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Conflict of Interest

Jinyong Lee is an employee of Samsung
Medison.

End-Stage Renal Disease Impairs the Multidirectional Movements of the Common Carotid Artery: Assessment Using Dimensional Speckle-Tracking Carotid Strain Ultrasonography

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ABSTRACT

BACKGROUND: Arterial stiffening is a major contributing factor in the development of cardiovascular disease in patients with end-stage renal disease (ESRD). However, there is no gold standard for evaluating arterial stiffness. This study aimed to evaluate the newly developed speckle-tracking carotid strain imaging method in assessing arterial stiffness in patients with ESRD.

METHODS: In total, 85 patients with normal renal function (controls) and 36 with ESRD were enrolled in this single-center study. Carotid B-mode ultrasonography was performed for all patients. Arterial stiffness indices and strain parameters of the common carotid arteries were analyzed. Values were compared between the groups, and multivariate linear regression analysis was performed to assess the impact of ESRD on carotid strain.

RESULTS: There were no differences in the intima-media thickness, β stiffness index, and arterial compliance, but arterial distensibility was lower, and the elastic modulus and pulse wave velocity β (PWV) were higher among patients with ESRD (all $p < 0.05$), whether assessed in the longitudinal or transverse plane. Both longitudinal and transverse strain rates were reduced in patients with ESRD (all $p < 0.05$). In multivariate analyses, ESRD independently reduced both transverse radial strain and strain rate (all $p < 0.05$), and the transverse circumferential strain and strain rate ($p < 0.05$). However, all conventional aortic stiffness indices and longitudinal strain parameters were not associated with ESRD.

CONCLUSIONS: Speckle-tracking carotid strain ultrasonography was successfully performed in both normal subjects and patients with ESRD. Multidirectional carotid strain analyses may provide more value than conventional aortic stiffness indices for risk stratification in patients with ESRD.

Keywords: Common carotid artery; Strain; End stage renal disease; Ultrasonography; Atherosclerosis

INTRODUCTION

Cardiovascular disease is one of the main causes of morbidity and mortality in patients with chronic kidney disease (CKD) and end-stage renal disease (ESRD). Prior studies have observed the subclinical damage to large arteries and have demonstrated that both structural and functional alterations of these large arteries are major contributing factors related to cardiovascular complications.^{1,2)}

Notably, arterial stiffening and remodeling in patients with ESRD are of multifactorial origin, with extensive arterial calcifications, and have been shown to be a strong and independent predictor for both all-cause and cardiovascular deaths.²⁻⁴⁾ It has been also observed that arterial stiffening occurs prior to the development of structural changes of the arterial wall.⁵⁾ Thus, arterial stiffness has emerged as a reliable feature for the early detection of atherosclerotic vascular changes and also as a potential novel therapeutic target for cardiovascular risk reduction in patients with ESRD. Therefore, the accurate non-invasive assessment of arterial stiffness has developed a considerable clinical interest as there is no gold standard for evaluating arterial stiffness thus far.⁶⁾

A recently developed multi-directional carotid artery strain assessment method using speckle-tracking ultrasonography is one of the most accessible and reproducible non-invasive imaging modality that can directly and quantitatively assess arterial stiffness, compared to conventional aortic stiffness measures such as distensibility, elastic modulus, β stiffness index, or carotid femoral pulse wave velocity.^{6,7)} The reliability and sensitivity of the speckle-tracking carotid strain imaging has been validated in both normal subjects and patients with hypertension and diabetes, but not in patients with ESRD so far.^{6,7)}

In this regard, we aimed to explore the difference in the multidirectional movements of the common carotid artery (CCA) as assessed by the speckle-tracking carotid strain imaging method between patients with ESRD and those with normal kidney function.

METHODS

Study population

From April 2014 to May 2015, 121 patients who underwent carotid ultrasonography for the evaluation of atherosclerosis status were enrolled in this single-center study. Patients were divided into the following groups according to their renal function and history of dialysis: normal renal function, defined as estimated glomerular filtration rate (eGFR) > 90 mL/min/1.73m², group (control) and the ESRD group according to the guidelines of the Kidney Disease: Improving Global Outcomes (KDIGO).⁸⁾ Patients with eGFR ≤ 90 mL/min/1.73m² but not undergoing dialysis, patients with significant CCA stenosis ($>50\%$), and patients with CCA stent or prior history of CCA surgery were excluded. The study design was approved by the institution's human research committee, and the study protocol conformed to the ethical guideline of the 1975 Declaration of Helsinki and informed consent was provided by the patients to participate in the study. Clinical characteristics and laboratory results at the time of carotid ultrasound were collected.

