A B S T R A C T

Idiopathic hypertrophic pachymeningitis is a rare inflammatory condition with diffuse thickening of the dura mater, which may cause a compressive effect or vascular compromise. We report on a 28-year-old Chinese woman with a history of granulomatous mastitis 7 years previously and oligomenorrhoea, headache, blurred vision, and raised prolactin level 2 years previously, that was diagnosed as prolactinoma and treated conservatively with bromocriptine. However, she had recurrent bilateral vision loss when the bromocriptine was stopped. Her symptoms were resolved by high-dose steroid injection but remained steroid-dependent. Serial magnetic resonance imaging scan showed progressive diffuse thickening of the pachymeningitis with disappearance of pituitary apoplexy. Lumbar puncture showed lymphocytosis with no organisms. Open biopsy of the meninges was performed and histology showed features of inflammatory infiltrates and vasculitis. This is an unusual presentation of a rare condition in this age-group, with co-existing granulomatous mastitis and chronic otitis media, and is a diagnostic challenge mimicking pituitary macroadenoma and meningioma in initial magnetic resonance imaging scans.

CASE REPORT

Idiopathic hypertrophic pachymeningitis mimicking prolactinoma with recurrent vision loss

Julie YC Lok, Nelson KF Yip, Kelvin KL Chong, CL Li, Alvin L Young *

Introduction

Idiopathic hypertrophic pachymeningitis is a rare inflammatory condition characterised by fibrosis and thickening of the dura mater. Diagnosis of idiopathic hypertrophic pachymeningitis requires a high index of suspicion, as its initial manifestation could be subtle clinically and radiologically. Idiopathic hypertrophic pachymeningitis has posed considerable diagnostic challenges to attending clinicians, including radiologists, neurologists, and ophthalmologists because of its highly variable presentation. Apart from clinical subtlety and variability, idiopathic hypertrophic pachymeningitis is also a great imposter because it can mimic other common and important neurological conditions such as prolactinoma.

Case report

A 28-year-old Chinese woman with a history of granulomatous mastitis 7 years previously was noted to have oligomenorrhoea, nausea, headache, and raised prolactin level of 1261.0 mIU/L (reference range [RR], 108.8-557.1 mIU/L) since March 2011. Her baseline hormone profile was otherwise normal. Plain magnetic resonance imaging (MRI) scan showed an enlarged pituitary gland of up to 1.6 cm (Figs 1a and 1b). At that time she was diagnosed with prolactinoma and treated with bromocriptine; her prolactin level was well-controlled subsequently.

After cessation of bromocriptine at the end of 2012, she developed periodic headache and flickering vision loss since May 2013. Her best-corrected visual acuities were 20/40 and 20/30 in her right and left eyes, respectively. She developed red colour desaturation of around 30%. Humphrey visual field 30-2 test showed superior field loss in the right eye, while the left eye was full field loss. Six weeks later her visual acuity decreased to 20/800 in the right eye and 20/400 in the left eye. She was given dexamethasone and her visual acuity increased to 20/20 and 20/16 in the right and left eyes, respectively. Other slit-lamp and fundal examinations were unremarkable. There was no papilloedema. There was no involvement of other cranial nerves. Glasgow Coma Scale score was 15/15 throughout. Magnetic resonance imaging with contrast showed increased contrast enhancement and inflammation over the dura of the sella and cavernous sinus (Figs 1c and 1d). Her symptoms resolved and visual function recovered quickly with high-dose steroid. However, she developed steroid dependency since repeated attacks developed when the steroid was stopped.

