

## Postoperative fluid management

Selami Ilgaz Kayilioglu, Tolga Dinc, Isa Sozen, Akin Bostanoglu, Mukerrem Cete, Faruk Coskun

Selami Ilgaz Kayilioglu, Tolga Dinc, Isa Sozen, Akin Bostanoglu, Mukerrem Cete, Faruk Coskun, Ankara Numune Training and Research Hospital, Department of General Surgery, 06100 Altindag, Ankara, Turkey

**Author contributions:** Kayilioglu SI, Dinc T and Coskun F designed the review; Kayilioglu SI, Dinc T, Sozen I, Bostanoglu A and Cete M conducted the literature review; Kayilioglu SI, Dinc T and Coskun F wrote the article; Cete M and Coskun F supervised all the process.

**Conflict-of-interest statement:** Authors have no conflict of interest.

**Open-Access:** This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

**Correspondence to:** Faruk Coskun, Professor of Surgery, Ankara Numune Training and Research Hospital, Department of General Surgery, Anafartalar Mah. Talatpasa Bul. No. 5, 06100 Altindag, Ankara, Turkey. [farukcoskun@mynet.com](mailto:farukcoskun@mynet.com)  
Telephone: +90-312-5085075  
Fax: +90-312-3103460

Received: November 28, 2014

Peer-review started: November 29, 2014

First decision: January 20, 2015

Revised: February 12, 2015

Accepted: April 1, 2015

Article in press: April 7, 2015

Published online: August 4, 2015

### Abstract

Postoperative care units are run by an anesthesiologist or a surgeon, or a team formed of both. Management of postoperative fluid therapy should be done considering both patients' status and intraoperative events. Types

of the fluids, amount of the fluid given and timing of the administration are the main topics that determine the fluid management strategy. The main goal of fluid resuscitation is to provide adequate tissue perfusion without harming the patient. The endothelial glycocalyx dysfunction and fluid shift to extracellular compartment should be considered wisely. Fluid management must be done based on patient's body fluid status. Patients who are responsive to fluids can benefit from fluid resuscitation, whereas patients who are not fluid responsive are more likely to suffer complications of over-hydration. Therefore, common use of central venous pressure measurement, which is proved to be inefficient to predict fluid responsiveness, should be avoided. Goal directed strategy is the most rational approach to assess the patient and maintain optimum fluid balance. However, accessible and applicable monitoring tools for determining patient's actual fluid need should be further studied and universalized. The debate around colloids and crystalloids should also be considered with goal directed therapies. Advantages and disadvantages of each solution must be evaluated with the patient's specific condition.

**Key words:** Body fluids; Body fluid compartments; Fluid therapy; Intensive care; Postoperative care

© **The Author(s) 2015.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Core tip:** Types of the fluids, amount of the fluid given and timing of the administration are the main topics that determine the fluid management strategy. Assessment of the patient's responsiveness to fluid resuscitation should determine the need of extra volume. Due to lack of evidence that supports central venous pressure (CVP) as an indicator of body fluid needs, we should not make our fluid resuscitation decisions based on CVP levels. On the other hand dynamic measures can be used to determine patient's fluid status. Among all fluid management strategies, goal directed strategy is the most rational approach to maintain optimum fluid balance.

Kayilioglu SI, Dinc T, Sozen I, Bostanoglu A, Cete M, Coskun F. Postoperative fluid management. *World J Crit Care Med* 2015; 4(3): 192-201 Available from: URL: <http://www.wjgnet.com/2220-3141/full/v4/i3/192.htm> DOI: <http://dx.doi.org/10.5492/wjccm.v4.i3.192>

## POSTOPERATIVE FLUID MANAGEMENT

Fluid management is an important part of overall surgical therapy. Proper administration of fluids is critical, especially in patients who undergo major surgeries such as emergency laparotomies, bowel resections and hepatectomy procedures. Body fluid composition may change in minutes or hours, resulting in impaired wound healing and homeostasis. Briefly, choice of strategy in intraoperative and postoperative fluid management may be significant.

We will examine different postoperative fluid management strategies in this review. Postoperative management of patients, who undergo surgery, is carried out by intensive care specialists, anesthesiologists and general surgeons in postoperative care units, in all over the world<sup>[1]</sup>. On the other hand, intraoperative management is a quite different expertise, which is totally put into practice by anesthesiologists only, and is not covered in this article. Although postoperative care units are mostly managed by a team of both anesthesiologists and surgeons or only by anesthesiologists in Europe and Japan, surgeons' presence and co-leadership is of great importance in postoperative care. Harmonious with this view, surgeons play the largest role in North America<sup>[1,2]</sup>.

Types of the fluids, amount of the fluid given and timing of the administration are the main topics that determine the fluid management strategy. Several debates have been continued about each of these topics. In early times of modern medicine, administering large amounts of fluids was favored, instead of facing the risk of hypovolemia<sup>[3]</sup>. In 1961, Shires *et al.*<sup>[4]</sup> defined the "third space" fluid deficit as nonfunctional fluid which can be accounted as fluid loss and they suggested use of large quantities of fluids to substitute this functional loss. After this strategy becomes popular, reports of adverse effects of high volume states induced by excessive saline use began to arise. Today, exact amount of fluid to maintain ideal homeostasis is still controversial. Similarly, there are varying types of intravenous fluids and all vary in their biological and chemical properties which results in varying distribution forms and varying effects on homeostasis, vascular integrity, and other hemodynamic variables. Apparently, fluid management is admitted to be an art of medicine and based on personal judgments. Although this approach may not be totally wrong, plenty of evidence acquired by large volume studies should be considered wisely.

Postoperative fluid management plays a key role in providing adequate tissue perfusion, stable

hemodynamics and reducing morbidities related with hemodynamics. Understanding body fluid physiology and possible outcomes of different fluid management strategies is crucial for all surgeons.

## BODY FLUID COMPARTMENTS

Total body water is approximately 60% of total body weight. One third of this water is extracellular and it can be divided to as intravascular (20%) and extravascular (80%). The remaining two-third of body water is intracellular, which also exists in intravascular and extravascular compartments. From another perspective, intravascular fluid contains of both intracellular (40%) and extracellular (60%) compounds and plasma is the intravascular-extracellular compound of total body water (approximately 4% of body weight; in example, about 2.8 L in a 70 kg individual).

The endothelium is the separating wall between intravascular and extravascular compartments, thus it is the cell wall that separates the intracellular and extracellular compartments. There are various control mechanisms on these separating walls that regulate volumes of each compartment. Cell membrane is completely permeable to water, whereas it is selectively permeable to ions and organic molecules. It has also the Na<sup>+</sup>/K<sup>+</sup>-adenosine triphosphatase enzyme that actively expels Na<sup>+</sup> ions and maintains the Na<sup>+</sup> gradient between compartments. There are also endocrine mechanisms that control the cellular intake of certain molecules, such as glucose.

