Novel missense mutation of the FAM83H gene causes retention of amelogenin and a mild clinical phenotype of hypocalcified enamel

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A B S T R A C T

Objective: Amelogenesis imperfecta (AI) is a group of clinically and genetically heterogeneous inherited conditions, causing alterations in the structure of enamel and chemical composition of enamel matrix during development. The objective of this study was to compare the clinical, radiographic, histological and immunohistochemical phenotypes of subjects affected with hypocalcified AI from three Chilean families and identify causal mutations in the FAM83H gene.

Design: The diagnosis was made using clinical, radiographic, histological and genealogical data from the patients, who were evaluated according to the classification criteria by Witkop. PCR and Sanger sequencing of the complete coding sequence and surrounding intron regions of the FAM83H gene were conducted. The structural study of the affected teeth was performed with light microscopy, scanning electron microscopy and immunohistochemistry.

Results: The probands of the three families were diagnosed with hypocalcified AI, but in only one of them the missense variant p.Gly557Cys was identified. This variant was not present in the SNP database or in 100 healthy controls and segregated with the disease in the affected family. Using light microscopy, a normal prismatic structure was observed in all three cases. However, the ultrastructure was found to be affected in two of the cases, showing persistence of organic matter including amelogenins.

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Conclusions: These results suggest that FAM83H missense mutation reported in one of the families analyzed in this study might cause a phenotype of hypocalcified enamel more attenuated with retention of amelogenin.

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1. Introduction

The process of forming the enamel (amelogenesis) occurs in two stages: the secretion of an organic matrix and the subsequent mineralization of that matrix. During the secretory stage, ameloblasts deposit a matrix layer rich in protein, partially mineralized, over the previously formed dentine until reaching the complete thickness of the future enamel. These proteins are 90% amelogenins, while the remaining 10% contains ameloblastin, enamelin, tuftelin, enamelysin, kalli- krein and other proteins. During the maturation stage, the removal of organic materials and water and the continuous deposition of phosphate and calcium convert enamel into the hardest tissue in the human body.1–3

The normal process of amelogenesis is regulated by the expression of multiple genes. Mutations in any of these genes may result in changes in tissue formation, consequently alter enamel phenotypes. Defective synthesis of enamel is called amelogenesis imperfecta (AI). AI is defined as a group of genetic disorders that affect, to varying degrees, the structure and appearance of the enamel of all or nearly all the teeth in either the primary or permanent dentition or both6,5 and may occur as an isolated condition that only affects the enamel or in association with abnormalities in other tissues.6,7

AI has been classified into three main types, whose defects are correlated with the stages of enamel formation. In hypoplastic AI, the enamel is thin or contains defects due to inadequate apposition in localized or generalized areas, but has a higher radiopacity than dentine. In hypocalcified AI, the enamel is of normal thickness, but there is a defect in the nucleation and mineralization of the prisms, which results in soft enamel that is easily removable and less radiopaque than dentine. In hypomature AI, although the enamel thickness is normal, there is a defect in protein removal and the growth of crystals during the maturation stage, resulting in enamel with abnormal hardness and transparency and a decreased radiopacity similar to that of dentine.8–10

The AI prevalence varies widely depending on the population studied; for example, the reported figures are 1:4000 in Sweden,11 1:8000 in Israeli Jews12 and 1:14,000 in the U.S.13 For Chile, there are no data on the prevalence and/or incidence of AI.

To date, mutations in genes encoding secreted proteins of the enamel matrix (e.g., AMELX, ENAM and AMBN), enamel proteases (e.g., MMP20 and KLK4) and proteins that participate in biomineralization (e.g., C4orf26), ion transport (e.g., SLC24A4 and STIM1) and the cytoskeletal organization of keratins (e.g., FAM83H), intracellular vesicular traffic (e.g., WDR72) and cellular adhesion proteins (e.g., LAMB3 and ITGB6) have been shown to cause non-syndromic AI, with various patterns of inheritance. Eighteen mutations in different positions on the AMELX gene are responsible for cases of hypoplastic or hypoplastic/hypomatured AI with patterns of X-linked inheritance.14–16 The molecular defects underlying the various clinical forms of autosomal dominant hypoplastic AI (ADAI) are ten and five mutations in the ENAM and LAMB3 genes, respectively.10,17–20 Moreover, two mutations in the ENAM gene,21,22 5 mutations in the C4orf26 gene23 and 4 mutations in the ITGB6 gene were recently reported to cause hypoplastic AI with an autosomal recessive pattern of inheritance (ARAI).24,25 Mostly recently, a deletion of ameloblastin exon 6 (AMBN) has been associated with hypoplastic ARAI.26

The ARAI form of hypomature type can be caused by seven mutations in the enamelysin (MMP20) gene,27,28 two mutations in the kallikrein 4 gene (KLK4),27,29 six mutations in the WDR72 gene,29,30 four mutations in the SLC24A4 gene31,32 and one mutation in the STIM1 gene.33 The autosomal dominant forms of hypocalcified AI are associated with 20 defects in exon 5 of the FAM83H gene.34–44

In this paper, we report the results of a clinical and molecular genetic study of three Chilean families with hypocalcified AI, focusing our efforts on the analysis of the complete coding sequence and exon–intron junctions of the FAM83H gene, the most likely candidate for autosomal dominant hypocalcified AI (ADHCAI).

