

Síndrome do Seio Cavernoso: Um diagnóstico Pouco Frequent

Cavernous Sinus Syndrome: An Unusual Diagnosis

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RESUMO

A síndrome do seio cavernoso caracteriza-se pela presença de oftalmoplegia, diplopia, hipostesia facial, síndrome de Horner e proptose. Os autores apresentam o caso clínico de uma mulher, 89 anos, que se apresentou no Serviço de Urgência com queixas de vômitos, cefaleia frontotemporal esquerda, fotocopia, ptose esquerda e diplopia com cerca de 24 horas de evolução. O exame neurológico revelou a presença de ptose e alteração nos movimentos oculares ao nível do olho esquerdo. A tomografia computorizada e ressonância magnética crânio-encefálica revelaram a presença de lesão expansiva na região supraselar e seio cavernoso esquerdo, sugestiva de macroadenoma com sinais de transformação hemorrágica. A síndrome do seio cavernoso relaciona-se com diversas etiologias, tais como, doenças cerebrovasculares, infecciosas e trauma. Para além da história clínica e exame objetivo, os exames imagiológicos são essenciais para o diagnóstico e determinação da etiologia desta síndrome.

PALAVRAS-CHAVE: Adenoma; Neoplasias Pituitárias; Seio Cavernoso

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Received/Received: 2022/11/13 - Aceite/Accepted: 2023/06/19 - Publicado online/Published online: 2023/09/06 - Publicado/Published: 2023/09/30
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ABSTRACT

Cavernous sinus syndrome is characterized by ophthalmoplegia, diplopia, facial hypoesthesia, Horner syndrome and proptosis.

The authors present a clinical case of an 89-year-old woman presented to the emergency department with complaints of vomiting, severe left frontotemporal headache, photophobia, left ptosis and diplopia with less than 24 hours of evolution. Neurological examination revealed ptosis and limitation of all eye movements in the left eye. Cranioencephalic computed tomography and magnetic resonance imaging showed an expansive lesion with expression in the suprasellar cistern and in the left cavernous sinus, suggestive of a macroadenoma with signs of hemorrhagic transformation.

Cavernous sinus syndrome has several etiologies, as cerebrovascular diseases, infectious diseases, and trauma. In addition to the clinical history and physical examination, imaging tests are essential to determine the etiology behind this syndrome.

KEYWORDS: Adenoma; Cavernous Sinus; Pituitary Neoplasms

An 89-year-old woman went to the emergency department with complaints of vomiting, severe left frontotemporal headache, photophobia, left ptosis and diplopia with less than 24 hours of evolution.

Neurological examination revealed ptosis and limitation of all eye movements in the left eye. She also presented hypoesthesia in the territory of the maxillary branch of the left trigeminal nerve, with no other focal alterations.

Cranioencephalic computed tomography (CT) and magnetic resonance imaging (MRI) showed an expansive lesion with expression in the suprasellar cistern and in the left cavernous sinus, suggestive of a macroadenoma with signs of hemorrhagic transformation." (Fig. 1).

The diagnosis of cavernous sinus syndrome (paresis of left cranial nerves III, IV and V2) was made, resulting from a macroadenoma with cavernous sinus invasion. The patient underwent endoscopic transphenoidal resection of the macroadenoma, with favorable clinical and imaging course and was later discharged from the hospital.

The cavernous sinus syndrome (CSS) is a set of signs and symptoms, including ophthalmoplegia, diplopia, facial hypoesthesia, Horner syndrome and proptosis.¹ Involvement of the III and IV cranial nerves can result in ophthalmoplegia and diplopia, while the involvement of the ophthalmic and maxillary divisions of the trigeminal nerve can result in altered sensitivity on the face.² CSS have several etiologies, as cerebrovascular diseases, infectious diseases, and trauma.¹

In addition to the clinical history and physical examination, imaging tests, especially MRI with a study aimed at cavernous sinus, are essential to determine the etiology behind this syndrome,³ as was the case with this patient.

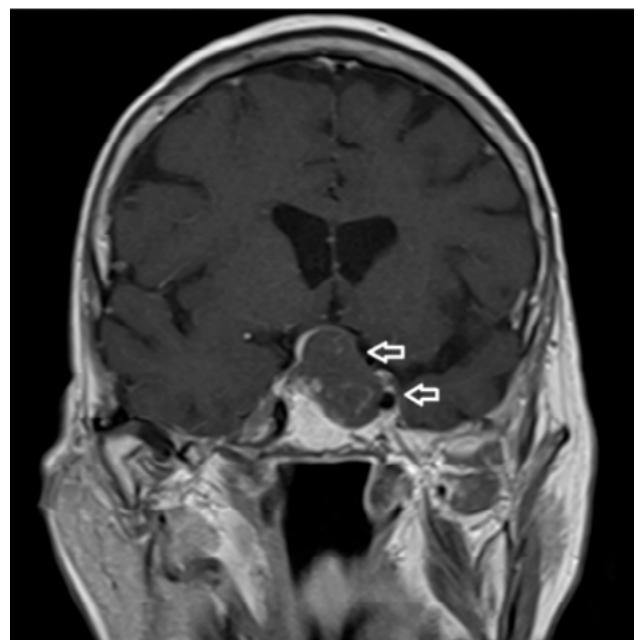


FIGURA 1. Brain magnetic resonance imaging (MRI) - T1-weighted coronal (B): "Voluminous pituitary adenoma with expression in the suprasellar cistern and left cavernous sinus and signs of hemorrhagic transformation (arrows)."

DECLARAÇÃO DE CONTRIBUIÇÃO/CONTRIBUTORSHIP STATEMENT

JA e MB: Escrita e revisão

AG: Pesquisa

LL: Revisão

JA and MB: Writing and review

AG: Research

LL: Review

RESPONSABILIDADES ÉTICAS

CONFLITOS DE INTERESSE: Os autores declaram a inexistência de conflitos de interesse na realização do presente trabalho.

FONTES DE FINANCIAMENTO: Não existiram fontes externas de financiamento para a realização deste artigo.

CONFIDENCIALIDADE DOS DADOS: Os autores declaram ter seguido os protocolos da sua instituição acerca da publicação dos dados de doentes.

CONSENTIMENTO: Consentimento do doente para publicação obtido.

PROVENIÊNCIA E REVISÃO POR PARES: Não comissionado; revisão externa por pares.

ETHICAL DISCLOSURES

CONFLICTS OF INTEREST: The authors have no conflicts of interest to declare.

FINANCING SUPPORT: This work has not received any contribution, grant or scholarship.

CONFIDENTIALITY OF DATA: The authors declare that they have followed the protocols of their work center on the publication of data from patients.

PATIENT CONSENT: Consent for publication was obtained.

PROVENANCE AND PEER REVIEW: Not commissioned; externally peer reviewed.

REFERENCES

1. Nambiar R, Nair SG. Cavernous sinus syndrome. Bayl Univ Med Cent Proc. 2017;30:455-456. doi: 10.1080/08998280.2017.11930227.
2. Munawar K, Nayak G, Fatterpekar GM, Sen C, Zagzag D, Zan E, et al. Cavernous sinus lesions. Clin Imaging. 2020;68:71-89. doi: 10.1016/j.clinimag.2020.06.029.
3. Bhatkar S, Mahesh KV, Sachdeva J, Goel A, Goyal MK, Takkar A, et al. Magnetic resonance imaging (MRI) versus computed tomographic scan (CT scan) of brain in evaluation of suspected cavernous sinus syndrome. Neuroradiol J. 2020;33:501-7. doi: 10.1177/1971400920970921.