

A randomised study to compare the effectiveness of hyoscine hydrobromide (Hyoscine) and glycopyrronium (Robinul) in the management of noisy breathing/retained secretions in cancer patients

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 04/01/2016	Condition category Cancer	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N0258107515

Study information

Scientific Title

A randomised study to compare the effectiveness of hyoscine hydrobromide (Hyoscine) and glycopyrronium (Robinul) in the management of noisy breathing/retained secretions in cancer patients

Study objectives

1. To compare the effectiveness of both hyoscine hydrobromide and glycopyrronium at relieving retained secretions at the time of death.
2. To test feasibility of the advanced 'consent process'.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Intentional

Study type(s)

Other

Health condition(s) or problem(s) studied

Cancer: Noisy breathing, retained secretions

Interventions

Randomised test intervention versus standardised intervention, non-blinded (Phase 3).

Intervention Type

Drug

Phase

Phase III

Drug/device/biological/vaccine name(s)

Hyoscine hydrobromide, glycopyrronium

Primary outcome(s)

1. Provide an evidence base for common practice
2. Provide a means of obtaining consent for research in dying patients

Key secondary outcome(s)

Not provided at time of registration

Completion date

28/02/2005

Eligibility

Key inclusion criteria

Total number of Royal Marsden Hospital patients 205

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

28/02/2002

Date of final enrolment

28/02/2005

Locations

Countries of recruitment

United Kingdom

Australia

Study participating centre

Mater Adult Hospital

South Brisbane

Australia

Qld 4101

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Charity

Funder Name

The Royal Marsden NHS Foundation Trust (UK) Charitable Funds

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration