Matrix Metalloproteinase-20 Over-Expression Is Detrimental to Enamel Development: A *Mus musculus* Model

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Abstract

**Background:** Matrix metalloproteinase-20 (Mmp20) ablated mice have enamel that is thin and soft with an abnormal rod pattern that abrades from the underlying dentin. We asked if introduction of transgenes expressing Mmp20 would revert this Mmp20 null phenotype back to normal. Unexpectedly, for transgenes expressing medium or high levels of Mmp20, we found opposite enamel phenotypes depending on the genetic background (Mmp20⁺⁻⁻ or Mmp20⁺⁺⁺) in which the transgenes were expressed.

**Methodology/Principal Findings:** Amelx-promoter-Mmp20 transgenic founder mouse lines were assessed for transgene expression and those expressing low, medium or high levels of Mmp20 were selected for breeding into the Mmp20 null background. Regardless of expression level, each transgene brought the null enamel back to full thickness. However, the high and medium expressing Mmp20 transgenes in the Mmp20 null background had significantly harder more mineralized enamel than did the low transgene expresser. Strikingly, when the high and medium expressing Mmp20 transgenes were present in the wild-type background, the enamel was significantly less well mineralized than normal. Protein gel analysis of enamel matrix proteins from the high and medium expressing transgenes present in the wild-type background demonstrated that greater than normal amounts of cleavage products and smaller quantities of higher molecular weight proteins were present within their enamel matrices.

**Conclusions/Significance:** Mmp20 expression levels must be within a specific range for normal enamel development to occur. Creation of a normally thick enamel layer may occur over a wider range of Mmp20 expression levels, but acquisition of normal enamel hardness has a narrower range. Since over-expression of Mmp20 results in decreased enamel hardness, this suggests that a balance exists between cleaved and full-length enamel matrix proteins that are essential for formation of a properly hardened enamel layer. It also suggests that few feedback controls are present in the enamel matrix to prevent excessive Mmp20 activity.

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Introduction

Dental enamel is the hardest tissue of the body, but it does not start that way. Enamel development (amelogenesis) can be defined as consisting of three stages: the secretory, transition, and maturation stages [1]. During the secretory stage the ameloblasts adjacent to the forming enamel elongate and secrete large quantities of protein into the enamel matrix. Approximately 90% of this protein is amelogenin [2]. Amelogenin has only one post-translational modification whereby Ser16 is phosphorylated [3,4], but its transcripts undergo extensive alternative splicing [5,6,7] to generate as many as 16 X-chromosomal murine amelogenin mRNAs [8,9,10]. The secretory stage is when thin crystallite ribbons begin growing in length [11,12,13] and they stop elongating once ameloblasts have defined the full thickness of the enamel layer. Secretory stage enamel is very soft and has a cheese-like consistency. Once the ameloblasts progress to the transition stage, they shorten and greatly reduce protein secretion. During the subsequent maturation stage, the short columnar ameloblasts reabsorb the proteins they had previously secreted. This is when the enamel ribbons grow to their greatest amount in width and thickness [1]. Enamel consists of mineralized rods that are sometimes entwined (gnarled) in human molars [14] or may form groups of rods that pass across each other (decussating) as is
observed in rodent incisors [15]. Each rod is formed by one ameloblast and contains approximately 10,000–40,000 crystalline ribbons [11]. Therefore, defective thin (hypoplastic) enamel suggests a developmental deficiency during the secretory stage when the crystalline ribbons are lengthening to establish the full thickness of the enamel layer. Erupted teeth with enamel that has reached full thickness, but is soft and not well mineralized (hypomatured) suggests a developmental deficiency during the maturation stage when the crystalline ribbons grow in width and thickness and interlock. Fundamental developmental deficiencies may also affect both stages of enamel development (hypocalcified) and these cases typically have the most severely dysplastic enamel phenotypes [16].

Matrix metalloproteinase-20 (MMP20; enamelysin) is a tooth specific MMP [17] that is secreted into the enamel matrix during the secretory and transition stages of enamel development [18-20]. DNA encoding the Mmp20 catalytic domain was previously deleted from the mouse genome to identify the role of MMP20 in enamel development [21]. Mmp20 null mice did not process amelogenin properly, had altered enamel rod patterns and had hypoplastic enamel that broke away from dentin. A subsequent study showed that relative to wild-type controls, Mmp20 null mice had an overall enamel mineral content that was reduced by 50% and an enamel hardness that was decreased by 37% [22]. Although Mmp20 is only expressed from the secretory through transition stages, it was concluded that protein processing by MMP20 is necessary so that the ameloblasts can efficiently remove proteins during the maturation stage. Kallikrein-related peptidase-4 (KLK4) is a serine proteinase secreted during the maturation stage to further cleave enamel matrix proteins prior to their export out of the hardening enamel [23]. However, KLK4 can apparently not compensate for a lack of prior MMP20 activity so the Mmp20 null mouse enamel retains matrix proteins and is therefore softer than normal. Seven different human MMP20 mutations are known to cause autosomal recessive non-syndromic amelogenesis imperfecta [24-30].

