Rare association of Visceral leishmaniasis with Hodgkin’s disease: A case report

Rakshit Kumar1*, Mradul K Daga1, Nitish L Kamble1, Arpit Sothwal1, Tejinder Singh2, Hemanta K Nayak1 and Nishant Raizada1

Abstract
We present here a case of young male with complaints of fever and swelling in the neck for eight months. History of progressive weakness associated with weight loss was present. Physical examination revealed pallor, multiple enlarged cervical lymph nodes and hepatosplenomegaly. Investigations showed pancytopenia, hyperglobinemia and Leishman-Donovan bodies on bone marrow aspiration. Serological test confirmed diagnosis of visceral leishmaniasis. However, cervical lymph node aspiration and biopsy were suggestive of Mixed cellularity Hodgkin’s disease. This made it a very rare case of Leishmaniasis as an opportunistic infection in a patient of pre-chemotherapy Hodgkin’s disease. There was marked improvement in haematological profile and regression of hepatosplenomegaly with Amphotericin B treatment followed by favourable response to chemotherapy. The case emphasizes the suspicion for leishmaniasis as a masquerader and as an opportunistic infection in haematological malignancies.

Keywords: Visceral leishmaniasis, Hodgkin’s disease

Leishmaniasis is frequently reported in patients with immunocompromised states due to HIV infection, post organ transplantation, solid tumors and chemotherapy [1,2]. However there are very few case reports of leishmaniasis with haematological malignancies, and even fewer with lymphoma, particularly Hodgkin’s disease [3-6]. To the best of our knowledge, there are three case reports of leishmaniasis in post-chemotherapy Hodgkin’s disease patients and only two case reports [7,8] of leishmaniasis in patients of Hodgkin’s disease, pre-chemotherapy [9]. The case presented herein is a rare case of association of visceral leishmaniasis and Hodgkin’s disease right from the beginning and delves into the pathological and practical implications of their coexistence [10,11].

The patient, 18-year-old male, presented with a history of fever and multiple swellings in left side of neck for last eight months. Fever was moderate-to-high grade, intermittent, associated with sweating, without any chills or rigors. The patient noticed progressively enlarging swellings in the left side of neck which were painless, without any change on the overlying skin. In the last two months, the patient also experienced easy fatigability, weakness and swelling in both feet. Patient’s father noticed weight loss which was not documented. The patient had been on irregular treatment from local practitioner without any relief in symptoms. The patient did not give any history of tuberculosis or any other systemic illness. There was no history of sexual contact, blood transfusion, or drug addiction and no significant family history was present.

On general physical examination, the patient was conscious, alert, and was of thin build. Vitals were stable and temperature was 39°C. Marked pallor was present with pitting pedal edema. There were multiple enlarged cervical lymph nodes in posterior triangle of the neck on left side. Lymph nodes were firm in consistency, varied in size (2-4 cm), were discrete and were mobile, with normal overlying skin and temperature. On systemic examination, only mild hepatomegaly and moderate splenomegaly were seen.

Laboratory reports revealed pancytopenia with haemoglobin of 2.7 gm/dl, platelet count of 1.31 lacs, leucocyte count of 3500, ESR-70, and peripheral smear showed...
microcytic hypochromic anaemia. Total serum proteins were raised with albumin being low (2 gm/dl). Ultrasonography of abdomen revealed hepatomegaly (liver span 18 cm) and splenomegaly (splenic span 20 cm). Other investigations including chest X-ray, electrocardiography, viral markers and HIV serology were normal. CECT chest and abdomen revealed hepatospleno megaly, minimal ascites and minimal pleural effusion. Echocardiography was essentially normal. FNAC of cervical lymph node was inconclusive and was repeated followed by excision biopsy. Meanwhile, bone marrow aspiration was done, which showed intracellular and extracellular Leishman-Donovan bodies (Figure 1). This was followed by a positive serology test for leishmaniasis. Patient was immediately started on Amphotericin B and given three units of blood transfusions. FNAC and excision biopsy reports confirmed the diagnosis of Mixed cellularity Hodgkin’s disease (Figure 2 and 3), making it an exceptional coexisting pathology. Marked clinical and hematological improvement was seen, with regression of hepatospleno megaly. The patient received three cycles of chemotherapy and showed positive response.

