



Corrigendum: Isoliquiritigenin Protects Against Pancreatic Injury and Intestinal Dysfunction After Severe Acute Pancreatitis *via* **Nrf2 Signaling**

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Isoliquiritigenin Protects Against Pancreatic Injury and Intestinal Dysfunction After Severe Acute Pancreatitis via Nrf2 Signaling

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In the original article, there was a mistake in **Figure 1**, **Figure 4G**, **Figure 5E** and **Figure 6E** as published. During the process of making substantial amendments to our manuscript, **Figures 1G**, **H** were replaced by **Figures 4D**, **E** by mistake. In **Figure 4G** and **Figure 5E**, the internal reference proteins, GAPDH were incorrect. The Sham group in **Figure 6E** was selected from another group by mistake. The corrected figures appear below.

The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

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FIGURE 1 | Isoliquiritigenin (ISL) treatment protects against combined cerulein plus LPS-induced severe acute pancreatitis (SAP) in pancreatic tissue. (**A**, **C**) Pancreatic morphological changes in the different groups at 24 h after SAP. (**B**) Pancreatic histological scores. (**D**, **E**) Expression of c-caspase-3 protein in pancreatic tissue from the different groups at 24 h after SAP. (**B**) Pancreatic histological scores. (**D**, **E**) Expression of c-caspase-3 protein in pancreatic tissue from the different groups at 24 h after SAP. GAPDH was used as the loading control and for band density normalization. (**F**) Statistical graph of c-caspase-3 proteins. (**G**, **H**) Immunofluorescence staining for c-caspase-3 (green) and DAPI (blue) in the different groups. Results are expressed as mean ± SEM. n = 5 per group. *P < 0.05 and **P < 0.01 when comparison was made in WT mice. *P < 0.05 and #*P < 0.01 when comparison was made between WT mice and Nrf2-/- mice.



of the different groups after SAP. GAPDH was used as the loading control and for band density normalization. (C) Statistical graph of Nrf2 and GAPDH protein in the different groups. (D, E) Immunofluorescence staining for Nrf2 (green) and DAPI (blue) in the different groups. (F, G) Protein expression of NF- κ B and I κ B in pancreatic tissue in the different groups after SAP. GAPDH was used as the loading control and for band density normalization. (H, I) Statistical graph of NF- κ B and I κ B in and GAPDH protein in the different groups. Results are expressed as mean \pm SEM. n = 5 per group. *P < 0.05, **P < 0.01, and ***P < 0.001 when comparison was made in WT mice. *P < 0.05 when comparison was made between WT mice and Nrf2-/- mice.







