

Oesophageal Doppler Monitor (ODM) guided individualised goal directed fluid management (iGDFM) in surgery - a technical review

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Abstract

Oesophageal Doppler Monitoring (ODM) is an easy to use, accurate and minimally invasive methodology for the optimisation of Stroke Volume. The predominant ODM device (CardioQ-ODM™) utilises a nomogram incorporating age, weight and height which calibrates descending aortic blood flow velocity directly against total cardiac output as measured by thermodilution, thus negating the need to adjust for upper body flow for accurate measurement of aortic diameter. The velocity-time integral generated by each left ventricular contraction thus equates to Stroke Volume. Extensive validation studies have shown ODM to be reliable in studies comparing it to simultaneous measurements made with a pulmonary artery catheter and other reference techniques, both in absolute values and for following trends. ODM directly measures blood flow velocity in the descending thoracic aorta by the change in frequency (Doppler shift) of a fixed 4.02 MHz frequency ultrasound beam emitted from a probe placed in the oesophagus of the patient. The probe can be placed orally or nasally, the latter approach being more comfortable in awake patients, for example when undergoing surgery with epidural anaesthesia alone.

Fluid management can be thus individualised to the Frank-Starling curve to safely deliver intravenous fluids to optimise Stroke Volume. This is facilitated by the additional and unique flow-based parameters offered by ODM, such as Stroke Distance, Flow Time corrected, Mean Acceleration and Peak Velocity, not available on pressure-based systems. These parameters provide invaluable information on left ventricular preload, afterload and contractility which, when combined with Stroke Volume, are highly effective in accurately identifying haemodynamic changes and guiding appropriate fluid and vasoactive drug treatment. ODM is particularly useful in haemodynamically unstable patients where it can track changes and therapeutic responses on a beat-by-beat basis. By measuring flow in a major central vessel (the aorta), its accuracy is not affected by changes in peripheral arterial compliance, resistance and impedance.

Individualised Goal Directed Fluid Management (i-GDFM) of intraoperative surgical patients guided by oesophageal Doppler monitoring has been proven in 9 prospective randomised clinical trials to significantly reduce complications, unplanned ICU admissions, hospital stay and costs. Patient populations include major abdomino-pelvic, cardiothoracic, orthopaedic and trauma. Individualised Doppler Guided Fluid Management (i-DGFM) is the only outcome validated methodology. Intraoperative delivery of fluid has been suggested to be of greatest benefit in the first quarter of the operative period. It is evident that not only does the right amount of fluid need to be administered to optimise Stroke Volume but that the timing of its delivery is of equal importance. Three separate meta-analyses have confirmed the benefit of

perioperative flow-based ODM guided goal directed fluid management. The US Agency for Healthcare Research and Quality; the US Centers for Medicare & Medicaid Services; the UK NHS Centre for Evidence based Purchasing; a subsequent UK NHS Health Technology Assessment; and the British Consensus Guidelines on Intravenous Fluid Therapy for Adult Surgical Patients have all endorsed the use of ODM-guided i-GDFM to improve patient outcomes after surgery.

Being minimally invasive, the ODM has relatively few limitations. Caution should be applied in patients with pathology of the oropharynx/oesophagus and undue force should not be applied during probe insertion. This is similar to practices with naso- and orogastric tubes. ODM can now be performed in awake patients utilising the dedicated probes for this application. Accuracy in measuring absolute values of cardiac output may be affected by patients undergoing epidural anaesthesia or in those with body metrics outside the nomogram range, however fluid management can be achieved using Stroke Distance changes as these will still be reliable.

Introduction

Patients undergoing moderate and major surgery can carry a significant risk of mortality and morbidity. The level of this risk is dependent on a number of factors which include the length and severity of the surgery, the age of the patient and the presence of co-existing disease. A high percentage of patients (one report of 63%) have clinically significant fluid depletion pre-operatively.^{1, 2} In 1988 Shoemaker demonstrated that patients who were unable to maintain adequate levels of oxygen delivery (DO₂) had higher levels of morbidity and mortality than those patients who received cardiovascular optimisation whilst undergoing major surgery.³ Since that time there have been a number of randomised controlled trials that have demonstrated reductions in morbidity, length of hospital stay and costs of treatment following perioperative Stroke Volume optimisation with ODM in various surgical procedures.⁴⁻¹⁶

Haemodynamic Monitoring

It is the primary function of the cardiovascular system to maintain a stable metabolic state at both the organ and cellular level. This is achieved by the delivery of adequate amounts of oxygen and other substrates to meet the demands of metabolism and the removal of metabolic waste products from all the cells of the body. Oesophageal Doppler haemodynamic monitoring enables the assessment of left ventricular output contributing to the assessment of oxygen delivery.

In the 1930's it was suggested that the measurement of arterial blood pressure would not necessarily give accurate assessments of circulating blood volume¹⁷ and since the 1940's it has been acknowledged that cardiac output may be significantly decreased without an associated fall in arterial blood pressure.¹⁸ From the late 1950's there have been a number of clinical studies reported where the authors have described an association between increased perioperative cardiac output and increased survival following major surgery.¹⁹⁻²¹

It has been demonstrated that routine physical assessment

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alone, which includes the measurement of blood pressure, heart rate and urine output, often fails to reveal the true haemodynamic status of the cardiovascular compromised patient. Several recent studies have demonstrated that clinicians were only able to accurately predict the haemodynamic status in approximately 50% of the patients monitored using physical assessment and clinical findings alone.²²⁻²⁶ Many factors may contribute to these findings, most notably the physiological cardiovascular compensatory mechanisms. These mechanisms divert blood from the peripheral and splanchnic circulations in response to hypoperfusion and consequently often mask the true nature of blood flow. Typically, while a patient may have a significant decrease in circulating blood volume and an associated decrease in Cardiac Output (CO), the initial compensatory response of peripheral and splanchnic vasoconstriction will result in an increase in systemic vascular resistance and thus the patient will have a relatively normal arterial blood pressure. The effects of these compensatory mechanisms inhibit the ability of pressure-based monitoring systems to accurately assess the decrease in blood flow and oxygen delivery.

After the introduction of the pulmonary artery flotation catheter (PAC) in the 1970's it became the primary technique used for monitoring the haemodynamic status of the critically ill patient. Use of the PAC was widespread with estimates of approximately 1 million catheters used in the United States in 1996 alone.²⁷ Despite this extensive use, an improvement in patient outcomes has not been demonstrated in a number of randomised clinical trials. Additionally, recent evidence would suggest that an increase in morbidity and mortality is associated with the use of the PAC.^{28, 29} These unfavourable outcomes may be associated with a lack of knowledge regarding the optimal use of the PAC among clinicians, reports of which have been well documented.³⁰⁻³² These findings, along with the known risks associated with the use of pulmonary artery catheters, have resulted in declining use of the PAC and to clinicians seeking alternative less invasive haemodynamic monitoring techniques.

The minimally invasive nature of the Oesophageal Doppler Monitor makes it a suitable method for haemodynamic assessment in the operating room. In contrast, traditional Fick based techniques, such as dye or thermodilution, are more difficult to apply in the intraoperative period due to their extended set up time, complex calibration requirements and the potential for drug interaction with the chemical agents used for calibration. The ODM is considered useful for most moderate to high-risk surgical patients who would not otherwise warrant the risk of the insertion of a PAC, arterial or CVP line.

The Doppler Principle

On the 25th May 1842, Christian Doppler presented a paper in Salzburg, in which he proposed that the velocity of a moving object is proportional to the shift in reflected frequency in an optical wave of known frequency.³³ The principle also applies to sound waves. The received frequency of sound or light waves emitted by or reflected from a moving object is proportional to the relative velocity between the object and the receiver.

Doppler's hypothesis was formulated in relation to light emitted by double stars and it was not until 1845 that Buys Ballot confirmed it empirically using sound waves.³⁴ The principle was finally revised by Edwin Hubble in 1929. For example, the pitch of the sound waves emitted by a car (object) moving towards you (receiver) will increase in

frequency, while the pitch decreases in frequency as the car moves away from you. This is known as the Doppler effect. Today the Doppler effect is widely used to measure the speed of motor vehicles, predict atmospheric events, and calculate the distance of celestial bodies. Applying the Doppler principle to sound waves, technologies have been developed that can measure blood flow velocities and other related haemodynamic variables.³⁵

The Doppler equation is as follows:

$$f_D = \frac{2 v f_T \cos \theta}{c}$$

this can be rearranged to measure velocity as:

$$v = \frac{c f_D}{2 f_T \cos \theta}$$

where v is the velocity of the red blood cells, c is the speed of the ultrasound waves through body tissues (1540ms^{-1}), f_D is the Doppler frequency shift, f_T is the transmitted frequency of the ultrasound and $\cos \theta$ is the cosine of the angle of insonation

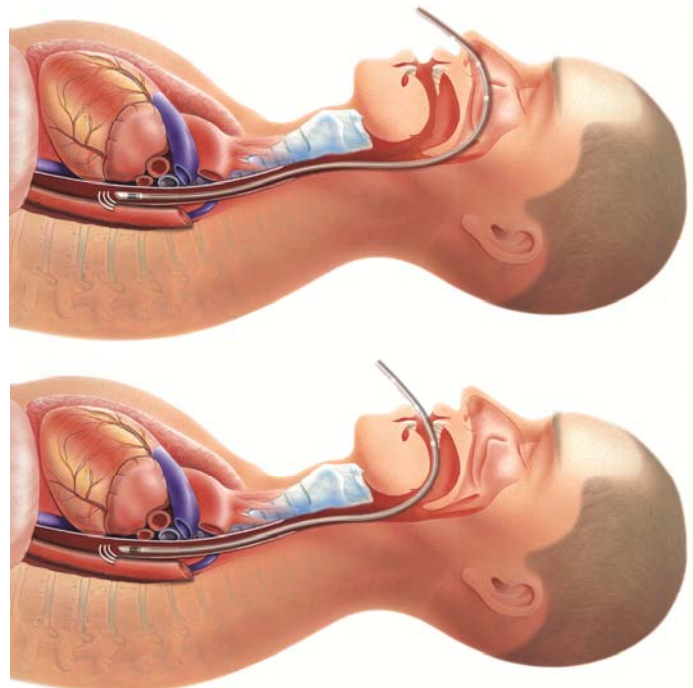


Figure 1. Nasal and oral positioning of oesophageal probe in relation to the aorta

between the sound beam axis and the direction of blood flow. The oesophagus is a convenient minimally invasive access point allowing ultrasound transmit and receive piezo-electric crystals to be located close to the descending aorta (Fig. 1). When using ultrasound Doppler to measure flow velocity, the angle of insonation is an important parameter (Figure 2). The angle of insonation is the angle between the Doppler ultrasound beam and the direction of blood flow in the vessel being examined. At large angles of insonation the errors in calculating blood flow velocity due to small errors in angle of measurement become unacceptably high (Figure 3). In fact ultrasound directed at 90° to the flow path does not undergo a

