INTRODUCTION

Transcranial Doppler (TCD) is frequently used as a substitute for brain angiography in a primary or outpatient setting. Practically, a number of TCD examinations are performed for explorative purpose in patients with complaint of headache. Of these, migraine patients frequently show abnormal Doppler findings before undergoing brain angiographies. In such cases, it is difficult to decide whether the abnormal Doppler findings indicate an undiagnosed intracranial arterial stenosis (ICAS) because many migraine patients exhibit an increase in flow velocity in the intracranial arteries without the presence of stenotic lesions. Apparently, most of the clinicians would obtain the angiographic data to determine the presence or absence of cerebrovascular abnormality in such patients.

To date, no study has compared the difference in abnormal Doppler findings between patients with migraine and those with ICAS. At present, it may be impossible to distinguish migraine from ICAS on the basis of Doppler abnormalities alone. Accordingly, in this study, we aimed to analyze the abnormal Doppler findings purely resulting from migraine pathophysi-
ology by comparing the features of TCD abnormality between migraine and ICAS.

PATIENTS AND METHODS

1. Study design and patients

From the TCD registry, we initially obtained the names and registry numbers of 72 migraine patients who underwent brain computed tomographic angiography (CTA) and TCD on the same day from May 2015 to June 2018. From the selected patients, 25 with normal TCD, four with any discernible stenosis on CTA, and two with transtemporal approach failure were excluded to analyze the abnormal Doppler findings purely arising from migraine. Consequently, 41 migraine patients (including seven with aura) with abnormal TCD but normal CTA were included in the analysis.

Thirty-five patients with TCD abnormalities due to any discernible ICAS were recruited by reviewing the TCD registry and their medical records. We tried to match their ages with those of migraine patients. All ICAS patients underwent TCD within 1 week after undergoing brain CTA or magnetic resonance angiography (MRA) during the same period. The ICAS group consisted of 27 patients with large artery atherosclerosis, five with moyamoya disease, two with intracranial vertebrobasilar dissection, and one with cerebral vasculitis. This study was approved by the Institutional Review Board of Soonchunhyang University Bucheon Hospital (SCHB 2018-09-021), and the requirement for informed consent was waived due to the retrospective nature of our study.

2. Clinical assessment

The following clinical data were obtained: age, gender, hypertension, diabetes mellitus, hyperlipidemia, current smoking, coronary artery disease, blood pressure at the TCD study, and complete blood count (white blood cell count, hemoglobin/hematocrit, and platelet count) within 1 week of the TCD study. Migraine was diagnosed according to the criteria of the International Classification of Headache Disorders 2nd edition.3

3. Imaging protocols

Brain CTA was performed with a 128-detector high-definition CT scanner (Discovery CT750 HD; GE Healthcare, Milwaukee, Wisconsin, USA) using standardized protocol (section thickness, 0.625 mm; tube voltage, 100 kV; and tube current, 300 mA). Images were scanned after a single bolus injection of 100 mL of nonionic contrast agent via an 18-gauge cannula inserted into an antecubital vein. Images were obtained in series from the aortic arch to the vertex. Brain MRA was performed on a 3.0 Tesla MRI scanner (MAGNETOM Skyra; Siemens, Erlangen, Germany), and RF coil for head and neck examination (head/neck 64 channel coil; Siemens) was utilized. To obtain MRA images, a three-dimensional time-of-flight pulse sequence method was used. The brain CTA and MRA were interpreted by two experienced neuroradiologists including AL.

4. TCD protocol and variables

TCD examination was performed in a quiet room according to previously recommended practice standards.4,5 Power M-mode TCD (ST3, Spencer Technologies, Seattle, WA, USA) was used for all examinations. Mean flow velocity (mFV) and pulsatile index (PI) of the siphon internal carotid artery (sICA) were recorded at a depth of 58–65 mm, middle cerebral artery (MCA) at 40–65 mm, anterior cerebral artery (ACA) at 60–75 mm, posterior cerebral artery (PCA) at 55–75 mm, vertebral artery (VA) at 40–75 mm, and basilar artery (BA) at 80–105 mm. Two different experienced sonographers were blinded to the clinical data while performing the TCD studies. The results were interpreted through consensus among two investigators (S.J.L. and H.K.) according to previously published interpretation criteria.5,6 For the bilateral arteries, side-to-side differences (%) in mFV were calculated by dividing the difference between the mFVs of bilateral arteries by the mFV of the lower value. Turbulent flow was defined as any definite increase in low-frequency components within the Doppler spectrum including bidirectional low-frequency noise and arterial wall covibration. For MCA, VA, and BA, focal increase in mFV was determined as more than 30 cm/s or 30% of the difference between arterial segments within the same artery (Fig. 1).
5. Statistical analysis

