Impaired cerebral autoregulation may be important in the pathogenesis of both stroke and global cerebral ischemia occurring during periods of hypoperfusion, such as intraoperatively. Risk factors for stroke, such as hypertension, are associated with altered cerebral autoregulation. An acute impairment of autoregulation is also found in a variety of disease states, including head injury, ischemic stroke, and vasospasm secondary to subarachnoid hemorrhage. A profound impairment of autoregulation is found in some patients with carotid artery stenosis or occlusion.

This may be associated with increased stroke risk and with increased stroke risk during hypotensive therapy or perioperatively. Conventionally static autoregulation has been measured. Cerebral blood flow itself, or an estimate such as cerebral blood flow velocity (CBFV), is measured during a large change in blood pressure that is usually induced pharmacologically. Such techniques are not suitable for many patients at risk of stroke because the blood pressure change induced could result in cerebral ischemia.

Therefore, most investigators have used an indirect measure of cerebral autoregulation such as the vasodilatory response to hypercapnia. Although this does correlate with cerebral autoregulation, it measures a slightly different physiological response, and the 2 may not always correlate.

More recently, noninvasive methods for measuring dynamic autoregulation have been proposed. These study the response of cerebral blood flow or CBFV to small changes in arterial blood pressure (ABP) and therefore are suitable for use in patients at risk of stroke. Aaslid et al suggested the use of bilateral leg cuffs inflated suprasystolically and then suddenly deflated to induce a transient fall in blood pressure, and they correlated the temporal pattern of the change in blood pressure with the change in middle cerebral artery (MCA) CBFV determined using transcranial Doppler. Such a technique may be more clinically relevant than static autoregulation in patients at risk of stroke because the transient blood pressure changes are likely to be more similar to those occurring in patients.

**Background and Purpose**—Assessment of cerebral autoregulation has been traditionally performed with static changes in arterial blood pressure. Newer dynamic methods require the induction of sudden drops in arterial blood pressure with the sudden release of bilateral thigh cuffs. An alternative method is proposed, based on the spontaneous variability of arterial blood pressure that does not require its manipulation. We compared this method with the established thigh cuff method in patients with carotid artery stenosis.

**Methods**—Cerebral blood flow velocity (determined by transcranial Doppler) and arterial blood pressure (determined by noninvasive servo-controlled plethysmograph) were recorded in 20 patients with carotid artery stenosis and 18 age-matched controls. At rest, grading of dynamic autoregulation was estimated from the impulse response of the blood pressure–velocity dynamic relationship. This was compared with the autoregulatory index (ARI) provided by the thigh cuff method and with the degree of stenosis. The critical closing pressure was derived from the fitted models and was also correlated with degree of stenosis.

**Results**—The 2 ARIs were significantly correlated ($r=0.76$) and reduced in subjects with carotid stenosis (baseline ARI, 3.65±3.11 versus 6.68±1.88, $P<0.0001$; thigh cuff ARI, 3.78±2.32 versus 6.35±1.06, $P<10^{-5}$). The critical closing pressure (relative to mean arterial blood pressure) was also significantly reduced ($−0.24±1.06$ versus $0.50±0.31$, $P<0.0001$) and correlated with the thigh cuff ARI ($r=0.68$). Both the baseline ARI and critical closing pressure were correlated with degree of stenosis ($P<10^{-6}$).

**Conclusions**—Grading of dynamic autoregulation with the use of undisturbed recordings of arterial blood pressure and cerebral blood flow velocity might provide a safer technique for assessment of patients in whom a sudden drop of arterial blood pressure is not desirable, such as patients with heart or autonomic failure. (Stroke. 1998;29:2341-2346.)

**Key Words:** autoregulation, blood pressure, carotid artery diseases, carotid endarterectomy, ultrasonography, Doppler, transcranial

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on a day-to-day basis than the large blood pressure changes induced during static autoregulatory testing. Tiecks et al have demonstrated an excellent agreement between this method and the classic assessment of static autoregulation.

