

## ความสัมพันธ์ระหว่างรูปแบบของ Circle of Willis และตำแหน่งของหลอดเลือดสมองโป่งพองที่แตก

ปัทมา อมาตยคง<sup>1\*</sup>, ปาริฉัตร ประจจะเนย์<sup>1</sup>, สิทธิชัย เอี่ยมสะอาด<sup>1</sup>, วรานนท์ มั่นคง<sup>2</sup>, สุกฤษฎ์ ตั้งมะโน<sup>3</sup>

<sup>1</sup>ภาควิชากายวิภาคศาสตร์<sup>2</sup>ภาควิชารังสีวิทยา คณะแพทยศาสตร์ มหาวิทยาลัยขอนแก่น จ.ขอนแก่น

<sup>3</sup>สำนักวิชาสาธารณสุขศาสตร์ มหาวิทยาลัยวลัยลักษณ์ จ.นครศรีธรรมราช

## The Association between Circle of Willis Patterns and Ruptured Aneurysm Sites

Pattama Amarttayakong<sup>1\*</sup>, Parichat Prajaney<sup>1</sup>, Sitthichai Iamsaard<sup>1</sup>, Waranon Munkong<sup>2</sup>, Sukrit Sangkhano<sup>3</sup>

<sup>1</sup>Department of Anatomy, <sup>2</sup>Department of Radiology, Faculty of Medicine, Khon Kaen University, Khon Kaen, Thailand.

<sup>3</sup>School of Public Health, Walailuk University, Nakhonsithammarat, Thailand

**หลักการและวัตถุประสงค์:** ภาวะเลือดออกใต้เยื่อหุ้มสมองชั้นกลางส่วนใหญ่เกิดจากการแตกของหลอดเลือดแดงสมองโป่งพองที่อยู่ใน circle of Willis (CW) หากตำแหน่งที่แตกได้รับการวินิจฉัยผิดพลาดจะทำให้เลือดออกซ้ำหลังการผ่าตัดจนผู้ป่วยอาจเสียชีวิตได้ CW ที่มีรูปแบบผิดปกติจะทำให้ผนังหลอดเลือดแดงที่เป็นองค์ประกอบมีความเครียดเพิ่มขึ้นและนำไปสู่การโป่งพองของหลอดเลือด ดังนั้นการศึกษานี้จึงมีวัตถุประสงค์เพื่อหาตำแหน่งของหลอดเลือดใน CW ที่มีการโป่งพองและแตกของผู้ป่วยที่มีเลือดออกใต้เยื่อหุ้มสมองชั้นกลาง ตลอดจนหาความสัมพันธ์ระหว่างรูปแบบของ CW และตำแหน่งที่แตกของหลอดเลือดสมองโป่งพอง

**วิธีการศึกษา:** ทำการศึกษาย้อนหลังในผู้ป่วยที่มีเลือดออกใต้เยื่อหุ้มสมองชั้นกลางจากการแตกของหลอดเลือดแดงสมองโป่งพองที่ได้รับการถ่ายภาพรังสีหลอดเลือด 2 มิติ (DSA) และ 3 มิติ (3DRA) เพื่อหาตำแหน่งของหลอดเลือดใน CW ที่มีการโป่งพอง และวัดเส้นผ่านศูนย์กลางของหลอดเลือดแดงที่เป็นองค์ประกอบ เพื่อจำแนกรูปแบบของ CW ออกเป็น 2 ชนิดหลัก คือชนิดตามแบบฉบับ และชนิดผิดปกติ และยังจำแนกชนิดผิดปกติออกเป็นรูปแบบย่อย 3 รูปแบบ คือ 1) atypical CW with transitional artery but without hypoplastic artery 2) atypical CW with hypoplastic artery และ 3) atypical CW with aplastic artery

**Background and Objectives:** Intracranial aneurysmal rupture is the major cause of subarachnoid hemorrhage (SAH). Misdiagnosis of the ruptured sites may result in postoperative rebleeding and the patients may eventually die. The variation of circle of Willis (CW) patterns could increase wall shear stress, leading to aneurysm formation. Therefore, this study aimed to investigate the sites of aneurysms inside the CW in aneurysmal SAH patients and the association between the CW patterns and ruptured aneurysm sites.

