

Review Article





# Endotracheal Cardiac Output Monitoring (ECOM)

#### **Abstract**

Cardiac output reflects the amount of blood expelled by the heart in one minute and calculated as the product of the heart rate by the stroke volume. The variables that affect systolic volume are preload, afterload, and contractile function. Cardiac output is the main determinant of oxygen transport and its monitoring helps to establish the differential diagnosis of the possible causes of shock and optimizes its treatment. A less invasive technique to monitor cardiac output in real time is always preferable as it can be evaluated more quickly and easily, even if it is slightly less accurate, especially in critical situations that require a rapid assessment of the patient's condition. There are a number of techniques that allow us to obtain measurements of cardiac output in a very effective and non-invasive way. The endotracheal cardiac output monitoring (ECOM) is an emerging technology for intubated patients enabling rapid real time and accurate data on the status of the cardiac output and various derived parameters. ECOM is poised to be the new gold standard in critical care management and an emerging toll in the armamentarium of intensivists.

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# Introduction

Cardiac output (CO) is the volume of blood ejected by each ventricle per minute and is the product of stroke volume and heart rate. CO is an integral part of hemodynamic monitoring during anesthesia for major and high risk surgeries as it serves as a marker of oxygen delivery to the tissues and flags patients at risk of significant morbidity, mortality or both. It also guides treatment, primarily for fluid resuscitation and the use of vasoactive and inotropic drugs. An ideal CO monitor should be minimally or non-invasive, continuous, accurate, operator independent, cost effective, reproducible, reliable during various physiological states, inexpensive, safe and have fast response time.<sup>1</sup>

The use of thermos dilution technique using Swan Ganz catheters remained a gold standard for cardiac output measurement for many years. The search for newer, less invasive methods of measuring CO led to the introduction of numerous devices like arterial pulse contour analysis, thoracic bio reactance, vascular unloading technique, pulse wave transit time, and radial artery application tonometry, aortic Doppler and thoracic electrical bio impedance into clinical practice.<sup>2,3</sup>

The term 'minimally invasive cardiac output monitors' collectively describes all devices that calculate CO without requiring insertion of a pulmonary artery catheter (PAC). Each of these devices, utilizes different parameters to determine CO. The latest endotracheal cardiac output monitor (ECOM), is based on the principle that blood is a charged solution with fluid motion and volumetric changes as it is pumped by the left ventricle. Smart sensors and transmitters affixed to the cuff of the endotracheal tube in direct contact with the ascending aorta measure impedance signals that use algorithms to determine the cardiac output and other derived parameters in real time.

#### **Evolution of co measurement**

The early methods of assessing cardiac ouput were all indirect and invasive measurements. Adolf Eugen Fick (1829-1901) was the first to measure cardiac output in 1870 by measurement of

oxygen concentration of arterial and venous blood and subsequent calculation of O2 consumption. In 1929, Werner Forssman inserted a ureteral catheter to perform right atrial catheterization on himself and documented this on x-ray film! In 1945, Coumand and Richards measured CO by direct Fick method. 1956: Forssmann, Cournand and Richards jointly won the Nobel Prize in Medicine. In 1970, Jeremy Swan and William Ganz developed their eponymous pulmonary artery (PA) catheter and revolutionized measurement of CO. Their invention enabled diagnostic measurements at the bedside and contributed to the birth of critical care medicine. In 1972, Forrester added a thermistor to the pulmonary artery catheter, allowing CO measurements by thermos dilution. Newer methods like bio impedance, Doppler and ECOM FOR non-invasive and direct, real time measurements of CO followed. The advantages and drawbacks of each technique is enumerated in Tables 1 and 2.4-6

#### ECOM:

The principle for ECOM is that blood is a charged solution with fluid motion and volumetric changes as it is pumped by the left ventricle. The electric resistance of blood changes when it moves or fluctuates in volume.



