



ISSN NO. 2320-5407

Journal homepage: <http://www.journalijar.com>

INTERNATIONAL JOURNAL
OF ADVANCED RESEARCH

RESEARCH ARTICLE

Non – Hodgkin's Lymphoma of Calcaneum, Inguinal & Popliteal region : A rare case report

Dr. Abhijit Sahu¹, Dr. Sohael Khan^{2*}, Dr. Shraddha Singhania³, Dr. Mahendra Gudhe⁴, Dr. Pradeep Singh⁵,
Dr. Karan Mane⁶, Dr. Mridul Arora⁷

1. Senior Resident, Orthopaedics Department, Bombay Hospital. Mumbai, Maharashtra

2. Assistant Professor, Orthopaedics Department, JNMC, Wardha, Maharashtra

3. Assistant Lecturer, Radiodiagnosis, JNMC, Wardha, Maharashtra

4. Assistant Professor, Orthopaedics Department, JNMC, Wardha, Maharashtra

5. Professor & Head, Orthopaedics Department, JNMC, Wardha Maharashtra

6. Consultant at Upasna Nursing Home, Mumbai, Maharashtra

7. Assistant Professor, Department of Orthopaedics, JNMC, Wardha, Maharashtra

Manuscript Info

Manuscript History:

Received: 15 August 2015

Final Accepted: 22 September 2015

Published Online: October 2015

Key words:

Non – Hodgkins Lymphoma,
Calcaneum, Popliteal Region,
Inguinal Region

*Corresponding Author

Dr. Abhijit Sahu

Abstract

Non-Hodgkin's lymphoma is cancer of the lymphoid tissue, which includes the lymph nodes, spleen, and other organs of the immune system^[6]. These are a heterogeneous group of B-cell and T-cell neoplasms that arise primarily in the lymph nodes.^[7] HIV and other infectious agents, such as human herpesvirus 8 and Epstein-Barr, appear to be associated with differing types of NHL, such as some B-cell lymphomas^[7]. The cancer may be low grade (slow growing), intermediate grade, or high grade (fast growing). Burkitt's lymphoma is an example of a high-grade lymphoma. Follicular lymphoma is a low-grade lymphoma. The cancer is further grouped by how the cells look under the microscope, for example, if there are certain proteins or genetic markers present.

Copy Right, IJAR, 2015,. All rights reserved

INTRODUCTION

The incidence of NHL is approximately 50% higher among men than women and 35% higher among white people than black people^[9]. Increase in NHL may be attributed to immunodeficiency, various infections, familial aggregation, blood transfusion, genetic susceptibility to NHL, diet, and chemical exposures to pesticides and solvents^[9]. Non-Hodgkin lymphoma resulted in 210,000 deaths globally in 2010 up from 143,000 in 1990.^[8]

Primary lymphoma of bone in general is a very rare condition which represents approximately 3% of all primary malignant bone tumors. Of these, less than 1% originate in the foot.^[1] Involvement of the ovary in NHL is well recognized^[11]. Male predominance, higher likelihood of early stage presentation, and a median age predominance of the fifth or sixth decade are usual features of primary bone lymphoma.^[2]

PLB constitutes 3-7% of all malignant bone tumors and approximately 3% of all extranodal lymphomas. It is found at all ages, being most frequently seen in adult life. Any part of the skeleton can be involved, but a trend exists in favor of bones with persistent bone marrow^[4]. Most tumors and tumor-like conditions initially present with pain as the patient's primary complaint. Therefore, any complaint of heel pain must be thoroughly evaluated and treated.^[5]

Although bony involvement of disseminated malignant lymphoma is not uncommon, primary lymphoma of bone is infrequent. Primary bone lymphoma is mostly large-cell or mixed small- and large-cell lymphoma of the B-cell lineage.^[3] It usually affects the long bones, indeed its very unusual presentation in this case report where it affects

the calcaneal bone. On the basis of radiological characteristics, primary bone lymphoma can be divided into three groups: (1) lytic destructive pattern, (2) blastic sclerotic pattern, and (3) subtle or near normal pattern.^[3]

In general, limb-sparing radiation therapy was considered as the gold standard in 1960s with a curing rate of 44-63%. But now patients with monostatic primary bone lymphoma treated with a combination of radiation therapy and chemotherapy do well^[2]. Most tumors and tumor-like conditions initially present with pain as the patient's primary complaint. Therefore, any complaint of heel pain must be thoroughly evaluated and treated.^[5]

CASE SUMMARY

A 22-year-old female patient was apparently alright till 3 months back when she developed pain over her right heel. It was gradually increasing and associated with swelling in the region. The pain and swelling increased over time and patient could not walk and needed support for ambulation. There was no history of trauma or fall.

