SHORT REPORT

Hereditary haemochromatosis: a case of iron accumulation in the basal ganglia associated with a parkinsonian syndrome

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Abstract
Hereditary haemochromatosis is characterised by excessive parenchymal iron deposition, particularly in the liver. Usually hereditary haemochromatosis is not associated with neurological symptoms and iron deposition in the brain has not previously been described as a pathological phenomenon. A patient is reported with hereditary haemochromatosis and a syndrome of dementia, dysarthria, a slowly progressive gait disturbance, imbalance, muscle weakness, rigidity, bradykinesia, tremor, ataxia, and dyssynergia. The findings on MRI of a large signal decrease in the basal ganglia, consistent with excessive iron accumulation, indicate a causal relation to the symptoms. Although the neurological symptoms did not improve in our patient, hereditary haemochromatosis should be considered in the differential diagnosis of parkinsonian syndromes, because complications of iron induced organ injury may be prevented by phlebotomy.

Keywords: hereditary haemochromatosis; basal ganglia iron; parkinsonian syndrome

Hereditary (or genetic) haemochromatosis is an autosomal recessive inherited iron overload disease, localised to chromosome 6p in proximity to the HLA-A3 locus. If the accumulation of excess iron is due to other conditions—for example, thalassaemia or sideroblastic anaemia—the term secondary iron overload syndrome is used. The genetic defect and the underlying metabolic error of hereditary haemochromatosis are unknown. Cellular accumulation of iron is thought to induce lipid peroxidation with the consequence of cell damage and cell death.1 The clinical manifestations are the result of the specific pattern of organs affected, and most often seen are weakness and weight loss, arthropaligae, abdominal pain, skin pigmentation, hepatomegaly, hepatic cirrhosis, diabetes mellitus, arthropathy, cardiac manifestations, and hypogonadism. The most common laboratory abnormalities are raised percentage saturation of transferrin and serum ferritin concentration, increased amounts of stainable iron in hepatocytes and an increased hepatic iron concentration. Most patients present with symptoms at the age of 40–50, but some may present earlier, in the second or third decade. There is a sex difference and the frequency of male to female haemochromatosis ranges from 2:1 to 18:1.2 Neurological features associated with hereditary haemochromatosis are rare. Lethargy, psychomotor retardation, fatigue, confusion, hearing loss, and polyneuropathy have been described.3

To the best of our knowledge, there have been no reports of excessive iron deposition in the CNS in hereditary haemochromatosis, an otherwise well known pathological feature of the Hallervorden-Spatz syndrome and familial hypoceruloplasminaemia.

We report on a patient with hereditary haemochromatosis with a parkinsonian syndrome, most likely caused by excessive iron accumulation in the basal ganglia and cerebellum.

Case report
The patient, a 50 year old man with a normal psychomotor development, without known predisposition of any kind and no family history of hereditary haemochromatosis, had been admitted to a neurology department when 29 years old, because of a slowly progressive gait disturbance with episodes of unsteadiness and imbalance since the age of 26. He noticed muscular weakness, rigidity, and slowness of voluntary movement of the left arm and leg. His gait was described as reeling with a deviation towards the left, dragging his left foot. Tendon reflexes were normal, except for a very brisk right sided biceps reflex. Otherwise the neurological examination was normal. A pneumoencephalogram showed slight central atrophy of the cerebrum. Cervical pneumomyllography was normal, as were radiographs of the chest, spine, and skull. Blood tests, examination of CSF, and EEG were all reported to be within the normal range. He was discharged without any definite diagnosis.

Subsequently he was referred for a specialist opinion concerning his application for a disablement pension. For 18 years he had
degree of pigmentation in the irides, the centres being brownish and hyperpigmented, was striking, but no Kayser-Fleischer ring was seen on slit lamp examination. Other cranial nerve functions were normal. Examination of the heart and the abdomen was normal, and there was no testicular atrophy.

