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FAM20B-catalyzed glycosaminoglycans control murine tooth number by restricting FGFR2b signaling

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Abstract

Background: The formation of supernumerary teeth is an excellent model for studying the molecular mechanisms that control stem/progenitor cell homeostasis needed to generate a renewable source of replacement cells and tissues. Although multiple growth factors and transcriptional factors have been associated with supernumerary tooth formation, the regulatory inputs of extracellular matrix in this regenerative process remains poorly understood.

Results: In this study, we present evidence that disrupting glycosaminoglycans (GAGs) in the dental epithelium of mice by inactivating FAM20B, a xylose kinase essential for GAG assembly, leads to supernumerary tooth formation in a pattern reminiscent of replacement teeth. The dental epithelial GAGs confine murine tooth number by restricting the homeostasis of Sox2(+) dental epithelial stem/progenitor cells in a non-autonomous manner. FAM20B-catalyzed GAGs regulate the cell fate of dental lamina by restricting FGFR2b signaling at the initial stage of tooth development to maintain a subtle balance between the renewal and differentiation of Sox2(+) cells. At the later cap stage, WNT signaling functions as a relay cue to facilitate the supernumerary tooth formation.

Conclusions: The novel mechanism we have characterized through which GAGs control the tooth number in mice may also be more broadly relevant for potentiating signaling interactions in other tissues during development and tissue homeostasis.

Keywords: Glycosaminoglycan, Proteoglycan, Fam20B, Kinase, Supernumerary teeth, Tooth renewal, Tooth replacement, Extracellular matrix, Stem cell, Sox2

Background
The ability to control stem/progenitor cell homeostasis is crucial for generating renewable source of replacement cells and tissues in regenerative medicine. A prerequisite for manipulating the renewal of stem cells is to understand the molecular mechanisms underlying the development of specific cell lineages and fates. The developing tooth organ is an excellent model system for studying the molecular mechanisms and signaling pathways that regulate organogenesis. The hierarchical interactions between the dental epithelium and underlying dental mesenchyme represent a common paradigm in the development of ectodermal placodes deployed in diverse types of epithelium organogenesis, such as salivary glands, lungs, kidneys, mammary glands, hair follicles, and limb buds [1]. Conserved signaling pathways, including those mediated by WNTs, bone morphogenetic proteins (BMPs), fibroblast growth factors (FGFs), and...
sonic hedgehog (SHH), are iteratively used in the cell-cell and cell-matrix communications during tooth development [2]. The secreted morphogens of these cascades interact with extracellular components, such as proteoglycans, to potentiate signal transduction. Proper crosstalk and a fine balance within these signaling pathways are critical for modulating the progressive temporal processes of tooth development, including tooth initiation, morphogenesis, and renewal. While proteoglycans have been identified in murine teeth at various embryonic stages [3–7], their precise inputs into tooth development remain poorly understood.

Proteoglycans consist of a core protein and one or multiple covalently attached glycosaminoglycan (GAG) chains. Based on the disaccharide structures, the polysaccharide GAGs can be classified into heparan sulfate (HS)/heparin, chondroitin sulfate (CS), dermatan sulfate (DS), keratan sulfate (KS), and hyaluronan (HA). Various sulfotransferases give rise to many sulfation patterns and modify the saccharide backbone of sulfated GAGs. The sulfated GAGs are among the most highly negatively charged biopolymers in nature, and variation in the sequence and length of the chains gives rise to enormous polydispersity. This rich structural diversity enables GAGs to interact with various proteins, including components of signaling cascades such as FGFs, WNTs, BMPs, and HHs that are involved in stem/progenitor cell homeostasis [8].

Family with sequence similarity member 20-B (FAM20B) is a kinase that specifically phosphorylates the xylose in the common linkage region of GAGs. The xylose phosphorylation is essential for the linkage region assembly and subsequent GAG elongation [9]. Inactivating FAM20B kinase leads to truncated polysaccharide chains that cannot be further elongated due to impaired function of galactosyl transferase II [10]. This fundamental property of FAM20B provides an useful tool for investigating the molecular functions of GAGs in organogenesis. Given that constitutive inactivation of Fam20B results in embryonic death at E13.5 [11], we generated a Fam20B-floxed allele in mice to facilitate conditional knockout studies. We found that inactivating Fam20B in the dental epithelium led to supernumerary incisors that were formed in a manner similar to replacement tooth formation [12], uncovering a previously unknown function of GAGs in the control of tooth number. Our results demonstrate that the FAM20B-catalyzed GAGs control the tooth number in mice by modulating the commitment of dental epithelial stem/progenitor cells through a mechanism involving the restriction of FGFR2b signaling at the initial stage of tooth development. Our findings provide novel insights into the molecular mechanism regulating tooth number and renewal in mice that may shed light on other GAG-mediated signaling events during organogenesis.

Results
GAG deficiency in the dental epithelium leads to supernumerary incisors in mice
It has been long known that proteoglycans are important molecules regulating signaling pathways during organogenesis. Decades ago, Thesleff et al. reported the expression of proteoglycans in developing murine teeth [13], and subsequent studies have identified multiple proteoglycans in both the dental epithelium and dental mesenchyme at various embryonic stages [6, 7, 14–16]. However, dissecting their mechanistic roles in tooth development has been challenging, because mice lacking individual proteoglycans or a particular type of GAGs did not show overt tooth phenotypes [17]. To explore this issue, we generated Fam20B-floxed mice to disrupt multiple GAGs in embryonic teeth in a tissue-specific manner.

Inactivating Fam20B in the dental epithelium (K14Cre/+; Fam20Bfl/fl) led to formation of duplicate incisors ensuing the native ones, while the molar and diastema regions were not affected [12]. Histology and lineage tracing analyses revealed that the duplicate incisors initiate from an outgrowth of the lingual/mesial side dental lamina of the native enamel organ at the late cap stage (~E15.0) (Figs. 1E–H and 2N–Q). Of note is that both the native and duplicate incisors retained the ability of continuous growth and did not show apparent difference in the growth rate.

Murine tooth number is specifically modulated by the GAGs in the dental epithelium
Supernumerary teeth have been implicated with alterations of the interactions between the dental epithelium and dental mesenchyme [18]. Signaling changes in the dental mesenchyme frequently cause extra tooth formation, such as those in Gas1−/−, Sostdc1−/−, Spry4−/−, Wnt1Cre; Polaris−/− and Osr2−/− mice [19]. To investigate whether GAGs in the dental mesenchyme also play a role in controlling the tooth number in mice, we generated Wnt1Cre; Fam20Bfl/fl mice to disrupt GAGs in the neural crest-derived mesenchymal cells. These mice did not show any changes in tooth number (Fig. 1I–L), although several mesenchyme-associated defects occurred as expected in their craniofacial complex [20]. This indicates that the murine tooth number is specifically modulated by the GAGs in the dental epithelium but not in the mesenchyme. Hence, we have focused on exploring its role in the dental epithelium.

