

# Early Liberal Fluid Therapy for Sepsis Patients Is Not Harmful: Hydrophobia Is Unwarranted but Drink Responsibly

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## EARLY INTERVENTIONS IMPROVE OUTCOME

Sepsis in the year 2016 remains the most expensive disease treated in hospitals and is the most common cause of in-hospital deaths in the United States (1). However, over the last 15 years, since the introduction of early goal-directed therapy (EGDT) and the Surviving Sepsis Campaign (SSC), there has been a consistent and historic reduction in mortality (2). The reduction from a historic mortality of 46.5% to less than 30% was validated when a trio of multinational trials named Protocolized Care for Early Septic Shock (ProCESS), Australasian Resuscitation in Sepsis Evaluation (ARISE), and Protocolized Management in Sepsis (ProMISe) “compared” various forms of resuscitation strategies (2, 3). This independently obtained historic mortality of 46.5% from an international task force of experts is identical to that of the original EGDT trial (2).

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Thus, it is absolutely clear that a protocolized approach consisting of early detection (lactate and fluid challenge), antibiotic therapy, source control, prevention of sudden cardiopulmonary events, and early hemodynamic optimization improves outcomes.

Even with unprecedented and replicated mortality benefit, many have proposed to disassemble the original EGDT trial and its components (4). ProCESS, ARISE, and ProMISe attempted to replicate and examine the efficacy of EGDT and have shown all time low mortalities, equal mortality reduction in all arms with no harm of EGDT. For some, these trials have made EGDT synonymous with an early liberal fluid strategy and its negative consequences (5–8). In rebuttal to our distinguished colleagues Genga and Russell (9); we advocate that treating early sepsis is not a time to be hydrophobic. Early fluid therapy in the context of a physiologically based protocol such as EGDT improves mortality for severe sepsis and septic shock.

## THE EBB AND FLOW PHASE OF FLUID MANAGEMENT

In 1942, Cuthbertson (10) described the metabolic response to inflammation, injury, and shock using the concept of the “ebb and flow” phase of critical illness.

“During the ebb-phase or resuscitation phase, there is low cardiac output, poor tissue perfusion and a cold and clammy patient. During the flow phase which is a staccato affair, the patient struggles to break from the grip of the ebb-phase which last about 3 days. Upon entering the flow-phase, the swollen patient has an increased cardiac output, normal tissue perfusion where diuresis occurs and body weight falls steady.”

This eloquent clinical description serves as the framework for the clinical principles of fluid management in early sepsis, **Figures 1 and 2**.

## MECHANISMS FOR THE DEVELOPMENT OF HYPOVOLEMIA IN SEPSIS

Sepsis-induced hypovolemia can be a result of vomiting (poor intake), diarrhea, sweating, edema, peritonitis, or

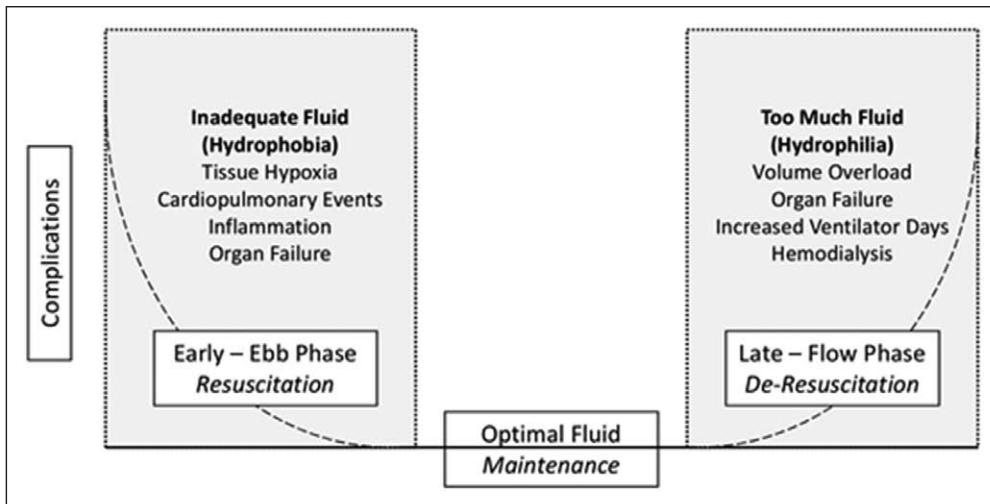


Figure 1. Optimal fluid therapy.

