Prevalence of dental abnormalities in different calcium metabolism disorders in a group of Iranian children.

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Abstract

Aim: Calcium metabolism disorders trigger to enhanced level (hypercalcemia) or lower level (hypocalcemia) of calcium. These alterations in calcium level leads to accumulation of poorly mineralized bone and dental abnormalities. The aim of this study was to assess the prevalence of dental abnormalities including taurodontism, enamel hypoplasia, dental abscess, dental caries, delayed teething, gingivitis in children with calcium metabolism disorders.

Design: Patients, who were suffering from calcium metabolism disorders, including renal osteodystrophy, hypoparathyroidism, and different types of rickets, participated in this study. All of them transferred into dental clinic of Children’s Medical Center to pass dental exams and preparing OPG images if it was necessary. Also, prior patient’s files were used; their demographic information and dental problems were recorded on designed checklists.

Results: We found that dental caries and delayed teething are the dominant abnormalities that are prevalent among patients with nutritional rickets and renal osteodystrophy.

Conclusions: According to the results these disorders are associated with vitamin D deficiency; it seems that treatment with supplements of vitamin D might be efficient method to prevent dental abnormalities.

Keywords: Calcium metabolism disorder, Nutritional rickets, Dental caries, Vitamin D, Children.

Introduction

In a healthy physiological condition, the actions of parathyroid hormone, vitamin D and the calcium sensing receptor maintain extracellular calcium levels within a tight range [1]. Disorders in calcium metabolism can lead to hypercalcemia, elevated plasma calcium levels or hypocalcemia, decreased plasma levels of calcium. Hypercalcemia has non-specific symptoms that if not recognized early could trigger patient deterioration and organ failure [2]. Serum calcium levels more than 12 mg/dL can lead to multiple organ dysfunctions and if its level exceeds 20 mg/dL, can even be fatal [2]. One of the responsible factors for hypercalcemia is decreased renal excretion of calcium which occurs through decreased glomerular filtration and increased renal tubular absorption of calcium [2,3].

On the other hand, other factors such as deficiency in active vitamin D or magnesium, or decreased secretion or action of Parathyroid Hormone (PTH) may result in hypocalcemia [4]. Total serum calcium concentration less than 7.5 mg/dL in neonates and below 8.7 mg/dL in children are defined as hypocalcemia [5]. Patients with renal insufficiency and those with extensive osteoblastic bone metastases have shown enhanced risk for hypocalcemia [6,7].

Hypoparathyroidism (HPT) is a disease caused by insufficient secretion of Parathyroid Hormone (PTH) and with unknown etiology. The reduced secretion of this hormone trigger to the deficiency in the synthesis of 1, 25-dihydroxyvitamin D and inhibition of calcium reabsorption in distal tubules and bone matrix which leads to hypocalcemia and hyperphosphatemia. Patients suffered with hypoparathyroidism commonly show cataracts, hair loss and dental abnormalities [8,9].

Hypophosphatemia, another disease associated with calcium metabolism, is caused by decreased intestinal absorption (such as phosphate binding antacids, malabsorption, and vitamin D deficiency) and excessive urinary excretion (including hyperparathyroidism, vitamin D deficiency, oncogenic osteomalacia, and familial hypophosphatemic rickets) [10-15].
Familial hypophosphatemic rickets is the common form of rickets that is inheritable and is characterized by features such as inappropriately low levels of serum concentration of 1, 25-dihydroxyvitamin D, renal phosphate wasting, short stature and radiographic evidence of rickets.

Although the autosomal dominant hypophosphatemic rickets has been characterized, it is usually transmitted as an X-linked dominant disorder [14]. Autosomal form of familial hypophosphatemic rickets is associated with hypercalciuria, a rare genetic disorder that is caused by a mutation in a gene that encodes sodium-phosphate transporter [16]. Recently, it has been suggested, in mutant forms, osteoblasts and odontoblasts secrete a protein that enters the circulation and acts on the kidney [17]. This abnormality leads to incoherent regulation between the production and the rate of mineralization of bone matrix, and subsequently the accumulation of poorly mineralized bone [18].

