



Discover Generics

Cost-Effective CT & MRI Contrast Agents



[VIEW CATALOG](#)

AJNR

Alberta Stroke Program Early CT Scoring of CT Perfusion in Early Stroke Visualization and Assessment

R.I. Aviv, J. Mandelcorn, S. Chakraborty, D. Gladstone, S. Malham, G. Tomlinson, A.J. Fox and S. Symons

This information is current as of September 2, 2025.

AJNR Am J Neuroradiol 2007, 28 (10) 1975-1980

doi: <https://doi.org/10.3174/ajnr.A0689>

<http://www.ajnr.org/content/28/10/1975>

ORIGINAL
RESEARCH

R.I. Aviv
J. Mandelcorn
S. Chakraborty
D. Gladstone
S. Malham
G. Tomlinson
A.J. Fox
S. Symons

Alberta Stroke Program Early CT Scoring of CT Perfusion in Early Stroke Visualization and Assessment

BACKGROUND AND PURPOSE: Qualitative CT perfusion (CTP) assessment by using the Alberta Stroke Program Early CT Score (ASPECTS) allows rapid calculation of infarct extent for middle cerebral artery infarcts. Published thresholds exist for noncontrast CT (NCCT) ASPECTS, which may distinguish outcome/complication risk, but early ischemic signs are difficult to detect. We hypothesized that different ASPECTS thresholds exist for CTP parameters versus NCCT and that these may be superior at predicting clinical and radiologic outcome in the acute setting.

MATERIALS AND METHODS: Thirty-six baseline acute stroke NCCT and CTP studies within 3 hours of symptoms were blindly reviewed by 3 neuroradiologists, and ASPECTS were assigned. Treatment response was defined as major neurologic improvement when a ≥ 8 -point National Institutes of Health Stroke Scale improvement at 24 hours occurred. Follow-up NCCT ASPECTS and 90-day modified Rankin score (mRS) were radiologic and clinical reference standards. Receiver operating characteristic curves derived optimal thresholds for outcome.

RESULTS: Cerebral blood volume and NCCT ASPECTS had similar radiologic correlations (0.6 and 0.5, respectively) and best predicted infarct size in the absence of major neurologic improvement. A NCCT ASPECT threshold of 7 and a cerebral blood volume threshold of 8 discriminated patients with poor follow-up scans ($P < .0002$ and $P = .0001$) and mRS ≤ 2 ($P = .001$ and $P < .001$). Only cerebral blood volume predicted major neurologic improvement ($P = .02$). Interobserver agreement was substantial (intraclass correlation coefficient, 0.69). Cerebral blood volume ASPECTS sensitivity, specificity, positive predictive value, and negative predictive value for clinical outcome were 60%, 100%, 100%, and 45%, respectively. No patients with cerebral blood volume ASPECTS < 8 achieved good clinical outcome.

CONCLUSION: Cerebral blood volume ASPECTS is equivalent to NCCT for predicting radiologic outcome but may have an additional benefit in predicting patients with major neurologic improvement.

The most widely used imaging technique for acute stroke remains noncontrast CT (NCCT). The reliability of early subtle ischemic sign detection in middle cerebral artery territory is improved with experience,¹ clinical history,² stroke window width and level,³ and the use of the Alberta Stroke Program Early CT Score (ASPECTS).⁴ NCCT ASPECTS allows a rapid and reliable assessment of stroke extent and has been shown to correlate with final infarct size.⁵ The combination of the ASPECTS, which is weighted for clinically significant regions, and CT perfusion (CTP), which can distinguish salvageable tissue from infarct core, suggests added benefit over NCCT alone for clinical outcome. Although multiple factors influence stroke outcome, knowledge of cerebral blood volume, cerebral blood flow, and mean-transit-time thresholds for outcome may influence the decision for thrombolysis treatment.

The aim of the study was to evaluate whether in the acute CT assessment of infarction, ASPECTS scoring of CTP has higher predictive value than NCCT for the final infarct size

and clinical outcome. We also sought to determine and test thresholds predictive of outcome in patients.

