Corrigendum

Corrigendum to “Andrographolide Enhances Proliferation and Prevents Dedifferentiation of Rabbit Articular Chondrocytes: An In Vitro Study”

Li-ke Luo,1,2 Qing-jun Wei,1,2 Lei Liu,1,2 Li Zheng,2,3 and Jin-min Zhao1,2

1Department of Orthopedic Trauma and Hand Surgery, The First Affiliated Hospital of Guangxi Medical University, Nanning, Guangxi 530021, China
2Guangxi Key Laboratory of Regenerative Medicine, Guangxi Medical University, Nanning, Guangxi 530021, China
3The Medical and Scientific Research Center, Guangxi Medical University, Nanning, Guangxi 530021, China

Correspondence should be addressed to Li-ke Luo; luolikegxnn@163.com

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In the article titled “Andrographolide Enhances Proliferation and Prevents Dedifferentiation of Rabbit Articular Chondrocytes: An In Vitro Study” [1], the authors informed the journal that Figures 3–6 are incorrect due to errors in the placement of samples. Initially, an expression of concern was published [2], but the authors have since been able to provide the raw data and corrected figures. The corrected figures are as follows.

![Figure 3: Safranin O staining showing the synthesis of extracellular matrix. Rabbit articular chondrocytes were cultured in vitro with 0 (Control), 1.5 (T1), 3 (T2), and 6 μM (T3) ANDRO for 2, 4, and 6 days. Cell seeding density: 2 × 10^4/mL (original magnification ×100). Scale bar = 200 μm.](image-url)
Figure 4: Confocal laser scanning microscopy images showing the viability of chondrocytes. Control, T1, T2, and T3 represent groups with 0 (Control), 1.5 (T1), 3 (T2), and 6 μM (T3) ANDRO, respectively, to be cultured in vitro for 2, 4, and 6 days. Cell seeding density: 2 × 10⁴/mL (original magnification × 100). Scale bar = 200 μm.

Figure 5: Hematoxylin-eosin staining images showing the morphology of chondrocytes. These chondrocytes were cultured in vitro with 0 (Control), 1.5 (T1), 3 (T2), and 6 μM (T3) ANDRO for 2, 4, and 6 days. Cell seeding density: 2 × 10⁴/mL (original magnification × 100). Scale bar = 200 μm.
References


Figure 6: Immunohistochemical staining images revealing the presence of collagen type II (COL2A1) and type I (COL1A1). Chondrocytes were cultured in vitro with 0 (Control), 1.5 (T1), 3 (T2), and 6 μM (T3) ANDRO for 2, 4, and 6 days. Cell seeding density: 2 × 10^4/mL (original magnification × 100). Scale bar = 200 μm.

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