Precursor B- Cell Lymphoblastic Lymphoma:

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Introduction

- Precursor B- cell Lymphoblastic Lymphoma (B-LBL) is a high grade malignancy characterized as neoplasm of precursor cells committed for B cell lineage.
- It is a rare subtype of non-Hodgkin Lymphoma and consists of about 10% of lymphoblastic lymphoma cases.
- Involvement of the oral cavity in middle aged woman is especially rare.

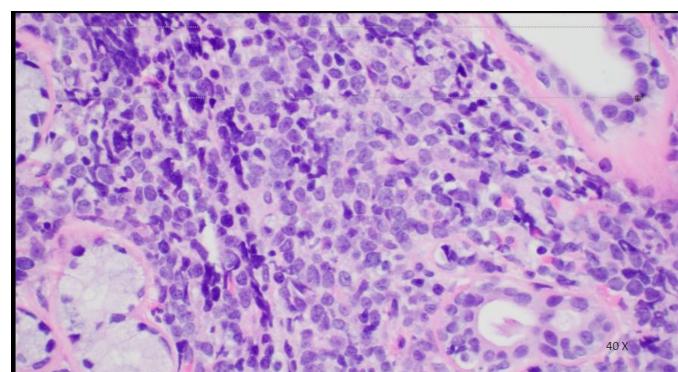
Presentation

 62 year old female with past medical history of Stage IIIB left breast cancer pT2N3a ER/PR positive, Her 2 neu negative s/p left lumpectomy with axillary node dissection s/p AACD, Abraxane, radiation to left breast on letrozole presented initially with left jaw swelling.

Figure 1. Left Posterior Mandibular biopsy demonstrates atypical lymphoid cells infiltrating into tissue parenchyma.

Work-Up

- CAT scan of the neck demonstrated lesion in the left body of the mandible and mandibular symphysis.
- She underwent direct laryngoscopy with left mandibular biopsy, and pathology revealed aggressive B- cell lymphoma
- Immunochemistry (IHC) from biopsy was positive for PAX5, CD10 and TdT and negative for CD20.
- Based on morphological and phenotypic profile, B- lymphoblastic lymphoma was diagnosed.
- Bone marrow biopsy was negative for lymphoma.
- Staging with PET scan showed new uptake in thoracic and lumbar vertebra as well as in spleen.
- Repeat mandibular and lumbar spine biopsy with CT guidance revealed similar findings
- Molecular study (FISH) is positive for mutation for BCR-ABL.



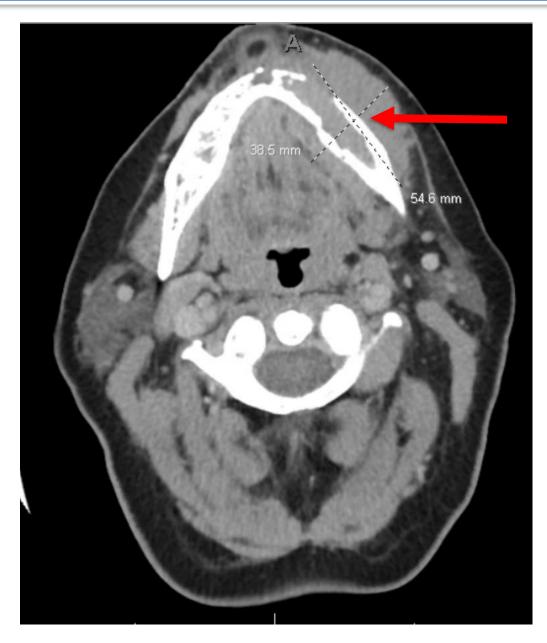


Figure 2. CAT scan of the neck which demonstrated a destructive lytic lesion in the left body of the mandible and mandibular symphysis (5.5 x 3.9 x 3.9 cm). The mass was noted to distort the buccal and lingual cortical margins of the mandible. There was increased density in the fat of the left retromolar trigone.

Outcome

Patient has completed 4 cycles of Hyper CVAD with intrathecal chemotherapy. However due to an increase in side effects, dosage was reduced. Patient is beginning stem cell transplant evaluation at Upenn.

Discussion

- This case highlights a unique presentation of B-LBL in a different demographic with a rare presentation.
- B-LBL is typically suspected in a child with painless lymphadenopathy in the setting of unexplained cytopenias, fatigue, infections, easy or spontaneous bruising/bleeding.

Conclusion

It is rare for B-LBL to present in the oral cavity without blood or bone marrow involvement. This case emphasizes the importance of recognizing rare presentations of aggressive malignancies.

References

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