# Nocturnal nasal postive pressure ventilation plus oxygen therapy versus oxygen alone in severe stable chronic obstructive pulmonary disease

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
10/10/2002	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
10/10/2002	Completed	Results
Last Edited	Condition category	[] Individual participant data
01/07/2009	Respiratory	<ul><li>Record updated in last year</li></ul>

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Dr DSC Hui

#### Contact details

Department of Medicine & Therapeutics Prince of Wales Hospital Chinese University of Hong Kong

Hong Kong

# -

# Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

# Study information

#### Scientific Title

#### Study objectives

Not provided at time of registration

#### 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)

Not specified

#### Study type(s)

**Not Specified** 

#### Participant information sheet

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

Respiratory diseases

#### **Interventions**

Patients will then be randomized into two groups receiving:

- 1. Non-invasive positive pressure ventilation (NPPV) plus LTOT, or
- 2. LTOT plus oral placebo over a 3-month period.

NPPV will be delivered via a portable bilevel positive airway pressure (BiPAP) Duet device via a nasal or oral-nasal mask.

#### Intervention Type

Drug

#### Phase

**Not Specified** 

# Drug/device/biological/vaccine name(s)

oxygen

#### Primary outcome measure

Not provided at time of registration

#### Secondary outcome measures

Not provided at time of registration

#### Overall study start date

01/01/2002

#### Completion date

31/12/2002

# **Eligibility**

#### Key inclusion criteria

Patients with severe stable chronic obstructive pulmonary disease with hypercapnic respiratory failure already on domiciliary long-term oxygen therapy (LTOT)

#### Participant type(s)

**Patient** 

#### Age group

**Not Specified** 

#### Sex

**Not Specified** 

# Target number of participants

Not provided at time of registration

#### Key exclusion criteria

Does not meet inclusion criteria

#### Date of first enrolment

01/01/2002

#### Date of final enrolment

31/12/2002

# Locations

#### Countries of recruitment

China

Hong Kong

# Study participating centre Department of Medicine & Therapeutics

-

Hong Kong

-

# Sponsor information

#### Organisation

Hong Kong Health Services Research Fund (Hong Kong)

#### Sponsor details

Health Welfare and Food Bureau Government Secretariat, HKSAR 20th floor Murray Building Garden Road

Hong Kong

-

+852 (0)2973 8288 hsrf@hwfb.gov.hk

#### Sponsor type

Government

#### Website

http://www.fhb.gov.hk/grants/english/funds/funds\_hhsrf/funds\_hhsrf\_abt/funds\_hhsrf\_abt.html

#### **ROR**

https://ror.org/03qh32912

# Funder(s)

# Funder type

Government

#### **Funder Name**

Hong Kong Health Services Research Fund (Hong Kong)

# **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