Approximately 50% of patients with AVMs will present with hemorrhage, resulting in morbidity in 30%–50% of cases and death in 10% of cases. Although hemorrhage is the most common presentation modality among patients with AVMs, hydrocephalus after AVM rupture, evaluated in the form of EVD placement and long-term shunt dependence, has not been well studied. In this study we evaluated the rates of EVD placement and shunt dependence, and risk factors for them, in a cohort of patients with ruptured AVMs.

Methods

With approval from the institutional review board of Brigham and Women's Hospital, the records of a consecutive series of patients with AVMs who were evaluated between January 2005 and November 2012 at our institution were reviewed. Patients with dural arteriovenous shunts, spinal arteriovenous shunts, and cavernous malformations were excluded, along with those with hemor-rheatic AVMs presenting with lethal, massive hemorrhage (GCS score of 3 and bilateral fixed and dilated pupils with no intervention performed).

Data Extraction

From a detailed chart review, the following information was extracted: patient age, sex, initial GCS score, hemorrhage pattern (intraparenchymal, subarachnoid, and/or intraventricular), AVM Spetzler-Martin grade (size, location, drainage), associated aneurysms, and whether the patient underwent early surgery to treat the AVM and/or evacuate the hemorrhage (within 2 weeks). Trace SAH in a single convexity sulcus was not included as a formal subarachnoid bleed for our analysis.

Our criteria for EVD placement mirror those for pa-
tients with aneurysmal SAH—nonpharmacological drowsiness and/or obtundation as a manifestation of hydrocephalus. The presence of headache or intraventricular blood were not, in themselves, indications for CSF diversion. We noted the frequency of EVD placement and resultant complications (infection, hemorrhage), permanent shunt dependence, and long-term outcome (measured as the patient’s mRS score at follow-up).

**Statistical Analysis**

The statistical program Stata version 12.0 was used to study potential risk factors for EVD placement and long-term shunt dependence. In addition, the impact of these factors on long-term follow-up mRS score was also evaluated. Univariate linear regressions were performed for the following factors: patient age, sex, initial GCS score, the presence of SAH, IVH, AVM grade, nidus size, deep venous drainage, AVM location (superficial vs deep supratentorial vs infratentorial), associated aneurysms, and early surgery (within 2 weeks). Subsequent multivariate linear regressions were performed, incorporating any factors nearing significance (p < 0.2), using patient age, sex, and initial GCS score empirically as covariates.

**Results**

**Patient Characteristics**

Of 142 patients with AVMs, 87 (61%) presented with hemorrhage. None of the 55 patients with unruptured AVMs required CSF diversion due to obstructive hydrocephalus or other pathophysiology associated with their vascular malformation. The remaining cohort of 87 patients with ruptured AVMs had a mean age of 46 ± 17.4 years. The male to female ratio was 1.2:1. Of 87 hemorrhages, 73 (84%) had an intraparenchymal component, 48 (55%) an intraventricular component, and 13 (15%) a subarachnoid component. Fifty-nine AVMs (68%) were supratentorial and superficial, 12 (14%) were supratentorial and deep, and 16 (18%) were infratentorial. The mean nidus size was 2.4 ± 1.5 cm. Thirty-three AVMs (38%) had associated deep venous drainage, and 23 (26%) had associated feeding artery or angiographically visualized intranidal aneurysms. The mean Spetzler-Martin grade was 2.3 ± 1.0. Fifty-eight patients (67%) underwent early surgical treatment of their AVM along with evacuation of their intraparenchymal hematoma, if present.

Thirty-eight patients (44%) required placement of an EVD, and 16 required permanent CSF diversion (18% of entire cohort, 42% of those undergoing initial ventriculostomy). Of 38 cases with ventriculostomy placement, 4 (11%) resulted in asymptomatic intraparenchymal hematomas, and 3 (8%) became infected. No asymptomatic hemorrhages occurred, and no infectious or hemorrhagic complications occurred after the 16 permanent shunt procedures.

**Univariate and Multivariate Analysis**

Univariate analysis of risk factors for EVD placement revealed GCS score (OR 0.86, 95% CI 0.78–0.95, p = 0.002), IVH (OR 6.46, 95% CI 2.44–17.1, p < 0.001), associated aneurysms (OR 4.86, 95% CI 1.74–13.53, p = 0.002), and early surgery (OR 2.22, 95% CI 0.87–5.69, p = 0.01) to be statistically significant (Table 1). Patient age, sex, associated SAH, AVM grade, AVM size, deep venous drainage, and AVM location were not significant. Multivariate analysis revealed only associated aneurysms to be a statistically significant risk factor for EVD placement (OR 20.96, 95% CI 2.41–181.9, p = 0.006).

