

# Effect of Occupational Therapy in patients with Chronic Obstructive Pulmonary Disease: a randomised controlled trial

<b>Submission date</b> 28/06/2007	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 27/09/2007	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 24/01/2020	<b>Condition category</b> Respiratory	<input type="checkbox"/> Individual participant data

## Plain English Summary

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Mrs Unni Martinsen

### Contact details

Diakonhjemmet Sykehus

Klinisk Aktivitetsavdeling

Pb 23, Vinderen

Oslo

Norway

0319

+47 22 45 15 00

unni.martinsen@diakonsyk.no

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

## Study information

### Scientific Title

Effect of Occupational Therapy in patients with Chronic Obstructive Pulmonary Disease: a randomised controlled trial

### Acronym

OT in COPD

### Study hypothesis

Chronic Obstructive Pulmonary Disease (COPD) has functional consequences in terms of activity limitations and participation restrictions. Although there are studies evaluating the effect of multidisciplinary rehabilitation in patients with COPD, there is a lack of high quality studies examining the effect of occupational therapy on enhancing activity and participation. The aim of this study is to evaluate the effect of occupational therapy for patients with COPD compared with a group not receiving such treatment.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Obtained in November 2005 from the regional Ethical Committee. Approval from Norwegian Social Science Data Service was obtained in May 2006.

### Study design

Randomised controlled trial

### Primary study design

Interventional

### Secondary study design

Randomised controlled trial

### Study setting(s)

Hospital

### Study type(s)

Quality of life

### Participant information sheet

### Condition

Chronic obstructive pulmonary disease

### Interventions

The control group receive treatment as usual; in this case no occupational therapy from the hospital. In the study the control group will be offered OT after the last measure at 12 months.

The intervention group will be offered two to three consultations of approximately 1 - 2 hours duration, depending on the kind and severity of functional limitations. The interventional group receive individual occupational therapy, including information and practical training in energy conserving methods, breathing techniques and use of assistive technology.

### **Intervention Type**

Procedure/Surgery

### **Primary outcome measure**

Activity and participation, measured by The Canadian Occupational Performance Measure (COPM) and Assessment of Motor and Process Skills (AMPS).

Outcomes will be measured at baseline, and after 4 and 12 months.

### **Secondary outcome measures**

1. Health-related quality of life, measured by St. George's Respiratory Questionnaire (SGRQ)
2. Lung function examined by spirometry and pulse oximetry

Outcomes will be measured at baseline, and after 4 and 12 months.

### **Overall study start date**

01/05/2007

### **Overall study end date**

30/06/2009

## **Eligibility**

### **Participant inclusion criteria**

Amendments as of 02/12/2008: please note that point two of the inclusion criteria below has been updated as follows:

2. Aged 20 - 80 years

Initial information at time of registration:

1. Patients with moderate to severe COPD (Global Initiative on Obstructive Lung Disease [GOLD])
2. Aged 20 - 75 years
3. Limited ability to perform daily activities
4. Ability to communicate in Norwegian

### **Participant type(s)**

Patient

### **Age group**

Adult

### **Sex**

Both

### **Target number of participants**

50

**Total final enrolment**

52

**Participant exclusion criteria**

1. Exacerbations for last three weeks
2. Cognitive or mental impairment
3. Co-morbidity with an impact on the ability to perform daily activity

**Recruitment start date**

01/05/2007

**Recruitment end date**

30/06/2009

**Locations****Countries of recruitment**

Norway

**Study participating centre**

Diakonhjemmet Sykehus

Oslo

Norway

0319

**Sponsor information****Organisation**

The Norwegian Foundation for Health and Rehabilitation (Helse og Rehabilitering) (Norway)

**Sponsor details**

Karl Johans gt 23 B

Oslo

Norway

0159

-

adm@helseogrehab.no

**Sponsor type**

Research organisation

**Website**

<http://www.helseogrehab.no/>

# Funder(s)

## Funder type

Research organisation

## Funder Name

The Norwegian Foundation for Health and Rehabilitation (Helse og Rehabilitering) (Norway) - <http://www.helseogrehab.no>

## Funder Name

The Federation of Norwegian Commercial and Service Enterprises (HSH) (Norway) - <http://www.hsh-org.no>

## Funder Name

The Norwegian Association of Occupational Therapists (NETF) (Norway) - <http://www.netf.no>

# 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

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/03/2017	24/01/2020	Yes	No