# Is a four minute wait required between transfer test measurements?

Submission date 28/09/2007	<b>Recruitment status</b> No longer recruiting	<ul> <li>Prospectively registered</li> <li>Protocol</li> </ul>
<b>Registration date</b> 28/09/2007	<b>Overall study status</b> Completed	<ul> <li>Statistical analysis plan</li> <li>Results</li> </ul>
Last Edited 09/08/2021	<b>Condition category</b> Respiratory	<ul> <li>Individual participant data</li> <li>Record updated in last year</li> </ul>

#### Plain English summary of protocol

Not provided at time of registration

# **Contact information**

**Type(s)** Scientific

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# Additional identifiers

EudraCT/CTIS number

**IRAS number** 

ClinicalTrials.gov number

Secondary identifying numbers N0226184632

# Study information

**Scientific Title** Is a four minute wait required between transfer test measurements?

#### **Study objectives**

Is a four minute wait required between transfer test measurements?
 Does airflow obstruction influence the results?

**Ethics approval required** Old ethics approval format

**Ethics approval(s)** Not provided at time of registration

**Study design** Randomised controlled trial

**Primary study design** Interventional

**Secondary study design** Randomised controlled trial

Study setting(s) Hospital

**Study type(s)** Treatment

#### Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

#### Health condition(s) or problem(s) studied

Respiratory: Function tests

#### Interventions

Prospective review of pulmonary function tests for patients referred to the pulmonary function laboratory. Participants will be randomised into the order of performing the test. This will either be a one minute wait followed by a four minute wait, or a four minute wait followed by a one minute wait.

**Intervention Type** Other

**Phase** Not Specified

Primary outcome measure

Whether the timing of subsequent TLco measurements influences the transfer factor measurement.

#### Secondary outcome measures

Whether the timing of subsequent TLco measurements influences the alveolar volume measurement.

#### Overall study start date

01/09/2006

#### **Completion date**

01/05/2007

# Eligibility

#### Key inclusion criteria

25 subjects from each of the following diagnostic groups:a. Healthy volunteers (no lung disease)b. Asthmatics (variable obstructive lung disease)c. COPD (obstructive lung disease)d. Fibrosis (restrictive lung disease)

Inclusion criteria:

- 1. Over 18 years of age
- 2. Subject must be able to understand and perform lung function test to national guidelines
- 3. Non or ex-smokers (>6 months)
- 4. Physician diagnosis of asthma, COPD or lung fibrosis

#### Participant type(s)

Healthy volunteer

Age group

Adult

**Lower age limit** 18 Years

**Sex** Not Specified

Target number of participants

25

#### Key exclusion criteria

- 1. Haemoptysis of unknown origin
- 2. Pneumothorax
- 3. Unstable cardiovascular status (recent MI or PE)
- 4. Thoracic, abdominal or cerebral aneurysms
- 5. Recent eye surgery
- 6. Nausea and vomiting

7. Recent thoracic or abdominal surgical procedures

8. Involved in other studies with licensed drugs or methodology studies within last weeks

Date of first enrolment 01/09/2006

Date of final enrolment 01/05/2007

## Locations

**Countries of recruitment** England

United Kingdom

**Study participating centre University Hospital of South Manchester NHS Foundation Trust** Manchester United Kingdom M23 9LT

## Sponsor information

**Organisation** Record Provided by the NHSTCT Register - 2007 Update - Department of Health

**Sponsor details** The Department of Health, Richmond House, 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type Government

Website http://www.dh.gov.uk/Home/fs/en

# Funder(s)

**Funder type** Government

**Funder Name** University Hospital of South Manchester NHS Foundation Trust (UK)

Funder Name NHS R&D Support Funding

# **Results and Publications**

#### **Publication and dissemination plan** Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

**IPD sharing plan summary** Not provided at time of registration