Relationship between Transcranial Doppler and CT Data in Acute Intracerebral Hemorrhage

Joan Martí-Fàbregas, Roberto Belví, Esteve Guàrdia, Dolores Cocho, and Josep-Lluis Martí-Vilalta

BACKGROUND AND PURPOSE: It would be useful to have a noninvasive test for correlation with CT findings in patients with intracerebral hemorrhage (ICH). We determined which transcranial Doppler (TCD) variables are related to which CT data in patients with ICH.

METHODS: We prospectively included 51 patients (age ± SD, 66.2 ± 12.4 years; 30 men, 21 women) with first-ever supratentorial, nontraumatic ICH. CT and TCD examination were performed in the acute stage (less than 12 hours from symptom onset). TCD recordings were obtained from the middle cerebral arteries, and the following variables were analyzed: systolic ($V_s$), diastolic ($V_d$), mean ($V_m$) velocities, and pulsatility index from the affected (a) and unaffected (u) hemispheres.

RESULTS: PIs obtained for both hemispheres were positively correlated with hematoma volume (aPI, $r = 0.43, P = .001$; uPI, $r = 0.44, P = .001$), volume of hypoattenuation (aPI $r = 0.64, P < .0001$; uPI, $r = 0.39, P = .005$), total volume (aPI, $r = 0.59, P < .0001$; uPI, $r = 0.48, P < .0001$), and midline shift (aPI, $r = 0.28, P = .04$; uPI, $r = 0.29, P = .03$). Both PIs were increased in patients with intraventricular hemorrhage (aPI, $P = .01$; uPI $P = .004$). No TCD parameter was correlated with ventricular size.

CONCLUSION: Most TCD parameters were correlated with CT data in the acute stage of ICH. An increase in PI probably reflects intracranial hypertension and mass effect. Further studies are needed to determine the clinical application of our findings.

Acute spontaneous intracerebral hemorrhage (ICH) is a disease with a 23–58% mortality rate within 6 months (1). Half of the deaths occur during the first 2 days (2) of the onset of the ICH and are attributable to intracranial hypertension and mass effect. The most important predictors of outcome are the Glasgow Coma Scale score and the volume of hematoma (1). Other CT abnormalities, such as ventricular enlargement and secondary intraventricular hemorrhage, have also been prognostic (3–5).

The sudden eruption of an intracranial mass destroys and displaces brain tissue and can induce an increase in intracranial pressure (ICP). Recent data indicate that ICH is a dynamic disease. During the first hours or days following appearance of the primary lesion, some patients have hematoma growth, perilesional edema, perilesional ischemia, hydrocephalus, or secondary intraventricular hemorrhage (6–8). All of these complications potentially increase ICP and mass effect, finally resulting in neurologic deterioration. However, the information that CT provides is static. As a result, it is not practical to perform frequent control CT studies in these patients. Although invasive devices can be used to monitor the ICP, the procedure has some risks (e.g., infection and tissue lesion) and pitfalls (e.g., those variability inherent to the technique, displacement, unilateral information). Moreover, the value of this approach in the management of ICH is equivocal (9–12). In view of these limitations, it would be useful to have a noninvasive test that can be used for correlation with CT and that provides dynamic information for reliable treatment and prognosis.

Increased ICP and decreased cerebral perfusion pressure (CPP) give rise to typical changes in the Doppler waveform obtained by transcranial sonation (i.e., a decrease of diastolic velocity and an increase in the pulsatility index [PI]) (13–15). Transcranial Doppler (TCD) could be useful for noninvasively and indirectly assessing mass effect and ICP in ICH. Data on the relation of radiologic data to one or

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From the Departments of Neurology, Cerebrovascular Unit (J.M.-F., R.B., D.C., J.-L.M.-V.), and Radiology, Section of Neuroradiology (E.G.), Hospital de la Santa Creu i Sant Pau, Universitat Autònoma de Barcelona, Spain.
Supported by a grant from Fondo de Investigaciones Sanitarias (00/0029), Ministerio de Sanidad, Spain.
Address reprint requests to Dr Joan Martí-Fàbregas, Servei de Neurologia, Hospital de la Santa Creu i Sant Pau, Avda Sant Antoni M Claret, 167, 08025 Barcelona, Spain.
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more specific TCD variables are sparse. In one study
(16), the PI obtained from the unaffected hemisphere
was an independent predictor of mortality in acute
ICH. In response to the lack of information, we stud-
ted the hematoma volume and other CT data, along
with TCD findings, in a prospective series of patients
in the acute stage of ICH.

