* loss-of-function models (Sun et al., 2016). Taken together, our data indicated a role for SOX2 in incisor morphogenesis and dental epithelium cell differentiation.

The laCL and its cell populations are established in *Shh-Cre; Sox2^{fl/fl}* mice

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Next, we investigated the structure of the laCL upon *Sox2* deletion in *Shh-Cre; Sox2^{fl/fl}* embryos. As expected from the epithelial reconstructions (Fig. 2A), laCL volume was drastically reduced in the *Sox2^{CKO}* (Fig. 3B), but the structure was not completely absent. Hence, in addition to *Sox2* we assessed via RT-qPCR the expression of *Sfp5*, which is a marker of early *Sox2*⁺ cell progeny (Juuri et al., 2012), *Shh*, which is expressed in TA cells, pre-ameloblasts and immature ameloblasts (Seidel et al., 2010), and *Lgr5*, which is expressed by a minor cell population in the laCL and marks SCs in several adult organs (Chang et al., 2013; Suomalainen and Thesleff, 2009; Yang et al., 2015) (Fig. 3C). *Sox2* expression was drastically reduced in the mutant incisors. *Sfp5* and *Shh* expression levels were also decreased, indicating defects in cell differentiation. Surprisingly, reduced SHH expression was found only in the lingual side of the incisor (Fig. S3). We did not detect an impact on *Lgr5*

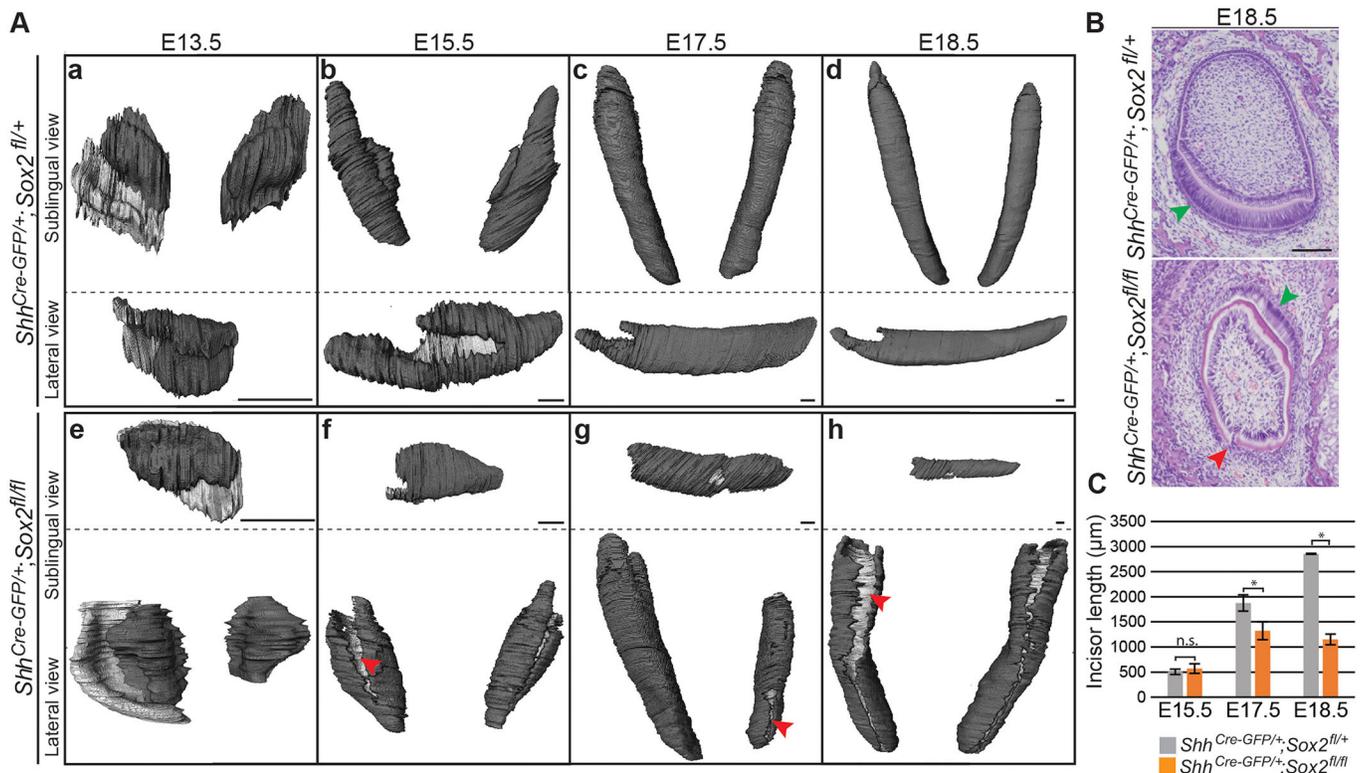


Fig. 2. Mouse lower incisor shape and length are regulated by *Sox2*. (A) 3D reconstructions from micro-CT scans showing the dental epithelium in dark grey. The internal layer of the dental epithelium (b-d,f-h) or vestibular lamina (a,e) appears in light grey. *Sox2^{CKO}* incisors exhibit an aberrant morphology at all embryonic stages. At E13.5, the tooth domain is broader, and at E15.5 clefts (red arrowhead) appear. The defects can vary within the same individual (g). (B) Histological staining of frontal sections shows that the well-organised ameloblast layer seen in control incisor (green arrowhead) is not visible in *Sox2^{CKO}* individuals. A cleft is visible on the labial side of the mutant incisor (red arrowhead). (C) The length of the epithelial compartment is similar in the control and *Sox2^{CKO}* at E15.5. The incisor length increases over 2 mm from E15.5 to E18.5 in controls, but only 0.6 mm in mutants. E17.5, **P*=0.016; E18.5, **P*=1×10⁻⁵; n.s., not significant. *n*=3. Scale bars: A, 50 µm; B, 100 µm.