**Methods:** This retrospective study was performed on the digital subtraction angiography (DSA) and three dimensional rotational angiography (3DRA) of patients with aneurysmal subarachnoid hemorrhage. The sites of aneurysms in the CW were assessed and the internal diameters of the component arteries were measured to classify the patterns of CW as the typical and the atypical patterns. Atypical patterns were divided into 3 subtypes: 1) atypical CW with transitional artery but without hypoplastic artery 2) atypical CW with hypoplastic artery and 3) atypical CW with aplastic artery.

**Results:** All 90 aneurysmal SAH patients enrolled in this study were 30% (27/90) males and 70% (63/90) females with the mean age of 55.93 ± 12.87 years. Atypical CW

\*Corresponding Author: Pattama Amarttayakong, Department of Anatomy, Faculty of Medicine Khon Kaen University, 123 Mittraphap Road, Muang District, Khon Kaen 40002, Thailand. E-mail: apatta@kku.ac.th

**ผลการศึกษา:** ผู้ป่วยที่มีภาวะเลือดออกใต้เยื่อหุ้มสมองชั้นกลางจากการแตกของหลอดเลือดแดงสมองโป่งพองที่ใช้ในการศึกษานี้มีจำนวนทั้งสิ้น 90 ราย เป็นเพศชาย 30% (27/90) และเพศหญิง 70% (63/90) มีอายุเฉลี่ย  $55.93 \pm 12.87$  ปี พบรูปแบบผิดปกติของ CW มากถึง 95.96% โดยชนิดย่อยที่มีอุบัติการณ์สูงสุดคือชนิด atypical CW with hypoplastic artery (54.44%) ตำแหน่งที่พบการแตกของหลอดเลือดโป่งพองมากที่สุดในรูปแบบผิดปกติทุกชนิดของ CW คือ หลอดเลือดแดง anterior communicating (ACoA)

**สรุป:** การแตกของหลอดเลือดแดงสมองที่ไม่ได้เกิดจากการบาดเจ็บมักเกิดขึ้นภายใน CW และตำแหน่งที่พบหลอดเลือดแดงสมองโป่งพองและแตกมากที่สุดคือ ACoA ผู้ป่วยส่วนใหญ่ที่มีการแตกของหลอดเลือดแดงสมองโดยเฉพาะอย่างยิ่งที่ ACoA มักมีรูปแบบ CW เป็นชนิดผิดปกติ บ่งชี้ว่ารูปแบบ CW ชนิดผิดปกติสัมพันธ์กับตำแหน่งการแตกของหลอดเลือดโป่งพอง

pattern was found as high as 95.96% of aneurysmal series with outnumbered hypoplastic subtype (54.44%). The highest incidence among the ruptured sites of all atypical subtypes appeared at the anterior communicating artery.

**Conclusion:** Most non-traumatic aneurysmal rupture in this study occurred within the CW and ACoA was the predominant aneurysm site. The prevalence of atypical pattern was very high among the aneurysmal SAH patients and ACoA was the predominated ruptured aneurysm site in all atypical subtypes, showing the association between the aberrant CW patterns and ruptured aneurysm sites.

**Keywords:** Subarachnoid hemorrhage, cerebral aneurysm, circle of Willis, atypical pattern, anterior communicating artery

ศรีนครินทร์เวชสาร 2561; 33(4): 314-9. • Srinagarind Med J 2018; 33(4): 314-9.

