PLEASE NOTE: This trial has been registered retrospectively.

Trial Description

Title

The "Beobachtungsnetz-Register"/Observation Network Registry

Trial Acronym

BeoNet-Register

URL of the trial

[---]*

Brief Summary in Lay Language

The so called "Beobachtungsnetzwerk-Register" should guarantee an interdisciplinary cooperation between scientists and physicians from outpatient care. It shall contribute to improve the care system and the quality management.

The BeoNet-Register is financially supported by the BMBF. The Deutsche Zentrum für Lungenforschung (DZL) is responsible for the implementation. The technical conditions are placed in Hannover, so Hannover is concerned with the technical developing of the BeoNet-Register.

The aim of this register is to presentet the care reality in a practical way ver a long-term periode. The long-term objective of the project is to get data in real time and in continuous monitoring in form of the electronic patients record and also through questionnaires.
As a result of these data collection it will be possible to get information regarding to aspects of monitoring and of individual courses of disease. Also systematical interventions can be illustrated and patient-centred evaluations can be established.

Brief Summary in Scientific Language

The aim of this register is to presentet the care reality in a practical way ver a long-term periode. The long-term objective of the project is to get data in real time and in continuous monitoring in form of the electronic patients record and also through questionnaires.
As a result of these data collection it will be possible to get information regarding to aspects of monitoring and of individual courses of disease. Also systematical interventions can be illustrated and patient-centred evaluations can be established.
Organizational Data

- **DRKS-ID:** DRKS00005822
- **Date of Registration in DRKS:** 2014/04/30
- **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.: 1481-2012, Ethikkommission der Medizinischen Hochschule Hannover

Secondary IDs

- Free text: All diseases

Health condition or Problem studied

- Free text: All diseases

Interventions/Observational Groups

- **Arm 1:** During and after the treatment of a patient by a doctor clinical and administrative data will be documented and stored electronically.

  Doctors will be informed about the progress and the planning process of the BeoNet-Register. If they would take part in the BeoNet-Register voluntarily, they store their documentation in the BeoNet-Register. All data will be passed in strictly pseudonymous form.

  In a second step, data will be anonymised. On request, research partners could get an extract.

  The aim is to develop a network based on general practitioners, paediatricians and specialists in pneumology. They should export their routinedata in the research registry daily. Based on disease-specific research questions there could be a possibility to define standards for treatment and care processes.

Characteristics

- **Study Type:** Non-interventional
- **Study Type Non-Interventional:** Observational study
- **Allocation:** Single arm study
- **Blinding:** [---]*
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

During and after the treatment of a patient by a doctor clinical and administrative data will be documented and stored electronically.

Doctors will be informed about the progress and the planning process of the BeoNet-Register. If they would take part in the BeoNet-Register voluntarily, they store their documentation in the BeoNET-Register. All data will be passed in strictly pseudonymous form.

In a second step, data will be anonymised. On request, research partners could get an extract.

On this basis a network related to general practioners could be implemented. The members of this network have allowed to put their routine data in a central research registry. With this data it will be possible to work an disease specific research questions, for example to create standards in public health research.

### Secondary Outcome

The available data can be processed for care-related question. Within the register data will collected first monthly, then daily.

### Countries of recruitment

- **DE Germany**

### Locations of Recruitment

- Doctor's Practice **deutschlandweit**
Recruitment

- Planned/Actual: Actual
- (Anticipated or Actual) Date of First Enrollment: 2013/02/01
- Target Sample Size: 100
- Monocenter/Multicenter trial: Multicenter trial
- National/International: National

Inclusion Criteria

- Gender: Both, male and female
- Minimum Age: 0 Years
- Maximum Age: no maximum age

Additional Inclusion Criteria

the doctors office is participating in the BeoNet-register

Exclusion criteria

the declaration of consent is canceled

Addresses

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Sources of Monetary or Material Support

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

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Fax: [---]*
E-mail: [---]*
URL: www.bmbf.de

Status

Recruitment Status: Recruiting ongoing
Study Closing (LPLV): [---]*
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.