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

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

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

921019

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