Transcranial Doppler measurement of MCA CBFV is only a suitable technique if there is no change in MCA diameter during the change in blood pressure. Newell et al compared internal carotid artery absolute flow values during this step change in blood pressure with transcranial Doppler MCA CBFV changes and found an extremely close correlation, suggesting that transcranial Doppler is an appropriate technique to use in this setting. We have previously shown that this method of assessing dynamic autoregulation may identify a subgroup of patients with carotid stenosis who have significant impairments of cerebral autoregulation. However, in some patients the use of thigh cuffs to induce a fall in blood pressure is not ideal for a number of reasons. First, in ~20% of cases, it is not possible to induce a sufficient fall in blood pressure with this method. Second, inflation of thigh cuffs may be uncomfortable in some patients. Third, particularly in patients with critical cerebrovascular hemodynamics, such as premature newborns and individuals suffering from heart or autonomic failure, there may be a risk associated with the induction of drops in blood pressure. For these reasons alternative methods of providing an assessment of dynamic autoregulation in humans, which do not require induction of ABP disturbances, would be highly desirable.

One attractive possibility is to explore the spontaneous variability in ABP that is observed in most individuals at rest. The feasibility of this approach was demonstrated by Panerai et al in neonates using a coherent averaging method to improve the signal-to-noise ratio of the CBFV response to transient elevations in ABP. A more general approach is the induction of drops in blood pressure. For these reasons,

Grading of Cerebral Dynamic Autoregulation

Subjects and Methods

Twenty subjects with carotid stenosis >60% were studied. Grading of stenosis was based on ultrasound Doppler velocities in combination with B-mode imaging with a color-flow duplex ultrasound system (Acuson XP). Eighteen healthy age-matched nonsmoking volunteers with carotid stenosis excluded on duplex ultrasound were also studied as the control group.

Measurements were performed with subjects in a supine position and with their heads slightly elevated. CBFV was recorded bilaterally simultaneously through the transtemporal window with 2-MHz transducers (DWL, Langerach). The MCA was imaged at a mean±SD depth of 50.2±3.5 mm for the control population and 52.6±3.4 mm for the carotid stenosis group. Continuous ABP recording was made with a noninvasive servo-controlled plethysmograph (Finapres 2300, Ohmeda), with the subject’s hand maintained at the same level as the head. Baseline measurement of resting ABP was made by automated arm cuff (Omega 1400 series, In Vivo Laboratories Inc). A sudden stepwise drop in ABP was induced by the rapid release of bilateral thigh cuffs that had been inflated suprasystolically for 3 minutes. Drops in ABP of <10 mm Hg were not accepted for analysis. Continuous recordings of bilateral CBFV and ABP were stored on the transcranial Doppler machine for a baseline period of 2 minutes preceding the cuff release and for another 1 minute after the sudden deflation. Five cycles of inflation/deflation were performed per subject with a 3-minute rest interval between cycles. Grading of autoregulation based on the CBFV response to the stepwise drop in ABP was provided by a software program supplied by the transcranial Doppler manufacturers, as described previously. For each integral value of ARICuff, ranging from 0 to 9, the predicted CBFV response for the mathematical model proposed by Tiecks et al was compared with the actual CBFV tracing, and the ARICuff value corresponding to the curve with the least square error over a 30-second interval was selected as the best estimate. The critical closing pressure (CCP) was manually selected to improve model fitting. We calculated mean values of ARICuff and CRCP cuff for each subject using the individual estimates from each acceptable inflation/deflation cycle.

Data Analysis

Estimation of the ARI with the use of the IRF method was obtained from the baseline recordings preceding the thigh cuff deflation. The 120-second-long records were transferred to a personal computer at a rate of 200 samples per second for subsequent analysis. Each record was inspected visually for the presence of artifact or ectopic beats. Recordings with >4 ectopic beats were rejected. Narrow spikes in the CBFV signals were removed by linear interpolation. Time series of mean values of ABP and bilateral CBFV were obtained by low-pass filtering these signals with a cutoff frequency of 1 Hz (8th order zero-phase Butterworth) and decimating the sampling rate to 5 samples per second.