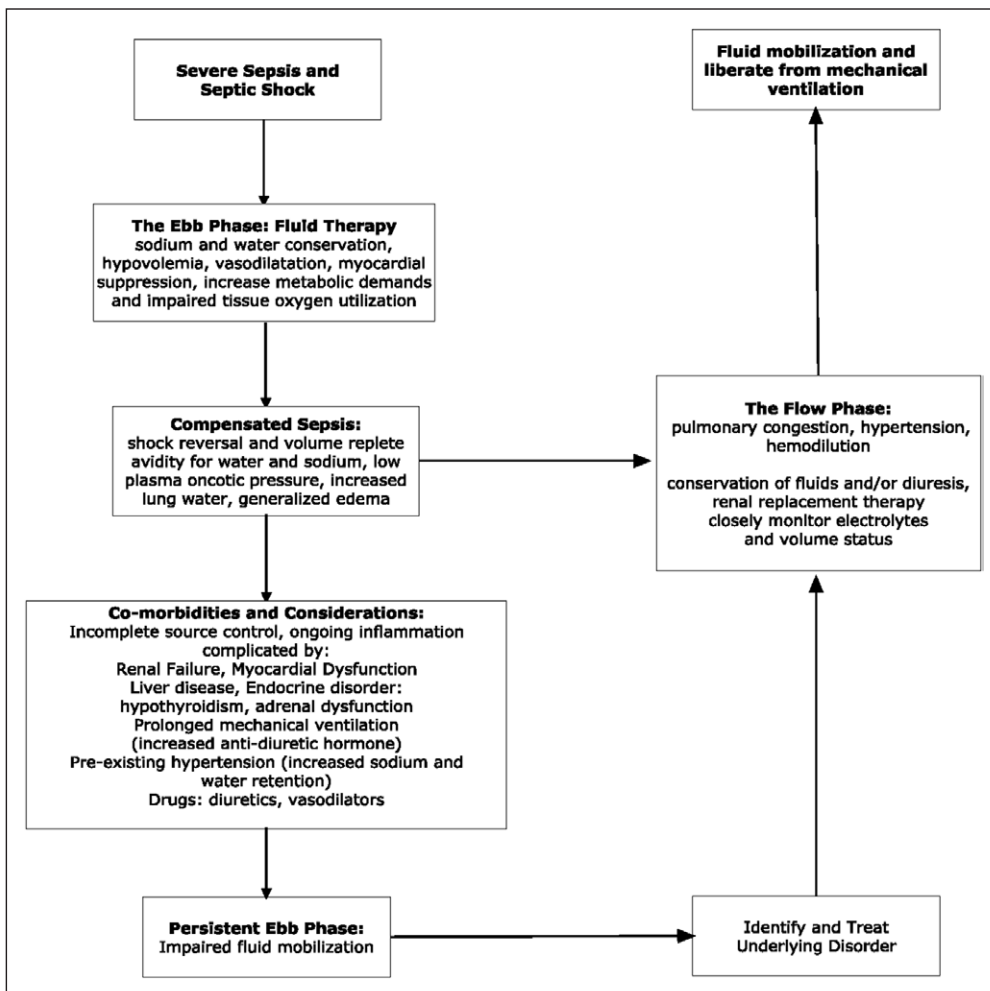


Figure 2. The dynamic landscape of fluid management. Reproduced from Schuller et al (11) and Rivers et al (12).

other exogenous losses. Further contributions to hypovolemia may result from vasodilatation, peripheral blood pooling, extravasation of fluid into the interstitial space, and increased capillary endothelial permeability. Hypovolemia

and water homeostasis and the dynamic “ebb and flow” landscape.

Microcirculatory changes observed in early sepsis such as acidosis, pyrexia, and increased RBC 2,3-diphosphoglycerate

and vasodilatation can be exaggerated in patients with comorbidities such as cardiovascular and renal disease where chronic therapy includes diuretics and afterload reduction. All of these mechanisms result in a decrease in intravascular volume which gives rise to a critical reduction in ventricular preload (central venous pressure [CVP]), ventricular diastolic pressure, stroke volume, cardiac output, systemic oxygen delivery, and an increase in systemic oxygen demands (decreased central venous oxygen saturation [ $ScvO_2$ ]) (13).

