Abstracts

**763** VITAMIN B12 AND FOLATE DEFICIENCY IN HEALTHY NEONATES AND THEIR MOTHERS

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F Guven, 1A Say, 1N Uygur Kulcu, 1S Nepesov, 1M Inalhan, 1S Degirmenci, 1TA Sabuncu. 1Zeynep Kamil Maternity and Children’s Disease Training and Research State Hospital, Istanbul, Turkey

**Background and Aim** We aimed to determine the incidence of vitamin B12 and folate deficiency in healthy neonates and their mothers; to show the correlation between maternal and neonatal B12 and folate levels.

**Method** The study consisted of 99 healthy neonates and their mothers who gave birth in Zeynep Kamil Maternity and Children Training and Research Hospital, Istanbul. Hospital records were reviewed and blood samples were obtained from both mothers and their babies in the 24 hours after birth. Serum vitamin B12 and folate levels were measured. SPSS 15.0 package program was used for statistical analysis.

**Results** The mean values were as follows: in mothers serum vitamin B12 level 180.84± 58.81 pg/dl and folate level 17.61±20.61 nmol/L; in neonates vitamin B12: 320.43±187.48 pg/dl and folate: 26.05±34.45 nmol/L. There was significant correlation between maternal and neonatal folate levels but a slight correlation between maternal and neonatal B12 levels. 97% of mothers and 25.3% of neonates had B12 deficiency when the threshold level is defined as 300 pg/ml for mothers and 200 pg/dl for neonates.

**Conclusion** Due to high vitamin B12 deficiency rate in mothers a universal B12 vitamin screening program and preventative measures of B12 deficiency in pregnant women can reduce the potential complications of vitamin B12 deficiency in mothers and their siblings.

**764** A STUDY ON THE OUTCOME OF DIRECT COOMBS TEST (DCT) POSITIVE NEONATES

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KM Upatissa, J Greenaway, S Mahadevan-Bava. Paediatrics, Russelle Hall Hospital, Dudley, UK

The management and follow up of neonates with DCT+ differ from one centre to another. At RHH, guidelines recommend twice weekly FBC and Bilirubin in the first 2 weeks, and review at 4–6weeks.

We retrospectively reviewed 52 DCT+ neonates born at RHH between May 2008 and June 2009 to assess their outcome and to evaluate the need to review the current guideline. We divided the sample into three groups. Group 1: Rh isoimmunisation due to RhD (39/52), 2: other Rh and non Rh alloimmunization (6/52) and 3: with ABO incompatibility (7/52).

In group 1, 38/39 mothers received anti-D prophylaxis (RAADP). 20/39 had weakly positive and 19/39 had moderately DCT+. None of these neonates developed anaemia or prolonged jaundice.

In group 2(n=6), 3/6 were strongly DCT+ and 2/3 developed jaundice < 24 hours requiring phototherapy. 3/6 showed moderately DCT+ and 2 required phototherapy in the first week. At 2 weeks 4/6 developed anaemia but did not warrant transfusion.

In group 3(n=7), 6 showed moderate DCT+ and 1 weakly DCT+. 5/7 in group 3 received phototherapy in the first week. 5/7 in group 2 developed prolonged jaundice. 1/7 at 2 weeks showed anaemia.

We conclude that

1. Neonates born to mothers who received RAPPD do not show significant post delivery haemolysis and may not require close monitoring
2. Those with non RhD alloimmunisation and ABO incompatibility require monitoring for haemolysis at 2 weeks and review at 6 weeks
3. Although small number in our study, strong positive DCT warrants close monitoring in the first 24 hours of age.

**765** LINEAR GROWTH AND CIRCULATING IGF-I CONCENTRATIONS IN CHILDREN WITH IRON DEFICIENCY ANEMIA AFTER TREATMENT

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A Soliman, M Eldabbagh, A Adel, A Sabt. Pediatrics, Hamad Medical Center, Doha, Qatar

**Objective** To assess linear growth of patients with Fe deficiency anemia (IDA) before and after in relation to their hematologic parameters and IGF-I concentration before and after treatment with iron.

**Methods** Forty children (aged 17.2 +/- 12.4) months with iron deficiency anemia were studied with 40 healthy normal age-matched children (controls). Patients were treated with iron syrup or drops to supply 6 mg/kg/day. Growth (weight, length and head-circumference) and hematological parameters were measured and IGF-I concentrations measured before and 3 and 6 months after treatment.

**Results** Growth parameters (weight, length and head-circumference) and hematologic parameters were studied for 6 months after iron therapy. At presentation, patients with IDA had low Hb (8.2 +/- 1.2 g/dl), hematocrit (29 +/- 2.3%), MCV (61.5 +/- 8.1), and MCH (19 +/- 3.2) which improved significantly after treatment to (11.2 +/- 1 g/dl), 70.6 +/- 6.8, 23.4 +/- 2.9 and 18.9 +/- 5 respectively. Before treatment children with iron deficiency they had length standard deviation score (LSDS) = -1.2 +/- 1, annualized growth velocity (GV) = 7.5 +/- -2.2, GV SDS = -1.42 +/- 0.6 and BMI = 13.5 +/- 1.2. After 6 months their LSDS = -0.6 +/- -0.9, annualized GV=13.2 +/- -4.4 cm/year, GVSDS = 1.7 +/- -0.5, and BMI = 14.2 +/- 1.1. Circulating IGF-I increased significantly after treatment (52 +/- 18.8 ng/ml) vs before treatment (26.5 +/- 4.2 ng/ml).

**766** A REVIEW OF TOLERATION OF PNEUMOCYSTIS CARINII PNEUMONITIS (PCP) PROPHYLAXIS TREATMENT IN CHILDREN WITH ACUTE LYMPHOBlastic LEUKAEMIA

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PM O’Hare, A Moody, T Zerb, D Lancaster. Paediatric Oncology, Royal Marsden Hospital, London, UK

**Background and Aims** According to the UKALL2003 protocol, co-trimoxazole should be used as first line therapy to prevent PCP...