Using independent t-test or chi-square test, the demographic variables, vascular risk factors, blood pressure, complete blood counts, and the TCD variables from the arteries with Doppler abnormality were compared between the migraine group and ICAS group. In addition, the absence of turbulent flow and mFV of <120 cm/s in any examined vessel were used to calculate the sensitivity and specificity for diagnosing migraine in the cohort. Statistical analyses were performed with SPSS software, version 21.0 (SPSS Inc., Chicago, IL, USA). p values <0.05 were considered significant.

RESULTS

A total of 76 patients (41 migraine patients and 35 ICAS patients) were included in the analysis. The migraine group consisted of an overwhelmingly higher proportion of women (92.7%), while the ICAS group had a higher proportion of men (62.9%). Expectedly, the
ICAS group had a significantly higher prevalence of vascular risk factors including hypertension, diabetes, hyperlipidemia, and current smoking than the migraine group. Coronary artery disease tended to be frequent in the ICAS patients (p=0.056). Additionally, age tended to be higher in the ICAS patients (p=0.099), but the age difference between the two groups was not statistically significant. White blood cell count, hemoglobin, hematocrit, and systolic and diastolic blood pressures were significantly higher in the ICAS group (Table 1).

In the 41 migraine patients, the most frequent vessel with TCD abnormality was MCA (32 patients, 78.0%), followed by VA (26 patients, 63.4%), sICA (10 patients, 24.4%), BA (eight patients, 19.5%), ACA (six patients, 14.6%), and PCA (0 patient). Among the 35 ICAS patients, the most frequent vessel with Doppler abnormality was MCA (27 patients, 77.1%), followed by sICA (15 patients, 42.9%), ACA (15 patients, 42.9%), VA (11 patients, 31.4%), BA (seven patients, 20.0%), and PCA (one patient, 2.9%) (Fig. 2). Therefore, PCA was excluded from the analysis.

Table 2 shows a comparative analysis of Doppler findings between patients with migraine and those with ICAS. For all the analyzed intracranial arteries, turbulent flow was significantly more frequent in the ICAS group than in the migraine group. The prevalence of side-to-side difference >30% was significantly higher in the MCA and VA of the ICAS patients than in those of migraine patients, but the prevalence of side to side difference >30% in the sICA and ACA was statistically similar between the two groups.

Focally increased mFV was more frequently observed in the MCA of ICAS patients than in that of migraine patients, but there was no significant difference in the BA and VA between the two groups. Compared with that of migraine patients, the mFV of ICAS patients not only had a wide range of distribution, but was also significantly higher in the MCA, VA, and BA, and was non-significantly (p<0.1) increased in the sICA and ACA.

### Table 1. Basic characteristics of patients with migraine vs. intracranial arterial stenosis

|                         | Migraine (n=41) | Intracranial arterial stenosis (n=35) | p-value |
|-------------------------|-----------------|--------------------------------------|---------|
| Age (years)             | 41.3±9.8        | 45.1±9.7                             | 0.099   |
| Female                  | 38 (92.7)       | 13 (37.1)                            | <0.001  |
| Hypertension            | 4 (9.8)         | 27 (77.1)                            | <0.001  |
| Diabetes                | 0 (0)           | 14 (40.0)                            | <0.001  |
| Hyperlipidemia          | 0 (0)           | 24 (68.6)                            | <0.001  |
| Current smoking         | 3 (7.3)         | 12 (34.3)                            | 0.003   |
| Coronary artery disease | 0 (0)           | 3 (8.6)                              | 0.056   |
| Systolic blood pressure (mmHg) | 120.6±21.5 | 150.1±22.2                          | <0.001  |
| Diastolic blood pressure (mmHg) | 76.9±12.0 | 91.0±17.5                           | <0.001  |
| Complete blood count    |                 |                                      |         |
| White blood cell (x10³/μL) | 7.0±2.4    | 8.6±2.7                              | 0.015   |
| Hemoglobin (g/dL)       | 13.1±1.4        | 14.0±1.4                             | 0.002   |
| Hematocrit (%)          | 38.7±4.4        | 41.0±3.6                             | 0.024   |
| Platelet count (x10³/μL) | 267.9±54.6   | 258.1±102.5                          | 0.659   |

Variables were analyzed using independent t-test (mean±standard deviation) or chi-square test (n [%]).