Compensatory responses as a reaction to decreased circulating blood volume are mediated by the activation of the neuroendocrine system. A redistribution of blood flow away from skeletal muscle beds and the splanchnic viscera supports vital organ blood flow to the heart and brain (14–16), with the augmentation of myocardial contractility leading to increased stroke volume (14). The constriction of arterial and venous capacitance vessels, particularly in the splanchnic bed, augments the venous return (14, 15). The activation of the renin-angiotensin axis releases aldosterone from the adrenal cortex. Changes in serum osmolarity lead to arginine-vasopressin release from the posterior pituitary. Both enhance fluid retention (15, 17–19). Comorbidities, such as congestive heart failure, renal failure, liver disease, and adrenal dysfunction, may modify this salt

occur, creating a local tissue environment to enhance the unloading of oxygen to tissues. Multiple factors may contribute to microvascular alterations, including driving pressure, alterations in RBC rheology, viscosity (local hematocrit), and leukocyte adhesion to endothelial cells, endothelial dysfunction, and interstitial edema. This is further complicated by the use of vasopressors. The use of techniques evaluating the microcirculation to optimize volume therapy remains to be validated (20, 21).

## EARLY FLUID THERAPY ATTENUATES THE EARLY PATHOGENESIS

The hemodynamic picture of early sepsis is hypotension, decreased CVP, decreased cardiac index, and decreased  $Scvo_2$  (13, 22). When the efficacy of antibiotics, cardiovascular support (fluids and dopamine titrated by intravascular monitoring to hemodynamic endpoints), and a combination of these two therapies in dogs with septic shock was compared, survival rates were 0%, 13%, 13%, and 43% in groups receiving no therapy (controls), antibiotics alone, cardiovascular support alone, or combined therapy, respectively (23). Although survivors and nonsurvivors in the combined therapy group required similar quantities of fluid therapy, nonsurvivors gained significantly more weight, suggesting abnormal vascular permeability with extravascular retention of fluids in the nonsurvivors indicating a more pronounced ebb phase. Thus, fluid overload is a result of the method of resuscitation, disease pathogenesis, and underlying comorbidities rather than the clinician's over prescription of fluid administration.

Early fluid therapy modulates initial inflammation. In human models of endotoxemia, isotonic prehydration significantly attenuates concentrations of proinflammatory cytokines (tumor necrosis factor- $\alpha$ , interleukin [IL]-8, and IL-1 $\beta$ ), while enhancing concentration of anti-inflammatory cytokines such as IL-10. This effect is associated with a reduction of endotoxin-induced symptoms and fever, while the endotoxin-induced changes in hemodynamic variables remain unchanged. More importantly, the peak activity of the inflammatory response is between 1 and 6 hours after introduction of the insult, which gives rise to the concept of early and late resuscitation as distinct therapeutic entities (24).

## DIAGNOSTIC, THERAPEUTIC, AND OUTCOME IMPLICATIONS OF FLUID THERAPY

Optimizing fluid therapy not only modulates inflammation and increases microvascular perfusion but also decreases the need for vasopressor therapy, steroid use, and more invasive monitoring (20, 24–26). When a clinician is confronted with a profoundly hypotensive patient with an infection, fluid therapy is indicated. However, the clinical assessment of volume status is insensitive, nonspecific, and one of the most challenging clinical assessments. In a post hoc analysis of the Fluids and Catheters Treatment Trial (FACTT) in treatment of acute

lung injury (ALI), this hypothesis was examined, which compared physical examination findings of ineffective circulation (capillary refill time > 2 s, skin mottling, and cool extremities) to variables obtained from pulmonary artery catheters. It was found that these physical examination findings are not useful predictors of a low cardiac index or low mixed venous oxygen saturation (27, 28).

