



Primary Splenic Diffuse Large B Cell Lymphoma: A Rare Case Report

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Introduction

Primary splenic lymphoma is a rare neoplasm of the spleen^{1,2}. Here we present a case of primary splenic diffuse large B cell lymphoma in a 69 year old male.

Case History

A 69 year old man presented with left side abdominal pain of 6 month duration. On examination spleen palpable 5cm below left costal margin, firm in consistency with rounded margin non tender. Right inguinal lymph node enlargement 2x1cm.

Peripheral Smear: Neutrophilic leukocytosis

FNAC lymph node shows reactive hyperplasia.

USG: Massive splenomegaly with multiple focal small sized lesions.

CECT: Multiple lesions in spleen with massive splenomegaly suggestive of splenic abscess.

Splenectomy was done. Gross examination show Spleen nodularly enlarged. Cut section shows multiple irregularly gray white nodules.

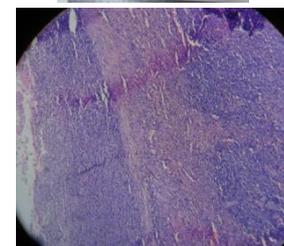


Fig 1 4x shows effacement of architecture by neoplasm arranged diffusely

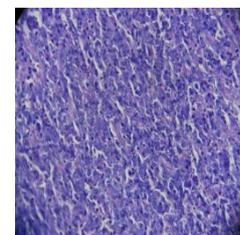
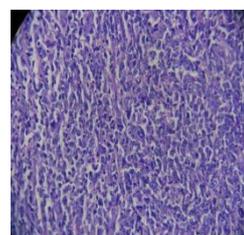


Fig 2 40X shows Individual cells are round to spindle with vesicular nucleus and distinct nucleolus.

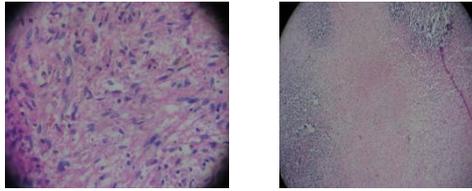
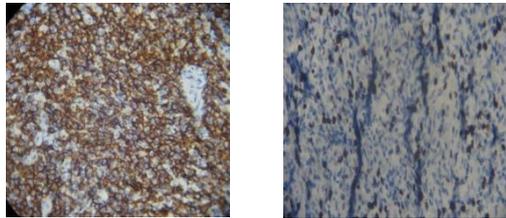


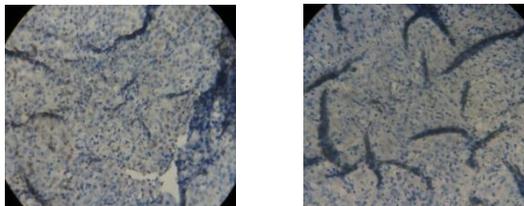
Fig 3 Granuloma and extensive areas of necrosis seen



CD 20

CD 3

Fig 4 IHC Tumor cells are positive for CD 20 and Negative for CD 3



CD 5

CD 23

Fig 5 IHC Tumor cells are negative for CD 5 and CD 23

Discussion

Primary splenic lymphoma comprises less than 2% of all the lymphoma³ and 1% of all the non Hodgkin lymphoma^{4,5}. Hodgkin and non Hodgkin lymphoma may affect spleen as a part of systemic involvement, where as primary splenic lymphoma is rather rare. Most often diagnosis is made after the histopathological evaluation of the splenectomy. A thorough clinical examination and investigation are mandatory to rule out disease in other organ including liver, Bone marrow and lymph node. Splenectomy is the most effective therapy for primary splenic lymphoma.

Conclusion

Primary splenic lymphoma is an extremely rare entity. Radiological and clinical appearance of primary splenic lymphoma may mimic splenic

abscess that may delay the proper diagnosis and management. Hence it is imperative that clinician keep this differential in their mind while dealing with similar cases and should take necessary steps like biopsy and immunohistochemical analysis to reach correct diagnosis.

References

1. Semra Ayturk et al, Primary Splenic DLBCL after splenectomy: A rare case with literature review, American Journal of Medical Case Reports, 2015, vol3 , No.8,p 265-268
2. Hanumanthappa M.B. et al, Primary Splenic Lymphoma: A rare clinical case report, Journal of Clinical and Diagnostic Research 2011 December , Vol-5(8) p1667-1669
3. Gobbi PG., Pozzetti U,et al. Primary splenic lymphoma ; does it exist? Hematologica 1994;79:286-293
4. Nuala A Conneely J B, Mahon S, O Riardon C, McAnena OJ. Primary splenic lymphoma presenting with ascities. Rare tumors 2011
5. Shao-Min H, Cheih-Lin T,Guan C, Che-An T Primary splenic lymphoma which was associated with haemophagocytic lymphohistiocytosis which was complicated with splenic rupture. J Chin Med Assoc 2008; 71(4):210-213.