CORRIGENDUM

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Pro-neurogenic effects of andrographolide on RSC96 Schwann cells *in vitro*

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Following the publication of the above article, an interested reader drew to the authors' attention that a pair of the data panels shown in Fig. 4 contained overlapping data. After having consulted their original data, the authors realized that a total of three panels featured in Figs. 4 and 6 had been erroneously selected. The image of hematoxylin and eosin (H&E) staining for RSC96 Schwann cells in the 1.5625 μ M andrographolide for 6 days data panel in Fig. 4 (bottom row, second panel from the left) was mistakenly submitted. In addition, in Fig. 6, the images of immunohistochemical staining for RSC96 Schwann cells in the 3.125 μ M for 4 days panel (middle row, second panel from the right) and the 6.25 μ M for 6 days panel (bottom row, furthest panel on the right) were mistakenly submitted.

The corrected versions of Figs. 4 and 6 are shown opposite. These corrections were approved by all authors. The authors regret that these errors were featured in the paper, even though they did not substantially alter any of the major conclusions reported in the study. Morever, the authors apologize to the Editor of *Molecular Medicine Reports* and to the readership for any inconvenience caused.



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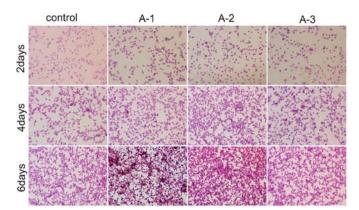


Figure 4. Hematoxylin-eosin staining showing the morphology of RSC96 Schwann cells cultured with $0 \,\mu$ M (control), 1.5625 μ M (A1), 3.125 μ M (A2) and 6.25 μ M (A3) andrographolide for 2, 4 and 6 days. Cell seeding density: 4x10³/ml (original magnification, x100).

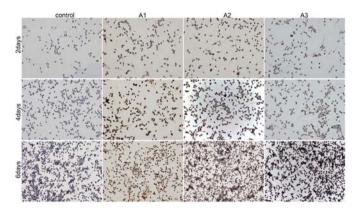


Figure 6. Immunohistochemical staining images showing the presence of S100 β . RSC96 Schwann cells were cultured with 0 μ M (control), 1.5625 μ M (A1), 3.125 μ M (A2) and 6.25 μ M (A3) andrographolide for 2, 4 and 6 days. Cell seeding density: 4x10³/ml (original magnification, x200).