CLINICAL TRIAL

PHENOROD2 - NCT04285398

Retinal RED

Observational

Prospective Natural History Study of Retinitis Pigmentosa (PHENOROD2)

Rare Eye Disease concerned by the trial :
Retinitis Pigmentosa

Status of the trial : recruiting

Orphan drug recognition : Yes

Inclusion criteria : RP with mutations affecting the RHO, PDE6A and PDE6B genes
Visual acuity ≥ 20/200 for at least one eye
Binocular Visual field diameter ≥ 5° as measured on the Goldmann III-4e isopter

Exclusion criteria : Patients with any other gene mutation known to be involved in RP
Patients with other ocular disorder likely to impact the visual function
Pregnant or breastfeeding women

Inclusion opening date : 01/02/2020

Inclusion closing date (previsional) : 01/03/2020

Children ☐

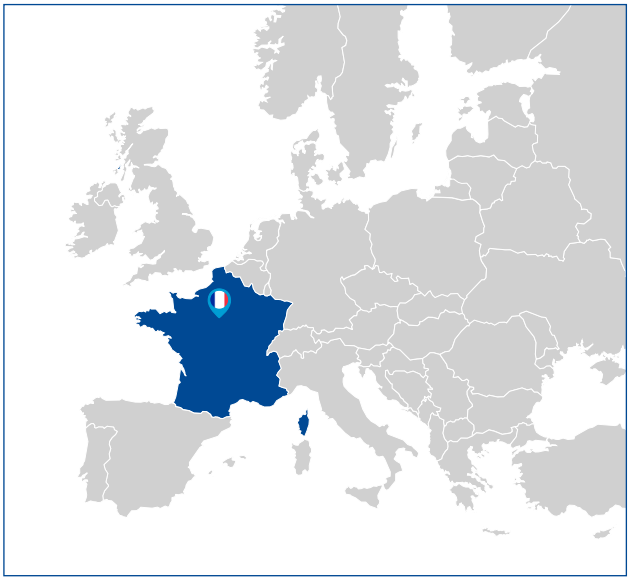
Adults ☑

Within ERN-EYE members

Location of the trial :

CHNO des XV XX - Centre Hospitalier National d'Ophtalmologie
28 rue de Charenton
75557 Paris Cedex 12
France

Contact details



Principal investigator name :
Dr Saddek Mohand Saïd

Other investigators :
Pr Isabelle Audo
Pr José-Alain Sahel

Funder type : Private

More informations