First DEGOS disease workshop

(Malignant atrohic papulosis) :

An ultra-rare disease that must be recognized

Friday 9th January 2026, virtual workshop :

https://us06web.zoom.us/j/83360069114?pwd=aYie5i0w68WeoHRu7ZMvlYH5wKxAPq.1

Scientific committee Christine BODEMER, Eric HACHULLA, Arnaud HOT, Axel VILLANI

15h00-15h30 : Physiopathological aspects Axel Villani (chair Emmanuel LEDOULT)

15h30-16h00 : Clinical aspects and therapeutic approaches Marie ROBERT (chair Arnaud HOT)

16h00-16h30 : Degos Disease Foundation: Nature of patient and physician inquiries, Study of long-term survivors. NIH: Degos research update (chair Eric HACHULLA)

16h30-17h00 : 2 Clinical cases Axel VILLANI/Marie ROBERT and Emmanuel LEDOULT (chair Christine BODEMER)

17h00-17h10 : Patient testimony

17h10-18h00 : Atrophic papulosis (Köhlmeier-Degos disease) in children and adolescents-A cross-sectional study and literature review by Christos ZOUBOULIS (chair Axel VILLANI)

Conclusions Christine BODEMER and Éric Eric HACHULLA