Carotid ultrasound image acquisition

High-resolution carotid ultrasound images were acquired using a linear array transducer (nominal bandwidth of 3–12 MHz) with an 8-MHz center frequency (RS80A and HS70A,

Samsung Medison, Seoul, Korea). To acquire a high frame rate (>120 fps), the view angle was reduced to ensure accurate measurements of CCA movements. Patients' heads were tilted slightly upward to the midline position, and both CCAs were examined using recordings at ≥ 2 consecutive beats.

Assessment of the multidirectional movements of the CCA

The images showing the multidirectional movements of the CCA were analyzed using commercially available software (Samsung Medison Co., Ltd., Seoul, Korea). The software assesses the functional property of CCAs by automatically calculating the displacement of the CCA walls (Figure 1).

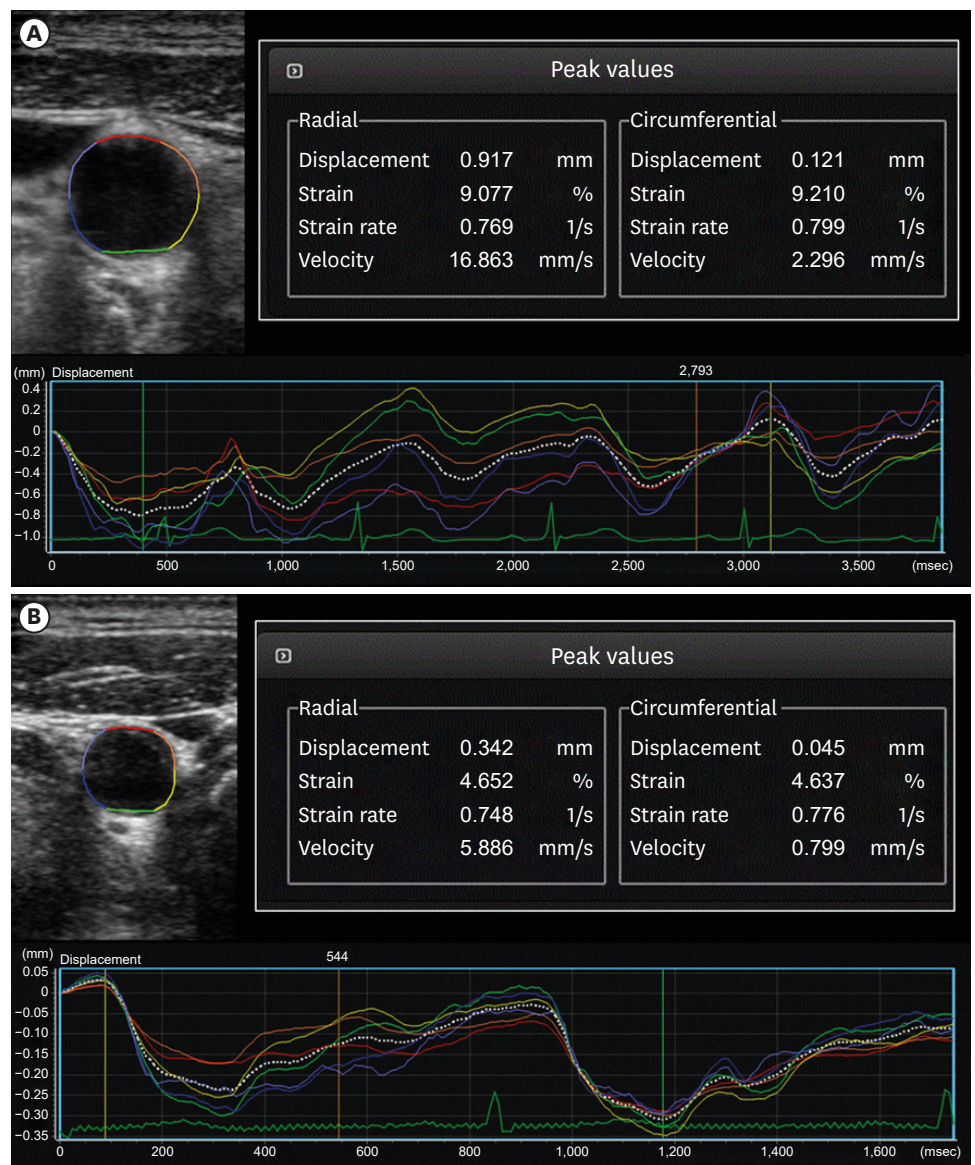


Figure 1. Strain analysis of the common carotid artery using the speckle tracking method for carotid B-mode ultrasonography in a 69-year-old normal control patient (A) and an age-matched ESRD patient (B). The strain and strain rate values were all decreased in the end-stage renal disease patient compared to the normal control patient.

For the analysis, ≥ 5 mm of the CCA below the origin of the carotid bulb was selected. The user-defined control points, the blood-intima boundaries of the CCA, were manually traced from a still frame image with the best image quality and were automatically tracked by the software based on an optical flow algorithm.⁹⁾ The control points are constrained to maintain the global shape and to move within a local range, to ensure stability in tracking the motion of vessels. At least four speckles on the CCA wall were identified, and the movements of each speckle during the cardiac cycle were traced at the near and far walls. Radial movement of the CCA in the longitudinal plane and radial circumferential movements in the transverse plane were automatically calculated using the speckle-tracking technique. Peak radial and circumferential displacements, strain, and strain rate were also calculated.

For the calculation of conventional arterial stiffness parameters,¹⁰⁾ the central blood pressure, derived from the aortic pulse wave, was measured after ≥ 5 min of resting in supine position by a trained technician with pulse wave analysis of the radial artery using a tonometry (SphygmoCor; AtCorMedical, Sydney, Australia). The SphygmoCor software calculated the aortic pulse wave by means of a generalized transfer function from the radial signal. The measured central blood pressure was entered into the software, and all arterial stiffness parameters were then automatically calculated