GAGs commit the cell fate of dental epithelium at the initial stage of tooth development
Tooth development in mice initiates from a thickening of the dental epithelium at E10.5 to form a placode. The epithelial placode then invaginates into the dental mesenchyme to form a tooth bud, which further folds into an enamel organ in the following stages. In K14Cre/+; Fam20Bfl/fl mice, the earliest sign of replacement tooth
formation appeared at the late cap stage (~E15.5) as an ectopic placodal thickening of the dental epithelium at the mesial-lingual side of the native enamel organ (Fig. 1E–H). TUNEL and EdU incorporation analyses on earlier stages showed overproliferation in both the dental epithelium and dental mesenchyme and reduced apoptosis in the dental epithelium at the lingual/mesial side of native enamel organs starting at ~E12.5 in the K14Cre/+;Fam20Bfl/fl mice (Fig. 2A–M). This suggests that the cell fate of the GAG-deficient dental epithelium had been changed prior to the cap stages. To determine a precise timing of the cell fate change, we employed a Tet-On system to knockout Fam20B from the dental epithelium at different embryonic stages. The results demonstrated that replacement tooth formation was induced only if Fam20B was inactivated within a time window between E10.5 and E12.5 (Additional File 1: Fig. S1) (Table 1). This implies that the cell fate of primary dental lamina was committed by FAM20B-catalyzed GAGs at the initial stage of tooth development [21, 22]. In cKO (K14Cre/+; Fam20Bfl/fl;Sox2GFP) mice, the Fam20B-deficient incisors showed progressively increasing expression of ectopic Sox2 at the lingual side of enamel organ from E13.5 to E15.5. In contrast, the Sox2 signal gradually vanished from the lingual side of the enamel organ in control incisors (K14Cre/+;Fam20Bfl/+;Sox2GFP) during these stages (Additional File 2: Fig. S2). At E16.5, the normal enamel organ completely lost Sox2(+) expression (Fig. 2P), while the cKO mice showed large amount of Sox2(+) cells in both the lingual side of the native enamel organ, the dental lamina and the enamel organ of the supernumerary teeth (Fig. 2Q). These results suggest that GAG deficiency in the dental epithelium leads to ectopic or extended renewal of Sox2(+) stem/progenitor cells.

Subsequently, we deleted Sox2 from the dental epithelium of K14Cre/+;Fam20Bfl/fl mice by introducing the Sox2fl/fl allele to examine the contribution of Sox2(+) progenitor cells to the supernumerary tooth formation in the GAG-deficient teeth. The resultant mice showed partial rescue of the supernumerary tooth phenotype as well as a reduced size of the native teeth (Additional File 3: Fig. S3). To further explore the regulatory input of GAGs on the renewal of Sox2(+) cells, we deleted Sox2-expressing stem/progenitor cells are believed to contribute to the whole enamel organ tissues during tooth development [21, 22].
Fig. 2 (See legend on next page.)
Fam20B from the Sox2(+) population at E11.5 using Sox2-CreER, the efficiency of which was validated by crossing with tdTomato indicator (Additional File 4: Fig. S4A). The Sox2CreER;Fam20Bfl/fl mice did not recapitulate the supernumerary tooth phenotype (Additional File 4: Fig. S4B), suggesting that FAM20B-catalyzed GAGs restrict the renewal of Sox2(+) cells in a non-autonomous manner.

GAGs determine the cell fate of dental epithelium by restricting FGFR2b signaling

Multiple signaling pathways, such as those mediated by WNTs, FGFs, BMPs, and HH, have been implicated in cell fate commitment in embryonic teeth [23, 24]. To begin to investigate the molecular mechanisms by which GAGs determine the cell fate of dental epithelium, we systematically screened signaling pathways potentially associated with the phenotype at the initial stage of tooth development using immunohistochemistry, in situ hybridization, and signaling reporter lines in mice. We did not detect significant changes in WNT and BMP signaling in the Fam20B-mutant incisors (Additional Files 5 and 6: Figs. S5 and S6). However, we identified a hyperactivation of FGF in the Fam20B-deficient incisors at E12.5 and E13.0, as indicated by a robust upregulation of Pax9, Etv-5, and p-ERK in the dental epithelium and/or dental mesenchyme (Fig. 3). In agreement with this, the transcription of Shh, a downstream gene of FGF signaling [25], showed an expanded scope of expression at the presumptive location of replacement teeth formation in the GAG-deficient dental epithelium. Accordingly, two SHH downstream markers in the dental mesenchyme, Gli1 and Patched1, showed broader responses to the epithelial HH signaling, as indicated by Gli1- and Ptch1-LacZ indicator mice (Fig. 4).

Several FGFs and their receptors are expressed in incisors during the critical time window (E11.5-E12.5) identified in the Fam20B-mutant mice for supernumerary tooth formation: FGF1, FGF2, and FGF9 are present in the dental epithelium, while their primary receptor, FGFr1c, is expressed in the dental mesenchyme. FGF10 is localized in the dental mesenchyme whereas its receptor, FGFr2b, is exclusively present in the dental epithelium [26] (Additional File 7: Fig. S7). The complementary expression pattern between FGF ligands and their receptors requires FGFs to diffuse to the counterpart location/tissue to perform their functions. We excluded several FGFs and their receptors based on their incongruent expression pattern and timing. For example, FGF8 was excluded for lacking an expression in incisors during this stage, while FGF3 and FGF4 were excluded for their expression timing later than E13.5, etc.

Table 1 Supernumerary teeth formed when Fam20B inactivation was induced between E10.5 and E12.5

| Induction time | Mice with Supernumerary teeth | Mice without Supernumerary teeth | Tested animals |
|---------------|-------------------------------|----------------------------------|----------------|
| E10.5         | 21 (100%)                     | 0                               | 21             |
| E11.5         | 17 (94%)                      | 1                               | 18             |
| E12.5         | 9 (31%)                       | 20                              | 29             |
| E13.5         | 2                             | 16                              | 18             |
| E14.5         | 0                             | 7                               | 7              |
It is well documented that FGFs in the dental epithelium and the dental mesenchyme stimulate one another via a positive feedback loop, which is negatively regulated by Sprouty [27, 28]. Previous studies indicated that inactivation of Sprouty led to supernumerary teeth in both diastema and incisor regions due to epithelium↔mesenchyme bidirectional hyperactivity of FGF, in which FGF10/FGF2b signaling appeared to play a major role compared with FGF3/FGF1c [27].

To determine whether the hyperactivated FGF signaling underlies the cell fate change of the GAG-deficient dental epithelium associated with the extra teeth formation, we employed a Tet-On system to inhibit FGFR2b signaling through overexpressing an FGFR2b inhibitor (dominant-negative FGFR-HFc protein) [29] in the Fam20B-deficient dental epithelium. Overexpression of the FGFR2b inhibitor successfully rescued the supernumerary tooth phenotype (Fig. 5) and clearly exhibited a trend whereby earlier
inhibition resulted in better rescue effects (Table 2). In agreement with this, inhibiting FGFr2b signaling in the Fam20B-deficient dental epithelium also reduced the expression scope of Shh back to the normal size (Additional File 8: Fig. S8), indicating that Shh is downstream to the FGF hyperactivation that initiates the supernumerary tooth formation. These results collectively indicate that FAM20B-catalyzed GAGs control the cell fate of dental epithelium by confining the FGF10/FGFR2b signaling at the initial stage of tooth formation. The confining effects are most likely associated with FGF10/FGFr2b reactivity/transmission but not their expression, because RNAScope assays of Fgf10 and Fgfr2b did not detect any changes in the Fam20B-deficient incisors (Additional File 7: Fig. S7).

