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ERRATUM

# Erratum to: Report of novel genetic variation in *NPHS2* gene associated with idiopathic nephrotic syndrome in South Indian children

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## Erratum to: Clin Exp Nephrol DOI 10.1007/s10157-016-1237-0

The original version of this article unfortunately contained errors.

In the "Results" section of the main text, under the heading "*NPHS2* gene mutation analysis", the sentence that begins with "The heterozygous mutations" should read:

The heterozygous mutations were observed one in exon 4 [nt21253 (G>A) in 2 %], one in exon 5 [nt23795 (C>T) in 2 %], one in exon 8 [nt29515 (C>T) in 3 %] and one in intronic region [nt21306 (A>G) in 1 %].

In Table 2, the values of "Amino acid change" for P23 and P24, and "Nucleotide change" for P46 and P4 were shown incorrectly. The corrected Table 2 is shown here.

The online version of the original article can be found under doi:10.1007/s10157-016-1237-0.

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**Table 2** Summary of *NPHS2* gene variations in cases (INS) and controls detected in the study

| Sample details | Gene location | Nucleotide change | Amino acid change | Mutation status |
|---|---|---|---|---|
| Control ($N = 100$) | All exons | No change | Not applicable | Not applicable |
| SSNS ($N = 100$) | All exons | No change | Not applicable | Not applicable |
| SRNS-number of patients ($P$) = 100 | | | | |
| P44 | Exon 4 | nt21237 (T>C) | L167P (novel) | Homozygous |
| P7, P73 and P96 | Exon 4 | nt21240 (G>A) | R168H (reported) rs ID: 530318579 | Homozygous |
| P23 | Intron | nt21306 (A>G) | No change (novel) | Heterozygous |
| P24 | Exon 5 | nt23771 (C>T) | R196* (reported) rs ID: 12568913 | Homozygous |
| P47, P3 and P76 | Exon 8 | nt29515 (C>T) | A297V (reported) rs ID: 199506378 | Heterozygous |
| P46 and P4 | Exon 5 | nt23795 (C>T) | L204L (reported) rs ID: 199837664 | Heterozygous |
| P43 and P48 | Exon 4 | nt21253 (G>A) | No change | Heterozygous |
| P51 | Exon 1 | nt5221 (T>C) | S46P (novel) | Homozygous |
| P62 | Exon 1 | nt5250 (G>A) | No change | Homozygous |
| P78 | Exon 5 | nt23841 (A>T) | Q219L (reported) rs ID: 113058664 | Homozygous |
| P93 | Exon 8 | nt29680 (C>T) | S192F (novel) | Homozygous |
| P98 | Exon 4 | nt21260 (C>T) | P175S (novel) | Homozygous |

*SSNS* steroid sensitive nephrotic syndrome, *SRNS* steroid resistant nephrotic syndrome, *NT* nucleotide