A NEW LEUKOENCEPHALOPATHY
WITH BRAINSTEM AND
SPINAL CORD INVOLVEMENT
AND HIGH LACTATE

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ABSTRACT
We identified eight patients with a distinct MRI pattern of inhomogeneous cerebral white matter abnormalities and selective involvement of brainstem and spinal tracts. Proton MRS showed increased lactate in the abnormal white matter. Clinically the patients had slowly progressive pyramidal, cerebellar and dorsal column dysfunction. The uniform, highly characteristic MRI pattern and the similarities in clinical and MRS findings provide evidence for a new disease entity. Autosomal recessive inheritance is likely.
INTRODUCTION

White matter disorders of unknown etiology constitute a considerable problem in childhood. MRI and MRS have been used to identify “new” diseases among them. In two MRI-defined disorders, megalencephalic leukoencephalopathy with subcortical cysts and vanishing white matter, the gene defects have recently been identified, confirming the value of this MRI-oriented approach.

We describe a novel distinct MRI pattern, shared by eight patients. Apart from cerebral and cerebellar white matter abnormalities, striking findings are the involvement of specific brainstem and spinal cord tracts and elevated lactate in the abnormal white matter. These findings are paralleled by a homogeneous clinical presentation, reinforcing the concept of a new syndrome.

PATIENTS AND METHODS

In all patients except patient 8, at least the last study was performed on the same 1.5 T MRI machine. The protocol included brain and spinal MRI, and short-echo time proton MRS of volumes of mid-parietal cortex (8 ml) and abnormal white matter in the centrum semiovale (5 ml), as described previously. Metabolite concentrations, calculated with LCModel in mmol/l, were compared with results of 12 controls (age range 9.3-26.3) using the two-tailed unpaired student T-test.

Diffusion tensor imaging (DTI) and magnetization transfer (MT) imaging were performed in patients 1-7 and 12 controls. For DTI a multi-slice EPI sequence was used. Mean diffusivity, <D>, and fractional anisotropy (FA) were calculated for the same white matter and cortex volumes as selected for MRS. MT imaging was performed with a 3D-FLASH sequence. MT ratio (MTR) maps were obtained from pairs of identical images acquired with and without MT saturation pulse. Mean MTR was calculated for again the same volumes. The DTI and MTR results of patients and controls were compared using the one-tailed unpaired student T-test.

RESULTS

Clinical profiles and laboratory examinations

Neurological deterioration started in childhood or adolescence (table 1). Spasticity and ataxia affected legs more than arms. Patients 1 and 2 are presently wheelchair-dependent and have severely decreased manual dexterity. Patients 3, 5, 6 and 7 have a spastic-ataxic gait, but can still walk unsupported and have moderately decreased manual dexterity.
Patients 4 and 8 show mild abnormalities on neurological examination but have no functional problems. Four patients have a distal decrease in position and vibration sense. Four patients have learning problems.

The following laboratory investigations were performed in all or multiple patients and unrevealing: in blood hematology and chemistry panels, blood gases, ammonia, creatine kinase, cholesterol, vitamin E, vitamin B\(_12\), folic acid, thyroid function, karyotype, amino acids, lactate, pyruvate, very long-chain fatty acids, phytic acid, transferrin, isoelectric focusing, carnitine, copper, ceruloplasmin, cholesterol, and lysosomal enzymes (arylsulfatase-A, galactocerebrosidase, beta-galactosidase and hexosaminidase-A); in urine organic acids, oligosaccharides, purines, pyrimidines, sialic acid and sulfatides; in CSF cell count, protein and lactate. Analysis of mitochondrial function in fresh mitochondria from muscle and mitochondrial DNA did not reveal abnormalities.

Somatosensory evoked potentials with stimulation of tibial and median nerves, obtained in five patients, were delayed in four and absent in one. Sensory and motor nerve conduction velocities (six patients) were normal. Sural nerve biopsy revealed no abnormalities in patient 4.

