EARLY SYSTOLIC TIME INTERVALS IN NEONATAL RESPI 51 RATORY DISTRESS

51 RATORY DISTRESS RATORY DISTRESS K.Heinonen and A.Hakulinen, Children's Hospital University of Kuopio, 70210 Kuopio 21, FINLAND In this study we prospectively evaluated cardiac perfor-mance in 184 neonates admitted consequtively to neonatal intensive care because of early-onset respiratory dis-order. We measured right and left ventricular systolic time intervals (RVSTI and LVSTI), blood pressure and heart rate at admission (2.9 + 0.8 h), at 5-7 h and at 24-26 h. <u>Group A</u> (N=161) consisted of term neonates with mild, moderate or severe transient tachypnoea (TTN), or with meconium aspiration (MA). Controls (N=14) were born through spontaneous vaginal delivery without medi-cations. <u>Group B</u> (N=23) consisted of pre-term infants with severe IRDS. Controls (N=10) were pre-term infants with severe IRDS. Controls (N=10) were pre-term infants without respiratory symptoms and increased FiO₂-require-ments. <u>adm 5-7 h</u> 24-26 h adm 5-7 h 24-26 h Group A Mild TIN RVSTI 0.42±0.04 0.46±0.03 0.34±0.05 0.32±0.06 0.38±0.04 0.31±0.04 LVSTI

Severe TIN+MA RVSTI 0.59±0.08 0.61±0.07 0.64±0.06 LVSTI 0.36±0.10 0.52±0.07 0.58±0.08 Group B Severe IRDS

Group B Severe IRDS RVSTI 0.36±0.05 0.37±0.09 0.38±0.07 LVSTI 0.49±0.07 0.53±0.09 0.58±0.10 ▲= prolonged over that observed in controls In severe respiratory distress of term meonates, pulmo-nary hypertension is an early and important factor; later at the age of 6 h and thereafter, signs of impaired left ventricular performance emerge. In pre-term infants, res-tricted left ventricular performance appear at early stages of respiratory disease, and may be associated with insufficient maintenance of systemic blood pressure.

 $52^{\mbox{Enzyme induction following prenatale exposure}\atop_{\mbox{to anticonvulsants measured by ^{13}C-breath tests}\limits_{\mbox{D.RATING* 1}, \mbox{H. NAU* 2}, \mbox{H. Helge1}}$ Department of Pediatrics] and 21nstitute of Embryonal-pharmacology, Free University of Berlin, GFR

Department of Pediatrics] and 2)Institute of Embryonal-pharmacology, Free University of Berlin, GFR Enzyme activity for demethylation processes can be esti-mated in vivo non-invasively by 13 C-breath tests (BT). After oral intake of stable isotope labeled 13 C-amino-pyrine (AP) (2 mg/kg) resp. 13 C-methacetin (MAC) (1,5 mg/kg) 13 CO₂-concentration in breath samples measured by ratio mass-spectrometry will reflect cytochrom P450 dependent AP-N-demethylation resp. P448 MAC-O-demethy-lation. Neonates of epileptic women exposed prenatally to anticonvulsants were studied by 13 C-AP-(n=25) and 13 C-MAC-BT (n=18) while 6 non-exposed newborns served as controls. Half life times of diaplacentally acquired anticonvulsants were determined in 14 resp. 7 of the exposed neonates. The 2h cumulative 13 C-dose)) was significantly (p<0.005) above those of non-exposed as significantly (p<0.005) above those of non-exposed as in 13C-MAC-BT (18.9: 13.8-25.8 versus 9.7: 5.3-18.0) (p<0.05). In normal neonates AP-N-demethylation amounts to only 15 % of those values found in older children aged 2 y or more, while MAC-O-demethylation at that time is sig-nificantly higher (30 % of older controls). Nowever, the intrauterine exposure to anticonvulsants will induce both demethylation processes to the same degree (60 % of older controls). Enzyme induction estimated by $^{13C-}$ BT did not correlate in all instances with half life time of anticonvulsants determined in the same indivi-dum,reflecting the multiplicity of enzyme systems and selectivity of the different tests.

53 EYE FINDINGS IN THE FETAL ALCOHOL SYNDROME (FAS)

Kerstin Strömland M.D, Department of Pediatric Ophtal-mology (sponsored by Ragnar Olegård M.D. Department of Pediatrics.) University of Göteborg, Östra sjukhuset 416 85 Göteborg, Sweden. 30 children with FAS were examined regarding eye find-ings. Typical facil features for FAS were noted, like ptosis, small palpebral fissures and strabismus. The most frequent findings were anomalies of the optic nerves and retinal vessels with more than 50 % of the children being affected . 29 % of the eyes had a visual acuity of 0.2 or less. Another 56 % had mode-rately reduced vision. A control study consisting of 22 children born by mothers who had not been abusing alcohol during pregnancy did not reveal any abnormali-ties of the eyes. Alcohol abuse during pregnancy seems to be a major cause of congenital anomalies of the optic nerves and retinal vessels.

