1177 THE RELATION OF FLUORESCENCE POLARIZATION VALUES TO LECITHIN SPHINGOMYELIN RATIOS (L/S) IN AMNI-OTIC FLUID (AF). Thomas A. Blumenfeld, Raymond I.Stark, Inge Dyrenfurth, Vincent J. Freda, L. Stanley James, Div. of Perinatal Med., Coll. of P & S, Columbia Univ., N.Y.

Fluorescence polarization (FP) measures the microviscosity of phospholipid micelles. The technique has been evaluated in both artificial solutions and in AF and its usefulness in determining gestational age and lung maturity examined.

We have shown that in lipid dispersions of known composition and in amniotic fluid the FP value correlates directly with the L/S ratio, Gestational age was studied in 97 serial samples from 19 isoimmunized pregnancies. There was a significant change in slope of FP values vs. gestational age occurring between 29 to 35 weeks (mean = 31.5 wks). In single samples from 120 pregnancies a significant correlation was also found

between FP and L/S.	Mature FP (く352)	Immature FP (>352)
Mature L∕S ( > 2)	88	0
immature L/S (< 2)	9 (7.5%)	23

Thus FP values correlate well with both gestational age and lung maturity. The determination of fluorescent polarization is precise with a coefficient variation of 0.48%. It can be performed rapidly (45 min) and is technically easy. This physical method therefore offers a great advantage over the currently available chemical methods. (Supported by NHLI HL-14218)

**1178** A COMPARISON OF MORTALITY AND MORBIDITY ASSOCIATED WITH PRESSURE-LIMITED AND VOLUME-LIMITED INFANT VEN-TILATORS IN THE TREATMENT OF SEVERE HYALINE MEMBRANE DISEASE - Stephen J. Boros, Children's Hospital, St. Paul, and Department of Pediatrics, University of Minnesota, Minneapolis, MN (Spon. by Edward L. Kaplan) The effect of pressure-limited (PL) and volume-limited (VL) infant ventilators on mortality and morbidity in severe hyaline membrane disease was examined in a prospective controlled study. Criteria for mechanical ventilation were CPAP failure (PaO<sub>2</sub> <50 mm. Hg. at FiO<sub>2</sub> >0.8 and CPAP >8 cm. H<sub>2</sub>O or PaCO<sub>2</sub> >70 mm. Hg.) or severe perinatal asphysia. Consecutive patients were alternately assigned to either PL or VL infant ventilators. 22 infants (900-2600 gms; 27-36 wks. gest.) were ventilated with PL machines assigned to either PL or VL infant ventilators. 22 infants (900-2600 gms; 27-36 wks. gest.) were ventilated with PL machines using low peak inspiratory pressures (mean Pmax = 29 cm. H<sub>2</sub>O), relatively low PEEP (mean = 5.4 cm. H<sub>2</sub>O) and prolonged inspira-tory times. 20 additional infants (600-3350 gms; 26-42 wks. gest.) were ventilated with VL machines using essentially unlim-ited peak inspiratory pressures (mean Pmax = 62 cm. H<sub>2</sub>O), PEEP (mean = 8.1 cm. H<sub>2</sub>O) and prolonged expiratory times. There were no significant differences in mortality (PL=37%; VL=40%), or in the incidence of pneumothoraces (PL=32%; VL=45%), or bronchopul-monary dysplasia (PL=4.6%; VL=10%). Intraventricular hemorrhages and pulmonary hemorrhages occurred with equal frequency in both groups. These data fail to demonstrate any significant advantage of one ventilator system over the other, and fail to confirm the impression that limiting peak inspiratory pressures to <35 cm. H<sub>2</sub>O significantly reduces the incidence of bronchopulmonary dysplasia.

BACTERIOLOGY OF ASPIRATION PNEUMONIA IN CHILDREN. I. 1179 Brook and S.M. Finegold. Fairview State Hospital, Costa Mesa; Wadsworth V.A. Hospital; and UCLA Medical Center, Los Angeles, Ca. (Spon. by J.D. Cherry) 1179 The bacteriology of aspiration pneumonia in institutionalized children and adults was determined by transtracheal aspiration.

Forty-three patients was determined by transtracted applration. Forty-three patients were studied, including 10 adults and 33 children ranging from 2 months to 18 years in age (average 8.5 years). Ignoring <u>Staphylococcus epidermidis</u>, the 43 aspirates yielded 85 aerobic and 86 anaerobic isolates, demonstrating the polybacterial etiology of the infection. Anaerobic bacteria were isolated from 30 children (91%) and from 9 of 10 adults. The predominant anaerobes were <u>Bacteroides melaninogenicus</u> (18 iso-lates), Fusobacterium sp. (10), and anaerobic and microaerophilic gram-positive cocci (21); there were 8 isolates of <u>B</u>. <u>fragilis</u>. These organisms were proportionately distributed between children and adults except for relatively more <u>B</u>. <u>melaninogenicus</u> in children and more <u>B</u>. <u>fragilis</u> in adults. Pneumococci (10 iso-lates), group A streptococci (8) and Haemophilus influenzae (2) were recovered only from children and among 11 Escherichia coli isolates, 10 were from children (mostly under 6 years of age) The Klebsiella-Enterobacter-Serratia group, Proteus, Pseudomonas and alpha-hemolytic streptococci were proportionately more common in adults than in children. All subjects recovered with appropriate therapy. Anaerobic bacteria clearly play a role in aspiration pneumonia in children as well as in adults. The differences in aerobic bacteriology cited require different therapeutic approaches.