On the other hand, the earliest theory on vascular barrier by Ernest Starling declared that the hydrostatic pressure gradient in blood vessels creates a flow and the oncotic pressure of interstitial tissue allows only reasonable amount of fluid to cross through endothelium<sup>[5]</sup>. Later studies showed the intravascular osmotic pressure is significantly higher than interstitial osmotic pressure, however this doesn't result in interstitial edema<sup>[6]</sup>. As a result, this unexplained situation led researchers to look for another actor in this fluid distribution balance. The endothelial glycocalyx is a carbohydrate-rich coating over endothelial surface which is supported by proteoglycans and glycoproteins. It is a dynamic formation, consisting of membrane-bound and soluble molecules<sup>[7]</sup>. Existence of this glycocalyx layer forms a distinct space in the interior neighborhood of the endothelium, and there develops a notable oncotic pressure in this particular protein-free space. This definition brings out the "double-layer concept" for the vascular barrier<sup>[6,8,9]</sup>. This concept is quite capable of clarifying oncotic pressure balance between two compartments.

## WHAT HAPPENS TO THE FLUID BALANCE IN SURGERY?

Homeostasis defines the tendency of the organism to maintain stability and balance. In this manner, body

fluid balance is controlled by previously described compartment mechanisms. On the other hand, any physical intervention may cause imbalance of the body fluids. During relatively long lasting major surgeries, which are performed with general anesthesia, whole intake is controlled by the anesthesiologist and fluid loss happens in numerous different ways such as bleeding, drainage of ascites, urination, insensible water loss and "third space losses". Intraoperative management of acute losses is not covered in this article. However, long term effects of these intraoperative events, such as possible over-hydrating by the anesthesiologist, dehydration, and bleeding should be considered in the postoperative care unit.

The third space is a term for spaces in which body fluids lose their function to affect fluid balance between intravascular and extravascular compartments. In other words, it can be called as non-functional extracellular volume. Bowel lumen, peritoneal and pleural cavities are thought to be the major examples of the third space. Studies that tried to explain the third space loss measured the extracellular volume (ECV) and functional ECV (fECV). fECV is defined as fluid accumulations within the interstitial space combined with plasma. Shires showed that, there is up to 28% loss in extracellular volume after two hours of operative time, during elective surgeries of thirteen adult patients<sup>[4]</sup>. Subsequent studies in 1960s support this finding and existence of the third space<sup>[10-12]</sup>. However, numerous trials with improved methodology proved that fECV levels do not decrease in or after surgery<sup>[13-16]</sup>. This correction of data couldn't be recognized well enough, but still, favored common belief is in the presence and importance of the third space. Current evidence supports that fECV is not negatively affected by surgery, however over-hydration with saline and surgical trauma cause endothelial dysfunction and interstitial edema due to fluid shift to ECV<sup>[13]</sup>. In conclusion, "the third space" term should only refer to anatomical cavities like bowel lumen, peritoneum and pleura, and should only be considered in certain cases. Moreover, possible endothelial glycocalyx dysfunction and fluid shift to ECV should be our guiding facts for determining the right strategy in postoperative fluid management.

## MONITORING BODY FLUID STATUS

Mostly, the main goal of fluid resuscitation is to provide adequate tissue perfusion without harming the patient. It can be also said that fluid resuscitation is generally the first step in patients with inadequate tissue perfusion. However, it should be kept in mind that infusion of large volumes of fluids to patients who don't have enough preload reserves may result in unbalanced fluid shift to interstitial tissue, having no useful effect on tissue perfusion. Intravenous fluid administration will have no effect on tissue perfusion, unless it increases the stroke volume. Studies show that nearly half of the unstable patients are not hemodynamically

responsive to fluid resuscitation<sup>[17,18]</sup>. This means that, fluid resuscitation may not always be the right way to provide adequate tissue perfusion, especially in unstable patients. Thereby, assessment of the patient's responsiveness to fluid resuscitation should determine the need of extra volume.

Thus, we need to determine the actual body fluid status of the patient and build a strategy accordingly. For this purpose, static measures of intravascular volume are being used for decades and central venous pressure (CVP) has been the most favorite tool<sup>[19,20]</sup>. CVP is widely believed to indicate general intravascular volume status of the patient. Moreover, many intensivists think that, CVP is directly correlated with right ventricle stroke volume and indirectly correlated with left ventricle stroke volume. However, a systematic review of 24 studies showed no relation between CVP and left ventricle stroke volume<sup>[21]</sup>. Due to lack of evidence that supports CVP as an indicator of body fluid needs, we should not make our fluid resuscitation decisions based on CVP levels. Similarly, pulmonary capillary wedge pressure is another static measure of intravascular volume and is incapable of predicting fluid responsiveness, in contrast to the common assumption<sup>[22]</sup>. Besides, the two even less favored static measures are left ventricular end-diastolic area and inferior vena caval diameter.

On the other hand, recent studies claim that monitoring of the interactions of heart and lung in mechanically ventilated patients, so called dynamic measures, can be used to determine patient's fluid status. According to Marik *et al.*<sup>[18]</sup>, non-invasive techniques such as the pulse pressure variation, the stroke volume variation, and systolic pressure variation can significantly predict fluid responsiveness in mechanically ventilated patients. These techniques are based on physiological facts. The patients, whose pulse pressures or stroke volumes are more dependent on intra-thoracic pressure variations provided by the ventilator, tend to be more responsive to fluid resuscitation.

The physiological principles underlying the pulse pressure variation (PPV) and the stroke volume variation (SVV) are based on the effects of increased pleural pressure. As the mechanical ventilator increases the pleural pressure, the increased resistance in the pulmonary system causes a decrease in the right ventricle preload and an increase in the right ventricle afterload. Meanwhile, the left ventricle preload and afterload are affected exactly the opposite way of right ventricle is: Left ventricle preload increases and afterload decreases at the end of inspiration. The pulse pressure and the left ventricle stroke volume are at their highest values at this moment. Afterwards, prolongation of blood transit time through pulmonary system results in a decrease in the left ventricle preload and reduction in the left ventricle stroke volume (and the pulse pressure) during expiratory period<sup>[23,24]</sup>. Echocardiographic evaluations of aortic flow velocity and stroke volume and vena caval diameter variation

are two other dynamic parameters based on similar physiological reactions.

Another technique for predicting fluid responsiveness is called the passive leg raising (PLR). While previously mentioned techniques are used for mechanically ventilated patients especially who has no spontaneous breathing, PLR can be used on any patient. Raising the legs to provide a better cardiac preload has been used for a long time in emergency patients. Recently PLR gained interest as a predictor for fluid responsiveness. Monnet pointed out that lifting the legs passively in a lying patient induces a significant blood flow towards the heart<sup>[25]</sup>. Therefore, Marik *et al.*<sup>[17]</sup> called this physiologic condition as "autotransfusion". In a study on mechanically ventilated patients, PLR-induced changes have been found to be strongly similar with the effects of 300 mL colloid infusion. As a result, PLR simulates the state after fluid administration. In other words, if the patient has enough preload reserve, PLR will increase left ventricle preload and stroke volume correspondingly. It is also been reported that, these effects are reversible, and when legs are returned to their horizontal positions, this preload increasing effect disappears<sup>[25]</sup>. Another important point is that PLR reaches its maximal effect in 1 min and its effects disappear gradually in time<sup>[26]</sup>. Accordingly, when PLR is used to predict fluid responsiveness changes in arterial pulse pressure<sup>[27]</sup>, descending aorta blood flow<sup>[28]</sup>, pulse contour-derived stroke volume, or pulsed Doppler-derived velocity-time integral<sup>[29]</sup> should be monitored closely at the first minute<sup>[25]</sup>.

