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CASE REPORT

Plasmablastic lymphoma of oral mucosa type: A case report

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Summary A case of the plasmablastic lymphoma of oral mucosa type is described. A 60-year-old white man presented an ulcerated and bleeding swelling in the upper lip. The right hard and soft palate showed a swelling covered by a yellowish membrane. Microscopical examination showed a monotonous diffuse lymphoid proliferation of large cells with plasmablastic differentiation. The cells were immunopositive for VS38c. A PCR demonstrated IgH with monoclonal pattern and EBV DNA. The lesions resolved after treatment with local radiotherapy and systemic chemotherapy. The patient remains free of the disease after twenty-three months of treatment.

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Introduction

Diffuse large B-cell lymphoma (DLBCL) is the most common type of lymphoma in western countries, representing about one third of these neoplasm.¹

The diversity of clinical presentations, genetic, and molecular characteristics suggests that these neoplasm represent a heterogeneous group. However, the World Health Organization (WHO, 2001) considers the DLBCL as a specific category.¹⁰

Plasmablastic lymphoma (PBL) was initially described as a rare variant of DLBCL. It's present in people infected by the human immunodeficiency virus (HIV).⁵ PBL has predilection for the oral cavity

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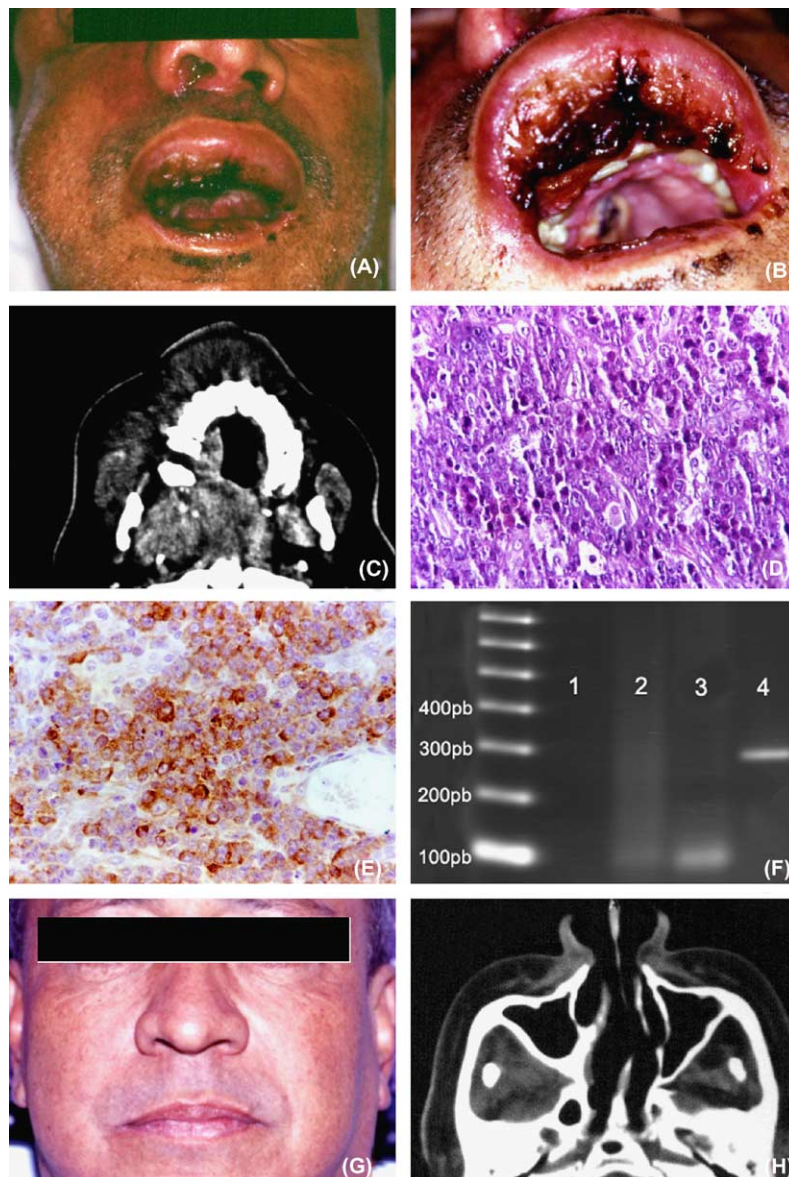


Figure 1 (A) Clinical appearance of the plasmablastic lymphoma of oral mucosa type show asymmetric face. (B) Intraoral view demonstrates an ulcerated and bleeding swelling in the upper lip. The right hard and soft palate showed a swelling covered by a yellowish membrane. (C) Computed tomography scan demonstrate a lesion infiltrating the parapharyngeal space, nasal fossae and right maxillary sinus. (D) Microscopic appearance of monotonous diffuse lymphoid proliferation composed by large cells with plasmablastic differentiation (hematoxylin-eosin, original magnification $\times 200$). (E) Lymphoid neoplastic cells immunopositive for VS38C (streptavidin-biotin technique, original magnification $\times 200$). (F) 2.0% agarose gel eletrophoresis after PCR amplification of DNA. Track 1 is negative control. Track 2 is DNA from tonsil. Track 3 is DNA from plasmablastic lymphoma of oral mucosa type with a discrete and homogeneity band (approximately 110 bases pare). Track 4 is DNA from plasmablastic lymphoma of oral mucosa type with presence of the EBV genome (band of 269 bases pares) (PCR technique, original size). (G) Clinical appearance of face after fifteen months of treatment. (H) Computed tomography scan after fifteen months of treatment within evidence of the lesion.

with local invasion and rapid dissemination to extraoral sites. PBL have a poor prognosis with average survival time of the six months.^{5,13,7} In addition, it exhibited an immunphenotypic profile with absent or weak expression of B-cell markers

and reactivity for plasma cell associated antigens, and is EBV-positive.⁴

We reported the clinical, microscopical, immunohistochemical, molecular features, EBV research and management of PBL of oral mucosa

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