T-Cell Lymphoma of the Oral Cavity: Case Report

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ABSTRACT
Lymphomas are heterogeneous malignancies of the lymphatic system characterized by lymphoid cell proliferation. They can broadly be divided into Hodgkin’s lymphoma (HL) and non- Hodgkin’s lymphoma (NHL). NHL can originate from B, T, or natural killer (NK) lymphocytes. Extra-nodal presentation of T-cell NHL is extremely rare, and is often seen in immunocompromised individuals. Here, we report a rare case of T-cell lymphoma of the oral cavity in an immune-competent patient. The patient was diagnosed to have T-cell NHL on the basis of biopsy and immunohistochemistry, and was referred to oncology department for chemotherapy for definite treatment.

CASE REPORT
A 55-year-old male patient presented to ENT out-patient department with the complaints of a growth on gums of right side of the lower jaw since three months. The growth had rapidly increased in size, and was associated with pain. There was no history of bleeding from the growth, nor did the patient complain of fever, night sweats or weight loss. The patient was a chronic tobacco chewer, however, there was no history of smoking or of alcohol addiction. On oral examination, there was an irregular proliferative mass involving the right lower gingivo-buccal sulcus, extending from the canine tooth till the third molar, and extending on to the retromolar trigone. The growth also involved the lingual surface of mandible and the adjacent floor of the mouth. The mass was tender and firm in consistency, and did not bleed on palpation. On examining the neck, there was no palpable lymphadenopathy.

The patient underwent a contrast enhanced computed tomography (CECT) scan of oral cavity and neck which revealed a heterogeneously enhancing soft tissue density lesion involving the right buccal mucosa and inferior gingivo-buccal sulcus on right side with involvement of the retromolar trigone. The lesion measured 3.8 x 3.6 x 2.3 cm and was seen to cause destruction of the alveolar process and the body of the underlying mandible. The lesion was found to be abutting the floor of the mouth.

The patient was taken up for an incisional biopsy of the growth under local anaesthesia. Histopathological examination revealed sheets of atypical lymphoid cells reaching till the overlying epidermis. On immunohistochemistry, the tumour cells were positive for leukocyte common antigen (LCA) and negative for CD20, B-cell, EMA, cytokeratin, anti-MPO, CD68 and CD20. Thus a diagnosis of T-cell NHL of oral cavity was established. In order to stage the disease and rule out any systemic manifestations, patient underwent CECT chest and abdomen, and a bone marrow biopsy. Both radiological and haematological investigations were found to be normal indicating no systemic infiltration of NHL cells. The patient was then referred to the oncologists where he was planned for six cycles of chemotherapy (CHOP regimen) with cyclophosphamide, hydrodaunorubicin, vincristine and prednisolone. Unfortunately, the patient received only 3 cycles of chemotherapy and died due to sepsis.

DISCUSSION
Lymphomas are heterogeneous malignancies characterized by proliferation of lymphoid cells or their precursors. They can be classified as HL or non-Hodgkin’s lymphoma (NHL) [1]. NHL can further be sub-classified into B-cell, T-cell, or NK/T cell types. NHL presents more commonly than HL, representing about 86% of all lymphomas, and B-cell lymphomas are seen more frequently than T-cell lymphomas [2]. HL rarely shows extra-nodal disease (1% cases), whereas NHL presents as extra-nodal disease in approximately 23-30% of cases [3]. Extranodal NHL is commonly seen to involve gastrointestinal tract, Waldeyer’s ring, skin, bones, and others. Oral cavity involvement is seen in only 2% of cases [4]. Due to rare presentation of extra-nodal T-cell NHL in oral cavity, clinicians often face difficulties in proper management of such patients. Various oral sub-sites involving T-cell NHL as mentioned in literature include labial commissure, gingiva, palate, maxilla, buccal mucosa, floor of the mouth, gingo-buccal sulcus, tongue and uvula [2,4,5].

These tumours show a wide spectrum of behavior ranging from relatively indolent to highly aggressive and potentially fatal course [6]. It has slowly grown from a rare cancer to the fifth most common cancer in the world over a period of 30 years [7]. Our patient expired...
after three cycles of chemotherapy due to the potentially fatal nature of the disease. On reviewing the literature, mixed treatment outcomes in oral T-cell NHL were seen, and it was observed to be unrelated to the site of the tumour. Similar fatal outcomes were seen in patients having manifestations of T-cell lymphoma as ulcerative, labial swelling, mouth ulcers, buccal mucosa and upper gingiva, within three months of commencement of chemotherapy [5,8]. Complete remission with chemotherapy or radiotherapy, though less commonly seen, has been reported in patients with involvement of buccal commissure and oral mucosa by Villa et al., and May et al. [2,9].

NHL has commonly been associated with HIV, and is considered the second most common HIV-associated malignancy after Kaposi’s sarcoma. The risk of NHL is 60 times greater in patients with HIV disease than in otherwise healthy persons [10]. T-cell lymphoma has occasionally been reported with systemic conditions like celiac sprue, crohn’s disease and several autoimmune diseases [11,12]. However, the patient in our case report was not found to be suffering from HIV-AIDS, any other systemic illness or immune compromising condition. Immunohistochemistry plays an important role in the diagnosis and classification of hematolymphoid neoplasms. The peripheral T-cell lymphomas are about 90% positive for CD3 and CD45RO while B-cell markers include CD19, CD20, CD30 and CD79. Newer agents effective in early stage of presentation. Most studies recommend which improves the cure rate. Radiotherapy alone is considered as a possible differential diagnosis of any growth in the oral cavity. They can have variable presentations and can easily be confused with other malignancy like squamous cell carcinoma or with pyogenic granuloma. Histopathology and immunohistochemistry play the most important role in the diagnosis of these disorders. Since T-cell lymphomas are associated with an aggressive course, early diagnosis and prompt treatment with chemotherapy and radiotherapy is the key for success in treatment of these disorders.

CONCLUSION

T-cell lymphomas present rarely in oral cavity and should be considered as a possible differential diagnosis of any growth in the oral cavity. They can have variable presentations and can easily be confused with other malignancy like squamous cell carcinoma or with pyogenic granuloma. Histopathology and immunohistochemistry play the most important role in the diagnosis of these disorders. Since T-cell lymphomas are associated with an aggressive course, early diagnosis and prompt treatment with chemotherapy and radiotherapy is the key for success in treatment of these disorders.

REFERENCES