CLINICAL TRIAL

GEPHIRD - NCT03662386

Retinal RED

Observational

Prospective Analysis of «Genotype-phenotype» Correlations Observed in a Large Cohort of Patients With Hereditary Retinal Dystrophies - GEPHIRD

Rare Eye Disease concerned by the trial :
Hereditary retinal dystrophies

Status of the trial : recruiting

Orphan drug recognition : Yes

Inclusion criteria : Patients hospitalized for suspicion of hereditary retinal dystrophy
Benefiting as part of the care of a genetic analysis

Exclusion criteria : Patient under a measure of legal protection

Inclusion
opening date : 01/09/2018

Inclusion closing
date (previsional) : 01/09/2032

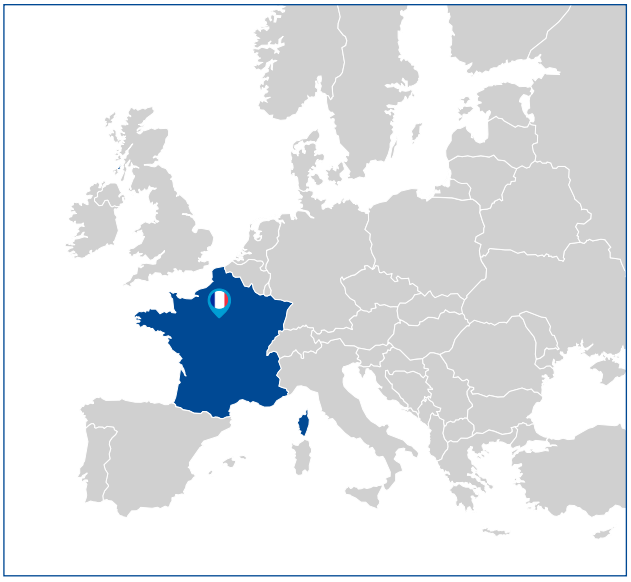
Children ☑

Adults ☑

Within ERN-EYE members

Location of the trial :

Fondation Ophtalmologique Adolphe de Rothschild
29 Rue Manin
75019 Paris



Principal investigator name :
Dr Elise Boulanger

Other investigators :
None

Funder type : Public

More informations