



Case Report: Crown Resorption in a Patient With Junctional Epidermolysis Bullosa and Amelogenesis Imperfecta With *LAMB3* Gene Mutations

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Background: Epidermolysis bullosa (EB) corresponds to a series of conditions characterized by extreme fragility of the skin and/or mucous membranes. Of the four main types of EB, junctional EB (JEB) is the most associated with alterations in the teeth. The purposes of this study were to determine the clinical, histopathological, and ultrastructural characteristics of teeth with amelogenesis imperfecta (AI) in a patient with JEB, and compare them with control teeth, and correlate the findings with the mutations present in the patient.

Case Report: The study was conducted on a 10-year-old patient with JEB carrier of two recessive mutations in the *LAMB3* gene and absence of the laminin-332 protein (LM-332), determined by immunofluorescence on a skin biopsy. The patient presents hypoplastic AI with very thin and yellow-brown colored enamel. Extraction of two permanent molars was performed due to pain and soft tissue covering the crown, resembling pulp polyp or hyperplastic gingiva. Light and scanning electron microscopy (SEM) revealed very thin enamel varying from complete absence to 60 µm, absence of normal prismatic structure, and presence of a cross-banding with a laminated appearance. The histopathological study revealed granulation tissue causing external crown resorption.

Conclusion: Although coronary resorption has been reported in patients with syndromic and non-syndromic AI, this is the first clinicopathological report of coronary resorption in partially erupted teeth in patients with JEB with mutations in the *LAMB3* gene and hypoplastic AI. In patients with this condition, the presence of partially erupted teeth with soft tissue covering part of the crown, without a periodontal pocket, and with a radiographic image of partial coronal radiolucency should lead to suspicion of external coronary resorption.

Keywords: amelogenesis imperfecta, junctional epidermolysis bullosa, crown resorption, laminin-332, rehabilitation of teeth, bullous genetic disease

INTRODUCTION

Among the genetic conditions that can affect tooth development is epidermolysis bullosa (EB), which corresponds to a series of conditions characterized by fragility of the skin and mucosa, where blisters are generated by minimal trauma. In this disease, mutations have been reported in the genes encoding proteins related to the basal lamina and epidermis, causing severe architectural alterations in tissues (1, 2).

Four types of EB are recognized: simplex (EBS), junctional (JEB), dystrophic (DEB), and Kindler epidermolysis bullosa (KEB). Patients with all four types of EB can present skin fragility and mucous membranes lesions such as oral blisters and ulcers. All patients diagnosed with JEB have alterations of the structures of the teeth. In JEB, the most frequently observed alteration of the teeth is a hypoplasia of the enamel, which can be extremely thin (2–5). EB can be caused by alterations in 1 of 16 genes encoding proteins related to the maintenance of dermo-epidermal junction proteins, which provide the skin with enormous stability (2).

Junctional EB occurs due to recessive mutations in the genes encoding the laminin-332 protein (*LAMA3*, *LAMB3*, and *LAMC2*), integrin $\alpha 6\beta 4$ (*ITGA6* and *ITGB4*), integrin $\alpha 3$ (*ITGA3*), and collagen type XVII (*COL17A1*). The three protein complexes are located in the lamina lucida, joining the basal keratinocytes with the lamina densa, thus giving stability and adhesion to the two regions (2, 3, 6).

Junctional EB is most frequently caused by recessive mutations in the *LAMB3* gene, which, in addition to the skin phenotype, also produces amelogenesis imperfecta (AI) in these patients (4, 5, 7). On the other hand, dominant mutations in *LAMB3* can cause non-syndromic AI (8–10). In these cases, the teeth present a thin enamel in the temporary and permanent dentition, but no skin conditions are described. Interestingly, a family has been described in which the proband presented JEB with a dominant mutation in the gene for collagen 17 (*COL17A1*; p.G627V) and hypoplastic tooth enamel in both dentitions. The mother carrying the same mutation had hypoplastic tooth enamel and an absence of skin fragility (11).

