

# Core Topics in Cardiothoracic Critical Care

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## Second Edition

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## Foreword

I am very pleased to be able to provide a brief introduction to the owner, borrower or reader of this text. This book is an update of the successful 2008 *Core Topics in Cardiothoracic Critical Care* text. When that book was published, it was the first to provide a detailed insight into the cardiothoracic critical care unit and was widely read and appreciated. Since then other authors have produced texts that explore this fascinating area of practice, but none have quite replicated that originality and quality... until now!

Cardiac critical care evolved quite separately from general intensive care. It essentially originated as a side room on the cardiac surgical ward in the 1950s where the patient who struggled after cardiac surgery was ventilated and cared for by the cardiac anaesthetist and surgeon. Today we have large multidisciplinary teams in large technology dominated purpose-built tertiary units. This has been a rapid and hugely successful evolution. Cardiothoracic critical care is now a full blooded and highly influential subspecialty in the ever expanding critical care field. Indeed I firmly believe that where cardiac intensivists tread today, general intensivists will follow tomorrow. This evolution has been accompanied by a vast expansion in research and regulation. No branch of medicine is so scrutinised and yet so open to new thinking and new solutions. The link between cardiothoracic anaesthesia and cardiothoracic critical care is vital in the joined up care of these complex patients, as is the close link with all the related specialties such as the surgeon, the cardiologist, the echocardiographer and so many more.

We are fortunate that the new generation of critical care doctors and authors from Papworth have stepped

up and, combined with a very eminent US academic, revisited, reorganised and rewritten the problems and solutions in this area of practice. Kamen Valchanov and Nicola Jones have taken over the authorship from their mentors at the world leading Papworth Hospital and have produced a book that retains the vision and wisdom of the original and added the significant advances in knowledge, technology and practice. A significant positive change is the addition of Professor Charles Hogue of Johns-Hopkins, Baltimore and Northwestern University, Chicago for a North American perspective. Knowing them all, it is not in the least surprising that they have produced a book of such scope and such high quality. The contributing authors are all experts in their fields and are drawn from a wide international base.

This book will prove invaluable to the critical care nurse, the trainee anaesthetist, surgeon and intensivist. It will also be of value to the new and established consultants who are involved with patients with cardiothoracic disease, which extends well beyond the bounds of surgery now. I feel proud to have been invited to write this foreword and I am proud to fully recommend this work.

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## Preface to the Second Edition

Why the second edition of *Core Topics in Cardiothoracic Critical Care*? The first edition of *Core Topics in Cardiothoracic Critical Care* was published in 2008. It has been a great success, providing a comprehensive text for the specialty and selling so many paper copies that Cambridge University Press had to reprint the book to meet the demands of the market. The first editors Dr Alain Vuylsteke, Dr Andrew Klein, and Mr Sam Nashef laid the foundation stone. However, a lot has happened in the world of medicine since 2008, not least in cardiothoracic critical care. Indeed practice has expanded so much that cardiothoracic critical care has been recognised as a separate sub-specialty by the Faculty of Intensive Care Medicine in the UK. Therefore, the current editors were tasked with providing an updated version of this textbook, which will hopefully offer to the reader state-of-the-art information on the current practice in cardiothoracic critical care.

### A Few Notes from the Editors

Different sources point to different events as the birth of our specialty of intensive care medicine. Most revolve around mechanical ventilation with some believing intensive care started in Boston in 1912 when a girl suffering from poliomyelitis received mechanical ventilation. Others feel that it is the organised care for polio victims in need of invasive ventilation that laid the foundations of the specialty. It is probably a little easier to define the birth of cardiothoracic critical care medicine as this was born when cardiac surgeons needed to leave patients who had undergone heroic operations in a place where they could recover. Similarly to general intensive care medicine we do not have a specific disease to treat, rather we have very sick patients with complex disorders of the cardiorespiratory system to care for.

How do we practise in this specialty? We provide organ support to patients who have undergone cardiothoracic surgery or who have failing cardiac or respiratory function, with the hope that they will respond to treatment and survive. However, these days with modern advances in life support technology, such as extracorporeal membrane oxygenation, death is no longer a binary phenomenon. As guardians of this technology we must be ever mindful of our patients' quality of life and the long-term outcome from our interventions. Importantly we must guard against sustaining life at all costs and offer patients and their loved ones, care which makes them happy, or at least acts in their best interests.

In 2018 a vast amount of evidence exists to guide this practice. However, it can be challenging to apply evidence from trials to the heterogeneous group of patients we treat in Cardiothoracic Critical Care each with unique, rapidly changing derangements of cardiorespiratory function. The world of evidence-based medicine is also riddled with problems of spurious evidence, and an ever-increasing number of articles describing scientific trials are being retracted by the publishers. In the end among a myriad of scientific and less scientific articles, guidelines and protocols, based on expert opinion, the patient has to be supported through their critical illness and recovery after surgery. In most cases good doctors, nurses and allied healthcare professionals use patient tailored approaches in their daily work to provide patients with the best possible care. We hope that the following text will offer ample and unbiased information to help us work in the best interest of each individual patient.