The IRF was estimated with a fast Fourier transform (FFT) algorithm. Before the direct FFT transform was computed, each signal was normalized by its mean value, and a cosine (Hanning) window was applied to the data. Two segments of data with 256 samples each were used to estimate the cross-spectra and the transfer function between the mean ABP (MABP) and CBFV signals with a frequency resolution of 0.0195 Hz. The amplitude spectra was smoothed with a 3-element triangular window, and the IRF was computed from the inverse FFT with a cutoff frequency of 0.5 Hz. Nyquist theorem states that this cutoff frequency is appropriate for signals that have been low-pass filtered at 1 Hz. The final IRF for each MCA was obtained as the average of all IRF available for each side, and it is termed IRFMCA. A numerical estimate of the baseline step response can be obtained by integration of the IRF. For technical reasons, however, grading of autoregulation based on baseline recordings was attempted by using the IRFMCA rather than the step response. The reasons for this choice will be given in the Discussion.

With the use of the model of Tiecks et al, for each step response a corresponding IRF (IRFmodel) was obtained by calculating the numerical derivative. To compare IRFMCA with IRFmodel it is necessary to take into account the parameter CrCP introduced by Tiecks et al in their original formulation. As discussed later, this parameter might not reflect the true critical closing pressure of the cerebral circulation. For simplicity, assume that autoregulation is impaired. In

\[
\text{ARI}_{\text{Cuff}} = \int_0^t \text{IRF}_{\text{base}}(t) \, dt
\]

\[
\text{ARI}_{\text{Cuff}} = \int_0^t \text{IRF}_{\text{model}}(t) \, dt
\]
this case, percent changes in ABP will induce velocity changes with unit gain, that is, $\Delta V_{\text{base}} = \Delta P$, where $\Delta V$ and $\Delta P$ are percent CBFV and ABP changes, respectively. The model of Tiecks et al., however, assumes that flow (or velocity) can become 0 for ABP values $0$ (i.e., when $ABP = CrCP$), as represented in Figure 4. In this case, percent changes in ABP will lead to percent changes in velocity, as shown in the following equation:

$$\Delta V_{\text{model}} = \frac{\Delta P}{1 - \frac{CrCP}{MABP}}$$

For an impaired autoregulation, the peak value of the IRF is given by the ratio $\Delta V/\Delta P$, and the relative amplitude ($A_{\text{rel}}$) between IRF model and IRFbase is then

$$A_{\text{rel}} = \frac{1}{1 - \frac{CrCP}{MABP}}$$

From Equation 2 it is possible to calculate the equivalent CrCP during baseline measurements, from the relative amplitude of the 2 impulse responses:

$$CrCP_{\text{base}} = MABP \left(1 - \frac{1}{A_{\text{rel}}}ight)$$

In summary, $A_{\text{rel}}$ was computed from the ratio of peak values of IRF model and IRFbase, leading to an estimate of the equivalent CrCPbase and also of the relative CrCPrel = CrCPbase/MABP. CrCPbase estimates were compared with values of CrCPcuff manually selected to improve model fitting for the thigh cuff data.

The temporal pattern of IRFbase was graded by identifying the best fit with 10 model curves, generated with the same set of parameters proposed by Tiecks et al., providing a value of $ARI_{\text{base}}$. Fractional values of $ARI_{\text{base}}$ were obtained by parabolic interpolation around the point with least square error.

Agreement between parameters derived from IRFbase and $ARI_{\text{cuff}}$ was assessed by the correlation coefficient and Bland-Altman plots. The Mann-Whitney $U$ test was used to test for differences in mean values. ANOVA with Scheffe’s test for post hoc analysis was used to test the relationship between model-derived parameters and the degree of stenosis. Linear regression analysis was performed to test for linear dependence between variables. A level of $P<0.05$ was considered significant.

