



A STUDY OF PROGNOSTIC VALUE OF CENTRAL VENOUS OXYGEN SATURATION IN PATIENTS WITH SEPTIC SHOCK AT A TERTIARY CARE HOSPITAL IN SOUTH GUJARAT

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ABSTRACT **Purpose of study :** Aim of the study is to measure ScvO₂ level at Baseline, 12 hours and 24 hours in patients presenting with septic shock & their correlation with mortality. **Introduction:** Severe sepsis and septic shock, is a common cause of emergency room admission and is associated with high morbidity and mortality worldwide. When oxygen delivery fails to meet tissue oxygen demand in critical illness, tissues maintain their oxygen utilization by extracting more oxygen which ultimately decreases central venous oxygen saturation (ScvO₂). As ScvO₂ decreased, the mortality tends to increase. This suggests that ScvO₂ is a useful prognostic marker in septic shock. **Methodology:** This is a prospective observational study of 30 patients with septic shock admitted in ICU of our tertiary care hospital. Patients are selected by SOFA score ≥ 2 and thorough general examination. 2nd step is to measure ScvO₂ at baseline and 12 hours apart. Final analysis has been done with the help of Open EPI and SPSS software. **Results:** Out of 30 patients, 13 (43%) did not survive for more than 7 days. Patients with ScvO₂ < 50 at any point of time was associated with poor prognosis in septic shock. **Conclusion:** Among patients with septic shock, it is better to monitor serial ScvO₂ rather than going for single value. Trend of ScvO₂ was also indicative of prognosis in septic shock. It is concluded that ScvO₂ can be used as an independent predictor of survival in patients with septic shock.

KEYWORDS : Septic shock, ScvO₂ : central venous oxygen saturation

INTRODUCTION:

Septicemia implies active replication of bacteria in the blood associated with systemic manifestations. Septic shock is characterized by hypotension (systolic BP < 90mmHg), hypoxia, increased serum lactate levels, high-anion-gap metabolic acidosis and oliguria with urine output < 30 ml/h[1]. In critical illness, the cardiovascular response to an increase in oxygen requirement is an increase in arterial oxygen content or an increase in CO (cardiac output). When these compensatory mechanisms fail to occur, CO and systemic oxygen delivery decreases; tissue extraction of oxygen will undergo a compensatory increase in order to sustain systemic oxygen consumption leading to a reduction in ScvO₂. Thus ScvO₂ can provide useful information regarding the adequacy of CO over time. So, ScvO₂ has long been studied as a prognostic marker and resuscitation endpoint in patients with septic shock[3]. Rivers et al.[4] showed that an early therapeutic strategy that includes aiming for the rapid normalization of ScvO₂ ($\geq 70\%$) in patients suffering from septicemia or septic shock at presentation to the emergency department could improve survival.[5-10]

METHODOLOGY:

The study was done at Surat Municipal Institute of Medical Education & Research (SMIMER) Hospital, Surat, Gujarat, from year June 2020 to Nov 2021. This is a prospective observation study of total 30 ICU patients of septic shock in our tertiary care hospital. Patients with presumed septic shock with SOFA score ≥ 2 ; age ≥ 18 yrs; willing for admission and willing to participate in study were included in the study after informed written consent. Patients having Cardiogenic shock, Neurogenic shock, Anaphylactic shock, Hypovolemic shock; Patients with chronic lung diseases, cyanotic heart diseases, not willing for admission and not willing to participate in study were excluded. Approval for this study was taken in institutional ethical committee. All necessary confidentiality of participants were maintained. Detail history, examination and necessary investigations were done for each participants. ScvO₂ at baseline and 12 hours apart done by ABL 800 basic analyzer. Data was entered in MS EXCEL spread sheet and was analyzed with the help of Open EPI and SPSS software. Statistical analysis was done by appropriate statistical method.

RESULT:

Out of total 30 patients, 6(20%) were having ARDS, 6(20%) were having community acquired pneumonia(CAP), 5(16.67%) were having septicemia, 5(16.67%) were having UTI+pyelonephritis, 1(3.33%) was having inflammatory bowel disease, 5(16.67%) were

having spontaneous bacterial peritonitis and 2(6.66%) were having meningitis. Most of the patients were having ARDS and CAP.

