

1134 DETECTION OF ENTEROVIRUSES BY DNA-RNA DOT HYBRIDIZATION--PROGRESS TOWARD A RAPID DIAGNOSIS. Harley A. Rotbart, Myron J. Levin, Luis P. Villarreal (Spon. by James K. Todd). Univ. of Colo. Health Sciences Center, Depts. of Pediatrics and Microbiology, Denver.

The antigenic heterogeneity among the more than 70 serotypes of enteroviruses has been the major obstacle to rapid diagnosis of infections due to these agents. A common antigen detectable by immunodiagnostic techniques has not been found. The enteroviruses do, however, share numerous physico-chemical and microbiological characteristics, suggesting some genetic conservation may exist across all of the subgroups. To test that hypothesis, we radioactively labeled fragments of DNA which are complementary (cDNA) to the genomic RNA of type 1 poliovirus. These cDNA sequences have been cloned into the pBR 325 plasmid of *E. coli* HB101. We then infected separate plates of LLC-MK2 tissue culture cells with type 1, type 2, and type 3 polioviruses, coxsackieviruses A9 and B1 and echovirus 11--representative members of the major enteroviral subgroups. Using standard techniques for dot hybridization, our type 1 poliovirus-derived molecular probes successfully detected all viruses tested. This hybridization technique is sensitive enough to follow the time course of viral replication in tissue culture. Pre-treatment of LLC-MK2 cells with 2.7 μ M Arildone (Sterling-Winthrop) ablates detectable hybridization, reflecting the anti-viral effect of this drug. Finally, preliminary trials of cerebrospinal fluid from patients with aseptic meningitis suggest that our molecular probes may be useful in a clinical setting for the rapid diagnosis of enteroviral infections.

1135 ROLE OF INTRAVASCULAR REPLICATION IN THE PATHOGENESIS OF EXPERIMENTAL HAEMOPHILUS INFLUENZAE B (Hib) BACTEREMIA. Lorry G. Rubin, Andre Zwahlen, E. Richard Moxon. SUNY at Stony Brook & Children's Hospital of Long Island Jewish-Hillside Medical Center, Dept. of Med., New Hyde Park, NY & Johns Hopkins U., Dept. of Peds. Baltimore, MD.

The occurrence of sustained bacteremia is a critical determinant in the pathogenesis of experimental Hib meningitis. We studied the potential role of (1) extravascular and (2) intravascular replication of Hib in the initiation of bacteremia. To identify tissues, in addition to nasopharynx (np), where Hib might replicate prior to seeding the bloodstream, rats (age 20 d.) were sacrificed 6-24 h. after intranasal (i.n.) inoculation with 10^5 Hib and whole organs (liver, spleen, etc.) and body fluids (CSF, joint fluid, etc.) were cultured to recover Hib. We were unable to recover Hib from any putative extravascular focus (other than np) prior to the development of bacteremia. To evaluate the potential contribution of intravascular replication in initiation of Hib bacteremia, rats were injected i.v. with a small inoculum (<100 Hib). Serial blood cultures showed a prompt (negligible lag) and exponential increase in bacterial counts in the blood in the ensuing 6 h. (mean doubling time ~ 47 min.). We noted a similar exponential increase in bacterial counts when serial blood cultures were performed 12-18 h. after intranasal inoculation with Hib. Thus, a few surviving Hib that enter the bloodstream from the np may undergo intravascular replication to initiate a sustained, high level (2×10^3 Hib per ml) bacteremia.

1136 α_2 -ACID GLYCOPROTEIN (AGP) AND C-REACTIVE PROTEIN (CRP) IN THE EVOLUTION OF BACTERIAL INFECTION OF NEONATES, Léon Sann, Françoise Bienvenu, Jacques Bienvenu, and Alistair G.S. Philip, Depts. of Pediatrics, Hôpital Debrousse, Lyon, France and Maine Medical Center, Portland, Maine.

AGP (orosomucoid) and CRP provide help in the diagnosis of bacterial infection in neonates. Their use in following the course of systemic neonatal infection was evaluated in 49 neonates (GA: mean 35 weeks, range 26-41w; BW: mean 2206g., range 880-4400) using sequential, quantitative (nephelometric) determinations. Early infection (<6 days of age) was observed in 28 patients; late in 21 patients. The outcome was favorable in all but 8 infants. Initial AGP and CRP concentrations were (mean \pm 1SD) 133 ± 75 mg/dl and 8.4 ± 7.5 mg/dl. After 5-6 days of favorable evolution, CRP decreased to a mean of 4.7 mg/dl ($<0.5-17.5$) and after 13-16 days to 2.0 mg/dl ($<0.5-18.8$). Serum AGP increased to 142 ± 73 mg/dl and then decreased to 116 ± 75 mg/dl after the same time intervals. Higher values were associated with meningitis. During the 3rd week, no CRP value was above 0.8 mg/dl, and AGP was normal at 86 ± 40 mg/dl. The normalization of serum AGP was similar to clinical healing. In contrast, a dramatic increase of CRP was observed in patients with a bad outcome, especially 3 patients who died and two with profound neurologic damage. In one patient with arthritis the recurrence of symptoms was observed with a re-elevation of serum AGP while serum CRP was normal. The data confirm the abnormal elevation of serum CRP and AGP in neonates with bacterial infection. They suggest that CRP is a reflection of the efficacy of the treatment and that the normalization of serum AGP coincides with recovery.

1137 COMPARISON OF NATIVE AND REDUCED AND ALKYLATED HUMAN IV IMMUNE SERUM GLOBULIN (ISG) IN EXPERIMENTAL H. INFLUENZAE TYPE B INFECTION. J. Schreiber, V. Barrus and G. Sibir. (Spon. K. McIntosh). Children's Hospital and Dana-Farber Cancer Inst., Dept. Inf. Dis., Boston, MA.

