Two families with autosomal recessive spastic paraplegia, pigmented maculopathy, and dementia

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Abstract

Objective—Two families with autosomal recessive hereditary spastic paraplegia and pigmented maculopathy are described.

Methods—All family members were examined by two neurologists. An assessment of cognitive function in affected members was made using the mini mental state examination (MMSE) or Cambridge cognitive examination (CAMCOG).

Results—Six patients from two families presented with a slowly progressive, autosomal recessive, spastic tetraplegia. Although they were always considered to be intellectually slower than their peers, further intellectual deterioration was noted during the second decade. Five had a pigmented maculopathy with mild decrease in visual acuity and all had distal amyotrophy, mild cerebellar signs, and developmental faecal and urinary incontinence late in the course of the disease.

Conclusion—The association of hereditary spastic paraplegia and pigmented maculopathy has rarely been described; only 11 families with 32 affected members have been reported, showing considerable heterogeneity in presentation. These described conditions may be allelic or more probably reflect mutations at different genetic loci.

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Keywords: hereditary spastic paraplegia; maculopathy; dementia; Kjellin's syndrome

Since the original description of hereditary spastic paraplegia, reports of complicated forms have been limited only by the finite way in which neurological disease can present. Until the underlying genetic and biochemical defects are established, classification must rely on accurate clinical description. The association of hereditary spastic paraplegia and pigmentation of the macula has rarely been described. Associated clinical features may include hearing loss, mental retardation, dementia, behavioural disturbance, cerebellar signs, distal amyotrophy, ichthyosis, and syndactyly.

Kjellin described two families with autosomal recessive inheritance, mental retardation, and onset of spastic paraplegia in the third decade, associated with macular pigmentation and distal amyotrophy. In this study two families are described with clinical features of Kjellin's syndrome but with several distinguishing points: the age of onset for spastic paraplegia was earlier and they had additional clinical features including sphincter disturbances, dysarthria, mild cerebellar ataxia of the upper limbs, and further intellectual deterioration in the second decade that were not described by Kjellin.

Methods

Members from both families were examined by two neurologists. An assessment of cognitive function in affected members was made using the mini mental state examination (MMSE) or Cambridge cognitive examination (CAMCOG). Scores in the CAMCOG are rated out of 107; >80 being considered normal, 60-79 mild dementia, 35-59 moderate dementia, and <35 severe dementia.

Affected family members had the following tests performed: a full blood count, blood film and differential, erythrocyte sedimentation rate and karyotyping, serum levels of urea and electrolytes, liver function tests, glucose, creatine phosphokinase, aldolase, thyroid function tests, vitamin B12 and folate, vitamin E, vitamin A, cholesterol and triglycerides, fasting pyruvate and lactate, copper, caeruloplasmin, lead, acanthocytes, immunoglobulins, syphilis serology, urinalysis for reducing sugars, protein, phenylketouria, acetate, amino acid chromatography, calcium, mucopolysaccharides, urbinilogen, porphyrins, indican, a chest radiograph and ECG, echocardiography (ECO), analysis of CSF including oligoclonal bands, EEG, visual evoked responses (VER), electroretinography (ERG), fundusphotography, nerve conduction studies, EMG, and neuroimaging, including MRI of the brain and spinal cord.

The Families

Both families (fig 1A and B) lived within 35 km of each other. In Family A, there were three affected (IV-2A, IV-5A, IV-6A) and four unaffected children (IV-1A, IV-3A, IV-4A, IV-7A). Their sister IV-1A was normal until she developed Rasmussen's encephalitis at 30 years of age. Their mother, III-12A, was normal and their father, III-11A who died aged 55, was reported to have been normal. Consanguinity was denied and there was no other affected member on either side of the family.

In Family B there were three affected (IV-1B, IV-2B, IV-3B) and three unaffected children (IV-4B, IV-5B, IV-6B). Their sister IV-4B had intellectual impairment after an episode of meningitis aged 4 months but was...
otherwise normal. Their parents were first cousins and were normal on examination. There was no history of similar clinical features on either side of the family.

CASE REPORTS
The affected members within each family had similar phenotype and clinical course (table 1). The following case reports are representative of each family.

Family A
Case IV-2A—This 39 year old man was born by forceps delivery. He was considered a “slow learner” and a psychologist reported borderline mental retardation when he was eight years old. His intellect showed further deterioration at 15 years and he developed a slowly progressive spastic paraplegia at 17 years. For the past year he has been confined to a wheelchair with both urinary and faecal incontinence. At 34 years he developed hallucinations, paranoid ideation, and mood swings. His present medications include thioridazine, benztropine, monthly injections of flupenthixol, and baclofen.