Materials and Methods

This was a retrospective study of consecutive tissue plasminogen activator (tPA)-treated patients presenting between January and June 2005. Patients were included if they presented with suspected acute stroke within 3 hours of onset, underwent a CT stroke protocol (including CT angiography [CTA] and CTP), were treated with intravenous thrombolysis, and were followed up at 5–7 days. A standard tPA administration regime was used (0.9 mg/kg dose, 10% bolus, 90% continuous infusion for 1 hour). Recanalization data were variably collected because of differences in practice of clinicians and did not exclude the patient from participation. To determine prediction of presentation perfusion parameters with radiologic outcome, we dichotomized patients into 2 groups: those with a major neurologic improvement and those without. Major neurologic improvement was defined in accordance with previous publications, as ≥ 8 -point National Institutes of Health Stroke Scale (NIHSS) improvement at 24 hours after tPA.^{6–8} Major neurologic improvement was used as a surrogate marker of response to treatment on the basis of the following rationale: Secondary analysis by the National Institute of Neurological Disorders and Stroke investigators demonstrated that this magnitude of improvement discriminated between the treatment and placebo group without overlap in the confidence intervals (CIs).⁶ This degree of improvement is an independent predictor of good 90-day outcome.⁷ Major neurologic improvement has, therefore, been interpreted in this study and others as a marker of treatment efficacy and has been suggested as a surrogate marker for clinical efficacy and a

Received November 13, 2006; accepted after revision April 2, 2007.

From the Division of Neuroradiology (R.I.A., J.M., S.C., S.M., A.J.F., S.S.), Neurology and Regional Stroke Center (D.G.), Sunnybrook Health Sciences Centre, Toronto, Ontario, Canada; and Department of Biostatistics (G.T.), University of Toronto, Toronto, Ontario, Canada.

Please address correspondence to: R. Aviv, MD, Division of Neuroradiology, Sunnybrook Health Sciences Centre, 2075 Bayview Ave, Toronto M4N 3M5, Ontario, Canada; e-mail: richardaviv@lincene.net

DOI 10.3174/ajnr.A0689

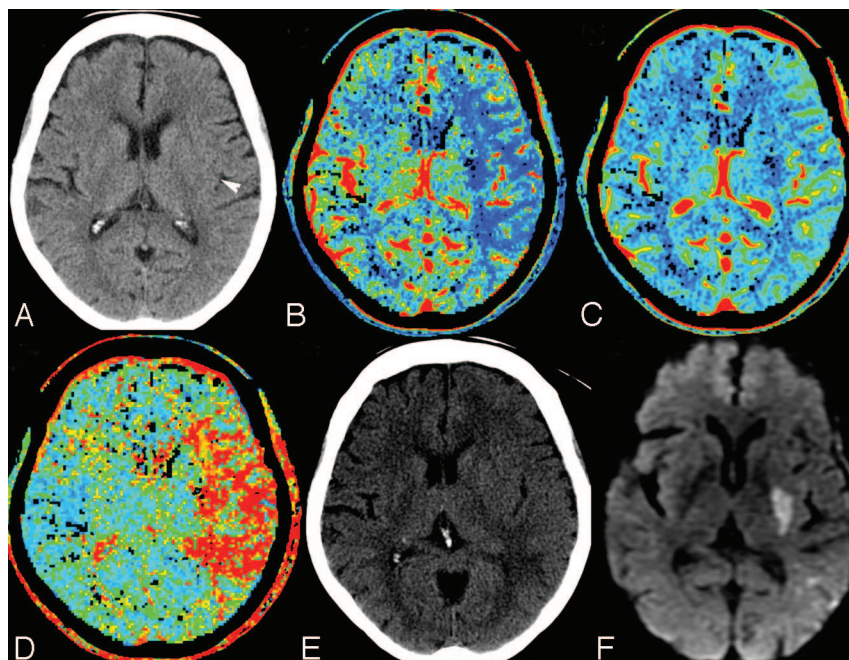


Fig 1. A 75-year-old man, within 2 hours of right-sided stroke and presentation of NIHSS 20. A, NCCT demonstrates subtle loss of the left posterior putamen, internal capsule, and posterior insular cortex (white arrowhead) (ASPECTS 7). B, Cerebral blood flow. C, Cerebral blood volume. D, Mean transit time. Cerebral blood volume demonstrates an abnormality confined to the posterior putamen and internal capsule (ASPECTS 8), with larger cerebral blood flow and mean-transit-time abnormalities corresponding to the left middle cerebral artery M1 segment occlusion (not shown). E, Follow-up NCCT at day 6 shows an indistinct posterior putamen confirmed on diffusion-weighted MR imaging (F). The patient recovered by 18 points with a final NIHSS of 2.

Image Analysis

Three reviewers (2 neuroradiologists and a neuroradiology fellow) read a number of mock cases and agreed by consensus on the method of ASPECTS scoring on CTP, according to a previously published method for NCCT.⁴ A qualitative approach to CTP was adopted, scoring any perfusion abnormality. Absolute perfusion values were not considered. By agreement, only images from the anterior commissure to the section above the lateral

ventricles were reviewed on NCCT. Readers were blinded to clinical information and randomly reviewed the studies on a PACS workstation (Version 3.1; Agfa, Waterloo, Ontario, Canada). Perfusion abnormality was scored by using ASPECTS for each of the cerebral blood volume, cerebral blood flow, and mean-transit-time maps; and the affected side was recorded. A delay of 1 week between each sequence read was instituted to avoid recall bias. The final NCCT ASPECTS was scored separately by 2 experienced readers without knowledge of previous imaging.