*Kamen Valchanov*  
*Nicola Jones*  
*Charles W Hogue*

# Link between Cardiothoracic Anaesthesia and Intensive Care: Which Patients are Admitted to Critical Care?

Andrew Klein

## Introduction

Admission to an intensive care area is undertaken for the diagnosis, management and monitoring of patients with conditions that require close or constant attention by a group of specially trained health professionals. Critical care encompasses all areas that provide level 2 and/or level 3 care as defined by the Intensive Care Society document 'Levels of Critical Care for Adult Patients, 2009' (Table 1). All level 2 and level 3 areas have higher staffing levels, specialist monitoring and more advanced treatment options available. Level 2 areas are commonly referred to as High Dependency Units (HDUs), while level 3 areas are Intensive Care Units (ICUs), and we will make this distinction in our text. In some hospitals, the two are separated geographically, whilst in others they co-exist in one area.

It is extremely common for patients undergoing cardiothoracic interventions under anaesthesia to be admitted to an ICU or HDU afterwards and this can often be a preplanned decision based on the potential for the patient to become more critically unwell or unstable. However, given the current pressures placed on the health service, in terms of both bed occupancy and finances, each individual case should be considered and a decision made as to whether such an admission will be necessary. These decisions can often be very difficult and must take into consideration a number of factors.

## Patient Related Factors

A patient's comorbidities, physiological reserve, prognosis and wishes should all be taken into account when planning their most appropriate postoperative destination. Prioritisation of patients for critical care beds should highlight only those patients likely to gain from an increased level of care and thus not those that are either too well or too sick to benefit.

It is clear that for some high-risk patients, such as those with known chronic organ failure undergoing

cardiac surgery, admission to an ICU will be mandatory after anaesthesia. It is reasonable to expect their condition to worsen following a period of cardiopulmonary bypass, and preparations should be made for any necessary organ support, for example use of inotropes or haemofiltration.

Consideration must also be taken as to whether the patient is appropriate for long-term management on an ICU. An example of this might be a palliative thoracic oncology patient undergoing a procedure for symptom relief; such a patient might be more appropriately placed in an HDU with a limit on the medical interventions that would be appropriate. This management plan should be discussed and formulated with the patient and relatives prior to the procedure itself.

## Diagnostic and Surgical Related Factors

A diagnostic model can be utilised in order to provide guidelines for admission, which identifies specific conditions and diseases where it is felt a higher level of care is always warranted. With respect to cardiothoracic intensive care, the majority of such conditions will fall under the umbrellas of the cardiac and/or respiratory systems. However, it is also possible for a patient to require admission on the basis of an additional diagnosis, such as sepsis or a neurological complication of surgery.

All patients undergoing sternotomy will mandate admission to either an ICU or cardiac recovery environment after their procedure. The differentiation between the two is discussed below. A number of cardiothoracic surgical procedures will always warrant ICU admission, due to the complex nature of the intervention and often long procedural times. Examples of these are repair of aortic dissection, or multiple valve procedures.

The majority of patients undergoing thoracic surgery will either be admitted to an HDU or discharged back to the ward following a period of close

### Which Patients are Admitted to Critical Care?

**Table 1** Levels of critical care

| Level of care             | Criteria for admission  | Examples   |
|---------------------------|---|--|
| 0<br>General ward         | <ul style="list-style-type: none"> <li>Requires hospitalisation but needs can be met through normal ward care</li> </ul>  | Intravenous drug administration<br>Observations needed less than 4 hourly  |
| 1<br>Coronary care unit   | <ul style="list-style-type: none"> <li>Recently discharged from higher level care</li> <li>In need of additional monitoring/intervention, clinical input or advice</li> <li>Requiring critical care outreach service support</li> </ul>   | Minimal 4 hourly observations<br>Continuous oxygen therapy, management of epidural, chest drain in situ<br>Risk of clinical deterioration, high early warning score  |
| 2<br>High dependency unit | <ul style="list-style-type: none"> <li>Requiring preoperative optimisation</li> <li>Requiring extended postoperative care</li> <li>Stepping down to level 2 from level 3 care</li> <li>Requiring single organ support</li> <li>Requiring basic respiratory plus basic cardiovascular support</li> </ul> | Invasive monitoring to optimise fluid balance<br>Major elective surgery, emergency surgery in unstable patient<br>Minimal hourly observations<br>Non-invasive ventilation, single intravenous vasoactive drug<br>Continuous oxygen therapy and intra-aortic balloon pump |
| 3<br>Intensive care unit  | <ul style="list-style-type: none"> <li>Requiring advanced respiratory support alone</li> <li>Requiring a minimum of two organ systems supported (except basic respiratory plus basic cardiovascular – level 2, as above)</li> </ul>   | Invasive mechanical ventilator support via endotracheal tube or tracheostomy<br>Acute renal replacement therapy and vasoactive medication  |

monitoring in recovery after surgery. An HDU bed may often be requested to ensure vigilance in the immediate postoperative period, and also to allow optimisation of pain control.

## Alternative Resources

Each individual institution will have slightly different facilities available for the care of their patients and these must be taken into consideration when planning postprocedural care. Early goal-directed therapy and utilisation of a ‘fast-track’ approach has been adopted successfully in many cardiothoracic centres and this may allow lower risk patients to be admitted to a cardiac recovery area as a temporary measure postoperatively, before being discharged back to a ‘stepdown’ unit or ward. For such systems to work and ensure safe patient care, there must be immediate access to critical care and adequate numbers of trained nursing staff. This model has been proven to be successful in some hospitals and can potentially improve patient flow. However, for many institutions the safest option remains to admit all cardiac surgical patients to the ICU postoperatively. The priority in such institutions is then to discharge out into a step-down unit as soon as possible after extubation and a period of stability.

