

Silvio Alencar Marques: Design and planning of the study; drafting and editing of the manuscript; effective participation in research orientation; intellectual participation in the propaedeutic and/or therapeutic conduct of the studied cases; critical review of the literature; critical review of the manuscript; approval of the final version of the manuscript.

## Conflicts of interest

None declared.

## References

1. Histiocytosis syndromes in children. Writing Group of the Histiocyte Society. *Lancet*. 1987;1:208–9.
2. Emile JF, Abla O, Fraitag S, Horne A, Haroche J, Donadieu J, et al. Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. *Blood*. 2016;127:2672–81.
3. Gianotti F, Caputo R, Ermacora E, Gianni E. Benign cephalic histiocytosis. *Arch Dermatol*. 1986;122:1038–43.
4. Polat Ekinci A, Buyukbabani N, Baykal C. Novel clinical observations on benign cephalic histiocytosis in a large series. *Pediatric Dermatol*. 2017;34:392–7.
5. Patsatsi A, Kyriakou A, Sotiriadis D. Benign cephalic histiocytosis: case report and review of the literature. *Pediatr Dermatol*. 2014;31:547–50.

Ana Flávia Teixeira de Abreu <sup>ID</sup>\*,  
Rebecca Perez de Amorim <sup>ID</sup>,  
Pedro Marciano de Oliveira <sup>ID</sup>,  
Marcelo Padovani de Toledo Moraes <sup>ID</sup>,  
Silvio Alencar Marques <sup>ID</sup>

*Department of Infectology, Dermatology, Imaging Diagnosis and Radiotherapy, Faculty of Medicine, Universidade Estadual Paulista, Botucatu, SP, Brazil*

\*Corresponding author.

E-mail: [anaflaviatabreu@hotmail.com](mailto:anaflaviatabreu@hotmail.com) (A.F. Abreu).

Received 29 September 2022; accepted 8 November 2022  
Available online 4 March 2024

<https://doi.org/10.1016/j.abd.2022.11.008>  
0365-0596/ © 2024 Sociedade Brasileira de Dermatologia.  
Published by Elsevier España, S.L.U. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

## Cutaneous plasmacytoma: a rare manifestation of multiple myeloma<sup>☆</sup>



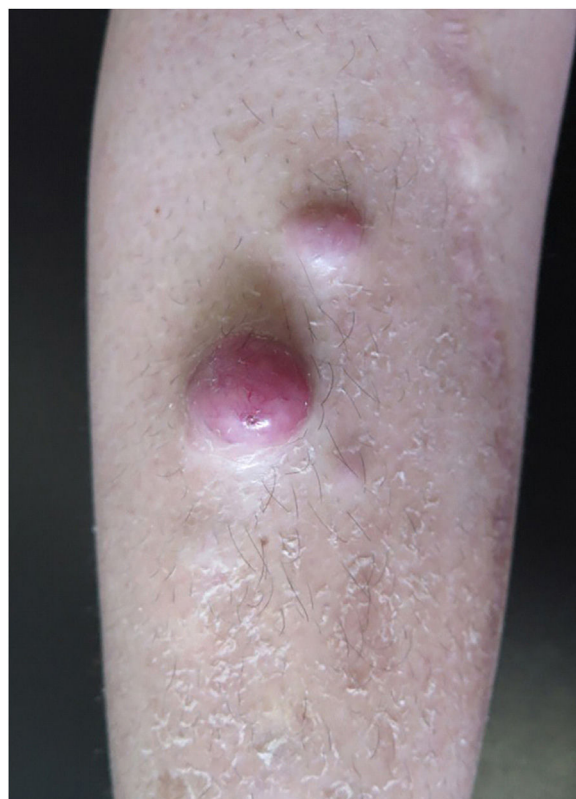
Dear Editor,

Cutaneous metastases result from the spread of a tumor to the skin through lymphatic or vascular embolization, direct implantation during surgery, or involvement of the skin through contiguity. Studies indicate a frequency of 0.7%–10.4%, mainly secondary to visceral neoplasms.<sup>1</sup> The primary neoplasms most often associated with skin metastasis include breast cancer, lung cancer, and melanoma.<sup>2</sup> There are few reported cases of cutaneous metastasis from multiple myeloma (MM), the main topic in this case report. Skin involvement associated with MM occurs in less than 10% of cases.

