Indirect revascularization is the most widely used treatment to induce angiogenesis in pediatric moyamoya disease (MMD). Molecular imaging methods targeted for angiogenesis have recently been developed. We performed angiogenesis imaging in indirect revascularization surgery for MMD to evaluate angiogenic activity and its correlation with treatment efficacy. **Methods:** Twelve patients with pediatric MMD were prospectively enrolled. Encephaloduroarteriosynangiosis surgery was conducted, and a \( ^{68}\text{Ga}-\text{Arg-Gly-Asp} \) (RGD) PET was performed at 3.7 \( \pm \) 1.0 mo after surgery. Basal perfusion and stress perfusion (PStr) in the middle cerebral artery territory were evaluated by acetazolamide-stress brain perfusion SPECT using statistical probabilistic anatomic mapping, at preoperative, early postoperative, and long-term follow-up states. Angiogenic activity was assessed on the images in terms of maximal uptake ratio, volume of increased uptake, and uptake-volume product. **Results:** Basal perfusion and PStr were significantly improved after surgery. Increased angiogenic activity was observed in the revascularized area, mainly around the bony flap. Angiogenic activity gradually decreased with time and significantly correlated with the postoperative time interval \( (P = 0.0015) \) for maximal uptake ratio and 0.0069 for volume of increased uptake. It was estimated to normalize at 6.3 mo after surgery. Uptake-volume product was inversely correlated with PStr improvement at the early postoperative state \( (r = -0.5960, P = 0.0409) \) and also weakly correlated with PStr improvement at long-term follow-up \( (r = -0.5010, P = 0.1165) \). **Conclusion:** Angiogenesis PET imaging with \( ^{68}\text{Ga}-\text{RGD} \) was successfully used for the assessment of angiogenic activity in indirect revascularization surgery for MMD, and angiogenic activation measured at approximately 3.7 mo after surgery was inversely correlated with perfusion improvement. The assessment of angiogenic activity using \( ^{68}\text{Ga}-\text{RGD} \) PET is expected to be effective for evaluating the mechanism or efficacy of revascularization treatment.
of pediatric MMD, the assessment of angiogenic activity by in vivo molecular imaging would be beneficial for investigating the revascularization mechanism and evaluating the efficacy of treatment.

In this study, we evaluated angiogenic activity using in vivo angiogenesis PET imaging in pediatric MMD patients who underwent indirect revascularization surgery. On the angiogenesis PET images, image characteristics, time course of angiogenic activity, and correlation between angiogenic activity and revascularization efficacy were analyzed.

MATERIALS AND METHODS

Patients and Study Design

Pediatric patients with newly diagnosed MMD were prospectively enrolled in this study. The following criteria were used for inclusion: MMD confirmed by clinical and radiologic findings; absence of other combined disorders suggesting moyamoya syndrome; absence of previous wide cerebral infarction; and eligibility for standard revascularization surgery and image studies including brain perfusion SPECT, MR imaging, cerebral angiography, and angiogenesis PET. The study design was approved by the Institutional Review Board (IRB) of Seoul National University Hospital (IRB no. H-0908-067-292), and signatures on written informed consent forms were obtained from all parents or legal guardians according to the Declaration of Helsinki.

The overall management methods and surgical outcome of our patients were reported previously (4). All patients underwent preoperative imaging studies including brain MR imaging, cerebral angiography, and brain perfusion SPECT. Revascularization surgery was performed using the unilateral encephaloduroarteriosynangiosis method in which the superficial temporal artery and galeal flap were laid on the surface of the exposed cortex and sutured to the incised edge of the dura mater after dissection of the arachnoid membrane. In some patients, the frontal or occipital area was simultaneously revascularized using the encephaloduroarteriosynangiosis method for the bifrontal area and the encephaloduroarteriosynangiosis method for the occipital area. Angiogenesis PET was performed more than 2 mo after surgery to minimize the effect of postoperative acute inflammation, before the next surgical revascularization for the contralateral side, if any. The patients were closely followed up, and early postoperative brain perfusion SPECT was performed 3 mo after surgery. To evaluate the long-term outcome, brain perfusion SPECT was again performed more than 1 y after surgery.

