

Outcomes in Patients with Vasodilatory Shock and Renal Replacement Therapy Treated with Intravenous Angiotensin II: Erratum

In the article beginning on page 949 in the June 2018 issue of Critical Care Medicine, there were several errors.

- 1. Figure 1 Survival through day 28 Replaced: HR 0.515 (0.304, 0.817) With: HR 0.515 (0.304, 0.871)
- 2. Page 952, Outcomes

Added: Using cumulative incidence estimates to adjust for death as a competing risk, patients in the Ang II group were more likely to discontinue RRT within 7 days (**un**adjusted HR, 2.90; 95%, 1.29–6.52; p = 0.007)

Supplemental digital content data corrections:

- 3. **Supplemental Table 5** Summary of post-hoc analysis baseline demographics and disease characteristics Replaced: Screening mean arterial pressure (mmHg), placebo median (IQR) 65.4 (68.9-67.5) With: Screening mean arterial pressure (mmHg), placebo median (IQR) 65.4 (61.9–67.5)
- Supplemental Table 7 Mean norepinephrine equivalent dose (µg/kg/min) Deleted superfluous data (last row):

ρ 0.65 (0.617)	0.49 (0.454)	0.0199ª
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REFERENCE

Tumlin JA, Murugan R, Deane AM, et al; on behalf of the Angiotensin II for the Treatment of High-Output Shock 3 (ATHOS-3) Investigators: Outcomes in Patients with Vasodilatory Shock and Renal Replacement Therapy Treated with Intravenous Angiotensin II. *Crit Care Med* 2018; 46:949–957

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