## **ERRATUM**

# Erratum to: Sphingosine-1-phosphate induces differentiation of cultured renal tubular epithelial cells under Rho kinase activation via the S1P2 receptor

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Figure 5e appeared incorrectly in the article cited above. The correct figure is shown here.

The online version of the original article can be found under doi:10.1007/s10157-014-0933-x.

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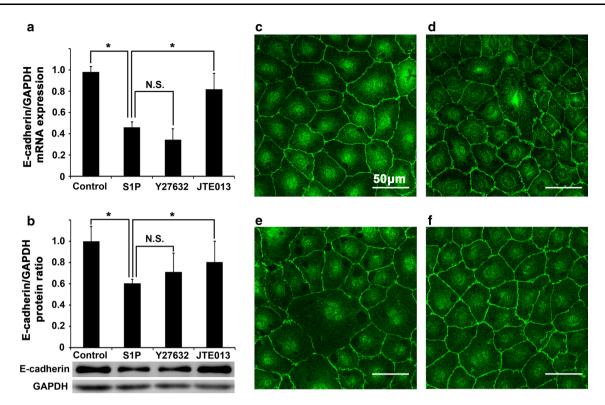
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**Fig. 5** Effect of Y-27632 and JTE013 on S1P-induced E-cadherin mRNA expression. After starvation in serum-free media for 24 h, NRK52E cells were stimulated with S1P (1  $\mu$ M) with or without pretreatment for 1 h with Y-27632 (10  $\mu$ M) or JTE013 (10  $\mu$ M). **a** After a 4-h stimulation with S1P, RNA was extracted, and E-cadherin mRNA was analyzed by real-time RT-PCR with GAPDH mRNA as the internal standard. **b** Total protein was prepared and subjected to SDS-PAGE and Western blot analysis with antibodies against E-cadherin or GAPDH as the internal standard. The results represent

the mean  $\pm$  SD of four separate experiments. \*P < 0.05. c-f Fluorescent immunocytochemistry for E-cadherin. c The cells were grown on coverslips to 80 % confluence then treated with BSA, d-f S1P (1  $\mu$ M) stimulation for 10 h, e Y27632 (10  $\mu$ M) and f JTE013 (10  $\mu$ M) pretreatment for 1 h before S1P stimulation. Immunofluorescence was performed using mouse monoclonal anti-E-cadherin and Alexa488-labeled goat anti-mouse antibodies. E-cadherin expression in the cells was visualized and photographed by fluorescence microscopy at a ×400 magnification

