

Review Article

Perioperative Variables Contributing to the Rupture of Intracranial Aneurysm: An Update

Tumul Chowdhury,¹ Ronald B. Cappellani,¹ Nora Sandu,²
Bernhard Schaller,² and Jayesh Daya¹

¹ Department of Anesthesiology, Health Sciences Center, University of Manitoba, 2nd Floor Harry Medovy House, 671 William Avenue, Winnipeg, MB, Canada R3E 0Z2

² Research, University of Southampton, University Road, Southampton SO17 1 BJ, UK

Correspondence should be addressed to Tumul Chowdhury; tumulthunder@gmail.com

Received 13 July 2013; Accepted 11 September 2013

Academic Editors: T. Hori and S. Kan

Copyright © 2013 Tumul Chowdhury et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Background. Perioperative aneurysm rupture (PAR) is one of the most dreaded complications of intracranial aneurysms, and approximately 80% of nontraumatic SAHs are related to such PAR aneurysms. The literature is currently scant and even controversial regarding the issues of various contributory factors on different phases of perioperative period. Thus this paper highlights the current understanding of various risk factors, variables, and outcomes in relation to PAR and try to summarize the current knowledge. **Method.** We have performed a PubMed search (1 January 1991–31 December 2012) using search terms including “cerebral aneurysm,” “intracranial aneurysm,” and “intraoperative/perioperative rupture.” **Results.** Various risk factors are summarized in relation to different phases of perioperative period and their relationship with outcome is also highlighted. There exist many well-known preoperative variables which are responsible for the highest percentage of PAR. The role of other variables in the intraoperative/postoperative period is not well known; however, these factors may have important contributory roles in aneurysm rupture. Preoperative variables mainly include natural course (age, gender, and familial history) as well as the pathophysiological factors (size, type, location, comorbidities, and procedure). Previously ruptured aneurysm is associated with rupture in all the phases of perioperative period. On the other hand intraoperative/postoperative variables usually depend upon anesthesia and surgery related factors. Intraoperative rupture during predissection phase is associated with poor outcome while intraoperative rupture at any step during embolization procedure imposes poor outcome. **Conclusion.** We have tried to create such an initial categorization but know that we cannot scale according to its clinical importance. Thorough understanding of various risk factors and other variables associated with PAR will assist in better clinical management as well as patient care in this group and will give insight into the development and prevention of such a catastrophic complication in these patients.

1. Introduction

Subarachnoid hemorrhage (SAH) is one of the most devastating neurological diseases. This condition not only produces the significant impact on mortality and morbidity but also imparts dire social consequences [1–3]. Perioperative aneurysm rupture (PAR) is one of the most dreaded complications of intracranial aneurysms and approximately 80% of nontraumatic SAHs are related to such ruptured intracranial aneurysms. PAR will depend upon the natural course of

disease, contributory risk factors, and time of intervention [4]. PAR will be influenced by different variables presented during the preoperative, intraoperative, and postoperative period. However, the literature is currently scant and even controversial regarding the issues of the various contributory factors during the different phases of the perioperative period. Thus this paper highlights the current understanding of various risk factors, variables, and outcomes in relation to the perioperative rupture of intracranial aneurysm and try to summarize the current knowledge.

2. Method

We have performed a PubMed search (1 January 1991–31 December 2012) using search terms including “cerebral aneurysm,” “intracranial aneurysm,” and “intraoperative/perioperative rupture.” Only papers in the English language that specifically discussed the relevant complication and various risk factors were included. The articles related to complications and management of complex giant aneurysms and pediatric aneurysm were not included in this review.

3. Risk Factors for Aneurysm Rupture and Outcome

The mechanism of cerebral aneurysmal rupture remains unknown at present. It is, however, known that a chronic inflammatory reaction is occurring within the aneurysmal wall, being associated with the degeneration of the aneurysmal wall and susceptibility of the aneurysm to bleeding [5]. However, screening or identification of bleeding-prone cerebral aneurysms cannot yet be performed, so that the knowledge of the various risk factors and variables is important and can be mainly divided into the three phases of the perioperative period. There exist many well-known preoperative variables which are responsible for the highest percentage of aneurysm rupture. The role of other variables in the intraoperative as well as postoperative period is not well known; however, these factors may have important contributory roles in aneurysm rupture.

