DIGITAL SUBTRACTION ANGLOGRAPHY IN SEVERE SEQUELAE OF VIRAL PNEUMONIA 138

D. Pariente, J. de Blic, <u>P. Scheinmann</u>, D. Lallemand D. Pariente, J. de Blic, <u>P. Scheinmann</u>, D. Lallemand Depts of Pediatric Radiology and Pediatrics, Univ. Paris V; Hopital des Enfants-Malades, Paris, France. 16 children presenting with severe viral pneumonia sequelae (VPS). were explored with digital subtraction angiography (DSA) in the last 2 vesse. The clinical matterns were severe perimeratenisodes of were explored with digital subtraction angiography (DSA) in the last 3 years. The clinical patterns were severe recurrent episodes of wheezing, significant dyspnea on exertion and/or productive cough. In all patients the initial episode of viral infection could be recognized. The age range was 18 months to 17 years and only the 3 younger patients were sedated for the procedure. The contrast medium injustion, was conformed providerable view in a large term. injection was performed peripherally via a large vein, Nc complication was observed except one mild episode of wheezing after complication was observed except one mild episode of wheezing after contrast injection. In all cases DSA showed perfusion defects which were peripheral. These anomalies were multiple in 14 cases, bilateral in 10 and unilateral in 4. In only 2 cases the defect was localized in one lobe. The proximal pulmonary arteries were always well seen, and of smaller size in the more affected lung in 5 cases. Correlations with isotopic studies were consistent, but DSA gave more information by demonstrating pulmonary vessels. Thus DSA, that is non annessive proredure, seems to be very valuable in the more information by gemonstrating pulmonary vessels. Hilds vow, that is non aggressive procedure, seems to be very valuable in the investigation of VPS. DSA displays an angiographic pattern that is strongly suggestive of this diagnosis. Moreover it has a pronostic value as it precises very accurately the extension of parenchymal lesions.

139 TRACHEAL DIAMETERS AND DYNAMICS IN INFANTS AND CHILDREN HEASURED BY CINE-CT. L. Newth, RC Brasch, RG Could, CA Gooding, MJ Lipton (Introduced by MM Grumbach, M.D.), Depts. of Pediatrics (Introduced by California, San Francisco, CA, USA and Radiology, University of California, San Francisco, CA, USA Nine children were evaluated for tracheal obstruction by having their intra- and extra-thoracic tracheal diameters measured using Cine-CT. The patients ranged in age from one month to four years, and had clin-ical signs and symptoms of respiratory distress with stridor. They entered a protocol in which there was full clinical evaluation, and in some, measurement of esophageal pressures and tidal breathing flow-volume loops. The patients were scanned without sedation using a cine-CT scanner (Imstron, Inc., CA) which provided eight level image acquisition in 240 mage without moving the patients. Using the scanner in the "Cine" mode, 13 to 20 slices obtained over 1 to 2 respiratory cycles were obtained at several levels of both the intra-and extra-thoracic trachea to detail dynamic changes in airway cali-ber. Images were obtained at relatively low dose and were recon-structed in 14 seconds. Cross-sectional areas of the tracheal diameters directly measured. The clinical and laboratory data correlated well with the imaging measurements, and were compared with Age-related normal Cine-CT tracheal diameters, and published post-mortem airway caliber data. We conclude that Cine-CT is a rapid method for tomographic evaluation of tracheal dynamics that can be employed for children without sedation-

EPRS—Abstracts for Poster Presentations

SLOW RELEASE THEOPHYLLINE IN PRE-SCHOOL ASTHMA: 140Loftus BG, Price JF, Dept. of Child Health, King's College Hospital, London, SE5 9RS.

Slow release theophylline is widely used in the management of few studies have assessed the drug in pre-school age group. The studies have assessed the drig in pre-school age group. Over a 9 month period, 49 pre-school children with chronic asthma (age range 1 year 6 months - 5 years 9 months) were started on slow release theophylline using a recommended regime. Mean initial dosage was 18.1 $\pm 2.3 \text{ mg/kg}$, day in 2 doses. Blood levels ranged from 3-21 mg/lt (mean \pm SD 9.1 ± 4.1). The dosage was increased in 23 children because of sub-therapeutic blood levels. 8 nationity were non-compliant = 4 denied non-compliance and 4 8 patients were non-compliant - 4 denied non-compliance and 4 patients were non-compliant - 4 deniet non-compliance and patients refused to take the preparation. 16 stopped therapy because of side effects. 6 had behavioural problems, 9 vomiting and 1 nightmares. Symptoms resolved when drug was withdrawn. 13 had satisfactory control of asthma. 12 were deemed treatment failures because of persisting symptoms, hospitalisation or repeated courses of steroids. Outcome might have been improved by a more gradual introduction of therapy and by use of an eight-buryly decare actually the the derived interaction theoremultion hourly dosage schedule, but the data indicates that theophylline usage in this age group needs to be critically re-evaluated.