**Fig. 4** SHH signaling was overactivated in the GAG-deficient incisors. A–D Whole mount in situ hybridization of E12.0 mouse embryos showed an expanded territory of Shh (white arrows) in both upper and lower incisor areas of the K14Cre+/Fam20Bfl/fl mice compared with the controls (Fam20Bfl/fl). Note that Shh signal in hair follicles can be used as internal references for the comparison between knockout and control mice. E,F ISH of Shh on the coronal sections of lower incisors showed an expanded expression of Shh in the GAG-deficient dental epithelium toward the mesial side (white arrow) compared with controls. G–J HH downstream markers, Gli1 and Patched1, showed broader reactivity in the dental mesenchyme of K14Cre+/Fam20Bfl/fl mice, as indicated by Gli1- and Pimch1-LacZ indicators (dashed lines plotted). Scale bars: 500 μm in A and B, 250 μm in C, D, and G–J, 50 μm in E and F.
GAGs may regulate FGFR2b signaling by confining the diffusion gradient of FGF10

In order to determine how FAM20B loss of function affects GAG assembly in the dental epithelium, we compared the GAG profile of Fam20B-deficient dental epithelium with WT using a newly developed GAG profiling method that relies on multiple reaction monitoring liquid chromatography mass spectrometry (MRM-LCMS) [30]. The amount of HS, CS, and total GAGs were remarkably reduced in the Fam20B-deficient dental epithelium (Fig. 6A), whereas their total composition did not show apparent changes (Fig. 6B). The HS composition in the Fam20B-deficient dental epithelium showed reduced NS and increased 0S (Fig. 6C), while the CS composition in Fam20B-deficient dental epithelium did not show apparent differences from the WT (Fig. 6D).

GAGs may regulate growth factor signaling in various manners, including shaping the diffusion gradients and mediating the interactions between the growth factors and their receptors [31]. In a recent study, we revealed

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**Table 2** The timing and effects of inhibiting FGFR2b in rescuing the supernumerary tooth formation in Fam20B-deficient mice

| Induction time | Fully rescued | Partially rescued | Not rescued | Tested animals |
|---------------|---------------|------------------|-------------|----------------|
| E10.5         | 6 (46%)       | 7 (53%)          | 0           | 13             |
| E11.5         | 3 (23%)       | 8 (69%)          | 1           | 13             |
| E12.5         | 2 (11%)       | 14 (77%)         | 3           | 18             |
| E13.5         | 0             | 1                | 6           | 7              |
| E14.5         | 0             | 0                | 6           | 6              |

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Fig. 5 Inhibiting FGFR2b rescued the supernumerary tooth phenotype. a The schematic depicts the rescue strategy of supernumerary tooth formation by inhibiting FGFR2b with a dominant-negative FGFR2b inhibitor through a tetO system. The K14rtTA, tetOFGFr2b/Igh, and K14CreK14rtTA;Fam20Bfl/fl mice were crossbred to ultimately generate K14rtTA;tetOFGFr2b/Igh;K14CreK14rtTA;Fam20Bfl/fl mice. Doxycycline was administrated to the resultant mice at 2 mg/kg at designated time points (E10.5–E14.5) to induce the expression of the dominant-negative FGFR2b inhibitor, FGFR2b/Igh, in the GAG-deficient dental epithelium. b–e Inhibiting FGFR2b in the dental epithelium fully (D) or partially (E) rescued the supernumerary tooth phenotype in the K14rtTA;Fam20Bfl/fl mice.
that the interactions between FGF10 and heparin are chain-length dependent, and the minimum binding size for the interactions is dp6 [32]. As cells lacking FAM20B cannot extend GAGs beyond the tetrasaccharide linkage and form very short saccharides [10], we estimate that the truncated saccharides of FAM20B-mutant GAGs cannot bind FGFs. In this case, we assayed the interactions between FGF10 and FAM20B non-mutant GAGs (heparin, HS, DiS HS, TriS HS, and CSA) at variable concentrations (0.00, 0.02, 0.08, 0.30, 1.25, and 5.00 μg/ml) in solution culture of BaF3 cells that had been engineered to report FGFR2b reactivity [33]. Heparin served as a positive control in the assays for its known synergistic effects on FGFR2b signaling [34]. Our results showed that none of the tested GAGs, except for the positive control heparin, had significant synergistic or inhibitory effects on FGFR2b signaling (Additional File 9: Figs. S9A and S9B), indicating that GAGs may not restrict FGF10-FGFR2b signaling through modulating ligand-receptor interactions.

Given the complementary expression pattern between FGF10 and FGFR2b, FGF10 needs to diffuse to the dental epithelium to perform its function. We built a hydrogel cell culture system mimicking the ECM environment to test if GAGs (CSA, CSB, HS, Des-2S heparin, and Des-6S heparin) confine the diffusion of FGF10 at certain concentrations (1 μg/ml, 3 μg/ml, and 5 μg/ml) (Fig. 7 and Additional File 10: Fig. S10). We focused on the 2S and 6S since these functional groups are critical in FGF interactions. The requirements of NS and NAc were examined in testing heparin and HS. GAGs at 1 μg/ml and 3 μg/ml displayed

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**Fig. 6** GAG profiling of Fam20B-deficient dental epithelium. A The amount of HS, CS, and total GAGs were remarkably reduced in the Fam20B-deficient dental epithelium compared to the WT (n = 1). B The HS and CS composition relative to the total GAGs were similar between KO and WT (n = 2). C Detailed HS composition showed reduced NS and increased 0S in the Fam20B-deficient epithelium (n = 2). D CS composition in the KO did not show apparent differences from the WT (n = 2). The analytical error was ~ 3% for these samples.
differential confining effects on FGF10 diffusion (Fig. 7c, d). In particular, 3 μg/ml GAGs confined FGF10 diffusion in a preference pattern: Des-2 heparin > Des-6 heparin > HS > CSB > CSA (Fig. 7d), and 1 μg/ml GAGs showed a similar preference pattern except for Des-2 heparin displaying a reduced confining effect (Fig. 7c). At 5 μg/ml, all tested GAGs showed almost equally strong inhibition on FGF10 diffusion (Fig. 7e). These data collectively suggest that GAGs differentially confine the diffusion of FGF10 based on their composition and sulfation states in a dose-dependent manner.

Hyperactivated WNT signaling serves as a relay cue in the GAG-mediated supernumerary tooth formation

Subsequent to the cell fate commitment of the GAG-deficient dental lamina, we identified an ectopic hyperactivation of canonical WNT signaling in the dental epithelium and adjacent dental mesenchyme at the lingual/mesial side of the native enamel organ starting at E14.5 (~1 day before the thickening of the extra dental lamina). This is demonstrated by an ectopic expression of LEF-1 (Fig. 8A–F) and BAT-GAL indicator (Fig. 8G, H) compared to the control mice. Conversely, a WNT inhibitor, Sostdc1, was downregulated in the dental follicle at the presumptive location of the supernumerary teeth (i.e., at the lingual/mesial side of native enamel organ) (Fig. 8I, J). The molar regions did not show WNT hyperactivity (data not shown).

We employed a Tet-On system to overexpress DKK1 [35], a WNT inhibitor in the Fam20B-deficient dental epithelium at E13.5 to determine the biological significance of the WNT hyperactivation in supernumerary tooth formation. The Dkk1 transgene fully rescued the tooth phenotype in the K14Cre/+;Fam20Bfl/fl mice (Fig. 8K–M), indicating that the ectopic hyperactivation of WNT signaling is essential for accomplishing the supernumerary tooth formation. However, the upregulation of WNT signaling is not a direct consequence of GAG deficiency, because inactivating Fam20B in the dental epithelium at E13.5 (~1 day before the overactivation of WNT in the K14Cre/+;Fam20Bfl/fl mice) did not cause...
supernumerary tooth formation or any WNT activity changes as indicated by BATGAL indicator (cKO mice K14rtTA;tetOCre;Fam20Bfl/fl;BATGAL versus control mice K14rtTA;tetOCre;Fam20Bfl/+;BATGAL) (data not shown). These results collectively suggest that the ectopic hyperactivation of WNT is a secondary reaction to the GAG deficiency in the dental epithelium and a relay cue for accomplishing the supernumerary tooth formation.