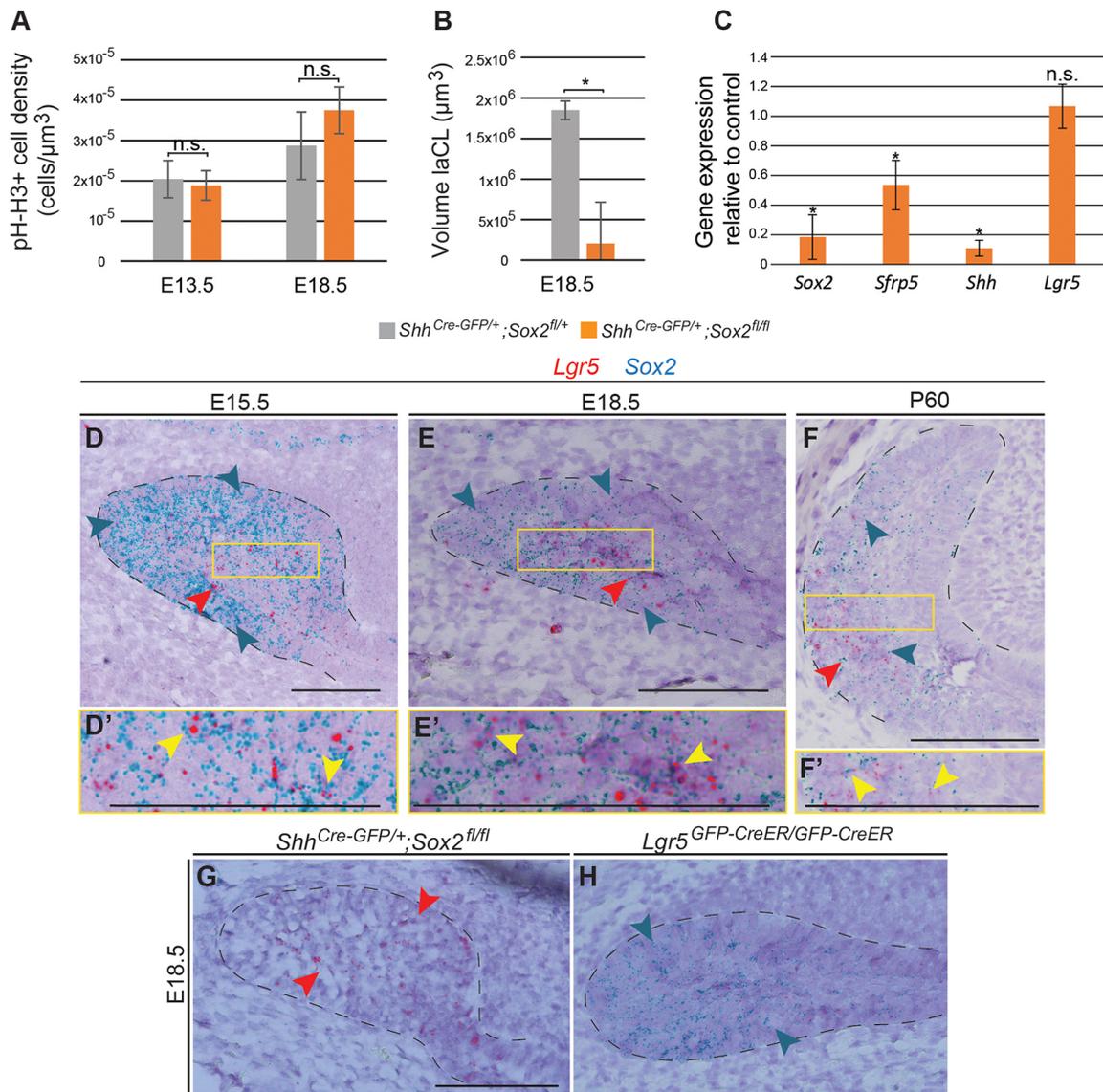


Fig. 3. *Sox2*^{KO} impacts laCL volume and the expression of different differentiation markers. (A) Quantification of phospho-histone H3 (pH-H3)⁺ cell density in the dental epithelium reveals no significant defect in cell proliferation in *Sox2*^{KO}. (B) The volume of the E18.5 laCL is drastically decreased in *Sox2*^{KO}. **P*=0.026. (C) qPCR analysis demonstrates that *Sox2*^{KO} induces a decrease in *Sfrp5* and *Shh* expression. *Lgr5* expression remains unaffected. **P*<0.05. (A-C) *n*=3. (D) *Sox2* transcripts (blue arrowheads in all images) at E15.5 are detected in all cells of the laCL (outlined). *Lgr5* transcripts (red arrowheads in all images) mark a subset of the *Sox2*⁺ population in the SR. (E) At E18.5 the *Sox2* expression domain is smaller than at previous stages. *Lgr5* expression is localised to the SR. The laCL houses the expression of both *Sox2* (EE and SR) and *Lgr5* (SR). (F) In the adult incisor, *Sox2* transcripts are localised in different areas of the laCL, while *Lgr5* expression is confined to the most proximal part. (D', E', F') Magnifications of the boxed regions in D, E, F. Yellow arrowheads indicate cells expressing both *Lgr5* and *Sox2* transcripts. (G, H) *Sox2*^{KO} shows no changes in *Lgr5* expression, as *Lgr5*^{KO} displays a normal *Sox2* expression pattern. Scale bars: 100 μm . qPCR.

expression, which has been reported at E14.5 in the molar bud (Kawasaki et al., 2014) but in the laCL incisor only after E16.5 (Suomalainen and Thesleff, 2009). However, the sensitivity of the RNAscope assay enabled the detection of *Lgr5* transcripts already at E15.5 in the incisor laCL (Fig. 3D-F) and, interestingly, the expression patterns of *Sox2* and *Lgr5* overlap (Fig. 3D-F', Fig. S4).

To further study the connection between *Sox2* and *Lgr5* expression, we analysed their expression patterns upon deletion of either gene (Fig. 3E, G, H). *Lgr5* null mice (*Lgr5*^{KO}) die neonatally due to gastrointestinal problems and ankyloglossia (Morita et al., 2004). These mice also exhibit cleft palate, but we did not identify morphological abnormalities in the laCL. In the control embryos, *Sox2* transcripts were found in the EE and SR, and *Lgr5* transcripts

were mainly restricted to the SR. Interestingly, in the *Sox2*^{KO} laCL, *Lgr5* expression was maintained, and the *Sox2* expression pattern did not change in the *Lgr5*^{KO} incisor (Fig. 3G, H). These observations highlighted the importance of *Sox2* for ameloblast lineage commitment, but not for establishment of the laCL.

***Sox2* expression is quick to recover after transient *Sox2* deletion in the adult incisor**

Having identified the importance of *Sox2* for dental epithelial cell differentiation during embryogenesis, we next analysed the effects of *Sox2* absence in the renewing incisor. As *Sox2*^{KO} mice die perinatally, we used *K14-CreER* mice (Huelsen et al., 2001; Järvinen et al., 2006; Vasioukhin et al., 1999) to delete *Sox2* in the

laCL. We investigated the morphology and *Sox2* expression pattern at 2 days, 11 days and 1 month after Cre activation. No obvious differences were observed in histological sections (Fig. S5A) nor in the SOX2 pattern (by immunofluorescence staining; data not shown). We also analysed the expression levels of *Sox2*, *Sfrp5*, *Lrg5* and *Bmi1* 24 h after induction via qPCR and found no statistically significant differences from controls (data not shown). This suggested a lack of recombination in the laCL, and our examination of *Sox2* and keratin 14 (*K14*) expression in the adult laCL at the protein and transcript levels (Fig. S5B,C) showed minimal overlap. *K14* was expressed in the SR region, but not in the most lingual part of the SR, nor in the IEE, where many of the *Sox2*⁺ cells reside; this is in line with the low recombination levels reported in the OEE with this Cre line (Hu et al., 2017).

