DISTRIBUTION OF ACUTE INTRACRANIAL LESIONS IN SMALL FOR GESTATIONAL AGE (SGA) INFANTS DURING NEONATAL 1357 1357 FOR GESTATIONAL AGE (SGA) INFARTS DURING NEOMATAL PERIOD. Jean Cacciabaudo, Mehmet Y. Dincsoy, Young <u>M. Kim, Syamala K. Naroji, Susan Tuck.</u> (Spon. by Norman L. Gootman). Health Sciences Center, SUNY at Stony Brook, Nassau County Medical Center, Dept. of Pediatrics, East Meadow, NY.

SGA infants are prone to subacute fetal distress which may produce growth retardation and wt loss. They are also subject to acute distress which may lead to prepartum and intrapartum cardiovascular instability, thus, to an acute intracranial bleed. Since periventricular (PVH), intraventricular (IVH) and parenchymal hemorrhage (PH) and ventricular dilatation (VD) can be detected by cranial ultrasound (US) examination, we studied 65 low birth weight (LBW) SGA infants and compared them with 65 weight matched controls. A separate gestation matched con-trol study is in progress. Following table depicts a compari-son (Number, X<sub>1</sub>'SEM) between the SGA and the weight matched con-trol groups which is based on first US:

trol groups which is based on first US: <u>n Gest(wk) Birth wt Apgar 5 PVH,IVH,PH VD Mortality</u> <u>SGA 65 35.240.4<sub>\*\*</sub> 1488 6.4±0.3 10(15%) <sub>\*\*</sub> 5(6%) 2(3%) Control 65 31.4±0.3 1503 6.6±0.3 3(4.6%) 11(17%) 6(9%) <sup>\*\*</sup>p<0.001, \*<0.025, (t test or  $X^2$  where appropriate) Excess number of VD in control group and excess of PVH,IVH and PH in SGA group persisted in the 2nd and 3rd US (p<0.05). There seems to be an increased incidence of PVH, IVH, PH in SGA in-</u> fants despite their more mature status as compared to the control infants who were gestationally less mature. Lesser inci-dence of VD in SGA infants may be related to an accelerated maturity in these infants.

**TI358** PRELIMINARY RESULTS OF A RANDOMIZED TRIAL OF HIGH-FREQUENCY JET VENTILATION(HFJV) VS CONVENTIONAL VENTILATION(CV) IN SEVERE RESPIRATORY DISTRESS SYM-DROME. WA Carlo, RL Chatburn, RJ Martin. CWRU, Dept. Peds, Cleve, OH Despite potential reduction of barotrauma, no controlled data are available on the incidence of adverse effects of long-term high frequency ventilation(HFV). To compare HFJV to pressure-

high frequency ventilation(HFV). To compare HFJV to pressure-limited time-cycled CV we screened 208 1-2 kg neonates and ran-domized those fulfilling all of 6 criteria by 24 hr: 1)Fi0\_2>.50; 2)peak insp. pressure(PIP)  $\geq$ 20cmH<sub>2</sub>0; 3)positive end exp. pres-sure(PEEP)  $\geq$ 4cmH<sub>2</sub>0; 4)frequency(f)  $\geq$ 20/min;5)PaO<sub>2</sub>  $\leq$ 90mmHg; 6) PaCO<sub>2</sub>  $\geq$ 35mmHg. We randomized 18 neonates to HFJV (BW 1.5±.3kg, CA 30±2 wks) at a f=250/min and I:E=1:3, or CV (BW 1.5±.3kg, CA 30±2 wks) at 2±6 hr and managed both groups with standardized ventilatory protocols for 48 hr. PIP, mean airway pressure(Paw) and blood gases(ABG) were averaged over 48 hr. Sequential anal-yses of development of air leaks(AL), intraventricular hemor-rhage(TWH) or death during the 48 hr.and chronic lung disease rhage(IVH) or death during the 48 hr, and chronic lung disease Subsequent assisted ventilation and O<sub>2</sub> suppl. were comparable. Bronchoscopies(n=5) revealed no evidence of necrotizing tracheo-bronchitis. We conclude that 48 hr of HFJV: 1) maintained or improved ABG at lower  $\overline{P}aw$ ; 2) did not eliminate AL or CLD; 3) caused no apparent increase in adverse effects. This first randomized HFV trial indicates a comparable outcome to CV; larger

trials will ultimately reveal if HFV is superior. ALA Ohio,ALANO

BILTRUBIN (BR) FLUX AND DISTRIBUTION IN THE BRAIN. + 1359 William J. Cashore and Philip Blazar. Brown Univ., Women & Infants Hosp., Dept. of Peds., Prov., R.I.

