The Presentation and Clinical Course of Intracranial Developmental Venous Anomalies in Adults
A Systematic Review and Prospective, Population-Based Study

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Background and Purpose—Reported risks of hemorrhage from intracranial developmental venous anomalies (DVAs) vary, so we investigated this in a systematic review and population-based study.

Methods—We systematically reviewed the literature (Ovid Medline and Embase to November 7, 2007) and selected studies of ≥20 participants with ≥1 DVA(s) that described their clinical presentation and/or their clinical course over a specified follow-up period. We also identified every adult first diagnosed with a DVA in Scotland from 1999 to 2003 and followed them in a prospective, population-based study.

Results—Of 2068 articles detected by the literature search, 15 met our inclusion criteria and described clinical presentation, 8 of which also described the clinical course of DVAs. In the 15 studies of 714 people first presenting with a DVA, 61% were incidental findings, the mode of presentation was unclear in 23%, 6% presented with nonhemorrhagic focal neurological deficit, 6% were associated with epileptic seizure, and <1% were associated with infarction. In studies of the clinical course of 422 people with a DVA, the hemorrhage rate after first presentation ranged from 0% to 1.28% per year. In the population-based study of 93 adults with DVAs, 98% were incidental, 1% presented with symptomatic hemorrhage, and 1% presented with an infarct, but there were no symptomatic hemorrhages or infarcts in 492 person-years of follow-up (0% per person-year; 95% CI, 0% to 0.7%).

Conclusions—Intracranial DVAs have a benign presentation and clinical course. (Stroke. 2009;40:00-00.)

Key Words: hemorrhagic • intracranial developmental venous anomaly • stroke • vascular malformations
quency of intracranial hemorrhage, infarction, and death attributable to DVAs at first presentation and during follow-up. We also sought to confirm or refute the findings of the review in a prospective, population-based study of DVAs.

Materials and Methods

Systematic Review

We systematically reviewed the literature on intracranial DVAs in November 2007 by running electronic search strategies (Supplemental Appendix, available online at http://stroke.ahajournals.org) in Ovid Medline (from 1950) and Embase (from 1980). The titles and available abstracts were read to identify studies of ≥20 participants with at least one intracranial DVA reporting original data on their clinical presentation and clinical course for objective outcome events over a specified follow-up period. Although a sample size of ≥20 participants represents an arbitrary cutoff, this is the minimum number required to reject a null hypothesis (at the P=0.05 level) and minimizes the influence of case reports and small case series, which tend to report unusual phenomena. If multiple publications arose from the same cohort, we selected the one with the largest sample size that was most pertinent to this review.

One author (J.M.L.H.) extracted the following data from each included study and crosschecked selected studies with another author (R.A.-S.S.). We derived the mode of clinical presentation for each participant from the published description. When associated CMs were thought to be responsible for the symptoms that led to DVA diagnosis, we regarded these DVAs as asymptomatic. We allocated a primary mode of presentation to each participant if >1 mode had been allocated by the study authors, in the following hierarchical order.

Symptomatic Hemorrhage

Symptomatic hemorrhage has evidence of recent hemorrhage close to the DVA (not due to an associated CM) on radiological or pathological examination, causing a symptomatic neurological event (any of the following: headache, seizures, global neurological deficit, or focal neurological deficit).

Symptomatic Infarction

Symptomatic infarction has radiological or pathological evidence of infarction close to the DVA causing rapidly developing clinical signs of focal or global neurological disturbance lasting ≥24 hours.

Nonhemorrhagic Focal Neurological Deficit

Nonhemorrhagic focal neurological deficit is a focal neurological deficit that was anatomically referable to the location of the DVA without radiological or pathological evidence of recent hemorrhage or infarction and that could not be due to an associated CM.

Seizure

A seizure is an epileptic seizure that was anatomically referable to the location of the DVA based on seizure semiology or electroencephalography.

Incidental

We regarded all other modes of presentation as incidental, including headache, asymptomatic hemorrhage, generalized seizure disorders, neurological deficits that were anatomically unrelated to the DVA, and deficits that were attributable to an associated CM.

We classified studies documenting the clinical course of DVAs as prospective, retrospective, or “lifetime” (if they used a retrospective approach to clinical events that occurred during each participant’s entire lifetime, i.e., from birth until the age reached at the study’s end point) or “not specified.” We extracted measures of follow-up duration (median or mean, range, and total person-years of follow-up), numbers of objective outcome events during follow-up, and calculated a symptomatic hemorrhage rate per person-year of follow-up with 95% CIs where possible.

Scottish Intracranial Vascular Malformation Study

Participants

The Scottish Intracranial Vascular Malformation Study (SIVMS) is a prospective, population-based cohort study based on an anonymized extract of data from the Scottish Audit of Intracranial Vascular Malformations (SAIVMs, www.saivms.scot.nhs.uk), which identified Scottish residents aged ≥16 years at the time they were first diagnosed with any type of intracranial vascular malformation in the years 1999 to 2003.13,14 SAIVMs uses multiple, overlapping sources of case ascertainment to identify incident adults and follows up those who do not opt out by medical records surveillance and annual postal questionnaires, which are sent to each adult’s general (family) practitioner. In this analysis, we included every adult in SAIVMs first diagnosed with an intracranial DVA during the years 1999 to 2003 and used follow-up data accrued until June 1, 2007. We defined a DVA as an intraparenchymal, purely venous variant of normal intracranial venous drainage, without an arteriovenous shunt, diagnosed by contrast-enhanced CT, MRI, or intra-arterial digital subtraction angiography.15

Data Collection

J.M.L.H. extracted the primary mode of DVA presentation (incidental, symptomatic hemorrhage, infarction, seizures, and focal neurological deficit [as defined previously]) for each SIVMS participant and double-checked with one of the authors (R.A.-S.S.) in case of doubt. During follow-up (defined as the period from the initial presentation that led to DVA diagnosis until the latest follow-up assessment or death), we specifically looked for the occurrence of any objective outcomes attributable to the DVA.

Ethical Approval

The Multicenter Research Ethics Committee for Scotland approved SIVMS (MREC/98/048).

Results

Systematic Review

Characteristics of Studies

The electronic searches yielded 1197 articles from Medline and 871 articles from Embase, of which 21 studies reported results on ≥20 participants with ≥1 DVA. Six of these studies did not match the selection criteria; one did not contain any information on clinical presentation or clinical course.13,14,16,22,23,27,29,32 Four were earlier, smaller versions of studies included in this systematic review,18-21 and one was a precursor of the SIVMS data presented in this article.7 The 15 remaining papers,11,13,16,22,23,27 reported on the clinical presentation of DVAs (Table 1), and 8 of these studies also described their clinical course (Table 3),12,13,16,22,24,26,28,29 The largest study included 100 participants23,12 (60%) studies of clinical presentation12,23,27, and 3 (18%) studies of clinical course12,24,26 included ≥50 participants. The proportion of DVAs with associated CMs ranged from 2% to 40% in the 11 papers that reported their occurrence.12,13,16,23–26,28–30,32 Radiological methods used to diagnose DVAs varied to such an extent within and between studies that we could not set a diagnostic threshold: methods included CT, CT venography, MRI, MR angiography/venography, intra-arterial digital subtraction angiography, postmortem, or biopsy. Of those studies using a consistent method of diagnosis for all participants, MRI was the most commonly used modality (60% of all studies).
Clinical Presentation

Fifteen studies reported the clinical presentation of a total of 714 participants (Table 1),11–13,16,22–32 of whom at least 10% had an associated CM. Some studies were not generalizable because, for example, they were restricted to cerebellar DVAs29 or included DVAs that had arterial components.22 There were insufficient details in several papers to adequately establish whether the DVA was symptomatic; in one paper, it was unclear whether “weakness” could be classified as focal neurological deficit22; in some cases, authors were not explicit about whether the hemorrhages were symptomatic, and in others, symptoms may have been related to an associated CM. In other studies, the distinction among prepresentation hemorrhages, presenting symptomatic hemorrhages, and those occurring after diagnosis was not always clear.12,13,24

