



OPEN

Author Correction: mRNA analysis identifies deep intronic variants causing Alport syndrome and overcomes the problem of negative results of exome sequencing

Xiaoyuan Wang, Yanqin Zhang, Jie Ding & Fang Wang

Correction to: *Scientific Reports* <https://doi.org/10.1038/s41598-021-97414-0>, published online 10 September 2021

The original version of this Article contained errors in the mRNA sequences.

As a result, in the Results section, under subheading ‘Analysis of urine NPHS2 and COL4A3-5 mRNAs of the control’,

“The sizes of all ten overlapping fragments covering the entire coding sequence of either *COL4A3*, *COL4A4*, or *COL4A5* mRNA were the same as originally conceived (Fig. 1C), and sequencing of these RT-PCR products confirmed that the amplified sequences mapped precisely to the published *COL4A3*, *COL4A4*, and *COL4A5* mRNA sequences (NM_000091.5, NM_000092.5, and NM_000495.5), respectively.”

now reads:

“The sizes of all ten overlapping fragments covering the entire coding sequence of either *COL4A3*, *COL4A4*, or *COL4A5* mRNA were the same as originally conceived (Fig. 1C), and sequencing of these RT-PCR products confirmed that the amplified sequences mapped precisely to the published *COL4A3*, *COL4A4*, and *COL4A5* mRNA sequences (NM_000091.5, NM_000092.5, NM_000495.5, and NM_033381), respectively.”

Additionally, Table 1 legend contained an error in the number of probands.

“*Age at which serum creatinine tests were carried out in the three probands was 25, 30, 10 and 14 years, respectively.”

now reads:

“*Age at which serum creatinine tests were carried out in the four probands was 25, 30, 10 and 14 years, respectively.”

Finally, in the Discussion section, the following sentence was erroneously included and has subsequently been removed.

“Genetic linkage analysis demonstrated co-segregation of the disease. the proband shared the haplotype of three microsatellite markers around the *COL4A5* gene with her affected daughter, analysis showed that the same haplotype was carried by all affected males and obligatory carrier females.”

Published online: 09 November 2021

The original Article has been corrected.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2021