

## Classification des Histiocytoses



Abdellatif Tazi  
Service de Pneumologie  
Hôpital Saint-Louis, Paris, France  
[abdellatif.tazi@aphp.fr](mailto:abdellatif.tazi@aphp.fr)

# Spectre nosologique des histiocytoses (1)

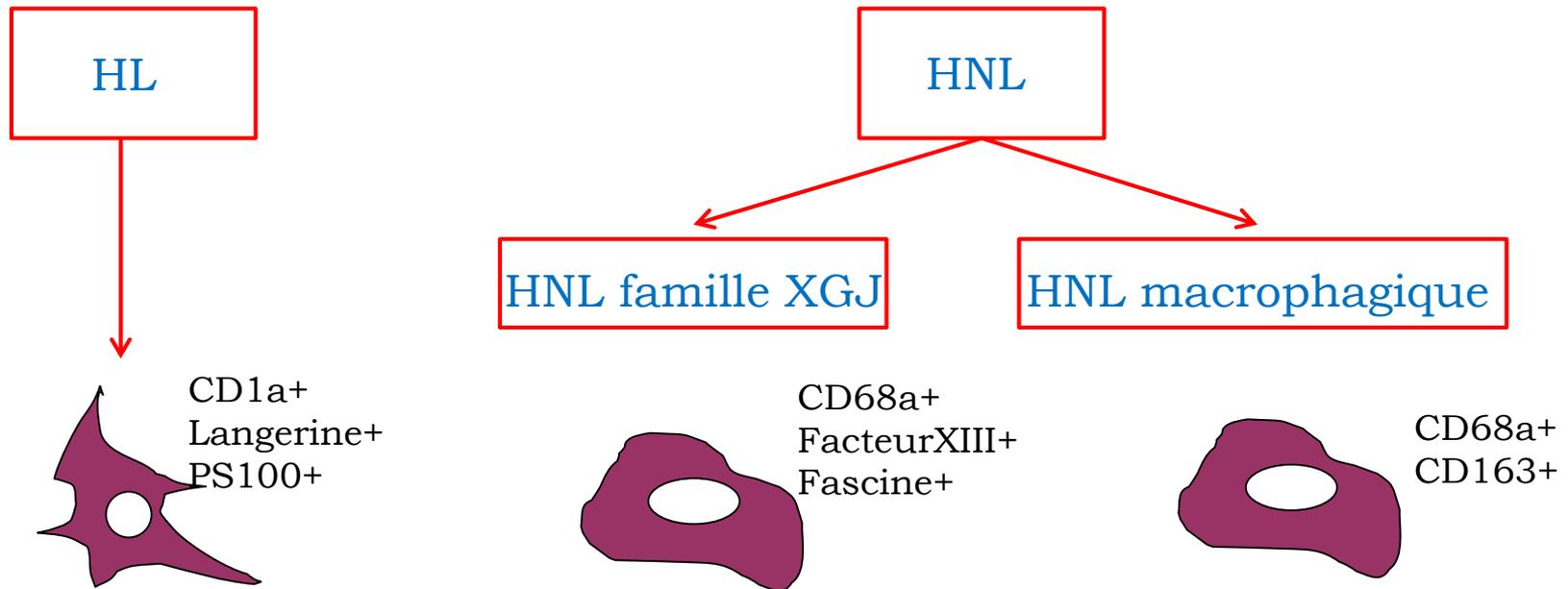
**HL**

**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)

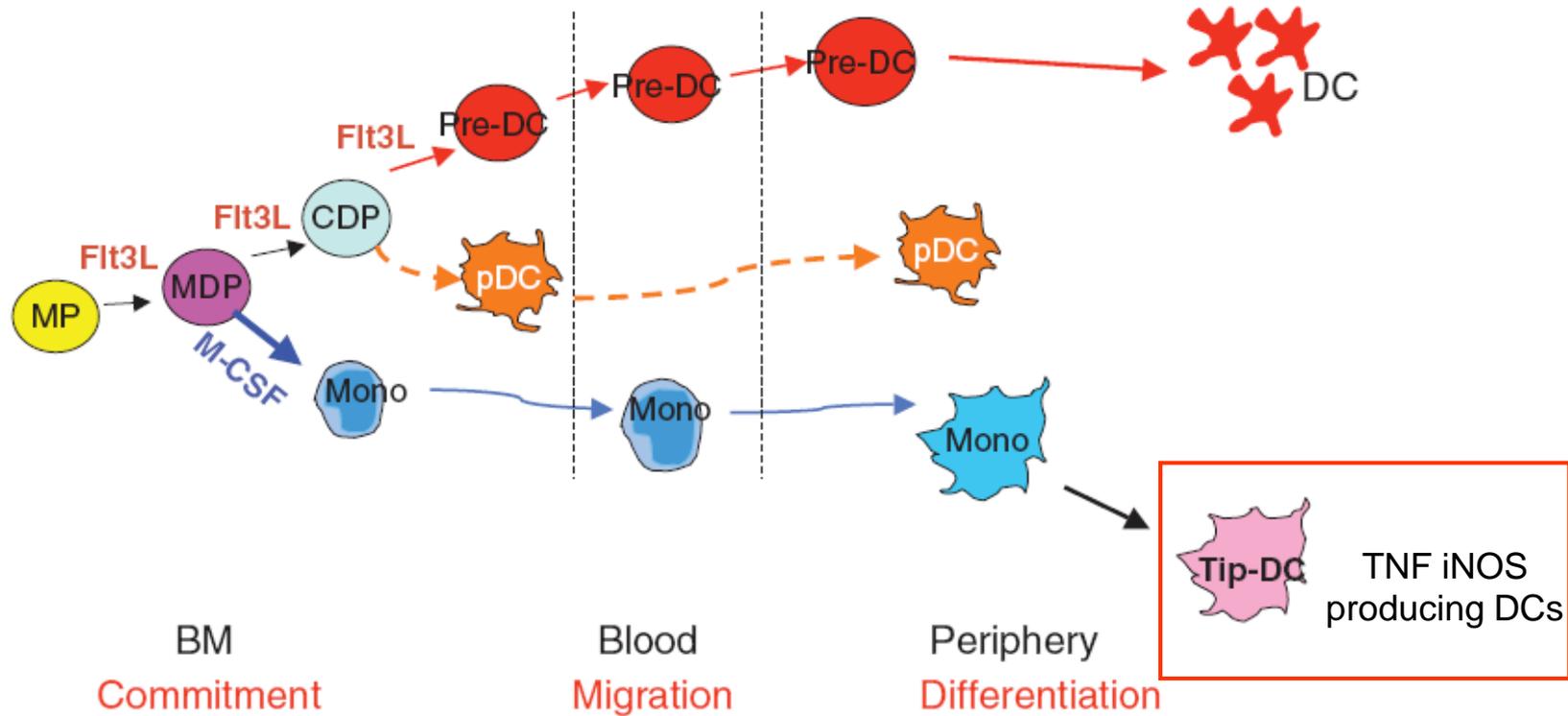


Histiocytoses mixtes, formes de passage

*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,<sup>1-3</sup> Jo-Anne Vergilio,<sup>4,5</sup> Barbara A. Degar,<sup>6-8</sup> Laura E. MacConaill,<sup>9</sup> Barbara Brandner,<sup>1-3</sup> Monica L. Calicchio,<sup>4</sup> Frank C. Kuo,<sup>5,10</sup> Azra H. Ligon,<sup>5,10,11</sup> Kristen E. Stevenson,<sup>12</sup> Sarah M. Kehoe,<sup>9</sup> Levi A. Garraway,<sup>1-3,9,13</sup> William C. Hahn,<sup>1-3,9,13</sup> Matthew Meyerson,<sup>1,2,9,13</sup> Mark D. Fleming,<sup>4,5</sup> and Barrett J. Rollins<sup>1-3</sup>

