Characteristics of Orofacial Amyloidosis: A Case Series

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Abstract

Amyloidosis derived from abnormal extracellular fibril deposits may contribute to multiple organ dysfunctions. The recognition of amyloidosis-associated orofacial changes may be beneficial for early diagnosis. This retrospective study determined the characteristics of orofacial amyloidosis to aid in recognition of this disease. The study included 11 patients who visited Peking University School of Stomatology from 1993 to 2015 and were diagnosed with orofacial amyloidosis. The median age at onset, most commonly affected site, predominant oral feature of amyloidosis, and complications were presented. We concluded that the recognition of amyloidosis-associated orofacial changes may be beneficial for the diagnosis of amyloidosis and the discovery of underlying disease.

Introduction

Amyloidosis is a cluster of heterogeneous diseases caused by the extracellular deposition of insoluble fibrillar proteins [1]. Amyloidosis is generally classified into three types: primary amyloidosis, secondary amyloidosis, and familial or hereditary amyloidosis [2]. Based on the site of fibrillar protein deposition, the disease can also be divided into localized or systemic amyloidosis. Primary systemic amyloidosis is often attributed to plasma cell dyscrasia arising from multiple myeloma or other clonal B cell diseases [3]. Secondary (or reactive) systemic amyloidosis is usually derived from inflammatory diseases such as rheumatoid arthritis, chronic suppuration tuberculosis, Hodgkin’s lymphoma, syphilis, and rickets [4]. The kidneys, liver, heart, and peripheral nervous system are affected most often, leading to nonspecific disorders such as proteinuria, liver enlargement and functional disorders, arrhythmia, heart hypertrophy, and cardiac insufficiency [5]. Gastrointestinal tract functional disturbances, bleeding, and blockage may also result from amyloidosis. Ultimately, single- or multiple-organ dysfunction develops. About 80% of the patients with primary systemic amyloidosis do not survive for 2 years [5].

Although systemic amyloidosis is incurable, an early accurate diagnosis may facilitate treatment to reduce the amyloid production and prevent the exacerbation of organ dysfunction [5]. The nonspecific manifestations of amyloidosis make the histopathological findings of amyloid materials in the affected tissues crucial for the diagnosis. However, the technical difficulty, bleeding, rare possibility of organ perforation, and patient discomfort limit biopsies of the liver, kidneys, and heart, which are important visceral organs [6]. Consequently, the British Committee for Standards in Hematology (BCSH) recommended a less invasive biopsy of an accessible site [7]. The orofacial region may be the best alternative because of its open nature, lower risk of biopsy-associated complications, and status as a commonly affected region in amyloidosis [8]. Therefore, the clinical characteristics of orofacial amyloidosis were summarized here for recognition.

Materials and Methods

This study included patients who visited the clinic of Peking University School of Stomatology from 1993 to 2015 and were diagnosed with orofacial amyloidosis based on the clinical manifestations and histopathological findings. Histological diagnoses were made by two pathologists, separately, based on the findings of a biopsy taken from the orofacial lesions. Both hematoxylin and eosin (H&E) and Congo red staining showed eosinophilic or orange-pink amorphous homogeneous amyloid deposits in the lamina propria (Figure 1). The demographic data, chief complaint, medical history, oral clinical assessment, finding of systemic diseases, and results of laboratory examinations were collected.

Results

During the past 21 years, 11 amyloidosis patients were initially diagnosed in our hospital based on orofacial abnormalities. The median age of onset of these patients was 62 (range 17–74 years) years, and the male-to-female ratio was 2:3. Before making the diagnosis, the oral clinical signs were
Amyloidosis is a rare disease mainly involving older populations [9,10] of systemic amyloidosis patients, 90% will develop amyloid deposits in the head, neck, or respiratory tract [11], and a previous study indicated that 65–70% of adults visit a dental clinic at least once a year [12]. Therefore, amyloidosis-associated oral mucosal changes may be recognized initially by oral health professionals, and this is beneficial for the discovery of underlying diseases. The results of this study indicate that about 82% of the included patients developing orofacial amyloidosis were around the age of 60 years, although one patient was diagnosed at 17 years of age and the orofacial abnormality had persisted for 13 years. The major disease duration was 0.5 to 2 years. These observations suggest that orofacial amyloidosis is usually asymptomatic and insidious over a long period. A lack of recognition of the disease leads to neglect by clinicians; in many cases, it is not discovered until autopsy [13].