Out of 30 patients, highest number of patients were in the age group of 21-40 years. Out of 30 patients, 13 (43.3%) expired within 7 days and 17 (56.7%) survived for more than 7 days. Highest mortality (16.7%) was seen in 21-40 years of age group.

In our study, male:female ratio was 8:7. Among 13 expired patients, 5(16.7%) were male and 8(26.6%) were female. Among 17 survived patients, 11(36.7%) were male and 6(20%) were female.

Table 1. The Sequential Organ Failure Assessment (SOFA) Score*

Variables	SOFA Score				
	0	1	2	3	4
Respiratory Pao ₂ /Fio ₂ , mm Hg	>400	≤ 400	≤ 300	≤ 200 †	≤ 100 †
Coagulation Platelets $\times 10^3/\mu\text{L}$	>150	≤ 150	≤ 100	≤ 50	≤ 20
Liver Bilirubin, mg/dL‡	<1.2	1.2-1.9	2.0-5.9	6.0-11.9	>12.0
Cardio-vascular Hypotension	No hypotension	Mean arterial pressure <70 mm Hg	Dop ≤ 5 or dop (any dose)§	Dop >5, epi ≤ 0.1 , or norepi ≤ 0.1 §	Dop >15, epi >0.1, or norepi >0.1§
Central nervous system Glasgow Coma Score Scale	15	13-14	10-12	6-9	<6
Renal Creatinine, mg/dL, or urine output, mL/§	<1.2	1.2-1.9	2.0-3.4	3.5-4.9 or <300	>5.0 or <200

*Norepi indicates norepinephrine; Dob, dobutamine; Dop, dopamine; Epi, epinephrine; and Fio₂, fraction of inspired oxygen.
†Values are with respiratory support.
‡To convert bilirubin from mg/dL to $\mu\text{mol/L}$, multiply by 17.1.
§Adrenergic agents administered for at least 1 hour (doses given are in $\mu\text{g/kg per minute}$).
¶To convert creatinine from mg/dL to $\mu\text{mol/L}$, multiply by 88.4.

Figure 1 : SOFA Score

Table-1 Trend of Central venous oxygen saturation (ScvO₂) in survived & expired patients:

Central venous oxygen saturation(ScvO ₂)	Survived	Expired
	Mean \pm SD	Mean \pm SD
0 Hour	64.91 \pm 1.74	47.64 \pm 2.03
12 Hour	65.3 \pm 0.934	44.84 \pm 1.49
24 Hour	68.21 \pm 2.47	42.2 \pm 0.744

ScvO₂ trend was decreasing in expired patient & increasing in survived patients, thus decreasing trend of ScvO₂ had bad prognosis in our study. We observed in our study that ScvO₂ < 50 at any point of time was associated with poor prognosis in septic shock. Trend of ScvO₂ was also indicative of prognosis in septic shock. Decreasing trend of ScvO₂ was usually associated with poor outcome in patients with septic shock.

DISCUSSION:

Central venous oxygen saturation has diagnostic, therapeutic as well as prognostic value in critically ill patients. As a diagnostic, derangement in ScvO₂ can ascertain the underlying etiology. The oxygen consumption by the tissues of whole body is usually independent of oxygen delivery (DO₂). ScvO₂ is a measure of the oxygen content of the blood returning to the right side of the heart after perfusing the entire body. When the oxygen supply is insufficient to meet the metabolic demands of the tissues, an abnormal ScvO₂ ensues and reflects an inadequacy in the systemic oxygenation. ScvO₂ is, therefore, dependent on oxygen delivery and oxygen extraction. In our study we had measured ScvO₂ in patients with septic shock. Mean value of ScvO₂ at 0 Hour, 12 Hours and 24 Hours in survived patients were 64.91, 65.3 and 68.21. Mean value of ScvO₂ at 0 Hour, 12 Hours and 24 Hours in expired patients were 47.64, 44.84 and 42.2. According to the study published in *Journal of anesthesia & intensive care medicine*¹¹, the overall mean ScvO₂ in non-survivors and survivors was 53.34 ± 4.08 and 73.33 ± 5.03 respectively, which is comparable to our study. On observing the ScvO₂ trend, Serial ScvO₂ levels were increasing in survived patients while serial ScvO₂ levels were decreasing in expired patients. There was correlation between ScvO₂ at 0, 12, and 24 hours with mortality in our study (P value <0.05). Present study had near perfect agreement with ScvO₂ level for mortality prediction (AUC >0.5). We can see that as ScvO₂ increased, the mortality tends to reduced. This suggests that ScvO₂ is a useful prognostic marker in patients with septic shock.