Methods

Patients

This study was conducted at a university hospital that
provides neurosurgical, neuroradiologic, and intensive care assis-
tance in addition to neurologic evaluation. Therefore, it is a
referral center for the care of patients with ICH.

We included patients whose symptoms began within 12
hours from the time they were included in the study. Their
symptoms were attributable to a first-ever supratentorial, sin-
gle, nontraumatic ICH, as diagnosed on cranial CT. We ex-
cluded patients who needed emergency surgical evacuation,
those with a nonsuitable temporal acoustic bony window, and
those with an imprecise time for the onset of symptoms. Also
excluded were patients with previous clinical cerebrovascular
disease (ischemic or hemorrhagic), epidural or subdural hema-
toma at initial CT, and those for whom the investigating team
was not contacted. As it is difficult to know the cause of such
hemorrhage in the first hours, all etiologies were included.

Our hospital Ethics Research Committee approved the
study protocol. In addition, patients or their legal representa-
tives gave written consent to participate. Patients were treated
according to a previously approved protocol written jointly by
neurologists, neurosurgeons, and intensive care specialists at
our hospital. Glasgow Coma Scale and National Institute of
Health and Stroke (NIHS) Scale scores were obtained.

CT Evaluation

For consistency and reliability, the same neuroradiologist
(E.G.) analyzed all of the CT scans blinded to the clinical and
TCD data. Axial 5-mm-thick contiguous sections were obtained
from the supratentorial region, while 3-mm-thick contiguous
sections were obtained from the infratentorial fossa. CT scans
were stored in the hard disk of the CT apparatus (Asion;
Toshiba, Nasu, Japan). Quantitative data were obtained with
the aid of computer-assisted measurements provided by the CT
apparatus.

In each CT examination, the following information was
obtained: 1) topography, which was classified as lobar or
deep; 2) hematoma volume, which was calculated according to
the \((A \times B \times C)/2\) method applied to the hyperattenu-
ating lesion (2), 3) perilesional hypoattenuation, which was
also calculated by applying the \((A \times B \times C)/2\) method to the
hypodensities surrounding the haemorrhage and by subtracting
from this value the hematoma vol-
"u
ume; 4) total volume, which was calculated as the sum of
the hyperattenuating and hypoattenuating volumes; and 5) mid-
line shift, which was assessed by calculating the lateral dis-
placement (in centimeters) from the midline of the septum
pellucidum, the third ventricle, or the pineal gland (A line
was traced from the anterior insertion of the falk cerebi
to the torcula, and the midline shift was calculated at the most
displaced midline structure; ) 6) ventricular size, which was
calculated according to the Evans ventricular ratio, or the
ratio between the maximum spread of ventricular horns to
the breadth of cranial cavity (17); and 7) intraventricular
hemorrhage. Irrespective of its amount or localization, any
intraventricular hyperattenuating image not attributable to
calcification or choroid plexus was considered evidence of
intraventricular hemorrhage.

TCD Protocol

We used a portable TCD instrument (Multi-Dop2, DWL
Elektroniche Systeme GmbH, Baden-Württenberg, Germany).
Both middle cerebral arteries (MCAs) were insonated. These
arteries were chosen because they provide 80% of the hemi-
spheric blood flow, and it is easy to detect the flow due to the
spatial orientation in relation to the probe. TCD examination
was performed as soon as possible after the CT diagnosis was
available, and always within 12 hours of the onset of symptoms.