Discussion
Interplay between growth factors is involved in the hierarchical and iteratively used signaling cascades that guide tooth development. These secreted proteins interact with extracellular components to transmit signaling intracellularly. Although accumulating evidence shows that proteoglycans in the extracellular matrix and on the cell surfaces are pivotal signaling regulators for the morphogenesis of multiple organs, their role in tooth development remains poorly understood. In this study, we demonstrate a novel molecular mechanism for how dental epithelium is pre-programmed by GAGs in the control of tooth number in mice. FAM20B-catalyzed GAGs control murine tooth number by committing the cell fate of dental epithelial stem/progenitor cells. This is achieved through restriction
of FGFR2b signaling at the initial stage of tooth development. At the cap stages, WNT signaling then relays relevant cues to complete the replacement tooth formation (Fig. 9A, B).

The supernumerary incisors in Fam20B-mutant mice initiate from an outgrowth of the lingual/mesial part of the native enamel organs at the late cap stage. This growth pattern is very similar to the supernumerary incisors in Sostdc1-mutant mice, which also originate from part of the native teeth, reminiscent of replacement tooth formation [36]. Of note is that Sostdc1-mutant mice also develop diastema teeth that are derived from revitalized diastema rudiments [37], and the gene profile change associated with the extra tooth formation is very different from that in the Fam20B-mutant mice. Previous studies showed that epithelial stabilization of canonical WNT signaling revitalizes the rudiments in molar and diastema regions [38–41]. In the Fam20B-knockout mice, the WNT hyperactivity is confined to the incisor regions starting at the late cap stage and appeared to be a secondary reaction to the hyperactivity of FGF signaling at the initial stage, because removing Fam20B from the dental epithelium at the cap stage failed to induce FGF/WNT hyperactivity and supernumerary teeth. These discrepancies suggest intrinsic differences between the development of molars and incisors and illustrate the complexity of the regulatory mechanism for the control of murine tooth number.

Sox2-expressing stem/progenitor cells in the dental epithelium are believed to hold the odontogenic potency in different modes of tooth development, including tooth initiation, tooth replacement, and the continuous growth of rodent incisors [19–23, 42]. The role of Sox2(+) cells in the labial cervical loop of murine incisors represents a paradigm of organ renewal derived by adult stem cells [21–23, 43–45]. FGF8 in the stellate reticulum of the cervical loop is believed to maintain the Sox2(+) population in the postnatal incisors [21]. However in embryonic teeth, it remains unclear if a similar molecular mechanism applies to the Sox2(+) population that initiates tooth replacement. Our genetic analyses reveal that the GA-deficient incisors revived tooth replacement-like capacity but retained the continuous growth, and there were no growth rate differences between the native and replacement incisors. This indicates that the Sox2(+) cells are differentially regulated for tooth renewal and tooth replacement in the cervical loop of postnatal incisors and in the dental epithelium of embryonic teeth.

FGF10-FGFR2b signaling has been associated with stem cell recruitment and maintenance in multiple tissues [46–49]. In embryonic teeth, FGF10 is dominantly present at the initial stage of tooth development [26]. Overexpressing FGF10 in zebrafish produces supernumerary and bicuspid teeth [28]. We found that FAM20B-catalyzed GAGs inhibit tooth replacement of murine incisors by confining the renewal of Sox2(+) cells in the dental epithelium via restricting FGFR2b signaling. This provides genetic evidence that the Sox2(+) cells in the embryonic teeth are differentially regulated by FGFs compared to those in the postnatal teeth. Hence, they modulate different tooth renewal patterns in the processes underlying tooth replacement versus continuous growth in mice.

Although the lineage tracing in this study clearly indicates that Sox2(+) cells contribute to the supernumerary tooth formation in GAG-deficient incisors, deleting Sox2 from Fam20B-deficient dental epithelium only partially rescued the phenotype. Sanz-Navarro et al. [50] also observed mild tooth phenotypes when they removed Sox2 from Shh(+)-population in the dental epithelium. They speculated that it may be related to the mosaic activation of Shh-Cre and/or redundancy from other Sox members. Together with our data, it appears that Sox2 is dispensable in the dental epithelium for odontogenesis, although it is an excellent marker for the dental epithelial stem/progenitor cells.

Previous studies have identified undersulfated GAGs on pluripotent embryonic stem cells (ESCs) whereas highly sulfated GAGs on differentiated cells, indicating that the sulfation pattern of GAGs are implicated with the progression of ESCs from self-renewal to a differentiated state [51, 52]. An explanation is that the stemness of ESCs is protected from the growth factor signaling by a shield of minimally sulfated GAGs [51]. However, this mechanism may not apply to the stem/progenitor cells in the dental epithelium of mice. The sulfation loss in the dental epithelial GAGs could be a secondary effect that is derived from the dramatic reduction of GAGs. More importantly, the undersulfated GAG remnants in the Fam20B-deficient dental epithelium appeared not to shield growth factors, because FGF signaling was upregulated rather than downregulated/abolished.

Accumulating evidence shows that HS/heparin 6-O- and 2-O-sulfo groups modulate FGF10-mediated epithelial morphogenesis and differentiation (such as those in submandibular and lacrimal glands) by increasing the affinity of FGF10 to FGFR2b, which forms an FGF10-FGFR2b-HS ternary signaling complex to result in diverse biological outcomes [53–55]. When Hs2st (heparan sulfate 2-O-sulfotransferase) and Hs6st (heparan sulfate 6-O-sulfotransferase 1) were both inactivated, the formation of FGF10-FGFR2b-heparan sulfate complex was disrupted on the cell surface and completely abolished lacrimal gland development [54]. In Fam20B-knockout mice, there was no apparent reduction of endogenous HS 6-O- and 2-O-sulfo groups in the Fam20B-deficient dental epithelium. An assumpive disruption of the FGF10-FGFR2b-HS complex in the Fam20B-deficient dental epithelium indeed promoted rather than attenuated FGF10-FGFR2b signaling. This in turn promoted the renewal but not differentiation of the Sox2(+) stem/progenitor cells, strongly suggesting an inhibitory role...
Fig. 9 (See legend on next page.)
of GAGs on FGF10-FGFR2b signaling. Our results suggest that GAGs in the dental epithelium are unlikely to inhibit FGF10-FGFR2b signaling by serving as co-receptors. Instead, they may regulate the signaling by sequestering FGF10 to restrict the diffusion gradient. The differential regulation of FGF10-FGFR2b signaling between the dental epithelium and other epithelium-derived organs also illustrates that GAG-mediated regulation of growth factors is highly dependent on the biological context.

It is interesting to note that the biological effects of GAG deficiency were very specific in Fam20B-deficient dental epithelium despite the broad spectrum of proteins potentially interacting with GAGs. Similar phenomenon has been observed in many other GAG-deficient animal models [17]. The specificity of GAG-growth factor interaction may be derived from a specific polysaccharide sequence, a polysaccharide conformation, an accurate control of enzymatic modification on saccharides, or a dominant presence of growth factors in certain biological context [56]. A prerequisite to answering these questions will require the development of oligosaccharide libraries with systematically varied structures and more sophisticated GAGosoma analyses, such as oligosaccharide mapping and sequencing, which are emerging technologies [57].