## TITRATION OF FLUIDS—A WORK IN PROGRESS

Titration of fluid therapy is performed with numerous methodologies. Frequently, more than one method is required to make an accurate assessment. History, physical examination, dynamic, static and volumetric devices, and ultrasound and metabolic variables can be used, each with certain limitations. Inherent in measuring intravascular pressures is the inference that pressure equals volume. There are many instances for which the pressure within an intracardiac chamber may be elevated and the intravascular volume status may be diminished. The goal is to infuse “adequate” volume to restore perfusion before the onset of irreversible tissue damage without raising cardiac filling pressure to a level that produces hydrostatic pulmonary edema (29, 30). Hemodynamic monitoring used to accomplish these goals can vary and have not been shown to have outcomes superiority (31–33). Although much maligned as a predictor of fluid responsiveness, the general use of CVP in the treatment of severe sepsis and septic shock has been associated with improved outcomes (34–36).

For example, a patient with a CVP of 30 mm Hg, mean arterial pressure (MAP) of 70 mm Hg, heart rate of 80 beats/min, lactate of 5 mM/L, and normal hemoglobin and arterial oxygen saturation and  $Scvo_2$  of 44% would get an inotrope. Interestingly, this scenario is associated with increased fluid administration because the CVP is lowered with improved ventricular compliance (37). A patient with a CVP of 4 mm Hg, MAP of 70 mm Hg, heart rate of 80 beats/min, lactate of 5 mM/L, and normal hemoglobin and arterial oxygen saturation and  $Scvo_2$  of 44% would get fluid therapy. If CVP was taken in isolation in the first scenario, one would not assume fluid therapy is indicated and some may mistakenly use a diuretic. An elevated CVP has been associated with increased mortality as an interpretation of volume overload when in reality it may be myocardial suppression (38). Thus, using all endpoints in an algorithmic approach is the foundation of a comprehensive resuscitation (39). The discussion regarding CVP is best summarized by Sondergaard et al (35) who stated:

“Knowledge of the CVP is essential for the measurement of the volume state, the performance of the heart and the SVR. It enters considerations of heart, volume and power efficiency. The CVP provides a floating ground for the differential measurement of intravascular pressures. It does not inherently measure preload or the volume state but its measurement is essential to their calculation. Once the above principles are understood, precise control of the circulation becomes a straightforward mathematically predictable process.”



## FLUID THERAPY—EARLY

A hypotensive episode is associated with an increased risk of death, and the response to an adequate fluid challenge improves upon this discriminatory value for risk stratification (40–43). Early fluid therapy targeted to endpoints is associated with a decrease in systemic inflammation, vasopressor use, and mortality and must be distinguished from late aggressive fluid therapy (24, 40, 44–48). Multiple studies have shown that the fluid challenge of 30 mL/kg fluid volume within 3 hours of presentation is associated with increased MAP, normalization of ScvO<sub>2</sub>, and decreased vasopressor use at 6 hours. There is also a 19% relative reduction in in-hospital 30-day mortality and hospital length of stay (40, 42, 43). This benefit is realized even in patients with a history of renal and heart failure (43). The benefits of fluid administration are maximal when initiated within 30 minutes, making the fluid challenge an important aspect of early sepsis care. Lee et al (40, 49) concluded that “earlier fluid resuscitation may account for the lack of outcome differences between the original EGDT publication and the more recent three trials (ProCESS, ARISE, and ProMISe) and may have contributed to the overall low 60-day in-hospital mortality rate of 19%.”

From hospital arrival (prerandomization period) to the end of the 6-hour study period, the total fluid volume was similar and ranged from 3.5 to 5.5 L for the original EGDT publication and ProCESS, ARISE, and ProMISe study groups. Because of the greater lead time prior to enrollment into these trials, between 2 and 2.6 L of fluid was given prior to randomization. There is striking similarity in the amount of fluid given in all trials that approximates the intravascular volume of 5 L. However, the comparative differences in fluid therapy was 1,482 mL (42.4%), 697 mL (16%), 175 mL (4.1%), and 229 mL (4.4%) between the EGDT and usual or control care treatment groups in the EGDT, ProCESS, ARISE, and ProMISe trials, respectively (Table 1).

In the EGDT study, this significantly greater volume therapy or treatment effect during the resuscitation (prerandomization and first 6 hr) was associated with a greater reduction (13.8%) in vasopressor therapy, less volume therapy (2 L or 23%), and lower mechanical ventilation rates (14.2%) between the EGDT and control group over the subsequent 6–72-hour time period. These results were observed in the absence of aggressive glucose control, steroid use, protective lung strategies, and conservative fluid management strategies. These were present during the conduction of the ProCESS, ARISE, and ProMISe trials.