Results

Of the 38 subjects, 1 patient and 1 control subject were rejected because of the high incidence of ectopic beats. As a result, 72 IRFbase were available for analysis. The mean±SD relative error for fitting the Tiecks model to IRFbase was $7.2\pm3.1\%$ of the IRFbase peak value. Figure 1 shows typical IRFbase from 1 subject and the corresponding IRF model best fits. The negative wave observed after $t=0$ in Figure 1A was present in all control subjects. Similarly, this negative wave was significantly reduced or absent in most patients with carotid artery disease (CAD) (Figure 1B).

Mean±SD values for the main variables studied are given in Table 1 for the control and patient groups, together with the $P$ values for the Mann-Whitney test. Highly significant differences were found between the 2 groups of individuals in relation to $ARI_{\text{cuff}}, ARI_{\text{base}}, CrCP_{\text{cuff}}, CrCP_{\text{rel}}$, and $CrCP_{\text{base}}$. The MABP was also significantly different for the 2 groups. For

### TABLE 1. Selected Variables in Control and Patient Groups

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control Group (n=34)</th>
<th>Patient Group (n=38)</th>
<th>$P$ (Mann-Whitney)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MABP, mm Hg</td>
<td>99.4±12.6</td>
<td>103.2±25.7</td>
<td>0.01</td>
</tr>
<tr>
<td>MCBFV, cm/s</td>
<td>51.1±13.1</td>
<td>57.1±15.0</td>
<td>0.06</td>
</tr>
<tr>
<td>$ARI_{\text{cuff}}$</td>
<td>6.35±1.06</td>
<td>3.78±2.32</td>
<td>$4\times10^{-6}$</td>
</tr>
<tr>
<td>$ARI_{\text{base}}$</td>
<td>6.68±1.88</td>
<td>3.65±3.11</td>
<td>0.0001</td>
</tr>
<tr>
<td>$CrCP_{\text{cuff}},$ mm Hg</td>
<td>62.9±9.36</td>
<td>36.8±27.7</td>
<td>$5\times10^{-6}$</td>
</tr>
<tr>
<td>$CrCP_{\text{rel}},$ relative units</td>
<td>0.50±0.31</td>
<td>−0.24±1.06</td>
<td>$5\times10^{-5}$</td>
</tr>
<tr>
<td>$CrCP_{\text{base}},$ mm Hg</td>
<td>46.7±35.5</td>
<td>−15.2±94.7</td>
<td>0.0003</td>
</tr>
</tbody>
</table>

Values are mean±SD.

![Figure 1. Representative IRF from a 44-year-old male CAD patient with no stenosis in the right carotid artery and 100% stenosis in the left carotid artery. A, IRF from the right MCA (continuous line) and best-fit model curve (dotted line) obtained with the model of Tiecks et al. The relative and absolute values of $CrCP$ are 0.46 and 47.6 mm Hg, respectively. The least square error is 6.4% of the IRF peak value. B, IRF from the left MCA. The corresponding relative and absolute values of $CrCP$ are 0.27 and 27.8 mm Hg, respectively. The least square error is 6.0% of the IRF peak value.](http://stroke.ahajournals.org/)

![Figure 2. ARI obtained from baseline recordings ($ARI_{\text{base}}$) vs corresponding values obtained from the thigh cuff test ($ARI_{\text{cuff}}$).](http://stroke.ahajournals.org/)
the control group, $ARI_{cuff}$ had a smaller coefficient of variation (16.7%) than $ARI_{base}$ (28.1%), but the difference was not significant.

Highly significant correlation coefficient values were obtained between the $ARI_{cuff}$ and the model-derived parameters $ARI_{base}$ ($r=0.764$), $CrCP_{rel}$ ($r=0.68$), $CrCP_{base}$ ($r=0.70$), and $A_{rel}$ ($r=0.69$).

Figure 2 presents a scatter diagram between $ARI_{base}$ and $ARI_{cuff}$. A linear regression between these 2 parameters had a highly significant slope ($P<0.0001$) and residuals that were normally distributed. Figure 2 suggests the presence of 2 distinct groups of arteries. When we performed a linear regression using only data from control subjects, the correlation coefficient was reduced to $r=0.406$, but the regression slope was still significant ($P<0.017$). Bland-Altman analysis of the agreement between $ARI_{base}$ and $ARI_{cuff}$ indicated a bias value of $-0.09$ and limits of agreement of $-3.89$ and $3.71$. For the control group, $CrCP_{rel}$ ($r=0.484$, $P=0.004$) and $CrCP_{base}$ ($r=0.45$, $P=0.007$) were also significantly correlated with $ARI_{cuff}$.

Patients were split into 4 subgroups according to the classification of stenosis with the following number of vessels in each subgroup: <60% stenosis (9 vessels), 60% to 79% (8), 80% to 99% (7), and 100% (12). One subject with a carotid artery bypass was not included. The mean ± SEM of $ARI_{base}$ and $CrCP_{base}$ are represented in Figure 3 for these 4 patient subgroups and for the control group, which contained 34 vessels.

ANOVA of the 5 patient subgroups represented in Figure 3 yielded very significant results for both $ARI_{base}$ and $CrCP_{rel}$ ($P<10^{-6}$). Scheffe’s test also indicated that these parameters can distinguish between some of these subgroups, with the $P$ values given in Table 2. Significant differences between the 5 subgroups were also obtained for the ANOVA of $CrCP_{base}$ ($P=1.5\times10^{-3}$) and $CrCP_{cuff}$ ($P=7\times10^{-6}$), but Scheffe’s test indicated inferior discrimination between individual subgroups compared with the results obtained for $CrCP_{base}$ (Table 2). On the other hand, ANOVA results were nonsignificant for mean CBFV (MCBFV), MABP, and fitting least square error for the different degrees of stenosis considered.

**Discussion**

The possibility of assessing the status of cerebral pressure autoregulation with the spontaneous variability of ABP as the stimulus to perturb CBFV has been demonstrated by previous studies, but hitherto it has been restricted to the dichotomous classification of normal/impaired autoregulation. In the present investigation we have demonstrated that the model proposed by Tiecks et al, to describe the CBFV response to ABP drops induced by the sudden deflation of thigh cuffs, can also be used to grade autoregulation using baseline recordings of ABP and CBFV. The $ARI$ obtained during undisturbed conditions ($ARI_{base}$) has shown a very high correlation with $ARI_{cuff}$ and considerable sensitivity to the degree of stenosis in a population of CAD patients (Figure 3). In a previous study, Zhang et al have also shown that baseline estimates of IRF can be used to predict CBFV changes after the sudden release of thigh cuffs in normal subjects, thus suggesting that the same IRF might apply to both situations.

Despite these initially encouraging results, several limitations of the methods adopted need to be kept in perspective. One major limitation of using baseline recordings to grade autoregulatory performance is the lack of ABP variability at the spectral frequency bands, which can stimulate an autoregulatory response. Although the pressure drop produced by the deflation of thigh cuffs is not a perfect “step function” and returns to its original value 10 to 15 seconds after the sudden release of the cuffs, it provides much more low-frequency power (<0.2 Hz) than normally available in a 120-second recording of undisturbed fluctuations in ABP. The amount of signal power available in the pertinent frequency band is important to overcome the noise levels, thus providing reliable estimates of the input-output relationship. This deficiency in low-frequency power becomes...
more of a problem if one is estimating the step response because of its higher relative content of low-frequency power compared with the IRF. For this reason, the analysis and fitting of the Tiecks et al\textsuperscript{6} model were performed on the IRF. Although the IRF provides a less intuitive “feeling” of the autoregulatory response corresponding to different values of ARI, its temporal pattern can also reflect grading of autoregulation. At \( t=0 \), when a hypothetical impulse-like disturbance in ABP takes place, there is an immediate direct response in CBFV. If the IRF remains flat, this means a lack of feedback reaction in CBFV that will then tend to follow ABP changes. This situation is characteristic of an absence of autoregulation. On the other extreme, if the positive immediate change in CBFV is counteracted by a negative wave in the IRF (Figure 1A), this will induce a return of CBFV to its original level, representing the case of a perfect autoregulation. Although the occurrence of this temporal pattern was in excellent agreement with the reference values of ARI\textsubscript{crit}, the limits of agreement between this index and ARI\textsubscript{base} (\( \sim 3.89 \) to 3.71) should be considered inadequate for clinical applications of the baseline method at this stage.

