Cutaneous Erysipeloid Leishmaniasis: An Unusual Presentation

Benahmed Jihane 1*, Hamich Soumaya 1, Kaoutar Znati 2, Meziane Mariam 1, Ismaili Nadia 1, Benzekri Laila 1 and Karima Senouci 1

1 Department of Dermatology, University Hospital Ibn Sina, Rabat, Morocco
2 Department of Histopathology, University Hospital Ibn Sina, Rabat, Morocco

*Corresponding author: Jihane Benahmed, Department of Dermatology, University Hospital Ibn Sina, Rabat, Morocco, Tel: 212651519363; E-mail: jihanebenahmed3@gmail.com

Received date: November 26, 2019; Accepted date: December 17, 2019; Published date: December 24, 2019

Abstract

Cutaneous leishmaniasis is a parasitic disease caused by Leishmania species that are transmitted through Phlebotomus vector. There are several atypical forms of CL that have been described, which may cause some difficulties in diagnosis and treatment. In this report, we present an 80-year-old male patient with erysipeloid leishmaniasis, a rare atypical form of cutaneous leishmaniasis, who was successfully treated.

Keywords: Leishmaniasis; Phlebotomus; Lutzomyia; Erysipeloid type

Introduction

Leishmaniasis is a tropical disease caused by an intracellular parasite transmitted to humans by the bite of a sand fly, mainly Phlebotomus and Lutzomyia. It is endemic in Asia, Africa, America, and in the Mediterranean region. Depending on parasite and host related factors, the disease can present in three ways: visceral, cutaneous or mucocutaneous forms. In Morocco, cutaneous leishmaniasis is caused by three important Leishmania species [Leishmania major, Leishmania Tropica and Leishmania infantum]. Different clinical presentations have been described. We report a case of an unusual presentation of CL: The erysipeloid type.

Case Report

An 80-year-old male patient with no medical history, presented with erythematous plaque on the left forearm. Initially, there was a painless papule (1 cm) on his wrist which slowly coalesced and formed a red infiltrative plaque in the past 2 months. The patient was treated by oral antibiotics without any improvement. Physical examination revealed an erythematous infiltrative plaque on the left forearm (Figure 1).

The patient was apyretic and no other systemic abnormalities were detected. Total blood count and inflammatory evaluation tests were normal. A biopsy revealed the presence of a non-caseating formation in the upper dermis and amastigotes forms of Leishmania (Figure 2) which were also found in the direct microscopic examination of smears. Subtype analysis couldn’t be performed. The patient was first treated with clarithromycin 15 mg/kg/j during 6 months with a small improvement. A treatment with systemic pentavalent antimony (10 mg/kg per day) administered intramuscularly was conducted. The patient received only a total of 6 injections. The treatment was interrupted because the evolution was marked by regression of symptoms within only 6 injections (Figure 3).

Figure 1: Erythematous infiltrative plaque on the forearm.

Figure 2: Histopathology showing granuloma formations and amastigotes forms of Leishmania.

Figure 3: Clinical appearance of the patient after treatment.
Discussion

Leishmaniasis is a disease affecting about 89 countries and is endemic in Asia, Africa, parts of North and South America and the Mediterranean region. Several forms of cutaneous leishmaniasis are described. They are mainly caused by Leishmania Major, L. infantum and L. Tropica.

After a bite by an infected female phlebotomine sand fly, cutaneous lesions occur at the site of inoculation after an incubation period. The most common presentation is noduloulcerative lesions localized on the exposed areas of the body. The lesions are usually painless. Regional lymphadenopathy may occur. In a few months, spontaneous healing can occur with filling of ulcer, leaving a clear or pigmented scar [1]. However, other clinical presentations of cutaneous melanomas have been reported such as psoriasiform, eczematous, varicelliform, zosteriform, erysipeloid, lupoid and sporotrichoid forms, depending on the host immune status and the subspecies of Leishmania.

Cases of cutaneous leishmaniasis in its erysipeloid form have been reported in Iran, Pakistan, Turkey and Tunisia [2-6]. This type usually affects elderly females and presents as erythematous infiltrative plaque over the face and resembling erysipelas. The etiology of this type is unknown although factors such as senility, specific species of leishmania, hormonal changes, and changes in skin barrier with ageing can cause such an unusual presentation. Our patient is from an endemic region of leishmaniasis and is elderly, thus, he presented with all the epidemiological features.

The diagnosis is confirmed by the demonstration of amastigotes in a Giemsa-stained smear. When the parasite cannot be detected on smear, histopathologic examination can be used.

In chronic lesions, there can be a variable degree of epithelial hypertrophy and the dermis may contain large number of small non-casing granulomas. Parasites are also present into macrophages and can be detected in early lesions. In our case, the histopathological study shows chronic inflammatory infiltration, non-casing granulomas in the dermis and amastigotes form of Leishmania, confirming the diagnosis.

Systemic treatment is recommended in cases of multiple or severe lesions (diameter >5 cm).

The parental options for systemic therapy include amphotericin B, pentavalent antimonial and pentamidine. There are also oral options such as azithromycine, miltefosine, ketoconazole and fluconazole [7]. The choice of antileishmanial agent, dose, and duration of therapy should be individualized. Our patient was first treated by clarithromycine for 6 months with small improvement. He received afterwards systemic treatment with parental pentavalent antimony (10 mg/kg per day dosage). However, regarding the biological adverse side effects observed (hepatotoxicity), the treatment was interrupted after a total of 6 days of systemic pentavalent antimony. Regression of the symptoms was seen one month later. In some settings, treatment with pentavalent antimony for as few as 10 days has been effective.

Although, pentavalent antimony has the disadvantages of multiple injections and mild-to-moderate clinical toxicity, it’s still an effective drug for cutaneous leishmaniasis and could be proposed for less than 10 days [7].

Conclusion

Cutaneous leishmaniasis is a parasitic disease; it shows a variety of morphologic pattern and may mimic other inflammatory, neoplastic and infectious diseases. Erysipeloid leishmaniasis is a rare presentation that typically occurs on the face or arms, looking like erysipelas, in patients who are native from an endemic region of cutaneous leishmaniasis. Pentavalent antimony is an effective drug for the treatment of cutaneous leishmaniasis but can cause adverse side effects therefore patients should be regularly monitored.

References