



Effect of Renal Vein Variations on Apparent Diffusion Coefficient in Asymptomatic Patients

Asemptomatik Hastalarda Renal Ven Varyasyonlarının Böbrek Görünür Difüzyon Katsayısı Değerlerine Etkisi

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Abstract

Aim: The aim of this study was to investigate whether renal vein variations had an effect on apparent diffusion coefficient (ADC) values in diffusion magnetic resonance imaging (MRI).

Methods: Images of 958 patients who underwent MRI between January 2017 and October 2018 were retrospectively evaluated for renal vein variations based on the records obtained from the image archive. Forty-six patients with renal vein variations and thirty patients of similar age and gender as the control group enrolled in the study. The ADC values of both kidneys were measured at low-high b values.

Results: The ADC values in the left kidney were lower than in the right kidney in both groups ($2.04 \times 10^{-3} \pm 0.22 \times 10^{-3}$ mm²/sec, $1.96 \times 10^{-3} \pm 0.17 \times 10^{-3}$ mm²/sec, for the renal vein variation group; $p=0.008$ and $2.08 \times 10^{-3} \pm 0.13 \times 10^{-3}$ mm²/sec, $1.94 \times 10^{-3} \pm 0.11 \times 10^{-3}$ mm²/sec, for the control group; $p=0.0001$). However, no significant difference was found between the renal vein variation and control groups in terms of the ADC values in both kidneys.

Conclusion: Renal vein variations had no effect on renal ADC values in asymptomatic patients. Further studies can provide additional information for symptomatic patient groups.

Keywords: Renal vein variations, diffusion magnetic resonance imaging, ADC

Öz

Amaç: Bu çalışmanın amacı, renal ven varyasyonlarının difüzyon ağırlıklı manyetik rezonans görüntüleme (MRG) görünür difüzyon katsayısına (GDK) etkisinin olup olmadığını araştırmaktır.

Yöntemler: Ocak 2017 ile Ekim 2018 tarihleri arasında abdominal MRG çekilmiş 958 hastanın tetkikleri renal ven varyasyonları açısından hastane görüntü arşivinden retrospektif olarak incelendi. Renal ven varyasyonu olan 46 hasta ve 30 hastadan oluşan kontrol grubunda yüksek ve düşük b değerlerinde her iki böbreğin GDK değerleri ölçüldü.

Bulgular: Renal ven varyasyonu olan grupta ve kontrol grubunda sağ ve sol böbrek GDK değerleri arasında fark saptandı [$2,04 \times 10^{-3} \pm 0,22 \times 10^{-3}$ mm²/sec, $1,96 \times 10^{-3} \pm 0,17 \times 10^{-3}$ mm²/sec; $p=0.008$ (varyasyonu olan grupta sağ ve sol böbrek, sırasıyla), $2,08 \times 10^{-3} \pm 0,13 \times 10^{-3}$ mm²/sec, $1,94 \times 10^{-3} \pm 0,11 \times 10^{-3}$ mm²/sec; $p=0,0001$, kontrol grubunda sağ ve sol böbrek, sırasıyla]. Sol böbrek GDK değerleri her iki grupta da sağ böbrekten daha düşüktü. Her iki böbrek GDK değerleri renal ven varyasyonu olan grupta ve kontrol grubunda benzerdi.

Sonuç: Renal ven varyasyonu asemptomatik hastalarda böbrek GDK değerleri üzerinde herhangi etki göstermemektedir. Gelecekte semptomatik hasta gruplarıyla yapılacak çalışmalar ek bilgi verebilir.

Anahtar Sözcükler: Renal ven varyasyonları, difüzyon manyetik rezonans görüntüleme, GDK

Introduction

Left renal vein and inferior vena cava variations are relatively frequent compared to those of the right renal vein due to the complexity of embryological development (1). The most common left renal vein variations are observed in the retroaortic and circumaortic renal veins. Renal vein variations are generally asymptomatic and frequently discovered incidentally (1,2). However, in some cases, increased venous pressure due to compression between the vertebrae and the aorta may cause symptoms, such as left side pain, hematuria, and proteinuria (3,4). It has also been reported that renal vein variations may be associated with left-sided varicoceles, pelvic congestion syndrome, dyspareunia, and dysmenorrhea (5,6). Recently, the prevalence of incidentally detected renal vein variations has increased due to the increased use of imaging techniques.