## Introduction

Subarachnoid hemorrhage (SAH) refers to extravasation of blood into the subarachnoid space. There are many causes of SAH such as trauma, hemorrhage from arteriovenous abnormality, hematologic disorders, and cerebral aneurysm ruptures. More than 80% of SAH are due to rupture of a cerebral aneurysm.<sup>1</sup> Cerebral aneurysm occurs in many areas of cerebral vasculatures. The incidence of aneurysmal SAH has been reported in Thai patients by digital subtraction angiography (DSA).<sup>2</sup> The cerebral aneurysm have a close relationship to the angle of branching and anastomosis in the circle of Willis (CW).<sup>3</sup> About 85% of aneurysms occurring in the CW is berry aneurysm which is the most common cause of SAH.<sup>4,5</sup> There are many risk factors of cerebral rupture such as female sex,<sup>6,7</sup> size and site of cerebral aneurysms,<sup>8,9</sup> hypertension,<sup>9,10</sup> smoking,<sup>9,11</sup> and elderly patients.<sup>8,9,12</sup> Previous studies reported that the aneurysm forming and rupture were associated with CW variations.<sup>13,14</sup> The CW anomalies can change blood flow.<sup>15,16</sup> In a recent year, Nam and his partners analyzed wall shear stresses related to aneurysm formation in CW variation by microfluidic system. They suggested that the variation of CW morphology increased wall shear stress and could lead

to aneurysm formation.<sup>17</sup> The CW variations produce hemodynamic changes.<sup>18,19</sup> The force from these changes acts on the vessel wall in collateral sites leading to aneurysm formation and rupture.<sup>18,19</sup> Emergency aneurysmal SAH and multiple cerebral aneurysms challenge the interventional radiologists and neurosurgeons. The physicians must determine which one is ruptured because it is impossible to treat all aneurysms in single craniotomy. Misdiagnosis of ruptured site may result in rebleeding and the patient will eventually die because the true rupture site is untreated.<sup>20</sup> Accurate localization of aneurysm ruptures in patient with multiple cerebral aneurysms is still a problem in clinical practice. Additionally, the CW anomalies in Thai SAH patients associated with aneurysmal rupture has not been reported by three dimensional rotational angiography (3DRA) which can assess very small arteries. Therefore, this study aimed to investigate the sites of aneurysm on the CW in aneurysmal SAH patients and the association between the CW patterns and ruptured aneurysm sites based on DSA and 3DRA.

## Materials and Methods

1. **Sample collection:** The DSA and 3DRA of aneurysmal SAH patients were retrospective studied.

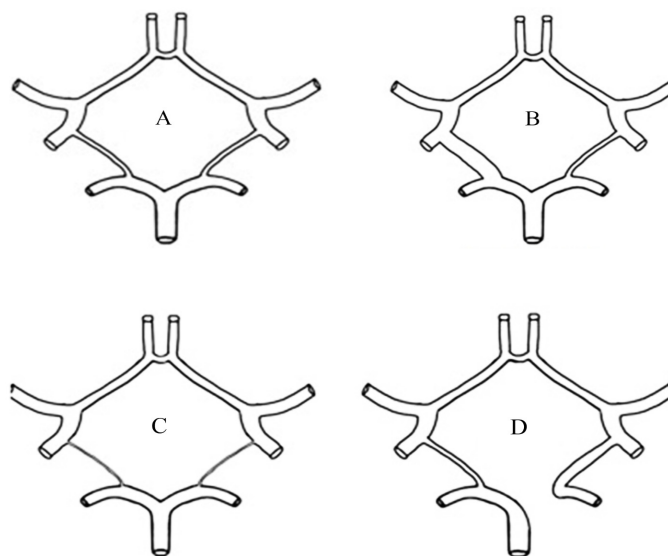
The patients underwent four vessels angiography at Interventional Radiology unit, Department of Radiology, Faculty of Medicine, Khon Kaen University since January 2015 to March 2016. The age, gender and sites of aneurysm were recorded. This study has been performed under the Khon Kaen University Ethics Committee in human research.

**2. Image technique:** The SAH angiography was conducted on a biplane system with 2 projections per vessel. Angiography was performed on a biplane neuroangiographic unit (Siemens Artis Zee Biplane). The DSA was conducted with a 1024 x 1024 matrix with a 17- to 20-cm field of view (FOV) and injection of 16 ml contrast material in the internal carotid and vertebral arteries in 2 projections. After processing, relevant images were sent to the picture archiving and communication system (PACS). A complete cerebral 3DRA is carried out on biplane system with 2 projections per vessel with an 8-second 180° rotational run, with acquisition of 200 images and injection of 3-4 milliliters contrast material per second in the internal carotid or vertebral arteries.

