

## Comment on: Evaluation of early biomarkers of renal dysfunction in diabetic patients

To the Editor

I have read with a great interest the article of "Evaluation of early biomarkers of renal dysfunction in diabetic patients" by Al-Hazmi et al.<sup>1</sup>

Diabetic nephropathy is one of the microvascular complications of diabetes mellitus. It has been linked to increase mortality in patients with diabetes mellitus.<sup>2</sup> The surge of mortality in these patients progresses with the advancement of diabetic nephropathy.<sup>3</sup> The standard biomarker of diabetic nephropathy is urine albumin/creatinine ratio. So, finding novel biomarkers that might detect the kidney damage early is a crucial advancement in medicine that might lead to early treatment and prevention of the disease progression.

The main result of the study was the significant increase of the studied biomarkers (urinary neutrophil gelatinase-associated lipocalin [uNGAL], urinary N-acetyl-beta-D-glucosaminidase [uNAG], and serum Cystatin C) in patients with diabetes mellitus and severely albuminuria as compared to those with moderately increased albuminuria, normal albuminuria, and normal control. This finding was similarly found in previous studies.<sup>4,5</sup> Additionally, the study came with a very interesting finding that the level of uNGAL was significantly higher in the group with diabetes mellitus and normal albuminuria as compared to normal control. The authors have indicated that such result might allude to early discovery of kidney injury in patients with diabetes. Such result is fascinating, and it might open the door for researches in the future regarding the early discovery and management of diabetic nephropathy.

There were several drawbacks on the study, first the sample size was small. Secondly, there was no matching point between the 4 groups. For example, all the groups had different age, duration of diabetes and different glycemic control which might contribute to differences in the results. Furthermore, the medications which

might affect the albumin secretion were not included in the study like ACEi, ARB, or sGLT2i. Finally, the issue of having a significant difference between a new biomarker in patients with normal albuminuria might not indicate an early disease since there no histological study to confirm that patients were suffering diabetic nephropathy. I agree such result creates a hypothesis, but on the other hand, it might simply mean that the biomarker is not specific.

In conclusion, diabetic nephropathy is a nasty complication of diabetes mellitus with an increase morbidity and mortality that necessitate early discovery of novel markers. The uNGAL might a be good biomarker to detect early renal dysfunction. However, a larger, and a well-organized study is needed in the future to validate their use in clinical practice.

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No reply was received from the Author.

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