



Get Clarity On Generics

Cost-Effective CT & MRI Contrast Agents

**FRESENIUS
KABI**

WATCH VIDEO

AJNR

This information is current as
of August 26, 2025.

**Quantitative Assessment of
Neovascularization after Indirect Bypass
Surgery: Color-Coded Digital Subtraction
Angiography in Pediatric Moyamoya Disease**

H.-H. Cho, J.-E. Cheon, S.-K. Kim, Y.H. Choi, I.-O. Kim,
W.S. Kim, S.-M. Lee, S.K. You and S.-M. Shin

AJNR Am J Neuroradiol 2016, 37 (5) 932-938

doi: <https://doi.org/10.3174/ajnr.A4614>

<http://www.ajnr.org/content/37/5/932>

Quantitative Assessment of Neovascularization after Indirect Bypass Surgery: Color-Coded Digital Subtraction Angiography in Pediatric Moyamoya Disease

 H.-H. Cho,  J.-E. Cheon,  S.-K. Kim,  Y.H. Choi,  I.-O. Kim,  W.S. Kim,  S.-M. Lee,  S.K. You, and  S.-M. Shin

ABSTRACT

BACKGROUND AND PURPOSE: For the postoperative follow-up in pediatric patients with Moyamoya disease, it is essential to evaluate the degree of neovascularization status. Our aim was to quantitatively assess the neovascularization status after bypass surgery in pediatric Moyamoya disease by using color-coded digital subtraction angiography.

MATERIALS AND METHODS: Time-attenuation intensity curves were generated at ROIs corresponding to surgical flap sites from color-coded DSA images of the common carotid artery, internal carotid artery, and external carotid artery angiograms obtained pre- and postoperatively in 32 children with Moyamoya disease. Time-to-peak and area under the curve values were obtained. Postoperative changes in adjusted time-to-peak (Δ TTP) and ratios of adjusted area under the curve changes (Δ AUC ratio) of common carotid artery, ICA, and external carotid artery angiograms were compared across clinical and angiographic outcome groups. To analyze diagnostic performance, we categorized clinical outcomes into favorable and unfavorable groups.

RESULTS: The Δ TTP at the common carotid artery increased among clinical and angiographic outcomes, in that order, with significant differences ($P = .003$ and $.005$, respectively). The Δ AUC ratio at the common carotid artery and external carotid artery also increased, in that order, among clinical and angiographic outcomes with a significant difference (all, $P = .000$). The Δ AUC ratio of ICA showed no significant difference among clinical and angiographic outcomes ($P = .418$ and $.424$, respectively). The Δ TTP for the common carotid artery of >1.27 seconds and the Δ AUC ratio of $>33.5\%$ for the common carotid artery and 504% for the external carotid artery are revealed as optimal cutoff values between favorable and unfavorable groups.

CONCLUSIONS: Postoperative changes in quantitative values obtained with color-coded DSA software showed a significant correlation with outcome scores and can be used as objective parameters for predicting the outcome in pediatric Moyamoya disease, with an additional cutoff value calculated through the receiver operating characteristic curve.

ABBREVIATIONS: AUC = area under the curve; CCA = common carotid artery; Δ AUC ratio = adjusted area under the curve changes; Δ TTP = difference between preoperative and postoperative time-to-peak; ECA = external carotid artery; EDAS = encephaloduroarteriosynangiosis; MMD = Moyamoya disease

Moyamoya disease (MMD) is a cerebrovascular disease causing progressive steno-occlusive changes at the terminal portion of the ICAs or proximal areas of the anterior or middle cerebral arteries with compensatory development of a fine vascular network, the so-called “Moyamoya” vessels.^{1–4} Surgical neo-

vascularization is considered the most successful means of improving cerebral hemodynamics and resolving clinical symptoms.^{1,2,4–7} These surgical procedures can be categorized into 3 types: direct, indirect, or combined bypass.^{1,2,5–9} Indirect bypass surgery using the parietal branch of the superficial temporal artery, encephaloduroarteriosynangiosis (EDAS), is preferred in pediatric patients with MMD because of the simple technique and the lower risk of temporary ischemia during the operation.^{5,10–12} Additional bifrontal encephalogaleo(perosteal)synangiosis can be performed to reinforce the anterior circulation.

After bypass surgery, it is essential to evaluate the degree of neovascularization via the external carotid artery (ECA) system and subsequent changes occurring in the ICAs. DSA is thought to be the most reliable imaging technique for these purposes and has been used for initial diagnosis and evaluation of neovascularization. However, evaluation of neovascularization after bypass sur-

Received July 27, 2015; accepted after revision September 15.

