

LETTER TO THE EDITOR

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Letter to editor

Antonio Messina^{1,2*} and Maurizio Cecconi^{1,2}



Dear Editor,

In a recent published paper, our group investigated the pharmacodynamic effect of a fluid challenge (FC) of a 4 ml/kg of crystalloids infused over 10 (FC₁₀) or over 20 min (FC₁₀), with the purpose of assess whether the fluid responsiveness [defined as an increase in the stroke volume index (SVI) \geq 10%] was affected by the time of infusion [1].

Considering the effect of the different time of infusion on both pressure and flow variables, in a secondary analysis of the paper, we calculated the percent changes of systolic arterial pressure (Δ SAP) and SVI (Δ SVI) from baseline to the half of the infusion of FC₁₀ and FC₂₀ and assessed the correlation by linear regression and the reliability of SAP changes in predicting fluid responsiveness by considering the area under (AUC_{ROC}) the receiver operating characteristic (ROC) [95% confidence interval (95% CI)]. The grey zones for all the statistically significant ROC curves were also calculated considering the low cut-off value including 90% of negative FC responses and a high cut-off value predicting positive FC in 90% of cases [2].

There was a significant positive correlation between ΔSAP and ΔSVI from baseline to ½ FC administration, during both FC₁₀ ($r^2=0.50$; p<0.0001; slope = 0.70 \pm 0.10) and FC₂₀ ($r^2=0.28$; p=0.01; slope = -0.53 ± 0.12) administrations (Fig. 1). However, the ROC curve of the changes in SAP after 1/2 FC₁₀ was significant (p=0.01) [AUC_{ROC} = 0.72 (95% CI 0.55–0.85); gray zone

10%/0%]. On the contrary, the ROC curve of the changes in SAP after ½ FC₂₀ was not statistically significant (p=0.11). Our results show a positive moderate linear correlation ($r^2=0.50$) between for Δ SAP and Δ SVI FC₁₀. The associated ROC curve constructed showed that Δ SAP > 10% is highly suggestive of FC response (i.e., sensitivity > 90%), whereas Δ SAP = 0 is highly suggestive of no response (i.e., specificity > 90%). On the contrary, the linear correlation of FC₂₀ was weak ($r^2=0.28$), and the ROC curve was insignificant.

The interplay between SAP and SVI is based on a complex balance between cardiac factors, arterial load, and resistance [3, 4]. In fact, the physiological relationship between pressure and flow variables is not linear, and SAP changes should not be used as a perfect surrogate for SVI to predict the effect of FC [5, 6]. Nevertheless, Δ SAP, only after FC₁₀, still maintains clinical utility, suggesting that no increase after ½ FC is associated with no fluid responsiveness, whereas an increase of at least 10% is associated to fluid responsiveness. In contexts of low resources or when a SVI monitoring is not available, our results may provide practical cut-offs to guide fluid optimization in elective surgical patients. This finding, however, could be partially dependent on the intrinsic mathematical coupling between pressure and flow variables changes after the fluid infusion, since the Most-Care® system is based on the high sample rate analysis of the arterial waveform.

Respectfully Yours.

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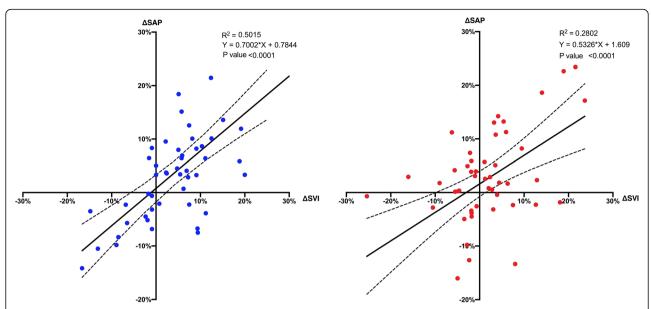


Fig. 1 Linear regression between the changes in SVI (Δ SVI) and in SAP (Δ SAP) after ½ fluid challenge administration in the FC₁₀ (blue dots, left panel) and FC₂₀ (red dots, right panel). Dashed lines represent 95% confidence intervals for the regression line (solid line)

Authors' contributions

Antonio Messina and Maurizio Cecconi conceived the idea for the manuscript and drafted, wrote, and approved the final version of the manuscript.

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Declarations

Competing interests

The authors declare that they have no competing interests.

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References

- Messina A, Palandri C, De Rosa S, Danzi V, Bonaldi E, Montagnini C et al (2021) Pharmacodynamic analysis of a fluid challenge with 4 ml kg(-1) over 10 or 20 min: a multicenter cross-over randomized clinical trial. J Clin Monit Comput 8:1–11. https://doi.org/10.1007/s10877-021-00756-3
- Cannesson M, Le Manach Y, Hofer CK, Goarin JP, Lehot JJ, Vallet B et al (2011) Assessing the diagnostic accuracy of pulse pressure variations for the prediction of fluid responsiveness: a "gray zone" approach. Anesthesiology 115(2):231–241. https://doi.org/10.1097/ALN.0b013e318225b80a
- O'Rourke MF (1982) Vascular impedance in studies of arterial and cardiac function. Physiol Rev 62:570–623. https://doi.org/10.1152/physrev.1982.62.2. 570
- Nichols WW, O'Rourke MF, Avolio AP, Yaginuma T, Murgo JP, Pepine CJ, Conti CR (1985) Effects of age on ventricular-vascular coupling. Am J Cardiol 55(9):1179–1184. https://doi.org/10.1016/0002-9149(85)90659-9
- Monge Garcia Ml, Saludes Orduna P, Cecconi M (2016) Understanding arterial load. Intensive Care Med 42(10):1625–1627. https://doi.org/10.1007/ s00134-016-4212-z
- Monge Garcia MI, Barrasa Gonzalez H (2017) Why did arterial pressure not increase after fluid administration? Med Intensiva 41(9):546–549. https://doi. org/10.1016/j.medin.2017.03.005

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