2. Materials and methods

2.1. Study subjects

Three unrelated Chilean families, including three probands affected with hypocalcified AI (two men and a woman) and eight healthy individuals (five women and three men), who are parents and siblings of the probands, participated in this study. The features and objectives of the research were explained to the eleven participants, who voluntarily signed a written informed consent form. This research was approved by the Ethical Scientific Committee (Act No.: 2013/06) and the Institutional Biosafety Committee of the Faculty of Dentistry (FDO No.: 2, June 24, 2013), University of Chile, and was conducted in full accordance with the ethical principles of the Declaration of Helsinki45 and local regulations.

2.2. Diagnosis of AI

The extra and intraoral clinical examination was conducted at the Diagnostic Service of Faculty of Dentistry, University of Chile, and was executed by two trained dental surgeons (AOP and DAF), who reached a consensus diagnosis. The participants were asked about systemic diseases, treatment with tetracycline, fluoride consumption and any other information that may affect the structure of their teeth during development.
The examination was performed with a No. 5 mirror and caries probe while air-drying the teeth being examined. Data were collected on anterior open bite, enamel hardness, changes in the shape and colour of teeth, such as the loss of contact points and alterations of the enamel surface, enamel wear and detachment from the underlying dentine. The affected and unaffected individuals were also clinically evaluated for the presence of abnormalities of the skin, hair, nails and bones, known to be a sign of syndromic conditions associated with AI. The clinical diagnosis were made based on historical, clinical, radiographic and genealogical data of the patients, who were evaluated according to the criteria of classification by Witkop. The following features were clinically evaluated in the affected individuals to determine whether a patient had hypocalcified AI: (1) soft, brittle enamel that is rapidly lost, revealing dentine, except in cervical areas and cusps tips, (2) tooth crowns having a cheesy or clay appearance often accumulate large deposits of calculus and their enamel radiographically less radiopaque than dentine, (3) enamel of unerupted or recently erupted teeth has a normal thickness, as determined by either radiography or clinical examination, (4) presence of a radiodense line of calcified enamel remaining in the cervical area and (5) yellowish-brown enamel that becomes brown or black due to extrinsic pigmentation acquired after the eruption.

2.3. Radiographic analysis

Periapical radiographs were taken (Heliodent DS, Siemens, Germany) of all affected individuals to detect the radiopacity of enamel and dentine, contrast between enamel and dentine, enamel thickness and to evaluate taurodontism. Additionally, left lateral cephalometric films were obtained (Orthophos DS, Siemens, Germany) from all affected participants in each family, and a cephalometric analysis with Björk-Jarabak and Steiner’s SNA and SNB traces (Quick Ceph 2.0) were conducted to assess skeletal open bite.

2.4. Molecular genetic analysis

Pedigree was constructed based on the family history of AI disease using interviews of the probands, their parents and relatives who provided informed consent to participate. For DNA purification, 10 ml of venous blood was taken from the patients and control subjects, and the genomic DNA was isolated from leukocytes by the salting-out method described by Miller et al. Complete sequencing of the coding regions and surrounding introns of the FAM83H gene in the probands of the three families was performed by amplification via polymerase chain reaction (PCR) with oligonucleotide primers designed based on the reference sequence of the FAM83H gene (NG_016652.1) using the Primer-BLAST software. The sequence and size of each amplified product and the associated annealing temperature are described in Table 1. The PCR products were extracted from agarose gels and purified using the Ultraclean® 15 DNA purification kit from Mo Bio Laboratories, Inc. (CA 92010, United States). The purified amplicons were sequenced in both directions by the DNA sequencing centre in Macrogen, Seoul, Korea. For computational sequence analysis, Vector NTI version 10.0 software and FAM83H gene reference sequences NG_016652.1 and NM_198488.3 were used. The detection of variant g.10953G–T in control subjects was conducted by direct sequencing and by Restriction Fragment Length Polymorphism-coupled PCR (RFLP-PCR), using primers that amplify a 332-bp fragment of exon 5 of the FAM83H gene (Table 1) and digestion with the Apal restriction enzyme.

