Pilotstudy early detection Angina Pectoris with Telecardiology and Biomarkers used by General Practitioners in The Netherlands

Published: 07-05-2007 Last updated: 30-11-2024

In this pilot study we investigate the use of these extra diagnostic tools. We also will use the results to see if these tools can diagnose cardiac events in patients that would otherwise be sent home

Ethical review Approved WMO Status Completed

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

Summary

ID

NL-OMON31379

Source

ToetsingOnline

Brief title

Telecardiology and Biomarkers for the GP in the Netherlands

Condition

Myocardial disorders

Synonym

Angina Pectoris

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Groningen

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Source(s) of monetary or material Support: IPT cardiodetect

Intervention

Keyword: Biomarkers, General Practitioner, Netherlands, Telecardiology

Outcome measures

Primary outcome

Extra necessary submissions to the hospital because of the extra diagnostic

tools.

Secondary outcome

Prevention events

Study description

Background summary

People above 55 years of age often have ECG changes, sometimes because of events, that were earlier not known.

Possibly we can detect these events by supplying the General Practitioner with extra diagnostic tools, the ECG and bio-assays (bio-markers).

Study objective

In this pilot study we investigate the use of these extra diagnostic tools. We also will use the results to see if these tools can diagnose cardiac events in patients that would otherwise be sent home

Study design

Patients that come to the GP with complaints possibly compatible with angina will be seen by the GP in the office. The diagnosis and other possible causes will be noted as well as the further strategy. (care as usual) Afterwards the patient will be offered the extra diagnostic tools: a tele-ECG, judged by a cardiologist, and a blood-sample to check the bio-markers.

Study burden and risks

No risks, 3 minutes to make the ECG, 1 blood sample

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Contacts

Public

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Scientific

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

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

Inclusion criteria

Angina Pectoris

Exclusion criteria

Insanity

Study design

Design

Study type: Observational invasive

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Diagnostic

Recruitment

NL

Recruitment status: Completed

Start date (anticipated): 01-04-2007

Enrollment: 50

Type: Actual

Ethics review

Approved WMO

Date: 07-05-2007

Application type: First submission

Review commission: RTPO, Regionale Toetsingscie Patientgebonden Onderzoek

(Leeuwarden)

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 CCMO

ID

NL16725.099.07