Admission to an ICU may also depend on the availability of a required specific treatment for an individual patient. Some centres provide specialised advanced organ support, such as extracorporeal membrane oxygenation. Also, cardiothoracic surgery is a high-risk specialty fraught with potential complications, some of which might require transfer out to an alternative centre, for example to access neurosurgical intervention.

## Time of Admission

A well-organised cardiothoracic surgical centre should incorporate a robust system of communication with both its ICU and HDU with respect to the daily admission requirements and bed availability. The majority of patients undergoing anaesthesia will require elective admission and surgical activity will be planned according to such requirements.

However, the ICU and HDU must also always take into account the potential for unplanned emergency admissions, either transferred in for surgical intervention, or due to unexpected complications intraoperatively. Patients should be admitted to the required higher level of care before their condition reaches a point from which recovery may be extremely difficult. In reality, it is often much better practice to assume



### Which Patients are Admitted to Critical Care?

a bed will be needed for your patient, than be left in a situation where the availability is not there and the patient is unstable. This could potentially lead to a worsened patient outcome, and may also put unnecessary pressure on the relevant intensive care unit to discharge prematurely.

### Conclusion

It is often assumed that all patients undergoing cardiothoracic surgery will warrant admission to either an ICU or HDU postoperatively and in many instances that remains the case. Cardiothoracic anaesthesia is a high-risk specialty and it is imperative that the postoperative care system in place in each institution is safe and robust.

However, variety in admission indications and rates does exist. In recent years there have been

advances in providing 'fast-track' surgery, and cardiac recovery units have become increasingly popular. In addition, thoracic surgery does not always necessitate an HDU bed and often an adequate level of care can be provided on general wards with critical care outreach support. Requirements for a higher level of care are by no means well defined and clinical practice will continue to evolve with time.

Given the current climate in the health care system, with a constant pressure for beds and a drive to improve patient flow, it is extremely important that each case undergoing cardiothoracic anaesthesia is considered individually and the safest care for that patient determined. Such planning will take into consideration patient related factors, their diagnosis and required surgery and the resources available in the institution.



# Scoring Systems and Prognosis

Allanah Barker and Sam Nashef

## Crystal Balls

Knowing the likelihood of survival after cardiac surgery is useful for multiple reasons including for weighing the potential risks versus benefits of surgery. Further, accurate predicting of outcome allows for comparison with the actual outcome and thus insight into the overall performance of the cardiac surgical unit. Knowledge of who is likely to develop major morbidity also has an impact on the use of valuable resources and may allow for sensible planning of operating lists. In addition, some believe that being able to predict mortality with some certitude may help clinicians to determine when further efforts are futile. Unfortunately, the perfect predictor – a crystal ball to foresee the future – has not yet been fully developed.

## Risk Models or Scoring Systems

Scoring systems allow reasonable prediction of outcome after cardiac surgery. Many models have been devised to work out the likelihood of survival, and these and others have also been shown to predict major morbidity, long-term survival and resource use with some accuracy. Models can be broadly divided into two groups:

- *Preoperative models*, applied before the operation, with no knowledge of intraoperative events; and
- *Postoperative models*, applied immediately after the operation on admission into the critical care unit, taking some account of what the operation did to the patient.

## Preoperative Models

These are most useful for

- Establishing the risk of surgery as an adjunct to surgical decision making (determining the indication to operate on the basis of risk-to-benefit assessment);
- Providing the patient with information, which is helpful in obtaining consent;

- Helping to measure the performance of the service by comparing actual and predicted outcomes; and
- Comparing the performance of different institutions, surgeons and anaesthetists by correcting for risk when outcomes are assessed.

Preoperative models take no account of what happens in the operating theatre and are therefore less useful in predicting which of a number of postoperative patients with complications are likely to emerge intact from the critical care unit.

There are probably more risk models in cardiac surgery than in any other branch of medicine. Most rely on a combination of risk factors, each of which is given a numerical ‘weight’. Weights are added, multiplied or otherwise mathematically processed to come up with a percentage figure to predict mortality or survival. In additive models, the weights given to the risk factors are simply summed to give the predicted risk. They are easy to use and can be calculated mentally or ‘on the back of an envelope’. They are less accurate than more sophisticated systems and have a tendency to overscore slightly in low-risk patients and to underscore considerably in very high-risk patients. Examples of such models are the Parsonnet (the pioneering heart surgery risk model) and the original additive EuroSCORE for cardiac surgery overall. Other models deal specifically with cardiac surgical subsets, like coronary surgery and valve surgery. Sophisticated models use Bayesian analysis, logistic regression or even computer neural networks. They do not allow easy bedside calculation, necessitating a computer for determining risk. They are, however, more stable than additive models across the risk range and slightly more accurate in exact risk prediction. Examples of such models are the Society of Thoracic Surgeons (STS) model, the logistic EuroSCORE and EuroSCORE II for overall cardiac surgery.

The widespread application of scoring systems in heart surgery has allowed robust performance

## Scoring Systems and Prognosis

measurement and probably contributed to the dramatic drop in cardiac surgical mortality seen in the last 15 years.

### Preoperative Model Risk Factors

Not surprisingly, several common risk factors are included in all models (age, gender and left ventricular (LV) function). Other risk factors are included in some models but not in others, such as hypertension, diabetes and obesity. Models also differ depending on whether they deal with all cardiac surgeries or a specific subset, such as coronary surgery or valvular surgery. They share many risk factors and it would be repetitive to list them all here, but the models are easily accessible and there are interactive calculators available online: [www.euroscore.org](http://www.euroscore.org) and <http://riskcalc.sts.org/stswebriskcalc/>. EuroSCORE II also offers a smartphone 'app' for use at the bedside.