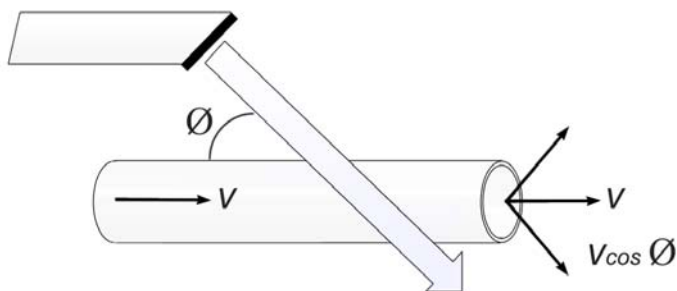


Figure 2. Derivation of the $V \cos \theta$ component of velocity away from the transducer

Doppler shift as the cosine of 90° is zero (Figure 3). At angles less than 45° , a 2° overestimation of the Doppler angle will give an overestimation of the true blood flow velocity of less than 5%. The percentage error resulting from a 2° uncertainty in the angle-of-insonation for angles from 0° to 90° is shown in Figure 3. The most commonly used ODM is designed to insonate at 45° with the probe shaft lying in the oesophagus and parallel to the aorta and it is generally recommended that devices use an angle of insonation of $<60^\circ$.

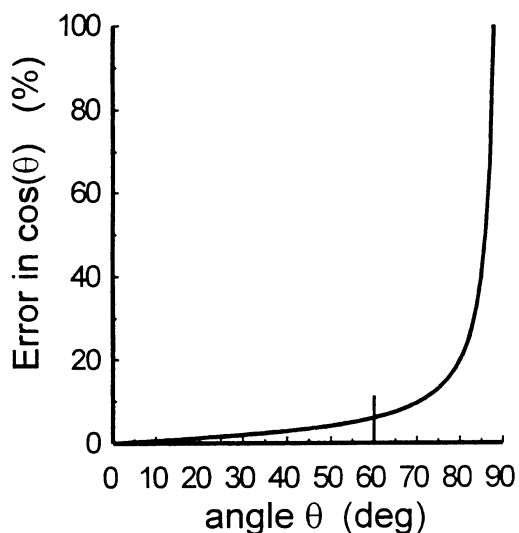


Figure 3. Potential for error due to a 2° uncertainty in the angle of insonation- redrawn from Cardiovascular and Haemodynamic Waveforms Explained - Oates 2008.³⁶

It was Satomura in 1957 and later Franklin in 1961 who were amongst the first to use the Doppler principle to measure red blood cell velocity in humans.^{37, 38} Since that time Doppler velocimetry has become a commonly used technique in many different medical applications and is utilised for the measurement of aortic and peripheral blood flow and intracardiac flow patterns.

During the intraoperative period anaesthetists can utilise oesophageal Doppler monitoring to measure descending thoracic aortic blood flow and derive cardiac output. The Oesophageal Doppler Monitor produces a velocity-time waveform that graphically displays the pulsatile blood flow in the descending thoracic aorta. This information can then be used as the basis for optimisation of Stroke Volume. ODM measurement requires the insertion of a lubricated probe into the

oesophagus of a patient. The angled tip of the probe contains ultrasound crystal transducers (piezo-electric) that are oriented towards the descending thoracic aorta (Figure 1). Red blood cells within the blood stream possess a higher density compared to that of the surrounding plasma. This density difference acts as an 'acoustic impedance mirror' to reflect the ultrasound emitted from the stationary probe.³⁶ The shift in the frequency of the reflected ultrasound waves returned from the moving red blood cells is converted into a 'beat to beat' real-time display of blood velocity against time. The Oesophageal Doppler Monitor waveform not only provides for the derivation of cardiac output, but also illustrates real-time changes in blood flow for qualitative assessment of preload, afterload, and contractility.



Figure 4. CardioQ-ODM™ Oesophageal Doppler Monitor.

The Oesophageal Doppler Monitor manufactured by Deltex Medical of Chichester UK (Figure 4) utilises a disposable probe, which emits continuous wave ultrasound. Ultrasound is emitted by one transducer in the probe tip while another transducer, also in the probe tip, continuously receives the ultrasound reflected by the moving blood (Figure 5). A proprietary nomogram is contained in the software of the ODM to convert the measured descending thoracic aortic blood flow velocity into total left ventricular Stroke Volume. The research for the nomogram was performed by Prof. Mervyn Singer of University

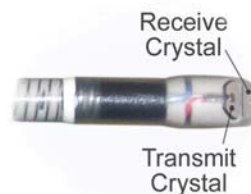


Figure 5. Receive and transmit ultrasound piezo-electric crystals

College London, and it uses the patient's age, weight, and height to generate a conversion factor and does not rely upon any other measurement.³⁹

There are some mis-understandings in the literature as to

how Oesophageal Doppler Monitors derive the total cardiac output whilst measuring flow only in the descending aorta.^{40,41} Approximately, 30% of blood flow leaves the aorta before this point to feed arteries supplying the heart, brain and limbs.⁴² Some reviews have made assumptions that ODM monitors make a 70:30 correction to derive the total cardiac output.^{40,41} This is not strictly accurate as the manufacturers have taken different approaches to derive the total cardiac output.^{42,43}

Table 1 Studies correlating ODM with alternative methodologies for Cardiac Output measurement – primarily the Pulmonary Artery Catheter (PAC)

Author	Compared to:	Population	n-data points	r	Bias/Precision
Shaw*	PAC	Critical Care	10/14	0.91 (r ²)	0.11/0.72
Seoudi*	PAC	Critical Care	15/187	0.89	NR
Di Corte*	PAC	Perioperative	34/160	0.77	NR
Tibby*	PAC	Paediatric	100/198	0.88 (r ²)	NR
Hersey*	Surface Echo	Paediatric	12	0.8	NR
Madan*	PAC	Critical Care	14/118	0.6	NR
Kincaid*	PAC	Critical Care	7/36	0.79	NR
Valtier*	PAC Fick	Critical Care	46/138 53	0.95/0.95	0.24/NR
LeFrant*	PAC	Critical Care	49/320	0.89	NR
Guzzetta*	PAC	Perioperative CABG	19/127	0.65	±2.9
Cuschien*	PAC	Critical Care	10	0.85 (r ²)	NR
	Fick			0.81 (r ²)	
	PAC vs Fick			0.72 (r ²)	
Cariou†	ABF vs	Critical Care	20/80	0.8	NR
Bernardin†	ABF vs	Critical Care	22/60	0.92	NR
Klein†	ABF vs	Perioperative	48/171	0.9	NR
Loik*	PAC	Critical Care	13/24	0.74 (r ²)	0.125/1.18
Catogni*	PAC	Preop CPB	14/40	N/A	-0.235
		Postop CPB	14/41		0.074
Nakatsuka*	PAC	Perioperative	7	0.96	0.296
		Liver transplant			
Sorohan*	PAC	Periop CPB	50/354	0.91	+0.2/0.5
Carrion*	Fick	Perioperative	15	0.72	NR
Carcellar*	PAC	Critical Care	15/176	0.78 (r ²)	0.196/2.188
Klotz*	PAC	Aortic Surgery			
		Pre-clamp	6/75	0.84	-0.96/NR
		Post-clamp	6/55	0.79	-1.51/NR
		Post-clamp	6/65	0.76	-1.47/NR
LeFrant*	PAC	Critical Care	11/85	0.93	NR
Belot*	PAC	Critical Care	44/127	0.86	NR
Muchada†	ABF vs	Perioperative	21/300	0.97	NR

*CardioQ Deltex Medical †Hemosonic Arrow International, ABF=Aortic Blood Flow, CPB=Cardiopulmonary bypass, NR=Not Reported, PAC=Pulmonary Artery Catheter, Table redrawn from Turner 2003⁶⁸.

The Hemosonic 100 device (formerly marketed by Arrow International, Reading, Pennsylvania, USA) utilizes M-mode echo to measure descending thoracic aortic diameter directly, and then uses a calibration factor to extrapolate this value to an estimate of total left ventricular Stroke Volume. M-mode measurement of the aortic diameter may potentially introduce large discrepancies, as an 8% error in diameter measurement during systole (e.g. 2mm variation from a true 25mm diameter) will generate an approximate 16% error in cross-sectional area. In comparison the CardioQ-ODM (Deltex Medical, Chichester, UK) incorporates a nomogram created by calibration of total left ventricular Stroke Volume as measured by the pulmonary artery catheter against descending aortic flow velocity and Stroke Distance as measured by the ODM. In this way any fluctuation of the aortic diameter is implicitly included in the calculation and as calibration is against total cardiac ejection there is no 70:30 proportioning required.⁴²

Accuracy of oesophageal Doppler monitoring

The reliability of the technology is widely supported with a large body of evidence demonstrating its accuracy for the measurement of cardiac output when compared to the “Clinical Standard” methodology of the PAC (Table 1).⁴⁴⁻⁶⁷

The Oesophageal Doppler Monitor measures blood flow velocity by highly accurate spectral analysis of the frequency-shifted (Doppler) signal. This real-time spectrum displays the distribution of red blood cell velocities at any given point in time. The ODM automatically traces the maximum velocity of the spectrum and, by calculating the area under this maximum-velocity curve during systole, a beat-to-beat value for Stroke

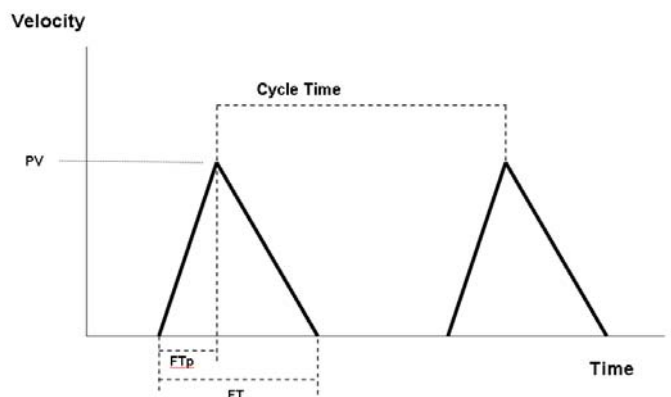


Figure 6. Stylised waveform of Velocity/Time display PV is the Peak Velocity measured during systole (in cm.s⁻¹). Flow Time (FT) is the time of systolic aortic blood flow (in ms). Flow Time to peak (FTp) is the time from the beginning of systole to the point when PV is detected (in ms). Cycle time is the time between identical successive points of the systolic waveform (in ms).