Progressive renal damages lead to impairment of the excretion of body metabolic waste products and also, affect the endocrine functions of the kidney (secretion of vitamin D and erythropoietin) [19,20]. Renal phosphate wasting can be divided into three types based on the serum calcium level: primary hyperparathyroidism (high serum calcium level), secondary hyperparathyroidism (low serum calcium level), and primary renal phosphate wasting (normal serum calcium level) [21]. Patients with renal failure represent various forms of oral and dental manifestation which of them gingival hyperplasia is one of the most widely documented [22].

The oral and dental manifestations of the calcium metabolism disorders such as hypophosphatemic vitamin D resistant rickets have been well described in the literature [23-25]. Enamel and dentin defects, usually enamel hypoplasia, interglobular dentinal masses with variable degrees of calcification, large pulp chambers, enlarged pulp horns associated with thin enamel, dentin defects, defects at the dentin enamel junction, and fissures that may extend from the enamel to the pulp are oral consequences calcium metabolism deficiency [23-26]. Due to the close relation between calcium metabolism disorders and oral manifestations, in this study we aimed to investigate dental abnormalities in children with calcium metabolism disorders including hypoparathyroidism, nutritional rickets, hypophosphatemic rickets, renal osteodystrophy and osteogenesis imperfecta. The result of this study could be useful to prevent dental abnormalities or to find therapeutic solution for them.

Materials and Methods

Patients

This cross-sectional study was performed on 100 patients who were suffering from calcium metabolism disorders and admitted at nephrology section, between 2005 and 2009, in Children’s Medical Center, Tehran, Iran.

Inclusion and exclusion criteria

Patients that had a laboratory record of at least one total serum calcium level evaluated at any time during their illness met the inclusion criteria of our study group.

Performance

Patients, who were suffering from calcium metabolism disorders, including renal osteodystrophy, hypoparathyroidism and different types of rickets, participated in this study. All of them transferred into dental clinic of Children’s Medical Center to pass dental exams and preparing OPG images if it was necessary. Also, prior patient’s files were used; their demographic information and dental problems were recorded on designed checklists.

Ethical consideration

The study was carried out in accordance with the Declaration of Helsinki, the ethics committee of the Tehran University of Medical Sciences approved the protocols of the study.

Data analysis

Qualitative data were analyzed and reported as frequency (%). All data were analyzed with IBM SPSS Statistics for Windows, version 21.0 (SPSS Inc., Chicago, Illinois, USA).

Results

As demography of patients has been presented in Table 1, the number of males is close to that of females. Also, the average of affected teeth has increased with age. Most of the patients were suffering from nutritional rickets and least of them from osteogenesis imperfecta. In addition, general oral health care was determined well.

Table 1. Demographic feature of patients with calcium metabolism disorders.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>42</td>
</tr>
<tr>
<td>Female</td>
<td>58</td>
</tr>
<tr>
<td>Average of affected teeth based on age</td>
<td></td>
</tr>
<tr>
<td>&lt;5</td>
<td>2.18</td>
</tr>
<tr>
<td>42865</td>
<td>2.88</td>
</tr>
<tr>
<td>&gt;10</td>
<td>4.53</td>
</tr>
<tr>
<td>Disease</td>
<td></td>
</tr>
<tr>
<td>HPT</td>
<td>12</td>
</tr>
<tr>
<td>NR</td>
<td>42</td>
</tr>
<tr>
<td>HR</td>
<td>19</td>
</tr>
<tr>
<td>RD</td>
<td>20</td>
</tr>
</tbody>
</table>
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Oral health status

<table>
<thead>
<tr>
<th>Status</th>
<th>Count</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well</td>
<td>77</td>
</tr>
<tr>
<td>Bad</td>
<td>23</td>
</tr>
</tbody>
</table>

HPT: Hypoparathyroidism; NR: Nutritional Rickets; HR: Hypophosphatemia Rickets; RO: Renal Osteodystrophy; OI: Osteogenesis Imperfecta.

