

Cardiopulmonary resuscitation (CPR) quality improvement programme

Submission date 30/11/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 05/07/2011	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 04/02/2016	Condition category Circulatory System	<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
PB-PG-1207-14246

Study information

Scientific Title
Cardiopulmonary resuscitation (CPR) quality improvement programme: a multicentre, prospective, cohort study

Study objectives

Implementing a CPR quality improvement initiative will improve the rate of return of spontaneous circulation and other outcomes in patients sustaining cardiac arrest

Ethics approval required

Old ethics approval format

Ethics approval(s)

Coventry Research Ethics Committee, 26/06/2009, REC ref: 09/H1210/65

Study design

Multicentre prospective cohort study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

In-hospital cardiac arrests

Interventions

1. Control
2. CPR feedback through Q-CPR® (Laerdal medical, UK)
3. CPR feedback through Q-CPR® (Laerdal medical, UK) plus team briefing / debriefing

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Survived event [defined as sustained (> 20 minutes) return of spontaneous circulation]

Key secondary outcome(s)

1. Patient outcomes
 - 1.1. Any return of spontaneous circulation
 - 1.2. Survival to hospital discharge
 - 1.3. Cerebral performance category of patients at discharge
2. Process outcomes
 - 2.1. Quality of emergency calls
 - 2.2. Chest compression depth, rate
 - 2.3. No flow time
 - 2.4. Duration of pre and post-shock pauses
 - 2.5. Ventilation rate
 - 2.6. time to first shock
 - 2.7. Appropriate decision to shock

- 3. Team factors
- 3.1. CPR knowledge
- 3.2. Confidence / preparedness

Completion date

31/01/2013

Eligibility

Key inclusion criteria

Patients have to be either 18 or older and suffer an in-hospital cardiac arrest where resuscitation is attempted (defined as loss of a pulse requiring the delivery of chest compressions).

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

1. Patient has a Do-not-attempt resuscitation order written and documented in their medical records
2. Cardiac arrest not attended by a resuscitation team
3. Out - of - hospital cardiac arrest
4. Previous participation in this study

Date of first enrolment

01/11/2009

Date of final enrolment

31/01/2013

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Warwick Medical School Clinical Trials Unit
Coventry
United Kingdom
CV4 7AL

Sponsor information

Organisation
Heart of England NHS Foundation Trust (UK)

Funder(s)

Funder type
Government

Funder Name
National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) programme (PB-PG-1207-14246)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2015		Yes	No
Protocol article	protocol	18/10/2011		Yes	No