Magnetic resonance imaging (MRI) can noninvasively assess the kidney structure and function in a single screening session. It is possible to evaluate the microstructure of the kidney using diffusion-weighted imaging (DWI), which evaluates the Brownian motion of water molecules in the tissue and allows the quantification of motion based on the apparent diffusion coefficient (ADC) (7,8). There are many human studies concerning the use of DWI in diffuse renal pathologies (9). There are also studies conducting DW MRI in patients with acute renal failure, acute pyelonephritis, acute graft dysfunction, polycystic disease, amyloidosis, diabetes, various glomerulonephritis, obstruction, renal artery stenosis, and other various etiologies (10). These studies revealed changes in renal parenchyma, such as edema and fibrosis using DW MRI and ADC values.

It is suggested that renal vein variations can cause venous hypertension, which may also be responsible for symptoms. If venous hypertension due to venous variation causes changes in the kidney ADC values, this parameter can be used in the diagnosis and follow-up of these patients. In this study, we aimed to evaluate whether left renal vein variations caused changes in the kidney ADC values.

Methods

Study Subjects

Ethics committee approval was obtained from Eskişehir Osmangazi University Faculty of Medicine (date: 27.11.2018, no: 25403353-050.99-E.128703) and the study was conducted in accordance with the Declaration of Helsinki. The study involved retrospective reevaluation of the upper abdominal MRI recorded between January 2017 and October 2018, obtained from the hospital image archive. Upper abdominal MRI performed in patients with non-urinary system indications (chronic liver

parenchymal disease, focal lesion in liver parenchyma, cholelithiasis, adrenal lesion, pancreatic lesions, etc.) were included in the study. Patients with renal dysfunction and solitary kidneys, and those with images that could not be evaluated for technical reasons (motion artefacts, MRI without diffusion examination at an appropriate value of b) were excluded from the study. As a result, the MRI images of 958 patients, comprising 498 (52%) females and 460 (48%) males were evaluated. The flowchart is presented in Figure 1.

None of the patients had any urinary symptom or pathological finding in urinalysis. Their serum urea and creatinine values were also normal. The presence and type (circumaortic, retroaortic) of left renal vein variations were recorded. Thirty patients of similar age and gender without renal vein variations were selected to form the control group. Renal function values were normal in the control group. The serum urea and creatinine values and urine analysis were also within the normal range.

Magnetic Resonance Imaging

MRI was performed using a 3-Tesla (General Electric, Milwaukee, WI) device. In all examinations, a 48-channel body coil was used. T2-weighted axial and coronal plane images, T1-weighted axial plane images, and diffusion-weighted echo planar images (DW-EPI) were obtained in each patient. The DW sections were obtained in the axial

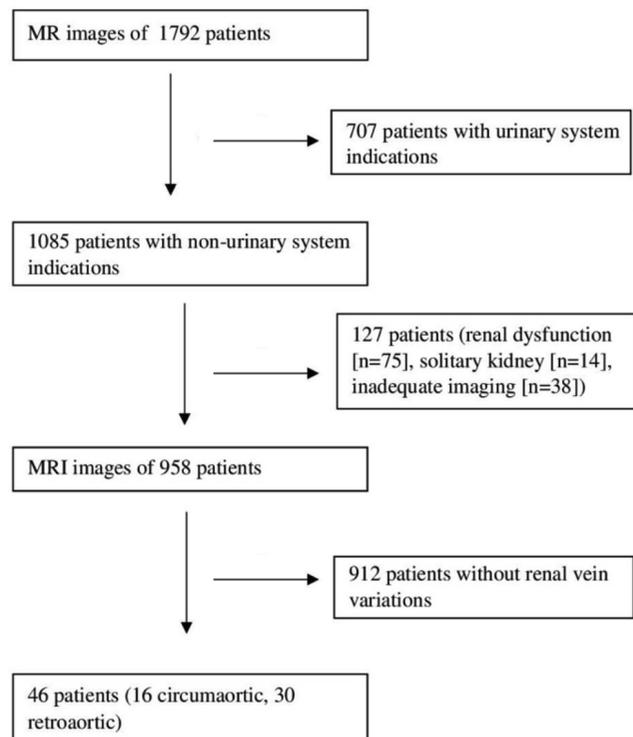


Figure 1. Recruitment schema of the participants
MR: Magnetic resonance

plane using the DW-EPI sequence at low ($b=0 \text{ s/mm}^2$) and high ($b=1000 \text{ s/mm}^2$) gradient values without breath-holding. The imaging parameters of the DW-EPI sequence were as follows: TR/TE, 9231/64.1; slice thickness, 5 mm; field of view, 42 cm; and matrix size; 98x128. The ADC value was automatically calculated by the device simultaneously. To prevent bowel movements causing artefacts, MRI was performed after four to six hours of fasting. However, the patients were not given any antispasmodic.