Briefly, fluid management must be done based on the patient's body fluid status. Patients who are responsive to fluids can benefit from fluid resuscitation, whereas patients who are not fluid responsive are more likely to suffer complications of over-hydration.

Therefore, common use of CVP, which is proved to be inefficient to predict fluid responsiveness, should be avoided and attempts should be made to extend the use of techniques like PLR, pulse pressure variation and the stroke volume variation. Practical tools should be manufactured and made available for common use.

## TYPES OF INTRAVENOUS FLUIDS: CRYSTALLOIDS AND COLLOIDS

Intravenous fluids are classified into two main types: Crystalloids and colloids. Each group has its very own characteristics and moreover, each particular solution has its unique properties.

### Crystalloids

Crystalloids consist of glucose or sodium chloride (saline) solutions. Osmolarity of the solution determines if the solution is hypotonic, isotonic or hypertonic. Isotonic solutions have the closest osmolarity to plasma and the other solution types are named comparing to plasma osmolarity. Saline solution containing 0.9 g of NaCl in

each liter of water is defined as isotonic saline, and it is the most popular intravenous fluid worldwide. Some widely used saline solutions also contain one or more of these components: potassium, calcium, bicarbonate, lactate, and glucose. Isotonic glucose solution contains 50 g glucose in each liter of water and it is defined as isotonic glucose. Glucose in these solutions is metabolized right after administration and solvent is mixed into total body water. On the other hand, saline solution's high NaCl concentration serves to keep its solvent water in the extracellular compartment. However, any crystalloid solution can freely pass through double barrier of endothelium. This condition causes up to four-fifth of the infused crystalloid to distribute directly into the interstitial compartment<sup>[13,30]</sup>. Accordingly, crystalloid infusion in high amounts is related with serious complications, such as pulmonary edema<sup>[31]</sup>, and hyperchloremic acidosis<sup>[32]</sup>. Despite that, colloid solutions are generally imprisoned in intravascular compartment, unless double-barrier of endothelium is impaired. Major advantage of crystalloids to colloids is containing only ions or small sized molecules which can easily be metabolized in reasonable amounts.

### Colloids

Colloids can be blood products, such as human albumin solution and fresh frozen plasma, or they can also be synthetic large molecules which are not able to distribute across vascular barrier such as gelatins, dextrans, and hydroxyethyl starches.

Colloids are, like crystalloids, widely used in fluid resuscitation<sup>[33]</sup>. Although colloids are thought to be more useful than crystalloids for increasing intravascular volume and providing osmotic pressure, they are both shown to be similarly effective on mortality<sup>[34,35]</sup>. Colloid solutions are prepared by dissolving colloid molecules in isotonic saline solutions, or more rarely in other crystalloids.

Endogenous albumin is primarily responsible for intravascular osmotic pressure in healthy subjects. Thus, albumin, as an intravenous colloid solution, makes perfect sense to maintain intravascular colloid pressure. However, like all blood products, it has significant disadvantages, like allergic reactions and (theoretically) infection risks, although it is generally considered safe. Molecular weight of albumin is around 69000 Dalton. Gelatins, dextrans and hydroxyethyl starches (HES) are other common colloid substances. Gelatins are products of biochemical processes executed on bovine collagen. Although there are some concerns about its relation with Creutzfeld-Jacob disease and bovine spongiform encephalitis, there is no solid evidence proving these concerns<sup>[36,37]</sup>. Dextrans are polysaccharides that can vary in size. Most common types of dextrans are dextran 70 and dextran 40, which are named after their average molecular weights: 70000 and 40000 Dalton, respectively. Lastly, HES is a nonionic starch derivative, which is synthesized from amylopectin. HESs

also vary in molecular weight, and can be classified as low (70000-130000 Dalton), medium, and high (450000-480000 Dalton) molecular weights. They are also classified by their molar substitution degree, which defines the proportion of glucose molecules that are replaced by hydroxyethyls. HESs are the most commonly used colloids in Europe. Commonly used examples of these colloids are Voluven® (Fresenius Kabi, Bad Homburg, Germany) which is a 130000 Dalton tetra starch, dissolved in saline with substitution degree of 0.4 and HAES-steril® (Fresenius Kabi, Bad Homburg, Germany) which is a 200000 Dalton pentastarch, dissolved in saline with substitution degree of 0.5.

Each type of colloid solution has its unique features. Effect on plasma volume and plasma viscosity, adverse reactions, and side effects on the system are the main concerns while choosing colloid solutions. Every colloid substance has a concentration decrease rate (half-life) in plasma by being metabolized, or by a loss through endothelial barrier and glomerular filtration. Half-life of a colloid determines the amount and the duration of plasma volume expansion. Higher molecular weight colloids tend to stay longer in the intravascular compartment. Besides, some studies point that the dextrans and the HESs provide significantly better expansion of plasma volume than the gelatins<sup>[38-40]</sup>. Whereas, some studies indicate that only albumin has significant advantage over other colloids and saline; and none of the other colloids is superior to others regarding plasma volume expansion<sup>[41-43]</sup>.

All colloids provide a level of expansion in plasma volume and this leads to hemodilution. Hemodilution causes a decrease in plasma viscosity. However, it is known that some colloids cause a total increase in viscosity due to red cell aggregation. High molecular weight dextrans and HESs cause a significant increase in viscosity, while low molecular weight dextrans HESs and albumin solutions decrease both red cell aggregation and plasma viscosity<sup>[44-47]</sup>. Colloids have various effects on hemostasis, such as impaired platelet function, decreased factor VIIIc and von Willebrand Factor levels, in addition to previously described hemodilution and altered red cell aggregation<sup>[44,48,49]</sup>. Particularly, dextrans are known with their significant antithrombotic effects<sup>[49-51]</sup>.

Accumulation of colloid substances in the body is possible. Dextrans and gelatins can be metabolized in humans. On the other hand, HESs may also accumulate. Metabolism and filtration of HES is relatively slow and storage in reticulo-endothelial system is not well recognized yet.

All colloids are large molecules and can trigger anaphylaxis of anaphylactoid events. Colloids also have minor anti-inflammatory effects.

Although it has been argued for a long time, there are still no definite rules on "crystalloid vs colloids" issue. There are studies that show crystalloid infusion is related with interstitial edema and worse anastomotic

healing<sup>[31,52,53]</sup>. On the other side, it is still arguable that colloid solutions are able to prevent consequences of these negative effects<sup>[54,55]</sup>. In a study on pancreaticoduodenectomy patients, who are resuscitated with lactated Ringer's solution (isotonic crystalloid solution; including lactate, potassium and calcium in addition to sodium chloride), the significantly increased interstitial edema in jejunum was shown<sup>[56]</sup>. However, colloid use has been reported to have an increasing effect on mortality, in some fairly criticized studies, especially on critically ill patients<sup>[57,58]</sup>. On the other hand, CRISTAL trial, which is a multicenter randomized study on critically ill patients, failed to demonstrate this effect on mortality. In contrast, fewer death rates were found within 90 d in colloids group<sup>[54]</sup>.

