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INSULIN-LIKE GROWTH FACTORS AND THEIR BINDING PROTEINS IN EARLY MILK FROM MOTHERS OF PRETERM AND TERM INFANTS AND THEI BIOLOGICAL REL EVANCE

EVANCE <u>A Louil</u>, M W Elmlinger<sup>2</sup>, F Hochhaus<sup>3</sup>, R Grund<sup>2</sup>, M Obladen<sup>1</sup>, M B Ranke<sup>2</sup> <sup>1</sup>Charité Virchow Hospital, Dpt. of Neonatology, Berlin, Germany; <sup>2</sup>University Children's Hospital, Dpt. of Pediatric Endocrinology, Tuebingen, Germany; <sup>3</sup>Ludvig-Maximilians-University. Dpt. of Ophthalmology, Munich, Germany **Background**: Breast fed preterm infants often show a better neurological outcome than formula-fed. The benefit of human milk is partly ascribed to it's content of the growth promoting insulin-like growth factors (IGFs). The aim of this study was to investigate, whether the content of IGFs und Their binding proteins (IGFBPs) is higher in human milk from methors. of neutron infants then of them is following in character and to text their isolation enlawance.

mothers of preterm infants than of term infants and to test their biological relevance. Methods: In a prospective study we investigated the concentrations of IGF-I, IGF-II and their binding proteins IGFBP-2 and IGFBP-3 in human milk from mothers of 30 preterm infants (<31 SSW) and of 19 term infants (>37 SSW) at day To and 21 after birth by means of an RIA, and the protein content of human milk by the Bichinonic acid (BCA) method. We analyzed the specific protoelysis of IGFBP-2 in human milk and the stability of 1251-IGF-1 and 125IGFBP-2 in the presence of gastric juice from neonates. **Results:** Mean concentrations ( $\pm$ SD) of IGF-I(2,8  $\pm$  0,2 vs. 2,3  $\pm$  0,1 ng/ml), IGF-II (12,0  $\pm$  0,4 vs. 12,2  $\pm$  0,5 ng/ml)

Results: Mean concentrations ( $\leq$  SDJ) of IGF-1(2,8  $\leq$  0.2 Vs. 2,3  $\leq$  0.1 ng/m), IGF-11(12,0  $\leq$  0.4 vs. 12,2  $\leq$  0.5 ng/m)) and IGFBP-3 (100,0  $\leq$  5,1 vs. 80,0  $\leq$  5,8 ng/m)) did not differ in human milk of preterm versus term infants on day 7. The content of IGFBP-2 (3144  $\leq$  172 vs. 2428  $\leq$  188 ng/ml) was higher in preterm human milk (p color), IGF-1, IGF-1, IGFBP-3 (correspondence) and IGFBP-2 (3144  $\leq$  172 vs. 2428  $\leq$  188 ng/ml) was higher in preterm human milk (p color), IGF1-1, IGFE-1 and IGFBP-2 (di sightly change until day 21. IGFBP-3 decreased from day 7 to day 21 by 41% (27% in human milk of preterm/ term infants (p<0,05). The protein content vas equal to term infants was on day 7 (2,5  $\leq$  0,1 g/dl) higher than on day 21 (p<0,05). On day 21 the protein content was equal to term infants (1,9  $\pm$  0,1 g/dl). Preterm human milk contained about 42% more IGFBP-2 framements of 14 und 25 kbal (p<-0,05). Incubation with gastric juice led to a complete digestion of 125IGFBP-2 and a partial digestion of 1251-IGF-II, while complexation with each other protected IGF-II and IGFBP-2 frame cleavage

UGEBP-2 from cleavage. Conclusion: We suggest that IGF-II and IGFBP-2 play a role in the nutrition of preterm infants. Both are found in a Conclusion: We suggest that IGF-II and IGFBP-2 play a role in the nutrition of preterm than in term human milk for the first constant level in human milk and IGF-2 showed a higher concentration in preterm than in term human milk for the first 3 weeks. IGFBP-2/IGF-II-complexes can partly pass the stomach and may exert their promotive effects in the intestine Taken together it can be concluded that preterm infants profits best from breast milk of it's own mother.

