

CLINICAL TRIAL

DIIO_EPPR_P13_03

Retinal RED
Low vision



Retinitis pigmentosa: molecular diagnosis by next generation sequencing

Rare Eye Disease concerned by the trial :
inherited retinal dystrophy

Status of the trial : Active, not recruiting

Orphan drug recognition : N/A

Inclusion criteria : probands or family pedigrees with inherited retinal dystrophy

Exclusion criteria : patients or family pedigrees with retinal abnormalities mimicking inherited retinal dystrophy



Inclusion
opening date : 18/06/2013



Inclusion
closing date : 19/09/2016

Children



Adults



Within ERN-EYE members



Principal location of the trial :

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Principal investigator name :

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Other investigators :

Francesco Parmeggiani

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Funder type : Patient organization

[More informations](#)