Objective: To identify specific angiographic factors associated with haemorrhagic presentation of brain arteriovenous malformation in Chinese paediatric patients.

Design: Retrospective cross-sectional observational study.

Setting: Four locoregional tertiary neurosurgical centres in Hong Kong: Queen Elizabeth Hospital, Tuen Mun Hospital, Kwong Wah Hospital, and Pamela Youde Nethersole Eastern Hospital.

Patients: Patients aged 18 years or younger who underwent pretreatment digital subtraction angiography for brain arteriovenous malformation between 1 January 2005 and 31 July 2013 were included. Patients were divided into haemorrhagic and non-haemorrhagic groups based on the initial presentation. Pretreatment digital subtraction angiographies were independently reviewed by two experienced neuroradiologists.

Main outcome measures: The following parameters were evaluated for their association with haemorrhagic presentation by univariate and multivariate analyses: nidus location, nidus size, nidus morphology (diffuse or compact); origin and number of arterial feeders; venous drainage; number of draining veins; presence of aneurysms, venous varices, and venous stenosis.

Results: A total of 67 children and adolescents (28 male, 39 female) with a mean age of 12 years were included. Of them, 52 (78%) presented with haemorrhage. Arteriovenous malformation size ($P=0.004$) and morphology ($P=0.05$) were found to be associated with haemorrhagic presentation by univariate analysis. Small arteriovenous malformation nidus size and diffuse nidal morphology were identified as independent risk factors for haemorrhage by multivariate analysis.

Conclusion: Smaller arteriovenous malformation size and diffuse nidal morphology are angiographic factors independently associated with haemorrhagic presentation. Bleeding risk is important in determining the therapeutic approach (aggressive vs conservative) and timeframe, particularly in paediatric patients.

New knowledge added by this study
- Studies on paediatric arteriovenous malformation (AVM) are scarce and mostly based in Caucasian populations. This multicentre study involving Chinese paediatric patients found that small AVM nidus size and diffuse nidal morphology are independent risk factors for haemorrhage.

Implications for clinical practice or policy
- These two angiographic features associated with haemorrhagic presentation can help local clinicians to assess bleeding risk and determine the therapeutic approach (aggressive vs conservative) and treatment timeframe in paediatric patients with cerebral AVM.
華籍兒童腦動靜脈畸形與出血相關的血管造影因素
霍泳珊、潘偉麟、謝健燊、劉顯宇、陳志軒、潘寧遠、曹慶恩、楊子慧、王耀忠、梁錦榮、邱麗珊、鄧國穎
目的：找出與腦動靜脈畸形華籍兒童患者出血相關的血管造影因素。
設計：回顧性橫斷面觀察研究。
安排：香港提供第三層醫療服務的四間腦外科中心：伊利沙伯醫院、屯門醫院、廣華醫院和東區尤德夫人那打素醫院。
患者：2005年1月1日至2013年7月31日期間因腦動靜脈畸形進行術前數字減影血管造影的18歲或以下患者均被列入研究範圍。根據患者病發時的情況把他們分為出血性和非出血性兩組，並由兩位經驗豐富的神經放射學醫生分別進行獨立審閱術前數字減影血管造影的影像。
主要結果測量：通過單變量和多變量分析為以下各項與出血相關的因素進行評估：病灶位置、大小和形態（瀰漫性或致密型）、供血動脈的來源和數量、引流靜脈的目的地和數量，以及是否出現動脈瘤、靜脈曲張和靜脈擴張。
結果：共67名兒童及青少年（28男39女；平均年齡12.2歲）被列入研究範圍。其中52人（78%）病發時有出血。單變量分析顯示腦動靜脈畸形的大小（P=0.004）和形態（P=0.05）與出血有關。多變量分析則顯示小型腦動靜脈畸形和其擴散形態為出血的獨立危險因素。
結論：小型腦動靜脈畸形和擴散形態均為血管造影中與出血相關的獨立因素。採用積極或保守的治療方法以及治療時間表均取決於患者的出血風險，尤以兒童患者為甚。

Introduction
Brain arteriovenous malformation (AVM) is a vascular abnormality that consists of multiple fistulous connections between arteries and veins without a normal intervening capillary bed. It is believed to be congenital in nature, and commonly presents in early adulthood.1 The usual clinical presentations of brain AVM include haemorrhage, seizures, headache, and progressive neurological deficit. About 52% to 77% of patients with AVM have initial haemorrhagic presentation,2-4 which is also associated with poorer prognosis. Various studies evaluating the history of AVM record an annual haemorrhage rate of about 2% to 4%.1,5

Computed tomography (CT) is the initial screening tool for identifying haemorrhage and demonstrating the location of the AVM. Subsequent angiographic evaluation is required for virtually all patients with suspected AVM, with digital subtraction angiography (DSA) being accepted as the gold standard for characterisation and grading. The information obtained from the angiogram is crucial in treatment decision-making and prognostication.

Brain AVM is an important cause of haemorrhagic stroke in children.6-4 Studies in adults have identified radiological features that are associated with haemorrhagic presentation and future haemorrhage.9-11 Similar studies on AVM in children are, however, scarce and mostly based on studies from Europe and North America.12,13 Whether those angiographic features that predict haemorrhage in Caucasian children with AVM similarly predict haemorrhage in Chinese children with AVM is unknown.

The objective of this multicentre study was to determine specific angiographic factors associated with haemorrhagic presentation in brain AVM in the Hong Kong Chinese paediatric population, with a view to assisting clinical decision-making regarding the optimal timing and type of treatment.

Methods
This was a multicentre retrospective cross-sectional observational study. We included patients aged 18 years or younger (at time of diagnosis) who underwent pretreatment cerebral DSA for a principal diagnosis of brain AVM from 1 January 2005 to 31 July 2013.

Patients were recruited from four locoregional tertiary neurological centres in Hong Kong: Queen Elizabeth Hospital, Tuen Mun Hospital, Kwong Wah Hospital, and Pamela Youde Nethersole Eastern Hospital. These are the major acute hospitals belonging to the catchment areas of Kowloon Central, New Territories West, Kowloon West, and Hong Kong East clusters, respectively, according to the geographical cluster designation by the Hospital Authority. These clusters serve approximately 4 million Hong Kong inhabitants. Consecutive patients were retrieved from the Clinical Data Analysis and Reporting System by entering the targeted date range (01 January 2005 to 31 July 2013, inclusive) and the following search parameters: age range (0-18 years); International Classification of Diseases, 9th Revision, diagnostic code (747.81, AVM); and procedure code (88.41, arteriography of cerebral arteries). Exclusion criteria included a lack of accessible pretreatment DSA, other angiographic diagnoses (eg spinal AVM, vein of Galen aneurysmal malformation, dural arteriovenous fistulae), and non-Chinese ethnicity based on data extracted from the electronic Patient Record (ePR) and radiology reports. Approval was obtained from the institutional ethics committee and patient consent was waived for this retrospective study.

Basic demographic factors, including age at presentation and clinical symptoms, were obtained from the ePR. Patients were divided into a haemorrhagic group (those presenting with intracranial haemorrhage) and a non-haemorrhagic group based on the CT of the brain at presentation. Pretreatment DSAs were independently reviewed by two experienced interventional neuroradiologists (with 7 years and 15 years of experience) who were blinded to the clinical presentation and provided with the same demographic data. Each brain AVM...
was evaluated for the following parameters: nidus location (deep: thalamus, basal ganglia, corpus callosum, or brain stem vs hemispheric: cerebral or cerebellar lobes), nidus size (small <3 cm vs medium 3-6 cm vs large >6 cm), nidus morphology (compact: little or no intervening brain within the nidus vs diffuse: presence of significant intervening brain within the nidus) [Figs 1 and 2], origin of arterial feeders (cortical vs deep), number of arterial feeders (single vs multiple), presence of either flow-related or intranidal aneurysms (yes vs no), venous drainage destination (superficial vs deep), number of draining veins (single vs multiple), presence of venous varices (yes vs no), and presence of venous stenosis (yes vs no). Any discrepancy in reviews between the two neuroradiologists was resolved by mutual consensus.

Association between the angiographic features and haemorrhage was analysed using Chi squared test and Fisher’s exact test for categorical variables, and Student’s t test for numerical variables in univariate analysis. Logistic regression (with “enter” strategy) was carried out for covariates with a P value of <0.15. All statistical calculations were performed using the Statistical Package for the Social Sciences (Windows version 16.0; SPSS Inc, Chicago [IL], US).

Results
The sample included 67 children and adolescents who were eligible for inclusion, of which 28 (42%) were boys and 39 (58%) were girls. Among the patients, 52 (78%) were in the haemorrhagic group and 15 (22%) were in the non-haemorrhagic group. The mean age at presentation was 12 years (range, 2-18 years). No significant differences in age (P=0.15) or sex (P=0.88) were demonstrated between the haemorrhagic and non-haemorrhagic groups. Of the 67 patients, one in the haemorrhagic group was known to have idiopathic thrombocytopenic purpura. The remaining 66 patients had no known medical condition predisposing to intracranial haemorrhage.

Of the 67 children, 25 (37%) presented with headache, 12 (18%) with hemiplegia, 11 (16%) with convulsion, seven (10%) with collapse, three (4%) with loss of consciousness, one (1%) with cerebellar signs, and eight (12%) had other features, including confusion, decreased responsiveness, numbness,
and restricted ocular motion. There were more asymptomatic patients in the non-haemorrhagic group (Fig 3). Three patients, all of whom were in the non-haemorrhagic group, were diagnosed incidentally with AVM during examination for precocious puberty, scalp haemangioma, and suspected neurofibromatosis type 1.

The frequency of haemorrhage of the 67 patients as a function of angioarchitectural features is shown in Table 1.

After univariate analysis, AVM size (P=0.004) and morphology (P=0.05) were the two factors found to be significantly associated with haemorrhagic presentation (Table 2). After multivariate analysis, small AVM size and diffuse nidal morphology were identified as independent risk factors for haemorrhage (Table 3). The odds of haemorrhagic presentation in patients with small AVM was about 9 times that of patients with medium-size AVM, whereas the odds for haemorrhagic presentation in patients with diffuse nidal morphology was approximately 12 times that of patients with compact AVM morphology. Factors found not to be statistically significantly associated with haemorrhagic presentation included location, origin of arterial feeders, number of arterial feeders, presence of related aneurysms, venous drainage, number of draining veins, presence of venous varices, and presence of venous stenosis.

**Discussion**

To date, DSA remains the gold standard for evaluating brain AVM owing to its superior temporal and spatial resolution, with the ability to provide dynamic information and allow accurate identification of supplying arteries and draining veins. Generally CT and magnetic resonance angiography studies do not provide important dynamic information on the arterial supply and venous drainage.

### TABLE 1. Frequency of haemorrhage as a function of angioarchitectural features in paediatric brain arteriovenous malformation (n=67)