Distance (SD) is given, being the distance a column of blood moves in the aorta during systole. This is shown graphically in Figure 6.

The calibration nomogram built-in to the devices software is accessed by entering the patient’s age, weight and height. From this the ODM calculates the Stroke Volume (SV) using the measured SD. Since the machine automatically calculates the patient’s Heart Rate (HR) from the spectrum, it can also provide a beat-to-beat measurement of Cardiac Output (SV x HR). The nomogram was constructed by correlating ODM

measurements of SD against simultaneous SV measurements made by PAC thermodilution, over a wide range of patients of differing ethnic origins varying in age, weight and height.³⁹

The accuracy of the SD calculation is dependent on the precision of both the velocity measurement (typically $\pm 0.25 \text{ cm.s}^{-1}$) and the flow-time measurement ($\pm 3 \text{ ms}$). Ultimately, the accuracy of the spectral analysis is dictated by the accuracy of the quartz crystal clocks (typically $\pm 0.005\%$). Repeatability for a known spectrum is better than $\pm 1\%$ for the measurements of SD, PV, FT and HR, upon which all other calculations are made.

Conversion of the Stroke Distance into Stroke Volume is dependent on the accurate measurements made of PV, FT and FTp. The Heart Rate is also recorded by the monitor. The adult ODM nomogram contains extensive patient data for patients from 16 to 99 years of age, 30 to 150 kg in weight and 149 to 212 cm in height. A paediatric and adult ODM is also available where the nomogram also contains patient data for patients from 0 to 15 years of age, 3 to 60 kg in weight and 50 to 170 cm in height.⁴⁷ Once the nomogram is accessed, the ODM monitor will continuously display the selected haemodynamic parameters.

Probe Placement and Focusing

The insertion of a disposable oesophageal Doppler probe is similar to the placement of an oro/naso-gastric tube and usually takes an average of 2-3 minutes.^{69,70} Both clinicians and nursing staff, depending on the protocols of individual institutions, may perform this simple procedure. Patients are usually sedated, however recent publications have described a protocol for placement of oesophageal Doppler probes in awake patients using a new more flexible probe.^{71,72}

Probes are inserted orally to a depth of approximately 35-40 cm from the incisors, or nasally to a depth of approximately 40-45 cm from the nasal septum. In either case this will place the tip of the probe in the region of the 5th or 6th thoracic vertebrae. At this level the oesophagus is typically parallel with and approximately 1cm from the descending thoracic aorta. The probe is manipulated by the operator adjusting depth and rotational position by using small movements until the characteristic descending thoracic aortic waveform shape is visualised and the distinctive Doppler "whip crack" sound associated with aortic blood flow is heard. An Oesophageal Doppler Monitor is not a "hands free" continuous monitor and a degree of refocusing will be required, particularly if the patient is moved significantly. ODM monitors do however offer continuously available, minimally invasive data.

Focusing the ultrasound beam into a position that maximizes the waveform display is straightforward. The blood flow velocity profile in the descending thoracic aorta is distinctive in that it differs from that typically found in most vessels. Due to the proximity of the heart and its pulsatile pumping action the flow velocity profile in the descending aorta is referred to as "Plug flow". Unlike a typical parabolic velocity profile where the flow is largest in the centre of the vessel and decreases to zero at the wall, in plug flow the peak velocity of flow is present across a large part of the internal diameter of the vessel⁷³ (Figure 7). An optimal pitch and a sharp visual image can therefore be accepted as evidence of correct probe positioning. As a result focusing on the peak flow in the aorta is considerably easier than would at first be thought. Additionally the ultrasound beam is approximately 5 mm in width and the large aortic luminal diameter results in a "go/no go" focusing result. The probe may require occasional adjustment to ensure an

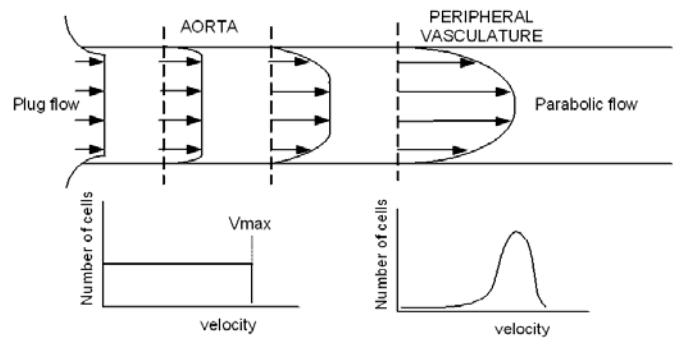


Figure 7. Vascular fluid dynamics – Peak Velocity flow profiles
Redrawn based on Caro et al⁷³

optimal signal. These adjustments typically take a few seconds.

Waveform and Parameters

The waveform provides valuable clinical information. The real-time waveform displayed on the monitor is the integral of the velocity of the blood passing the tip of the probe. A normal waveform will be triangular in shape (Figure 8). The peak of the triangle represents the Peak Velocity detected during systole. The upslope of the triangle depicts the acceleration of blood as it is ejected down the descending thoracic aorta at the beginning of systole. The area under the velocity-time waveform is the Stroke Distance (SD). SD is the distance a column of blood will travel down the aorta with each left ventricular contraction.

Any change in the left ventricular output will cause a proportional change in the descending thoracic aortic blood flow, which will in turn cause a change in the size and shape of the waveform. This assumes fixed proportionality of upper and lower body blood flow and that the aortic diameter remains constant during the period of the measurements. An increase in flow will result in the waveform increasing in size;

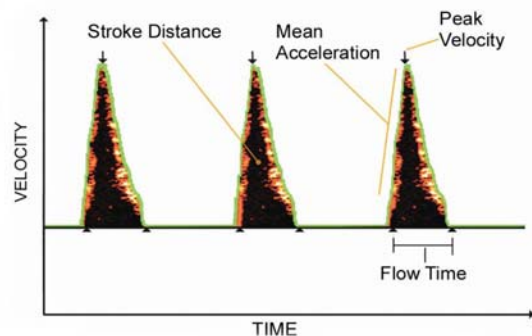


Figure 8. Waveform displaying Stroke Distance, Peak Velocity, Mean Acceleration and Flow Time measurements

consequently the area under the curve will also be increased. Conversely when left ventricular output is decreased, the blood flow decreases and the area under the curve will be less. This demonstrates a proportional relationship between Stroke Volume and the area under the systolic portion of the waveform. The velocity-time waveform display provides real-time information on changes in blood flow and left-ventricular function. Examination of the shape and size of the waveform

can be used to identify specific haemodynamic changes.

The ODM monitor has the facility to store 'snapshots' of patient data at various stages of therapy. A "snapshot" captures the image of the waveform and a display of all twelve of the monitors recorded parameters. Snapshots are a useful adjunct to the assessment of haemodynamic changes in the patient as treatment proceeds.

Oesophageal Doppler monitoring enables several measured and derived parameters to be assessed. Three of the primary measurements are, Stroke Distance, Flow Time and Peak Velocity. It is important to note that a true measurement of Stroke Volume and Cardiac Output does not exist. Therefore the absolute value of Stroke Volume and Cardiac Output may be subject to some imperfection. This holds true for all technologies providing Stroke Volume and Cardiac Output. However with the ODM it is extremely easy to identify a high, adequate or low cardiac output state. Trending of the derived parameters is of greater clinical value and is extremely useful in guiding therapies.⁴²

Stroke Volume and Cardiac Output

The area under the systolic portion of the waveform is defined as Stroke Distance (Figure 8). The Stroke Volume (SV) is calculated from the measured SD and a calibration constant derived from the nomogram. CO is then calculated by multiplying the SV by the Heart Rate (HR).

Preload

Preload is understood clinically as the degree of ventricular filling such that a low preload equates with under filling due to for example, hypovolaemia or an obstruction in the circulation (e.g. pulmonary embolus) whereas an excessive preload is associated with intravascular fluid overload. The width of the base of the waveform represents the systolic ejection time and is expressed as Flow Time corrected (FTc). This is the measured flow time corrected for heart rate in the same way as the Q-T interval of an ECG is corrected. This is achieved by using Bazett's equation.⁷⁴ The measured Flow Time (Figure 8) is divided by the square root of the cardiac cycle time thus adjusting the heart rate to 60 beats per minute. This results in one corrected cardiac cycle per second. Normally systole is approximately one-third of the cardiac cycle at a heart rate of 60 bpm, i.e. a third of a second. The FTc is displayed on the ODM in milliseconds (ms) and a number in the region of 330ms to 360ms is considered normal. However in patients undergoing regional or general anaesthesia there may be cases when their SVR is altered by the anaesthetic agents. Consequently caution should be observed in using FTc alone to assess hypovolaemia in surgical patients. Stroke Volume and Stroke Distance are the preferred index for guiding and monitoring haemodynamic optimisation.