#### Age

There is an increased risk above the age of 60 years.

#### Gender

Females have a higher operative mortality than males, possibly because of smaller coronary artery size, smaller blood volume predisposing to risks associated with perioperative anaemia and transfusion, although the definitive reason for the difference is unknown.

#### Left Ventricular Function

As estimated by echocardiography or angiography, LV function is a good measure of cardiac status, but determination can be operator dependent and it is difficult to produce an accurate and reproducible percentage ejection fraction. Thus, LV function is generally classified as 'good', 'moderate' or 'poor'; EuroSCORE II has an additional category of 'very poor'.

#### Type of Surgery

General cardiac risk models take into account patients that undergo different surgeries – the risk for coronary artery bypass graft (CABG) surgery is less than for valve surgery, which in turn is less than that for surgery of the thoracic aorta. Combined procedures like valve with CABG carry a higher risk than single procedures.

### Extent of Cardiac Disease

The severity of coronary disease is subjective and therefore not included in surgical risk scores. The Syntax score allows for a measure of the severity of disease, but is time consuming and partly subjective. Left main stem disease may be associated with more risk. Objective measures of cardiac disease include recent myocardial infarction (MI), unstable angina and mechanical complications of MI such as acute rupture of the mitral valve or ventricular septum.

### Repeat Operation

Previous cardiac surgery (or previous sternotomy) increases difficulty of access and prolongs operative time. These patients therefore carry an increased risk of bleeding as well as possibly having more advanced disease than those undergoing their first cardiac procedure.

### Lung Disease

The presence of chronic pulmonary disease such as chronic obstructive pulmonary disease (COPD) has a large impact on how a patient is managed in anaesthetic and ventilatory terms. After cardiac surgery, patients with concurrent lung disease are more likely to require extended ventilation and to develop pulmonary complications, such as chest infections. Lung function is difficult to quantify with a single test and severity is based partly on subjective judgements. However, chronic pulmonary disease is taken into account in the EuroSCORE and STS.

### Renal Disease

Renal dysfunction, as evidenced by dependence on dialysis, increases mortality by as much as 40%, but the spectrum of renal failure is wide and difficult to quantify. Creatinine levels are easy to measure, but are not always an accurate measure of true kidney function. The original EuroSCORE uses grossly deranged serum creatinine ( $>200 \mu\text{mol/l}$ ) as a measure of significant renal impairment. Other scores use dialysis dependence. The best measure is probably creatinine clearance (CC), and this now features in EuroSCORE II, where the categories of renal dysfunction have expanded into four: normal function (CC  $> 85 \text{ ml/minute}$ ), moderate (CC  $50\text{--}85 \text{ ml/minute}$ ), severe (CC  $<50 \text{ ml/minute}$ ) and on dialysis (regardless of CC). Interestingly, patients with severe dysfunction but not on dialysis yet fare worst.

## Other Risk Factors

These include peripheral vascular disease, neurological dysfunction, degree of urgency, diabetes, hypertension and degree of pulmonary hypertension. In addition, various scoring systems give weight to the type of operation performed.

## Postoperative Models

These models benefit from information that is only available after the completion of the operation, such as the physiological parameters on admission to critical care. Many have been devised for critically ill patients outside the cardiac surgical specialty, but have been used and validated in cardiac surgery. The most well-known models are the Acute Physiology and Chronic Health Evaluation (APACHE) and the Sequential Organ Failure Assessment (SOFA) (Table 1). The APACHE score is used on admission to critical care to assess the risk of in-hospital death, whereas the SOFA was developed to quantify the severity of a patient's illness using the degree of organ dysfunction at any one time. The BRiSc score is specifically aimed at predicting patients likely to bleed excessively after heart surgery.

## Postoperative Model Risk Factors

Postoperative risk scores look at each organ system systematically and score according to derangement of function. Basically, the more organ dysfunction, the poorer the prognosis.

### Respiratory

Oxygenation and the requirement for ventilatory support are used as measures of respiratory function.

### Circulatory

Most scores which are applied postoperatively use mean arterial pressure as an easily measured and monitored parameter. However, whereas APACHE concentrates on derangement of normal physiology, SOFA concentrates on the need for (and level of) inotropic support.

### Neurological

Trends are more useful than a snapshot at a particular point in time, but the Glasgow Coma Scale is easily measured and provides an easily reproducible measure of neurological status.

## Renal

As preoperatively, the mainstay of renal function is serum creatinine level as it is easily measured and a relatively inexpensive test; this variable can be used to monitor changes in renal function and to compare current with preoperative function.

## Gastrointestinal/Hepatic

Both APACHE and SOFA use bilirubin levels as a measure of liver function. APACHE is used more widely in general critical care units and includes many more variables, such as amylase, albumin (as a rough measure of nutritional status) and other liver function tests. The APACHE score also contains variables to measure metabolic function and septic status. These criteria are less relevant in cardiac surgery.

## Thoracic Surgery

Risk modelling is not as developed in thoracic surgery, although recently some attempts have been made to produce models for predicting mortality after lung resection. The most important risk factors associated with a poor outcome are age (older people do less well) and how much functioning lung remains long after the resection (the more, the better).

