## Abstract

**Introduction:** The early identification and scrupulous monitoring of tissue dysoxia can improve the management of critically-ill patients. In this light, the final product of aerobe and anaerobe metabolism (that is, carbon dioxide) can provide useful information on adequacy of tissue perfusion and metabolism (9,10). *The aim of our study* was to evaluate whether the venous-arterial PCO2 gradient provides useful information on tissue dysfunction in patients admitted to the ICU

<u>Methods</u>: We prospectively studied 50 patients admitted to ICU in 2012/2013 with length of stay (LOS) >24 hours. A sample of arterial and venous blood was taken for gas analysis at admission. Venous-arterial PCO2 gradient ( $\Delta$ pCO2), organ dysfunction in the first 24 hours and ICU mortality were collected. Organ dysfunction was defined as a SOFA score  $\geq 2$  for each organ. The patients, whether ventilated or not, were subdivided and compared on the basis of  $\Delta$ pCO2 value:  $\Delta$ pCO2  $\geq 6$  mmHg (*Higher* group) and  $\Delta$ pCO2 <6 mmHg (*Normal* group).

**<u>Results:</u>** Twenty-nine patients (58%) showed a  $\Delta pCO_2 \ge 6$  mmHg (<u>*Higher*</u> group) and twenty-one patients (42%) showed a  $\Delta pCO_2 \le 6$  mmHg (<u>*Normal*</u> group). The *Higher* group showed a larger rate (34%) of cardiovascular dysfunction than the *Normal* group (8%)(*P*< 0.05). Respiratory dysfunction was observed in 54% of the patients of the High group and only in 32% of the Normal group. Similarly, renal dysfunction was also slightly larger in the Higher group (26%) than in the Normal group (20%) (*P* > 0.05). As expected, patients of *Higher* group showed more complications (52%) than *Normal* group (32%) (*P* > 0.05) and ICU mortality (40%) three times larger than *Normal* group (12%) (*P* < 0.05).

<u>Conclusions & Recommendations</u>: Despite its limitations, The above data support the hypothesis that  $\Delta pCO_2$  can provide useful information on the tissue perfusion and metabolism in ICU patients and can be used as a reliable biomarker for early prediction of organ dysfunction and outcome in critically-ill patients. But, further studies on a larger number of patients are needed to confirm its reliability.

*Limitations*: Our study was done on a small sample size and based on a pre-defined set of study parameters, which might not have reflected the true nature of general changes observed in sepsis.

<u>*Key words:*</u> Dysoxia - Shock - Delta PCO2 - Haemodynamics - Scoring systems - Organ dysfunction - Clinical outcome.

