Glycogenic acanthosis on mouth clinically present as white plaque

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ABSTRACT

Glycogenic acanthosis is a benign condition, commonly observed during endoscopic procedures in older patients, which present as slightly elevated whitish plaques often on the lower third of the oesophagus. Microscopically, glycogenic acanthosis is composed of hyperplastic squamous epithelium with intracytoplasmic glycogen deposits. The extraoesophageal glycogenic acanthosis is extremely rare, with only three case reports in the English-language literature. We report a white lesion showing glycogenic acanthosis-like features located on the left posterosilateral border of the tongue, affecting a 56-year-old male patient. The medical history was non-contributory and the patient did not show any lesions during endoscopic examination of the oesophagus, stomach, and upper duodenum. Glycogenic acanthosis is a benign condition, which should be included in the differential diagnosis when assessing oral white lesions. It is important also to recognize this benign condition early and rule out the possibility of other more severe diseases, but further studies were necessary for better define their potential for persistence or recurrence, as observed in the current case.


RESUMO

A acantose glicogênica é uma condição benigna, comumente observada durante procedimentos endoscópicos em pacientes idosos, e se apresenta como placas brancas levemente elevadas, frequentemente encontrada no terço inferior do esôfago. Microscopicamente, a acantose glicogênica é composta por epitélio escamoso hiperplásico com depósitos de glicogênio intracitoplasmático. A acantose glicogênica extraesofágica é extremamente rara, com apenas três relatos de casos na literatura em língua inglesa. Apresentamos um caso com uma lesão branca diagnosticada como acantose glicogênica, localizada na borda póstero-lateral esquerda da língua, afetando um paciente do sexo masculino de 56 anos. A história médica não foi contributiva e o paciente não apresentou lesões durante o exame endoscópico do esôfago, estômago e duodeno superior. A acantose glicogênica é uma condição benigna, que deve ser incluída no diagnóstico diferencial na avaliação de lesões brancas orais. É importante também reconhecer precocemente essa condição benigna e descartar a possibilidade de outras doenças mais graves, porém mais estudos são necessários para melhor definir seu potencial de persistência ou recorrência, como observado no presente caso.


INTRODUCTION

Glycogenic acanthosis (GA) is a benign lesion characterized by multifocal white plaques usually subtle, rounded elevations, often involving the lower oesophagus [1]. Microscopically, it is composed of hyperplastic squamous epithelium with abundant intracellular glycogen deposits. As any white lesions affecting the oral mucosa, this needs to be carefully investigated with all attention towards

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to oral leukoplakia (OL) since it is considered a potentially malignant disorder [2]. The diagnosis of OL is established by exclusion, eliminating all the others possible conditions such as frictional keratosis, lichenoid lesion, candidiasis, leukoedema, graft-versus-host disease, white sponge nevus, among others [3]. Mostly, the histopathological analysis becomes very important, which in close correlation with the clinical findings, allows establishing a diagnostic conclusion.

The extraoesophageal manifestation of GA is extremely rare, and to the best of our knowledge, there are only four extraoesophageal case reports, three cases involving the oral cavity and one involving the larynx [4-6]. In this study, our aim was to report a rare case of white lesion showing GA-like features affecting the lateral border of the tongue and emphasizing the differential diagnosis.

**CASE REPORT**

A 56-year-old white man was referred to the Oral Medicine Service for evaluation of a white patch in the oral cavity. The patient denied smoking and alcohol consumption or other habits and his medical history were non-contributory. According to the patient, a similar lesion had been removed 4 years ago in the same location. This previous excisional biopsy did not reveal any sign of dysplasia, being only followed up. On intraoral examination, there was a well-defined and asymptomatic white plaque with a fluffy surface, measuring 3.0 cm in diameter, located on the left posterolateral border of the tongue (figure 1A). There was no sign of chronic local trauma. Although it has an uncommon presentation, OL was the main hypothesis of clinical diagnosis. Our conduct was to perform an excisional biopsy. The microscopic analysis showed a hyperplastic squamous epithelium, showing prominent acanthosis in close transition with parakeratosis, which through Periodic acid–Schiff (PAS) with and without diastase stain revealed abundant intracytoplasmatic glycogen deposits (figure 2B-E). No cellular pleomorphism, nuclear hyperchromatism, mitotic figures or abnormalities were visualized. However, different from typical OL, the acanthosis and irregular parakeratosis were suggestive of a reactive process. The endoscopy examination did not show any alteration on the oesophagus, stomach, and upper duodenum. Considering the clinical aspect and the microscopical features, the diagnosis of glycogenic acanthosis was established. After 16 months of follow up, no signs of recurrence were observed.

**DISCUSSION**

Glycogenic acanthosis was a term adopted by Rywlin and Ortega [7] in 1970 to refer to areas of mucosal thickening, usually seen on the lower oesophagus, characterized microscopically by epithelial hyperplasia with an accumulation of intracytoplasmatic glycogen. Instead of OL, it does not show any potential for malignancy. The oesophageal GA has become increasingly know on endoscopy examination, with some recent evidence to its association to the gastroesophageal reflux disease [8-10]. In this study, we reported a rare case of white lesion showing GA-like features on the tongue and we review the extraoesophageal case reports of GA published in the English-language literature (table 1).

Among all cases found, including our case, the median age at the time of diagnosis was 65 years, ranging from 53 to 79 years. Thus, an age-related, degenerative process should be considered in their etiopathogenetic mechanisms. Tongue involvement was observed in only two patients, being the lesion size not greater than 4.0 cm in diameter. Other oral sites may be affected by the glycogenic acanthosis, as the buccal mucosa [11].