A number of different reasons might contribute to explain this poor agreement. First, it is appropriate to question the accuracy and precision of ARI\textsubscript{crit} as a “gold standard” for assessment of dynamic autoregulation. More studies on the reproducibility of the thigh cuff technique are undoubtedly needed, but initial results suggest a possible problem in this area.\textsuperscript{18} Second, the transcranial Doppler equipment allows the operator to select values of CrCP\textsubscript{crit} to improve the fitting between model and data, but different values of ARI\textsubscript{crit} could have been obtained if an objective and automatic procedure was adopted for this purpose, such as we have used to fit the model during baseline recordings. Third, the model of Tiecks et al\textsuperscript{6} is an obvious first choice to fit and grade the IRF\textsubscript{base}, but we have not explored other alternatives that could lead to better fitting and reduced least square errors. Future research on this topic is important to arrive at more accurate and sensitive mathematical models to describe the dynamic pressure-velocity relationship during baseline recordings. Fourth, and related to this previous point, is the problem of performing the FFT analysis on segments of data that are long enough to yield accurate estimates of IRF and to maximize the low-frequency power spectral content. IRF estimates with higher signal-to-noise ratios than hitherto available are necessary to allow further investigation of alternative mathematical models to fit and grade the IRF\textsubscript{base}. To summarize this point of the discussion, it is important that more work is performed to understand the sources of variability of both ARI\textsubscript{crit} and ARI\textsubscript{base}, before either is prematurely discarded or uncritically accepted.

Newell et al\textsuperscript{5} have shown that CBFV changes after the sudden deflation of thigh cuffs provide an accurate estimate of the corresponding changes in cerebral blood flow. Their results suggest that the diameter of the MCA does not change significantly during the thigh cuff test, but it is not possible to assume that the same holds true for the spontaneous variability of CBFV recorded during baseline measurements. If the diameter of the MCA remains constant, the observed variability of CBFV is a true reflection of spontaneous or pressure-induced fluctuations in flow. In this case, if autoregulation is intact, the negative transient of the IRF is a reflection of adjustments in small-vessel resistance, possibly involving a metabolic mechanism.\textsuperscript{2,13,14} On the other hand, as observed by Zhang et al\textsuperscript{15} if flow is constant during baseline and the velocity fluctuations are due to small changes in MCA diameter, then the IRF might be reflecting a myogenic mechanism involving the direct action of ABP variability on large-vessel diameter. Methods based on the power of the reflected Doppler signal might be able to shed light on this problem in the near future.\textsuperscript{5,19}

The CrCP parameter, as introduced by Tiecks et al\textsuperscript{6} is important to explain the finding that IRFs can have markedly different amplitudes. As shown by Figure 4, a sudden change in ABP (\( \Delta P \)) can produce different \( \Delta V \) transients and therefore distinct peak values of the IRF, depending on the CrCP parameter. The peak value of the IRF, represented at \( t=0 \) in Figure 1, reflects the immediate perturbation of CBFV to a sudden change in ABP, before the few seconds required for the autoregulatory response to be manifested.\textsuperscript{2,5,13,14}

