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CLINICAL CASE

Multiple recurrent eccrine porocarcinoma with inguinal metastasis. A case report[☆]



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KEYWORDS

Porocarcinoma;
Lymphadenectomy;
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Abstract

Background: Eccrine porocarcinoma, first described in 1963, is a rare malignant lesion arising from the eccrine sweat glands. It is usually a primary tumour, or even more common, a malignant degeneration of an eccrine poroma. It usually affects older persons and is located most commonly on the lower extremities. About 20% of eccrine porocarcinoma will recur after treatment. The treatment is wide local excision of the primary lesion. This uncommon skin tumour has a locally aggressive behaviour and a high recurrence rate.

Clinical case: An 82 year-old man presenting with multiple recurrent eccrine porocarcinoma with inguinal metastasis. The treatment was a radical excision and inguinal lymphadenectomy. There were no postoperative complications, but there was local recurrence after six months.

Conclusion: Early diagnosis and wide excision is the best way to achieve a good prognosis, due to the aggressiveness of this tumour.

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PALABRAS CLAVE

Porocarcinoma;
Linfadenectomía;
Recurrente

Porocarcinoma ecrino multirrecidivante con metástasis inguinal. Caso clínico

Resumen

Antecedentes: Descrito por primera vez en 1963, el porocarcinoma ecrino es un raro tumour maligno que se desarrolla de las glándulas ecrinas, presentándose como tumour primario, o más

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frecuentemente mediante transformación maligna de un poroma eccrino. Afecta a pacientes de edad avanzada y la localización más frecuente es en extremidades inferiores. Se observa una recurrencia local del 20% después del tratamiento. La escisión quirúrgica es el tratamiento de elección. Esta infrecuente neoplasia cutánea tiene un comportamiento localmente agresivo y un alto índice de recidiva.

Caso clínico: Presentamos el caso de un varón de 82 años con porocarcinoma eccrino multirrecidivado y metástasis ganglionar, tratado con extirpación amplia y linfadenectomía inguinal. Buena evolución postoperatoria. Múltiples recidivas locales a los 6 meses.

Conclusión: Debido a su carácter potencialmente agresivo, el diagnóstico precoz y su rápido tratamiento quirúrgico es la forma adecuada de conseguir un pronóstico favorable frente a este tipo de tumor.

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Background

According to Marone et al.,¹ eccrine porocarcinoma was described in 1963 by Pinkus and Mehregan as epidermotropic eccrine carcinoma. It is a rare type of skin cancer (0.005–0.01% of all skin tumours), which arises from the sweat glands. It can be a primary or a transformation of an eccrine poroma. It has the same incidence in both genders, and principally affects people of advanced age. Fewer than 300 cases have been reported in worldwide medical literature,^{1,2} and its incidence in Europe is rare. Twenty percent of these tumours recur, 10% metastasise to solid organs, and 20% metastasise to regional lymph nodes, with a mortality of almost 70% in the latter. It is difficult to establish an accurate prognosis as there is little follow-up of the cases described because the tumour is so rare.

Clinically it manifests as a solitary lesion, which can be ulcerated nodule in type, or a papule or verrucous lesion. Differential diagnosis can be made with basal cell carcinoma, epidermoid carcinoma, seborrheic keratosis, etc.

The most common location is on the lower limbs, head and neck, and it is less frequent on the trunk, vulva, chest, nail bed, and upper limbs.

Histological diagnostic criteria for eccrine porocarcinoma are an accumulation of atypical cells, some with tubular structures, which emigrate from the epidermis, and are glycogen rich and PAS positive.

The treatment for this type of tumour is by broad surgical excision, Mohs surgery or radiotherapy. Mohs surgery is a surgical technique which enables the removal in layers of selected skin tumours and achieves the highest cure rates. Likewise it enables maximum saving of healthy tissue. All of this is possible through microscopic supervision, which enables 100% analysis of the tumour edges of each of the layers, and thus guide the surgeon through the successive lines of tumour until it is completely removed.

Surgical treatment is curative in 80% of cases. The role of prophylactic radical lymphadenectomy is debatable.³

Clinical case

An 82-year old male patient with a personal history of arterial hypertension, diabetes mellitus and benign prostatic

hypertrophy. The patient began with a papulous lesion 4 years previously, which was circular, 1 cm in diameter, in the area of the right hip, which was removed with an anatomopathological diagnosis of porocarcinoma, for which he received radiotherapy. He was referred to the plastic surgery unit with local recurrence in the form of a 1 cm papule next to the scar, which was surrounded by an area of major radio dermatitis of 8 cm × 5 cm. The patient underwent computed tomography on which right inguinal adenopathies were identified radiologically suspicious of metastasis. A fine-needle aspiration was undertaken (FNA) of these adenopathies which were palpable on physical examination, and were reported as positive malignancies; the remainder of the extension study reported no evidence of distant metastasis.

The patient was operated under general anaesthesia, and an en bloc excision of the area of radio dermatitis was performed including recurrence with more than a 2 cm margin of healthy tissue (Fig. 1); cover was made using a large rotation flap of the right flank with medial base (Fig. 2); and right inguinal lymphadenectomy (Fig. 3). The anatomopathological report was compatible with recurrence of porocarcinoma, radiodermal changes and broad surgical margins; porocarcinoma metastasis in 2 of the isolated lymph nodes.



Figure 1 Area of recurrence surrounded by radiodermatitis. Marking of broad excision, and design of flap for cover.

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