The role of the microbiota in the systemic immune response

Published: 05-11-2012 Last updated: 15-05-2024

To investigate the role of the gut microbiota in the systemic priming of immune effector cells

Ethical review Approved WMO

Status Recruitment stopped

Health condition type Hepatobiliary neoplasms malignant and unspecified

Study type Interventional

Summary

ID

NL-OMON36859

Source

ToetsingOnline

Brief title

MISSION-1

Condition

Hepatobiliary neoplasms malignant and unspecified

Synonym

sepsis

Research involving

Human

Sponsors and support

Primary sponsor: Academisch Medisch Centrum

Source(s) of monetary or material Support: ZonMw Klinische Fellowship beurs

Intervention

Keyword: antibiotics, gut microbiota, immune response

Outcome measures

Primary outcome

Side effects, laboratory parameters for inflammatory responses, functional assays (ex vivo inflammatory responses) and gut microbiota composition

Secondary outcome

Not applicable

Study description

Background summary

Sepsis ranks among the top ten leading causes of death worldwide. Most nonsurvivors die in a state of immunosuppression. The gut microbiota exerts numerous beneficial functions in the host response against infections. Gut flora components express microorganism-associated molecular patterns (MAMPs) such as lipopolysaccharide (LPS), which are recognized by pattern recognition receptors (PRRs) expressed by neutrophils and macrophages. MAMPs from the intestinal microbiota constitutively translocate to the circulation and prime bone marrow neutrophils via PRRs. Antibiotic treatment, which is standard of care for all patients with sepsis, depletes the gut microbiota and leads to a diminished release of MAMPs. This may attribute to sepsis associated immunosuppression. Our ultimate aim is to develop new therapeutic strategies to restore immunity, such as faeces transplantation or administration of selective components of the microbiota.

Study objective

To investigate the role of the gut microbiota in the systemic priming of immune effector cells

Study design

Within-subject-controlled intervention study in human volunteers, n=12

Intervention

Subjects self-administer antibiotics for seven days:

- ciprofloxacin 500mg 2dd1 (Gram negatives and positives)

- vancomycin 250mg 3dd2 (Gram positives)
- metronidazol 500mg 3dd1 (anaerobes) Blood and feces will be collected before taking antibiotics and 24 hours and 6 weeks after taking antibiotics.

Study burden and risks

Volunteers may experience side effects when using antibiotics, mostly gastrointestinal complaints like diarrhoea. Subjects known with any allergic reaction to any previously administered antibiotic will be excluded from participation, but novel allergic reactions are possible. The burden also includes 3 visits - after the initial screening for eligibility visit * to draw blood and collect faeces (subject himself at home). Subjects are not allowed to smoke, drink (until 48 hours after the last dose of antibiotics) or travel to tropical countries.

Contacts

Public

Academisch Medisch Centrum

Meibergdreef 9 Amsterdam 1105 AZ NI

Scientific

Academisch Medisch Centrum

Meibergdreef 9 Amsterdam 1105 AZ NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

18-35 years of age male healthy (medical history, physical exam, no medications, laboratory screening) non- smoking normal defecation pattern (<3x/ day, >3x/week)

Exclusion criteria

allergic to antibiotics (any kind) recent use of antibiotics (<12 maanden) difficulty swallowing pills

Study design

Design

Study type: Interventional

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Basic science

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 13-03-2012

Enrollment: 12

Type: Actual

Ethics review

Approved WMO

Date: 05-11-2012

Application type: First submission

Review commission: METC Amsterdam UMC

Approved WMO

Date: 15-02-2013

Application type: Amendment

Review commission: METC Amsterdam UMC

Approved WMO

Date: 03-12-2015

Application type: Amendment

Review commission: METC Amsterdam UMC

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

ID: 27808 Source: NTR

Title:

In other registers

Register ID

CCMO NL42072.018.12 OMON NL-OMON27808