The FTc is inversely correlated with the systemic vascular resistance and can be expressed as:

$$FTc \propto \frac{1}{SVR}$$

Therefore a narrow waveform base (<330ms) is an indicator of vasoconstriction, of which hypovolaemia is the commonest cause.

Studies have reported that changes in FTc are as good or better than pulmonary artery wedge (or occlusion) pressures (PAWP/PAOP) for indicating changes in preload.⁷⁵⁻⁷⁸ FTc has also been reported as useful in predicting fluid responsiveness

particularly when used in conjunction with other clinical information and CVP measurements.⁸⁰ Pulmonary artery occlusion pressure is still widely used to assess intravascular volume status, despite it having been shown to be a relatively poor indicator of preload.^{80,81}

Contractility

Contractility is the inotropic status of the myocardium. The Peak Velocity (PV) measured by the amplitude of the waveform, is a marker of contractility. Contractility decreases with age from typically 90 to 120 cm.s⁻¹ in a 20 year old to 30 to 60 cm.s⁻¹ in a 90 year old. In a hypo-contractile state such as left ventricular dysfunction, the waveform will appear dampened with decreased amplitude, resulting in a low PV. If the left ventricle is stimulated with a positive inotrope, the amplitude of the waveform will increase. Mean Acceleration, the average acceleration of blood from the start of systole to detected Peak Velocity, is also a marker of contractility and can be used to guide inotropic therapy.

Afterload

The afterload is the resistance or 'load' against which the heart has to eject blood. It affects the relationship between the width and amplitude of the waveform. An increase in afterload may be noted by a simultaneous reduction in both the FTc and Peak Velocity, resulting in a narrow waveform with decreased amplitude. This may be seen with any condition causing vasoconstriction, e.g. hypovolaemia, flow obstruction, excess vasopressor, hypotension or hypothermia. A reduction in afterload will result in an increase in amplitude (increased PV) and a widening at the base (increased FTc), as the left ventricle has less resistance to pump against. It is important to remember that any changes in the size of the waveform represents proportional changes in Stroke Distance and therefore Stroke Volume.

Individualised Goal Directed Fluid Management

Enhanced Recovery Programmes (ERP) are gaining increasing importance in surgery. The challenge to the modern clinician is the improvement of patient care combined with a reduction in hospital costs.⁸²⁻⁸⁴ Whilst many ERP are still in their infancy and yet to fully introduce the complete package of possible care management initiatives major advances have been made in numerous centres within Europe. Colorectal surgery has been a leading discipline in the implementation of fluid management as a major component of enhanced recovery and many have reported significant improvements in patient outcome and reductions in costs.⁸⁵⁻⁸⁸

Individualised Goal Directed Fluid Management (iGDFM) is based on optimisation of Stroke Volume. Thus in its simplest form iGDFM is achieved through the administration of fluid, guided by an algorithm to enhance Stroke Volume without the risk of fluid overload. Numerous authors have developed algorithms based on the measurement of Stroke Volume, central venous pressure and FTc.⁴⁻¹⁶

These algorithms utilise the Frank-Starling law, which established the relationship between left ventricular stroke volume and left ventricular end diastolic volume (Figure 9). The Frank-Starling law states that: "Within limits, the greater the heart muscle is stretched during filling, the greater will be the force of contraction and the greater the quantity of blood pumped into the receiving vessels".

When a fluid challenge (see Figure 9) is given to a patient who has a low left ventricular end diastolic volume a significant

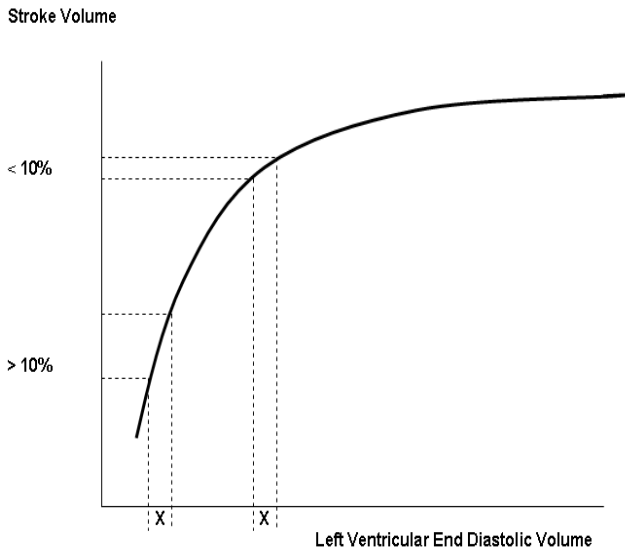


Figure 9. Frank-Starling Curve

rise in Stroke Volume (SV) is expected. Conversely if the patient has a high left ventricular end diastolic volume, little or no increase in stroke volume is expected for the same challenge volume. A typical perioperative Stroke Volume optimisation algorithm is shown in Figure 10. As described earlier the calculation of Stroke Volume by an ODM is based on the highly accurate measurement of Stroke Distance. As Stroke Volume is Stroke Distance multiplied by a constant provided by the built-in calibration nomogram of the ODM Stroke Distance and Stroke Volume are directly related. Stroke Distance is a useful parameter for guiding therapy in any circumstances where the patient parameters are outside the

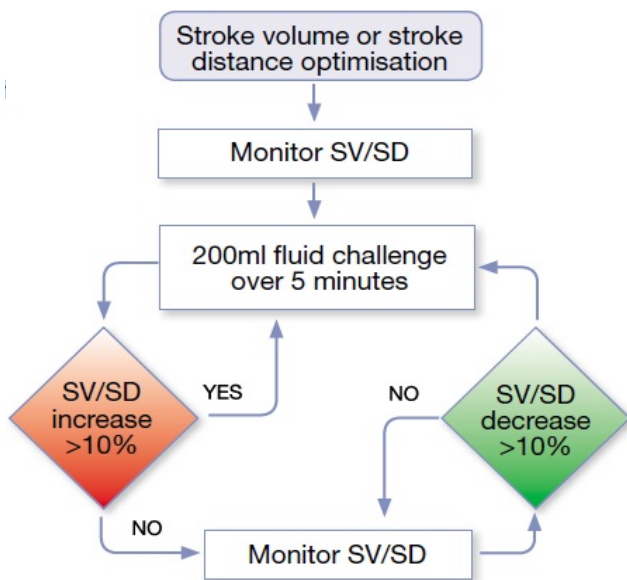


Figure 10. Typical perioperative Stroke Distance or Stroke Volume optimisation algorithm

nomogram limits (e.g. due to obesity), during periods of aortic cross clamping or where epidural use alters the upper/lower flow proportionality. Stroke Volume Index may also be used if preferred.

Patients are challenged with 200 ml of fluid over 5 minutes.

Following the challenge the result is monitored using an ODM. If the Stroke Distance or Stroke Volume increases 10% or more this indicates that the patient's left ventricular end diastolic volume is not optimal and that a further fluid challenge should be delivered.

Fluid challenges are repeated until the change in Stroke Distance or Stroke Volume as monitored by the ODM is less than 10%. A change of less than 10% either suggests the left ventricular end diastolic volume has been optimized or that the patient is losing fluid at a rate equal to or greater than that of infusion. Stroke Distance or Stroke Volume are then monitored and if a decrease of 10% or more is observed or if there is excessive fluid loss then a further fluid challenge is given. Fluid challenges given as part of perioperative Stroke Volume optimisation are additional to the normal fluid maintenance. A number of algorithms have utilised FTc as an indicator of hypovolaemia. Flow Time corrected (FTc) is often used as an indicator of hypovolaemia and fluid responsiveness, however during anaesthesia the vasodilatory effects of anaesthetic agents should be considered. Anaesthetics and other vasoplegic agents may create a decrease in left ventricular afterload such that the baseline FTc may be elevated above the normal range of 330ms to 360ms. Clinicians need to be aware of these effects and to take other parameters into consideration so as not to rely solely upon single parameters for fluid management.

It is recommended therefore that Stroke Distance, Stroke Volume or Stroke Volume index be used as the primary guiding parameter for individualised Doppler guided fluid management.

Clinical Application

Oesophageal Doppler monitoring may be used in many clinical settings, including the operating room, the intensive care unit, the post-anaesthesia recovery unit and the emergency or trauma department. Many patient groups will benefit from the use of oesophageal Doppler monitoring including those undergoing surgery, particularly moderate to high-risk patients, patients with large volume blood loss, the elderly, critically ill patients with an unstable haemodynamic status and those at risk of hypoperfusion or fluid overload due to inadequate left ventricular function.

In the context of ODM as applied to surgery, oesophageal Doppler monitoring has been proven to be beneficial in patients undergoing operations of longer than 1 hour duration and/or where the surgery involves entry into a body cavity or during orthopaedic surgery.⁴⁻¹⁶ Table 2 summarises the surgical criteria for use of ODM as reported in the literature. The ODM is also useful in detecting changes that might otherwise go unnoticed. The haemodynamic data can also

ASA Grade	Anaerobic threshold mL/min/kg	Body cavity exposed/penetrated	Orthopaedic surgery
2 or more ^{4,5,7,11,12,14,15}	<13.9 ⁸⁹	Yes ^{4,7,12,14,15}	Yes ^{5,11}

Table 2. Summary of criteria for use of ODM in surgery

guide the titration of vasopressors and inotropes as well as fluids.

In recent years Stroke Volume optimisation using the Oesophageal Doppler Monitor has been used to enhance DO₂ in the perioperative period. A significant contributor to tissue hypoxia is hypovolaemia. Any degree of hypovolaemia jeopardises oxygen transport and increases the risk for tissue injury or death.

