Optic neuropathy secondary to cat scratch disease: case report

Neuropatia óptica secundária a doença da arranhadura do gato: relato de caso

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ABSTRACT
Optic neuropathy due to cat scratch disease is a relatively infrequent occurrence associated with macular star formation and is characterized by sudden painless loss of vision mostly unilateral. Bartonella henselae is well recognized as the etiologic agent in cat scratch disease. Ocular complications of the disease occur in up to 10% of patients and include neuroretinitis. Ocular bartonellosis is usually self-limited with complete or near-complete recovery of vision in otherwise healthy patients. A case of a boy with neuroretinitis caused by B. henselae is reported.

Keywords: Optic nerve diseases; Cat-scratch disease; Macula lutea; Macular edema; Papilledema; Retinitis; Case reports

RESUMO
Neuropatia óptica associada à doença da arranhadura do gato é relativamente infrequente, está associada com formação de estrela macular e caracteriza-se por perda visual súbita sem dor na maioria das vezes unilateral. Bartonella henselae é o agente etiológico da enfermidade. As complicações oculares da doença ocorrem em mais de 10% dos casos, entre elas está a neuroretinitis. Bartonelose ocular é uma doença autolimitada com completa ou quase completa recuperação da acuidade visual em pacientes imunocompetentes. Relatamos o caso de um adolescente com neuroretinitis causada por B. henselae.

Descritores: Doenças do nervo óptico; Doença da arranhadura de gato; Mácula lutea; Edema macular; Papiledema; Retinite; Relatos de casos

INTRODUCTION
Optic neuropathy refers to a clinical disorder characterized by sudden-to-chronic loss of vision in one or both eyes due to optic nerve dysfunction that might be idiopathic, ischemic, primary demyelinating, infectious, or inflammatory in etiology[1]. Neuroretinitis is a form of optic neuropathy, characterized by acute painless visual loss in association with optic disc swelling and macular stellated exudate. It is typically unilateral and has an excellent prognosis regardless of treatment. Optic neuropathy due to cat scratch disease is a relatively infrequent occurrence caused by Bartonella henselae. Other infectious causes include syphilis, Lyme disease, viruses and diffuse unilateral subacute neuroretinitis. Cat scratch disease has been reported in the literature for more than half a century as a syndrome of regional lymphadenopathy and fever. However, it has been only a quarter of a century since Bartonella henselae was identified as the etiologic agent. They are small, fastidious, intracellular Gram-negative bacilli that are aerobic and oxidase-negative[2]. Cats are the major reservoir for B. henselae, with up to half of domestic cats having antibodies to B. henselae. The cat flea is responsible for a relatively infrequent occurrence caused by Bartonella henselae. Other infectious causes include syphilis, Lyme disease, viruses and diffuse unilateral subacute neuroretinitis. The infection response to infection depends on the immune status of the infected host. The typical cat scratch disease is characterized by lymphadenopathy with fever, benign and self-limited, with no other signs or symptoms[3]. Other manifestations are less common (hepato-esplenomegaly, granulomatous hepatitis, osteomyelitis, encephalitis, glomerulonephritis, endocarditis, and pneumonia). The ocular involvement occurs in 10% of the cases[4]. The most common ocular manifestations is the Parinaud syndrome that consist of fever, regional lymphadenopathy, and follicular conjunctivitis, followed by the neuroretinitis[5]. We describe a case of ocular bartonellosis in one adolescent, with unilateral sudden painless vision loss and neuroretinitis.

CASE REPORT
A 17-year-old healthy boy suddenly noticed a painless vision loss in his right eye. He denied pain on eye movements and had no systemic or neurological complaints. Measured visual acuity was counting fingers at 1 meter in the right eye (OD) and 20/20 in the left eye (OS). The swinging flashlight test showed a prominent right relative afferent pupillary defect. The fundoscopy in the left eye was unremarkable but in the right eye revealed a diffusely swollen, hyperemic optic disc, serous detachment extending from the disc to the macula, and macular exudates in a stellate pattern (Figure 1 OD). The patient referred scratches on the left arm by a kitten two weeks before his eyesight was affected. Dyschromatopsia was seen in right eye. Chest x-ray, complete blood count with differential was normal. Erythrocyte sedimentation rate was 4 mm. Rheumatoid factor, Veneral Disease Research Laboratory test (VDRL), and fluorescent treponemal antibody absorption test (FTA-ABS) were negative. Tuberculin skin test was 6 mm. Serologic tests to toxoplasmosis, cytomegalovirusis and HIV 1 and 2 were negative. Cerebrospinal fluid studies revealed normal cell count, normal chemicals, and absent oligoclonal bands. Magnetic resonance images of the head and orbits were normal.

Study carried out at Ophthalmology Service, Hospital Universitário Walter Cantídio, Universidade Federal do Ceará - UFC - Fortaleza (CE), Brazil. 1Physician, Ophthalmology Service, Hospital Universitário Walter Cantídio, Universidade Federal do Ceará - UFC - Fortaleza (CE), Brazil. Correspondence address: Ricardo E. Marrocos de Araújo. Rua Oswaldo Cruz, 2.335 - Fortaleza (CE) - CEP 60125-151 - E-mail: ricardomarrosoc@yahoo.com

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The patient started a 3-week course of oral 100 mg Doxycycline once daily.

*Bartonella henselae* IgG was positive at a titer of 1:1024. Three weeks after the presentation, visual acuity improved to 20/160 OD. The prominent right relative afferent pupillary defect persisted. The optic disc swelling was markedly reduced and the macula stellate exudates remained.

Six weeks after the initial presentation, his visual acuity improved to 20/25. The pupillary response was normal. The optic disc swelling diminished and the macula stellate remained in the temporal side.

**DISCUSSION**

Optic neuropathy can be due to various causes that show significant overlap in clinical presentation and clinical examination[2]. The Optic Neuritis Treatment Trial has shown that intravenous administration of high-dose steroids decreases the risk of progression of optic neuropathy to multiple sclerosis within 2 years of follow-up[5]. The diagnosis of the underlying cause of optic neuropathy should be established as soon as possible.

The diagnosis of *B. henselae* infection is clinical, with laboratory evaluation to confirm the initial suspicion[2].

The most common posterior segment ocular complication of *B. henselae* infection is the neuroretinitis[2]. Neuroretinitis is characterized by acute visual loss due to optic nerve edema associated with macular exudates in a starlike pattern. The macular star formation may not develop until 1 to 2 weeks after the initial presentation[5]. The exact prevalence of neuroretinitis by *B. henselae* infection is unknown[7]. On MRI, the optic-globe nerve junction is highly specific for *B. henselae* as cause of optic neuropathy. The macular exudates take months to resolve, and even after resolution, patients may experience abnormal color vision and evoked potentials, subnormal contrast sensitivity, residual disc pallor, afferent papillary defects, retinal pigmentary changes[2]. Although the prognosis is on the whole good there may be residual loss of visual acuity, disc pallor, afferent papillary defects and retinal pigmentary changes.

Early signs that help to distinguish *B. henselae*-associated papillitis and/or neuroretinitis from other forms of anterior optic neuropathy are therefore important for both diagnosis and therapy[2,10].

**REFERENCES**