Histogenetic Study of Verruciform Xanthoma of the Gingiva

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Abstract

Verruciform xanthoma is characterized macroscopically by papillomatous or verrucous hyperplasia of the mucosal epithelium and histopathologically by papillary epithelial hyperplasia and foam cell accumulation in the lamina propria between epithelial processes. This relatively rare lesion does not appear to be a true tumor, but rather an inflammatory event. Here, verruciform xanthoma of the gingiva was subjected to histopathological and immunohistochemical analyses. Notably, the cortical layer of the lesion exhibited a verrucous and granular outer surface, similar to a papilloma. Within the lesion, keratinized stratified squamous epithelium and epithelial process extensions were observed, and the connective tissue between clubbed epithelial processes was filled with foam cells. These foam cells exhibited strong cytoplasmic and membrane expression of CD68, c1–anti-trypsin, and macrophage scavenger receptor-1 (MSR-1), as well as human leukocyte antigen (HLA)-DR and oxidized low-density lipoprotein cholesterol (ox-LDL). The epithelial cells also expressed HLA-DR in the cytoplasm and cell membrane. By contrast, the expression of S-100 and CD1a in Langerhans cells was clearly reduced in the epithelium of the verruciform xanthoma, while the inflammatory infiltrating cell population comprised of mainly CD3- or CD8-positive cells, with few CD20- or CD4-positive cells. The increased lipid content of the cell membrane and concomitant epithelial hyperplasia causes cellular injury and leakage into the connective tissue consistent with dysregulated cellular immunity in the stratified squamous epithelium. Accordingly, it may be concluded that macrophages phagocytose these lipids and differentiate to foam cells.

Keywords: Verruciform xanthoma; Foam cells; CD68; Epithelial hyperplasia; Inflammatory cell infiltration

Introduction

Xanthomas are dermal lesions attributed to lipid or lipoprotein disorders. Within this category, lesions of the oral region that exhibit papillomatous hyperplasia are classified as verruciform xanthoma [1]. These relatively rare lesions are characterized histologically by epithelial papillary hyperplasia and foam cell accumulation in the lamina propria between epithelial processes [2-8].

Two theories have been proposed to describe the histogenetic mechanism underlying verruciform xanthoma. In the first, epithelial papillary hyperplasia is triggered by an inflammatory response (e.g., chronic stimulation), and underlying macrophages subsequently phagocytose lipids from the cell membranes of the modified epithelial cells and differentiate to foam cells [2,9]. In the second hypothesis, a lipid metabolic abnormality occurs consequent to delayed inflammation or repetitive mechanical stimulation, after which macrophages in the connective tissue differentiate to foam cells; in this scenario, papillary hyperplasia of epithelium is a secondary event [10]. To date, neither hypothesis has been assessed the epithelium and differentiated.

In addition to the mechanistic uncertainties, few reports have addressed the epithelium and inflammatory cells associated with verruciform xanthoma, although an understanding of these factors is needed to elucidate the guidelines for and characteristics of this disease. Therefore, this report describes a histogenetic study of verruciform xanthoma of the gingiva based on histopathological and immunohistochemical analyses.

Materials and Methods

Clinical information

A 29-year-old man presented to our dental office in December 2015 for an evaluation of a painless swelling on the labial side of the marginal gingiva around the lower right medial incisor. He had long noticed the small lesion but had not previously sought consultation at a medical institution and was referred to our clinic by his regular dentist. The lesion measured 5 mm × 2 mm and matched the color of the gingiva. Additionally, it was sharply marginated, with a granular surface and soft elasticity at the affected gingiva. Although the associated incisor had a normal tooth crown, the surrounding gingiva exhibited slight redness and swelling. X-ray findings showed no abnormality. The tumor was resected en bloc under local anesthesia.

Histological analysis

The specimens were fixed in 10% neutral buffered formalin, embedded in paraffin, and cut into 7-mm-thick sections. The sections were stained with hematoxylin and eosin according to a routine procedure and observed with a light microscope (CX41; Olympus Corp., Tokyo, Japan).

