Letters to the Editor

protected disease course and a failure to improve.1 Acute inflammatory demyelinating neuropathy in association with acute leukaemia has been reported in only a few instances.2,3 However, in contrast to our patient, GBS in these cases was rapid, progressive, and the patient died in a few weeks without haematological remission.1,3 This case is remarkable because the associated neurological symptoms apparently did not adversely affect the therapeutic response and the prognosis of the myeloproliferative disorder. The reasons for this difference are not clear. The neuropathy in these cases is usually related to leukaemic infiltrations.1,3 However, remote effects probably related to transient immunosuppression in acute leukaemia could serve as one contributory cause.4

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Polyunsaturated fatty acids in peroxisomal disorders: a hypothesis and a proposal for treatment

Defects of peroxisomal β-oxydation of fatty acids are characteristic of a group of congenital disorders with severe disturbances of brain function, or of the spinal cord, such as the cerebro-hepato-renal syndrome of Zellweger, infantile Refsum’s disease, neonatal adrenoleukodystrophy, X-linked adrenoleukodystrophy and adrenoleucodystrophy, hyperleptic acidosis, isolated absence of an enzyme of the β-oxydation sequence, and several variants described clinically as “Zellweger-like” and “neonatal adrenoleukodystrophy-like”.1 In all these syndromes very long chain fatty acids (C24-C26) accumulate in plasma. Medical treatment at present succeeds in normalising their level, by suppressing endogenous synthesis by glycerol trioleate and glycerol trierucate administration; but it is doubtful whether this leads to clinical improvement.

It has been shown recently that brain, retina, liver, kidney and erythrocytes of patients with several peroxisomal disorders are depleted of docosahexaenoic acid C 22:6o3 (DHA), a component which normally rises in the brain and retina during prenatal and postnatal development.2 Depletion of DHA occurs in the monkey as well as in children causes disturbances of photoreceptor function (ERG, visual acuity) and peripheral neuropathy.3 Martinez4 orally administered 250 mg of DHA ethyl ester daily for one year to one child with a NALD-like syndrome. DHA erythrocyte levels normalised but in addition there was improvement of psychomotor development and visual acuity.1 It is worth noting that the relationship between peroxisomal β-oxydation and DHA levels is not fortuitous, but is established by the results of Voss et al.5 These authors have shown that in the rat a deficiency of C 22:6o3 proceeds by β-oxydation of C 24:6o3. Obviously, oxidation of C 24 is impaired in patients with a peroxisomal β-oxydation defect and this must result in shortage of DHA. The established block mechanism is confirmed by most recent, and unexpected, data from patients with schizophrenia treated with high doses of pheno-thiazines.6 These drugs are in vivo inhibitors of peroxisomal β-oxydation. The patients' thymocytes became deficient in PUFA, especially arachidonic acid and DHA; while the ratio C 26:0/22:0 increased.4 We propose that patients with an impaired peroxisomal function should be assayed for PUFA levels in erythrocytes or platelets, and in biopsied liver. Brain and spinal cord will still be more informative, as their content is not necessarily reflected in blood cells. Data on X-ALD and AMN, which form the largest group of peroxisomal diseases, are still fragmentary. Although erythrocyte levels were normal in several patients, Martinez et al.5 have shown that DHA in the brain of a single case to be below those of extreme nutritional deprivation; the brain of a patient with adrenoleucodystrophy had normal PUFA; b) patients with a lowered DHA should be treated by DHA supplementation. Such treatments have been initiated by Dr M Martinez, under a European project "Pathogenesis, prevention and treatment of peroxisomal disorders of leukodystrophy", in which more than 30 centres in 13 countries take part. For a correct evaluation it is necessary that objective tests (MRI, neuropathological and other tests) be carried out before treatment starts. The evolution of the PUFA level should be monitored.