**3. Measurement and classification of CW patterns:** The internal diameters of component vessels of CW including supraclinoid part of internal carotid arteries (ICA), A1 segment of anterior cerebral arteries, posterior communicating arteries (PCoA), P1 segment of

posterior cerebral arteries, basilar artery (BA), and anterior communicating artery (ACoA) were measured at the proximal and distal parts to calculate the average diameters. The internal diameters of anterior and posterior communicating arteries can be detected and measured at only proximal part. Measurement was performed to classify the types of arteries as normal, hypoplastic and aplastic. The CW arteries with the internal diameter larger than 1 millimeter were classified as normal arteries. Any arteries with the diameter less than 1 millimeter were previously described as hypoplastic artery and the vessels invisible on angiogram were classified as an aplastic artery.<sup>21</sup> The term “transitional vessel” is used to describe PCoA with the diameter equal or larger than P1, but P1 was not hypoplastic.<sup>22</sup>

According to the internal diameter of CW arteries described above, types of CW were classified into 2 groups: typical and atypical groups. Typical group possesses normal diameter of all components of CW. Atypical group comprised of the anomalies of some component arteries in CW. Atypical group was classified into 3 subtypes including; 1) atypical CW with transitional artery but without hypoplastic artery, 2) atypical CW with hypoplastic artery, and 3) atypical CW with aplastic artery. (Figure 1)



**Figure 1** The example of CW pattern A; Typical CW pattern, B-D; atypical CW pattern (B; with transitional artery but without hypoplastic artery, C; with hypoplastic artery and D; with aplastic artery)

### Results

Among SAH patients enrolled in this study, there were 90 aneurysmal subjects with males in 30% (27/90) and females in 70% (63/90). The mean age of all

aneurysmal subjects was  $55.93 \pm 12.87$  years (range 1-70). Of the 90 aneurysmal SAH patients, 60% (54/90) had aneurysms inside the CW and 40% (36/90) outside. (Table 1)

**Table 1** The distribution of intracranial aneurysmal subjects relating with the CW

	Pattern of CW	Aneurysmal subjects		
		Inside CW	Outside CW	total
Typical 4 (4.44%)	Without hypoplastic or aplastic artery	4	0	4 (4.44%)
	With transitional artery but without hypoplastic artery	2	1	3 (3.33%)
Atypical 86 (95.56%)	With hypoplastic artery	27	22	49 (54.44%)
	With aplastic artery	21	13	34 (37.78%)
<b>Total</b>		54 (60%)	36 (40%)	90

The typical CW pattern was found in 4.44% among 90 aneurysmal patients, whereas that of atypical pattern was 95.96% as shown in Table 1. The atypical pattern of CW with hypoplastic vessels was the subtype

outnumbered with 54.44% of aneurysmal group. The lowest incidence (3.33%) was found in the subtype of atypical pattern with transitional artery but without hypoplastic artery.

**Table 2** The association between the CW patterns and the ruptured aneurysm sites

Patterns of CW	Aneurysms inside CW								total	
	Anterior circulation					Posterior circulation				
	Rt ICA	Lt ICA	Rt A1	Lt A1	ACoA	Rt PCoA	Lt PCoA	BA tip		
Typical 4 (4.44%)	Without hypoplastic / aplastic artery	-	-	1	-	1	-	2	-	4 (7.41%)
	With transitional artery but without hypoplastic artery	-	-	-	-	1	1	-	-	2 (3.70%)
Atypical 86 (95.56%)	With hypoplastic artery	1	1	-	-	9	8	4	4	27 (50.00%)
	With aplastic artery	-	-	1	-	10	5	4	2	21 (38.89%)
<b>Total</b>		1 (1.85%)	1 (1.85)	1 (1.85%)	0	21 (38.89%)	14 (25.93%)	10 (18.53%)	6 (11.11%)	54 (88.88%)