Analysis of Images

The images of the patients with renal vein variations and the control group were evaluated using the dedicated workstation (GE, Advantage Workstation 4.3, USA) by two radiologists (one experienced in abdominal imaging) based on consensus. The T1- and T2-weighted images were evaluated for focal kidney lesions. There was no solid renal tumor in patients with renal vein variations. In patients with simple cysts, the levels including the cysts were not included in the measurement. The circular-shaped regions of interest (ROI) with a diameter of 1 cm were placed in the corticomedullary area in both renal parenchyma (Figure 2, 3). Circular ROI was placed in three regions in the upper, middle and lower sections of the posteromedial of both kidneys. All measurements were undertaken by a single radiologist experienced in abdominal imaging. The measurements were performed twice, and the mean ADC values were used for further evaluation. The diameter of the left renal vein in both the control and renal vein variation groups was measured by the same radiologist

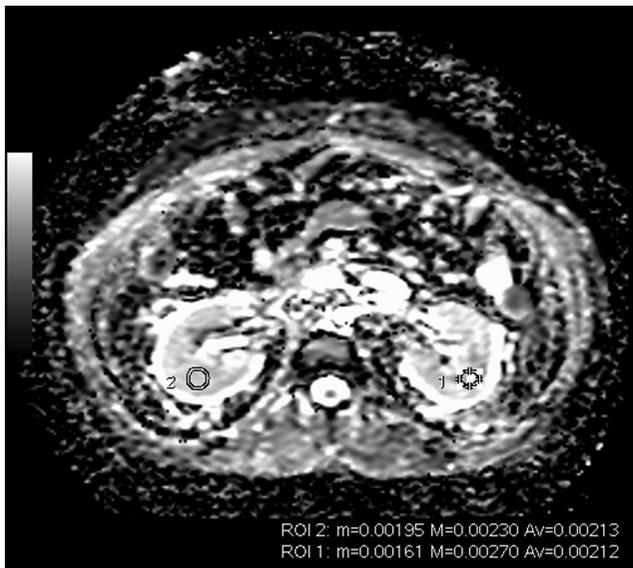


Figure 2. ADC maps ($b=1000 \text{ s/mm}^2$) in of a 52-year-old male in the control group. Regions of interest for the ADC measurements of the right and left kidneys are indicated by white circles
ADC: Apparent diffusion coefficient

based on the midpoint between the abdominal aorta and the left renal hilum in the axial plane (Figure 4, 5).

Statistical Analysis

SPSS software v.22.0 (Chicago, IL) was used for statistical analysis. The Shapiro-Wilk test was used for normality testing. Quantitative variables were shown as arithmetic mean \pm standard deviation, and qualitative variables as numbers and percentages. The right and left kidney ADC values in both the control and renal vein variation groups were compared using the Student's t-test. The same test was used to compare the left renal vein diameter between the two groups. The paired samples t-test was used to investigate whether there

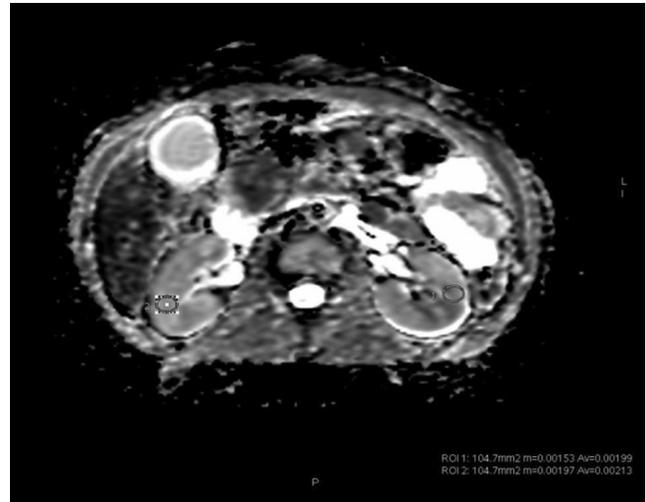


Figure 3. ADC maps ($b=1000 \text{ s/mm}^2$) of a 61-year-old male in the renal vein variation group. Regions of interest for the ADC measurements of the right and left kidneys are indicated by white circles
ADC: Apparent diffusion coefficient