These findings suggest that the dental enamel would be more prone than the skin to express clinical changes due to the mutations in the genes that synthesize proteins of the lamina lucida (11). However, there has been only limited information regarding dental structural alterations associated with the different mutated genes responsible for JEB. The main objective of this research was to compare the clinical, radiographic, histopathological, and ultrastructural characteristics of teeth with AI in one patient with JEB with the unaffected teeth of the control subject and to correlate these findings with the mutations found in *LAMB3*.

The results of this study are of great importance for patients since they contribute to an accurate diagnosis of each of these pathologies and their clinical variants. Several findings, in this case, are unique, such as the histopathological diagnosis of coronary resorption that clinically resembled a pulp or gingival polyp and the reduction in tooth sensitivity after crowning the remaining teeth affected by AI. These findings allow us to understand the pathogenesis of these conditions and contribute

to undertaking preventive measures and providing specific dental therapies to protect and/or restore altered dental structures in these patients with special needs.

CASE DESCRIPTION

A 10-year-old female patient with severe JEB presented with dental pain to the Special Care Clinic at the Faculty of Dentistry of the University of Chile. The patient has been on a preventive dental program since the age of 3 years. The care has been based on education and prevention using fluoride varnish applications. Clinically, very thin yellowish enamel was observed, both in the temporary and permanent dentition.

Her EB diagnosis was based on clinical, histopathological, and molecular findings. Immunofluorescence mapping showed the absence of the $\beta 3$ chain of laminin 332 in the zone of the basement membrane, and two recessive mutations in *LAMB3*: c.957_958dup77 and c.3228+1G>A (NM_000228.2) were identified and previously published in Fuentes I. (12).

Despite the generalized enamel hypoplasia, the patient had remained free of decay. At the dental checkup at age 10, she reported pain in the first permanent mandibular molars: teeth 3.6 and 4.6. Clinically, soft tissue was covering the crown of both permanent mandibular molars, compatible with pulp polyp or hyperplastic gingiva. The patient refused to receive restorative treatment because her perioral granulation tissue areas caused too much pain when opening the mouth for long periods, and she was concerned about the slow healing rate of these wounds. Therefore, a surgical approach was chosen (extraction of both molars). Both sockets healed uneventfully. The remaining teeth were restored using fixed crowns. Ceramic fixed crowns were performed on all eight incisors that were already fully erupted, and provisory acrylic crowns were used to cover the canines, premolars, and molars, which were partially erupted.

Surprisingly, the greatest benefit of covering the teeth was the reduction of tooth sensitivity while eating reported by the patient. She had never complained of sensitivity before because she has always had the same discomfort since her teeth erupted.

The morphological study of the two extracted molars presented in this article was part of the REGULAR FONDECYT project number 1140905, which was approved by the Ethics Committee of the Faculty of Dentistry at the University of Chile (Act No. 2013/06). The mutations of the patient were detected under research funded by INITIATION FONDECYT project number 11140440. Informed consent was obtained from all individual participants included in the study.

DIAGNOSTIC ASSESSMENT

Mutational Study

To detect mutations causing the disease, DNA from peripheral blood was isolated with a DNA extraction kit (Thermo Fisher Scientific, USA). The DNA was used as a template to amplify the exons of 16 genes related to EB. The resulting library was sequenced in the Ion PGM Sequencer (Life Technologies), and massive sequencing data were analyzed with Ion Reporter (Life Technologies Thermo Fisher, USA) software. To confirm

the findings with massive sequencing, bidirectional Sanger sequencing was used. Finally, the pattern of recessive inheritance of both mutations found in the patient was confirmed by analyzing the genomic sequence of the parents and a healthy sister of the patient and was previously published in Fuentes I (12).