from the software according to the pre-defined formula for each parameter. β stiffness index was defined as the ratio of the natural logarithm of central systolic/diastolic pressure to relative change in diameter. Arterial compliance was defined as the absolute change in CCA diameter for a given change in pressure (e.g., difference between the diastolic and systolic diameters $[\Delta D]$ /difference between the systolic and diastolic blood pressures $[\Delta P]$). Distensibility refers to the relative change in vessel diameter for a given change in pressure (e.g., $\Delta D/[\Delta P \times \text{CCA diameter in the diastolic phase (D)}]$ in mmHg). The elastic modulus index was defined as the pressure change required for the theoretical stretch from the resting vessel diameter (e.g., $[\Delta P \times D]/\Delta D$ in mmHg). (e.g., $\log [\text{systolic blood pressure/diastolic blood pressure}]/[\Delta D/D]$). Pulse wave velocity β (PWV) was defined as the speed with which the pulse wave travels along the length of the artery (e.g., stiffness index \times diastolic blood pressure/ $2 \times$ blood density]), while assuming the blood density to be 1.050 g/cm^3 .

The presence of carotid plaque was defined as a focal structure that protrudes into the arterial lumen and measured ≥ 1.5 mm or $\geq 50\%$ of the surrounding intima-media thickness (IMT). Additional images were acquired at another location close to the CCA that does not contain the plaque for patients with carotid plaques for the analysis. Carotid IMT was measured in a semiautomatic manner in the end-diastole phase for bilateral CCAs in the longitudinal plane, and the mean value of the right and left IMTs was used in the analysis. In patients with carotid plaque, IMT was measured at a location where the plaque was not included.

Statistical analysis

Parametric continuous variables were presented as mean \pm standard deviation, and categorical variables were expressed as count (%). Differences between continuous variables were compared using Student's t-test or Mann Whitney U test, as appropriate, whereas the Pearson's chi-squared or Fisher's exact test was employed, as appropriate, for categorical variables. To assess the impact of ESRD on the carotid strain values and conventional aortic stiffness parameters, multivariate linear regression analyses were constructed and adjusted for age, male sex, hypertension, diabetes mellitus, body mass index, creatinine level, high-density lipoprotein level, the presence of carotid plaque, and the maximal diameter of CCA. A two-tailed p -value < 0.05 was considered statistically significant. All analyses were performed using SPSS (Version 23.0, IBM Corp., Armonk, NY, USA).