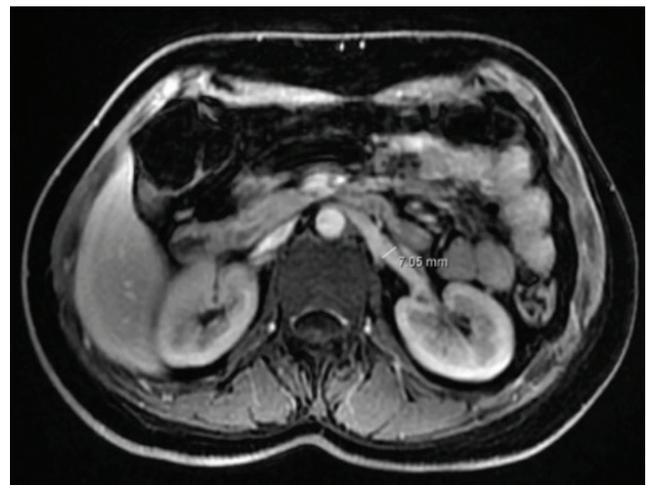


Figure 4. Measurement of the left renal vein diameter on the axial plane images in a 45-year-old female in the control group

was a difference in the right and left kidney ADC values between the two groups.

Results

Left renal vein variations were found in 46 (4.80%) of 958 patients [circumaortic in 16 (Figure 6a and 6b) and retroaortic in 30 (Figure 7)]. There were 24 women (age range: 23-88 years; mean age: 58.1±14.6 years) and 22 men (age range: 30-92 years; mean age: 57.7±16.8 years) with renal vein variations and 18 women (age range: 27-69 years; mean age: 56.2±13.6 years), 12 men (age range: 34-77 years; mean age: 60.5±14.3 years) in the control group. There was no difference in age between the groups.

Table 1 presents the comparison of the data between the two groups. The mean ADC of the left kidney was

lower than that of the right kidney in control group (p=0.0001). The mean ADC of the left kidney was lower than that of the right kidney in renal vein variation group (p=0.008). In the comparison of the right and left kidney ADC values between the renal vein variation and control groups, no significant difference was found.

Parameters	Renal vein variations group	Control group	p value
Number of patients (n)	46	30	-
Male age/ female age (years)	57.7±16.8/58.1±14.6	60.5±14.3/56.2±13.6	0.12
Male/female (n)	22/24	12/18	-
ADC of right kidney (mm ² /sec)	2.04x10 ⁻³ ±0.22x10 ⁻³	2.08x10 ⁻³ ±0.13x10 ⁻³	0.26
ADC of left kidney (mm ² /sec)	1.96x10 ⁻³ ± 0.17x10 ⁻³	1.94x10 ⁻³ ±0.11x10 ⁻³	0.71
Left renal vein diameter (cm)	7.39±0.90	7.32±0.81	0.75