Therefore, we generated *Sox2*^{CreER/fl} mice to specifically delete *Sox2* in the *Sox2*⁺ cells upon Tamoxifen administration. Two-

month-old *Sox2*^{CreER/fl} mice were administered Tamoxifen for three consecutive days. We then examined the laCL at 3 days, 1 week and 4 weeks chase (Fig. 4A), and *Sox2*^{fl/+} littermates were used as controls. We also injected adult *Sox2*^{CreER/fl} mice with corn oil and observed no aberrant phenotype (Fig. S6). After 3 days of chase, we observed a large decrease in the number of *Sox2* transcripts (Fig. 4B,C). Moreover, the spherical shape of the laCL was lost in the mutants, which instead exhibited an elongated SC niche (Fig. 4D,E). However, the transient loss of *Sox2* expression did not lead to cell death (Fig. S7), and after 5 days without Cre activation some *Sox2* transcripts were detected, although the expression was fainter than in the control laCL (Fig. 4F,G). Interestingly, by this time point, laCL shape was restored (Fig. 4H,I). At 1 month chase, we did not detect any differences in laCL morphology or *Sox2* expression pattern between control and mutant (Fig. 4J,K).

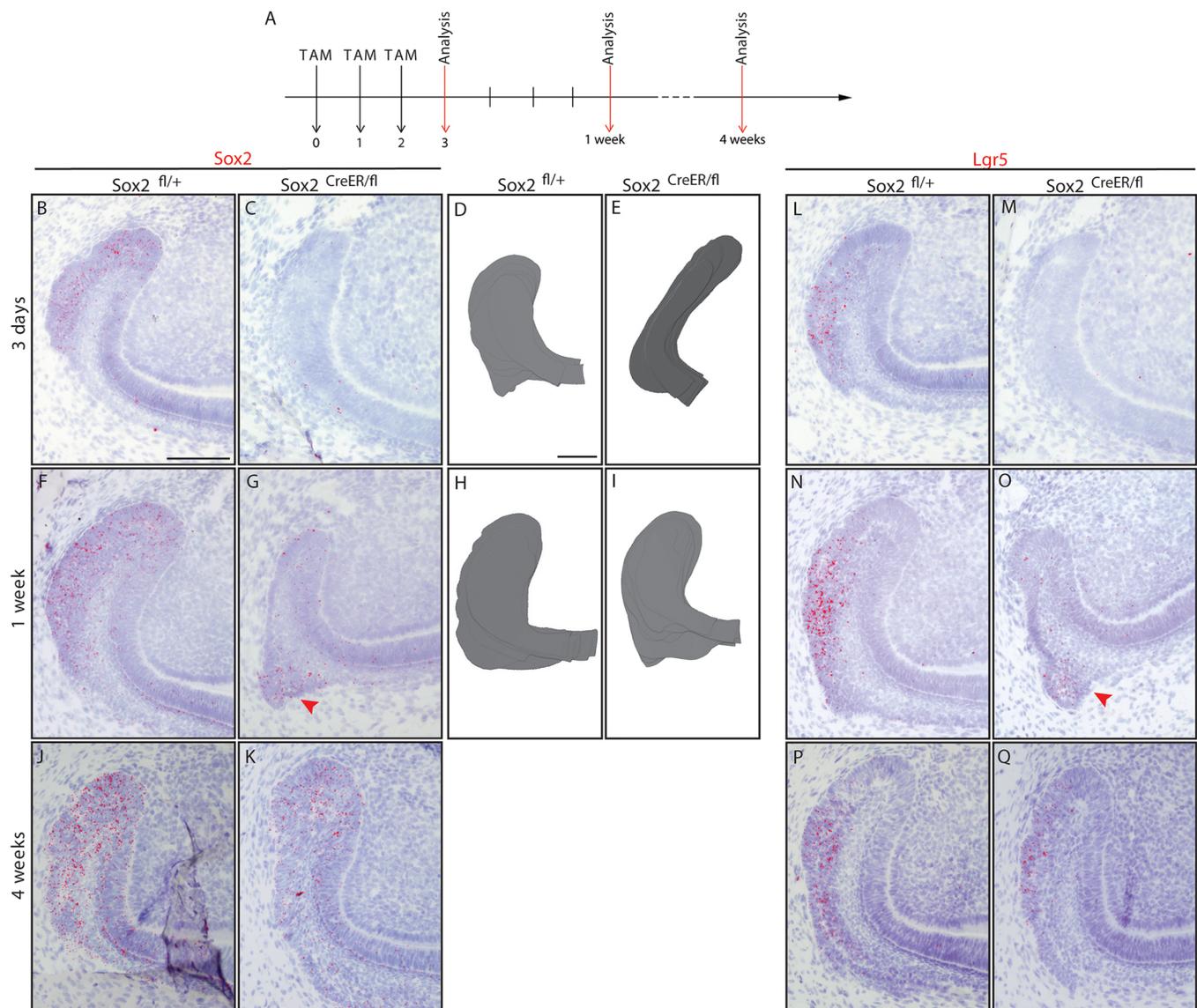


Fig. 4. *Sox2* and *Lgr5* expression are lost then restored in *Sox2*^{CreER/fl} laCL. (A) The experimental setup. (B,C) *Sox2* and (L,M) *Lgr5* expression are almost abolished after three Tamoxifen injections in *Sox2*^{CreER/fl} mice. (D,E) The *Sox2*^{CreER/fl} laCL is narrower than that of controls. One week after the first Tamoxifen administration, a faint *Sox2* signal is detected in the laCL (F,G, arrowhead), while the *Lgr5* expression pattern appears to be normal in the mutant (N,O, arrowhead). (H,I) At this stage, the morphology of the laCL appears to be normal. (J,K,P,Q) One month after Cre recombinase activation, the mouse incisor SC niche of *Sox2*^{CreER/fl} mice is indistinguishable from that of control littermates. Scale bars: 100 μ m.

Lgr5 marks a small cell population in the laCL (Suomalainen and Thesleff, 2009; Yang et al., 2015), and its expression pattern overlaps with that of *Sox2* (Fig. 3D-F, Fig. S3). Therefore, we investigated the effect of *Sox2* loss on *Lgr5* expression. After 3 days of Tamoxifen administration, *Lgr5* transcripts were greatly decreased (Fig. 4L,M), similarly to *Sox2*. Four days later, *Lgr5* expression levels were similar to those of the control (Fig. 4N,O) and returned to normal after 1 month of chase (Fig. 4P,Q).

To evaluate the effect of long-term *Sox2* loss, we administered Tamoxifen to *Sox2^{CreER/fl}* mice seven times over 11 days. Incisors were collected 1 day after the last injection (Fig. 5A). As expected, very few *Sox2* transcripts were detected in the laCL, whereas controls exhibited expression of *Sox2* as previously reported (Fig. 5B-C'). *Lgr5* expression was faint as well (Fig. 5D-E'). Very few *Sox2* transcripts were detected in the area where *Lgr5* expression was localised. Moreover, the morphology of the laCL was not affected, similar to results obtained with another *Sox2^{cKO}*

