

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
Pasquale Aragona
Birgit Lorenz
Tunde Peto
Cecília Martinho

EDUCATIONAL PROGRAMME



TAKE ADVANTAGE OF WEBINARS
ON OPHTHALMOLOGY
CLINICAL RESEARCH

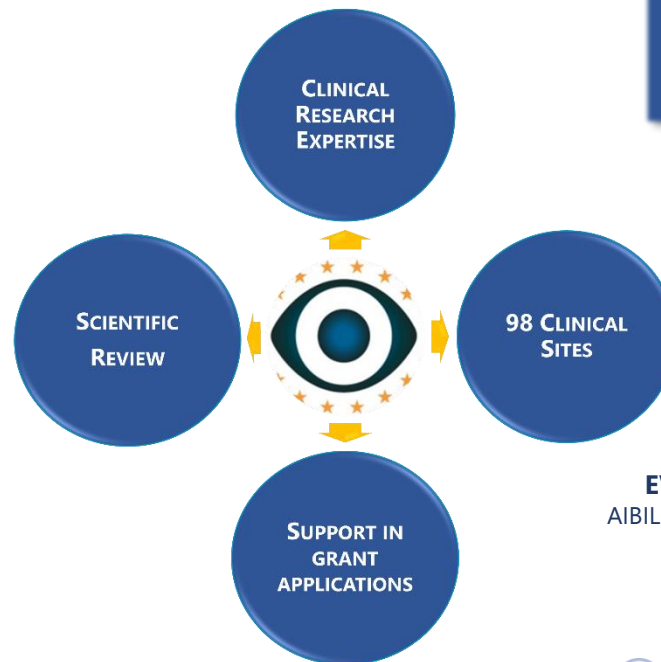


EVICR.net

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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A NETWORK OF 98 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**
Pasquale Aragona

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
(MEMBERS RESTRICTED AREA)



If you are not an EVICR.net member yet:

- Please contact the Coordinating Centre: evicrnet@aibili.pt