From the Department of Radiology (H.-H.C., J.-E.C., Y.H.C., I.-O.K., W.S.K., S.-M.L., S.K.Y.), Seoul National University Hospital, Seoul, Korea; Department of Radiology (J.-E.C., Y.H.C., I.-O.K., W.S.K.), Seoul National University College of Medicine, Seoul, Korea; Institute of Radiation Medicine (J.-E.C., I.-O.K., W.S.K.), Seoul National University Medical Research Center, Seoul, Korea; Division of Pediatric Neurosurgery (S.-K.K.), Seoul National University Children's Hospital, Seoul, Korea; and Department of Radiology (S.-M.S.), Seoul Metropolitan Government–Seoul National University, Boramae Medical Center, Seoul, Korea.

Please address correspondence to Jung-Eun Cheon, MD, Department of Radiology, Seoul National University Hospital, 101 Daehak-ro, Jongno-gu, Seoul, 110–744, Korea; e-mail: cheonje@snu.ac.kr

<http://dx.doi.org/10.3174/ajnr.A4614>

gery with DSA is usually subjective and could not provide the objective information for the postoperative revascularization status.

Recently, a color-coded postprocessing software program for the quantitative analysis of DSA was introduced¹³ and used as guidance for thrombolytic therapy or embolization and for perfusion analysis in adult patients with MMD.^{14–16} The purpose of this study was to quantitatively assess the neovascularization status after indirect bypass surgery in children with MMD by using postprocessing software.

MATERIALS AND METHODS

Patients

The institutional review board of Seoul National University Hospital approved this retrospective study; informed consent was waived. From August 2012 to March 2014, 198 patients underwent DSA for underlying MMD in our institution. We included patients with both preoperative and postoperative DSA after an initial unilateral EDAS. Ninety-one patients were excluded because there were no available preoperative ($n = 23$) or postoperative ($n = 68$) DSA images. Seventy-five patients were excluded because DSA images were obtained with a different protocol between the pre- and postoperative DSAs (no available common carotid artery [CCA] angiogram, $n = 28$; different contrast media injection rate and volume, $n = 35$; different magnification, $n = 12$).

Finally, 32 patients (13 male and 19 female; mean age, 8.0 years; range, 3–18 years) were included. Initial symptoms included motor TIA in 22 patients (68.8%), seizures in 6 patients (18.8%), headache in 2 patients (6.3%), and sensory TIA in 2 patients (6.3%). Among these, 23 patients (71.9%) underwent EDAS and 9 patients (28.1%) underwent unilateral EDAS and bifrontal encephaloplexy (periosteal) synangiosis simultaneously. Selection of the side for the initial operation was determined on the basis of the dominant hemisphere of the patient, symptom severity, and perfusion alteration based on MR imaging. None suffered from postoperative complications, including hemorrhage or infarction involving the ipsilateral brain hemisphere, seen on the postoperative imaging studies such as CT or MR imaging.

Conventional DSA

All enrolled patients underwent both preoperative and postoperative cerebral angiography in a dual-plane angiographic machine (Axiom Artis; Siemens, Erlangen, Germany) performed by 3 pediatric radiologists (Y.H.C., H.-H.C., and S.-M.L., with 10, 6, and 6 years of experience, respectively). A 4F or 5F Davis angiographic catheter (A&A Medical Device, Gyeonggi-do, Korea) was used according to the patient's femoral artery diameter. All procedures were performed with same catheter diameter for both preoperative and postoperative cerebral angiography. Conventional DSA images included anteroposterior and lateral ICA, ECA, and CCA angiograms. Diluted contrast media (ioditridol; Xenetix 350; Guerbet, Aulnay-sous-Bois, France) with a contrast-to-normal-saline ratio of 1:1 was used with an automatic injector (Angiomat Illumena; Liebel-Flarsheim, Cincinnati, Ohio). The injection rate and amount were determined by the operator on the basis of the

size of each vessel. The mean injection rates for patients younger than 5 years of age were 2 mL/s for the CCA, 1 mL/s for the ICA, and 1.5 mL/s for the ECA. The rates for patients older than 5 years were approximately 3 mL/s for the CCA, 1.5 mL/s for the ICA, and 2 mL/s for ECA angiography, and they were adjusted for each patient's circumstances. The injection duration of contrast material was fixed for 2 seconds. The same contrast injection rate and amount were used for pre- and postoperative DSA in each patient. The mean time interval between pre- and postoperative DSA was approximately 82.9 days (range, 72–103 days), and the mean time interval between the operation and postoperative DSA was approximately 76.2 days (range, 64–96). The stages of preoperative DSA were classified by using the criteria of Suzuki and Kodama¹⁷: stage 1 ($n = 4$), stage 2 ($n = 9$), and stage 3 ($n = 19$).