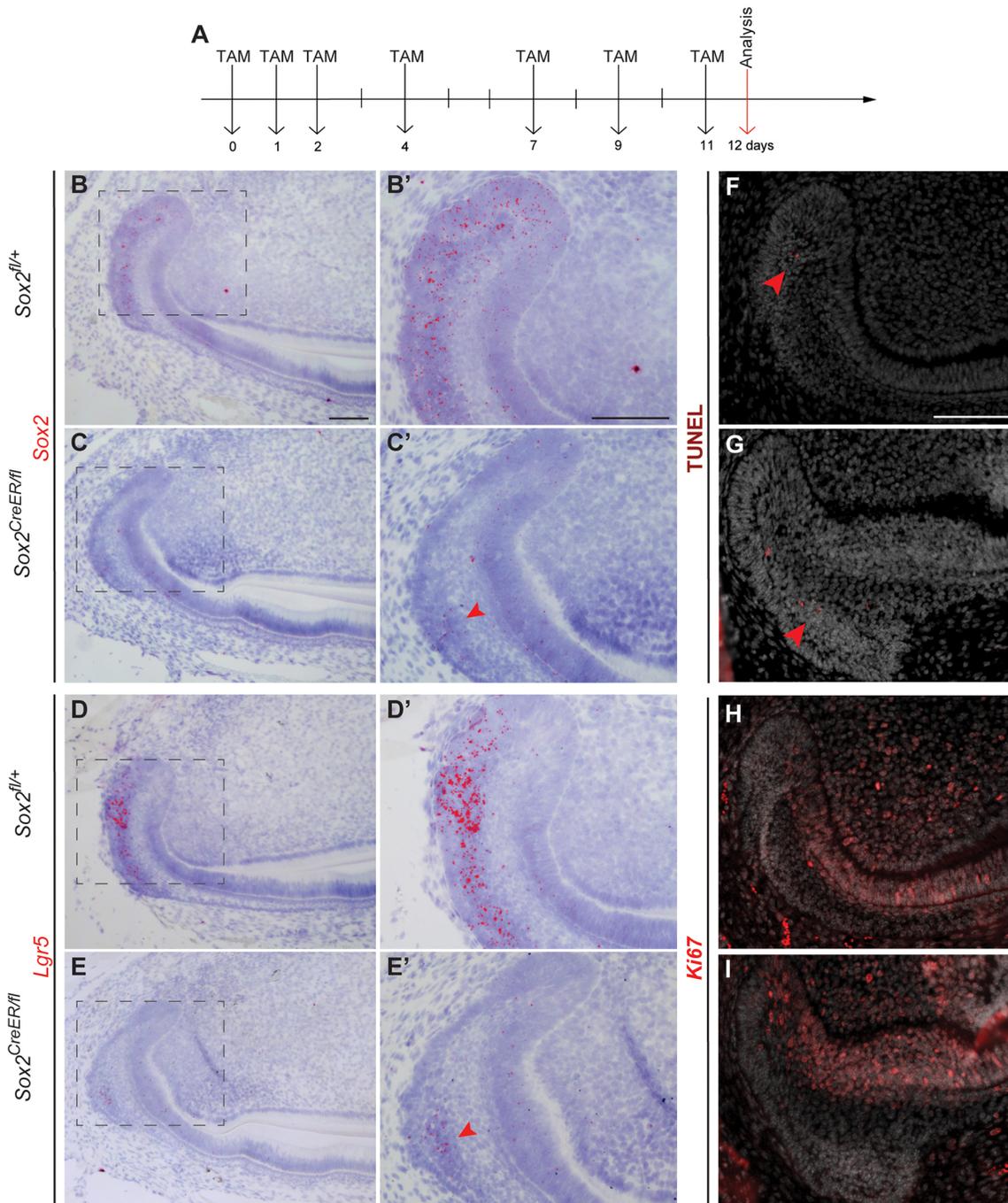


Fig. 5. A day after an 11-day *Sox2* ablation, the *Sox2* expression pattern, proliferation and cell differentiation appear disturbed. (A) The experimental setup. (B-E') A small amount of *Sox2* transcripts is detected in the proximal area of the SR (arrowhead), where faint expression of *Lgr5* is seen (arrowhead). (F,G) TUNEL assay confirms that there is no increase in apoptosis in the laCL, where only a few positive cells were found (arrowheads). (H,I) The domain of proliferating cells, visualised with Ki67 staining, appears similar to the control. Scale bars: 100 μ m.

model *Rosa26^{CreER/+}; Sox2^{fl/fl}* (Sun et al., 2016). We further examined the cellular response to the deletion of *Sox2* in the *Sox2⁺* SCs and did not observe any obvious increase in cell death or aberrant cell proliferation (Fig. 5F-I).

The *Sox2⁺* population is regenerated from the SR

The restoration of a *Sox2⁺* cell population days after deleting *Sox2* suggested a high degree of plasticity within the laCL. To understand the origin of the newly generated *Sox2⁺* population, we studied the dividing cells during laCL restoration using EdU incorporation. Under normal renewal conditions, dividing cells are localised to the lingual and distal part of the laCL, the IEE, the TA cells, and the distalmost SR cells (Hu et al., 2017) (Fig. 6A). In comparing this pattern with *Sox2* expression, it is apparent that most of the proliferating cells in the laCL do not strongly express *Sox2* (Fig. 6B). In addition, the cells in the proximal region of the laCL are quiescent (Seidel et al., 2010). We marked the proliferative cells 3 days after the first Tamoxifen administration and analysed their position and quantity 4 days later (Fig. 6C). In the control, the EdU⁺ cells were located in the central and proximal sections of the SR, where *Sox2* is faintly expressed (Fig. 6D-E'). By contrast, in the *Sox2^{CreER/fl}* mice, these cells were displaced to the proximalmost part, close to the *Lgr5⁺* cell area (Fig. 6F-G'). Furthermore, the percentage of EdU⁺ cells was significantly increased in the mutant SR compared with the control (Fig. 6H). This reflected an increase in proliferation in the SR at the beginning of the rescue period.

Modulation of the *Sox2⁺* cell signature in embryonic versus adult incisors

The observation that almost all laCL cells are *Sox2⁺* at E15.5 (Fig. S3), when *Lgr5* expression appears, suggested that embryonic *Sox2⁺* cells might give rise to the *Lgr5⁺* subpopulation during incisor formation. However, during laCL regeneration after *Sox2* ablation, the *Sox2⁺* cells seemed to partially arise from the subpopulation expressing both *Sox2* and *Lgr5* (Fig. 4F,G,N,O). This observation raised the question: how similar are the *Sox2⁺* populations in the forming and renewing incisor? We therefore compared the transcriptome of the *Sox2⁺* cells in the early incisor (E14.5) with that of the *Sox2⁺* cells in the renewing incisor (P30) using gene expression microarrays. We first extracted the transcripts that were similarly expressed in embryonic and adult *Sox2⁺* cells ($-2 < \text{fold change} < 2$) and compared them with the transcriptome of mouse embryonic stem cells (mESCs) as a naïve cell reference (Fig. 7A). Then, we compared the transcriptomes of the embryonic and adult *Sox2⁺* cells (Fig. 7B). We selected the transcripts that were statistically significant ($P < 0.05$, ANOVA) and exhibited a consequential fold change (fold change > 2) (Table S1).

