PLEASE NOTE: *This trial has been registered retrospectively.*

Trial Description

Title

Extension of the EUROBservational Research Programme Atrial Fibrillation General Registry (EORP AFIB registry) for Germany

Trial Acronym

AFNET-EORP registry Atrial Fibrillation Germany

URL of the trial

[---]*

Brief Summary in Lay Language

The research programme is a large, scientific patient registry conducted by the Competence Network AFNET association in coorperation with the German Society of Cardiology and the European Societey of Cardiology in Germany. This registry is conducted throughout Europe in a comparable manner to document the management of atrial filbrillation. The objective of the german patient registry is to collect and analyse medical data in Germany in hospitals and outpatient institutions. Information serves for implementation in guidelines to improve the management of patients with atrial fibrillation.

Brief Summary in Scientific Language

The research programme is a large, scientific patient registry conducted by the Competence Network AFNET association in coorperation with the German Society of Cardiology and the European Societey of Cardiology in Germany. This registry is conducted throughout Europe in a comparable manner to document the management of atrial filbrillation. The objective of the german patient registry is to collect and analyse medical data in Germany in hospitals and outpatient institutions. Information serves for implementation in guidelines to improve the management of patients with atrial fibrillation.

Organizational Data

- DRKS-ID: **DRKS00006219**
- Date of Registration in DRKS: **2014/07/23**
- Date of Registration in Partner Registry or other Primary Registry: **[---]***
- Investigator Sponsored/Initiated Trial (IST/IIT): **yes**
- Ethics Approval/Approval of the Ethics Committee: **Approved**
- (leading) Ethics Committee Nr.: **2013-533-f-s , Ethik-Kommission der Ärztekammer**

DRKS-ID: **DRKS00006219**

Date of Registration in DRKS: **2014/07/23**

Date of Registration in Partner Registry or other Primary Registry: **[---]***

Investigator Sponsored/Initiated Trial (IST/IIT): **yes**

Ethics Approval/Approval of the Ethics Committee: **Approved**

Westfalen-Lippe und der med. Fakultät der Westfälischen Wilhelms-Universität Münster

Secondary IDs

Health condition or Problem studied

- ICD10: **I48.9 - [generalization I48: Atrial fibrillation and flutter]**

Interventions/Observational Groups

- Arm 1: **patient registry: collection of data concerning management of treatment in patients with artrial fibrillation in in- and outpatient manner**

Characteristics

- Study Type: **Non-interventional**
- Study Type Non-Interventional: **Observational study**
- Allocation: **Single arm study**
- Blinding: **[---]***
- Who is blinded: **[---]***
- Control: **Uncontrolled/Single arm**
- Purpose: **Health care system**
- Assignment: **Single (group)**
- Phase: **N/A**
- Off-label use (Zulassungsüberschreitende Anwendung eines Arzneimittels): **N/A**

Primary Outcome

The aim of the registry is the improvement of the management of patients with

atrial fibrillation

Secondary Outcome

none

Countries of recruitment

- DE **Germany**

Locations of Recruitment

Recruitment

- Planned/Actual: **Actual**
- (Anticipated or Actual) Date of First Enrollment: **2014/05/14**
- Target Sample Size: **3500**
- Monocenter/Multicenter trial: **Multicenter trial**
- National/International: **National**

Inclusion Criteria

- Gender: **Both, male and female**
- Minimum Age: **18 Years**
- Maximum Age: **no maximum age**

Additional Inclusion Criteria

- Patients will be officially enrolled in the study only if ECG diagnosis of AF has been made.
- The qualifying episode of AF should have occurred within one year before the date of baseline.
- AF is the primary or secondary diagnosis, i.e. the current admission / visit may be due to other reasons.
- Patients need not be in AF at the time of enrolment.
- Signed Patient Inform Consent
- Age <= 18 years

Exclusion criteria

Patients who are currently or are planned to be taking part in a cardiac clinical trial.
Please note that patients can participate concomitantly in other registries.

Addresses

- **Primary Sponsor**

  Die Register-Studie wird durchgeführt vom: Kompetenznetz Vorhofflimmern AFNET e.V.
  Mendelstraße 11
  48149 Münster
  Germany

  Telephone: **0049-251-980 1343**
  Fax: **0049-215-980 1349**
  E-mail: **info at kompentenznetz-Vorhoffllimmern.de**
  URL: **www.kompetenznetz-vorhofflimmern.de**

- **Contact for Scientific Queries**

  Universitätsklinikum Großhadern Medizinische Klinik I
  Mr. Professor Michael Näbauer
  Marchioninistraße 15
  81377 München
  Germany

  Telephone: **0049-89-4400 73060**
  Fax: **[---]***
  E-mail: **michael.nabauer at med.uni-muenchen.de**
  URL: **[---]***

- **Collaborator, Other Address**

  - Deutsche Gesellschaft für Kadiologie- European Society of Cardiology

  Telephone: **[---]***
  Fax: **[---]***
  E-mail: **[---]***
  URL: **[---]***

Sources of Monetary or Material Support

- **Commercial (pharmaceutical industry, medical engineering industry, etc.)**

  Bristol Myers Squibb GmbH & Co kGaA
  Arnulfstr. 29
  80636 München
  Germany

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Telephone: **[---]***
Fax: **[---]***
E-mail: **[---]***
URL: **[---]***

- **Public funding institutions financed by tax money/Government funding body (German Research Foundation (DFG), Federal Ministry of Education and Research (BMBF), etc.)**

Deutsches Zentrum für Herz-Kreislauf-Forschung e.V.
Oudenarder Str. 16
13347 Berlin
Germany

Telephone: **[---]***
Fax: **[---]***
E-mail: **[---]***
URL: **http://dzhk.de/**

Status

- Recruitment Status: **Recruiting complete, follow-up complete**
- Study Closing (LPLV): **2018/12/31**

Trial Publications, Results and other documents

** This entry means the parameter is not applicable or has not been set.*
**** This entry means that data is not displayed due to insufficient data privacy clearing.*