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
Jorge Alió
Birgit Lorenz
Tunde Peto
Cecília Martinho

Educational programme

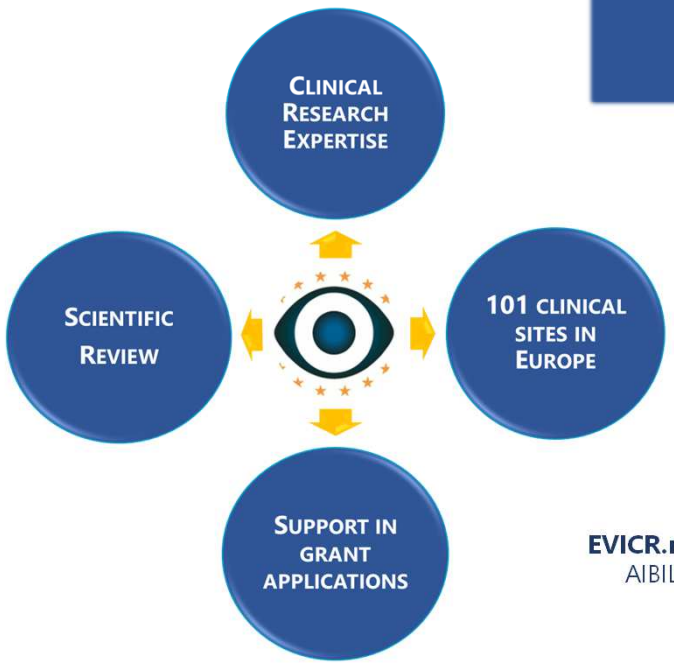
Take advantage of webinars on ophthalmology clinical research

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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EVICR.net
EUROPEAN VISION CLINICAL RESEARCH

A NETWORK OF 101 CLINICAL RESEARCH SITES FROM 15 EUROPEAN 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

- **MEDICAL DEVICES** - *Jorge Alió*
- **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