CORRECTION





Correction: Whole-exome sequencing identifies a novel CCDC151 mutation, c.325GT (p.E109X), in a patient with primary ciliary dyskinesia and situs inversus

Weizhi Zhang^{1,2} · Dongping Li^{1,2} · Shijie Wei^{1,2} · Ting Guo¹ · Jian Wang¹ · Hong Luo¹ · Yifeng Yang^{1,2} · Zhiping Tan^{1,2}

Published online: 10 June 2019

© The Author(s), under exclusive licence to The Japan Society of Human Genetics 2019

Correction to: Journal of Human Genetics

https://doi.org/10.1038/s10038-018-0540-x published online 30 November 2018

In the originally version of this article, the author affiliation 1 was incomplete. Clinical Center for Gene Diagnosis and Therapy isn't an independent institution

and belongs to The Second Xiangya Hospital of Central South University. The full name of affiliation 1 is Clinical Center for Gene Diagnosis and Therapy, The Second Xiangya Hospital of Central South University, Changsha, China.

The authors would like to apologize for any inconvenience caused.

[☑] Zhiping Tan zhipingtan@csu.edu.cn

Clinical Center for Gene Diagnosis and Therapy, The Second Xiangya Hospital of Central South University, Changsha, China

Department of Cardiovascular Surgery, The Second Xiangya Hospital of Central South University, Changsha, China