## Learning Points

- Many models help to predict the outcome of cardiac surgery, and these can be applied before or after the operation.
- Preoperative models help in the decision making, consent and assessment of clinical performance.
- Postoperative models can help to plan resource use and provide information to relatives.
- Models devised specifically for mortality have also been found to be useful in predicting major morbidity, resource use and long-term outcomes.
- No amount of risk modelling can predict with certainty which patient will live and which will die and they should be used as an adjunct rather than as a replacement for sound clinical judgement.

## Scoring Systems and Prognosis

**Table 1** Postoperative cardiac surgery risk assessment scores

| Organ system               | SOFA  | APACHE   |
|----------------------------|---|--|
| Respiratory                | Oxygenation (PaO <sub>2</sub> /FiO <sub>2</sub> )<br>Respiratory support                            | Respiratory rate non-ventilated<br>PaO <sub>2</sub> with FiO <sub>2</sub> 1.0<br>PaCO <sub>2</sub>   |
| Coagulation/haematological | WCC   | WCC<br>Haematocrit<br>Platelet count<br>Prothrombin time   |
| Circulatory                | Mean arterial pressure<br>Dopamine dose<br>Adrenaline dose<br>Norepinephrine dose<br>Dobutamine use | Mean arterial pressure<br>Heart rate ventricular response<br>Central venous pressure<br>Evidence of acute MI<br>Arrhythmia<br>Serum lactate<br>Arterial pH |
| Neurological               | Glasgow Coma Scale  | Glasgow Coma Scale   |
| Renal                      | Creatinine<br>Urine output/24 hour  | Creatinine<br>Urine output/24 hour<br>Blood urea nitrogen  |
| Gastrointestinal/hepatic   | Bilirubin   | Amylase<br>Albumin<br>Bilirubin<br>Alkaline phosphatase<br>Liver enzymes<br>Anergy by skin testing   |
| Septic                     |   | Cerebrospinal fluid positive culture<br>Blood culture positive<br>Fungal culture positive<br>Rectal temperature  |
| Metabolic                  |   | Calcium level<br>Glucose<br>Sodium<br>Potassium<br>Bicarbonate<br>Serum osmolarity   |

Abbreviations: APACHE, Acute Physiology and Chronic Health Evaluation; FiO<sub>2</sub>, fraction of inspired oxygen; MI, myocardial infarction; PaCO<sub>2</sub>, partial pressure of carbon dioxide in arterial blood; PaO<sub>2</sub>, partial pressure of oxygen in arterial blood; SOFA, Sequential Organ Failure Assessment; WCC, white cell count.

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## Abbreviations

|        |   |         |   |
|--------|---|---------|---|
| AC     | Assist-Control ventilation                        | BLUE    | Beside Lung Ultrasound in<br>Emergency                                |
| ACBT   | Active Cycle of Breathing Technique               | BNP     | B-type Natriuretic Peptide  |
| ACEI   | Angiotensin Converting Enzyme<br>Inhibitor        | BPF     | Bronchopleural Fistula  |
| ACLS   | Advanced Cardiac Life Support                     | BPS     | Behavioural Pain Scale  |
| ACT    | Activated Clotting Time                           | BTC     | Bridge to Candidacy   |
| AD     | Advanced Directive                                | BTS     | British Thoracic Society  |
| AEDs   | Automated External Defibrillators                 | BTT     | Bridge to Transplant  |
| AEG    | Atrial Electrocardiogram                          | BURP    | Backwards, Upwards and Rightward<br>Pressure on the thyroid cartilage |
| AEP    | Auditory Evoked Potentials                        | CABG    | Coronary Artery Bypass Grafting                                       |
| AF     | Atrial Fibrillation                               | CAM-ICU | Confusion Assessment Method for the<br>ICU                            |
| AFE    | Amniotic Fluid Embolism                           | CAP     | Community Acquired Pneumonia  |
| AKI    | Acute Kidney Injury                               | CC      | Creatinine Clearance  |
| ALG    | Anti-human Lymphocyte Globulin                    | CCA     | Critical Care Area  |
| ALS    | Advanced Life Support                             | CCS     | Canadian Cardiovascular Society                                       |
| AMP    | Adenosine Monophosphate                           | ccTGA   | Congenitally Corrected Transposition<br>of the Great Arteries         |
| APACHE | Acute Physiology and Chronic Health<br>Evaluation | CCU     | Coronary Care Unit  |
| APRV   | Airway Pressure Release Ventilation               | CDC     | Centers for Disease Control   |
| aPTT   | Activated Partial Thromboplastin<br>Time          | cEEG    | Continuous Electroencephalography                                     |
| AR     | Aortic Regurgitation                              | CF      | Cystic Fibrosis   |
| ARB    | Angiotensin Receptor Blockers                     | CHD     | Congenital Heart Disease  |
| ARDS   | Acute Respiratory Distress<br>Syndrome            | CHF     | Congestive Heart Failure  |
| ARF    | Acute Respiratory Failure                         | CICO    | 'Can't Intubate, Can't Oxygenate'                                     |
| ASD    | Atrial Septal Defect                              | CIN     | Contrast Induced Nephropathy  |
| ATG    | Anti-human Thymocyte Globulin                     | CI      | Cardiac Index   |
| ATLS   | Advanced Trauma Life Support                      | CIS     | Clinical Information Systems  |
| AVNRT  | Atrioventricular Node Re-entrant<br>Tachycardia   | CK      | Creatinine Kinase   |
| AVSD   | Atrioventricular Septal Defect                    | CKD     | Chronic Kidney Disease  |
| BAL    | Bronchoalveolar Lavage                            | CLABSI  | Central Line Associated Bloodstream<br>Infections                     |
| BALF   | Bronchoalveolar Lavage Fluid                      | CLAD    | Chronic Lung Allograft Dysfunction                                    |
| BIPAP  | Biphasic or Bilevel Positive Airway<br>Pressure   | CMR     | Cardiac Magnetic Resonance  |
| BIPDs  | Bilateral Independent PDs                         | CMV     | Continuous Mandatory Ventilation                                      |
| BIS    | Bispectral Index                                  | CMV     | Cytomegalovirus   |
| BiVAD  | Bilateral Ventricular Assist Device               | CNI     | Calcineurin Inhibitors  |
| BLS    | Basic Life Support                                | CO      | Cardiac Output  |

