Hyperplastic Dental Follicle – Review of Literature and Report of Two Cases in One Family

Chun Xiao SUN¹, Craig RIRIE¹, Jeffrey M HENKIN¹

Hyperplastic dental follicle is an extremely rare lesion. The practitioner should be able to differentiate it from a dentigerous cyst. The present article will review related literature and report on two cases in one family. A 12-year-old white female and her 15-year-old brother were referred for exposure of unerupted canines. No systemic diseases or syndromes were present. Intra-oral examinations were unremarkable, except for the absence of all eight canines. Radiographic examinations revealed impacted canines with each unerupted tooth surrounded by a well-demarcated radiolucency, which passed beyond the cementoenamel junction. The teeth were surgically exposed and tissue specimens surrounding the unerupted teeth were analysed histologically. Histology revealed fibrous connective tissue with areas demonstrating some ground substance and multiple odontogenic epithelial rests. Some surfaces were partially lined by reduced enamel epithelium. A diagnosis of hyperplastic dental follicle was made. Sometimes, it is difficult to differentiate hyperplastic dental follicle from odontogenic fibroma, both simple and central forms. A correct diagnosis should be based on clinical, radiographic and histological findings.

Key words: hyperplastic dental follicle, unerupted canine

Hyperplastic dental follicle or peri-follicle fibrosis, is an unusual lesion often confused with odontogenic fibroma¹,². Gardner was the first one who attempted to clarify these similar lesions¹. He pointed out that three different lesions may resemble each other. The first one is hyperplastic dental follicle, which is mainly composed of fibrous connective tissue and wavy collagen fibres. It may contain strands and rests of odontogenic epithelium. Calcified materials may be present in the connective tissue, which is then called calcifying hyperplastic dental follicle, and was first described by Sandler et al².

The second form of the lesion has been called simple odontogenic fibroma. This lesion resembles hyperplastic dental follicle histologically, but differs in its size and location. The simple odontogenic fibroma is usually much larger than hyperplastic dental follicle, and is invariably associated with crowns of unerupted teeth, which is not always the case with odontogenic fibroma. The third lesion has been named central odontogenic fibroma (WHO type). This lesion has some unique histological features such as dysplastic dentin, cementum-like calcified tissues and marked epithelial proliferation. The last two lesions are considered true neoplasms, whereas the hyperplastic dental follicle is considered a hamartomatous lesion.

Multiple hyperplastic dental follicles is an extremely rare condition. To date, there are only six cases reported in the literature (Table 1)²-⁶. None of these reported any symptoms, and they mainly affect unerupted posterior teeth, which accounts for 30 out of 36 teeth involved. All

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### Table 1  Reported cases of multiple hyperplastic dental follicles.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Patients</th>
<th>Teeth affected</th>
<th>Defective affected teeth</th>
<th>Calcification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sun, et al (current report)</td>
<td>15-year-old white male</td>
<td>6, 11, 22, 27</td>
<td>Not mentioned</td>
<td>No</td>
</tr>
<tr>
<td></td>
<td>12-year-old white female</td>
<td>6, 11, 22, 27</td>
<td>Not mentioned</td>
<td>Foci of irregular calcification</td>
</tr>
<tr>
<td>Walker, et al (reference 6)*</td>
<td>6-year-old female</td>
<td>T</td>
<td>Not mentioned</td>
<td>Type A and B calcification in fibrous soft tissue and dystrophic calcification</td>
</tr>
<tr>
<td>Gomez, et al (reference 4)</td>
<td>15-year-old male</td>
<td>1, 2, 5, 6, 7, 11, 12, 15, 22, 27, 32</td>
<td>Unremarkable</td>
<td>Type A and B calcification in fibrous soft tissue and dystrophic calcification in the pulp tissue and dentine matrix</td>
</tr>
<tr>
<td>Gardner, et al (reference 5)</td>
<td>26-year-old white male</td>
<td>17, 18</td>
<td>Not mentioned</td>
<td>Type A calcification dominant</td>
</tr>
<tr>
<td></td>
<td>40-year-old male</td>
<td>7 teeth</td>
<td>Hypoplastic enamel and defective crowns and roots</td>
<td>Basophilic, concentric, calcified bodies, type A calcification</td>
</tr>
<tr>
<td>Lukinmaa, et al (reference 3)</td>
<td>24-year-old male</td>
<td>18, 20</td>
<td>Hypoplastic enamel and defective crowns and roots</td>
<td>Basophilic, concentric, calcified bodies, type A calcification</td>
</tr>
<tr>
<td>Sandler, et al (reference 2)</td>
<td>15-year-old black male</td>
<td>1, 2, 13, 15, 16, 17, 18, 20, 21, 28, 29, 31, 32</td>
<td>Unremarkable</td>
<td>Abundant calcified materials in the whorled areas of the connective tissue and in the hyalinised tissue surrounding the odontogenic epithelium</td>
</tr>
</tbody>
</table>

* The diagnosis was hyperplastic dental follicle with an incipient adenomatoid odontogenic tumour

**Fig 1** Panoramic radiographs of the 12-year-old female patient 6 months after tooth 13 extraction. Radiolucency around each unerupted canine was well demarcated.