Statistically significant risk factors for shunt placement included initial GCS score (OR 0.84, 95% CI 0.75–0.94, p = 0.003), IVH (OR 7.62, 95% CI 1.61–36, p = 0.01), deep supratentorial location (OR 4.16, 95% CI 1.12–15.46, p = 0.034), and associated aneurysms (OR 3.44, 95% CI 1.11–10.6, p = 0.03; Table 1). Patients with ruptured, supratentorial, superficial AVMs had a lower risk of shunt dependence (OR 0.28, 95% CI 0.093–0.87, p = 0.028). Patient age, sex, associated SAH, AVM grade, AVM size, deep venous drainage, and early surgery were not associated with long-term shunt dependence. Multivariate analysis revealed initial GCS score to be a statistically significant risk factor (OR 0.76, 95% CI 0.59–0.99, p = 0.041) as well as a strong trend for associated aneurysms (OR 6.31, 95% CI 0.93–43.1, p = 0.06).

**Impact on mRS Score**

A univariate and multivariate analysis was performed to evaluate the potential impact of each risk factor on follow-up mRS score (Table 2). Shunt placement was also included as a means to evaluate its relationship to long-term outcome. Statistically significant risk factors from univariate analysis were increasing age (p < 0.001), lower initial GCS score (p < 0.001), and the placement of a shunt (p = 0.022). Multivariate analysis continued to demonstrate older age as a significant risk factor (p = 0.02), and a trend for the placement of a shunt (p = 0.26).

**Discussion**

Cerebral AVMs are a considerable source of neurological morbidity as a result of hemorrhages, seizures, and/or steal phenomena. In contrast to aneurysm rupture in which morbidity and mortality are largely attributable to hydrocephalus and infrequently from local mass effect on eloquent parenchyma from intraparenchymal hemorrhage, both mechanisms are at play with AVM rupture. Prospective studies evaluating AVM rupture generally cite annual rates from 2.4% to 4.6%, with prior hemorrhage consistently serving as a significant risk factor for hemorrhage along with deep location, deep venous drainage, and associated aneurysms. A systematic approach to evaluating morbidity from AVM hemorrhage, requiring a stratification of morbidity resulting from hydrocephalus and morbidity resulting from local mass effect of hemorrhage on eloquent parenchyma, has not been conducted.

This analysis quantifies the frequency of hydrocephalus after AVM rupture and risk factors for its occurrence. Overall, 44% of patients required the placement of an EVD and 18% required permanent ventriculoperitoneal shunt placement. Intuitively, initial GCS score and the presence of IVH were significant risk factors for EVD placement.
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Initial GCS score, IVH, and associated aneurysms were also significant factors influencing shunt dependence as they were for EVD placement, with the addition of supratentorial deep AVMs. On multivariate analysis, however, only initial GCS score continued to be a significant risk factor, with a strong trend for AVMs with associated aneurysms (p = 0.06). Other AVM morphological factors (size, location, and drainage) were not associated with the risk of hydrocephalus.

The rate of long-term shunt dependence from AVM rupture in this study (18%) is comparable to the rates found in several studies evaluating hydrocephalus after aneurysm rupture. These studies primarily focused on a comparison of the risk of shunt dependence after coil- ing as compared with clipping of ruptured aneurysms and demonstrated no significant difference. In a somewhat analogous fashion, in this study early surgery was not associated with a lower risk of shunt dependence. The interesting finding in our study that associated aneurysms increase this risk has important therapeutic implications because it suggests that rupture from these AVMs may be more morbid. This data adds to the current literature that already reinforces the fact that the risk of hemorrhage is higher for these lesions. Taken together, these findings encourage the timely treatment of these AVMs to not only combat the greater risk of hemorrhage from these lesions, but also potentially greater associated morbidity from hemorrhage. Importantly, our final analysis reinforced the fact that hydrocephalus and shunt placement after AVM rupture has significant clinical implications, as shunt dependence was associated with worse long-term outcome as measured by mRS score.

**Conclusions**

The overall rate of early EVD placement was 44% and the rate of shunt dependence was 18% after AVM rupture. Significant risk factors for both were initial GCS score, IVH, and associated aneurysms. Arteriovenous malformations with associated aneurysms thus not only have a greater risk of rupture but also a greater potential risk of permanent morbidity from rupture as a result of hydrocephalus, encouraging their timely treatment.
Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper.

Author contributions to the study and manuscript preparation include the following. Conception and design: Du, Gross. Acquisition of data: Gross, Lai. Analysis and interpretation of data: all authors. Drafting the article: Gross. Critically revising the article: all authors. Reviewed submitted version of manuscript: all authors. Approved the final version of the manuscript on behalf of all authors: Du. Statistical analysis: Lai. Administrative/technical/material support: Du. Study supervision: Du.

References


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