Dubniks in a thesis entitled "Aspects of Fluid Therapy" has stated that hypovolaemia implies a reduced circulating blood volume and is one of the most common reasons of circulatory instability in surgical and critically ill patients.⁹⁰ Hypovolaemia can be absolute or relative. Absolute hypovolaemia is the result of haemorrhage, external or internal fluid losses. Internal fluid losses due to increased microvascular permeability is a common reason for hypovolaemia in critically ill patients suffering from sepsis, shock or systemic inflammatory reaction syndrome (SIRS). Relative hypovolaemia can result from vasoplegia, which can be caused by pharmacologically induced vasodilation during anaesthesia.⁹¹⁻⁹⁵ Dubnik further states that hypovolaemia leads to reduced venous return and inadequate cardiac preload, decreased cardiac output and insufficient oxygen delivery to the tissues. The decrease in circulating volume triggers activation of the baroreflex originating from stretch receptors in the central veins, the right atrium, in the carotid sinus and in the aortic arch. This leads to increase an in sympathetically mediated vasomotor tone in venous system aimed to preserve central blood volume, cardiac preload, cardiac output and systemic arterial pressure. Unloading of the arterial baroreceptors results in arterial vasoconstriction, which is selective to maintain perfusion in the vital organs such as the brain and the heart. Simultaneously, however, it will result in hypoperfusion in regional beds, such as the splanchnic area, skin, and muscle.^{90,96,97} The consequences of decreased oxygen delivery and impaired microcirculatory flow are tissue hypoxia and oxygen debt⁹⁸ which, if not corrected early,⁹⁹ leads to cell damage, organ dysfunction, multiple organ failure (MOF), and death.^{94, 95, 97, 100} It is important to also recognize that this covert tissue hypoxia in various organ beds may not be detected by conventional means as blood pressure may frequently remain stable.

Hamilton-Davies evaluated commonly measured cardiovascular parameters after removing 25% of circulating volume from healthy volunteers, and found no significant changes.¹⁰¹ Price reported similar findings in that a 40% reduction in splanchnic blood volume resulted from only a 10-15% reduction in blood volume of healthy volunteers. This further demonstrated that systemic circulation was maintained at the expense of the splanchnic bed.¹⁰² Fluid therapy aimed at restoring and maintaining circulating blood volume is, therefore, an important part of the complex circulatory management of perioperative patients.⁹⁰

ODM can be used to correct hypovolaemia in the patient who may be at risk. In an observational study, ODM has been shown to detect subsequent complications in the critically ill.¹⁰³

The goal of the study was to compare oesophageal Doppler parameters with standard haemodynamic variables (mean arterial pressure, central venous pressure, heart rate, arterial base-deficit, and urine output) used to predict complications after cardiac surgery. The results showed that ODM monitoring of Stroke Volume was the best marker for predicting postoperative complications during the initial postoperative period. Furthermore, those patients who developed complications received less volume in the first four hours postoperatively.

Several studies have demonstrated that intraoperative Stroke Volume optimisation guided by ODM significantly improves outcomes as evidenced by a decreased length of hospital stay ranging from 30-40%.⁴⁻¹⁶ These studies were performed in a wide range of surgical populations, such as cardiac, orthopaedic, colorectal and general surgery. All of the studies used similar algorithms to guide volume administration as previously described (Figure 10). These same studies have indicated that oesophageal Doppler guided intraoperative Stroke Volume optimisation resulted in improvement in patient outcome, as measured by reduction in post operative morbidity, reduction in time spent in the intensive care unit and overall hospital stay.⁴⁻¹⁶ These improvements also present the possibility of significant cost, equipment and resource savings.

Mythen et al in 1995 were the first to demonstrate that perioperative expansion of plasma volume using a colloid reduced the incidence of gastric hypoperfusion and directly associate this with improved outcome and reduction of major complications. Only 7% of patients in the flow-based oesophageal Doppler guided fluid management group showed gastric hypoperfusion as measured by tonometry by pHi <7.32 compared to 56% in the control group.⁴

Noblett et al has indicated that it is not necessarily the amount of fluid that is the determining factor in improved outcomes but the timing of the fluid optimisation within the operative period.¹⁵ This study also reported that no differences were found in the overall volume of fluid administered between their two study groups. However, the intervention group had higher FTc, SV, CO and CI at the end of the procedure; 46% of the fluid boluses, accounting for more than 50% of additional protocol volume given in the intervention group, were administered within the first quarter of the operating time. This suggests that the early delivery of fluid challenges in the intraoperative period, rather than the overall fluid volume was significant in optimising cardiovascular parameters.¹⁵

It has long been established that one of the initial responses to a reduction in circulating volume is the redirection of blood away from the splanchnic bed in favour of more vital organs. The gut mucosa is particularly susceptible to hypoxia. Gut mucosal hypoperfusion may lead to bacterial translocation, endotoxaemia and activation of inflammatory cascades, all of which may contribute to the systemic inflammatory response after surgery.^{104,105}

Peak systemic inflammatory cytokine (IL-6) levels have been shown to be reduced in fluid optimised patients, suggesting that individualised Doppler guided fluid management with early achievement of a higher Stroke Volume may have reduced the systemic inflammatory response to surgical trauma.¹⁵ Perioperative vasoconstrictors may affect splanchnic blood flow and oxygen supply/uptake ratios.¹⁰⁶⁻¹⁰⁸

Early achievement and maintenance of a normovolaemia reduces splanchnic hypoperfusion, reduces inflammatory mediator release and has beneficial effects on patient outcome. It is apparent that, although excess fluid administration may lead to complications, the inadequate treatment of occult hypovolaemia is also a factor in postoperative morbidity.

Studies validating the outcome benefit of individualised Doppler guided fluid management suggest that it is event that fluid management is essentially the right amount of fluid at the right time and that early in the intraoperative period provides the best outcomes⁴⁻¹⁶.

Haemodynamic monitoring is rarely available in the emergency or trauma department, yet this is where many critically ill patients are often admitted. Clinicians must rely on

assessment skills; along with all other clinical findings to assess the haemodynamic status of the patient and make appropriate treatment decisions. However, predicting haemodynamic status without the aid of additional monitoring is quite challenging and often inaccurate. The goal of a study done in a large urban emergency department was to determine how well physicians could accurately predict haemodynamic parameters. The study showed that the agreement between the physician and an ODM (CardioQ, Deltex Medical) was 48% for volume status, 50% for cardiac output, 39% for contractility, and 48% for afterload, requiring a change in therapy in approximately 40% of the cases.¹⁰⁹ Rivers et al reported that early goal-directed therapy in the emergency department, requiring haemodynamic monitoring, results in reduced morbidity, mortality, and hospital costs.⁹⁹ These findings suggest that oesophageal Doppler monitoring may also have a role to play in the emergency department, allowing for early and accurate assessment of patient haemodynamic status, leading to more appropriate and timely interventions.

Chytra et al have reported on trauma patients in a randomized controlled trial. Eighty multiple trauma patients with blood loss of more than 2,000 ml admitted to the intensive care unit (ICU) were randomly assigned to the protocol group with the ODM monitoring (Hemosonic 100, Arrow International Reading, Pennsylvania, USA) and to the control group. Fluid resuscitation in the Doppler group was guided for the first 12 hours of ICU stay according to the protocol based on data obtained by oesophageal Doppler, whereas control patients were managed conventionally. Fewer patients in the ODM group developed infectious complications 15 (18.8%) versus 28 (34.1%). ICU stay in the Doppler group was reduced from a median of 8.5 days to 7 days, and hospital stay was decreased from a median of 17.5 days to 14 days.¹⁶

Latterly as the clinical evidence from randomised clinical trials utilising ODM has expanded meta-analyses have been published.¹¹⁰⁻¹¹³ Two of these meta-analyses have specifically examined the evidence for use of ODM-guided fluid management in abdominal surgery.^{110,111}

Abbas and Hill analysed data from five randomized controlled trials which had recruited 420 patients undergoing major abdominal surgery. The patients had received either intravenous fluid treatment guided by ODM or fluid administration according to conventional parameters. Their analysis showed a reduced hospital stay in the ODM treatment group. Overall, there were fewer complications and ICU admissions, and less requirement for inotropes in the ODM treatment group. Return of normal gastro-intestinal function was also significantly faster in the ODM treatment group.¹¹⁰

Walsh et al investigated the potential of intraoperative fluid therapy guided by oesophageal Doppler to improve postoperative outcome. Their searched identified four randomised controlled trials comparing Doppler-guided intraoperative fluid management to standard practice in patients undergoing major abdominal surgery. Analysis of the 393 patients demonstrated fewer postoperative complications and shorter hospital stays where ODM was used to guide fluid management.¹¹¹

Phan et al have reported on meta-analysis of the use of ODM across all clinical settings. Nine clinical trials were included in the analysis seven using ODM in surgery and two in ICU. The primary outcome of this meta-analysis was length of hospital stay (LOS), defined as the number of postoperative days in an acute care hospital setting. Use of the ODM resulted in a significant reduction in LOS of 2.34 days. The group also analysed the colorectal surgery studies separately and concluded the LOS was reduced by 2.17 in this subset.

They concluded that using ODM to guide intraoperative IV fluid therapy increased the administration of intraoperative IV fluid, reduced length of hospital stay, time to resume full oral diet, and postoperative morbidity or complications.¹¹²

Hamilton et al have reported a meta-analysis of 7 studies totalling 618 patients. The meta-analysis showed that there was no significant difference in mortality between control and treatment groups. However there was a significant reduction in the length of hospital stay for patients receiving Doppler guided fluid management of 2.98 days.¹¹³

The National Health Service for the UK and the Centres for Medicare and Medicaid (CMS) for the USA have commissioned Health Technology Assessments (HTA) to examine the value of wider introduction of ODM-guided fluid management. In the USA the Emergency Care Research Institute (ECRI) reported their HTA findings to Agency for Healthcare Research and Quality (AHRQ) in 2007.¹¹⁴ The findings resulted in CMS revising their financial coverage of ultrasound diagnostic procedures to include ODM for ventilated patients in the ICU and surgical patients with a need for intraoperative fluid optimization.¹¹⁵

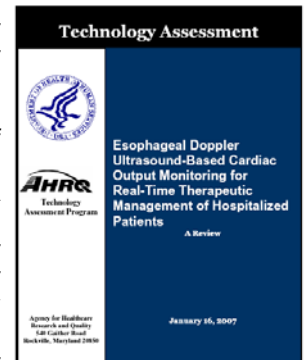


Figure 11. AHRQ HTA 2007
ODM for ventilated patients in the ICU and surgical patients with a need for intraoperative fluid optimization.¹¹⁵

