A Rare Case of Multiple Talon Cusps in Three Siblings

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INTRODUCTION

Talon cusp is a rare entity in which a prominent wisplike structure originates from the cervical area of lingual or labial surfaces of the anterior teeth, occurring lingually with the most frequency (1). Mitchell was the first to recognize this anomaly in 1892 (2), which was later named talon by Ripa and Mellor due to its resemblance to an eagle’s talon (3). It is most often seen in the permanent dentition, predominantly in the lateral maxillary incisor followed by the central maxillary incisor and maxillary canine (1,4,5). Both males and females can be affected, with the former being more involved (4,5).

Hattab et al. (4) classified talon cusps as: type I (talon), a morphologically well-delineated additional cusp that extends at least 50% of the inciso-cervical length of the tooth; type II, (semitalon), a morphologically well-defined additional cusp that extends 25-50% of the tooth length; and type III (trace talon), an enlarged or prominent cingulum, which occupies less than 25% of the distance from the cementoenamel junction to the incisal edge (4). Talon cusp can be accompanied by syndromes such as Berardinelli-Seip (6), Mohr, Rubinstein and Taybi, Ellis-van Creveld, Sturge-Weber and incontinentia pigmenti achromians (7). The co-occurrence of talon cusp and other dental anomalies such as dens in dente and tooth germination has also reported (8,9).

INTRODUCTION

Talon cusp is an uncommon anomaly, whose etiology may be disturbances in the morphodifferentiation stage. Dens in dente is also a rare anomaly that is challenging in clinic as it may cause pulp necrosis or periapical lesions due to the communication with the oral cavity. This article reports multiple talon cusps on permanent maxillary and mandibular incisors and molars in 3 siblings. A 9-year-old boy presented with structures like nodules, shaped as cylindrical cones with a sharp point or a raindrop with deep developmental fissures on the palatal aspect of the maxillary central incisors, lingual aspect of the mandibular central incisors and labial aspect of both mandibular first molars. The buccal surface of the maxillary right central incisor was also affected. Some of them exhibited dens in dente. His 15-year-old sister had prominent talon cusps on the palatal surface of maxillary central incisors and buccal surface of the mandibular first molars and mandibular left second molar. His 7-year-old brother had only one trace talon on the maxillary left central incisor. No syndrome was identified in the patients. In conclusion, genetic inheritance may be a causative factor in talon cusp.

Key Words: case report, talon cusp, sibling, dens in dente.

The aim of the present study was to report a complicated case of a boy with several developmental anomalies along with his two siblings, who presented talon cusps. This is thought to be the first report of the occurrence of multiple talon cusps in three siblings.
CASE REPORT

A 9-year-old boy was referred to the Department of Pediatric Dentistry of Mashhad Dental School. His medical history was uneventful. Oral hygiene was poor with multiple carious lesions. The chief complaint was the presence of a fistula with a history of 7-8 months on the buccal aspect of the maxillary left central incisor. The tooth had undergone an emergency treatment and the access cavity was sealed with a temporary filling material. There were structures like nodules, shaped as cylindrical cones with a sharp point or a raindrop, with deep developmental fissures on the palatal aspect of both maxillary central incisors. The buccal surface of the maxillary right central incisor was also affected (Fig. 1). Hence, talons were found on both labial and lingual surfaces of this tooth. Same structures were found on the lingual surface of mandibular central incisors and the labial surface of both mandibular first molars (Fig. 2). Radiographs showed dens in dente in the maxillary central incisors and left lateral incisor (Fig. 3). Periapical radiolucency was observed in the maxillary left incisor. Oral examination of his parents was not contributory, but his two siblings presented similar structures.

His 15-year-old sister had prominent talon cusps on the palatal surface of the maxillary central incisors and buccal surface of the mandibular first molars and left second molar (Fig. 4). His 7-year-old brother had only one trace talon on the maxillary left central incisor (Fig. 5), but he still had several unerupted teeth. Due to
multiple teeth with carious lesions and pulp necrosis, all siblings were referred to the Departments of Endodontics and Pediatric Dentistry for appropriate endodontic, restorative and preventive treatments.

**DISCUSSION**

The accessory cusp-like structure, talon cusp, originates from disturbances in the morphodifferentiation stage of tooth development, as does any other anomaly in the size or shape of the tooth (14). The real cause of this condition is still unknown, but genetic and environmental factors are thought to be involved in the etiology. Sporadic occurrence of this abnormality is not uncommon (1,7).

_Dens in dente_ results from the invagination of the affected tooth crown before the mineralization (10,15). It is usually detected in routine radiographs. Dealing with a _dens in dente_ is challenging in clinic as it may cause pulp necrosis or periapical lesions due to the communication with the oral cavity and influx of the irritants to the pulp (10,12).

There are very few reports of multiple siblings affected by talon cusps. Meon reported talon cusps in a 10-year-old boy and his 12-year-old sister (16). Segura and Jimenez-Rubio reported 2 cases of talon cusp affecting consanguineous first cousins, and affirmed that family involvement and the association of the talon cusp with other dental abnormalities suggest genetics as a probable major causative factor (17). Hattab et al. (18) and Solanki et al. (6) reported siblings with talon cusps

![Figure 3. Periapical radiograph of maxillary incisors. _Dens in dente_ of both maxillary central incisors and left lateral incisor is evident. Due to the access cavity preparation, the _dens in dente_ of the left central incisor is not clear. Note the radiopacity of buccal talon cusp on right central incisor (Case 1).](image)

![Figure 4. Intraoral clinical images. A: Prominent talon cusps on the maxillary central incisors. B: Talon cusps on the mandibular first molars and mandibular left second molar (Case 2).](image)

![Figure 5. Intraoral clinical image. A trace talon cusp on the maxillary right central incisor. (Case 3).](image)
but these patients were carriers of Ellis-van Creveld and Berardinelli-Seip syndromes, respectively. Unlike these cases (6,18), none of our patients presented talon cusp associated with any known systemic syndrome.

The talon cusp may cause clinical problems related to caries or occlusal interferences (7). Treatment objectives for talon cusps are aimed at the management of deep caries-susceptible pits and fissures, occlusal interferences, esthetic problems, soft tissue irritation, interference with the tongue space, and pulpal involvement (14). The treatment options for the mentioned problems are dependent upon the presence of a pulp horn in the cusp and include: placement of pit-and-fissure sealants or restorations, periodic gradual occlusal grinding, root canal therapy and extractions (1).

Mandibular talon cusps are less common than maxillary ones (8,19). Talon cusp on the occlusal surface is subjected to direct occlusal forces, causing attrition or fracture at a much higher rate than the talon cusp on lingual surface (20). Early diagnosis and treatment of talon cusps is strongly recommended.

Due to the presence of a fistula in the open apex maxillary left central incisor, the first patient was referred for endodontic treatment of the tooth (root canal therapy with an apical MTA plug). Other teeth of this patient and the other cases underwent restorative and preventive therapies. Early oral examination could facilitate early diagnosis of the talon cusp and prevent occlusal problems and dental caries.

Based on this report and other similar anomalies in the family, it may be suggested that genetic inheritance may be a causative factor. However, genetic research needs to be carried out in order to prove this statement.

REFERENCES


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