Imaging

The CT stroke protocol using a 4-section CT scanner (LightSpeed; GE Healthcare, Milwaukee, Wis) comprised pre- and postcontrast head scans from the skull base to the vertex. Imaging parameters were the following: 120 kVp, 340 mA, 4 × 5 mm collimation, 1 s/rotation, and table speed of 15 mm/rotation. CTP parameters were the following: 45-second cine scanning (80 kVp, 190 mA), 3- to 5-second delay, injection of 0.5-mL/kg (30–50 mL) iohexol (300 mg/mL; Omnipaque, Nycomed, Princeton, NJ) at a rate of 4 mL/s. CTP studies covered a 20-mm slab with four 5-mm sections carefully positioned to allow visualization of the sections required for ASPECTS scoring from the basal ganglia to the centrum. This was achievable in all patients (Fig 1). CTA covered carotid bifurcations to the vertex with the following parameters: 0.7-mL/kg iohexol (maximum, 90 mL); 5- to 10-second delay; 120 kVp; 270 mA; 1 s/rotation; 1.25-mm-thick sections; table speed, 3.75 mm/rotation. For each CTP study, a time attenuation curve was made, displaying the change in Hounsfield units for a specified region for the duration of the scanning. Arterial and venous input functions were obtained from the ipsilateral anterior cerebral artery and from the superior sagittal sinus, respectively. CT Perfusion 3 (GE Healthcare) was used to calculate parametric maps of cerebral blood flow and cerebral blood volume by deconvolution of tissue-enhancement curves and arterial time-attenuation curve in 2 × 2 pixel blocks. Mean-transit-time maps were also calculated. Partial volume averaging of the arterial input curve was corrected by using the venous time-attenuation curve.

ventricles were reviewed on NCCT. Readers were blinded to clinical information and randomly reviewed the studies on a PACS workstation (Version 3.1; Agfa, Waterloo, Ontario, Canada). Perfusion abnormality was scored by using ASPECTS for each of the cerebral blood volume, cerebral blood flow, and mean-transit-time maps; and the affected side was recorded. A delay of 1 week between each sequence read was instituted to avoid recall bias. The final NCCT ASPECTS was scored separately by 2 experienced readers without knowledge of previous imaging.

Statistical Analysis

Data were transferred to and analyzed in Statistical Package for the Social Sciences (Version 10; SPSS, Chicago, Ill). Median ASPECTSs for the presentation and follow-up NCCT and CTP parameters were compared by using paired *t* tests, dichotomizing patients according to recanalization implied by NIHSS improvement as discussed previously. Pearson correlation coefficients were computed between ASPECTS for NCCT, cerebral blood volume, cerebral blood flow, mean transit time, and radiologic NCCT outcome. Median ASPECTSs for each CT parameter were compared for patients having good and bad clinical outcome (mRS, ≤2 or ≥3, respectively) by using unpaired *t* tests. For each sequence and follow-up NCCT, receiver operating characteristic (ROC) curves were computed with the dichotomized mRS as the reference outcome; the optimal threshold for predicting outcome was defined as that which gave maximum accuracy (sensitivity plus specificity). Using these optimal thresholds in our patient group, we calculated rate ratios and reported the optimal positive and negative predictive values for identifying good clinical outcomes. To assess agreement among readers for ASPECTS as a continuous variable, we computed the intraclass correlation coefficient (ICC). For all analyses, statistical significance was determined as *P* < .05.

Results

There were 18 female and 18 male patients. The median age was 75 years (range, 35–95 years). Median NIHSS was 15

Table 1: Mean ASPECTS values and 95% CIs for differences in ASPECTS at baseline for NCCT and CTP compared with final NCCT ASPECTS

	NC-CT	CBV	CBF	MTT	Final NC-CT ASPECTS
MNI (<i>n</i> = 7) ^a					
Mean	8.5	9	5.166667	5	9
NCCT		−2.3–1.4 ^{NS}	0.06–3.8 ^S	0.11–3.88 ^S	−2.1–1.7 ^{NS}
CBV			0.5–4.3 ^S	0.5–4.4 ^S	−1.6–2.2 ^{NS}
CBF				−1.8–1.9 ^{NS}	−4.0 to −0.3 ^S
MTT					−4.0 to −0.3 ^S
No MNI (<i>n</i> = 29)					
Mean	6.8	6.5	4.0	4.0	5.4
NCCT		−0.4–1.1 ^{NS}	1.7–4.0 ^{S**}	S 1.7–4.0 ^{S**}	0.6–2.2 ^S
CBV			1.3–3.6 ^{S**}	S 1.4–3.6 ^{S**}	0.07–2.3 ^S
CBF				−1.3–1.4 ^{NS}	−2.6–0.05 ^{NS}
MTT					−2.7–0.01 ^{NS}

