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LETTER TO THE EDITOR

## Insertion/Deletion (I/D) Polymorphism of Angiotensin-Converting Enzyme Gene in Steroid-Resistant Nephrotic Syndrome for Children: An Updated Meta-analysis

## Sirs

Steroid-resistant nephrotic syndrome (SRNS) is a risk disease in children and is prone to develop into endstage renal disease. Some investigations suggested that the genetic factor might play an important role in the susceptibility of SRNS. We performed a metaanalysis to explore the association between angiotensinconverting enzyme (ACE) insertion/deletion (I/D) gene polymorphism and SRNS risk and conducted the search on 1 September 2010.<sup>1</sup> We found that D allele and DD genotype were not associated with the risk of SRNS. However, Prasun et al.<sup>2</sup> reported an investigation on the association of ACE I/D gene polymorphism with SRNS risk in 2011, but we did not include this study for our meta-analysis. They reported that the frequency distribution of the DD homozygous was markedly high in the SRNS group compared with that in the control subjects.

The investigation of Prasun et al. <sup>2</sup> was performed in Asian children. As mentioned above, we included the study of Prasun et al.<sup>2</sup> and reran the meta-analysis in overall populations and for Asian children. We found that D allele was associated with SRNS risk in overall populations [odds ratios (OR) = 2.12, 95% confidence intervals (CI): 1.05–4.25; p = 0.04]. This result was not consistent with our previous result for overall populations (OR = 1.69, 95% CI: 0.88–3.25; p = 0.12). Furthermore, the pooled OR for DD genotype was more favorable to SRNS group in overall populations (OR = 2.18, 95% CI: 0.74–6.47), although the difference was not statistically significant (p = 0.16). We also found that the pooled OR for D allele or DD genotype was more favorable to SRNS group compared with the relevant results in our previous meta-analysis, although all the differences were not statistically significant.

The results presented above indicated that the D allele or DD genotype might be a risk factor for SRNS in overall populations and for Asians. The conclusion from rerun meta-analysis might be more robust than that in the previous meta-analysis. It indicated that more case–control association investigations to further clarify the role of the ACE I/D gene polymorphism in SRNS susceptibility were much necessary. So, we call for that more studies on the association between ACE I/D gene polymorphism and SRNS susceptibility be performed in the future.

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