theory remains speculative as the animal studies have shown no evidence of strings or bands as the cause of the limb deficiency. If the vascular compromise occurred at the time of premature rupture of the membranes there would be adequate time for healing of lesions and resorption of amputations. Healing in the fetus is particularly fast and relatively non-scarring.⁷ It is likely that this child's limb defects were related to amnion rupture, and are not associated with Van der Woude syndrome.

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Fetal valproate phenotype is recognisable by mid pregnancy

SIR,

In the November 1987 issue of your journal, Winter *et al*¹ reported four infants who were exposed to sodium valproate or valproic acid during pregnancy. Three surviving infants showed the characteristic facial features of the fetal valproate syndrome described by DiLiberti *et al*² and confirmed by several authors.³ We report here a 22 week fetus with a large myelomeningocele and similar facial abnormalities.

The mother of this subject, a 23 year old primigravida of Iberian origin, attended a psychiatric clinic and suffered from grand mal epilepsy. She received sodium valproate (500 mg twice a day) from the age of 18 years in combination with clobazam (10 mg, three times a day). When she became pregnant, she was referred to us for prenatal diagnosis of open neural tube defect.

At 18 weeks of gestation, serum and amniotic α fetoprotein (AFP) concentrations were evaluated by

radioimmunoassay. Maternal serum AFP was raised ($2.9 \times \text{MoM}$). The amniotic fluid, obtained by amniocentesis, was clear and blood free, with an AFP concentration within the normal range ($<3 \text{ MoM}$). Ten ml of amniotic fluid were centrifuged at 14 000 g for five minutes to remove any red cell membrane contamination. Quantitative measurements of acetylcholinesterase on unfrozen material

FIG 1 Dorsal view of the fetus.

FIG 2 Fetal spine after dissection.

FIG 3 Front and side view of the fetal face.

analysed within 24 hours showed a normal activity: 5.5 U/l (normal 5.8 U/l). Isozymes of acetylcholinesterase, examined by polyacrylamide gel electrophoresis, gave two gel bands; the extra band of acetylcholinesterase had the same appearance and position as acetylcholinesterase from cerebrospinal fluid and was inhibited by means of the specific inhibitor BW284C51.

The mother was advised to have an ultrasonic study of the fetal spine. This was done one month later and showed a lumbosacral meningocele. Consent to terminate the pregnancy was obtained. Immediate examination of the female fetus showed no growth retardation: weight 450 g, total length 28 cm, head circumference 18 cm, foot 41 mm.

There was an elongated spina bifida aperta (30 mm) involving five lumbar and two sacral vertebrae with spinal cord showing in the upper part. The spinal cord was open and spread out. In the lower part, only the filum terminale was visible (figs 1 and 2). There was a brachycephalic skull and facial peculiarities (fig 3) including bulging ocular globes, infraorbital grooves, broad and flat nasal bridge, increased inner canthal distance (20 mm), very long upper lip, thin vermillion borders, small oral opening, and prominent antihelices of the ears. No visceral abnormality was found. The cerebellum was small and partially penetrating the foramen magnum, simulating an Arnold-Chiari malformation. The fresh brain weighed 51 g. Frontal sections of the brain after fixation showed enlargement of the lateral and third ventricles.

An open neural tube defect was detected in the fetus by positive maternal serum AFP levels and by two bands on acetylcholinesterase electrophoresis of the amniotic fluid. It should be noted that the amniotic fluid concentration of AFP and acetylcholinesterase activity were within normal limits.

Owing to the difficulty of evaluating a fetal face, the existence of a valproate phenotype could be questioned in our case. However, there was such a markedly typical abnormal facial appearance that it was immediately attributed to the sodium valproate therapy taken by the mother. Thus, as might be logically expected with a teratogen, the valproate phenotype is already apparent by mid pregnancy.

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