The primary mode of DVA clinical presentation was insufficiently described (“unclear”) for 23% overall; 61% presented with symptoms entirely incidental to their DVA, 6% presented with symptomatic hemorrhage, 4% presented with an epileptic seizure, and ≤1% presented with symptomatic infarction (Table 1). Symptomatic hemorrhages were uncommon at presentation overall (6%), and 3 studies—in which 18%,27 37%,22 and 43%,11 of participants presented with symptomatic hemorrhage—heavily influenced this pooled estimate (Table 1). One of these studies reported a subarachnoid hemorrhage caused by a DVA,27 whereas the other bleeds (including one due to a DVA with an arterial component22) were intraparenchymal. Two studies found that infratentorial DVAs were more likely to present with intracranial hemorrhage22,33; this association remained when we used all the available data to test it (OR, 2.9; 95% CI, 1.4 to 5.9; Table 2), but the data were heavily influenced by just one study22 whose omission removed any apparent association (OR, 1.4; 95% CI, 0.5 to 3.6).

Clinical Course

Eight studies reported the clinical course of a total of 422 participants with DVAs (Table 3),12,13,16,22,24,26,28,29 Only 2 studies were prospective,24,26 and the remainder were retro-
or the method of follow-up was not specified; the 3 largest studies also calculated “lifetime” hemorrhage rates. The mean length of follow-up ranged from 2.5 to 4.2 years (the median of the available means was 3.6 years). One study quoted neither average nor total person-years of follow-up. None of the studies reported cerebral infarction or death attributable to a DVA. Retrospective “lifetime” hemorrhage rates ranged from 0.15% to 0.61% per year, but asymptomatic hemorrhages were included among these. Symptomatic hemorrhages occurred during follow-up in only 2 of the 8 studies (and in both, undocu-
mented CMs may have been the cause); 3 (0.7%) of all 422 participants experienced symptomatic hemorrhage for annualized hemorrhage rates of 0.34% per year to 1.28% per year in the individual studies (a pooled rate could not be calculated because only 2 studies documented their total person-years of follow-up).

### Table 2. Exploration of the Association Between DVA Location and Symptomatic Hemorrhage at Presentation

<table>
<thead>
<tr>
<th>Study</th>
<th>Total in Each Study</th>
<th>No. Causing Symptomatic ICH</th>
<th>OR (Infratentorial Versus Supratentorial) (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fuji et al22</td>
<td>25</td>
<td>4 (16)</td>
<td>1.6 (2.6–52.0)</td>
</tr>
<tr>
<td>Malik et al11</td>
<td>11</td>
<td>5 (45)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>Hendrich et al27*</td>
<td>36</td>
<td>6 (17)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>Naff et al26</td>
<td>45</td>
<td>1 (2)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>Garner et al12</td>
<td>83</td>
<td>1 (1)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>Buhl et al13</td>
<td>23</td>
<td>1 (4)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>Total</td>
<td>223</td>
<td>18 (8)</td>
<td>1.4 (0.5–3.6)</td>
</tr>
<tr>
<td>SIVMS</td>
<td>56</td>
<td>0</td>
<td>OR incalculable</td>
</tr>
</tbody>
</table>

*Figures only available for intraparenchymal hemorrhages. ICH indicates intracranial hemorrhage.