The National Institute for Health Research (NIHR) released its systematic review of the randomised controlled trials published on the use of ODM in January 2009. The UK review was based on the systematic review conducted by the AHRQ with supplementary evidence from additional studies identified in a web based search. The effectiveness of ODM was compared with standard care, use of pulmonary artery catheters (PAC), pulse contour analysis monitoring and lithium or thermodilution cardiac monitoring. Data were extracted on mortality, length of stay overall and in critical care, complications and quality of life. The HTA compared ODM with conventional clinical assessment and reported that ODM is likely to be cost-effective since the initial cost of ODM is compensated by reduced complications and shorter length of stay in hospital. The report assessed the potential economic impact of ODM for surgical patients under the Quality Adjusted Life Year (QALY) methodology used by the National Institute for Health and Clinical Excellence (NICE). The analysis showed ODM to be both more effective and less costly under virtually every scenario modelled and that the NHS would need to spend between £642 and £4,441



Figure 12. NIHR HTA 2009

extra on each additional survivor of surgery before ODM would no longer be considered cost effective. The report concluded that available evidence suggested that the addition of ODM-guided fluid administration to CVP monitoring plus conventional assessment during surgery resulted in fewer major and total complications, a shorter length of hospital stay and

possibly fewer deaths. Pooled estimates for all outcomes showed a statistically significant difference in favour of the ODM group.¹¹⁶

In parallel with the preparation of the NIHR HTA report the NHS Centre for Evidence-based Purchasing (CEP) utilised the

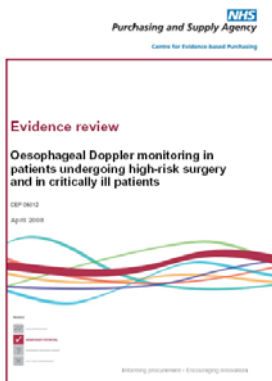


Figure 13. Centre for Evidence-based Purchasing report 2008

results to make recommendations for patients undergoing high risk surgery. CEP concluded that compared with CVP monitoring plus conventional clinical assessment, addition of ODM-guided fluid administration probably results in fewer deaths, fewer complications, and shorter length of hospital stay. CEP also concluded that the cost of ODM is likely to be offset by reductions in both complications and length of stay.¹¹⁷

The British Consensus Guidelines on Intravenous Therapy for Adult Surgical Patients (GIFTASUP) were released for dissemination to members of participating professional bodies late in 2008. The guidelines were developed on behalf of BAPEN Medical; the Association for Clinical Biochemistry; the Association of Surgeons of Great Britain and Ireland; the Society of Academic and Research Surgery; the Renal Association; and the Intensive Care Society. The guidelines represent the latest clinical thinking on fluid management and contain a number of recommendations for IV fluid management with oesophageal Doppler monitoring. Each recommendation has been given an evidence level grade from 1-5 in accordance with the Oxford Centre for Evidence-based Medicine Levels of Evidence, with a score of 1a representing the highest possible level of supporting clinical evidence. In the context of ODM recommendation 14 was based on level 1a evidence for abdominal surgery and level 1b for orthopaedic surgery. Recommendation 14 is solely based on evidence from studies using oesophageal Doppler monitoring.

CEP also concluded that the cost of ODM is likely to be offset by reductions in both complications and length of stay.¹¹⁷

The British Consensus Guidelines on Intravenous Therapy for Adult Surgical Patients (GIFTASUP) were released for



Figure 14. GIFTASUP 2009

Recommendation 13: *In patients undergoing abdominal surgery, intraoperative treatment with intravenous fluid to achieve an optimal value of stroke volume should be used where possible as this may reduce postoperative complication rates and duration of hospital stay.*

Further recommendations at Evidence level 1b also apply to optimisation of stroke volume. The beneficial effect is likely to relate to the early tailoring of fluid administration to the requirements of the individual patient rather than the blanket administration of extra fluid.

Recommendation 13: *In patients undergoing some forms of orthopaedic surgery, intraoperative treatment with intravenous fluid to achieve an optimal value of stroke volume should be used where possible as this may reduce postoperative complication rates and duration of hospital stay.*

Orthopaedic surgery: Evidence level 1b

Recommendation 14: *Patients undergoing non-elective major abdominal or orthopaedic surgery should receive intravenous fluid to achieve an optimal value of stroke volume during and for the first eight hours after surgery. This may be supplemented by a low dose dexepamine infusion.*

Evidence level 1b

The recommendations also call for wherever possible that preoperative or operative hypovolaemia should be diagnosed by flow-based measurements. The clinical context should also be taken into account as this will provide an important indication of whether hypovolaemia is possible or likely.¹¹⁸

Limitations

ODM can be used with both awake and sedated patients but as with the use of naso- or orogastric tubes, or any other tube/probe placed into the oesophagus, caution or special consideration should be given to patients where there may be an increased risk of bleeding, trauma or misplacing of the tube/probe. There may be circumstances where signal acquisition is not possible or very difficult due to coarctation of the aorta or during use of intra-aortic balloon pumps or in patients with thoracic aortic aneurysms. Users need to be aware that under epidural anaesthesia the proportionality of upper and lower body flow may alter, in these circumstances relative changes can be monitored using Stroke Distance. Additionally where patients fall outside the nomogram limits (i.e. obese) or during periods of cross clamping of the aorta; Stroke Distance (SD) changes can be used to guide fluid management.

Conclusions

An Oesophageal Doppler Monitor provides real-time flow-based measurements and visualisation of blood flow from the left side of the heart. Individualised Doppler guided fluid management (iDGFM) is a minimally invasive method of optimising Stroke Volume with a very low risk to the patient. The procedure requires significantly less insertion time than current standard methodologies, carries less risk of complications than invasive technologies requiring arterial catheter insertion. Current evidence from randomised clinical trials, meta-analyses and systematic reviews suggests that iDGFM to achieve Stroke Volume optimisation offers the potential for cost savings, decreased complications, decreased nursing and physician time, decreased hospital length of stay, and improved patient outcomes.