Image Acquisition

For angiogenesis PET, 68Ga-RGD was prepared in our institution as described previously (7). In brief, a cyclic RGD (cRGDyK) was combined with a derivative of 1,4,7-triazacyclononane-1,4,7-triacetic acid (NOTA). This compound was added to 68Ga eluted from a 56Ge/68Ga generator, and 68Ga-RGD was purified from the mixture using high-performance liquid chromatography and alumina sorbent cartridges. 68Ga-RGD PET images were acquired using a whole-body PET/CT device equipped with a 40-slice CT scanner (Biograph 40; Siemens Healthcare), which has a PET spatial resolution of 4.2 mm. Patients were injected with 68Ga-RGD (5.5 MBq/kg), and images were acquired 20 min after the injection. A low-dose CT scan was obtained first for attenuation correction, and PET images were acquired for 30 min using 3-dimensional mode. PET images were reconstructed using an iterative algorithm (order-subset expectation maximization; 4 iterations and 8 subsets) on 256 × 256 matrices.

Basal/acetazolamide brain perfusion SPECT was performed using a triple-head γ camera (Triad XLT 9; Trionix Research Laboratory) equipped with low-energy ultrahigh-resolution fanbeam collimators. Basal SPECT images were acquired 5 min after intravenous injection of 99mTc-hexamethylpropyleneamine oxime (9.25 MBq/kg). Forty step-and-shoot images were acquired for 20–25 s per step, with intervals of 3°. Ten minutes before the end of the basal SPECT, acetazolamide (20 mg/kg) was injected intravenously for vasodilatation, and 99mTc-hexamethylpropyleneamine oxime (18.5 MBq/kg) was injected again at the end of the basal SPECT acquisition. Five minutes later, a second acquisition of SPECT was started without position change. Acetazolamide-stress SPECT images were derived by decay-corrected subtraction of the basal images from the second SPECT images. All SPECT images were reconstructed on 128 × 128 matrices using a filtered backprojection method with a Butterworth filter.

Brain MR imaging (including MR angiography and perfusion imaging) and cerebral angiography were performed in routine clinical practice before and after the surgery.

Image Analysis

Angiogenic activity was assessed on PET images in terms of 68Ga-RGD uptake. For quantification, the contralateral temporoparietal brain parenchyma was adopted as a reference region, in which a spheric volume of interest (VOI; 4.04 cm³) was drawn to measure the mean and SD of uptake counts. In the revascularized temporoparietal area, an isocontour VOI for increased uptake was defined as the maximal uptake ratio (URmax), volume of increased uptake (V IU), and uptake-volume product (UVP), which were defined as follows: URmax = (maximal count in the VOI for increased uptake)/(mean count in the reference VOI), V IU = (size of the VOI for increased uptake in mm³), and UVP = (V IU) × (mean count ratio in the VOI). URmax was deemed as the maximal angiogenic activity, V IU was deemed as the involved area by angiogenic activation, and UVP was deemed as the overall angiogenic activity. The frontal or occipital area was not included for the analysis even when the area was revascularized during surgery.

The regional brain perfusion in the middle cerebral artery (MCA) territory was assessed on perfusion SPECT. Each SPECT image was spatially normalized to the ethnically matched standard perfusion SPECT template (15) using SPM2 software (University College London) implemented in MATLAB 6.5 (The MathWorks). The probabilistic VOIs for the MCA territory and cerebellum were automatically
defined using a statistical probabilistic anatomic map (SPAM) (16), and perfusion values of these VOIs were calculated. The perfusion value of the MCA territory was normalized to the mean value of both cerebellar lobes and expressed as a percentage. Basal perfusion (P\textsubscript{Bas}) and perfusion values of these VOIs were calculated.

Revascularization surgery was performed using encephaloduroar-teriosynangiosis for the right temporoparietal area and left tempoparietal area in 6 patients each. The frontal area was also revascularized using encephaloduroar-teriosynangiosis in 2 patients. All patients underwent a second revascularization surgery for the contralateral side 4.0±1.2 mo (range, 2.4–5.8 mo) after the first surgery. Three patients exhibited postoperative complications of focal infarction or a small amount of subdural hemorrhage (in the right anterior frontal lobe in 2 patients, and in the left high frontal lobe in 1 patient). In the follow-up after revascularization surgery, no patient presented with overt ischemic symptoms related to the operated area.

**Perfusion Change**

Preoperative perfusion SPECT was performed 0.6±0.8 mo (range, 0.0–2.3 mo) before revascularization surgery, and early postoperative SPECT was performed 3.0±0.9 mo (range, 1.9–4.6 mo) after surgery. Long-term follow-up SPECT was performed 31.7±7.2 mo (range, 14.4–40.8 mo) after surgery. In 1 patient, long-term follow-up SPECT was not performed because of intolerance to acetazolamide stress. P\textsubscript{Bas} and P\textsubscript{Str} in the operated MCA territory were 94.3±9.0 and 86.0±9.5 before revascularization surgery, and they significantly improved to 103.2±8.8 and 95.6±9.5 at the first postoperative follow-up (P = 0.0047 and 0.0064, respectively). At the long-term follow-up, they were 103.8±18.2 and 103.7±17.3 (P = 0.0560 and 0.0020, respectively, in comparison with preoperative values; Fig 1).

**Angiogenic Activity After Revascularization Surgery**

\(^{68}\)Ga-RGD PET was performed 3.7±1.0 mo (range, 2.1–5.3 mo) after revascularization surgery. On PET images, increased uptake was observed in the operated area, mainly around the bony flap. The uptake was most evident in the craniotomy margin area, and mild uptake increase was also observed on the inner surface (brain side) of the bony flap (Fig 2).

\(^{68}\)Ga-RGD uptake was time-dependent and gradually decreased with time after revascularization surgery. UR\textsubscript{max} and V\textsubscript{IU} showed significant correlations with the postoperative time interval between revascularization surgery and \(^{68}\)Ga-RGD PET (P = 0.0015 and 0.0069, respectively; Fig 3). When the linear regression model was applied for extrapolation of V\textsubscript{IU}, the uptake was estimated to normalize (reach zero) at 6.3 mo after surgery.

**Correlation Between Angiogenic Activity and Revascularization Efficacy**

The interval between \(^{68}\)Ga-RGD PET and early postoperative perfusion SPECT was 17±20 d (range, 0–54 d). On \(^{68}\)Ga-RGD PET, UVP was significantly correlated with ΔP\textsubscript{Bas} and ΔP\textsubscript{Str}, with negative correlation coefficients (r = −0.7189, P = 0.0084, and r = −0.5960, P = 0.0409, respectively; Figs. 4A and 4B). UR\textsubscript{max} and V\textsubscript{IU} showed a tendency of correlation with ΔP\textsubscript{Bas} and ΔP\textsubscript{Str}, although the correlations were not statistically significant (P = 0.0555 and 0.0706, respectively; Supplemental Fig 1). When the patients were classified by ΔP\textsubscript{Str} (above and below the median value), UVP was significantly lower in the group with high ΔP\textsubscript{Str} (P = 0.0390, Table 1). When the correlation between postoperative time interval and ΔP\textsubscript{Str} was assessed for consideration of the influence of time, no significant correlation existed between them (r = 0.4294, P = 0.1637; Fig 4C). Additionally, similar correlation patterns were observed between \(^{68}\)Ga-RGD uptake and ΔP\textsubscript{Bas} or

**Statistical Analysis**

All results were expressed as mean ± SD. Comparison of 2 group values was performed using the Student t test, and correlation between 2 parameters was evaluated using the Pearson correlation test. P values of less than 0.05 were regarded significant.

Statistical analyses were performed using a commercial software package (MedCalc, version 9.5.0.0; MedCalc Software).

**RESULTS**

**Patients**

Twelve patients were included in this study (9 males and 3 females; mean age ± SD, 7.9±3.7 y; range, 2–15 y). The patients preoperatively presented with symptoms of transient ischemic attack (n = 11), headache (n = 5), and seizure (n = 1). Four patients had preoperative cerebral infarction in the frontal or occipital area. Revascularization surgery was performed using encephaloduroarteriosynangiosis for the right temporoparietal area and left temporoparietal area in 6 patients each. The frontal area was also revascularized using encephalodurolesionsynangiosis in 7 patients, and the occipital area was revascularized using encephaloduroarteriosynangiosis in 2 patients. All patients underwent a second revascularization surgery for the contralateral side 4.0±1.2 mo (range, 2.4–5.8 mo) after the first surgery. Three patients exhibited postoperative complications of focal infarction or a small amount of subdural hemorrhage (in the right anterior frontal lobe in 2 patients, and in the left high frontal lobe in 1 patient). In the follow-up after revascularization surgery, no patient presented with overt ischemic symptoms related to the operated area.

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Fig. 6, although the correlation was not statistically significant. Representative cases with high and low $^{68}$Ga-RGD uptake are shown in Figure 5 (also in Supplemental Fig. 2).

On comparing $^{68}$Ga-RGD uptake and long-term perfusion change, UVP also showed a tendency of correlation with $\Delta P_{\text{Str}}$ (A and B), with statistical significances. When patients were classified into groups of relatively shorter ($\bullet$) and longer (○) postoperative time interval, similar correlation patterns were observed in both groups. $\Delta P_{\text{Str}}$ had no significant correlation with time interval (C).