Due to the rarity of this manifestation, as well as the importance of its correct diagnosis, the present report describes a patient with MM and cutaneous metastasis after disease recurrence.

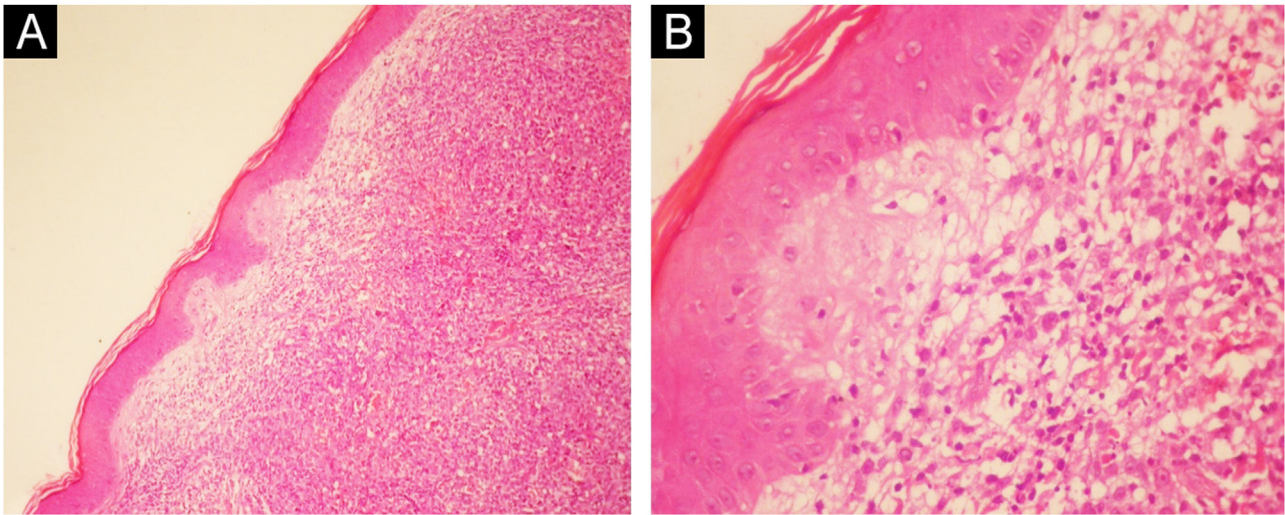
A 49-year-old female patient had been diagnosed with MM 12 years before. She underwent several treatments, including a bone marrow transplant. She had a painless lesion on her right leg that had been developing for three months. She had a history of excision of a tumor in the right tibia with prosthetic reconstruction in the previous year. On examination, she had two well-defined, erythematous tumors with regular contours, located on the right pre-tibial region, measuring up to 3 cm (Fig. 1). At the site

of the orthopedic prosthesis scar, she had an erythematous, hardened, and painless nodule measuring approximately 2 cm, adhered to deep planes (Fig. 1). The pathological analysis of an incisional biopsy was compatible with a

