

Classification des Histiocytoses



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Spectre nosologique des histiocytoses (1)

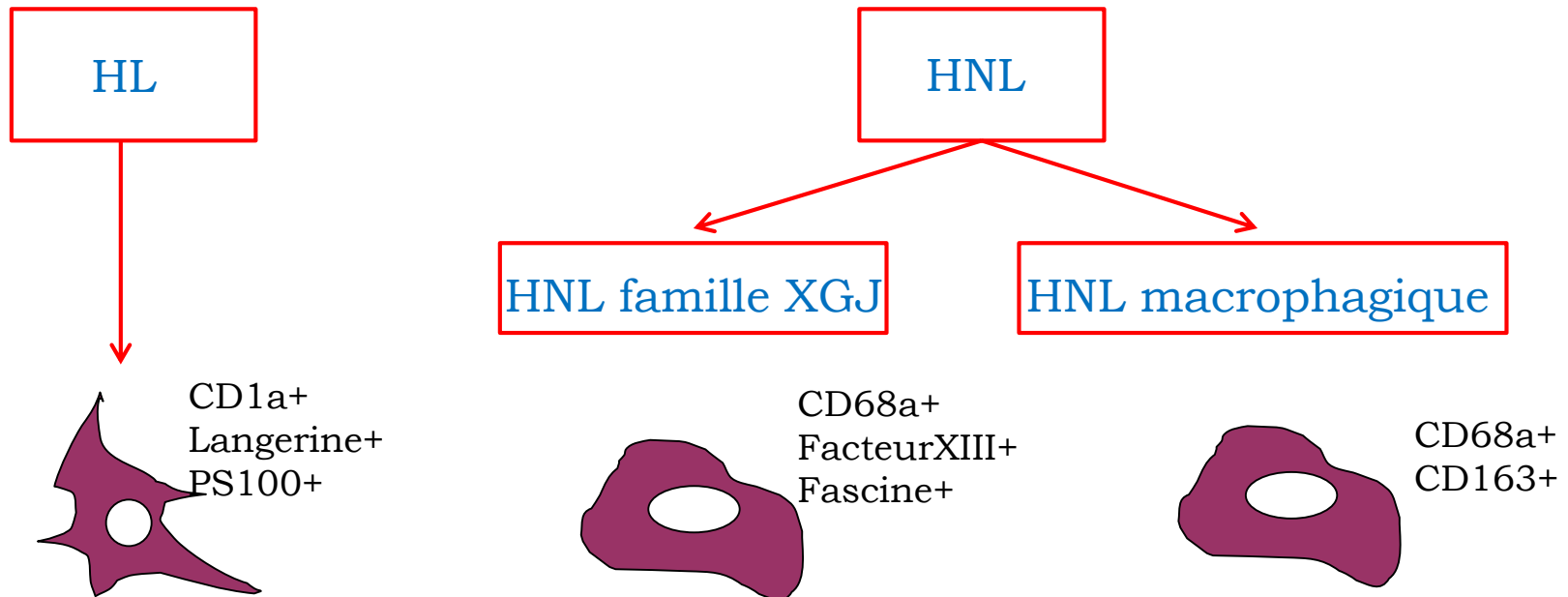
HL

HNL de la famille du XGJ

HNL macrophagiques

	XGJ	H. de surcharge exogènes
	XG de type adulte	H. métaboliques
	Xanthoma disseminatum	H. infectieuses
	H. bénigne céphalique	H. hémophagocytaires avec activation macrophagique
	H. généralisée éruptive	Maladie de Destombes-Rosai-Dorfman
	H. progressive nodulaire	Réticulohistiocytose multicentrique
	Maladie d'Erdheim-Chester	
	<i>H. à cellules indéterminées (?)</i>	

Spectre nosologique des histiocytoses (1)