Note:—MNI indicates major neurologic improvement; CBV, cerebral blood volume; CBF, cerebral blood flow; MTT, mean transit time; NS, nonsignificant; S, significant (level of significance determined by paired *t* test, *P* < .05, or *S***, *P* < .01).

^a In patients with MNI, mean CBV and NCCT were not significantly different from mean final NCCT ASPECTS, whereas in the absence of MNI, CBF, and MTT most closely approximated final NCCT ASPECTS.

Table 2: Performance measures of NCCT, CTP, final ASPECTS, and 24-hour NIHSS change for clinical outcome

Threshold ^a	Good Outcome, mRS ≤2 (% Patients)	RR (95% CI)	<i>P</i> *	Sensitivity (95% CI)	Specificity (95% CI)	Accuracy
NCCT >7 vs ≤7	44 (6)	8.0 (1.1–57.6)	.018	0.63 (0.4–0.8)	0.89 (0.47–0.99)	0.69
CBV ≥8 vs <8	45 (0)	15.4 (0.96–246)	.002	0.60 (0.4–0.8)	100 (0.66–100)	0.69
Final NCCT >6 vs ≤6	47 (0)	17 (1.1–27.3)	.001	1.00 (0.7–1.0)	0.63 (0.44–0.78)	0.72
CBF/MTT >5 vs ≤5	46 (13)	3.54 (1.1–11.8)	.046	0.74 (0.5–0.9)	0.67 (0.35–0.88)	0.69
Change NIHSS 24 hours ≥8	86 (10)	8 (2.7–25.2)	.0001	0.96 (0.8–1.0)	0.67 (0.3–0.9)	0.89

Note:—RR indicates relative risk; CBV, cerebral blood volume; CBF, cerebral blood flow; MTT, mean transit time.

^a Dichotomized thresholds were determined by ROC analysis. Percentage of patients presenting with good outcome above and below the selected threshold is illustrated. The best early predictors of clinical outcome were 24-hour NIHSS and presentation CBV.

* *P* value was calculated using Fisher exact test.

(range, 6–30) at presentation. The median time to presentation from symptom onset was 165 minutes (range, 84–180 minutes). Median NIHSS 24-hours post-tPA was 4 (range, 0–46). Median hospital length of stay was 11 days (range, 1–110). The location of thrombus demonstrated on presentation CTA was the M1 segment in 16 patients and the M1–4 segments in 20 patients. No inferior cerebellar artery, posterior inferior cerebellar artery, or anterior cerebellar artery infarcts were present. Left-sided infarcts occurred in 16 patients (44%). Hemorrhagic transformation (type 1 or 2) and parenchymal hematoma (PH 1 or 2) were seen in 8 (22%) and 6 (16%) patients, respectively. Major neurologic improvement was present in 7 patients.

Radiologic Outcome

The median ASPECTS and level of significance for baseline NCCT, final NCCT ASPECTS, and CTP are given in Table 1. The mean cerebral blood volume and NCCT ASPECTS at presentation most closely predicted final NCCT ASPECTS in patients with major neurologic improvement. In the absence of such improvement, cerebral blood flow and mean-transit-time values most closely agreed with final NCCT ASPECTS.

A correlation matrix was determined for each CTP parameter predicting final NCCT ASPECTS. NCCT and cerebral blood volume ASPECTS (*r* = 0.6 and 0.5, respectively) were superior to mean transit time or cerebral blood flow (*r* = 0.4 each) in patients who showed major neurologic improvement. In the absence of such NIHSS improvement, cerebral blood flow and mean-transit-time ASPECTS (*r* = 0.6 each) were superior to cerebral blood volume and NCCT (*r* = 0.5).

An ASPECT threshold of 7 and cerebral blood volume

threshold of 8 at presentation discriminated patients with poor follow-up findings on scans (*P* < .0002 and *P* = .0001, respectively). In patients with no major neurologic improvement, cerebral blood flow and mean-transit-time thresholds of 5 were predictive of radiologic outcome (*P* < .0001 each).