For the first analysis (*Sox2⁺* versus mESCs), we observed that 927 transcripts (2.34% of the signature) were enriched in *Sox2⁺* cells, independent of their stage, and 1583 transcripts (4% of the signature) were downregulated. This observation reflected that ~6.34% of the signature was differentially regulated in *Sox2⁺* cells compared with mESCs (Fig. 7A, Table S2). We compared the gene ontology processes (GOPs) between these samples and found that 187 were activated in *Sox2⁺* cells, including those specific to ectodermal organ formation and regulation, and odontogenesis (Table S3). Also, both canonical and non-canonical Wnt signalling were activated. Among the enriched genes, we examined *Vangl2* as a test case because it is a member of the planar cell polarity signalling pathway (non-canonical Wnt). Moreover, its expression has previously been reported in the ameloblasts and odontoblasts of embryonic molars, where it regulates cell alignment (Obara et al., 2017; Wu et al., 2016). We found *Vangl2*

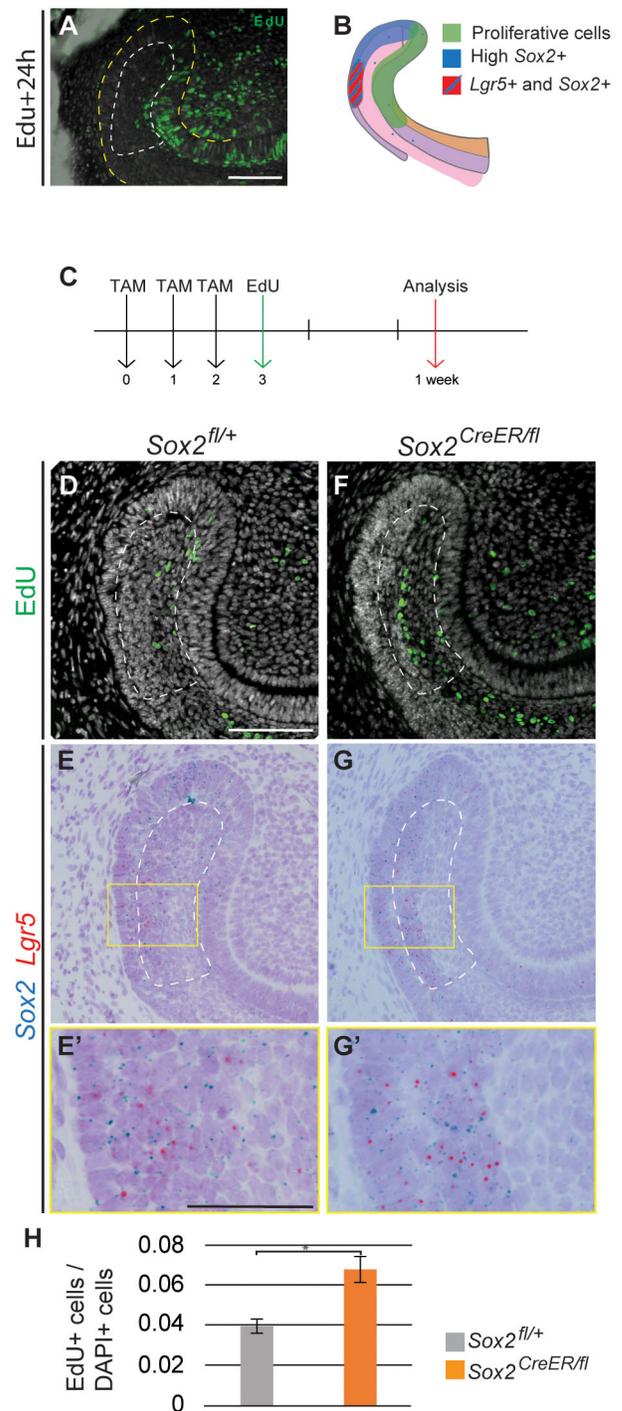


Fig. 6. Expression of *Sox2* and *Lgr5* is rescued from the SR. (A) Pattern of proliferative cells (EdU⁺) 24 h after being administered to the mouse. (B) Schematic representation of the proliferative region and the *Sox2⁺* and *Lgr5⁺* domains. (C) The experimental setup. EdU⁺ cells (D,F) and *Sox2* and *Lgr5* mRNA detection (E,G) performed in identical sections. (E',G') Higher magnifications of the boxed regions in E,G. (H) Quantification of EdU⁺ cells in the SR of control and *Sox2^{CreER/fl}* mice. The area quantified is delineated by the dashed line in D-G. Results are expressed as the fraction of EdU⁺ cells among total cells (number of nuclei, DAPI) compared with the control. * $P=0.007$. Scale bars: D-G, 100 μm ; E',G', 50 μm .

transcripts in the incisor at E14.5 and in the adult laCL. However, in the adult stage, most of the transcripts were found in TA cells and ameloblasts, where there are fewer *Sox2⁺* cells (Fig. 7C-D').