Postprocessing of DSA

Postprocessing of DSA images was performed by using syngo iFlow (Siemens). With this program, we could obtain a color-coded (range from red to blue) single image showing the history of contrast material passage through vessels, which reflected the time from injection. This color map provided flow velocity and total amount of injected contrast material passing through the vessels. We also obtained signal-intensity curves of the selected ROIs, adjusted with a reference ROI. Two pediatric radiologists (H.-H.C. and S.-M.L. with 6 years of experience each) drew a rectangular ROI (approximately 2500 mm²) to cover the EDAS flap in consensus. A round reference ROI was drawn within the proximal input artery (approximately 10 mm²) at the level of the atlas for maintenance of the identical location between the pre- and postinterventional DSAs (Fig 1A). A time-attenuation intensity curve was obtained automatically and summarized the maximum concentration of contrast passing through the ROI at that point (Fig 1B). Using the curve, the program automatically calculated the “peak time of contrast concentration,” which corresponds to time-to-peak of perfusion MR imaging. The calculated area under the curve (AUC) is the total amount of contrast material passed, which corresponds to the cerebral blood volume of the perfusion MR imaging. The TTP difference (Δ TTP) in seconds was calculated by subtracting the TTP of the preoperative DSA from the TTP of the postoperative DSA. For the AUC, the adjusted area under the curve changes (Δ AUC ratio) in percentages were calculated by dividing the difference between post- and preoperative AUCs by the preoperative AUC values (Fig 1C):

$$\Delta\text{TTP} = \text{TTP}_{\text{preoperative}} - \text{TTP}_{\text{postoperative}}$$

$$\Delta\text{AUC Ratio} = (\text{AUC}_{\text{postoperative}} - \text{AUC}_{\text{preoperative}}) / \text{AUC}_{\text{preoperative}}$$

In this study, postprocessing of DSA was performed by using only a lateral projection because detection of the EDAS flap site was easier than in anteroposterior images. Likewise, the widely used angiographic staging method for MMD, the Matsushima classification, also used lateral images for evaluating neovascularization degree.¹⁸

Angiographic Outcome Group

To classify postoperative neovascularization status, we used the Matsushima 3-grade method.¹⁸ The “good group” was defined as

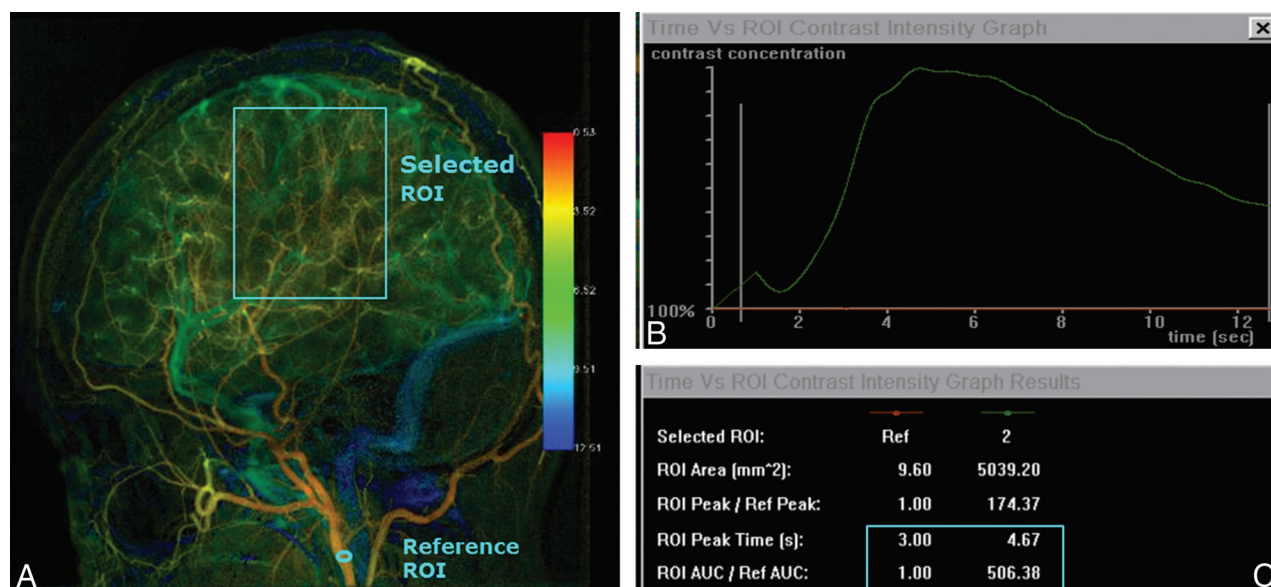


FIG1. Postprocessing cascade of the DSA image. *A*, A color-coded single image is obtained from DSA images of CCA angiography; then, we draw selected ROIs along the estimated EDAS flap site. A reference ROI is drawn within the proximal input artery. *B*, The time-attenuation intensity curve is obtained automatically. The superior green line represents the time-attenuation intensity curve within the selected ROI, and the inferior red line represents that of the reference ROI. *C*, Using the time-attenuation intensity curve, we calculated the TTP and AUC within the selected ROI, adjusted by the reference ROI.

more than two-thirds of the MCA distribution supplied by neovascularization after surgery. The “fair group” was defined as a neovascularization area covering between two-thirds and one-third of the MCA distribution. The “poor group” was defined as only 1 cortical branch of the MCA covered through the bypass or no collateral circulation observed. Angiographic outcomes were classified by a radiologist after each procedure.