Here we sought to determine if introduction of transgenes that express low, medium or high levels of MMP20 in the Mmp20 null background would revert the murine Mmp20 enamel phenotype back to normal. The high and medium expressing transgenics did result in a marked improvement in enamel development over that of the Mmp20 null mice. Unexpectedly, the high or medium expressing transgene caused malformed enamel when present in the wild-type background suggesting that a critical balance exists between cleaved and full-length enamel matrix proteins that are essential for proper hardening of the enamel layer. Beyond a specific threshold, too much MMP20 activity disrupts this balance and high quantities of cleavage products disrupt normal enamel development.

**Materials and Methods**

**Ethics Statement**

All animals used in this study were housed in an Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC) accredited facilities (animal welfare assurance number: A3051-01) and were treated humanely based on a protocol 11-021 approved by the Institutional Animal Care and Use Committee (IACUC) at The Forsyth Institute. Experimental protocols were designed along institutional and National Institutes of Health guidelines for the humane use of animals.

**Construction of the Mmp20 Transgene**

C57BL/6 mice were sacrificed on post-natal day 5, and first molars were removed. Total RNA was extracted with Trizol (Invitrogen, Carlsbad, CA, USA) and converted into cDNA with SuperScript II reverse transcriptase (Invitrogen). Mmp20 cDNA was amplified for ligation into the pCR2.1-TOPO vector (Invitrogen). Primers used for this amplification were: forward, 5'-TTG CGC GCC AGA GGA GTG GTG GCC ACC TCG C3' and reverse, 5'-AGC GAT GAG CAC TGC AGG TGC TAC CAG GAA GTA GG-3'. These primers amplified 2278 bp of the Mmp20 cDNA that included the start codon and 821 bp of 3' non-coding region. The region of the forward primer contained an Ascl restriction enzyme site and the beginning of the reverse primer contained a SgfI restriction enzyme site to enable the PCR product's directional ligation into the vector. This vector was previously constructed to contain 4639 bp of mouse AmelX promoter region including the first exon (non-coding) and first intron. Immediately 3' to the Mmp20 cDNA insertion site was 1127 bp of AmelX 3' non-coding region that included amelogenin polyadenylation signals [31].

**Generation of Mmp20 Transgenic Mice and Breeding with Mmp20+/− Mice**

The Amelx promoter-Mmp20 cDNA with the 3'-Amelx non-coding region was excised from the vector by restriction digestion with NotI-SfiI, purified with a Qiagen gel extraction kit (Qiagen, Germantown, MD, USA) and microinjected into fertilized oocytes for surgical transfer to recipients. Germline transmission was determined by PCR analyses of genomic DNA obtained from tail biopsies. PCR primers used to identify the presence of the transgene were: 5'-GAA AAT GGT TGG CAG CAT CA-3', and 5'-CIT GCC ACC ATT TGG CCA GCC-3'. For mouse genotyping, genomic DNA was isolated from tails and the Extract-Amp Kit (Sigma) was used for PCR reactions. PCR primer sequences for determining Mmp20 genotypes were: 5'-CTG CCC CAG ACT TTT GAG TT-3', and 5'-GCT TTT CAT GGC CAG AAT GCT CT-3', to detect the ablated allele and 5'- AAG TAG ACT GAA GTC AGG AGA GCC-3', and 5'-CTG TAG TGG TGA CCC TAG TGA TCT T-3', to detect the wild-type allele. Offspring carrying the Mmp20 transgene (Tg) were mated with Mmp20 ablated mice [22] to produce Tg positive Mmp20+/− and these mice were bred to generate both Tg positive Mmp20+/− and Tg positive Mmp20−/− mice. Therefore, each transgenic founder mouse line had a similar genetic background.

