

Reliable Identification of Benign Clinical Course in Aneurysmal Subarachnoid Hemorrhage: A Simple and Qualitative Clinical Algorithm

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Introduction

While a number of grading systems have been previously described to help guide clinical decision making in the setting of aneurysmal subarachnoid hemorrhage (aSAH), a method to reliably predict low vasospasm risk in aSAH patients has not been proposed. We developed a simple qualitative clinical algorithm that combines admission clinical severity, defined by Hunt Hess grade, and subarachnoid blood distribution, based on the Hijdra sum scoring system, to reliably identify patients at low risk of clinical vasospasm.

Objective

We developed a simple qualitative clinical algorithm that combines admission aSAH clinical severity, defined by Hunt Hess grade, and subarachnoid blood distribution, based on the Hijdra sum scoring system, to reliably identify patients at low risk of clinical vasospasm.

Hypothesis

We hypothesized that a simple to use qualitative clinical algorithm to reliably identify low risk of clinical vasospasm could be developed by analyzing clinical and systematically evaluating radiographic risk factors on admission.

Methods

Clinical severity, admission non-contrasted head computed tomography scans (CTH), and incidence of radiographic and clinical vasospasm among 214 aSAH patients treated at our institution were evaluated. Admission CTH's were systematically assessed for several different distributions of cisternal and ventricular blood. A final clinical algorithm was developed. Patients who satisfied all of the following 4 criteria experienced considerably lower risk of vasospasm. 1) Hunt Hess grade 1-2. 2) Lack of thick subarachnoid blood filling two adjacent cisterns. 3) Lack of thick interhemispheric blood. 4) Lack of biventricular IVH.

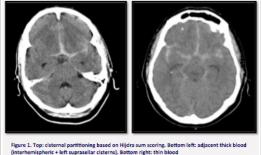
Results

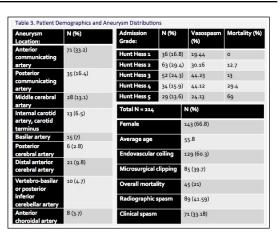
Eighty-nine patients (41.6%) developed clinically silent vasospasm, seventy-one patients (33.2%) developed vasospasm with neurological deficit, and forty-five patients expired (21%). Adjacent cistern blood (OR 4.13, 95% confidence interval [CI] 2.1-8.09), interhemispheric thick blood (OR 6.39, 95% CI 3.17-12.87), and biventricular IVH (OR 2.05, 95% CI 1.04-4.04) were all statistically significant risk factors. Retrospective application of our proposed clinical algorithm yielded a sensitivity of 40% (95% CI 28.47-52.41%), specificity of 100% (95% CI 83.89-100%), positive predictive value of 100% (95% CI 87.66-100%), and negative predictive value of 33.3% (95% CI 21.95-46.34%). Inter-observer variability was substantial at k0.79.

Aneurysmal subarachnoid hemorrhage	
Yes	
Hunt Hess Grade 1 or 2	
Yes	No
Presence of any of the following: 1) Interhemispheric thick* blood 2) Adjacent cistern thick* blood 3) Bilateral IVH**	Routine vasospasm surveillance
No	
Early transfer from ICU to step down unit ^a	
Hyperdense filling of any of the following cisterns based on the	
nt suprasellar, interhemispheric, left or right ambient, and quadri pendicular to the direction of the course of eistern. Biventricular IVH was defined as presence of blood in both later - 48 hours after aneurysm secured.	

Risk Factor:		Spasm	No spa:	-	Odds rat	la (arth	P-value	
KISK PACCOT		opesm	но зра	-11	confiden	ce interval)	Pevalue	
Both suprasellar thick		22	27		1.9 (0.96	3.77)	0.066	
Not both suprasellar thick		36	84					
Adjacent thick		38	35		4.13 (2.1-	8.09)	<0.001	
Not adjacent thick		20	76					
Interhemispheric thick		37	24		6.39 (3.17	-12.87)	<0.001	
Not interhemispheric thick		21	87					
Bilateral IVH		41	60		2.05 (1.0)	L-4.04)	0.038	
Not Bilateral IVH		17	51					
Either adjacent or interhemispheric thick		47	42		7.02 (3.28		<0.001	
Neither adjacent nor interhemispheric thick		11	69					
Either adjacent or interher or IVH	nispheric	56	75		23.44 (3.2	-58.18)	<0.001	
Neither adjacent nor interhemispheric nor IVH		2	36					
Table 2. Multivariate	Logistic	Regress	ion					
Parameter	Coe	fficient		Odds Ratio)	95% Confide		p-value
Intercept	-0.7	98						
Adjacent thick	0.8	32		2.29		1.07-4.9	4	0.03
rugueene ennen								
Interhemispheric thick	1.43	36		4.2		1.94-9.1		<0.01







Conclusions

Application of the final clinical algorithm produced successful identification of aSAH patients who experience effectively zero risk of clinical vasospasm. Our algorithm is simple to apply with high reliability. Prospective application of our algorithm has considerable clinical and economic implications.

Learning Objectives

By the conclusion of this session, participants should be able to 1) Understand the derivation of the proposed clinical algorithm and 2) Apply the proposed clinical algorithm at home institutions for identification of subgroups of low-risk vasospasm patients.