Factors associated with epileptic seizure of cavernous malformations in the central nervous system in West China

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ABSTRACT
Objective: To explore the factors associated with preoperative and postoperative epileptic seizure in patients with cavernous malformations (CMs).
Methods: A total of 52 consecutive patients from January 2009 to June 2011 who underwent surgical treatment in West China Hospital of Sichuan University due to CMs and confirmed by histopathology were retrospectively reviewed. Patients were divided into two groups (epilepsy-group and non-epilepsy group) according to clinical presentation. Other clinical data, treatment procedure, and follow-up information were collected. Engel classification was used to evaluate seizure outcome.
Results: Low birth weight, temporal lobe involvement and cortical lesion showed significant difference between two groups (p=0.017, 0.003 and 0.025 respectively). Cortical lesion highly increased risk for preoperative epileptic seizure (OR=10.48; 95% CI 1.61-68.23). After a mean follow-up of 2.1 years, 77.8% of epileptic patients achieved Engel class I. Temporal lobe involvement, lesion size < 2.5 cm and surgery within one year of symptom onset were found associated with better seizure outcome (p=0.016, 0.012 and 0.050). Temporal lobe involvement significantly decreased the risk for postoperative epileptic seizure (OR=0.038; 95% CI 0.002-0.833). Application of ECoG made no significant difference to seizure outcome (p=0.430). Most patients need continuing medication therapy after surgery.
Conclusion: Surgical treatment of patient with CMs is satisfactory in most cases and temporal lobe involvement usually predict favourable postoperative seizure outcome whether under the monitoring of ECoG or not. Thus, epileptic patients with CMs should be considered for surgical treatment especially when cortical brain layer or temporal lobe was involved.

KEY WORD: Cerebral cavernous malformation, Epilepsy, Neurosurgery, Risk factors.

INTRODUCTION

Cavernous malformations (CMs) are common vascular malformations in the central nervous system which usually consist of enlarged thin-wall blood vessels without neural tissue. The prevalence of CMs in population is around 0.4-0.8% and seizures is common clinical presentation (23-50% of cases).¹ Other main symptoms include headache (6-52%), neurological deficits (20-40%) and hemorrhage (9-56%).¹ There are a number of studies discussing clinical manifestation, diagnostic procedure, treatment and surgical outcome of CMs.²-⁴
However, these studies occasionally distinguished arteriovenous malformations (AVMs) and CMs and usually discussed about the global risk brought by CMs.\(^3\)\(^5\) Few studies considered about the specific risk factors for developing epileptic seizure in patients with CMs especially preoperative. Age, cortical site, family history, pre-, peri- and postnatal factors were reported as independent predictors of symptomatic seizures.\(^5\) Other factors such as alcohol and tobacco addiction may increase the risk of epileptic seizure as well according to clinical practice.\(^5\) Factors influencing postoperative seizure occurrence are still in dispute. Thus, in order to explore the factors associated with preoperative and postoperative epileptic seizure in patients with CMs, we retrospectively analyzed the clinical data of 52 consecutive patients diagnosed with CMs and surgically treated in West China Hospital of Sichuan University during the period from January 2009 to June 2011.

**METHODS**

A total of 52 consecutive patients diagnosed with CMs who underwent surgical treatment in Department of Neurosurgery, West China Hospital of Sichuan University from January 2009 to June 2011 were retrospectively studied. Inclusion criteria were patients with CMs confirmed by pathology after surgery. Patients who were clinically diagnosed with CMs on the evidence of neuroimaging but confirmed as other type of vascular malformations such as AVMs were excluded.

We assigned patients with preoperative epilepsy into epilepsy group (E-Group) and those with headache, focal neurological deficits (FND) or intracranial hemorrhage (ICH) into non-epilepsy group (NE-Group). We defined epilepsy as presentation of seizure not symptomatic of ICH caused by CMs at least two times, and ICH as symptomatic events associated with evidence of intracranial blood on neuroimaging and FND as symptoms of neurologic dysfunction related to the anatomical site of CMs but without presence of epileptic seizure. We grouped CMs locations into 4 types: supratentorial lobar (frontal, parietal, temporal, occipital), supratentorial deep area (limbic, thalamus, hypothalamus, callosal, basal ganglia, and choroidal), infratentorial (brain-stem and cerebellum), spine, or multiple (if the patient had at least two CMs). The size of CMs was measured as the maximum diameter of the CM.