In summary, this study reveals that Fam20B-catalyzed GAGs determine the monophyodont phenotype in mice by restricting the capacity for renewal of dental epithelial stem/progenitor cells through inhibition of FGFR2b signaling at the initial stage of tooth development. The GAGs interact with FGF ligands to shape a restricted diffusion gradient of FGF10, which maintains a subtle balance between the renewal and differentiation of Sox2(+) cells in the dental epithelium. Disrupting GAG assembly breaks this balance by overactivating FGFR2b signaling, which overweights the renewal of Sox2(+) cells and initiates the replacement tooth formation in monophyodont mice (Fig. 9C, D).

Conclusion
In conclusion, this study demonstrates that the Fam20B-catalyzed GAGs control the number of murine teeth by regulating the commitment of dental epithelial stem/progenitor cells through a mechanism involving the restriction of FGFR2b signaling at the initial stage of tooth development. This novel mechanism may also be more broadly relevant for potentiating signaling interactions in other tissues during development and tissue homeostasis.

Materials and methods

Animals
All of the animal experiments were carried out according to the protocol approved by the Institutional Animal Care and Use Committee of Texas A & M University College of Dentistry (Dallas, TX, USA) and performed in accordance with the NIH Guide for the Care and Use of Laboratory Animals.

Fam20Blox/lox mice and TetOΔK1 mice were generated as previously described [12, 35]. BRE-LacZ mice were kindly gifted by Dr. Leif Oxburgh (Maine Medical Center Research Institute). Mice purchased from Jackson Laboratory (Bar Harbor, ME, USA): KI4Cre (stock #004782), Wnt1Cre2 (stock #022137) [58], KI4Fta (stock #008099) [59], TetOCre (stock #006224) [60], TetOFgfy2b/fgfy2b (stock #025672) [29], Rosa26Tmato (stock #007909) [61], BAT-Gal (stock #005317) [62], Gl(1lacZ (stock #008211) [63], Pch1lacZ (stock #003081) [64], Sox2GFP (stock #017592) [65], Sox22creR (stock #017593) [65], and Sox22fl (stock #013093) [66]. Genotyping were carried out as previously described [12] or following Jackson Lab’s instructions.

Pregnant mice were fed with Doxycycline chow (1 g/kg, Bio-Serv, NJ, USA) and water (2 mg/ml, Sigma, St. Louis, MO, USA) at designated time points for 3 consecutive days to induce TetO transgene expression. Tamoxifen (T5648, Sigma, St. Louis, MO, USA) was injected to pregnant mice intraperitoneally at 50 mg/kg at designated time points for 3 consecutive days to induce CreER expression.

Tissue preparation
Embryonic stage was determined according to the vaginal plug (day 0.5) and confirmed by morphological criteria. Mouse embryos of desired stages were harvested in diethyl pyrocarbonate (DEPC)-treated Dulbecco’s phosphate-buffered saline (PBS). Dissected heads from the embryos were fixed with 4% paraformaldehyde (PFA) in 0.1% diethyl pyrocarbonate(DEPC)-treated PBS at 4°C overnight and decalcified in 0.1% DEPC-treated 15% EDTA (pH 7.4) at 4°C for 1 to 2 days as needed.
then dehydrated in a serial gradient of ethyl alcohol solutions followed by paraffin embedding.

Tissue clearing and 3D imaging
Mandibles for 3D-image reconstruction were collected from 6 embryos of KO or control mice at each desired time point. Tissue clearing was performed on the mandibles as previously described [67]. Fluorescent images were acquired with Zeiss LSM780 two-photon microscopy (Visible laser lines: 488,633 nm). Image processing and 3D rendering were performed with Imaris 9.0 (Bit-plane) as previously described [67].

Immunohistochemistry (IHC)
Immunohistochemistry staining was performed on 4-μm-thick coronal sections prepared from 6 mandibles of KO or control mice using a DAB substrate kit (Vector Laboratories, Burlingame, CA, USA) following the manufacturer’s instruction. The primary antibodies used for immunohistochemistry are: anti-phospho-ERK (9101, 1:200, Cell Signaling, Danvers, MA, USA), anti-Lef1 (C12A5, 1:200, Cell signaling, Danver, MA, USA), anti-β-catenin (sc-7963, Santa Cruz), anti-Sox2 (ab97959, Abcam, Cambridge, MA, USA), and anti-phospho-SMAD1/5 (9516, Cell Signaling). Immunohistochemistry staining was performed on 4-μm-thick sections prepared from 6 mandibles of KO or control mice according to the manufacturer’s instructions. Slides were baked for 1 h at 60 °C prior to use. After dehydration, the tissues were air dried and treated with peroxidase blocker before boiling at 100–104 °C in target retrieval reagents for 15 min. Protease was then applied for 30 min at 40 °C. Target probes Fgf10, Fgfr2b, and Fgf9 (446371, 806301, 499811, Advanced Cell Diagnostics) were hybridized for 2 h at 40 °C, followed by a series of signal amplification and washing steps. All hybridizations at 40 °C were performed in a HybEZ Hybridization System. RNA staining signal was identified by DAB as brown chromogenic dots. Following the RNAscope assay, samples were counterstained for 2 min with hematoxylin. Each sample was quality controlled for RNA integrity with a probe specific to the housekeeping gene cyclophilin B (PPIB); only samples with an average of > 4 dots per cell were included for analysis. Negative control background staining was evaluated using a probe specific to the bacterial dapB gene; only samples with an average of < 1 dot per 10 cells were included for analysis. Bright field images were acquired by Olympus CKX41 inverted microscope using a × 40 objective.

For semi-quantitation analysis, the RNAscope signal is scored on the basis of number of dots per cell as follows: 0 = 0 dot/cell, 1 = 1–3 dots/cell, 2 = 4–10 dots/cell, 3 = 10–15 dots/cell, and 4 = > 15 dots/cell with > 10% of dots in clusters. To evaluate heterogeneity in marker expression, H-score analysis is performed. The H-score is calculated by adding up the percentage of cells in each scoring category multiplied by the corresponding score, so the scores are on a scale of 0–400. The RNAscope signal area proportion of each probe is quantified by Image J based on × 20 tooth bud images (250 × 180 μm).

Cell proliferation assay (EdU) and TUNEL staining
Timed pregnant mice were injected intraperitoneally with EdU at 15 μg/kg in PBS (C10352, Invitrogen, Carlsbad, CA, USA). After 1 h of injection, embryo heads were collected and processed for paraffin embedding and section. EdU incorporation was detected on 5-μm-thick paraffin sections using a Click-iT Kit (Invitrogen) following the manufacturer’s protocol. Apoptotic cells were identified on the sections by TUNEL staining using an ApopTag Plus In Situ Apoptosis Fluorescein Detection Kit (S7111, Millipore, Burlington, MA, USA) according to the manufacturer’s instruction. DAPI was used as counterstaining. Mounted sections were examined and photographed using an SP5 confocal microscope (Leica, Buffalo Grove, IL, USA).

X-Gal staining
The embryos for whole-mount X-Gal staining were fixed with 0.2% glutaraldehyde in PBS at 4 °C for 30 min. After
three wash in 0.005% NP-40 and 0.01% sodium deoxycholate, the embryos were incubated in staining solution (5 mM potassium ferrocyanide and potassium ferricyanide, 2 mM MgCl2, 0.4% X-Gal in dimethylformamide) at 37 °C for 3–24 h, followed by post-fixation with 4% paraformaldehyde (PFA) in PBS at room temperature for 1 h. Embryos for cryosection were fixed in 4% PFA for 1–2 h at room temp and dehydrated in 30% sucrose at 4 °C overnight, then embedded in OCT for cryosection. X-Gal staining was performed on the cryosections, and nuclear fast red was used for counterstaining.