Clinical Outcome Group

Postoperative clinical outcomes were classified according to ipsilateral MCA symptoms (such as TIAs or seizures) by the clinician at the time of admission for the postoperative angiography, blinded to the results of imaging studies, including MR imaging, perfusion MR imaging, or DSA.^{6,19} The “excellent group” represented total disappearance of symptoms (such as TIAs or seizures) without fixed neurologic deficits. The “good group” represented total disappearance of symptoms, but with neurologic deficits remaining. The “fair group” represented persistent symptoms, albeit less frequently. The “poor group” represented remaining unchanged or worsened symptoms. The excellent and good groups were considered favorable outcomes, and the fair and poor groups were considered unfavorable outcomes in receiver operating characteristic curve analysis. Anterior cerebral artery symptoms, such as paraparesis or urinary incontinence, were not considered in this study because it was designed for the evaluation of the neovascularization status of EDAS sites.

Statistical Analysis

Comparisons between AUC differences in CCA, ECA, and ICA angiograms and mean TTP differences in the CCA between pre- and postoperative studies in each clinical and angiographic outcome group were analyzed by using the Jonckheere-Terpstra test by using PASW software for Windows (Version 18.0; IBM, Ar-

monk, New York). To determine the diagnostic performance of parameters in each angiogram, we performed receiver operating characteristic analysis by using MedCalc for Windows (MedCalc Software, Mariakerke, Belgium). *P* values < .05 were statistically significant.

RESULTS

DSA Parameters and Clinical Outcome

Sixteen patients showed excellent; 4 patients, good; 8 patients, fair; and 4 patients, poor clinical outcomes. After neovascularization surgery, postoperative TTP decreased compared with preoperative TTP in CCA angiography in the excellent, good, and fair clinical outcome groups. In the poor clinical outcome group, postoperative TTP increased compared with preoperative TTP. The Δ TTP values in CCA angiography were significantly different among clinical outcome groups (*P* = .003), and a gradual decrease of Δ TTP was noted from the excellent-to-poor groups.

Calculated Δ AUC ratios in CCA angiography showed a significant increase among clinical outcome groups in sequence from poor to excellent (*P* = .000). Mean Δ AUC ratios in CCA angiography were 68.1% for excellent, 31.5% for good, 9.1% for fair, and -11.3% for poor. The Δ AUC ratios of ECA angiography also showed a significant increase among clinical outcome groups in sequence from poor to excellent (*P* = .000). The Δ AUC ratios in ECA angiography were 1119.8% for excellent, 768.2% for good, 279.5% for fair, and 248.7% for poor. Figures 2 and 3 show representative cases with pre- and postoperative DSA images in the excellent and poor clinical outcome groups. The Δ AUC ratios in ICA angiography showed no significant difference among the clinical outcome groups (*P* = .418). The Δ AUC ratios in ICA angiography were -41.0% for

