

## ADVANTAGES TO CLINICAL SITE MEMBERS

- **Participation in subspecialty Scientific Sessions**
- Support for **multinational Investigator-Initiated Research**
- **Receive for free EVICR.net organisational SOPs** and become a certified Clinical Site
- Opportunity to participate in **new multinational clinical research studies**
- Support for **EU Clinical Research applications**
- Support in **grant applications to industry**
- Access to **educational programme and training**
- **Promotion** of your Clinical Research Centre as member of EVICR.net
- **Networking** with other European and USA networks

### BECOME A MEMBER



### STEERING COMMITTEE

Hendrik Scholl (Chairman)  
Francesco Bandello  
José Cunha-Vaz  
Francesca Cordeiro  
Marie-José Tassignon  
Robert Finger  
Birgit Lorenz  
Tunde Peto  
Cecília Martinho

### EDUCATIONAL PROGRAMME



TAKE ADVANTAGE OF WEBINARS  
ON OPHTHALMOLOGY  
CLINICAL RESEARCH

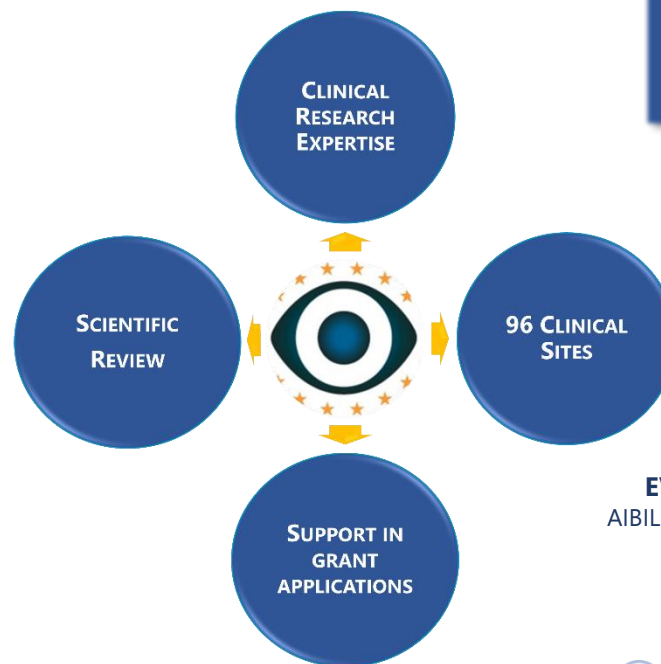


# EVICR.net

EUROPEAN VISION  
CLINICAL RESEARCH

Your partner in  
**INVESTIGATOR-  
INITIATED RESEARCH  
(IIR)**

IF YOU ARE AN  
**INVESTIGATOR**  
AND YOU HAVE AN  
IDEA FOR AN  
**OPHTHALMOLOGY**  
**MULTINATIONAL CLINICAL**  
**STUDY**, EVICR.net  
CAN HELP YOU



### VISIT US



### CONTACT US

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www.evicr.net

## A NETWORK OF 96 CLINICAL RESEARCH SITES FROM 17 COUNTRIES

### WE SUPPORT:

- Design and implementation of multinational clinical research activities in ophthalmology
- Grant applications to Pharmaceutical industry or to European Union

## SCIENTIFIC REVIEW AND EXPERTISE



### EXPERT COMMITTEES AND COORDINATORS

- **AGE-RELATED MACULAR DEGENERATION**  
*Francesco Bandello*
- **RETINAL DYSTROPHIES**  
*Hendrik Scholl*
- **DIABETIC RETINOPATHY & VASCULAR DISEASES**  
*José Cunha-Vaz*
- **GLAUCOMA**  
*Francesca Cordeiro*
- **ANTERIOR SEGMENT**  
*Marie-José Tassignon*
- **OCULAR SURFACE, INFLAMMATION, DRY EYE & ALLERGIES**  
*Robert Finger*

### TRANSVERSAL SECTIONS AND COORDINATORS

- **RARE DISEASES** - Birgit Lorenz
- **READING CENTRES** - Tunde Peto

## WE PROVIDE SUPPORT TO

### PRE-STUDY SERVICES

- Study design
- Study Protocol development
- Informed Consent Form development
- Case Report Form design
- Database validation and implementation
- eCRF management and support
- Clinical Sites feasibility
- SOP development
- Study Documents development
- Regulatory affairs  
(submission and reports)
- Contracts

### IN-STUDY SERVICES

- Study management
- IMP management
- Monitoring
- Data management
- Pharmacovigilance

### POST-STUDY SERVICES

- Biostatistics
- Final Study report
- Publication
- Medical Writing

## HOW TO PROCEED?

### If you are a member:

- Submit your idea by filling the abstract available at [www.evicr.net](http://www.evicr.net)  
(MEMBERS RESTRICTED AREA)



### If you are not an EVICR.net member yet:

- Please contact the Coordinating Centre: [evicrnet@aibili.pt](mailto:evicrnet@aibili.pt)