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

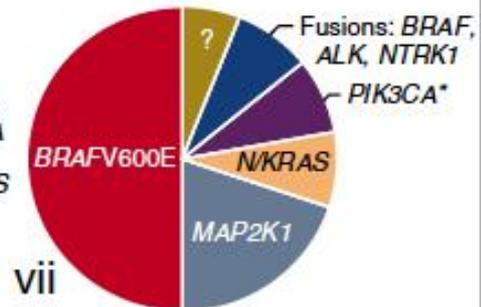
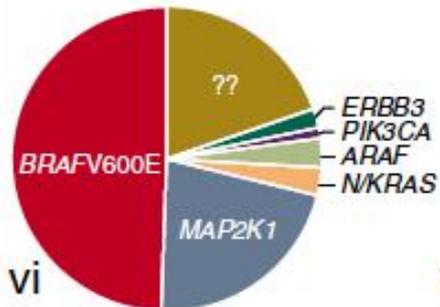
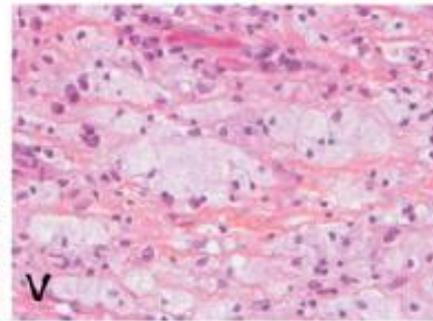
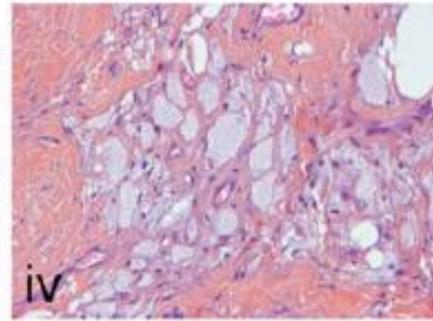
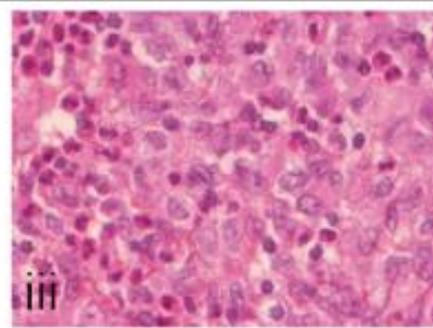
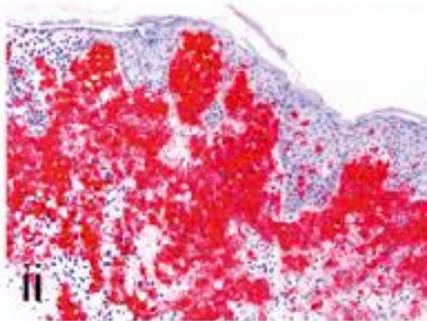
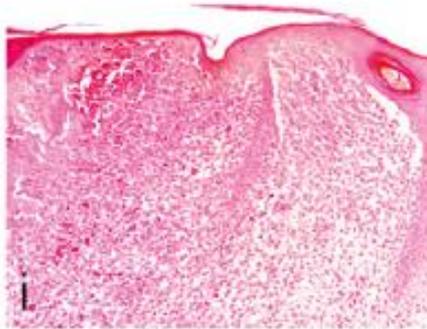
- Familial Rosai-Dorfman Disease (RDD)
- Sporadic RDD
  - Classical RDD
  - Extra-nodal RDD
  - RDD with neoplasia or immune disease
  - Unclassified