2.5. Histological analysis

Ground sections. For the histological analysis of AI-affected teeth, three teeth were examined: one deciduous second molar exfoliated physiologically (proband III.5 of the FA12 family) and two permanent third molars extracted for orthodontic purposes (proband II.1 of the FA15 and FA14 families, respectively). Additionally, two teeth donated voluntarily by healthy individuals not related with the families were used as controls (deciduous molar and third molar). These teeth were subjected to wear with a diamond-cutting saw and were mounted for examination by light microscopy.

Scanning electron microscopy (SEM). The teeth were cleaned with 5% sodium hypochlorite, fractured and dehydrated in ascending acetone concentrations of 70%, 90% and 100%. The metallic finish and drying were performed at the Laboratory of Scanning Electron Microscopy in a Zeiss microscope model DMS 940 from the Faculty of Medicine, University of Chile.

Immunohistochemistry. For antigen retrieval, the cuts were exposed to 0.01 M citrate buffer (pH 6) in a heat source at 95 °C for 20 min. To block endogenous peroxidase activity, the sections were placed in 3% hydrogen peroxide in methanol for 10 min at room temperature. Sections of 5 μm were incubated with anti-human-amelogenin (Anti-AMELX produced in rabbit, polyclonal, diluted 1:200; Sigma HPA-005988, Sigma–Aldrich®) and Amelogenin code F-11 produced in mouse, monoclonal, diluted 1:2500; sc-365284 code, Santa Cruz Biotechnology®) for 20 min at room temperature. Subsequently, the cuts were washed in PBS and incubated with the biotinylated secondary antibody for 20 min at 37 °C. The avidin–biotin–peroxidase complex was used to bind the primary antibody using VECTASTAIN® (ABC, CA 94010, United States). The reaction products were visualized by immersion of the histopathological sections in diaminobenzidine (DAB) in the case of anti-AMELX and aminoethylcarbazole (AEC) for amelogenin antibody F-11 for 3–5 min. Nuclear counterstaining was performed with Harris haematoxylin. To study the negative controls for the antibodies, sections were treated with phosphate-buffered saline instead of primary antibody, and the absence of staining was confirmed. The positive controls used tooth germ in the advanced bell stage from a tumour resection and an included third permanent molar obtained from a dentigerous cyst.

2.6. Bioinformatic analysis

The effect of the sequence variant found in the FAM83H gene of the proband in the family FA14 was predicted using the computer programmes Polymorphism Phenotyping v2 (PolyPhen), Protein ANnotation Through Evolutionary Relationship (PANTHER) and Pathological Mutations (FMut). The analysis of the pattern of disulfide bonds was performed with the server DiAminoacid Neural Network Application (DiA-NNA), and the analysis of the interactions with other
Table 1 - Primers designed with the Primer-BLAST programme to amplify and sequence the FAM83H gene in probands and controls of the study.

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proteins used the Database of Interacting Proteins (DIP)52 and Human Protein–Protein Interaction Prediction (PIPs)53. Sequence information for the variants was submitted to a SNP database (dbSNP)54 and was assigned the identification numbers of ss715578273 and rs312262803.

3. Results

3.1. Clinical and radiographic findings

In this study, three unrelated Chilean families were diagnosed with hypocalcified AI. The clinical, radiographic and cephalometric parameters of two of the probands (FAI2, subject III.5 and FAI5, subject II.1) were described in detail in a previous study.52 As seen in the clinical picture presented in Fig. 1A, the teeth of the proband III.5 of family FAI2 have a yellowish-brown colouration and rough surface. This enamel was soft, which was evident by the wear of cusps and incisal edges and enamel largely missing from the crown, except in the cervical region, which was relatively better preserved (Fig. 1A, clinical picture, black arrowheads). The patient had dental hypersensitivity to heat stimuli, generalized gingivitis and multiple composite resins restorations, some of which were fractured (Fig. 1A, clinical picture, white arrowhead). The proband also had delayed eruption, and cephalometrically, he presented balanced mesomorphic facial profile. Radiographically, there was a similar contrast between enamel and dentine and generalized detachment of the enamel from the erupted teeth, similar to adits were observed (Fig. 1A, upper radiography, white arrowheads); in contrast, those teeth that had not erupted exhibited normal crown anatomy, with enamel of normal thickness but lacking contrast from the dentine (Fig. 1A, lower radiograph, white arrowhead). The intraoral examination of the parents and two brothers noted no deviation from normal enamel with respect to colour, surface, hardness or thickness.

Similar to the previous case, proband II.1 of family FAI5 had yellowish-brown enamel on all teeth. The enamel had a rough surface and was detached and worn widely, except in the
cervical region (Fig. 1B, photo clinic). This patient had generalized gingivitis, as noted by the characteristic erythematos interdental papillae and increased volume (Fig. 1B, black arrowhead). This proband also had dental hypersensitivity to thermal stimuli and previous composite restorations, which became detached or fractured shortly after being installed. The radiographic analysis showed that the enamel and dentine were similarly contrasted in both unerupted and
erupted teeth. The erupted teeth had thinner enamel with a tendency of fracture (Fig. 1B, X-ray, white arrowhead), in contrast to the unerupted pieces, which had enamel with an approximately normal thickness on a preserved and fairly normal crown anatomy (Fig. 1B, X-ray, red arrowhead). The parents of this patient could not be involved, but the patient indicated that they did not present similar abnormalities.