3.1. Preoperative Variables. Though the overall risk of rupture in unruptured cerebral aneurysm is low (<0.5% per year) except for giant aneurysms, the morbidity and mortality associated with rupture can be high [6]. These unruptured cerebral aneurysms often remain asymptomatic, or sometimes, the smaller aneurysms can produce also compressive symptoms (involvement of cranial nerves mainly third and fourth) or ischemic symptoms due to thromboembolic episodes [7, 8]. Thus, it is imperative to know about the natural course (age, gender, and familial history) as well as the pathophysiological factors (size, type, location, comorbidities, procedure) which would govern cerebral aneurysm rupture perioperatively [9]. In addition, postrupture of cerebral aneurysm certain preventable medical conditions further adds to overall morbidity and mortality. In addition, one study has highlighted that comorbidities (such as arterial hypertension, congestive heart failure, and electrolyte disturbances) were associated with increase in the likelihood of longer median length of stay in all patients with cerebral aneurysms [10, 11].

Prevalence of unruptured cerebral aneurysms is increasing worldwide and is estimated to be approximately 0.65%–3.2%. The two most common risk factors associated with SAH are patients with adult polycystic kidney (ADPKD) and a positive family history of intracranial aneurysm of SAH [12]. Interestingly, in some countries like Japan and Finland, higher incidence of SAH is not found to be correlated with higher prevalence of unruptured aneurysms (UIA) and thus

suggests the existence of other cofactors responsible for causing SAH in this population [13].

Female gender represents also a significant risk factor. However, the influence of female hormones on the pathogenesis of aneurysm formation and rupture is inconclusive. In the affected families, the most frequent relationship exists between the siblings. The genetic predisposition is a significant contributory factor for multiplicity as well as location of aneurysm. In addition, familial SAH was found to be associated with larger aneurysm at the time of rupture [13, 14].

A Japanese study group revealed that more than 90% of cerebral aneurysm cases were incidentally detected and warrants the need for preventive as well as screening methods to detect unruptured aneurysm. Most common aneurysms detected in this study were in the middle cerebral arteries (MCA) and the internal carotid arteries (ICA); however, aneurysms located in anterior (ACoA) and posterior communicating artery (PCoA) were more likely to be ruptured in comparison to MCA aneurysms [15]. Familial intracranial aneurysm (FIA) study also showed the preponderance of MCA location as opposed to International Study of Unruptured Intracranial Aneurysms (ISUIA) which had more patients with PCoA aneurysms (patients who had a family history of intracranial aneurysm or subarachnoid hemorrhage were excluded) [14]. The effect of geographical area on incidence and type of aneurysm rupture is inconclusive at present; however, Japanese study showed that the large, posterior-circulation, and symptomatic aneurysms were associated with significantly higher rates of rupture. The selection bias could not be eliminated from this study and could possibly account for the higher incidence of rupture. In contrast, US based study found that more than 90% of the unruptured aneurysms were located in the anterior circulation and a location was also similar for patients with ruptured aneurysms. However, the lack of correlation of size and age at rupture (exposure to risk factors) suggests that the size at rupture is more dependent on hemodynamic stress [16, 17].

According to one of the largest studies, age is the most significant risk factor associated with surgical outcome; however, the size and location of an aneurysm influence both surgical as well as endovascular outcomes [18]. Similarly, the International Study of Unruptured Intracranial Aneurysms (ISUIA) found that a higher rupture rate was reported to be associated with larger aneurysms, located in the posterior circulation, and previous history of SAH.

Biochemical factors linked together with ICA rupture were also investigated but not well understood. Complement dysregulation is the major mechanism which in turn increases the susceptibility to complement activation, inflammation, and tissue damage within the intra-arterial wall. The role of gene variations of endothelial nitric oxide synthase (eNOS), a vasomodulatory protein to assist in identifying patients with a high likelihood of rupture, has also been highlighted [19, 20]. Apoptosis is the described major mechanism responsible for the process of aneurysm rupture.

The geometrical as well as biophysical characteristics were also studied for the future prediction of cerebral aneurysm rupture and include but are not limited to three-dimensional

diameter (D-max), aspect ratio, flow angle, parent-daughter angle, wall shear stress, and oscillatory shear index (OSI). However, none of these factors have been found to be superior to one another [21–23]. The predictors of aneurysm growth were also studied and included as a diameter of at least 10 mm and a location at the basilar artery (BA) bifurcation or the ICA [24].

Some of the rare types of cerebral aneurysms also have an effect on the periprocedural rupture rate and thus increase the overall morbidity and mortality. In this regard, intrasellar aneurysms can impose grave complications. Most of them are associated with either compressive symptoms or endocrinopathy and often mimic as pituitary tumors [25, 26]. The other rarer type of aneurysm often described as blood-blister like aneurysm was also very notorious to be ruptured acutely and often requires internal carotid artery trapping combined with high-flow extracranial-intracranial (trapping/EC-IC) bypass during the acute period following SAH [27–30]. The management of this type of aneurysm is often challenging for both the surgeon and interventional neuroradiologist. Pericallosal aneurysms, which comprise another rarer group, often have an increased chances of intraoperative rupture [26].