Carotid artery injection of Evans Blue or albumin-bound bili-rubin (ALB+BR) for 60 sec. in adult rats produced no cortical staining with an intact blood-brain barrier (BBB) and non-uniform "tufted" perivascular staining after barrier opening to the "tufted" perivascular staining after barrier opening to ALB by 1.8 Molar Urea x 30 sec. Unbound BR given for 10-60 sec. to 25 rats without barrier opening, followed by carotid perfusion with saline, produced uniform staining, and the uptake of BR measured by chloroform extraction was dose-dependent (r=0.641, p<0.001). Retained BR was 11.6±6.2% (MEAN±S.D.) of the dose injected, and the permeability-surface area (P.S) product for BR was  $0.13\pm0.05$  ml min<sup>-1</sup> g<sup>-1</sup>. The estimated volume of distribution for unbound BR at 60 sec. was  $195\pm72 \ \mu$ l/g. When injection of unbound BR was followed by re-infusion with 5% ALB, residual brain BR decreased as shown:

Group	N	Brain BR.	% BR	p-value
RP + Saline	11	μg/gm 7.3+2.8	Retained 9.7±3.1	(unpaired t-test)
DK · Darine		/.52210	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	<0.01

1.9±1.4 1.8±1.3 11 BR + ALB BR + ALB 11 1.8±1.3 1.9±1.4 Bound BR crossing a disrupted BBB and unbound BR crossing an in-tact barrier appear to have different extravascular patterns of distribution, consistent with the greater lipid solubility and capillary permeability of unbound BR, versus the tendency of ALB + BR to cross the barrier only at osmotically opened capillaries. Because ALB decreases residual brain BR, we speculate that BR is initially distributed in a "shallow" compartment and is not imme-diately strongly bound or irreversibly precipitated in the brain.

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1360 THE APGAR SCORE REVISITED: DEVELOPMENTAL PATTERNS IN "OPTIMAL" NEWBORNS. Elizabeth A. Catlin, Marshall W. Carpenter, Benjamin S. Brann IV, Steven R. Mayfield, Philip W. Shaul, Marshall Goldstein, William Oh, Brown Univ, Women & Infants Hosp, Dept of Ped, Providence, RI The five objective criteria used in the Apgar score to assess the newborn include four signs (tone, reflex irritability, color, respiratory drive) that depend upon developmental maturity. The immature infant, with typically flaccid muscle tone is penalized by the Apgar method for developmental immaturity. We hypothe-sized that in nonasphysiated fetuses, Apgar scores should corresized that in nonasphyxlated fetuses, Apgar scores should corre-late directly with gestational age (lower scores with decreasing naturity). 62 parturients (gestational age (lower scores with decided and maturity). 62 parturients (gestation: 23-42 weeks) were prospec-tively enrolled. By objective documentation of pregnancy his-tory, labor complications, fetal heart rate tracings, cord blood pH & BE, their fetuses were judged as normal at the time of deli-very. Direct relationship between Apgar scores and gestational age following a linear regression (y=.34x-4.8 at 1', r=0.82; y=.28x-1.6 at 5', r=0.77) was observed. Analysis of the total score into components reveals heart rate to be least influenced by maturity. Respiratory effort and muscle tone increase with increasing maturity. Skin color improves with development at the S' score. Reflex irritability trends upward but with more varia-bility at 1' than 5'. These data suggest that 1) the standard Agar score may not be appropriate for the assessment of fetal well-being in premature infants, 2) scores thought representative of asphyxia in more mature newborns may not represent asphyxia in immature infants, and 3) these normative data are useful in the assessment of fetal status at various levels of maturity.