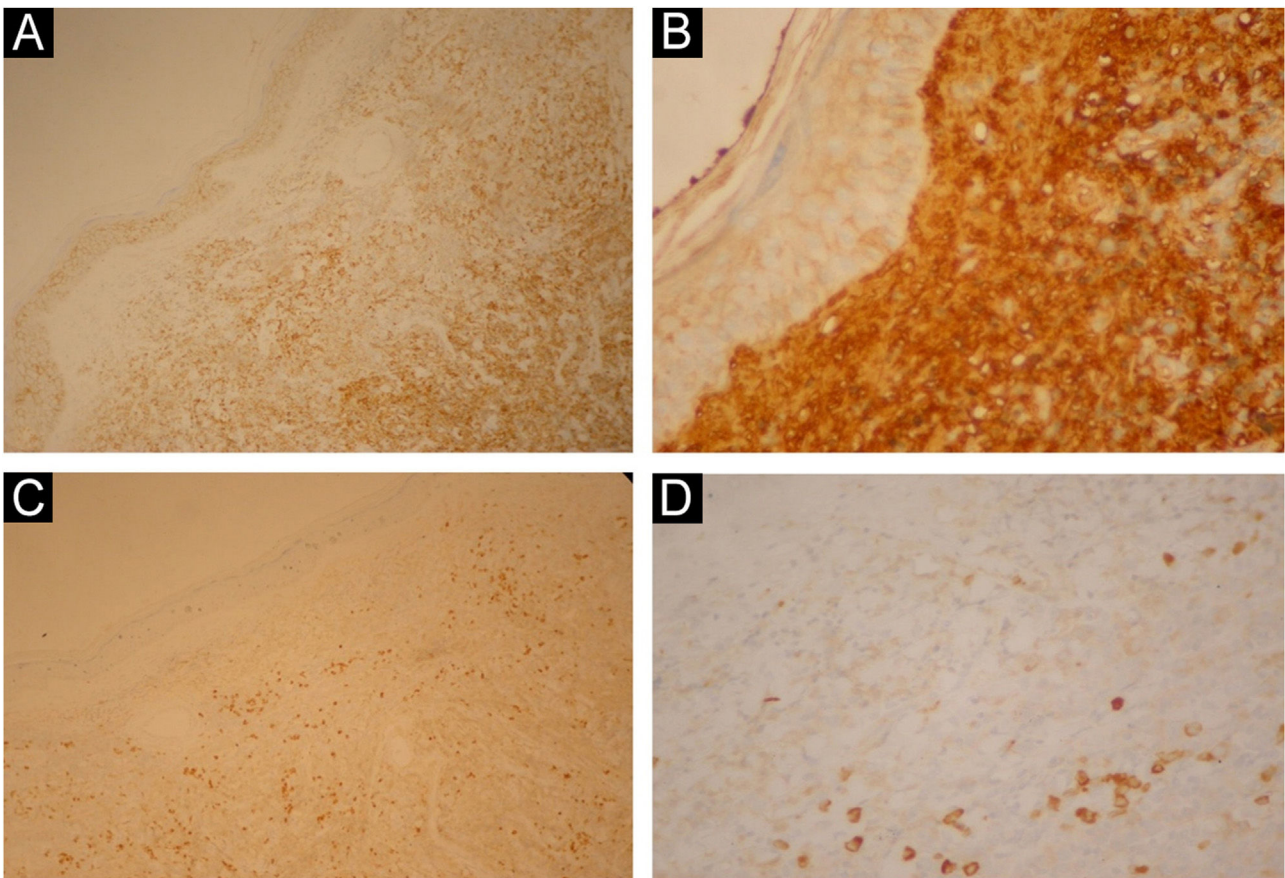


**Figure 1** Erythematous tumors on the right pre-tibial region.

<sup>☆</sup> Study conducted at the Pontifícia Universidade Católica de Campinas, Campinas, SP, Brazil.



**Figure 2** Immunohistochemistry. Diffuse infiltration of the dermis by atypical cells (A); dermal infiltration by large, poorly differentiated cells, suggestive of plasma cells – greater magnification (B).



**Figure 3** Diffuse positivity for C138 in tumor cells (A); diffuse intense KAPPA positivity in the lesion (B); focal positive CD79A (C); focal positive CD79A – higher magnification (D).

neoplasm of large, poorly differentiated cells, of probable metastatic origin (Fig. 2). Immunohistochemistry was positive for CD79a, CD138 and Kappa, confirming the diagnosis of MM skin metastasis (Fig. 3). In a joint decision with hematology, radiotherapy was chosen due to the poor prognosis.

However, the patient moved to another city and was lost to follow-up at the service.

Cutaneous involvement by MM is a rare event, and cutaneous metastasis can appear in any area of the skin, most frequently the trunk, extremities, and face.<sup>3</sup> Frequently,

multiple lesions are observed, although solitary lesions have also been recorded.<sup>4</sup>

Cutaneous metastatic lesions of MM are classified into non-specific ones, which are more common: secondary amyloidosis, alopecia, pyoderma gangrenosum, flat xanthomas, anhidrosis, sclerodermiform lesions, lichen myxedematosus, among others, and specific ones, which represent the spread of multiple myeloma in the final stage of the disease: secondary plasmacytomas that occur by direct extension to the skin from underlying bone lesions, such as in the case described in the present report, or by lymphatic and/or hematogenous spread.<sup>3,4</sup> They present as erythematous nodules, ulcerated or not, or plaques measuring up to 5 cm in diameter. Around 50% of the patients die within six months of the diagnosis. Cutaneous plasmacytomas can also appear in patients without a previous diagnosis of MM and are then called primary cutaneous plasmacytomas.