RESULTS

Baseline characteristics of the study population

A total of 85 control patients (61±12 years old, 53% were male) and 36 with ESRD (55±3 years old, 61% were male) were enrolled (**Table 1**). There was no difference in the percentage of male sex ($p = 0.431$), but patients with ESRD were younger and had more hypertension than control patients (89% vs. 47%, $p < .001$). Central systolic and diastolic blood pressures were also higher in the ESRD group (all $p < 0.05$), but there were no differences in body mass index and the presence of diabetes mellitus or hyperlipidemia (all $p > 0.05$). The level of total cholesterol was higher in the ESRD group (180.8±40.5 mg/dL vs. 160.8±40.0 mg/dL, $p = 0.016$), but there were no differences in the level of high-density lipoprotein, low-density lipoprotein, and triglyceride (all $p > 0.05$).

Carotid ultrasonographic findings

Carotid ultrasonography and subsequent strain analysis were successfully performed on all participating patients. Atherosclerotic carotid plaques were detected in 82.4% of the controls and 31.4% of patients with ESRD ($p = 0.002$) (**Table 2**). The mean maximal

Table 1. Baseline characteristics and laboratory results of the controls and patients with ESRD

| | Control (n = 85) | ESRD patients (n = 36) | p |
|---|------------------|------------------------|---------|
| Age (years) | 61 ± 12 | 55 ± 3 | 0.008 |
| Sex (male) | 45 (52.9) | 22 (61.1) | 0.431 |
| Heart rate (/min) | 69 ± 14 | 73 ± 12 | 0.084 |
| Central systolic blood pressure (mmHg) | 127 ± 19 | 149 ± 30 | < 0.001 |
| Central diastolic blood pressure (mmHg) | 78 ± 11 | 85 ± 17 | 0.025 |
| Body mass index (kg/m ²) | 24.9 ± 4.0 | 24.4 ± 4.1 | 0.542 |
| Hypertension | 40 (47.1) | 32 (88.9) | < 0.001 |
| Diabetes mellitus | 40 (47.1) | 18 (50.0) | 0.843 |
| Coronary artery disease | 1 (1.2) | 3 (8.3) | < 0.001 |
| Hyperlipidemia | 43 (50.6) | 17 (47.2) | 0.843 |
| Blood urea nitrogen (mg/dL) | 14.5 ± 4.6 | 60.2 ± 18.3 | < 0.001 |
| Creatinine (mg/dL) | 0.73 ± 0.14 | 10.06 ± 3.61 | < 0.001 |
| Total cholesterol (mg/dL) | 160.8 ± 40.0 | 180.8 ± 40.5 | 0.016 |
| High-density lipoprotein (mg/dL) | 46.0 ± 12.7 | 55.2 ± 54.5 | 0.369 |
| Low-density lipoprotein (mg/dL) | 91.5 ± 36.6 | 92.4 ± 30.6 | 0.909 |
| Triglyceride (mg/dL) | 121.3 ± 80.3 | 158.5 ± 157.4 | 0.112 |

Values are presented as means ± standard deviation or number (%).
ESRD: end-stage renal disease.

Table 2. Comparison of ultrasonographic parameters in the longitudinal plane between the controls and patients with ESRD

| | Control (n = 85) | ESRD patients (n = 36) | p |
|--------------------------------|------------------|------------------------|---------|
| Carotid plaque | 70 (82.4) | 11 (31.4) | 0.002 |
| Intima-media thickness (mm) | 0.81 ± 0.17 | 0.76 ± 0.15 | 0.121 |
| Minimal diameter (mm) | 6.09 ± 0.87 | 7.01 ± 1.23 | < 0.001 |
| Maximal diameter (mm) | 6.54 ± 0.91 | 7.54 ± 1.32 | < 0.001 |
| β stiffness index | 7.6 ± 4.3 | 8.5 ± 4.3 | 0.309 |
| Arterial compliance (mm/kPa) | 0.097 ± 0.044 | 0.104 ± 0.057 | 0.892 |
| Arterial distensibility (/kPa) | 0.0016 ± 0.0008 | 0.0013 ± 0.0007 | 0.022 |
| Elastic modulus (kPa) | 766.5 ± 433.1 | 979.6 ± 590.5 | 0.022 |
| Pulse wave velocity (m/s) | 281.0 ± 152.6 | 342.0 ± 191.1 | 0.041 |
| Strain parameters (radial) | | | |
| Displacement (mm) | 0.449 ± 0.203 | 0.504 ± 0.238 | 0.357 |
| Strain (%) | 7.24 ± 3.27 | 6.84 ± 3.06 | 0.454 |
| Strain rate (1/s) | 1.005 ± 0.508 | 0.711 ± 0.298 | 0.001 |