### Table. Clinical History and Findings

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<th>Characteristics</th>
<th>Patient 1</th>
<th>Patient 2</th>
<th>Patient 3</th>
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<td>(Patient 6)</td>
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n = normal; mo = months; yr = years; + = present; ++ = serious; ↓ = decrease; ↓↓ = serious decrease; ↑ = increase; ↑↑ = serious increase.
Thirty brain MRIs were obtained (for ages see table 1). The full-blown pattern consisted of extensive signal changes within the periventricular and deep cerebral white matter, relatively or completely sparing the temporal lobes and the U-fibers throughout (Fig. 1, 2). There was some increase in extent over time in three of the patients. The signal changes were inhomogeneous with a spotty aspect. The spots had a lower signal intensity on FLAIR images suggesting focal rarefaction. The posterior corpus callosum was always affected; the anterior part was not involved or less seriously. The posterior limb of the internal capsule was abnormal. The cerebellar white matter became involved later, initially in the subcortical regions (Fig. 1d) and subsequently diffusely (Fig. 2d). The oldest patient had some cerebellar atrophy. Within the brainstem, the superior and inferior cerebellar peduncles were involved early, whereas the middle cerebellar peduncles became affected in a late stage. Within the midbrain, the pyramidal tracts and medial lemniscus were abnormal. Within the pons, the structures typically involved were the pyramidal tracts, medial lemniscus, intraparenchymal trajectories of the trigeminal nerves and mesencephalic trigeminal tracts. In late stages, the transverse pontine fibers became abnormal. Within the medulla, signal changes were typically present in the pyramids, medial lemniscus, inferior cerebellar peduncles and anterior spinocerebellar tracts. After contrast (two patients) no enhancement was found.

The clinical severity correlated with the severity of disease on brain MRI. Patient 4 had the mildest clinical picture and MRI abnormalities; patient 1 had the most severe clinical symptoms and MRI abnormalities.

Fifteen spinal MRIs were obtained. They showed abnormalities in the dorsal columns and lateral corticospinal tracts uniformly over the entire length in all patients (Fig. 1). There were no changes on follow-up. There was no evident atrophy.

White matter \( <D > \) was significantly higher in patients (mean 1.07 \( \times 10^{-3} \) mm\(^2\)/s; s.d. 0.30) than in controls (mean 0.81 \( \times 10^{-3} \) mm\(^2\)/s, s.d. 0.07). However, there was a considerable variability and in some patients \( <D > \) was normal. The increase in cortical \( <D > \) was also significant but less pronounced (1.07 \( \times 10^{-3} \) mm\(^2\)/s, s.d. 0.05 in patients; 0.94 \( \times 10^{-3} \) mm\(^2\)/s, s.d. 0.03 in controls). White matter FA was significantly lower in patients (mean 0.22, s.d. 0.04) than in controls (mean 0.37, s.d. 0.04), while cortical FA was normal (mean 0.17, s.d. 0.01 in patients; mean 0.18, s.d. 0.01 in controls). MTR was significantly lower in the white matter of patients (mean 20.5%, s.d. 1.5) than controls (mean 31.8%, s.d. 1.5), and normal in the cortex (mean 24.6%, s.d. 1.5 in patients; mean 25.4%, s.d. 1.2 in controls).

