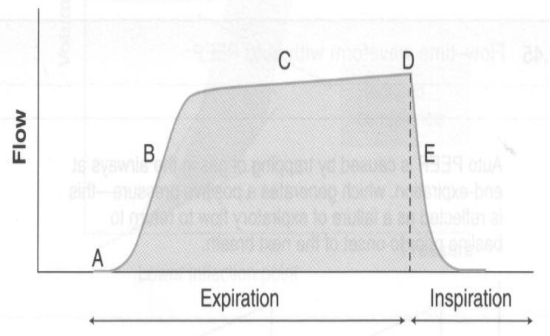


capnography

general

- Capnometry consists of the measurement and numeric display of expired carbon dioxide (CO₂) at the patient's airway opening. When a waveform plotting CO₂ against time or volume is also displayed, it is referred to as capnography, and the waveform is referred to as a capnogram.

- Capnometry works by passing infrared light through a sample chamber to a detector on the opposite side. More infrared light passing through the sample chamber (i.e., less CO₂) causes a larger signal in the detector relative to the infrared light passing through a reference cell.



- A = Zero baseline
- B = Rapid sharp rise at start of expiration
- C = Alveolar plateau
- D = End-tidal CO₂ value
- E = Rapid sharp downstroke with inspiration

normal capnography waveform

capnography traces

Waveform characteristic	Significance
Sudden drop in ETco ₂ to zero or near zero	Circuit disconnection Dislodged, occluded or kinked endotracheal tube Oesophageal intubation Massive CO ₂ embolism during laparoscopy
Sudden drop in ETco ₂ to lower levels without well-defined normal plateaus	Circuit leak Partial airway occlusion
Persistent low ETco ₂ with normal plateaus	Hypothermia Hypoventilation Deep anaesthesia/sedation Increased dead-space ventilation
Persistent low ETco ₂ without normal plateaus	Incomplete expiration related to partial airway occlusion, bronchospasm, mucus plugging or poor stream sampling
Marked decrease in ETco ₂ with normal plateaus	Reduced pulmonary blood flow—pulmonary embolism (clot, fat, air), profound hypotension
Elevated ETco ₂ with normal plateaus	Inadequate minute volume but adequate tidal volumes Increased CO ₂ production (e.g. hyperthermia)
Gradually increasing ETco ₂ with normal plateaus	Hypoventilation Sudden increase in metabolic rate (e.g. malignant hyperthermia)

interpretation

PaCO₂-PetCO₂ GRADIENT:

- Normal subjects have a PaCO₂-PETCO₂ gradient of 4 to 5 mm Hg,
- In critically ill patients, the PaCO₂-PETCO₂ gradient can be markedly elevated
- During expiration, lung regions with high ventilation-to-perfusion ratios dilute the mixed CO₂ concentration so that PETCO₂ is usually lower than PaCO₂
- When CO₂ production is elevated (or expiration is prolonged), PETCO₂ more closely resembles mixed venous PCO₂, as a higher amount of CO₂ diffuses into a progressively smaller lung volume.
- Thus, the PaCO₂-PETCO₂ gradient can be affected by changes in respiratory rate and tidal volume (VT), owing to alterations in expiratory time, and by CO₂ production and mixed venous CO₂ content.
- Inotropic or vasoactive drugs may affect the PaCO₂-PETCO₂ gradient in an unpredictable manner, either by increasing cardiac output and pulmonary perfusion (thereby reducing alveolar deadspace) or by reducing pulmonary vascular resistance and magnifying intrapulmonary shunt by countering hypoxic pulmonary vasoconstriction.

PaCO₂-PETCO₂ GRADIENT, PEEP, AND LUNG RECRUITMENT:

- PEEP recruits collapsed alveoli, improves ventilation-perfusion matching, and reduces alveolar deadspace, although excessive levels cause overdistention and increased alveolar deadspace.
- Because the PaCO₂-PETCO₂ gradient correlates strongly with the physiologic deadspace-to-tidal volume ratio (VD/VT), it may be useful in titrating PEEP in acute respiratory distress syndrome (ARDS) although data in humans are limited.

PetCO₂ MONITORING DURING CARDIOPULMONARY RESUSCITATION:

- Monitoring end-tidal CO₂ concentration is a reliable method for evaluating the effectiveness of cardiopulmonary resuscitation with changes in PETCO₂ directly proportional to changes in cardiac output.
- PETCO₂ during precordial compressions can distinguish successful from unsuccessful resuscitation, with values greater than 10 mm Hg or greater than 16 mm Hg depending on the study associated with successful resuscitation.

MEASUREMENT OF DEADSPACE VENTILATION

- Ventilation-perfusion abnormalities are the primary physiologic disturbance in nearly all pulmonary diseases and the principal mechanism for elevated PaCO₂.
- Dead-space ventilation (VD), the portion of VT that does not encounter perfused alveoli, directly impacts CO₂ excretion and is used as an indirect measure of ventilation-perfusion abnormalities.
- Physiologic VD/VT is used to assess the severity of pulmonary disease and the efficacy of ventilator manipulations. It may be particularly useful in patients with suspected PE
- Expired gas collection with a Douglas bag is the classic method for measuring VD/VT.
- Metabolic monitors produce equally accurate, reliable results and are less cumbersome.
- In addition, newer monitors incorporating capnography and pneumotachygraphy now provide accurate single-breath determinations of VD/VT.

use

- (i) the diagnosis of pulmonary embolism,
- (ii) determination of lung recruitment response to positive end-expiratory pressure (PEEP),
- (iii) detection of intrinsic PEEP,
- (iv) evaluation of weaning,
- (v) indirect marker of elevated deadspace ventilation,
- (vi) assessment of cardiopulmonary resuscitation,
- (vii) indirect determination of cardiac output through partial CO₂ rebreathing,
- (viii) verification of endotracheal cannulation,
- (viii) detection of airway accidents, and
- (ix) verification of feeding tube placement.