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<td><strong>Author(s)</strong></td>
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<td><strong>Citation</strong></td>
<td>Journal of Dentistry: Oral Health &amp; Cosmesis, 2016, v. 1 n. 1, article no. 003</td>
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<tr>
<td><strong>Issued Date</strong></td>
<td>2016</td>
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<tr>
<td><strong>URL</strong></td>
<td><a href="http://hdl.handle.net/10722/224879">http://hdl.handle.net/10722/224879</a></td>
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Williams Syndrome - Review of Clinical Features and It’s Medical Considerations in Dental Treatment

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Abstract

Williams syndrome is a congenital disorder that involves multiple system. The condition is characterized by a number of developmental and physical abnormalities. The cause of this disorder is a micro deletion from the long arm of chromosome 7 at 7q11.23. The deleted portion of the chromosome 7q11.23 includes the ELN gene that codes for the structural protein elastin and therefore in the affected individuals arteries may become narrowed. Hypocalcaemia is also found in the affected individuals. Typical facial, oral, and dental features are characteristic of this syndrome. Presence of congenital heart defects and the risk of developing cardiac arrest make dental treatment challenging for such patients. This article narrates the etiology, pathogenesis, and clinical features including oral features of patients with Williams syndrome. A note on role of dental evaluation in diagnosis of Williams syndrome and medical consideration in the dental treatment of patients with Williams syndrome has also been presented.

Keywords: Dental features; Dental treatment; Oral features; Review; William syndrome

Introduction

Williams Syndrome (WS) was described by JCP Williams and AJ Beuren. This condition is also known by other names such as “Williams-Beuren syndrome”, “elfin-facies syndrome” and “idiopathic hypercalcemia-supravalvular aortic stenosis syndrome”. This is a congenital disorder characterized by several developmental and physical abnormalities [1-3]. The prevalence of this condition is 1 in 7,500 individuals [4]. Individuals affected with Williams syndrome has typical facial, and dental features that are relevant to dentists. Additionally, due to multisystem disorders including congenital cardiac defects affected individuals pose a challenge for dental treatment. This article narrates the clinical features including oral, dental features and the significance of dental evaluation in the diagnosis of patients with Williams syndrome. The article also presents medical considerations in the dental treatment of patients with Williams syndrome.

Etiology

Williams Syndrome (WS) is caused by a micro deletion from the long arm of chromosome 7 at 7q11.23 [3]. The disorder is sporadic. However, an autosomal dominant inheritance pattern has also been established in which the affected individuals have a 50% chance of transmitting the disorder to their offspring. The disorder has no gender or ethnic predilection [1].

Pathogenesis

The deleted portion of the chromosome 7q11.23 includes the ELN gene that codes for the structural protein elastin, an essential component of the elastic fibres present in the connective tissue of several organs. The deletion of elastin explains few features of WS, such as part of the facial features, hoarse voice, bladder and bowel diverticula, cardiovascular disease, and orthopaedic problems. The pathogenesis of other features, such as hypercalcemia, mental retardation, and distinctive personality traits, are still unexplained. One possibility is that the loss of 1 or more genes contiguous to the ELN gene contributes to the phenotype. Since the elastin protein is an essential component of elastic fibres in the arterial wall, arteries (in general) become narrowed. Idiopathic infantile hypercalcemia is a feature of WS that can lead to severe irritability, vomiting, constipation, and muscle cramps associated with this condition [5]. Symptomatic hypercalcemia commonly does not manifest beyond childhood, but lifelong abnormalities of calcium and vitamin D metabolism may continue to exist. Hypercalciuria is common and predisposes to nephrocalcinosis. The reasons of the abnormality in calcium metabolism are unknown.

Clinical Features

WS is a multisystem disorder. The cardiovascular system of affected individuals presents with supravalvular aortic stenosis, pulmonary artery stenosis, atrial septal defect, ventricular septal defect, aplasia of the portal vein, arterial hypertension and heart murmur. The affected individuals have mental deficiency with IQ between 41 and 80. Patients with this syndrome have typical facial features described as “Elfin face features” with reduced circumference of skull, craniosynostosis, broad forehead, short palpebral fissures, epicanthal folds, medial eyebrow flare, convergent strabismus, depressed nasal bridge, stellate pattern in iris, long philtrum, anteverted nostrils, wide mouth (Figure 1), drooping lower lip, hypoplastic mandible [6-8].