On examination he had pronounced intellectual impairment and his speech was dysarthric, impoverished, and lacked spontaneity. Lateral eye movements were slightly reduced. At 39...
years, corrected visual acuity was 6/60 in the right eye and 6/24 in the left eye. Formal assessment of visual fields was not possible. Fundoscopy at 32 years had been normal. Repeat fundoscopy at 39 years showed about 20 discrete areas of pigment aggregation 0.1-0.5 mm in diameter and each aggregation was surrounded by a yellow atrophic halo (fig 2). Although largely confined to the macula in both eyes these lesions were also seen medial or inferior to the optic disc. His arms showed fasciculations but no wasting, power was reduced for finger and thumb abduction, elbow extension and neck flexion, he was unable to sit without support, reflexes were brisk, tone was mildly increased, there was an intention tremor bilaterally, and sensation was normal. His legs showed no wasting or fasciculations. There was a flexion deformity of the right hip and knee and shortening of the achilles tendon. The left hip, knee, and ankle were fixed in extension. There was no voluntary movement of either leg, spontaneous clonus was present, plantar responses were extensor bilaterally, and sensation was normal. His feet had pes cavus deformity.

Family B
Case IV-2B—This 32 year old man developed a slowly progressive spastic paraplegia at 14 years and was confined to a wheelchair at 21 years. In the past two years he has developed faecal and urinary incontinence. He was always considered by his parents to be intellectually slower than the other children and had further cognitive deterioration in the second decade.

On examination he sat slumped forward in his wheelchair, being unable to support himself in a sitting position. He had pronounced kyphosis, rounded shoulders, and had difficulty keeping his head up. There was no spontaneous speech and responses were slow, monosyllabic, and dysarthric with a nasal quality. His mother thought that his vision was normal but formal assessment was not possible. At 24 years ophthalmological examination disclosed clumps of macular pigmentation bilaterally, each surrounded by areas of atrophic change. His arms showed fasciculations with considerable wasting of the thenar and to a lesser extent the hypothenar eminence. He had mild contracture deformity at both elbows, tone was mildly increased on the left side, and reflexes were brisk bilaterally. He had reduced power and lack of spontaneous movement in both arms. Although he reached out for objects in front and to the side of him without any tremor he was unable to lift his arms above his head. There was wasting with fasciculations in his calf muscles, tone was considerably increased with adductor spasm, and power was absent. Both achilles tendons were shortened. Reflexes were brisk at the knees but absent at the ankles. His planter responses were extensor and his feet were flat.

Results

Family A

The CAMCOG score was 35 in IV-2A, 46 in IV-5A, and 56 in IV-6A. Brain MRI showed
diffuse atrophy of the cerebral hemispheres, corpus callosum, and brain stem in all three affected members. The EEG was normal in IV-2A but showed non-specific voltage changes and slow wave activity in IV-5A and IV-6A. Measurements in CSF including oligoclonal banding was normal in IV-6A. The ERG was normal in IV-5A but showed increased latency in IV-2A and IV-5A. The VER was normal in IV-2A but the rod isolated responses were delayed and the amplitude was reduced in IV-5A and IV-6A and the cone responses were normal but the amplitude was reduced in IV-6A. Nerve conduction studies were normal but the EMG showed fasciculations in the small muscles of the hand in IV-6A. The ECG was normal in IV-5A but there was right axis deviation with incomplete right bundle branch block in IV-2A and generalised T wave flattening in IV-6A. An ECHO was normal in IV-6A. A muscle biopsy, in IV-6A, showed random fibre atrophy and fibre type grouping indicative of denervation and associated re-innervation. There were no histological features to support a mitochondrial disorder. All other tests were normal.

**Discussion**

Six members from two families have an autosomal recessive syndrome characterised by mental retardation, further intellectual deterioration in the second decade, spastic tetraparesis, distal amyotrophy, macular pigmentation, mild reduction in visual acuity, dysarthria, mild cerebellar signs, and faecal and urinary incontinence (table 1).

These two families, although unknown to each other, are probably related, as they have an identical, rare, autosomal recessive clinical syndrome and have lived within 35 km of each other for over five generations.

All affected members were considered by their parents to be intellectually slow as children and to have further cognitive deterioration in the second decade. The earlier the age of onset and the longer the duration of spastic tetraparesis the more severe was the cognitive impairment.

They all developed an abnormal gait between 13 and 23 years of age and progressed to using a wheelchair within three to 21 years. All had a similar pattern of muscle weakness in the upper body, being unable to support themselves in a sitting position and having reduced power for neck flexion, elbow extension, and finger and thumb abduction. All affected members had evidence of a lower motor neuron lesion in the arms, probably of anterior horn cell origin.

