disorder. The combination of HLA types in these patients may be a marker for a predisposition to the Guillain-Barré syndrome.

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Adult on set acid maltase deficiency associated with epilepsy and dementia: a case report

Adult onset acid maltase deficiency (AMD) (glycogenosis type II b) characteristically presents after the age of 20 years with a slowly progressive limb-girdle muscle weakness. The onset of muscle involvement occurs in a third of cases and may be the presenting feature. Although CNS involvement is seen in the more severe infantile form of AMD (Glycogenosis type IIa), it has not been described in adults. We report a case of adult onset AMD associated with epilepsy and dementia.

A 65 year old woman was referred with intractable epilepsy, the onset of which was at 20 years with a generalised tonic-clonic seizure. Seizures had continued without remission despite treatment with phenobarbital, phenytoin, and sodium valproate. The patient's history consisted of a secondarily generalised tonic-clonic seizure twice a month and a complex partial seizure approximately monthly. At the age of 29 years she required psychiatric admission because of attempted suicide and hysterical behaviour. She was thought to have an immature personality and to lack foresight and judgement. From the age of 52 years she had been in residential care, and anti-social behaviour, incontinence of urine and impaired cognition had been noted. From the age of 57 years she developed progressive limb weakness.

She had five siblings. One was reported to have epilepsy but the details were not available. Three of the other siblings had adult onset AMD and their histories have been published previously. The parents had no history of any neurological disorder and there was no consanguinity.

On examination cooperation was limited. She was unable to stand. Fundoscopy and eye movements were normal. There was a mild weakness of neck flexion. Wasting of the shoulder girdle muscles and small muscles of the hand was present bilaterally. Wasting was difficult to assess in the legs because of obesity and oedema. There was a grade 4 weakness proximally in the arms, grade 4+ distally, and a grade 2–3 weakness in the legs, more marked proximally. Coordination was normal. Bilateral grasp reflexes were elicited. Upper limb tendon reflexes were normal but knee and ankle jerks were absent bilaterally. Plantar responses were flexor. No sensory deficit was found and discrimination was unremarkable except for obesity.

The following investigations were normal or negative: full blood count, electrolytes, thyroid function, liver function tests, coagulation profile, serum electrophoresis, autoimmune profile, blood film for acanthocytes, sphingolipid, plasma and urinary amino acids, white cell lysosomal enzymes and nerve conduction studies. T2 was 45 mmol/l in 1 hour and the creatine kinase was raised at 405 IU/L (0–243 IU/L). Initially serum lactate was slightly elevated at 1.83 mmol/l (0.5–1.65 mmol/l), but was normal on two subsequent occasions. Serum pyruvate was normal. An electrocardiogram showed poor R wave progression but was otherwise normal. Chest radiography revealed a raised right diaphragm. Lying and standing vital capacity were 1.051 and 1.51 respectively. Arterial blood gases were normal. During a sleep study lasting 9-18 hours oxygen saturation was 70–80% for 2-3 minutes, 80–90% for 35 minutes, and 90% for the remainder of the study. Electrographic examinations were moderately myopathic without any specific features. Histological appearances of a quadriceps muscle biopsy were indicative of AMD, with a high proportion of the muscle fibres containing multiple vacuoles packed with glycogen. Acid maltase activity in the muscle was reduced to 0·01 (0·07–0·29) nmol maltose hydrolysed per gram of muscle (wt weight) per minute. Oxidative enzymes, phosphorylase, and adenylate deaminase were normal. The diagnosis was supported by finding a significant number of lymphocytes containing small discrete glycogen filled vacuoles.

CT brain scan showed cerebellar atrophy but the patient was unable to comply with an MRI scan. An electroncephalogram showed an irregular dominant rhythm of 8 Hz, widespread theta and delta activity and superimposed bilateral spike and spike wave discharges. Neuropsychological assessment revealed that the patient had a severe disturbance of frontal function, in the absence of cognitive impairment. The patient's performance was consistent with a mild degree of frontal dysfunction.

The progressive weakness with evidence of respiratory muscle involvement, muscle histology, and lymphocyte appearances in this patient were all typical of adult onset AMD. Serum lactate was raised on one out of three measurements, which although not a usual feature of AMD may have been secondary to hyperventilation. Additional features in this case were epilepsy, psychiatric disturbance and impaired cognition, for which no other aetiology was identified. Her three siblings with AMD showed a spectrum of clinical and pathological findings, but did not have epilepsy or dementia, and an elder sister with no history of muscle weakness, had epilepsy. It is possible that, in this family there is a tendency to epilepsy independent of the AMD. An alternative explanation is that the epilepsy, psychiatric disturbance and impaired cognitive function seen in this patient are a manifestation of central AMD.

