

## ASSESSMENT OF MICROCIRCULATION USING REMOTE PHOTOPLETHYSMOGRAPHY AND AUTOMATED CAPILLARY REFILL TIME IN CRITICALLY ILL PATIENTS

Mara Klibus<sup>1</sup>, Veronika Eunapu<sup>2</sup>, Uldis Rubins<sup>3</sup>, Zbignevs Marcinkevics<sup>3</sup>, Andris Grabovskis<sup>3</sup>, Indulis Vanags<sup>4</sup>, Oļegs Sabeļņikovs<sup>5</sup>

<sup>1</sup>Department of Doctoral Studies, Rīga Stradiņš University; Clinic of Anaesthesiology and Intensive Care, Pauls Stradins Clinical University Hospital; Department of Clinical Skills and Medical Technologies, Rīga Stradiņš University

<sup>2</sup>Department of Clinical Skills and Medical Technologies, Rīga Stradiņš University

<sup>3</sup>University of Latvia

<sup>4</sup>Clinic of Anaesthesiology and Intensive Care, Pauls Stradins Clinical University Hospital

<sup>5</sup>Clinic of Anaesthesiology and Intensive Care, Pauls Stradins Clinical University Hospital; Department of Clinical Skills and Medical Technologies, Rīga Stradiņš University

**Objectives.** Microcirculation assessment during fluid resuscitation of septic shock is challenging due to lack of objective clinical tests. New methods for evaluation of microcirculation monitoring have been developed—remote photoplethysmography (rPPG) and automated objective capillary refill time measurement technique (aCRT).

**The aim:** Assess rPPG and aCRT methods as an alternative method for microcirculation evaluation.

**Materials and Methods.** patients with positive passive leg raising test (PLRT) were initially resuscitated with crystalloid. Patients were divided into 2 groups: COVID-19 (n = 18) and bacterial septic shock (BSS) (n=16). Hemodynamic variables, *manual capillary refill time* (mCRT) and *aCRT parameters* (T90 – time when 90% of capillary refill is over, Tst – time when capillary refill is fully over), peripheral perfusion index (PPI) detected using rPPG were collected before and after PLRT and after volume expansion (VE).

**Results.** In COVID-19 mean PPI increased during PLRT by 7% (from  $43 \pm 27$  to  $46.5 \pm 29.1$ ), by 15% after VE (from  $43.0 \pm 27.8$  to  $49.5 \pm 22.6$ ), while in BSS PPI increased during PLRT by 18% (from  $28.3 \pm 20.9$  to  $33.6 \pm 25.3$ ), by 28% after VE (from  $28.3 \pm 20.0$  to  $36.3 \pm 25.8$ ). Mean mCRT in COVID-19 decreased by 22% during PLRT ( $2.57 \pm 0.59$  to  $1.98 \pm 0.68$ ), by 22% after VE (from  $2.57 \pm 0.59$  to  $1.98 \pm 0.78$ ), while in BSS decreased by 31% during PLRT (from  $1.85 \pm 0.64$  to  $1.29 \pm 0.38$ ), by 32% after VE (from  $1.85 \pm 0.64$  to  $1.26 \pm 0.29$ ). Mean aCRT T90 in COVID-19 decreased by 32% during PLRT (from  $1.74 \pm 1.16$  to  $1.17 \pm 0.79$ ), by 17% after VE (from  $1.74 \pm 1.16$  to  $1.45 \pm 1.06$ ), while in BSS decreased by 41% during PLRT (from  $1.93 \pm 1.03$  to  $1.38 \pm 0.79$ ), by 8% after VE (from  $1.93 \pm 1.03$  to  $1.78 \pm 0.66$ ). Mean Tst in COVID-19 decreased by 21% during PLRT (from  $3.33 \pm 1.59$  to  $2.63 \pm 1.37$ ), by 10% after VE (from  $3.33 \pm 1.59$  to  $3.03 \pm 1.44$ ) while in BSS decreased by 25% during PLRT (from  $3.74 \pm 1.24$  to  $2.81 \pm 1.22$ ) by 2% after VE (from  $3.74 \pm 1.24$  to  $3.69 \pm 1.12$ ).

**Conclusions.** This study results shows that rPPG and aCRT are potentially applicable to assess microcirculation in critically ill patients.