

CORRESPONDENCE

Prediction and prevention of hypertensive disorders of pregnancy: a methodological mistake

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I was interested to read the paper by Ohkuchi *et al.*¹ published in *Hypertens Res* Jan 2017. The most common classifications of hypertensive disorders of pregnancy are chronic hypertension, gestational hypertension, pre-eclampsia (PE) and superimposed PE. The most successful translational research model for explaining the development of PE is the angiogenic/angiostatic balance theory, which involves soluble fms-like tyrosine kinase-1, placental growth factor and soluble endoglin. The aim of the authors was to predict early-onset PE in the beginning of the third trimester. In addition, the authors suggested that an onset threshold or a serial approach appeared to be clinically useful for predicting the imminent onset of PE. The study suggested that onset occurring <4 weeks after blood sampling in the second or early-third trimesters may be predictable because the observed positive likelihood ratio was >10 and the positive predictive value was >20%.¹

However, this result has nothing to do with prediction. First, positive likelihood ratio and

positive predictive values are estimates that are used to evaluate the diagnostic accuracy of a single test compared to a gold standard. Moreover, for prediction studies, we need data from two different cohorts or at least from one cohort divided into two to first develop a prediction model and subsequently validate it. Misleading results are generally the main outcome of research that fails to validate its prediction models.^{2–6}

Finally, in prediction studies, we must assess the interactions between important variables. Final results can be impacted markedly when qualitative interactions are present.^{2–6} This means that most of the time, without assessing the interaction terms, prediction studies will mainly produce misleading messages.

CONFLICT OF INTEREST

The author declares no conflict of interest.

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