Tumor-educated platelets in myositis patients: screening for malignancies

Published: 18-10-2021 Last updated: 05-04-2024

We aim to investigate in a pilot study the blood platelet mRNA profiles in patients with IIM

and a known malignancy.

Ethical review Approved WMO

Status Pending

Health condition type Miscellaneous and site unspecified neoplasms malignant and

unspecified

Study type Observational invasive

Summary

ID

NL-OMON50252

Source

ToetsingOnline

Brief title

TEPs in myositis patients: malignancy screening

Condition

- Miscellaneous and site unspecified neoplasms malignant and unspecified
- Neuromuscular disorders

Synonym

cancer, Myositis

Research involving

Human

Sponsors and support

Primary sponsor: Academisch Medisch Centrum

Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: Malignancies, Myositis, TEPs, Tumor educated platelets

Outcome measures

Primary outcome

blood platelet mRNA profile

Secondary outcome

na

Study description

Background summary

The screening for malignancies in patients with myositis may be improved and less invasive by novel techniques.

Study objective

We aim to investigate in a pilot study the blood platelet mRNA profiles in patients with IIM and a known malignancy.

Study design

Pilot study - cross sectional cohort study in ten patients

Study burden and risks

The overall risk of this study is considered low and relate to two extra tubes during a single episode of blood sampling which is part of clinical care. Potential benefits include an improvement of cancer screening methods for patients with myositis.

Contacts

Public

Academisch Medisch Centrum

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

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

Inclusion criteria

- 1. Patients with dermatomyositis (DM) and immune mediated necrotizing myopathy (IMNM) with or without a myositis specific antibody, with a known malignancy (out of a list of 16 malignancies).
- 2. Informed consent.

Exclusion criteria

No informed consent Women who are pregnant or breastfeeding

Study design

Design

Study type: Observational invasive

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Diagnostic

Recruitment

NL

Recruitment status: Pending

Start date (anticipated): 26-09-2021

Enrollment: 10

Type: Anticipated

Ethics review

Approved WMO

Date: 18-10-2021

Application type: First submission

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

No registrations found.

In other registers

Register ID

CCMO NL78849.018.21