$\Delta P_{\text{Str}}$, in both groups when the patients were grouped by the postoperative time interval (above and below the median value; Figs. 4A and 4B). Representative cases with high and low $^{68}$Ga-RGD uptake are shown in Figure 5 (also in Supplemental Fig. 2).

On comparing $^{68}$Ga-RGD uptake and long-term perfusion change, UVP also showed a tendency of correlation with $\Delta P_{\text{Str}}$ (Fig. 6) and $\Delta P_{\text{Str}}$ (Fig. 6), although the correlation was not statistically significant ($r = -0.5447$, $P = 0.0832$, and $r = -0.5010$, $P = 0.1165$, respectively).

**DISCUSSION**

Angiogenesis and collateral vessel formation is the key mechanism in indirect revascularization surgery for MMD. In the present study, we evaluated angiogenic activity using in vivo angiogenesis PET in revascularization treatment. With the use of $^{68}$Ga-RGD PET, the pattern of angiogenic activation was visualized, and angiogenic activity was shown to correlate with the efficacy of revascularization.

In indirect revascularization surgery for MMD, vasculature-rich tissue is placed on the surface of the ischemic brain. Angiogenesis is induced in association with wound-healing and the repair process (17), which is presumably related to the underlying high levels of various growth factors (18). Multiple cranial burr-hole surgery is also performed as an effective adjunct using the same mechanism (19). The efficacy of these treatments is currently assessed by the final outcome of perfusion improvement or new vessel formation using perfusion SPECT, cerebral angiography, or MR imaging (6). However, direct evaluation of angiogenesis would be beneficial for investigating the mechanism of the revascularization process and improving the treatment methods. Recently, radiolabeled RGD peptides were developed for in vivo PET imaging of angiogenesis (8,9,11). Among them, $^{68}$Ga-labeled RGD has good accessibility because $^{68}$Ga can be obtained easily from a generator system, and it has been used for evaluating angiogenic activity in cancer (11), ischemic heart disease (13), and cerebral infarction (12).

In the present study, $^{68}$Ga-RGD PET was performed in indirect revascularization surgery for MMD and exhibited increased activity mainly around the bony flap including adjacent tissue such as galea. The bony flap that covers the operation field is a part of the red marrow–rich skull, and it is in the same vascular territory as that of the revascularized area. Thus, it is speculated that angiogenic activity of the bony flap and adjacent tissue closely correlates with angiogenic activity of the revascularized area. In accordance with this speculation, it has been reported that larger craniotomy is related to better prognosis (20).

It is well known that the uptake of RGD peptides is also enhanced by inflammation (21), because integrin $\alpha_v \beta_3$ is expressed not only on endothelial cells of newly formed vessels but also on activated immune cells such as macrophages (22). In the present study, $^{68}$Ga-RGD uptake possibly correlated with postoperative inflammation, although we performed $^{68}$Ga-RGD PET more than 2 mo after surgery to remove the effect of acute inflammation. The inverse linear correlation between postoperative time interval and $^{68}$Ga-RGD uptake may have been partially influenced by possible inflammation. However, inflammation is also one of the main processes involved in angiogenesis. In an animal experiment, a fibrous coat enriched with immune cells such as macrophages was the initial pathologic finding in indirect revascularization surgery (17). Also in atherosclerosis, activated monocytes are intimately involved in angiogenesis (23). Thus, increased $^{68}$Ga-RGD uptake was analyzed without strict discrimination between inflammation and angiogenesis in this study.

With regard to the time course of angiogenic activity, it was estimated to be active until approximately 6 mo from the extrapolation of $V_{\text{HU}}$. Previous studies have reported that new activity correlated with perfusion improvement and $^{68}$Ga-RGD uptake (A and B), and correlation between postoperative time interval and perfusion improvement (C). UVP was inversely correlated with $\Delta P_{\text{Bas}}$ (A) and $\Delta P_{\text{Str}}$ (B), with statistical significances. When patients were classified into groups of relatively shorter ($\bullet$) and longer (○) postoperative time interval, similar correlation patterns were observed in both groups. $\Delta P_{\text{Str}}$ had no significant correlation with time interval (C).