The role of diurnal variation on cerebral aneurysm rupture has not been well studied; however, nocturnal rupture associated SAH has been found to be an independent risk factor for developing cerebral ischemia and poor outcome [31]. The relation between sexual intercourse and chances of aneurysm rupture was also highlighted in a review and could be one of the possibilities of higher incidence of aneurysm rupture among young individuals [32].

Cerebral aneurysms which are present at bifurcations tend to be more at risk for rupture due to shear wall stress; however, in the study of 77 PCoA aneurysms, true PCoA aneurysms might be more prone for rupture than junctional aneurysms of similar size [33]. Similarly lower risks were observed for the rupture of carotid bifurcation aneurysms and further support the hypothesis that the anatomical geometry of the bifurcations and concomitant hemodynamic stress are not the sole factors which govern the overall risk of rupture [34].

3.2. Intraoperative Variables

3.2.1. Preprocedural Factors. Though rare, the preprocedural rupture of cerebral aneurysms can produce catastrophic complications with high mortality. Anesthesia related factors contribute to this phase of intraoperative rupture. There exists a fine balance between mean arterial pressure, ICP and transmural pressure (TMP). The acute changes in hemodynamics during the periods of induction, surgical incision, and skull pin fixation may increase the transmural pressure and may be responsible for intraoperative aneurysm rupture [35, 36]. Sudden coughing or gagging during any point of time can evoke acute rise in blood pressure and intracranial pressure that, hence, leads to rupture.

Raised intracranial pressure (ICP) is usually associated with poor grade SAH but may present in good grade too.

TABLE 1: Intraoperative variables contributing intracranial aneurysm rupture.

	Outcome
<i>(1) Preprocedural factors (same for coiling procedures)</i>	
Anesthesia related	Poor
Pain	
Light plane of anesthesia	
Intubation response	
Acute fluctuations in TMP	
High blood pressure (induction, skull pin fixation, and incision)	
Coughing/gagging	
Raised ICP	
Large and rapid bolus mannitol	
Acute hyperventilation	
<i>(2) Intraprocedural factors</i>	Variable
Increased vessel wall fragility (same for coiling procedures)	Poor
Smoking/COPD	
Hyperlipidemia/CAD	
Lower Hunt and Hess grade	
Predissection phase	Poor
(Dural opening and arachnoid opening, hematoma removal, and brain retraction)	
Dissection phase	Variable
Type of artery (PICA, ACA, and PCA)	
Previously ruptured aneurysm	
Surgeon's experience	

Raised ICP can again affect the TMP and can precipitate intraoperative aneurysm rupture (IOAR). However, data do not support significant link between raised ICP and intraoperative rupture of aneurysm [37]. The other variables like rapid boluses of mannitol and excessive hyperventilation before the dural opening may produce IOAR [35].

3.2.2. Intraprocedural Factors. The intraprocedural rupture (IPAR) mainly depends upon vessel wall fragility which in turn may be modified by the several comorbidities including coronary artery disease, hyperlipidemia, race, COPD, and lower Hunt and Hess grade (Table 1) [38].

In study of 1010 patients with cerebral aneurysm surgery (299 coiled, 711 clipped), 14.6% developed intraprocedural aneurysm rupture (IPAR) (19% with clipping and 5% with coiling). There was significant higher mortality in coiling related IPAR [38]. Predissection phase variables which are responsible for IOAR usually include dural opening and arachnoid opening, hematoma removal, and brain retraction. In study of 398 patients with ruptured aneurysm, 6% showed premature (before securing the parent vessel) rupture of aneurysm and significantly contributed to brain swelling [39]. The partial resection of brain could give the space for temporarily clipping and could be found to be related

to poor neurological outcome. In another study in 170 patients with ACA and MCA rupture aneurysm, there was 9% incidence of intraoperative rupture due to the procedure itself. Intraoperative aneurysmal rupture usually occurred during dissection of the aneurysm, dissection of the artery adhering to the aneurysm, or during clip application [40].

In a retrospective study of 1269 surgically treated patients, 113 IOAR events occurred. The posteroinferior cerebellar artery and ACoA and PCoA aneurysms were more liable to rupture intraoperatively. The IOR rate was greater in ruptured than unruptured aneurysms [41].

Experience of the surgeon certainly plays a crucial role during the intraprocedural period. Data suggests that increase in surgical experience reduces the number of intraprocedural ruptures, duration of temporary clipping, and the surgical mortality rate and hence favors better outcome [42]. This conclusion also exists for the neurointerventional surgeon/radiologist. The association between the types of surgical approaches and intraoperative rupture requires further attention and is described in more detail elsewhere.

