

Section 06

Cardiac arrest data

Context

6.1

Time of arrest to nearest hour (24-hour clock) (Time Span)

6.2

Perioperative phase (Pick One)

- Pre-induction
- At induction
- Transfer to theatre/procedure location
- After induction but before surgery/procedure started
- During surgery/procedure under GA
- During surgery/procedure under LA/RA
- Conversion of LA/RA to GA
- Emergence/extubation
- Post-operative transfer to recovery
- Post-operative in recovery room (or other immediate location)
- Post-operative after discharge from recovery room or equivalent (or in critical care area if recovery not used)
- N/A: special inclusion criteria

6.3

Location at time of cardiac arrest (Pick One)

- Anaesthetic room
- Theatre: main theatre suite
- Theatre: day surgery unit
- Theatre: obstetrics
- Theatre: other
- Labour ward
- Neuroradiology
- Cardiac catheter lab
- Pacing room
- Interventional radiology
- MRI
- CT
- Endoscopy
- ECT
- Ward
- Recovery
- Emergency Department
- Critical care area
- Other

If other:

6.4

Anaesthetic presence

At time of cardiac arrest Consultant

At any time during resuscitation Consultant

At time of cardiac arrest SAS doctor

At any time during resuscitation SAS doctor

At time of cardiac arrest Post CCT or CESR doctor

At any time during resuscitation Post CCT or CESR doctor

At time of cardiac arrest ST5+ or equivalent

At any time during resuscitation ST5+ or equivalent

At time of cardiac arrest ST3-4 or equivalent

At any time during resuscitation ST3-4 or equivalent

At time of cardiac arrest CT2 or equivalent

At any time during resuscitation CT2 or equivalent

At time of cardiac arrest CT1 or equivalent after Initial Assessment of Competence

At any time during resuscitation CT1 or equivalent after Initial Assessment of Competence

At time of cardiac arrest CT1 or equivalent before completion of Initial Assessment of Competence

At any time during resuscitation CT1 or equivalent before completion of Initial Assessment of Competence

At time of cardiac arrest Anaesthesia Associate

At any time during resuscitation Anaesthesia Associate

At time of cardiac arrest Nurse specialist

At any time during resuscitation Nurse specialist

At time of cardiac arrest Other

At any time during resuscitation Other

6.5

Was the cardiac arrest witnessed ? (Pick One)

Yes

No

6.6

Was a 2222 call activated? (Pick One)

Yes

No

If yes by whom? (Pick One)

Anaesthetist or anaesthesia associate

Anaesthetic assistant / ODP

Nurse

Midwife

- Surgeon
- Healthcare assistant / runner
- Other

6.7

What forms of monitoring were already in place when need for chest compressions and/or defibrillation was first recognised? (Pick One)

- Same monitoring as previously detailed for case
- Other
- None

Pulse oximetry
(True / False)

Non-invasive blood pressure
(True / False)

ECG
(True / False)

End tidal CO₂ / Capnography
(True / False)

End tidal anaesthetic agents
(True / False)

FiO₂
(True / False)

Airway pressure
(True / False)

Intra-arterial blood pressure
(True / False)

Central venous pressure
(True / False)

Cardiac output
(True / False)

Transthoracic echocardiography
(True / False)

Transoesophageal echocardiography
(True / False)

Continuous temperature measurement
(True / False)

Intermittent temperature measurement
(True / False)

Non-quantitative neuromuscular monitoring
(True / False)

Quantitative neuromuscular monitoring
(True / False)

Raw or processed EEG
(True / False)

Near-infrared Spectroscopy
(True / False)

Arterial blood gas
(True / False)

Point of care coagulation
(True / False)

Antecedents

6.8

Was there a change in patient position immediately before cardiac arrest recognised? (Pick One)

Yes
 No

What was the change?

6.9

Was there a change in location/transfer immediately before cardiac arrest recognised? (Pick One)

Yes
 No

What was the change?

Pre-arrest data

6.10

Observations most proximate to starting chest compressions and/or defibrillation (Time 0) and at previous timepoints as available.

We appreciate that data may not be available at each timepoint. Please only enter available values.

Please Fill in: **Section PreArreset**
Multiple copies may be included.

