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7 **Erythematous Plaque in the Left Axillary Region**

8 *A diagnostic challenge where dermoscopy can help*

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16 A 70-year-old male patient was attended in our outpatient dermatological clinic for a
17 plaque on left axillae of unknown time of evolution. Physical examination revealed an
18 erythematous plaque with no desquamative component (Fig. 1A). Dermoscopy showed
19 thick reticular white lines, white clods, and structureless areas (Fig. 1B), dotted and linear
20 vessels, and a sessile projection in the lower part of the plaque. Histological examination
21 using a 4 mm punch biopsy revealed a hyperkeratotic epidermis without dyskeratosis, with
22 intraepidermal infiltration of dysplastic cells distributed singly and in clusters (Fig.1C).
23 Immunohistochemistry staining of this tumoral component was positive for GCDFP-15,
24 mammaglobin and CK7, and negative for HMB45 and S-100 protein, which was
25 consistent with the diagnosis of extramammary Paget's disease (EMPD). Complementary
26 examinations (body-TC, a complete blood test with hemogram, liver and renal function,
27 ions, protein S100b and PSA; colonoscopy and mammary ecography) ruled out the
28 presence of concomitant tumour. The lesion was completely removed through wide local
29 excision and primary closure, with no local recurrence or distant metastasis during the
30 following 12-months. He is currently under long-term follow-up with periodic revisions
31 every 6 months.

32

33 **Comment**

34 EMPD is an infrequent cutaneous adenocarcinoma with unknown incidence, which
35 predominantly affects postmenopausal Caucasian women and Asian men aged between 60
36 and 80 years old. It is usually presented as a slow-growing rounded or oval erythematous
37 plaque, asymptomatic or pruritic, located in genital, perianal or, less frequently, axillary
38 regions.¹ Dermoscopy is a non-invasive technique which raises suspicion of EMPD.
39 Pigmented, whitish and reddish structures represent the main group of dermoscopy
40 patterns. Glomerular and dotted vessels are the most common vascular patterns. Lava lake
41 structures (BAW) and cloud-like structureless areas (WAW) were described by
42 Payapvipapong *et al.*⁴ None of the patients had axillary lesions, practically all of them
43 were located in the ano-genital region. This could explain the fact that we have not
44 observed BAW and WAW in our patient. Histology allows to distinguish between EMPD
45 and other diagnoses such as Langerhans cell histiocytosis, Bowen's disease, amelanotic
46 melanoma and mycosis fungoides (Table 1).¹
47 The most relevant prognostic factor is the presence of underlying malignancy, either a
48 contiguous tumour (23% of cases) or a distant one (8%-46% of cases). Underlying cancer
49 screening might include a complete physical examination with lymph node and breast
50 examination. Complementary tests may include urine cytology and colonoscopy for both
51 genders, PSA blood test and digital rectal examination in male patients, and Papanicolaou
52 test and mammography in female patients.²
53 Primary cases, as the one we present, are usually indolent, with good survival rates. Wide
54 local excision is the gold standard treatment, although imiquimod, photodynamic therapy
55 or radiotherapy could be considered in non-surgical cases.^{1,3} Recurrence rates are
56 considerably elevated regardless of the treatment, so a long-term follow-up based on a
57 careful physical examination is highly recommended.¹
58 The authors have obtained patient consent for publication purposes.

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60 **Authors' Contribution**

61 All authors contributed equally in the conceptualization, design, data acquisition, analysis
62 and write up of the manuscript.

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79 **Figure 1:** (A) Round erythematous plaque with slightly raised edges without scaling or
80 ulceration, located on the left axilla (black arrow: biopsy area); (B) Dermoscopy showed
81 milky red structureless areas, reticular white lines (red arrow) and white clods (black
82 arrow). Dotted and linear vessels (asterisk) were the vascular pattern identified; (C)
83 Epidermis with hyperkeratosis without dyskeratosis, with a component of dysplastic cells
84 distributed singly and in clusters, with the following features (black arrow): large cells
85 with pale cytoplasm and irregular nuclei with visible nucleoli, most of them just above the
86 epidermal basal layer, with others ascending to intermediate epidermal strata (H&E 4x).