Values are presented as means ± standard deviation or number (%).
ESRD: end-stage renal disease.

Table 3. Comparison of ultrasonographic parameters in the transverse plane between the controls and patients with ESRD

| | Control (n = 85) | ESRD patients (n = 36) | p |
|-------------------------------------|------------------|------------------------|---------|
| Minimal diameter (mm) | 7.22 ± 1.11 | 7.76 ± 1.29 | 0.022 |
| Maximal diameter (mm) | 7.79 ± 1.12 | 8.39 ± 1.39 | 0.013 |
| β stiffness index | 7.1 ± 3.3 | 8.5 ± 4.3 | 0.105 |
| Arterial compliance (mm/kPa) | 0.145 ± 0.071 | 0.129 ± 0.066 | 0.150 |
| Arterial distensibility (/kPa) | 0.0018 ± 0.0010 | 0.0013 ± 0.0006 | 0.027 |
| Elastic modulus (kPa) | 724.4 ± 393.0 | 995.4 ± 700.0 | 0.027 |
| Pulse wave velocity (m/s) | 264.4 ± 135.9 | 346.1 ± 219.9 | 0.045 |
| Strain parameters (radial) | | | |
| Displacement (mm) | 0.562 ± 0.223 | 0.601 ± 0.342 | 0.905 |
| Strain (%) | 7.6 ± 3.2 | 7.2 ± 3.8 | 0.355 |
| Strain rate (1/s) | 0.97 ± 0.45 | 0.72 ± 0.47 | < 0.001 |
| Strain parameters (circumferential) | | | |
| Displacement (mm) | 0.074 ± 0.029 | 0.079 ± 0.045 | 0.916 |
| Strain (%) | 7.7 ± 3.3 | 7.3 ± 4.0 | 0.350 |
| Strain rate (1/s) | 0.99 ± 0.47 | 0.74 ± 0.54 | < 0.001 |

Values are presented as means ± standard deviation.
ESRD: end-stage renal disease.

diameter of CCAs in the longitudinal plane were greater in patients with ESRD than in the controls (7.54±1.32 mm vs. 6.54±0.91 mm, $p < .001$), but the IMT was not different (0.76±0.15 mm vs. 0.81±0.17 mm, $p = 0.121$). There were no differences in the β stiffness index and arterial compliance (all $p > 0.05$) between groups, but the arterial distensibility was lower and the elastic modulus and PWV were higher in the ESRD group (all $p < 0.05$). When the strain parameters were compared, the radial strain rate in patients with ESRD was significantly lower than that in the controls (0.711±0.298 /s vs. 1.005±0.508 /s, $p = 0.001$), although there were no differences in displacement and strain.

The results of the assessment of the transverse plane were also consistent (**Table 3**). The minimal and maximal CCA diameters were greater in patients with ESRD (all $p < 0.05$). β stiffness index and arterial compliance were not different, but the patients with ESRD exhibited lower arterial distensibility and higher elastic modulus and PWV compared with the controls (all $p < 0.05$). The speckle-tracking carotid strain analysis revealed that while there were no differences in displacement and strain between both the radial and circumferential methods, both radial and circumferential strain rates were significantly reduced in patients with ESRD (0.72±0.47 /s vs. 0.97±0.45 /s and 0.74±0.54 /s vs. 0.99±0.47 /s, all $p < .001$).

Impact of ESRD on strain parameters

In the multivariate linear regression analyses adjusted for age, male sex, hypertension, diabetes mellitus, body mass index, creatinine level, high-density lipoprotein level, the presence of carotid plaque, and the maximal diameter of CCA, ESRD had no impact on both conventional aortic stiffness parameters, including distensibility, elastic modulus, and PWV, and strain parameters assessed in the longitudinal plane (all $p > 0.05$) (**Table 4**). ESRD also had no impact on distensibility, elastic modulus, and PWV assessed in the transverse plane (all $p > 0.05$).

