



Segmental vitiligo-like hypopigmentation on the right lower cheek in a patient with metastatic melanoma (A); Wood lamp examination highlighted the achromic areas (B).

prognostic factor² and correlates with a better therapeutic outcome in patients undergoing treatment with biotherapy.⁵

In most cases, the onset of achromic lesions follows the diagnosis of melanoma. Hypopigmentation appears on average 4.8 years after the initial diagnosis and approximately 1 to 2 years after lymph

node or distant metastasis.¹ In our case, it occurred 12 years after the initial diagnosis and 2 years after metastatic disease was diagnosed.

Despite having widespread metastatic melanoma, our patient only developed achromic patches on the area near the prior melanoma. However, most affected patients present with hypopigmented patches in a bilateral symmetric distribution pattern similar to common vitiligo. No correlation has been found between the hypopigmentation distribution and the location of the primary tumor.¹

Because fotemustine is not likely to induce hypopigmentation, we believe that the vitiligolike hypopigmentation in our patient was related to an immune-mediated response associated with melanoma. To help explain our findings, one hypothesis considered was that cutaneous mosaicism may be involved in segmental vitiligo.⁶ The tumor may have triggered an immune response that affected a close susceptible area of mosaic vitiligo, leading to these clinical findings.

REFERENCES

1. Hartmann A, Bedenk C, Keikavoussi P, et al. Vitiligo and melanoma-associated hypopigmentation (MAH): shared and discriminative features. *J Dtsch Dermatol Ges.* 2008;6:1053-1059.
2. Quaglino P, Marengo F, Osella-Abate S, et al. Vitiligo is an independent favourable prognostic factor in stage III and IV metastatic melanoma patients: results from a single-institution hospital-based observational cohort study. *Ann Oncol.* 2010;21:409-414.
3. Taïeb A, Picardo M, VETF Members. The definition and assessment of vitiligo: a consensus report of the Vitiligo European Task Force. *Pigment Cell Res.* 2007;20:27-35.
4. Becker JC, Guldberg P, Zeuthen J, et al. Accumulation of identical T cells in melanoma and vitiligo-like leukoderma. *J Invest Dermatol.* 1999;113:1033-1038.
5. Boasberg PD, Hoon DS, Piro LD, et al. Enhanced survival associated with vitiligo expression during maintenance biotherapy for metastatic melanoma. *J Invest Dermatol.* 2006;126:2658-2663.
6. Van Geel N, Speeckaert R, Melsens E, et al. The distribution pattern of segmental vitiligo: clues for somatic mosaicism. *Br J Dermatol.* 2013;168:56-64.