CONJUGATED HYPERBILIRUBINEMIA (CH): A PREDICTOR OF PERINATAL MORTALITY & POOR NEUROLOGIC OUTCOME. S.S. Usmani, C.G. Sia, R.G.

Harper, E. Carrera, F. Daum. Cornell U. Med. Coll., North Shore U. Hosp. (NSUH), Depts. of Ped. & Ob/Gyn, Manhasset. Thirty-six of 1236 high-risk newborns transported to NSUH between 1/73 and 12/82 developed CH (direct

bilirubin >2.0 mg/dl or >50% of total); 36 transports matched for sex, race, gest. age + 2 wk, weight + 200 gm, birthdate + 1 yr, served as controls. All charts were reviewed to determine association between

CH and various factors.

	# Study Pts	# Controls	P-Value*
Perinatal Asphyxia	6	5	NS
Shock	14	2	<0.05
C.H.F.	7	2	NS
Hemolytic Disorders	15	1	<0.01
Multiple $T_X$ (>4 $T_X$ )	18	4	<0.02
Exchange T <sub>X</sub>	13	3	<0.05
T.P.N.	8	8	NS
Infections	13	5	NS
Mortality	19	3	<0.01
Poor Neuro. Outcome	9/17	1/33	<0.02

\*Chi square with Yates correction

These results indicate that CH is significantly associated with shock, hemolytic disorders, multiple Tx, exchange Tx, higher mortality and subsequent poor neurologic outcome. T.P.N. was not significantly associated with CH. The outlook for surviyal and normal development for the neonate with CH is poorer than has been previously recognized.

IN VIVO EFFECT OF FREE FATTY ACIDS (FFA) ON BILIRUBIN 

of Pediatrics.

In healthy adults a wide range (0.28 to 5.34mMo1/L) of plasma FFA levels were generated by the fat meal-heparin regimen (Schalch DS, Kipnis DM, J.Clin.Invest.44:2010,1965). In each experiment 4-5 blood samples were collected, albumin (Alb) and FFA concentrations were measured and BR binding assessed using: (a) BR titration of sera using Sephadex (S) Column Chromatography (Valaes T, Hyte M, Pediatrics 59:881,1977); (b) the Undiluted Peroxidase (P) method (Ahlfors CE, Clin.Chem.27:692,1981); and (c) the automated Bilirubin Hematofluorometer (HMF) (Wells R et al Clin.Chem.28:432,1982). With S a significant negative correlation was found between the BR/Alb (molar ratio) at the BR titration point ("BR binding capacity") and the FFA/Alb (Y=0.883 -0.088·X, r=-0.90, p(0.001, n=38). Similarly, after "spicing" the sera to BR/Alb of 0.367±0.029(SD) the Apparent Unbound BR (Mol/L) by the P method was significantly related to the FFA/Alb (Y=13.5+41.66.X, r=0.957, p<0.001, n=28). The BR/Alb at BR binding capacity by HMF was only weakly related to FFA/Alb (Y=0.711-0.009·X, r=-0.353, p<0.05, n=34). However, "spicing" of the blood samples to BR/Alb of 0.54±0.054(SD) improved the correlation (Y=0.029=0.001.X, r=-0.025.000.001.X).

tion (Y=0.929-0.041-X, r=-0.835, p<0.001, n=12). The reasons for this effect of "spicing" are not clear.

Previous conclusions of no effect up to FFA/Alb of 4, derived from studies with Alb solutions and in vitro additions of FFA, should be revised.

CRITICAL SERUM CA FOR CARDIAC FUNCTION: ECHO-CARDIOGRAPHIC CORRELATIONS, STUDIES IN IONIZED CA (ica), EFFECT OF THERAPY IN PROFOUND EARLY NEONATAL HYPOCALCEMIA (HC). P. Venkataraman, R. Sheldon, D. Wilson (spon. O.M. Rennert). U. Okla., Dept. Pediatr. Okla. City.

