

Case Report





Chronic unilateral leg warty lesion: clinical image and literature review

Abstract

54 years old diabetic male presented with history of with painless multiple progressive warty lesions on the right lower two third of the left leg remain undiagnosed for twelve years. It was spreading and growing relentlessly, not responding to medication prescribed by multiple doctors and patient thought of a uncurable disease. The biopsy came confirmatory. Antifungal therapy along with surgical excision hasten up the healing. An adequate biopsy is a must in undiagnosed skin lesion. Specific medical therapy and adjuvant surgery give early relief.

Keywords: leg lesion, painless warty lesion, long-duration lesion, chromoblastomycosis

Volume 6 Issue 2 - 2022

Aswini K Pujahari

Department of Surgical Gastroenterology, Saptagiri Institute of Medical Sciences, India

Correspondence: Aswini K Pujahari, Prof and head, Department of Surgical Gastroenterology, Saptagiri Institute of Medical Sciences, Bangalore 560090, India, Tel 09632469433, Email akpuja@rediffmail.com

Received: February 25, 2022 | Published: March 22, 2022

Case report

A 54-year-old male, a paddy field worker getting often gets minor injuries on the legs, had presented with progressive cutaneous warty lesions on his lower right leg of 12 years duration. The lesion was exophytic, painless, progressive and spreading in nature. On pressure during examination, thick white discharge was noted, however there was no evidence of blood or pus discharge. He was diagnosed to be diabetic seven years back and was on oral hypo-glycaemic agents. It started as an elevated reddish nodular lesion on the lateral margin of the right foot. There was no pain or itching, he felt that it was oozing watery fluid. The swelling kept on increasing in number and size on the leg with time. He consulted multiple doctors, who prescribed oral medicines but it had no effect and it kept on increasing in numbers as well as size of the warty lesion. There were no constitutional symptoms during whole period.

On examination, the limb was hyper pigmented and swollen compared to the other limb. There were multiple warty lesions, many of them had conglomerated to form a large cauliflower like masses at places. It was involving only the lower two third of the leg without involving the weight bearing part of the sole. The lesions towards the knee were healing at places (Three views of the leg-Figure 1). There was no local rise of temperature, firm to feel, non-tender and on pressure thick, white sticky fluid was oozing out (Arrow in the Figure 1). There was no significant inguinal lymph node enlargement. He had superficial hypo-pigmented lesion on his back that of Taenia furfur (Figure 2). On investigation, the haematological and biochemical parameters including renal, and liver functions were normal except for hyper-glycemia with HbA1C of 7.9gm%. He was negative for retroviral serology. The micro photograph of the same (Figure 3) showing area with fungi (white arrow) surrounded by inflammatory tissue in the form of a granuloma and was reported as Chromoblastomycosis.



Figure I The lesions towards the knee were healing at places (Three views of the leσ).



Figure 2 Superficial hypo-pigmented lesion on his back that of Taenia furfur.

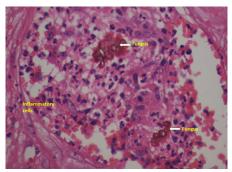


Figure 3 The micro photograph of the same (Figure 3) showing area with fungi (white arrow) surrounded by inflammatory tissue in the form of a granuloma and was reported as Chromoblastomycosis.

Discussion

Chromoblastomycosis is a chronic fungal skin and subcutaneous tissue infection. It is seen worldwide but more common in the tropics. These are group of melanized fungi and cause pigmentation on the skin. Present case had black pigmentation on the affected right leg clearly visible in comparison to the left (Figure 1). The main pathogens are Fonsecaea pedrosoi, Phialophora verrucosa, Cladosporium carrionii, or Fonsecaea compacta. It is relatively rare clinical condition and hence clinical diagnosis is difficult. It is mostly nonfatal and continues for long duration up to 36 years before starting the therapy. Besides the typical presentation like the present patient,