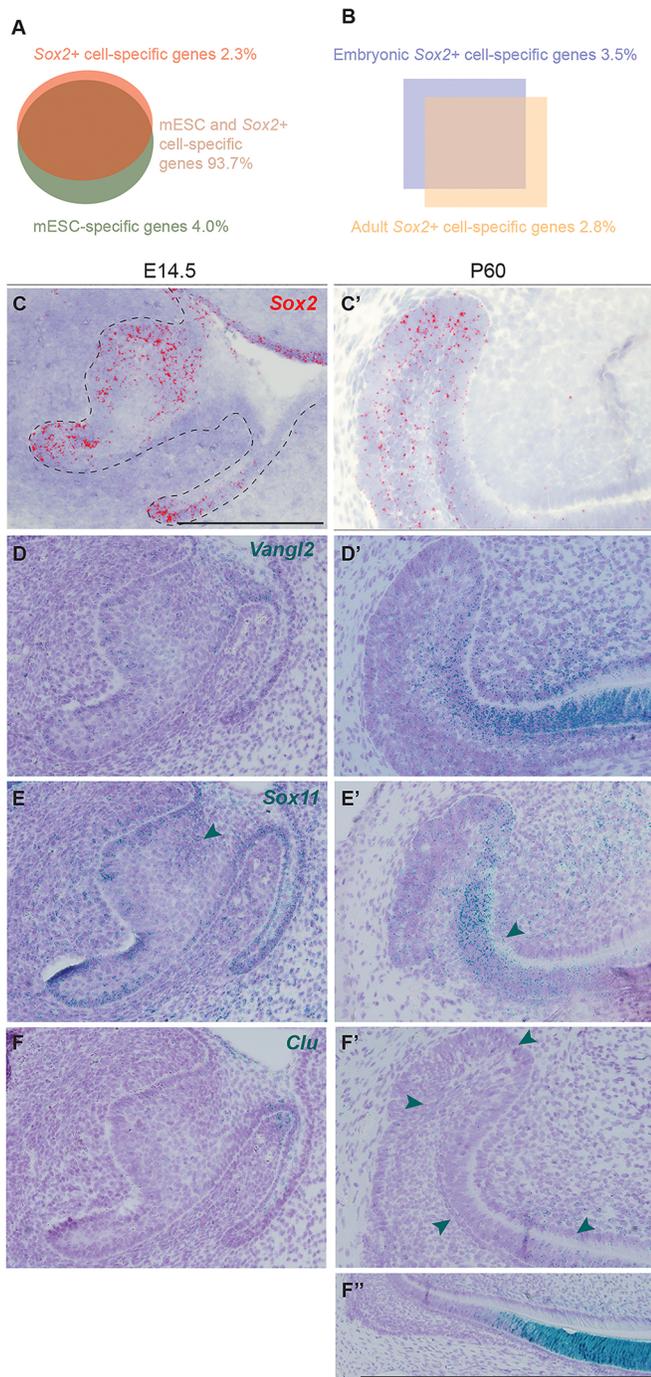


Fig. 7. Transcriptomic changes between *Sox2*⁺ embryonic progenitors and *Sox2*⁺ dental SC. (A) Signatures of *Sox2*⁺ cells and mESCs overlap by 93.7%. (B) Comparison of embryonic and adult *Sox2*⁺ cells: 3.5% of genes are specific to embryonic *Sox2*⁺ cells, and 2.8% specific to the renewing incisor *Sox2*⁺ SCs. (C,C') *Sox2* expression pattern at E14.5 and adult stage. (D,D') *Vangl2* is enriched in *Sox2*⁺ cells compared with mESCs. It is expressed in the embryonic incisor and in the adult tooth. (E,E') *Sox11* is highly expressed in the embryonic incisor (arrowhead) and in the surrounding mesenchyme. In the adult, expression is mostly localised to the TA cells (arrowhead). (F,F') *Clu* expression is seen in the adult incisor (arrowheads), with the majority of transcripts found in the differentiated epithelial cells (F''). Scale bars: 100 μ m.

When comparing embryonic and adult *Sox2*⁺ cells, we observed that 3.54% of the signature (1400 hits) was enriched in embryonic cells (Fig. 7B, Table S1). Among these, 143 GOPs were enriched

over 2.5 fold, including cell division regulation processes (e.g. mitotic DNA replication, DNA replication initiation) (Table S4). We also found a number of genes important for the mineralisation of forming teeth, such as *embigin* (Xie et al., 2015), *Six4* (Nonomura et al., 2010) and *Cxcr4* (Juuri et al., 2013b), the last of which is thought to be important for the migration of epithelial progenitors in adult laCL (Yokohama-Tamaki et al., 2015). Also, *Sox11*, which is involved in palate development (Sock et al., 2004; Watanabe et al., 2016) and expressed in the mouse embryonic molar (Dy et al., 2008; Hargrave et al., 1997), was enriched by 4.40 fold. *Sox11* transcripts were abundant in the mouse incisor epithelium at E14.5, with fewer transcripts in the adult laCL. In adults, we found *Sox11* expression also in pre-ameloblasts and in the mesenchymal compartment (Fig. 7E,E').

Similarly, 2.75% (1089 transcripts) of the signature was enriched in the adult *Sox2*⁺ population (Fig. 7B); 153 GOPs were enriched in adult *Sox2*⁺ cells, with those specific for the immune response well represented (e.g. antigen processing, macrophage activation, Toll-like receptor signalling) (Table S5). The adult *Sox2*⁺ population signature contained mineralisation markers (e.g. *Dspp*, enamel, amelogenin, ameloblastin) and metalloproteinases (e.g. *Mmp13*, *Mmp14*, *Mmp20*) (Table S1). We also detected expression of *Barx2*, a gene involved in cell migration and differentiation (Juuri et al., 2013b). Moreover, clusterin (*Clu*), a stress-activated and apoptosis-associated chaperone, which has been found in embryonic and postnatal mouse molars (Chou et al., 2009; Khan et al., 2013; Shiota et al., 2012), was enriched in adult *Sox2*⁺ cells by a fold change of 187.08. In the embryonic (E14.5) incisor we found almost negligible amounts of *Clu* expression, and few transcripts were found in the vestibular lamina at this stage. In the adult, low *Clu* expression levels were found in the laCL; higher expression levels were evident in the pre-ameloblasts and ameloblasts (Fig. 7F-F''), where *Sox2* transcripts were also found (Fig. 7C,C').

DISCUSSION

We have previously demonstrated the role of *Sox2*⁺ SCs in incisor renewal (Juuri et al., 2012) and in successional tooth formation in the mouse (Juuri et al., 2013a). More recently, we showed that early deletion of *Sox2* in the dental epithelium led to the absence of the incisor at E18 (Sun et al., 2016). As this prevented analysis during late time points of embryonic development, here we used the *Shh*^{GFP^{Cre}/+} allele to delete *Sox2* at the dental placode stage. *Sox2* deletion using this driver resulted in growth and shape irregularities, including a curved incisor phenotype, which, together with the drastic decrease of *Shh* and *Sfrp5* expression, pointed to improper differentiation of cell lineages. Interestingly, SHH expression (Fig. S6) and ameloblast-like cells (Fig. 2B) were found on the lingual side of the incisor. SHH expression adjacent to the lingual cervical loop (liCL) has previously been linked to an expanded liCL and to ectopic lingual ameloblasts (Klein et al., 2008). However, the laCL structure was preserved, but its morphology and the cell arrangement were affected. These observations suggest an essential role for *Sox2* in lineage commitment and early differentiation towards the enamel-secreting ameloblast fate. We have previously reported that the number of proliferative cells is reduced after early *Sox2* deletion (Sun et al., 2016). However, we did not detect a significant reduction in proliferation in the *Sox2*^{ckO} model used here; instead, we propose that slower renewal is caused by defects in cell differentiation. We conclude that *Sox2* plays a key role in the maintenance of enamel organ morphology and proper cell differentiation during incisor morphogenesis.

Lgr5 marks intestinal and skin SCs (Barker et al., 2007; Haegebarth and Clevers, 2009; Jaks et al., 2008), and it also

marks a small epithelial cell population in the mouse incisor laCL (Chang et al., 2013; Suomalainen and Thesleff, 2009; Yang et al., 2015). Our identification of *Lgr5*⁺ cells within the *Sox2*^{CKO} laCL indicates that other potential laCL SC populations can be present in the absence of *Sox2*. Therefore, we propose that *Sox2* expression is of importance for initiating differentiation towards ameloblast fate, but not for establishing the SC niche.

In the adult incisor, *Sox2* deletion in the *Sox2*⁺ cells (*Sox2*^{CreER/Fl}) caused a drastic, but temporary change in laCL shape, and the loss of *Lgr5* expression. We showed that *Lgr5* and *Sox2* expression overlap during embryonic and postnatal stages. Therefore, we conclude that *Lgr5*⁺ cells represent a subpopulation of *Sox2*⁺ cells in the developing and renewing laCL. This situation differs from other SC niches, such as the stomach, where *Lgr5* and *Sox2* mark distinct cell populations (Arnold et al., 2011).

To ensure tissue homeostasis, the number of SCs in a niche is kept stable, but tissue damage can trigger an SC increase (Fuchs and Chen, 2012). If the damage is too great, the niche cells can display signs of transient plasticity in order to replenish the SC compartment, as shown in skin (Rompolas et al., 2013) and intestine (Tian et al., 2011). Such intra-organ plasticity (Blanpain and Fuchs, 2014) requires cell de-differentiation or transdifferentiation to ensure the maintenance of organ integrity. While this mechanism is well studied in other SC niches, it has not yet been documented in the dental context. The loss of *Sox2* expression in *Sox2*^{CreER/Fl} mice led to a morphologically thinner laCL depleted of *Sox2*⁺ and *Lgr5*⁺ cells. This phenotype was rapidly rescued, and the *Lgr5*⁺ subpopulation was the first to emerge, from the distal section of the laCL. Moreover, our EdU incorporation experiment demonstrated that some SR cells were plastic enough to regenerate the lost cell populations within the laCL.

