MMP20 Cleaves E-Cadherin and Influences Ameloblast Development

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Key Words

Matrix metalloproteinases · Adhesion · Cell junctions

Abstract

Dental enamel development occurs in stages as observed by the changing morphology of the ameloblasts that are responsible for enamel formation. During the secretory stage of development, proteins including MMP20 are secreted into the enamel matrix. MMP20 is required for proper enamel formation as mutation of the *Mmp20* gene causes autosomal recessive amelogenesis imperfecta. Here, we examined in detail the morphology of the *Mmp20*-null ameloblast cell layer. Intriguingly, we found that the *Mmp20*-null mouse secretory stage ameloblasts retract their Tomes' processes as if preparing to enter the maturation stage but later reextend their Tomes' processes as if resuming the secretory stage. We also demonstrated that MMP20 cleaves epithelial cadherin, i.e. E-cadherin. Cadherins are transmembrane proteins with extracellular domains that provide adhesive contacts between neighboring cells. Their intracellular domains bind to the cell cytoskeleton through catenins, including β -catenin. When specific MMPs cleave the cadherin extracellular domain, β -catenin is released and may locate to the cell nucleus as a transcription factor. Therefore, MMP20 may influence ameloblast developmental progression through hydrolysis of cadherin extracellular domains with associated release of transcription factor(s).

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Introduction

The two predominant stages of enamel development are the secretory and maturation stages. The secretory stage is when the ameloblasts of the enamel organ develop protruding Tomes' processes at their apical ends and begin moving laterally in rows to form the rod and interrod enamel. These Tomes' processes are retracted as the ameloblasts transition to the maturation stage of development. Significantly, adherens junctions (AJ) were identified along the possible sliding interface between adjacent rows of secretory stage ameloblasts [Nishikawa et al., 1990].

AJ are composed of several classes of proteins including cadherins and catenins. Cadherins are transmembrane proteins with extracellular domains that provide important adhesive contacts between neighboring cells. The extracellular domains connect through homotypic trans-pairing between cadherins on adjacent cells, and the intracellular domains of cadherins are linked to the actin cytoskeleton by the catenins [reviewed in Hartsock and Nelson,

Abbreviations used in this paper

AJ	adherens junction
JMD	juxtamembrane domain
p120	p120-catenin

2008]. Ameloblasts express E-, N-, and P-cadherins as well as β -catenin and p120-catenin (p120), during dental enamel development [Palacios et al., 1995; Fausser et al., 1998; Obara et al., 1998; Terling et al., 1998; Sorkin et al., 2000; Heymann et al., 2002; Bartlett et al., 2010]. A major pathway for signal transduction by AJs involves the regulation of β -catenin, which can act as either a structural protein at cell-cell junctions or a transcription factor in the cell nucleus [reviewed in Munshi and Stack, 2006]. When MMPs cleave a cadherin's extracellular domain, β -catenin is removed from its position near the cell membrane and in most cases will translocate to the cell nucleus thereby promoting cell migration, cell invasion, and/or cell proliferation [Lochter et al., 1997; McGuire et al., 2003; Sanceau et al., 2003; Cowden Dahl et al., 2008; Lynch et al., 2010].

MMP20 is required for healthy dental enamel development. People and mice with homozygous *MMP20* mutations have soft discolored enamel that may be hypoplastic and that easily abrades from the dentin surface [Caterina et al., 2002; Bartlett et al., 2004; Kim et al., 2005; Ozdemir et al., 2005; Bartlett et al., 2006; Papagerakis et al., 2008; Lee et al., 2010]. Herein we demonstrate that *Mmp20*-null mouse ameloblasts improperly progress through the secretory stage of development and that MMP20 cleaves the E-cadherin extracellular domain. These results are consistent with MMP20 playing a role in ameloblast development via cadherin hydrolysis and the subsequent release of intracellular transcription factors such as β -catenin.