## M Group

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

## L Group

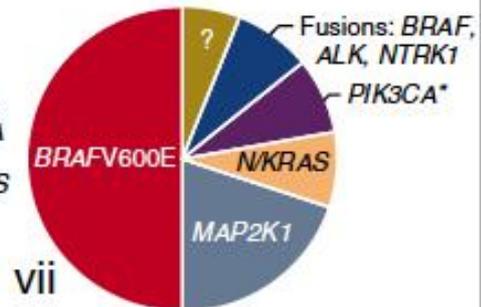
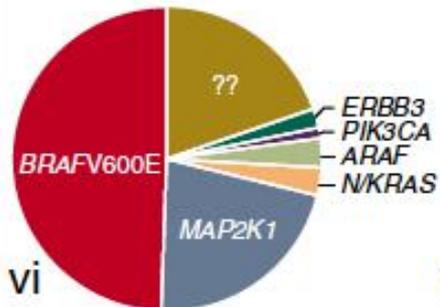
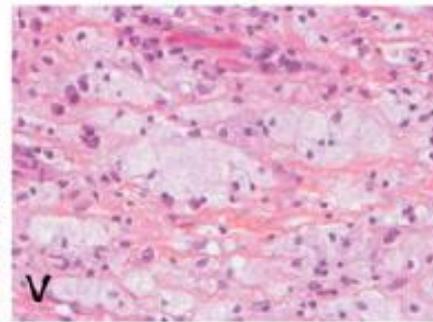
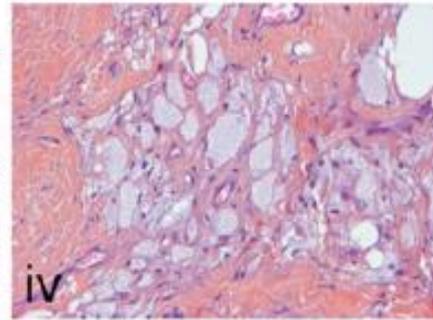
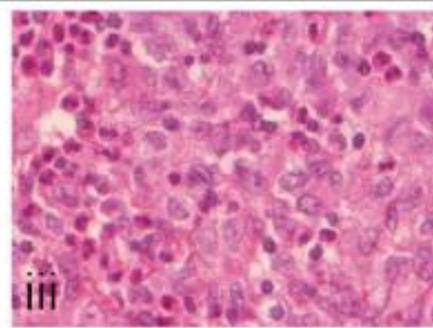
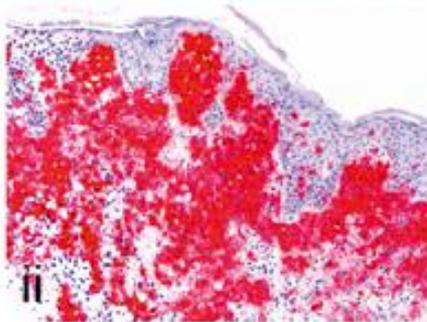
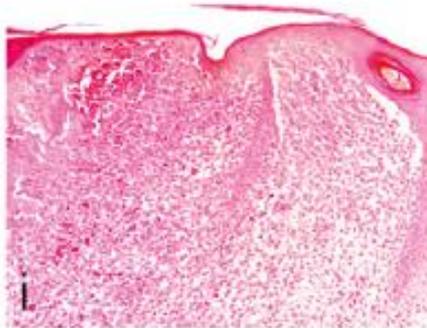
- LCH
- ICH
- ECD
- Mixed LCH/ECD



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

## L Group

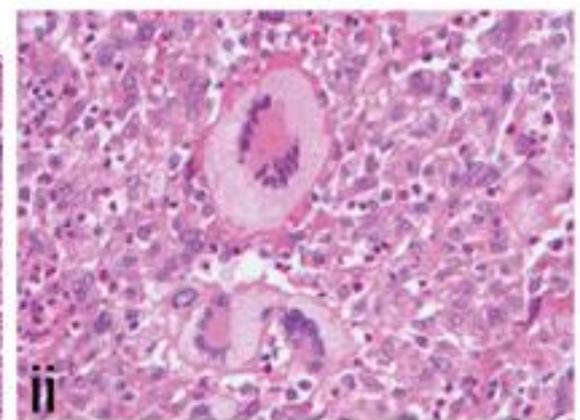
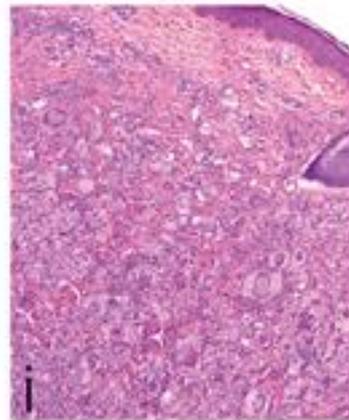
- LCH
- ICH
- ECD
- Mixed LCH/ECD



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

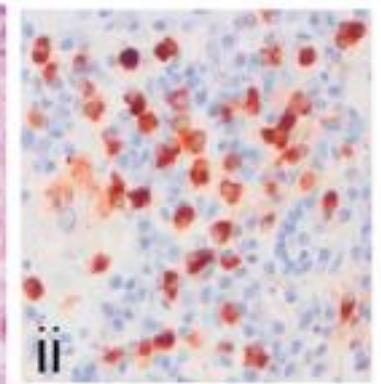
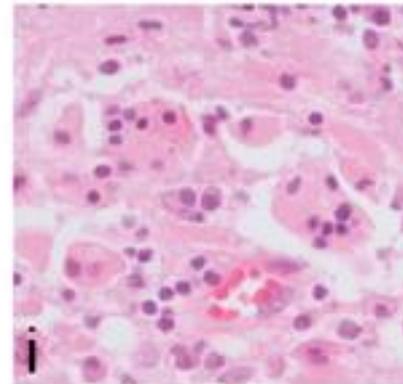
## C Group

- 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*