The number and percentage of dental abnormalities have been presented in Table 2. Gingivitis was at least frequency and its highest number was observed among patients with renal osteodystrophy with 20%. Taurodontism and dental abscess were not prevalent among patients and the highest abundance was observed among hypoparathyroidism group with 33.3% and 41.7% respectively. Enamel hypoplasia and delayed teething were more prevalent among patients with osteogenesis imperfecta and nutritional rickets, respectively. Dental caries was observed in 65% of patients which its occurrence was observed in 90% of patients with renal osteodystrophy.

Dental caries and delayed teething are more prevalent abnormalities in this study, and occurrence of them was more common among patients with nutritional rickets (Figure 1).

Table 2. Representation of dental abnormalities in relation with different types of calcium metabolism disorder.

<table>
<thead>
<tr>
<th></th>
<th>Hypoparathyroidism</th>
<th>Nutritional rickets</th>
<th>Hypophosphatemic rickets</th>
<th>Renal osteodystrophy</th>
<th>Osteogenesis imperfecta</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Taurodontism</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Negative</td>
<td>8 (66.7)</td>
<td>39 (92.9)</td>
<td>16 (84.2)</td>
<td>15 (75)</td>
<td>6 (85.7)</td>
<td>84 (84)</td>
</tr>
<tr>
<td>Positive</td>
<td>4 (33.3)</td>
<td>3 (7.1)</td>
<td>3 (15.8)</td>
<td>5 (25)</td>
<td>1 (14.3)</td>
<td>16 (16)</td>
</tr>
<tr>
<td>Total</td>
<td>12 (100)</td>
<td>42 (100)</td>
<td>19 (100)</td>
<td>20 (100)</td>
<td>7 (100)</td>
<td>100 (100)</td>
</tr>
<tr>
<td>Enamel hypoplasia</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Negative</td>
<td>6 (50)</td>
<td>38 (90.5)</td>
<td>11 (57.9)</td>
<td>11 (55)</td>
<td>3 (42.9)</td>
<td>69 (69)</td>
</tr>
<tr>
<td>Positive</td>
<td>6 (50)</td>
<td>4 (9.5)</td>
<td>8 (42.1)</td>
<td>9 (45)</td>
<td>4 (57.1)</td>
<td>31 (31)</td>
</tr>
<tr>
<td>Total</td>
<td>12 (100)</td>
<td>42 (100)</td>
<td>19 (100)</td>
<td>20 (100)</td>
<td>7 (100)</td>
<td>100 (100)</td>
</tr>
<tr>
<td>Dental abscess</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Negative</td>
<td>7 (58.3)</td>
<td>41 (97.6)</td>
<td>17 (89.5)</td>
<td>10 (50)</td>
<td>7 (100)</td>
<td>82 (82)</td>
</tr>
<tr>
<td>Positive</td>
<td>5 (41.7)</td>
<td>1 (2.4)</td>
<td>2 (10.5)</td>
<td>10 (50)</td>
<td>0 (0)</td>
<td>18 (18)</td>
</tr>
<tr>
<td>Total</td>
<td>12 (100)</td>
<td>42 (100)</td>
<td>19 (100)</td>
<td>20 (100)</td>
<td>7 (100)</td>
<td>100 (100)</td>
</tr>
<tr>
<td>Dental caries</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Negative</td>
<td>6 (50)</td>
<td>15 (35.7)</td>
<td>10 (52.6)</td>
<td>2 (10)</td>
<td>2 (28.6)</td>
<td>35 (35)</td>
</tr>
<tr>
<td>Positive</td>
<td>6 (50)</td>
<td>27 (64.3)</td>
<td>9 (47.4)</td>
<td>18 (90)</td>
<td>5 (71.4)</td>
<td>65 (65)</td>
</tr>
<tr>
<td>Total</td>
<td>12 (100)</td>
<td>42 (100)</td>
<td>19 (100)</td>
<td>20 (100)</td>
<td>7 (100)</td>
<td>100 (100)</td>
</tr>
<tr>
<td>Delayed teething</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Negative</td>
<td>8 (66.7)</td>
<td>16 (38.1)</td>
<td>10 (52.6)</td>
<td>15 (75)</td>
<td>5 (71.4)</td>
<td>54 (54)</td>
</tr>
<tr>
<td>Positive</td>
<td>4 (33.3)</td>
<td>26 (61.9)</td>
<td>9 (47.4)</td>
<td>5 (25)</td>
<td>2 (28.6)</td>
<td>46 (46)</td>
</tr>
<tr>
<td>Total</td>
<td>12 (100)</td>
<td>42 (100)</td>
<td>19 (100)</td>
<td>20 (100)</td>
<td>7 (100)</td>
<td>100 (100)</td>
</tr>
</tbody>
</table>

