

Effects of nurse-delivered psychological support on reducing anxiety and satisfying the needs of families with relatives in intensive care: a randomised controlled trial

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/09/2014	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0201119628

Study information

Scientific Title

Study objectives

The effectiveness of nurse-led intervention to provide psychological support to relatives and patients in intensive care.

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

Health condition(s) or problem(s) studied

Mental and Behavioural Disorders: Anxiety

Interventions

1. Control group will receive the usual psychological support provided on AICU
2. Intervention group will require additional psychological support tailored to the individual's needs.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/03/2003

Completion date

01/04/2004

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

272 relatives and patients over 12 months and 400 for the duration of the study

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

30/03/2003

Date of final enrolment

01/04/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Anaesthetic & Critical Care

London

United Kingdom

SW3 6NP

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

Royal Brompton and Harefield NHS Trust (UK)

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