## VASOPRESSOR THERAPY

ProCESS, ARISE, and ProMISe trials used vasopressor therapy twice as much compared to the EGDT trial. The more frequent administration of vasopressors in ProCESS, ARISE, and ProMISe may result in a hemodynamic phenotype of “vasodilatory septic shock” which is associated with a lower mortality risk as described by Hernandez et al (50). These findings may further indicate that early vasopressor administration instead of continued fluid therapy may be beneficial once the resuscitation is

complete (51). Thus, it seems that ProCESS, ARISE, and ProMISe showed us that 4–6 L of fluid is generally required for early hemodynamic optimization.

The introduction of early vasopressor use is intriguing but challenging (51). Vasodilatory shock after adequate volume resuscitation is the requisite for the physiologic use of corticosteroids. This definition will be altered when vasopressors are introduced at an earlier stage before adequate fluid resuscitation. Waechter et al (52) reports that vasopressor use in the first hour may be associated with increased mortality in patients with greater illness severity. Vasopressor use also increases CVP which may impact volume therapy when this endpoint is artificially elevated (53).

Vasopressor therapy elevates the intensity of patient care because vasopressors require monitoring (i.e., intraarterial pressure monitoring). By placing a patient on vasopressors when adequate volume therapy may have reversed the hypotension, one is placing the patient in a higher level of care and potentially increasing healthcare resource consumption. With EGDT, vasopressor use diminishes by 13.8% over the subsequent 72 hours.

## FLUID THERAPY—LATER

When comparing the original EGDT to ProCESS, ARISE, and ProMISe between 6 and 72 hours, significantly more volume therapy (almost two-fold) was administered in the EGDT study in both treatment groups compared to the three follow-up trials. There was an approximate two-fold increase in vasopressor therapy in ProCESS, ARISE, and ProMISe compared to the EGDT trial during this time period.

ProCESS, ARISE, and ProMISe had mechanical ventilation rates approximately half that of the EGDT and the reference literature (54–56). For example, the SSC database from 2005 to 2008 reports a mechanical ventilation rate of 52.4% (7,877/15,022 of patients), which is almost identical to the EGDT study. Mortality rates in this report were 48.3%, 45.7%, and 33.0% in mechanically ventilated patients with ALI, without ALI, and without mechanical ventilation, respectively. The use of protective lung strategies and conservative fluid management strategies for ALI was not used in the EGDT study as it preceded these trials.

The FACTT trial isolated the manipulation of volume therapy as a controlled intervention which began an average of 43 hours after ICU admission and 24 hours after the establishment of ALI (11). Although there was no difference in 60-day mortality, patients in the conservative strategy group had significantly improved lung and CNS function (decreased need for sedation) along with mechanical ventilation and thus ICU care. There was a statistically significant 0.3-day increase in cardiovascular failure free days in the liberal compared to the conservative fluid group. The increased early volume in the EGDT led to a decreased need for mechanical ventilation over the first 72 hours of hospitalization. This may be due to the modulating effects on IL-8, which has been identified as an ALI culprit within the first 72 hours of presentation (57, 58).

The findings of the FACTT trial brought attention to the negative consequences of the overuse of fluid administration.