In MRS a significant decrease in N-acetylaspartate (NAA) and increase in myo-inositol (mIns) were found in the white matter of patients (mean NAA 4.6 mmol/l, s.d. 0.8; mean mIns 5.9 mmol/l, s.d. 0.7) as compared to controls (mean NAA 7.7 mmol/l, s.d. 1.0; mean mIns 3.6 mmol/l, s.d. 0.3). The increase in white matter choline (Cho) was also statistically significant, but less pronounced (in patients mean 1.8 mmol/l, s.d. 0.3; in controls mean 1.3 mmol/l; s.d. 0.2). White matter creatine (Cr) was normal (in patients...
Figure 1 shows T2-weighted images of the brain (a - e) and spinal cord (f - h) in patient 6. There are irregular, spotty signal abnormalities in the periventricular white matter (a, b), extending into the posterior limb of the internal capsule (b). Within the brainstem, the intraparenchymal trajectories of the trigeminal nerves (arrow head in c), the mesencephalic trigeminal tracts (white arrow in c), the superior cerebellar peduncles (black arrow in c), the inferior cerebellar peduncles (bold arrow in e) and the pyramids (small arrow in e) are involved. The subcortical cerebellar white matter contains signal abnormalities (d). The sagittal spinal image (f) shows that the cord contains signal abnormalities over its entire length. Transverse images at the cervical (g) and thoracic (h) levels show that the dorsal columns (black arrow) and lateral corticospinal tracts (white arrow) have an abnormal signal. The low signal behind the spinal cord (h) represents flow artefacts.
Figure 2 shows the corresponding images in the more severely affected patient 3. The periventricular white matter changes have an inhomogeneous, spotty appearance (a,b). The posterior part of the corpus callosum is more severely involved than the anterior part (b). The posterior limb of the internal capsule is abnormal. Within the brainstem, the intraparenchymal trajectories of the trigeminal nerves (white arrow head in c), the mesencephalic trigeminal tracts (white arrow in c), medial lemniscus (black arrow in c, white arrow in e), pyramidal tracts (c), transverse pontine fibers (c), superior cerebellar peduncles (black arrow head in c), inferior cerebellar peduncles (bold black arrow in e), pyramids (small black arrow in e), and anterior spinocerebellar tracts (arrow head in e) are abnormal in signal. All white matter of the cerebellar hemispheres is abnormal (d).
mean 4.6 mmol/l, s.d. 0.8; in controls mean 4.5 mmol/l, s.d. 0.6). White matter lactate was elevated in all patients except one (mean 2.4 mmol/l, range 0.5-4.1, s.d. 1.1; in controls 0.2 mmol/l, s.d. 0.3). MRS of white matter in patient 8 in another center showed increased lactate. Cortical spectra were normal.

DISCUSSION

Definition of the syndrome

A novel leukoencephalopathy is described. It is clinically characterized by slowly progressive pyramidal and cerebellar dysfunction, often with concomitant dorsal column dysfunction. The MRI pattern is distinct and includes inhomogeneous cerebral white matter abnormalities and strikingly selective tract involvement. The pyramidal tracts extending downwards through the internal capsule and the brainstem into the lateral corticospinal tracts are affected over their entire length. The sensory tracts, including dorsal columns, medial lemniscus up to the level of the thalamus, and corona radiata above that level are also affected over their entire length. Cerebellar connections are selectively involved. The consistent involvement of the intraparenchymal trajectories of the trigeminal nerve and mesencephalic trigeminal tracts is remarkable. This MRI pattern is uniform among the patients and different from patterns in both classical and recently defined leukoencephalopathies. It does not resemble the pattern of pontocerebellar atrophy observed in the familial spinocerebellar ataxias or the white matter changes observed in some hereditary spastic paraparesis syndromes. At present the diagnosis is MRI-based and requires the presence of typical spinal cord abnormalities in combination with spotty cerebral white matter abnormalities and specific brainstem abnormalities.

Pathophysiology

MRI is very sensitive in detecting white matter abnormalities but lacks specificity with respect to underlying pathology. Other MR techniques may contribute in this respect. In MRS, the abnormal cerebral white matter showed normal to mildly elevated Cho concentrations, suggesting low-grade demyelination. The decrease in NAA and increase in mIns were more pronounced, indicating considerable axonal damage or loss and gliosis, respectively. Gliosis can contribute to Cho elevations. DTI and MTR provide information about tissue microstructure. Within the abnormal white matter of our patients, \( <D > \), a measure for isotropic water diffusivity, was increased in most, whereas FA, a measure for the degree of diffusion anisotropy, was decreased in all; these changes indicate an enhanced mobility of water molecules in all directions as a result of damage to the tissue matrix. The MTR of the abnormal white matter was decreased, probably reflecting loss of axons and myelin sheaths.

Overall, our findings suggest considerable axonal damage or loss in the white matter. The involvement of entire tracts argues in favor of primary axonal degeneration.
There is an element of myelin loss, which could be secondary. It is difficult to explain the early discrepancy between MRI abnormalities and clinical problems. At present the pathologic basis of the disease remains unsolved.

Extensive laboratory investigations did not reveal a cause. Considering the presence of two affected sib-pairs among our patients, an autosomal recessive mode of inheritance is likely. We will initiate a genetic linkage study as soon as sufficient informative families are available.

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REFERENCES