Materials and Methods

All experimental procedures involving the use of animals were reviewed and approved by the Forsyth Institute and the University of Michigan Institutional Animal Care and Use Committees.

Morphological Analyses

Adult C57BL/6 mice were anaesthetized with chloral hydrate and fixed by intravascular perfusion with 1% glutaraldehyde buffered with 0.08 M sodium cacodylate containing 0.05% CaCl_2 , pH 7.2. The hemimandibles were removed and decalcified at 4°C in 4.13% EDTA. The samples were washed and postfixed in osmium tetroxide reduced with potassium ferrocyanide, dehydrated in graded alcohols, and embedded in LR White acrylic resin (London Resin, Berkshire, UK) as previously described [Zalzal et al., 2008]. Semithin sections of incisor segments were cut with glass knives and stained with toluidine blue for examination by light microscopy (400 $\times$ magnification).

Preparation of Enamel Extracts

Tooth germs of permanent molars were surgically extracted from the maxillae and mandibles of 6-month-old pigs at the

Michigan State University Meat Laboratory (East Lansing, Mich., USA). The soft, cheese-like enamel was separated from the crown with a spatula. The neutral extract (supernatant) was obtained by homogenization of the enamel samples in Sørensen buffer (made by mixing Na_2HPO_4 and KH_2PO_4 to achieve a final phosphate concentration of 50 mM and a pH of 7.4), followed by centrifugation. The pellet was resuspended in carbonate buffer (made by mixing NaHCO_3 and Na_2CO_3 to a final concentration of 50 mM and a pH of 10.8), homogenized, and then centrifuged. The supernatant was designated as alkaline extract.

Purification of Porcine MMP20

The soft enamel-alkaline extract was fractionated by ion exchange chromatography in a Q-SepharoseTM Fast Flow column (1.6 $\times$ 20 cm; GE Healthcare Biosciences, Little Chalfont, UK) equilibrated with 50 mM Tris-HCl/6 M urea (pH 7.4) and eluted with a linear NaCl gradient (0–0.5 M). MMP20 eluted in the fourth peak was desalted and concentrated by ultrafiltration (YM-3; Millipore Corp.) and lyophilized. The lyophilized sample was dissolved with 20 mM sodium citrate/6 M urea (pH 7.4) and applied to a Heparin Sepharose 6 Fast Flow column (1.6 $\times$ 20 cm, GE Healthcare) equilibrated with the same buffer and eluted with a step NaCl gradient (0.02, 0.05, and 0.5 M at a flow rate of 0.2 ml/min). MMP20 eluted in the second peak, which was buffer exchanged with 50 mM Tris-HCl (pH 7.4) with a YM-3 membrane.

Digestion of the E-Cadherin Extracellular Domain by MMP20

The human E-cadherin extracellular domain (100 μg) linked to the Fc IgG fragment (648-EC; R&D Systems) was incubated in 45- μl reactions with approximately 10 μg of MMP-20 for 20 h at 37°C in 50 mM Tris-HCl/10 mM CaCl_2 /150 mM NaCl (pH 7.4) and analyzed by SDS-PAGE and immunoblotting as described previously [Nagano et al., 2009] with a polyclonal antibody against E-cadherin (AF648, R&D Systems) or the Fc fragment (I-9135, Sigma).

Results

Mmp20-Null Mouse Ameloblasts Can Initiate, Terminate, and Reinitiate the Secretory Stage of Enamel Development

Secretory stage ameloblasts have a conical shaped apical protrusion termed the 'Tomes' process' that lengthens the enamel rods. Once the rods are full length and the enamel has reached its final thickness, the Tomes' process retracts and the ameloblasts progress to the maturation stage of development. Surprisingly, we found that ameloblasts in *Mmp20*-null mice will retract their Tomes' process during development and then later reform Tomes' processes once again. We observed this phenomenon in all 4 incisors examined which were each from 4 different *Mmp20*-null mice. This was not observed in incisors from *Mmp20* heterozygous (1 mouse) or wild-type mice (2 mice). Figure 1a–d shows consecutive sections of an

Fig. 1. *Mmp20*-null mouse ameloblasts can initiate, terminate, and reinitiate the secretory stage of enamel development. **a–d** Successive sections of an *Mmp20*-null mouse incisor. White arrows point to prominent Tomes' processes that disappear in **b** and **c**. **d** The arrowhead indicates the return of the Tomes' processes, with arrows indicating prominent processes. It appears that signals necessary for the ameloblasts to progress through the secretory stage are absent. Am = Ameloblast; d = dentin; e = enamel.