ECOM is dramatically different from other cardiac output monitors and based on bio impedance technology. The ECOM system uses an adaptive multi-parameter algorithm which allows for the reduction of between-subject variability. The endotracheal tube has 7 silver doped plastic electrodes. A low intensity high frequency alternating current (4 milliamperes at 20–100 kiloHertz) with input





86

impedance of 1 mOhms applied to aspecially designed 7.5-mm internal diameter endotracheal tube designed with six electrodes with conducting ink on a plastic back ingglued on the cuff and one on the shaft. The electrode surface is smooth and atraumatic to the tracheal mucosa and the shaft electrode is a 15-mm-wide band that serves as a ground. Three orthogonal pairs of sensing transmitting and receiving electrodes (1 and 2, 4 and 5, 1 and 6) glued on the cuff is positioned immediately proximal to the tracheal bifurcation and in direct contact with the ascending aorta. Three impedance signals ( $DZ^x$ ,  $DZ^y$ ,  $DZ^z$ ) are measured without any interference or anomalous signals from heart structures, lungs or the great vessels (Figures 1-3). A threedimensional impedance field is derived from these three inputs. Since the electrodes are positioned in close proximity to the ascending aorta the inputs are highly accurate. The electrode arrays and ECOM software have inbuilt software that automatically compensates for any changes in the positioning of the endotracheal tube and also that of the patient's body, and displays a beat to beat measurement of the CO and derived parameters non invasively. The application of current to the electrodes on the endotracheal tube do not cause any injury to the tracheal mucosa even after 24 h of continuous use, as all circuits are electrically isolated.7-10

Table I

Continuous CO by PAC   Invasive PAC   Continuous CO measurement   Catheter related complications Cost	1	PAC	Invasive	Gold standard	Catheter related complications
invasive measurements Measure SV and SVV Lithium therapy  A Picco Minimally invasive during hemodynamic instability  FloTrac Minimally invasive  Measure flow only in descending thoracic aorta Assumptions about aortic size may not be accurate  Minimally invasive  Minimally invasive  Measure flow only in descending thoracic aorta Assumptions about aortic size may not be accurate  Minimally invasive  Measure flow only in descending thoracic aorta Assumptions about aortic size may not be accurate  Affected by changes in dead space or VQ matching invasive systems  Thoracic Non Continuous CO measurement bioimpedance invasive  Thoracic Non Continuous CO measurement invasive hemodynamic stability Not useful in dysrhythmias  The ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	2		Invasive	Continous CO measurement	Catheter related complications Cost
invasive during hemodynamic instability  5 FloTrac Minimally Continuous CO measurement No calibration invasive  6 PRAM Minimally No calibration Still not validated invasive  7 ED Minimally Simple to use Reliable Useful in GDT Measure flow only in descending thoracic aorta Assumptions about aortic size may not be accurate  8 TEE Minimally Evaluate cardiac anatomy preload and invasive ventricular function  9 Partial non-rebreathing invasive systems  10 Thoracic Non Continuous CO measurement bioimpedance invasive invasive hemodynamic stability. Not useful in dysrhythmias  11 ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	3	LiDCO		•	
invasive  7 ED Minimally invasive  8 TEE Minimally Evaluate cardiac anatomy preload and invasive ventricular function  9 Partial non-rebreathing systems  10 Thoracic Non Continuous CO measurement bioimpedance invasive invasive bioimpedance invasive invasive hemodynamic stability Not useful in dysrhythmias  11 ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	4	PiCCO			Requires good arterial waveform Requires calibration
invasive  TED Minimally Simple to use Reliable Useful in GDT Measure flow only in descending thoracic aorta Assumptions about aortic size may not be accurate  TEE Minimally Evaluate cardiac anatomy preload and ventricular function  Partial non-rebreathing invasive systems  Thoracic Non Continuous CO measurement bioimpedance invasive invasive invasive hemodynamic stability Not useful in dysrhythmias  Thoracic Non Continuous CO measurement hemodynamic stability Not useful in dysrhythmias  Coronary blood flow not recorded Electrocautery produces interference	5	FloTrac		Continuous CO measurement No calibration	Requires good arterial waveform
invasive size may not be accurate  8 TEE Minimally Evaluate cardiac anatomy preload and cost Skilled personnel invasive ventricular function  9 Partial non- Non Ease of use Continuous CO measurement rebreathing invasive systems  10 Thoracic Non Continuous CO measurement hemodynamic stability Not useful in dysrhythmias  11 ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	6	PRAM		No calibration	Still not validated
invasive ventricular function  9 Partial non- rebreathing invasive systems  10 Thoracic Non Continuous CO measurement bioimpedance invasive bioimpedance invasive  TCOM Non Continuous CO measurement hemodynamic stability Not useful in dysrhythmias  Coronary blood flow not recorded Electrocautery produces interference	7	ED		Simple to use Reliable Useful in GDT	
rebreathing invasive systems  10 Thoracic Non Continuous CO measurement Affected by electrical noise, movement, temperature and humidity Requires bioimpedance invasive hemodynamic stability Not useful in dysrhythmias  11 ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	8	TEE			Cost Skilled personnel
bioimpedance invasive hemodynamic stability Not useful in dysrhythmias  11 ECOM Non Continuous CO measurement Coronary blood flow not recorded Electrocautery produces interference	9	rebreathing		Ease of use Continuous CO measurement	Affected by changes in dead space or V/Q matching
,,,,,	10			Continuous CO measurement	
	11	ECOM		Continuous CO measurement	Coronary blood flow not recorded Electrocautery produces interference