**TABLE 1. Comparison of Treatments in Sepsis Trials**

Therapeutic Intervention	Protocolized Care for Early Septic Shock					Australasian Resuscitation in Sepsis Evaluation		Protocolized Management in Sepsis	
	EGDT		Protocol-Based Standard Therapy			EGDT	UC	EGDT	UC
	EGDT	Control	EGDT	Protocol-Based Standard Therapy	UC				
Fluid from emergency department arrival to 6 hr <sup>a</sup> , mL	4,981	3,499	5,059	5,511	4,362	4,479	4,304	4,216	3,987
Difference between groups <sup>b</sup> , mL	1,482		-452 and 667			175		229	
Difference between groups (%)	42.4		16			4.1		4.4	
Fluids 6–72 hr, mL	8,625	10,602	4,458	4,918	4,354	4,274	4,382	4,215	4,366
Total fluids 0–72 hr, mL	13,443	13,358	7,253	8,193	6,663	6,906	6,672	5,946	5,844
Vasopressor 0–6 hr, %	27.4	30.3	54.9	52.2	44.1	66.6	57.8	53.3	46.6
Vasopressor 6–72 hr, %	29.1	42.9	47.6	46.6	43.2	58.8	51.5	57.9	52.6
Vasopressor 0–72 hr, %	36.8	51.3	60.4	61.2	53.7			60.5	55.0
Inotrope 0–6 hr, %	13.7	0.8	8.0	1.1	0.9	15.4	2.6	18.1	3.8
Inotrope 6–72 hr, %	14.5	8.4	4.3	2.0	2.2	9.5	5.0	17.7	6.5
Mechanical ventilation 0–6 hr, %	53.0	53.8	26.4	24.7	21.7	34.8 <sup>c</sup>	32.9 <sup>c</sup>	20.2	19.0
Mechanical ventilation 6–72 hr, %	2.6	16.8	33.7	31.4	27.9	38.6 <sup>c</sup>	40.6 <sup>c</sup>	24.4	25.4
Any mechanical ventilation, %	55.6	70.6	36.2	34.1	29.6	30.0	31.5	27.4	28.5
Steroids prerandomization, %	None	None	9.3	9.4	8.3			5	4
Steroids 0–6 hr, %	None	None	12.3	10.8	8.1			11.7	11.5
Any steroids 72 hr, %	None	None				36.9	35.9	21.9	21.1

EGDT = early goal-directed therapy, UC = usual care.

<sup>a</sup>The prerandomization period refers to a time frame prior to the time informed consent for study enrollment. Interventions were initiated as indicated.

<sup>b</sup>Difference between groups are early goal-directed therapy minus the treatment group in each trial; Prerandomization and 6 hr of study.

<sup>c</sup>Combined invasive and noninvasive mechanical ventilation.

In order to generalize these results and avoid mitigating the salutary findings, multiple variables must be considered when applying a conservative fluid management approach. The exclusion of patients on hemodialysis, overt renal insufficiency, heart failure, and the relatively young age of the patients studied (age, 50 yr) make the FACTT trial a departure from the reality that many clinicians will face in the treatment of severe sepsis and septic shock.

## FLUID REMOVAL-OPTIMAL TIMING

Although pathogenically well described, the clinical landmark that separates the ebb from flow phase is frequently indistinct and complex. It requires meticulous attention to past medical history (i.e., renal failure and congestive heart failure), cumulative fluid balance, relevant hemodynamic

variables, laboratory findings (hemodilution and renal function), and physical examination findings of fluid overload, Figure 2. In the absence of early recognition of the flow phase, the complications of pulmonary edema, myocardial complications, respiratory insufficiency, and the continued need for ventilatory support results. Fluid conservation, diuretic therapy, and the institution of renal replacement therapy are clinical decisions made in the flow phase especially in patients with renal insufficiency and cardiac disease (59). When renal replacement therapy is required in the treatment of septic shock, mortality approaches 50%. The optimal timing of initiating renal replacement therapy in the presence of acute kidney injury is not clearly established (60–62).

The body of observational evidence over the last 10 years points towards the deleterious effects of persistently positive

fluid balances in septic shock. Nevertheless, only one large clinical trial has demonstrated the efficacy and safety of fluid removal in ICU patients without cardiovascular dysfunction. The best intervention for active fluid management may be the avoidance of unnecessary fluid loading with the best evidence-based approaches, whereas active fluid withdrawal may have a later role. More evidence from randomized clinical trials is needed to guide physicians in the art and science of late fluid management in septic shock (63).

## HYDROPHOBIA IS UNWARRANTED IN EARLY SEPSIS CARE

- Early titrated fluid administration modulates inflammation, improves microvascular perfusion, organ function, and outcomes.
- Fluid therapy is determined by the method of resuscitation along with various clinically measured variables, disease pathogenesis, and underlying comorbidities rather than the clinician's over prescription of fluid administration.
- Contrary to popular belief, recent EGD<sup>T</sup> septic shock trials used similar amounts resuscitation fluids in the acute phase when compared to the original EGD<sup>T</sup> trial. This was associated all time lows in mortality.
- Patients who suffer the negative consequences of fluid overload are already at high risk because of preexisting and acquired comorbidities.
- A comprehensive understanding of "ebb and flow" combined with closely monitored early fluid therapy, accumulation and timely removal is of most importance.
- Fluid therapy should be treated as any drug which involves consideration of the type, dose, duration of treatment, and toxicity (64).
- Hydrophobia may be synonymous with rabies but should not be with early sepsis management.