Moreover, although colloids are proved to be capable of maintaining efficient plasma volume, they do not appear to have positive effects on renal function. Contrarily, reports had shown significant harmful effects of dextran 40 use on kidney function in the second half of 20<sup>th</sup> century<sup>[59-61]</sup>. Some of the subsequent studies on HESs also revealed negative effects of these solutions on kidneys<sup>[62,63]</sup>. Schortgen *et al*<sup>[64]</sup> also reported that the use of hyperoncotic colloids and human albumin is significantly associated with renal dysfunction. However, in a multicenter study on over 3000 intensive care patients, no significant relation was detected between HES use and renal dysfunction<sup>[65]</sup>. Similarly, in a review of studies with different HES products, no adverse effects on kidneys were reported<sup>[66]</sup>. In a randomized clinical multicenter trial, 6997 critically ill patients were randomized into two groups. One group was assigned to receive 4% of albumin and the other group was assigned to receive saline for intravenous resuscitation during 28 d. There was no significant difference between two groups, regarding to mortality, days spent in intensive care unit, days of mechanical ventilation, or days of renal replacement therapy<sup>[67]</sup>. In addition to all of these results, it should be taken into consideration that none of the colloid solutions is proved to be directly toxic to the kidneys<sup>[68]</sup>.

Considering all pros and cons of each solution family, it is still not possible to make a strict evidence based statement about how to use colloids and crystalloids<sup>[57,69]</sup>. It should be kept in mind that, crystalloids have less negative effects on hemostasis, immune system and kidneys; whereas colloids may provide a better plasma volume expansion with less interstitial edema in elective surgery patients<sup>[69]</sup>.

## FLUID RESUSCITATION STRATEGIES

Although there has been various different strategies defined in literature in decades, none has been adopted alone by most of the clinicians as the superior strategy. We think that many clinicians tend to keep their accustomed strategy, despite the evidences in the literature. There are studies that compare outcomes

of different strategies of fluid management. Lately, “crystalloids vs colloids” debates are fading, while recent studies mostly focus on the amount of fluid given perioperatively.

Traditional approach to determine the fluid amounts is more likely to generate formulas based on parameters such as patients’ body weights and duration of surgeries. However, there is an evidence that each patient has his/her own body fluid status depending on the type of surgery, comorbid conditions, fluid already administered before, and various other factors. In addition, each patient should be considered as unique and his/her unique status should be monitored closely in the correct ways. As stated before, the main goal of fluid management is to maintain adequate tissue perfusion, with minimized risks of complications of over-hydration, such as pulmonary edema, cerebral edema, and intestinal edema. Both inadequate and excessive fluid administration may increase the stress on the circulatory system, and can affect tissue healing after surgery. From this perspective, without decent monitoring of patient’s current status, any strategy may fail.

Debates about fluid management strategies are gathered around liberal strategy, restricted (conservative) strategy and goal-directed strategy so far. Liberal and restricted strategies are defined by different authors with variable volume ranges. For example, in one study, restricted fluid volume is defined as 1000 mL plus loss through drains<sup>[70]</sup>, while in another study, patients in restricted fluid volume group were subjected to over 2000 mL fluid on the day of surgery<sup>[71]</sup>. These variances make it difficult to consider these studies as a whole. Still, majority of authors studying this subject point out that restrictive strategy has positive effects on gastrointestinal function, wound healing and pulmonary function<sup>[44,70,72-74]</sup>. Brandstrup *et al.*<sup>[70]</sup> stated that, excessive hydration with crystalloids is related with increased major complications, such as leakage, peritonitis, sepsis, pulmonary edema and bleeding in patients who underwent elective colorectal surgery. Also, intestinal edema is known to be related with increased bacterial translocation and multiple organ dysfunction syndrome rates<sup>[75,76]</sup>. It can be concluded that, staying closer to the dehydration level is more reasonable, because it is safer and more efficient than administering large volumes to avoid dehydration. On the other hand, the liberal strategy is superior to the restricted strategy for reducing postoperative nausea, headache, dizziness and vomiting<sup>[77,78]</sup>.

However, the goal directed strategy (GDS) is totally based on patient’s current data, obtained from monitoring methods (See section: Monitoring body fluid status). Rivers and colleagues, one of the pioneers of this strategy, monitored CVP, mean arterial pressure, serum lactate, and mixed venous oxygen saturation in order to manage therapy in sepsis patients<sup>[79]</sup>. Later studies were focused on monitoring hemodynamics,

and the effects of administered fluids on patients. Now, GDS can be defined as an individualized fluid therapy, based on patient’s fluid responsiveness; in other words, “fluid need”. The extra volume, which won’t be able to affect the left ventricle stroke volume is regarded as unnecessary; and as a matter of fact, hazardous. It makes perfect sense to totally evaluate patient’s needs and replace what is needed. Still, efficiency of GDS is limited with the power of our monitoring tools, which is determined by accessibility, applicability of the tools and the quality of information we acquire from them.

PPV and SVV are defined to monitor the fluid need of the patient dynamically as it is stated above<sup>[18]</sup>. Esophageal Doppler monitoring of cardiac volumes and aortic flow are also one of the helpful tools in GDS. In a systematic review of esophageal Doppler guided GDS studies; reduced hospital stay, fewer ICU admissions, and less inotropes usage were detected in GDS group<sup>[80]</sup>. In a single center, blinded, prospective controlled trial, 128 patients who underwent colorectal resection were randomized into two groups. Each group was managed with esophageal Doppler or CVP guided fluid therapy during surgery. Intraoperative Doppler guided fluid management was associated with decrease in the duration of hospital stay<sup>[81]</sup>. A randomized controlled study on 108 elective colorectal surgery patients also showed shorter hospital stay and decreased morbidity in GDS group<sup>[82]</sup>. GDS is also advantageous in patients who undergo major surgery<sup>[79]</sup>. A systematic review and meta-analysis studies by Hamilton on major surgery patients state that preemptive hemodynamic monitoring reduces mortality and morbidity<sup>[83]</sup>. Similarly, Poeze *et al.*<sup>[84]</sup> showed that efforts to achieve an optimized hemodynamic condition resulted in a decreased mortality rate, in their meta-analysis study in 2005. Another meta-analysis also shows that GDS reduces both major and minor gastrointestinal complications after surgery<sup>[85]</sup>.

In contrast with these studies, in a multicenter study, which included 762 high risk patients in 56 intensive care units, no significant effects of GDS were found. In this study, patients were randomly assigned to cardiac-index group, mixed venous oxygen-saturation group and standard therapy group. Predetermined hemodynamic targets were reached significantly better in the control group. There were no significant differences among the three groups, regarding mortality at six months. Even the subgroup analysis of patients, whose predetermined hemodynamic targets have been reached successfully, showed similar mortality rates among the three groups. Moreover, the number of dysfunctional organs and the duration of stay in the intensive care unit were similar in all groups<sup>[86]</sup>.

