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



INITIATED RESEARCH (IIR)

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



EDUCATIONAL PROGRAMME



TAKE ADVANTAGE OF WEBINARS ON OPHTHALMOLOGY CLINICAL RESEARCH

MAY 2020



A NETWORK OF 99 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

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- **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

POST-STUDY SERVICES

- Biostatistics
- Final Study report
- Publication
- Medical Writing

How to proceed?

IN-STUDY SERVICES

Study management

IMP management

Data managementPharmacovigilance

Monitoring

If you are a member:

 Submit your idea by filling the abstract available at <u>www.evicr.net</u> (MEMBERS RESTRICTED AREA)



If you are not an EVICR.net member yet:

Please contact the Coordinating Centre: evicrnet@aibili.pt