Neurovascular management of very small aneurysms often imposes challenges for the both the surgical as well as interventional procedures [43]. Interestingly in one study small cerebral aneurysms were associated with larger volume of SAH; however the other factors including patient sex and age, intraparenchymal or intraventricular hemorrhage, multiple aneurysms, history of hypertension, and aneurysm location were not associated with a larger volume of SAH [44, 45]. The chances of intraprocedural aneurysm rupture were found to be relatively higher during embolization especially in very small aneurysms. The main causes were coil protrusion and microcatheter perforation. Angiographic demonstration of deterioration in cerebral hemodynamics was found to be an independent detriment of poor outcome. The study on endovascular management (CLARITY trial) revealed that a higher proportion of patients with intraoperative rupture were found to be younger (age less than 65 year) as well as hypertensive and more commonly associated with MCA aneurysm [46]. Furthermore, authors concluded that it is the location of aneurysm not the size which predicts the chances of intraoperative aneurysm rupture. On the other hand, the same study highlighted the significance of both location and size of aneurysm neck in causing thromboembolic complications. Another study revealed that there were 5 times greater chances of intraprocedural (endovascular) rupture in previously ruptured very small aneurysm in comparison with large ruptured aneurysm (Table 2). In this regard, balloon assisted hemostasis was associated with a better outcome. Furthermore, when coiling alone was compared with remodeling methods, the intraprocedural ruptures were found to be similar with comparable overall mortality and morbidity [47, 48].

On the other hand, management of very large or even giant cerebral aneurysms is also very complex and associated with significant risk [29]. Surgical indications are usually reserved for younger patients; however, depending upon risk and benefit ratio, elderly age group patients warrant considerations for endovascular management. Minimization

TABLE 2: Intraoperative variables contributing intracranial aneurysm rupture (coiling only).

	Outcome
Preprocedural	
Hypertensive patients	Poor
Angiography in acute phase of SAH	
Contrast injection induced hypertension	
Intraprocedural factors	
Age younger than 65 years	Poor
Very small aneurysms	
Wide neck > 4 mm	
Type of artery (MCA and posterior circulation)	
Previously ruptured aneurysm	
Experience of neuroradiologist	

of temporary occlusion and the use of intraoperative angiography have been shown to favor the surgical outcome in these cases [49].

The present literature is also variable regarding the intraoperative rupture and long-term neurological outcome. Some studies have concluded that intraoperative rupture adversely affects the outcome; however, these data should be interpreted in relation to various factors including size, location, procedure (clipping versus coiling), age, management strategies, and experience of surgeon [50]. The outcome is usually worse in cases of IOR associated with coiling. Even in the hybrid suite, the intraoperative rupture during coiling was associated with 40% mortality and 20% long-term morbidity. On the other hand, outcome was found to be better after IOR of small aneurysm during coiling in comparison with a large one [51].

Temporarily arterial occlusion at the time of the clipping is now well-established modality, and still may impose many complications. In a retrospective study of 112 patients, 17% of the patients were found to have symptomatic stroke and 26% had radiological evidence of stroke attributable to temporary arterial occlusion [52]. Advancing age and poor clinical grade were the risk factors significantly associated with symptomatic stroke. Intraoperative cerebral aneurysm rupture and duration of temporary clip application that lasted more than 20 minutes were found to be independent predictors of perioperative stroke outcome. In this regard, duration of single clipping seems to be more important risk factors rather than the placement of multiple clips for the short period of time [53].

3.3. Postoperative Variables. There are only few reports available regarding the postoperative cerebral aneurysm rupture in which residual aneurysm or arterial dissection was highlighted, and the factors contributed were intracranial hypotension due to excessive CSF drainage, hemodynamic stress caused by hypertension, and trauma due to surgical manipulation [54, 55].

There have been reports of aneurysm recurrence and subsequent SAH from small residual necks (1-2 mm) after clipping [56]. On the other hand, in cases of incomplete

TABLE 3: Perioperative variables contributing intracranial aneurysm rupture.

Perioperative variable	Outcome
(1) <i>Preoperative factors</i>	Variable
Age	
Female gender	
Size (large aneurysm) and location (posterior circulation aneurysm)	
Genetic factors/familial factors	
Hypertension	
Smoking	
Ruptured aneurysm	Poor
(2) <i>Intraoperative factors</i>	Poor
Preprocedural factors	
Anesthesia related	
Factors affecting TMP	
Predissection variables	
Intraprocedural variables	
Predissection variables	Poor
Dissection variables	
Coiling versus surgery	Poor/variable
Size and location	
Comorbidities	
(3) <i>Postoperative factors</i>	Poor
Emergence/extubation related	
Surgical dissection/trauma	
Incomplete obliteration by coiling or clipping	

occlusion after coiling, this may impose more chances of rerupture risks; however, the data is scant at present so the contributory factors are yet to be studied.

