WALL-EYED BILATERAL INTERNUCLEAR OPTHALMOPLEGIA (WEBINO SYNDROME) AND MYELOPATHY IN PYODERMA GANGRENOsum

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SUMMARY — A 35-year-old female with pyoderma gangrenosum developed paraparesis with a sensory level at L1. Three months later she complained of diplopia and was found to have bilateral internuclear ophthalmoplegia with exotropia and no ocular convergence. The term Webino syndrome has been coined to design this set of neuro-opthalmologic findings. Although it was initially attributed to lesions affecting the medial longitudinal fasciculus and the medial rectus subnuclei of the oculomotor complex in the midbrain the exact location of the lesion is still disputed. In the present case both myelopathy and Webino syndrome were probably due to vascular occlusive disease resulting from central nervous system vasculitis occurring in concomitance to pyoderma gangrenosum.

Oftalmoplegia internuclear bilateral e exotropia (síndrome de Webino) e mielopatia no pioderma gangrenosum.

RESUMO — O termo Webino é formado pelas iniciais dos componentes da síndrome (wall-eyed bilateral internuclear ophthalmoplegia), havendo também perda da convergência ocular. Relatamos o caso de uma paciente de 35 anos de idade, com pioderma gangrenosum, que desenvolveu subitamente mielopatia com nível sensitivo em L1 e, três meses depois, quadro súbito de oftalmoplegia internuclear bilateral, exotropia e perda de convergência. Há poucos casos de síndrome de Webino relatados na literatura, a maioria deles secundária a esclerose múltipla e a doença vascular cerebral. A localização da lesão responsável pela síndrome ainda não está bem estabelecida mas, acredita-se que os fascículos longitudinais e os subnúcleos dos retos mediais do complexo oculomotor no mesencéfalo sejam afetados, embora não haja ainda confirmação anatomo-patológica. O pioderma gangrenosum é condição caracterizada por úlceras de crescimento rápido e de bordas elevadas com halo eritematoso localizadas preferencialmente na cabeça, tronco e membros inferiores. Embora sua natureza ainda seja desconhecida, parece ser secundário a vasculite subjacente, estando associado a grande variedade de doenças sistêmicas. No caso presente, a ocorrência da síndrome de Webino e de mielopatia sugere que as lesões no sistema nervoso sejam de natureza vascular, provavelmente secundárias a vasculite. Este é o primeiro relato na literatura de complicações neurológicas do pioderma gangrenosum e da associação de mielopatia à síndrome de Webino.

Wall-eyed bilateral internuclear ophthalmoplegia (Webino syndrome) is a variant of Cogan’s anterior internuclear ophthalmoplegia with exotropia in primary position. It has been attributed to lesions affecting the medial longitudinal fasciculus and the medial rectus subnuclei of the oculomotor complex. Only a few cases of Webino syndrome have been reported in the literature. We describe a patient with pyoderma gangrenosum who developed bilateral internuclear ophthalmoplegia with exotropia, loss of convergence and myelopathy in the course of the disease. To our knowledge neither neurological complications of pyoderma gangrenosum nor association of Webino syndrome to myelopathy have been described previously.

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CASE REPORT

MSL, a 35-year-old female was admitted to the hospital with history of recurrent ulcers in left arm and thigh since the age of 18 years. In the last two years she developed chronic urinary infection and had been treated with antibiotics. She had also been given talidomide, chlorphazimine and DDS as treatment for pyoderma gangrenosum in another hospital, where the diagnosis was made through a skin biopsy. On admission the patient presented a large painful necrotic ulcer surrounded by an erythematous halo in the left thigh (Fig. 1) and signs of old inactive dermatological lesions on the left arm. There was no neurological or ophthalmological complaint. Physical examination was unrevealing except for the skin lesions. Admission laboratory work-up was as follows: red blood cell count 3,500,000 per cubic millimeter, hemoglobin 10.5 g/dL, hematocrit 35%, and white blood cell count 5,300 per cubic millimeter. Serum glucose, creatinine, BUN, thyroid and liver function studies and protein electrophoresis showed normal values. Serum VDRL, FTA-ABS, antinuclear antibody titers LE cell preparation were also negative as were roentgenograms of the chest and gastrointestinal series. Biopsy specimens taken from the erythematous area at the border of the lesion on patient's left thigh disclosed lymphocytic vasculitis with endothelial swelling, fibronoid necrosis and extravasation of erythrocytes and lymphocytes (Fig. 1). As the patient was being treated with azathioprine and steroids she suddenly developed paraparesis and urinary retention. Neurological consultation at this time disclosed paraparesis with a sensory level at L1. The deep tendon reflexes were hypoactive and Babinski sign could not be elicited. Lumbar puncture showed an opening pressure of 120 mmH₂O, clear cerebrospinal fluid with 3 lymphocytes per cubic millimeter, glucose content of 100 mg/dL and protein content of 12 mg/dL with normal electrophoretic fractions. Three
months later while on physical therapy she suddenly experienced left hemiparesis and diplopia. Neurological re-evaluation showed exotropia in primary position, bilateral medial rectus palsy on attempted horizontal gaze on either direction with nystagmus of the abducting eye (Fig. 2). Vertical gaze was normal and there was a loss of ocular convergence. The pupils were 3 mm in size, round and fully reactive to light. Fundoscopic examination and visual fields were normal. There were a left central facial weakness, moderate left hemiparesis and a residual weakness on the right lower limb. Muscle stretch reflexes were decreased on the left upper limb but brisk on both lower extremities. Bilateral Babinski sign was present and sensory level at L1 persisted unchanged. High quality computed tomographic scan of the head exhibited no abnormalities.