On examination: - There was diffuse swelling over her 1>right heel 6 x 5 cms. Soft, tender 2>popliteal 5 x 3 cms soft, tender 3> inguinal region, which was soft in consistency and tender with local rise in temperature. Overlying skin was normal and distal arterial pulsations were normal. Active toe movements were present but ankle movements were restricted due to severe pain. (Figure 1). On investigation the all the blood counts except for TLC were under normal limits which was 10,100/cumm Granulocytes-76%, lymphocytes- 21%, Eosinophils-2% monocytes-1%. The patient was non-diabetic and the urine examination was also within normal limits.

Radiograph of the right heel was done which was suggestive of diffuse Sclerotic lesion was seen; Expansile; Osteolytic; Permeated appearance of calcaneum; however other bones of foot- normal (Figure 2). Patient was then planned for an FNAC from the calcaneum region (Figure 3 a, b, c)

Monotonous population of lymphoid cells; Atypical large non-cleaved cells ; Plentiful smudging of nuclei ; Conspicuous nucleoli ; Repetitive occurrence of the same cells ; Absence of other cell types except for small lymphocytes

Patient was then planned for an Open Biopsy from the site which had a consistent cellular and morphological appearance as of cytology. (Figure 4 a, b)

DISCUSSION

Primary Non-Hodgkin's lymphoma of bone constitutes about 3% of malignant bone tumors^[1] and 3% of extra nodal NHL. Occurrence in the small bones are rare. The common type of NHL affecting bones are small cleaved, Burkitt's type, ALCL, Lymphoblastic, Large cell type, Immunoblastic, and Hodgkin's lymphoma. It occurs at all bones, more in the bones which are known to carry haemopoiesis^[12]. The world literature search for primary NHL of the bone rarely reported primary NHL of calcaneum. The sites commonly affected are the long bones. The highlight of present case is addition of rare occurrence of primary NHL of calcaneum to the literature. Recently, Beal *et al.*^[10] studied the clinical characteristics of 82 PLB patients and reported the most common sites to be involved as the femur (27%), pelvis (15%), tibia (13%), humerus (12%), spine (9%), mandible (2%), skull (1%), scapula (1%), radius (1%), and ulna (1%). The differential diagnosis for this patient included 1) Ewing's sarcoma 2) Primary Non-Hodgkin's lymphoma 3) Small cell variant Osteosarcoma 4) Primary malignant lymphoma 5) Non-Hodgkin's lymphoma Intermediate grade, Diffuse large cell type Treatment with cyclophosphamide, doxorubicin, vincristine, and prednisone (CHOP) plus etoposide established complete remission.