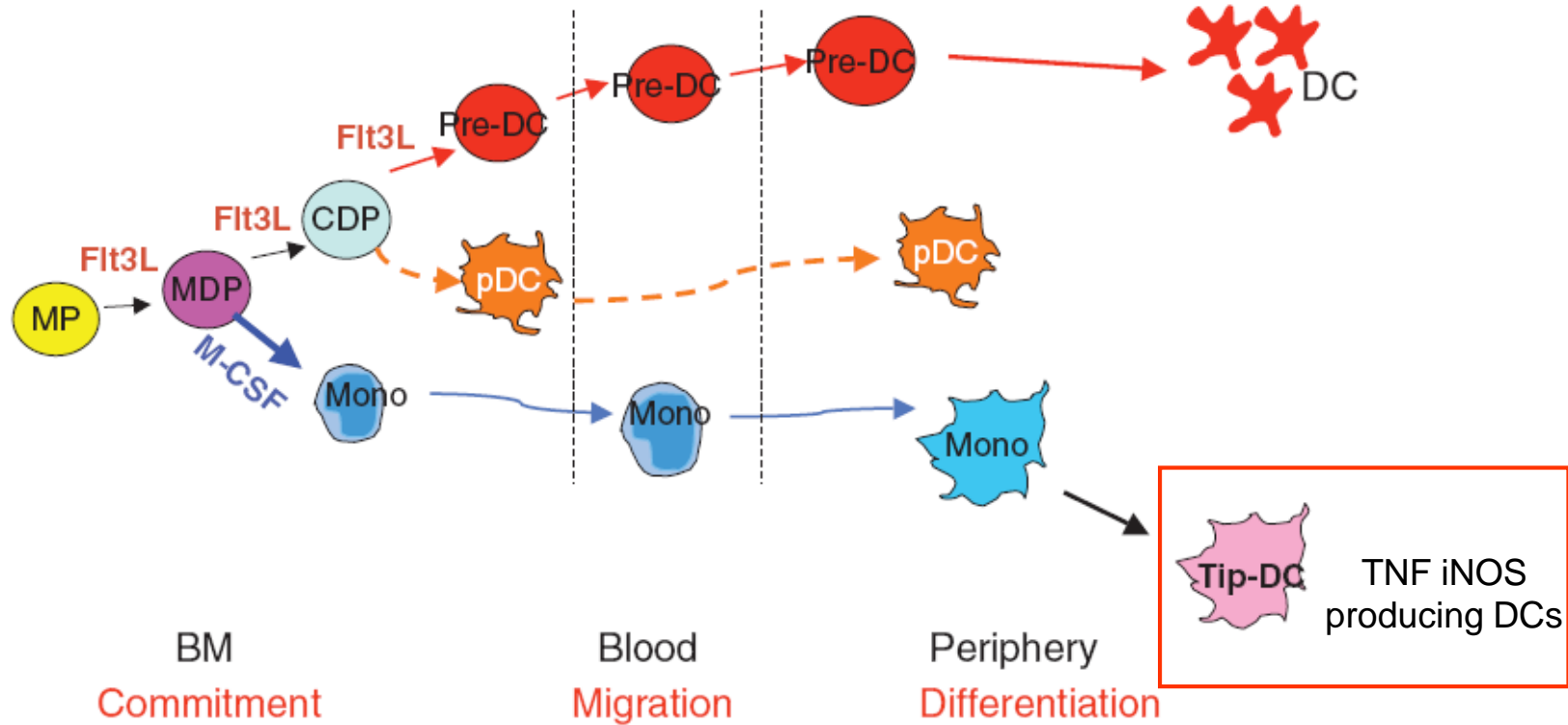


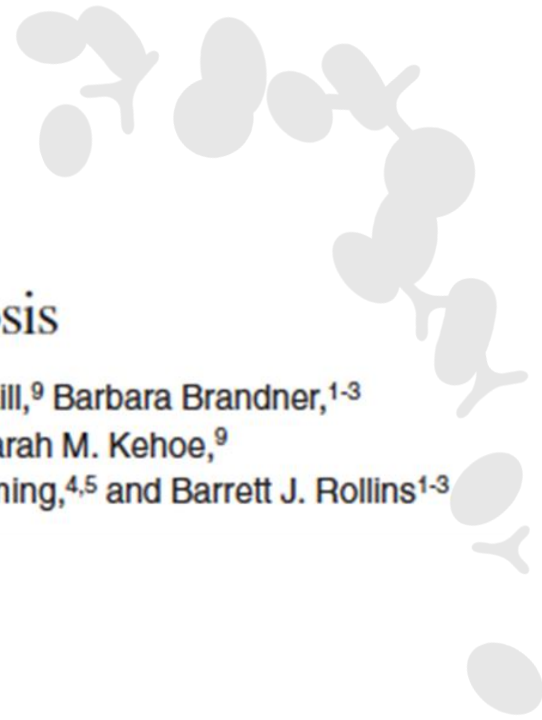
Histiocytoses mixtes, formes de passage

Hervier et al Blood 2014

Kang Liu
Michel C. Nussenzweig

Origin and development of dendritic cells





Brief report

Recurrent *BRAF* mutations in Langerhans cell histiocytosis

Gayane Badalian-Very,¹⁻³ Jo-Anne Vergilio,^{4,5} Barbara A. Degar,⁶⁻⁸ Laura E. MacConaill,⁹ Barbara Brandner,¹⁻³ Monica L. Calicchio,⁴ Frank C. Kuo,^{5,10} Azra H. Ligon,^{5,10,11} Kristen E. Stevenson,¹² Sarah M. Kehoe,⁹ Levi A. Garraway,^{1-3,9,13} William C. Hahn,^{1-3,9,13} Matthew Meyerson,^{1,2,9,13} Mark D. Fleming,^{4,5} and Barrett J. Rollins¹⁻³

BLOOD, 16 SEPTEMBER 2010 • VOLUME 116, NUMBER 11

High prevalence of *BRAF V600E* mutations in Erdheim-Chester disease but not in other non-Langerhans cell histiocytoses

Julien Haroche, Frédéric Charlotte, Laurent Arnaud, Andreas von Deimling, Zofia Hélias-Rodzewicz, Baptiste Hervier, Fleur Cohen-Aubart, David Launay, Annette Lesot, Karima Mokhtari, Danielle Canioni, Louise Galmiche, Christian