

DEVELOPMENTAL CHANGES IN VENTRICULAR FUNCTION IN THE ●91 CHICK EMBRYO

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City, IA How ventricular function changes during embryonic development is unknown. We correlated measures of ventricular physiology is unknown. We correlated measures of ventricular physiology with ventricular dry weight and myocyte myofibril and mitochondrial volume in stage 18 (3 day), 21 (3½ day), 24 (4 day), 27 (5 day) and 29 (6 day) chick embryos. In ovo measurements of ventricular pressure and dorsal aortic blood velocity were made with a servo null system and 20 MHz pulsed-Doppler respectively. dP/dt and dV/dt were electronically derived. Percent cell volume of mitochondria and myofibrils were dotomicad from electors mismorabe of the ventricular walls were derived. Percent cell volume of mitochondria and myotibrils were determined from electron micrographs of the ventricular wall. From stage 18 to 29, ventricular weight increased from 28 ± 2 µg ($\bar{x}\pm$ SEM) to 140±6 µg while embryo weight increased from 1.02±.05 mg to 15.36±.30 mg. Ventricular systolic and end-diastolic pressure, and dP/dt increased from 1.31±.05 mmHg, .23±.03 mmHg and 23±1 mmHg/s. Dorsal aortic dV/dt increased from 878±17 mm/s² to 2076±65 mm/s². Relative myofibril volume increased from 17% at stage 18 to 24% at stage 27 but decreased to 18% at stage 29, while mitochondria volume remained constant at 11%. With while mitochondria volume remained constant at 11%. With development, heart:embryo weight ratio and function measures indexed for myocardial weight or myofibril percent decreased as much as 50%. We speculate that the decrease in indexed function may be due to changes in energy utilization, available energy, or geometric shape of the developing heart.

CARDIOPULMONARY EFFECTS OF A PATENT DUCTUS •92 ARTERIOSUS. Ronald I. Clyman, Michael A., Heymann, Francoise Mauray, CVRI and Dept. of Pediatrics, University of California - San Francisco and Mt. Zion Hospital,

San Francisco, CA San Francisco, CA The premature newborn's response to increasing degrees of $Lt \rightarrow RF$ PDA shunt is unknown. In 14 fetal lambs at 128d gestation (term 147d), we infiltrated the ductus with formalin and placed a mechanical occluder about it so that its patency could be regulated. 3d later the lamb was delivered, given sheep surfactant, paralyzed and mechanically ventilated. Microsphere measurements of left ventricular output (LVO) and Microsphere measurements of left Ventricular output (LVO) and its distribution were made prior to and 1.25 h after a change in ductus patency. Twenty four changes from open-closed or closed->open were assigned randomly from 3 to 7 h after delivery. Lambs with ductus shunts < 50% LVO (38%) had a significant increase in LVO (167 vs. 268 ml/kg/min:closed vs. significant increase in LVO (167 vs. 200 mir/kg/min/closed vs. open) with no change in effective systemic blood flow or BP. There was a significant increase in coronary flow but no change in flow to the kidneys, GI tract, brain, or carcass (m1/100 gm/min). Although lambs with shunts > 50% LVO (65%) had a significant increase in LVO (164 vs. 300 m1/kg/min), there was a significant decrease in effective systemic blood flow and BP. There were significant decreases in flow to the kidneys, GI tract, brain and carcass. Coronary flow was unchanged but there was a decrease in subendocardial flow. There was no change in HR or pulmonary compliance at either degree of shunt. Preterm lambs are capable of compensating for moderate degrees of PDA shunt by increasing preload and increasing stroke volume; highe degrees of shunt significantly decrease organ blood flow.

NONINVASIVE ESTIMATION OF PULMONARY ARTERY (PA) NONLIVASIVE ESTIMATION OF PULMONARY ARTERY (PA)
PRESSURE IN CHILDREN BY DOPPLER ULTRASONOGRAPHY, Allan G. Cougle, John M. Eckerd, Gerald A. Serwer,
Duke University Medical Center, Dept of Pediatrics, Durham, NC Noninvasive means for PA pressure assessment has been sought without a reliable method found. Doppler interrogation of PA flow offers the potential for an easily performed method of es-timating PA pressure from changes in velocity patterns. 34 chil-dren, ages 1 day-23 yrs (mean 5.3 yrs) with a variety of cardiac abnormalities, underwent simultaneous PA Doppler study and PA pressure measurement during the course of routine cardiac cath-eterization. 14 had peak systolic pressures greater than 40 torr. abnormal releases, under which similar tank tank course of proutine cardiac catheterization. 14 had peak systolic pressures greater than 40 torr. Data was collected by an on-line computer with analysis being performed by computer. Time from onset of ejection to peak velocity (TPV) was measured and compared to that of a separate group of 33 nl pts, similarly studied. As TPV varied inversely with heart rate (HR) in nls, TPV was divided by predicted TPV (TPVN) as determined by linear regression equation (TPVN=182-0.79 KHR) derived from nl pt data. The ratio of TPV/TPVN was HR independent. TPV/TPVN was linearly related to peak PA pressure (r=0.88, p<.01), and less well to PA mean (r=0.85) and PA diastolic pressure (r=0.77). From nl pt data TPV/TPVN ratios <0.74 were considered abnormal. No pt with a PA peak pressure \leq 40 torr had an abnormal ratio (100% specific). 13 of 14 with PA pressures > 40 torr had ratios <0.74 (93% sensitive). Thus TPV/TPVN was an easily measured noninvasive parameter which is a highly sensitive and specific predictor of pts with significant PA systolic hypertension. sion。