CONCLUSION:

Our study confirms the prognostic value of serial ScvO₂ monitoring and its trends for prediction of mortality. If ScvO₂ levels are static or increasing then prognosis is good and prognosis is grave if it is decreasing. Our study indicates that it is better to monitor serial ScvO₂ level rather than going for single value. We concluded that ScvO₂ can be used as an independent predictor of survival in patients with septic shock.

REFERENCES:

1. Lee SM, An WS. New clinical criteria for septic shock: Serum lactate level as new emerging vital sign. *J Thorac Dis.* 2016;8(7):1388–90.
2. Gil Cebrian J, Bello Cámara MP, Diaz-Alersi R. Apache Ii. *Intensive Care Med.* 1987;13(2):143.
3. Rady MY, Rivers EP, Nowak RM: Resuscitation of the critically ill in the ED: responses of blood pressure, heart rate, shock index, central venous oxygen saturation, and lactate. *Am J Emerg Med* 1996, 14:218–225
4. Rivers E, Nguyen B, Havstad S, Ressler J, Muzzin A, Knoblich B, Peterson E, Tomlanovich M, Early Goal-Directed Therapy Collaborative Group: Early Goal-Directed Therapy Collaborative Group: early goal directed therapy in the treatment of severe sepsis and septic shock. *N Engl J Med* 2001,345:1368–1377.
5. Dellinger RP, Levy MM, Rhodes A, Annane D, Gerlach H, Opal SM, Sevransky JE, Sprung CL, Douglas IS, Jaeschke R, Osborn TM, Nunnally ME, Townsend SR, Reinhart K, Klempell RM, Angus DC, Deutschman CS, Machado FR, Rubenfeld GD, Webb SA, Beale RJ, Vincent JL, Moreno R, Surviving Sepsis Campaign Guidelines Committee including the Pediatric Subgroup: Surviving Sepsis Campaign Guidelines Committee including the Pediatric Subgroup. Surviving sepsis campaign: international guidelines for management of severe sepsis and septic shock: 2012. *Crit Care Med* 2013,41:580–637.
6. Bellomo R, Reade MC, Warrillow SJ: The pursuit of a high central venous oxygen saturation in sepsis: growing concerns. *Crit Care* 2008, 12:130.
7. Perel A: Bench-to-bedside review: the initial hemodynamic resuscitation of the septic patient according to Surviving Sepsis Campaign guidelines—does one size fit all? *Crit Care* 2008, 12:223.
8. Peake S, Webb S, Delaney A: Early goal-directed therapy of septic shock: we honestly remain skeptical. *Crit Care Med* 2007, 35:994–995.
9. Textoris J, Fouché L, Wirusus S, Antonini F, Tho S, Martin C, Leone M: High central venous oxygen saturation in the latter stages of septic shock is associated with increased mortality. *Crit Care* 2011, 15:R176
10. van Beest PA, Hofstra JJ, Schultz MJ, Boerma EC, Spronk PE, Kuiper MA: The incidence of low venous oxygen saturation on admission to the intensive care unit: a multi-center observational study in The Netherlands. *Crit Care* 2008, 12:R33.
11. Central Venous Oxygen Saturation as a Surrogate Marker for Outcome in Critically ill Patients - A Prospective Observational Cohort Study. jupiperpublishers.com/jaicm/JAICM.MS.ID.555698.php