Uncommon skin cancer: pleomorphic dermal sarcoma

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DESCRIPTIONS
An 86-year-old man presented to the dermatology clinic with a 10-week history of lesion on the vertex of his scalp. This had been intermittently crusting and bleeding but was non-tender and not enlarging. The patient had a background of two previous squamous cell skin carcinomas of his right shoulder and left ear. With regards to his sun exposure, he had previously served in the armed forces and had been posted to hot countries. On physical examination there was a 9×13 mm raised lesion with an overlying crust and rolled edges on the scalp vertex (see figure 1A,B). There was no evidence of cervical lymphadenopathy. Clinical suspicion was that of a keratoacanthoma or squamous cell carcinoma, given the patient’s background. A 4 mm cutaneous punch biopsy was organised on the fast-track pathway.

Histological analysis of the scalp lesion revealed a highly infiltrative tumour composed of small clusters of atypical plump spindle cells. The deeper aspect was composed of epithelioid cells with large, pleomorphic nuclei. There were mitoses present within the tumour, including atypical forms. There was no evidence of perineural or lymphovascular invasion. The tumour invaded the subcutaneous adipose tissue with a maximum depth of 4.5 mm. The malignant cells showed weak to moderate immunostaining for smooth muscle actin and did not stain for AE1/AE3, S-100, desmin, CD31, CD34 or p63 (see figures 2 and 3).

The clinical, morphological and immunophenotype was in keeping with a pleomorphic dermal sarcoma (PDS). The patient was reviewed in clinic and the results discussed. Subsequent excision biopsy was performed with an overlying skin graft under the plastic surgery team with additional findings of actinic keratosis on histology. Our patient is currently awaiting further review.

PDS are rare and not well documented in the literature. They are thought to be closely related to atypical fibroxanthomas (AFX), sharing similar clinical, histological and immunophenotypic features. Clinically, both AFX and PDS affect elderly males on sun-damaged skin, with a strong predilection for the scalp.1 They typically have a rapid growth and ulcerate and bleed. Histologically, AFX are well-circumscribed growths, confined to the dermis. They contain pleomorphic, epithelioid, spindle and multinucleated tumour giant cells.1 PDS, however, display additional features such as infiltrative growth, invasion of the subcutis, tumour necrosis as well as lymphovascular invasion, suggesting that they may have a more aggressive course than AFX. Although a rare skin cancer and

Learning points

► Pleomorphic dermal sarcoma and atypical fibroxanthomas are closely related and likely part of the same disease spectrum.
► Although a rare cancer, it is important not to miss this diagnosis and follow-up patients due to their potential metastatic risk.

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their metastatic potential estimated at <5%, it is important to not miss this diagnosis.\(^2\)

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**REFERENCES**