Histopathological Examination

The two extracted molars were sent to the laboratory in 10% buffered formalin. The molar with the greatest amount of soft tissue adhered to the crown was decalcified for routine histopathology in 10% ethylene diamine tetra-acetic acid (EDTA) for 4 weeks, processed by routine histopathological technique, and stained with hematoxylin and eosin. The second tooth was segmented for ground section light microscopy and scanning electron microscopy (SEM). The ground sections were made in a Buehler cutting machine IsoMet® 1000, Illinois, USA. Cuts of 0.6 mm were obtained, which were then thinned with Arkansas stone to achieve samples smaller than 0.4 mm. The samples were dehydrated in ascending alcohols and xylol and mounted for histopathological examination. The histopathological study was performed using an optical microscope (Olympus CX 21, Japan) with traditional and polarized light.

Scanning Electron Microscopy

For this study, the samples were fractured, washed in 7% EDTA to remove the smear layer, and dehydrated in ascending concentrations of alcohol. The preparation of samples was conducted in the electron microscopy laboratory of the Faculty of Medicine at the University of Chile (CESAT). These samples were analyzed with a Zeiss model DMS 940, German SEM.

RESULTS

Mutational Study

The results of the analysis allowed the identification of two recessive mutations in *LAMB3*: c.957_958dup77 and c.3228+1G>A (NM_000228.2) as previously published in Fuentes I (12). The pedigree, chromatogram, and agarose gel electrophoresis are shown in Supplementary Figure 1 (Not shown in previous reports).

Clinical and Radiographic Features

During the clinical examination of the patient, it was observed that the teeth had a very thin yellowish enamel. Clinically, soft tissue covered the crown of the lower first permanent molars, compatible with pulp polyp or hyperplastic gingiva (Figures 1A,B). Figure 1C shows the restored incisor teeth. Panoramic x-ray showed very thin, almost undetectable enamel. First mandibular molars presented partial coronal radiolucencies (Figure 1D).

Histopathologic Features

In the ground sections of an affected molar, extremely thin enamel was observed, varying from complete absence to very thin enamel. The thicker enamel was adjacent to the root cementum zone while it was absent in some coronary cusp

areas (Figures 2A,B). The enamel in the cervical area showed an absence of normal prismatic structure and the presence of a cross-banding with a laminated appearance compatible with cross-striations (Figure 3A, black arrow). Despite the marked thinness or absence of enamel, the dentin and cementum had a normal appearance (Figures 2A,B) except for the cementum at the apical level, which had an increased thickness (not shown). In contrast, control teeth presented an enamel more than 10 times thicker, with normal prisms (Figures 2C,D, 3C,D).

In the demineralized segment treated with routine histopathological technique, crown and cervical dentin resorption were observed, with the presence of odontoclasts, apposition of cement-like material, and slight chronic inflammatory infiltrate in the resorption connective tissue (Figures 4A,B). Figures 4C,D show a normal permanent tooth.

Ultrastructural Features

When analyzing the affected molars with SEM, the absence of enamel was observed in the cusp areas. In the cervical area, very thin enamel, up to 60 μm thick was measured. The cross-banding with a laminated appearance observed in the ground sections became more recognizable, with a width of 4 μm. The prisms remained undetectable (Figure 3B, black arrow). In the control teeth, the presence of normal prismatic structures was observed (Figure 3D).

DISCUSSION

Of the four types of EB recognized today, only patients with JEB presents significant alterations of the enamel structure (13). Among the defects of the dental structure, the most frequently reported are enamel hypoplasia, pits, and irregular surface (2, 13, 14). Other features described in patients with JEB are generalized failure of dental eruption and rarely resorption of unerupted teeth (15). Although dental resorption has been described in non-syndromic AI hypoplastic type associated with mutations of the *AMELX*, *ENAM*, or *FAM20A* genes (16–18) and syndromic types like enamel renal syndrome (19), to our knowledge, no histopathological and ultrastructural description of dental resorption has been reported in patients with JEB with *LAMB3* gene recessive mutations.