CO: Cardiac output; LiDCO: Lithium dilution CO; PiCCO and FloTrac: Pulse contour analysis; PRAM: Pressure recording analytic method; ED: Esophgeal Doppler; TEE: Transesophgeal echocardiography; ECOM: Endotracheal cardiac output monitor; PAC: Pulmonary artery catheter; SV: Stroke volume; SVV: SV variation; GDT: Goal directed therapy.

Table 2

Overview of	characteristic	s for differe	nt method	s to measu	are cardiac output
PAC CCO	+++	Continuous	Moderate	Moderate	Requires a PAC and triplicate measurements
TP Td bolus	++	Intermittent	High	High	Requires a PAC and triplicate measurements
TP Li bolus	++	Intermittent	Moderate	Moderate	Requires only arterial catheter but needs triplicate measurement for sufficient agreement with reference methods
PiCCO	++	Beat-to- beat	Moderate	Moderate	Requires frequent calibration with independent (other) method
LiDCO	++	Beat-to- beat	Moderate	Moderate	Requires frequent calibration with independent (other) method or lithium indicator method
Vigileo	++	Beat-to- beat	Moderate	High	Needs specific sensor
Modelflow	++	Beat-to- beat	High	High	Needs femoral or radial arterial catheter
TOD	+	Continuous	High	Low	Not well tolerated in awake subjects and transducer position difficult
TTE	-	Continuous	Moderate	Low	Large inter-operator variability
Bioimpedance	-	Continuous	Low	Low	Artifacts due to anatomic variations, shunt, movement, electrical noise

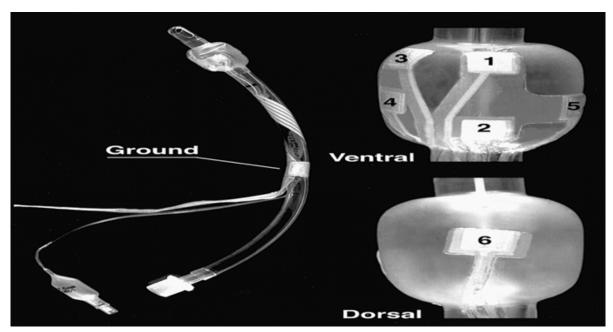


Figure I Model 6-3D endotracheal tube with six electrodes on the balloon and one ground electrode on the shaft. (Left ) The entire endotracheal tube showing the ground electrode on the shaft of the tube. (Top right ) Ventral surface of the balloon with electrodes I –5. (Bottom right ) Dorsal surface of the balloon with electrode 6. Electrode 3 is the current source. Electrode pairs I and 2, 4 and 5, and I and 6 give the three orthogonal signals.

