# Influence of pre-operative vaccination on immune response and post-operative infection rate

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>		
11/08/2006	No longer recruiting	☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
29/01/2007	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
16/05/2016	Cancer			

## Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

**Prof Claudia Spies** 

#### Contact details

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# Additional identifiers

Clinical Trials Information System (CTIS)

2004-004352-40

#### Protocol serial number

N/A

# Study information

#### Scientific Title

Influence of pre-operative vaccination on immune response and post-operative infection rate

#### **Study objectives**

Pre-operative vaccination with influenza antigen improves post-operative immune suppression.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethics board committee Berlin, Landesamt für Gesundheit und Soziales (LaGeSo), Berlin, 11/03/2005, ref: EA 1/048/04

#### Study design

Prospective randomised double-blind placebo-controlled clinical trial

#### Primary study design

Interventional

#### Study type(s)

Treatment

#### Health condition(s) or problem(s) studied

Upper aerodigestive tract cancer undergoing surgery

#### **Interventions**

Vaccination with influenza haemagglutinin antigen versus a placebo control group.

# Intervention Type

Biological/Vaccine

# Primary outcome(s)

Influence of pre-operative vaccination on immune response.

# Key secondary outcome(s))

Influence of pre-operative vaccination post-operative infection rate.

# Completion date

01/01/2009

# **Eligibility**

# Key inclusion criteria

Tumour resection of upper aerodigestive tract.

# Participant type(s)

Patient

## Healthy volunteers allowed

No

#### Age group

Adult

#### Sex

All

#### Key exclusion criteria

Initial information at time of registration:

- 1. No consent
- 2. Under 18 years
- 3. Liver insufficiency (more than Child B)
- 4. Acute Coronary Syndrome (less than 8 weeks)
- 5. Human Immunodeficiency Virus (HIV) positive
- 6. Corticosteriodal treatment
- 7. After organ transplantation
- 8. Current infection
- 9. Pregnancy

#### Added as of 30/03/2009:

- 10. Diabetes mellitus
- 11. Status post splenectomy
- 12. Protein deficiency
- 13. Known allergy to components of the vaccine
- 14. Immunosuppression

#### Date of first enrolment

01/11/2005

#### Date of final enrolment

01/01/2009

# Locations

#### Countries of recruitment

Germany

## Study participating centre Charite Universitaetsmedizin Berlin

Berlin Germany 13353

# Sponsor information

#### Organisation

Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

#### **ROR**

https://ror.org/001w7jn25

# Funder(s)

#### Funder type

University/education

#### **Funder Name**

Charité Universitätsmedizin Berlin

## Alternative Name(s)

Medical School - Charité - University Medicine Berlin

# **Funding Body Type**

Private sector organisation

#### **Funding Body Subtype**

For-profit companies (industry)

#### Location

Germany

# **Results and Publications**

Individual participant data (IPD) sharing plan

# IPD sharing plan summary

Not provided at time of registration

# Study outputs

Output type	Details	Date created Date added	Peer reviewed?	Patient-facing?
Results article	results	01/08/2015	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes