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EVALUATION OF A NEW INDIRECT BLOOD PRESSURE SYSTEM FOR NEONATES. Leonard L. Fox, Dale L. Phelps, (Spon. by Cynthia T. Barrett), UCLA School of Medicine, Dept. of Pediatrics, Los Angeles, California.

Infrasonde<sup>R</sup> (Sphyngometrics), a small, lightweight (6.2 lbs), battery-powered, indirect blood pressure system was tested under a variety of clinical conditions in neonates. This system consists of a passive transducer (1x1 cm) placed under a disposable cuff which electronically senses arterial wall oscillations on a narrow subaudible (infrasonic) frequency band thereby improving accuracy by increasing the signal to noise ratio.

Ten neonates (1360 to 4500 gms), catheterized for medical reasons (RDS, shock, sepsis), were studied. Approximately 20 readings each by experienced and inexperienced personnel were made on the right arm, left arm, and one leg, simultaneously recording the direct transducer pressure from an arterial catheter attached to a KDC monitor at the exact moment that the Infrasonde reading was taken. Comparing absolute differences between the Infrasonde and the direct arterial pressure over a wide range (diastolic 18 to 60 mm Hg, systolic 38 to 105 mm Hg), we were unable to detect any difference between experienced and inexperienced personnel, right and left arms, and arms and legs; therefore all the readings were combined. The correlation coefficient between direct arterial and Infrasonde pressures was  $r = .900$ ,  $n = 113$  diastolic and  $r = .940$ ,  $n = 113$  systolic.

These data indicate that Infrasonde correlates well with direct arterial pressures and could be especially useful when there is no indwelling catheter present.

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THYMUS SIZE AND ITS RELATIONSHIP TO RESPIRATORY DISTRESS SYNDROME. I.H. Gewolb, R.L. Lebowitz, and H.W. Tausch, Jr., Dept. of Pediatrics and Radiology, Harvard Medical School, Boston, MA.

In an attempt to correlate the thymolytic effect of glucocorticoids with their role in the induction of surfactant synthesis and the prevention of RDS, the size of the thymus on the initial chest radiographs of newborns who required intensive care was reviewed. Utilizing a ratio of the cardiomyic silhouette at the carina to the thoracic width (C-T/T) we found an increase in C-T/T with increasing gestational age. Newborn infants with RDS had a significantly elevated mean C-T/T (0.40 vs 0.36;  $p < 0.001$ ) an increased percentage of 'enlarged' (C-T/T > 0.45) thymuses (28% vs 8% in non-RDS babies), and a decreased percentage of 'small' (C-T/T < 0.30) thymuses (11% vs. 26%). Moreover, in a series of 8 multiple gestations that were discordant for RDS, the larger thymus was seen in those with RDS in 6 of 8, while in only one did the converse hold true. Surprisingly, babies whose mothers had received prepartum dexamethasone to prevent RDS showed no difference in thymic size when compared to a control population. These data are consistent with the theory that endogenous glucocorticoids play a physiologic role in the prevention of RDS. The initial size of the thymus may also be of predictive value in differentiating between the diverse causes of respiratory distress in newborn infants.

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EFFECTS OF INTRALIPID INFUSIONS ON ARTERIAL BLOOD GASES AND PULMONARY FUNCTION IN SMALL PREMATURE NEONATES. William W. Fox, Gilberto R. Pereira, Jacob G. Schwartz. (Spon. by Jean A. Cortner), Univ. of Pa. Sch. of Med., Dept. of Peds. and The Children's Hosp. of Phila., Phila., PA.

Intralipid is widely used for parenteral nutrition of infants with respiratory distress but hypoxemia has been reported in adults receiving IV intralipid infusions. To determine changes in pulmonary function during a 4 hr. intralipid infusion, 1 gm/kg intralipid was administered to 7 small premature neonates with mean wt. 1.16 kg (range 0.88-1.50) and mean gest. age 29 weeks. Arterial blood gases, func. resid. cap. (FRC), tidal volume (TV), resp. rate (RR), lung compliance (CL) and insp. (R<sub>I</sub>) and exp. (R<sub>E</sub>) pulmonary resistance, serum triglycerides (TG) and free fatty acids (FFA) were determined at control (pre-infusion), at 4 hours and at followup (4 hours after infusion). Control (mean ± SEM) FFA (0.58 ± .17) and TG (54.1 ± 6.3) increased at 4 hours to 2.0 ± 0.37 and 230.0 ± 66.2. Control P<sub>O</sub>2 (66.4 ± 11.1) decreased post infusion to 54.1 ± 2.1 mm Hg. At followup, both mean FFA and TG were higher than control values and less than post infusion. Mean ± SEM followup P<sub>O</sub>2 (62.6 ± 6.4) was higher than post infusion but lower than controls. There were no significant differences in pCO<sub>2</sub>, pH, base excess, RR, VT, min. vent., CL, R<sub>I</sub>, R<sub>E</sub>, or FRC for all 3 parts of the study. This study indicates that intralipid infusion in premature neonates affects P<sub>O</sub>2 but does not cause quantitative changes in other blood gas parameters or pulmonary function.