In contrast, ESRD was negatively and independently associated with both transverse radial strain and strain rate (β : -6.087 (95% confidence interval [CI]: -10.790 to -1.384) and β : -0.833 (95% CI: -1.390 to -0.275), respectively, all $p < 0.05$). ESRD also independently reduced the transverse circumferential strain and strain rate (β : -6.771 (95% CI: -11.525 to -2.018) and β : -0.976 (95% CI: -1.582 to -0.371), respectively, all $p < 0.05$). There was no correlation between male sex and strain values in all models.

Table 4. Multivariate linear regression analysis for the impact of ESRD on aortic stiffness and strain parameters*

| | β | 95% CI | SE | p |
|--|---------|-------------------|-------|-------|
| Longitudinal distensibility | 0.000 | -0.001 to 0.001 | 0.001 | 0.324 |
| Longitudinal elastic modulus | 35.8 | -551.8 to 623.4 | 295.4 | 0.904 |
| Longitudinal pulse wave velocity | 23.4 | -184.7 to 231.6 | 104.6 | 0.823 |
| Longitudinal radial strain | 0.582 | -4.067 to 5.230 | 2.337 | 0.804 |
| Longitudinal radial strain rate | 0.419 | -0.259 to 1.097 | 0.341 | 0.222 |
| Transverse distensibility | -0.001 | -0.002 to 0.001 | 0.001 | 0.276 |
| Transverse elastic modulus | 451.8 | -98.8 to 1,002.4 | 276.8 | 0.106 |
| Transverse pulse wave velocity | 162.8 | -28.1 to 353.7 | 96.0 | 0.094 |
| Transverse radial strain | -6.087 | -10.790 to -1.384 | 2.364 | 0.012 |
| Transverse radial strain rate | -0.833 | -1.390 to -0.275 | 0.280 | 0.004 |
| Transverse circumferential strain | -6.771 | -11.525 to -2.018 | 2.389 | 0.006 |
| Transverse circumferential strain rate | -0.976 | -1.582 to -0.371 | 0.305 | 0.002 |

*Adjusted for age, gender, hypertension, diabetes mellitus, body mass index, creatinine level, high-density lipoprotein level, the presence of carotid plaque, and the maximal diameter of the common carotid artery.

CI: confidence interval, ESRD: end-stage renal disease, SE: standard error.

DISCUSSION

In the present study, speckle-tracking strain ultrasonography of the CCA was successfully performed in both normal subjects and patients with ESRD. When compared with patients with normal renal function, the CCA diameter was greater and the overall conventional aortic stiffness indices and strain parameters were reduced in patients with ESRD. In multivariate analyses, ESRD independently reduced both radial and circumferential strain parameters assessed on the transverse plane, while not exhibiting an impact on the longitudinal strain values and conventional aortic stiffness parameters.

Arterial remodeling in patients with ESRD is characterized by increased arterial diameters, outward remodeling, and arterial stiffening.^{11,12} These arterial remodeling also affects the arterial wall viscoelastic properties.¹¹ The alterations of the intrinsic elastic properties of arterial walls, caused by fibroelastic intimal thickening, calcification of vessel walls, inflammation, and increase in collagen, promote the hardening and stiffening of arteries in patients with ESRD.^{2,3} This changes in arterial viscoelastic properties have shown to be associated with the development of left ventricular hypertrophy and congestive heart failure.¹¹ In hemodialyzed patients, arterial stiffening was associated with calcification of the arteries and worsened as calcification increased.^{13,14} Recent studies have also emphasized the role of arterial stiffness as an independent cardiovascular risk factor in various disease populations.² In this regard, various indices that either directly or indirectly measure aortic stiffness have been developed and validated.

Conventional indices for aortic stiffness require the measurement of the central blood pressure and certain assumptions to be calculated. PWV, the most widely adopted technique for the assessment of aortic stiffness so far, assesses the propagation velocity of the pressure wave from the proximal aorta toward the peripheral vessels. PWV can be measured on any arteries, but only the carotid to femoral PWV has been proven to have a predictive value for clinical outcomes.¹⁵ PWV assesses the stiffness of an artery as a hollow structure, and is dependent on the artery geometry, the intrinsic properties of the arterial wall biomaterials, and blood density, which is difficult to measure in clinical setting and has to be assumed.

