Trial Description

Title

Evaluation of the quality of supply of patients with PAD (intermittent claudication) stage II in Germany

Trial Acronym

[---]*

URL of the trial

[---]*

Brief Summary in Lay Language

Supervised exercise programme is part of the guidelines for the treatment of intermittent claudication. This study will try to detect insufficient offer of walking training. On the basis of a survey the treatment of patients with intermittent claudication in Germany will be examined. The patients will be recruited in 10 different vascular centres in Germany.

Brief Summary in Scientific Language

Supervised exercise programme is part of the guidelines for the treatment of PAD. This study will try to detect insufficient offer of walking training. On the basis of a survey the treatment of patients with PAD in Germany will be examined. The patients will be recruited in 10 different vascular centres in Germany.

Organizational Data

- DRKS-ID: **DRKS00004964**
- Date of Registration in DRKS: **2013/05/15**
- 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.: **183/13 , Ethik-Kommission der Albert-Ludwigs-Universität Freiburg**

Secondary IDs

Health condition or Problem studied

- ICD10: **I73.9 - Peripheral vascular disease, unspecified**

Interventions/Observational Groups

- Arm 1: **Patients with PAD will be questioned once with a self-developed survey about the treatment of PAD and their estimate of success.**

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

Assessment of of guideline-based therapy of PAD on the basis of a survey.

Secondary Outcome

none

Countries of recruitment

- DE **Germany**

Locations of Recruitment

- other **Gefäßzentrum des Universitäts-Herzzentrums, Freiburg im Breisgau**

Recruitment

- Planned/Actual: **Actual**
- (Anticipated or Actual) Date of First Enrollment: **2013/08/30**
- Target Sample Size: **1000**
- Monocenter/Multicenter trial: **Monocenter trial**
- National/International: **National**

Inclusion Criteria

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

Additional Inclusion Criteria

Patients with intermittent claudication (PAD)

Exclusion criteria

- pain in the hip or knee joints
- stroke or cardiac infarction
- visual disorders

Addresses

- **Primary Sponsor**

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- **Contact for Scientific Queries**

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Sources of Monetary or Material Support

- **Institutional budget, no external funding (budget of sponsor/PI)**

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Status

- Recruitment Status: **Recruiting complete, follow-up complete**
- Study Closing (LPLV): **2014/10/08**

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.*