

14 PARTNERS ACROSS EUROPE: A collaborative research effort to find a cure for AMD



Eberhard Karls Universität Tübingen (EKUT)

Tübingen, Germany



Erasmus Universitair Medisch Centrum Rotterdam (EMC)

Rotterdam, Netherlands



Fundacio Centre de Regulacio Genomica (CRG)

Barcelona, Spain



Stichting Katholieke Universiteit (RUNMC)

Nijmegen, Netherlands



Université de Bordeaux (UB)

Bordeaux, France



University College London (UCL)

London, UK



Barcelona Macula Foundation Research for Vision (BMF)

Barcelona, Spain



PRO RETINA Deutschland (PROR)

Bonn, Germany



AYOXXA Biosystems GmbH (AYOX)

Cologne, Germany



F. Hoffmann-La Roche AG (ROCHE)

Basel, Switzerland



Moorfields Eye Hospital NHS Foundation Trust (MEH)

London, UK



Fundacion Publica Andaluza Progreso y Salud (FPS)

Sevilla, Spain



Queen's University of Belfast (QUB)

Belfast, UK



University College Dublin (NUID UCD)

Dublin, UK

COORDINATOR

Prof. Marius Ueffing. Institute for Ophthalmic Research. Centre for Ophthalmology. University of Tübingen.

CO-COORDINATOR

Prof. Caroline Klaver. Erasmus Medical Center. Dept. of Ophthalmology and Epidemiology.

CONTACT

coordinator@eyerisk.eu www.eyerisk.eu @EyeRiskEU



EYE-RISK is funded by the Horizon 2020 program of the European Union. Funding is provided in the period of 2015-2019 under Grant Agreement number 634479.



Exploring the combined role of genetic and non-genetic factors for developing Age-Related Macular Degeneration:

A systems-level analysis of disease subgroups, risk factors and pathways

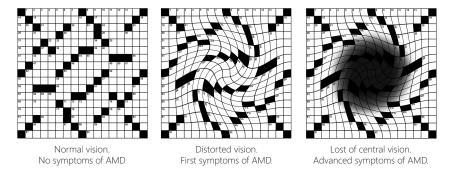




TOWARDS A MODEL THAT DESCRIBES MOLECULAR DRIVERS AND THE INTERACTION OF RISK FACTORS IN AGE-RELATED MACULAR DEGENERATION

People above 60 have an elevated risk to lose eyesight by Age-related macular degeneration.

Age-related macular degeneration (AMD) is a persistent, progressive and incurable disease leading to declining sight that progresses to complete loss of vision. Patients suffering from AMD lose vision in the central part of the retina that is critical for reading, driving a car and recognizing faces.



More than 17.7 million EU inhabitants are currently affected by AMD, with an expected rise to 21 million based on population growth and ageing in the next decades.*

People above 60 have an elevated risk for AMD. Susceptibility is determined by a combination of genes, environment, lifestyle and age (see figure). The interaction of these risk factors during disease onset and progression is not understood and the alterations of cell signalling in AMD are not known.

The EYE-RISK consortium addresses the challenges of risk prediction and pathway identification in AMD.

EYE-RISK research will specify who is at risk of developing AMD, who is at risk for progression, why and how risks combine to advance progression in specific patients and what we can do to lower their risk. The project will also identify molecular drivers for AMD. This will allow better diagnosis, better risk-based prevention strategies and better development of therapies.

AMD RISKS Genes **Environment** Lifestyle Phenotype Age severity **Black Box** onset combined risk progression Web-based risk prediction for patients and physicians **Precision** Medicine

EYE-RISK implements a multi-disciplinary approach.

The approach integrates clinical phenotyping and diagnosis, genotyping, next-generation targeted re-sequencing, bioinformatics and statistics, clinical data analysis, computational biology, systems-biology oriented pathway analysis and modelling.

Find a cure for the AMD disease can save over 50 billion € in health care cost in 10 years.

5 majors goals of EYE-RISK

- 1 Robust algorithms to identify personalised risks of development of advanced AMD, and progression of dry AMD.
- 2 Novel biomarkers for further stratification of disease risk.
- 3 Molecular drivers/ biological pathways relevant for onset and progression of advanced AMD.
- 4 Clinical guidelines for individuals at risk of developing AMD.
- 5 Criteria of inclusion and stratification for patients entering clinical trials.