The recognition of a similar standard of epithelial response in larynx and oesophagus suggest that epithelial hyperplasia associated with glycogen
accumulation might be a non-specific pattern of epithelial response linked to site-specific injury changes [4]. In a previously published GA case, during a 5-year follow-up, no sign of malignant transformation was observed [6]. Curiously, in the current case, acanthosis and parakeratosis were observed. This latter may be due to the location of the lesion and the possible effects of chewing and swallowing, suggesting a reactive process.

Mucosal lesions showing similar features with GA have been reported in patients with Cowden’s Syndrome [5], a genetic disorder characterized by multiple hamartomas, which affect multiple organs [12]. However, the current patient did not show any mucous or cutaneous lesions that would indicate a syndromic condition.

Oesophageal GA is found in 3.5% of the oesophageal endoscopic examinations [8]. Despite the name, there is no association between GA and abnormalities of glucose metabolism, such as diabetes9. During the examination, these lesions appear as white-grey plaques, slightly elevated, usually measuring 2-10 mm of diameter [10]. Interestingly, different from previously published GA cases, the current case had a recurrence. Nevertheless, since this condition did not require any treatment, or it is reported in autopsies and postmortem studies, the rate of recurrence was not established yet.

In the differential diagnoses, the frictional keratosis was excluded since there was no sign of chronic local trauma. We considered OL as the principal clinical hypothesis since it is the most common oral potentially malignant disorder with a frequency of malignant transformation of 17-24% [13]. However, there was no associated risk factor, such as smoking and alcohol consumption, mainly associated with OL. Also, the histopathological analysis was not consistent with typical OL, which usually shows in hyperplastic lesions, acanthosis, hypergranulosis, and hyperorthokeratosis, maintaining the cellular architecture, different from the current case. A specific clinical subtype of OL is verruciform or verrucous leukoplakia (VL). This condition is characterized by numerous pointed projections developed on the surface and may be similar with the current case [14]. However, different from GA, VL microscopically reveals varying degrees of epithelial dysplasia and hyperkeratosis. Oral verrucous hyperplasia (VH) is a distinctive entity, which shows considerable clinical and histopathological resemblance to verrucous carcinoma (VC); however, different from VC, in oral VH its base lies above the line connecting the bases of the normal-appearing epithelium on both sides of the lesion. Oral VH has been recognized as a potentially malignant disorder, with a malignant transformation rate estimated of 10-20% [15,16]. VC, a low-grade variant of squamous cell carcinoma, is often on the oral cavity. It is characterized by a diffuse, largely exophytic, superficial spreading highly keratinized lesion. It occurs mostly in elderly male, presenting slow growth and localized invasion, without metastatic potential [17]. Lastly, verrucous epidermal nevus (VEN), a congenital

Table 1. Extraesophageal case reports of glycogenic acanthosis.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Year</th>
<th>Gender</th>
<th>Age</th>
<th>Country</th>
<th>Oral site</th>
<th>Size</th>
<th>Other findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fyfe &amp; Garcia [4]</td>
<td>1998</td>
<td>Male</td>
<td>79</td>
<td>USA</td>
<td>Larynx</td>
<td>2cm</td>
<td>Hypertension and atherosclerosis</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Cowden syndrome: Colon polyposis, lipoma, ovarian cyst, thyroid adenoma and breast fibroma</td>
</tr>
<tr>
<td>Nishizawa et al. [5]</td>
<td>2009</td>
<td>Female</td>
<td>53</td>
<td>Japan</td>
<td>Gingiva</td>
<td>-</td>
<td>Cowden Syndrome: Colon polyposis, lipoma, ovarian cyst, thyroid adenoma and breast fibroma</td>
</tr>
<tr>
<td>Montebuognoli et al. [6]</td>
<td>2010</td>
<td>Male</td>
<td>72</td>
<td>Italy</td>
<td>Tongue</td>
<td>4cm</td>
<td>No other findings</td>
</tr>
<tr>
<td>Present case</td>
<td>2017</td>
<td>Male</td>
<td>56</td>
<td>Brazil</td>
<td>Tongue</td>
<td>3cm</td>
<td>No other findings</td>
</tr>
</tbody>
</table>

malformation or hamartoma derived from embryonic ectoderm, should be also considered. Oral involvement by VEN is a rare manifestation, described as unilateral or midline papules or nodules, with a papillary or verrucous surface, usually affecting the tongue, lips, gingiva, buccal mucosa and palate [18].

CONCLUSION

Glycogenic acanthosis is not yet widely recognized in the field of dentistry. However, it is necessary to include it in the differential diagnosis list for white lesions of the oral mucosa. The clinicopathological presentation of the current case suggests a reactive or age-related process.

REFERENCES


Collaborators

MK SCHULZ, MR BIANCARDI, FERNANDES and LY ALMEIDA acquisition of data: laboratory or clinical/literature search, analysis and interpretation of data collected and drafting of article and/or critical revision. D, conception and design of study/review/case series, acquisition of data: laboratory or clinical/literature search, analysis and interpretation of data collected and drafting of article and/or critical revision. A BUFALINO and JE LEÓN, conception and design of study/review/case series, acquisition of data: laboratory or clinical/literature search, analysis and interpretation of data collected and drafting of article and/or critical revision.