1180 CARDIOVASCULAR EFFECTS OF PROLONGED INSPIRATORY TIMES WITH CONTINUOUS POSITIVE PRESSURE VENTILATION (CPPV). David L. Brown, Philip J. Lipsitz, Joy S. Hagelberg, Norman Gootman, SUNY at Stony Brook, Health Sciences Center, Long Island Jewish-Hillside Medical Center, Division of Neonatology, New Hyde Park. New York

Norman Gootman, SUNY at Stony Brook, Health Sciences Center, Long Island Jewish-Hillside Medical Center, Division of Neonatology, New Hyde Park, New York. The use of prolonged inspiratory time with CPPV has been re-commended for the management of infants with the Respiratory Distress Syndrome. The cardiovascular effects of such inspira-tory to expiratory (I:E) ratios were investigated in 5 piglets, (age 6 to 10 days), under 0.25% halothane anesthesia and decame-thonium paralysis. Heart rate (HR), aortic blood pressure (AoP), renal (RF), femoral (FF), and carotid (CF) blood flows were mea-sured at three I:E ratios (1:2, 2:1, and 3:1), at each of three levels of positive end expiratory pressure (PEEP), (2,5 and 10 cm H<sub>2</sub>O). The cardiovascular parameters at the 2:1 and 3:1 ratios were compared to those of the control of 1:2 while PEEP, peak pressure, respiratory rate, and ventilation remained constant. HR increased at both I:E ratios. Mean AoP exhibited variable changes at the 2:1 ratio, while at 3:1 there was a consistent de-crease in AoP with maximum decrease of 17.8% at 10 cm PEEP. At the 3:1 ratio both RF and FF decreased at 5 and 10 cm PEEP. At the 3:1 ratio with 5 and 10 cm PEEP there was a decrease in CF ranging from 4.5% to 22.0%. CF decreased at 3:1 with all levels of PEEP with a mean decrease of 11.3%  $\pm$  1.6 at 5 cm and 13.3%  $\pm$ 2.8 at 10 cm PEEP. These preliminary results indicate that ad-verse cardiovascular effects occur with the addition of prolonged inspiratory times to CPPV.

1181 ARTERIAL OXYGEN TENSION (PaO.) MEASUREMENTS-SAFETY AD ACCURACY OF A NEW CATHETER SENSOR. Edwin G. Brown, Chung C. Liu, Prancis E. McDonnell, Avron Y. Swet and Sidney K. WOlfson. Dept. of Pediatrics, Mt. Sinsi School of Med., New York, and Dept. of Surgery and Chemical geneering. University of Pittsburgh, Pittsburgh. The most reliable method requires intermittent sampling from an umbilical artery catheter or from a peripheral artery for analysis with a micro blood gas analyzer. Methods to continu-ouly measure PaO. using indwelling catheters have been developed but may of them receive limited use because of inaccuracy, tissue injury, clotting and interference with the function for which catheters are intended. Oxygen measuring method. We have devised a polarographic catheter electrode which is simply devised a polarographic catheter electrode which is simply devised a polarographic catheter sealibration at 6-12 hour intervals. Micro clot analysis, sortic angiography, scanning electron microscopy of sensors and vessels, and haptoglobin and red cell injury. The sensor responds rapidly and accurately ochanges in PaO. in newborn infants medults. It reveals changes in PaO. is an even infants and dults. It reveals changes in PaO. is an even infants and dults. It reveals changes in PaO. is an even infants and dults. It reveals content or respiratory equipment. A technique for continu-one and red cell injury. The sensor responder apidly and accurately changes in PaO. in mewborn infants and dults. It reveals

THE RELATIONSHIP BETWEEN THE CHEST RADIOGRAPH, REGION AL LUNG FUNCTION USING  $^{13}$ N, EXERCISE TOLERANCE AND CLINICAL CONDITION IN CHILDREN WITH CYSTIC FIBROSIS. 1182

A.L. Coates, P. Boyce, D.G. Shore, M. Mearns and S. Godfrey, (In-tr. by K.N. Drummond), Institute of Child Health, Hammersmith Hospital, University of London, England. We evaluated the role of the chest radiograph interpretation

in children with cystic fibrosis in delineating localised abnormalities of ventilation and perfusion as well as the overall se-verity in relation to maximal mid expiratory flow rate (MMEF), alveolar-arterial oxygen gradient, exercise tolerance (blcycle ergometer), and clinical severity. The upper and lower region of each chest radiograph was scored separately and compared with regional lung function tests (RLF) using the arrival and disappear-ance of boluses of radioactive nitrogen  $(^{13}N_2)$  given by inhala-tion and infusion. Whilst the worse areas on the chest radiograph were found to correlate with similar regions on the RLF, (r=0.58) errors occurred in half of the cases when the radiograph alone was used to judge regional physiological derangement. Despite this, the total radiographic score correlated well with the total regional function, r = 0.62, p: 0.01. Of the other parameters, the radiographic score correlated well with the MMEF, (r = 0.69, p = .01), maximum work expected, (r = 0.64, p = .01) and clinical grading, (r = 0.79, p = .001), but not with the (A-a) 0, gradient. From this, we have concluded that the chest radiograph, whilst not an accurate index of local physiological abnormalities, does a solute the meril event of the local denormalities. reflect the overall severity of the lung disease, exercise tolerance and clinical condition in cystic fibrosis.