Note: Data are presented as frequency (%).
Discussion

Hypercalcemia, as a calcium metabolism disorder, is the most common life-threatening metabolic complication and its prevalence is about 5%-20% in the adult population. In paediatric cancers it is equally as life-threatening but not as common, but must be corrected especially prior to induction chemotherapy [2]. Also, acute leukaemia, which is the most common paediatric cancer, almost associated with hypercalcemia [27]. Also defects in the secretion or action of parathyroid hormone result in hypocalcemia and a relative inability to excrete phosphorus [28].

Nutritional rickets is mainly due to deficiencies of sunlight, malabsorption of vitamin D, failure to activate vitamin D or unresponsiveness tissue to vitamin D that can predispose the child to hypocalcemia. Since vitamin D is the initial regulator of phosphorus homeostasis, insufficient vitamin D synthesis or action, that cause to hypocalcemia, is usually accompanied by a decreased serum phosphorus concentration [28,29]. According to the results of present study showed that 42% of patients were suffering from nutritional rickets which was the majority among patients. Of these patients 64.3% and 61.9% were positive to dental caries and delayed teething which indicated to the relation between nutritional rickets and dental anomalies and the need for more attention children nourishment. Hard dental tissue alterations, mainly enamel hypoplasia and dentin defects, has been ascribed to Vitamin D insufficiency should benefit from oral 1, 25-dihydroxyvitamin D supplements [37]. Also, prophylactic antibiotics have been recommended prior every invasive dental protocol [33]. In addition, it has been suggested that patients with renal failure require special considerations in relation to dental treatment, not only because of the conditions inherent to the disease and its multiple oral manifestations, but also because of the side effects and characteristics of the treatments they receive [33].

Hypophosphatemic rickets presents with significantly depressed phosphate levels and normal calcium levels, whereas 1, 25 (OH) 2 vitamin D levels are low or in the low normal range [25]. In the present study, dental abnormalities such as dental caries and delayed teething, same as nutritional rickets and renal osteodystrophy, were prevalent. For treatment of hypophosphatemic rickets, vitamin D and hydroxylated forms of vitamin D have been used [38]. Vitamin D is intimately involved in a wide variety of biologic processes, including calcium homeostasis, bone formation, and cellular differentiation [39,40].

Conclusions

In conclusion, our results showed dental caries and delayed teething as more dominant dental abnormalities among children with calcium metabolism disorders. Also, nutritional rickets and renal osteodystrophy were more prevalent disorders which both of them are associated with vitamin D deficiencies. Thus, as aforementioned, addition of vitamin D supplements to children’s diet could be an effective solution to prevent dental abnormalities.

References


Figure 1. Frequency of dental abnormalities in different calcium metabolism disorders. HPT: Hypo Parathyroidism; NR: Nutritional Rickets; HR: Hypophosphatemia Rickets; RO: Renal Osteodisthrophia; OI: Osteogenes Imperfecta.
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