4. Future Directions

To really better understand the risk factors that influence cerebral aneurysmal rupture, we need first to better understand the mechanism of rupture. Imaging methods may assist us to gain better insights into these underlying mechanisms that are clinically important. It is surprisingly that until now very few relevant studies have been performed in this context. Such created knowledge would assist to better categorize the numerous risk factors that are currently known and discussed within this paper. We have tried to create such an initial categorization (Table 3) but know that we cannot scale according to its clinical importance. However, deeper analysis shows that these situations are anything but simple and are often additionally influenced by dogmas in treatment that makes it not easier to compare the treatment between different institutions. But to go ahead a further step includes openly describing also errors and adverse effects that have been seen during the treatment of cerebral aneurysm, a fact that lacks generally in medicine.

In conclusion, thorough understanding of various risk factors and other variables associated with perioperative aneurysm rupture will assist in better clinical management as well as patient care in this group and will give insight into the development and prevention of such a catastrophic complication in these patients.

Conflict of Interests

There is no conflict of interests, and Bernhard Schaller is the editorial member of Scientific World Journal (molecular imaging).

Authors' Contribution

Tumul Chowdhury is responsible for data acquisition, data interpretation, drafting and writing the paper. Tumul Chowdhury approved the final paper. Ronald B. Cappellani helped in designing and writing the paper and approved the final paper. Nora Sandu edited and revised the paper and approved the final paper. Bernhard Schaller helped write the paper and approved the final paper. Jayesh Daya edited and revised the paper and approved the final paper.

References

- [1] F. Siddiq, S. A. Chaudhry, R. P. Tummala, M. F. Suri, and A. I. Qureshi, "Factors and outcomes associated with early and delayed aneurysm treatment in subarachnoid hemorrhage patients in the United States," *Neurosurgery*, vol. 71, no. 3, pp. 670–677, 2012.
- [2] R. J. McDonald, J. S. McDonald, J. P. Bida, D. F. Kallmes, and H. J. Cloft, "Subarachnoid hemorrhage incidence in the United States does not vary with season or temperature," *American Journal of Neuroradiology*, vol. 33, pp. 1663–1668, 2012.
- [3] N. M. van der Kolk, A. Algra, and G. J. E. Rinkel, "Risk of aneurysm rupture at intracranial arterial bifurcations," *Cerebrovascular Diseases*, vol. 30, no. 1, pp. 29–35, 2010.
- [4] J. G. Guimond, P. M. Chagnon, and M. W. Bojanowski, "Clipping vs. coiling in acute aneurysmal subarachnoid haemorrhage: should the patient's medical condition influence treatment modality?" *Neurochirurgie*, vol. 58, no. 2-3, pp. 115–119, 2012.
- [5] D. O. Wiebers, D. G. Piepgras, F. B. Meyer et al., "Pathogenesis, natural history, and treatment of unruptured intracranial aneurysms," *Mayo Clinic Proceedings*, vol. 79, no. 12, pp. 1572–1583, 2004.
- [6] B. Weir, L. Disney, and T. Karrison, "Sizes of ruptured and unruptured aneurysms in relation to their sites and the ages of patients," *Journal of Neurosurgery*, vol. 96, no. 1, pp. 64–70, 2002.
- [7] J. A. Friedman, D. G. Piepgras, M. A. Pichelmann, K. K. Hansen, R. D. Brown Jr., and D. O. Wiebers, "Small cerebral aneurysms presenting with symptoms other than rupture," *Neurology*, vol. 57, no. 7, pp. 1212–1216, 2001.
- [8] T. W. Malisch, G. Guglielmi, F. Viñuela et al., "Unruptured aneurysms presenting with mass effect symptoms: response to endosaccular treatment with Guglielmi detachable coils. Part I. Symptoms of cranial nerve dysfunction," *Journal of Neurosurgery*, vol. 89, no. 6, pp. 956–961, 1998.