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References

- Ackland GL, Singh-Ranger D, Fox S, McClaskey B, Down JF, Farrar D, Sivaloganathan M, Mythen MG. Assessment of preoperative fluid depletion using bioimpedance analysis. *Br J Anaesth*. 2004 Jan; 92 (1):134-136.
- Mythen MG, Webb AR. Intraoperative gut mucosal hypoperfusion is associated with increased post-operative complications and cost. *Intensive Care Med*. 1994; 20:99-104.
- Shoemaker WC, Appel PL, Kram HB, Waxman K, Lee TS. Prospective trial of supranormal values of survivors as therapeutic goals in high-risk surgical patients. *Chest* 1988; 94(6): 1176-86.
- Mythen MG, Webb AR. Perioperative plasma volume expansion reduces the incidence of gastric mucosal hypoperfusion during cardiac surgery. *Arch Surg* 1995; 130: 423-429.
- Sinclair S, James S, Singer M. Intraoperative intravascular volume optimisation and length of stay after repair of proximal femoral fracture: randomised control trial. *BMJ* 1997; 315: 909-912.
- Gan TJ, Wakeling H, Hardman HM, et al. Intraoperative volume expansion guided by esophageal Doppler reduces the incidence of gastric mucosal hypoperfusion and may be associated with improved outcomes following surgery. *Anesthesiology* 1997; 87(3 suppl): A391.
- Gan TJ, Horacek A, Maroof M, El-Maolim H, Bell E, et al. Intraoperative volume expansion guided by esophageal Doppler improved postoperative outcome and shorten hospital stay. *Anesth Analg* 1999; 88(S1-424): S179.
- Shi C, Morse L, Downing LK, et al. Optimizing intraoperative volume management during coronary bypass surgery. *Anesthesiology* 2000; 93, no.3(suppl): A347.
- Saberi D, Caudwell I, McGloin H, Singer M. Proactive circulatory management in the first 4 hours post cardiac surgery: interim analysis of a nurse-led, oesophageal Doppler-guided protocol. *Intensive Care Med* 2000; 26, no.3 (suppl): S220.
- Gan TJ, Soppitt A, Maroof M, El-Maolim H, Robertson KM. Goal-directed intraoperative fluid administration reduces length of hospital stay after major surgery. *Anesthesiology* 2002; 97: 820-826.
- Venn R, Steele A, Richardson P, et al. Randomized controlled trial to investigate influence of the fluid challenge on duration of hospital stay and perioperative morbidity in patients with hip fractures. *Br J Anaesth* 2002; 88: 65-71.
- Conway DH, Mayall R, Abdul-Latif MS, et al. Randomised controlled trial investigating the influence of intravenous fluid titration using oesophageal Doppler monitoring during bowel surgery. *Anaesthesia* 2002; 57: 845-849.
- McKendry M, McGloin H, Saberi D, Caudwell L, Brady AR, Singer M. Randomised controlled trial assessing the impact of a nurse delivered, flow monitored protocol for optimisation of circulatory status after cardiac surgery. *BMJ* 2004; 329:258-262
- Wakeling HG, McFall MR, Jenkins CS, Woods WGA, Barclay GR, Fleming SC. Intraoperative oesophageal Doppler guided fluid management shortens postoperative hospital stay after major bowel surgery. *Br J Anaesth* 2005; 95: 634-642.
- Noblett SE, Snowden CP, Shenton BK, Horgan AF. Randomized clinical trial assessing the effect of Doppler-optimized fluid management on outcome after elective colorectal resection. *Br J Surg* 2006; 93: 1069-1076.
- Chytra I, Pradl R, Bosman R, Pelná P, Kasal E, Židková A. Esophageal Doppler-guided fluid management decreases blood lactate levels in multiple-trauma patients: a randomized controlled trial. *Crit Care* 2007; 11:R24.
- Blalock A. Experimental shock. The cause of the low blood pressure produced by muscle injury. *Archives of Surgery* (Chicago) 1930; 20:959-996.
- Blalock A. A consideration of the present status of the shock problem. *Surgery* 1943; 14: 487-508.
- Boyd AR, Tremblay RE, Spencer FC, Bahnson HT. Estimation of cardiac output soon after cardiac surgery with cardiopulmonary bypass. *Annals of Surgery* 1959; 150: 613-625.
- Clowes G, Del Guercio L. Circulatory response to trauma of surgical patients. *Metabolism* 1960; 9(1):67-81.
- Shoemaker WC, Montgomery ES, Kaplan E, Elwyn DH. Physiology patterns in surviving and non-surviving shock patients. Use of sequential cardiorespiratory parameters in defining criteria for therapeutic goals and early warning of death. *Archives of Surgery* 1973; 106: 630-636.
- Connors A, McCaffrey D, Gray B. Evaluation of right heart catheterization in the critically ill patient without acute myocardial infarction. *N Engl J Med* 1983; 308: 263-267.
- Eisenberg PR, Jaffe AS, Schuster DP. Clinical evaluation compared to pulmonary artery catheterization in the hemodynamic assessment of critically ill patients. *Crit Care Med* 1984; 12: 549-553.
- Tuchschmidt J, Sharma O. Impact of hemodynamic monitoring in a medical intensive care unit. *Crit Care Med* 1987; 15: 840-843.
- Stiengrub JS, Celoria G, Vickers-Lathi M, et al. Therapeutic impact of pulmonary artery catheterization in a medical/surgical ICU. *Chest* 1991; 99: 1451-1455.
- Connors AF Jr, Dawson NV, Shaw P, et al. Hemodynamic status in critically ill patients with and without acute heart disease. *Chest* 1990; 98:1200-1206.
- Dalen JP, Bone RC. Is it time to pull the pulmonary artery catheter? *JAMA* 1996; 276: 916-918.
- Connors AF Jr, Speroff T, Dawson NV, et al. The effectiveness of right heart catheterization in the initial care of the critically ill patients. *JAMA* 1996; 276: 889-897.
- Polaczky CA, Rohde LE, Golman L, et al. The effectiveness of right heart catheterization and cardiac complications in patients undergoing non cardiac surgery; an observational study. *JAMA* 2001; 286(3): 309-314.
- Iberty JT, Daily EK, Leibowitz AB, et al. Assessment of critical care nurses knowledge of the pulmonary artery catheter. *Crit Care Med* 1994; 22: 1674-1678.
- Gnaegi A, Peihl F, Perret C. Intensive care physicians' knowledge of right-heart catheterization at the bedside; time to act. *Crit Care Med* 1997; 25: 213-220.
- Iberty TJ, Fischer EP, Leibowitz AB, et al. A multicenter study of physicians' knowledge of the pulmonary artery catheter. *JAMA* 1990; 264: 2928-2932.
- Doppler JC, Ueber das farbige Licht der Doppelsterne und einiger anderer Gestirne des Himmels. *Abhandl Konigl Bohm Ges Ser* 1843; 2:465-482.
- Ballot CHDB, Akutische Versuche auf der Niederlandischen Eisenbahn nebst gelegentlichen Bemerkungen zur Theorie des Hrn. Prof. Doppler. *Pogg Ann* 1845; B66: 321-351.
- Payen DM. Oesophageal Doppler monitoring: history, physics principles, and clinical applications. *International Proceedings Journal: Int Care Med* 1994; 1:3-9.
- Oates C, Cardiovascular Haemodynamics and Doppler Waveforms Explained. Cambridge University Press 2008 ISBN 9780521734.
- Satomura S. (1957) Ultrasonic Doppler method for the inspection of cardiac function, *J Acoust Soc Am* 29, 1181-1185.
- Franklin D, Schlegel W, Rushmer R. Blood flow measured by Doppler ultrasound frequency shift of back-scattered ultrasound. *Science* 1961; 134: 546-565.
- Singer M. Continuous Haemodynamic Monitoring by Oesophageal Doppler. *A dissertation presented to the University of London in application for the degree of Doctor of Medicine*. April 1989.
- Jonas M, Fennel J, Brudney CS. Haemodynamic optimisation of the surgical patient revisited. *Anaesthesia International* Spring 2008; 2: No 1.
- Venn R, Rhodes A, Bennett ED. The esophageal Doppler. In: Vincent JL, (ed). Yearbook of Intensive Care and Emergency Medicine. Berlin: Springer-Verlag. 1999: 482-493.
- Singer M. Oesophageal Doppler. *Curr Opin Crit Care*. 2009; 15:244-248.
- Schober P, Loer SA, Schwarte LA. Perioperative Hemodynamic Monitoring with Transesophageal Doppler Technology. *International Anaesthesia Research Society* 2009; 2:340-353.
- Shaw AD, Weavind LM, Palmly L. Comparison of thermodilution, esophageal Doppler and transesophageal echocardiograph data in the hemodynamic assessment of critically ill cancer patients. *Crit Care Med* 2000; 28(12 suppl): A73.
- Seoudi H, Perkal M, Hanrahan A, Angood PB. The esophageal Doppler monitor in mechanically ventilated surgical patients; does it work? *J Trauma* 1999; 47(6): 1171. abstract.
- Dicorte CJ, Latham P, Greilich P, et al. Esophageal Doppler Monitor determinations of cardiac output and preload during cardiac operations. *Ann Thorac Surg* 2000; 69:1782-1786.
- Tibby SM, Hatherill M, Murdoch IA. Use of transoesophageal Doppler ultrasonography in ventilated paediatric patients: Derivation of cardiac output. *Crit Care Med* 2000; 28: 2045-2050.
- Hersey SL, Taylor M, Brock J. Trans-esophageal Doppler measurement of cardiac output in children. *Anesthesiology* 1999; 91(3suppl) A1312.
- Madan AK, HyBaretta VV, Shaghayegh A, et al. Esophageal Doppler ultrasound monitor versus pulmonary artery catheter in the hemodynamic management of critically ill surgical patients. *J Trauma Injury Infect Crit Care* 1999; 46(4): 807-811.
- Kincaid H, Fly M, Chang M. Noninvasive measurements of preload using esophageal Doppler are superior to pressure-based estimates in critically ill patients. *Crit Care Med* 1999; 27(1): A111.
- Valtier B, Chollet BP, Belot JP, et al. Noninvasive monitoring of cardiac output in critically ill patients using transesophageal Doppler. *Am J Respir Crit Care Med* 1998; 158: 77-83.
- LeFrant JY, Bruele P, Aya AGM, et al. Training is required to improve the reliability of esophageal Doppler to measure cardiac output in critically ill patients. *Intensive Care Med* 1998; 24: 347-352.
- Guzzetta N, Ramsay J, Baily JM, et al. Clinical evaluation of the esophageal Doppler monitor for continuous cardiac output monitoring. Poster presentation at SCA meeting, 1998.
- Cuschieri J, River M, Caruso J, et al. A comparison of transesophageal Doppler, thermodilution and Fick cardiac output measurements in critically ill patients. *Crit Care Med* 1998; 26(1 suppl)A62.
- Cariou A, Monchi M, Joly IM, Bellenfant F, Claessens YEI. Noninvasive hemodynamic monitoring by aortic blood flow determination: evaluation of the Somotec DYNEMO 300 system. *Crit Care Med* 1998; 26(12): 2066-2072.
- Bernardin G, Tiger F, Fouche R, Mattei M. Continuous non-invasive measurement of aortic blood flow in critically ill patients with a new esophageal echo-Doppler system. *J Crit Care* 1998; 13(4): 177-183.
- Klein G, Emmerich M, Maisch O, Dummmler R. Clinical evaluation of non-invasive monitoring Aortic Blood Flow (ABF) by a transesophageal echo-Doppler device. *Anesthesiology* 1998; 89: 3A, A953.
- Loik P, Andrews A, Folk L, et al. Comparison of esophageal Doppler and thermal dilution cardiac output in critical care patients. *Respir Care* 1997; 12(11): 1073.
- Catogni P, Provenchere S, Philip I, Daccache P, Depoix P, et al. Does esophageal Doppler accurately assess hemodynamic parameters in patients with aortic stenosis? *Anesthesiology* 1997; 87(3 suppl) A339.
- Nakatsuka M, Fisher RA, Ham JM, Seaman DS, Posner MP. Validation of the esophageal Doppler cardiac function monitor with the standard thermodilution method during liver transplantation. *Anesth Analg* 1997; 84: SCA51.
- Sorohan J, Gilbert HC, Atwell D, et al. Dual-center validation of an esophageal Doppler continuous cardiac output monitor. *Crit Care Med* 1997; 25(1 Suppl)A50, no27.
- Carrion MS, Polo A, Vazquez E, Prieto M, de las Nieves HV. Comparison of cardiac output measurement techniques: Doppler versus Fick method. *Br J Anaesth* 1996; 76(2Suppl) A42.
- Carceller J, Langunilla J, Rídriguez J, Compana O, Rodríguez V, Alvarez J. Clinical evaluation of esophageal Doppler monitoring (ODMH) versus continuous cardiac output/SvO₂ monitoring system in cardiac surgery patients. *Intensive Care Med* 1995; 21 (1 suppl): 885.