Hypercalciemias not uncommon in the affected individuals and when present appears until the second year of life causing hypotonia and anorexia. The skeletal abnormalities include, growth deficiency (prenatal and postnatal), craniosynostosis (secondary to microcephaly), retarded bone age, pectusexcavatum, hallux valgus and fifth finger clinodactyly [9]. Abnormalities of the renal system include...
Williams Syndrome (WS) is associated with an abnormal growth pattern. The incidence of pre-natal growth deficiency is approximately 50-70% [5,11]. Feeding problems, prolonged colic, gastroesophageal reflux, and constipation lead to failure to thrive in 80% of infants with WS [12]. Until age four, weight gain and linear growth are poor. During childhood the rate of linear growth is 75% of normal [13]. In mid-childhood besides improvement in growth, 70% remain below the third percentile for mid parental height and are thus shorter than the children of similar age from their genetic background. Puberty often occurs early and is associated with a shorter growth spurt. These combined factors leads to an average final adult height that is below the third percentile [11].

Supravalvular Aortic Stenosis in WS

Supravalvular Aortic Stenosis (SVAS) is a rare but significantly characterized form of left ventricular outflow obstruction. The lesion involves the ascending aorta and may frequently be observed with associated pulmonary arterial stenosis or stenosis of other arteries, particularly at major branch points [14]. The onset and severity of disease varies but, if left untreated, may lead to heart failure, myocardial infarction, and death. Surgical treatment is the only option in symptomatic cases. Pathological studies of patients with SVAS recognized abnormalities in elastin structure and function. A quantitative deficiency of elastic fibres was also noted, resulting in an excess of smooth muscle wall and bundles in the diseased media when compared with the normal aorta in which elastic fibres are in large number. The physiology of the vascular system, particularly the aorta, suggests that abnormalities in elastin could cause SVAS. Decreased vascular elasticity would have an adverse effect on vascular resistance and since hemodynamic stress is significant in the ascending aorta, this structure would be extremely affected [15].

Renal Abnormalities in WS

The spectrum of abnormal renal findings ranged from minor abnormality such as bladder diverticula to major structural malformations such as renal agenesis or renal hypoplasia. The prevalence of renal agenesis is 0.2% in the normal population [16] and about 3.8% in WS patients. Burn [17] proposed the hypothesis that a delayed manifestation of renal diseases and urinary tract infections could be the result of hypercalcemia in early life. Like other arterial stenosis, incidence of stenosis in renal artery could be higher in WS. This hypothesis is supported by the findings that hypoplasia of the ascending aorta and the aortic loop and even long aortic stenosis was seen in a comparatively high number of patients [18]. Considering all the potential renal abnormalities sonographic examination of the kidneys and urinary tract should be included in the standard diagnostic work-up once the initial diagnosis of WS has been made.

Hypercalcemia in WS

The importance of hypercalcemia related to WS has been a subject of debate; absolute hypercalcemia is routinely found only in infancy, if at all. Nearly all patients have symptoms of hypercalcemia in both infancy (vomiting, constipation, and irritability) and adult-hood (constipation, peptic ulcer disease, and frequent urination). Culler et al. [19] have suggested that an abnormality in calcitonin production or release could result in the relative hypercalcemia seen in WS. This relative hypercalcemia could result in an increased calcium load to the kidneys and in polyuria. The relationship of the abnormality in calcium metabolism to the pathogenesis of the other features of the syndrome is not known.

Hypertension in WS

Renal artery stenosis in WS has been correlated with hypertension [20]. Systolic blood pressure is raised in 25% of individuals with WS, diastolic blood pressure is raised in 9%, and both systolic and diastolic blood pressures are raised in 47% [21]. The presence of hypertension in WS increases with age. Cardiovascular disease accounts for nearly all of the morbidity and mortality of WS.

Oral and Dental Features

Hypoplasia of the mandible, high-arched palate, macroglossia, frenulum hyperplasia, hypodontia, microdontia, abnormal incisor morphology, excessive inter dental spacing, anterior cross bite, anterior deep bite or open bite, small slender roots, dens invaginatus, hypoplastic bud-shaped maxillary primary second molars and mandibular permanent first molar, osteosclerotic changes in the lamina dura, delayed mineralization of the teeth, delayed eruption of teeth, hypoplastic defects of the primary and permanent dentition [6-9,22-24].

Hertzberg et al. [22] reported 40.7% abnormal incisor morphology in individuals with WS and microdontic maxillary lateral incisors (peg-shaped) were the most common. Thus, the altered incisor tooth morphology is an important dental feature of WS. Malocclusion occurs in majority of individuals with WS and this could possibly be due to the combined effects of hypodontia, tongue thrust, and connective tissue abnormality. Generalized widened periodontal ligament with vital teeth was recently reported [25] in a patient diagnosed with WS.
Diagnosis

Diagnosis is generally made during mid-childhood when the typical facial features, cognitive profile, and cardiac findings become more apparent [1]. One of the first potential indicators that a neonate may have WS is “prolonged gestation”, classically more than 42 weeks. Other presentations are feeding difficulties, profuse vomiting, and failure to thrive. Due to variability in the clinical findings, diagnosis is routinely confirmed by chromosomal Fluorescent In-Situ Hybridization (FISH) analysis, detecting submicroscopic deletions of 7q11.23 [3] or by polymerase chain reaction [2]. Prenatal diagnosis of the deletion is obtainable for families at risk, using chorionic villous sampling or amniocentesis.

