Sjögren's Syndrome in Children: About 15 Cases in Guinea Conakry

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Case report

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Abstract

Objectives- Sjögren's syndrome is difficult to diagnose in the African context. It is rarely reported in children in Black Africa. We report a series of 15 cases of Sjögren's syndrome in order to clarify the particularities of this condition in children.

Methods- This 2-year retrospective study focused on children under 16 years of age with a male predominance who were followed for AS in the rheumatology and pediatrics departments. Patient data were collected and analyzed using STATA/SE version 11.2 software. Anonymity and respect for ethical rules were the norm. There was no link between patients and researchers.

Results- The average age of the patients was 11 years with extremes of 5 to 15 years. An anamnesis revealing dry mouth was found in more than half of the cases, i.e. in 10 (66.7%) patients. The clinical examination found oral ulceration and periodontitis in equal proportions, i.e. 6 (40%). The immunological assessment and the biopsy of the accessory salivary glands were used as diagnostic evidence in the 15 patients according to the American-European criteria of 2002.

Conclusion- Sjögren's syndrome is a rare entity in pediatrics. It is difficult to diagnose in pediatrics and its severity is linked to the occurrence of visceral and lymphomatous late dry syndrome. Rapid diagnosis and the use of a synthetic antimalarial drug (Hydroxychloroquine) increases the hope of a cure.

Introduction

Sjögren's syndrome (SS) is a rare autoimmune disease, most often secondary in children associated with other autoimmune diseases [1]. It affects girls much more than boys [2] and is characterized by a dry eye, mouth and in about one third of patients, the pathology is more systemic and can affect various organs [3]. Diagnosis is difficult in children because of their inability to accurately list their symptoms and the difficulty in obtaining reliable data on their personal and family history. Diagnosis is based on a combination of biological and immunological clinical signs and biopsy of accessory salivary glands [4]. Prompt diagnosis is necessary to prevent early complications, especially functional ones related to dry and late syndrome such as visceral and lymphomatous [5].

Frequently encountered clinical manifestations are related to autoimmune exocrinopathy with xerostomia; xerophthalmia; dental caries; periodontitis; dysphagia; dysgeusia and keratoconjunctivitis [6].

In spite of the numerous diagnostic criteria of SS, a controversy persists as to the choice of a precise diagnostic criterion especially in children. Adapting the American-European consensus group (AECG) classification, Bartoukova and Al proposed a new scheme for the diagnosis of AS in children [7, 8]. We report 15 cases of AS in children.

Patients And Methods

This was a retrospective study conducted over a 2-year period from August 15, 2018 to August 15, 2020. We targeted patient records (246) followed in the rheumatology and pediatrics department of the Ignace Deen National Hospital, Conakry for an SS. The study focused exclusively on children under the age of 16 years who were followed for an SS. The diagnosis of AS was selected in accordance with the American-European consensus criteria and Vitali's criteria for secondary AS. The data analyzed were sociodemographic (age, sex), clinical (dry oral, ocular, skin, bronchial, ENT), joint, peripheral and axial involvement. Clinical forms: primary and secondary SS (associated with another autoimmune disease). Paraclinical data: immunological (anti-nuclear antibodies, anti-SSA and SSB antibodies). Biopsy of accessory salivary glands according to the Schis olm and Mason classification). Patient data were collected and analyzed using STATA/SE version 11.2 software. All parents and, if possible, capable children (16 years of age) signed consent forms before being included in the study. It was made clear that if patients did not wish to participate in the study, this would not affect the quality of their care.

Results

Of the 15 patients followed in this study, 8 were male and 7 were female with an M/F sex ratio equal to 1.14. The mean age of the patients was 11 years with extremes of 5 to 15 years. The anamnesis revealed dry mouth, nose and eyes in 10 (66.7%), 4 (6.27%) and 3 (20%) of the patients respectively. Ocular pain, hypo agueusia, foreign body sensation in the eyes and ocular burning were found in the following proportions: 2 (13.3); 3 (20); 8 (53.3%); 6 (40%) (Table I). The clinical examination found oral ulceration in 6 (40%); periodontitis in 6 (40%); parotidomegaly in 5 (33.3%) (Table I). Ocular redness and photosensitivity were observed in comparable proportions (66.7%) (Table I). Biology showed a normal blood count, an inflammatory syndrome with a mean sedimentation rate of 55 mm/h (extreme 5 and 130), and a positive C-reactive protein with a mean of 35 mg/l (6 and 78). The immunological check-up routinely performed in all patients showed anti-SSA antibody in 6 children (40%) and anti-SSB antibody in 8 children (53.3%) (Table I). Six (6) children (40%) were diagnosed with a Schis olm and Mason grade IV classification (Table II). In addition, 4 children presented with SS with a grade III Schis olm and Mason classification (Table II). Only three children (6.27%) had an SS with a normal Schis olm-Mason classification. The clinical forms were dominated by primary AS in 9 cases, while secondary AS was present in 6 cases (rheumatoid arthritis 4 cases, lupus 2 cases). After an ophthalmological examination the patients were put on hydroxychloroquine 500mg/24 hours.

