



# Pachyonychia Congenita Project

*Fighting for a cure. Connecting & helping patients. Empowering research.*



## **International Patient Meeting - Pachyonychia Congenita.**

### **Patients, Family Members, Physicians and Scientists**

Organizers: *Mary E Schwartz, Fernando Larcher y Marcela Del Río*

*Madrid, October 22nd*

CIEMAT, Avenida Complutense 40, Madrid

9:00: Welcome: Fernando Larcher and Marcela Del Río (CIEMAT-CIBERER and IIS-Fundación Jiménez Díaz  
"The World of Genodermatoses")

9:30: About PC Project, Cindy Atha (with translator)

10:00: What is PC?: My Story, Pamela Ibáñez Triguero

10:30 Break

11:00 Genetics of PC Frances Smith

11:30: Specialists Panel - Question and Answer Session:

Eulalia Baselga, MD, Jessica Gonzalez, PhD, Alain Hovnanian, MD, PhD, Roger Kaspar, PhD, Fernando Larcher, PhD. Vanesa Lopez

Gonzalez, MD, Frances Smith, PhD and Marcela Del Río, PharmD/PhD

12:15: PC Research, Roger Kaspar

1:00 Lunch

2:30 Breakout Sessions (children will join)

Experts GROUP 1 PC-K6a and PC-K17 Patients

GROUP 2 PC-K16 Patients

Specialists TEAM A Roger Kaspar (rotate 15 mins)

Specialists TEAM B Frances Smith (rotate 15 mins)

3:00 Care Tips Discussion, Claudia Avella

3:30 Closing Remarks, Cindy Atha

INSCRIPTIONS: <https://www.surveymonkey.com/r/PC2016SPAIN>