**Fig 2** Radiographs showed radiolucencies passed beyond the cementoenamel junction (CEJ). Periapical radiographs of teeth 22 (right) and 27 (left) are shown. Radiolucencies passed beyond the CEJ and no obvious resorption was noticed on adjacent teeth.
of these cases presented with calcification. In the present report, the authors present two cases; each has four unerupted canines associated with non-calcified hyperplastic dental follicles, and occur in the same family.

Case reports

In January 2009, a 12-year-old Caucasian female was referred to the Department of Periodontics, Loma Linda University School of Dentistry for exposure of unerupted teeth 6, 11, 22 and 27. The patient had unerupted tooth 13 extracted by an oral surgeon 6 months prior to the present consultation. There were no associated systemic diseases or syndromes present. Intraoral examination revealed that teeth 6, 11, 22 and 27 were not clinically present, and the patient was undergoing orthodontic treatment. Swelling on the buccal sides of teeth 22 and 27, depressible on palpation, were detected. A panoramic radiograph showed a unilocular, well-circumscribed radiolucency around the crowns of each impacted tooth (Fig 1). Teeth 1 and 16 were still forming while teeth 17 and 32 appeared to be congenitally missing. Periapical radiographs (Fig 2) and cone beam 3-D images (data not shown) showed that the radiolucencies were beyond the cementoenamel junction.

Impacted teeth (6, 11, 22 and 27) were surgically exposed via a closed flap technique to preserve keratinised tissue, and abundant dense fibrotic tissues were found surrounding the impacted teeth, extending into the areas of the roots (Fig 3). The tissues around the impacted cuspsids were collected and sent to a private laboratory for pathological analysis. Under light microscopy, the pathologist found considerable ground substance and multiple odontogenic epithelial rests, with some exhibiting juxta-epithelial hyalinisation. The histological diagnosis was hyperplastic dental follicles (peri-follicular fibrosis) without calcification (Fig 4).

Three months later, the patient’s 15-year-old brother was referred to the department of periodontics for the same reason. This patient had almost identical clinical findings as his younger sister, except that all of his third molars were forming. Following his surgery, the pathological diagnosis was also reported as hyperplastic dental follicle (Fig 5).

Discussion

The most frequently impacted teeth are mandibular third molars, followed by maxillary canines, mandibular canines, premolars and incisors. The incidence of maxillary canine impaction ranges from 1 to 3%, and palatal impaction is more often observed than labial impaction.
(the ratio is approximately 2:1 to 6.6:1). The aetiology of tooth impaction is still not very clear. Possible primary causes are the rate of root resorption of deciduous teeth, trauma of the deciduous tooth bud, disturbances in tooth eruption sequence, availability of space in the arch, rotation of tooth buds, premature root closure and canine eruption into the cleft area in persons with cleft palate. The proposed secondary causes are abnormal muscle pressure, febrile disease, endocrine disturbances and vitamin D deficiency.

The normal pericoronal radiolucency is considered to be in the range of 2 to 3 mm, and there is no definitive explanation of the mechanism causing hyperplastic dental follicle. Kim et al reported that the expression of several matrix metalloproteinases is down-regulated in hyperplastic dental follicles; moreover, several collagen genes are up-regulated. Previous reports have described only male patients (Table 1). The present report is the first case report involving permanent teeth in a female patient. This is also the first report of hyperplastic dental follicle involved in only canines. To the authors’ knowledge, this report is also the first one to report two cases in siblings, which suggests that the condition may have a genetic predilection.

Tooth impaction caused by hyperplastic dental follicle may be associated with defective tooth formation, and it also may be associated with other symptoms, such as amelogenesis imperfecta, rough hypoplastic type and gingival hyperplasia. It is very important for dentists to recognise these possible associated lesions.

Hyperplastic dental follicle may involve just one tooth or may be present in multiple teeth. Sometimes, it is difficult to differentiate hyperplastic dental follicle from odontogenic tumours, especially when calcification is present. Calcification was estimated to be present in one-third of dental follicles. The final diagnosis must consider the clinical, radiographic and histological features of the lesion.

Acknowledgements

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Fig 4. Histological observation with hematoxylin and eosin (H&E) staining. Abundant fibrous collagen tissue interspersed with some ground substances (arrowheads in a), and odontogenic epithelial rests (arrows in a). Some surfaces were partially lined by reduced enamel epithelium (red arrows in b). No calcification was found in either section. Original magnification ×20.

Fig 5. The radiographs and histology of the 15-year-old male patient. This patient had almost identical lesions as his sister described in Figures 1 to 3. Panoramic radiograph (a), 1 year after extraction of deciduous teeth in impacted canine positions. Histology (b) with H&E staining at magnification ×20.

References