Discussion

Sjögren's syndrome is an adult condition with a high frequency in the fourth and fifth decade. It is rare and most often secondary in children and adolescents [6]. We report 15 cases of Sjögren's syndrome in children with a predominance of primitive forms and also noted a male predominance. However, in the literature there is a predominance of female and secondary forms [1]. This finding in our study could be explained by the size of our (smaller) sample.

Diagnostic research is difficult in children because of the often atypical clinical presentation. The clinical manifestations listed in this study are the same ones commonly cited in the literature [7]. The American-European consensus criteria and that of Vitali allowed us to establish the diagnosis in the patients studied in this work. Houghton et al [9] had found 72.5% of recurrent periodontitis, indeed 40% of our patients had already presented a periodontitis. The occurrence of periodontitis is a fundamental element that increases the sensitivity of the European diagnostic criterion of Vitali et al [9].

The immunological assessment often reports the presence of antinuclear antibodies. Anti-SSA antibodies were positive in 40% of children and anti-SSB antibodies in 53.3%. Hamzaoui et al [10]. report that anti-SSA antibodies are most common in 54–75% of cases. Although AS is rare in children, and most often secondary, this study found more cases of primary AS than secondary AS. However, we have not found any data or causal link justifying this etiological disproportion. The schirmer test was positive (15mm of humidification after 5min) in more than half of the patients (60%). The treatment is primarily symptomatic with prevention of dental carries and ulcerative keratitis. Even if no systemic therapy has proven to date to be really effective, the synthetic antimalarial drug (Hydroxychloroquine) used in our patients had a satisfactory result. Corticotherapy and immunosuppressive treatment can be initiated in case of multi-visceral disease

Conclusion

Sjögren's syndrome is a rare entity in pediatrics, we report 15 cases with male predominance and secondary forms. Its diagnosis can be difficult to establish in children because of their inability to describe their symptoms accurately. And its severity is linked to the occurrence of dry and late-onset syndromes such as visceral and lymphomatosis. Prompt diagnosis and the induction of effective treatment will help avoid complications.

Abbreviations

AECG: American-European consensus group

Declarations

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Contributions of the authors

Kaba Condé was the lead author and designed the research; **Carlos Othon Guelngar** recruited the data, followed up with patients and performed the data and statistical analyses. **Mamadou Ciré Barry**, **Hugues Ghislain Atakla** wrote the first draft of the manuscript, which was critically reviewed by **Awada Mohamed**, **Fodé Abass Cissé**. All authors have read and approved the final manuscript.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethical approval and consent to participate

Ethics Committee of the Ignace DEEN, Academic Hospital, chaired by Doctor **Awada Mohamed**, Director General of the hospital.

Consent for publication

Not applicable.

Competing interests

The authors declare no conflict of interest.

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Tables

Table I: Socio-demographic characteristics of patients, clinical manifestations and paraclinical data of dry syndrome

Variables	Effective (%) N=15
Socio-demographic characteristics	
Sex	
Male	8 (53,3%)
Female	7 (46,7 %)
Average age (extreme)	11 years (5 et 16 years)
Clinical events	
Dry mouth	10 (66,7%)
Oral ulceration	6 (40%)
Mouth-sticky foods	4 (26,7%)
Oral pain	2 (13,3%)
Hypo agueusia	3 (20%)
Periodontitis	6 (40%)
Parotidomegaly	5 (33,3%)
Foreign body sensation in the eyes	8 (53,3%)
Eye burn	6 (40%)
Eye redness	9 (60%)
Photosensitivity	10 (66,7%)
Absence of tears	3 (20%)
Nasal dryness	4 (26,7%)
Schimer test	9 (60%)
Immunological assessment	
Anti-SSA antibodies	6 (40%)
Antibodies to SSB	8 (53,3%)

Table II: Schisolm and Mason classification

Stades	Effective	Percentage
Normal	3	20%
Grade I	2	13,3%
Grade II	0	0%
Grade III	4	26,7%
Grade IV	6	40%
Total	15	100%