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FETAL PROLACTIN (PRL) AND THE RESPIRATORY DISTRESS SYNDROME (RDS). P.D. Gluckman, P.L. Ballard, J.A. Kitterman, S.L. Kaplan, and M.M. Grumbach. Dept. of

Pediat., Univ. of California San Francisco, San Francisco, Ca. There is a temporal relationship between plasma PRL, cortisol (F) and pulmonary surfactant production in the fetus. PRL administered to fetal rabbits is reported to increase pulmonary surfactant content. To explore these relationships we have measured PRL in umbilical cord sera from 78 infants of gestational age 28 to 40 weeks. A positive relationship to gestational age was demonstrated. Between 30 and 36 weeks gestation, analysis of covariance of PRL versus gestational age showed that infants who did not develop RDS had significantly higher cord PRL concentrations ( $p < 0.05$ ) than infants who developed RDS. Between 32 and 35.5 weeks the mean ± SEM cord PRL of normal infants (161.8 ± 18.9 ng/ml) is higher ( $p < 0.025$ ) than in those who developed RDS (101.7 ± 9.5 ng/ml). Growth hormone concentrations in the same samples did not show any relationship to the incidence of RDS. Cord PRL did not correlate with cord F or dehydroepiandrosterone sulphate (DHEAS) concentration. Serum PRL was not suppressed in a second group of 114 infants whose mothers were treated prenatally with betamethasone, but F and DHEAS were. The finding of low cord PRL in infants who develop RDS is consistent with the possibility that PRL may influence fetal lung maturation. To exclude the possibility that this potential association is due to a postulated adrenocorticotrophic effect of PRL, we infused ovine PRL (1 mg/day) into fetal sheep for periods of 6 to 8 days in late gestation. No effect on F, DHEAS, or F binding capacity was noted in the fetus.

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BICARBONATE (B) AND INTRACRANIAL HEMORRHAGE (ICH) IN HYALINE MEMBRANE DISEASE (HMD). Zvi Friedman, Curtis A. Cammarata, Keith H. Marks, M. Jeffrey Maisels.

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We studied the relationship between B administration and ICH in 53 infants with HMD. 27 had no ICH (Group I). 26 had ICH of whom 10 received B prior to the diagnosis of ICH (Group IIa) and 16 received B after ICH was diagnosed (Group IIb).

Groups	No. Infants	B meq/kg	Gest Age (wks)	B. Wt. (gms)	Apgar	
I	27	2.4 ± 1.4 <sup>a</sup>	32.4 ± 2.5 <sup>c</sup>	1860 ± 640 <sup>c</sup>	6 ± 2 <sup>c</sup>	7 ± 2 <sup>c</sup>
IIa	10	2.5 ± 1.6 <sup>b</sup>	29.6 ± 1.8	1320 ± 255	3 ± 1	4 ± 2
IIb	16	6.5 ± 2.2				

- a - Group I vs IIb  $p < 0.01$
- b - Group IIa vs IIb  $p < 0.01$
- c - Group I vs IIa + b  $p < 0.01$

There were no significant differences between the groups in the level of serum sodium, osmolality, inborn/outborn ratio and age of admission to NICU.

We conclude that a direct relationship exists between ICH, perinatal asphyxia and prematurity. No correlation was found between B administration and ICH.

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PNEUMOPERICARDIUM: AN APPROACH TO DIAGNOSIS AND TREATMENT IN THE NEONATE. Ronald N. Goldberg, Luis A. Cabal, Bijan Siassi and Joan E. Hodgman. Children's

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Pneumopericardium (PPC) developed in 16 of 873 (2%) neonates receiving assisted ventilation; 14 were prematures with RDS requiring assisted ventilation within 12 hours. Pneumothorax (PNX) developed coincidentally in all but one. Pericardiocentesis (PCX) was performed in 11 of whom 7 (42%) survived. Cardiovascular parameters of heart rate (HR), arterial blood pressure (ABP), central venous pressure (CVP) and ECG were measured during 10 episodes of PPC. Early evidence of cardiac tamponade (CT) was moderate tachycardia and a sharp rise in CVP. This was followed by bradycardia and an abrupt decrease in pulse and ABP. Marked ST-segment depression with AV dissociation and 2° heart block were noted at this time. Following PCX, a rebound tachycardia, widening of pulse pressure, return of ABP to baseline and a gradual fall in CVP occurred in survivors while in those who died, bradycardia and hypotension persisted. PPC as a complication of assisted ventilation should be expected in a small number of infants with RDS. If the infant develops a pneumothorax, this risk is significantly increased. The diagnosis can be made before death by cardiovascular changes described. The condition can be successfully treated and adequacy of treatment can be assessed by the same factors used for diagnosis.