Rose, Marc Schmalzing, Sandra Croockewit, Marianne Kambouchner, Marie-Christine Copin, Sylvie Fraitag, Felix Sahm, Nicole Brousse, Zahir Amoura, Jean Donadieu and Jean-François Emile

Blood. 2012



Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages

Jean-François Emile,^{1,2} Oussama Abla,³ Sylvie Fraitag,⁴ Annacarin Horne,⁵ Julien Haroche,^{6,7} Jean Donadieu,^{1,8} Luis Requena-Caballero,⁹ Michael B. Jordan,¹⁰ Omar Abdel-Wahab,¹¹ Carl E. Allen,¹² Frédéric Charlotte,^{7,13} Eli L. Diamond,¹⁴ R. Maarten Egeler,³ Alain Fischer,^{15,16} Juana Gil Herrera,¹⁷ Jan-Inge Henter,¹⁸ Filip Janku,¹⁹ Miriam Merad,²⁰ Jennifer Picarsic,²¹ Carlos Rodriguez-Galindo,²² Barret J. Rollins,^{23,24} Abdellatif Tazi,²⁵ Robert Vassallo,²⁶ and Lawrence M. Weiss,²⁷ for the Histiocyte Society

BLOOD, 2 JUNE 2016 • VOLUME 127, NUMBER 22

L Group

- LCH
- ICH
- ECD
- Mixed LCH/ECD

C Group

- Cutaneous non-LCH
 - XG family: JXG, AXG, SRH, BCH, GEH, PNH
 - Non-XG family: cutaneous RDD, NXG, other NOS
- Cutaneous non-LCH with a major systemic component

