



Challenging Cases

3 6 Persistent Erythema and Thickening of the Nasal 8 Skin

9 Clinical history

Q2 A 72-year-old man with a history of alcohol use and stable chronic
11 lymphocytic leukemia (CLL) not requiring treatment presented to der-
12 matology with a 3-year history of lesions located on the nasal dorsum
13 for evaluation. The patient had previously been diagnosed with rosacea
14 and rhinophyma and had received topical and oral antibiotic therapy
15 without improvement.

16 Physical examination

17 Physical examination revealed evident swelling along with erythema
18 and soft-tissue enlargement throughout the nasal dorsum, with marked
19 induration on palpation (Fig. 1).

20 Histopathology

21 Biopsy showed an unremarkable epidermis and a lymphocytic infil-
22 trate in the dermis composed of small lymphocytes with hyperchromatic
23 nuclei and scant cytoplasm (Fig. 2). Immunohistochemistry demon-
24 strated positivity for CD20, CD23, and BCL-2 (Fig. 3A–C).



Fig. 1. Erythema and soft-tissue enlargement of the nasal dorsum.

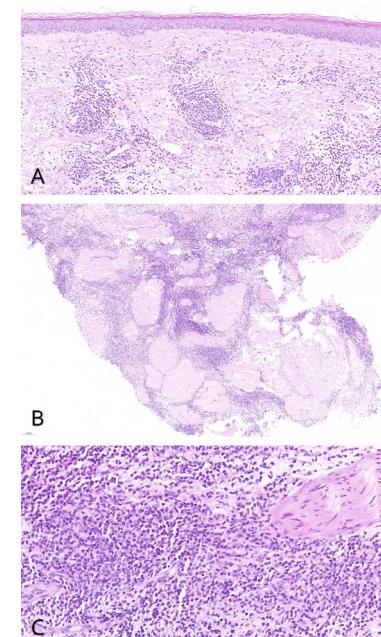


Fig. 2. (A–C) Hematoxylin–eosin stain: dermal lymphocytic infiltrate without epidermotropism.

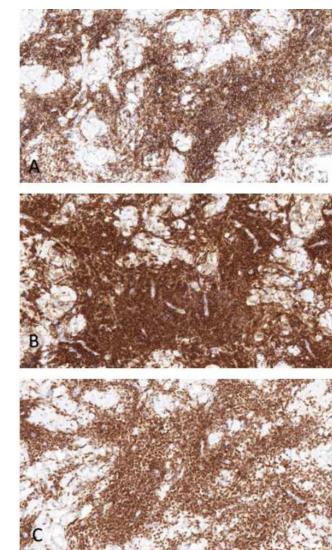


Fig. 3. (A) Immunohistochemistry positive for CD20; (B) CD23; (C) BCL-2.

What is your diagnosis?

<https://doi.org/10.1016/j.ad.2025.104580>

0001-7310/© 2025 AEDV. Published by Elsevier España, S.L.U. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

26 Diagnosis

27 CLL-induced cutaneous infiltration.

28 Course of the disease and treatment

29 Given progression of the CLL, with progressive enlargement of
 30 cervical, axillary, and inguinal lymph nodes, the Hematology unit
 31 started ibrutinib therapy. The patient showed a good hematologic
 32 response, along with notable parallel improvement of the cuta-
 33 neous lesions, with a marked reduction in nasal induration and
 34 erythema 1 month into therapy.

35 Comment

36 CLL is the most common type of leukemia in adults and is char-
 37 acterized by the accumulation of monoclonal B lymphocytes in the
 38 blood, bone marrow, spleen, and lymph nodes. Cutaneous involve-
 39 ment can be observed in approximately 25% of cases, mainly as
 40 nonspecific signs, such as infectious complications, exaggerated
 41 reactions to insect bites, or vasculitis, among others.¹ In addition,
 42 specific lesions due to cutaneous infiltration by tumor cells may
 43 occur, a condition known as leukemia cutis (LC).¹