Most cases of non-syndromic hypoplastic AI with coronary resorption have been described in non-erupted teeth and are asymptomatic (20). In the present case, the resorption occurred in a partially erupted molar and was associated with pain. It is most likely that the pain is due to the extreme thinness of the enamel since the dental resorptions are generally painless (21). Additionally, the anterior teeth were sensitive, which disappeared when covering them with crowns.

Although tooth resorptions have been reported in up to 41% of patients with AI (22), the reason for the occurrence of coronary resorptions in teeth with AI is unknown. It has been suggested that it can occur due to an alteration in the surface texture of the enamel (16), or poor recognition of self, causing the body to set up an autoimmune response to the teeth (20).

Wright et al. (13) studied teeth from patients with different forms of EB with SEM technique and found that teeth from

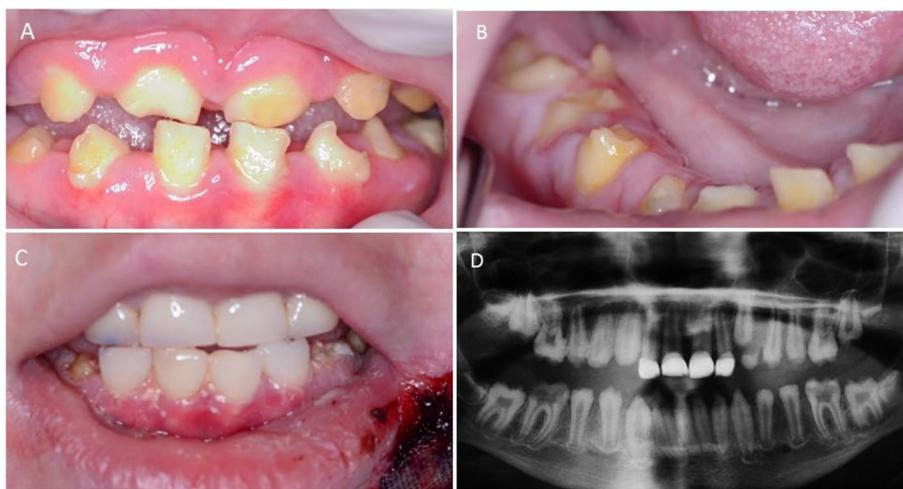


FIGURE 1 | Clinical and radiographic images of a patient with amelogenesis imperfecta (AI) and junctional epidermolysis bullosa (JEB), with *LAMB3* gene mutations. **(A)** Permanent incisors at 10 years of age in were thin, yellow-brown colored enamel is observed. **(B)** Lower right molar with soft tissue over the crown, premolars, and canine have partially erupted. **(C)** Incisors restored with ceramic crowns. **(D)** Panoramic radiograph, almost undetectable, very thin enamel throughout the dentition. Mandibular first molars with coronal radiolucencies and maxillary incisors rehabilitated with ceramic crowns.