FIG. 2. Prevalence of arteries with Doppler abnormality in patients with migraine and intracranial arterial stenosis (ICAS). sICA; siphon internal carotid artery, MCA; middle cerebral artery, ACA; anterior cerebral artery, PCA; posterior cerebral artery, VA; vertebral artery, BA; basilar artery.
On the contrary, the mFV in migraine patients was only moderately increased: the maximum value was below 95 cm/s in the sICA and ACA, 120 cm/s in the MCA, and 85 cm/s in the BA and VA. PI was nonsignificantly higher in the VA of ICAS patients, but was significantly increased in the ACA of ICAS patients compared with their counterpart.

The absence of turbulent flow and mFV <120 cm/s in any examined vessel were used to calculate the sensitivity and specificity for the diagnosis of migraine in our cohort. The absence of turbulence flow had a sensitivity of 92.5% and a specificity of 88.0%, while mFV <120 cm/s had a sensitivity of 74.5% and a specificity of 100% to diagnose migraine.

### Table 2. Comparison of Doppler findings between patients with migraine vs. ICAS

| Vessels with Doppler abnormality | Migraine | ICAS | p-value |
|---------------------------------|----------|------|---------|
| Number of sICA                  | 13       | 21   | 0.097   |
| mFV (cm/s)*                     | 78.5±6.3 (72-92) | 87.6±22.7 (53-146) | 0.097   |
| Pulsatile index                 | 0.68±0.15 (0.42-0.98) | 0.67±0.16 (0.40-0.93) | 0.825   |
| Side to side difference >30%   | 3 (23.1) | 6 (28.6) | 1.000   |
| Turbulent flow                  | 2 (15.4) | 17 (81.0) | <0.001 |

| Number of MCA                   | 54       | 44   | <0.001 |
| mFV (cm/s)*                     | 90.4±7.5 (81-116) | 140.8±62.7 (67-299) | <0.001 |
| Pulsatile index                 | 0.65±0.11 (0.44-0.93) | 0.68±0.18 (0.20-1.08) | 0.337   |
| Side to side difference >30%   | 0 (0)    | 32 (72.7) | <0.001 |
| Turbulent flow                  | 2 (3.7)  | 36 (81.8) | <0.001 |
| Focal increase in mFV†          | 6 (11.1) | 25 (56.8) | <0.001 |

| Number of ACA                   | 7        | 21   | 0.059   |
| mFV (cm/s)*                     | 86.3±4.7 (81-94) | 107.5±28.0 (79-199) | 0.059   |
| Pulsatile index                 | 0.58±0.10 (0.43-0.74) | 0.75±0.17 (0.54-1.25) | 0.017   |
| Side to side difference >30%   | 3 (42.9) | 8 (38.1) | 1.000   |
| Turbulent flow                  | 0 (0)    | 15 (71.4) | 0.001   |
| Number of VA                    | 39       | 18   | 0.002   |
| mFV (cm/s)*                     | 58.5±7.2 (50-83) | 108.6±58.3 (51-251) | 0.002   |
| Pulsatile index                 | 0.66±0.12 (0.46-0.91) | 0.72±0.10 (0.61-0.95) | 0.075   |
| Side to side difference >30%   | 3 (7.7)  | 9 (50.0) | 0.001   |
| Turbulent flow                  | 0 (0)    | 15 (83.3) | <0.001 |
| Focal increase in mFV†          | 21 (53.8) | 12 (66.7) | 0.362   |

| Number of BA                    | 8        | 7    | 0.030   |
| mFV (cm/s)*                     | 71.9±7.4 (63-81) | 103.3±29.3 (64-142) | 0.030   |
| Pulsatile index                 | 0.69±0.9 (0.56-0.81) | 0.76±0.19 (0.52-1.00) | 0.378   |
| Turbulent flow                  | 0 (0)    | 4 (57.1) | 0.026   |
| Focal increase in mFV†          | 2 (25.0) | 4 (57.1) | 0.315   |

Variables were analyzed using independent t-test (mean±standard deviation [range]) or chi-square test (n [%]).