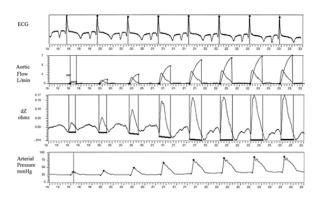


Figure 2 The top trace is the electrocardiogram derived from the endotracheal tube electrodes. The R wave is identified by a square and a vertical line. The second trace is aortic flow (liters/minute) measured by the transit time flow probe. The curve ending in a small square is the integral of transit time flow probe flow and gives stroke volume. The third trace is the change in impedance with time (DZ) measured in ohms. The trace between the two vertical lines corresponds to the aortic flow. The last trace is arterial pressure in millimeters mercury.11

#### Comparison of ECOM and TTFP Cardiac **Output Measurements**

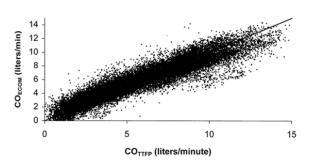


Figure 3 Plot of cardiac output calculated from the endotracheal cardiac output monitor system and the transit time flow probe-The relationship is linear.

## Limitations

The ECOM electrodes must stay in contact with the tracheal mucosa at all times and, if they lose this, there will be loss of signal. However, the ECOM algorithm can compensate for any loss of contact of one or two of the sensing electrodes. This can also occur in the event of high airway pressures or poorly inflated endotracheal cuff, when several electrodes may lift off the tracheal mucosal surface. The signal will return if these situations are rectified.

The ECOM system applies the current and measures the resulting voltage directly from the tracheal mucosa. Hence the signal obtained closely matches the shape and appearance of the ascending aortic flow waveform avoiding flow signals from the superior vena cava, right atrium, left atrium, and pulmonary artery. It designed to maximize the aortic signal and minimize impedance signals from other vascular structures. There is no interference with other electrical equipment on the patient (e.g., ECG, pacemaker, diathermy etc.)

#### Hemodynamic parameters measured and calculated by the **ECOM System:**

Cardiac output and derived parameters like cardiac index, stroke volume and systemic vascular resistance are all displayed on the

#### **Indications for ECOM:**

Indications for ECOMcardiac output monitoring	Influence on management	
Deranged cardiac function in shock states	Titration of fluids, inotropes	
Goal-directed resuscitation of complex shock states	and vasopressors	
Continuous monitoring following cardiac and non-cardiac surgery for high risk patients	Early intervention for altered cardiac function	
Protocol-driven management of hemodynamically unstable patients	Standardised management algorithms driven by cardiac output monitoring can support management decisions	

## Summary

The ECOM system provides a continuous real time display of CO derived from impedance measurements sensed by its array of electrodes. A linear relationship between the ECOM and CO is ensured between 0 and 15 l/min and bias is estimated at 0.15 l/min and the standard deviation being 1.34 l/min. The limits of agreement are -2.53 to 2.82 l/min.11,12

#### **Conclusion**

The ECOM system is able to measure stroke volume and CO accurately without being influenced by variations in systolic pressure, hematocrit, blood conductivity, r time. The signal corresponds to the shape and appearance of the waveform in the ascending aorta and the electrode design filters flow signals from the superior vena cava, right atrium, left atrium, and pulmonary artery. A variety of factors (institutional, device related, and patient specific) influence the selection of a cardiac output monitoring devices and clinicians need to understand the underlying principles and the inherent limitations of these devices. Endotracheal CO monitor is a promising technology that will give a quantum leap to critical patient management strategies in the future.

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