**Quantitative Real-time PCR (qPCR)**

RNA extracted from incisor enamel organ of adult mice or first molar enamel organs of 5 day-old mice were used to determine relative expression levels of Mmp20 as a function of a stably expressed internal reference control gene (18S rRNA) as previously described [32,33]. Primers were: Mmp20 forward 5'-GCC TCT TCG CAG GTG AACCCA-3', Mmp20 reverse (5'-AGC CAT GCA GGG CGC CCT GT-3'); 18S forward (5'-GTA ACC CTA TGA ACC CCA TT-3'), 18S reverse 5'-CCA TCG AAT CCG TAG TAG CG-3'. Reactions were performed on a Roche LightCycler 480 using the following program: 3 min at 95°C for initial denaturation, and 95°C 15 sec, 58°C 15 sec, 72°C 15 sec for 40 cycles, followed by a melting curve. Each time point was obtained by triplicate qPCR analysis, and all expression levels are presented as relative ratios to the wild-type incisor or day 5 first molar data.
Protein Gels, Immunoblotting and Zymography

Mandibular incisors were extracted from adult mice and first molars were removed from 5-day-old mouse pups. The mineral was rapidly dissolved by submerging the teeth in 2 mL of 0.17 M HCl/0.98% formic acid for 2 h at 4°C. Undissolved material was removed by centrifugation at 3,500 g for 5 min at 4°C. The samples were then dialyzed against water overnight and lyophilized for immunoblot or zymography analysis. The lyophilized proteins were weighed and eluted into sample buffer. For immunoblots, an equal amount of protein was loaded in each lane and was run on SDS-PAGE and transferred to a membrane. Immunoblots were performed with antisemur specific for the N-terminus of active MMP20 (Abcam, Cambridge, MA, USA; Ab39037, 1:2500 in TBST) or with antisemur for amelogenin (rabbit anti-recombinant antibody [34], 1:3000 in TBST). Casein zymogram gels were purchased from Invitrogen and electrophoresis was performed at a constant voltage of 120 V for 1 h. After electrophoresis, gels were washed twice for 30 min in 2.5% TritonX-100. Gels were incubated for 2 days at 37°C in 30 mM Tris-HCl buffer (pH 7.2) containing 10 mM CaCl2 and were then stained with Coomassie Brilliant Blue (CBB) R-250 solution (0.1% CBB R-250, 10% acetic acid, and 50% methanol) for 15 min and destained with 30% methanol and 10% acetic acid until clear bands of substrate lysis were observed. Silver staining was performed according to the manufacturer’s protocol (Amersham Biosciences, NJ, USA).

Photographs of Mouse Teeth

Soft tissues were removed from right half-mandibles of adult mice. Photographs of the incisors and lingual views of molars were taken using a Nikon SMZ745T microscope and Leica DFC400 digital camera at 30x magnification.

MicroComputed Tomography (µCT)

Adult mouse incisor enamel was assessed for mineralization levels by µCT. Hemi-mandibles with soft tissues removed were immersed in saline and scanned in a µCT-40 (Scanco Medical, Wayne PA, USA) with the following settings: 70 kV, 114 mA, and 0.01 mm isotropic voxels. Images were processed with ImageJ (http://rsb.info.nih.gov/ij/) on TIFF images that were normalized to have the same background. ImageJ was used to orient the incisors so that enamel layer mineralization could be clearly observed.

Scanning Electron Microscopy (SEM) of Mandibular Incisors

Adult mouse mandibular incisors with soft tissue removed were fixed in 5% glutaraldehyde overnight with rotation, washed in PBS x 3 and were embedded in Epon. Perpendicular cuts through incisors were made 8 mm from their apical ends. Incisors were then re-embedded in Castolite AC (Eager polymers, Chicago IL, USA) with the cutting plane face down and were allowed to harden at room temperature overnight. The surfaces were then ground with silicon carbide (400, 800 & 1200 grit) prior to polishing with a 1 μm diamond polish paste. The surfaces were sputter-coated with gold for conventional SEM or were carbon coated for backscattering SEM. SEM was performed as described previously [31]. Five-level pseudo-color mapping was performed as described previously [35] using ImageJ [http://rsb.info.nih.gov/ij/] on TIFF images that were normalized to have the same mean gray-level intensities for mineralized dentin (mature dentin mineral shows no significant changes with or without MMP20 expression). For pseudo-colorization of images, gray levels 1–57 were assigned as black, 58–97 as white, 98–162 as blue, 163–192 as orange, and 193–255 as red. These settings were saved in a look-up table (LUT) and applied to selected SEM images.

Vickers Microhardness Testing

Erupted portions of mandibular incisors from wild-type, Mmp20−/− and Mmp20 transgenic mice were washed and dehydrated with graded alcohol and acetone. Incisors were embedded sagittally in hard-formulation epoxy embedding medium (EpoFix, EMS, Hatfield, PA, USA). Samples were ground and polished to 0.25 mm with diamond suspensions (EMS). The polished samples were tested for enamel microhardness on an M 400 HI testing machine (Leco, St. Joseph, MI, USA). Testing was performed with a load of 25 g for 5 sec with a Vickers tip. Twenty indentations per sample were performed on at least 4 teeth per group and averaged.