## Abbreviations

|                     |   |                  |   |
|---------------------|---|------------------|---|
| COAD                | Chronic Obstructive Airways Disease, same as COPD | ELSO             | Extracorporeal Life Support Organisation        |
| COPD                | Chronic Obstructive Pulmonary Disease             | EMR              | Electronic Medical Records                      |
| CP                  | Constrictive Pericarditis                         | ERP              | Enhanced Recovery Programmes                    |
| CPAP                | Constant Positive Airway Pressure                 | ESBL             | Extended Spectrum Beta-Lactamases               |
| CPAx                | Chelsea Critical Care Physical Assessment Tool    | ESG              | Endovascular stent graft                        |
| CPB                 | Cardiopulmonary Bypass                            | ETT              | Endotracheal Tube                               |
| CPE                 | Carbapenemase Producing Enterobacteriaceae        | EVLWI            | Extravascular Lung Water Index                  |
| CPOT                | Critical Care Pain Observation Tool               | EWMA             | Exponentially Weighted Moving Average           |
| CPP                 | Cerebral Perfusion Pressure                       | EWS              | Early Warning Scores                            |
| CRP                 | C-Reactive Protein                                | FAC              | Fractional Area Change                          |
| CT                  | Computerised Tomography                           | FALLS            | Fluid Administration Limited by Lung Sonography |
| CTCA                | Computerised Tomography Coronary Angiogram        | FAM              | Functional Assessment Measure                   |
| CTEPH               | Chronic Thromboembolic Pulmonary Hypertension     | FB               | Flexible Bronchoscopy                           |
| CV                  | Stroke Volume                                     | FBC              | Full Blood Count                                |
| CVC                 | Central Venous Catheter                           | FDO <sub>2</sub> | Fraction of Oxygen Delivered                    |
| CVD                 | Cardiovascular Disease                            | FEV1             | Forced Expiratory Volume for 1 second           |
| CVP                 | Central Venous Pressure                           | FFP              | Fresh Frozen Plasma                             |
| CXR                 | Chest X-Ray                                       | FIM              | Functional Independence Measure                 |
| DAG                 | 1,2-Diacylglycerol                                | FIRDA            | Frontal IRDA                                    |
| DBD                 | Donation after Brain Death                        | FOUR             | Full Outline of Unresponsiveness                |
| DBexs               | Deep Breathing Exercises                          | FRC              | Function of Residual Capacity                   |
| DCD                 | Donation after Circulatory Death                  | FS               | Fraction of Shortening                          |
| DD                  | Diastolic Dysfunction                             | FVC              | Forced Vital Capacity                           |
| DNAR                | Do Not Attempt Resuscitation Order                | GBS              | Guillain-Barré Syndrome                         |
| DOLS                | Deprivation of Liberty Safeguards                 | GCS              | Glasgow Coma Score                              |
| DSI                 | Daily Sedation Interruption                       | GEDVI            | Global End-Diastolic Volume Index               |
| DT                  | Destination Therapy                               | GICS             | Gastrointestinal Complication Score             |
| DTI                 | Direct Thrombin Inhibitor                         | GPCR             | G Protein Coupled Receptors                     |
| DVT                 | Deep Venous Thrombosis                            | GUCH             | Grown-Up Congenital Heart disease               |
| EACA                | Epsilon Aminocaproic Acid                         | HD               | Haemodialysis                                   |
| ECC                 | Emergency Cardiovascular Care                     | HDF              | Haemodiafiltration                              |
| ECCO <sub>2</sub> R | Extracorporeal Carbon Dioxide Removal             | HDU              | High Dependency Unit                            |
| ECG                 | Electrocardiography                               | HES              | Hydroxyethyl Starch                             |
| ECLS                | Extracorporeal Life Support                       | HF               | Haemofiltration                                 |
| ECMO                | Extracorporeal Membrane Oxygenation               | HFV              | High Frequency Ventilation                      |
| ECPR                | Extracorporeal Cardiopulmonary Resuscitation      | HIT              | Heparin Induced Thrombocytopenia                |
| EDA                 | End-Diastolic Area                                | HIV              | Human Immunodeficiency Virus                    |
| EEG                 | Electroencephalography                            | HLHS             | Hypoplastic Left Heart Syndrome                 |
| EF                  | Ejection Fraction                                 | HOCM             | Hypertrophic Obstructive Cardiomyopathy         |
| ELISA               | Enzyme-Linked Immunosorbent Assay                 | HSV              | Herpes Simplex Virus                            |
|                     |   | HTEA             | High Thoracic Epidural Analgesia                |
|                     |   | IABP             | Intra-aortic Balloon Pump                       |
|                     |   | ICD              | Implantable Cardioverter-Defibrillators         |
|                     |   | ICP              | Intracranial Pressure                           |



