Management of intracranial aneurysm in children: clipped and coiled

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Abstract

Objectives The objective was to describe the clinical features of intracranial aneurysm in children and evaluate the efficacy and safety of different treatment approaches for children with intracranial aneurysms.

Materials and methods Clinical data of 13 child patients with intracranial aneurysms that were diagnosed and treated in our hospital in the past 16 years were reviewed. Long-term follow-up data were also collected and reported.

Results Direct surgical treatment was performed on eight out of all 13 patients; among them, seven achieved good recovery and one died. The remaining five patients were treated with Guglielmi detachable coils (GDC). The aneurysm was successfully sealed off in one case and achieved good postoperative recovery. Residual aneurysm was detected in two cases; a second resection surgery was performed on one patient who died after the operation and on another with neck occlusion who recovered well. Two cases experienced aneurysm recanalization after intervention therapy; among them, one was treated with a second embolotherapy but had recanalization once again and was finally cured by aneurysm resection surgery.

Conclusion Aneurysmal surgery is proved to be a safe and effective treatment for children with intracranial aneurysm. Although requiring a long-term follow-up period to determine the clinical outcome, GDC is also shown to be safe for a child having acute bleeding. Once a residual aneurysm or recanalization occurs, microsurgery is recommended.

Keywords Intracranial aneurysm · Guglielmi detachable coils

Introduction

In child population, intracranial aneurysm has a very low incidence and the clinical manifestation is usually atypical. Additionally, the location and the size of intracranial aneurysm in children are often different from those of adult patients [1–8, 17]. In this paper, we reviewed the clinical data of 13 child intracranial aneurysm patients that were treated in our hospital between January 1990 and December 2006.

Materials and methods

Clinical data

Patients A total of 13 child patients (nine males and four females) in the age range of 3–15 years old, five cases ranging from 3 to 6 years old and eight cases ranging from 6 to 15 years old, respectively, were included in the current study.

Clinical symptoms Six were acute cases with chief complaint of headache and/or dizziness. Their clinical manifes-
tations suggested intraventricular hemorrhage. Younger children could not make clear chief complaints but were dysphoric and irritable, whereas the most frequent chief complaint of older children was persistent and increasingly severe headache. Other symptoms were most frequent chief complaint of persistent headache (four cases), vomiting (six cases), nausea (two cases), neck stiffness and neck pain (five cases), fatigue (one case), asthenia (two cases), muscular spasm of limbs (four cases), and low-grade fever (three cases). The remaining seven were chronic cases. Five of them had symptoms of monolateral asthenia, stagger, or holding immobilizing disability; one case had barylalia; one case had reduction in visual capacity with ptosis; and two cases experienced epileptic seizure. All 13 patients did not have an identified head injury or infection history. No family history of intracranial aneurysm or intracranial arteriovenous malformation (AVM) was reported.

Main physical signs Out of all 13 patients, six had neck rigidity and five had meningsismus. One case reported monolateral oculomotor paralysis. Six patients had conscious disturbance of different degrees; among them, one was in a state of unconsciousness. Monolateral asthenia with Babinski’s sign was found in seven patients. In the scale of Hunt and Hess, one patient was grade I, three were grade II, and two were grade IV.

Auxiliary examination

Lumbar puncture Bloody lumbar puncture was reported in five patients.

Imageology Cranial computed tomography (CT) scan was performed on all patients. Subarachnoid hemorrhage (SAH) was detected in five cases and intracranial hematoma was detected in one case. Magnetic resonance angiography (MRA) examination was performed on one case and an intracranial arterial deformation was detected (Fig. 1). Intracranial anticipate pathological changes were revealed in seven cases. Magnetic resonance imaging (MRI) examination was performed on six patients; among them, four patients were suggested to have intracranial aneurysms. Intracranial giant aneurysms were confirmed in four patients by CT angiography (CTA) examination.

Digital substract angiography (DSA) or 3D-DSA DSA was conducted in 13 patients within 1–3 weeks after onset of the
Among them, five cases were examined within 1 week and eight were examined within 2–3 weeks. All the patients were diagnosed as having an intracranial aneurysm. DSA or 3D-DSA also revealed abnormal bulking of arterial walls (Figs. 2, 3, 4, 5, 6, and 7). Thromb formation, adjacent artery angiospasm or steno sis and occlusion, and formation of compensatory circulation were observed in some of these patients. Complication of AVM was confirmed in one case by postoperative pathological examination.