Statistics

Unpaired t-tests were performed using GraphPad Prism version 5.0 (GraphPad Software, San Diego, CA) to analyze the significance of differences in enamel thickness and enamel hardness. T-tests were also performed for analyzing the significance of qPCR results.

Results

Production of Transgenic Mice

A transgene was designed such that 4639 bp of the mouse amelogenin promoter was inserted 5′ to the mouse Mmp20 cDNA and 1127 bp of amelogenin 3′ non-coding region that included amelogenin polyadenylation sites was inserted immediately 3′ to the Mmp20 cDNA. The construct was sequenced in its entirety to confirm its integrity. Three transgenic founder mice were selected based on the ability of their incisor enamel organs to express low (Tg42), intermediate (Tg6) or high (Tg24) levels of Mmp20 transgene transcripts in the Mmp20 null background (Fig. 1A). We also performed qPCR on incisor enamel organs to confirm that the total level of wild-type and transgene Mmp20 transcripts were increased when the transgenes were present in the wild-type background (Fig. 1B). Interestingly, transgene expression in first molar enamel organs from 5 day-old mice that are predominately in the secretory stage of enamel development, displayed expression levels that were different from incisors. In molars, Tg6 transgenic mice had the highest level of expression followed by the Tg24 transgene and the Tg42 transgene was expressed in mice at lower than wild-type endogenous levels (Fig. 1C). Note that minimal Mmp20 null (KO) amplification products may have been generated because the qPCR primers located upstream of the deleted region (exon 5) encoding the Mmp20 catalytic domain.

To confirm that transcript expression paralleled MMP20 protein levels, we performed immunoblots on proteins extracted from molars or incisors and performed zymography on 5 day-old first molar enamel matrix proteins. Consistent with the qPCR results for 5 day-old molars (m), the Tg6 transgenic mice expressed the most MMP20 protein (H) whereas Tg42 transgenic mice expressed the least (L). The wild-type (+/+ ) mice expressed more MMP20 protein than the Mmp20 heterozygous (+/− ) mice while the null (−/− ) mice expressed no MMP20 (Fig. 2A). The zymography results supported the immunoblot results by demonstrating that more active MMP20 was present in enamel from Tg6m (H) transgenic 5 day-old mouse first molars than was present in the Tg24m (M) (medium expression) or Tg42m (L) (lowest expression) transgenic mouse molars (Fig. 2B). Conversely, in extracted incisor (i) enamel the Tg24i (H) transgene was expressed at the highest level followed by the Tg6i (M) and Tg42i
transgenes respectively. When the transgenes were expressed in wild-type mice that express endogenous MMP20, the quantities of MMP20 extracted from incisor enamel from each of the transgenic mice was increased over the levels observed when each transgene was present in the *Mmp20* null background (Fig. 2C). Therefore, the order of transgene expression level in mouse incisors was Tg24i (H) > Tg6i (M) > Tg42i (L) and was Tg6m (H) > Tg24m (M) > Tg42m (L) for molars. The highest levels of total MMP20 expression (endogenous and transgenic) were observed in wild-type mice expressing the Tg24 transgene in incisors or the Tg6 transgene in molars.

**Examination of Transgenic Mouse Teeth**

Mandibular incisors and molars from *Mmp20* transgenic mice were examined under a light microscope (Fig. 3). Each transgene was present in the wild-type or *Mmp20* null background. The wild-type incisor (Tg– *Mmp20*+/+) had the characteristic yellow-brown color and had a sharp tip. The wild-type molars had distinctive cusp tips that were thick with a full thickness enamel layer. In contrast, *Mmp20* null incisors (Tg– *Mmp20*–/–) showed no yellow-brown color, had enamel that peeled away from the underlying dentin and had a blunted tip. *Mmp20* null molars had worn cusp...
tips or had thin cusp tips because the enamel had abraded away. As expected, expression of the transgenes in the Mmp20 null background did recover some (Tg24, Tg42 molars) or nearly all (Tg6, Tg24 incisors) of the otherwise dysplastic enamel phenotype. In the wild-type background, although the teeth from the lowest expressing transgene (Tg42) appeared normal, the teeth from the two highest expressing transgenes (Tg6, Tg24) unexpectedly had blunted incisor tips and had molars with worn cusp tips. The enamel phenotype in the presence of these two transgenes was much worse in the wild-type background when compared to the enamel phenotype when these transgenes were expressed in the Mmp20 null background.