### Table 3. The Clinical Course of DVAs in Studies With >20 Participants

**Clinical (Symptomatic) Outcomes During Follow-Up Attributable to DVA**

<table>
<thead>
<tr>
<th>Study</th>
<th>Study Design</th>
<th>Participants With &gt;1 DVA</th>
<th>Associated CM</th>
<th>Follow-Up Duration</th>
<th>ICH</th>
<th>Hemorrhage Rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Garner et al12</td>
<td>Retrospective and lifetime</td>
<td>100</td>
<td>2 (2)</td>
<td>2.5 (7)</td>
<td>0</td>
<td>0.22</td>
</tr>
<tr>
<td>McLaughlin et al24</td>
<td>Prospective and lifetime</td>
<td>80</td>
<td>3 (4)</td>
<td>3.6 (7)</td>
<td>1* (1.3)</td>
<td>0.34 (0.1–1.9)</td>
</tr>
<tr>
<td>Naff et al26</td>
<td>Prospective and lifetime</td>
<td>63</td>
<td>8 (13)</td>
<td>4.2 (1.0–7)</td>
<td>0</td>
<td>0.15†</td>
</tr>
<tr>
<td>Huber et al28</td>
<td>Retrospective</td>
<td>43</td>
<td>17 (40)</td>
<td>7 (2.0–10.0)</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Fujii et al27</td>
<td>Not specified</td>
<td>40†</td>
<td>7</td>
<td>3.9 (0.0–16.0)</td>
<td>2§ (5.0)</td>
<td>1.28 (0.4–4.5)</td>
</tr>
<tr>
<td>Buhl et al13</td>
<td>Not specified</td>
<td>35</td>
<td>4 (11)</td>
<td>2.8 (0.3–6.0)</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Kovacs et al29</td>
<td>Retrospective</td>
<td>32†</td>
<td>2 (6)</td>
<td>3.5 (2.0–9.0)</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Rigamonti et al16</td>
<td>Retrospective</td>
<td>29</td>
<td>2 (7)</td>
<td>3.8 (1.5–8.7)</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Totals</td>
<td></td>
<td>422</td>
<td>38 (9)</td>
<td>3.5 (0.0–16.0)</td>
<td>3 (0.7)</td>
<td>0.</td>
</tr>
<tr>
<td>SIVMS</td>
<td>Prospective</td>
<td>93</td>
<td>19 (20)</td>
<td>5.3 (0.0–8.8)</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

*This may have been due to an undocumented CM.
†Rate includes “asymptomatic” hemorrhages.
‡Three DVAs had arterial components.
§Both were recurrent bleeds; it is unclear whether one of these DVAs may have had an arterial component.
¶This was calculated based on an estimated total person years of follow-up (157 years) deduced from the mean length of follow-up multiplied by the no. of participants.
††Every DVA was located in the cerebellum.
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Most of the infratentorial DVAs were located in the cerebellum (n=31 of 38 [82%]), and the remaining 7 were in the brainstem. The modalities used for DVA diagnosis were MRI (62%), contrast-enhanced CT (29%), intra-arterial digital subtraction angiography (6%), postmortem (2%), and biopsy (1%). Nineteen adults (20%) had CM in close proximity to the DVA diagnosed by MRI (n=17) or pathological examination (n=2).

Clinical Presentation

Ninety-one of the 93 participants had presented with medical attention with symptoms that appeared to be entirely incidental to the DVA: headache (n=23), neurological symptoms unrelated to a DVA or CM (n=20), nonneurological symptoms (n=14), epileptic seizure(s) (n=10), symptoms attributable to an associated CM (n=7), detected during cancer staging imaging (n=5), unrelated cerebral infarction (n=4), tinnitus (n=4), unrelated intracranial hemorrhage (n=2), and cognitive impairment (n=2). The 2 remaining participants presented with symptoms that appeared to be anatomically referable to their DVA.

A 49-year-old woman (SIVMS ID 292) presented with sudden loss of balance, diminished sensation around her upper lip and nose, left facial weakness, and dysesthesia of her right thigh; MRI with gadolinium contrast administration (Figure 1) revealed a pontine hemorrhage, originally thought to be due to a CM, but surgical evacuation of the hematoma (Figure 1) revealed a pontine hemorrhage due to DVAs (Table 1),11,22,27 their findings may be explained by selection, investigation, and reporting biases (for example, none of them used MRI and consequently did not report whether there were associated CMs and one of them included DVAs with arterial components22).