6.11

How were these data collected

Recall
(True / False)

Anaesthetic chart - manual
(True / False)

Anaesthetic chart - electronic
(True / False)

Anaesthetic machine
(True / False)

Not applicable - unavailable
(True / False)

6.12
Were any of the following already in place?
Vasopressors (continuous infusion)
(True / False)

Inotropes (continuous infusion)
(True / False)

Mechanical ventilation
(True / False)

Non-invasive ventilation (including HFNO)
(True / False)

VV ECMO
(True / False)

None
(True / False)

6.13

At the time of cardiac arrest, was the patient supported by any form of Ventricular Assist Device (VAD)? (Pick One)
 Yes
 No

6.14

At the time of cardiac arrest, did the patient have a pre-existing internal or external cardioverter defibrillator? (Pick One)
 Yes
 No

6.15
At the time of cardiac arrest, were any of the following in place?
Patient docked to surgical robot
(True / False)

Patient head in pins
(True / False)

Specialist table/mattress (e.g. Montreal)
(True / False)

In scan/procedure room remote from anaesthetist (e.g. in CT scanner, MRI)
(True / False)

Other
(True / False)

None
(True / False)

Was there a significant delay in undocking patient? (Pick One)
 Yes
 No
 Unknown

Pins removed
(True / False)

Clamp released
(True / False)

Head attachment removed and held (by hand)
(True / False)

Head placed on head-ring/theatre table
(True / False)

Other
(True / False)

Cardiac Arrest Process

6.16

Which initial patient condition best describes the event? (Pick One)

- Pulseless
- Pulse but poor perfusion
- Systolic blood pressure <50 mmHg - non-invasive
- Systolic blood pressure <50 mmHg - invasive
- Unknown

6.17

What triggered the start of cardiopulmonary resuscitation (chest compressions and/or defibrillation)?

Impalpable pulse
(True / False)

Unrecordable blood pressure
(True / False)

Severe hypotension
(True / False)

Severe bradycardia
(True / False)

Monitored cardiac rhythm
(True / False)

Reduction in ETCO2
(True / False)

Other
(True / False)

Please specify

6.18

Did the patient receive chest compressions (includes open cardiac massage)? (Pick One)

- Yes - ≥ 5
- Yes - <5
- No

If patient prone at the time of cardiac arrest, were compressions started in the prone position? (Pick One)

- Yes
- No
- Unknown
- N/A - not prone at time of cardiac arrest

6.19

What was the initial cardiac arrest rhythm? (Pick One)

- Ventricular fibrillation (VF)
- Pulseless ventricular tachycardia (pVT)
- Pulseless electrical activity (PEA)
- Asystole
- Bradycardia
- AED used - shockable
- AED used - non-shockable
- Unknown

6.20

Did the patient receive a precordial thump? (Pick One)

- Yes - successful at achieving ROSC at next rhythm check
- Yes - unsuccessful
- No
- Unknown

6.21

Was an automated external defibrillator (AED) or manual defibrillator in AED/Shock Advisory mode applied? (Pick One)

- Yes
- No
- Unknown

6.22

Did the patient receive defibrillation for ventricular fibrillation (VF) or pulseless ventricular tachycardia? (Pick One)

- Yes
- No
- Unknown

What was the total number of defibrillatory shocks delivered? (Pick One)

- 1
- 2
- 3
- 4
- >4
- Unknown

6.23

Was Extracorporeal Cardiopulmonary Resuscitation (ECPR) with Venoarterial extracorporeal membrane oxygenation (VA-ECMO) attempted during cardiac arrest? (Pick One)

- Yes
- No
- Unknown

6.24

Was a mechanical chest compression device used? (Pick One)

- Yes
- No
- Unknown

6.25

Was adrenaline given by intravenous or intraosseous route during the resuscitation event?

Yes - initial 1mg bolus (or 10mcg/kg paediatric dose)
(True / False)

Yes - initial titrated IV aliquots (e.g. 10-50mcg intermittent boluses)
(True / False)

Yes - infusion
(True / False)

No
(True / False)

Unknown
(True / False)

Time between cardiac arrest and first dose

No of doses

Total dose

At time of cardiac arrest Unit (Pick One)

- mcg
- mg
- ml

6.26

Other drugs

Please Fill in: **Section CAPDrugs**

Multiple copies may be included.