Clinical Outcome

ROC curves were drawn, and the areas calculated as 0.8 each for cerebral blood volume, cerebral blood flow, mean transit time, and final NCCT ASPECTS; and 0.7 for baseline CT, respectively. Optimal thresholds were determined and are presented in Table 2.

A cerebral blood volume threshold of 8 was the only presentation parameter to predict patients with major neurologic improvement (*P* = .02) (Fig 2). NCCT (*P* = .08), cerebral blood flow (*P* = .22), and mean transit time (*P* = .20) did not reach significance. The mean NIHSS at 24 hours in patients presenting with cerebral blood volume ≥8 was 7.3 (SD, ±6.7) versus 16.4 (SD, ±6) in patients with cerebral blood volume <8 (*P* = .0004). Major neurologic improvement predicted a good clinical outcome in 86% of patients with sensitivity, specificity, positive predictive value, and negative predictive value of 96%, 67%, 89%, and 86%, respectively.

Figure 3 demonstrates that the greatest discriminator of clinical outcome was the follow-up CT ASPECTS (*P* < .001), followed by cerebral blood volume (*P* < .001). Sensitivity, specificity, positive predictive value, and negative predictive value for cerebral blood volume ASPECTS was 60%, 100%, 100%, and 45%, respectively. Values for initial NCCT ASPECTS were 63%, 89%, 94%, and 44%, respectively. No patients with a cerebral blood volume threshold of

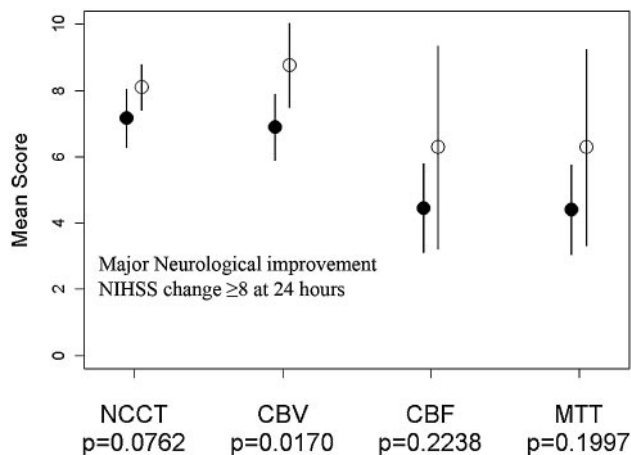


Fig 2. A plot of mean baseline ASPECTS NCCT, cerebral blood volume (CBV), cerebral blood flow (CBF), and mean transit time (MTT) against major neurologic improvement, demonstrating that only CBV is predictive of 24-hour NIHSS change.

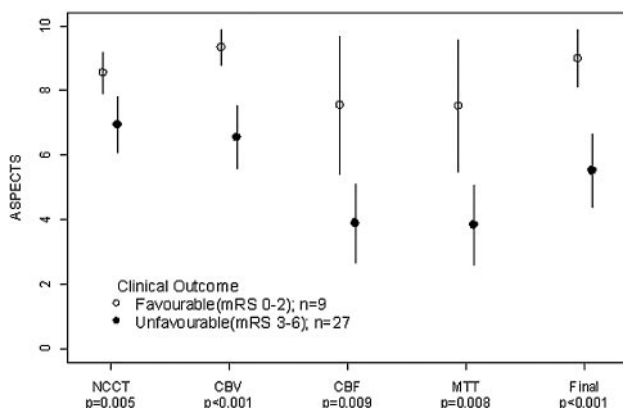


Fig 3. A plot of mean baseline ASPECTS NCCT, cerebral blood volume (CBV), cerebral blood flow (CBF), and mean transit time (MTT). Results are dichotomized for clinical outcome by using 2-sample *t* tests. Favorable outcome was defined as mRS ≤ 2 . Comparison of mean clinical outcomes was based on dichotomized mRS (0–2 versus 3–6), by using unequal variance 2-sample *t* tests.

< 8 had an mRS ≤ 2 at follow-up. However, the prognosis for patients with cerebral blood volume ≥ 8 was more variable, with only 45% achieving an mRS ≤ 2 . However, 70% achieved an mRS ≤ 3 with only 1 false-negative (6%) (Table 3). There was no difference between mean transit time and cerebral blood flow for clinical outcome ($P = .008$ and $.009$). The sensitivity and specificity were 67% and 70%, respectively.

Interobserver Agreement

The ICC for each CT parameter is given in Table 4. Cerebral blood volume agreement was slightly higher than NCCT agreement, but both demonstrated comparable CIs. The ICC for cerebral blood flow and mean transit time was high, with narrow CIs.