44 The leukemia subtype most frequently associated with LC is
 45 acute myeloid leukemia, especially monocytic and myelomonocytic
 46 variants. Clinically, LC typically presents as solitary or
 47 grouped papules, plaques, or nodules, often arising in areas of skin
 48 previously affected by herpetic lesions.¹

49 Cases have been reported in the literature of LC presenting
 50 as rosacea-like eruptions.²⁻⁵ It has been proposed that the devel-
 51 opment of LC in the same location as prior rosacea lesions may
 52 represent a form of Wolf's isotopic response, similar to what occurs
 53 after prior varicella-zoster or herpes simplex infection.² Chronic
 54 stimulation by *Demodex folliculorum* of skin-associated lymphoid tis-
 55 sue has also been suggested as a potential pathway leading to
 56 lymphoma development, analogous to the relationship between
 57 *Helicobacter pylori* and gastric MALT lymphoma.^{2,3}

58 In our patient, the striking swelling confined exclusively to the
 59 nasal area and the absence of the sebaceous skin quality charac-
 60 teristic of rosacea prompted us to reconsider the diagnosis and
 61 perform a biopsy. Histology revealed the presence of a superfi-
 62 cial and deep perivascular dermal lymphocytic infiltrate that may

62 resemble the perivascular cuffing seen in tumid lupus.⁶ However,
 63 the absence of interstitial mucin deposition, the patient's hema-
 64 tologic history, and immunohistochemistry findings allowed us to
 65 reach the diagnosis of CLL-induced LC.

66 This case illustrates a very unusual clinical presentation of
 67 LC and underscores the importance of including this entity in
 68 the differential diagnosis of atypical rosacea-like lesions that are
 69 refractory to standard treatments.

70 Conflict of interest

71 The authors declare no conflict of interest. Q3

72 References

1. Morozova EA, Olisova OY, Nikitin EA. Cutaneous manifestations of B-cell chronic lymphocytic leukemia. *Int J Hematol.* 2020;112:459-465, <http://dx.doi.org/10.1007/s12185-020-02978-8>.
2. Vázquez-Osorio I, Chamorro-Chamorro P, Gonzalvo-Rodríguez P, Rodríguez-Díaz E. Leucemia cutis simulando una rosácea granulomatosa: presentación clínica de una leucemia linfóide crónica. *Actas Dermosifiliogr.* 2022;113:326-328, <http://dx.doi.org/10.1016/j.ad.2020.03.018>.
3. di Meo N, Stinco G, Trevisan G. Cutaneous B-cell chronic lymphocytic leukaemia resembling a granulomatous rosacea. *Dermatol Online J.* 2013;19, <http://dx.doi.org/10.5070/d31910020033>, 2003.
4. Murad A, Fortune A, O' Keane C, Ralph N. Granulomatous rosacea-like facial eruption in an elderly man: leukaemia cutis. *BMJ Case Rep.* 2016;2016, <http://dx.doi.org/10.1136/bcr-2016-215568>, bcr2016215568.
5. Ng E, Patel V, Engler D, Grossman M. Chronic lymphocytic leukemia associated leukemia cutis presenting as acne rosacea. *Leuk Lymphoma.* 2012;53:2304-2306, <http://dx.doi.org/10.3109/10428194.2012.676171>.
6. Thomas RM, Harrell JE, Rudnick E, et al. Leukemia cutis mimicking tumid lupus as the presenting sign in a case of mixed T/B-cell acute lymphoblastic leukemia. *JAAD Case Rep.* 2020;6:598-602.

73 B. Rebollo Caballero  ^{a,*}, S. Mallo García ^a, B. Ocaña Castillo ^b Q1

74 ^a Servicio de Dermatología, Hospital General Universitario Reina Sofía
 75 Murcia, Murcia, Spain

76 ^b Servicio de Anatomía Patológica, Hospital General Universitario Reina
 77 Sofía, Murcia, Spain

78 * Corresponding author.

79 E-mail address: blanca.rebollo97@gmail.com

80 (B. Rebollo Caballero).

81 93
 82 94
 83 95
 84 96
 85 97
 86 98
 87 99
 88 99
 89 99
 90 99
 91 99

92 Q1