Proband II.1 of family FAI14 had grey-brown enamel with a rough texture, some detached on the labial surfaces (Fig. 1C, clinical picture, black arrowheads) and worn cusps and incisal edges. The enamel in the cervical region maintained an average thickness with a less intense colour. The patient presented tooth hypersensitivity to thermal stimuli, generalized gingivitis and numerous composite restorations in poor condition. Radiographically, in contrast to the previous cases, a somewhat greater contrast in the enamel, especially at the lower molar (Fig. 1C, radiograph, white arrowheads) was observed. These molars, in turn, had near intact clinical crown. The intraoral examination of the mother and two half-sisters did not reveal any alterations in enamel colour, surface hardness or thickness.

3.2. Molecular genetic analysis

According to the pedigree shown in Fig. 1A, individual III.5 of family FAI2 was found to be affected, and individual IV.2 was reported to be affected by the mother of the proband, but he was unwilling to be evaluated clinically. In the pedigree shown in Fig. 1B, the parents of the proband were reported to be unaffected. A similar situation was observed in the FAI14 family, whose pedigree (Fig. 1C) shows that the proband is the only affected subject. In this case, we were only able to examine her mother and two half-sisters, who did not have the trait. In addition, the proband’s mother reported that neither parent had dental problems. Thus, the data collected from these three families indicate either a recessive inheritance pattern or sporadic cases due to new mutations.

The sequencing analysis of gene FAM83H performed in the probands of the three families revealed the presence of a mutation in exon 5 in only one of the probands; subject II.1 of family FAI14 (Fig. 1C, chromatogram of the subject II.1). The other two probands were negative for this analysis. The variant identified corresponds to a transversion substitution that changes a guanine-to-thymine at position 10953 of the genomic DNA (NG_016652.1 reference sequence) and position 1669 of the cDNA (NM_198488.3 reference sequence). A missense change occurs at the protein level involving the replacement of a nonpolar amino acid (glycine) by a polar (neutral or uncharged) amino acid (cysteine) at position 557 (NP_940890.3). The co-segregation analysis in the family showed that the mother and older half-sister had no such change in the sequence (Fig. 1C, chromatograms of the subjects I.2 and II.2) and that only the proband carried the variant in a heterozygous state, corroborating that this mutation is dominant. The absence of this variant in the SNP database, the 1000 genomes project database, in 100 healthy control subjects analyzed by PCR-RFLP and 15 controls analyzed by Sanger sequencing was confirmed. Thus, we believe that this variant could correspond to the causal mutation in the family studied.

3.3. Bioinformatic analysis

The programme PolyPhen 2.0 predicted that the substitution was most likely pathogenic, with a score of 0.992. The PANTHER programme indicated that the mutation was deleterious with a probability of 0.851, and the PMut programme suggested that the variant was pathological with a probability of 0.841 and a reliability value of 6. These results support the pathogenic role of the mutation found in the proband of the FAI14 family.

The mutation reported in this study involves the substitution of glycine at position 557 by a cysteine (p.Gly557Cys). Glycine is an amino acid with a hydrogen side chain, which does not interact with the environment, whereas cysteine, with its sulphydryl group side chain, can form disulfide bonds with metals. As the wild type FAM83H protein has nine cysteine residues in its amino acid sequence at positions 57, 196, 218, 237, 550, 590, 789, 1090 and 1140, determining whether this substitution may alter the pattern links in the mutated protein compared with the wild-type would be interesting. The prediction analysis performed with the DiANNA programme suggested that the pattern of disulfide bonds in the mutant protein changes. In wild-type FAM83H, C590 likely shares a disulfide bond with C1090 (score 0.89631). In the mutant protein, C557 could establish a disulfide bond with C1090 (score 0.99824) or C590 (score 0.99436) and is more likely to form a new bond between C557 and C590. Moreover, with respect to the formation of bonds with metals, the predictions indicate that only C550 might be a possible target for metals and that the remaining cysteines, including the substitution to C557, may not form bonds with metals.

Additionally, the analyses performed with the DIP and PIPS programmes, which predict interactions with other proteins, showed that both the wild-type protein and the mutant had no significant interactions with other proteins.