Despite these evidences, low accessibility and applicability of esophageal Doppler are the major disadvantages of this method. This leads researchers to search for a more accessible and applicable method for common use in postoperative care unit, such as non-

invasive pulse oximetry and invasive arterial pressure measurement. Thus, predictive value of pulse pressure variation, systolic pressure variation and stroke volume variation tests for fluid responsiveness are defined<sup>[17]</sup>. All of these tests are applicable in an average postoperative care unit. However, the true value of these tests should be evaluated by larger studies. After that, optimization of patient monitoring devices should be done accordingly. Moreover, even PLR alone can provide important information about fluid responsiveness and lead the intensivists for GDS.

Since there is still insufficient number of randomized controlled trials with standardized criteria, the fluid management debates are going on. A consensus on criteria for each fluid management strategy should be made. We think that the related studies from all around the world with defined criteria are going to reveal the true value of each strategy.

Each surgeon should keep in mind that the patient is totally managed by the anesthesiologist during the surgery, so depending on the anesthesiologist's preference on fluid strategy, patient's fluid status after surgery may vary widely. Besides, intraoperative bleeding and other causes of surgical fluid loss should also be considered. During or after the surgery, the blood loss in patients with low hemoglobin levels is generally managed with erythrocyte suspensions. However, in patients with reasonable hemoglobin levels, appropriate fluid strategy should be chosen to avoid complications of transfusion. We think that determining the actual fluid status and the needs of a postoperative patient, by using monitoring tools and examining the report of the anesthesiologist, is of great importance.

## CONCLUSION

Postoperative care units can be managed by an anesthesiologist, a surgeon or a team composed of both. Management of postoperative fluid therapy should be done considering both patients' unique status and intraoperative events. Thus, surgeons must be aware of pros and cons of current fluid management strategies and their effects on surgical outcome. Although there has been a significant progress on fluid status monitoring and fluid management strategies, most clinicians still prefer their traditional approaches for postoperative fluid management. This tendency towards empirical fluid management can be replaced by evidence based strategies, only if significant benefits of new strategies are proved with multicenter randomized controlled trials which use standardized criteria. GDS is the most rational approach to assess the patient and maintain optimum fluid balance. However, accessible and applicable monitoring tools for determining patient's actual fluid need should be further studied and universalized. The debate around colloids and crystalloids should also be considered with goal directed therapies. Advantages and disadvantages of each solution must be evaluated with

the patient's specific condition.

## REFERENCES

- 1 **Linke GR**, Mieth M, Hofer S, Trierweiler-Hauke B, Weitz J, Martin E, Büchler MW. Surgical intensive care unit - essential for good outcome in major abdominal surgery? *Langenbecks Arch Surg* 2011; **396**: 417-428 [PMID: 21369847 DOI: 10.1007/s00423-011-0758-y]
- 2 **Johnson JL**, Moore EE, Aasen AO, Rogy MA, Wang JE, Alsanea O, Aikawa N, Neira JA, Tisminetzky GJ. The role of the surgeon as intensivist: an international perspective. *Curr Opin Crit Care* 2006; **12**: 357-369 [PMID: 16810049 DOI: 10.1097/01.ccx.0000235215.71612.a9]
- 3 **Bamboatz ZM**, Bordeianou L. Perioperative fluid management. *Clin Colon Rectal Surg* 2009; **22**: 28-33 [PMID: 20119553 DOI: 10.1055/s-0029-1202883]
- 4 **Shires T**, Williams J, Brown F. Acute change in extracellular fluids associated with major surgical procedures. *Ann Surg* 1961; **154**: 803-810 [PMID: 13912109]
- 5 **Starling EH**. On the Absorption of Fluids from the Connective Tissue Spaces. *J Physiol* 1896; **19**: 312-326 [PMID: 16992325]
- 6 **Adamson RH**, Lenz JF, Zhang X, Adamson GN, Weinbaum S, Curry FE. Oncotic pressures opposing filtration across non-fenestrated rat microvessels. *J Physiol* 2004; **557**: 889-907 [PMID: 15073281 DOI: 10.1113/jphysiol.2003.058255]
- 7 **Reitsma S**, Slaaf DW, Vink H, van Zandvoort MA, oude Egbrink MG. The endothelial glycocalyx: composition, functions, and visualization. *Pflugers Arch* 2007; **454**: 345-359 [PMID: 17256154 DOI: 10.1007/s00424-007-0212-8]
- 8 **Strunden MS**, Heckel K, Goetz AE, Reuter DA. Perioperative fluid and volume management: physiological basis, tools and strategies. *Ann Intensive Care* 2011; **1**: 2 [PMID: 21906324 DOI: 10.1186/2110-5820-1-2]
- 9 **Rehm M**, Zahler S, Lötsch M, Welsch U, Conzen P, Jacob M, Becker BF. Endothelial glycocalyx as an additional barrier determining extravasation of 6% hydroxyethyl starch or 5% albumin solutions in the coronary vascular bed. *Anesthesiology* 2004; **100**: 1211-1223 [PMID: 15114220]
- 10 **Carrico CJ**, Coln CD, Lightfoot SA, Allsman A, Shires GT. Extracellular fluid volume replacement in hemorrhagic shock. *Surg Forum* 1963; **14**: 10-12 [PMID: 14064470]
- 11 **Shires T**, Coln D, Carrico J, Lightfoot S. Fluid therapy in hemorrhagic shock. *Arch Surg* 1964; **88**: 688-693 [PMID: 14107023]
- 12 **Fukuda Y**, Fujita T, Shibuya J, Albert SN. The distribution between the intravascular and interstitial compartments of commonly utilized replacement fluids. *Anesth Analg* 1977; **48**: 831-838 [PMID: 4897746]
- 13 **Jacob M**, Chappell D, Rehm M. The 'third space'--fact or fiction? *Best Pract Res Clin Anaesthesiol* 2009; **23**: 145-157 [PMID: 19653435]
- 14 **Gumpert JR**, Zollinger RM, Riddell AG. Proceedings: the measurement of extracellular fluid volume with radiobromide simultaneous plasma and lymph disappearance in man. *Br J Surg* 1973; **60**: 903 [PMID: 4584778]
- 15 **Breckenridge IM**, Digeress SB, Kirklín JW. Validity of concept of increased extracellular fluid after open heart surgery. *Surg Forum* 1969; **20**: 169-171 [PMID: 4910576]
- 16 **Nielsen OM**, Engell HC. Extracellular fluid volume and distribution in relation to changes in plasma colloid osmotic pressure after major surgery. A randomized study. *Acta Chir Scand* 1985; **151**: 221-225 [PMID: 3892993]
- 17 **Marik PE**, Monnet X, Teboul JL. Hemodynamic parameters to guide fluid therapy. *Ann Intensive Care* 2011; **1**: 1 [PMID: 21906322 DOI: 10.1186/2110-5820-1-1]
- 18 **Marik PE**, Cavallazzi R, Vasu T, Hirani A. Dynamic changes in arterial waveform derived variables and fluid responsiveness