**Fig. 2 — Case MSL. Bilateral medial rectus palsy on attempted horizontal gaze and exotropia in primary position. Vertical gaze is normal.**

**COMMENTS**

Internuclear ophthalmoplegia is a well-characterized clinical syndrome manifested by paresis of adduction on attempted horizontal gaze associated with jerk nystagmus of the abducting eye. Vertical nystagmus with attempted upward gaze and loss of convergence may be concomitant features. Although the clinical description of the syndrome is attributed to Wilson in 1906, some authors date back its recognition to about 1864. The role of the medial longitudinal fasciculus (MLF) in mediation of conjugate eye movements was first suspected by Muskens as early as 1914. This suspicion was confirmed ten years later through the first pathological study of a case of internuclear ophthalmoplegia. Since then it has been well demonstrated that lesions involving the MLF connecting the pontine paramedian reticular formation (PPRF) and the ipsilateral abducens nucleus with the contralateral medial rectus subnucleus in the midbrain produce failure of the medial rectus on the side of the lesion to act in horizontal gaze and nystagmus of the abducting eye. Weakness of adduction may range from complete inability to adduct the eye beyond midline to only a mild decrease in velocity of adducting saccades without any limitation of adduction. The origin of the contralateral abducting nystagmus is still a matter of controversy. Some authors believe that it may reflect the effects of convergence, impaired inhibition of the contralateral medial rectus, interruption of descending internuclear fibers projecting to the abducens nucleus, a gaze-evoked nystagmus, or even adaption to the contralateral medial adduction weakness.

Internuclear ophthalmoplegia has been divided into anterior and posterior types. When convergence is absent concomitantly with adductor paralysis it is designated «anterior internuclear ophthalmoplegia», the lesion involving the rostral portion of the MLF conducting the impulses for convergence from the pretectal region to the third nerve nucleus. This convergence tract is indeed located anterior to the third nerve nucleus. On the other hand in «posterior internuclear ophthalmoplegia» there is preservation of convergence despite absence of voluntary adduction, and a caudal lesion sparing the medial rectus subnucleus of the oculomotor nuclear complex can be assumed. It has been established however that while the posterior internuclear ophthalmoplegia is clinically useful as a localizing sign consistently pointing to a caudal lesion, the absence of convergence in the setting of an internuclear ophthalmoplegia (the anterior type) does not necessarily imply a rostral lesion involving the medial rectus nuclear subdivision. Unilateral internuclear ophthalmoplegia is usually due to an infarct of the midbrain tegmentum; demyelinating disease, tumors
and infections are less frequent causes. On the other hand the most common cause of bilateral internuclear ophthalmoplegia is multiple sclerosis, although posterior fossa tumors, brainstem infection, Arnold-Chiari malformations with hydrocephalus, head trauma, basilar artery disease, carcinomatous meningitis, cryptococcal meningitis, systemic lupus erythematosus, intrathecal chemotherapy and cranial irradiation, syphilis, periarteritis nodosa, spinocerebellar degenerations and branched-chain ketoaminociduria. Myasthenia gravis, abetalipoproteinemia and coma caused by overdose of phenytoin, narcotics or barbiturates may produce ocular findings similar to bilateral internuclear ophthalmoplegia.

The term Webino syndrome (wall-eyed bilateral internuclear ophthalmoplegia) was coined by Lubow to describe patients with bilateral adduction paresis, absence of ocular convergence and exotropia. Lubow believed that the syndrome was caused by lesions involving the medial longitudinal fasciculus bilaterally in the midbrain together with the medial rectus subnuclei of the oculomotor complex. There have been only a few cases of Webino syndrome reported in the literature, most of them caused by occlusive vascular or demyelinating diseases. An additional case was due to a tuberculous granuloma in the midbrain that presumably affected the MLF and the medial rectus subnuclei of the oculomotor complex by mass effect. In a recent reported case of Webino syndrome caused by multiple sclerosis autopsy did not reveal any lesion in the medial rectus subnuclei of the oculomotor complex leading their authors to suppose that lesions in more than one locus may be responsible for eliciting the syndrome. Others have attributed the syndrome to lesions affecting the PPRF and the MLF but this hypothesis has not been confirmed by pathological studies. As loss of convergence is concerned a supplemental anatomic explanation may not be necessary as it commonly occurs in long-standing disruptions of ocular fusion. It has been proposed however that interruption of corticofugal fibers from frontal, parietal or occipital eye fields may play a role in the convergence dysfunction.

Pyoderma gangrenosum is an uncommon ulcerative skin condition with distinctive clinical appearance comprising a destructive burrowing ulcer with an irregular margin and a ragged purple-red overhanging edge. It typically starts as an isolated sterile pustule or nodule, often on one of the lower extremities but rapidly breaks down and evolves into an ulcer that is painful, enlarging and necrotic. It has been reported in association with a wide variety of systemic diseases including ulcerative colitis, Crohn's disease, rheumatoid arthritis, chronic myeloid leukemia, agammaglobulinemia, respiratory infections and post-surgical interventions. Histopathologic and immunopathologic study of pyoderma gangrenosum has shown the most characteristic changes are those of lymphocytic vasculitis, mild fibrinoid necrosis of the blood vessel walls, disruption of the blood vessel structure and thromboses of the lumen of the vessels.

It seems reasonable to assume therefore that the lesions involving brainstem structures causing Webino syndrome and that causing myelopathy in the present case probably resulted from vasculitis and occlusive vascular disease.

REFERENCES