

Melanoma in Organ Transplant Recipients: A Case Series[☆]



Melanoma y trasplante: serie de casos

To the Editor:

Malignant melanoma is a highly immunogenic cancer¹ and immunosuppression may affect its incidence and outcome. We describe a retrospective case series of 12 adult patients with a diagnosis of melanoma following solid-organ transplant from 1996 to 2017 in the dermatology department. During the study period, we found more than 1650 solid-organ transplant patients in follow-up, most of whom had received a kidney transplant. Twelve cases of melanoma were diagnosed, 8 of which were men and 4, women. The mean age at transplant was 55.5 years (interval, 29–73 y) and the mean age at melanoma diagnosis was 69 years (interval, 45–78 y). The mean time from transplant to diagnosis of melanoma was 8.5 years (interval, 1–22 y). The majority (9 patients) received a kidney transplant, and the most frequent location of the melanoma was the face. Three cases were *in situ* melanoma and, of the patients with invasive melanoma, the Breslow thickness was 0.65 mm (interval, 0.30–1.24 mm). The mean follow-up time was 6.5 years (interval, 0.12–17 y). Only 1 patient presented recurrence of the disease, with lymph-node infiltration 14 years after the initial diagnosis (Patient 8). No deaths due to melanoma were observed; however, Patient 11 was in an advanced state on diagnosis and died a month-and-a-half after diagnosis of the melanoma, due to a massive acute myocardial infarction, before chemotherapy was instated.

Skin cancer is the most common neoplasia after transplant. Regarding the association between melanoma and transplant, 3 main scenarios should be analyzed:

- The first scenario refers to patients with a history of melanoma. These patients should be ruled out as donors, in accordance with the consensus report of the Spanish National Transplant Organization (May 2006) (**Table 1**).
- The second scenario involves patients with melanoma diagnosed before transplant. To date, no evidence exists to suggest an increase in the risk of recurrence or higher mortality in patients with a history of melanoma prior to transplant.^{2–4} After reviewing 68 patients with melanoma, who subsequently received a transplant, and comparing them with 141,441 patients with melanoma in the general population, Arron et al³ observed no statistically significant differences in mortality specifically due to melanoma; these data support the current practice of considering patients with a personal history of melanoma as candidates for transplant. Guidelines regarding how to act to select patients with a history of melanoma as transplant recipients are scarce; in 2016, Zwald et al⁵ proposed the recommendations for action shown in **Table 2**.
- Finally, in relation to patients in whom melanoma is diagnosed after transplant, an increase in the incidence of melanoma is known to exist in transplant patients. Garrett et al studied 10,649 transplant patients and found a melanoma incidence of 125 per 100,000 people/year, compared to between 9 and 18 per 100,000 people/year in the general population. Those authors identified advanced age, Caucasian race, male sex, and having received a thoracic organ as risk factors.⁶ Ascha et al identified use of sirolimus and cyclosporin as immunosuppressants

Table 1 Characteristics of Study Patients.

[0,2-4]Patient				[0,5-7]Transplant			[0,8-10]Melanoma		
Nº	Sex	Year of birth	Year	Organ	Year of diagnosis	Location	Breslow thickness	Recurrence	Mortality due to MM
1	M	1938	2011	Kidney	2013	Face	0.30	No	No
2	F	1930	1992	Kidney	2004	Face	0.50	No	No
3	M	1941	2007	Kidney	2010	Face	0.40	No	No
4	F	1938	1998	Kidney	2012	Extremity	1.24	No	No
5	M	1953	2007	Kidney	2011	Trunk	0.60	No	No
6	M	1963	2013	Kidney	2014	—	0.88	No	No
7	M	1957	1986	Kidney	2007	Face	NM	No	No
8	M	1957	1997	Liver	2002	Trunk	0.85	Yes	No
9	M	1936	1992	Kidney	2014	Face	0.70	No	No
10	F	1945	2000	Liver	2014	Trunk	NM	No	No
11	M	1963	2014	Lung	2017	Extremity	— ^a	—	No
12	F	1940	2004	Kidney	2017	Face	NM	No	No

MM, malignant melanoma; NM, not measurable because they were *in situ* melanomas.

^a The Breslow thickness was not available as the patient presented metastatic disease on diagnosis (lymph node and liver involvement) and complete excision of the primary lesion was not performed.

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Table 2 Recommended Waiting Times Before Transplant for Patients With a History of Melanoma.

MM (AJCC staging)	Time Between Melanoma Diagnosis and Transplant
In situ	Wait not necessary
Stage IA	2 y
Stage IB/IIA	2-5 y
Stage IIB/IIC	5 y
Stage III/IV	Not a transplant candidate

AJCC, American Joint Committee on Cancer; MM, malignant melanoma.

Source: Zwald et al.⁵

among the risk factors for melanoma in patients who had undergone a kidney transplant. The mean time between transplant and diagnosis of melanoma was 1.45 years,⁷ which was less than in our series. The outcome of post-transplant melanoma appears to be determined by the thickness of the tumor on diagnosis and is similar to that of the general population when the Breslow thickness is 1.5 mm or less.^{4,8} None of our patients presented a Breslow thickness of greater than 1.5 mm. It is important to point out that, although the change from immunosuppressive therapy to mTOR inhibitors is associated with a lower incidence of skin cancer, the antitumor effect of these agents for melanoma alone has only been shown in mouse models.⁹

In conclusion, in our study, we observed no mortality due to melanoma in patients with post-transplant melanoma, although 1 patient developed a recurrence of the disease and another patient presented metastatic disease at the time of diagnosis. This study is a descriptive, retrospective, observational study and the existence of an association may be fortuitous. We aim to provoke reflection on a circumstance that is rare, but which is of transcendent importance to the life of the patient, as malignant melanoma may be an event that leads to a life change, but denying an organ transplant is a death sentence.

Conflict of interests

The authors declare that they have no conflict of interest.

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