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#### THE "NEW"BRONCHOPULMONARY DYSPLASIA. EPIDEMIOLOGICAL STUDY ON THE EURONEONET DATABASE

THE EURONEONET DATABASE <u>B Loureiro</u>, K Latorre, S Páramo, A Cotero, C Centeno, Y de Carlos, A Valls-i-Soler, E N N Spanish Collaborative Group <u>Cruces Hospital</u>, Department of Paediatrics. Neonatal Unit. University of Basque Country., Bilbao, Spain **Background**: Bronchopulmonary dysplasia (BPD) has classically been related to the use of high oxygen concentrations and mechanical ventilation in infants with respiratory distress syndrome (RDS). However, an increasing number of very low birth weight infants (VLBW1) develop BPD without having acute RDS, which has been termed "new" BPD. **Aim:** To identify the characteristics of VLBWI who develop this "new " form of BPD. **Methods:** This is a multicenter, prospective, longitudinal study of a cohort of 1798 VLBWI born in 2001 and 2002, at one of the 30 Spanish NICU included in EuroNeoNet initiative. The database, including demographic characteristics, risk factors, interventions and outcomes, was searched to identify babies alive at 36 weeks PMA (n=1549). BPD was diagnosed (need of oxygen at 36 PMA) in 221(142<sup>6</sup>), babies 32 of who did not have RDS. An univariate and logistic repression (need of oxygen at 36 PMA) in 221(14%) babies, 32 of who did not have RDS. An univariate and logistic regression analysis was used to asses the risk factors for BPD **Results:** In comparison to the classical BPD, babies with the "new" BPD were more mature, had a higher birthweight,

needed less resuscitation manoeuvres and respiratory support, had a lower incidence of PCA, but a higher one of early sepsis (7.4 vs. 12.5%) and NEC (6.9 vs. 21.9 %). Multifactorial analysis in infants without RDS (n=804), showed an increased risk for BPD related to use of CPAP (0R 3.6; 95% IC 1.5–8.8) and mechanical ventilation (4.8; 1.9–11.8), and to the presence of both, early sepsis (9.1; 1.9–42.4) and late sepsis (4.8; IC 2–11.1). However, in patients with RDS (n=745) we showed an increased risk related to CPAP, mechanical ventilation, late sepsis and PDA (1.7, 1.1-2.6), but not with early onset sepsis

Conclusion: As previously reported (1), we showed risk factors of developing DBP in infants with and without RDS are CPAP, mechanical ventilation, and late sepsis. Nevertheless, in babies without RDS, BDP is associated with early sepsis, and in those with RDS with PDA. Despite the small number of patients without RDS who developed BPD in our cohort, the higher incidence of early sepsis and NEC point out toward prenatal factors related to fetal inflammatory disease (2). (1) Rojas MA et al. J Pediatr 1995;126:605–10. (2) Lyon A. Eur J Pediatr 2000;159:798–802.

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## COMPARISON OF RESUSCITATION OF PIGLETS WITH 21% AND 100% OXYGEN

COMPARISON OF RESUSCITATION OF PIGLETS WITH 21% AND 100% OXYGEN AFTER EXPOSURE TO INFLAMMATION AND HYPOXIA K\_Lyng<sup>1</sup>, M readmin<sup>2</sup>, J F Froer<sup>2</sup>, B Stray-Pedersor<sup>2</sup>, O S Dagustad<sup>2</sup> Hiskshopitalet, Department of Gynaecology and Obstetrics, Department of Pediatric Research, Oslo, Norway: <sup>2</sup>Riskshopitalet, Department of Pediatric Research, Oslo, Norway: <sup>2</sup>Riskshopitalet, Department of Gynaecology and Obstetrics, Oslo, Norway Objective: To assess if endotoxin increases vulnerability to hypoxia and if resuscitation with 10% O2 increases brain damage in the presence of inflammation. Method: After anaesthesia and surgery, newborn pielets (n=31) wer randomized to four interventional groups. Two groups received pretreatment with saline for 60 min followed by hypoxia and resuscitation with 21% (Group 1, n=8) or 100% oxygen (Group II, n=8). The other two groups received pretreatment with endotxin for 60 min followed by hypoxia and resuscitation with 21% (Group II, n=7) or 100% oxygen (Group IV, n=8). Hypoxia was administered for the first 30 minutes, followed by 21% oxygen for 150 min for all groups. During the experiment, we measured extracellular brain tissue glucose, glycerol, and lactate/pyruvate by microdialysis, brain tissue oxygen tension and laser doppler flow. laser doppler flow.