- [9] C. L. Taylor, Z. Yuan, W. R. Selman, R. A. Ratcheson, and A. A. Rimm, "Cerebral arterial aneurysm formation and rupture in 20,767 elderly patients: hypertension and other risk factors," *Journal of Neurosurgery*, vol. 83, no. 5, pp. 812–819, 1995.
- [10] N. J. Solenski, E. C. Haley Jr., N. F. Kassell et al., "Medical complications of aneurysmal subarachnoid hemorrhage: a report of the multicenter, cooperative aneurysm study. Participants of the Multicenter Cooperative Aneurysm Study," *Critical Care Medicine*, vol. 23, no. 6, pp. 1007–1017, 1995.
- [11] G. A. Smith, P. Dagostino, M. G. Maltenfort, A. S. Dumont, and J. K. Ratliff, "Geographic variation and regional trends in adoption of endovascular techniques for cerebral aneurysms," *Journal of Neurosurgery*, vol. 114, no. 6, pp. 1768–1777, 2011.
- [12] M. H. M. Vlak, A. Algra, R. Brandenburg, and G. J. E. Rinkel, "Prevalence of unruptured intracranial aneurysms, with emphasis on sex, age, comorbidity, country, and time period: a systematic review and meta-analysis," *The Lancet Neurology*, vol. 10, no. 7, pp. 626–636, 2011.
- [13] Y. M. Ruigrok, G. J. E. Rinkel, A. Algra, T. W. M. Raaymakers, and J. van Gijn, "Characteristics of intracranial aneurysms in patients with familial subarachnoid hemorrhage," *Neurology*, vol. 62, no. 6, pp. 891–894, 2004.
- [14] J. Mackey, R. D. Brown Jr., C. J. Moomaw et al., "Unruptured intracranial aneurysms in the Familial Intracranial Aneurysm and International Study of Unruptured Intracranial Aneurysms cohorts: differences in multiplicity and location," *Journal of Neurosurgery*, vol. 117, no. 1, pp. 60–64, 2012.
- [15] The UCAS Japan Investigators, "The natural course of unruptured cerebral aneurysms in a Japanese cohort," *The New England Journal of Medicine*, vol. 366, pp. 2474–2482, 2012.
- [16] A. Morita, S. Fujiwara, K. Hashi, H. Ohtsu, and T. Kirino, "Risk of rupture associated with intact cerebral aneurysms in the Japanese population: a systematic review of the literature from Japan," *Journal of Neurosurgery*, vol. 102, no. 4, pp. 601–606, 2005.
- [17] H. Richard Winn, J. A. Jane Sr., J. Taylor, D. Kaiser, and A. G. W. Britz, "Prevalence of asymptomatic incidental aneurysms: review of 4568 arteriograms," *Journal of Neurosurgery*, vol. 96, no. 1, pp. 43–49, 2002.
- [18] D. O. Wiebers, J. P. Whisnant, J. Huston III et al., "Unruptured intracranial aneurysms: natural history, clinical outcome, and risks of surgical and endovascular treatment," *The Lancet*, vol. 362, no. 9378, pp. 103–110, 2003.
- [19] V. G. Khurana, I. Meissner, Y. R. Sohni et al., "The presence of tandem endothelial nitric oxide synthase gene polymorphisms identifying brain aneurysms more prone to rupture," *Journal of Neurosurgery*, vol. 102, no. 3, pp. 526–531, 2005.
- [20] R. Tulamo, J. Frösen, A. Paetau et al., "Lack of complement inhibitors in the outer intracranial artery aneurysm wall associates with complement terminal pathway activation," *American Journal of Pathology*, vol. 177, no. 6, pp. 3224–3232, 2010.
- [21] I. Shiue, H. Arima, G. J. Hankey, and C. S. Anderson, "Location and size of ruptured intracranial aneurysm and serious clinical outcomes early after subarachnoid hemorrhage: a population-based study in Australasia," *Cerebrovascular Diseases*, vol. 31, no. 6, pp. 573–579, 2011.
- [22] T. Kawaguchi, S. Nishimura, M. Kanamori et al., "Distinctive flow pattern of wall shear stress and oscillatory shear index: similarity and dissimilarity in ruptured and unruptured cerebral aneurysm blebs," *Journal of Neurosurgery*, vol. 117, no. 4, pp. 774–780, 2012.