COMPARISON OF A STANDARD AND MICROVOLUME BLOOD AND 1361 MEDIA SYSTEM ON THE RATE OF DETECTION OF BACTEREMIA IN PREMATURE INFANTS. <u>Subhash C. Chaudhary</u>, J. Strano, (Sponsored by Robert E. Merrill), SIU School of Medicine, Department of Pediatrics and St. John's Hospital, Department of Microbiology.

A microvolume blood and media system was compared to a standard BACTEC anaerobic vial for its efficacy to detect microorganisms in the small volume of blood obtainable from premature infants. Each culture, when possible, included the inoculation of 0.5 ml. of blood into a BACTEC anaerobic vial (BAN) and 0.3 ml. of blood into each of an aerobic (MA) and an anaerobic (MAN) vial containing 2.7 ml. of trypticase soy broth. Following inoculation, all vials were processed identically. Of a total Due to the occasional low volume of blood collected in some of bue to the occasional low volume of blood collected in some of the 39 positive cultures, 8 BAN and 2 MAN vials were not inoculated. Twenty cultures were positive for the same organism in all three vials. When both anaerobic cultures were done, 26 BAN and 23 MAN were positive. In the microvolume system, 31 MA and 28 MAN were positive and both were positive in 23. Differences were not statistically significant. Also, four iso-lates of <u>Candida albicans</u> grew in MA and MAN vials only. The results demonstrate that microvolume blood and media systems These can be reliable.

EFFECT OF CAFFEINE, THEOPHYLLINE, AND PHENOBARBITAL 1362 ON OCULAR BLOOD FLOW IN THE NEWBORN FIGLET. S. Chem-tob, K. Beharry, N. Laudignon, J. Rex, J.V. Aranda. Depts. of Peds. & Pharmaco.; McGill Univ - Montreal Child Hosp; Montreal, Canada.

The effect of commonly used drugs in the newborn (NB) period The effect of commonly used drugs in the newborn (NB) period on ocular blood flow (OBF) was evaluated in 6 groups of piglets (1-4 do). Caffeine (C) and theophylline (T) were administered at 5 and 20 mg/kg IV bolus (N= 7/grp). Phenobarbital (P) (15 and 45 mg/kg IV bolus) were administered to 2 groups of piglets (N=9/grp). Control groups were obtained for each drug evaluated. OBF was measured by radiolabelled microsphere (1<sup>41</sup>Ce <sup>46</sup>Sc <sup>8</sup> Sr <sup>9</sup> Nb); arterial and venous blood gases, hemoglobin, and drug levels were done at times 0, 15, 30, and 60 minutes.

Drug	Dose	0'	15'	30'	60'	
	5 mg/kg	96.8±62.0	59.4±19.2	83.5 ± 100.9	126.4 ± 76.3	
20	0 mg/kg	67.2±46.6	53.1 ± 21.2	54.2 ± 12.7	62.9±37.5	
	5 mg/kg	$49.9 \pm 25.4$	47.4 ± 9.6	66.9± 14.7	55.1 ± 22.1	
1 20	0 mg/kg	45.0±18.2	46.6±11.8	36.9± 9.1	$51.7 \pm 16.1$	
	5 mg/kg	$48.2 \pm 12.8$	*40.1± 9.3	43.6± 15.8	49.4±14.6	
P 20	0 mg/kg	47.4 ± 22.3	$39.0 \pm 14.4$	*35.5 ± 16.8	$45.5 \pm 16.1$	
Values are OBF $\overline{x} \pm S.D.$ (m1/min/100g); *p <0.01 compared to t=0						
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Therapeutic doses of C produced fluctuations in OBF, an effect not observed in hypercaffeinema. Similarly both doses of T pro-duced fluctuations in OBF. P decreased OBF transiently. A con-trol group of animals did not show any changes in OBF. Blood pressure and blood gases remained constant, with no correlation to OBF. These suggest that C, T, and P probably exert direct pharmacologic effects on OBF. Their possible role on retinopathy of prematurity needs further studies.