Figure 1 – Suggestive of swelling over the ankle



Figure 2 – Lateral and Axial view of the Right Calcaneum

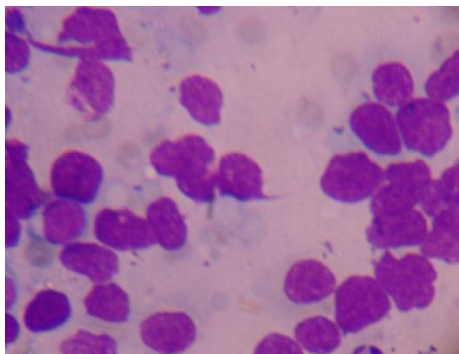


Figure 3 a - FNAC from calcaneum, scanner

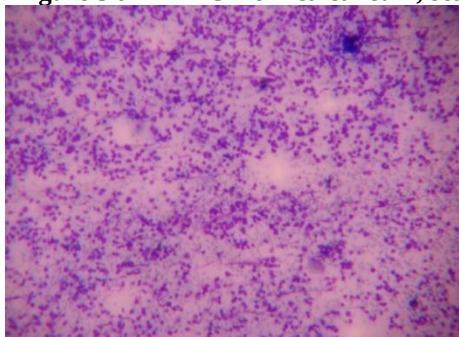
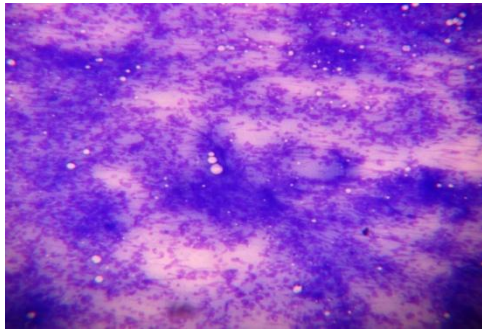
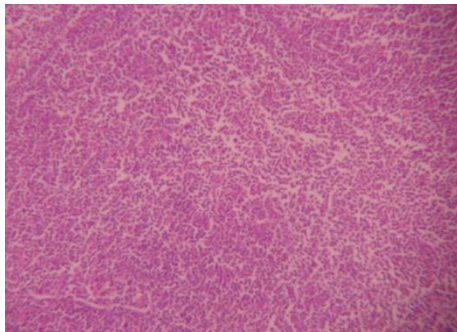
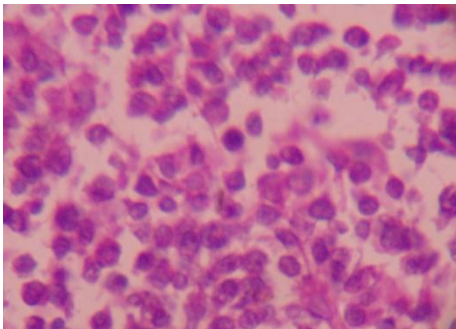


Figure 3 b - FNAC from calcaneum, 40X**Figure 3 c FNAC From popliteal swelling, scanner.****Figure 4 a Histopathology, scanner****Figure 4 b Histopathology, 40X****REFERENCE :-**

1. Skorman SE, Martin R. Primary lymphoma of the calcaneus with recurrence in the distal tibia: a case report. *Foot Ankle Surg.* 1999 Jul-Aug;38(4):278-82 PMID:10464723.
2. Beal K, Allen L, Yahalom J. Primary bone lymphoma: Treatment results and prognostic factors with long-term follow-up of 82 patients. *Cancer* 2006;106:2652-6.
3. Krishnan A, Shirkhoda A, Tehranzadeh J, Armin AR, Irwin R, Les K. Primary bone lymphoma: Radiographic-MR imaging correlation. *Radiographics* 2003;23:1371-83; discussion 1384-7.
4. Pinheiro RF, Filho FD, Lima GG, Ferreira FV. Primary non-Hodgkin lymphoma of bone: An unusual presentation. *J Can Res Ther* 2009;5:52-3
5. Berlin SJ, Mirkin GS, Tubridy SP. *Clin Podiatr Med Surg.* 1990 Apr;7(2):307-21. Tumors of the heel.
6. National Cancer Institute: PDQ Adult Non-Hodgkin Lymphoma Treatment. Bethesda, MD: National Cancer Institute. Date last modified 09/28/2012. Available at <http://www.cancer.gov/cancertopics/pdq/treatment/adult-non-hodgkins/HealthProfessional>. Accessed 01/04/2013.

7. Int J Cancer. 2007;120 Suppl 12:1-39.The non-Hodgkin lymphomas: a review of the epidemiologic literature. Alexander DD, Mink PJ, Adami HO, Chang ET, Cole P, Mandel JS, Trichopoulos D. Exponent Health Sciences Group, Exponent, Inc., Wood Dale, IL, USA. PMID:17405121
8. Lozano, R (2012 Dec 15). "Global and regional mortality from 235 causes of death for 20 age groups in 1990 and 2010: a systematic analysis for the Global Burden of Disease Study 2010.". Lancet380 (9859): 2095–128. [PMID 23245604](#).
9. Clin Lymphoma. 2003 Dec;4(3):161-8.An update of the epidemiology of non-Hodgkin's lymphoma. Chiu BC, Weisenburger DD. PMID:14715098
- 10.Beal K, Allen L, Yahalom J. Primary bone lymphoma: Treatment results and prognostic factors with long-term follow-up of 82 patients. Cancer 2006;106:2652-6.
- 11.Chorlton I, Norris HJ, King FM. Malignant reticuloendothelial disease involving the ovary as a primary manifestation: a series of 19 lymphomas and 1 granulocytic sarcoma. Cancer 1974;34:397–407.
- 12.Pinheiro RF, Filho FD, Lima GG, Ferreira FV. Primary non-Hodgkin lymphoma of bone: An unusual presentation. J Can Res Ther 2009;5:52