- [23] N. Lin, A. Ho, B. A. Gross et al., "Differences in simple morphological variables in ruptured and unruptured middle cerebral artery aneurysms," *Journal of Neurosurgery*, vol. 117, no. 5, pp. 913–919, 2012.
- [24] S. Matsubara, H. Hadeishi, A. Suzuki, N. Yasui, and H. Nishimura, "Incidence and risk factors for the growth of unruptured cerebral aneurysms: observation using serial computerized tomography angiography," *Journal of Neurosurgery*, vol. 101, no. 6, pp. 908–914, 2004.
- [25] B. W. Hanak, G. Zada, V. V. Nayar et al., "Cerebral aneurysms with intrasellar extension: a systematic review of clinical, anatomical, and treatment characteristics," *Journal of Neurosurgery*, vol. 116, no. 1, pp. 164–178, 2012.
- [26] T. N. Nguyen, J. Raymond, D. Roy et al., "Endovascular treatment of pericallosal aneurysms," *Journal of Neurosurgery*, vol. 107, no. 5, pp. 973–976, 2007.
- [27] K. Kamijo and T. Matsui, "Acute extracranial-intracranial bypass using a radial artery graft along with trapping of a ruptured blood blister-like aneurysm of the internal carotid artery," *Journal of Neurosurgery*, vol. 113, no. 4, pp. 781–785, 2010.
- [28] B. Schaller, "Extracranial-intracranial bypass to reduce the risk of ischemic stroke in intracranial aneurysms of the anterior cerebral circulation: a systematic review," *Journal of Stroke and Cerebrovascular Diseases*, vol. 17, no. 5, pp. 287–298, 2008.
- [29] B. Schaller and P. Lyrer, "Anticoagulation of an unruptured, thrombosed giant intracranial aneurysm without hemorrhage or recanalization in the long-term follow-up," *European Journal of Neurology*, vol. 10, no. 3, pp. 331–332, 2003.
- [30] E. F. Hauck, B. Wohlfeld, B. G. Welch, J. A. White, and D. Samson, "Clipping of very large or giant unruptured intracranial aneurysms in the anterior circulation: an outcome study," *Journal of Neurosurgery*, vol. 109, no. 6, pp. 1012–1018, 2008.
- [31] S. Juvela, J. Siironen, J. Varis, K. Poussa, and M. Porras, "Risk factors for ischemic lesions following aneurysmal subarachnoid hemorrhage," *Journal of Neurosurgery*, vol. 102, no. 2, pp. 194–201, 2005.
- [32] M. R. Reynolds, J. T. Willie, G. J. Zipfel, and R. G. Dacey, "Sexual intercourse and cerebral aneurysmal rupture: potential mechanisms and precipitants," *Journal of Neurosurgery*, vol. 114, no. 4, pp. 969–977, 2011.
- [33] W. He, J. Hauptman, L. Pasupuleti et al., "True posterior communicating artery aneurysms: are they more prone to rupture? A biomorphometric analysis," *Journal of Neurosurgery*, vol. 112, no. 3, pp. 611–615, 2010.
- [34] S. Omodaka, S. Sugiyama, T. Inoue et al., "Local hemodynamics at the rupture point of cerebral aneurysms determined by computational fluid dynamics analysis," *Cerebrovascular Diseases*, vol. 34, pp. 121–129, 2012.
- [35] J. Guy, B. J. McGrath, C. O. Borel, A. H. Friedman, and D. S. Warner, "Perioperative management of aneurysmal subarachnoid hemorrhage: part I. Operative management," *Anesthesia and Analgesia*, vol. 81, no. 5, pp. 1060–1072, 1995.
- [36] J. B. Bederson, E. S. Connolly Jr., H. H. Batjer et al., "Guidelines for the management of aneurysmal subarachnoid hemorrhage: a statement for healthcare professionals from a special writing group of the Stroke Council, American Heart Association," *Stroke*, vol. 40, no. 3, pp. 994–1025, 2009.
- [37] G. G. Heuer, M. J. Smith, J. P. Elliott, H. R. Winn, and P. D. LeRoux, "Relationship between intracranial pressure and other clinical variables in patients with aneurysmal subarachnoid hemorrhage," *Journal of Neurosurgery*, vol. 101, no. 3, pp. 408–416, 2004.