6.27

Airway interventions

Airway interventions In Situ (Pick One)

- None
- Oxygen mask or nasal specs
- Face mask (+/- Guedel)
- Supraglottic airway (1st generation)
- Supraglottic airway (2nd generation)
- Tracheal tube (oral or nasal)
- Tracheostomy
- Emergency front of neck airway
- High flow nasal O2/THRIVE
- Rigid bronchoscope
- Other

Please specify

None
(True / False)

Oxygen mask or nasal specs
(True / False)

Face mask (+/- Guedel)
(True / False)

Supraglottic airway (1st generation)
(True / False)

Supraglottic airway (2nd generation)
(True / False)

Tracheal tube (oral or nasal)
(True / False)

Tracheostomy
(True / False)

Emergency front of neck airway
(True / False)

High flow nasal O2/THRIVE
(True / False)

Rigid bronchoscope
(True / False)

Other
(True / False)

Please specify

6.28

Method(s) of confirmation used to ensure correct placement of tracheal tube or tracheostomy tube
Waveform capnography
(True / False)

Capnometry (numeric ETCO2)
(True / False)

Exhaled CO2 colorimetric monitor
(True / False)

Oesophageal detection device
(True / False)

Ultrasound
(True / False)

Revisualisation with direct laryngoscopy
(True / False)

Flexible optical bronchoscope
(True / False)

None of the above
(True / False)

Unknown
(True / False)

N/A - no tracheal tube used
(True / False)

6.29

Were any of the following mechanisms or processes in place during the resuscitation to measure the quality of CPR being delivered?
Waveform capnography (ETCO2)
(True / False)

Arterial waveform
(True / False)

Diastolic pressure
(True / False)

CPR mechanics device (e.g. accelerometer, force transducer, Triaxial Field Induction device)
(True / False)

CPR quality coach
(True / False)

Metronome
(True / False)

Other
(True / False)

Unknown
(True / False)

None
(True / False)

Please specify

6.30

Were any of the following additional resuscitative procedures undertaken?

Cardiac Pacing
(True / False)

Cardio-pulmonary bypass
(True / False)

Chest drain (any)
(True / False)

DC Cardioversion (unplanned)
(True / False)

Embolectomy
(True / False)

Front of neck airway
(True / False)

Hyperkalaemia management
(True / False)

Intra-arterial balloon pump (IABP)
(True / False)

Needle decompression of chest
(True / False)

Precordial thump
(True / False)

Resternotomy
(True / False)

Thoracostomy
(True / False)

Thoracotomy
(True / False)

Thrombolysis
(True / False)

Transfusion of blood products
(True / False)

Other
(True / False)

None
(True / False)

Please specify

6.31

Was echocardiography used during resuscitation (Pick One)

Yes - transthoracic

Yes - transoesophageal

No

Unknown

ALS/APLS cardiac arrest protocol
(True / False)

Association of Anaesthetists (AoA) Quick Reference Handbook
(True / False)

Resuscitation Council UK (RCUK) anaphylaxis guideline
(True / False)

Association of Anaesthetists anaphylaxis guideline
(True / False)

RCUK cardiac arrest during neurosurgery
(True / False)

Association of Anaesthetists local anaesthetic toxicity
(True / False)

Association of Anaesthetists malignant hyperthermia guideline
(True / False)

Cardiac advanced life support (CALs)
(True / False)

Other
(True / False)

Unknown
(True / False)

No
(True / False)

Smartphone
(True / False)

Laminate
(True / False)

In treatment pack
(True / False)

Printed copy in theatre
(True / False)

Computer/tablet
(True / False)

Memory
(True / False)

Other
(True / False)

6.32

What was the most likely cause of cardiac arrest

6.33

What was the time interval from onset of presenting clinical feature to the start of chest compressions and/or defibrillation? (Pick One)

- 0-1 mins
- 2-3 mins
- 4-5 mins
- 6-10 mins
- 11-15 mins
- 16-30 mins
- 31-60 mins
- 61-120 mins
- >120 mins

6.34

Was there a delay in the treatment of cardiac arrest? (Pick One)

- Yes
- No

Slow to diagnose
(True / False)

Appropriate assistance not available
(True / False)

Drugs not available
(True / False)

Equipment not available
(True / False)

No or limited PPE available
(True / False)

Requirement to change patient position to start CPR
(True / False)

Other
(True / False)

Please specify

6.35

Was additional anaesthetic assistance summoned? (Pick One)

Yes

No

Shout for help
(True / False)

Emergency bell
(True / False)

Phone 2222
(True / False)

Phone for assistance - on site staff (not 2222)
(True / False)

Send a runner
(True / False)

Bleep for assistance- on site
(True / False)

Phone for assistance - off site staff
(True / False)

Unknown
(True / False)

Other
(True / False)

Please specify

How long until assistance arrived (Pick One)

0-1 mins

2-3 mins

4-5 mins

6-10 mins

11-15 mins

16-30 mins

31-60 mins

61-120 mins

>120 mins

6.36

Did theatre team contribute effectively to the management of the incident? (Pick One)

- Yes, all
- Yes, some
- No
- Unknown

6.37

Were there multiple cardiac arrests? (Pick One)