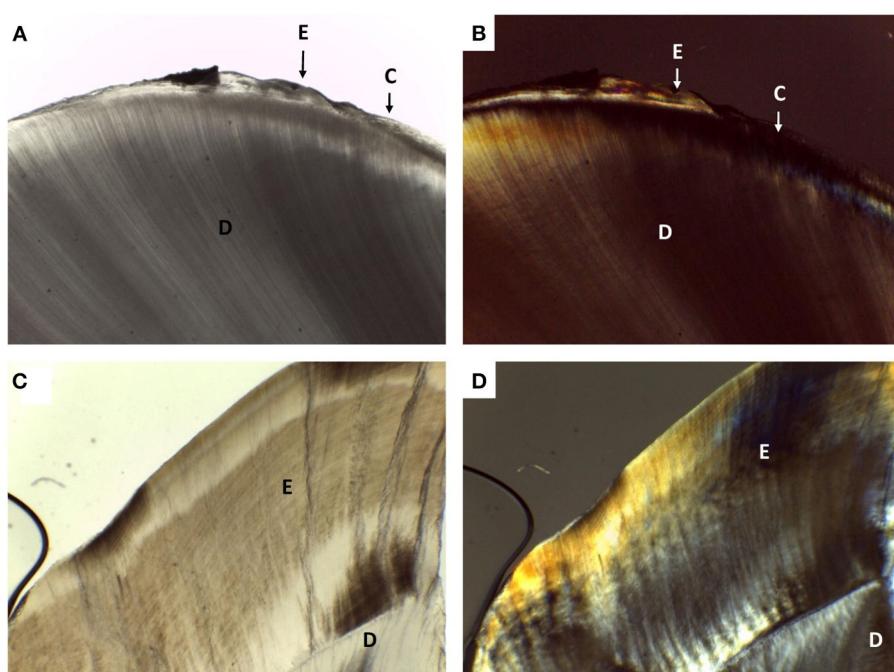


FIGURE 2 | Molar ground section from a patient with AI and JEB visualized with light microscopy. **(A)** Photomicrograph with transmitted light of the coronary area of the permanent molar. **(B)** The same section was studied with polarized light. **(C)** Transmitted light photomicrograph of a permanent molar from a healthy control patient. **(D)** Healthy control molar visualized with polarized light. Original magnification $\times 4$. E, Enamel; D, Dentin; C, Root cement.

patients with the severe, generalized form of JEB (formerly called the Herlitz type) had very thin enamel of $\sim 40\text{ }\mu\text{m}$ thick. These changes were considered a direct consequence of genetic alterations in laminin-332 (13). Currently, it is known that JEB can be caused by mutations in the genes encoding laminin-332, type XVII collagen, and integrins (2). It is unclear whether mutations of these different genes involved in the structure of the

lamina lucida generate ultrastructural differences in the enamel specific to each mutated gene.

In the present study, enamel was observed both in the clinical and microscopic examination as extremely thin or absent in some areas. The zone in which a greater amount of enamel was observed in ground sections was near the cementoenamel junction. Better preservation of enamel in the cementoenamel