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Hepers simplex encephalitis in a patient with complex partial epilepsy: confirmation by the polymerase chain reaction with necropsy studies

Hepers simplex virus 1 (HSV) is a relatively common and eminently treatable cause of encephalitis. Despite its moderately distinctive clinical presentation,1 the clinical diagnosis of herpes simplex encephalitis (HSE) is unreliable,2 and until recently laboratory diagnosis of HSE has been unsatisfactory: isolation of the virus from CSF is rarely achieved; ampicillin therapy has not proved problematic, diagnosis by serology or immunoblotting must rely on the evolving immune response and is therefore delayed and indirect, and brain biopsy is highly invasive. The polymerase chain reaction (PCR) is a method for the amplification of DNA with exquisite sensitivity. In cases of suspected HSE it holds out the promise of early and accurate diagnosis.

We describe a case in which the eventual diagnosis of HSE by PCR was confirmed at necropsy. It illustrates the potential difficulty of making this diagnosis in a patient with pre-existing epilepsy and also exemplifies the occasional occurrence of a false negative result from PCR. This implies that negative results should be assessed critically and that repeated sampling may be worthwhile in difficult cases.

A 73 year old woman was admitted "drowsy but responsive" to her District General Hospital in 1991 following "multi- ple fits". In 1982 she had been investigated because of a five year history of depression, depersonalisation, micropsia, formal visual hallucinations and, rarely, loss of consciousness. Successive EEGs revealed only moderate bitemporal abnormalities, but a CT scan demonstrated a right petrous meningioma with associated intracerebral oedema. The possibility of surgery was considered but rejected. Her epilepsy was controlled by anticonvulsants, and she became prone to periods of prolonged postictal confusion. On admission in 1991 she was drowsy but alert and there were no focal neurological signs. Her level of alertness improved over the following few days, although she remained confused, beligerent and possibly dysphasic. Five days after admission she had a flurry of complex partial seizures, became drowsy and was treated with intravenous antibio-otics. Three days later she was transferred to our care. On arrival she was cyanosed, febrile at 40·5°C, and deeply unconscious. She was severely hypoxic and was ventilated. A chest radiograph suggested left basal consolidation. An EEG showed rhythm-
mic slow wave discharges on the right at 1–2 Hz, occasionally associated with spikes, and more clearly epileptiform activity on the left. Intravenous diazepam abolished the activity on the left, but spared that on the right. The initial diagnosis was of pneumo-

nia with complex partial status epilepticus. On the following day the EEG record was dominated by widespread, irregular, theta wave activity with a period of approximately two seconds. Unenhanced CT scan appearances had not, however, changed since 1988. A lumbar puncture was performed but the cerebrospinal fluid pressure had failed to improve. The CSF was under a pressure of 24 cm and contained 52x10^6/L leucocytes (93% lymphocytes) with normal CSF protein and CSF glucose ratio. A further lumbar puncture two days later showed worsening abnormality and treatment was started with intravenous Acyclovir, rifampicin, isoniazid and pyrazinamide. Over the following nine days headache, reflux responses deteriorated and she died.

A polymerase chain reaction for the amplification of HSV-1 DNA was carried out post-mortem using the primer sequence described by Aurelius et al, with some modifications. Reaction conditions for first and second round PCR were as described previously. The patient’s second CSF sample and the positive control contained 500 copies of HSV DNA after two rounds of PCR (fg). The first CSF sample, taken two days before, was negative. Neither oligonucleotide bands of total IgG nor antigen specific oligonucleotide bands were detected in either CSF sample. Viral culture and viral titres were negative in the first CSF sample; in the second sample viral culture was again negative but a weak IgG reactivity to herpes simplex virus 1 was detected by ELISA.

Post mortem examination confirmed the presence of a right petrous meningioma. The brain was swollen and soft, with uncal and cerebellar tonsillar herniation. Temporal and insular cortex were involved in a marked meningocoecephalitis, with necrotising venulitis and perivascular infil-

tration by lymphocytes and macrophages. Immunoperoxidase staining revealed a very striking positive reaction for herpes simplex virus antigens in neurons, macrophages, and many cells whose nature could not be identified.

This complex case emphasises the importance of prompt treatment with Acyclovir where there is clinical suspicion of the diagnosis of HSE. In general the diagnosis should be considered in any patient with fever and depression of consciousness: suspicion should be heightened by accompanying abnormalities of behaviour, focal seizures or signs, especially dysphasia, evi-

dence of acute temporal lobe pathology from neuroradiology or EEG, and a CSF lymphocy-

tosis. A number of factors conspired to reduce clinical suspicion in the present case, in particular the patient’s long history of epilepsy with pronounced post-ictal con-

fusion and the demonstration of a severe pneumonia. In retrospect, the onset of encephalopathy with drowsiness, seizures, fever and new EEG disturbance was fully in keeping with the eventual diagnosis of HSE. It is of course always important, even in the context of a patient with a chronic disorder, to assess an acute illness on its own merits.