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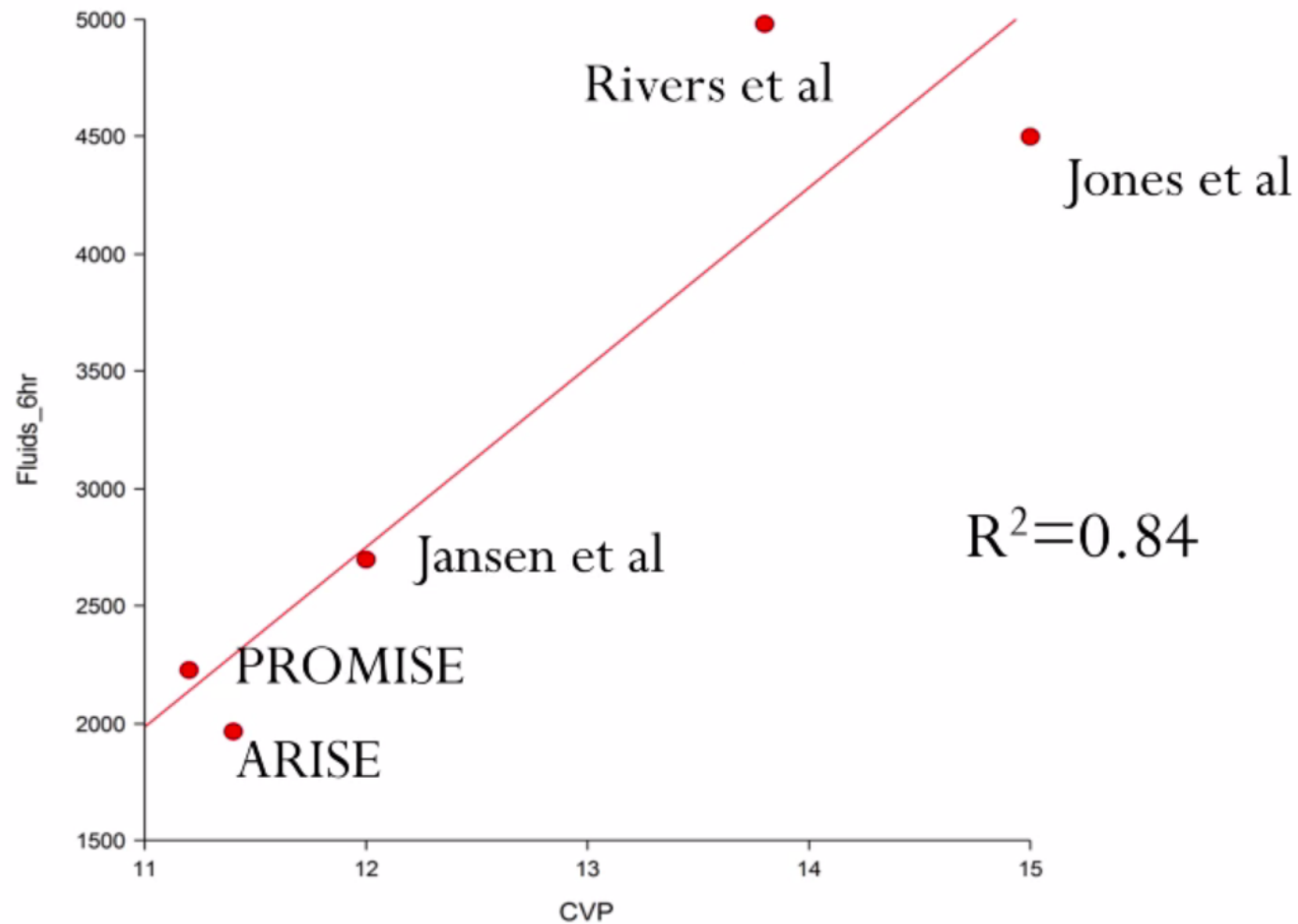
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# Fluids administered 0-6 hr and CVP at 6 hours

## EGDT Arm 2001-2015





# Fluids administered 0-72 hr and 90 day mortality

## Control arm of EGDT Studies 2001-2015