The MCA were examined through the temporal acoustic
window at a depth of 45–55 mm with the patient in the supine
position. The MCA from the unaffected hemisphere was ex-
"a
"amined first, and then the MCA ipsilateral to the hematoma
was examined. When a clearly readable waveform was ob-
tained, it was used in the calculations. The spectra cycles
contained in a 6-second frame were analyzed. If all of the
spectra were technically interpretable, the time-averaged re-
sults automatically calculated by the machine were considered
reliable.

The following variables were analyzed: systolic (\(V_s\)) di-
astolic (\(V_d\)), mean (\(V_m\)) velocities, and PI from the affected (a)
and unaffected (u) hemispheres. However, when results of one
or more cycles were not interpretable, the measurements were
calculated from the best spectra on the printed paper or on the
monitor screen. \(V_m\) was calculated according to the equation
\(V_m = (V_s - V_d)/3 + V_d\), and PI was calculated according to
the formula \(PI = (V_s - V_d)/V_m\). Normal values for PI \(\pm SD\)
were 0.98 \(\pm 0.2\) (18).

Statistical Analysis

The statistical analyses included the following CT data: he-
matoma volume, volume of hypoattenuation, total volume,
midline shift, ventricular ratio, and intraventricular hemor-
 rhage (yes/no). Continuous variables were correlated with the
eight TCD variables: four on the affected side, or a\(s\), a\(m\), a\(d\), and aPI, and four on the unaffected side, or u\(s\), u\(m\), u\(d\),
and uPI. Pearson correlation coefficients were obtained.

For binary variables, the mean and SD of the eight TCD
variables were compared in the two groups with the Student t
test. Because of the large SD detected in the CT data, we
performed the Kolmogorov-Smirnov test to determine if these
variables were normally distributed. In addition, we analyzed
the same variables with nonparametric statistical tests (Spear-
man correlation and Mann-Whitney test). A result was consid-
ered statistically significant when \(P < .05\). The statistical anal-
"y
"yses were performed with the aid of SPSS version 10 software
(SPPS, Chicago, IL).

Results

During a 2-year period, 138 consecutive patients
received a diagnosis of spontaneous supratentorial
ICH. Patients were excluded for the following rea-
sons: emergency evacuation of hematoma (\(n = 7\)),
admission after 12 hours from the onset of symptoms
or an unknown time of onset (\(n = 29\)), deficient
acoustic window (\(n = 5\)), previous cerebrovascular
disease (\(n = 11\)), or contact with the investigator team
after the 12-hour window (\(n = 17\)). In addition, 16
patients were excluded because they died before TCD
study was performed, and two patients were excluded
because of diagnostic errors. Therefore, this study
consisted of 51 patients (30 men, 21 women), whose
mean age was 66.2 \(\pm 12.4\) years.

First CT was performed a mean of 190 \(\pm 175\)
minutes (range, 15–702 minutes) after the onset of
symptoms, while the first TCD study was performed
after a mean of 340 ± 219 minutes (range, 60–720 minutes) after the onset of symptoms. Suspected etiologies were high blood pressure (n = 29), anticoagulant therapy (n = 2), other coagulation abnormalities (n = 2), arteriovenous malformation (n = 1), tumor (n = 1), other etiologies (n = 2), and unknown (n = 14). The median Glasgow Coma Scale score was 14, while the median NIH Scale score was 17. The mortality rate at 30 days was 33% (n = 17).

Data from acute CT are shown in Table 1. About 25% of the patients had hematomas of more than 50 mL, and the range of hematoma volume was wide (0.05–344 mL). Secondary intraventricular hemorrhage was present in 55% of the patients. The TCD data are shown in Table 2. Values were comparable in the affected and unaffected hemispheres (Student’s t test).