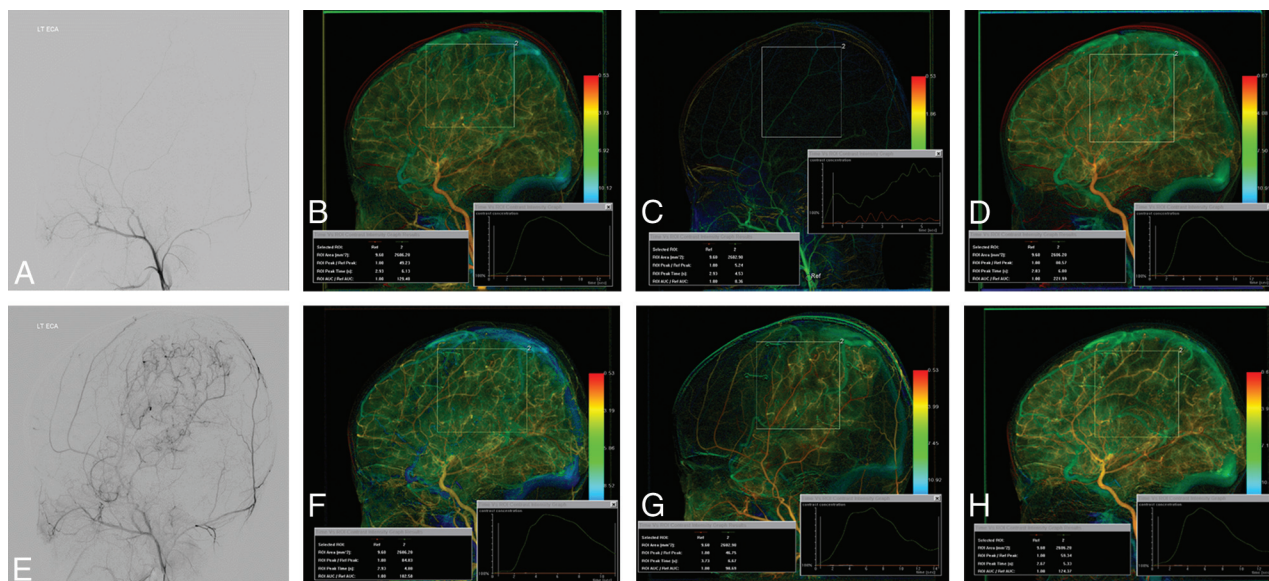


FIG 2. A 4-year-old boy with excellent clinical outcome after EDAS neovascularization. Comparing pre- (A) and post- (E) operative ECA angiograms shows good neovascularization after EDAS. Postprocessed pre- (B) and post- (F) operative CCA angiograms show shortening of TTP of approximately 2.3 seconds and an increase of the percentage of AUC difference of about 51.0% after the operation. Approximately a 1080.5% increment of the percentage of AUC difference after surgery is noted when comparing pre- (C) and post- (G) operative ECA angiograms, and an approximate 44.0% of AUC difference decrease is noted when comparing pre- (D) and post- (H) operative ICA angiograms.

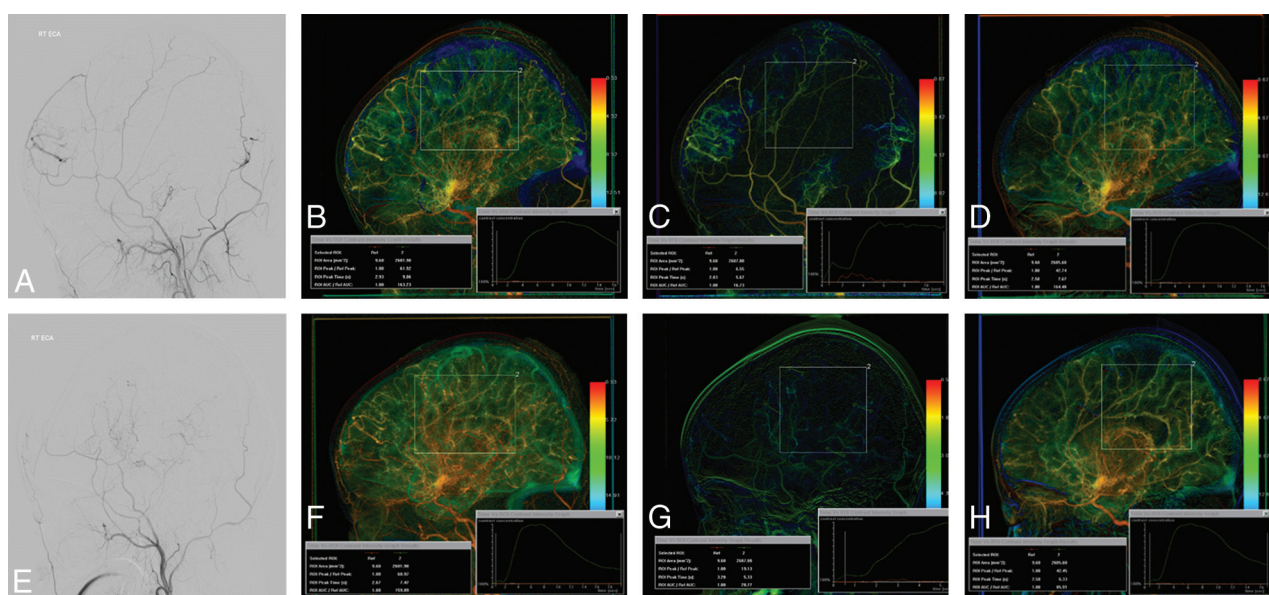


FIG 3. A 5-year-old boy with poor clinical outcome after EDAS neovascularization. Comparing pre- (A) and post- (E) operative ECA angiograms shows poor neovascularization after EDAS. Postprocessed pre- (B) and post- (F) operative CCA angiograms show no definite shortening of the TTP (0.0 seconds), with a decrease of the percentage of AUC difference of approximately 2.0% after the operation. About a 41.2% increase of the percentage of AUC difference after the operation is noted when comparing pre- (C) and post- (G) operative ECA angiograms, and about a 13.7% of AUC difference increase is noted when comparing pre- (D) and post- (H) operative ICA angiograms.

Table 1: Relationships between DSA parameters and clinical outcomes

Group	Excellent (n = 16)	Good (n = 4)	Fair (n = 8)	Poor (n = 4)	P Value
Δ TTP in CCA (sec)	2.02	1.0	0.80	−0.13	.003
Δ AUC ratio in CCA (%)	68.1	31.5	9.1	−11.3	.000
Δ AUC ratio in ECA (%)	1119.8	768.2	279.5	248.7	.000
Δ AUC ratio in ICA (%)	−41.0	−35.6	−44.2	−22.3	.418

excellent, −35.6% for good, −44.2% for fair, and −22.3% for poor (Table 1).

DSA Parameters and Angiographic Outcome

Angiographic outcome by using the Matsushima grade was good in 16, fair in 7, and poor in 9 patients. Δ TTP showed a statistically significant difference among angiographic outcome groups in CCA angiography ($P = .005$), and a gradual decrease of Δ TTP was noted from good to poor.

Table 2: Relationships between DSA parameters and angiographic outcomes

Group	Good (n = 16)	Fair (n = 7)	Poor (n = 9)	P Value
ΔTTP in CCA (sec)	2.01	0.97	0.49	.005
ΔAUC ratio in CCA (%)	64.5	29.8	0.4	.000
ΔAUC ratio in ECA (%)	1065.9	753.1	210.5	.000
ΔAUC ratio in ICA (%)	−41.0	−42.4	−32.2	.424

Table 3: Cutoff values between favorable and unfavorable groups

Parameter	Criterion	Sensitivity (%)	Specificity (%)
ΔTTP in CCA (sec)	>1.27	81.4	91.7
ΔAUC ratio in CCA (%)	>33.5	81	100
ΔAUC ratio in ECA (%)	>504	85.3	92
ΔAUC ratio in ICA (%)	≤20.2	85.9	33.4

The ΔAUC ratios in CCA angiography showed a statistically significant difference among angiographic outcome groups in sequence from poor to excellent ($P = .000$), being approximately 64.5% for good, 29.8% for fair, and 0.4% for poor. The ΔAUC ratios of ECA angiography also showed a statistically significant difference among the angiographic outcome groups in sequence from poor to excellent ($P = .000$). The ΔAUC ratios in ECA angiography were noted as approximately 1065.9% for good, 753.1% for fair, and 210.5% for poor. The ΔAUC ratios of ICA angiography showed no significant difference among the angiographic outcome groups ($P = .424$). The ΔAUC ratios in ICA angiography were approximately −41.0% for good, −42.4% for fair, and −32.2% for poor groups (Table 2).

The ΔTTP and ΔAUC ratio showed no significant difference between EDAS only and unilateral EDAS in the bifrontal encephalogleo(perosteal)synangiosis groups ($P = .76$).

Cutoff Value between Favorable and Unfavorable Groups

According to receiver operating characteristic analysis, an optimal cutoff value of 1.27 seconds was obtained for the difference between pre- and postoperative adjusted TTP in CCA angiography, with 81.4% sensitivity and 91.7% specificity between favorable and unfavorable groups. The optimal cutoff value for the difference ratio of the AUC in CCA angiography was 33.5%, with 81% sensitivity and 100% specificity, while that in ECA angiography was 504%, with 85.3% sensitivity and 92% specificity (Table 3).

DISCUSSION

The optimal goal for EDAS in pediatric patients with MMD is to re-establish cerebral blood flow to hypoperfused areas by using angiogenesis from parietal branches of the superficial temporal artery.²⁰ After the operation, collateral vessels from the superficial temporal artery develop at the flap site, along with simultaneous and sequential progression of steno-occlusive changes of the ICA and disappearance or diminishing of Moyamoya vessels.^{18,21} Hemodynamic changes after the operation might be difficult to understand; likewise, estimating postoperative neovascularization status is challenging. Noninvasive imaging modalities such as perfusion MR imaging, single-photon emission CT, and positron-emission tomography

can demonstrate secondary hemodynamic changes.^{1-4,12,21-27} However, these methods show only overall cerebral perfusion changes and cannot directly visualize fine parenchymal neovascularization itself following the operation.

Concerning the evaluation of postoperative neovascularization status, DSA is still the most reliable imaging technique and is widely used for the initial diagnosis and follow-up of MMD.^{1,9,19} DSA also provides anatomic information of vessels and direct visualization of steno-occlusive changes and visualizes neovascularization status after the operation. However, it cannot provide the objective perfusion parameters. It might be difficult to compare the change in the DSA during longitudinal or postoperative follow-up. The decision for the postoperative stage might be made only by the operator.

In this study, to supplement these shortcomings of DSA, we used the postprocessing software syngo iFlow. This software calculates opacity changes within each pixel at each moment and then summarizes these to make a color-coded map of contrast flow. With this software, additional hemodynamic information can be obtained from serial DSA images without additional radiation exposure or contrast material, with a simplified method with little effort.

Although this study included only hemodynamic changes within the operative bed, we predicted that the increase of AUC with shortening TTP might be noted on CCA and ECA angiography. We also predicted that a decrease of AUC in ICA angiography might be noted after neovascularization surgery because ICA flows, including basal collaterals (Moyamoya vessels), are no longer required to function as collateral pathways. The actual results showed patterns similar to our estimations, except in the poor clinical outcome group. The largest ΔTTP, which represents the degree of shortening-delayed TTP due to neovascularization, was noted in the excellent clinical outcome group on CCA angiography. The largest increment of the ΔAUC ratio, which represents increased CBV through neovascularization, was also noted in the excellent clinical outcome group in CCA and ECA angiography. We suggest that the values obtained from CCA angiography provide real hemodynamic changes because those values might include both increased blood flow through neovascularization of the superficial temporal artery and diminished blood flow through the ipsilateral ICA.

With receiver operating characteristic analysis, improved TTP delay of >1.27 seconds and improved percentage AUC difference of >33.5% for CCA and 504% for ECA angiography are optimal cutoff values between favorable and unfavorable groups with reliable sensitivity and specificity. Using these cutoff values, we could provide additional information to predict patient outcomes; and for the unfavorable group, an additional operation for the contralateral side or bifrontal encephalogleo(perosteal)synangiosis was considered earlier than in the favorable group. In addition, in these cases, shorter imaging and clinical follow-up were recommended.

The mean time interval between the operation and postoperative DSA was approximately 76.2 days (range, 64~96). Moreover, neovascularization after indirect bypass surgery was known to become well-developed after 3~6 months postoperatively.^{20,28-31}

There are several study limitations. In some cases, postoperative DSA data might not be enough to provide real hemo-

dynamic status after neovascularization. However, previous studies explained that the neovascularization begins 2 weeks after surgery,^{21,32,33} so the obtained data in this study could provide the near-complete status of the neovascularization. In addition, although the data obtained from the postoperative DSA might provide the earlier neovascularization status before completion, we thought they might provide the later outcome because the data showed good correlation with the outcomes in this study.

Although neovascularization after indirect bypass surgery develops along the flap site, accurate flap site drawing was difficult; thus, another limitation is that some neovascularization developing outside the flap site was not included in this study. However, the portion of neovascularization developing outside the flap site might be small and therefore negligible. We used a manufactured head fixator for maintenance of the head position during the study, so another limitation of the study could be that there was little difference between the pre- and postoperative DSA. We suggest that the standardized positioning might be helpful for a more exact comparison.

CONCLUSIONS

Using simple postprocessing software might provide objective quantitative perfusion values based on obtained DSA images with significant correlation with clinical and angiographic outcomes for pediatric patients with MMD without additional radiation. Furthermore, the cutoff values obtained with receiver operating characteristic analysis may help in predicting patient outcomes after the operation and could affect the establishment of further additional treatment or follow-up plans.

ACKNOWLEDGMENTS

We thank Hyung Bin Yu, the project manager of Siemens, Korea, for technical support.

REFERENCES

- Kuroda S, Houkin K. **Moyamoya disease: current concepts and future perspectives.** *Lancet Neurol* 2008;7:1056–66 Medline
- Kim SK, Cho BK, Phi JH, et al. **Pediatric moyamoya disease: an analysis of 410 consecutive cases.** *Ann Neurol* 2010;68:92–101 CrossRef Medline
- Suzuki J, Takaku A. **Cerebrovascular “moyamoya” disease: disease showing abnormal net-like vessels in base of brain.** *Arch Neurol* 1969;20:288–99 Medline
- Fukui M. **Guidelines for the diagnosis and treatment of spontaneous occlusion of the circle of Willis (‘moyamoya’ disease): Research Committee on Spontaneous Occlusion of the Circle of Willis (Moyamoya Disease) of the Ministry of Health and Welfare, Japan.** *Clin Neurol Neurosurg* 1997;99(suppl 2):S238–40 Medline
- Kim SK, Wang KC, Kim IO, et al. **Combined encephaloduroarteriosynangiosis and bifrontal encephalogaleo (periosteal) synangiosis in pediatric moyamoya disease.** *Neurosurgery* 2008;62:1456–64 Medline
- Kim CY, Wang KC, Kim SK, et al. **Encephaloduroarteriosynangiosis with bifrontal encephalogaleo(periosteal)synangiosis in the pediatric moyamoya disease: the surgical technique and its outcomes.** *Child Nerv Syst* 2003;19:316–24 Medline
- Mesiwala AH, Sviri G, Fatemi N, et al. **Long-term outcome of superficial temporal artery-middle cerebral artery bypass for patients with moyamoya disease in the US.** *Neurosurg Focus* 2008;24:E15 CrossRef Medline
- Houkin K, Kuroda S, Ishikawa T, et al. **Neovascularization (angiogenesis) after revascularization in moyamoya disease: which technique is most useful for moyamoya disease?** *Acta Neurochir (Wien)* 2000;142:269–76 Medline
- Goda M, Isono M, Ishii K, et al. **Long-term effects of indirect bypass surgery on collateral vessel formation in pediatric moyamoya disease.** *J Neurosurg* 2004;100:156–62 Medline
- Yun TJ, Cheon JE, Na DG, et al. **Childhood moyamoya disease: quantitative evaluation of perfusion MR imaging—correlation with clinical outcome after revascularization surgery.** *Radiology* 2009;251:216–23 CrossRef Medline
- Choi JU, Kim DS, Kim EY, et al. **Natural history of moyamoya disease: comparison of activity of daily living in surgery and non-surgery groups.** *Clin Neurol Neurosurg* 1997;99(suppl 2):S11–18 Medline
- Kim SK, Wang KC, Oh CW, et al. **Evaluation of cerebral hemodynamics with perfusion MRI in childhood moyamoya disease.** *Pediatr Neurol* 2003;38:68–75 Medline
- Strother CM, Bender F, Deuerling-Zheng Y, et al. **Parametric color coding of digital subtraction angiography.** *AJNR Am J Neuroradiol* 2010;31:919–24 CrossRef Medline
- Lin CJ, Yu M, Hung SC, et al. **In-room assessment of cerebral blood volume for guidance during intra-arterial thrombolytic therapy.** *Interv Neurol* 2012;18:463–68 Medline
- Zhang XB, Zhuang ZG, Ye H, et al. **Objective assessment of transcatheter arterial chemoembolization angiographic endpoints: preliminary study of quantitative digital subtraction angiography.** *J Vasc Interv Radiol* 2013;24:667–71 CrossRef Medline
- Hung SC, Liang ML, Lin CF, et al. **New grading of moyamoya disease using color-coded parametric quantitative digital subtraction angiography.** *J Chin Med Assoc* 2014;77:437–42 CrossRef Medline
- Suzuki J, Kodama N. **Moyamoya disease: a review.** *Stroke* 1983;14:104–09 Medline
- Matsushima T, Inoue T, Suzuki SO, et al. **Surgical treatment of moyamoya disease in pediatric patients: comparison between the results of indirect and direct revascularization procedures.** *Neurosurgery* 1992;31:401–05 Medline
- Kim SK, Wang KC, Kim IO, et al. **Combined encephaloduroarteriosynangiosis and bifrontal encephalogaleo(periosteal)synangiosis in pediatric moyamoya disease.** *Neurosurgery* 2002;50:88–96 Medline
- Matsushima T, Fujiwara S, Nagata S, et al. **Surgical treatment for paediatric patients with moyamoya disease by indirect revascularization procedures (EDAS, EMS, EMAS).** *Acta Neurochir (Wien)* 1989;98:135–40 Medline
- Houkin K, Nakayama N, Kuroda S, et al. **How does angiogenesis develop in pediatric moyamoya disease after surgery? A prospective study with MR angiography.** *Childs Nerv Syst* 2004;20:734–41 Medline
- Honda M, Kitagawa N, Tsutsumi K, et al. **Magnetic resonance angiography evaluation of external carotid artery tributaries in moyamoya disease.** *Surg Neurol* 2005;64:325–30 Medline
- Ikezaki K, Matsushima T, Kuwabara Y, et al. **Cerebral circulation and oxygen metabolism in childhood moyamoya disease: a perioperative positron emission tomography study.** *J Neurosurg* 1994;81:843–50 Medline
- So Y, Lee HY, Kim SK, et al. **Prediction of the clinical outcome of pediatric moyamoya disease with postoperative basal/acetazolamide stress brain perfusion SPECT after revascularization surgery.** *Stroke* 2005;36:1485–89 Medline
- Hoshi H, Ohnishi T, Jinnouchi S, et al. **Cerebral blood flow study in patients with moyamoya disease evaluated by IMP SPECT.** *J Nucl Med* 1994;35:44–50 Medline
- Mountz JM, Foster NL, Ackermann RJ, et al. **SPECT imaging of moyamoya disease using 99mTc-HM-PAO: comparison with computed tomography findings.** *J Comput Tomogr* 1988;12:247–50 Medline
- Fujimura M, Mugikura S, Shimizu H, et al. **Diagnostic value of perfusion-weighted MRI for evaluating postoperative alteration of cerebral hemodynamics following STA-MCA anastomosis in patients with**

- moyamoya disease** [in Japanese]. *No Shinkei Geka* 2006;34:801–09 [Medline](#)
28. Fung LW, Thompson D, Ganesan V. **Revascularisation surgery for paediatric moyamoya: a review of the literature.** *Childs Nerv Syst* 2005;21:358–64 [Medline](#)
 29. Sakamoto H, Kitano S, Yasui T, et al. **Direct extracranial-intracranial bypass for children with moyamoya disease.** *Clin Neurol Neurosurg* 1997;99(suppl 2):S128–33 [Medline](#)
 30. Togao O, Mihara F, Yoshiura T, et al. **Cerebral hemodynamics in moyamoya disease: correlation between perfusion-weighted MR imaging and cerebral angiography.** *AJNR Am J Neuroradiol* 2006;27:391–97 [Medline](#)
 31. Kassner A, Zhu XP, Li KL, et al. **Neoangiogenesis in association with moyamoya syndrome shown by estimation of relative recirculation based on dynamic contrast-enhanced MR images.** *AJNR Am J Neuroradiol* 2003;24:810–18 [Medline](#)
 32. Kinugasa K, Mandai S, Kamata I, et al. **Surgical treatment of moyamoya disease: operative technique for encephalo-duro-arterio-myo-synangiosis, its follow-up, clinical results, and angiograms.** *Neurosurgery* 1993;32:527–31 [Medline](#)
 33. Kim SK, Seol HJ, Cho BK, et al. **Moyamoya disease among young patients: its aggressive clinical course and the role of active surgical treatment.** *Neurosurgery* 2004;54:840–44; discussion 844–46 [Medline](#)