**GAG profiling**

GAG profile of E11.5 dental epithelium in WT and Fam20B-deficient (KO) mice were characterized regarding the GAG type, amount, sulfation, and disaccharide composition using a recently developed method MRM-LCMS (multiple reaction monitoring liquid chromatography mass spectrometry) [70]. Briefly, the dental epithelium of lower incisors was isolated from the mandibles of E11.5 KO and WT embryos after dispase digestion (1.8 U/ml in Ca- and Mg-free PBS, Gibco) at 37 °C for 30 min. The epithelium pooled from 6 embryos of each group were lysed in digestion buffer (50 mM ammonium acetate, 2 mM calcium chloride) and digested by cocktail of GAG-lyases (heparin lyase I, II, III, and chondroitin lyase ABC (10 mU each), then placed in 37 °C incubator overnight. The resulting disaccharides were recovered by centrifugal filtration, labeled with 2-aminoacridone (AMAC), and analyzed by liquid chromatography mass spectrometry (LC-MS/MS, Thermo Inc.) running at multiple reaction monitoring mode. The separation was carried out with an Agilent 1200 HPLC separation system on an Agilent Poroshell 120 ECC18 column (3.0 × 150 mm, 2.7 μm, Agilent, USA) at 45 °C. The analytical error for GAG profiling was < 3%.

**Cell culture**

The BaF3 cells used in this study were engineered to stably express FGFR2b [71]. The cells were maintained in RPMI 1640 culture media (Gibco Life Science, Gaithersburg, MD, USA) supplemented with 10% newborn calf serum (Gibco), 0.5 ng/ml murine recombinant interleukin-3 (Gibco), 2 mM L-glutamine, penicillin/streptomycin (P/S), and 50 nM β-mercaptoethanol (Gibco). The cells were treated with G418 (600 μg/ml, Gibco) for 2 weeks before being used for the subsequent assays.

**Cell proliferation assay**

BaF3-FGFR2b cells (4 × 10⁴) were seeded in each well of 96-well plates in culture medium without interleukin-3. The culture medium was supplemented with or without 1.5 μg/ml heparin (as co-receptor for FGF signaling) for the assays of inhibitory or synergistic effects of GAGs on FGF10-FGFR2b reactivity. FGF10 (1000 pM) in 10% BSA and 0–5 μg/ml (0, 0.02, 0.08, 0.30, 1.25, and 5.00 μg/ml) of HS, DiS HS, TriS HS, CSA or heparin were added to each well in duplicates for each group. The cells were incubated at 37 °C for 36–48 h before being assayed for cell number/viability with the CCK-8 kit (Sigma) following the manufacturer’s instruction. Two-way ANOVA shows that heparin is different from all other GAGs, p < 0.001. DiS HS was synthesized from N-sulfo heparosan with modification using C5-epimerase and 2-O-sulfotransferase (2OST) following our previously reported procedures [72]. TriS HS (NS2S6S) was synthesized from N-sulfo heparosan with subsequent modification with C5-epimerase, 2-O-sulfotransferase, and 6-O-sulfotransferases (6OST1/6OST3) [72].

**3D hydrogel cell culture**

A hydrogel cell culture system was used to mimic the extracellular matrix context in the dental epithelium (Fig. 7a). Briefly, 1.0 μg/ml, 3.0 μg/ml, or 5.0 μg/ml of each GAGs (CSA, CSB, HS, Des-2, and Des-6) in 10 × RPMI1640 and 2.0 × 10⁻³/ml BaF3-FGFR2b cells were mixed with premade collagen solution (2 mg/ml, C4243, Sigma). In each well of 24-well plates, 500 μl of such mixture was dispensed and incubated at 37 °C for 1 h to allow gelation. The hydrogel cylinders were then maintained in RPMI culture media supplemented with 100 ng/ml recombinant human FGF-10 (Invitrogen) for further analyses.

**Cell viability assay**

Hydrogel was collected for cell viability assay on day 2 of culture using a Live/Dead Viability Kit (Invitrogen). After washing in PBS for 5 min, the hydrogel was incubated in 2 μM calcein AM and 4 μM ethidium homodimer-1 prepared in PBS in the dark at 37 °C for 40 min, then washed in PBS for imaging. Fluorescent images were obtained with a Zeiss LSM 880 two-photon microscopy (Visible laser lines: 406 and 488 nm). The number of live cells in each sample was counted in 3D hydrogel chips reconstructed from 6 randomly selected areas using Imaris 9.0 (Bitplane) and subjected to statistical analysis.

**Statistics**

The data was expressed as mean ± SD of at least 6 determinations in all experiments unless otherwise indicated. We used 2-sample t tests to evaluate the pairs of samples and independent-samples T test for the independent samples. Before performing t test, normal distribution was verified by Levene’s test using SPSS (IBM, NY, USA). Two-way ANOVA was used to compare multiple groups. A P value of < 0.05 was considered to indicate statistically significant differences.
Additional file 1: Figure S1. GAGs commit the cell fate of dental epithelium at the initial stage of tooth development. A K14Cre, Rsa26-DT-Tomato mice were crossed to ultimately generate K14CreERT2Rosa26-DT-Tomato mice. B Doxycycline (Dox) was administrated to the transgenic mice at 2 mg/kg/day for one week, followed by 5 mg/ml FGF10 for 45 h. Heparin (positive control) showed > 0.05). A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q, R, S, T, U, V, W, X, Y, Z. Additional file 2: Figure S2. The dynamics of Sox2 expression during supernumerary incisor formation. H&E staining of K14Cre; Sox2fl/fl; Rosa26-Tomato mice was performed on sagittal sections of lower incisors. Yellow arrows indicate the orientation of labial and lingual sides. A, B At E12.5, the Fam20B-deficient dental epithelium started showing more Sox2 expression than normal. C, D At E13.5, the control incisors showed reduced expression of Sox2 in the dental epithelium, while the Fam20B-deficient dental epithelium had strong expression of Sox2 in the lingual side of the enamel organ (arrow). E-H At E14.5 and E15.5, the control incisors gradually lost Sox2 expression from the lingual side of enamel organ. In contrast, the Fam20B-deficient incisors showed strong ectopic Sox2 expression at the lingual side of enamel organ (arrows). Scale bars, 50 μm in A and B, 100 μm in C and D, 200 μm in E and F, 400 μm in G and H.

Additional file 3: Figure S3. Inactivation of Sox2 from the dental epithelium partially rescued the supernumerary tooth phenotype in K14CreERT2; Fam20Bfl/fl; Fgfr2bfl/fl mice. We introduced Sox2CreER to activate both Sox2 and Fam20B in the dental epithelium. The double knockout (K14CreERT2; Fam20Bfl/fl; Sox2fl/fl) mice showed reduced number and size of supernumerary teeth (black arrows) compared with Fam20B single knockout (K14CreERT2; Fam20Bfl/fl; Sox2+/+) mice. The native teeth lacking Sox2 allele (K14CreERT2; Fam20Bfl/fl; Sox2+/+) displayed a smaller size than normal (white arrows).

