

Neurology

Multicenter observational prospective study on thrombectomy device.

International observational multicenter studies.

Ophtalmology

Observational multicenter studies (lenses, multifocal implants).

Respiratory

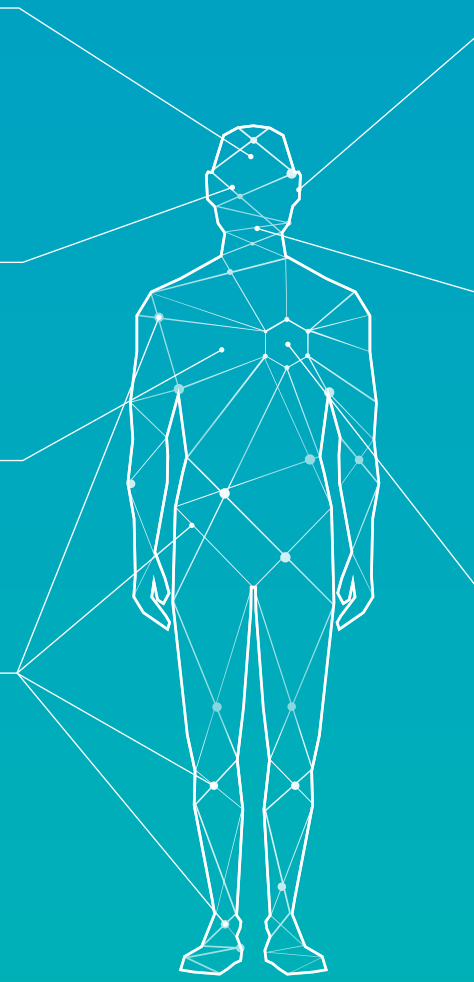
Interventional and observational prospective multicenter studies.

Orthopaedics

Interventional and observational multicenter studies (knees, hips, shoulders, ankles, feet).

Aesthetics

Interventional prospective study on reducing superficial fats.



Auditory

Observational multicenter study on audio prosthesis with bone anchor.

Dental

Observational retrospective and multicenter study on bone filter membrane.

Multicenter and retrospective study on dental implants.

Cardiology

Interventional study (first implantation in humans) pre-market study.

Prospective clinical registry on the treatment of aortic arch for 6 years.

Textile

Interventional study in the treatment of Ehlers-Danlos syndromes.



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Guarantee the validity of your clinical results

Evamed is a clinical research organization, specialized in the evaluation and clinical investigation of medical devices.

Founded in 2005, Evamed helps manufacturers, learned societies and public organisms with their interventional and observational studies and registries. With a strong experience in the medical devices field, Evamed is a trusted partner for your clinical study projects.

Our team of experts is committed to ensuring the validity and confidentiality of your clinical results since 2005. Evamed's clinical investigations range from 3 months to 20 years, and involve up to 2500 patients per study, using our eCRF in more than 25 countries : Germany, Italy, Belgium, England, Switzerland, Japan, Canada, China, Slovakia...

Evamed supports you with your clinical strategy

Support on clinical strategy

- Definition of objectives
- Needs analysis
- Co-construction of the clinical development

Regulatory Submissions

- Regulatory files preparation
- Organisms submissions (CPP, ANSM, CNIL, CERES...)
- Support in case of substantial modifications

Clinical Evaluation Report

- Methodology
- State of art
- Risks analysis
- CER writing

Data Collection tool (eCRF)

- Data Collection software developed by Evamed
- High security hosting on health data licensed servers
- Online tools to follow-up your clinical investigation
- ePRO add-on
- Patient data entry
- Direct access to the eCRF for the patients

Data-Management

- Data-Management plan writing
- Control consistencies programming
- Inconsistencies listing generation
- Listing injection in the eCRF (automatic queries generation)

Study Report

- Protocol, monitoring, investigation plan and statistical analysis integration
- Results interpretation
- Revision by medical director

Clinical Investigation Plan

- Synopsis writing
- Clinical investigation plan writing
- Sample size calculation
- Patients documents writing
- Consent form
- Information note

Study Management

- Monitoring plan writing
- Site setting (remote or on site)
- Monitoring (remote or on site)
- Investigators training
- Study coordination
- Sites closure (remote or on site)

Statistical Analysis

- Statistical analysis plan writing
- Realization of descriptive, comparative, and multivariate statistical analysis
- Statistical analysis report writing

Publication

- Medical publication writing
- Medical Director expertise

EVAMED IN A FEW FIGURES



RIPH 1 & 2
30%

RIPH 3
60%

REGISTER
10%

DISTRIBUTION OF THE NUMBER OF PROJECTS BY THERAPEUTIC AREA