The following information was collected during the review of medical records: gender, age, disease course, neurological manifestation at admission (seizure, headache, FND or ICH), diagnostic procedures, neuroimage character, anatomic location, size of CMs, surgical procedures, histopathology results. In order to explore the risk factors of seizure in patients with CMs, we also spared no efforts to collect following data: family history of epileptic seizure, pre- and peri- and postnatal factors, history of tobacco and alcohol use.

Follow-up information was obtained by outpatient and telephone interviews. The surgical outcome of patients with epilepsy was classified according to the Engel’s classification into class I, completely seizure free, seizure free for at least two years, auras only, or convulsions with drug withdrawal only; class II, rare seizure (≤2 seizures per year); class III, worthwhile improvement; class IV, no significant improvement or worse. For patients without seizure, the surgical outcome was classified into following four classes according to patients’ subjective feelings: class I, significant improvement; class II, improvement; class III, no significant improvement; class IV, worsen.

The analysis of risk factors for epilepsy in patients with CMs was estimated by calculating odds ratios (OR), with 95% confidence intervals (95% CI). The chi-square test and two-tailed t-test were performed when appropriate. The significance level was set at 0.05. Variables, which would potentially predict increasing risk for epilepsy, were then evaluated by logistic regression model.

**RESULTS**

The study consisted of 52 patients: 31 (59.6%, 31/52) in E-Group and 21 (40.4%, 21/52) in NE-Group (Table-I). Of the 31 patients with epilepsy, the predominant seizure type was secondarily generalized seizure (74.2%, 22/31), followed by complex partial seizure (29.0%, 9/31), simple partial

| Table-I: General information of patients with CMs (52 cases). |
|-----------------------------|-----------------------------|-----------------------------|
| E-Group | NE-Group | p value |
|---|---|---|
| Number | 31 | 21 | - |
| Gender (M:F) | 2.44 | 1.10 | 0.242 |
| Male | 22 | 11 | - |
| Female | 9 | 10 | - |
| Age (y, ± SD) | 36.2 ± 13.4 | 36.3 ± 12.1 | 0.963 |
| Mean age at onset, y, ± SD | 31.3 ± 13.4 | 34.0 ± 13.8 | 0.473 |
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Factors associated with preoperative epileptic seizure: In our study, the factors seen significantly associated with preoperative epilepsy were low birth weight (p=0.017), temporal lobe involvement (p=0.003) and cortical lesion (p=0.025) (Table-II). No significant association was found between other pre-, peri- and post-natal factors (complication of pregnancy, low gestational age, complication of delivery, febrile seizure, psychomotor retardation, history of encephalitis) and preoperative epilepsy separately (shown in Table-II). However, there were more patients with at least one of these risk factors in E-Group than NE-Group (14 vs. 3) (p=0.020). Mean age at onset, family history of epilepsy, history of injury, operation, tobacco and alcohol using were not seen associated with an increased risk for epilepsy (shown in Table-II). In the multivariate analysis, the cortical lesion still showed a high risk for epilepsy (OR=10.48; 95% CI 1.61-68.23) while the pre- and peri- and postnatal factors (OR=2.85; 95% CI 0.17-48.86), low birth weight (OR=4.99; 95% CI 0.14-176.31), involvement of temporal lesion (OR=4.63; 95% CI 0.72-29.86) were no longer significant (Table-III).

Table-II: Factors for preoperative seizure due to CMs in central nervous system (52 cases).

| Features                                      | E-Group (n=31) | NE-Group (n=21) | p value |
|-----------------------------------------------|----------------|-----------------|--------|
| Mean age at onset                             | 31.3           | 34.0            | 0.473  |
| Family history of epilepsy                    | 1              | 0               | 0.406  |
| Pre-, peri- and post-natal factors            | 14             | 3               | 0.020  |
| Complication of pregnancy                     | 1              | 0               | 0.406  |
| Low gestational age                           | 3              | 0               | 0.142  |
| Low birth weight                              | 10             | 1               | 0.017  |
| Complication of delivery                      | 1              | 1               | 0.798  |
| Febrile seizure                               | 3              | 1               | 0.514  |
| Psychomotor retardation                       | 0              | 0               | -      |
| History of encephalitis                       | 1              | 0               | 0.406  |
| History of head trauma                        | 6              | 1               | 0.130  |
| History of operation                          | 3              | 4               | 0.331  |
| History of tobacco addiction                  | 11             | 7               | 0.873  |
| History of alcohol addiction                  | 10             | 9               | 0.436  |
| Multiple vs. single lesion                    | 2              | 0               | 0.235  |
| Supratentorial vs. infratentorial              | 31             | 12              | -      |
| Cortical vs. subcortical                      | 24             | 5               | 0.025  |
| Frontal lobe                                  | 7              | 5               | 0.918  |
| Parietal lobe                                 | 7              | 2               | 0.222  |
| Temporal lobe                                 | 17             | 3               | 0.003  |
| deep brain area                               | 0              | 3               | 0.030  |
| Side (right vs. left)                         | 14/17          | 6/15            | 0.599  |
| Size (diameter ≥ 2.5 cm)                      | 12             | 10              | 0.084  |

