

Hemorrhagic adenoma mimicking anterior communicating artery aneurysm

Ícaro Araújo-de Sousa, Arthur de Oliveira-Veras, Octávio Marques-Pontes-Neto, Elizeu Pereira dos Santos-Neto

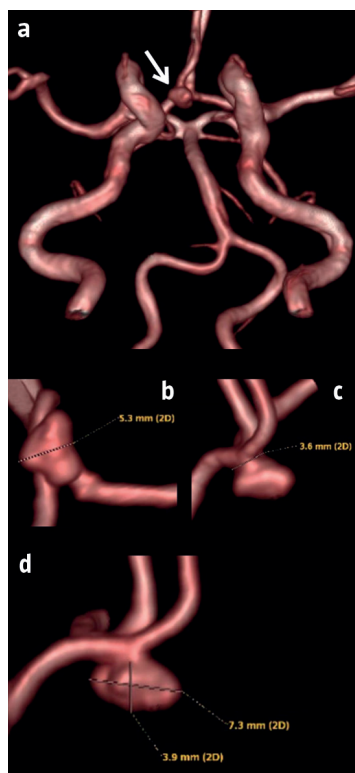


Figure 1. a) Magnetic resonance angiography showing an image suggestive of aneurysm in the anterior communicating artery complex (white arrow); b-d) Dimensions of the presumed aneurysm and its neck.

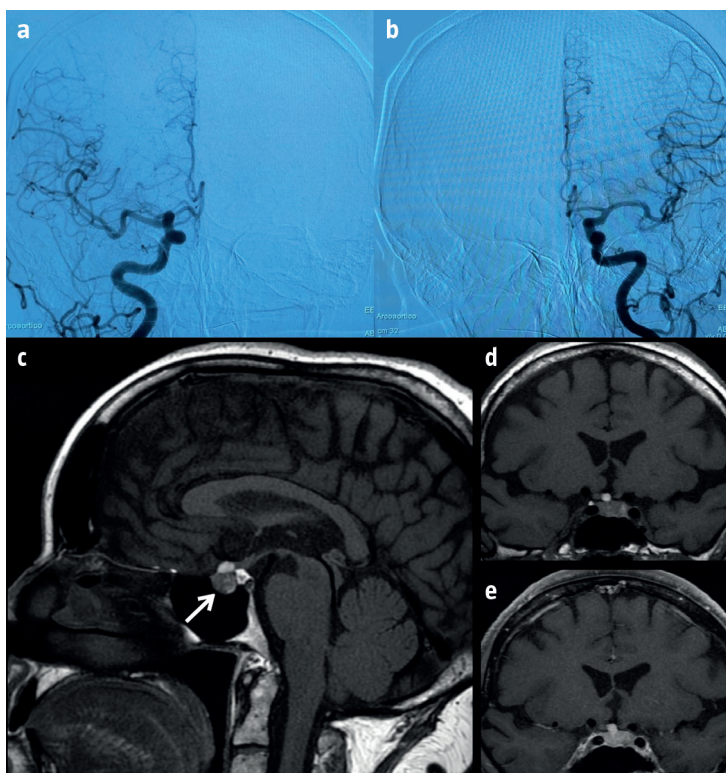


Figure 2. Digital Angiography subtraction. A and b) Posteroanterior projection showing no evidence of aneurysm. Thin-slice pituitary magnetic resonance imaging; c) Sagittal projection demonstrating a lesion (white arrow) with hypersignal on T₁-weighted image in the projection of the pituitary stalk adjacent to the A2 segments of the anterior cerebral arteries; d and e) Coronal projection before and after contrast enhancement showing hemorrhagic adenoma.

A patient in their 50s with no previous comorbidities and a significant family history of cerebral aneurysms in three first-degree relatives underwent magnetic resonance angiography, which revealed an image suggestive of an anterior communicating artery aneurysm (Fig. 1). A digital subtraction angiography was performed for therapeutic planning. The exam was conducted with 3D-studies of the carotid

arteries to increase sensibility but did not show any evidence of an aneurysm (Fig. 2). A contrast-enhanced fine-cut magnetic resonance imaging of the sella turcica showed a hyperintense lesion on the T₁-weighted image in the adenohypophysis, consistent with hemorrhagic adenoma diagnosis (Fig. 2). A comprehensive hormonal screening was performed, and no alterations were found.

Differentiating between a pituitary adenoma and an aneurysm is vital, as these two have different management options. In contrast to most cases described in the literature, where internal carotid aneurysms mimic pituitary adenomas, our report presented a distinct pattern in which a hemorrhagic adenoma initially simulated an aneurysm in the anterior communicating artery complex [1-3]. Sellar

Department of Neuroscience and Behavior Sciences. Medical School of Ribeirão Preto. Universidade de São Paulo. Ribeirão Preto (Í. Araújo-de Sousa, A. de Oliveira-Veras, O. Marques-Pontes-Neto). Institute of Radiology. Hospital das Clínicas. School of Medicine. Universidade de São Paulo. São Paulo (E. Pereira dos Santos-Neto). Department of Neurology. Hospital Getúlio Vargas. Teresina, Brazil (E. Pereira dos Santos-Neto).

Correspondence:

Dr. Ícaro Araújo de Sousa. Department of Neuroscience and Behavior Sciences. Medical School of Ribeirão Preto. University of São Paulo. Av. Bandeirantes, 3900. Ribeirão Preto, São Paulo, Brazil. CEP 14048-900.

E-mail:

icaroas@hcrp.usp.com.br

ORCID:

0000-0002-9452-1207 (Í.A.S.).

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magnetic resonance imaging is considered the imaging modality of choice for the evaluation of lesions in the sella turcica and can provide the size and shape of the adenomas, the presence of cysts or bleeding within the tumor, its relationship with the optic pathways and surrounding structures and differential diagnosis with other sellar lesions. Furthermore, a fluid-fluid level, a hypointense rim on T₂-weighted images, septation, and an off-midline location were more specific for pituitary adenomas [4].

Specifically, regarding the sellar lesions with high T₁ signal, the range of differential diagnosis is narrow and can be divided according to the substance causing the hyperintensity. Thus, clotting of blood can cause a high signal in hemorrhagic pituitary adenoma, like ours, pituitary apoplexy, and thrombosed aneurysm. Furthermore, cystic

sellar lesions can be enhanced due to the high protein concentration, like Rathke cleft and pars intermedia cysts. The first has a closer relationship with the infundibulum and sometimes can mimic pituitary adenoma with suprasellar extension. The second usually has a posterior location and inferoposterior relationship to the anterior communicating artery [5]. Moreover, fat (e.g., dermoid cyst) and calcification (e.g., chondroid tumor) can cause high T₁ signal as well; and some tumors may have more than one source for its signal change (e.g., craniopharyngioma).

Thus, this atypical presentation highlights the diagnostic complexity associated with bright lesions on sellar T₁-weighted scans and emphasizes the importance of detailed and differential assessment to ensure accurate diagnoses and adequate clinical management.

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