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EDITED AND REVIEWED BY Bo Liu, University of Wisconsin-Madison, United States

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SPECIALTY SECTION This article was submitted to Atherosclerosis and Vascular Medicine, a section of the journal Frontiers in Cardiovascular Medicine

RECEIVED 05 July 2022 ACCEPTED 18 July 2022 PUBLISHED 05 August 2022

CITATION

Yuan B, Liu H, Dong X, Pan X, Sun X, Sun J and Pan L-L (2022) Corrigendum: A novel resveratrol analog upregulates SIRT1 expression and ameliorates neointima formation. *Front. Cardiovasc. Med.* 9:986353. doi: 10.3389/fcvm.2022.986353

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Corrigendum: A novel resveratrol analog upregulates SIRT1 expression and ameliorates neointima formation

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KEYWORDS

(*R*)-TML104, neointima formation, nicotinamide adenine dinucleotide phosphate oxidase 4, nuclear factor- κ B, vascular smooth muscle cells, reactive oxygen species, SIRT1

A corrigendum on

A novel resveratrol analog upregulates SIRT1 expression and ameliorates neointima formation

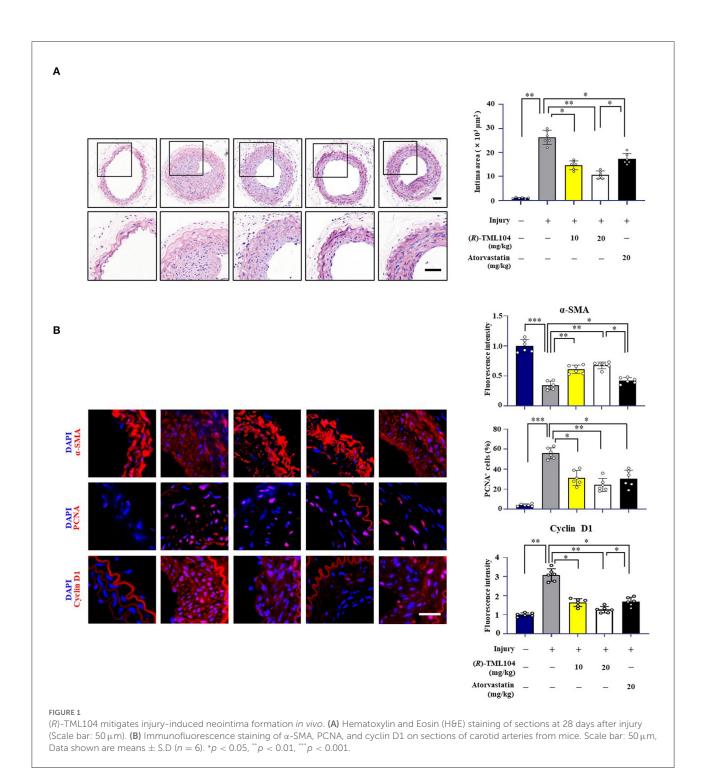
by Yuan, B., Liu, H., Dong, X., Pan, X., Sun, X., Sun, J., and Pan, L.-L. (2021). Front. Cardiovasc. Med. 8:756098. doi: 10.3389/fcvm.2021.756098

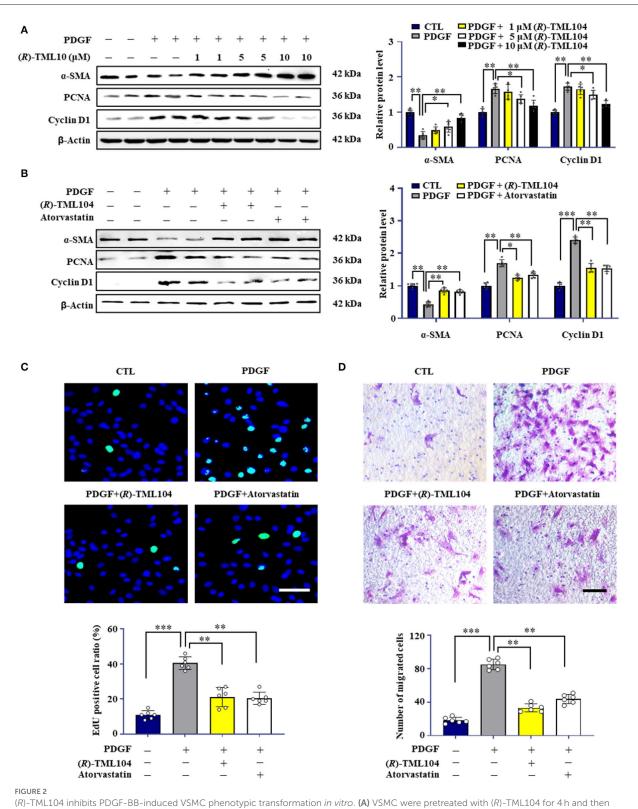
In the published article, there was an error in Figures 1, 2 as published. Due to our mistake in combining images, two graphs in Figures 1B, 2D were misused. The corrected Figures 1, 2 appear below.

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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(*R*)-TML104 inhibits PDGF-BB-induced VSMC phenotypic transformation *in vitro*. (**A**) VSMC were pretreated with (*R*)-TML104 for 4 h and then stimulated with PDGF-BB (20 ng/mL) for 24 h. The protein levels of α -SMA, PCNA, and cyclin D1 were determined by western blotting. (**B**) The protein levels of α -SMA, PCNA, and cyclin D1 were determined by western blotting. (**C**) DNA synthesis in VSMC was determined with an EdU incorporation assay. Blue fluorescence (Hoechst 33342) showed cell nuclei and green fluorescence (EdU) stands for cells with DNA synthesis. (**D**) Transwell assay was performed to determine the migration of VSMC. Scale bar: 50 μ m, Data shown are means \pm S.D (n = 6). *p < 0.05, **p < 0.01, **p < 0.001.