



OPEN ACCESS

APPROVED BY

Frontiers Editorial Office, Frontiers Media SA, Switzerland

*CORRESPONDENCE

Sheng Ding,

to this work

□ doctordingsheng@163.com
 Ping Jing,

ing51118@sina.com

ing51118@sina.com

†These authors have contributed equally

RECEIVED 05 August 2023 ACCEPTED 07 August 2023 PUBLISHED 15 August 2023

CITATION

Li J, Luo T, Wang X, Wang M, Zheng T, Dang X, Deng A, Zhang Y, Ding S, Jing P and Zhu L (2023), Corrigendum: A heterozygous mutation in *NOTCH3* in a Chinese family with CADASIL. *Front. Genet.* 14:1273023. doi: 10.3389/fgene.2023.1273023

COPYRIGHT

© 2023 Li, Luo, Wang, Wang, Zheng, Dang, Deng, Zhang, Ding, Jing and Zhu. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY).

The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Corrigendum: A heterozygous mutation in *NOTCH3* in a Chinese family with CADASIL

Juyi Li^{1†}, Tao Luo^{2†}, Xiufang Wang^{3†}, Mengjie Wang^{4†}, Tao Zheng⁵, Xiao Dang⁶, Aiping Deng¹, Youzhi Zhang⁷, Sheng Ding^{4*}, Ping Jing^{2*} and Lin Zhu^{8*}

¹Department of Pharmacy, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei, China, ²Department of Neurology, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei, China, ³Department of Pain, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei, China, ⁴Department of Endocrinology, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei, China, ⁵Department of Pharmacy, Taihe Hospital, Hubei University of Medicine, Shiyan, Hubei, China, 6Department of Prenatal Diagnostic Center, Guangzhou Women and Children's Medical Center, Guangzhou Medical University, Guangdong, China, ⁵School of Pharmacy, Hubei University of Science and Technology, Xianning, China, ⁵Department of Pediatrics, Tongji Hospital, Huazhong University of Science and Technology, Wuhan, Hubei, China

KEYWORDS

CADASIL, whole-exome-sequencing, heterozygous, Notch3, treatment scheme

A Corrigendum on

A heterozygous mutation in NOTCH3 in a Chinese family with CADASIL

by Li J, Luo T, Wang X, Wang M, Zheng T, Dang X, Deng A, Zhang Y, Ding S, Jing P and Zhu L (2022). Front. Genet. 13:943117. doi: 10.3389/fgene.2022.943117

In the published article, there was an error in the **Funding** statement. "A fund (Central Guiding Local Science and Technology Development Special Project, No. 2022BGE272) which supports our work was not included in the article." The correct **Funding** statement appears below.

"This study was supported by the Health and Family Planning Commission of Wuhan City (WX18M02) and Central Guiding Local Science and Technology Development Special Project (No. 2022BGE272)."

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.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.