



Pachyonychia Congenita Project

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

International Patient Meeting - Pachyonychia Congenita Patients, Family Members, Physicians and Scientists

Saturday, October 22th, 2016

Meeting sessions from 9.00 am to 5.30 pm

LOCATION: CIEMAT. Avenida Complutense 40. Madrid, Spain

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

- 9:00 Welcome: Fernando Larcher, “The World of Genodermatoses”.
- 9:30 “About PC Project”, Cindy Atha.
- 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
 - Marcela Del Río, PhD
- 12:15 “PC Research”, Roger Kaspar.
- 1:00 Lunch.
- 2:30 Breakout Sessions :
 - 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.



Universidad
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