54 PLASMA VITAMEN A AND ZINC IN CYSTIC FIEROSIS (CF) M.2. MICHAL, F.E. WELLS. G.H. HAMPLETON ROYAL MANCHESTER CHILDREN'S HOSPITAL and BOOTH HALL CHILDREN'S HOSPITAL, MANCHESTER, UK

Zinc is required for the synthesis of Vitamin A (Vit.A) carrier proteins - retinol binding protein (RBP) and prealbumin (PA). Low plasma zinc and Vit.A have been observed in CF. We have studied the relationship between plasma levels of Vit:A, RBP and PA with plasma zinc in 37 patients with CF and 25 similar aged controls. There was no statistically significant difference between CF patients and controls for plasma Vit.A and zinc concentrations. In CF patients controls for plasma vit.A and zinc concentrations. In CP patients there was a significant correlation between Vit.A and REP (r = 0.69, p < 0.001) and PA (r := +0.80, p < 0.001). In control subjects there was a significant correlation between Vit.A and REP (r = 0.56, p < 0.01) but not with PA (r := +0.77, NS). In CP patients REP and PA were correlated with the following

	RBP		PA	
	r	p	r	p
Shwachmann score	+0.48	<0.01	+0.50	<0.01
Ht. Velocity (kg/yr)	+0.46	<0.02	+0.48	<0.01
Wt. Velocity (cm/yr)	+0.05	NS	+0.38	€0.05
Serum zinc	+0.03	NS	+0.19	NS

Thus we were unable to find evidence of low plasma Vit. A or plasma zinc concentrations in a fairly healthy (median Stwachmann score 82) group of patients with CT. There was no correlation between plasma zinc and Vit.A carrier proteins.

Sudden infant death (SID): Histological study 55 of the external arcuate nucleus (EAN) of the

5.5 brainstem S.PUGATSCH* and G.MOLZ*(sponsored by R.ILLIG) Department of Anatomy University of Zürich CH8057 Zürich/Switzerl.

A current hypothesis linking SID to a central nervous system dysfunction due to absence of the EAN prompted the present study. Histological examinations of the brainstem were carried

The present study. Histological examinations of the brainstem were carried out on 13 term and 3 preterm infants who had suddenly died at home between the third and 29.week of life. The control group consisted of two term infants, age 5 and 9 months who had died due to self-strangulation, two term newborns, age 3 and 18 days, dying from respiratory dis-tress and peritonitis, respectively and in addition of 8 fetuses at 17 to 33 weeks gestational age. Each brainstem was sectioned serially from the caudal pons to the end of the medulla at 6 -8 um intervals. Every fourth section was stained with hematoxylin and eosin and in each case 140-160 sections were analyzed. Neurons were counted separately in two areas in which the central chemoreceptors may exist, one located ro-strally of the hypoglossus root(field 1), the other me-dially to the rostral part of the hypoglossus root (field 2).

(field 2). Results were as follows:1)all SID-infants demonstrated the EAN. 2) number of neurons per area decreased with infants growth; by_20 weeks we found 1.390x10'/mm , by birth 73x10'/mm _in SID-infants and in controls 50-30x 10'/mm _ 3) cell distribution was irregular, es-pecially in field 2. (Supported by SWISS NATIONAL SCIENCE FOUNDATION grant 3.997.82)

56 EFFECT OF BANKED HUMAN BREAST MILK(BHEM)ON THE POST-NATAL EVOLUTION OF MIDARM CIRCUMFERENCE (MAC), TRICIPITAL(T), AND SUBSCAPULAR(S) SKIN-FOLD (SKF)THICKNESS, ARM MUSCLE(AMA)AND FAT AREAS(AFA)IN

FOLD (SKF)THICKNESS, ARM MUSCLE(AMA)AND FAT AREAS(AFA)IN APPROPRIATE(AGA)AND SMALL FOR GESTATIONAL AGE(SGA)VERY LOW BIRTH-WEIGHT INFANTS(VLBW). J.L.Excler; L.Sann;Y.Las-ney J.Picard; Höpital Debrousse, Lyon, France. We studied 28 newborn infants : 20 AGA(mean[±]SD)GA:30[±] 1,47weeks(range 28-32) BW: 1301[±]256g(890-1890) and 85GA VLBW:CA=34[±]2,4 (29-37), BW=1137[±]185g (890-1890) and 85GA ULBW:CA=34[±]2,4 (29-37), BW=1137[±]185g (890-1890) and 85GA ULBW:CA=34[±]2,4 (29-37), BW=1137[±]185g (890-1360). We fol-lowed them during 3 to 10 weeks, measuring each week MAC T and SSKF at 60 seconds, AFA and AMA. They were fed BH-EM at 40 cc/Kg/d at day 1 to reach 200 cc/Kg/d between day 15 and day 21. NaCl(2mg/Kg/d)was added for 3weeks when BW:1300g.

day 15 and day 21. NaCl(2mg/kg/a) was dured for smears when BW(1300g, Compared with the fetus values previously described, at 33,36,and40weeks of fetal age(FA)in AGA,MAC were res-pectively 7,2±0,7, 7,63±1 and 8±1,41cm, vs(mean)7,8,8,85 and 10,3cm in utero(lower limit of prediction 8,9cm at 40w);in SGA:6,26±0,64, 7,2±0,6, 7,7±0,3cm, vs 6,2, 7,1, and 8,15cm in utero. The same evolution was found with T and 8,15cm in utero. The same evolution was found with T and SSKF.In AGA,MAM were respectively 333,7595,5,351,31 \pm 80,6,366,8 \pm 118,4, vs 375,485 and 647,8mm²in utero; in SGA, 263,4 \pm 68,7, 339,5 \pm 60,2 and 365,8 \pm 49, vs 255,330 and 425mm²in utero.In AGA,AFA were 86,3 \pm 19,9, 116 \pm 37,9 and 150,5 \pm 61,6,vs 105, 145and 200mm²in utero; in SGA, AFA were 51,34 \pm 14,6, 75 \pm 18,9 and 106,8 \pm 7,34, vs 50,75and 105 mm²in utero. After 36weeks of FA,AGA are below the fetus values of AGA for MAC,SKF,AFA and AMA, whereas SGA,VLBW follow the fetus values of SGA. These results suggest that when VLBW are fed exclusively BHEM, 4 weeks after birth, AFA and AMA are below the fetus values in utero.