- Yes
- No

Please describe

6.38

Were there any issues related to Personal Protective Equipment (PPE)? (Pick One)

- Yes
- No
- Unknown

No or limited PPE available
(True / False)

Delay in starting anaesthetic care
(True / False)

Delay in starting CPR
(True / False)

Resuscitation hindered by PPE after starting CPR: technical aspects (e.g., unable to perform effective chest compressions)
(True / False)

Resuscitation hindered by PPE after starting CPR: non-technical aspects (e.g., unable to communicate effectively with team members)
(True / False)

Other/additional comments
(True / False)

Please specify

Post-resuscitation data

6.39

Was coronary angiography undertaken? (Pick One)

- Yes - with ongoing CPR
- Yes - urgent (within 2 hours)

- Yes - delayed (during same hospital admission)
 No

6.40

Was coronary reperfusion attempted? (Pick One)

- Yes - PCI
 Yes - Thrombolysis
 Yes - Coronary artery bypass grafts
 No

Please specify timing (Pick One)

- Intra-arrest
 Within 24 hours of ROSC
 >24 hours but pre-discharge
 Unknown

6.41

Was treatment for massive pulmonary embolism attempted? (Pick One)

- Yes - embolectomy
 Yes - thrombolysis
 No

Please specify timing (Pick One)

- Intra-arrest
 Within 24 hours of ROSC
 >24 hours but pre-discharge
 Unknown

6.42

Where was the patient first transferred after resuscitation? (Pick One)

- Operating Theatre
 Recovery Room
 ICU/HDU
 Ward
 Other

Please specify

6.43

Time between cardiac arrest and patient transfer to HDU/ICU?

(True / False)

6.44

Unplanned post-operative admission to high-dependency area? (Pick One)

- Yes
 No

If Yes (Pick One)

- Coronary Care Unit
 HDU
 ICU
 Other

Please specify

6.45

What was the additional unplanned length of stay in days?

Level 2 days (Pick One)

- 0
- <1
- 1-3
- 3-5
- 5-7
- 7-10
- 10-14
- 14-21
- 21-28
- >28
- N/A

Is the level 2 stay ongoing? (Pick One)

- Yes
- No

Level 3 days (Pick One)

- 0
- <1
- 1-3
- 3-5
- 5-7
- 7-10
- 10-14
- 14-21
- 21-28
- >28
- N/A

Is the level 3 stay ongoing? (Pick One)

- Yes
- No

Hospital stay (Pick One)

- 0
- <1
- 1-3
- 3-5
- 5-7
- 7-10
- 10-14
- 14-21
- 21-28
- >28
- N/A

Is the hospital stay ongoing? (Pick One)

- Yes
- No

6.46

Was transfer to a different hospital required for critical care? (Pick One)

- Yes
- No

If Yes: (Pick One)

- No HDU/ICU in hospital where event occurred
- No HDU/ICU capacity
- Higher level of care required than could be provided in current hospital

6.47

Was the patient transferred to a specialist hospital (e.g. providing 24/7 percutaneous coronary intervention, targeted temperature management, post-arrest haemodynamic support) for further treatment? (Pick One)

- Yes
- No

6.48

Was the procedure significantly modified, abandoned or postponed as a result of the cardiac arrest? (Pick One)

- Yes, abandoned before procedure started
- Yes, abandoned after procedure started
- Yes, procedure modified
- Yes, additional unplanned return to theatre
- No

6.49

Was the theatre list or anaesthetic on-call shift terminated early? (Pick One)

- Yes
- No
- Unknown

6.50

Did any members of the team stand down from clinical activity immediately after the event? (Pick One)

- Yes
- No

Took a short break (e.g. <1 hour)
(True / False)

Took a sustained break* (e.g. >1 hour)
(True / False)

Theatre list terminated early
(True / False)

Anaesthetic on-call shift terminated early
(True / False)

Other
(True / False)

Please specify

6.51

Was there direct communication with the patient's relatives / NOK following the event? (Pick One)

- Yes
- No
- Unknown

Consultant anaesthetist
(True / False)

Trainee anaesthetist
(True / False)

SAS anaesthetist
(True / False)

Consultant surgeon
(True / False)

Trainee surgeon
(True / False)

SAS surgeon
(True / False)

Consultant from ICU
(True / False)

ICU Trainee
(True / False)

SAS from ICU
(True / False)

Anaesthesia associate
(True / False)

Nursing staff
(True / False)

Physician
(True / False)

Other
(True / False)

Unknown
(True / False)