## Abbreviations

|              |  |       |  |
|--------------|--|-------|--|
| ICSD         | Intensive Care Delirium Screening Checklist                                  | MDT   | Multidisciplinary Team                             |
| ICU-AW       | Intensive Care Unit Acquired Weakness  | MET   | Medical Emergency Teams                            |
| IE           | Infective Endocarditis   | MHI   | Manual Hyperinflation                              |
| IJV          | Internal Jugular Vein  | MI    | Myocardial Infarction                              |
| IMCA         | Independent Mental Capacity Advocate   | MIC   | Minimum Inhibitory Concentration                   |
| IMV          | Invasive Mechanical Ventilation  | MMF   | Mycophenolate Mofetil                              |
| INR          | International Normalised Ratio   | MMV   | Mandatory Minute Ventilation                       |
| INTERMACS    | Interagency Registry for Mechanically Assisted Circulatory Support           | mPAP  | Mean Pulmonary Arterial Pressure                   |
| IPF          | Idiopathic Pulmonary Fibrosis  | MR    | Mitral Regurgitation                               |
| IR           | Interventional Radiology   | MRSA  | Methicillin Resistant <i>Staphylococcus aureus</i> |
| IRDA         | Intermittent Rhythmic Delta Activity   | MSE   | Myoclonic Status Epilepticus                       |
| IRV          | Inversed Ratio Ventilation   | MSSA  | Methicillin-Sensitive <i>Staphylococcus aureus</i> |
| IS           | Incentive Spirometry   | mTOR  | Mammalian Target of Rapamicin Inhibitors           |
| ISHLT        | International Society for Heart and Lung Transplantation                     | MUST  | Malnutrition Universal Screening Tool              |
| ITBVI        | Intrathoracic Blood Volume Index   | MV    | Mitral Valve                                       |
| IUGR         | Intrauterine Growth Retardation  | NAAT  | Nucleic Acid-Based Amplification Technologies      |
| IVC          | Inferior Vena Cava   | NAP4  | Fourth National Audit Project                      |
| IVS          | Interventricular Septum  | NAVA  | Neurally Adjusted Ventilatory Assist               |
| JET          | Junctional Ectopic Tachycardia   | NCS   | Non-convulsive Seizures                            |
| LAD          | Left Anterior Descending artery  | NCSE  | Non-convulsive Status Epilepticus                  |
| LAS          | Lateral Amiotrophic Sclerosis  | NHSBT | National Health Service Blood and Transfusion      |
| LBBB         | Left Bundle Branch Block   | NI    | Narcotrend Index                                   |
| LCx          | Left Circumflex Artery   | NICE  | National Institute for Clinical Excellence         |
| LMA          | Laryngeal Mask Airway  | NIPPV | Non-invasive Positive Pressure Ventilation         |
| LMCA         | Left Main Coronary Artery  | NIRS  | Near Infrared Spectroscopy                         |
| LMWH         | Low Molecular Weight Heparin   | NIV   | Non-invasive Ventilation                           |
| LTACH        | Long-Term Acute Care Hospitals   | NMDA  | <i>N</i> -Acetyl-D-Aspartate receptor              |
| LV           | Left Ventricle   | NOAC  | Newer Oral Anticoagulants                          |
| LVAD         | Left Ventricular Assist Device   | NRS   | Nutritional Risk Screening                         |
| LVEDV        | Left Ventricular End-Diastolic Volume  | NVE   | Native Valve Endocarditis                          |
| LVESV        | Left Ventricular End-Systolic Volume   | NYHA  | New York Heart Association                         |
| LVOT         | Left Ventricular Outflow Tract   | OD    | Optical Density                                    |
| LVOTO        | Left Ventricular Outflow Tract Obstruction                                   | OHCA  | Out-of-Hospital Cardiac Arrest                     |
| LVSF         | Left Ventricular Systolic Function   | OIRDA | Occipital IRDA                                     |
| MACE         | Major Adverse Cardiac Events   | OpCAB | Off pump Coronary Artery Bypass                    |
| MALDI TOF MS | Matrix Assisted Laser Desorption/Ionisation Time-of-Flight Mass Spectrometry | PAC   | Pulmonary Artery Catheter                          |
| MAO          | Monoamine Oxydase  | PAH   | Pulmonary Arterial Hypertension                    |
| MAP          | Mean Arterial Pressure   | PAP   | Pulmonary Arterial Pressure                        |
| MCCD         | Mechanical Chest Compression Devices   | PAWP  | Pulmonary Arterial Wedge Pressure                  |
| MCFP         | Mean Circulatory Filling Pressure  | PBM   | Patient Blood Management                           |
| MDR          | Multidrug Resistance   | PBW   | Predicted Body Weight                              |
|              |  | PCAS  | Post-Cardiac Arrest Syndrome                       |
|              |  | PCI   | Percutaneous Coronary Intervention                 |