3.4. Histological, ultrastructural and immunohistochemical analysis

Light microscopy analysis with 100X magnification of the cuts by wear observed in those affected with hypocalcified AI; second deciduous molar of proband III.5 from the FAI2 family (Fig. 2A); third molar of proband II.1 from the FAI5 family (Fig. 2B) and third molar of proband II.1 of the FAI14 family (Fig. 2C) revealed no alterations in the structure of the enamel at this level, with the prism structure provided in the three cases being similar to that of the control tooth (Fig. 2D). The SEM analysis enabled the study of the ultrastructure of the teeth of the three affected individuals. In the cases of the second molar of proband III.5 from the FAI2 family (Fig. 2E) and the third molar of proband II.1 from the FAI5 family (Fig. 2F), both teeth had a prismatic structure similar to that of their respective controls (Fig. 2H, third molar). Furthermore, in the third molar of proband II.1 from family FAI14, areas in which the prisms do not maintain a constant thickness, similar to rosette beads, with upraised projections were observed (Fig. 2G), and in other sections of the same piece, a slightly irregular amorphous prismatic structure was observed (not shown).

The histopathological and Immunohistochemical analysis using the anti-amelogenin antibody showed the presence of amelogenin-like proteins through an intense brown staining
Fig. 2 – Images A–D: histological, E–H: ultrastructural, I: histopathological and J–L: immunohistochemical analysis of the enamel of teeth affected with hypocalcified AI. Enamel ground sections of the deciduous second molar of proband FAI2-III.5 (A), the third molar of proband FAI5-II.1 (B), the third molar of proband FAI14-II.1 (C) and the third molar of a normal subject (D). Photographs of SEM of the enamel from deciduous second molar of proband FAI2-II.1 (E), the third molar of proband FAI5-II.1 (F), the third molar of proband FAI14-II.1 (G) and the third molar of a normal subject (H). Haematoxylin and Eosin staining of the enamel from deciduous second molar of proband FAI2-III.5, showing enamel that retains a lot of organic matter (I). Photomicrographs of the immunohistochemical technique using the anti-amelogenin antibody in enamel of the third molar of proband FAI5-II.1 (J), the third molar of proband FAI14-II.1 (K) and a included third molar as positive control (L).
in the sample from proband II.1 of family FAI5 (Fig. 2J) and through red staining in proband II.1 of family FAI14 (Fig. 2K). Moreover, in both preparations, areas of prismatic enamel (Fig. 2J, black arrow and Fig. 2K, white arrow) and amorphous areas of apcrismatic enamel (Fig. 2J, black arrowhead and Fig. 2K, white arrowhead) could be observed. With respect to the areas of prismatic enamel, brown staining was found in both the periphery and interior of the prisms in the sample from the proband II.1 of family FAI5 (Fig. 2J, black arrow), whereas the reddish staining was confined to the periphery of the prisms in the sample from proband II.1 of family FAI14 (Fig. 2K, white arrow). Unfortunately, due to the exhaustion of the study material, this technique could not be performed in the sample from proband III.5 of family FAI2, however the histopathological analysis using H&E staining, showed persistence of organic matter in the affected enamel (Fig. 2I).

Moreover, the study of the negative controls for the teeth of probands II.1 of family FAI14 and II.1 of family FAI5 using phosphate-buffered saline instead of the anti-amelogenin antibody showed no staining (not shown), whereas the analysis of the positive controls resulted in intense labelling. In the case of the advanced bell stage tooth germ, intense brownish staining that was relatively abundant compared with the matrix secreted by the ameloblasts was observed (not shown); in the case of the third molar, intense reddish staining was observed in the sample (Fig. 2I).

4. Discussion

This study characterized patients with AI from three unrelated Chilean families from clinical, radiographic, histological, immunohistochemical and genetic perspectives. As the FAM83H gene is the only gene to date in which causal mutations have been linked to hypocalcified AI, it was decided to molecularly analyze this gene in the recruited probands. Also teeth had been voluntarily donated by the probands and their parents, which allowed for a macro and ultrastructural study, using light microscopy and SEM. Moreover, in two of the cases studied (II.1 FAI5 and II.1 FAI14), the presence or absence of amelogenin-like proteins was determined by immunohistochemistry on tooth sections.

4.1. Clinical and radiographic analysis

The clinical features of the three patients are most consistent with hypocalcified AI (Fig. 1A–C), exhibiting characteristics similar to those proposed in the classification by Witkop4 and described in other studies with the same type of AI, such as the following: yellowish-brown staining, rough texture, wear or loss of enamel upon eruption and a similar contrast between the dentine and enamel.7,34,35,42,43