Discussion

We demonstrated that a cerebral blood volume ASPECTS threshold of 8 discriminates patients who experience major neurologic improvement and have good clinical outcome at 3 months (mRS ≤ 2). The ability of a cerebral blood volume ASPECTS threshold at presentation to predict major neurologic improvement has not previously been described, to our

knowledge, and has important clinical implications. Major neurologic improvement is defined as NIHSS improvement ≥ 8 points. It is usually determined by subtracting the post-tPA treatment NIHSS from the presentation score. Major neurologic improvement is considered a marker of treatment response and thought to reflect recanalization.^{8,9} Major neurologic improvement and 24-hour NIHSS are known to strongly discriminate 90-day outcome with an area under the curve for 24-hour NIHSS of 0.71.⁸⁻¹⁰ In our study, baseline cerebral blood volume ASPECTS was the only parameter to discriminate patients who showed major neurologic improvement. This improvement was also associated with the highest accuracy for prediction of 3-month clinical outcome. Our findings suggest that ASPECTS scoring of cerebral blood volume may provide additional clinical prognostic information not available on NCCT.

In this series, patients with a cerebral blood volume ASPECTS < 8 treated with thrombolysis did not have a good clinical outcome. The prognosis is more guarded when the cerebral blood volume is ≥ 8 , with an outcome of mRS ≤ 2 in only 45% of patients, and is consistent with a recent report of ASPECTS in the Canadian Alteplase for Stroke Effectiveness Study dataset.¹⁰ The rate of patients' achieving mRS ≤ 3 was 70%, also consistent with the previously mentioned study with 1 false-negative (6%).

CTP has a spatial resolution ($1.5 \times 1.5 \times 3$ cm) similar to that of MR perfusion but is limited by spatial coverage, ionizing radiation, and iodinated contrast injection. Despite limited 2-cm coverage on a 4-section scanner, a carefully positioned slab ensures that the ASPECTSs were highly correlated with final infarct volume and clinical outcome. A recent study has evaluated the dose obtained at CTP. With commonly used parameters, the dose of 1.1–1.3 mSv is lower than that in NCCT. The dose was recorded for a 64-section scanner over 2.4 cm, which is greater than the 2 cm achievable on a 4-section scanner.¹¹ In clinical practice, we do not repeat the CTP study if movement occurs. It is often possible for the CTP raw data to be edited and the sections with movement to be eliminated. Nevertheless, 3 studies in this consecutive series were uninterpretable. The score is robust and unaffected by scanning angle or interpretation of extent of middle cerebral artery territory.¹² In comparison with more elaborate and time-consuming perfusion threshold or volume measures¹³⁻¹⁷ for determining final infarct size, the ASPECTS scoring of CTP may seem relatively crude. We believe that this simplicity is its strength, and we have shown that CTP ASPECTS correlated well with final NCCT ASPECTS. The extended coverage that a 64-section scanner or other techniques (such as table toggle¹⁸ or double injection¹⁹) provide, however, may allow greater final infarct volume correlation.

At acute stroke presentation, limited data are available to guide thrombolysis decision-making or to provide prognostic information. Conflicting data exist as to whether patients with early ischemic signs are at higher risk for poor outcome and thrombolysis-related complications, based on NCCT interpretation.²⁰⁻²⁸ Although there is a general pattern of worse outcome when the hypoattenuation is widespread (ASPECTS > 7 or 1/3 middle cerebral arteries¹⁹⁻²³), only 3 studies have looked directly for an association between such patients and thrombolysis treatment and have found no such interac-

Table 3: Relationship between 3-month functional outcome (mRS) and dichotomized baseline CBV ASPECTS

Baseline CBV ASPECTS	mRS					Total
	0–2 (%)	3 (%)	4 (%)	5 (%)	6 (%)	
CBV < 8	0 (0)	1 (6.2)	5 (31.2)	7 (43.8)	3 (18.8)	16
CBV ≥ 8	9 (45)	5 (25)	4 (20)	1 (5)	1 (5)	20

Note:—CBV indicates cerebral blood volume.

Table 4: Interobserver variability

	ICC	CI
NCCT	0.65	0.49–0.78
CBV	0.69	0.54–0.81
CBF	0.82	0.72–0.89
MTT	0.81	0.7–0.89

Note:—CBV indicates cerebral blood volume; CBF, cerebral blood flow; MTT, mean transit time.

tion.^{25–27} Early ischemic changes are difficult to detect on NCCT, with low interobserver agreement for presence and extent.^{29,30} Whereas loss of gray-white differentiation is consistent with infarction,³¹ cerebral swelling may be intermixed, extending the apparent boundary of infarct. Cerebral swelling is associated with a variable and lower stroke risk³² and may account for overestimation of initial infarct size on NCCT. Use of ASPECTS on CTA signal intensity allows easier detection of the infarct extent and provides additional information.⁵ When compared with NCCT, the median cerebral blood volume ASPECTS was higher and lower in patients with good and bad outcome, respectively. A single previous study in the neurology literature has looked at ASPECTS interpretation of CTP and reported benefit of cerebral blood volume ASPECTS over NCCT.³³ However, a lower threshold of 6–7 was used, resulting in a false-negative rate of 13%–18%.