The clinical forms of leishmaniasis are visceral leishmaniasis (VL), cutaneous leishmaniasis (CL) and mucocutaneous leishmaniasis (ML). Out of 1.5-2 million infected patients worldwide, 90% of CL cases are in the endemic zone of Afghanistan, Pakistan, Middle East, Brazil and Peru. However, 90% of VL cases are seen to occur in the endemic zone of India, Bangladesh, Nepal, Sudan and Brazil. Cervical lymphadenopathy is uncommon in India though common in Mediterranean countries, Africa and China. On the other hand, incidence of malignancy is 11 million new cases per year, worldwide, with many cases occurring in the endemic zone of leishmaniasis. Moreover, the medical management of malignancy induces further immunosuppression, a condition which is already present in the setting of lymphoma and leukaemia, and becomes a breeding ground for opportunistic infections like leishmaniasis.

Leishmaniasis has emerged as a severe opportunistic infection in endemic areas. In immunocompromised hosts, it can be asymptomatic, often with no splenomegaly, and may be present in unusual location and run a severe and refractory course with frequent relapse. Although leishmaniasis and cancer are very common diseases, little attention has been paid to the pathophysiological and practical implications of their co-existence. Leishmaniasis is recognized as an opportunistic infection in HIV and malignancy patients [1,2], but reports with lymphoma are scarce [3-5]. There are three case reports of leishmaniasis in post chemotherapy Hodgkin’s disease patients, including the recently published study on coexistence of both in a single lymph node, after multiple cycles of chemotherapy [6]. To the best of our knowledge, there are only two case reports [7,8] (Benhamou et al [7] and Magnan et al [8]) of

![Figure 1 Bone marrow slides showing Leishman-Donovan bodies (arrow)](image1)

![Figure 2 Mixed cellularity seen in Hodgkin’s disease](image2)

![Figure 3 Immunohistochemistry showing CD30 marker (arrow) in the lymph node](image3)
leishmaniasis with pre-chemotherapy Hodgkin’s disease, similar to our case.

In a significant review involving 37 case studies and 44 patients, Kopterides et al (2007)[9] found four types of association between leishmaniasis and malignant disorders: 1) Leishmaniasis masquerading as a malignant disorder, 2) Leishmaniasis developing as a difficult-to-diagnose/treat infection among patients receiving chemotherapy for various malignant disorders; 3) Simultaneous diagnosis of leishmaniasis and a neoplastic disorder in the same tissue samples of immunocompromised patients, and 4) Direct involvement of Leishmania spp. in the pathogenesis of cancerous lesions. Mangoud et al [10] reported that dysplasia was detected surrounding the leishmanial ulcer in 5 of 35 CL cases. Leishmaniasis adversely affects the activation and function of macrophages and dendritic cells [11], permitting the escape of malignant cells from immune destruction. Further, chronic leishmanial infections cause CD4 lymphopenia and low CD4/CD8 ratio. Similarly, Hodgkin’s/Reed-Stenberge cells induce regulatory T cells that impair adaptive immune responses by releasing IL-10 and TNF B. Both the cytokines block the activation of naïve and effector T cells, and have been reported to suppress the immune response to leishmania. Hence, mutual immunomodulation of leishmaniasis and Hodgkin’s disease is highlighted in this patient and requires further epidemiological and pathology studies to understand the exact association or causal relation.

Consent
Written informed consent was obtained from the patient for publication of this case report. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

List of abbreviations
HIV: Human Immunodeficiency Virus; ESR: Erythrocyte Sedimentation Rate; CECT: Contrast Enhanced Computed Tomography; FNAC: Fine Needle Aspiration Cytology; CL: Cutaneous Leishmaniasis; IL-10: Interleukin-10; TNF B: Tumor Necrosis Factor Beta.

