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BONE MINERAL CONTENT AND LINEAR GROWTH IN INFANTS TREATED WITH DEIAMETHASONE FOR BRONCHOPULMONARY DYSPLASIA Gopi Menon<sup>1</sup>, Jeremy R Williams<sup>2</sup>, Breda McLoughlin<sup>1</sup>, Fauxia Davidson<sup>2</sup> and Neil McIntosh<sup>1</sup>. Dept. of Child Life and Health<sup>1</sup> and Hedical Physics<sup>2</sup>, University of Edinburgh, U.K.

Arterial & venous cerebral blood flow velocity (CBFV) in newborn infants during continuous positive airway pressure breathing (CPAP) & following hypoxic-ischemic encephalopathy(HIE) C. Pacioni, F. Allemand, P.Tusim-Cottafavi, S. Arachi, R. Lucchon, G.I. Sallustio, C. Tozzi, M. Orzalesi, Institute of Child Health, University "La Sapienza", Rome, Italy

Ortalexi, Institute of Child Health, University "La Sapienta", Rome, Italy Mean arterial and venous CBFV (em/sec) were measured simultaneously by transcranad Doppler Velorimetry of the America Cerebral Artery and Internal Cerebral Vern in 3 groups of newborn infants of various Gestational Ages(GA,w), Birth Weights(BW,g) and Post-Natal Ages(PNA,d). Group 1, 53 normal infants, GA 26-41w, BW 950-3890 g, PNA 1-60kl. Group 2, 27 infants with RDS incated with different levels of CPAP, GA 26-34w, BW 830-3130g, PNA 1-30kl. Group 3, 18 infants with HIE, GA 32-42w, BW 1180-3750g, PNA 1-30k diagnosis of HIE was bosed on the following criteria need for vigorous resiscination at birth; severe acidosis (BD 12 and pH 7.20) within one from the mean contrast and pH 7.20. of age; clinical evidence of Grade 2 or 3 HE by Sarnat classification (2)by infants of Group 1, mean arterial and venous CBFV were GA and PNA dependent, in they increased significantly (psd 03) with anction and Chiny Chry Witter (A) and 1988 at the process CHP V (but not arterial CHPV) decreased significantly (p.0.03) with mercasing CPAP by approximately 1 timsec per cond. D. CPAP increment in infants of <u>Group 3</u>, mean arterial and venous CHPV were considered "abnormal" when higher or lower than 2.80 from the average normal values obtained in infants of Group 1. By these criteria, 6 infants with both "abnormal" arterial and venous CHPV all had a severe outcome, i.e. 4 ded and 2 had severe neurological abnormalines at follow-up; none of the 12 infants with both "normal" (2) infants with both "abnormal" arterial and venous CHPV all had a severe neurological abnormalines at follow-up; none of the 12 infants with both "normal" (2) infants with both "normal" (2). or just one "abnormal" (5) CBFV either died or became severely handicapped. The present study provides normal reference values of arterial and venous CBFV in the newborn; it suggests that monitoring of mean venous CBFV in infants with RDS could provide useful information on the possible interference of excessive CPAP on the cerebral circulation; finally, it suggests that the simultaneous evaluation of both arterial and venous CBFV is a better indicator of severe outcome prinfants with IHE than the measurement of arterial CBFV alone (\*)Sarnat H B and Samat N S Arch Neurol 1976:33 696-705

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RETINAL AND CHOROIDAL BLOOD FLOW DURING HYPOXEMIA AND

RETINAL AND CHOROIDAL BLOOD FLOW DURING HYPOXEMIA AND REOXYGENATION WITH 21% AND 100% O, IN THE NEWBORN PIGLET. J-P.Odden, T.Rootwelt, C.Hall, D.Bratlid. Depts. Fed. Res. and Surg.Res., Rikshospitalet, University of Oslo, Norway. The effects of hypoxemia (HO, 8% O,) followed by resuscitation with 21% or 100% O, on retinal (RBF) and choroidal blood flow (ChBF) were studied in 19 newborn piglets with the microsphere method. When base excess reached - 20 mmol/L or systolic blood pressure fell below 30 mmHg, the piglets were randomly resuscitated with either 21% O, (n=10) or 100% O, (n=9) for 25 min followed by 21% O, in both groups. RBF and ChBF were measured at baseline (B1), at the end of HO, and 5 min, 20 min and 60 min of reoxygenation. Flow values were (ml/min/100g, mean:SD):

BL HO 5 min 20 min 60 min 60 min RBF 21% 38±11 34±23 150±73\* 89±20\* 56±23 100% 44±18 43±26 189±83\* 81±40\* 56±19

RBF 21% 38±11 34±23 150±73\* 89±28\* 56±23 100% 44±18 43±26 189±83\* 81±40\* 56±19 ChBF 21% 2272±1000 1046±723\* 3584±2272# 2304±1059 1724±970\* 100% 1914±823 1110±698\* 3287±1571\* 2107±966 1622±743 \*p< 0.05 from Bl, #p= 0.06 from Bl. HO significantly decreased ChDF, but not RBF. RBF significantly increased during 5 and 20 min of reoxygenation, while in ChBF this increase was only seen at 5 min of

reoxgenation.