The speckle-tracking carotid strain ultrasonography enables the direct measurement of arterial dimension changes during the cardiac cycle. This straightforward method can

provide the stress-strain relationship and does not require the value of central blood pressure or any assumptions for the calculation, hence can be yielded at any given blood pressure level. Our group recently has shown the feasibility and value of carotid strain analysis in patients with stroke.¹⁶⁾ In the current study, the analysis of carotid strain was also easily and successfully completed in patients with ESRD.

ESRD is characterized by a high prevalence of cardiovascular risk factors, such as hypertension, diabetes mellitus, dyslipidemia, inflammation, and metabolic disturbances.¹⁷⁾ Exposure of arteries to these environments might induce arterial remodeling and stiffening. Previous studies emphasized the role of arterial stiffness in the development of cardiovascular diseases, and the parallel increases of the risk of cardiovascular disease and aortic stiffness along with the deterioration of CKD have already been observed.¹⁸⁾¹⁹⁾ In this, only circumferential wall stress was independently associated and occurred in parallel with the deterioration of CKD.¹⁸⁾ The aortic stiffness was greater in CKD patients even after age and blood pressure were adjusted for, when compared with healthy subjects and hypertensive patients.¹³⁾¹⁸⁾

In the current study, while the CCA diameter was greater in patients with ESRD, there were no significant differences in IMT, β stiffness index, and arterial compliance. This observation is in line with a recent study reporting that carotid IMT declined during the worsening of CKD.¹⁸⁾ Although arterial distensibility, elastic modulus, and PWV, among conventional aortic stiff indices, showed a significant difference between the patients with normal renal function and patients with ESRD, these parameters also have failed to demonstrate an independent association with ESRD in the multivariate analyses. In contrast, ESRD independently reduced both transverse radial and circumferential strain and strain rate. These results suggest that the carotid strain analysis in the transverse plane, which can be easily and readily be performed, might be a better parameter for the evaluation of aortic stiffness that has an independent relationship with ESRD.

This study had several limitations. First, due to the study's cross-sectional design, the prognostic value of the carotid strain parameters on clinical outcomes could not be evaluated. Second, because only patients with ESRD were enrolled, the feasibility and validity of the strain values in patients with CKD were not assessed. However, given the fact that strain values were the only parameters that were independently associated with ESRD in this study, prospective studies enrolling patients with both CKD and ESRD to determine the relationship between CKD deterioration and carotid strain, and the predictive value of strain analyses, now seem warranted.

In conclusion, we have demonstrated the feasibility of carotid strain analyses in patients with ESRD and its value in the assessment of aortic stiffness. ESRD had an independent and negative relationship only with transverse radial and circumferential strain and strain rate, but not with the conventional aortic stiffness indices. Multidirectional carotid strain analyses in patients with ESRD have the potential to provide useful information for risk stratification.