**TABLE 1**

<table>
<thead>
<tr>
<th>Image index</th>
<th>High $\Delta P_{\text{Str}}$ ($n = 6$)</th>
<th>Low $\Delta P_{\text{Str}}$ ($n = 6$)</th>
<th>$P$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\Delta P_{\text{Str}}$</td>
<td>19.4 ± 6.8</td>
<td>1.7 ± 5.2</td>
<td></td>
</tr>
<tr>
<td>$U_{\text{HU}}$</td>
<td>2.71 ± 0.28</td>
<td>3.20 ± 0.58</td>
<td>0.0897</td>
</tr>
<tr>
<td>$V_{\text{HU}}$</td>
<td>21.9 ± 15.0</td>
<td>41.2 ± 15.5</td>
<td>0.0530</td>
</tr>
<tr>
<td>UVP</td>
<td>40.5 ± 27.1</td>
<td>84.1 ± 35.9</td>
<td>0.0390</td>
</tr>
</tbody>
</table>

Data are mean ± SD.
vessels are observed at 3–4 mo after revascularization surgery (24). However, our results suggest that some delayed angiogenesis may occur by 6 mo after surgery. Accordingly, P$_{_{\text{Bas}}}$ and P$_{_{\text{Str}}}$ were slightly higher on long-term follow-up SPECT than on immediate postoperative SPECT. The actual time course of angiogenic activity can be assessed by serial follow-up study using noninvasive $^{68}$Ga-RGD PET, and such repetitive evaluation of angiogenic activity would be available within a tolerable range of radiation exposure (25).

One of the intriguing results of our study is that postoperative angiogenic activity was inversely correlated with perfusion improvement. It did not appear to be affected by the time course of angiogenic activity because perfusion improvement did not correlate with the postoperative time interval. In addition, this correlation pattern was observed in both groups when the patients were grouped by the postoperative time interval (Fig. 4). The inverse correlation suggests that the prolonged angiogenic activity may be a result of functionally insufficient revascularization, because $^{68}$Ga-RGD PET was performed 3.7 ± 1.0 mo after surgery. The angiogenic activity was also weakly correlated with the long-term P$_{_{\text{Bas}}}$ and P$_{_{\text{Str}}}$ improvement despite statistical insignificance (Fig. 6). Thus, $^{68}$Ga-RGD PET could be used as a prognostic factor for the outcome. However, further studies are required for the image pattern and clinical significance of early angiogenic activity after revascularization surgery.

We analyzed volume-related indices of V$_{_{1U}}$ and UVP as well as the maximal uptake of UR$_{_{\text{max}}}$ on $^{68}$Ga-RGD PET. In molecular imaging methods, the overall presence, expression, or activation of the imaging target should be quantified. However, the target region could not be specifically defined and we measured the entire volume that showed uptake above the threshold. Additionally, UVP was calculated to consider both degree and volume of uptake. This approach is widely used in oncologic PET studies to quantify the overall metabolic burden of tumors (26,27). In the present study, UVP that reflects the overall angiogenic activity showed the most significant correlation with P$_{_{\text{Bas}}}$ and P$_{_{\text{Str}}}$ improvement among all the indices. Thus, UVP may be regarded as the most sensitive index to measure angiogenic activity on $^{68}$Ga-RGD PET.

In brain perfusion SPECT, SPAM analysis has been used as an objective and effective method to measure regional perfusion, with an ethnically matched brain template and regional map (16). It may be a limitation of our study that we applied the brain template and probabilistic map obtained from adult brain data to pediatric patients. However, the probabilistic brain map has been applied to pediatric patients without significant errors (28,29). In addition, the VOI of the MCA territory is relatively large; thus, the results would not have been significantly affected by possible minor missegmentation.

There are limitations that should be considered in the present study. First, the case number was small, although we observed statistically significant results. However, multivariate analyses that could show the specific influences of time interval and perfusion improvement on $^{68}$Ga-RGD uptake were unavailable in this study because of the small case number. Second, the cerebellum was used as a reference tissue for perfusion normalization. It should be considered that the cerebellar perfusion may be affected by crossed cerebellar diaschisis, although the effect is deemed to be minimized with acetazolamide-stress. Additionally, $^{68}$Ga-RGD PET was performed 1 time at relatively variable time points after the acute phase. These were inevitable because
clinical conditions and radiation safety should have been considered for pediatric patients, and serial image acquisition is limited. More sophisticated designs are required in further studies in which angiogenesis imaging is used in pediatric MMD patients.

CONCLUSION

In the present study, 68Ga-RGD PET was successfully used for the assessment of angiogenic activity in indirect revascularization surgery for MMD. Angiogenic activity gradually decreases in a time-dependent manner and was estimated to normalize at approximately 6 mo after surgery. Angiogenic activation measured at approximately 3.7 mo after surgery was inversely correlated with perfusion improvement. The assessment of angiogenic activity using 68Ga-RGD PET is expected to be effective for evaluating the revascularization mechanism or the efficacy of new treatment.

DISCLOSURE

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