Therefore, a thorough dermatological examination is essential for the early diagnosis of cutaneous metastases from multiple myeloma. Thus, it becomes possible not only to optimize patient treatment but also to corroborate the importance of dermatologists because of their responsibility in the diagnosis and follow-up of patients with severe systemic diseases.

### Financial support

None declared.

### Authors' contributions

Larissa Helena Marques Carrai: Design and planning of the study; drafting and editing of the manuscript; collection, analysis and interpretation of data; critical review of the literature.

Elaine Cristina Faria Abrahão Machado: Design and planning of the study; drafting and editing of the manuscript; collection, analysis and interpretation of data; critical review of the literature.

Livia Matida Gontijo: Approval of the final version of the manuscript; effective participation in research orientation; drafting and editing of the manuscript; collection,

analysis and interpretation of data; critical review of the manuscript.





Luiza Castro: Design and planning of the study; drafting and editing of the manuscript; collection, analysis and interpretation of data; critical review of the literature.

### Conflicts of interest

None declared.

### References

1. Marques SA, Shibata AS, Martins DS, Miot HA, Marques MEA. Metástase cutânea de câncer de mama: relato de caso e revisão da literatura. *Diagn Tratamento*. 2008;13:164–8.
2. Machado DSB, Borges GS, Felipe GC, Ribeiro M, Zamboni F, Siqueira K, et al. Metástases cutâneas como apresentação de câncer de pulmão: relato de caso. *Revista Brasileira de Oncologia Clínica*. 2010;7:124–5.
3. Araújo C, Marques H, Fernandes JC, Pardo A, Brito C. Cutaneous plasmacytomas secondary to nonsecretory multiple myeloma. *J Dermatol Clin Res*. 2014;2:1022.
4. Souza DAF, Freitas THP, Helena P, Paes RAP, Müller H, Hungria VT. Mieloma múltiplo com plasmocitomas cutâneos. *An Bras Dermatol*. 2004;79:581–5.

Larissa Helena Marques Carrai \*,  
Elaine Cristina Faria Abrahão Machado ,  
Luiza Castro , Livia Matida Gontijo 

*Pontifícia Universidade Católica de Campinas, Campinas, SP, Brazil*

\* Corresponding author.

E-mail: [dralarissa.carrai.dermatologia@gmail.com](mailto:dralarissa.carrai.dermatologia@gmail.com)  
(L.H. Carrai).

Received 5 October 2022; accepted 27 October 2022  
Available online 23 February 2024

<https://doi.org/10.1016/j.abd.2022.10.018>  
0365-0596/ © 2024 Published by Elsevier España, S.L.U. on behalf of Sociedade Brasileira de Dermatologia. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

## High-resolution ultrasound with Doppler as a confirmatory diagnostic method in retronychia<sup>☆</sup>



Dear Editor,

We present the case of a 45-year-old female patient who was referred to the dermatologist with erythema, pain, and discharge in the left hallux. This condition had been present

for four months and was resistant to topical and oral antibiotics. She had active secretion and complete loss of the union of the proximal fold in the affected nail, henceforth chronic paronychia was the original diagnosis (Fig. 1). A high-resolution ultrasound with Doppler analysis of the nail apparatus was requested. The findings established the diagnosis of rethonychia (Figs. 2–4). Then we decided to perform a surgical intervention.

Retronychia is a disorder of the nail apparatus in which there is an abnormal growth of the nail plate within the proximal fold, leading to the formation of several generations of misaligned nail plates under the fold.<sup>1,2</sup>

Retronychia affects middle-aged adults, mostly females.<sup>3</sup> It is characterized by unilateral involvement and affects almost exclusively the hallux.<sup>4</sup> Repeated trauma, pregnancy,

<sup>☆</sup> Study conducted at the Sonoderma Institution, Medellín, Colombia.