Trial Description

Title

Analysis of the immune-status of patients with lymphoma

Trial Acronym

Immulymph

URL of the trial

[---]*

Brief Summary in Lay Language

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Brief Summary in Scientific Language

The project aims at investigating the immune status of patients with lymphoma in the peripheral blood, on tumor samples or apharesis products.

Organizational Data

- DRKS-ID: **DRKS00010064**
- Date of Registration in DRKS: **2016/02/16**
- 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.: **15-320 , Ethik-Kommission der Medizinischen Fakultät der Universität zu Köln**

Secondary IDs

Health condition or Problem studied

- Free text: **Lymphoma**
- Free text: **Probandes without lymphoma**

Interventions/Observational Groups

- Arm 1: **The project aims at investigating the immune status of patients with lymphoma in the peripheral blood, on tumor samples or apharesis products.**

Characteristics

- Study Type: **Non-interventional**
- Study Type Non-Interventional: **Observational study**
- Allocation: **Other**
- Blinding: **[---]***
- Who is blinded: **[---]***
- Control: **Other**
- Purpose: **Basic research/physiological study**
- Assignment: **Other**
- Phase: **N/A**
- Off-label use (Zulassungsüberschreitende Anwendung eines Arzneimittels): **N/A**

Primary Outcome

Basic research project:
-Sterility of samples
- FACS analysis of cellular composition

Secondary Outcome

Basic research project:
- Ability for expansion of T cells
- Ability for transduction with chimeric antigene of T T cells

Countries of recruitment

- DE **Germany**

Locations of Recruitment

- University Medical Center **Köln**

Recruitment

- Planned/Actual: **Planned**

Planned/Actual: **Planned**

- (Anticipated or Actual) Date of First Enrollment: **2016/02/16**
- Target Sample Size: **200**
- 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

1. Patients with Lymphoma
2. Probands without lymphoma
3. Age of consent

Exclusion criteria

Portends who are not able to consent.

Addresses

- **Primary Sponsor**

Uniklinik Köln
Mr. Prof. Dr. Michael Hallek
Kerpenerstraße 62
50924 Köln
Germany

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

- **Contact for Scientific Queries**

Uniklinik Köln
Ms. Dr. Natali Pflug
Kerpenerstraße 62
50924 Köln
Germany

Contact for Scientific Queries

Uniklinik Köln
Ms. Dr. Natali Pflug
Kerpenerstraße 62
50924 Köln
Germany

Telephone: **0221-478-0**
Fax: **[---]***
E-mail: **Natali.Pflug at uk-koeln.de**
URL: **[---]***

- **Contact for Public Queries**

Uniklinik Köln
Ms. Dr. Natali Pflug
Kerpenerstraße 62
50924 Köln
Germany

Telephone: **0221-478-0**
Fax: **[---]***
E-mail: **Natali.Pflug at uk-koeln.de**
URL: **[---]***

Sources of Monetary or Material Support

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

Uniklinik Köln
50924 Köln
Germany

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

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.*