Acknowledgements
The authors are grateful to Department of Pathology, Maulana Azad Medical College for their contribution in diagnostic evaluation.

Author details
1Department of Medicine, Maulana Azad Medical College, Bahadur Shah Zafar Marg, New Delhi, 110002, India. 2Department of Pathology, Maulana Azad Medical College, Bahadur Shah Zafar Marg, New Delhi, 110002, India.

Authors’ contributions
RK: Conceptualization of the manuscript, search of literature, and drafting and writing the manuscript, head of the resident team managing the patient
MKD: Vital inputs in drafting and editing of the manuscript, supervised clinical management of the patient being the head of the medical unit
NLK: Helped in literature search, editing and proofreading the manuscript
AS: Helped in literature search, editing and proofreading the manuscript

TS: Provided pathological insights in the case, with his vast experience on past such cases
HKN: Part of the investigative team involved in clinical management
NR: Part of the investigative team involved in clinical management
All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

Received: 10 June 2011 Accepted: 20 October 2011
Published: 20 October 2011

References
1. Frances C, Merle Beral H, Franceschini P, Lessana-Leibowitch M, Escande JP: Kalazar in immunosuppressed patients. Presse Med 1984, 13(40):2433-6.
2. Fernandez-Guerrero ML, Aguado JM, Buzon L, Barros C, Montalbán C, Martin T, Bouza E: Viscera leishmaniasis in immunocompromised hosts. Am J Med 1987, 83(6):1098-1102.
3. Derêu J, Duong Thanh H, Lavabre-Bertrand T, Carton G, Bastides F, Richard-Lenoble D, Dedet JP: Visceral leishmaniasis: Persistence of parasites in lymph nodes after clinical cure. J Infect 2003, 47:77-81.
4. Kae J, Norgiaard P, Himmelsrump B: Visceral leishmaniasis diagnosed in a patient with MALT lymphoma. Eur J Intern Med 2007, 18:235-237.
5. Boutous N, Hawkins D, Nelson M, Lampert IA, Naresh KN: Burkitt lymphoma and leishmaniasis in the same tissue sample in an AIDS patient. Histopathology 2006, 48:880-881.
6. Domingues M, Menezes Y, Ostronoff F, Calixto R, Florencio R, Sucupira A, Souzo-Maier AP, Ostronoff M: Coexistence of Leishmaniasis and Hodgkin’s Lymphoma in a lymph node. Journal of Clinical Oncology 2009, 27(32):e184-e185.
7. Benhamou E, Ferrand B, Chiche JC, Sudaka C, Maladie de Hodgkin et Kala-Azar, Remarques cliniques, biologiques, the’rapeutiques et anatomo-pathologiques. Presse Méd 1955, 86:1778-1781.
8. Magnan A, Jacqueme R, Sudan N, Redding E, Blanc AP, Poirier R: Visceral Leishmaniasis associated with Hodgkin’s lymphoma: diagnostic difficulties. Rev Prat 1991, 41(4):188-91.
9. Kopterides P, Mourtzoukou EG, Skopelites E, Tsavaris N, Falagas ME: Aspects of the association between leishmaniasis and malignant disorders. Transactions of the Royal Society of Tropical Medicine and Hygiene 2007, 101:1181-1188.
10. Mangoud AM, Sanad EM, Fouad MA, Morsy TA: Proliferative changes of epidermal cells in lesions of cutaneous leishmaniasis. J Egypt Soc Parasitol 2003, 33:61-72.
11. Brandonisio O, Spinelli R, Pepe M: Dendritic cells in Leishmania infection. Microbes Infect 2004, 6:1402-1409.

doi:10.1186/1750-9378-6-17
Cite this article as: Kumar et al.: Rare association of Visceral leishmaniasis with Hodgkin’s disease: A case report. Infectious Agents and Cancer 2011 6:17.