Although AI corresponds to an alteration in the structure and appearance of the enamel, other dental anomalies may coexist,52 such as the delayed eruption in proband III.5 of the FAI2 family. In addition, the three probands were diagnosed with generalized gingivitis, which was most likely due to the rough surface of the enamel that allowed for the retention of plaque, which adds dental hypersensitivity to thermal stimuli hindering proper oral hygiene. Additionally, the three affected individuals had composite restorations that were fractured or failed after a period of time, which may be due to poor adherence of the material to the affected enamel surface. Notably, proband II.1 of the FAI14 family (Fig. 1C) showed some particular clinical features, such as average enamel thickness in the middle region and less intense discoloring of the enamel in the cervical region. Some of their teeth, especially the lower molars, were relatively intact, enabling their crown anatomy to be distinguished on radiographs with a slightly higher contrast in the enamel. Other studies37,39 have also described distinctive phenotypes in some families affected with hypocalcified AI, with the diagnoses confirmed by the finding of mutations in the FAM83H gene. Wright et al.36 found that affected individuals in two families had hypocalcified enamel exclusively in the cervical third of the teeth. Song et al.7 found that the affected members of a family had the classic features of hypocalcified enamel but a polished and shiny surface. These observations confirm the great clinical and radiographic variability that may exist not only between different types of AI but also within the same clinical type.

4.2. Genetic and molecular analysis

The pedigree of family FAI2 revealed that subject III.5 was the only one affected (Fig. 1A). While individual IV.2 was reported to be affected, he was not examined, suggesting recessive inheritance or sporadic-case in all three families. The pedigrees comprise only two generations, and thus an inheritance pattern cannot be established with certainty (Fig. 1B and C) nor can sporadic cases corresponding to spontaneous or de novo mutations be confirmed.

The mutational analysis of gene FAM83H involved the study of the entire coding sequence and intron–exon neighbourhoods using PCR and direct sequencing. No mutation in this gene was found in the DNA from two probands; FAI2-II.5 and FAI5-II.1. However, in proband II.1 of family FAI14, the variant g.10953G>T (c.1669G>T; p.Gly557Cys) was found (Fig. 1C), which has not been reported as a SNP or as a previously described causal mutation when it was discovered. Co-segregation analysis by direct sequencing using DNA from the mother, half-sister and 15 healthy controls without AI revealed that none of the relatives or controls had that variant. Furthermore, a PCR-RFLP analysis designed in the laboratory for the presence of variant g.10953G>T was performed in 100 healthy controls, including 15 subjects previously verified by direct sequencing. These results and the absence of the variant in the SNP database and the 1000 genomes project support the finding that this sequence change is the cause of the clinical phenotype in the FAI14 family.

The paralogous sequence alignment of eight family members of the FAM83 family (FAM83 A–H) with orthologous sequences in seven vertebrate species performed by Ding et al.39 showed that C557 corresponds to a relatively conserved amino acid; four members of the FAM83 family and three species of mammals, which are closer on the evolutionary scale (human, mouse and bovine), had a conserved glycine in the aforementioned position. The missense change of p.Gly557Cys likely affects the function of the gene product as cysteine is able to form disulfide bonds, which may alter the three-dimensional structure of the protein and thereby affect...
its function. The in silico prediction of bond formation with metals and formation of disulfide bridges showed that C557 did not produce metal bonds but did alter the pattern of disulfide bonds within the mutated protein compared with the wild-type because the latter may generate disulfide bonds with C590 and C1090 whereas the mutant can form bonds with C557, C590 or C1090. The PolyPhen 2.0, PANTHER and PMut programmes showed that the p.Gly557Cys substitution is most likely pathogenic with scores of 0.992, 0.851 and 0.841, respectively.

All alterations reported to date in the FAM83H gene correspond to nonsense and frameshift mutations (Table 2) that introduce a premature stop codon within amino acids S287 and E394, resulting in a truncated protein.7,36,43 These truncated proteins will interfere with the normal protein and produce the phenotype through a dominant negative effect on the wild-type protein.40 Although the mutation reported in this study (g.10953C>T, c.1669G>T and p.Gly557Cys) is located in the same region of exon 5 of FAM83H where all known mutations have been reported, this report is the first of a missense mutation causing an amino acid change in the protein that is associated with a phenotype of generalized enamel, although the phenotype is attenuated or less severe than those described above (Table 2).

At present, at what stage of the process of amelogenesis does the defect that causes hypocalcified AI occurs is unknown; if this occurs in the primary mineralization during the secretary stage, or if this happens in the secondary mineralization during the maturation stage, or another instance between or throughout the two stages.

Until recently, the role of FAM83H was largely unknown. Ding et al.39 reported that mouse Famm3h is located in the intracellular environment and is associated with perinuclear vesicles in the vicinity of the Golgi apparatus. Subsequently, Lee et al.40 demonstrated in vitro that the mutations p.Q677*, p.W460* and p.R325* altered the localization of the protein, increasing its concentration within the nucleus.