The association between the CW patterns and ruptured aneurysm sites was shown in Table 2. There were 88.88% of patients with aneurysms at the anterior circulation and 11.11% at the posterior circulation. The aneurysmal rupture were found at ACoA in 38.89%, the right PCoA in 25.93% and the left PCoA in 18.53%. Of 54 patients with aneurysms inside the CW, hypoplastic vessels and aplastic vessels subtypes appeared most frequently with 50% and 38.89%, respectively. ACoA was the ruptured aneurysm site with the highest incidence in all subtypes of atypical pattern: with transitional artery but without hypoplastic artery in 50% (1/2) with hypoplastic vessels in 33.33% (9/27) and with aplastic vessels in 47.62% (10/21).

### Discussion

The mean age of aneurysmal SAH patients in the present study was similar to that of a previous work.<sup>23</sup> In the present study, the prevalence of atypical pattern was as high as 95.96% of the aneurysmal SAH patients. These variations were previously reported by Kapoor and partners in 40% investigated in the Indian population. They proposed that usual causes for such anomalies were the persistence of some embryonic vessel that normally disappear, disappearance of vessels that would normally persist or sprouting of new vessels due to hemodynamic and genetic factors. Most aneurysms in this study occurred within the CW. This is supported that the aneurysms frequently develop at the sites of arterial branching which was mostly found on CW.<sup>3</sup> Predominant aneurysm site was ACoA, corresponding to a previous study.<sup>2</sup> Kasuya and coworkers described that ACoA site had smaller A1-A2 junction, leading to higher hemodynamic stress and aneurysm formation.<sup>24</sup> The efficacy of ACoA vasodilatation was also limited, comparing to the rest of intracranial arteries, because it is the only cerebral artery developing from the plexiform blood vessels.<sup>13</sup> ACoA and PCoA aneurysms with CW irregularities were associated with rupture because both sites seemed most relevant to hemodynamic changes due to CW

anomalies.<sup>14</sup> Ruptured aneurysms occurred more frequently in patients with atypical pattern of CW because the CW anomalies altered blood flow in the circle.<sup>15</sup> The CW variations generated hemodynamic changes with the force acting on the vessel wall in collateral sites, leading to aneurysm formation and rupture.<sup>19</sup>

In conclusions, most non-traumatic aneurysmal rupture in this study occurred within the CW and ACoA was the predominant aneurysm site. The prevalence of atypical pattern was very high among the aneurysmal SAH patients and ACoA was the predominated ruptured aneurysm site in all atypical subtypes, showing the association between the aberrant CW patterns and ruptured aneurysm sites. These findings should be imperative for the diagnosis and the surgical approaches of the intracranial aneurysm.

### Acknowledgements

This study was supported by the Department of Anatomy and the Department of Radiology, Faculty of Medicine, Khon Kaen University.

### References

1. Ishibashi T, Murayama Y, Urashima M, Saguchi T, Ebara M, Arakawa H, et al. Unruptured intracranial aneurysms: incidence of rupture and risk factors. *Stroke J Cereb Circ* 2009; 40: 313-6.
2. Kitkhuandee A, Thammaroj J, Munkong W, Duangthongpon P, Thanapaisal C. Cerebral angiographic findings in patients with non-traumatic subarachnoid hemorrhage. *J Med Assoc Thai* 2012; 95: 121-9.
3. Crawford T. Some observations on the pathogenesis and natural history of intracranial aneurysms. *J Neurol Neurosurg Psychiatry* 1959; 22: 259-66.
4. Gasparotti R, Liserre R. Intracranial aneurysms. *Eur Radiol* 2005; 15: 441-7.
5. D'Souza S. Aneurysmal subarachnoid hemorrhage. *J Neurosurg Anesthesiol* 2015; 27: 222-40.
6. Waaijer A, van Leeuwen MS, van der Worp HB, Verhagen HJ, Mali WP, Velthuis BK. Anatomic variations in the circle of Willis in patients with symptomatic carotid artery stenosis assessed with multidetector row CT angiography. *Cerebrovasc Dis* 2007; 23: 267-74.

