(3) In considering the dosage of pyridoxine used in treatment, in mg/kg, it is not valid to compare a healthy adult to a child with proved pyridoxine dependency, a condition requiring higher than physiological doses of pyridoxine to maintain normal concentrations.12

(4) There is no doubt that the minimum effective dose is best. We did not investigate the efficiency of maintenance doses of less than 75 mg pyridoxine/day because of evidence from the published reports that the effective dose is 40 to 100 mg/day.10-12 Doses as high as 200 mg/day have been required9 to control seizures. In considering the optimal dosage, the potential hypoxic damage from recurrence of seizures cannot be ignored.

The conclusion of Dr Reynolds that our 'treatment may, in fact, have contributed substantially to the impaired mental development noted ...' is not supported by our cases. The findings of others in man were of peripheral neuropathy, and 'the central nervous system was spared'1 in man and in beagle dogs.2

We encourage the diagnostic use of pyridoxine in a baby with intractable seizures. Once a definite clinical response has been established treatment should be continued indefinitely: the optimal daily dosage, however, may need to be reviewed.

References


Metabolic alkalosis in a neonate after frusemide

Sir,

Acute metabolic alkalosis is uncommon in newborns. When it occurs it is usually secondary to electrolyte imbalance, fluid depletion, chronic hypercapnoea or, more rarely, exchange transfusion with citrated blood.

We describe a term neonate of birthweight 2.52 kg who developed acute profound metabolic alkalosis after intravenous frusemide. At the time the baby was on nasal constant positive airways pressure for pulmonary hypoplasia secondary to chronic amniotic liquor loss during the pregnancy. Frusemide (1 mg/kg) was administered intravenously on two occasions (four hours apart) because of chest x ray changes suggestive of interstitial pulmonary oedema. After the second dose of frusemide the baby developed a metabolic alkalosis with pH 7.66, HCO₃ 58.5 mmol/l, base excess 28 mmol/l, PCO₂ 50 mm Hg. Her plasma electrolytes were urea 3.5 mmol/l, potassium 5.2 mmol/l, and sodium 135 mmol/l. The frusemide was stopped and the alkalosis corrected spontaneously in 14 hours. When a further dose of frusemide was administered two days later a similar episode of metabolic alkalosis occurred which again resolved spontaneously when the drug was stopped. Chlorothiazide was subsequently prescribed without side effects.

Frusemide is highly bound to albumin—more than 99% in adults and at least 95% in neonates. In the first week of life frusemide clearance is only 10% of the adult rate.1 Thus, the prolonged half life may potentiate its action, particularly if doses are given less than 12 hours apart. De Rubentis et al2 suggest three mechanisms that may contribute to its production of alkalosis, namely potassium depletion, renal loss of hydrogen ion, and extracellular fluid space contraction without accompanying bicarbonate loss. The latter, contraction alkalosis, may have played an important role in our patient. In patients at risk from a metabolic derangement it has been suggested that the dose of frusemide should be administered every other day or is combined with a potassium sparing agent.3

We suggest that metabolic alkalosis should be anticipated and looked for when neonates are given frusemide, particularly if the dosage interval is less than 12 hours.

References


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