<table>
<thead>
<tr>
<th>Feature</th>
<th>Total No.</th>
<th>No. (%) of patients with haemorrhage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Location</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Deep</td>
<td>22</td>
<td>19 (86.4)</td>
</tr>
<tr>
<td>Hemispheric</td>
<td>45</td>
<td>33 (73.3)</td>
</tr>
<tr>
<td>Size</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;3 cm</td>
<td>45</td>
<td>40 (88.9)</td>
</tr>
<tr>
<td>3-6 cm</td>
<td>22</td>
<td>12 (54.5)</td>
</tr>
<tr>
<td>Morphology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Compact</td>
<td>48</td>
<td>34 (70.8)</td>
</tr>
<tr>
<td>Diffuse</td>
<td>19</td>
<td>18 (94.7)</td>
</tr>
<tr>
<td>Venous drainage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Superficial</td>
<td>40</td>
<td>29 (72.5)</td>
</tr>
<tr>
<td>Deep</td>
<td>27</td>
<td>23 (85.2)</td>
</tr>
<tr>
<td>Draining vein</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single</td>
<td>39</td>
<td>32 (82.1)</td>
</tr>
<tr>
<td>Multiple</td>
<td>28</td>
<td>20 (71.4)</td>
</tr>
<tr>
<td>Intranidal aneurysm</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>18</td>
<td>13 (72.2)</td>
</tr>
<tr>
<td>Absent</td>
<td>49</td>
<td>39 (79.6)</td>
</tr>
<tr>
<td>Venous varix</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>6</td>
<td>3 (50.0)</td>
</tr>
<tr>
<td>Absent</td>
<td>61</td>
<td>49 (80.3)</td>
</tr>
<tr>
<td>Venous stenosis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>4</td>
<td>2 (50.0)</td>
</tr>
<tr>
<td>Absent</td>
<td>63</td>
<td>50 (79.4)</td>
</tr>
<tr>
<td>Feeding artery</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single</td>
<td>15</td>
<td>13 (86.7)</td>
</tr>
<tr>
<td>Multiple</td>
<td>52</td>
<td>39 (75.0)</td>
</tr>
<tr>
<td>Origin of feeding artery</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cortical</td>
<td>49</td>
<td>37 (75.5)</td>
</tr>
<tr>
<td>Deep</td>
<td>18</td>
<td>15 (83.3)</td>
</tr>
</tbody>
</table>

![FIG 3. Presenting symptoms in the haemorrhagic and non-haemorrhagic groups](image)
or treatment-related morbidity) may have lifelong and profound sequelae.14

To the best of our knowledge, our study is the first to evaluate the risk factors for brain AVM haemorrhage in Chinese paediatric patients. Ethnic differences exist in the incidence and haemorrhagic risk of AVM, and data from western populations are not routinely generalisable to the Chinese population. A cohort study of 1028 adult patients with AVM in the US has established a role of ethnic differences in brain AVM, with excess incidence in Asians, blacks, and Hispanics compared with Caucasians.15 The analysis reveals a statistically significant increased risk for subsequent AVM haemorrhage among Hispanics compared with Caucasians, and an insignificant trend for blacks and Asians.

Studies in adults have identified specific angiographic features of AVM that are associated with haemorrhagic presentation and future bleeding, including small size (<3 cm), deep location, deep venous drainage, single draining vein, intranidal aneurysms, and associated venous ectasia or stenosis.5,14,16,17

Our study identified small AVM size (<3 cm) to be an independent risk factor for haemorrhage. Small AVM size has been identified as a risk factor for haemorrhage in multiple adult studies,5,16-20 which is also demonstrated in a western paediatric population.12 Although the underlying pathophysiological mechanism is uncertain, some authors have postulated a relationship between AVM size and feeding artery pressures.5,12 Spetzler et al17 found a higher rate of haemorrhagic presentation among smaller AVMs and noted that smaller AVMs were associated with higher feeding artery pressures at the time of surgical management as well as larger haematoma sizes.

Diffuse AVM nidal morphology was identified as another independent risk factor for haemorrhage in our study. Although a similar relationship between morphology and haemorrhage was not demonstrated in a Caucasian paediatric population,16 diffuse AVM nidal morphology has been demonstrated in adults as a risk factor for haemorrhage.21 The underlying pathophysiological mechanism is uncertain. More information is needed to determine whether diffuse morphology is associated with haemodynamic aberrations such as increased pressure in the feeding artery or draining vein to account for the observed increased risk of haemorrhage.

Our study has several limitations. First, owing to the retrospective nature of this study, AVM patients with poorer clinical presentation who are unfit for DSA were not included. Second, although this is a multicentre study, the sample size was relatively small owing to the small number of paediatric patients undergoing DSA for AVM. Our study has also underestimated the haemorrhagic proportion of the study population, thus any potential associations between other angiographic features with haemorrhagic presentation that are more subtle to detect would remain undetected. Third, variations exist in the quality and amount of available angiographic images, as well as in the level of experience of the angiographers among the various centres; these may affect the radiological interpretation. Presence of intracranial haemorrhage can be inferred from the pretreatment DSA due to presence of blood vessel displacement, which is an inherent limitation of this study. Fourth, as presence of haemorrhage may render an originally compact nidus into a diffuse morphology, this is a limiting factor in determining the association between diffuse morphology and haemorrhagic presentation. Fifth, we were unable to control for the timing of DSA following the onset of presentation owing to the retrospective nature of this study. Lastly, the extent to which certain angiographic risk factors existent at the time of haemorrhagic presentation can be extrapolated as predictors of future haemorrhage in AVM is controversial. In other words, factors present at the time of presentation are not necessarily accurate predictors of future risk. For instance, several adult-based studies have identified a higher incidence of haemorrhagic presentation in small