Conclusion: We could not demonstrate any significant differences in RBF or ChBF between two groups of hypoxemic newborn piglets reoxygenated with 21% or 100% O<sub>2</sub>.

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#### BRAIN LACTATE INCREASED WITH IMMATURITY OF HEALTHY NEONATES

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Glucose is the predominant cerebral fuel energy under physiologic conditions although other substrates may support the cerebral metabolism. The present study was undertaken to determine 1) whether lactate is present in the immature human brain, and if so 2) whether concentrations of lactate differ between infants who are light-for-gestational-age (LGA) or appropriate-for-gestational-age (AGA). Twenty-one stable and healthy infants with normal brains were investigated. All received milk enterally, and they had normal blood glucose levels. The gestational age averaged 36 completed weeks (range 28-41), and mean birth weight was 2350 g (range 855-4100). Proton NMR spectra from corpus striatum were obtained while the infants were sleeping quietly. Lactate was present in 8 preterm LGA and 3 preterm AGA infants, and the concentration was inversely related to the postmenstrual age (P=0.0043). Thus, lactate could not be demonstrated in infants with a post-menstrual age above 40 weeks. The lactate concentration was identical among LGA and AGA infants (P=0.15). Apparently, the immature brain consumes lactate, and the preferential substrate utilisation changes gradually with increasing age.

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Morphine and Stress Hormone Levels in Ventilated Newborns Quinn MW, Wild J, Dean HG, Hartley R, Puntis J, Levene MI Academic Unit of Paediatrics, Leeds General Infirmary, UK.

In a previously reported uncontrolled study we have shown that morphine reduces stress hormone levels in ventilated preterm babies. We now report a randomised double blind placebo controlled trial

41 preterm babies who qualified for Curosurf (A/a ratio  $\leq$  0.22) were enrolled in the study. 20 received 5% dextrose infusion (group P) whilst 21 received morphine (100 ugs/kg/hr x 2 hrs then 25 ugs/kg/hr infusion) in 5% dextrose (group M). Plasma catecholamine levels were measured on entry and after 24 hours. Blood pressure was documented on entry and after 6 hours. The two groups showed no differences with respect to method of delivery, Apgar scores, birth weight, gestation and catecholamine levels on enrolment. Group M showed a small but significant (P=0.01) reduction in adrenaline levels ([median (range) change -0.4 nmols/L (-34.7 to +0.06)], compared with group P [median (range) change +0.2 (-13.5 to +19.0)]. There were no significant changes in noradrenaline levels. Group M treated babies showed a slight reduction (median: 4 mmHg) in blood pressure. There were no differences between the groups for the incidence of intraventricular haemorrhage, patent ductus arteriosus, pneumothorax, the number of ventilator days and death.

We conclude that morphine reduces adrenaline levels in ventilated preterm babies and appears to have no significant adverse effects

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SCHOOL PERFORMANCE IN VERY PRETERM CHILDREN A report from the Collaborative Project On Preterm and Small for gestational age (POPS) infants in the Netherlands

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A virtually complete birthcohort of infants <32 weeks and/or <1500 g (n=1338) was followedup until 9 years of age. In 88% of survivors (n=813) data on schoolperformance were available.

	totai	mainstream	remedial teaching	retention	education	
all	813	324 (40)	120 (15)	213 (26)	155 (19)	-
non-handicapped	708	319 (61)	115 (16)	192 (27)	71 (10)	

Of the children in mainstream education 38% had remedial teaching (27% at appropriate level, 60% below level). The need for remedial teaching was not different for children who were not handicapped at 5 years of ago. Logistic regression for non-handicapped children with any schoolfailure as dependent factor showed significantly higher OR's for SGA (1.8) boy's (1.9) low or middle SES (5.4 and 2.1) mild or severe developmental delay (5 years (3.6) When special education was the dependent factor higher OR's were found for boy's (2.0) twins (1.9) low and middle SES (7.1 and 3.0) solvere montal delay (10.1) mild speech delay (3.3) high inattention and hyporactivityscore (2.9) and low schoolresults at 5 years (2.8). Perinatal factors (gestational age, birthweight and neonatal illness) were not significant.