Mmp20-null secretory stage incisor. White arrows point to prominent Tomes' processes and an arrowhead in figure 1d denotes the beginning of the reextension of Tomes' processes. Note that the dentin progressively thickens while the enamel does not.

MMP20 Cleaves E-Cadherin

β -catenin is part of the structure that links cadherins to the cytoskeleton. When a cadherin is cleaved by an MMP, it can release β -catenin so that it may translocate to the cell nucleus as a transcription factor. We therefore asked if MMP20 can cleave the epithelial cadherin, i.e. E-cadherin, expressed by ameloblasts during enamel formation. Purified porcine MMP20 was incubated with the recombinant human E-cadherin extracellular domain linked to the IgG Fc fragment. After 20 h of incubation, immunoblots specific for E-cadherin or the Fc fragment were performed. The results definitively demonstrated that MMP20 cleaves the extracellular domain of E-cadherin (fig. 2). Thus, the possibility exists that MMP20 releases intracellular signaling molecules through hydrolysis of the ameloblast E-cadherin extracellular domain.

Discussion

We provide evidence suggesting that MMP20 cleaves the extracellular domain of cadherins which may release transcription factors such as β -catenin that are important for enamel development. Faulty cell signaling may be the reason *Mmp20*-null ameloblasts abnormally extend, retract, and later reextend their Tomes' processes during enamel development.

Previously, we demonstrated that conditional deletion of the cadherin stabilizing molecule, p120, from epithelial tissues had a profound effect on mouse dental enamel development. p120 binds to an intracellular cadherin domain near the cell membrane termed the 'juxtamembrane domain' (JMD). Mutation of the E-cadherin JMD demonstrated that p120-E-cadherin interaction was required for high-level cell adhesiveness [Thoreson et al., 2000; Ferber et al., 2002], and it was also shown that the binding of p120 to the JMD prevents cadherins from becoming internalized and degraded [Davis et al., 2003; Xiao et al., 2003; Maeda et al., 2006]. When p120 was deleted from epithelial tissues, the general shape of the teeth was normal. However, the resulting dental enamel had no rod pattern

Fig. 2. Immunoblot showing that MMP20 cleaves the E-cadherin extracellular domain. The recombinant human E-cadherin/Fc chimera that links the E-cadherin extracellular domain to an Fc fragment was incubated with MMP20 that was purified from porcine enamel. Incubations were for 0 or 20 h followed by immu-

noblotting with primary antisera specific for E-cadherin or the linked Fc fragment. The secondary antiserum was from rabbit (for IgG-Fc Ab) or goat (for E-Cadherin Ab). Also shown is an SDS-PAGE gel stained for protein and 2 negative control immunoblots that were treated with only the secondary antisera.

and had a very rough surface composed of mineralized spherical structures, and this enamel easily abraded from the tooth surface [Bartlett et al., 2010]. Therefore, stabilization of cadherins to the cell surface is essential for dental enamel formation and this suggests that the hydrolysis of cadherin contacts necessary for ameloblast migration and signaling may also be of importance in enamel formation.

Here we demonstrate that *Mmp20*-null ameloblasts have abnormal extension and retraction of their Tomes' processes and therefore seem unable to maintain normal

linear continuity of appositional growth during the secretory stage of amelogenesis. We also show that MMP20 can cleave an important epithelial cadherin and that this hydrolysis may have important cell signaling functions.

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