R Group

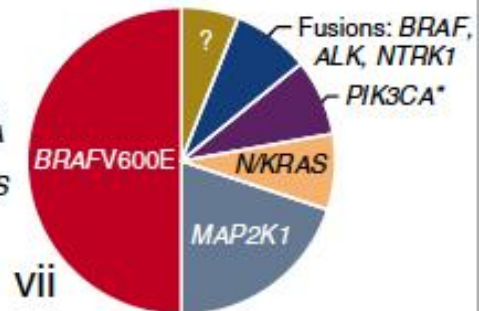
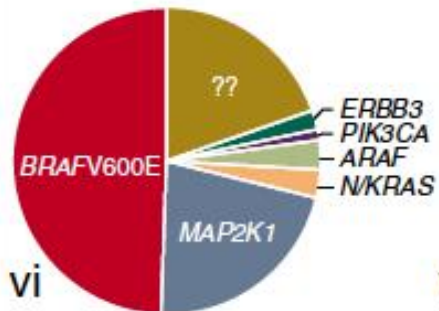
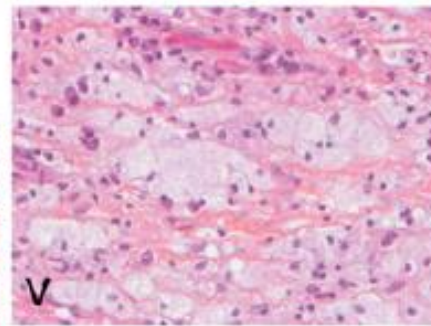
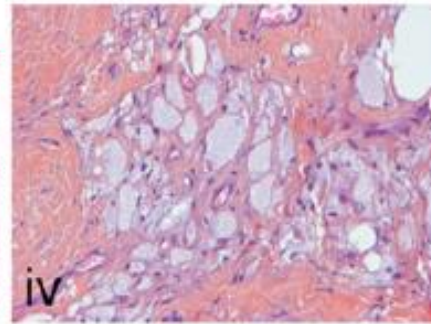
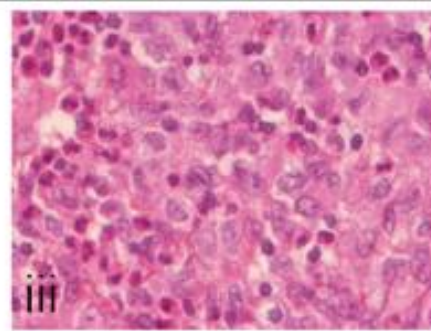
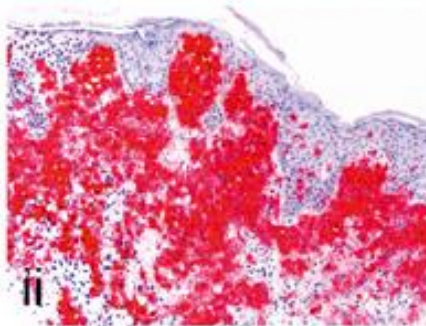
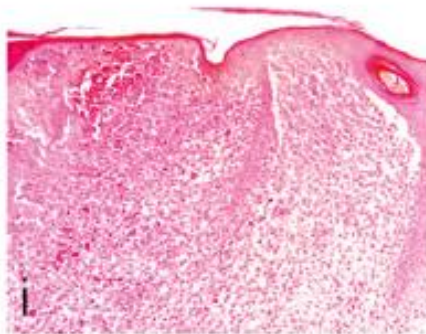
- Familial Rosai-Dorfman Disease (RDD)
- Sporadic RDD
 - Classical RDD
 - Extra-nodal RDD
 - RDD with neoplasia or immune disease
 - Unclassified

M Group

- Primary Malignant Histiocytoses
- Secondary Malignant Histiocytoses (following or associated with another hematologic neoplasia)
Subtypes: Histiocytic, Interdigitating, Langerhans, Indeterminate Cell