Reduced and alkylated (RA) ISG for intravenous use contains IgG with disrupted inter-heavy chain disulfide bonds. This change alters Fc function, particularly the ability to fix complement. We compared protective activity of RA and native (N) ISG's in infant rats passively immunized with ISG and then challenged with 10^3 Hib ip. Both ISG's contained 20,000 ng anti-PRP Ab/ml and were cidal *in vitro* in the presence of complement. Cidal activity was blocked by absorption with PRP. Cumulative bacteremia, meningitis and death 5 days after challenge were:

	N		RA		Saline	
PRP Ab dose (ng)	3,000	1,500	375	0	0	0
Bacteremia (%)	0	17	0	40	100	100
Meningitis (%)	0	0	0	40	33	100
Death (%)	0	0	0	20	0	73
Dl serum [PRP Ab]	771	804	395	513	160	130
\bar{X} cfu $\times 10^3$ /ml	0	0.1	0	5.4	2.2*	40*

High doses of both preparations of anti-PRP Ab were protective. At lower doses, bacteremia, meningitis and mortality rates as well as the magnitude of bacteremia ($p = .025$)* were higher in rats treated with RA ISG than N ISG. We conclude that RA ISG was less efficacious than N ISG in protecting infant rats from Hib infection. The mechanism of this difference may be related to altered Fc effector function of RA ISG. Differences in protective Ab specificities other than anti-PRP, however, cannot be excluded.

1138 ALTERED HELPER: SUPPRESSOR LYMPHOCYTE RATIO DURING CYTOMEGALOVIRUS INFECTION OF MICE. Matthew S. Sell, James F. Bale, Jr. and Roger Giller. (Spon. by Ronald G. Strauss), University of Iowa College of Medicine, Department of Pediatrics, Iowa City.

Cytomegalovirus (CMV) infections of humans and experimental animals have been associated with altered cell mediated immunity. In a murine model, we have observed an increased susceptibility to secondary infection during acute murine CMV (MCMV) infection. To further investigate the effect of MCMV infection upon host defense, we studied helper: suppressor lymphocyte (H:S) ratios in Balb/c mice undergoing sublethal MCMV infection. On days 1, 3, 5 and possibly 9, H:S ratios were reduced. On day 3 the H:S ratio was 0.82 in MCMV-infected mice vs 1.78 in controls ($p < 0.01$) and on day 5, 0.59 vs 3.85 ($p < 0.01$). On days 3 and 5, reduced H:S ratios were attributable to decreased absolute numbers of helper lymphocytes. Thereafter, suppressor cell numbers increased. Infectious MCMV was recovered from blood lymphocytes and bone marrow on days 3, 5 and 9 and from spleen on days 1 through 16. Alterations in H:S ratio appeared to correlate most closely with recovery of infectious virus from spleen ($p = 0.07$). Examination of bone marrow preparations suggested diminished cellularity on days 1 through 5. These results indicate that acute MCMV infection is associated with reduced H:S ratios and suggest that altered H:S ratios may correlate with enhanced susceptibility to secondary infection. These observations may be relevant to the association between CMV and opportunistic infections in human organ transplant recipients as well as in patients with the acquired immune deficiency syndrome.

1139 EFFECTS OF HYDROGEN PEROXIDE (H_2O_2) ON PROSTACYCLIN AND OTHER VASCULAR ARACHIDONIC ACID METABOLITES. B.N.Y. Setty, Elizabeth Jurek, Carolyn Ganley, and Marie Stuart, Dept Pediatrics, SUNY, Upstate Med Ctr, Syracuse, N.Y.

Local production of H_2O_2 by granulocytes during phagocytosis affects the function of red cells and platelets. We evaluated the effects of H_2O_2 on vascular arachidonic acid (AA) metabolism and report that H_2O_2 treatment inhibits prostacyclin (PGI_2), PGE_2 , and $PGF_{2\alpha}$ production. Short term exposure of human umbilical arteries to H_2O_2 (25 to 200 μ M) resulted in a concentration dependent inhibition in the ability of vessels to produce $6KPGF_{1\alpha}$ (the stable metabolite of PGI_2) either from endogenous or exogenously provided substrate. Following exposure to 200 μ M H_2O_2 , vessels produced 1.45 ± 0.24 (1SE) pmol $6KPGF_{1\alpha}$ per mg (by RIA) compared to 2.08 ± 0.29 in paired control segments, an inhibition of 33% ($n=8$; $p < 0.001$). Mean inhibitions of 15% and 21% were observed at H_2O_2 concentrations of 50 and 100 μ M. Inhibition was present at 5' post addition of H_2O_2 , with maximal effects occurring by 15'. Production of $6KPGF_{1\alpha}$, PGE_2 and $F_{2\alpha}$ from exogenously provided ^{14}C AA was inhibited to a similar degree in microsomes from these vessels, suggesting that the effect occurred at the cyclo-oxygenase level. Following incubation with 200 μ M H_2O_2 , mean ^{14}C metabolite production was 449 ± 134 pmol per mg microsomal protein compared to paired control values of 612 ± 146 pmol, ($n=5$; $p < 0.01$). Catalase (250U/ml) abolished the effect, whereas SOD (2.5 μ g/ml) did not. PGI_2 , PGE_2 , and $F_{2\alpha}$ have been shown to affect leucocyte adherence to vascular endothelium, and to play a role in edema formation. Release of H_2O_2 from phagocytes may play a role in the inflammatory process by modulating prostaglandin biosynthesis.