Asymmetrically Pigmented Patch on the Vulvo-perineal Area: A Quiz

Maria Lentini and Maria Le Donne

Departments of Human Pathology and Gynaecological, Obstetric Sciences and Reproductive Medicine, University of Messina, IT-98124 Messina, Italy.
E-mail: maria.lentini@unime.it

A 50-year-old Caucasian woman presented with a 4-month history of a vulvo-perineal lesion, occasionally accompanied by day-time pruritus, especially before menstruation. At its onset, the lesion was pink-to-white in colour, but recently the patient noted an alarming change to dark brown (Fig. 1). She was in otherwise good health and denied taking any medications. Her medical history was notable for the surgical excision of a pancreatic adenoma two years before. A gynaecological examination and colposcopy showed normal findings. PAP smear and HPV-DNA tests were negative.

Dermoscopic analysis revealed an equivocal pigmented lesion. An incisional biopsy showed atypical neoplastic cells, positive for periodic acid-Schiff and cytokeratin 7 (Fig. 2) and surrounded by numerous reactive dendritic HMB-45-positive melanocytes. No dermal invasion was observed.

What is your diagnosis? See next page for answer.
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Diagnosis: Pigmented extramammary Paget’s disease.

The patient was subjected to blood chemistry and serum carcinoembryonic antigen tests, the results of which were all negative. Due to the previous surgical excision of a pancreatic adenoma, she also underwent abdominal and pelvic computer tomography and gastrointestinal endoscopy. Again the results were negative, as were those of a mammography. The patient underwent a partial left hemivulvectomy with deep resection to the superficial fascia and a Z-plastic procedure. Histological examination of a surgical specimen confirmed the diagnosis and excluded the presence of an underlying invasive neoplasm. At 28-months post-surgery the patient remains disease free.

Pigmented extramammary Paget’s disease (PEMPD) is a very rare clinico-pathological variant of extramammary Paget’s disease (EMPD) (1). In a small proportion of cases, EMPD is associated with underlying in situ or invasive neoplasia, or with distant internal organ neoplasms, most frequently gastrointestinal or urothelian (2, 3). Vulval localisation is rare, but typically occurs in postmenopausal Caucasian women. Clinically, it manifests as a uni- or multi-focal erythematous and/or eczematous lesion (2). Occasionally the lesion is pigmented (1). Dermoscopy is insufficient to produce a correct diagnosis, so differential diagnosis requires a histological biopsy to exclude malignant melanoma, heavily pigmented seborrhoeic keratosis and vulvar intraepithelial neoplasia. The histological typical features include reactive dendritic melanocytes in a process that is probably mediated by tumour-derived factors (4).

The standard treatment for primary vulval PEMPD does not differ from that for classical EMPD, i.e. surgical excision of the visible lesion with resection to the fascia and 2-cm gross margins (5). Although the prognosis following surgical resection is generally favourable, long-term monitoring of patients is recommended because recurrence is common and it is often necessary to repeat the surgical excision (6). For the preservation of the vulval anatomy, alternative non-surgical treatments have been used with conflicting results (7, 8).

The topical immunomodulatory cream imiquimod can be useful in cases in which surgery would be problematic because of the location and size of the tumour, as well as for recurrences (9). However, when we tested it in another EMPD-affected patient, control skin biopsies taken after the completion of therapy revealed residual tumour cells, although there was no clinical evidence of residual EMPD and pruritus was absent.

Photodynamic therapy based on the application of 5-aminolevulinic acid and subsequent irradiation with red light seems to have a place in treating recurrent forms of EMPD and in controlling symptoms (10), as well as in enhancing the efficacy of carbon dioxide laser ablation (11).

REFERENCES