The fact that CrCP values required to fit the Tiecks model to IRF\textsubscript{base} are spread over a relatively wide range and show extremely high correlation with the degree of stenosis (Figure 3) was not expected at the outset of our study. From the work of Burton,\textsuperscript{26} the concept of CrCP has been formulated as the point at which reductions in perfusion pressure lead to a stagnation in blood flow or velocity. For a number of reasons, however, the use of this concept in the model of Tiecks et al\textsuperscript{6} can be misleading, and it is not possible to assume that the resulting values reflect the true critical closing pressure of the cerebral circulation. Values of CrCP \( >0 \) have been obtained in animals\textsuperscript{21} and humans\textsuperscript{10,22} by using the instantaneous pressure-velocity relationship for a complete cardiac cycle.\textsuperscript{10} Unfortunately, in humans it is not yet possible to measure the ABP of large cerebral vessels, and this raises questions about the accuracy of CrCP estimates based on ABP measurements in the aorta or radial arteries. This problem becomes worse when the Finapres is used to record the ABP waveform. Note that the CrCP parameter in the formulation of Tiecks et al\textsuperscript{6} bears no relationship to estimates based on instantaneous pressure-velocity relationships, because the signals in Equation 1 represent mean values of CBFV and ABP for each cardiac cycle. In addition, this parameter has not been estimated by the extrapolation of the linear regression of velocity against ABP, as adopted by other studies.\textsuperscript{10,21,22}

Despite these differences and the risk of misleading interpre-
Grading of Cerebral Dynamic Autoregulation

...tions, pressure-velocity relationships, as depicted in Figure 4, can still provide a useful model to interpret the role of the CrCP parameter, as introduced by Tiecks et al.6 particularly regarding the unusual finding of large negative values of CrCP (Figure 3). For a step or impulselike change in ABP of amplitude \( \Delta P \), in the case of CrCP, \( \Delta P > 0 \), the relative change will result in \( (\Delta V_{A} /\text{MCBFV}) = (\Delta P /\text{MABP}) \), as observed in control subjects and CAD patients with stenosis \( < 60\% \) (Figure 3). On the other hand, for CrCP, \( \Delta P < 0 \), the same change \( \Delta P \) will lead to much smaller relative changes in velocity and will result in \( (\Delta V_{A} /\text{MCBFV}) = (\Delta P /\text{MABP}) \), as exemplified by the patients with \( > 80\% \) stenosis. The most likely explanation for the latter is that negative CrCP values are an illusion resulting from significant degrees of stenosis causing downstream falls in perfusion pressure. As a consequence, the “true” perfusion pressure change at the MCA will be \( \Delta P' \ll \Delta P \) (Figure 4), and the relative changes in velocity can then be compatible with positive, albeit small, values of CrCP. On the other hand, the reduction in CrCP with diminishing autoregulatory capacity seems to be a real phenomenon since it is also observed in the control group.

Further discussion about the physiology of elevated CrCP, as observed in some control subjects, is beyond the scope of this report. Nevertheless, the fact that CrCP is correlated with ARIbase and the degree of stenosis and the large negative values encountered in some patients deserve further investigation.

Grading of IRFbase can be obtained with empirical parameters, such as the area of the negative wave in the interval 0 to 5 seconds, but this could lead to confounding of the differential effects of cerebral autoregulation and CAD on the IRF. By separating changes in amplitude (ie, CrCP) from the morphology of the IRF (ie, ARIbase), it was possible to confirm the hypothesis initially explored by White and Markus3 that CAD has a significant depressing effect on cerebral autoregulation. The combined use of the 2 parameters (ARIbase and CrCP) seems to allow a better noninvasive discrimination of degree of stenosis (Table 1) than obtained in their original study with ARIbase.3

Although further studies are necessary, the initial results described above suggest that system analysis of spontaneous fluctuations in ABP might be useful as a convenient and trouble-free test of dynamic autoregulation. The relative simplicity of this approach might facilitate more comprehensive studies of several patient subgroups in which cerebral autoregulation is disturbed, particularly for critically ill patients in whom inducing a blood pressure reduction might be inadequate. Such patient groups might include acute cerebral ischemia, head injury, and neonates.

Acknowledgments

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