Additional file 4: Figure S4. Fam20B-catalyzed GAGs regulate the homeostasis of Sox2+ cells in a non-autonomous manner. A To validate the efficiency of the Sox2CreER, we crossed the CreER line with Rosa26-DT-Tomato indicator mice and induced the Cre expression with single IP. injection of tamoxifen at E11.5. The embryos were collected at E12.5 and subjected to cys ysosagen for fluorescence assay. On the same cryosections, Sox2-expressing cells were labeled by immunofluorescence using anti-Sox2 antibody and EGFP-conjugated secondary antibody. The Cre activity indicated by Tomato fluorescence (red) was strongly present in the dental epithelium (arrow) and oral epithelium, as well as nasal mucosa and palatal epithelium. The antibody-labeled Sox2-expressing cells (green) mostly overlapped with CreER active cells (red) and showed yellow on the merged channel. CreER-active cells showed overall broader range than the antibody-labeled Sox2+ cells in the dental epithelium (especially in the distal side) and nasal mucosa, indicating that the efficiency of Sox2CreER was strong enough for deleting floxed alleles from the Sox2-expressing cells. n, nose; p, palate; m, mandible. B To determine the regulatory manner of GAGs on Sox2+/+ cell homeostasis, we inactivated Fam20B from Sox2+/+ lineage using Sox2CreER. Tamoxifen was administered to mice by ip injection at E11.5 and E12.0. Sox2CreER; Fam20Bfl/fl mice did not recapitulate the replacement tooth phenotype, suggesting that GAGs regulate the homeostasis of Sox2+/+ cells in a non-autonomous manner.

Additional file 5: Figure S5. WNT signaling was not changed in the Fam20B-deficient incisors at the early stage of tooth development. A, B Immunohistochemistry staining of LEF1 on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5. D→m indicates the orientation of distal and mesial sides. C, D Whole-mount staining of β-Gal on E12.5 mandibles showed no differences between the Fam20B-deficient and control incisors at E12.5. E, F Immunohistochemistry staining of β-Catenin on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5. In situ hybridization staining of Msx-1 on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5 and E13.5. Yellow dotted lines in E-H indicate the boarder line between the dental epithelium and the dental mesenchyme. Scale bars, 100 μm.

Additional file 6: Figure S6. BMP signaling was not changed in the Fam20B-deficient incisors at the early stage of tooth development. A-D In situ hybridization staining of Msx-1 on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5 and E13.5. Yellow dotted lines in E-H indicate the boarder line between the dental epithelium and the dental mesenchyme. E, F Whole-mount staining of BRE-LacZ indicator on E12.5 mandibles showed no differences between the incisors of Fam20B-mutants and controls. The yellow dotted lines plotted the areas of LacZ positive staining in the lower incisors. G, H Whole-mount (ISH staining of Bmp4 on E12.5 mandibles showed no differences between the incisors of Fam20B-deficient and control incisors at E12.5 and E13.5. Scale bars, 100 μm.

Additional file 7: Figure S7. The transcriptional expression of Fgf10, Fgfr2b, and Fgf9 was not changed in the Fam20B-deficient incisors at the early stage of tooth development. A-F RNAscope staining of Fgf10, Fgfr2b, and Fgf9 on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5. The dotted lines indicate the boarder line between the dental epithelium and the dental mesenchyme. G, H Whole-mount (ISH staining of Msx-1 on the coronal sections of lower incisors showed no differences between the Fam20B-deficient and control incisors at E12.5 and E13.5. Scale bars, 100 μm.

Additional file 8: Figure S8. Inhibition of Fgf2b in the Fam20B-deficient dental epithelium reduced the expanded expression scope of Shh back to the normal size. Scale bars, 250 μm.

Additional file 9: Figures S9A-S9B. Fig. S9A-[GAGs did not show synergistic effects on FGF10-FGFR2b signaling]. Fig. S9B-[GAGs did not show inhibitory effects on FGF10-FGFR2b signaling]. HS and CSA did not show significant synergistic or inhibitory effects on FGF10-FGFR2b signaling in BaF3 cells. A BaF3 cells expressing FGF2b were cultured in RPMI 1640 media supplemented to the culture media did not show any inhibitory effects on FGFR2b signaling (P > 0.05). B BaF3-FGF2bRb cells were cultured in RPMI 1640 media supplemented with 1000 pM FGF10 and 0.5 ng/ml heparin to boost the base-line of FGFR2b signaling activity (indicated by cell viability OD value). GAGs (i.e., HS/HS2S/HS6S/CSA/heparin) supplemented to the culture media did not show any inhibitory effects on FGFR2b signaling (P > 0.05). Two-way ANOVA was used to evaluate the differences among groups.

Additional file 10: Figure S10. Live image of BaF3 cells in the hydrogel. After 48 h of culture, BaF3-FGFR2b cells were stained with Live/Dead Staining. Calcein AM in Green represents live cells. DAPI was used for counter staining. Live images of the cells in hydrogel were acquired by confocal microscopy and reconstructed with Imaris 9.0. The number of live cells in each sample was counted from 3D hydrogel chips reconstructed from 6 randomly selected areas for statistical analysis.

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Authors’ contributions
The hypotheses and experimental design for this research was developed by J.W., R.L., D.O., and X.W. Experiments and data analysis were undertaken by J.W., Y.T., L.H., C.L., T.S., L.L., Y.Y., B.L., and H.Z. J.W., L.H., R.D., S.M., R.K., J.F., D.O., OK, F.Z., R.L., and X.W. were involved in figure production and drafting the manuscript. All authors read and approved the final manuscript.
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Availability of data and materials
All data generated during this study are included in this published article and its additional files. Raw data and materials will be available at request.

Ethics approval and consent to participate
All experimental procedures were conducted with the approval of Texas A&M University Animal Care and Ethics Committee and conform to the National Health and Medical Research guidelines for animal handling.

Competing interests
The authors declare that they have no competing interests.

