# the role of the goodpasture antigen binding protein in autoimmunity

Published: 29-12-2010 Last updated: 15-05-2024

Primary Objective: characterization of GPBP expression on different subsets of human leukocytesSecondary Objective: functional characterization of GPBP on these cells

**Ethical review** Approved WMO **Status** Recruiting

**Health condition type** Autoimmune disorders **Study type** Observational invasive

## **Summary**

### ID

NL-OMON34337

Source

ToetsingOnline

**Brief title** 

GPBP in autoimmunity

#### **Condition**

Autoimmune disorders

**Synonym** 

autoimmunity

Research involving

Human

### **Sponsors and support**

**Primary sponsor:** Medisch Universitair Ziekenhuis Maastricht

Source(s) of monetary or material Support: Ham Foundation; Prof. Dr. M. De Baets

#### Intervention

**Keyword:** autoimmunity, goodpasture antigen binding protein, inflammation, leukocytes

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#### **Outcome measures**

### **Primary outcome**

Characterization of human leukocyte subsets that express GPBP.

Differences in GPBP expression in different inflammatory conditions (autoimmune

vs. healthy)

### **Secondary outcome**

Determination of the function of GPBP on human leukocytes.

# **Study description**

### **Background summary**

The Goodpasture antigen-binding protein (GPBP) is a serine/threonine kinase that binds and phosphorylates collagen type IV, which has a major role in the organisation of the glomerular basement membrane and is the target of autoantibodies mediating glomerulonephritis in Goodpasture syndrome. Several studies have suggested a link between GPBP and the immune system. GPBP has been shown to be upregulated in autoimmune conditions, to transport the signalling lipid ceramide, to be regulated by immune signalling molecules and to interact with pentraxins, the key activators of the complement system. However, the exact role of GPBP in regulation of the (auto)immune response still remains unclear. Therefore, we intend to investigate the exact role of GPBP in the activation and modulation of the immune system.

### Study objective

Primary Objective: characterization of GPBP expression on different subsets of

human leukocytes

Secondary Objective: functional characterization of GPBP on these cells

### Study design

10 ml of venous blood will be obtained from patients with an autoimmune disease or another inflammatory disease. Leukocytes will be isolated from the blood to study GPBP expression and function on these cells. For comparison, it is necessary to study normal, healthy blood cells as well. This will be obtained

from healthy controls.

### Study burden and risks

Patients will be asked to give blood during a regular outpatient clinic visit, so it will demand no extra time investment from the patient. For control subject, the most important burden will be time investment. A local bruise is possible on the site of puncture.

### **Contacts**

#### **Public**

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# **Trial sites**

### **Listed location countries**

**Netherlands** 

# **Eligibility criteria**

#### Age

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

### Inclusion criteria

Autoimmune patients: 18 - 50 years old, diagnosed with myasthenia gravis or multiple

sclerose

Healthy controls: 18 - 50 years old

### **Exclusion criteria**

no informed consent minors or incapacitated

healthy controls: previous inflammation related disease

# Study design

### **Design**

Study type: Observational invasive

Intervention model: Other

Allocation: Non-randomized controlled trial

Masking: Open (masking not used)

Control: Active

Primary purpose: Basic science

### Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 07-02-2011

Enrollment: 100

Type: Actual

### **Ethics review**

Approved WMO

Date: 29-12-2010

Application type: First submission

Review commission: METC academisch ziekenhuis Maastricht/Universiteit

Maastricht, METC azM/UM (Maastricht)

# **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: 27227

Source: Nationaal Trial Register

Title:

### In other registers

Register	ID
CCMO	NL33086.068.10
Other	nog niet gekend
OMON	NL-OMON27227