64. Klotz K-F, Klingsick S, Singer M et al. Continuous measurement of cardiac output during aortic cross-clamping by the oesophageal Doppler monitor ODM 1. *Br J Anaesth* 1995; 74:655-660.
65. LeFrant JY, Ayal G, de la Coussaye JE, Bassout B, Auffray JP, Eledjam JJ. Comparison of cardiac output measured by esophageal Doppler vs thermodilution. *Intensive Care Med* 1992; 18 (2 suppl): S177. P238.
66. Belot JP, Valtier B, de la Coussaye JE, Morrín D, Payen D. Continuous estimation of cardiac output in critically ill mechanically ventilated patients by a new transesophageal Doppler probe. *Intensive Care Med* 1992; 18 (2 suppl) S178. P211.
67. Muchcada R, Cathignol D, Lavender B, Lamazou J, Haro D. Aortic blood flow measurement. *Am J Non-invasive Cardiol* 1998; 2: 24-31.
68. Turner M A, Doppler-based Hemodynamic Monitoring. *AACN Clinical Issues* Vol. 14, No. 2, 220-231.
69. Singer M, Clarke J, Bennett ED. Continuous hemodynamic monitoring by esophageal Doppler. *Crit Care Med* 1989; 17:447-452.
70. Gan TJ. The esophageal Doppler as an alternative to the pulmonary artery catheter. *Curr Opin Crit Care* 2000; 6: 214-221.
71. Atlas G, Morr T. Placement of the esophageal Doppler ultrasound probe in awake patients. *Chest* 2001; 119:319.
72. English JD, Moppett IK. Feasibility of Performing Transoesophageal Doppler Measurements in Awake, Unpremedicated Healthy Volunteers. Poster presented at the Association of Anaesthetists of Great Britain and Ireland. Cardiff 2004.
73. Caro CG, Pedley TJ, Schroter RC and Seed WA. *The Mechanics of the Circulation*. Oxford University Press, UK, 1978.
74. Bazett MC. An analysis of the time-relations of electrocardiograms. *Heart* 1920; 7:355-364.
75. Seoudi II, Perkal M, Hanrahan A, Angood PB. The esophageal Doppler monitor in mechanically ventilated surgical patients; does it work? *J Trauma* 1999; 17(6):1171 Abstract.
76. Dicorte CJ, Latham P, et al. Esophageal Doppler monitor determinations of cardiac output and preload during cardiac operations. *Ann Thoracic Surg* 2000; 69:1782-1786.
77. Madan AK, UyBarreta VV, Shaghayegh A, et al. Esophageal Doppler ultrasound monitor versus pulmonary artery catheter in the hemodynamic management of critically ill surgical patients. *J Trauma Injury Infect Crit Care* 1999; 46(4):807-811.
78. Kincaid H, Fly M, Chang M. Noninvasive measurements of preload using esophageal Doppler are superior to pressure-based estimates in critically injured patients. *Crit Care Med* 1999; 27(1): A111.
79. Lee J-H, Kim J-T, Yoon S Z, Lim Y-J, Jeon Y, Bahk J-H and Kim C S. Evaluation of corrected flow time in oesophageal Doppler as a predictor of fluid responsiveness. *Br J Anaesth* 2007; 99:343-348.
80. Calvin JE, Driedger AA, Sibbald WJ. Does the pulmonary capillary wedge pressure predict left ventricular preload in critically ill patients. *Crit Care Med* 1981; 9:437-443.
81. Raper R, Sibbald WJ, Misled by the wedge? The Swann-Ganz catheter and left ventricular preload. *Chest* 1986; 89:427-434.
82. Fearon KCH, Ljungqvist O, Von Meyenfeldt M, Revhaug A, Dejong CHC, Lassen K, Nygren J, Hausel J, Soop M, Andersen J, Kehlet H. Enhanced recovery after surgery: A consensus review of clinical care for patients undergoing colonic resection *Clinical Nutrition* 2005; 24:466-477.
83. Modernising Care for Patients Undergoing Major Surgery-Improving Patient Outcomes and Increasing Clinical Efficiency. A report by the *Improving Surgical Outcomes Group* June 2005.
84. Modernising Care for Patients Undergoing Major Surgery-Implementation Guide. A report by the *Improving Surgical Outcomes Group*. June 2007.
85. Improving Surgery. *The Clinical Services Journal* September 2007; 90-91.
86. Parker M, Recovery times slashed by three weeks. *Health Director* February 2008.
87. Windsor A, Improving surgical outcomes, reducing length of stay. *Health Director*. October 2007.
88. Wakeling H & Jenkins C. Oesophageal Doppler monitoring saves lives. *The Clinical Services Journal*. June 2009.
89. Older P., Hall A., Hader R., Cardiopulmonary exercise testing as a screening test for perioperative management of major surgery in the elderly. *Chest* 1999; 116: 355-362.
90. Dubniks M. Aspects of Fluid Therapy-An experimental study of the effects of systemic inflammation, microvascular permeability, blood pressure and plasma volume. Lund University, Faculty of Medicine Doctoral Dissertation Series, 2008:23 ISSN 1652-8220/ISBN 978-91-85897-76-6.
91. Mythen MG, Salmon JB, Webb AR. The rational administration of colloids. *Blood Rev* 1993; 7(4): 223-228.
92. Imm A, Carlson RW. Fluid resuscitation in circulatory shock. *Crit Care Clin* 1993; 9:313-333.
93. Fleck A, Raines G, Hawker F, Trotter J, Wallace PI, Ledingham IM, Calman KC. Increased vascular permeability: a major cause of hypoalbuminaemia in disease and injury. *Lancet* 1985; 1:781-783.
94. Groeneveld AB, Bronsveld W, Thijs LG. Haemodynamic determinants of mortality in human septic shock. *Surgery* 1986; 99:140-153.
95. Morisaki H, Sibbald WJ Issues in colloid and transfusion therapy of sepsis. In: Vincent JL editor. Yearbook of intensive care and emergency medicine. Berlin. Springer. 1993;357-372.
96. Ba ZF, Wang P, Koo DJ, Cioffi WG, Bland KI, Chaudry IH. Alterations in tissue oxygen consumption and extraction after trauma and hemorrhagic shock. *Crit Care Med* 2000; 8:2837-2842.
97. Pastores SM, Katz DP, Kvetan V. Splanchnic ischemia and gut mucosal injury in sepsis and the multiple organ dysfunction syndrome. *Am J Gastroenterol* 1996; 91:1697-1710.
98. Shoemaker WC, Appel PL, Kram HB. Role of oxygen debt in the development of organ failure, sepsis, and death in high risk surgical patients. *Chest* 1992; 102:208-215.
99. Rivers E, Nguyen B, Havstad S, et al. Early goal-directed therapy in the treatment of severe sepsis and septic shock. *N Eng J Med* 2001; 345:1368-1377.
100. Shoemaker WC, Appel PL, Kram HB. Tissue oxygen debt as a determinant of lethal and non lethal postoperative organ failure. *Crit Care Med* 1988; 16:1117-1120.
101. Hamilton-Davies C, Mythen MG, Salmon JB, et al. Comparison of commonly used clinical indicators of hypovolaemia with gastrointestinal tonometry. *Int Care Med* 1997; 23: 276-281.
102. Price HL, Deutsch S, Marshall BE, et al. Haemodynamic and metabolic effects of haemorrhage in man with particular reference to the splanchnic circulation. *Circ Res* 1966; 18:469-474.
103. Poeze M, Ramsey G, Greve JM, Singer M. Prediction of postoperative cardiac surgical morbidity and organ failure within 4 hours of intensive care admission using esophageal Doppler ultrasonography. *Crit Care Med* 1999; 27:1288-1294.
104. Fiddian-Green RG. Splanchnic ischaemia and multiple organ failure in the critically ill. *Ann R Coll Surg Engl* 1988; 70:128-134.
105. Deitch EA. The role of intestinal barrier failure and bacterial translocation in the development of systemic infection and multiple organ failure. *Arch Surg* 1990; 125:403-404.
106. Jakob SM, Takala J. Splanchnic hemodynamics in critical illness. *Curr Opin Crit Care* 2000; 6: 123-129.
107. Priebe HJ, Noldge GFE, Armbruster K, Geiger K. Differential effects of dobutamine, dopamine, and noradrenaline on splanchnic haemodynamics and oxygenation in the pig. *Acta Anaesthesiol Scand* 1995; 39:1088-1096.
108. De Backer D, Vincent JL. Pharmacological modulation of splanchnic blood flow. *Curr Opin Crit Care* 1998; 4:104-110.
109. Urrunaga JJ, Rivers E, Mullen M, et al. Hemodynamic evaluation of the critically ill in the emergency department: a comparison of clinical impression versus transesophageal Doppler measurement. *Ann Emerg Med* 1999; 34(4): A176.
110. Abbas SM, Hill AG. Systematic review of the literature for the use of oesophageal Doppler monitor for fluid replacement in major abdominal surgery. *Anaesthesia* 2008; 63:44-51.
111. Walsh SR, Tang T, Bass S, Gaunt ME. Doppler-guided intra-operative fluid management during major abdominal surgery: systematic review and metaanalysis. *Int J Clin Pract* 2008; 62:466-470.
112. Phan TD, Ismail H, Alex Heriot AG, Ho KM. Improving Perioperative Outcomes: Fluid Optimization with the Esophageal Doppler Monitor, a Metaanalysis and Review. *J Am Coll Surg* 2008; 207(6): 935-941
113. Hamilton M, Grocott PW, Mythen M, Bennett. Does oesophageal Doppler guided goal directed therapy reduce surgical mortality and length of stay? *Intensive Care Med* 2006; 32(1) 0442.
114. Esophageal Doppler Ultrasound-Based Cardiac Output Monitoring for Real-Time Therapeutic Management of Hospitalized Patients - A Review. Agency For Health Research and Quality(AHRQ) January 16, 2007. Available at: www.cms.hhs.gov/mcd/viewtechassess.asp?where=index&tid=45
115. Decision Memo for Ultrasound Diagnostic Procedures (CAG-00309R). Centers for Medicare and Medicaid Services (CMS) May 22, 2007. Available at: www.cms.hhs.gov/mcd/viewdecisionmemo.asp?id=196
116. Mowatt G, Houston G, Hernández R, de Verteuil R, Fraser C, Cuthbertson B and Vale L. Systematic review of the clinical effectiveness and cost-effectiveness of oesophageal Doppler monitoring in critically ill and high-risk patients. *Health Technology Assessment* 2009; Vol.13: No.7.
117. Evidence review: Oesophageal Doppler monitoring in patients undergoing high-risk surgery and in critically ill patients. CEP08012. NHS Purchasing and Supply Agency; 2008. Also available from: www.pasa.nhs.uk/PASAWeb/NHSprocurement/CEP
118. Powell-Tuck J, Gosling P, Lobo D N, Allison S P, Carlson G L, Gore M, Lewington A J, Pearse R M and Mythen M G. British Consensus Guidelines on Intravenous Fluid Therapy for Adult Surgical Patients. GIFTASUP. On behalf of BAPEN Medical-core group of BAPEN, the Association for Clinical Biochemistry, the Association of Surgeons of Great Britain and Ireland, the Society of Academic and Research Surgery, the Renal Association and the Intensive care Society. <http://www.asgbi.org.uk>

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