Traditionally, in infants, serum Ca<7.0 mg/dl is considered to impair cordiac function.

cardiac function, while in adults cardiac function is unimpaired with iCa of 2.4 mg/dl. In VLBW infants, we studied the thesis that decline in serum Ca to 6.0 mg/dl: 1) would not impair cardiac function, 2) iCa would be > 3.0 mg/dl, and 3) evaluated effect of Ca infusion on cardiac function. 13 normokalemic, normonatremic AGA infants, birth wt 998±198 gms (mean ±s.d.) gestation < 32 wks were studied. When serum Ca< 6.0 mg/dl, the an  $\pm 3.4$ , gestation  $\pm 32$  was were studied. When serum Ca,  $\pm 3.6$  in g/kg of Ca as  $\pm 10\%$  Ca gluconate was infused over  $\pm 10$  min. Serum Ca, iCa, Mg, P, EKG and M mode echo were obtained on entry ( $\pm 10^{\circ}$  ± 2 h), when HC ( $\pm 40^{\circ}$  ± 9 h), immediately post Ca, and  $\pm 8$  hrs post Ca. In all infants, serum Ca declined to  $\pm 7.0$  and in 7 to  $\pm 6.0$  mg/dl, Mg and P did not change. In profound HC (mean  $\pm 8.6$ ):

change. In protound i	C (Incan - 3.C.	/·		
* (p < 0.025)	Baseline	Pre Ca	Post Ca	+ 8 Hrs
Serum Ca mg/dl	$7.9 \pm 0.7$	5.1±0.2*	$9.2 \pm 0.7$	$7.1 \pm 0.3$
Serum iCa mg/dl	4.9±0.3	3.8±0.2*	$6.8 \pm 0.4$	4.2±0.2
Heart rate/min	152±4	155±5	157±3	161±3
Systolic BP mmHg	44±3	49±3	51±4	47±4
LVSTI	$.30 \pm .02$	$.31 \pm .02$	.27±.02	$.31 \pm .01$
RVSTI	$.35 \pm .02$	$.30 \pm .03$	.29± .03	.29±.02
Fiber shortening %	$33 \pm 1.5$	35.4 ±2	34.4±1.8	35±2
VcF circ/s	2.15 ±0.16	2.15 ±0.16	2.2 ±0.19	2.39 ±0.32
In VLBW infants, seru	m Ca low as	5.0 mg/dl 1)	does not imp	air cardiac
function, 2) iCa is > 3.				

NEONATAL RISKS OF A GROUP OF SGA INFANTS. Patricio 1544 Ventura-Junca, Gabriela Juez, Eduardo F. Lucero, (Spon. by Ivan D. Frantz III). Catholic University School of Medicine, Dept.of Pediatrics, Santiago, Chile.

A preceeding study demonstrated an important difference between Lubchenco's intrauterine growth curve, and a curve designed by us from normal chilean newborns. A significant increase in the number of SGA infants resulted from the use of our curve.

The purpose of this study is to determine, if this additional group of SGA infants, has higher neonatal risks than the AGA infants. Both curves were applied to 9537 consecutive live births older than 35 weeks of gestation occurred in our Hospital between Jan.1/78 and Dec.31/82. Data obtained from the comparison of the two curves were as follows:

		LGA	AG	A SG	A
Lubchenco 17%		80	8 28	(192)	
Our	curve	9,5	<b>₹</b> 79	,5% 11%	(1024)
Mo	rtality,	hospitaliza	ation and	low Apgar so	ore(<4) of the
addit	ional 84	2(9%) SGA ir	fants wer	e determined	and compared to
those	of AGA	infants. Res	sults (%,	x2 test) in	table below:
	N	Neo.Mort.	Hospit.	Apg.1min.<	4 Apg.5min.<4
AGA	7551	0,17%	10,3%	2,4%	0,15%
SGA	842	0,70%	14,6%	3,9%	0,47%
g		<.003	<.001	<.01	<.05

From these results we conclude, that the additional group of SGA infants identified with our curve, has a significant higher risk of mortality and morbidity, stressing the clinical importance of using this curve to diagnose SGA newborns in Chile.

EXOGENOUS SURFACTANT THERAPY IN INFANTS WITH RDS: COM-EXOGENOUS SURFACTANT THEKAPY IN INFANTS WITH RDS: COMPARISON OF EARLY VS LATE TREATMENT. Tetsuro Fujiwara,
Mineo Konishi, Shoichi Chida, Senji Shimada, Haruo
Maeta, Kotaro Oyama (Spon. by D. Vidyasagar). Department of Pediatrics, Iwate Medical University, Morioka, Japan.