Recent reports attributed FAM83H a role in the development of various cancers.58-60 The study by Kuga et al.61 conducted on cell lines and tumour tissues of colorectal cancer demonstrated in vitro and in vivo that FAM83H associates with casein kinase 1 alpha (CK-1α) and with keratins 8 and 18. In this association, FAM83H participates as a linker protein

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**Table 2 – Reported mutations to date in exon 5 of the FAM83H gene.**

<table>
<thead>
<tr>
<th>No.</th>
<th>cDNA</th>
<th>Protein</th>
<th>Family ID</th>
<th>Ethnicity</th>
<th>Pattern of inheritance</th>
<th>References</th>
<th>Clinical phenotype</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>c.973C&gt;T</td>
<td>p.R325*</td>
<td>F1ADHC1A</td>
<td>Korean</td>
<td>AD</td>
<td>Kim JW, 200874</td>
<td>Generalized hypocalcified</td>
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<tr>
<td>2</td>
<td>c.1192C&gt;T</td>
<td>p.Q398*</td>
<td>F2ADHC1A</td>
<td>Korean</td>
<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>3</td>
<td>c.1243G&gt;T</td>
<td>p.E415*</td>
<td>F1(AI#8)</td>
<td>Asian</td>
<td>AD</td>
<td>Lee SK, 20088</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>4</td>
<td>c.891T&gt;A</td>
<td>p.Y297*</td>
<td>F2(AI#12)</td>
<td>Asian</td>
<td>AD</td>
<td>Lee SK, 20088</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>5</td>
<td>c.1380G&gt;A</td>
<td>p.W460*</td>
<td>F3(AI#21)</td>
<td>Caucasian</td>
<td>AD</td>
<td>Lee SK, 20088</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>6</td>
<td>c.1209G&gt;T</td>
<td>p.Q677*</td>
<td>F4(AI#24)</td>
<td>Caucasian</td>
<td>AD</td>
<td>Chan HC, 20118</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>7</td>
<td>c.1330C&gt;T</td>
<td>p.Q444*</td>
<td>Family 1,2,3</td>
<td>Turkish</td>
<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>8</td>
<td>c.1366C&gt;T</td>
<td>p.Q456*</td>
<td>Family 1,2,3</td>
<td>Turkish</td>
<td>AD</td>
<td>Ding Y, 20098</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>9</td>
<td>c.1408C&gt;T</td>
<td>p.Q470*</td>
<td>A1C5, A1C33</td>
<td>Caucasian</td>
<td>AD</td>
<td>Wright JT, 20098</td>
<td>Generalized hypocalcified</td>
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<tr>
<td>11</td>
<td>c.923_924delTC</td>
<td>p.L308fs*323</td>
<td>A1C38</td>
<td>Caucasian</td>
<td>AD</td>
<td>Wright JT, 20098</td>
<td>Generalized hypocalcified</td>
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<tr>
<td>13</td>
<td>c.1872_1873delCC</td>
<td>p.L625fs*703</td>
<td>A1C46</td>
<td>Caucasian</td>
<td>AD</td>
<td>Wright JT, 20098</td>
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<td>14</td>
<td>c.2080G&gt;T</td>
<td>p.E594*</td>
<td>A1T0-002</td>
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<td>15</td>
<td>c.1354C&gt;T</td>
<td>p.Q452*</td>
<td>Family 39</td>
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<td>AD</td>
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<td>16</td>
<td>c.1993C&gt;T</td>
<td>p.Q565*</td>
<td>Family 4, 4.4</td>
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<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
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<tr>
<td>18</td>
<td>c.906T&gt;G</td>
<td>p.Y302*</td>
<td>Fam 1, IV.12</td>
<td>Danish</td>
<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>19</td>
<td>c.1289C&gt;A</td>
<td>p.S430*</td>
<td>Family 2, 3.1</td>
<td>Chinese</td>
<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>20</td>
<td>c.924dupT</td>
<td>p.V309Kfs*324</td>
<td>Fam 1, IV.14, FA14LP</td>
<td>Chinese</td>
<td>AD</td>
<td>Song YL, 20127</td>
<td>Generalized hypocalcified</td>
</tr>
<tr>
<td>21</td>
<td>c.1669G&gt;T</td>
<td>p.G557C</td>
<td>Fam 1, III.1</td>
<td>Korean</td>
<td>AD</td>
<td>Lee SK, 20118</td>
<td>Generalized hypocalcified</td>
</tr>
</tbody>
</table>
between CK-1α and keratins. FAM83H interacts with CK-1α by means of its amino terminal region and the amino acid sequence FDEEFRILF and its carboxyl terminus binds keratins. Thus, FAM83H can recruit CK-1α to keratin filaments, enabling CK-1α to modulate the filamentous status of keratins by phosphorylation/dephosphorylation. Under normal conditions, the levels of FAM83H expression orchestrate the subcellular distribution and function of CK-1α, thereby allowing FAM83H to regulate the keratin cytoskeleton organization by recruiting CK-1α to the filaments. These observations allow us to suggest that the g.10953C>T mutation, acting via a mechanism of haploinsufficiency, could disrupt the cytoskeletal keratins and partially affect the function of the ameloblasts and enamel synthesis, thus explaining the more attenuated clinical phenotype observed in the proband. Moreover, the phenotype associated with the same case, i.e., the retention of amelogenin in the periphery of the prisms, which was observed in the tooth of the proband, could be explained by the simultaneous gain of a regulatory function of FAM83H on the expression and/or degradation of amelogenin during amelogenesis.

