

Intensive care unit outcome study

Submission date 14/03/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 02/05/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 30/03/2017	Condition category Other	<input type="checkbox"/> Individual participant data

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
Version 7.0

Study information

Scientific Title
Intensive care unit outcome study

Acronym

ICON (Intensive Care Outcome Network)

Study objectives

The primary purpose of the study is to generate a register of patients discharged from intensive care units (ICUs). The registry will be used to study ICU treatment and its relation to mortality and long-term psychological morbidity, as well as changes in health-related quality of life over time after discharge from ICU.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Oxford Research Ethics Committee B, 12/04/2006, ref: 06/Q1605/17

Study design

Prospective longitudinal questionnaire-based study

Primary study design

Observational

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Patients admitted to intensive care units

Interventions

Questionnaires regarding health-related quality of life (HRQOL).

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

To start and maintain a register of patients discharged from hospital after being treated for greater than 24 hours in a participating ICU.

Key secondary outcome(s)

To evaluate in survivors of intensive care treatment:

1. Health-related quality of life
2. Risk of post-traumatic stress disorder (PTSD)
3. Major depression and anxiety incidence
4. Quality-adjusted life-years (QALYs)

Completion date

28/03/2013

Eligibility

Key inclusion criteria

1. All patients that survive to discharge from a participating ICU and remained on the ICU for greater than 24 hours
2. Patients aged 16 years or older on date of ICU admission

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Patients in non-participating ICUs
2. Patients admitted to a participating ICU who stayed less than 24 hours
3. Patients that do not survive to ICU discharge
4. Patients who are under 16 years old on date of ICU admission
5. Patients readmitted to an ICU and already entered in the ICON study will not have their new admission recorded

Date of first enrolment

01/03/2006

Date of final enrolment

28/03/2013

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

John Radcliffe Hospital

Oxford

United Kingdom

OX3 9DU

Sponsor information

Organisation

Oxford Radcliffe Hospitals NHS Trust (UK)

ROR

<https://ror.org/03h2bh287>

Funder(s)**Funder type**

Research organisation

Funder Name

Intensive Care Society

Alternative Name(s)

The Intensive Care Society, ICS

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary****Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	27/03/2017		Yes	No
Protocol article	protocol	17/06/2008		Yes	No