- in mechanically ventilated patients: a systematic review of the literature. *Crit Care Med* 2009; **37**: 2642-2647 [PMID: 19602972 DOI: 10.1097/CCM.0b013e3181a590da]
- 19 **McIntyre LA**, Hébert PC, Fergusson D, Cook DJ, Aziz A; Canadian Critical Care Trials Group. A survey of Canadian intensivists' resuscitation practices in early septic shock. *Crit Care* 2007; **11**: R74 [PMID: 17623059 DOI: 10.1186/cc5962]
  - 20 **Kastrup M**, Markewitz A, Spies C, Carl M, Erb J, Grosse J, Schirmer U. Current practice of hemodynamic monitoring and vasopressor and inotropic therapy in post-operative cardiac surgery patients in Germany: results from a postal survey. *Acta Anaesthesiol Scand* 2007; **51**: 347-358 [PMID: 17096667 DOI: 10.1111/j.1399-6576.2006.01190.x]
  - 21 **Marik PE**, Cavallazzi R. Does the central venous pressure predict fluid responsiveness? An updated meta-analysis and a plea for some common sense. *Crit Care Med* 2013; **41**: 1774-1781 [PMID: 23774337 DOI: 10.1097/CCM.0b013e31828a25fd]
  - 22 **Solus-Biguenet H**, Fleyfel M, Tavernier B, Kipnis E, Onimus J, Robin E, Lebuffe G, Decoene C, Pruvot FR, Vallet B. Non-invasive prediction of fluid responsiveness during major hepatic surgery. *Br J Anaesth* 2006; **97**: 808-816 [PMID: 16980709 DOI: 10.1093/bja/ael250]
  - 23 **Michard F**, Teboul JL. Using heart-lung interactions to assess fluid responsiveness during mechanical ventilation. *Crit Care* 2000; **4**: 282-289 [PMID: 11094507 DOI: 10.1186/cc710]
  - 24 **Theres H**, Binkau J, Laule M, Heinze R, Hundertmark J, Blobner M, Erhardt W, Baumann G, Stangl K. Phase-related changes in right ventricular cardiac output under volume-controlled mechanical ventilation with positive end-expiratory pressure. *Crit Care Med* 1999; **27**: 953-958 [PMID: 10362419]
  - 25 **Monnet X**, Teboul JL. Passive leg raising. *Intensive Care Med* 2008; **34**: 659-663 [PMID: 18214429 DOI: 10.1007/s00134-008-0994-y]
  - 26 **Monnet X**, Rienzo M, Osman D, Anguel N, Richard C, Pinsky MR, Teboul JL. Passive leg raising predicts fluid responsiveness in the critically ill. *Crit Care Med* 2006; **34**: 1402-1407 [PMID: 16540963 DOI: 10.1097/01.CCM.0000215453.11735.06]
  - 27 **Boulaïn T**, Achard JM, Teboul JL, Richard C, Perrotin D, Ginies G. Changes in BP induced by passive leg raising predict response to fluid loading in critically ill patients. *Chest* 2002; **121**: 1245-1252 [PMID: 11948060]
  - 28 **Lafanechère A**, Pène F, Goulenok C, Delahaye A, Mallet V, Choukroun G, Chiche JD, Mira JP, Cariou A. Changes in aortic blood flow induced by passive leg raising predict fluid responsiveness in critically ill patients. *Crit Care* 2006; **10**: R132 [PMID: 16970817 DOI: 10.1186/cc5044]
  - 29 **Lamia B**, Ochagavia A, Monnet X, Chemla D, Richard C, Teboul JL. Echocardiographic prediction of volume responsiveness in critically ill patients with spontaneously breathing activity. *Intensive Care Med* 2007; **33**: 1125-1132 [PMID: 17508199 DOI: 10.1007/s00134-007-0646-7]
  - 30 **Kinsella SM**, Pirllet M, Mills MS, Tuckey JP, Thomas TA. Randomized study of intravenous fluid preload before epidural analgesia during labour. *Br J Anaesth* 2000; **85**: 311-313 [PMID: 10992845]
  - 31 **Stein L**, Beraud JJ, Morissette M, Luz PD, Weil MH, Shubin H. Pulmonary edema during volume infusion. *Circulation* 1975; **52**: 483-489 [PMID: 1157248]
  - 32 **Waters JH**, Gottlieb A, Schoenwald P, Popovich MJ, Sprung J, Nelson DR. Normal saline versus lactated Ringer's solution for intraoperative fluid management in patients undergoing abdominal aortic aneurysm repair: an outcome study. *Anesth Analg* 2001; **93**: 817-822 [PMID: 11574339]
  - 33 **Yim JM**, Vermeulen LC, Erstad BL, Matuszewski KA, Burnett DA, Vlasses PH. Albumin and nonprotein colloid solution use in US academic health centers. *Arch Intern Med* 1995; **155**: 2450-2455 [PMID: 7503604]
  - 34 **Alderson P**, Bunn F, Lefebvre C, Li WP, Li L, Roberts I, Schierhout G. Human albumin solution for resuscitation and volume expansion in critically ill patients. *Cochrane Database Syst Rev* 2004; **(4)**: CD001208 [PMID: 15495011 DOI: 10.1002/14651858.CD001208.pub2]
  - 35 **Roberts I**, Alderson P, Bunn F, Chinnock P, Ker K, Schierhout G. Colloids versus crystalloids for fluid resuscitation in critically ill patients. *Cochrane Database Syst Rev* 2004; **(4)**: CD000567 [PMID: 15495001 DOI: 10.1002/14651858.CD000567.pub2]
  - 36 **Taylor DM**. Inactivation of TSE agents: safety of blood and blood-derived products. *Transfus Clin Biol* 2003; **10**: 23-25 [PMID: 12668184]
  - 37 **Grobben AH**, Steele PJ, Somerville RA, Taylor DM, Schreuder BE. Inactivation of the BSE agent by the heat and pressure process for manufacturing gelatine. *Vet Rec* 2005; **157**: 277-281 [PMID: 16157568]
  - 38 **Lamke LO**, Liljedahl SO. Plasma volume changes after infusion of various plasma expanders. *Resuscitation* 1976; **5**: 93-102 [PMID: 69313]
  - 39 **Mortelmans YJ**, Vermaut G, Verbruggen AM, Arnout JM, Vermylen J, Van Aken H, Mortelmans LA. Effects of 6% hydroxyethyl starch and 3% modified fluid gelatin on intravascular volume and coagulation during intraoperative hemodilution. *Anesth Analg* 1995; **81**: 1235-1242 [PMID: 7486110]
  - 40 **Van der Linden PJ**, De Hert SG, Deraedt D, Cromheecke S, De Decker K, De Paep R, Rodrigus I, Daper A, Trenchant A. Hydroxyethyl starch 130/0.4 versus modified fluid gelatin for volume expansion in cardiac surgery patients: the effects on perioperative bleeding and transfusion needs. *Anesth Analg* 2005; **101**: 629-634, table of contents [PMID: 16115963 DOI: 10.1213/01.ANE.0000175216.53374.27]
  - 41 **Dubniks M**, Persson J, Grände PO. Plasma volume expansion of 5% albumin, 4% gelatin, 6% HES 130/0.4, and normal saline under increased microvascular permeability in the rat. *Intensive Care Med* 2007; **33**: 293-299 [PMID: 17119921 DOI: 10.1007/s00134-006-0454-5]
  - 42 **Persson J**, Grände PO. Volume expansion of albumin, gelatin, hydroxyethyl starch, saline and erythrocytes after haemorrhage in the rat. *Intensive Care Med* 2005; **31**: 296-301 [PMID: 15609019 DOI: 10.1007/s00134-004-2510-3]
  - 43 **Beyer R**, Harmening U, Rittmeyer O, Zielmann S, Mielck F, Kazmaier S, Kettler D. Use of modified fluid gelatin and hydroxyethyl starch for colloidal volume replacement in major orthopaedic surgery. *Br J Anaesth* 1997; **78**: 44-50 [PMID: 9059203]
  - 44 **Grocott MP**, Mythen MG, Gan TJ. Perioperative fluid management and clinical outcomes in adults. *Anesth Analg* 2005; **100**: 1093-1106 [PMID: 15781528 DOI: 10.1213/01.ANE.0000148691.33690.AC]
  - 45 **Freyburger G**, Dubreuil M, Boisseau MR, Janvier G. Rheological properties of commonly used plasma substitutes during preoperative normovolaemic acute haemodilution. *Br J Anaesth* 1996; **76**: 519-525 [PMID: 8652324]
  - 46 **Korosue K**, Heros RC, Ogilvy CS, Hyodo A, Tu YK, Graichen R. Comparison of crystalloids and colloids for hemodilution in a model of focal cerebral ischemia. *J Neurosurg* 1990; **73**: 576-584 [PMID: 1697903 DOI: 10.3171/jns.1990.73.4.0576]
  - 47 **Neff TA**, Fischler L, Mark M, Stocker R, Reinhart WH. The influence of two different hydroxyethyl starch solutions (6% HES 130/0.4 and 200/0.5) on blood viscosity. *Anesth Analg* 2005; **100**: 1773-1780 [PMID: 15920212 DOI: 10.1213/01.ANE.0000149326.45137.9A]
  - 48 **de Jonge E**, Levi M. Effects of different plasma substitutes on blood coagulation: a comparative review. *Crit Care Med* 2001; **29**: 1261-1267 [PMID: 11395618]
  - 49 **Aberg M**, Hedner U, Bergentz SE. Effect of dextran on factor VIII (antihemophilic factor) and platelet function. *Ann Surg* 1979; **189**: 243-247 [PMID: 426556]
  - 50 **Jones CI**, Payne DA, Hayes PD, Naylor AR, Bell PR, Thompson MM, Goodall AH. The antithrombotic effect of dextran-40 in man is due to enhanced fibrinolysis in vivo. *J Vasc Surg* 2008; **48**: 715-722 [PMID: 18572351 DOI: 10.1016/j.jvs.2008.04.008]
  - 51 **Salemark L**, Wieslander JB, Dougan P, Arnljots B. Studies of the antithrombotic effects of dextran 40 following microarterial trauma. *Br J Plast Surg* 1991; **44**: 15-22 [PMID: 1704269]