A limitation of this study is the absence of direct evidence of recanalization; however, the purpose was to investigate what prognostic baseline data the CTP study offers independent of other factors. Furthermore, unless continuous transcranial Doppler is used, the exact moment of recanalization and duration of ischemia cannot be known. A post-thrombolysis NIHSS improvement of ≥8 points (major neurologic improvement) was considered a surrogate marker of recanalization, supported by recent publications.^{6–9} In support of major neurologic improvement as a marker of treatment response, 86% of patients in this series fulfilling these criteria achieved a good outcome. This is similar for published data of outcome in patients with successful recanalization following intra-arterial therapy³⁴ and in agreement with outcomes in other studies using major neurologic improvement.⁸

In conclusion, a threshold of 8 applied to the ASPECTS interpretation of cerebral blood volume is equivalent to that of NCCT for radiologic outcome and may confer some advantage over NCCT at predicting 24-hour improvement and final clinical outcome. The results of our series indicate that patients treated with thrombolysis presenting with a cerebral blood volume ASPECTS of <8 have a poor outcome.

References

- von Kummer R, Nolte PN, Schnittger H, et al. Detectability of cerebral hemisphere ischaemic infarcts by CT within 6 h of stroke. *Neuroradiology* 1996;38:31–33
- Mullins ME, Lev MH, Schellingerhout D, et al. Influence of availability of clinical history on detection of early stroke using unenhanced CT and diffusion-weighted MR imaging. *AJR Am J Roentgenol* 2002;179:223–28