Incisor μCT Analysis
Rodent incisors have enamel on only their labial sides. Presented in the micrographs of Figure 4 are incisors with their labial sides oriented to the left hand side of the incisor. On the labial side of the wild-type incisors (Tg− Mmp20+/+, top right panel) a translucent line (layer) of enamel is present that begins at the apical end (bottom arrow) of the incisor encased in bone (arrowheads) and extends up to the incisal tip (top arrow). This layer of enamel is very thin, spotty or absent along the labial edge of the transgene negative Mmp20 null (Tg− Mmp20−/−, top left panel) mouse incisors. An enamel layer was present on each transgene positive incisor in the Mmp20 null background and in the wild-type background an enamel layer was present on incisors of mice expressing the Tg6i (M) or Tg42i (L) transgenes. However, wild-type mice with the highly expressing Tg24i (H) transgene showed very little enamel on their incisors. These μCT data confirm the light microscope results demonstrating that high levels of MMP20 over-expression in normal mice results in dysplastic enamel.

Backscattered SEM
The more highly mineralized the tissue, the more calcium there is to scatter electrons. Therefore, to characterize the degree of enamel mineralization for each of the transgenoses in the wild-type and Mmp20 null backgrounds, we performed backscattered SEM imaging and used computer software to assign different colors to images that fell within one of five ranges of grey level intensity [35]. In Figure 5, the top panel is the backscattered image and the panel beneath it is the same image that has been pseudo-colorized for assessment of mineralization levels. The most highly mineralized tissues appear red and less mineralized areas are in blue color.

The first column of Figure 5 shows incisor cross sections from Mmp20−/− mice and the second column shows incisor cross sections from Mmp20+/+ mice. For wild-type mice (Tg− Mmp20+/+), the colorized sections demonstrate that enamel is more highly mineralized than dentin. As was previously observed in Mmp20−/− mice (Tg− Mmp20−/−) [35], the outer enamel layer is more mineralized than the inner enamel layer and the characteristic nodules present on Mmp20 null enamel can be observed protruding from the outer enamel layer. The Tg+ Mmp20−/− column shows that Tg6i (M) transgenic mice increased the degree of enamel mineralization to almost normal levels. However, the lateral sides of the enamel layer appeared slightly softer than normal. The Tg24i (H) transgenic mice also reverted the Mmp20 null enamel phenotype almost completely back to normal. Although mineralization near the dentin-enamel junction (DEJ) was slightly less than normal and the enamel surface on the lateral side was not completely smooth. In contrast, Tg42i (L) transgenic mice expressed the least amount of MMP20 and had enamel that was not fully mineralized with a prominent line of

Image 315x362 to 554x730

Figure 3. Assessment of control and Mmp20 transgenic enamel by light microscopy. Wild-type control (Tg− Mmp20+/+) incisors had a sharp incisal tip and the characteristic yellow-brown coloration. The molars contained a fully thick enamel layer and the cusp tips were well defined. In contrast, Mmp20 null (Tg− Mmp20−/−) incisors had a blunted tip and showed little or no enamel and no yellow-brown color. The molars were worn and the remaining cusp tips appeared thin from abrasion and absence of a full thickness enamel layer. For the transgenoses in the Mmp20 null background (Tg+ Mmp20−/−): The Tg6 transgenic enamel had a sharp incisal tip, but the yellow-brown color was mostly missing while the molars appeared fully recovered from the null phenotype. The Tg24 transgenic animals had incisors that appeared sharp and somewhat yellow-brown in color, but the enamel surface was rough and appeared slightly chalky rather than translucent. The molar cusp tips were worn and the enamel layer appeared hypoplastic. The Tg42 transgenic mice had incisors that were blunted with a rough enamel surface that was mildly yellow-brown color with a chalky appearance like the Tg24 transgenic incisors. The molars appeared well formed and fully recovered from the null phenotype. For the transgenoses in the Mmp20 wild-type background (Tg+ Mmp20+/+): The Tg6 transgenic mice had well pigmented but blunted incisors with a chalky-white appearance and rough enamel surface. The Tg24 transgenic enamel appeared almost identical to the Tg6 enamel in the wild-type background. In contrast, Tg42 transgenic incisors had a sharp tip and smooth enamel, but with almost no yellow-brown color. The Tg42 transgenic molars appeared no different than the wild-type molars. The observed Tg6 transgenic enamel was the best example of an overall recovery (both incisors and molars) from the Mmp20 null phenotype.

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The transgenic incisors in the null background (Tg–) recovered some or most of the enamel layer along the labial surface. The presented longitudinally oriented incisors transgenics and controls.