7. Bor AS, Velthuis BK, Majoie CB, Rinkel GJ. Configuration of intracranial arteries and development of aneurysms: a follow-up study. *Neurology* 2008; 70: 700-5.
8. Nahed BV, DiLuna ML, Morgan T, Ocal E, Hawkins AA, Ozduman K, et al. Hypertension, age, and location predict rupture of small intracranial aneurysms. *Neurosurgery* 2005; 57: 676-83.
9. Wermer MJ, van der Schaaf IC, Algra A, Rinkel GJ. Risk of rupture of unruptured intracranial aneurysms in relation to patient and aneurysm characteristics: an updated meta-analysis. *Stroke* 2007; 38: 1404-10.
10. Suarez JI, Tarr RW, Selman WR. Aneurysmal subarachnoid hemorrhage. *N Engl J Med* 2006; 354: 387-96.
11. Juvela S, Poussa K, Porras M. Factors affecting formation and growth of intracranial aneurysms: a long-term follow-up study. *Stroke* 2001; 32: 485-91.
12. Juvela S, Porras M, Poussa K. Natural history of unruptured intracranial aneurysms: probability of and risk factors for aneurysm rupture. *J Neurosurg* 2008; 108: 1052-60.
13. Stojanovic N, Stefanovic I, Randjelovic S, Mitic R, Bosnjakovic P, Stojanov D. Presence of anatomical variations of the circle of Willis in patients undergoing surgical treatment for ruptured intracranial aneurysms. *Vojn Pregl* 2009; 66: 711-7.
14. Lazzaro MA, Ouyang B, Chen M. The role of circle of Willis anomalies in cerebral aneurysm rupture. *J Neurointerv Surg* 2012; 4: 22-6.
15. Hendrikse J, van Raamt AF, van der Graaf Y, Mali WP, van der Grond J. Distribution of cerebral blood flow in the circle of Willis. *Radiology* 2005; 235: 184-9.
16. Nakayama Y, Tanaka A, Kumate S, Tomonaga M, Takebayashi S. Giant fusiform aneurysm of the basilar artery: consideration of its pathogenesis. *Surg Neurol* 1999; 51: 140-5.
17. Nam SW, Choi S, Cheong Y, Kim YH, Park HK. Evaluation of aneurysm-associated wall shear stress related to morphological variations of circle of Willis using a microfluidic device. *J Biomech* 2015; 48: 348-53.
18. Burleson AC, Strother CM, Turitto VT. Computer modeling of intracranial saccular and lateral aneurysms for the study of their hemodynamics. *Neurosurgery* 1995; 37: 774-82.
19. Ujiie H, Liepsch DW, Goetz M, Yamaguchi R, Yonetani H, Takakura K. Hemodynamic study of the anterior communicating artery. *Stroke* 1996; 27: 2086-93.
20. Lee KC, Joo JY, Lee KS. False localization of rupture by computed tomography in bilateral internal carotid artery aneurysms. *Surg Neurol* 1996; 45: 435-40.
21. Siddiqi H, Tahir M, Lone KP. Variations in cerebral arterial circle of Willis in adult Pakistani population. *J Coll Physicians Surg Pak* 2013; 23: 615-9.
22. Karatas A, Coban G, Cinar C, Oran I, Uz A. Assessment of the circle of Willis with cranial tomography angiography. *Med Sci Monit* 2015; 21: 2647-52.
23. Nabaweesi-Batuka J, Kitunguu PK, Kiboi JG. Pattern of cerebral aneurysms in a Kenyan population as seen at an urban hospital. *World Neurosurg* 2016; 87: 255-65.
24. Kasuya H, Shimizu T, Nakaya K, Sasahara A, Hori T, Takakura K. Angles between A1 and A2 segments of the anterior cerebral artery visualized by three-dimensional computed tomographic angiography and association of anterior communicating artery aneurysms. *Neurosurgery* 1999; 45: 89-93.