Results: Administration of endotoxin caused a reduction in the time to reach BE -20 by median 31.5 minutes compared The second seco

Conclusion: Endotoxin and hypoxia acted synergistically in inducing arterial acidosis. In the presence of experimental inflammation, 21% seems as safe as 100% O2 in reoxygenating newborn piglets.

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# BABYLINK - A WEB BASED TOOL TO IMPROVE COMMUNICATION BETWEEN CLI-

**BABTIANCE AND PARENTS IN A NEONATAL UNIT** <u>AJ Lyon</u>, Y Freer, C Coyle, B Stenson Royal Infirmary of Edinburgh, Neonatal Unit, Edinburgh, United Kingdom **Background**: There is an ever-increasing demand for improved communication with parents and their families. At present, parents gather information from multiple sources, which can lead to incorrect and/or inconsistent advice and information being received. The aim of this project was to develop, and evaluate a system that gives parents secure access via the intermet to clinical information, extracted automatically from the electronic patient record and presented in a format appropriate to parents. **Method**: Software has been written that searches the electronic particle resolution in proceeding of the provide the process of the automatically to a library of short parent information templates and creates a parent report that is clinically relevant to the problems of the individual baby. The administrator can define the format of the reports, as well as the frequency and time interval sampled. Reports are encrypted and sent to an internet server. Parents access the reports on the Babylink website (www.babylink.info/edinburgh), using their own log-in and password. They can also view their baby's diary, which contains photographs and non-clinical messages written by the staft. The web-site has an open area providing general details about the neonatal unit and generic information about common neonatal problems. As part of data validation, reports have been checked manaully for accuracy. Parents can comment or ask specific questions about their baby on-line. This feedback has been used to develop the website and the content of the reports. The project was approved by the local ethies committee and parents give written, informed consent before information is forwarded to the website. **Results:** The reports and diaries have been successfully implemented. To date the system has been live for 6 weeks. 23 families have been accessing reports on the internet. Two refused to be involved. There has been no negative feedback, with harents expressing very nositive views. Many constructive comments have helped develop the website feedback, with the arent

parents expressing very positive views. Many constructive comments have helped develop the website and the parent reports. Conclusions: This system offers the facility for parents to access specific, up to date information about their baby. It can

be used with any electronic record system from which data can be exported. Further developments include prospective tailored information about discharge planning and care of the baby at home. The effect of the system on parental perceptions of communication is being evaluated.

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#### ASYMPTOMATIC CONGENITAL CYTOMEGALOVIRUS INFECTION

ASI INTOMATICO CONTRACTOR DOMESSION OF A CONTRACTOR OF THE CHONAL ALL AND A CONTRACTOR OF A CONTRACT AND A C Amadora, Portugal

Background: Cytomegalovirus (CMV) is the most common congenital infection in developed countries with reported incidences varying from 0.15% to 2.0%. Congenital CMV infection may be a cause of a congenital syndrome or run without symptoms. Approximately 90% of congenitally infected infants are asymptomatic at birth but may still present handicaps at a later age.