**Treatment**

After the identification of diagnosis and obtaining of informed consent from the patients’ family members, neurosurgeons and neurointerventional therapists cooperated in conducting the treatment programs. Surgical approaches included aneurysm neck occlusion, aneurysm neck ligation, occlusion of the carrying artery of aneurysm, aneurysm resection surgery, aneurysm wrapping with muscle, and carotid artery ligation. Guglielmi detachable coil (GDC) embolization was conducted as the following procedure: After heparinization, a Fasguide guiding catheter was placed in the patient’s femoral artery. An Excel 14-microcatheter/Mizzen 10 microguide wire system (Boston Scientific/Target) was advanced through the guiding catheter and the tip of the microcatheter was positioned at the center of the aneurysm. All aneurysms were densely packed using GDC coils selected according to the size of the aneurysm. Angiograms were repeatedly performed during and after operation to obtain information about the positions of the coils and the degree of embolization (Figs. 8, 9, and 10).

**Results**

Follow-up visits have been conducted in all patients for more than 1 year. The Glasgow Outcome Scale standard was used for the evaluation of curative effect. Postoperative DSA was performed on three cases for aneurysm neck occlusion and three cases for aneurysm resection with aneurysm neck occlusion among the eight patients undergoing direct surgical treatment, which demonstrated a complete disappearance of the aneurysm. One patient treated with muscle wrapping and right internal carotid artery (ICA) ligation also recovered well and postoperative DSA revealed diminution in aneurysm size. Hematoma removal was performed on one case with aneurysm carrying artery occlusion and the patient died after the operation. Pathological examination demonstrated a complication of AVMs in this case. Among the five patients treated with GDC embolization, residual aneurysm was found in two cases; isolated aneurysm resection surgery.
was performed on one patient who died after the operation and aneurysm resection surgery was performed on another with aneurysm neck occlusion and the patient recovered well. The aneurysms were successfully sealed off in three cases with a 100% degree of embolization. Follow-up visit at postoperative 1 year showed good recovery was achieved in one case but aneurysm recanalization occurred in the other two cases. A second embolization treatment was performed on one of the two cases but postoperative residual aneurysm was detected. Then the two cases received aneurysm resection surgeries and finally achieved satisfactory recovery. DSA examination demonstrated the complete disappearance of the aneurysm. All children with good recovery were able to undertake normal activities of daily living (Table 1).

Discussion

A child patient with intracranial aneurysm is relatively rare in clinical practice, usually having an incidence of 1–2% in all intracranial aneurysm cases [1, 4, 5]. The 13 child cases reported in the present study were about 1.4% of all intracranial aneurysm cases treated in our hospital during the same period. In 1849, Hare firstly reported an adolescent case with spontaneous SAH caused by a rupture of the posterior communicating aneurysm. Then in 1871, Eppinger reported a boy 15 years of age that fell ill during exercises and died 3 days later. Autopsy revealed a ruptured aneurysm of the anterior cerebral artery complicated with aortic stenosis. Therefore, although not so common in childhood and adolescence, SAH is a severe disease and should be managed carefully in clinical practice. In recent years, with the development of DSA technique and the clinical application of CTA and MRA techniques, the detection rate and also our understanding of aneurysm in children have been greatly improved.

In adult patients, aneurysm is the main cause of SAH, having a high death rate of 14% to 34% within 48 h after onset. In contrast to this, childhood aneurysm usually is symptomatic and often detected in DSA examination or even autopsy. Aneurysm has a higher incidence in boys than in girls and is mostly located at the furcation of ICA or vertebra-basilar artery posterior circulation. The incidence of large or giant aneurysms in child patients is higher than in adult patients whereas occurrence of aneurysm rupture is less, thus leading to a relative low death rate within 48 h after onset of the disease [1, 5, 7, 8]. Out of the nine male cases and four female cases reviewed in this study, nine (69%) had large or giant aneurysms. Among them, six

![Fig. 5 DSA and 3D-DSA obtained before second embolization (August 7, 2003). Left: DSA showing aneurysm recanalization and persisting overexpansion, oblique view. Right: 3D-DSA image of the left ICA demonstrating the presence of the aneurysm of the left middle cerebral artery and persisting overexpansion](image1)