SYSTOLIC LEFT VENTRICULAR AND AORTIC BLOOD FLOW RE-SPONSE TO INCREASING SYSTEMIC VASCULAR RESISTANCE 94

24 <u>Christine M. Donnelly, Linda J. Addonizio, Fredrick</u> Z. Bierman, <u>Lynne L. Johnson</u> Dept. of Peds. Columbia University College of Physicians and Surgeons, New York, N.Y.

Classical indices of myocardial performance may be insensitive to changes in myocardial reserve when native afterload is reduced The objective of this study was to examine the response of left ventricular and regional acrtic blood flow to changes in systemic vascular resistance(SVR) in a canine model using combined gated pulsed doppler echocardiography(GPDE) and a non-imaging car-diac probe(NICP). SVR was augmented by continuous methoxamine infusion and calculated by standard thermodilution techniques in 5 animals (3-5kg). Aortic blood flow was evaluated by GPDE interrogation of the thoracic aorta. Systolic aortic blood flow was described by the mean velocity and the frequency time integral partitioned into the first and latter two-thirds of systole. The left ventricular ejection rate and stroke volume were measured simultaneously by NICP. Increasing SVR resulted in a pansystolic mean systolic aortic flow velocity. A normal reciprocal relation-ship was present for the first and latter two-thirds of left ventricular and aortic systolic blood flow as measured by percent change of stroke volume. This baseline relationship persisted with increasing SVR. In conclusion, combined GPDE and NICP can provide an effective means to noninvasively evaluate left ventri-cular performance and regional aortic blood flow changes with augmentation of afterload.

95 GATED PULSED DOPPLER ASSESSMENT OF SIMPLE COARCTATION OF THE AORTA IN PEDIATRIC PATIENTS. Christine M. Donnelly, Fredrick Z. Bierman. Columbia University,

Dept. of Pediatrics, NY, NY. Pulsatile flow dynamics are altered distal to an area of obstruction. To investigate the effects of simple coarctation of the aorta (CAo) on descending aortic blood flow patterns, doppler interrogation of the thoracoabdominal aorta was performed in 12 infants with GAO less than 18 months of age, and 15 age matched controls. All patients with CAO demonstrated continuation of high forward flow throughout the cardiac cycle with prolongation of both acceleration and deceleration phases. Indexing the ratio of acceleration and deceleration phases of flow with acceleration time, measured as the time from on-set of flow to peak frequency corrected for cycle length, separated CAo patients from normal subjects. In contrast, dopler frequency distributions from the ascending aorta did not disdoppler examination of the thoracoabdominal aorta displayed a frequency distribution specific for coarctation of the aorta. The discrepancy between ascending and descending frequency distributions represents systolic loading and diastolic decompression of the obstructed proximal thoracic aorta. This method can provide a useful adjunct to serial follow-up of coarctation patients and early detection of recoarctation.

INFLUENCE OF α -BLOCKING DRUGS PRAZOSIN (Pz), PHENTO-1960 RESPONSE TO DOPAMINE IN AKAKE LAMSS <u>Willa H. Drum</u>-montal Becky Williams, Hugh H. Shrager, Wendy A. Daily. Univer-sity of Florida College of Medicine, Gainesville, H. Combination vasodilator and cardiotonic therapy may help fiber in the pulmonary vasospasm and low cardiac output. To test with effectiveness of α -blockers with different receptor pressure in the pulmonary varery (PAP), aorta (SAP) and left atrium (LAP). The ductus arteriosus was tied and a flow seponse to 2.7, 27 and 270 ug/kg/min was measured then repeated with Phe (1 mg/kg, n=5), Pz (1 mg/kg, n=6), or T (5 mg/kg/hr, n=3). Dopamine raised SAP, PAP, LAP, systemic vascular resis-sponse to 2.7, 27 and 270 ug/kg/sin was measured then repeated with Phe (1 mg/kg, n=5), Pz (1 mg/kg, n=6), or T (5 mg/kg/hr, n=3). Dopamine raised SAP, PAP, LAP, systemic vascular resis-pulmer vascular resistance (PK) orso at the top dose (p².01) while QP didn't change and PVK/SVR fel1 (p².01). Only Phe raising HR (208420, to 256428, p².01). All drugs blocked dopar interinduced SAP and SVR rise and allowed increased QP, HR and population with systemic α -receptors, all increased HR dopamine induced PAP increase, which Phe did not; PVR was lower was for twas coupled with dopamine (p².05). With Pz and Phe, PVR was dopamine interaction with systemic α -receptors, all increased HR docardiac output when coupled with dopamine. None completed bi butted it. Thus, all α -blockers can facilitate the cadiotonic provided dopamine's pulmonary circulatory effect although the pulmet of the dose dopamine, but none completely ablates to pulmet it, perhaps because of its non-adrenergic effect.