The tongue was the most frequently affected site in this study, followed by the buccal and labial mucosae, gingiva, and parotid gland. Multiple, painless, waxy nodules were the predominant manifestations on the oral mucosa. One patient also exhibited oral purple nodules, which were the same as those reported by Babburri et al. [14]. Two patients had mucocutaneous ecchymosis, which was thought to be associated with multiple myeloma [15]. Macroglossia was found occasionally, although it is generally considered to be most common in the oral cavity of systemic amyloidosis patients [11,15].

Amyloid infiltration in the major salivary glands is rare, and may be localized or secondary to systemic amyloidosis, presenting with even or lobulated gland enlargement [16-18]. In this study, enlargement of the parotid gland was seen with multiple nodule-like lesions. The patient with parotid gland amyloidosis had a high serum level of rheumatoid factor (RF) and suffered from concomitant cryoglobulinemia. Cryoglobulinemia is characterized by the presence of cryoglobulins in the serum at low temperatures. The marked increase in RF in subject DXE may be ascribed to the deposition of cryoglobulins [19], of which types II and III have RF activity [20]. Both amyloid and cryoglobulins may be derived from lymphoproliferative disorders such as multiple myeloma [21,22]. In this regard, lymphoma resulting from monoclonal lymphocyte proliferation should be excluded. No histopathological evidence was found in the present study for lymphoproliferative disorders.

Primary systemic amyloidosis may arise from multiple myeloma or other clonal B cell diseases [3]. Approximately 15–20% of primary systemic amyloidosis patients have multiple myeloma and vice versa [12]. Amyloidosis may develop before or after multiple myeloma, and physicians should be aware of the close relationship between these disorders. Osteoporosis and even osteolysis may occur secondary

Figure 1: a) Photomicrograph of the lesion showing the surface epithelium and homogenous eosinophilic area in the connective tissue (H&E, 40×). b) Photomicrograph of the lesion showing the surface epithelium; the amyloid appears orange-pink in the connective tissue (Congo red, 40×). (The photomicrographs were taken from the pathological slices from subject YJ.)

Figure 2: a) Intraoral photograph showing multiple, painless, waxy, well-circumscribed hard nodules involving the lateral side of the tongue in subject YJ. b) Intraoral photograph showing multiple, painless, waxy, well-circumscribed hard nodules accompanied by purple nodules without fading under pressure involving the tongue in subject ZSX. c) Intraoral photograph showing the lesion involving the buccal mucosa in subject ZSX. d) Intraoral photograph showing diffuse, hard enlargement of tongue (macroglossia) in subject ZGQ.

Table 1

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to multiple myeloma, as observed in the present study. With the expansion of neoplastic plasma cells within the bone marrow in multiple myeloma, normal bone homeostasis maintained by activated osteoblasts and osteoclasts is disrupted. Osteoclast activity is promoted by proteins secreted from stromal cells, while osteoblast activity is inhibited [23,24]. Primary cutaneous nodular amyloidosis was histologically considered to be identical to myeloma-associated systemic amyloidosis with monoclonal immunoglobulin light chain deposits, and may be complicated with psoriasis [25]. Similarly, both psoriasis and cutaneous amyloidosis developed in one patient with multiple myeloma (subject FP). This suggests that patients with monoclonal immunoglobulin light chain-associated amyloidosis are susceptible to psoriasis.

In summary, orofacial amyloidosis may present with macroglossia and diffuse lingual enlargement or with asymptomatic multiple, waxy, or purple nodules on the lingual and buccal mucosae, gingiva, and parotid gland. It may occur secondary to multiple myeloma or be complicated with cryoglobulinemia, psoriasis, osteoporosis, osteolysis, or other nonspecific disorders of the liver, kidneys, heart, and neurological system. The recognition of amyloidosis-associated orofacial changes and relevant systemic diseases by oral clinicians may be of benefit in the diagnosis of amyloidosis and the discovery of underlying diseases; moreover, it may limit further disease progression. However, the high frequencies of nonspecific complications in older populations, together with the limitations of disease progression. However, the high frequencies of nonspecific complications in older populations, together with the limitations of disease progression.

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