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References
1. Vainio S, Lin Y. Organogenesis: coordinating early kidney development: lessons from gene targeting. Nat Rev Genet. 2002;3(7):533.
2. Balic A, Thesleff I. Tissue interactions regulating tooth development and renewal. Curr Top Dev Biol. 2015;117:157–186.
3. Thesleff I, Barach HJ, Foord JM, Vainio S, Pratt RM, Martin GR. Changes in the distribution of type IV collagen, laminin, proteoglycan, and fibronectin during mouse tooth development. Dev Biol. 1981;81(1):182–92.
4. Thesleff I, Hurmeirina K. Tissue interactions in tooth development. Differentiation. 1981;18(1–3):75–88.
5. Thesleff I, Vahhatoki A, Partanen AM. Regulation of organogenesis. Common molecular mechanisms regulating the development of teeth and other organs. Int J Dev Biol. 1995;39(3–5):35–50.
6. Bai XM, Van der Schueren B, Cassaman JJ, Van den Berge H, David G. Differential expression of multiple cell-surface heparan sulfate proteoglycans during embryonic tooth development. J Histochem Cytochem. 1994;42(8):1043–54.
7. Wu J, Li H, Han L, Sun T, Tian Y, Wang X. The spatiotemporal expression pattern of Syndecans in murine embryonic teeth. Gene Expr Patterns 2020;36:119109. https://doi.org/10.1016/j.gep.2020.119109.
8. Mythreye K, Blake GC. Proteoglycan signaling co-receptors: roles in cell adhesion, migration and invasion. Cell Signal. 2009;21(11):1548–58.
9. Koke T, Irimikawa T, Tamura J, Kitagawa H. FAM20A is a kinase that phosphorylates xylose in the glycosaminoglycan–protein linkage region. Biochem J. 2009;421(2):157–62.
10. Wen J, Xiao J, Rahbar M, Choudhury BP, Cui J, Taylor GS, Esko JD, Dixon JE. Xylose phosphorylation functions as a molecular switch to regulate proteoglycan biosynthesis. P Natl Acad Sci USA. 2014;111(48):15723–8.
11. Vogel P, Hansen GM, Read RW, Vance RB, Thiel M, Liu J, Wronske TJ, Smith DD, Jeter-Jones S, Brommage R. Aromelogenesis imperfecta and other bimolecular defects in Fam20a and Fam20c null mice. Vet Pathol. 2012;49(6):998–1017.
12. Tian Y, Ma P, Liu C, Yang X, Crawford DM, Yan W, Bai D, Qin C, Wang X. Inactivation of Fam20B in the dental epithelium of mice leads to supernumerary incisors. Eur J Oral Sci. 2015;123(6):396–402.
13. Thesleff I, Jalkanen M, Vainio S, Benfield M. Cell surface proteoglycan expression correlates with epithelial-mesenchymal interaction during tooth morphogenesis. Dev Biol. 1988;129(2):565–72.
14. Vainio S, Jalkanen M, Thesleff I. Syndecan and tenasin expression correlates with mesenchymal–epithelial interaction during tooth morphogenesis. J Cell Biol. 1989;108(5):1945–53.
15. Salmivirta M, Elenius K, Vainio S, Hofer U, Chihurst-Rhimsson R, Thesleff I, Jalkanen M. Syndecan from embryonic tooth mesenchyme binds tenasin. J Biol Chem. 1991;266(12):7733–9.
16. Vainio S, Jalkanen M, Vahhatoki A, Sahlberg C, Mali M, Benfield M, Thesleff I. Expression of syndecan gene is induced early, is transient, and correlates with changes in mesenchymal cell proliferation during tooth organogenesis. Dev Biol. 1991;147(2):322–33.
17. Mizumoto S, Yamada S, Sugahara K. Human genetic disorders and knockout mice deficient in glycosaminoglycan. Biomed Res Int. 2014;2014:495764.
18. Wang XP, Fan J. Molecular genetics of supernumerary tooth formation. Curr Opin Genet Dev. 2009;19(5):504–10.
19. Liu X, Li N, Zhang H, Liu J, Zhou N, Ran C, Chen X, Lu Y, Wang X, Qin C, Xiao J. Inactivation of Fam20B in the neural crest-derived mesenchyme of mouse causes multiple craniofacial defects. Eur J Oral Sci. 2018;126(5):433–6.
20. Juuri E, Saito K, Ahtiainen L, Seidal K, Tummers M, Helloedinger K, Klein OD, Thesleff I, Michon F. Sox2+ stem cells contribute to all epithelial lineages of the tooth via Sfrp5+ progenitors. Dev Cell. 2012;23(2):317–28.
21. Juuri E, Jussila M, Seidal K, Holmes S, Wu P, Richman J, Heikinheimo K, Chuong CM, Arnold K, Helloedinger K, Klein O. Sox2 marks epithelial competence to generate teeth in mammals and reptiles. Development. 2013;140(7):1424–32.
22. Thesleff I, Nieminen P. Tooth morphogenesis and cell differentiation.Curr Opin Cell Biol. 1996;8(8):844–50.
23. Jussila M, Thesleff I. Signaling networks regulating tooth organogenesis and regeneration, and the specification of dental mesenchyme and epithelial cell lineages. CSH Persp Biol. 2012;4(4):a008425.
24. Lan Y, Ja S, Jiang R. Molecular patterning of the mammalian dentition. Semin Cell Dev Biol. 2014;25:61–70.
25. Du W, Du W, Yu H. The role of fibroblast growth factors in tooth development and incisor renewal. Stem Cell Int. 2018;2018:7549160.
26. Klein OD, Mounowada G, Petekova R, Kangas A, Yu BD, Lesot H, Petekova M, Jernvall J, Martin GR. Sproutsy genes control diastema tooth development via bidirectional antagonism of epidermal-mesenchymal FGF signaling. Dev Cell. 2006;11(2):181–90.
27. Jackman WR, Davies SH, Lyons DB, Stauder CK, Denton-Schneider BR, Jowdyr A, Aglier SR, Vogel SA, Stock DW. Manipulation of Fgf and Bmp signaling in teleost fishes suggests potential pathways for the evolutionary origin of multicusp teeth. Evol Dev. 2013;15(2):107–18.
28. Hokuto I, Perl AK, Whitsett JA. Prenatal, but not postnatal, inhibition of fibroblast growth factor receptor signaling causes embryophya. J Biol Chem. 2002;278(1):415–21.
29. Li G, Li L, Tian F, Zhang L, Xue C, Linhardt RJ. Glycosaminoglycanomics of cultured cells using a rapid and sensitive LC-MS/MS approach. ACS Chem Biol. 2015;10(5):1303–10.
30. Xu D, Esko JD. Demystifying heparan sulfate–protein interactions. Annu Rev Biochem. 2014;83:129–57.
31. Zhang F, Zheng L, Cheng S, Peng Y, Fu L, Zhang X, Linhardt RJ. Comparison of the interactions of different growth factors and glycosaminoglycans. Molecules. 2019;24(18):3360.
32. Omitz DM, Xu J, Colvin JS, McEwen DG, MacArthur CA, Coulier F, Gao G, Goldfarb M. Receptor specificity of the fibroblast growth factor family. J Biol Chem. 1996;271(25):15292–7.
33. Omitz DM, Itoh N. The fibroblast growth factor signaling pathway. Wiley Interdiscip Rev Dev Biol. 2015;4(3):215–66.
34. Chu EY, Hens J, Andl T, Kairo A, Yamaguchi TP, Briskin C, Glick A, Wysockimski JJ, Millar SE. Canonical WNT signaling promotes mammary gland development and is essential for initiation of mammary gland morphogenesis. Development. 2004;131(19):4819–29.
36. Munne PM, Tummers M, Järvinen E, Thesleff I, Jernvall J. Tinkering with the inductive mesenchyme. Sostdc1 uncovers the role of dental mesenchyme in limiting tooth induction. Development. 2009;136(3):393–402.

37. Ayn H, Sanderson BW, Klein OD, Krumlauf R. Inhibition of Wnt signaling by Gsk3 and negative feedback from Shh controls tooth number and patterning. Development. 2010;137(19):3221–31.

38. Järvinen E, Birchmeer W, Taketo MM, Jernvall J, Thesleff I. Continuous tooth generation in mouse is induced by activated epithelial Wnt/β-catenin signaling. Proc Natl Acad Sci U S A. 2006;103(49):18627–32.

39. Wang XP, O’Connell DJ, Lund JJ, Saadi I, Kuraguchi H, Turbe-Doan A, Michon F. Plasticity within the niche ensures the maintenance of a Sox2+ stem cell population in the mouse incisor. Development. 2018;145(1):4115–26.

40. Harada H, Mitsuyasu T, Toyono T, Toyoshima K. Epithelial stem cells in teeth. Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

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37. Ayn H, Sanderson BW, Klein OD, Krumlauf R. Inhibition of Wnt signaling by Gsk3 and negative feedback from Shh controls tooth number and patterning. Development. 2010;137(19):3221–31.

38. Järvinen E, Birchmeer W, Taketo MM, Jernvall J, Thesleff I. Continuous tooth generation in mouse is induced by activated epithelial Wnt/β-catenin signaling. Proc Natl Acad Sci U S A. 2006;103(49):18627–32.

39. Wang XP, O’Connell DJ, Lund JJ, Saadi I, Kuraguchi H, Turbe-Doan A, Michon F. Plasticity within the niche ensures the maintenance of a Sox2+ stem cell population in the mouse incisor. Development. 2018;145(1):4115–26.

40. Harada H, Mitsuyasu T, Toyono T, Toyoshima K. Epithelial stem cells in teeth. Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

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