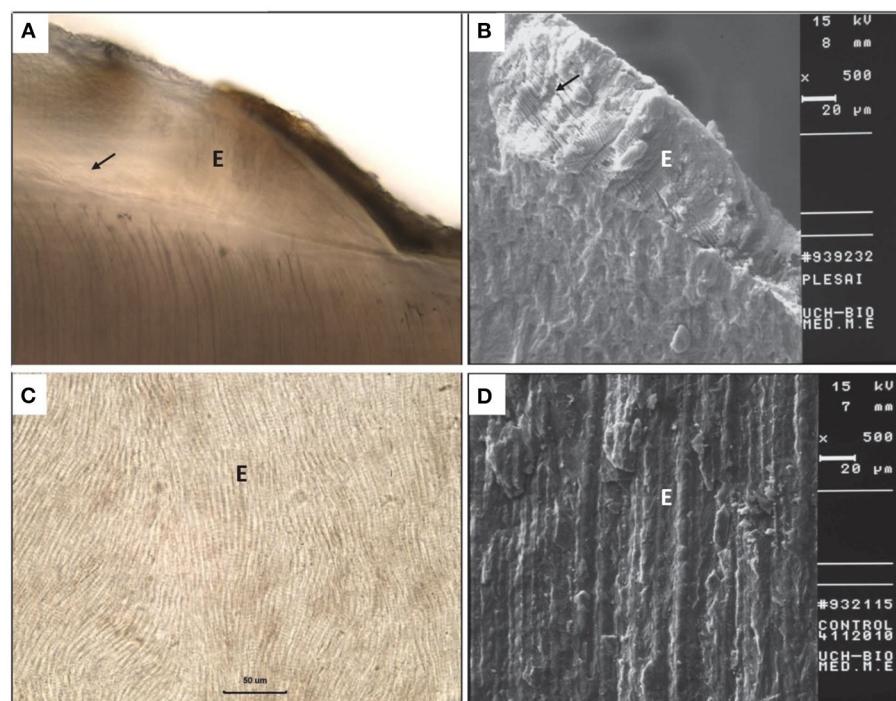


FIGURE 3 | Optical microscopy and scanning electron microscopy (SEM) of a molar from the patient with AI and JEB and molar from a healthy control patient. **(A)** Ground section of a permanent molar from the patient with AI and JEB. In the cervical area, enamel bands semi-parallel to the dentin-enamel junction (black arrow) ($\times 40$) are observed. **(B)** SEM of a molar from the patient with AI and JEB. Cervical enamel, with transverse striations indicated by black arrows ($\times 500$). **(C)** Ground section of a permanent molar from a healthy control patient showing normal enamel prisms ($\times 40$). **(D)** SEM of a normal permanent molar ($\times 500$). E, Enamel.

junction has been reported in other forms of AI (23, 24), and a protective role of the gingival fluid has been suggested.

When analyzing the structure of the affected enamel with SEM, absence of prisms, and the presence of laminar structure in the cervical area, semi-parallel to the surface of the dentin-enamel junction, exhibiting periodicity of $\sim 4 \mu\text{m}$ were observed. In 2009, Antoine et al. (25) recorded a similar frequency of cross-striations in normal human teeth, so these laminar structures could correspond to these striations. To our knowledge, the presence of prominent cross-striations in the enamel has not been previously described in patients with JEB.

During odontogenesis, the three subunits of laminin-332 $\alpha 3$, $\beta 3$, and $\gamma 2$ are expressed in ameloblasts. In the presecretory and maturation stages, they are part of the basal lamina and pre-ameloblasts and disappear at the early bell stage. Therefore, it has been suggested that laminin-332 would play an important role in differentiating ameloblasts (26).

The presence of severe hypoplastic enamel abnormalities as a manifestation of AI should alert dentists, and ideally, mutation analysis of the genes encoding proteins of the lamina lucida should be carried out. Murrell et al. (27) reported a family in which two children died before the first year of life with the development of multiple bullae on the skin. When studying the parents and the older brother of these children, it was found that the parents and the living child had heterozygous mutations in the *COL17A1* gene in addition to the presence of AI. Both parents presented pits on their teeth and the brother presented white spots on the teeth, none of them had skin changes. Thus,

a retrospective diagnosis of JEB in the two deceased children was carried out (27). Moreover, Pasmooy et al. reported a genotype-phenotype correlation analysis performed in 12 patients with JEB who were deficient in collagen XVII. The DNA analysis showed five different deletions. These patients had phenotypes ranging from very mild to severe forms of bullae in the skin and mucous membranes, accompanied by alopecia and nail dystrophy. Despite the different expressions of lesions on the skin, mucous membranes, and nails, AI was observed in all 12 patients (28).

Although the literature has described patients with this type of enamel as having a high tendency to develop caries (7), the patient, in this case, remained without decay until the age of 13, when she passed away. The main dental problems of the patient were tooth sensitivity, aesthetic changes, attrition, and occlusal alterations due to very thin enamel and crown resorption. Almost all the features listed were successfully managed when complete crown treatment was performed at the age of 10.

It is of utmost importance that patients with this condition are looked after by a multidisciplinary medical and dental team, to manage timely each complication that can arise, thus offering them a better oral health-related quality of life. The preventive approach allowed this patient to remain free from decay throughout her life, and rehabilitation based on the crowns solved her tooth sensitivity. Suggestions for the dental management of patients with EB can be found in the Clinical Guidelines published in 2020 (5).