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REFERENCES

1. Lindner A, Charra B, Sherrard DJ, Scribner BH. Accelerated atherosclerosis in prolonged maintenance hemodialysis. *N Engl J Med* 1974;290:697-701.
[PUBMED](#) | [CROSSREF](#)
2. Briet M, Boutouyrie P, Laurent S, London GM. Arterial stiffness and pulse pressure in CKD and ESRD. *Kidney Int* 2012;82:388-400.
[PUBMED](#) | [CROSSREF](#)
3. Georgianos PI, Sarafidis PA, Lasaridis AN. Arterial stiffness: a novel cardiovascular risk factor in kidney disease patients. *Curr Vasc Pharmacol* 2015;13:229-38.
[PUBMED](#) | [CROSSREF](#)
4. London GM, Guérin AP, Marchais SJ, Métivier F, Pannier B, Adda H. Arterial media calcification in end-stage renal disease: impact on all-cause and cardiovascular mortality. *Nephrol Dial Transplant* 2003;18:1731-40.
[PUBMED](#) | [CROSSREF](#)
5. Kim SA, Park SM, Kim MN, et al. The relationship between mechanical properties of carotid artery and coronary artery disease. *Eur Heart J Cardiovasc Imaging* 2012;13:568-73.
[PUBMED](#) | [CROSSREF](#)
6. Bjällmark A, Lind B, Peolsson M, Shahgaldi K, Brodin LA, Nowak J. Ultrasonographic strain imaging is superior to conventional non-invasive measures of vascular stiffness in the detection of age-dependent differences in the mechanical properties of the common carotid artery. *Eur J Echocardiogr* 2010;11:630-6.
[PUBMED](#) | [CROSSREF](#)
7. Yang EY, Brunner G, Dokainish H, et al. Application of speckle-tracking in the evaluation of carotid artery function in subjects with hypertension and diabetes. *J Am Soc Echocardiogr* 2013;26:901-9.e1.
[PUBMED](#) | [CROSSREF](#)
8. Kidney Disease: Improving Global Outcomes (KDIGO) CKD-MBD Work Group. KDIGO clinical practice guideline for the diagnosis, evaluation, prevention, and treatment of Chronic Kidney Disease-Mineral and Bone Disorder (CKD-MBD). *Kidney Int Suppl* 2009:S1-130.
[PUBMED](#)
9. Veronesi F, Corsi C, Caiani EG, Sarti A, Lamberti C. Tracking of left ventricular long axis from real-time three-dimensional echocardiography using optical flow techniques. *IEEE Trans Inf Technol Biomed* 2006;10:174-81.
[PUBMED](#) | [CROSSREF](#)
10. Gamble G, Zorn J, Sanders G, MacMahon S, Sharpe N. Estimation of arterial stiffness, compliance, and distensibility from M-mode ultrasound measurements of the common carotid artery. *Stroke* 1994;25:11-6.
[PUBMED](#) | [CROSSREF](#)
11. Zoungas S, Cameron JD, Kerr PG, et al. Association of carotid intima-medial thickness and indices of arterial stiffness with cardiovascular disease outcomes in CKD. *Am J Kidney Dis* 2007;50:622-30.
[PUBMED](#) | [CROSSREF](#)
12. London GM, Guerin AP, Marchais SJ, et al. Cardiac and arterial interactions in end-stage renal disease. *Kidney Int* 1996;50:600-8.
[PUBMED](#) | [CROSSREF](#)
13. Sigrist MK, Taal MW, Bungay P, McIntyre CW. Progressive vascular calcification over 2 years is associated with arterial stiffening and increased mortality in patients with stages 4 and 5 chronic kidney disease. *Clin J Am Soc Nephrol* 2007;2:1241-8.
[PUBMED](#) | [CROSSREF](#)
14. Raggi P, Bellasi A, Ferramosca E, Islam T, Muntner P, Block GA. Association of pulse wave velocity with vascular and valvular calcification in hemodialysis patients. *Kidney Int* 2007;71:802-7.
[PUBMED](#) | [CROSSREF](#)
15. Blacher J, Guerin AP, Pannier B, Marchais SJ, Safar ME, London GM. Impact of aortic stiffness on survival in end-stage renal disease. *Circulation* 1999;99:2434-9.
[PUBMED](#) | [CROSSREF](#)
16. Yoon JH, Han D, Kim S, et al. Assessment of multidirectional movements of the common carotid artery in atherothrombotic stroke using dimensional speckle tracking carotid ultrasonography: A prospective, controlled cohort study. *Echocardiography* 2018;35:957-64.
[PUBMED](#) | [CROSSREF](#)
17. Palmer SC, Hayen A, Macaskill P, et al. Serum levels of phosphorus, parathyroid hormone, and calcium and risks of death and cardiovascular disease in individuals with chronic kidney disease: a systematic review and meta-analysis. *JAMA* 2011;305:1119-27.
[PUBMED](#) | [CROSSREF](#)

18. Briet M, Collin C, Karras A, et al. Arterial remodeling associates with CKD progression. *J Am Soc Nephrol* 2011;22:967-74.
[PUBMED](#) | [CROSSREF](#)
19. Go AS, Chertow GM, Fan D, McCulloch CE, Hsu CY. Chronic kidney disease and the risks of death, cardiovascular events, and hospitalization. *N Engl J Med* 2004;351:1296-305.
[PUBMED](#) | [CROSSREF](#)