* Corresponding author: youngla@ha.org.hk
Serial MRI scan showed progressive diffuse thickening of the pachymeningitis with disappearance of pituitary apoplexy, together with chronic otitis media (Figs 1c and 1d). Lumbar puncture showed 13 cm H₂O and lymphocytosis without organisms. Open biopsy of the meninges was performed and histology showed features of inflammatory infiltrates and vasculitis, but was negative for malignancy (Fig 2). Gene rearrangement polymerase chain reaction assay for immunoglobulin (Ig) heavy chain, T-cell receptor (TCR)–beta and TCR-gamma all showed no clonal peak. Polymerase chain reaction for Mycobacterium tuberculosis and culture for dural biopsy were also negative; IgG4 was 975 mg/L (RR, 61-1214 mg/L). Complete blood count showed normal haemoglobin, white blood cell, and platelet levels. Liver and renal function tests, serum calcium, creatine kinase, and lactate dehydrogenase levels were normal. Erythrocyte sedimentation rate was raised to 113 mm/h (RR, 0-20 mm/h). C-reactive protein was raised at 67.4 mg/L (reference level, <9.9 mg/L). In coagulation profile, activated partial thromboplastin time was slightly prolonged to 40.0 seconds (RR, 28.2-37.4 seconds). Autoimmune markers, including anti-nuclear antibodies, antineutrophil cytoplasmic antibodies, anti-DNA immunofluorescence test, anti-extractable nuclear antigen, anti-cardiolipin antibodies, and rheumatoid factors are all negative except for the presence of lupus anticoagulant. Complement C3 was normal while complement C4 was slightly raised to 0.42 g/L (RR, 0.1-0.4 g/L). Serum vitamin B12 level was 552 pmol/L (RR, 156-698 pmol/L) and serum folate level was 29.3 nmol/L (RR, 10.4-42.4 nmol/L); IgG4 was 975 mg/L. Virology screening—including human immunodeficiency virus, hepatitis B virus, and hepatitis C virus—was negative. Serum and cerebrospinal fluid Venereal Disease Research Laboratory tests were negative. Chest X-ray was clear with no features of tuberculosis or sarcoidosis.

The patient was initially treated with pulse steroid and was well-controlled by low-dose steroid. She remained symptom-free 6 months after biopsy.

Discussion

Idiopathic hypertrophic pachymeningitis is a rare inflammatory condition involving focal and/or diffuse thickening and fibrosis of the dura mater. Thickened dura mater with local mass effect may be pathognomonic of this condition. This pressure effect may serve as a mechanistic explanation of the observed neurological defect. As almost every part of the dura mater can be affected focally and/or diffusely, there is a highly variable clinical picture. Diagnosis is almost always made by exclusion of a large number of aetiologies, for example, infectious causes such as Lyme disease, syphilis, and M tuberculosis; inflammatory causes such as Wegener granulomatosis, rheumatoid arthritis, Behçet disease, and sarcoidosis (which is rare in Chinese people); and malignancy. In this patient, these tests were all negative, so the diagnosis of idiopathic hypertrophic pachymeningitis was made.
Demographically, the median age of patients with idiopathic hypertrophic pachymeningitis is 58.3 years (standard deviation, 15.8; range, 37-88 years). Only a few paediatric patients have been reported, with the youngest age being 2 years and 11 months in India. Our patient had early-onset disease. There are few data on ethnicity due to the rarity of the disease. Idiopathic hypertrophic pachymeningitis is extremely rare in Chinese people. The exact aetiopathophysiology of idiopathic hypertrophic pachymeningitis is not known. It is believed to be autoimmune in origin. Lupus anticoagulant is a type of autoantibody that binds to phospholipid and protein, which is commonly associated with autoimmune diseases such as systemic lupus erythematosus and antiphospholipid syndrome. Our patient’s symptoms and signs did not fit into any diagnostic category of autoimmune disease. Only the presence of lupus coagulants might suggest that idiopathic hypertrophic pachymeningitis is a form of vasculitis, which might share some common phenomenon with other autoimmune diseases, although the true relationship is controversial.

Clinically, headache is by far the most common feature and the optic nerve is one of the most common cranial nerves to be involved, which was the case for this patient. Multiple cranial nerve involvement, ataxia, cortical blindness, psychosis, motor function disturbance, fever, convolution, and/or loss of consciousness have all been reported. Yamada et al thought that the inflammatory thickening of the dura may cause damage to the superior hypophyseal artery resulting in subarachnoid haemorrhage and apoplexy in the anterior lobe of the pituitary gland. The posterior lobe was spared in their patient, who had a normal hormonal profile, unlike our patient. The initial enlarged pituitary gland with raised prolactin was more likely to result from the stalk effect than from true prolactinoma.

Granulomatous mastitis is a rare idiopathic chronic benign breast condition, which is believed to be autoimmune in origin, and mainly affects women of child-bearing age. Granulomatous mastitis is mainly diagnosed by exclusion of other diagnoses. To the best of our knowledge, this is the first case of idiopathic hypertrophic pachymeningitis reported with the association of granulomatous mastitis, possibly related to the scarcity of cases affecting menstruating women.

Idiopathic hypertrophic pachymeningitis is a rare condition with a highly variable clinical presentation making accurate and timely diagnosis difficult. Therefore the attending clinician should maintain high vigilance in the event of an atypical presentation of a presumably typical disease. Early diagnosis and prompt therapeutic intervention such as high-dose steroid may be the key to preserving vision as well as life.

References