Taken together, our data suggest that after damage the SR cells regenerate first a *Sox2*⁺ *Lgr5*⁺ double-positive cell population, and then a *Sox2*⁺ *Lgr5*⁻ population (Fig. 8A-D). Also, these results suggest that *Sox2* marks a heterogeneous population, where different lineage specificities exist. However, long-term ablation of *Sox2* did not lead to laCL shape malformation, and the laCL maintained a very small *Sox2*⁺ *Lgr5*⁺ cell population (Fig. 8E).

We have previously reported that the global deletion of *Sox2* (*Rosa26*^{CreER/+}; *Sox2*^{fl/fl}) in adult mice leads to a reduction in incisor renewal rate (Sun et al., 2016). In the *Sox2*^{CreER/Fl} mice, the proliferation pattern in the laCL is maintained after prolonged *Sox2* ablation. Therefore, we hypothesize that the incisor growth

defect reported earlier (Sun et al., 2016) is caused by defective cell differentiation, similar to the embryonic scenario.

Finally, an important question that remains largely unanswered in the stem cell field is that of the origin of adult SCs. We found that the transcriptomic signatures of embryonic and adult *Sox2*⁺ cells are very similar. Moreover, they express a number of genes differently compared with naïve mESCs. From this observation, we propose that dental SC identity is represented either in the embryonic and adult *Sox2*⁺ cell overlapping transcriptomes or in the genes enriched in adult *Sox2*⁺ cells. Although our results alone are not enough to distinguish between these possibilities, an early generation of the dental SC signature would require the formation of the niche microenvironment early on. Interestingly, *Notch1*, a marker of the dental epithelial SC niche, is expressed in the mouse incisor at E14.5 (Felszeghy et al., 2010; Mucchielli and Mitsiadis, 2000). Therefore, we postulate that SC niche and *Sox2*⁺ cell dental fate are already established by E14.5 in the mouse incisor. The corollary of such a conclusion would be that the genes enriched in embryonic or adult *Sox2*⁺ cells should be related to the role of cells within the organ at this stage. For instance, *Sox11*, a transcription factor involved in epithelial-mesenchymal interactions (Hargrave et al., 1997), was enriched in the forming incisor. On the other hand, we found an enrichment of *Clu* in adult *Sox2*⁺ cells. This chaperone has been reported to play a role in secretory odontogenesis, an important function in the adult incisor (Khan et al., 2013), and was highly enriched in the incisor ameloblasts. These observations strengthen our hypothesis that the differences between embryonic and adult *Sox2*⁺ transcriptomes are inherent to the temporal role of the cell population (morphogenesis versus ameloblast lineage renewal). From these data, we conclude that the differences in gene expression that we observed reflect cues from the microenvironment and minor changes in the role of the *Sox2*⁺ cells.

Collectively, our data demonstrate the importance of a *Sox2*⁺ cell population for incisor renewal and cell differentiation. We also observed an impressive cellular plasticity in the laCL that acts to maintain the *Sox2*⁺ population. We propose that *Lgr5*⁺ cells are a subpopulation of the *Sox2*⁺ cells, and are the first cells to reappear in the event of transient *Sox2* loss. This indicates the existence of a complex relationship between the *Lgr5*-expressing and *Sox2*-expressing cells.

MATERIALS AND METHODS

Mouse lines

The stage of the embryos was determined according to morphological criteria and noon of plug day was counted as E0.5. All animals are available from The

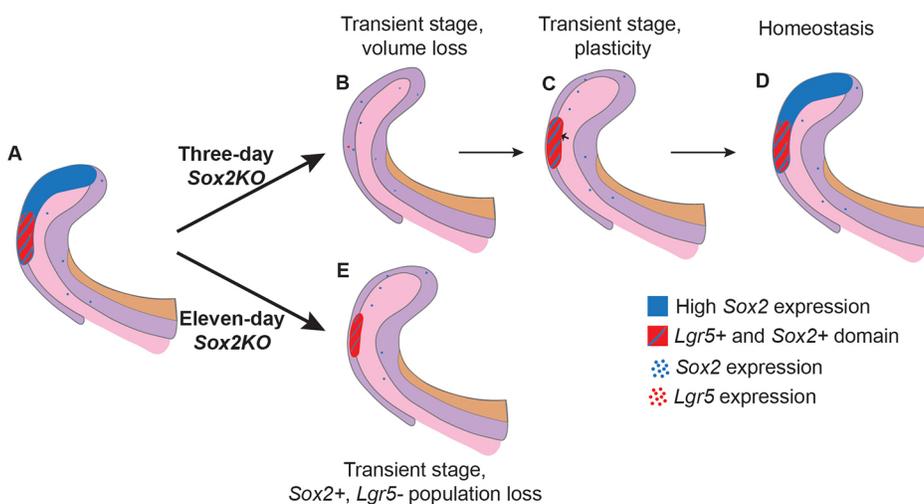


Fig. 8. Model for the effects of short- and long-term *Sox2* ablation. (A) Summary of *Sox2* and *Lgr5* expression domains in normal conditions within the laCL. (B) Upon conditional deletion of *Sox2* in *Sox2*-expressing cells for 3 days, the laCL becomes narrower, and almost all *Sox2* and *Lgr5* transcripts are lost. (C) Shortly after, the volume of the laCL is back to normal due to an increase in cell proliferation in the SR (small arrow). Overlapping expression of *Sox2* and *Lgr5* is found in the distal side of the laCL. (D) Eventually, the laCL reaches homeostasis and returns to its original state. (E) In the event of *Sox2* ablation for 11 days, the laCL maintains a small *Lgr5*⁺ *Sox2*⁺ cell population.

Jackson Laboratory. *Shh^{GFP-Cre/+}; Sox2^{fl/fl}* were used as *Sox2^{ckO}* (Juuri et al., 2013a). *K14-CreER* males [Tg(KRT14-cre/ERT)20Efu/J, stock 005107] were crossed with *Sox2^{tm1.1Lan/J}* females (*Sox2^{tm1.1Lan/J}*, stock 013093) to generate *K14-CreER; Sox2^{fl/fl}* mice. To generate the *Sox2^{CreER/fl}* mice (inducible *Sox2* cKO), *Sox2^{fl/fl}* females were crossed with *Sox2^{CreER/+}* males [*Sox2^{tm1(cre/ERT2)Hoch1/J}*, stock 017593]. *Lgr5^{GFP-CreER/+}* [B6.129P2-*Lgr5^{tm1(cre/ERT2)Cle1/J}*, stock 008875] animals were crossed to generate *Lgr5^{GFP-CreER/GFP-CreER}* embryos (*Lgr5^{KO}*) embryos. *Sox2^{GFP}* males (B6;129S-*Sox2^{tm2Hoch1/J}*, stock 017592) were crossed with NMRI females to produce *Sox2^{GFP}* embryos. Mice were genotyped using the primers listed in Table S6. All aspects of mouse care and experimental protocols were approved by the Finnish National Board of Animal Experimentation.

