

Desmoplastic melanoma component (Core)

Desmoplastic melanoma (DM) is a rare subtype of melanoma characterized by malignant spindle cells separated by prominent fibrocollagenous or fibromyxoid stroma. Primary melanomas may be entirely or almost entirely desmoplastic (“pure” DM) or exhibit a desmoplastic component admixed with a non-desmoplastic component (“mixed” DM).¹ Spindle (or epithelioid) melanoma cells not separated by desmoplastic stroma are not regarded as desmoplastic melanoma but may form the non-desmoplastic component of a mixed desmoplastic/non-desmoplastic melanoma. In 2004, Busam *et al* reported a clinicopathologic study of DM patients in which subdividing the tumours into “pure” and “mixed” subtypes correlated with clinical outcome.² In that study, the authors classified melanomas as “pure” DM if “the overwhelming majority (≥90%) of invasive tumour was desmoplastic”, or “mixed” DM if “typical features of DM were mixed with densely cellular tumour foci without fibrosis and desmoplasia” and the DM areas involved <90% and >10% of the invasive melanoma. Similar findings have since been reported by others.²⁻¹⁵ Improved disease-specific survival is seen in patients with “pure” DM, when compared with patients with “mixed” DM and those with melanomas lacking a desmoplastic component.²⁻¹⁵ Furthermore, regional nodal metastasis (including that detected by sentinel lymph node biopsy) is less common in patients presenting with clinically localized pure DM compared with those who had mixed DM or conventional melanomas.²⁻¹⁵

References

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