Although PCR is highly sensitive and specific, the negative result from the first CSF sample reminds us that all laboratory tests give rise to occasional false negative results: it is advisable to test more than once in difficult cases, and a negative PCR result should not preclude the use of acyclovir where HSE is suspected clinically.

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4 Wakefield AJ, Fox JD, Sawyer AM, et al. Detection of herpes virus DNA in the large intestine of patients with ulcerative colitis and Crohn’s disease using the nested poly-


Matters arising

Accuracy of clinical diagnosis of idiopathic Parkinson’s disease

I read the paper by Hughes et al in the journal with much interest and wish to compliment them for their work.

The percentage of inaccurate clinical diagnosis of idiopathic Parkinson’s disease (IPD) in their study is identical to that which we reported last year.2 Their observations, however, are different from our study in several respects. For example, the largest subgroup of patients (5 cases) which were erroneously diagnosed as having IPD by Hughes et al had progressive supranuclear palsy (PSP), in contrast, all the necropsy proven PSP cases in our study were recog-
nised before death.2 The reasons for that difference are unclear. The final clinical diagnosis in all our patients was made by the same neurologist.1 Hughes et al did not indicate the number of neurologists and the geriatricians involved in the evaluation of their patients and those contributing to the brain bank. The larger the number of clinicians assessing the patients, the greater would be the variability and the chance of error.

The second possibility for the misdiagno-
sis is that the patients may have been evalu-
ated during an early stage of illness, before the features characteristic of PSP were evi-
dent. The clinical data in our cases were collected prospectively and we were able to assess the issue of diagnostic accuracy based on each, the initial and the final clinical diagnosis before death. While only 65% of our cases whose death did not have PSP diagnosis were retested, we had Lewy body pathology, the diagnostic accuracy increased to 76% by the time the final assessment was done—mean 12 years after onset. Most cases who had other vari-

ants of Parkinson’s syndrome (PS) were recognised within 5 years of onset. It is unclear in the report if they relied on initial or the final clinical diagnosis.

Another aspect that we made a note of but is not clarified in their study1 is the method used to assign one diagnosis to a patient. We considered the clinical diagno-
sis correct only if it was the sole diagnosis or the pathological diagnosis was stated at the top of diagnostic possibilities.2 Several variants of PS, including PSP,1 were clinically and pathologically identified in the early 1960s. Those patients who were evaluated before 1983 were more likely to have a larger proportion of inaccurate diagnosis by contemporary standards. Their paper does not indicate if some of the errors in diagnosis may be related to the calendar year of the patient assessment.

In addition to our paper, there is a small study by Forno1 which addressed the issue of the accuracy of clinical diagnosis in IPD. Hughes et al retrospectively analysed patients using the UK Parkinson’s Disease Society Brain Bank (PDSBB) clinical diagno-
sic criteria. By those criteria, 11 patients did not have IPD, yet 3 (27%) of those turned out to have Lewy body pathology. They have not discussed the reasons for this significant error. Whatever the reason, it illustrates that no diagnostic criteria are fool-proof.

I suggest a minor amendment to the PDSBB clinical criteria for consideration by the committee. Postural instability which has been recommended as one of the two major manifestations necessary for making the diagnosis of PS should be deleted. The diagnosis of PSP is frequently present in a large segment of normal elderly people. Most clinicians looking after elderly people recognise that individuals in this age group lose balance rather easily compared with younger age groups. Postural instabil-

ity, as evaluated in the PS, was carefully studied by Weiner et al in the elderly. When all the neurological, mechanical and other possible causes for postural reflex-related impairment were excluded, they noted that postural instability was an age-related pheno-

menon.6 While 43% of those between age 60–69 years had impaired postural reflexes, 70% between 80–89 and 100% of those between age 90–99 years had impaired post-

ural reflexes.7

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