In patient III.5 of the FAI2 family and patient II.1 of the FAI5 family, no mutations in FAM83H explain their phenotype, similar to the findings in other studies on hypocalcified AI. For example, Lee et al. found mutations in only four of seven families; three had AD inheritance and the other had a de novo mutation. Haubek et al. found mutations in two of six families, one with AD inheritance and the other with a de novo mutation. Finally, Song et al. found causal mutations in five of six families, with all showing AD inheritance. From our study, we cannot rule out the existence of causal mutations in non-coding or regulatory regions of the FAM83H gene that were not analyzed in this study or previous studies or the presence of other yet to be discovered enamel genes that may also be responsible for AI in families where causal mutations have not been determined.

4.3 Histological, ultrastructural and immunohistochemical analyses

Previous studies in hypocalcified AI teeth using light microscopy and SEM reported diverse results. Some studies have described areas of prismatic enamel with non-crystalline amorphous material, possibly proteins; crystals with a normal shape, size and orientation but of higher porosity and that are more labile; prismatic structure with cracks or crevices that could not have any mineral content; incomplete malformed prisms with presence of amorphous material, which may indicate organic material retention. Similarly, in this study, diverse results were also found. The light microscopy and SEM analyses showed that the enamel prismatic structure of the teeth of proband III.5 of FAI2 family and proband II.1 of family FAI5 appeared similar to that observed in the controls. However, in the tooth of proband II.1 of family FAI14, the prism structure did not maintain a constant diameter under SEM, with upraised projections, and some sectors that maintain irregular prismatic structure.

The immunohistochemical analysis detected the presence of amelogenin, demonstrated by an intense brown labelling, both inside and on the periphery of the enamel prism of the tooth of the proband II.1 of family FAI5. In the sample from proband II.1 of family FAI14, bright red labelling was observed only in the periphery of the prisms, suggesting that in the latter case, the organic component of the interior of the prisms is removed in a timely manner. Unfortunately, which of the two samples had less remaining amelogenin could not be determined, which would have been interesting because if proband II.1 of family FAI14 had less, the phenotype could be related to his crown anatomy more conserved and the largest contrast between the enamel and dentine in some of his teeth. Other studies using the teeth of subjects affected with hypocalcified AI have also found a high protein content and in some cases, reactivity has been demonstrated with anti-amelogenin antibody by western blot analysis. Because the average protein content of mature enamel is less than 1%, the higher protein content, combined with a lower mineral content, is postulated to cause the enamel to wear easily and serve as a poor insulator against thermal changes, causing hypersensitivity.

5. Conclusions

These results suggest that FAM83H missense mutation reported in one of the families analyzed in this study might cause a phenotype of hypocalcified enamel more attenuated with retention of amelogenin. Additional experimental studies and new approaches to elucidate the effects of this mutation on the clinical phenotype of the enamel and to further define the mechanisms that simultaneously affect tissue formation and degradation as well as amelogenin expression during the formation thereof are required.

Authors’ contributions

B.U. conceived the study and its design, recruited and interviewed patients, coordinated, designed, conducted various experiments and supervised all molecular genetic study, drafted and wrote the manuscript, leads the research project and funding. C.M. participated in part of molecular genetic and part of clinical and histological analysis. A.O.F. participated in clinical examination and assessment of patients, interpretation of histological results, and contributed in writing and revising the manuscript. D.A. performed clinical examination and assessment of patients. A.P. helped with the molecular studies. I.M.B., L.J., C.L. and C.L. participated in data analysis and critical review of the manuscript. G.R. performed the bioinformatic analysis. M.R. performed the histological and immunohistochemical analysis.

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Competing interests

The authors declare that they have no competing interests.
Ethical approval

This project was approved by Scientific Ethical Committee, Faculty of Dentistry of the University of Chile (CEC-FOUCH). Informed consent/assent was obtained from all participants and their parents.

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REFERENCES

34. Kim JW, Lee SK, Lee ZH, Park JC, Lee KE, Lee MH, et al. FAM83H mutations in families with autosomal-dominant...
49. Protein ANotation TH rough Evolutionary Relationship software (PANTHER). http://www.pantherdb.org/.
52. Database of Interacting Proteins software (DIP). http://dip.doe-mbi.ucla.edu/dip/Main.cgi/.
53. Database of Human Protein-Protein Interaction Prediction software (PPIs). http://www.compbio.dundee.ac.uk/www-pips/.