J. Bauer, <u>H. Lindemann</u>, G. Hüls, H.J. Schwandt Children's Hospital, Justus-Liebig-University, Giessen

This study was performed to evaluate the suitability of monitoring theophylline concentrations in the lacrimal fluid for therapy control in asthmatic children.

Samples of plasma and lacrimal fluid of 38 children (5 - 16yrs) were collected simultaneously and measured by means of high-performance liquid chromatography. Lacrimal fluid was collected using a small strip of filter paper placed into the conjunctival sac of the lower eyelid.

Comparison of theophylline concentrations in plasma and lacrimal samples showed a significant difference. Analysis of the data, howsamples showed a significant difference. Mulysis of the data, how ever, revealed a close linear correlation (r = 0,83) between theo-phylline concentrations of the two fluids. The intercept of the regression line is very small ($\gamma = 0,55x + 0,26$). The mean plasma/lacrimal fluid ratio was 1,79 ($\frac{1}{2}$ 0,40).

It is concluded, that monitoring theophylline concentrations by means of lacrimal fluid analysis is a valuable tool for therapy control.

THEOPHYLLINE CONCENTRATION MEASUREMENTS IN 142 SERUM OR SALIVA FOR THERAPY MONITORING IN BRONCHIAL ASTHMA.

HRONCHIAL ASTHMA. W. Leupold, K. Feller, E. Paditz, K. Richter (Introduced by H. von der Hardt). Depts. Paed. and Clin. Pharmacology, Medical Academy Dresden, GDR Theophylline concentrations in serum and in saliva obtained without and with stimulation (dil. citric acid) of 20 patients in the age range of 5-19 years on a long term theophylline therapy were simultaneously assessed within 6 hours after drug intake; at the same time seve-ral parameters of the lung function were taken. The theo-phylline concentration measurements were performed gasphylline concentration measurements were performed gas phylline concentration measurements were performed gas-chromatographically. The saliva concentrations obtained without stimulation very strongly correlated with the serum concentrations (r=0,98; mean deviation from the regression line 0.70/ug/ml). The saliva concentrations obtained with stimulation revealed a significantly weaker correlation with the serum concentrations (r=0,82). A decondernee of the concentrations (r=0,82). A correlation with the serum concentrations (r=0,82). A dependence of the concentrations from the salivation rate (ml/2 minutes) could be identified (r=0,66) in the saliva obtained with as well as in that obtained without stimulation; this dependence exists both intra- and inter-individually. Reliability and evidence of the saliva concentrations obtained without stimulation can be further increased by measurement of the secreted saliva volume per period of time.

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NEBULISED SALBUTAMOL IN ACUTE ASTHMA - EFFECT ON OXYGENATION. Loftus BG, <u>Price JF</u>, Dept. of Child Health, King's College Hospital, London, SE5.

The effect of oxygenation of nebulised salbutamol, given with compressed air, was studied in 18 patients (aged 18 months to 9 years) with acute asthma. A Radiometer TCM2 transcutaneous oxygen monitor was used, preliminary studied having shown that this accurately reflects trends in arterial oxygen tension. Heart and accurately reflects trends in arterial oxygen tension. Heart and respiratory rates were measured at 10 minute intervals and transcutaneous oxygen (TcO₂) continuously. Values of TcO₂ were averaged over 10 minute periods. Given when baseline TcO₂ was stable, nebulised salbutamol was associated with a rise of TcO₂ in 9 children (range 1.1 - 1.3 mmHg, mean 5.2) and a fall in 4 (range 1.1 - 7.3 mmHg, mean 4.7) at 10-20 minutes after nebulised therapy. 5 children fell asleep during or shortly after treatment, and all of these showed a fall in TcO₂ (Range 5.4 - 19, mean 9.8) with the onset of sleep. Heart rate increased in all, this effect being maximal immediately after nebulisation. The changes in TcO₂ were not predictable on the basis of age, initial TcO₃. Heart or respiratory rate, though children who fell asleep tended to have higher initial pulse and respiratory rates. These results emphasise the need for caution in the use of home nebulised therapy in acute the need for caution in the use of home nebulised therapy in acute asthma.