- 52 **Baum TD**, Wang H, Rothschild HR, Gang DL, Fink MP. Mesenteric oxygen metabolism, ileal mucosal hydrogen ion concentration, and tissue edema after crystalloid or colloid resuscitation in porcine endotoxic shock: comparison of Ringer's lactate and 6% hetastarch. *Circ Shock* 1990; **30**: 385-397 [PMID: 1693551]
- 53 **Marjanovic G**, Villain C, Timme S, zur Hausen A, Hoepfner J, Makowicz F, Holzner P, Hopt UT, Obermaier R. Colloid vs. crystalloid infusions in gastrointestinal surgery and their different impact on the healing of intestinal anastomoses. *Int J Colorectal Dis* 2010; **25**: 491-498 [PMID: 19943164 DOI: 10.1007/s00384-009-0854-4]
- 54 **Annane D**, Siami S, Jaber S, Martin C, Elatrous S, Declère AD, Preiser JC, Outin H, Troché G, Charpentier C, Trouillet JL, Kimmoun A, Forceville X, Darmon M, Lesur O, Reignier J, Abroug F, Berger P, Clec'h C, Cousson J, Thibault L, Chevret S. Effects of fluid resuscitation with colloids vs crystalloids on mortality in critically ill patients presenting with hypovolemic shock: the CRISTAL randomized trial. *JAMA* 2013; **310**: 1809-1817 [PMID: 24108515 DOI: 10.1001/jama.2013.280502]
- 55 **Perel P**, Roberts I. Colloids versus crystalloids for fluid resuscitation in critically ill patients. *Cochrane Database Syst Rev* 2007; **(4)**: CD000567 [PMID: 17943746 DOI: 10.1002/14651858.CD000567.pub3]
- 56 **Prien T**, Backhaus N, Pelster F, Pircher W, Bunte H, Lawin P. Effect of intraoperative fluid administration and colloid osmotic pressure on the formation of intestinal edema during gastrointestinal surgery. *J Clin Anesth* 1990; **2**: 317-323 [PMID: 1702977]
- 57 **Choi PT**, Yip G, Quinonez LG, Cook DJ. Crystalloids vs. colloids in fluid resuscitation: a systematic review. *Crit Care Med* 1999; **27**: 200-210 [PMID: 9934917]
- 58 **Schierhout G**, Roberts I. Fluid resuscitation with colloid or crystalloid solutions in critically ill patients: a systematic review of randomised trials. *BMJ* 1998; **316**: 961-964 [PMID: 9550953]
- 59 **Mailloux L**, Swartz CD, Capizzi R, Kim KE, Onesti G, Ramirez O, Brest AN. Acute renal failure after administration of low-molecular weight dextran. *N Engl J Med* 1967; **277**: 1113-1118 [PMID: 6054998 DOI: 10.1056/NEJM196711232772103]
- 60 **Diomi P**, Ericsson JL, Matheson NA, Shearer JR. Studies on renal tubular morphology and toxicity after large doses of dextran 40 in the rabbit. *Lab Invest* 1970; **22**: 355-360 [PMID: 5429535]
- 61 **Biesenbach G**, Kaiser W, Zazgornik J. Incidence of acute oligoanuric renal failure in dextran 40 treated patients with acute ischemic stroke stage III or IV. *Ren Fail* 1997; **19**: 69-75 [PMID: 9044453]
- 62 **Legendre C**, Thervet E, Page B, Percheron A, Noël LH, Kreis H. Hydroxyethylstarch and osmotic-nephrosis-like lesions in kidney transplantation. *Lancet* 1993; **342**: 248-249 [PMID: 7686994]
- 63 **Brunkhorst FM**, Engel C, Bloos F, Meier-Hellmann A, Ragaller M, Weiler N, Moerer O, Gruendling M, Oppert M, Grond S, Korthoff D, Jaschinski U, John S, Rossaint R, Welte T, Schaefer M, Kern P, Kuhnt E, Kiehnopf M, Hartog C, Natanson C, Loeffler M, Reinhart K. Intensive insulin therapy and pentastarch resuscitation in severe sepsis. *N Engl J Med* 2008; **358**: 125-139 [PMID: 18184958 DOI: 10.1056/NEJMoa070716]
- 64 **Schortgen F**, Girou E, Deye N, Brochard L. The risk associated with hyperoncotic colloids in patients with shock. *Intensive Care Med* 2008; **34**: 2157-2168 [PMID: 18685828 DOI: 10.1007/s00134-008-1225-2]
- 65 **Sakr Y**, Payen D, Reinhart K, Sipmann FS, Zavala E, Bewley J, Marx G, Vincent JL. Effects of hydroxyethyl starch administration on renal function in critically ill patients. *Br J Anaesth* 2007; **98**: 216-224 [PMID: 17251213 DOI: 10.1093/bja/ael333]
- 66 **Boldt J**, Priebe HJ. Intravascular volume replacement therapy with synthetic colloids: is there an influence on renal function? *Anesth Analg* 2003; **96**: 376-382, table of contents [PMID: 12538180]
- 67 **Finfer S**, Bellomo R, Boyce N, French J, Myburgh J, Norton R. A comparison of albumin and saline for fluid resuscitation in the intensive care unit. *N Engl J Med* 2004; **350**: 2247-2256 [PMID: 15163774 DOI: 10.1056/NEJMoa040232]
- 68 **Roche AM**, James MF. Colloids and crystalloids: does it matter to the kidney? *Curr Opin Crit Care* 2009; **15**: 520-524 [PMID: 19829107 DOI: 10.1097/MCC.0b013e3283232f686]
- 69 **Velanovich V**. Crystalloid versus colloid fluid resuscitation: a meta-analysis of mortality. *Surgery* 1989; **105**: 65-71 [PMID: 2911805]