Table-III: Logistic regression analysis of risk factors for seizure due to CMs.

| Risk factors                                      | Odds ratio | 95% CI   | p value |
|---------------------------------------------------|------------|----------|---------|
| Pre-, peri- and post-natal factors                | 2.85       | 0.17-48.86 | 0.470  |
| Low birth weight                                  | 4.99       | 0.14-176.31 | 0.377  |
| Temporal lobe                                     | 4.63       | 0.72-29.86 | 0.107  |
| Cortical involvement                             | 10.48      | 1.61-68.23 | 0.014  |
Factors associated with postoperative seizures:
There were 21 cases that had clearly clarified the data of size of CMs during surgical process. Intraoperative electrocorticography (ECoG) was performed in 19 cases and additional resection was performed as indicated by the ECoG findings. Follow-up information of 46 (88.5%, 46/52) patients was collected while the other 6 patients were lost to follow-up (4 of E-Group and 2 of NE-Group) (Table-IV). The mean follow-up time was 24.8 months (range 14-41 months) in E-Group and 26.6 months (range 14-43 months) in NE-Group. In E-Group, each received regular medication after surgery following doctor’s advice. There were 20 (74.1%, 20/27) received monotherapy while 7 (25.9%, 7/27) duotherapy after surgery. Continued AEDs therapy lasted for less than 3 months in 7 out of 20 patients on monotherapy while the other 20 patients went on medication for a longer period of time. At our final follow-up, 19 (70.4%, 19/27) patients had withdrawal medication. Most (77.8%, 21/27) patients were seizure free and classified as Engel class I while the other 6 (22.2%, 6/27) patients still had seizure now or then. In NE-Group, 18 (94.1%, 18/19) patients in total got symptom-free. In our study, no mortality occurred.

Patients with temporal lobe CMs were found more likely to be seizure free after surgery (p=0.016) (Table-V). The maximum diameter of CMs longer than 2.5 cm and disease course longer than one year predicated unfavorable outcome (p=0.012, 0.050). Other factors such as mean age at onset, mean disease course, gender, family history of epilepsy, pre-, peri-, and post-natal factors were not associated with seizure outcome. Moreover, application of ECoG, surgical strategy and postoperative medication therapy did not make significant difference to seizure outcome. Of the 8 cases where no ECoG were applied, 6 (75%, 6/8) were temporal lobe involved. In the multivariate

Table-IV: Postoperative follow-up outcome of 44 patients with CMs.

| Group   | No. of patients | Signs and symptoms, n (%) | Mean follow-up, month, (range) |
|---------|-----------------|---------------------------|-------------------------------|
|         |                 | Engel I or free | Engel II or improvement | Engel III or no change | Engel IV or Worsen |
| E-Group | 27              | 21(77.8)         | 5(18.5)                     | 0                  | 1(3.7)               | 24.8(14-41) |
| NE-Group| 19              | 18(94.7)         | 0                           | 0                  | 1(5.3)               | 26.6(14-43) |
| Total   | 46              | 37(84.1)         | 5(11.4)                     | 0                  | 2(4.5)               | 25.5(14-43) |

Table-V: Factors associated with postoperative seizure of CMs.

| Features                        | Seizure free (n=21) | Not seizure free (n=6) | p value |
|--------------------------------|---------------------|------------------------|---------|
| Gender ration (M:F)             | 3.2                 | 1.0                    | 0.215   |
| Mean age at onset, y            | 30.7                | 31.0                   | 0.976   |
| Disease course (≥1y vs.<1y)     | 12/9                | 6/15                   | 0.050   |
| Family history of epilepsy     | 1                   | 0                      | 0.586   |
| Pre-, peri- and post-natal factors | 11                  | 2                      | 0.410   |
| Multiple vs. single lesions     | 2                   | 0                      | 0.432   |
| Frontal lobe                    | 4                   | 3                      | 0.127   |
| Parietal lobe                   | 3                   | 2                      | 0.289   |
| Temporal lobe                   | 15                  | 1                      | 0.016   |
| Cortical vs. subcortical        | 16                  | 5                      | 0.711   |
| Side (right vs. left)           | 8                   | 3                      | 0.601   |
| Size (diameter ≥2.5 cm vs. not) | 6/15                | 6/0                    | 0.012   |
| Seizure type(generalized vs. partial) | 16/5                | 3/3                    | 0.215   |
| Total vs. subtotal resection    | 20/1                | 5/1                    | 0.326   |
| Perilesional hemosiderin removal | 9                  | 3                      | 0.756   |
| Intraoperative ECoG             | 14                  | 5                      | 0.430   |
| Monotherapy vs. duotherapy      | 15/6                | 5/1                    | 0.557   |
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**DISCUSSION**