Figure 4. μCT analyses of incisor enamel from Mmp20 transgenics and controls. Enamel on rodent incisors is present only on the labial side (arrows). The presented longitudinally oriented incisors were reconstructed from μCT images. The transgenic incisors (Tg– Mmp20+/−, top right panel) have a bright line of mineralized enamel that extends from the apical region (bottom arrow) to the labial incisal tip (top arrow). This mineralized enamel was mostly missing from the Mmp20 null incisors (Tg– Mmp20−/−, top left panel). The transgenic incisors in the null background (Tg+ Mmp20+/−) recovered some or most of the enamel layer along the labial surface. When the transgenes were present in the wild-type background (Tg+ Mmp20+/+), the enamel layer seemed relatively normal on incisors from mice transgenic for Tg6i (M) or Tg24i (L), but it was severely disrupted on the Tg24i (H) Mmp20+/− incisors. For incisors, Tg24 was the highest, Tg6 the middle and Tg42 the lowest expressing transgene. doi:10.1371/journal.pone.0086774.g004

softer enamel near the DEJ. This incisor also had a rough enamel surface.

The degree of enamel mineralization in wild-type mice expressing various transgenes was striking because it was reversed from what occurred in the Mmp20 null background. The low transgene expression level in Tg24i (L) transgenic mice resulted in enamel that was of normal thickness. In the incisor enamel organ, the Tg24i (H) transgene expresses the highest levels of Mmp20 and this combined with the endogenously expressed Mmp20 resulted in a thinner than normal enamel layer (P<0.001) (Fig. 6B). Therefore, low levels of Mmp20 expression are sufficient to achieve normally thick enamel, but high levels interfere with the developmental processes necessary to achieve a fully thick enamel layer.

Quantification of Enamel Hardness

Incisors were embedded in epoxy resin and were ground and polished in preparation for microhardness testing. Each bar represents hardness measurements for incisors from at least 4 different mice. The Tg– Mmp20+/− and the Tg– Mmp20−/− samples were each pooled separately so these bars represent hardness measurements for incisors from at least 8 different mice. Vickers hardness values plus or minus the standard error of the mean for wild-type (Tg– Mmp20+/+) and Mmp20 null (Tg– Mmp20−/−) mice (440.1±52.77 N = 4); were 530.2±15.34 N = 8 and 164.9±8.063 N = 11 respectively. Enamel hardness was not significantly different between incisors from wild-type and Tg24i (H) Mmp20+/− mice (440.1±52.77 N = 4). However, Vickers harness values demonstrated that wild-type enamel (Fig. 7A) was significantly harder than Tg6i (M) Mmp20+/− enamel (451.9±21.93 N = 4; P<0.05) and also significantly harder than Tg42i (L) Mmp20+/− enamel (291.3±26.48 N = 7; P<0.0001). Therefore, each transgene in the Mmp20 null background increased the level of enamel hardness over that of the null mouse incisors, but only the highly expressed Tg24i (H) transgenic mice had enamel hardness values comparable to wild-type controls.

Microhardness results for transgenes expressed in Mmp20+/+ background were strikingly opposite to the results observed in the Mmp20−/− background. The only commonality between the groups was that transgene expression resulted in harder enamel than was present on the Mmp20 null (Tg– Mmp20−/−) incisors (Fig. 7B). In the wild-type background the Tg24i (L) transgenic mice had Vickers hardness values (540.2±9.192 N = 4) that were not significantly different from wild-type enamel. However, wild-type enamel was significantly harder than Tg6i (M) Mmp20+/+ enamel (393.7±39.45 N = 4; P<0.01) and was also significantly harder than Tg24i (H) Mmp20+/− enamel (271.1±48.71 N = 4;
Therefore MMP20 must be expressed within a tight range for the development of enamel with optimal hardness.

Enamel Matrix Protein and Amelogenin Banding Patterns

To determine if enamel matrix proteins are processed differently from normal when high levels of MMP20 are present, we performed an enamel matrix total protein analysis and performed immunoblotting for amelogenin. Five day-old first molars were collected and proteins were extracted from their developing enamel. The enamel proteins were run on an SDS-PAGE gel (Fig. 8A) and were subjected to immunoblotting for amelogenin.
amelogenin (Fig. 8B). As was previously observed [22], the Mmp20 null enamel has a prominent amelogenin band of approximately 27 kDa that was weakly present in enamel from wild-type mice indicating that the 27-kDa band is cleaved by MMP20. A band below 20 kDa was present in the wild-type enamel, but not in the Mmp20 null mouse enamel suggesting that it was an MMP20 cleavage product. This lower band was present in greatest quantities in the Tg6i (H) Mmp20+/− (N = 4) enamel and the higher molecular weight bands were less abundant for these two transgenes. In contrast, the low level expression of the Tg42m (L) transgene precluded the detection of the band below 20 kDa in the null background and the higher molecular weight bands were present at approximately wild-type levels in the Tg42m (L) Mmp20+/− mouse enamel. Apparently the band below 20 kDa is an MMP20 cleavage product that accumulates in greater quantities at the expense of the higher molecular weight bands when more MMP20 is present.