Our study is the only prospective population-based analysis of DVAs, uses multiple overlapping sources of case ascertainment in a population of 5.1 million,15 and has recruited the second largest cohort of DVAs with the longest duration of follow-up (Table 3). Our finding of a benign prognosis is consistent with most of the large, published studies (Table 3), but even then, we are likely have overestimated the severity of DVA prognosis because of the asymptomatic reservoir of DVAs in the population.9 It is impossible to rule out a bias toward underreporting of symptomatic DVAs due to the incomplete investigation of intracranial hemorrhage in everyday clinical practice, the occasional difficulty of establishing a DVA diagnosis with certainty (Figure 1), or the obliteration of a DVA by the hemorrhage it caused (although this would not alter the low risk of recurrent hemorrhage because there would be nothing left to bleed from).

Neurological events that have been attributed to DVAs may have been due to venous hypertension,5 undetected cavernous malformations,5 rare “mechanical” or flow-related complications,10 or they may have been completely coincidental. There is anecdotal experience in SIVMS of an apparently symptomatic infarct later turning out to be incidental; one patient’s infarct initially appeared to be due to an...
anatomically related DVA, but follow-up revealed a different mechanism of infarction to be more likely (Figure 2); another patient presented with adult-onset focal epileptic seizures and an anatomically related DVA was considered as the cause, but at postmortem, a white matter hamartoma was discovered. Furthermore, the prevalences of DVAs and spontaneous intracerebral hemorrhages without an apparent cause (so-called “primary”) make a chance association a possibility.

**Summary**

Although this systematic review will have been affected by the selection, investigation, and reporting biases in this area of clinical research, its summary data are in agreement with the largest and longest prospective population-based study on DVAs. Solitary DVAs are very rarely associated with symptoms, which may be a chance association. When identified, DVAs appear to have a benign short-term prognosis.

**Acknowledgments**

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**Disclosures**

None.

**References**

Supplemental Appendix

Electronic Search Strategies

**OVID MEDLINE From 1950**
1. central nervous system venous angioma/
2. Cerebral Veins/ab, pp [Abnormalities, Physiopathology]
3. *(hemangioma/ or veins/)
4. ((vein$ or venous) adj5 (anomal$ or angioma$ or hemangioma$ or hemangioma$ or malformation$ or abnormal$)).tw.
5. 3 or 4
6. exp brain/ or central nervous system/ or exp cerebral veins/ or exp cerebrovascular disorders/
7. (brain$ or cerebral or intracerebral or central nervous system or intracranial or cerebellar or intraventricular or supratentorial or medullary or cerebrovascular$).tw.
8. 6 or 7
9. 5 and 8
10. 1 or 2 or 9
11. limit 10 to humans
12. (exp child/ or exp infant/ or adolescent/) and exp adult/
13. exp child/ or exp infant/ or adolescent/
14. 13 not 12
15. 11 not 14

**EMBASE From 1980**
1. brain vein/
2. brain hemangioma/ or vein malformation/ or brain malformation/ or cerebrovascular malformation/ or congenital blood vessel malformation/ or central nervous system malformation/
3. 1 and 2
4. (hemangioma/ or angioma/) and brain vein/
5. (*hemangioma/ or *angioma/) and *vein/
6. *vein malformation/
7. ((vein$ or venous) adj5 (anomal$ or angioma$ or hemangioma$ or hemangioma$ or malformation$ or abnormal$)).tw.
8. 5 or 6 or 7
9. exp brain/ or central nervous system/ or exp cerebrovascular disease/
10. (brain$ or cerebral or intracerebral or central nervous system or intracranial or cerebellar or intraventricular or supratentorial or medullary or cerebrovascular$).tw.
11. 9 or 10
12. 8 and 11
13. 3 or 4 or 12
14. limit 13 to human
15. (exp child/ or exp adolescent/ or newborn/) and (adult/ or middle aged/ or exp aged/)
16. exp child/ or exp adolescent/ or newborn/
17. 16 not 15
18. 14 not 17
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