ADC: Apparent diffusion coefficient, n: Number

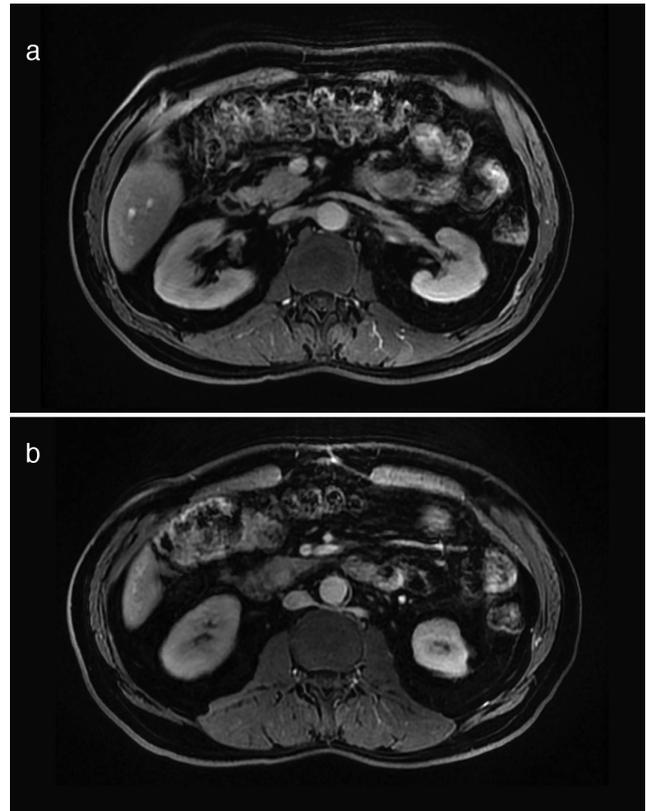


Figure 6. Axial plane MRI showing the circumaortic position of the left renal vein. The left renal vein reaches both the anterior (a) and posterior (b) of the abdominal aorta
MRI: Magnetic resonance imaging

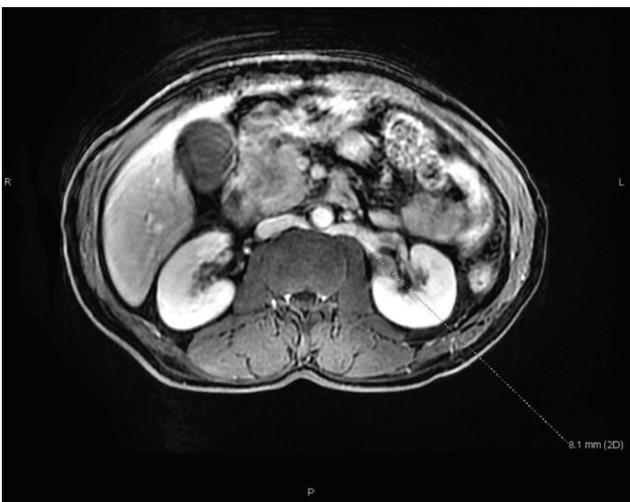


Figure 5. Measurement of the left renal vein diameter on the axial plane images in a 61-year-old male with (retroaortic) renal vein variations

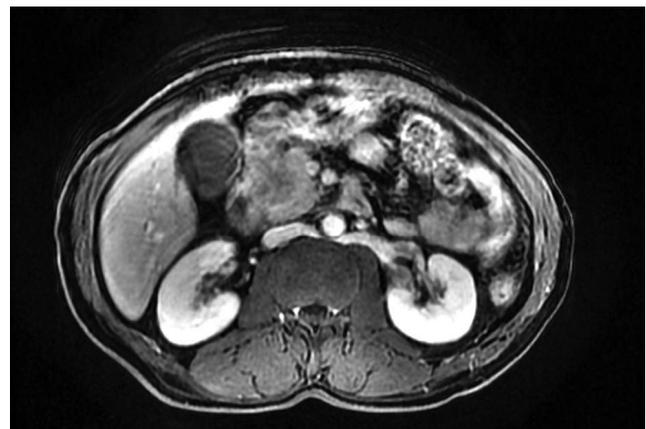


Figure 7. Axial plane MRI showing the retroaortic position of the left renal vein
MRI: Magnetic resonance imaging

Lastly, the mean left renal vein diameter in the renal vein variation group did not differ from that of the control group.

Discussion

Numerous radiological (with computed tomography or MRI), surgical and post-mortem studies have been conducted to investigate renal vein variations. In these studies, the incidence of retroaortic and circumaortic renal vein variations was reported to be 0.5-17% and 0.3-6.8%, respectively (3). In the current study, the incidence of left renal vein variations was similar to the values reported in previous studies.

In recent years, DWI has gained increased interest in the evaluation of chronic kidney disease due to the correlation between reduced tissue water diffusion and fibrosis development (11-14). There are many studies in the literature on the ADC values obtained from both patients with diffuse renal parenchymal diseases and individuals with normal renal parenchyma. These studies generally showed that the ADC values in patients with diffuse renal diseases were decreased, and the glomerular filtration rate and renal ADC values were positively correlated in patients with renal dysfunction (15-17). In patients with both acute and chronic renal failure, ADC values are known to decrease. The relationship between reduced ADC and pathological fibrosis development seems to primarily relate to the renal cortex (11-14). Although the literature contains several studies on the renal functions and diffusion values in patients with diffuse renal diseases, to the best of our knowledge, no study has been undertaken to evaluate the effect of renal vascular structures and renal anomalies on ADC values. Therefore, our study is the first and will contribute to the literature in this regard.