We report successful treatment of RDS with exogenous surfactant We report successful treatment of RDS with exogenous surractant (TA). We studied the clinical course of three groups of infants with RDS. Control Grp. (C,nlO) did not receive TA, early Grp. (E,nlO) received TA at a x age of 3½ hrs. Late Grp. (L,nlO) received surfactant at x=8½ hrs. of age. TA surfactant dispersed in saline was given via endotracheal tube. Results of sequential MAP and a/APO2 are shown below. There were no differences in B.Wt. and GA between

Before 1 hr. after 12 hrs. after 72 hrs. after

E(10) a/APO <sub>2</sub>	0.29±0.11	0.58±0.12	0.63±0.16	0.72±0.7	
E(10)	MAP 2	10.0±2.3	$6.3 \pm 1.2$	4.32±0.5	4.1 ± 0.7
L(10)	a/APO2	0.26±0.09	0.51±0.13	0.62±0.13	0.54±0.15
L(10)	MAP 2	11.6±1.3	$8.5 \pm 1.9$	5.5 ± 1.2	5.4 ± 2.0
C(10)	a/APO2	0.19±0.09	0.19±0.10	0.19±0.11	0.46±0.15
C(10)	MAP 2	11.7±2.6	11.1±2.6	10.3±3.0	6.0 ± 1.3
the G	rps. Bef	ore treatme	ent MAP and a	APO2 were the	same in all
Grps.	Followi	ng therapy	, MAP dropped	significantly	both in E & L
Grps.	(p<.01)	by 1 hr. :	It decreased	steadily in Gr	p. E. The dif-
feren	ces betw	een Grps. 1	E and L were	also significa	nt (p<.01).
Simil.	arly, a/	APO2 improv	ved significa	ntly (p<.01)	in both Grps. E
and L	1 hr. f	ollowing to	reatment. Che	est x-rays clea	ared rapidly in

E & L Grps., but not in Grp. C. However, treated Grp. had high incidence of silent PDA. There were no deaths in any group. We conclude that: a) TA treatment rapidly improves the course of RDS; b) E treatment rapidly decreases MAP than Grp. L; and c) both E & L treatment are equally beneficial.

A CONTROLLED STUDY OF SURFACTANT TA IN PRETERM BABOONS 1546 WITH HMD. Haruo Maeta, Tonse Raju, Rama Bhat, Eunice John, Margaret Go, Abolhassan Yamin, Michael Evans, Dharmapuri Vidyasagar. Department of Pediatrics, University of Illinois Hospital, Chicago.

We studied the effects of exogenous surfactant (S-TA, Fujiwara)

the course of HMD in baboons delivered at 76% of term, 100 mg/kg of S-TA in saline was instilled via the E-T tube in 5 animals (Group S) at 2.0 hrs. of age; 3 animals were controls. Blood gases, lung compliance (CL) and mean airway pressure (MAP) were measured for 16 hrs. (Table:x±SD, \*p<0.02, \*\*p<0.01). Variables

Time	(age)	l hr.	2.5 hrs.	8 hrs.	16 hrs.	
CL	C Group	0.18±0.05	0.20±0.05*	0.16±0.07**	0.13±0.04*	
	S Group	0.14±0.05	0.34±0.05	0.30±0.04	0.31±0.07	
a/APO <sub>2</sub>	C Group	0.20±0.09	0.25±0.07*	0.16±0.02**	0.23±0.05*	
	2 S Group	0.19±0.11;	<0.45±0.15	0.44±0.17	0.45±0.17	
Fi02	C Group	1.0 ± 0	0.93±0.05	0.93±0.06**	0.90±0.07**	
	S Group	$1.0 \pm 0$	0.84±0.14	0.46±0.12	0.41±0.16	
MAP	C Group	15.0±0.64	14.9±0.70	16.2±1.36*	18.4±3.77**	
	S Group	14.7±1.22	13.3±1.76	12.0±3.30	11.7±1.89	
were	identical before therapy. Pre-treatment values of CL 0.14,					

a/APO2 0.19 increased in S. Grp. within 30 minutes (p<.001), to improve and were increased to 0.34 and 0.45, respectively at 16 hrs.; in C Group they were 0.13 and 0.23 at 16 hrs. (p<0.02). Pulmonary pressure volume curves obtained at autopsy showed: in S Group at 5, 10 and 30 cms of  $\rm H_2O$  distending pressure 12,15 and 20 ml volumes were achieved (deflation) but in C Group they were only 1.2, 2.5 and 5.7 mls. (p<0.001). Our results suggest S-TA significantly improves CL, blood gases, and x-ray changes of HMD.