Multidetector Computed Tomography Features of Plasmablastic Lymphoma of the Oral Cavity in an HIV-Positive Patient

Case Report

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Abstract

Overview: Plasmablastic lymphoma (PBL) is an uncommon aggressive malignancy that is associated with patients with human immunodeficiency virus (HIV), other immunosuppression conditions, or in elderly patients, that tends to appear in the oral cavity. PBL accounts for 2.6% of all HIV-associated non-Hodgkin lymphomas.

Case Report: We describe the case of a chemotherapy-treated patient with HIV-associated plasmablastic lymphoma who presented with a three-month history of left facial swelling. Features seen on different imaging modalities including panoramic and multi-detector computed tomography (MDCT) were suggestive of a malignancy and/or an inflammatory lesion. However, histopathological examination revealed an aggressive immunoblastic neoplasm, consistent with AIDS-associated PBL.

Conclusion: PBL is hard to diagnose, rapidly progressive, with overall poor prognosis among HIV-infected patients, and often rapidly fatal. The maxillofacial radiologist must be aware of the existence of this relatively newly described lymphoma as it is associated with immunosuppression and may be misdiagnosed as other pathoses.

Keywords: Plasmablastic Lymphoma; PBL; HIV; AIDS; Multidetector Computed Tomography.

Introduction

Lymphoma is defined as a group of tumors that develop from lymph nodes [1]. Patients with lymphomas may experience fever, weight loss, night sweats, lymphadenopathy, and fatigue [1]. Hodgkin lymphomas (HL) and non-Hodgkin lymphomas (NHL) are the two primary kinds of lymphoma [2]. Plasmablastic lymphoma (PBL) is classified as a distinct entity of B-cell lymphoma in the recent WHO classification of lymphoid neoplasms [2]. Other names of this variety of lymphoma include anaplastic lymphoma kinase-positive diffuse large B-cell lymphoma (ALK-DLBCL), plasmablastic plasmacytoma, and the human herpesvirus 8 (HHV8) - related primary effusion lymphoma (PEL) [2].

PBL is considered a rare lymphoma that may present in the oral cavity [3]. It was first noted by Delecluse et al., in 1997 and is known to have a strong association with human immunodeficiency virus (HIV) and Epstein-Barr virus (EBV) infections [4, 5]. Tyrosine kinases also displayed a pathogenic role in lymphomas in which anaplastic lymphoma kinase (ALK) is the most transforming gene of the majority of anaplastic large cell lymphomas (ALCL) [6].

The estimated frequency of PBL is about 2.6% of NHL cases with acquired immune deficiency syndrome (AIDS) [7]. 92% of oral PBL cases depicted in the English medical literature were HIV-positive patients [8]. However, PBL is also known to affect...
HIV-negative patients [9]. This case report describes the Multi-Detector Computed Tomography (MDCT) imaging features of oral PBL in an HIV-positive patient and discusses the progression of this condition.

**Case Report**

A 57-year-old male patient with a medical history of HIV and pulmonary embolism presented with a three-month history of left facial swelling to the Emergency Room. The patient was given vancomycin and clindamycin before transferring to the emergency department, however, his left facial swelling continued to progress. Clinical examination revealed intraoral swelling in the left mandibular posterior region associated with left inferior alveolar nerve paresthesia (Figure 1).

A panoramic study (Orthopantomograph® OP100 D digital panoramic X-ray unit, Instrumentarium Dental, Tuusula, Finland), at 70 kVp, 12 mA, for 17.6 seconds shows an ill-defined, mixed-density area with loss of cortical outline of the inferior alveolar canal in the left mandibular posterior region. Based on the radiographic features and patient history, differential diagnoses of malignancy and inflammatory lesion were made. The study was interpreted by a board-certified oral and maxillofacial radiologist (Figure 2).

Furthermore, the patient underwent an MDCT examination (Aquilion One, Toshiba America Medical Systems, Tustin, California, USA) that also revealed an ill-defined, mixed-density lytic area in the left posterior mandible with an aggressive soft-tissue mass centered within the alveolar ridge, and involving the masticator, sublingual and left parapharyngeal spaces. A floating tooth is not centered within the alveolar ridge, and involving the masticator, sublingual and left parapharyngeal spaces. A floating tooth is noted as well. The inferior alveolar nerve is involved and displaced to the right. A board-certified neuroradiologist evaluated the dataset and maintained the differential diagnosis (Figure 3-5).

Histopathologic evaluation was performed with histochemical processing and interpretation. Biopsy revealed malignant plasmablastic cells with markedly increased cellular marker (Ki-67) proliferation index (90%), immuno-reactive for transmembrane heparan sulfate proteoglycan (CD138), and diffusely reactive for Epstein-Barr Encoded RNA (EBER). Moreover, malignant plasmablastic cells lacked significant immunoreactivity for B-cell markers (CD20, CD79a), paired Box 5 (PAX5), common acute lymphoblastic leukemia antigen (CALLA)/CD10, B-cell lymphoma 6 (BCL6), neural cell adhesion molecule (NCAM)/CD56, stem cell growth factor (CD117), stem cell marker (CD34), S100, Lysozyme, and T-cell marker (CD3). Overall, these findings were consistent with AIDS-associated PBL (Figure 6).