In the literature, different ADC values for the right and left kidneys were found in normal kidneys in studies using different devices with different magnetic field strength (1.5 and 3 Tesla) (18-22). However, in some of these studies, the right and left kidney values were not separately reported (19). Kim et al. (18) found that the ADC values of the left kidney were lower than those of the right kidney at high b values using 3 Tesla MRI devices. However, the authors did not provide any information on whether these differences were statistically significant. Similar to our study, Yoshikawa et al. (22) found that the left kidney ADC values were lower than the right kidney values using a 1.5 Tesla MRI device. However, they did not present any information about the statistical significance of their results. In contrast, Song et al. (23) determined that the ADC values of the left kidney were higher than those of the right kidney in individuals with healthy renal

function. In our study, a difference was found between the right and left kidney ADC values in both the control and renal vein variation groups. However, similar ADC results between these two groups led us to conclude that renal vein variations had no effect on kidney ADC values.

Variations do not cause pressure changes in the renal vein as long as they are asymptomatic. Therefore, we may not have detected the differences in the ADC value between the control and renal vein variation groups in our study. Even if there is a renal vein pressure change that is not causing deterioration in the renal microstructure, may explain why ADC value is not affected. According to our results, it is possible to conclude that renal vein variations do not cause changes in the kidney microstructure as long as they are asymptomatic.

DWI measures the random motion of water molecules, which can be free or restricted by cellular membranes or other barriers (17). It provides microstructural information about tissue microstructure by using the movement of water to probe extracellular and intracellular extravascular spaces (24). However, the ADC value is affected by not only true water diffusion but also microperfusion and tubular flow in the renal tissue (25-27). In the literature, the use of novel DWI models, e.g. intravoxel incoherent motion (IVIM) and diffusion tensor imaging (DTI), in the assessment of diffuse renal pathologies has been investigated. IVIM could provide more accurate information on pseudo diffusion and true diffusion DTI, an advancement of DW MRI, can offer an insight into the structural properties of tissue by assessing the directionality of water diffusion, which is quantified as the percentage of spatially oriented diffusion. Diffusion anisotropy is related to structural organization, and therefore can be compromised in a pathological process (28). The evaluation of the effects of variations using DTI and IVIM can give more accurate information about the pure diffusion effect. In addition, directional information in DTI is an advantage for vascular structures.

Study Limitations

One of the limitations of our study is that all patients in the group with renal vein variations were asymptomatic. Hematuria and proteinuria were not present in either group. The absence of clinical symptoms in our patients may explain the similarity of the results between the renal vein variation and control groups. Clinically, nutcracker syndrome is observed in a small proportion of patients with renal vein variations. Studies conducted with symptomatic group scan provide more accurate results. There is no consensus on the treatment method for symptomatic patients or the selection of appropriate cases for treatment. Further studies evaluating ADC parameters

in both symptomatic and asymptomatic groups based on the severity of findings can provide more information concerning the kidney microstructure. ADC values can be one of the parameters that can be considered in treatment selection by providing information about the renal microstructure. The other limitations of our study can be regarded as the relatively low number of patients with renal vein variations. However, this was inevitable considering the low incidence of such variations.

Conclusion

The patients with renal vein variations and the control group had similar ADC values for both the right and left kidneys. Thus, it can be stated that renal vein variations have no effect on kidney ADC values. However, right and left kidney ADC values being different in groups with and without the renal vein variation should be kept in mind when evaluating this parameter in other conditions.

Authorship Contributions

Concept: E.G., E.E., M.K. Design: E.G., M.O., M.K. Data Collection or Processing: E.E., M.O. Analysis or Interpretation: E.G., M.K. Literature Search: E.G., E.E., M.O. Writing: E.G.

Conflict of Interest: The authors declared no conflicts of interest concerning the authorship and publication of this article.

Financial Disclosure: The authors received no financial support for the research and/or authorship of this article.

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