CLINICAL TRIAL

2013-005393-22

Retinal RFD





A Study of the Efficacy and Safety of QLT091001 in Subjects with Inherited Retinal Disease (IRD) Caused by Mutation in Retinal Pigment Epithelium Protein 65 (RPE65) or Lecithin: Retinal Acyltransferase (LRAT)

Rare Eye Disease concerned by the trial:

Retinitis pigmentosa; Inherited Retinal Disease caused by Mutation in RPE65 or LRAT genes

Inclusion criteria: click here to see detailed criteria

Exclusion criteria: click here to see detailed criteria

Inclusion

opening date: 06/09/2016



Inclusion closing

date (previsional): 06/09/2019

Status of the trial: not yet recruiting

Orphan drug recognition: Yes







Within ERN-EYE members



Principal location of the trial:

Center for Ophthalmology, University of Tübingen

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Contact details



Principal investigator name:

Eberhart Zrenner



Other investigators:

not known yet



Funder type: Drug company - Industry