

# CLINICAL TRIAL

**XOLARIS - NCT03314207**

Retinal RED



## Natural history of the progression of X-linked retinitis pigmentosa.

Rare Eye Disease concerned by the trial :  
X-linked retinitis pigmentosa with a mutation  
in RPGR gene

Status of the trial : recruiting  
Orphan drug recognition : N/A

Inclusion criteria : Visual Acuity from 20/200 to 20/20

Exclusion criteria : Age < 18 years



Inclusion  
opening date : 01/01/2018



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

Children



Adults



## Within ERN-EYE members



Principal location of the trial :

**Maolya - National Rare Disease center**  
Centre national de référence affections  
sensorielles génétiques  
CHU Gui de Chauliac  
80, avenue Augustin Fliche  
34 295 Montpellier Cedex 5

[Contact details](#)

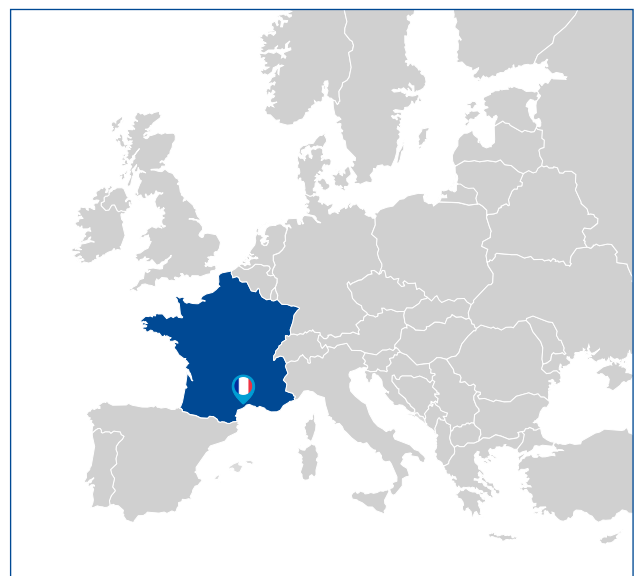


Principal investigator name :  
Isabelle Meunier



Other investigators :  
None

Funder type : Drug company - Industry



[More informations](#)

# CLINICAL TRIAL

## XOLARIS

Retinal RED  
Registries



### Multi-centre, prospective, observational study of the Natural History of the Progression of X-Linked Retinitis Pigmentosa

Rare Eye Disease concerned by the trial :  
X-linked RP (RPGR)

Status of the trial : recruiting  
Orphan drug recognition : N/A

Inclusion criteria : Molecular confirmed mutation in RPGR

Exclusion criteria : Visual Acuity worse than 6/60



Inclusion  
opening date : 01/11/2017



Inclusion closing  
date (previsional) : 01/08/2018

Children



Adults



### Within ERN-EYE members



Principal location of the trial :

**Leeds Teaching Hospital (St.James's Hospital)**

Beckett Street  
Leeds LS9 7TF  
West Yorkshire  
UK

[Contact details](#)



Principal investigator name :

Kamron Khan

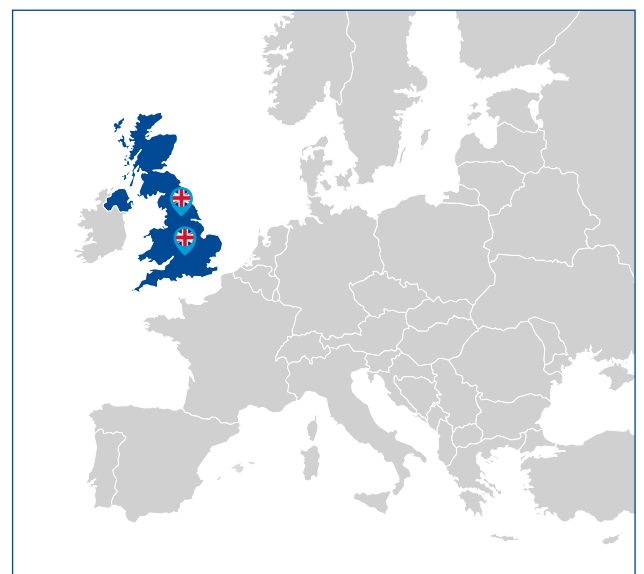


Other investigators :

Robert MacLaren (Chief Investigator)

**Oxford Eye Hospital**

Level Lg1 John Radcliffe Hospital  
Headley Way, Headington  
Oxford OX3 9DU  
UK



Funder type : Drug company - Industry

[More informations](#)

# CLINICAL TRIAL

## XOLARIS

Research



### Natural History of the Progression of X-Linked Retinitis Pigmentosa

Rare Eye Disease concerned by the trial :  
X-Linked Retinitis Pigmentosa

Status of the trial : not yet recruiting  
Orphan drug recognition : N/A

**Inclusion criteria :** Patients are willing and able to provide informed consent for participation in the study, are male  $\geq 16$  years of age, have a genetically confirmed diagnosis of XLRP (with retinitis pigmentosa GTPase regulator [RPGR] mutation), have active disease clinically visible within the macular region, are willing and able to undergo ophthalmic examinations once every 3 months for up to 24 months, have an ETDRS BCVA better than or equal to 34 letters in at least one eye.

**Exclusion criteria :** [click here to see detailed criteria](#)



Inclusion opening date : 01/12/2017



Inclusion closing date (previsonal) : 01/12/2018

Children



Adults



### Within ERN-EYE members



Principal location of the trial :

**Center for Ophthalmic Research, University Hospital of Tübingen**  
Elfriede-Aulhorn-Str. 7  
72076 Tübingen  
Germany

[Contact details](#)



Principal investigator name :

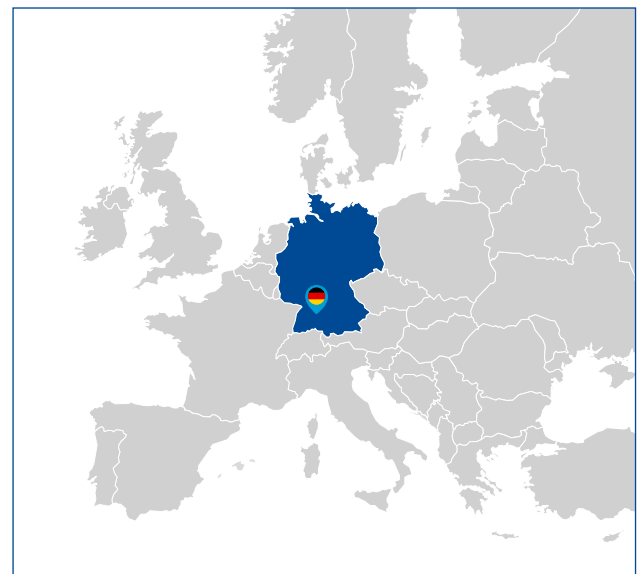
Prof. Dr. Dr. Dominik Fischer



Other investigators :

Prof. Dr. med. Karl Ulrich Bartz-Schmidt

**University Hospital of Tübingen**  
Tübingen, Germany



Funder type : Drug company - Industry

[More informations](#)