Teaching NeuroImages:

A neuroendocrine rarity

Wolfram syndrome

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Figure Wolfram syndrome type 1 in a 32-year-old man with deafness and insulin-dependent diabetes mellitus since infancy and a 4-year progressive visual loss and episodes of hyponatremia due to polyuria and polydipsia

(A) Axial T1-weighted brain MRI shows bilateral optic nerve (red arrows) and pontine and middle cerebellar peduncle atrophy (white arrow). (B) Sagittal T1-weighted brain MRI shows brainstem (particularly the pons; red arrow) and cerebellar atrophy.

Wolfram syndrome 1 (WS1) is an autosomal recessive disorder characterized by diabetes insipidus, diabetes mellitus, optic atrophy, and deafness (DIDMOAD syndrome). It is caused by a mutation in the *WFS1* gene (chromosome 4p16.1, involved in regulation of intracellular calcium) that encodes wolframin, a transmembrane protein of pancreatic β cells. WS1 is a rare disorder characterized by the development of insulin-dependent diabetes mellitus at an average age of 6 years, followed by optic atrophy (figure) at an average age of 11 and partial central diabetes insipidus and deafness in adolescence. Additional neurologic manifestations include truncal ataxia, myoclonus, epilepsy, nystagmus, and hyposmia.^{1,2}

AUTHOR CONTRIBUTIONS

Thiago Cardoso Vale: drafting/revising the manuscript, study concept or design, analysis or interpretation of data, accepts responsibility for

conduct of research and final approval, acquisition of data, study supervision. Francisco Otaviano Lima Perpétuo: drafting/revising the manuscript, study concept or design, analysis or interpretation of data, accepts responsibility for conduct of research and final approval, acquisition of data, study supervision.

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REFERENCES

1. Rigoli L, Di Bella C. Wolfram syndrome 1 and Wolfram syndrome 2. *Curr Opin Pediatr* 2012;24:512–517.
2. Ito S, Sakakibara R, Hattori T. Wolfram syndrome presenting marked brain MR imaging abnormalities with few neurologic abnormalities. *AJNR Am J Neuroradiol* 2007;28:305–306.

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