Correspondence

Congenital nephrogenic diabetes insipidus

Sir,

Schreiner and co-workers reported a baby girl with congenital nephrogenic diabetes insipidus (NDI) (Archives, 1978, 53, 906). I should like to contribute a case diagnosed as having partial NDI.

This boy had been transferred from a local hospital at age 9 months because of high fever with high serum Na and Cl concentrations, initially noticed in the first week of life. His mother showed polydipsia and polyuria: her urine osmolality after water deprivation for 12 hours was 270 mmol/kg. Investigation of the patient showed high serum osmolality (315 mmol/kg), low urine osmolality (100 mmol/kg), serum Na 172 mmol/l, and serum Cl 126 mmol/l.

Subsequently clearance studies were performed while the boy was receiving both a normal and a salt-restricted diet, after water deprivation, and after intravenous DDAVP (Aronson and Svenningsen, 1974). The main results were: (1) On water deprivation the patient was able to concentrate urine up to 600 mmol/kg simultaneously with reduction of urine volume; after DDAVP (0.4 ml IV) urine volume decreased further; osmolar clearance, which had increased during water deprivation, decreased after DDAVP. (2) On normal diet and on salt-restricted diet DDAVP resulted in increasing osmolar and free water clearance (and urine volume).

We made the following interpretations on the effect of DDAVP (a synthetic vasopressin analogue) on our patient’s kidneys: renal resistance to DDAVP was incomplete; there was an adequate response after giving DDAVP during water deprivation, whereas a paradoxical effect on osmolar and free water clearance (Brodehl et al., 1965) was apparent during the control periods.

Partial NDI (McConnell et al., 1977) was diagnosed and treated by salt restriction and frusemide (15 mg/day). Follow-up 3 years later shows uncomplicated physical and psychological development of the patient with normal or almost normal values for serum Na, Cl, urea nitrogen, osmolality, and bicarbonate. Urine osmolality ranged from 200 to 265 mmol/kg.

I wonder if Schreiner and co-workers would like to comment on our patient, particularly with regard to their own therapeutic experience.

References


Myotonic dystrophy and bonding failure

Sir,

In their report of 5 cases of the neonatal form of dystrophia myotonica (Archives, 1979, 54, 331), Pearse and Höweler emphasise the ethical problems facing the clinician once the diagnosis has been made, and they discuss the need for genetic counselling. We should like to add support to their concern for earlier diagnosis by drawing attention to another important and potentially lethal hazard run by babies with this condition—that is bonding failure. If they survive the neonatal period, these babies must always be considered to be at increased risk of rejection and neglect. Not only are they likely to have accumulated a number of the features commonly associated with child abuse—for example, neonatal separation, early ill health, maternal physical and emotional illness (Lynch, 1975; Lynch and Roberts, 1977)—but both mother and child are further handicapped by the inability to use facial expression effectively as a means of communication.

This was well illustrated by a family referred to the Park Hospital in Oxford. The parents were both very young and came from unhappy and divided homes, thus the potential for abuse was high. The mother’s relationship with her own mother was hostile yet dependent, and the father had experienced a rigid authoritarian upbringing. When we met the mother she was just 20 and had had two caesarean sections within 13 months. Her second son had been very ill in the neonatal period, spending the first 7 weeks of his life in the special care baby unit. He had required assisted ventilation for some days. During his stay in the unit the diagnosis of myotonic dystrophy was made both in himself and his mother. A distaff family history of dystrophia myotonica was discovered, with the mother’s own mother, the grandmother, and a great aunt suffering from cataracts and facial diplegia. It was also found that the elder brother had a milder form of the disease.

When finally discharged home, the baby had a floppy, expressionless face; he made none of the grimaces and little noises a normal baby makes. His abnormal cry was quiet and ineffective; his smile did not appear. Feeding was a nightmare. Much of his time he slept ‘like a dead
Correspondence

Early use of sodium nitroprusside in respiratory distress syndrome

Sir, Beverley et al. (Archives, 1979, 54, 403) reported the response to sodium nitroprusside in an infant with hyaline membrane disease, and I should like to make some comments on their patient.

The course of the illness, with the infant not requiring added oxygen and only very low ventilator pressure by 24 hours of age, must cast doubt on the diagnosis of surfacant-deficient hyaline membrane disease, and suggests that the baby was suffering from the after effects of severe birth asphyxia, the early problems of intrapartum pneumonia, or perhaps from some iatrogenic disease.

During the time sodium nitroprusside was being infused, 3 other major changes in treatment took place which may have been responsible for improving the baby's condition. Firstly, his pH was corrected. The effect of this on pulmonary perfusion and pulmonary vascular tone is well known, and could itself be responsible for the improved oxygenation in the infant. Secondly, the inspiratory to expiratory ratio was reduced from 4:1 to 1:5:1 and while we do not know the ventilator pressures that were sustained during this period, it seems likely that this would be associated with a considerable fall in the mean airways pressure. Particularly in infants without hyaline membrane disease, a high mean airways pressure can be responsible for serious pulmonary underperfusion, and lowering the airways pressure as was done in this patient, can be associated with a large increase in arterial P02. Thirdly, they transfused their patient, and while we are given no blood pressure data before this was started, the fact that they were able to increase the baby's blood volume by more than double suggests that, irrespective of the hypotension-producing effect of the nitroprusside, the infant was previously hypovolaemic or hypotensive, or both. Correction of that could also be responsible for an improvement in the infant's condition and oxygenation.

I would submit therefore that Beverley et al. have provided absolutely no evidence for a beneficial effect from sodium nitroprusside in their patient, and that before this comparatively dangerous drug is used, evidence that it improves oxygenation while other factors in the treatment are held constant is required. This, for example, has been provided for tolazoline (McIntosh and Walters, 1979).

Reference


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Lord Justice Scott and the test of guilt

The letter from Beverley et al. (Archives, 1979, 54, 403), and the report by Abbott et al. (1978) to which they refer, should both be treated with extreme caution. In neither is a good case made for the use of sodium nitroprusside to reduce pulmonary vascular resistance (PVR) in severely hypoxic newborn babies. While Beverley and colleagues refer to the known hazards of this hitherto untied drug, they give no details of the state of the circulation or of the Paco2 during their infant's first few hours. These are important since poor tissue oxygenation and hypercapnia may both contribute to acidosis, and PVR in the newborn may be expected to increase as pH falls (Rudolph, 1977). The extreme acidosis at 2-4 hours of age was presumably the legacy of severe birth asphyxia.

Nevertheless, the information provided suggests that after a blood transfusion (which should have improved tissue oxygenation) and during the slow correction of the severe acidosis, there was initially a steady fall in right-to-left shunt, followed at the age of 11 hours by a more rapid improvement in both the arterial pH and PaCO2.