they might have, painless papules or nodules appear in the affected area progressing to scaly and verrucous plaques. Once the fungal agents enter the body part, small lesions comes up and it grows slowly for long time. The response to a single antifungal agent is not adequate and hence combination therapy is preferred.7 As the present case came after twelve years of the disease, we considered him as a severe case and advised surgical excision and electrocautery application to the subcutaneous tissue and started on two antifungal therapies. The drugs are itraconazole 200 mg twice daily and terbinafine 500mg per day initially. After a month the itraconazole was reduced to 200 mg daily once he responded to the treatment and advised to continue after 3 months of disappearance of all lesion. Besides the antifungal agents he has under gone two sessions of surgical excision and electro cauterization. On follow up, 3 months after the combined therapy showed good results, with reduction in the number of lesions (Figure 4). The follow up haematological and biochemical parameters has not shown any drug induced toxicities. In a small group of six patients pulse therapy of 400mg of itraconazole once a day and terbinafine alternate week were found to be effective and with less side effect and found no haematological and biochemical changes.^{7,8} The optimum duration of antifungal therapy is not fixed. It is given as per the response of the patients. But as per the literature antifungal therapy is recommended to continue till the tissue culture/ histology shows no fungi.8 The most common complications of chromoblastomycosis are ulceration, secondary bacterial infection, lymphedema that leads to elephantiasis and myiasis.9 Rare cases of malignant transformation to squamous cell carcinoma have been documented.10



Figure 4 On follow up, 3 months after the combined therapy showed good results, with reduction in the number of lesions.

Conclusion

Biopsy is diagnostic in long standing undiagnosed skin lesion. Chromoblastomycosis showing the fungal agent confirm the diagnosis. Two oral fungal agents give good response. Surgical excision restores the leg earlier.

Acknowledgments

None.

Conflicts of interest

The author declares there is no conflict of interest.

References

- Queiroz-Telles F, Esterre P, Perez-Blanco M, et al. Chromoblastomycosis: an overview of clinical manifestations, diagnosis and treatment. *Med Mycol*. 2009;47(1):3–15.
- Ameen M. Chromoblastomycosis: clinical presentation and management. Clin Exp Dermatol. 2009;3498):849–854.
- Najafzadeh MJ, Rezusta A, Cameo MI, et al. Successful treatment of chromoblastomycosis of 36 years duration caused by Fonsecaea monophora. *Med Mycol*. 2010;48(2):390–393.
- López Martínez R, Méndez Tovar LJ. Chromoblastomycosis. Clin Dermatol. 2007;25(2):188–194.
- Pindycka-Piaszczyńska M, Krzyściak P, et al. Chromoblastomycosis as an endemic disease in temperate Europe: first confirmed case and review of the literature. Eur J Clin Microbiol Infect Dis. 2014;33(3):391–398.
- Chen Y, Yin S, Li M, et al. A case of chromoblastomycosis by Fonsecaea nubica indicating a possible insect route of transmission. *Mycoses*. 2016;59(10):662–667.
- Gupta AK, Taborda PR, Sanzovo AD. Alternate week and combination itraconazole and terbinafine therapy for chromoblastomycosis caused by Fonsecaea pedrosoi in Brazil. Med Mycol. 2002;40(5):529–534.
- Ungpakorn R, Reangchainam S. Pulse itraconazole 400 mg daily in the treatment of chromoblastomycosis. *Clin Exp Dermatol*. 2006;31(2):245– 247.
- Slesak G, Inthalad S, Matthias Marschal, et al. Chromoblastomycosis after a leech bite complicated by myiasis: a case report. BMC infect dis. 2011;12:11:14.
- Rojas OC, Gonzalez GM, María Moreno-Treviño, et al. Chromoblasyomycosis by Cladophialophora carrionii associated with squamous cell carcinoma and review of published reports. *Mycopathologia*. 2015;179(1–2):153–157.