Cavernous malformations (CMs) are common vascular malformations in the central nervous system which usually consist of enlarged thin-wall blood vessels without neural tissue. The prevalence of CMs in population is around 0.4-0.8% and epileptic seizures are common clinical patients with seizures caused by CMs usually achieved seizure control by antiepileptic drugs in about 60% while surgical resection seemed to be a superior method with satisfactory outcome (symptom free or Engel I) after surgical in up to 80% of all patients. However, factors associated with surgical outcome remains controversial.11-13 In our study, 77.8% epileptic patients achieved seizure free after surgery. Temporal lobe involvement and maximum diameter of lesion less than 2.5 cm predicted better outcome and this finding was in accordance with those reported by Baumann.11 This might suggest that CMs located in different site may have different type of functional connectivity and involvement with surrounding brain area to generalize epileptic discharge. However, it needs further physiological and functional investigations.

Several studies have showed patients with shorter epileptic duration and lower number of preoperative seizures were associated with better seizure control.14 A lack of seizure control might be due to poor surgical strategy such as subtotal resection or persistence of surrounding hemosiderin deposits.8,11,13 However, several studies had demonstrated pure lesionectomy might eliminate epileptic seizure.14,16 This might attribute to the pathological differentiation characteristic of CMs which are commonly well-circumscribed lesions. It was in accordance with our findings which showed no significant difference between the different surgical strategies. It is still controversial whether the pure lesionectomy is acceptable to achieve satisfactory surgical outcome and further studies with larger sample sizes needed.

Intraoperative ECoG is a commonly used technique in epilepsy surgery to detect epileptic discharge and help identifying the epileptogenic zone. However, due to its own limitations such as short duration, lack of ictal records and anesthesia disturbance, its application in epileptic surgery is controversial. In our study, we found no significant association between application of ECoG and postoperative seizure occurrence as reported before.11,17 In the 8 (25.8%, 8/31) cases who had achieved favorable seizure outcome without applying ECoG, most were temporal lobe involved and this may explain the results. The ratio of ECoG usage is 74.2% (23/31) in our hospital considering epileptic patients due to CMs and it is much lower than those of hospitals in developed countries and areas partly due to financial status of patients.17,18

Continued AEDs treatment after surgery is still controversial. Ferroli et al found that antiepileptic drugs were not necessary in most cases and part of patients with long clinical history of seizures might need continued AEDs.19 In our study, all patients received AEDs postoperatively and there were about 30% patients on medication at final follow up. Parts of patients with CMs might require AEDs after surgery so that it might be reasonable to prescribe medicine for patients especially with high risk for postoperative seizures.

Our study has several limitations. The primary limitation was its retrospective nature which might lead to a possible selection bias. Another limitation was limited sample size. However, as relatively low incidence of CMs with presentation of seizure, it is sometimes reasonable to explore related factors through retrospective study. The follow-up was not long. All these factors should be considered in the future studies.

**CONCLUSION**

Our study suggests that the cortical involvement is the risk factors for preoperative epilepsy in patients with CMs. Surgical treatment of patient with CMs is satisfactory in most cases and temporal lobe involvement usually predict favorable postoperative seizure outcome whether under the monitoring of ECoG or not. Continued AEDs were recommended after successful surgical resection. Thus, epileptic patients with CMs should be considered for surgical treatment especially when cortical brain layer or temporal lobe was involved.

| Risk factors | Odds ratio | 95% CI | pvalue |
|--------------|------------|--------|--------|
| Temporal lobe | 0.040 | 0.002-0.833 | 0.038 |
| Size (diameter ≥2.5 cm) | <0.001 | - | 0.999 |

Table VI: Logistic regression analysis of risk factors for postoperative seizure.
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We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