A slight muscle atrophy was evident in the lower limbs, mainly distally on the left side. Muscular strength was moderately impaired in the upper limbs and in the lower limbs on the left in a pyramidal distribution. The muscular tone was increased in both upper and lower limbs in the form of rigidity, mainly on the left side, where there was also a discrete cogwheel phenomenon. Cutaneous sensation, joint position, and vibration senses were normal. Tendon reflexes were easily obtained in the upper limbs and were very brisk in the lower limbs, particularly on the left, with clonus at the ankle. The right plantar reflex was normal, whereas a Babinski's reflex was found on the left. Coordination was influenced by ataxia and severe dysdiadochokinesia. A combined static and postural tremor was noted, as well as truncal instability, with a tilting to the left. The gait was characterised by spasticity with circumscription of the left leg, holding the left arm in an adducted, flexed position; the step length and pace was normal. No involuntary movements were noted, but bradykinesia was seen. The skin was bronzed. The patient started treatment with levodopa. This resulted in an immediate improvement.

Normal findings included an ECG, blood cell counts, haemoglobin, erythrocyte sedimentation rate, serum concentrations of creatinine, carbamide, sodium, potassium, calcium, albumin, carbon dioxide, transaminase, alkaline phosphatase, total bilirubin, IgA, IgG, IgM, urate, a negative serum antinuclear antibody titre, and iron, plasma concentrations of fasting glucose and ceruloplasmin, and plasma prothrombin time. Urinary copper excretion was normal and the copper content in the liver (5.2 mg/kg) was also within the normal range (5.15 mg/kg).

Serum ferritin (1313 pmol/l (normal 33–666 pmol/l) and transferrin saturation (53% (normal <50%)) were high. Serum transferrin (25 \(\mu\)mol/l (normal 31–57 \(\mu\)mol/l)) was low. A liver biopsy showed a heavy iron overload, but liver architecture was normal. Brain MRI (fig 1) showed a pronounced signal decrease in the caudate nucleus, the inner segment of the globus pallidus, the red nucleus, substantia nigra, and the dentate nucleus, consistent with excessive iron accumulation. No atrophy or focal abnormalities were seen. Liver MRI (fig 2) showed a pronounced signal intensity decrease, corresponding to a hepatic iron concentration of 70 \(\mu\)mol/g wet weight (normal range 1–9 \(\mu\)mol/g). HLA typing showed heterozygosity for the HLA system (HLA-A2,3,B7,44).

Phlebotomy was started. After eight treatments the fatigue, the headache, and the skin bronzing had disappeared. There were no other improvements.
Discussion

The diagnosis of hereditary haemochromatosis was based on the presence of biotically confirmed excessive parenchymal iron accumulation without any indication of secondary iron overload, the increased percentage saturation of transferrin, and the increased serum ferritin concentration. In accordance with the recessive mode of inheritance, and maybe the sex difference also, his parents as well as his two sisters and half-brother had no symptoms of hereditary haemochromatosis. The currently applied definition of hereditary haemochromatosis also requires homozygosity for the haemochromatosis alleles. Our patient is heterozygous for the HLA-A and B haplotypes, but presumably homozygous for the haemochromatosis allele, which, owing to family relations is not accessible for investigation. The two most common HLA alleles associated with hereditary haemochromatosis in Denmark are A3 and B7, alleles which were both found in our patient. HLA-A3 occurred in about 75% of patients with hereditary haemochromatosis compared with 28–30% in controls in Europe, the United States, and Australia. Usually those who are heterozygous for the haemochromatosis allele do not present with clinical disease or develop massive body iron overload, but a proportion may display biochemical abnormalities (most commonly increased transferrin saturation).

Excessive metal accumulation in the CNS is known to cause neurological symptoms. Wilson’s disease (hepatolenticular degeneration) is an autosomal recessive condition, located on chromosome 13q with abnormal copper deposition in the brain (particularly the basal ganglia), liver, cornea, and other organs. Usually the neurological onset is in adolescence, with symptoms such as clumsiness, personality change, reduced mental performance, tremor, rigidity, bradykinesia, dystonia, chorea, incoordination, ataxia, dysarthria and, rarely, seizures. Almost always a Kayser-Fleischer ring at the edge of the cornea is seen. Confirmation of the diagnosis by DNA analysis is possible. Biochemically the diagnosis is confirmed by a low plasma ceruloplasmin, a low serum copper, a high urinary copper excretion, and a high concentration of copper in the liver. Thus this condition was ruled out in our patient.