![Fig. 6 DSA and 3D-DSA obtained after second embolization (August 7, 2003). Left: DSA anteroposterior view showing residual aneurysm of the left middle cerebral artery. Right: 3D-DSA image of the left ICA revealing the residual aneurysm of the left middle cerebral artery](image2)
(46%) patients initially suffered from hemorrhage and seven (54%) had intracranial anticipate pathological changes. Evidence shows that multiple aneurysms seldom occur in child patients. The previous study indicated that the incidence of multiple aneurysms is in the range of 6–8% for child patients but was about 29% for adult patients. Taken together, these data suggest that there may be a difference in the pathological formation procedure of aneurysm between children and adults [1–6].

It has been confirmed that many factors including insufficiency or breakage of endovascular elastic layer and/or smooth muscle layer, either congenital structural defect or posterior degeneration (i.e., atherosclerosis, hypertension, or hemodynamic alterations), are implicated in the genesis of cystic intracranial aneurysm. The similarity of aneurysmal histology between children and adults indicates that the development of childhood aneurysm also involves these factors. Congenital factor is more prevalent than the others [1, 5, 7–9] because other congenital diseases or vascular malformations in brain or peripheral tissues are often complicated in child patients. Stehbens [9] considered that the main cause of aneurysm is posterior degenerative changes; however, this is not the case in children because inflammatory changes or atherosclerosis is seldom observed in children and this may have been the explanation of the relative low incidence of childhood aneurysm. Some other studies also presumed that insufficiency of types III and IV collagen may be concerned with the development of childhood aneurysm [10]. Some viral or bacterial infections had also been confirmed in childhood aneurysms. All factors described above constitute the distinct characteristics of childhood aneurysm in size, location, and complications [1–11].

Due to the inadequate understanding of courses of childhood aneurysm rupture with profuse bleeding, the treatment selection was mainly based on knowledge coming from adult patients [7, 8]. Direct occlusion of the aneurysm neck is the optimal treatment prescription. It is noteworthy that childhood aneurysm of posterior circulation is often giant and fusiform in shape, making it life-threatening to occlude the aneurysm directly. The best treatment approach would be gradually occluding the aneurysm carrying the vertebral artery. In Amacher and Drake’s opinion [12], children in general have good tolerance to the operative procedure. Previous studies have demonstrated that children show good tolerance to the SAH with lower rates of mortality and deformity than adult
patients. Cerebral infarction seldom occurs in child patients, even for those having severe cerebral angiospasm, whereas in adult patients the incidence of angiospasm-induced cerebral infarction is about 75%. As reported by Amacher and Drake, the overall post-surgery mortality of child patients of aneurysm, including those with giant aneurysm, is less than 5%.

GDC, which was designed and firstly used by Guglielmi (an Italian neurointerventional therapist), is a novel technique of aneurysm embolization. The invention and application of GDC greatly improved the treatment of intracranial aneurysm. It can occlude the aneurysm and ensure the unobstruction of the carrying artery simultaneously and finally achieve the outcome proximal to that of aneurysm occlusion surgery. According to Sluzewski's study [13], during the GDC embolization of intracranial aneurysm, the occurrence rate of aneurysm rupture and death rate were 2.5% and 1%, respectively. Therefore, GDC embolization has been considered as a less invasive treatment and is becoming more acceptable than other approaches. Currently, GDC embolization has been commonly used and is a main approach in the treatment of adult patients with intracranial aneurysm. However, enough data about its safety and efficacy in child patients are still not available. Yu and his colleagues have reported successful application of GDC in the treatment of three children with intracranial aneurysms [14]. All three cases (two for posterior communicating aneurysms and one for anterior communicating aneurysms) had acute SAH caused by rupture of the aneurysm. The diameters of the aneurysm necks were in the range of 2.2–2.4 mm and the largest diameter of the aneurysm was less than 5.0 mm. In the scale of Hunt and Hess, one patient was grade I, three were grade II, and two were grade IV. A 100% degree of embolization was achieved and clinical symptoms disappeared completely after the operation without reoccurrence of bleeding or any other complications. Follow-up visits have been carried out for all three patients for more than half a year (including two cases which had been followed up for more than 1 year). They demonstrated a good recovery and could undertake normal daily activity. DSA examination of two cases did not reveal residual aneurysm or recanalization. We consider that GDC embolization is safe for the treatment of child patients with intracranial aneurysms. It provides an alternative approach especially for those with acute rupture of aneurysm.

