Microfilaria in Association with Adrenal Lymphoma Diagnosed on Cytology: An Extremely Rare Case Report

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ABSTRACT
Filariasis is a common public health problem in Asian countries. In this report, the authors have described detection of microfilaria in the cytological specimen of adrenal lymphoma. To the best of our knowledge, this is the first reported case in the literature where microfilaria was seen in an adrenal gland involved by lymphoma. The literature is briefly reviewed.

Key Words: Microfilaria, Adrenal, Lymphoma, Fine-needle biopsy

INTRODUCTION
Filariasis is a common health problem in tropical countries like India (1). In India, it is caused mainly by Wuchereria bancrofti and Brugia malayi (2,3). Culex mosquito is the most important (>50%) vector of Wuchereria bancrofti (4). The literature has few reports of incidental detection of microfilaria in many of the commonly evaluated cytological specimens such as thyroid, breast, pancreas and in body fluids associated with neoplastic lesions (6,7,8). However, to the best of our knowledge, microfilaria associated with lymphoma of the adrenal gland has never been reported. The present report is the first case in the literature where microfilaria was seen in the adrenal gland involved by lymphoma.

CASE REPORT
A 55-year-old diabetic male presented with complaints of fever, generalized weakness, weight loss and pain in the abdomen for the last 2 months. On physical examination there was blackish pigmentation over the face, palm, knuckles, feet and oral mucosa. There was no cervical, axillary and inguinal lymphadenopathy, hepatomegaly or splenomegaly. However, ultrasonography (USG) revealed enlargement of adrenal glands as well as retroperitoneal lymph nodes. The patient underwent USG-guided fine needle aspiration (FNA) of an adrenal gland and retroperitoneal lymph nodes.

Smears prepared from USG-guided FNA using a 21 gauge needle from both retroperitoneal lymph nodes and adrenal gland were stained with air dried May-Grunwald Geimsa and ethanol-fixed Papanacolaou stains.

Smears from retroperitoneal lymph node and adrenal glands were suggestive of lymphoma. The FNA smears from the left adrenal were paucicellular and showed the presence of few scattered atypical lymphoid cells in the hemorrhagic background along with a few microfilariae. These microfilaria showed nuclei in the centre and a tip free of nuclei, and there was a sheath projecting beyond the body of the larvae (Figure 1). Laboratory investigation revealed a hemoglobin of 9.4 gram %, total leukocyte count of 9,700/mm3 and platelet count was 250000 per cmm. Peripheral blood smear examination showed a differential of polymorphs 70% and lymphocytes 30%. There was no eosinophilia, parasites or atypical cells. Bone marrow aspiration and biopsy did not show any evidence of lymphomatous infiltration and were within normal limits.

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Histopathological and immunohistochemical confirmation of Non-Hodgkin’s B cell lymphoma was done as advised on cytology. Although no microfilaria was seen in the paraffin-embedded sections, the sections from both the lymph node and adrenal biopsy showed tumor composed of sheets of atypical lymphoid cells displaying round to oval nuclei, dispersed chromatin, inconspicuous nucleoli and scant cytoplasm (Figure 2). Mitotic activity was occasional. On immunohistochemistry, the tumor cells were positive for Leucocyte common antigen (CD45) (Figure 3) and CD20 and were negative for pancytokeratin and CD3.

**DISCUSSION**

Microfilaria is a nematode belonging to the superfamily Filarioidea and is found in the blood and skin. Lymphatic filariasis is a major health problem in tropical and subtropical countries in Asia and the Pacific Island (1). It is estimated that about 600 million people live in areas endemic for lymphatic filariasis in Southeast Asia (2). Eight filarial species infect the human but serious infestation is caused by only four, i.e. *Wuchereria bancrofti*, *Brugia malayi*, *Onchocerca volvulus* and *Loa loa* (1,3).

*Wuchereria bancrofti* is the most common causative organism, accounting for about 95% of all filarial infections. It was first reported in 1876 by Bancroft in Brisbane and the term filarial bancrofti was given in 1977. The life cycle is completed in two hosts. Man is the definitive host and the female *Culex quinquefasciatus*, the most important (>50%) vector of *Wuchereria bancrofti*, acts as an intermediate host (4).

When a Culex mosquito bites an infected individual, the sheathed microfilarias are ingested by the mosquito. These microfilarias transform into first, second and third staged larvae. These infected third-stage larva are inoculated in uninfected humans by a mosquito bite and develop into the adult worm in the lymphatic system. This prepatent period of growth of third-stage larvae to adult worm lasts for about one to one and half years and is seldom symptomatic. The chronic stage may develop in some patients, while others may remain asymptomatic for many years (5). The chronic stage usually develops 10 to 15 years later, in persons exposed continuously to reinfection. The most commonly...
involved lymphatics are those of lower limb, retroperitoneal tissue, spermatic cord, epididymus and mammary gland. The diagnosis of microfilaria is conventionally made by demonstration of microfilaria in the peripheral blood smear. It causes a spectrum of disease including lymphangitis, lymphadenitis, hydrocele, chyluria and elephantiasis. In occult filariasis, adult microfilarias are not present in the blood but are seen in the affected tissue. The diagnosis in asymptomatic and occult filariasis is therefore difficult by blood examination.

A review of the literature reveals incidental detection of microfilaria in many of the commonly performed cytological specimens and has been described in almost all tissues of the body. Thyroid, breast, pancreas and body fluids associated with neoplastic lesions are a few examples (6,7,8,9). The adrenal gland is another rare site that is also involved by lymphoma. To the best of our knowledge, this is the first case reported in the literature where microfilaria were seen in adrenal gland involved by lymphoma. Their presence in association with tumors involving lymph nodes and lymphatics can be explained by their being normal habitation for the filarial organism. However, this view does not completely explain their association with tumors from other sites. As the retroperitoneal lymph node also revealed lymphomatous infiltration in this case, the possible explanation is that it might have infiltrated from its normal habitant lymph node via the vascular supply to the adrenal gland along with the atypical lymphoid cells. Increased concentration at the tumor site and subsequent rupture and hemorrhage causes release of parasite into the tumor tissue. However, the retroperitoneal lymph node showed no evidence of microfilaria on cytology as well as on biopsy.

In conclusion; Fine needle aspiration cytology (FNAC) is not the method of choice for diagnosis of microfilaria unlike the peripheral blood smear examination. However, cases have been detected as a co- incidental finding on cytology like the index case where the patient was subjected for FNAC to diagnose or confirm some other lesion. The illustrated case is the first case reported in the literature to the best of our knowledge where microfilaria is associated with adrenal lymphoma and highlights the value of proper and careful screening of the cytological smears in the detection of asymptomatic and clinically unsuspected cases of Bancrofti filariasis even if smears are hypocellular, hemorrhagic or display atypical cells. Timely institution of the specific therapy obviates the severe manifestation of lymphatic obstruction.

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