Discussion

Mmp20 transgenes driven by the amelogenin promoter and amelogenin downstream non-coding sequence were introduced into Mmp20 ablated mice to determine if the severe Mmp20 null phenotype could be reversed back to normal. Low, medium and high expressing transgenic mouse lines were selected to assess the level of Mmp20 activity necessary to revert the null phenotype and for their effect on enamel when present in the wild-type background. These experiments were performed on enamel from adult mouse incisors and on enamel from 5 day-old mouse pup first molars. Since rodent incisors continuously erupt, every stage of enamel development is present along the erupting incisor. Although the secretory stage of enamel development is only approximately 2 mm long near the basal end of the mouse incisor [36], extraction of total incisor enamel contained detectable MMP20 on immunoblots (Fig. 2). However, enamel matrix protein profiles from total incisor protein contain both MMP20 and KLK4 cleavage products. Therefore to assess total protein or amelogenin MMP20 cleavage profiles (Fig. 8), we used 5 day-old mouse pup first molars that were predominantly in the secretory stage of development when MMP20 is the only proteinase present in the enamel matrix.

It was surprising to find that transgene expression levels differed depending on whether expression was assessed in incisors or molars. Perhaps the localization of a given transgene within the mouse genome will favor expression in the continuously erupting incisor or favor expression in developing pup molars. Interestingly, the immunoblots for MMP20 were different depending on if the molar or incisor was probed. The wild-type molars (Fig. 2A, lane 3) show two MMP20 bands as indicated by the arrows while the wild-type incisors (Fig. 2C, lane 1) show three bands. Both bands identified by the arrows are catalytically active [37]. Therefore, the higher molecular weight MMP20 band observed from extracted incisor enamel may be the inactive zymogen. The results for Tg24m (M) transgene expression in molars were difficult to interpret. The qPCR, immunoblot and zymography results suggest that mice transgenic for Tg24 expressed this transgene at levels higher than the endogenous wild-type levels. However, the enamel present on molars in the Mmp20+/− background appeared malformed (Fig. 3) and although we tried several times, we had difficulty extracting total enamel proteins from Tg24m (M) Mmp20+/− mouse molars (Fig. 8). It appeared that amelogenin in these molars was not prevalent. Perhaps enamel formation was delayed in these mice so that their molars had a thin enamel layer with little protein. Paradoxically, amelogenins were successfully
Figure 8. Enamel matrix protein profile and immunoblotting of amelogenin splice and cleavage products. Total extracted enamel matrix proteins (mostly amelogenins) from 5-day old molars were run on an SDS PAGE gel (A). Comparison of the first two lanes containing extracted wild-type or Mmp20 null enamel matrix proteins demonstrate that the null enamel has a prominent band at approximately 27 kDa that is only weakly present in the wild-type lane indicating that this band is cleaved by MMP20. Also a doublet just above 20 kDa was present in the wild-type lane but was a single band in the null lane indicating that one of the wild-type bands is an MMP20 cleavage product. The protein profile for the Tg6m (H) transgene in the Mmp20+/– background was similar to that of the protein extracted from wild-type enamel. However, the protein profile for Tg6m (H) and Tg24m (M) in the Mmp20+/– background had a strong band below the 20 kDa marker and a weaker band beneath that indicating that more cleavage products than normal were present. Both transgenes in the wild-type background had bands above 20 kDa that were less prominent than the bands observed from wild-type enamel with no transgene. In contrast, regardless of the Mmp20 background, the Tg42m (L) positive mice had protein banding patterns that looked substantially like the wild-type results. Amelogenin immunoblot results from the various genotypes (B) were similar to the protein gel results. An approximate 27-kDa amelogenin band was present in the Mmp20 null but not wild-type lanes. In the wild-type background, both Tg6m (H) and Tg24m (M) transgenes had prominent amelogenin bands that located below 20 kDa and the bands above 20 kDa appeared less prominent than those observed in the Tg– wild-type lane. Also the amelogenin profile of Tg42m (L) in the wild-type background looked similar to that of wild-type mice.

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extracted from Tg24m (M) Mmp20+/+ molar enamel suggesting that transgenic MMP20 may lack an essential function that the endogenous MMP20 provides.