**TABLE 2.** Univariate analysis of the angiographic features associated with haemorrhagic presentation

<table>
<thead>
<tr>
<th>Clinical/radiological factor</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Location (hemispheric vs deep)</td>
<td>0.351*</td>
</tr>
<tr>
<td>Size (small &lt;3 cm vs medium 3-6 cm)</td>
<td>0.004†</td>
</tr>
<tr>
<td>Morphology (diffuse vs compact)</td>
<td>0.050*</td>
</tr>
<tr>
<td>Venous drainage destination (deep vs superficial)</td>
<td>0.222†</td>
</tr>
<tr>
<td>No. of draining veins (1 vs &gt;1)</td>
<td>0.306†</td>
</tr>
<tr>
<td>Related aneurysm (no vs yes)</td>
<td>0.524*</td>
</tr>
<tr>
<td>Varix (no vs yes)</td>
<td>0.121*</td>
</tr>
<tr>
<td>Venous stenosis (no vs yes)</td>
<td>0.214*</td>
</tr>
<tr>
<td>No. of feeding arteries (1 vs &gt;1)</td>
<td>0.490*</td>
</tr>
<tr>
<td>Origin of feeding artery (deep vs superficial)</td>
<td>0.742*</td>
</tr>
</tbody>
</table>

* Fisher’s exact test
† Chi-squared test

**TABLE 3.** Multivariate analysis of the angiographic features associated with haemorrhagic presentation

<table>
<thead>
<tr>
<th>Feature</th>
<th>B value</th>
<th>Exp</th>
<th>P value</th>
<th>95% Confidence interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Size (small vs medium)</td>
<td>2.22</td>
<td>9.186</td>
<td>0.002</td>
<td>2.33-36.12</td>
</tr>
<tr>
<td>Morphology (diffuse vs compact)</td>
<td>2.46</td>
<td>11.74</td>
<td>0.031</td>
<td>1.26-109.76</td>
</tr>
</tbody>
</table>
AVMs, but failed to find an association between AVM size and future haemorrhage.22,23

Unlike in adults, large-scale prospective studies aiming to study the natural course of paediatric AVMs are unlikely to take place owing to the relatively strong argument against conservative treatment, according to the prevailing view that ruptured paediatric AVMs should be treated aggressively owing to the significant risk of recurrent haemorrhage and subsequent morbidity and mortality.12,24 The recent controversial ARUBA (A Randomised trial of Unruptured Brain Arteriovenous malformations) in adults has demonstrated that medical management alone is superior in patients with unruptured AVMs,25 but there is insufficient scientific evidence to justify extrapolation of these results to a paediatric population. Moreover, while it has been shown that paediatric AVMs with haemorrhagic presentation do not necessarily have a higher risk of future haemorrhage nor a higher annualised bleeding risk than adults,26 their greater cumulative risk given their longer remaining life expectancy may be an argument for more aggressive treatment of paediatric AVMs. Choice of treatment for a small, unruptured paediatric AVM is therefore complex and should involve thorough consideration of other angioarchitectural factors on a case-by-case basis.

Despite these limitations, our study provides useful initial insights to the angiographic features associated with haemorrhagic presentation of AVMs in Chinese paediatric patients from multiple locoregional neurosurgical centres. These features may assist in stratifying risk of haemorrhage and assign priority for intervention, although data from future larger-scale studies may be needed before such features can be robustly applied as haemorrhagic risk predictors in Chinese children with AVM.

References