CLINICAL TRIAL

VITAL

Low vision

Observational

Prospective study of visual function in USHER syndrome induced by MYO7A mutation and retinotosis pigmentosa

Rare Eye Disease concerned by the trial :
USHER syndrome induced by MYO7A mutation and and pigmentary retinopathy

Status of the trial : recruiting

Orphan drug recognition : N/A

Inclusion criteria : USH1B group: genotype with two pathogenic mutations of the MYO7A gene
RP group: a genotype with a pathogenic mutation identified.
For all: Diagnosis of confirmed retinopathy pigmentosa

Exclusion criteria : Persons who have previously participated in a clinical trial of gene therapy or with ocular disease or ocular opacity. History of amblyopia

Inclusion opening date : 03/11/2017

Inclusion closing date (previsional) : 04/11/2019

Children ☑

Adults ☑

Within ERN-EYE members

Principal location of the trial :

REFERET - Centre de référence des Maladies Rares neuro-rétiniennes
CHNO des XV XX
28 rue de charenton
75012 Paris
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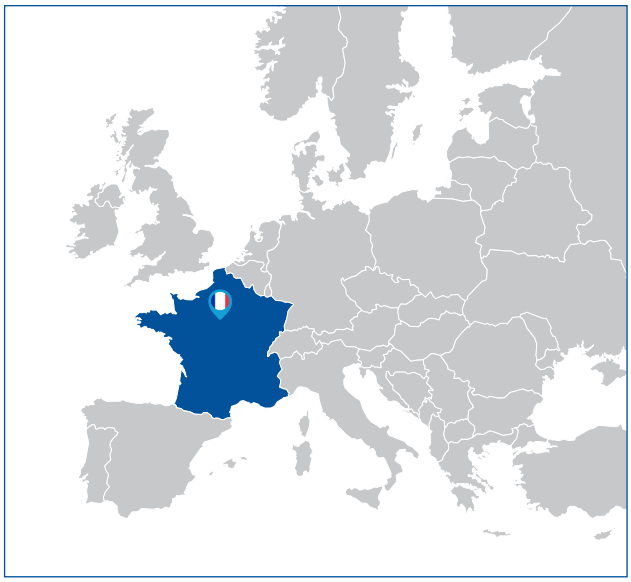
Contact details



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Funder type : Drug company - Industry
Public Foundation