Acute HTLV-1 leukemia/lymphoma in a 33 year old grenadian migrant: A case report

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ABSTRACT

We present a case of an acute HTLV-1 leukemia in a 33-year-old Grenadian immigrant. Her diffuse skin nodules, and her coming from an area endemic for HTLV-1, namely the Caribbean made the clinical diagnosis of HTLV-1 leukemia/lymphoma more feasible. Her disease was rapidly progressive, she having survived for one month, and fits into the acute classification of ATLL as proposed by Shimoyoyama M. Immigrants from areas endemic for HTLV-1 who presents with skin rashes should be investigated for HTLV-1 neoplasms.

Key words: HTLV-1 leukemia/lymphoma; Skin manifestation; Endemic areas; Migrants

INTRODUCTION

HTLV-1 is a retrovirus which is associated with HAM/TSP [1-3] and which causes leukemia/lymphoma [4].

The neoplastic process presents in the early stages with a skin lesion as the primary site [5,6] and progresses to the advance stages with multiple organs involvement of the reticuloendothelial system such as the liver, spleen, and, bone marrow. There are indeed other presentations as in the case in discussion that presented as an acute leukemia with diffuse skin involvement.

CASE REPORT

A 33-year-old Grenadian national was on a six months visit to Trinidad when she developed a generalized pruritic skin rash and abdominal pain following consumption of shrimps. She visited the General Hospital for the above complain three days following the above symptoms.

On examination she was found to be icteric and with pale mucous membrane. Her pulse was 80/min, blood pressure 90/60 mm Hg, respiratory rate 15/min and temperature 36.5°C.

She was not in discomfort. She had generalized nodules distributed on the upper and lower limbs, chest and face (Figs. 1 and 2). They were non-erythematous, and non-tender. No cervical, axillary or inguinal lymph nodes were palpable.

She was mildly tender in the left hypochondrium, with a markedly palpable spleen.

Her genitourinary and central nervous systems examination were essentially normal. She was started on analgesic and antipruritic oral medications and was admitted to the medical ward for further investigation.

Laboratory investigations revealed an Hb of 7.5 g/dl (11.7-15.5), hematocrit of 27.1%. (9 34.5-46.30), MCHC 27.6 g/dl (342.5-35.2), MCV 83 fl (80-100-95.5) MCH 22.9 pg. (27.2-33.5), WBC 131.6 10^3/UL (4.10-11.20), with 83.6% Lymphocytes. (18.8-50.8), 13.3% (39.9-73.9) Neutrophils, 7.9% (0.3-1.8) Basophils, 0.1% (0.9-6.0) Eosinophils, 2.9% (0.3-0.9) Monocytes, platelets 263 x 10^3/UL. (159-388.0).

There were polylobated lymphocytes, “flower cell” in the peripheral blood smear, but no hairy cells or blasts. She was hypercalcemic, calcium 13.1 mg/dl, Na+ 127 mmol/L., K+ 4.8 mmol/L,
Chloride 90 mmol/L., AST 91 IU/L., ALT 16 IU/L., LDH 2901 U/L., GGT 68 U/L., Alkaline phosphatase 123 U/L., Total protein 6.0 g/dl, Albumin 3.3 g/dl, Globulin 2.7 g/ dl (3.5-5.5), Total Bilirubin 4.6 mg/dl, direct bilirubin 1.2 mg/dl, indirect bilirubin 3.4 mg/dl.

A bone marrow aspirate and trephine biopsy showed an infiltrate of malignant T-lymphocytes. There were no blast cells.

A skin biopsy of the nodule revealed a moderate infiltrate of small to medium sized lymphocytes with pleomorphic nuclei and sparse minute eosinophilic nucleoli arranged in a nodular fashion involving the dermis and subcutaneous tissue (Fig. 3). Admixed with this lymphocytic infiltrate were histiocytic cells (Fig. 4). Epidermotropism was absent.

The lymphocytes expressed T-cell markers with CD3, CD5, and CD4 positivity and were negative for CD7 and CD34.

Her HTLV-1 was positive. She was started on hydroxyurea and referred to the oncologist for further management.

The patient died one month after diagnosis while undergoing chemotherapy.

DISCUSSION

Human T-cell Lymphotrophic virus-1 is endemic in the Caribbean, Japan [7,8] and is seen now in most countries amongst immigrants from endemic countries [9,10].

The virus causes HAM/TSP and is the causative agent of HTLV-1 leukemia/lymphoma [1].

The acute, chronic, smoldering and lymphoma are the four different clinical manifestations of the neoplasm that have been identified [2].
We describe here a Grenadian national who presented in Trinidad W.I with splenomegaly and an acute HTLV-1 leukemia, having a white blood cell count of 131 x 10^3/UL with 83.6% atypical lymphocytes within the peripheral blood and a diffuse nodular skin lesion which histologically contained malignant HTLV-1 infiltrate. Hairy cell leukemia and chronic myeloid leukemia both have splenomegaly of the magnitude seen in this patient. However these two leukemia were eliminated from the diagnosis on account of the HTLV-1 positivity, the immuno-histology of the malignant cells, the absence of hairy cells and myeloblast in the peripheral blood and bone marrow aspirate. The skin nodules were composed of malignant infiltrate of HTLV-1 cells, which had no correlation with the consumption of shrimps. In addition to the lymphocytosis, she also presented with hypercalcaemia and a markedly elevated LDH, which are poor prognostic markers. This patient only survived one month after diagnosis.

HTLV-1 skin manifestation are varied and includes infective dermatitis, maculo-papular hyper pigmented or hypo pigmented rashes, nodules or ill-defined rashes, which mimics Mycosis fungoides, Psoriasis, Eczema and a list of common dermatoses, [11] and may be an early presentation of this neoplastic disease entity [5,6].

Clinicians should have a high index of suspicion of HTLV-1 related disease in patients from endemic regions who presents with a skin rash.

CONCLUSION

HTLV-1 is endemic in the Caribbean and Japan. Immigrants presenting with a skin rash from endemic regions should be investigated for HTLV-1 Leukemia/ Lymphoma and it’s other related diseases [12].

REFERENCES