Locally increased iron concentration in the basal ganglia, not related to any abnormality in the general iron metabolism, with axonal swellings and neuronal degeneration, is pathognomonic for the Hallervorden-Spatz syndrome; characterised by onset in childhood, progressive dementia, bradykinesia, rigidity, spasticity, dystonia, and involuntary movements. The inheritance is recessive, and a genetic localisation is not established. Eidelberg et al reported on three adult onset single cases, all mentally retarded women, in whom symptoms started between 31 and 42 years of age. Three sibs, with late age of onset from early adolescence to 55 years of age, with Hallervorden-Spatz disease presenting as familial parkinsonism, were described by Jankovic et al. A biochemical analysis of the brain of the oldest man affected showed considerable loss of dopamine in the nigrostriatal areas, a feature which may explain the beneficial effect of levodopa seen in our patient.

Miyajima et al described a 52 year old woman with familial hypocuprolasminemia, blepharospasm, and retinal degeneration. This disorder is dominantly inherited, located on chromosome 3q, and caused by apoceruloplasmin deficiency, which is thought to be causally linked to the iron deposition in the basal ganglia and other organs, shown by kinetics, CT, and histochemical studies.

Parkinson-like syndromes have also been reported after chronic manganese and thallium intoxication, but never in association with hereditary haemochromatosis. Dementia, ataxia, rigidity, and myoclonic jerks were described in two patients with idiopathic haemochromatosis. Both exhibited slight abnormalities of liver function. Patient 1 had normal brain CT. A postmortem micronodular cirrhosis was found, but no specific changes were noted on neuropathological investigation. A small amount of iron in astrocytes and in the basal ganglia was not thought to represent a pathognomonic change. Patient 2 had a history of alcohol abuse. Cerebrum CT and postmortem examination were not performed. The authors concluded that cirrhosis of the liver, although clinically occult, were the cause of the syndrome of chronic hepatocerebral degeneration.

Furthermore, it has been proposed that a selective increase of iron in the substantia nigra might be linked to the neurodegenerative aspects of Parkinson’s disease, involving iron and iron-melanin induced membrane lipid peroxidation, but this issue still remains to be solved.

Brain MRI showed pronounced signal decrease in the caudate nucleus, the inner segment of the globus pallidus, the red nucleus, substantia nigra, and the dentate nucleus. Several MR studies performed in extrapyramidal movement disorders including Parkinson’s disease, multiple system atrophy, Huntington’s disease, and Hallervorden-Spatz disease have shown a close correlation.
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between brain iron concentration and T2 relaxation time within these regions.\textsuperscript{14,15}

A recent study by Chen et al on post-mortem brain tissue from patients with Parkinson's disease and Huntington's disease showed a trend towards T2 shortening with increasing iron concentration within both the putamen and the globus pallidus in Parkinson's disease and within the globus pallidus in Huntington's disease. In Huntington's disease the highest iron concentration and the longest T2 values were in the putamen, and it was concluded that signal intensity of the basal ganglia on T2 weighted images may be determined by several factors; however, the fact that iron deposition causes a reduction in T2 relaxation was not disputed.\textsuperscript{16}

In the present patient extremely low signal intensity was found in the liver on gradient echo MRI consistent with the finding of biologically confirmed excess iron accumulation in the liver. A strong correlation has previously been shown between signal intensity and iron concentration in the liver.\textsuperscript{17} This finding and that of reduced signal intensity within the mentioned brain areas in our patient suggests that increased iron accumulation may take place in the brain of patients with hereditary haemochromatosis.

In conclusion, it is therefore likely that excessive iron deposition in the basal ganglia and cerebellum, as shown on MRI, is causally linked to the parkinsonian syndrome in our patient. Although he has presently not improved neurologically on the phlebotomy treatment there is still a possibility that he may do so, because serum ferritin is not yet within the normal range. Hereditary haemochromatosis is not usually considered in the differential diagnosis of parkinsonian syndromes, but because complications of iron induced organ injury may be prevented by phlebotomy, we draw attention to hereditary haemochromatosis as a possible explanation when facing an otherwise "idiopathic" extrapyramidal syndrome.

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