In our study, GDC embolization treatment was performed on a total of five child patients. The aneurysm was successfully sealed off in one male patient with acute aneurysm rupture (scale of Hunt and Hess, grade II). DSA
examination revealed a posterior communicating aneurysm on the right side; the aneurysm neck had a diameter of 2.4 mm and the aneurysm size was 5.0×3.6 mm. A 100% degree of embolization was obtained. DSA recheck did not find any residual aneurysm or recanalization. In one case with a giant vertebral aneurysm, post-embolizing residual aneurysm was detected. Isolated aneurysm resection surgery was then performed on the patient who died after the operation. Post-embolizing residual aneurysm was also detected in one case with a giant posterior cerebral aneurysm on the left side. For the remaining two cases, the first embolization surgery succeeded but follow-up visits revealed aneurysm recanalization. A second embolization treatment was performed on one of the two cases but postoperative residual aneurysm was reported again. Then, the two cases received aneurysm resection surgeries. DSA examination demonstrated that the aneurysm disappeared completely and good recovery was achieved.

Hayakawa [15] has reported the application of GDC in the treatment of 455 cases of intracranial aneurysms. The occurrence of postoperative residual aneurysm stem was 39% (178 cases). Of all patients with narrow stem small aneurysms, gradual thromb formation and aneurysm recanalization were reported in six cases (25%) and ten cases (42%), respectively. In patients with large and giant aneurysms, the occurrence of recanalization was reported to be 87% (13 cases) and 90% (nine cases), respectively. Overall occurrence of gradual thromb formation and recanalization was respectively 25% (18 cases) and 49% (36 cases). Therefore, postoperative recanalization of residual aneurysm, especially for patients with broad stem large or giant aneurysms, was a very common event. As that child patient is more likely to have large or giant aneurysms, we consider GDC embolization to be more suitable and safe for children with small- or medium-sized aneurysms with acute rupture. However, to evaluate the long-term outcome of GDC treatment on children with large or giant aneurysms requires a longer follow-up period. Ausman [16] considered that occlusion of aneurysm by using a clip could help to draw close the aneurysm neck elastic layer which is the most burdening structure of the arterial wall, whereas embolization treatment could not do this at all. Therefore, the selection of treatment approach for child patients with large or giant aneurysms relies on the close cooperation of neurosurgeons and neurointerventional therapists. Follow-up DSA examination shall be enhanced for the cases having residual aneurysm necks. Once the residual aneurysm increases or recanalization occurs, microsurgical treatment is recommended.

| Patient’s code | Aneurysm position | Aneurysm size | Clinical signs | Management | Outcome |
|----------------|-------------------|---------------|---------------|------------|---------|
| 1              | LACoA             | Small         | SAH           | Aneurysm neck occlusion | Excellent |
| 2              | LICAC2            | Small         | SAH           | Aneurysm neck occlusion | Excellent |
| 3              | LPCoA             | Medium        | SAH           | Aneurysm neck occlusion | Excellent |
| 4              | LMCA              | Large         | SAH           | Aneurysm resection with aneurysm neck occlusion | Excellent |
| 5              | RMCA              | Large         | SAH           | Aneurysm resection with aneurysm neck occlusion | Excellent |
| 6              | RPICA             | Large         | SAH           | Aneurysm resection with aneurysm neck occlusion | Excellent |
| 7              | RICAC2            | Large         |SAH           | Muscle wrapping, right ICA ligation | Good |
| 8              | LMCA and AVMS     | Giant         | Intracranial hematoma | Hematoma removal, aneurysm carrying artery occlusion | Dead |
| 9              | RPCoA             | Medium        | SAH           | Interventional embolization | Excellent |
| 10             | LPCA              | Large         | SAH           | Residual aneurysm after interventional embolization, aneurysm resection with aneurysm neck occlusion | Good |
| 11             | LMCA              | Giant         | SAH           | Second interventional therapy, aneurysm resection with aneurysm neck occlusion | Good |
| 12             | RMCA              | Large         |SAH           | Recanalization after interventional therapy, aneurysm resection with aneurysm neck occlusion | Good |
| 13             | RVA                | Giant         |SAH           | Residual aneurysm after interventional embolization, isolated aneurysm resection surgery | Dead |
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