DOES A HYPOALLERGENIC MILK IMPROVE SLEEP DISTURBANCE IN YOUNG INFANTS?

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We report a prospective double-blind randomised cross over trial designed to compare the effect of a hypoallergenic formula NAN-HA with a standard cows' milk protein (whey dominant) formula NAN on 67 infants (45 M, 22 F) aged 6 weeks to 6 months. Records of sleep, crying, diarrhoea, colic, regurgitation, skin rashes and coughs and wheezing were recorded for 2 three-week periods (one on each formula) following an initial one week

Night sleep (between 8pm and 8am) was confirmed to be less initially in the "sleep disturbed" infants compared to controls (p < 0.01). Sleep improved in the infants whether allocated HA or NAN first and improved throughout the 6 weeks of the trial to be no different from control infants at the end. There was no significant difference between the two groups with respect to any of the symptoms studied except that loose stools/diarrhoea were significantly increased (p < 0.01) on HA. 22 mothers completed the cross over design; 14 expressed a preference for the HA formula and only 8 for NAN. We cannot therefore confirm that the hypoallergenic milk (HA) improves sleep in a group of sleep disturbed infants.

ALLERGY PREVENTION IN CHILDREN OF PARENTS WITH ATOPIC DERMATITIS (AD)

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The effects of 4 feeding regimes on the development of atopic disease and sensitization against cow's milk proteins were compared in a prospective study of 522 children with a 2 year follow-up. Healthy newborns were assigned to 4 groups: (Br)Exclusively breast fed; (Br+HA)Breastfed and hypoallergenic formula; (Br+CM)Breastfed and cow's milk; (CM)Cow's milk. Serological investigations included IgG-,IgA- and IgE-antibodies against 4 CM-proteins interpreted by a geometrical mean titer (gmt)(Bürgin-Wolff, Eur J Ped 1980:133). Results: The percentage of children developing clinical signs of atopic disease is given in the table.

Office the parental allergy— and AD-status.CM-antibodies in breastmilk were not protective.Children of parents with AD developed allergy most frequently.Feeding HA formula is only of limited benefit.

CEREBRAL BLOOD FLOW/CIRCULATION

COMPARISON BETWEEN 133XENON CLEARANCE AND NEAR INFRARED SPECTROSCOPY FOR ESTIMATION OF CEREBRAL BLOOD FLOW

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Estimations of cerebral blood flow (CBF) were attempted by both ¹³³Xe clearance and near infrared spectroscopy (NIRS) on 11 occasions in 9 infants (4 girls and 5 boys). The median (range) gestational age was 27 (26-29) weeks and postnatal age was 3 (1-10) days. All infants were receiving mechanical ventilation and cranial ultrasound or autopsy revealed that while 1 infant had no evidence of cerebral injury, 3 infants had intraparenchymal lesions, and 5 had intraventricular haemorrhage; 6 infants died.

In each study one estimation of CBF by ¹³³Xe was attempted; the measurement was technically inadequate in 2 cases. Within 1-9 (median 5) hours 3-8 estimations by NIRS were made, 26/61 measurements were technically inadequate, but at least 2 were possible in every infant. Comparisons were thus possible on 9 occasions. ¹³³Xe measurements ranged from 4.6-13.2 ml100g ¹min ¹ and mean NIRS measurements ranged from 8.6-20.4 ml100g ¹min ¹. The mean difference between the methods was 2.3, and the limits of agreement were -5.0 to +9.7 ml100g ¹min ¹.

Considering the different principles involved and the time gap between estimations, this study shows reasonable agreement between the methods.

ESTIMATING CEREBRAL BLOOD FLOW IN NEWBORN INFANTS: COMPARISON OF NEAR INFRARED SPECTROSCOPY AND 133XE-CLEARANCE.

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A new method of measuring cerebral blood flow in newborn infants by mean of near infrared spectroscopy (CBFnirs) was compared with the intravenous '''Xe clearance technique (CBFxe). Forty CBFnirs measurements were obtained during 19 '''Xe measurements in 16 infants. The test-retest variation of repeated NIR-measurements during each '''Xe-clearance was 17.5%. CBFnirs was closely related to CBFxe ($r^2\!=\!0.77,~p\!<\!0.0001)$ slope $0.75\pm0.064,~intercept of 1.55\pm0.54~ml/100g/min. We found the best agreement between the two methods in the low range of CBF, whereas the NIRS method underestimated CBF in the high range of CBF due to methodological constrains. The NIRS method may have limitations of application in clinical research, but it has the advantage of being non-invasive and does not involve ionizing radiation.$

CEREBRAL BLOOD FLOW VELOCITY IN THE HIGH RISK NEWBORN - AN INDICATOR OF EARLY PROGNOSIS

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In 175 high risk neonates (87:>2500 g; 50:2500-1500 g; 38:<1500 g) cerebral blood flow velocities (CBFV) were measured by transcranial Doppler in the basal cerebral arteries on the 1st, 3rd to 5th and 8th to 10th day of life. They were compared with values of healthy neonates. Cranial ultrasound was performed on all babies. On the 1st day 66% of the babies had normal, 23% decreased, and 12% increased CBFV. On day 8 to 10 87% had normal, 6% decreased, and 7% increased CBFV. In babies below 1500 g, normal CBFV were associated with the lowest incidence of cerebral hemorrhage (h) and death (d) (6/23 h; no d) while increased CBFV indicated poor prognosis (8/12 h; 7/12 d). Babies with decreased CBFV showed more hemorrhages and deaths than those with normal CBFV. A similar incidence of hypoxic ischemic encephalopathies, hemorrhages and deaths was found in neonates more than 1500 g with both normal or abnormal CBFV. Conclusion: CBFV measured by Doppler are useful as an early prognostic parameter in infants below 2500 g; in higher birth weights they may be useful for

COLOR DOPPLER ECHOENCEPHALOGRAPHY

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From January to December 1990, 100 color doppler echoencephalograms were obtained in 40 infants without cardiopulmonary affections and with a mean birth weight of 2940g (range 1760-3420 g), a mean gestational age of 37.2 weeks (range 34-40w). The examinations were performed using a real-time 2-D Doppler Scanner and a 3,75 MHz sector transducer. Serial scans were obtained in sagittal, coronal and axial plans. Detection rate of intracranial vessels was: internal carotid, anterior cerebral, pericallosal, basilar, vertebral artery (100%); callosomarginal artery (90%); frontopolar artery (82%); medial and lateral striate artery (17%); internal, great cerebral vein(GCV) and straight sinus (\$\$100%); superior sagittal sinus (69%); occipital sinus (39%). Flow mean velocity of GCV and SS was 6-10 and 15-24 cm/sec respectively. Usually vein curve has a continuous profile (therefore we can calculate only mean velocity) while sometimes, expecially in the biggest vein (\$\$S\$), we observe particular aspects of vein flow similar to artery's one with a sort of systolic and diastolic phase. Probably this waving venous curve is within respiratory control becouse the "squeeze" of superior cava vein may induce an increase of vein flow velocity during inspiration.