## Abbreviations

|      |   |       |   |
|------|---|-------|---|
| PCP  | <i>Pneumocystis jirovecii</i> Carinii<br><i>Pneumonia</i> | ROTEM | Rotational Thromboelastometry   |
| PCR  | Polymerase Chain Reaction                                 | RRT   | Renal Replacement Therapy   |
| PCT  | Procalcitonin   | RV    | Right Ventricle   |
| PCWP | Pulmonary Capillary Wedge Pressure                        | RVP   | Right Ventricular Pressure  |
| PD   | Peritoneal Dialysis                                       | RVAD  | Right Ventricular Assist Device   |
| PDA  | Posterior Descending Artery                               | RWMA  | Regional Wall Motion Abnormalities  |
| PDE  | Phosphodiesterase Inhibitors                              | SACP  | Selective Antegrade Cerebral<br>Perfusion                                 |
| PDR  | Posterior Dominant Rhythm                                 | SAH   | Subarachnoid Haemorrhage  |
| PDs  | Periodic Discharges                                       | SAM   | Systolic Anterior Motion  |
| PE   | Pulmonary Embolism  | SAS   | Sedation Agitation Scale  |
| PEA  | Pulmonary Endarterectomy                                  | SDD   | Selective Digestive Decontamination                                       |
| PEEP | Positive End Expiratory Pressure                          | SE    | Status Epilepticus  |
| PF   | Pulmonary Fibrosis  | SGA   | Subjective Global Assessment  |
| PF4  | Platelet Factor 4   | SIMV  | Synchronised Intermittent Mandatory<br>Ventilation                        |
| PFIT | Physical Functional Intensive Care<br>Test                | SLED  | Slow Low-Efficiency Dialysis  |
| PGD  | Primary Graft Dysfunction                                 | SMR   | Standardised Mortality Ratio  |
| PH   | Pulmonary Hypertension, same as<br>PAH                    | SOFA  | Sepsis Related Organ Failure<br>Assessment                                |
| PKC  | Protein Kinase C  | SR    | Sarcoplasmic Reticulum  |
| PLC  | Phospholipase C   | SRA   | Serotonin Release Assay   |
| PPCs | Postoperative Pulmonary<br>Complications                  | SSEP  | Somatosensory Evoked Potentials   |
| PPCI | Primary Percutaneous Coronary<br>Intervention             | SSRI  | Selective Serotonin Reuptake Inhibitor                                    |
| PPCM | Peripartum Cardiomyopathy                                 | SVC   | Superior Vena Cava  |
| PPHN | Persistent Pulmonary Hypertension of<br>the Newborn       | SVCS  | Superior Vena Cava Syndrome   |
| PPV  | Pulse Pressure Variation                                  | SVR   | Systemic Vascular Resistance  |
| PRC  | Post-resuscitation Care                                   | TAA   | Thoracic Aortic Aneurysm  |
| PRES | Posterior Reversible Encephalopathy<br>Syndrome           | TAH   | Total Artificial Heart  |
| PRVC | Pressure Regulated Volume<br>Controlled Ventilation       | TAPSE | Tricuspid Annular Plane Systolic<br>Excursion                             |
| PSI  | Patient State Index                                       | TAPVD | Total Anomalous Pulmonary Venous<br>Drainage                              |
| PT   | Prothrombin Time  | TCPC  | Total Cavopulmonary Connection  |
| PTE  | Pulmonary Thromboendarterectomy,<br>same as PEA           | TEG   | Thromboelastography   |
| PTLD | Post-transplantation<br>Lymphoproliferative Disorder      | TETS  | Transcutaneous Energy Transfer<br>Systems                                 |
| PVE  | Prosthetic Valve Endocarditis                             | TEVAR | Thoracic Endovascular Aortic Repair                                       |
| PVR  | Pulmonary Vascular Resistance                             | TGA   | Transposition of the Great Arteries                                       |
| RAP  | Right Atrial Pressure                                     | TnC   | Troponin C  |
| RASS | Richmond Agitation Sedation Scale                         | TNF   | Tumour Necrosis Factor  |
| RBBB | Right Bundle Branch Block                                 | TOE   | Transoesophageal Echocardiography   |
| RCM  | Restrictive Cardiomyopathy                                | TOF   | Tetralogy of Fallot   |
| RCT  | Randomised Controlled Trial                               | TPG   | Transpulmonary Gradient   |
| ROC  | Receiver Operating Characteristic                         | TR    | Tricuspid Regurgitation   |
| ROSC | Return of Spontaneous Circulation                         | TRALI | Transfusion Related Lung Injury   |
|      |   | TTE   | Transthoracic Echocardiography or<br>Thoracic Expansion Exercises as TTEs |
|      |   | TTM   | Targeted Temperature Management   |
|      |   | TXA   | Tranexamic Acid   |

**Abbreviations**

|      |                                   |      |  |
|------|-----------------------------------|------|--|
| URR  | Urea Reduction Ratio              | VT   | Ventricular Tachycardia                  |
| VALI | Ventilator Associated Lung Injury | VTI  | Velocity-Time Integral                   |
| VAP  | Ventilator Associated Pneumonia   | VTM  | Viral Transport Media                    |
| VATS | Video Assisted Thoracic Surgery   | vWF  | von Willebrand Factor                    |
| VF   | Ventricular Fibrillation          | WCRS | Withdrawal of Cardiorespiratory Supports |
| VHI  | Ventilator Hyperinflation         | WOB  | Work of Breathing                        |
| VRE  | Vancomycin Resistant Enterococci  | WPW  | Wolff–Parkinson–White syndrome           |
| VSD  | Ventricular Septal Defect         |      |  |