- 70 **Brandstrup B**, Tønnesen H, Beier-Holgersen R, Hjortso E, Ording H, Lindorff-Larsen K, Rasmussen MS, Lannig C, Wallin L, Iversen LH, Gramkow CS, Okholm M, Blemmer T, Svendsen PE, Rottensten HH, Thage B, Riis J, Jeppesen IS, Teilmann D, Christensen AM, Graungaard B, Pott F. Effects of intravenous fluid restriction on postoperative complications: comparison of two perioperative fluid regimens: a randomized assessor-blinded multicenter trial. *Ann Surg* 2003; **238**: 641-648 [PMID: 14578723 DOI: 10.1097/01.sla.0000094387.50865.23]
- 71 **Mackay G**, Fearon K, McConnachie A, Serpell MG, Molloy RG, O'Dwyer PJ. Randomized clinical trial of the effect of postoperative intravenous fluid restriction on recovery after elective colorectal surgery. *Br J Surg* 2006; **93**: 1469-1474 [PMID: 17078116 DOI: 10.1002/bjs.5593]
- 72 **Lobo SM**, Ronchi LS, Oliveira NE, Brandão PG, Froes A, Cunrath GS, Nishiyama KG, Netinho JG, Lobo FR. Restrictive strategy of intraoperative fluid maintenance during optimization of oxygen delivery decreases major complications after high-risk surgery. *Crit Care* 2011; **15**: R226 [PMID: 21943111 DOI: 10.1186/cc10466]
- 73 **Nisanovich V**, Felsenstein I, Almogy G, Weissman C, Einav S, Matot I. Effect of intraoperative fluid management on outcome after intraabdominal surgery. *Anesthesiology* 2005; **103**: 25-32 [PMID: 15983453]
- 74 **Rahbari NN**, Zimmermann JB, Schmidt T, Koch M, Weigand MA, Weitz J. Meta-analysis of standard, restrictive and supplemental fluid administration in colorectal surgery. *Br J Surg* 2009; **96**: 331-341 [PMID: 19283742 DOI: 10.1002/bjs.6552]
- 75 **Baker JW**, Deitch EA, Li M, Berg RD, Specian RD. Hemorrhagic shock induces bacterial translocation from the gut. *J Trauma* 1988; **28**: 896-906 [PMID: 3294427]
- 76 **Wilmore DW**, Smith RJ, O'Dwyer ST, Jacobs DO, Ziegler TR, Wang XD. The gut: a central organ after surgical stress. *Surgery* 1988; **104**: 917-923 [PMID: 3055397]
- 77 **Maharaj CH**, Kallam SR, Malik A, Hassett P, Grady D, Laffey JG. Preoperative intravenous fluid therapy decreases postoperative nausea and pain in high risk patients. *Anesth Analg* 2005; **100**: 675-682, table of contents [PMID: 15728051 DOI: 10.1213/01.ANE.0000148684.64286.36]
- 78 **Moretti EW**, Robertson KM, El-Moalem H, Gan TJ. Intraoperative colloid administration reduces postoperative nausea and vomiting and improves postoperative outcomes compared with crystalloid administration. *Anesth Analg* 2003; **96**: 611-617, table of contents [PMID: 12538221]
- 79 **Rivers EP**, Nguyen HB, Huang DT, Donnino M. Early goal-directed therapy. *Crit Care Med* 2004; **32**: 314-315; author reply 315 [PMID: 14707615 DOI: 10.1097/01.CCM.0000104937.09370.53]
- 80 **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 [PMID: 18086070 DOI: 10.1111/j.1365-2044.2007.05233.x]
- 81 **Wakeling HG**, McFall MR, Jenkins CS, Woods WG, Miles WF, 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 [PMID: 16155038 DOI: 10.1093/bja/ael223]
- 82 **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 [PMID: 16888706 DOI: 10.1002/bjs.5454]
- 83 **Hamilton MA**. Perioperative fluid management: progress despite lingering controversies. *Cleve Clin J Med* 2009; **76** Suppl 4: S28-S31 [PMID: 19880832 DOI: 10.3949/ccjm.76.s4.05]
- 84 **Poeze M**, Greve JW, Ramsay G. Meta-analysis of hemodynamic optimization: relationship to methodological quality. *Crit Care* 2005; **9**: R771-R779 [PMID: 16356226 DOI: 10.1186/cc3902]
- 85 **Giglio MT**, Marucci M, Testini M, Brienza N. Goal-directed haemodynamic therapy and gastrointestinal complications in major

surgery: a meta-analysis of randomized controlled trials. *Br J Anaesth* 2009; **103**: 637-646 [PMID: 19837807 DOI: 10.1093/bja/aep279]

86 **Gattinoni L**, Brazzi L, Pelosi P, Latini R, Tognoni G, Pesenti

A, Fumagalli R. A trial of goal-oriented hemodynamic therapy in critically ill patients. SvO<sub>2</sub> Collaborative Group. *N Engl J Med* 1995; **333**: 1025-1032 [PMID: 7675044 DOI: 10.1056/NEJM199510193331601]

**P- Reviewer:** Gurjar M, Wheeler DS **S- Editor:** Ma YJ **L- Editor:** A  
**E- Editor:** Wu HL





Published by **Baishideng Publishing Group Inc**

8226 Regency Drive, Pleasanton, CA 94588, USA

Telephone: +1-925-223-8242

Fax: +1-925-223-8243

E-mail: [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com)

Help Desk: <http://www.wjgnet.com/esps/helpdesk.aspx>

<http://www.wjgnet.com>