Mean PI in the acute stage was clearly above normal values in both hemispheres. Table 3 and Figure 1 shows the relationships between quantitative CT variables and TCD data obtained with Pearson correlation coefficient. We found a positive correlation between the PI from both hemispheres and the volume of hematoma (aPI, r = 0.43, P < .001; uPI, r = 0.44, P = .001), the hypotauenating volume surrounding the hematoma (aPI, r = 0.64, P < .0001; uPI, r = 0.39, P = .005), total volume (aPI, r = 0.59, P < .0001; uPI, r = 0.48, P < .0001), and midline shift (aPI, r = 0.28, P = .04; uPI, r = 0.29, P = .03). Vd from the affected hemisphere was negatively correlated with the volume of hypotauenation (r = −0.32, P = .02) and the total volume (r = −0.32, P = .02). No statistically significant correlations were found between systolic or mean velocities and any of the CT data. No TCD variable was correlated with the ventricular ratio, and TCD results were equivalent in patients with deep or lobar ICH. Level of consciousness measured with the Glasgow Coma Scale was negatively correlated with aPI (r = −0.53, P < .0001) and uPI (r = −0.61, P < .0001). Neurologic severity measured with the NIH scale showed a positive correlation with the PI from both hemispheres (aPI, r = 0.52, P < .0001; uPI, r = 0.55, P < .0001).

Table 4 shows the comparison of TCD variables between the group with and the group without intraventricular hemorrhage. Both aPI and uPI were significantly higher in the group with intraventricular hemorrhage (aPI, P = .01; uPI P = .004). Vd from the affected hemisphere was lower in patients with intraventricular hemorrhage (P = .03). Mean and systolic velocities did not show differences in patients with or without intraventricular hemorrhage. Figure 2 presents an example of the TCD and CT results in one representative patient.

The Kolmogorov-Smirnov test showed that the distribution of all CT data were not normal. Therefore, we evaluated the data by using nonparametric analyses that also showed the following correlations: volume of the hematoma (aPI, r = 0.61, P < .0001; uPI, r = 0.56, P < .0001); volume of hypoattenuation (aPI, r = 0.56, P < .0001; uPI, r = 0.45, P = .01; and aPI, r = −0.31, P = .02); the total volume (aPI, r = 0.66, P < .0001; uPI, r = 0.58, P < .0001; aVd, r = −0.36, P = .008; and uVd, r = −0.30, P = .03); and the midline shift (aPI, r = 0.50, P < .0001; uPI, r = 0.50, P < .0001; and aVd, r = −0.34, P = .01). Patients with intraventricular hemorrhage had significantly higher aPI (P = .01) and uPI (P = .03) than those of patients without intraventricular hemorrhage.

### Discussion

By insonating the MCA in patients during the acute stage of ICH, we found that TCD data are related to CT data. Considering all of the patients, we found an increase of mean PI in both hemispheres in the acute stage. We also found that the PI obtained from the affected and unaffected hemispheres were correlated with those CT signs associated with mass effect: volume of the hematoma, volume of surrounding edema, total volume, midline shift, and intraventricular hemorrhage. TCD measurements that were most often correlated with the CT data were either an increase of the PI (aPI or uPI) or a decrease in the Vd. In fact, as the PI was obtained by the ratio (Vd − Vd/Vm, the decrease in Vd made the numerator higher and the denominator lower, increasing the value of the PI.

Early mortality in patients with ICH is sometimes related to an increase of ICP and mass effect, whereas death in the chronic stage is often attributed to respiratory infections or other consequences of immobility (19, 20). Accordingly, hematoma volume (the main cause of increased ICP) and decreased level of consciousness (a consequence of increased ICP) are the main predictors of outcome in ICH. Experimental studies have found immediate and sustained increases in ICP after an ICH (21, 22). These findings have

### Table 1: CT data from the acute stage of ICH

<table>
<thead>
<tr>
<th>A: Topography and hemorrhage</th>
<th>n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Topography</td>
<td></td>
</tr>
<tr>
<td>Lobar</td>
<td>19 (37)</td>
</tr>
<tr>
<td>Deep</td>
<td>32 (63)</td>
</tr>
<tr>
<td>Intraventricular hemorrhage</td>
<td>28 (55)</td>
</tr>
<tr>
<td>B: Volume, shift, and ratio</td>
<td></td>
</tr>
<tr>
<td>Hematoma volume (mL)</td>
<td>51.7 ± 70.4 (0.05–344 mL)</td>
</tr>
<tr>
<td>Edema volume (mL)</td>
<td>28.4 ± 38.1 (0–210.5 mL)</td>
</tr>
<tr>
<td>Total volume (mL)</td>
<td>80.1 ± 93.7 (0.28–348 mL)</td>
</tr>
<tr>
<td>Midline shift (cm)</td>
<td>0.56 ± 0.79 (0–4 cm)</td>
</tr>
<tr>
<td>Ventricular ratio</td>
<td>0.23 ± 0.04 (0.11–0.33)</td>
</tr>
</tbody>
</table>

