



# ACTAS Dermo-Sifiliográficas

Full English text available at  
[www.actasdermo.org](http://www.actasdermo.org)



## RESIDENT'S FORUM

### RF-Nevi With Moderate Dysplasia Excised With Positive Surgical Margins: Should We Re-Excise?☆



### FR-Nevus con displasia moderada extirpados con márgenes quirúrgicos afectados. ¿Se deben reextirpar?

D. Morgado-Carrasco,\* H. Feola, X. Fustà-Novell, N. Martínez

Servicio de Dermatología, Hospital Clínic de Barcelona, Universitat de Barcelona, Barcelona, Spain

Received 23 December 2018; accepted 1 January 2019

#### KEYWORDS

Nevus;  
Dysplastic nevi;  
Melanoma;  
Positive histologic margins;  
Moderate dysplasia

#### PALABRAS CLAVE

Nevus;  
Nevus displásico;  
Melanoma;  
Márgenes afectos;  
Displasia moderada

The concept of dysplastic nevus (DN) is somewhat controversial. While DN is not a proven precursor of cutaneous melanoma, several studies suggest that it may be a risk factor. The management of DN is also a topic of some debate.

There is no consensus on the optimal approach (observation or re-excision) when a nevus with moderate dysplasia has been excised and the margins are affected. A survey conducted in North America found that 48% of those surveyed would re-excite the lesion. In 2015, the North American Pigmented Lesion Subcommittee suggested that observation may be a reasonable option, although the evidence to support this recommendation was limited.<sup>1</sup>

Recently, Kim et al<sup>2</sup> published the results of a retrospective multicenter study that examined the development of melanoma in 438 patients with excised nevi with moderate dysplasia and compromised margins (467 lesions). Of these, 50.4% had undergone previous biopsies of dysplastic nevi, 33.2% had a history of melanoma, and 23.8% (83 of 349 patients) had a family history of melanoma. The biopsy methods used were shave (46.4%), punch (45.3%), and elliptical (7.7%) excision. Dysplasia was independently assessed by an expert. During a mean follow-up period of 6.9 years (range, 3–21.3 y), none of the participants developed melanoma at the site of the excised DN. However, 22.8% developed a primary melanoma at a different site. Of these, 3 patients had metastatic melanoma (2 with invasive melanoma prior to biopsy). Comparison of those that developed a subsequent cutaneous melanoma with those that did not identified the following risk factors: older age (mean age of 49.7 vs 45.9 years;  $P = 0.04$ ); history of previous melanoma (49% vs 9.9%;  $P < 0.001$ ); and a previously biopsied DN (33.5% vs 13.2%;  $P < 0.001$ ). No association was observed for sex or family history of melanoma. Multivariate regres-

☆ Please cite this article as: Morgado-Carrasco D, Feola H, Fustà-Novell X, Martínez N. FR-Nevus con displasia moderada extirpados con márgenes quirúrgicos afectados. ¿Se deben reextirpar? Actas Dermosifiliogr. 2020;111:688–689.

\* Corresponding author.

E-mail address: [\(D. Morgado-Carrasco\).](mailto:morgadodaniel8@gmail.com)

**Table 1** Summary of Studies on the Follow-up of Excised Dysplastic Nevi with Affected Margins

Author, Year	Type of Study	No. of Dysplastic Nevi	Type of Dysplasia	Follow-up Duration, y	Melanoma on Site of Excised Dysplastic Nevus	Melanoma Elsewhere
Cockerell and Berson, 1985 <sup>4</sup>	Retrospective	28	ND	5	0	0
Kmetz et al, 2009 <sup>5</sup>	Retrospective	26	ND	6.12 (minimum 5 years)	0	Not evaluated
Goodson et al, 2010 <sup>6</sup>	Retrospective	65	Mild	≥ 2	0	Not evaluated
		4	Moderate			
Hocker et al, 2013 <sup>7</sup>	Retrospective	66	Mild	17.4 (range, 0–29.9)	0	Not evaluated <sup>b</sup>
		42	Moderate			
		7	Severe			
Hiscox et al, 2017 <sup>3</sup>	Retrospective	147	Moderate	3.6 (range, 1.2–8.7)	0	Not evaluated
Kim et al, 2018 <sup>2</sup>	Retrospective	467	Moderate	6.9 (range, 3–21.3)	0	100 (22.8%)

Abbreviation: ND, not described.

<sup>a</sup>Studies based exclusively on the follow-up of nevi with severe dysplasia are not included in this table.

<sup>b</sup> No patient presented metastatic melanoma.

sion analysis revealed that a history of melanoma (odds ratio [OR], 11.74; 95%CI, 5.71–24.15;  $P < 0.001$ ) and previous (histologically confirmed) DN (OR, 2.55; 95%CI, 1.23–5.28;  $P = 0.01$ ) were independently associated with the development of melanoma at a site other than that of the DN biopsy, even after adjusting for age, sex, and family history of melanoma.

The results of the study by Kim et al, the largest cohort described to date, are consistent with the findings of previous publications (Table 1),<sup>3</sup> suggesting that observation is a reasonable option in these cases and supporting the hypothesis that DN is not a precursor lesion of melanoma, but an important risk marker. It should be noted that around 70% of cutaneous melanomas develop de novo and not on a pre-existing nevus.

In patients with nevi with moderate dysplasia and affected margins after excision, clinical observation may be a valid option. Patients with a previous history of melanoma or 2 DNs (one of which shows moderate dysplasia) should be considered at high risk and followed closely.

## References

1. Kim CC, Swetter SM, Curiel-Lewandrowski C, Grichnik JM, Grossman D, Halpern AC, et al. Addressing the knowledge gap

in clinical recommendations for management and complete excision of clinically atypical nevi/dysplastic nevi: Pigmented Lesion Subcommittee consensus statement. JAMA Dermatol. 2015;151:212–8.

2. Kim CC, Berry EG, Marchetti MA, Swetter SM, Lim G, Grossman D, et al. Risk of subsequent cutaneous melanoma in moderately dysplastic nevi excisionally biopsied but with positive histologic margins. JAMA Dermatol. 2018;154:1401–8.
3. Hiscox B, Hardin MR, Orengo IF, Rosen T, Mir M, Diwan AH. Recurrence of moderately dysplastic nevi with positive histologic margins. J Am Acad Dermatol. 2017;76:527–30.
4. Cockerell CJ, Berson DS. A retrospective look at dysplastic nevi. What were they in 1978 and how have they fared since? Am J Dermatopathol. 1985;7 Suppl:93–7.
5. Kmetz EC, Sanders H, Fisher G, Lang PG, Maize JC. The role of observation in the management of atypical nevi. South Med J. 2009;102:45–8.
6. Goodson AG, Florell SR, Boucher KM, Grossman D. Low rates of clinical recurrence after biopsy of benign to moderately dysplastic melanocytic nevi. J Am Acad Dermatol. 2010;62:591–6.
7. Hocker TL, Alikhan A, Comfere NI, Peters MS. Favorable long-term outcomes in patients with histologically dysplastic nevi that approach a specimen border. J Am Acad Dermatol. 2013;68:545–51.