

Opsoclonus investigation: how to proceed?

Investigação de opsoclonus: como proceder?

Roberto Battistella¹

1. Setor de Neuroftalmologia, Clínica Oftalmológica, Hospital das Clínicas, Faculdade de Medicina, Universidade de São Paulo, São Paulo, SP. Brazil.

A 4-month-old child presented with abnormal eve movements for 1 week and muscle twitching. Cerebrospinal fluid (CSF) analysis revealed increased cellularity and positive bacterioscopic test and negative culture results. Other complementary exam results were negative. The patient was diagnosed with parainfectious Kinsbourne syndrome (or opsoclonus--myoclonus-ataxia syndrome). Ceftriaxone and intravenous immunoglobulin administration resulted in gradual improvement.

Parainfectious Kinsbourne syndrome is a rare disorder characterized by opsoclonus (involuntary anarchic conjugate multidirectional binocular movements) and cerebellar ataxia, sometimes with a myoclonic component. The etiology varies and includes idiopathic or parainfectious (viral and bacterial) causes and metabolic, toxic, or paraneoplastic syndromes. Neuroblastoma is the primary consideration in children^{1,2}.

In adults, parainfectious Kinsbourne syndrome may continue to represent paraneoplastic disease associated with cancer (lung, breast, and ovary), brainstem hemorrhage, or multiple sclerosis.

The diagnostic investigation should include a search for tumors, paraneoplastic syndrome, and infections by performing complete ophthalmological and neurological examinations. Magnetic resonance imaging or computed tomography of the head, thorax, and abdomen; full-body positron emission tomography; CSF examination; and urinary catecholamine measurements may be necessary³.



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Treatment is directed at the underlying etiology. Immunomodulation with corticosteroids, plasmapheresis, or intravenous immunoglobulin can help improve the clinical condition¹⁻³.

REFERENCES

- Bhatia P, Heim J, Cornejo P, Kane L, Santiago J, Kruer MC. Opsoclonus-myoclonus-ataxia syndrome in children. J Neurol. 2022 Feb;269(2):750-757.
- Blaes F, Dharmalingam B. Childhood opsoclonus-myoclonus syndrome: diagnosis and treatment. Expert Rev Neurother. 2016 Jun;16(6):641-8.
- Rossor T, Yeh EA, Khakoo Y, Angelini P, Hemingway C, Irani SR, Schleiermacher G, Santosh P, Lotze T, Dale RC, Deiva K, Hero B, Klein A, de Alarcon P, Gorman MP, Mitchell WG, Lim M; OMS Study Group. Diagnosis and Management of Opsoclonus-Myoclonus-Ataxia Syndrome in Children: An International Perspective. Neurol Neuroimmunol Neuroinflamm. 2022 Mar 8;9(3):e1153.

AUTHOR INFORMATION



» **Roberto Battistella** https://orcid.org/0000-0002-3768-2004 https://lattes.cnpq.br/0709371452223526