### Table 2: TCD results in the acute stage of ICH

<table>
<thead>
<tr>
<th>Result</th>
<th>Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>aVd (cm/s)</td>
<td>79.2 ± 36.0</td>
</tr>
<tr>
<td>aVf (cm/s)</td>
<td>19.7 ± 11.1</td>
</tr>
<tr>
<td>aVm (cm/s)</td>
<td>35.6 ± 16.5</td>
</tr>
<tr>
<td>aPI</td>
<td>1.85 ± 0.87</td>
</tr>
<tr>
<td>uVd (cm/s)</td>
<td>81.2 ± 34.2</td>
</tr>
<tr>
<td>uVf (cm/s)</td>
<td>22.8 ± 13.6</td>
</tr>
<tr>
<td>uVm (cm/s)</td>
<td>39.6 ± 18.0</td>
</tr>
<tr>
<td>uPI</td>
<td>1.62 ± 0.93</td>
</tr>
</tbody>
</table>
been confirmed in clinical studies in which high ICP and mass effect were recorded after stroke (12, 23). Our results agree with those of several authors who found that the progressive increase in ICP and decrease in CPP induce characteristic changes in the Doppler waveform (10, 13–15, 24, 25): a progressive decrease of $V_d$ and a progressive increase in PI, which is more marked with ICP above 60 mmHg (15). The reduction in $V_d$ is accounted for by the increase in peripheral cerebrovascular resistance, which is determined mainly by the ICP and the diameter of the arterioles (26). As a consequence, TCD has been used to indirectly estimate ICP and CPP in patients with severe head injuries (10, 14, 25–29). These experimental studies demonstrate a high correlation between ICP and PI, either linear or exponential, and an inverse correlation between CPP and PI (14, 15, 29). Therefore, some authors have used TCD to monitor the effect of different therapies (30, 31). An alternative explanation for an increase in PI is vasospasm, but this is unlikely in spontaneous ICH. In our study, the PI from both hemispheres was correlated with the hypoattenuating area surrounding the hematoma. Although the exact composition of this perihematoma hypoattenuation in the acute stage is controversial, our finding indicates that it also contributes to intracranial hypertension (as it is correlated with an increase in PI).

Few groups have analyzed the correlations between TCD and CT data in spontaneous ICH. Mayer et al (32) examined 30 patients with supratentorial ICH and found an elevation of aPI, the aPI/uPI ratio, and uPI in hematomas larger than 25 mL. We also found that the PI obtained from both hemispheres were directly correlated with the volume of the hematoma, the volume of edema, and the total volume and that the PI was higher in patients with secondary intraventricular hemorrhage. Our study is not entirely comparable to that of Mayer et al, because they included patients admitted within 24 hours of symptom onset (vs. 12 hours in our series), they excluded patients with Glasgow Coma Scale scores between 3 and 5, and TCD examination was performed after considerable delay (on either the first or second day of hospitalization). Moreover, our series included larger hematomas, and we did not dichotomize the hematoma volumes. Despite these differences, it is conspicuous that our results agree with those of Mayer et al.

Our study has some limitations. We studied only the MCA, but we do not know how relevant the TCD findings from other arteries of the anterior or posterior circulation are. We chose the MCA because findings were easily obtainable and because MCA carries 80% of the blood flow to the hemispheres of the brain. TCD values change depending on the hematocrit, pCO$_2$, systemic blood pressure, and autoregulatory capacity. However, these variables affect mainly flow velocities and their ratios (such as PI), although to a lesser degree. Recent findings suggest that autoregulation is preserved in ICH (33). Our data are based on a single TCD examination, but because of the broad interindividual variability of measured TCD parameters, it would be interesting to follow up individual subjects over time to see if a change in any CT and clinical parameter is correlated with a change in a TCD parameter. We assumed that the TCD changes provoked by ICH were related to intracranial hypertension. However, we did not mea-