Macular pigmentation was seen in five affected members and although lesions predominantly involved the macula, in two members more widespread retinal pigmentation was seen. In the four cases presented by Kjellin extension of pigmentation outside the macula was not seen at fundoscopy but in all cases an

**Table 2** Autosomal recessive hereditary spastic paraparesis (HSP) and macular pigmentation—summary of previous reports

<table>
<thead>
<tr>
<th>Author (ref)</th>
<th>Pedigree</th>
<th>HSP onset (y)</th>
<th>Mental retardation</th>
<th>Dementia</th>
<th>Fundal lesions</th>
<th>Amyotrophy of arms</th>
<th>Cerebellar signs</th>
<th>Other features</th>
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<tr>
<td>Webb et al, present study</td>
<td>1 17 18 23</td>
<td>Yes Yes Yes Yes Yes</td>
<td>Yes Yes Yes Yes Yes(mild)</td>
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<td>Nil</td>
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<tr>
<td>2 13 14 16</td>
<td>Yes Yes Yes Yes Yes(mild)</td>
<td>Yes Yes Yes Yes Yes(mild) Electrical myoclonus</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Focal dystonia</td>
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<tr>
<td>Farmer et al</td>
<td>1 13</td>
<td>No Yes No Yes No Yes(mild)</td>
<td>No Yes No No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Nicotine addiction</td>
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<tr>
<td>Sjaastad et al</td>
<td>1 5 16</td>
<td>No Yes No Yes Yes(mild)</td>
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<td>No</td>
<td>No</td>
<td>No</td>
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<tr>
<td>Mahlouji and Chuke and Farmer et al</td>
<td>1 30 35 35 30's</td>
<td>Yes No No Yes No No</td>
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<td>No</td>
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<td>No</td>
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<td>Ledic and Van Bogaert</td>
<td>1 10 12</td>
<td>No Yes Yes Yes No Hearing loss</td>
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<tr>
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<td>No</td>
<td>No</td>
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The association of hereditary spastic paraplegia and pigmented maculopathy has rarely been described, yet there is considerable clinical heterogeneity among the few reported cases (table 2). The two families reported in this study are similar on clinical grounds but differ from all previously reported cases, including those described with autosomal recessive hereditary spastic paraplegia. The onset of spastic paraplegia may range from three to 35 years but within each family the onset is tightly clustered (table 2). The presence or absence of specific clinical features varies between families but is relatively constant within each family. Although the macula is predominantly affected by disease, there is often evidence of more widespread retinal involvement both clinically and on ERG. The maculopathy usually occurs late and is slowly progressive or static. Except for the reports by Leduc and Van Bogaert and by Schaffer,10,11 the effect on visual acuity is usually mild. Amyotrophic changes are common and affect distal musculature, more in the arms than in the legs and particularly the thenar eminence. Cerebellar disturbance is mild or absent. Dementia has been described before in some of these families but has not been recognised as a feature of Kjellin’s syndrome. Spastic paraparesis, pigmented maculopathy, and dementia may occur together in other diseases (for example, Hallervorden-Spatz disease and Lawerence-Moon-Biedl disease). Although these conditions should be included in any differential diagnosis, they are characterised by predominant clinical and pathological features other than spastic paraparesis and have therefore been excluded from table 2. Hallervorden-Spatz disease is an autosomal recessive condition in which spastic paraparesis, pigmented maculopathy, mental retardation, and dementia may occur together. The usual presentation of clinical features are an akinetic-rigid and dystonic syndrome beginning between 7 and 12 years of age. Brain MRI may also show the presence of iron in the globus pallidus and substantia nigra. Lawerence-Moon-Biedl disease is an autosomal recessive condition characterised by retinitis pigmentosa, mental retardation, obesity, hypogonadism, and polydactyly; spastic paraparesis may be a variable, late finding. Batten’s disease is an autosomal recessive condition characterised by a progressive visual loss, pigmented retinopathy, and dementia with periodic acid Schiff positive material in neurons. Spastic paraparesis, and extrapyramidal and cerebellar signs may occur late in this condition. Subacute necrotising encephalomyelopathy (Leigh’s disease) is a mitochondrial disorder of pyruvate metabolism and may present with developmental arrest, retinal pigmentation, optic atrophy, loss of vision, dementia, ophthalmoplegia, nystagmus, dystonia, seizures, respiratory abnormalities, and spastic or flaccid paraparesis. Brain MRI may show areas of necrosis in the basal ganglia and putamen.

In summary, two families have been described with autosomal recessive hereditary spastic paraplegia and a pigmented maculopathy. All had mental retardation and developed a progressive spastic paraplegia in the second decade associated with further cognitive deterioration, a pigmented maculopathy, and amytrophic changes. This condition may be easily missed, unless specifically sought, as only a mild reduction in visual acuity is produced and the pigmented maculopathy may occur several years after the onset of the spastic paraplegia. Therefore, we suggest that all autosomal recessive and sporadic cases of hereditary spastic paraplegia should have an ophthalmological examination performed with dilatation of the pupils.

References:
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