Tamoxifen administration

A working solution of Tamoxifen (Sigma-Aldrich, T5648) in corn oil (Sigma-Aldrich, 47112-U) was prepared at 50 mg/ml. Tamoxifen solution was sonicated for 15 min and kept at -20°C . Mice were administered 10 mg Tamoxifen solution via oral gavage.

Tissue processing, histology, immunofluorescence, RNAscope, and TUNEL assay

For histology, tissues were fixed in 4% paraformaldehyde (PFA) at 4°C overnight, dehydrated, and embedded in paraffin. Adult samples were decalcified for 2 weeks in 0.5 M EDTA pH 7.5 after fixation. Samples were processed into 5 μm -thick sagittal sections. Hematoxylin-Eosin (H&E) staining was performed as previously described (Juuri et al., 2012).

For RNAscope *in situ* hybridisation (Advanced Cell Diagnostics, ACDbio), we used both the red channel and duplex kits. Mouse tissues were processed into 5 μm sections as described above. Sections were processed using an optimised protocol (as detailed in the supplementary Materials and Methods). Probes were purchased from ACDbio.

TUNEL assay was performed using the *In Situ* Cell Death Detection Kit, Fluorescein (Roche, 11684795910) according to the manufacturer's protocol.

We used the following antibodies in immunofluorescence assays: SOX2 (goat, Santa Cruz, SC-17320; 1:200), keratin 14 (rabbit, NeoMarkers, RB-9020-P; 1:200), P-cadherin (cadherin 3) (goat, R&D Systems, AF761; 1:500), phospho-histone H3 (rabbit, Abcam, ab5176; 1:200), Ki67 (rabbit, Abcam, ab16667; 1:200) and GFP (chicken, Abcam, ab13970; 1:200). As secondary antibodies, we used Alexa 488 goat anti-rabbit (Life Technologies, A11057; 1:500) and Alexa 568 donkey anti-goat (Life Technologies, A11008; 1:500). Nuclei were stained with Hoechst 33342 (Thermo Fisher Scientific, H3570). For DAB immunostaining we used SHH primary antibody (mouse, R&D Systems, AF464) with HRP anti-goat secondary antibody (rabbit, Jackson ImmunoResearch, 305-035-003). Detailed protocols are provided in the supplementary Materials and Methods.

EdU labelling assay

Mice were intraperitoneally injected with EdU (25 $\mu\text{g/g}$ body weight; Thermo Fisher Scientific, A10044) 3 days after the first Tamoxifen administration. Samples were collected 4 days later and processed into 5 μm paraffin sections. Detection was performed using the Click-iT EdU Alexa Fluor 488 Imaging Kit (Thermo Fisher Scientific, C10337) according to the manufacturer's instructions.

Data acquisition and processing

Samples were imaged with a Zeiss Axio Imager M2 microscope and further processed with Adobe Photoshop. RNAscope signal was enhanced by selecting the red or blue-green pixels of the image using the 'select color range tool'. The selected areas were false-coloured using the 'brush tool'.

pH-H3⁺ cell, EdU⁺ cell and laCL volume quantifications

Samples were imaged with a Zeiss Axio Imager M2 microscope and processed with Adobe Photoshop. For embryonic stages, pH-H3⁺ cells were quantified for every second 5 μm -thick section, and the incisor area was drawn by hand using Zen 2011 software (Zeiss). For adult stages, EdU⁺ cells and laCL area were quantified from every second 5 μm -thick section from the central region of the incisor (a total of 70 μm). Volumes were calculated by taking into account the thickness of the sections.

Multiplex quantitative real-time (qRT) PCR

The proximal end of the incisor was dissected out from E18 embryos (approximately one-third of the total length) and stored at -80°C . RNA was extracted using the RNeasy Micro kit (Qiagen, 74004) and reverse transcribed using the QuantiTect Reverse Transcription Kit (Qiagen, 205310).

Multiplex qRT-PCR (CFX96 Touch Real-Time PCR Detection System, Bio-Rad) was performed using iTaq Universal Probe Super Mix (Bio-Rad, 1725130) and 10 ng cDNA per reaction. Probe combinations (PrimePCR Probe Assay, Bio-Rad) are provided in the supplementary Materials and Methods.

Micro-CT and 3D reconstructions

Samples were fixed in 4% PFA, rinsed with PBS, dehydrated in an ethanol series, and stained for 2 weeks in 0.1% phosphotungstic acid (PTA) in 70% ethanol. Samples were scanned with a micro-CT scanner (Phoenix Nanotom). Three mutant and two control lower jaws were scanned for each stage. 3D reconstructions were prepared with Avizo software (Thermo Fisher Scientific) from micro-CT scans (embryonic stages) and from H&E-stained sagittal sections (adult stages).

Microarray

Incisor buds and laCL were microdissected from *Sox2^{GFP}* animals and then incubated with 0.2 U/ml dispase for 10 min at room temperature and the mesenchymal tissue removed. The epithelial explants were incubated with StemPro accutase (Thermo Fisher Scientific, A1110501) for 45 min at 37°C . Samples were passed through a cell strainer (pore size 35 μm) and 7AAD (Thermo Fisher Scientific, A1310) was added to mark dead cells. 7AAD-negative and GFP-positive cells were collected by fluorescence-activated cell sorting (FACS). RNA was extracted using the RNeasy Micro Kit (Qiagen, 74004). Total RNAs were provided to the Functional Genomics Unit, University of Helsinki, Finland, for hybridisation on MTA Affymetrix Arrays. Samples were amplified and labelled using the Affymetrix WT Pico Reagent Kit (Thermo Fisher Scientific, 902622) (see the supplementary Materials and Methods). Three replicates were prepared for each sample, and mESCs were used as a non-dental cell reference. For further analysis, we discarded the non-significant hits ($P > 0.05$, ANOVA).

Statistical analysis

For each experiment $n=3$, except for gene expression analysis in *K14-CreER; Sox2^{fl/fl}* mice where $n=6$ (data not shown). Each incisor is considered one biological replicate (only one incisor was analysed per animal), except for gene expression in *Sox2^{ckO}*, where one litter was considered as a biological replicate ($n=3$). Data are shown as mean \pm s.d. (equal variances not assumed). Unpaired, two-tailed *t*-test was used to test statistical significance. $P < 0.05$ was used as significance threshold.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: M.S.-N., Z.S., B.A.A., O.D.K., F.M.; Methodology: M.S.-N., K.S., O.D.K., F.M.; Validation: M.S.-N., K.S., F.M.; Formal analysis: M.S.-N., O.D.K., F.M.; Investigation: M.S.-N., K.S., L.B.-B.; Resources: M.S.-N., O.D.K., F.M.; Data curation: M.S.-N., F.M.; Writing - original draft: M.S.-N., O.D.K., F.M.; Writing - review & editing: M.S.-N., K.S., Z.S., L.B.-B., B.A.A., O.D.K., F.M.; Visualization: M.S.-N., F.M.; Supervision: F.M.; Project administration: F.M.; Funding acquisition: M.S.-N., O.D.K., F.M.

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Data availability

The microarray data have been deposited at Gene Expression Omnibus under accession number GSE104808.

Supplementary information

Supplementary information available online at <http://dev.biologists.org/lookup/doi/10.1242/dev.155929.supplemental>

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