The most surprising finding was that for incisor enamel, the mid-level expressing transgene Tg6i (M) and the high-level expressing transgene Tg24i (H) almost completely recovered the Mmp20 null enamel phenotype. But, when these same transgenes were present in the wild-type background, the incisor enamel was severely compromised and enamel from mice transgenic for high-level expressing Tg24i (H) was more severely compromised than mice expressing the mid-level Tg6i (M) transgene (Fig. 5). Therefore, the endogenously expressed MMP20 combined with transgenic MMP20 activity resulted in poor quality enamel. This has striking implications. The results suggest that enamel matrix proteins have primary and secondary MMP20 cleavage sites and imply that if the secondary sites are cleaved too early due to much higher than normal MMP20 activity, malformed enamel will result.

In support of this interpretation, previous studies have shown that MMP20 rapidly cleaves amelogenin near its C-terminus and that progressive amelogenin degradation occurs with longer MMP20 incubation times [38]. A more detailed study demonstrated that MMP20 processes amelogenins into groups of cleavage products that accumulate and that are only slowly degraded further by MMP20 [39]. Ameloblastin [40] and enamelin [41,42] are the two other non-protease components of the enamel matrix. Although, these proteins are present in the matrix in far less abundance compared to amelogenin, they are nonetheless essential for murine enamel development [43,44]. Enamelin is so quickly hydrolyzed by MMP20 [45] that apparently all of its cleavage sites are primary MMP20 sites. Ameloblastin, however, is cleaved early by MMP20 at its N-terminus and secondary cleavages occur near the C-terminus [46,47]. Furthermore, during the secretory stage the enamel crystallites predominantly grow in length. However, over time MMP20 degrades enamel proteins and allows some growth of crystals in width and thickness, particularly in the deeper enamel [48]. MMP20 is the only known secretory stage enamel matrix proteinase. So, prolonged incubation with MMP20 may allow this growth. Again, this suggests that MMP20 has secondary enamel matrix protein cleavage sites that permit growth in width and thickness. Therefore, it is likely that a delicate balance exists in the enamel matrix among full-length proteins, proteins that are cleaved quickly, and proteins that are cleaved slowly and that too much MMP20 activity disrupts this balance.

Although transgene expression levels in the Mmp20 null background correlated positively with enamel quality, no difference among transgenic and wild-type mice existed with regard to incisor enamel thickness. With the exception of Tg24i (H) transgenic mice, the same was true for the transgenes expressed in the wild-type background. Therefore, for all but the most highly expressing transgene in the wild-type background, the enamel crystallites were able to extend to their full length to form a normally thick enamel layer. This suggests that enamel matrix protein cleavage products are more important for achieving fully mineralized enamel than they are for achieving a fully thick.
enamel layer. This fits well with current theories stating that full-length enamel proteins are responsible for crystallite elongation and that their cleavage products are more important for providing a supporting structure for the lengthening crystallites [19,49]. Perhaps very low Mmp20 expression as observed in the Tg42i (L) Mmp20<−/− enamel leaves behind un-cleaved proteins that interfere with proper mineralization and perhaps high Mmp20 expression levels cause early cleavage of secondary sites that compromise the crystallite support structure. When Mmp20 is expressed at very high levels such as in Tg24i (H) Mmp20<+/+ incisor enamel, the full-length enamel matrix proteins may be cleaved so quickly that they cannot adequately support continued lengthening of the crystallites. In any case, achievement of a normally thick enamel layer may occur over a wide range of Mmp20 expression levels, but achievement of normal enamel hardness has a much narrower range.

A prior study demonstrated that Mmp20 heterozygous mice expressed half the quantity of Mmp20 transcripts and their enamel contained approximately half the quantity of MMP20 protein as did wild-type mice [50]. This same study showed that no difference existed in enamel hardness between wild-type and heterozygous mice. Therefore, half the normal level of MMP20 activity does not significantly affect enamel mineralization. But from the present transgene results, it appears that if the level of MMP20 activity falls much below 50%, enamel mineralization will be adversely affected.

We conclude that enamel matrix proteins must be cleaved by MMP20 at a rate that is permissible for optimal enamel development. If the rate of cleavage is too fast or too slow, the enamel will be defective. Also, although the chances for under expression of MMP20 are much greater in humans than for its over expression, the finding that too high a level of MMP20 activity leads to enhanced degradation of enamel matrix proteins implies that there are likely few inhibitors or feedback controls over this protease once it is secreted.

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Author Contributions

Conceived and designed the experiments: JPS JDB. Performed the experiments: MS YC XG CCD JVA. Analyzed the data: MS CES JPS JDB. Contributed reagents/materials/analysis tools: CES JPS. Wrote the paper: MS CES JDB. Designed the look-up table (LUT) used in the pseudo-coloring analysis: CES.

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