- [38] L. Eljovich, R. T. Higashida, M. T. Lawton et al., "Predictors and outcomes of intraprocedural rupture in patients treated for ruptured intracranial aneurysms: the CARAT study," *Stroke*, vol. 39, no. 5, pp. 1501–1506, 2008.
- [39] K. Houkin, S. Kuroda, A. Takahashi et al., "Intra-operative premature rupture of the cerebral aneurysms. Analysis of the causes and management," *Acta Neurochirurgica*, vol. 141, no. 12, pp. 1255–1263, 1999.
- [40] T. Inagawa, "Dissection from fundus to neck for ruptured anterior and middle cerebral artery aneurysms at the acute surgery," *Acta Neurochirurgica*, vol. 141, no. 6, pp. 563–570, 1999.
- [41] T. J. Leipzig, J. Morgan, T. G. Horner, T. Payner, K. Redelman, and C. S. Johnson, "Analysis of intraoperative rupture in the surgical treatment of 1694 saccular aneurysms," *Neurosurgery*, vol. 56, no. 3, pp. 455–468, 2005.
- [42] M. T. Lawton and R. Du, "Effect of the neurosurgeon's surgical experience on outcomes from intraoperative aneurysmal rupture," *Neurosurgery*, vol. 57, no. 1, pp. 9–15, 2005.
- [43] S. M. Russell, K. Lin, S. A. Hahn, and J. J. Jafar, "Smaller cerebral aneurysms producing more extensive subarachnoid hemorrhage following rupture: a radiological investigation and discussion of theoretical determinants," *Journal of Neurosurgery*, vol. 99, no. 2, pp. 248–253, 2003.
- [44] I. Ioannidis, S. Laloo, R. Corkill, W. Kuker, and J. V. Byrne, "Endovascular treatment of very small intracranial aneurysms," *Journal of Neurosurgery*, vol. 112, no. 3, pp. 551–556, 2010.
- [45] T. N. Nguyen, J. Raymond, F. Guilbert et al., "Association of endovascular therapy of very small ruptured aneurysms with higher rates of procedure-related rupture," *Journal of Neurosurgery*, vol. 108, no. 6, pp. 1088–1092, 2008.
- [46] L. Pierot, C. Cognard, R. Anxionnat, and F. Ricolfi, "Ruptured intracranial aneurysms: factors affecting the rate and outcome of endovascular treatment complications in a series of 782 patients (CLARITY study)," *Radiology*, vol. 256, no. 3, pp. 916–923, 2010.
- [47] L. Pierot, L. Spelle, X. Leclerc, C. Cognard, A. Bonafé, and J. Moret, "Endovascular treatment of unruptured intracranial aneurysms: comparison of safety of remodeling technique and standard treatment with coils," *Radiology*, vol. 251, no. 3, pp. 846–855, 2009.
- [48] A. Gil, P. Vega, E. Murias, and H. Cuellar, "Balloon-assisted extrasaccular coil embolization technique for the treatment of very small cerebral aneurysms," *Journal of Neurosurgery*, vol. 112, no. 3, pp. 585–588, 2010.
- [49] B. Schaller and P. Lyrer, "Focal neurological deficits following spontaneous thrombosis of unruptured giant aneurysms," *European Neurology*, vol. 47, no. 3, pp. 175–182, 2002.
- [50] C. B. Luo, M. Mu-Huo Teng, F. C. Chang, C. J. Lin, W. Y. Guo, and C. Y. Chang, "Intraprocedure aneurysm rupture in embolization: clinical outcome with imaging correlation," *Journal of the Chinese Medical Association*, vol. 75, no. 6, pp. 281–285, 2012.
- [51] K. Yamakawa, S. Kiyama, Y. Murayama, and S. Uezono, "Incidence and neurological outcomes of aneurysm rupture during interventional neuroradiology procedures in a hybrid operating suite," *Journal of Anesthesia*, vol. 26, no. 4, pp. 592–594, 2012.
- [52] R. Ferch, A. Pasqualin, G. Pinna, F. Chioffi, and A. Bricolo, "Temporary arterial occlusion in the repair of ruptured intracranial aneurysms: an analysis of risk factors for stroke," *Journal of Neurosurgery*, vol. 97, no. 4, pp. 836–842, 2002.
- [53] C. S. Ogilvy, B. S. Carter, S. Kaplan, C. Rich, and R. M. Crowell, "Temporary vessel occlusion for aneurysm surgery: risk factors for stroke in patients protected by induced hypothermia and hypertension and intravenous mannitol administration," *Journal of Neurosurgery*, vol. 84, no. 5, pp. 785–791, 1996.
- [54] Y. Hashimoto, Y. Horita, T. Imaizumi, and J. Niwa, "Post-operative rupture of an untreated aneurysm on the 3rd day after subarachnoid hemorrhage surgery," *Neurologia Medico-Chirurgica*, vol. 45, no. 5, pp. 249–252, 2005.
- [55] Y. Goto, M. Hojo, S. Yamagata, and K.-I. Kikuta, "Fatal bleeding from arterial dissection after clipping of a ruptured aneurysm—case report," *Neurologia Medico-Chirurgica*, vol. 43, no. 12, pp. 608–611, 2003.
- [56] N. Pouratian, R. J. Oskouian Jr., M. E. Jensen, N. F. Kassell, and A. S. Dumont, "Endovascular management of unruptured intracranial aneurysms," *Journal of Neurology, Neurosurgery and Psychiatry*, vol. 77, no. 5, pp. 572–578, 2006.



Hindawi

Submit your manuscripts at
<http://www.hindawi.com>