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### TABLE 3: Relationship between TCD and quantitative CT variables

<table>
<thead>
<tr>
<th>Variable</th>
<th>Hematoma Volume</th>
<th>Hypoattenuating Volume</th>
<th>Total Volume</th>
<th>Midline Shift</th>
<th>Ventricular ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>$aV_d$ (cm/s)</td>
<td>$r$</td>
<td>$P$ Value</td>
<td>$r$</td>
<td>$P$ Value</td>
<td>$r$</td>
</tr>
<tr>
<td>$aPI$</td>
<td>0.25</td>
<td>.07</td>
<td>0.32</td>
<td>.02</td>
<td>0.32</td>
</tr>
<tr>
<td>$uV_d$ (cm/s)</td>
<td>0.43</td>
<td>.001</td>
<td>0.64</td>
<td>&lt;.0001</td>
<td>0.59</td>
</tr>
<tr>
<td>$uPI$</td>
<td>0.19</td>
<td>.16</td>
<td>0.24</td>
<td>.08</td>
<td>0.24</td>
</tr>
</tbody>
</table>

Note.—Data were analyzed by using the Pearson correlation coefficient.

### TABLE 4: Comparison of mean ± SD TCD parameters with binary CT variables

<table>
<thead>
<tr>
<th>Intraventricular Hemorrhage</th>
<th>Yes ($n = 28$)</th>
<th>No ($n = 23$)</th>
<th>$P$ Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>$aV_d$ (cm/s)</td>
<td>16.7 (10.7)</td>
<td>23.4 (10.6)</td>
<td>.03</td>
</tr>
<tr>
<td>$aPI$</td>
<td>2.12 (0.96)</td>
<td>1.53 (0.62)</td>
<td>.01</td>
</tr>
<tr>
<td>$uV_d$ (cm/s)</td>
<td>20.9 (15.2)</td>
<td>25.2 (11.2)</td>
<td>&gt;.05</td>
</tr>
<tr>
<td>$uPI$</td>
<td>1.94 (1.13)</td>
<td>1.24 (0.33)</td>
<td>.004</td>
</tr>
</tbody>
</table>

FIG 1. Scatterplot shows the correlation between hematoma volume, volume of hypoattenuation, and total volume with PI in the affected and unaffected hemispheres of the brain.
sure the ICP, realizing that TCD really measures the flow dynamics related to intracranial compliance. Finally, we acknowledge that the variance of most CT data explained by the TCD results is rather low.

Our results suggest that TCD indirectly assesses intracranial hypertension and mass effect. Except for ventricular size, all CT parameters indicating mass effect or intracranial hypertension were correlated to some degree with TCD measurements. Obviously, TCD is not a substitute for urgent scanning in patients in deteriorating condition, but it has the advantage of high temporal resolution, as well as noninvasiveness. It is relatively easy to perform at the bedside and at frequent intervals (or with continuous monitoring) without interfering with the treatment of the patient. However, because of the low variance explained by the TCD data and because we did not perform longitudinal studies, our results should be viewed as preliminary and without definitive implications regarding the treatment of patients with ICH. Further studies with serial examinations should evaluate whether TCD is useful to monitor ICP and CPP, to evaluate the effect of therapy (e.g., surgical evacuation [34] or osmotic therapy [31]), and to analyze its prognostic value (16) (as suggested by the high correlation found between PI and Glasgow Coma Scale and NIH scale scores). Increased PI could lead to placement of an ICP measurement device a patient who is clinically deteriorating. Studies that include simultaneous recording of ICP and TCD at frequent intervals and that correlate the findings with corresponding CT results are likely to provide important information.

Conclusion

In acute ICH, the TCD waveform is affected by many factors that are components of the CT study. TCD results do not replace information provided on CT, but TCD gives complementary data. Although our results are not definitive, they do provide a firm background for future studies. We hope that these findings will further the understanding of the relation between TCD and CT and ultimately translate to clinical practice.

Acknowledgment

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References


