Biochemical vitamin deficiencies in Friedreich's ataxia

Sir: In their studies of patients with Friedreich's ataxia, Purkiss et al. recorded that five of 18 patients "showed enhanced response to thiamine stimulation of red cell transketolase" and two patients had "very low values for red cell transketolase". Prompted by these findings, we determined the vitamin status in two patients with Friedreich's ataxia.

Our patients were a 23-year-old coloured male (GI) and his 22-year-old sister (CI). Each patient fulfilled recommended clinical criteria for the diagnosis of Friedreich's ataxia. Their clinical deficits were similar in severity (Group 1 of Dyck and Lambert) and extent. Symptoms of neurologic disorder had been noticed by GI since the age of 14 years and by CI since the age of 15. In both patients, electrophysiological studies confirmed the presence of peripheral neuropathy and revealed abnormalities of visual and brainstem auditory evoked potentials. Red cell transketolase, TPP effect, nicotinic acid, vitamin B_{6} and ascorbic acid were determined according to previously published methods. Serum and red cell folate and vitamin B_{12} were determined by the Amersham Combination Radioassay Kit and riboflavin coefficient by the method of Nichoals. Vitamin profiles of the two patients on admission and after 16 days in hospital are recorded in the table. In both patients, the following investigations were normal on admission: Serum albumin, transferrin, folate, vitamin B_{12} and fasting cholesterol; red cell folate; and fasting plasma glucose and triglycerides. On admission, red cell transketolase was in the low range of normal in both patients and patient GI had a markedly raised TPP effect. These findings are in accord with those of Purkiss et al. and, on the basis of the levels of transketolase activity, indicate chronic thiamine deficiency in patient CI and acute-on-chronic deficiency in patient GI. Since neither patient received any medication throughout their admission the correction of these deficiencies after 16 days on hospital diet is compatible with a dietary origin for their deficiency, rather than one of abnormal thiamine metabolism. Further support for this concept is provided by the concomitant tendency towards normalisation of riboflavin and ascorbic acid values, which were also indicative of deficiency at the time of admission. In addition, dietary analysis revealed that vitamin intake in both patients was less than that recommended. (Details are obtainable from Dr D Labadarios, PO Box 63, Tygerberg, 7505, South Africa).

Whether dietary deficiency was operative in the patients of Purkiss et al. cannot be judged from their published data. At all events, their findings and those reported here would seem sufficient reasons for further studies on the vitamin status of patients with Friedreich's ataxia. The detection of thiamine deficiency and its correction may be of special importance in these patients, given the cofactor function of thiamin for pyruvate dehydrogenase, the role of this enzyme in energy metabolism, and Barbeau's proposal that defective energy metabolism may be a key factor in the pathogenesis of Friedreich's ataxia.

References


Purkiss et al reply

Thank you for the opportunity to reply to the comments of Drs Gledhill and Labadarios on the vitamin status in Friedreich's ataxia arising from our paper (J Neurol Neurosurg Psychiatry 1981;44:574-80). In our group of patients, direct blood vitamin assays were not performed nor was a dietary vitamin analysis made. Thiamine status was assessed by measurements of erythrocyte transketolase activity and the effect of thiamine on this enzyme.

Our in vivo data indicated that there was...