Immunohistochemical analysis

For the immunohistochemical analysis, the specimens were fixed, embedded, and cut as described above, and labeled using an indirect peroxidase-labeled streptavidin-biotin technique (DAKO, Glostrup, Denmark). The antibodies used for immunohistochemistry, Negative controls were incubated with phosphate-buffered saline (PBS) instead.
of antibodies, Normal tissue adjacent to the lesion (i.e., without inflammation) was used as the control tissue.

Results

Histological analysis

In hematoxylin and eosin-stained sections of the verruciform xanthoma, the outer side of the cortical layer exhibited a verrucous and granular appearance similar to a papilloma. Stratified squamous epithelium with keratinization and extended epithelial processes were also observed. The connective tissue between the clubbed epithelial processes was filled with foam cells, and the extended connective tissue exhibited densely aggregated foam cells and an increase in the number of enlarged capillary vessels. Furthermore, lymphocytic inflammatory cell infiltration occasionally with the focal region was also observed there (Table 1 and Figures 1-3).

Table 1: Immunohistochemical analysis.

<table>
<thead>
<tr>
<th>Antibodies</th>
<th>Clone</th>
<th>Source</th>
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<tbody>
<tr>
<td>CD68</td>
<td>PGM-1</td>
<td>Dako/Agilent Technologies</td>
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<tr>
<td>α1-antitrypsin</td>
<td>Polyclonal</td>
<td>Dako/Agilent Technologies</td>
</tr>
<tr>
<td>HLA-DR</td>
<td>TAL-1B5</td>
<td>Dako/Agilent Technologies</td>
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<tr>
<td>ox-LDL</td>
<td>Polyclonal</td>
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<td>S-100</td>
<td>Polyclonal</td>
<td>Roche Diagnostics</td>
</tr>
<tr>
<td>CD1a</td>
<td>EP3622</td>
<td>Cell Marque/Sigma-Aldrich</td>
</tr>
<tr>
<td>CD3</td>
<td>2GV6</td>
<td>Roche Diagnostics</td>
</tr>
<tr>
<td>CD20</td>
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<td>Roche Diagnostics</td>
</tr>
<tr>
<td>CD8</td>
<td>C8/144B</td>
<td>Nichirei Biosciences</td>
</tr>
<tr>
<td>CD4</td>
<td>SP35</td>
<td>Nichirei Biosciences</td>
</tr>
</tbody>
</table>

Figure 1: Clinical findings, A: The tumor was 5 × 2 mm, gingival color, sharply margined, granular surface, and soft elasticity at the affected gingiva of lower right medial incisor. Although the tooth crown of the incisor was normal, the surrounding gingiva exhibited slight redness and swelling. B: X-ray findings. No abnormality was shown.

Figure 2: Histological analysis, the sections were stained with hematoxylin and eosin. Stratified squamous epithelium with keratinization and extended epithelial processes were observed. Connective tissue between epithelial processes was filled densely aggregated foam cells. Lymphocytic inflammatory cell infiltration was also observed in extended connective tissue. A; ×40, B; ×100.

Figure 3: Immunohistochemical analyses, the foam cells exhibited strong cytoplasmic and cell membrane expression of CD68 (A), α1-antitrypsin (B), HLA-DR (C), and ox-LDL (D) were also detected on foam cells. A-D; ×200. A clear reduction in the population of S-100 protein- (E) and CD1a- (F) positive cells was observed. E, F; ×200. The inflammatory infiltrating cells mainly comprised CD3- (G) or CD8- (H) positive cells, with few CD20- (I) or CD4- (J) positive cells. G-J; ×200.