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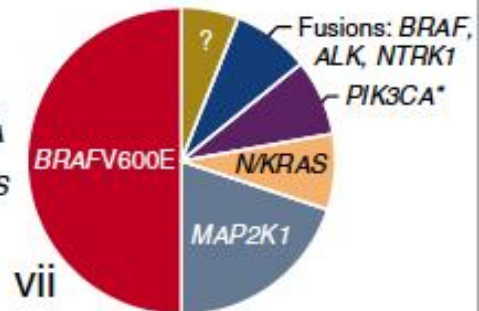
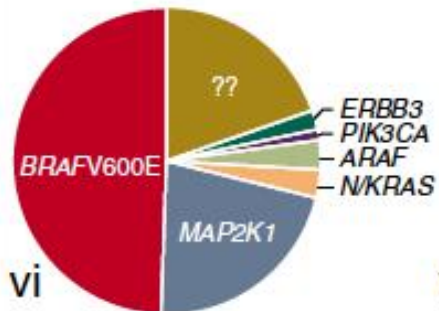
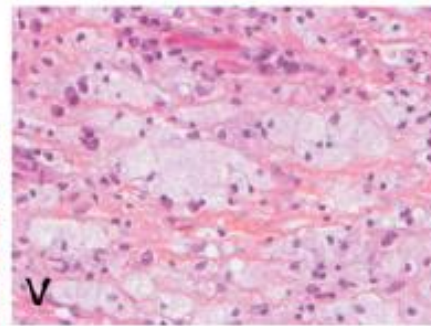
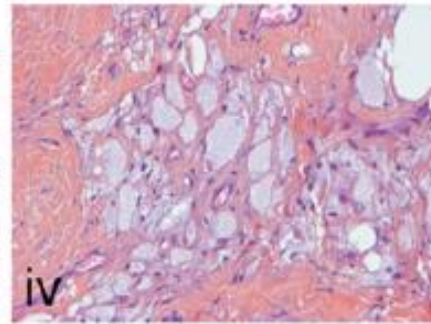
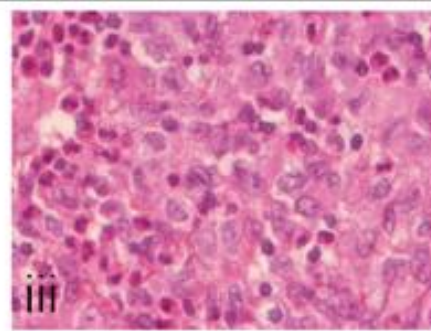
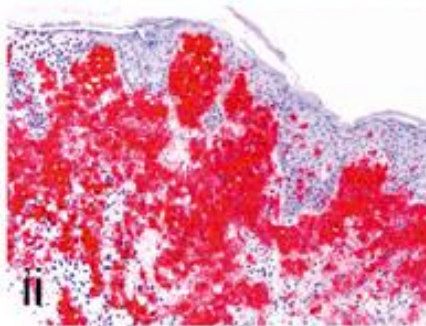
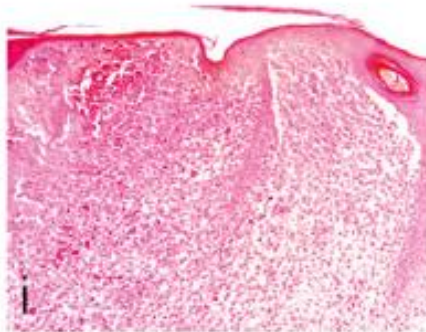
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* A proportion of *PIK3CA* mutant patients have concomitant *BRAFV600E* mutations.

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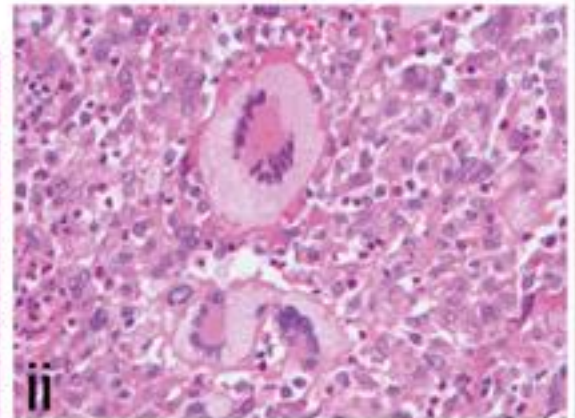
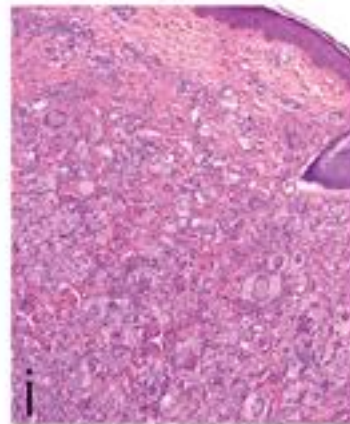
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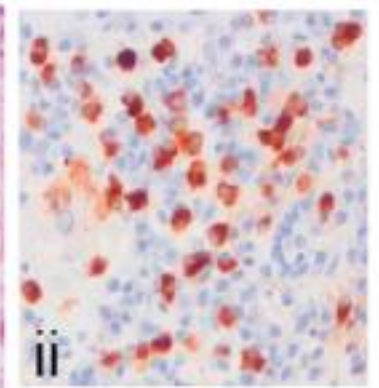
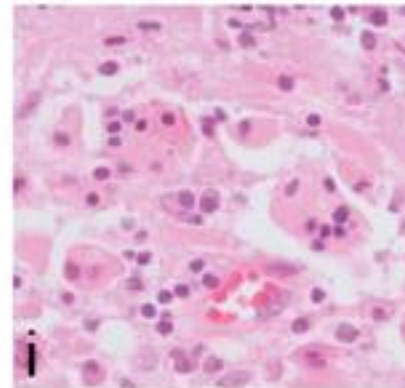
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