CLINICAL TRIAL

NCT01235624

Retinal RED

Interventional

Autosomal Dominant Retinitis Pigmentosa: Prevalence of Known Genes Identification of New Loci / Genes

Rare Eye Disease concerned by the trial :
Autosomal Dominant Retinitis Pigmentosa

Status of the trial : completed

Orphan drug recognition : N/A

Inclusion criteria : Click here to see detailed criteria

Exclusion criteria : Click here to see detailed criteria

Inclusion opening date : 01/03/2009

Inclusion closing date (previsional) : 01/04/2013

Children ☑

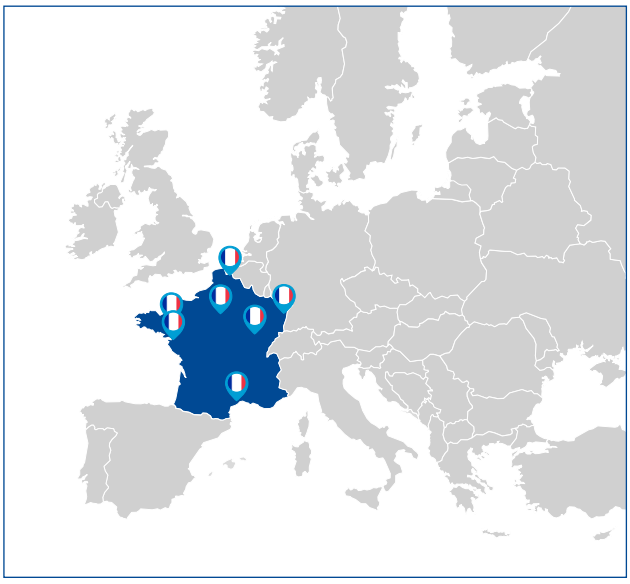
Adults ☑

Within ERN-EYE members

Principal location of the trial :

Maolya - National Rare Disease Center
CHU de Montpellier
191 avenue du Doyen Gaston Giraud
34295 Montpellier cedex 05

Contact details



Principal investigator name :
Christian HAMEL, Isabelle Meunier

Other investigators :
J-A SAHEL, CHNO des XV-XX, Paris
Jean Louis DUFIER, Hôpital Necker, Paris
Hélène DOLLFUS, CHU de Strasbourg
Sylvie ODENT, CHU de RENNES
Sabine DEFOORT, CHRU de Lille
Michel WEBER, CHU de Nantes
Xavier ZANLONGH, Nantes
L. LO OLIVIER-FAIVRE, Children Hospital of DIJON

Funder type : Other

More informations