Treatment

Once a diagnosis of WS is made, the multisystem nature of the disorder requires continued treatment. Due to co-existence of multiple problems, many subspecialists including cardiologists, geneticists, ophthalmologists, neurologists, audiologists, dentist (with orthodontist), psychologists and occupational and physiotherapist may be involved in the care of these children. Psycho educational testing should be done routinely so that appropriate early intervention and school placement are provided. Whenever clinical symptoms appear in the urinary tract appropriate referral is indicated [5].

Sudden Death in WS

Sudden death has been reported in many patients with WS. Significant, symptomatic obstruction may develop with any artery [5]. Coronary artery stenosis was found to have occurred in most of the cases that faced sudden death [26]. Abnormalities of the neurovascular lumen leading to stroke have also been reported [27].

Suggestions of Early Diagnosis

Early diagnosis depends on a careful assessment of infants who fail to thrive and are irritable, particularly when these factors are associated with a cardiac murmur, hernias, delayed developmental milestones, or facial features that are different from those of the parents or siblings [5]. In fact, the Committee on Genetics from the American Academy of Pediatrics has established a series of evaluations (Table 1) to assist the clinicians for the early diagnosis of WS [28].

Reasons for Delay in Diagnosis

Despite frequent visits to physicians for treatment of multiple medical problems, the diagnosis of WS is often delayed. The first reason for the delay is that the characteristic facial features may not be obvious at birth but become more obvious with time. Second, although the infants have symptoms that may be related to hypercalceemia, the symptoms are fairly nonspecific and serum calcium levels are not performed usually. Third, despite the fact that 80% of the children with WS have cardiac murmurs, most are asymptomatic in childhood and their murmurs are discharged as functional. Fourth, the developmental delays are often attributed to the chronic medical problems, and the small size and infantile facial features of children with WS make the patients look younger than their chronological age, thus further hide the developmental delays. Parents also may neglect to seek a diagnosis or may disagree with an early diagnosis for similar reasons. Morris et al., [5] describes that the affected children may seem to be “bright” because they are friendly, talkative, and “cute”.

The Role of Dental Evaluation and Cephalometric Analysis in the Diagnosis of WS

The clinical complexity WS is familiar, making it difficult to diagnose on the clinical picture, while genetic testing is not cheap and it is not cost effective to screen all patients. Therefore, dentists can be of help in diagnosing WS early. Tarjan et al., [29] performed Cephalometric analysis and dental examination of 33 patients with genetically confirmed diagnosis of WS. Cephalometric analysis of soft tissues presented that with normal SNA, SNB and ANB angles, the lips were in front of the line of harmony. This observation was present in all WS patients (n=33) but in none of the age-matched controls (n=100). No other variations were found between WS and control patients. This Cephalometric presentation is specific and sensitive for WS and can be applied in the diagnostic procedure.

Medical Considerations in the Dental Treatment of Children with WS

Cardiovascular and renal impairments tend to advance with age. Therefore, before performing any dental treatment, the extent of the disease should be assessed. Myocardial infarction or cardiac arrest has been reported after dental treatment under General Anesthesia (GA) in infancy [26]. Congenital heart defects usually appear in 53%-80% of the affected patients. This phenomenon requires both reduction of anxiety (from dental treatment) and administration of prophylactic antibiotics for bacterial endocarditis. In younger children, when bearing in mind the tendency to anorexia and vomiting, early dental examination and parental counseling are important. Preventive dental regimens and dietary counseling must be custom planned and implemented [30]. In children with WS, orthodontic assessment is generally recommended as soon as the patient's co-operative level allows treatment [22]. Mucogingival problems resulting from anterior cross-bite and pseudo class III malocclusions should also be treated in primary and mixed dentition stages. The apprehension and hyperactive behaviour of children and adolescents with WS should be taken into consideration before planning any dental treatment. Seventy-five percent to 95% of children with WS have mild to moderate mental retardation, and about 80% of patients exhibit generalized anxiety disorder and are more likely to have Attention Deficit and Hyperactivity Disorder (ADHD). Large number (74%-95%) of children with WS is sensitive to sounds [31]. Oral,
References