The patient received a chemotherapy regimen consisting of dose-adjusted etoposide, prednisone, oncovin (vincristine), cyclophosphamide and hydroxydaunorubicin (doxorubicin) (DA-EPOCH). He completed four cycles of DA-EPOCH; however, the left jaw mass increased in size. Therefore, chemotherapy regimen was adjusted to ifosfamide, carboplatin, etoposide (ICE). After he completed four cycles of ICE, he was scheduled for radiation therapy. He missed his follow-up appointments.

**Discussion**

PBL has been well known for its frequency in the oral cavity of HIV-positive individuals [10]. However, HIV-positive patients may experience PBL as an extraoral involvement, such as skull, paranasal sinuses, bone, skin, subcutaneous tissues, gastrointestinal tract (GIT), central nervous system, lungs, mediastinum, and various lymph nodes [5]. Lymph nodes, skin and GIT are the most affected extraoral sites [5, 10]. Similarly, oral cavity and GIT are the most involved sites in HIV-negative PBL patients [5]. The estimated frequency of oral involvement in HIV-positive is 58% whereas in HIV-negative patients is 16% [5]. Oral PBL shows a male predilection with 54.1 average age [9]. Plasmablastic lymphoma has been reported in the elderly patients who are not immuno-compromised [11, 12].

Clinically, oral cavity compromise is the most notable characteristic of PBL in AIDS patients, as noted in our patient who had a large facial mass.

Radiographically, PBL is characterized by an expansive, highly destructive, lytic lesion in the affected areas with overlying swelling of the soft tissues [10]. Moreover, MDCT shows perineural extension, lymphadenopathy as well as infiltration to the adjacent facial spaces as shown in the present case.

Histopathologically, PBL is formed by sheets of large atypical lymphoid cells [10]. The background infiltrate contains small mature lymphocytes mixed with macrophages, which gives an appearance of a ‘starry-sky’ pattern [9]. Immuno-histochemically, the PBL tumor cells are characterized by expression of the plasma cell markers VS38c, CD38, CD138, and multiple myeloma onco-

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**Figure 1. Clinical photo shows intraoral swelling in the left mandibular posterior region.**
Figure 2. Panoramic radiograph depicting a mixed density, ill-defined entity at the left mandibular posterior region. Note loss of cortical outline of the left inferior alveolar canal.

Figure 3. Bone window axial MDCT: Aggressive mixed-density osteolytic area centered within the left mandibular alveolar ridge. Note the soft tissue swelling on the affected side.

Figure 4. Bone window para-sagittal MDCT shows destruction of the left mandibular body with a floating mandibular left first molar and rampant osteolysis. Inferior alveolar neurovascular bundle is not traceable at this region.
gene-1 (MUM-1) [8]. However, there is an absence of pan B-cell markers CD20 and PAX5 [5, 8]. There is also lacking of T cell marker CD3.8 There is weak expression of leukocyte common antigen (LCA) or CD45, and CD79a [5, 8]. Moreover, the Ki67 index reveals a high proliferation rate [8]. The findings of the present case were consistent with plasmablastic lymphoma.

Clinical and histopathological features of PBL may mimic other malignancies, such as plasmablastic plasmacytoma, immunoblastic diffuse large B-cell lymphoma (immunoblastic-DLBCL), PEL and ALK-DLBCL [5].

Plasmablastic plasmacytoma is inactive neoplasm and most likely localized [13]. It lacks a starry sky pattern and has low rate of Ki67 index [13]. Patients with this neoplasm have responded well to radiation therapy and/or chemotherapy [13]. Positive CD20 and CD45 immunoreactivity exclude immunoblastic DLBCL from the differential diagnosis [5]. Primary effusion lymphoma
Plasmablastic lymphoma (PBL) represents around 4% of all HIV-related non-Hodgkin's lymphomas [14]. PBL is characterized by positive HHV8 while it involves plasmablastic lymphoma if only associated with Castleman disease [14]. Plasmablastic lymphoma most likely occurs in the oral cavity of the HIV patients [10]. It is positive-EBER and negative-ALK [15]. In contrast, ALK-DLBCL has positive ALK but does not seem to be associated with EBV and tends to occur in younger immuno-competent patients [2, 6].

PBL has a poor prognosis with or without treatment [16]. Death may occur in 1-24 months with a 6 months average survival time [10]. The treatment of PBL is mainly through chemotherapy and radiotherapy, with or without surgical excision [5]. PBL's prognosis is improved by adding highly active antiretroviral therapy (HA ART) to the treatment [16].

**Conclusion**

The present case reports radiographic findings of a case of PBL in an immunocompromised patient. Although the definitive diagnosis of PBL relies on biopsy with accurate pathologic and immunohistological testing, awareness of possible diverse findings on MDCT can prevent misdiagnosis and delayed treatment. An oral and maxillofacial radiologist must be aware of this lesion, as it is associated with immune-suppression and could be misconstrued for other pathoses [10]. Immediate referral to oncology is critical for appropriate intervention.

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