

The prevalence and significance of chronic obstructive pulmonary disease in subjects with acute coronary syndrome

Submission date 20/03/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 04/05/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 14/02/2017	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Chronic obstructive pulmonary disease (COPD) is a group of lung conditions that cause breathing difficulties. COPD shares a common risk factor, tobacco, with heart disease, which is currently the leading cause of death globally. Studies have found that patients with COPD have an increased risk of heart disease. However, these studies have several limitations. Firstly, the diagnosis of COPD has been based purely on medical records. Secondly, patients with asthma have sometimes also been included. As about 80% of patients with COPD are undiagnosed and 20-50% of patients with COPD do not fulfil the diagnostic criteria, the prevalence of COPD in heart disease has not been reliably assessed. Spirometry is a simple test that can be used to diagnose COPD by measuring how much air patients can breathe out in one forced breath. The aims of this study are to estimate the true prevalence of COPD in heart disease using spirometry, and to find out whether spirometry-verified COPD predicts heart disease prognosis.

Who can participate?

Patients hospitalised with heart disease (myocardial infarction or unstable angina), living in the county of Jämtland, Sweden

What does the study involve?

Participants' lung volumes are measured (spirometry) during hospitalisation or at the first follow-up outpatient visit after discharge. The prevalence of spirometry-detected COPD is compared with the prevalence of COPD based on medical records.

What are the possible benefits and risks of participating?

If COPD is detected, participants receive disease information and, if needed, are offered treatment. The test requires forced expirations (breathing out). Some may experience slight physical discomfort during the test.

Where is the study run from?

Östersund Hospital (Sweden)

When is the study starting and how long is it expected to run for?
January 2010 to December 2014

Who is funding the study?

1. Jämtland County Council (Sweden)
2. The Swedish Heart and Lung Association (Sweden)

Who is the main contact?

Dr Nikolai Stenfors
nikolai.stenfors@medicin.umu.se

Contact information

Type(s)

Scientific

Contact name

Dr Nikolai Stenfors

Contact details

Jamtland County Council
Research and Development Unit
Building 10, 5th floor
Östersund Hospital
Östersund
Sweden
83183
+46 63 153417
nikolai.stenfors@medicin.umu.se

Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

The prevalence and significance of Chronic Obstructive Pulmonary disease in subjects with Acute Coronary Syndrome: an observational study

Acronym

COPACS

Study objectives

1. Spirometry screening after acute coronary syndrom (ACS) will reveal that the majority of subjects with chronic obstructive pulmonary disease (COPD) are undiagnosed.
2. Spirometry-verified COPD is an independent predictor of death, reinfarction and stroke in subjects with Acute Coronary Syndrome (ACS).

Ethics approval required

Old ethics approval format

Ethics approval(s)

Regional Ethical Review Board, Umeå University, Umeå, Sweden, 16/12/2009, ref: Dnr 09-142M

Study design

Single-centre observational study

Primary study design

Observational

Study type(s)

Screening

Health condition(s) or problem(s) studied

Chronic obstructive pulmonary disease, acute coronary syndrome

Interventions

The study subjects will be screened with spirometry to detect presence of chronic airflow limitation. Lung volume measurements (spirometry) will be done during the hospitalisation or at the first follow-up out-patient visit after discharge. Vital capacity (VC), forced vital capacity (FVC) and forced expiratory volume in one second (FEV1) will be measured before and after bronchodilation. FEV1/FVC <0.7 post bronchodilation will be considered as COPD.

The prevalence of spirometry-detected COPD will be compared with the prevalence of COPD based on medical records in the study population. We will evaluate whether spirometry-verified COPD predicts prognosis after ACS. Inclusion between 01 January 2010 and 30 June 2012, follow-up until 31 December 2014.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

1. The prevalence of spirometry-verified COPD versus medical-record based COPD in subjects with ACS
2. The adjusted hazard ratio of spirometry-verified COPD for death, reinfarction and stroke in subjects with ACS. Adjustment for age, gender, renal function, C-reactive protein, Killip class, previous myocardial infarction (MI), atrial fibrillation, coronary intervention, secondary preventive medication, diabetes, and smoking history.

Key secondary outcome(s)

In subjects with ACS, spirometry-verified COPD is an independent predictor of hospital admissions (due to coronary angiography, percutaneous coronary intervention, coronary artery bypass surgery, angina, heart failure) and length of hospital stay.

Completion date

31/12/2014

Eligibility

Key inclusion criteria

1. All patients living in the county of Jämtland, Sweden, and hospitalised with a diagnosis of myocardial infarction or unstable angina will be assessed for inclusion
2. Östersund hospital is the only hospital in the county and all patients, terminal care excluded, with symptoms of a suspected acute coronary syndrome are referred for diagnostic evaluation
3. Study subjects are recruited in conjunction with the study "Secondary preventive, nurse based, telephone follow-up for risk factor control after an acute coronary syndrome" (<http://www.isrctn.com/ISRCTN96595458/>)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Dementia
2. Patients with severe disease
3. Subjects unable to perform adequate spirometry

Date of first enrolment

01/01/2010

Date of final enrolment

31/12/2014

Locations

Countries of recruitment

Sweden

Study participating centre

Jamtland County Council

Östersund

Sweden

83183

Sponsor information

Organisation

Jämtland County Council (Sweden)

Funder(s)

Funder type

Government

Funder Name

Jämtland County Council (Sweden)

Funder Name

Hjärt-Lungfonden

Alternative Name(s)

Swedish Heart-Lung Foundation

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Sweden

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