

Trial Description

Title

Adjunctive use of essential oils following Scaling and root planing

Trial Acronym

essential oils

URL of the trial

http://nicht vorhanden

Brief Summary in Lay Language

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Brief Summary in Scientific Language

The study deals with scaling and root planing in periodontitis patients. In a randomized double-blind study we want to check if there is a benefit of the adjunctive use of essential oils during and after scaling and root planing for the improvement of clinical variables as well as at periodontopathogenic bacteria.

Organizational Data

- DRKS-ID: **DRKS00009387**
- Date of Registration in DRKS: **2015/09/30**
- Date of Registration in Partner Registry or other Primary Registry: **2012/09/24**
- Investigator Sponsored/Initiated Trial (IST/IIT): **yes**
- Ethics Approval/Approval of the Ethics Committee: **Approved**
- (leading) Ethics Committee Nr.: **332-12-24092012 , Ethikkommission an der Medizinischen Fakultät der Universität Leipzig**

Secondary IDs

Health condition or Problem studied

- ICD10: **K05.3 - Chronic periodontitis**

Interventions/Observational Groups

- Arm 1: **scaling and root planing with essential oils**
- Arm 2: **scaling and root planing without essential oils**

Characteristics

- Study Type: **Interventional**
- Study Type Non-Interventional: **[---]***
- Allocation: **Randomized controlled trial**
- Blinding: **[---]***
- Who is blinded: **investigator/therapist, assessor, data analyst**
- Control: **Active control (effective treatment of control group)**
- Purpose: **Treatment**
- Assignment: **Parallel**
- Phase: **N/A**
- Off-label use (Zulassungsüberschreitende Anwendung eines Arzneimittels): **N/A**

Primary Outcome

probing depth measured with a manual periodontometer at baseline as well as after three and six months

Secondary Outcome

attachment level, bleeding on probing, periodontopathogenic bacteria

Countries of recruitment

- DE **Germany**

Locations of Recruitment

- Doctor's Practice **Hamburg**

Recruitment

- Planned/Actual: **Actual**

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- (Anticipated or Actual) Date of First Enrollment: **2013/04/02**
- Target Sample Size: **50**
- Monocenter/Multicenter trial: **Monocenter trial**
- National/International: **National**

Inclusion Criteria

- Gender: **Both, male and female**
- Minimum Age: **40 Years**
- Maximum Age: **65 Years**

Additional Inclusion Criteria

generalized moderate periodontitis, no antibiotic use within the 1st six months

Exclusion criteria

Pregnancy, breastfeeding and allergy to the ingredients of the herbal distillate products

Addresses

- **Primary Sponsor**

**Institut für Mikroökologie und SymbioVaccin
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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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■ **Commercial (pharmaceutical industry, medical engineering industry, etc.)**

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Status

- Recruitment Status: **Recruiting complete, follow-up complete**
- Study Closing (LPLV): **2014/09/28**

Trial Publications, Results and other documents

** This entry means the parameter is not applicable or has not been set.*