- Lev MH, Farkas J, Gemmete JJ, et al. Acute stroke: improved nonenhanced CT detection—benefits of soft-copy interpretation by using variable window width and center level settings. *Radiology* 1999;213:150–55
- Barber PA, Demchuk AM, Zhang J, et al. Validity and reliability of a quantitative computed tomography score in predicting outcome of hyperacute stroke before thrombolytic therapy: ASPECTS Study Group—Alberta Stroke Programme Early CT Score. *Lancet* 2000;355:1670–74
- Coutts SB, Lev MH, Eliasziw M, et al. ASPECTS on CTA source images versus unenhanced CT: added value in predicting final infarct extent and clinical outcome. *Stroke* 2004;35:2472–76
- Haley EC Jr, Lewandowski C, Tilley BC. Myths regarding the NINDS rt-PA Stroke Trial: setting the record straight. *Ann Emerg Med* 1997;30:676–82
- Saposnik G, Di Legge S, Webster F, et al. Predictors of major neurologic improvement after thrombolysis in acute stroke. *Neurology* 2005;65:1169–74
- Brown DL, Johnston KC, Wagner DP, et al. Predicting major neurological improvement with intravenous recombinant tissue plasminogen activator treatment of stroke. *Stroke* 2004;35:147–50
- Alexandrov AV, Demchuk AM, Felberg RA, et al. High rate of complete recanalization and dramatic clinical recovery during tPA infusion when continuously monitored with 2-MHz transcranial Doppler monitoring. *Stroke* 2000;31:610–14
- Weir NU, Pexman JH, Hill MD, et al. CASES investigators: how well does ASPECTS predict the outcome of acute stroke treated with IV tPA? *Neurology* 2006;67:516–18
- Cohnen M, Witsack HJ, Assadi S, et al. Radiation exposure of patients in comprehensive computed tomography of the head in acute stroke. *AJNR Am J Neuroradiol* 2006;27:1741–45
- Pexman JH, Barber PA, Hill MD, et al. Use of the Alberta Stroke Program Early CT Score (ASPECTS) for assessing CT scans in patients with acute stroke. *AJNR Am J Neuroradiol* 2001;22:1534–42
- Schaefer PW, Roccatagliata L, Ledezma C, et al. First-pass quantitative CT perfusion identifies thresholds for salvageable penumbra in acute stroke patients treated with intra-arterial therapy. *AJNR Am J Neuroradiol* 2006;27:20–25
- Wintermark M, Flanders AE, Velthuis B, et al. Perfusion-CT assessment of infarct core and penumbra: receiver operating characteristic curve analysis in 130 patients suspected of acute hemispheric stroke. *Stroke* 2006;37:979–85
- Murphy BD, Fox AJ, Lee DH, et al. Identification of penumbra and infarct in acute ischemic stroke using computed tomography perfusion-derived blood flow and blood volume measurements. *Stroke* 2006;37:1771–77
- Schaefer PW, Roccatagliata L, Ledezma C, et al. First-pass quantitative CT perfusion identifies thresholds for salvageable penumbra in acute stroke patients treated with intra-arterial therapy. *AJNR Am J Neuroradiol* 2006;27:20–25
- Wintermark M, Reichhart M, Thiran JP, et al. Prognostic accuracy of cerebral blood flow measurement by perfusion computed tomography, at the time of emergency room admission, in acute stroke patients. *Ann Neurol* 2002;51:417–32
- Roberts HC, Roberts TP, Smith WS, et al. Multisection dynamic CT perfusion for acute cerebral ischemia: the “toggling-table” technique. *AJNR Am J Neuroradiol* 2001;22:1077–80
- Wintermark M, Smith WS, Ko NU, et al. Dynamic perfusion CT: optimizing the temporal resolution and contrast volume for calculation of perfusion CT parameters in stroke patients. *AJNR Am J Neuroradiol* 2004;25:720–29
- Adams HP Jr, Adams RJ, Brott T, et al. Guidelines for the early management of patients with ischemic stroke: a scientific statement from the Stroke Council of the American Stroke Association. *Stroke* 2003;34:1056–83
- Moulin T, Cattin F, Crepin-Leblond T, et al. Early CT signs in acute middle cerebral artery infarction: predictive value for subsequent infarct locations and outcome. *Neurology* 1996;47:366–75
- von Kummer R, Bourquain H, Bastianello S, et al. Early prediction of irreversible brain damage after ischemic stroke at CT. *Radiology* 2001;219:95–100
- Hill MD, Rowley HA, Adler F, et al, and the PROACT-II Investigators. Selection of acute ischemic stroke patients for intra-arterial thrombolysis with pro-urokinase by using ASPECTS. *Stroke* 2003;34:1925–31. Epub 2003 Jul 3
- Hacke W, Kaste M, Fieschi C, et al. Intravenous thrombolysis with recombinant tissue plasminogen activator for acute hemispheric stroke: + The European Cooperative Acute Stroke Study (ECASS). *JAMA* 1995;274:1017–25
- Dzialowski I, Hill MD, Coutts SB, et al. Extent of early ischemic changes on computed tomography (CT) before thrombolysis: prognostic value of the Alberta Stroke Program Early CT Score in ECASS II. *Stroke* 2006;37:973–78

26. Cornu C, Boutitie F, Candelise L, et al. Streptokinase in acute ischemic stroke: an individual patient data meta-analysis—the Thrombolysis in Acute Stroke Pooling Project. *Stroke* 2000;31:1555–60
27. Patel SC, Levine SR, Tilley BC, et al. Lack of clinical significance of early ischemic changes on computed tomography in acute stroke. *JAMA* 2001;286:2830–38
28. Hill MD, Demchuk AM, Tomsick TA, et al. Using the baseline CT scan to select acute stroke patients for IV-IA therapy. *AJNR Am J Neuroradiol* 2006;27:1612–16
29. Tissue plasminogen activator for acute ischemic stroke: The National Institute of Neurological Disorders and Stroke rt-PA Stroke Study Group. *N Engl J Med* 1995;333:1581–87
30. Wardlaw JM, Dorman PJ, Lewis SC, et al. Can stroke physicians and neuroradiologists identify signs of early cerebral infarction on CT? *J Neurol Neurosurg Psychiatry* 1999;67:651–53
31. Sobesky J, von Kummer R, Frackowiak M, et al. Early ischemic edema on cerebral computed tomography: its relation to diffusion changes and hypoperfusion within 6 hours after human ischemic stroke—a comparison of CT, MRI and PET. *Cerebrovasc Dis* 2006;21:336–39
32. Na DG, Kim EY, Ryoo JW, et al. CT sign of brain swelling without concomitant parenchymal hypoattenuation: comparison with diffusion- and perfusion-weighted MR imaging. *Radiology* 2005;235:992–48
33. Parsons MW, Pepper EM, Chan V, et al. Perfusion computed tomography: prediction of final infarct extent and stroke outcome. *Ann Neurol* 2005;58:672–79
34. Jahan R, Duckwiler GR, Kidwell CS, et al. Intraarterial thrombolysis for treatment of acute stroke: experience in 26 patients with long-term follow-up. *AJNR Am J Neuroradiol* 1999;20:1291–99