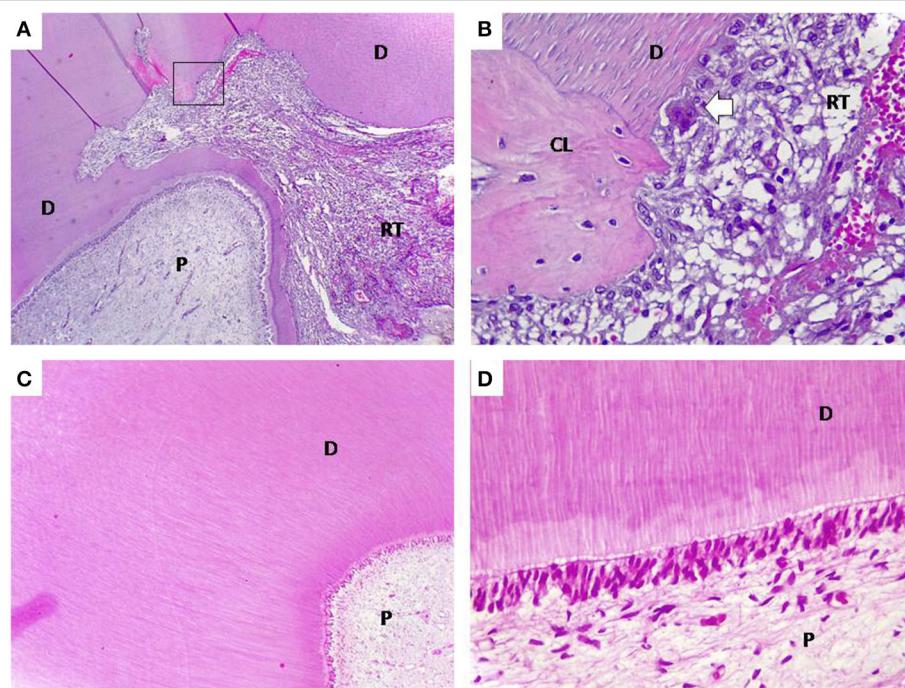


FIGURE 4 | Routine histopathology of a demineralized molar from a patient with AI and JEB and molar from a control patient. **(A)** Dental crown area where pulp (P), dentin (D), and resorptive connective tissue (RT) are observed, (Original magnification $\times 4$). **(B)** Magnification of the area of the rectangle in **(A)**. Odontoclast (white arrow) and cement-like material (CL) (Original magnification $\times 40$). **(C)** Control molar, note the continuity of the dentin around the pulp. (Original magnification $\times 4$). **(D)** Control molar, note the presence and continuity of odontoblasts (Original magnification $\times 40$) (Hematoxylin-eosin staining).

As this is a case report of a very rare condition, its main limitation is that it is not possible to know whether coronary resorption occurs in all patients with AI associated with JEB. We hope that new reports will provide more information and complement the guidelines for the care of these patients.

To conclude, this is the first report of coronary resorption in partially erupted teeth in patients with JEB with mutations in the *LAMB3* gene and hypoplastic AI. In patients with this condition, the presence of partially erupted teeth with soft tissue covering part of the crown, without a periodontal pocket, and with a radiographic image of a partial crown radiolucency should lead to suspicion of coronary resorption. In addition to the skin and mucous membrane bullae, patients with this disorder have teeth defects, such as an extremely thin enamel and absence of prismatic structure. In this patient, conservative treatment based on education, prevention, and restorations with dental crowns reduced tooth sensitivity and improved oral health-related quality of life.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Comité de Ética, School Dentistry, and University

of Chile for the INITIATION FONDECYT project no. 11140440. The study was reviewed and approved by the Ethics Committee at Facultad de Odontología, Universidad de Chile. Written informed consent to present this case description and clinical images was provided by the participant's legal guardian.

AUTHOR CONTRIBUTIONS

AO-P and BU helped with drafting and critical analysis of the general article. SK helped with clinical dental evaluation, prevention and treatment, and analyzed clinical dental data. AO-P and CC helped with histopathological and ultrastructural analysis. BU and IM-B performed the critical analysis of the article regarding amelogenesis imperfecta (AI). MY, FP, and IF performed analysis of *LAMB3* gene mutations. IF performed the critical analysis of the article regarding epidermolysis bullosa (EB). All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fdmed.2021.704423/full#supplementary-material>

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