REFERENCES

1. Batra S, Lin D, Recinos PF, Zhang J, Rigamonti D. Cavernous malformations: natural history, diagnosis and treatment. Nat Rev Neurol. 2009;5(12):659-670.
2. Arita K, Kurisu K, Iida K, Hanaya R, Sugiyama K, Akimitsu T, et al. Surgical treatment for intractable epilepsy caused by cavernous angioma in the temporal lobe of the dominant hemisphere—three case reports. Neurol Med Chir (Tokyo). 2000;40(8):439-445.
3. Josephson CB, Leach JP, Duncan R, Roberts RC, Counsell CE, Al-Shahi Salman R, et al. Seizure risk from cavernous or arteriovenous malformations: prospective population-based study. Neurology. 2011;76(18):1548-1554.
4. Moran NF, Fish DR, Kitchen N, Shorvon S, Kendall BE, Stevens JM. Supratentorial cavernous haemangiomas and epilepsy: a review of the literature and case series. J Neurol Neurosurg Psychiatry. 1999;66(5):561-568.
5. Leone MA, Ivashynka AV, Tonini MC, Bogliun G, Montano V, Ravetti C, et al. Risk factors for a first epileptic seizure symptomatic of brain tumour or brain vascular malformation. A case control study. Swiss Med Wkly. 2011;141:w13155.
6. Del Curling O, Jr., Kelly DL, Jr., Elster AD, Craven TE. An analysis of the natural history of cavernous angiomas. J Neurosurg. 1991;75(5):702-708.
7. Noto S, Fujii M, Akimura T, Imoto H, Nomura S, Kajiwara K, et al. Management of patients with cavernous angiomas presenting epileptic seizures. Surg Neurol. 2005;64(6):495-498, discussion 498-499.
8. Stavroul, Baumgartner C, Frischer JM, Trautnig S, Knop E. Long-term seizure control after resection of supratentorial cavernomas: a retrospective single-center study in 53 patients. Neurosurg. 2008;63(5):888-896; discussion 897.
9. Brunon J, Nuti C. Natural history of cavernomas of the central nervous system. Neurochirurgie. 2007;53(2-3 Pt 2):122-130.
10. Siegel AM, Andermann E, Badhwar A, Rouleau GA, Wofford GL, Andermann F, et al. Anticipation in familial cavernous angioma: a study of 52 families from International Familial Cavernous Angioma Study. IFCAS Group. Lancet. 1998;352(9141):1676-1677.
11. Baumann CR, Acciarri N, Bertalanffy H, Devinsky O, Elger CE, Lo Russo G, et al. Seizure outcome after resection of supratentorial cavernous malformations: a study of 168 patients. Epilepsia. 2007;48(3):559-563.
12. Betej M, Czepko R, Lopatka P, Danilewicz B, Uhl H. Diagnosis and operative treatment cavernous angiomas of the central nervous system. Przegl Lek. 2006;63(2):61-63.
13. Rassi-Neto A, Ribeiro PR, Prates MA, Muszkat M, de Campos CJ, Ferraz FA. Surgical treatment of cerebral vascular pathologies in epileptic patients. Arq Neuropsiquiatr. 1997;55(3A):408-412.
14. Cohen DS, Zubay GP, Goodman RR. Seizure outcome after lesionectomy for cavernous malformations. J Neurosurg. 1995;83(2):237-242.
15. Baumann CR, Schuknecht B, Lo Russo G, Cosso M, Citterio A, Andermann F, et al. Seizure outcome after resection of cavernous malformations is better when surrounding hemosiderin-stained brain also is removed. Epilepsia. 2006;47(3):563-566.
16. Casazza M, Broggi G, Franzini A, Avanzini G, Spreatico R, Bracci M, et al. Supratentorial cavernous angiomas and epileptic seizures: preoperative course and postoperative outcome. Neurosurg. 1996;39(1):26-32; discussion 32-24.
17. Van Gompel JJ, Rubio J, Cascino GD, Worrell GA, Meyer FB. Electrocorticography-guided resection of temporal cavernoma: is electrocorticography warranted and does it alter the surgical approach? J Neurosurg. 2009;110(6):1179-1185.
18. Siegel AM, Roberts DW, Harbaugh RE, Williamson PD. Pure lesionectomy versus tailored epilepsy surgery in treatment of cavernous malformations presenting with epilepsy. Neurosurg Rev. 2001;24(2):80-83.
19. Ferroli P, Casazza M, Marras C, Mendola C, Franzini A, Broggi G. Cerebral cavernomas and seizures: a retrospective study on 163 patients who underwent pure lesionectomy. Neurological Sci. 2006;26(6):390-394.