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Corrigendum

Safety of HES 130/0.4 (Voluven®) in patients with preoperative renal dysfunction undergoing abdominal aortic surgery: a prospective, randomized, controlled, parallel-group multicentre trial

G. Godet, J.-J. Lehot, G. Janvier, A. Steib, V. De Castro, P. Coriat

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Several incorrect references were given in the Discussion section of the paper by G. Godet, first published online 20 May 2008 [1].

The authors apologise for these errors.

Discussion

Paragraph 4, line 10; reference to [14,24] should be [15,24].

Paragraph 5, line 4; reference to [19] should be [20].

Paragraph 5, line 13; reference to [22] should be [23].

Paragraph 5, line 17; reference to [18] should be [19].

Reference

1. Godet G, Lehot J-J, Janvier G, Steib A, De Castro V, Coriat P. Safety of HES 130/0.4 (Voluven®) in patients with preoperative renal dysfunction undergoing abdominal aortic surgery: a prospective, randomized, controlled, parallel-group multicentre trial. *Eur J Anaesthesiol* Published by Cambridge University Press, 20 May 2008. doi:10.1017/S026502150800447X.

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