Assessment of Collaborative Requesting

Submission date	Recruitment status No longer recruiting	Prospectively registered		
22/03/2007		Protocol		
Registration date	Overall study status	Statistical analysis plan		
16/07/2007	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
12/10/2009	Other			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Duncan Young

Contact details

c/o Miss Sarah Edwards Kadoorie Centre for Critical Care Research and Education Level 3 John Radcliffe Hospital Oxford United Kingdom OX3 9DU

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Protocol v5.2

Study information

Scientific Title

Acronym

ACRE

Study objectives

Relatives of patients in whom death has been determined using brain stem death criteria are more likely to consent to organ donation when interviewed by the clinical team if a donor transplant co-ordinator is present, compared with relatives interviewed by the clinical team alone.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved by Oxfordshire Research Ethics Committee A (Ref: 06/Q1604/119)

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Relatives of brain stem dead patients; Organ donation - collaborative requesting

Interventions

The intervention is collaborative requesting which is the addition of a donor transplant coordinator to the team requesting organ donation. The routine arm is the relatives interviewed by the clinical team alone.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Proportion of relatives refusing consent for organ donation.

Secondary outcome measures

- 1. Number of patients in whom the family consent to beating heart organ donation
- 2. Solid organs retrieved by type
- 3. Solid organs transplanted by type
- 4. Tissues retrieved by type
- 5. Consent rates by predefined subgroup (relative relationship to donor; ethnic group and sex of donor and next of kin; clinician seniority and sex; sex of donor transplant co-ordinator)

Overall study start date

01/04/2007

Completion date

01/08/2009

Eligibility

Key inclusion criteria

Relatives of patients in whom death has been determined using brain stem death criteria

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

946

Key exclusion criteria

- 1. Relatives who do not understand written or verbal infromation for whom an interpreter is not available
- 2. Relatives who decline discussion on organ donation

Date of first enrolment

01/04/2007

Date of final enrolment

01/08/2009

Locations

Countries of recruitment

England

United Kingdom

Study participating centre c/o Miss Sarah Edwards Oxford United Kingdom OX3 9DU

Sponsor information

Organisation

Oxford Radcliffe Hospitals NHS Trust (UK)

Sponsor details

Research and Develpment Office Manor House John Radcliffe Hospital Oxford England United Kingdom OX3 9DU

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/03h2bh287

Funder(s)

Funder type

Government

Funder Name

UK Transplant

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	08/10/2009		Yes	No