ICAS; intracranial arterial stenosis, sICA; siphon internal carotid artery, mFV; mean flow velocity, MCA; middle cerebral artery, ACA; anterior cerebral artery, VA; vertebral artery, BA; basilar artery
*Maximum mean flow velocity at each vessel was used.
†Focal increase in mean flow velocity was defined as 30 cm/s or 30% of the difference between arterial segments within the same artery.
DISCUSSION

For all the analyzed arteries, turbulent flow was the persistent feature of ICAS distinct from migraine. It can be explained by the fact that TCD directly insonates stenosed large arteries in the ICAS patients. The stenosis geometry is prone to cause disturbed flow rather than smooth laminar flow. Furthermore, the localized and lateralized nature of stenotic lesions is more likely to cause a higher prevalence of side-to-side difference and focal increase in mFV in ICAS. The features, in particular, were well presented in the MCAs of ICAS patients, which are not only less variant, but also have a straight course allowing a favorable angle of insonation.

In contrast, increased flow velocity in migraine patients is assumed to be attributable to arteriolar vasodilatation rather than reduced lumen diameter in the insonated large arteries. This stands to reason in our study because we recruited migraine patients with normal CTA, which was performed on the same day with TCD. As previously reported, increased neuronal excitability and neurovascular coupling in migraine could be linked with reduced arteriolar resistance and increased regional blood flow, which can result in the increased flow velocity in the insonated large arteries. From the pathophysiological aspect, even increase in flow velocity along the large arteries is anticipated, and thus turbulent flow and focal increase in mFV might well be less prevalent in migraine patients.

The mFV was significantly or non-significantly higher in ICAS patients. According to a simple model of the continuity principle, flow velocity increases by an amount inversely proportional to the reduction of arterial lumen area if other factors (e.g., collateral channel, blood viscosity, blood pressure) are excluded. Therefore, the flow velocity in ICAS can markedly increase with a wide distribution of values as the stenosis progresses. By contrast, the mFV in migraine was only modestly increased in line with previous studies (Fig. 1).

Focally increased mFV was more frequently found in the MCA of ICAS patients than in that of migraine patients. However, there was no significant difference in the prevalence in the VA and BA between the two groups. Particularly, 53.8% of VA in migraine patients had a focal increase in mFV. It may be because of the frequent tortuous courses with morphological variations in posterior circulation arteries in addition to the incapability of angle correction in the TCD machine used in the study. For this reason, VA is one of the arteries with the lowest sensitivity for TCD diagnosis of its stenosis.

In summary, turbulent flow with a marked elevation, focal variation, and asymmetry of mFV favor the diagnosis of ICAS, while a relatively uniform and mild to moderate increase in mFV favor the diagnosis of migraine if other influencing factors are excluded. The results seem to reflect the different locations where the luminal diameter of vessels changes in each disease entity.

To the best of our knowledge, this is the first study to compare the abnormal TCD findings of migraine patients with those of ICAS patients. This approach can provide a useful insight to interpret the Doppler findings of patients with migraine from the pathophysiological aspect and thus bring out a further clinical use of TCD in real-world headache practice. However, this study has some limitations. It was based on a small number of data obtained from a single center. Therefore, the statistical power is too low for multivariable analysis. Besides, age could not be completely matched in the process of patient selection. That is to say, the patients in the ICAS group tended to be older (p<0.05). This is because the ICAS patients in the Soonchunhyang University Bucheon Hospital are mostly older individuals. The incompleteness in age matching might have led to the differences in the results for some measurements (e.g., increased PI) in the present study. Additionally, there were differences in blood pressure and blood constituents (hemoglobin and hematocrit) between the two groups. It could have affected flow velocity and PI in our data. In addition, we could not determine if it was ictal or inter-ictal phase when TCD was performed in the migraine patients because of the retrospective nature of our study. As previous studies have demonstrated an increased flow velocity during the headache-free period, in the migraine status of most of our patients are believed to have corresponded to the inter-ictal phase of migraine.

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Conflicts of Interest
No potential conflicts of interest relevant to this article were reported.

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