Studies in previous cases of adult onset AMD clinical involvement has been restricted to the skeletal muscles, necropsy studies have revealed widespread disease with microscopic abnormalities in all skeletal muscles, whether weak or not, and variable involvement of smooth muscle, cardiac muscle, the tongue and the spleen. In these two studies of adult onset AMD no microscopic abnormalities were found in the brain. Acid maltase activity is lowest in skeletal muscle, but also reduced in the liver, heart and brain. Cerebral acid maltase activity has been recorded as 6-5% of normal. There is variation in the degree to which various organs are affected. For example in one post mortem study, acid maltase activity in the myocardium was as low as that in skeletal muscle and this was reflected by a vacuolar cardiac myopathy.

In cases of infantile AMD there is more widespread involvement, with both cardiac and central nervous system manifestations being more prominent, with abnormal glycogen storage in most cells of the brain and spinal cord.

In the few adult onset cases that have been studied, acid maltase activity in the brain has been low. It is possible that this patient with adult onset AMD has very low levels of cerebral acid maltase activity and that this is the cause of her disturbance of cerebral function. The association of adult onset AMD with epilepsy, cognitive impairment and psychiatric disturbance, however, may be fortuitous and does not appear to have been drawn from a single patient. We draw attention to the association in case similar patients have been seen by others.

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A reappraisal of “direction of scratch” test: using somatosensory evoked potentials and vibration perception

We would like to comment on the article by Hankney and Edin which proposed “direction of scratch” test for the clinical examination of posterior column function. That method is easy to perform, and the errors in the detection of direction of scratch on the skin were reported to be correlated with impaired position and/or vibration sense tested using 128 Hz tuning fork. Since somatosensory evoked potential (SEP) and vibration sense was thought to be mediated through the posterior column and lemniscal pathways, reduction of the useful-ness of their test together with SEPs and quantitative vibration sense.

Fifty six patients, mean age 43.5, were studied. They consisted of 43 patients with...
Seventh nerve palsy as a false localising sign

Benign intracranial hypertension (BIH) usually produces a clinical picture of head-ache, visual disturbances and bilateral papilloedema. Abducens nerve palsies have been reported in between 10–30% of cases. Other cranial nerve palsies have rarely been reported in this condition and those that have occurred were nearly always in children. One case of facial diplegia occurring in an adult with this condition has been documented.1 We report a patient with BIH who presented with bilateral sixth nerve palsies and a right sided seventh nerve palsy all of which resolved after a lumbar puncture.

A twenty five year old obese woman on the contraceptive pill presented with a three week history of bifrontal headaches, nausea and vomiting. The headache was not made worse by coughing or stooping and neither was it relieved by analgesics. One week before admission the patient developed diplopia on horizontal gaze and visual obscuration. Examination revealed mild bilateral sixth nerve palsies and a right sided facial nerve palsy of lower motor neuron type with a reduced blink reflex. The visual acuity was 6/6-2 on the right and 6/5-1 on the left. The blind spots were enlarged but the visual fields were otherwise normal. Marked bilateral papilloedema was observed. An ophthalmological examination was normal. A CT head scan, including a contrast examination, was normal. Lumbar puncture revealed clear and colourless CSF with an opening pressure of 300 mm CSF. The fluid was acellular with a protein of 230 mg/l and a glucose of 4.0 mmol/l (plasma glucose 5.2 mmol/l). Gram stain and cytology for malignant cells were both negative. The following investigations were normal: CXR, FBC, ESR, ANF, VDRL, clotting studies, lupus anti-coagulant, fibrinogen level, immunoglobulins, C3, C4, serum osteoitin, androstenedione, serum DHA - sulphate, LH and FSH. On the day following lumbar puncture, the diplopia had resolved and facial weakness improved. By the second day, the cranial nerves examinations was normal except for papilloedema. The patient was started on a weight reducing diet and has required no further treatment. On review six weeks after discharge, the visual acuity was 6/4 bilaterally and mild papilloedema was present.

Identifying a neurological sign as being falsely localising rests on two important considerations. Firstly, that an accurate knowledge of the anatomy is attained and secondly, that the sign itself must arise in an anatomical area that is remote from the site of the original pathology. A sixth nerve palsy is a classic example in patients with raised intracranial pressure from a brain tumour distant from the course of the nerve. The long intracranial course of the abducens nerve is thought to be responsible for its predilection compared to other cranial nerves. However, Collier favours an alternative explanation based on the direction in which the nerve emerges from the brainstem. The abducens nerve emerges in a straightforward direction, whilst all other cranial nerves emerge transversely or obliquely. Therefore, backward displacement of the brainstem by an intracranial space occupying lesion would exert its maximum effect on the sixth nerve. Either way the patho-physiological mechanism is thought to involve displacement of the brain by enlarging intracranial mass and traction upon cranial nerves. False localisation from other cranial nerve palsies have also been described.2 BIH offers a very good model to study the effect of raised pressure without focal mass effect suggesting that the cranial nerve palsies could be the result of general pressure related effect without implying a specific directional intracranial pressure forces. BIH was first described by Quincke in 1893.3 At that time ostitis media was thought to be the most common aetiology and there has been renewed interest in sinus occlusion in the genesis of BIH.4 Many other mechanisms have been implicated including obesity5 and the contraceptive pill6 both of which were involved in our patient.

Whilst abducens nerve palsies are not uncommonly seen as a false localising sign of...