Immunohistochemical analysis

The foam cells exhibited strong cytoplasmic and cell membrane expression of CD68, α1-antitrypsin, and macrophage scavenger receptor-1 (MSR-1). Human leukocyte antigen (HLA)-DR and oxidized low-density lipoprotein (ox-LDL) were also detected on foam cells (Figure 3A-3D, and data not shown). The epithelial cells also expressed HLA-DR and ox-LDL in the cytoplasm and cell membrane. However, a clear reduction in the population of S-100 protein- and CD1a- positive Langerhans cells was observed (Figure 3E and 3F). The inflammatory infiltrating cells mainly comprised CD3- or CD8- positive cells, with few CD20- or CD4- positive cells (Figure 3G-3J). An abundance of infiltrating cells was observed in the epithelium of the verruciform xanthoma epithelium compared to that of normal tissue.

Discussion and Conclusion

Verruciform xanthoma, a relatively rare condition, most frequently affects the oral mucosa and genital skin. As noted above epithelial
lymphocyte-mediated immunological activation, which may aggravate xanthoma, and skin verruciform xanthomas have also reported to occur concomitantly with oral verruciform involving a decrease of Langerhans cells and increase in subepithelial boundary, localized proliferation, and a diameter consistent with planus, leukoplakia, pemphigus vulgaris, and carcinoma in situ, have been reported to occur concomitantly with oral verruciform xanthoma, and skin verruciform xanthomas have also reported as concomitant symptoms of many epithelial proliferative diseases. Therefore, some researchers insist that verruciform xanthoma is a secondary disease [13].

In addition to histological features, verruciform xanthoma can also be identified using immunohistochemical characteristics. For example, the foam cells detected in the connective tissue appear to originate from migrating/inflammatory macrophages and can be detected immunohistochemically using antibodies specific for CD68 and α1-antitrypsin [2-8]. Additionally, the detection of HLA-DR suggests T-lymphocytes, particularly CD8-positive T-lymphocytes, which are infiltrated with superficial localized lymphocytes and neutrophils [1-5]. These typical histopathological findings facilitated a diagnosis of verruciform xanthoma.

Various local epithelial proliferative diseases, including lichen planus, leukoplakia, pemphigus vulgaris, and carcinoma in situ, have been reported to occur concomitantly with oral verruciform xanthoma, and skin verruciform xanthomas have also reported as concomitant symptoms of many epithelial proliferative diseases. Therefore, some researchers insist that verruciform xanthoma is a secondary disease [13].

In contrast to the above-described markers, the foam cells in this case did not express S-100 protein or CD1a, which are markers of epithelial tissue-resident macrophages or Langerhans cells, which act as antigen presenting cells in the mucosa. This result corresponds to previous reports, wherein many positive cells were detected in the normal epithelial layers [7,16-18]. The reason for this result is not clear, although some researchers reported a counter-regulatory mechanism involving a decrease of Langerhans cells and increase in subepithelial macrophages [19].

In the present study, ox-LDL was strongly expressed in the epithelium of the verruciform xanthoma. In inflammatory diseases, epithelial hyperplasia of the stratified squamous epithelium is often associated with an increase in membrane lipid levels [20,21]. Using electron microscopy, Suka [15] reported the loss of basement membrane, degeneration of the basal cells, and disruption of the intracellular junction, which led to the leakage of lipids from the broken epithelium [20]. In turn, these lipids were oxidized by macrophages, and the resulting ox-LDL is incorporated into the cells via MSR-1. Finally, the macrophages become foam cells [22,23]. The foam cells in this case expressed both ox-LDL and MSR-1, indicating that MSR-1-positive macrophages accumulated in the papillary area of the connective tissue, where they phagocytosed lipids released from epithelial or basal membranes and transitioned to foam cells within the verruciform xanthoma.

As noted in the introduction, there are two theories regarding the histogenetic of verruciform xanthoma. As the present case was unlikely to have involved long-term mechanical stimulation, it is more likely to follow the process of the first theory due to an inflammatory reaction such as periodontitis. It is appropriate to be concluded that increasing the lipid content while epithelial hyperplasia leaks into the connective tissue caused by cellular injury, consistent with dysregulated cellular immunity in the stratified squamous epithelium, and the phagocytosis of these lipids by macrophages was occurred, which then differentiate into foam cells.

Consent

Written informed consent was obtained from the patient for publication of this report and any accompanying figure.

References