Mucaps at a fater age. Aims: The objectives of the present study were to evaluate the importance of this agent as cause of morbidity and long m sequelae, and to assess the utility of the different methods of congenital CMV infection screening Methods: Between May and August 2002, 757 consecutive newborns were screened for CMV infection by Polymerase

Methods: Between May and August 2002, 757 consecutive newborns were screened for CMV infection by Polymerase Chain Reaction (PCR) in corb blood and through the isolation of CMV by *shell-vial* from urine on the first two days of life. Epidemiological characteristics of the mothers and newborn were analysed from medical records. Follow-up was done at 3, 6, 9, 12, 15 and 18 months with neurologic, audiologic, ophthalmologic and development assessments. **Results**: Asymptomatic congenital CMV infection was diagnosed in 5 out of 757 (0.7%) by means of a virus culture in urine. PCR in cord blood was negative in all cases. In three newborns, within the first 3 weeks of life, the clinical findings included: pneumonia (1); bone marrow failure (1); hepatil gailig avascultits (2). At 18 months, follow-up of two infants showed short stature (1), minor abnormal neurodevelopment (1) and partial hearing loss (1). **Conclusion**: The incidence of congenital CMV infection was similar to that reported in other studies on highly immune populations. Urinary CMV culture is a reliable and convenient method, and may be a screening tool for the detection of congenital CMV infection. Infants with asymptomatic congenital CMV infection may have sever diseases that are not clinically evident at birth. In our series of studies, secuelae were found in 3 out of 5 (60%) children. Funded by Science clinically evident at birth. In our series of studies, sequelae were found in 3 out of 5 (60%) children. Funded by Science and Technology Foundation

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EARLY FEEDING TOLERANCE IN VERY LOW BIRTH WEIGHT INFANTS WITH IN-

 
Exact Control Contrecontrol Control Control Control Control Control Con growth retradition (IUGR) should be considered an important factor in decisions about their enteral nutrition. We conducted a retrospective cohort study to evaluate whether IUGR (birth weight and/or length at least 1.28 standard deviations (SD) below the mean for gestational age) was independently associated with early feeding intolerance in VLBW infants

Methods: All infants admitted during 2000–2003 to our NICU who lived for at least 28 days were included in the study if they had BW  $\leq$ 1500 g and no major congenital anomalies. They were fed following a standardized protocol beginning within 48 hours of age with  $\leq$ 20 ml/kg daily increments. Feeding tolerance was assessed as the age at which full enteral feeds (150 mL/kg/d) were achieved. Multiple regression analysis was used to identify factors that might influence early

feeds (150 mL/kg/d) were achieved. Multiple regression analysis was used to identify factors that might influence early feeding tolerance during hospitalization **Results**: 78 SGA and 143 AGA infants met the inclusion criteria. SGA infants had a similar BW (1188 $\pm$ 246 vs 1138 $\pm$ 227 g), higher GA (31.7 $\pm$ 1.7 vs 28.6 $\pm$ 1.8 wks; p<0.001), lower birth Z scores (ZzW +1.65 $\pm$ 0.64 vs -0.06 $\pm$ 0.74, ZsHC-1.54 $\pm$ 0.74 vs -0.16 $\pm$ 0.67, ZsL -1.82 $\pm$ 0.89 vs -0.13 $\pm$ 0.65; p<0.0001) and lower CRIB score (1.0 $\pm$ 1.7 vs 2.6 $\pm$ 2.9; p<0.0001) than AGA infants. Thirty-nine (50%) SGA infants had all the 3 Zscores <-1.28 D0 (Symmetric SGA) and 20 (16.7%) had 2 Zscores <-2 SD (Symmetric SGA) and 20 (16.7%) had 2 Zscores <-2 SD (Symmetric SGA) and 3bowed shorter time to achieve full feeds after initiation of milk feeds (17.0 $\pm$ 9.0 vs 21.4 $\pm$ 11.8 days; p = 0.048). No difference was found in percentage of human milk intake from the beginning of milk feeding until full enteral feeds were achieved (59.4 $\pm$ 32,2 vs 52.6 $\pm$ 31.8%). When confounding variables were controlled in a multivatiate regression model (70.027). Zscores at birth and simmetric or severe IIIGM were not beginning of min requirement for the first sector (37.4) and (37.

Conclusion: In our population IUGR did not influenced early feeding tolerance of VLBW infants