National Prescription Register. Cox proportional hazard regression models were used to calculate the hazard of miscarriage in women with a partner exposed to methotrexate. The study was approved by the Danish Data Protection Agency (2015-41-4309). **Results** We identified 1,364,063 registered pregnancies with known paternity, of whom 520 fathers were exposure to methotrexate within the three months before conception to the end of the first trimester. Among these, 46 (8.9%) experienced a miscarriage compared to 122,926 (9.0%) among the unexposed.

There was no increased risk of experiencing a miscarriage in pregnancies to men exposed to methotrexate before pregnancy compared to unexposed (adjusted hazard ratio 0.99 (CI95% 0.67- 1.46)). Furthermore, we found no increased risk of experiencing a miscarriage in pregnancies to men exposed to methotrexate during first trimester compared to unexposed (adjusted hazard ratio 0.90 (CI95% 0.61-1.32)).

Conclusion We found no association between paternal exposure to methotrexate before and during early pregnancy and miscarriage. Available data suggest that paternal methotrexate exposure should not be of major concern. Multinational recommendations could be changed accordingly.

Disclosure(s) Nothing to disclose

P07 ROXITHROMYCIN IN EARLY PREGNANCY AND THE RISK OF MAJOR CONGENITAL MALFORMATION: A REGISTER BASED NATIONWIDE COHORT STUDY

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Background Medicine use during pregnancy often causes concern for fetal harm. Roxithromycin, a macrolide antibiotic, is regarded as inadvisable to use during pregnancy due to lack of safety data. However, alternative macrolides have been associated with adverse outcomes in pregnancy. We conducted a register-based nationwide cohort study testing the hypothesis that use of roxithromycin in the first trimester is associated with major congenital malformations.

Methods We included all Danish women giving live birth from 1997 to 2012. Women with at least one redeemed receipt of roxithromycin during first trimester were regarded as exposed. Multivariable logistic regression adjusting for maternal age, multiple birth, parity, year of conception, smoking, educational length, and household income was performed, supplemented by sensitivity analyses comparing unexposed with exposure to increasing accumulated doses of roxithromycin.

Results The study included 966,372 pregnancies of which 2,430 children were born to an exposed mother, 78 (3.34%) of the exposed children were diagnosed with a major congenital malformation compared with 33,609 (3.49%) among children born to unexposed mothers. The odds ratio for the occurrence of a major congenital malformation after exposure to roxithromycin was 0.96 (95% CI 0.76-1.20) and multivariationally adjusted 0.94 (0.74-1.18). Sensitivity analyses comparing unexposed with exposure to increasing accumulated doses of roxithromycin showed no dose response relationship. Further, no differences in the type of major malformation according to the EUROCAT subgrouping system were seen.

Conclusions We found no association between exposure to roxithromycin in the first trimester of pregnancy and major congenital malformations.

Disclosure(s) Nothing to disclose

P08 IS THERE A DIFFERENCE IN THE FREQUENCY OF MANIPULATED ORALLY AND RECTALLY ADMINISTERED MEDICINES TO PAEDIATRIC IN-PATIENTS IN SWEDEN IN THE YEAR 2009 AND 2018 RESPECTIVELY? A REGISTER STUDY

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Background Since there is a lack of drugs in suitable strengths and child-friendly dosage forms, manipulation is sometimes necessary in paediatrics. A manipulation is the physical alteration of a drug dosage form with the purpose to extract and administer the prescribed proportion of a drug dose.

The purpose of this study was to calculate the frequency of manipulated medicines administered to paediatric in-patients at a large Children's Hospital for two separate years and compare whether there has been a change in practice.

Material and Methods Data were collected for all administered doses during 2 separate years (2009 and 2018) at the paediatric wards at our Children's Hospital, from a hospital-based electronic register. All administered doses where the number of tablets or suppositories were decimal were added and calculated as a percentage of the total number of oral and rectal administrations. Data are anonymous but information regarding gender, age, hospital ward and number of drugs per patient were available and were analysed.

Results In one year, approximately 450 000 doses of medicine are administered to paediatric patients in our Children's hospital.

The results will be analysed with regards to differences between patient age, gender, prescribing year and drug substance. A pilot study showed that 10% of all solid oral administrations to patients 6 - 12 years old, were part of a tablet. For patients 0 - 2 years over 20% of all solid rectal administrations were part of a suppository.

The extent of manipulation is affected by a lot of factors, where the most prominent is whether there are strengths suitable for that age-group available on the market or not.

Conclusion Most often there is a lack of knowledge how manipulation of medicines influences the dosing accuracy and often we do this to our most vulnerable patients.

Disclosure(s) Nothing to disclose

P09 SERUM NEUROFILAMENT LIGHT CHAIN LEVELS ARE AN INDEPENDENT PREDICTOR OF NEURODEVELOPMENTAL OUTCOME IN PRETERM INFANTS

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Background and Purpose Serum neurofilament light chain (sNfL) has recently emerged as a promising biomarker reflecting structural neuro-axonal damage in different neurological diseases. Our study aimed at assessing whether sNfL can predict the functional outcome in preterm infants who suffered from neonatal haemorrhagic brain injury.

Methods In this prospective observational study, we used an ultrasensitive single-molecule array assay to measure serum and cerebrospinal fluid (CSF) concentrations of NfL in preterm infants diagnosed with intraventricular haemorrhage (IVH) in the first few days of life. We determined the temporal profile of serum and CSF NfL levels from first diagnosis of IVH until term equivalent age, their association with cerebral imaging markers, and with clinical and functional outcome until 2 years of age assessed by Bayley Scales of Infant Development. We fitted univariable and multivariable logistic regression models to determine risk factors for low motor and cognitive development. Longitudinal mixed effects models modelled NfL levels using cubic spline smoothers to track the trajectory over time.

Results The study included 48 infants born with less than 32 weeks of gestation. At the time point of IVH diagnosis, NfL median levels were 271.9 pg/mL (IQR 151.2–389.7), and strongly decreased until term equivalent age to 15.7 pg/mL (IQR 11.1–32.2). CSF values were 113-fold higher (IQR 40–211) than corresponding serum values. Additional cerebral infarction (n=23) but not post-haemorrhagic hydrocephalus with permanent external ventricular drainage (n=29) or other diseases independently determined sNfL levels. In multivariate logistic regression models, the only significant predictor of poor motor outcome at 2 years or death was sNfL level (p=0.02). There was a clear difference between the NfL trajectory for those with poor motor outcome at 1 year.

Conclusions This study shows that early sNfL is an independent prognostic biomarker for motor functional outcome in preterm infants after IVH.

Disclosure(s) Nothing to disclose

P10 REPORTING OF OFFSPRING DATA IN DIABETES, HIV INFECTION AND HYPERTENSION TRIALS DURING PREGNANCY: A SYSTEMATIC REVIEW

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Background According to international guidelines clinical trials are conducted during pregnancy to evaluate benefits and risks of medicines for both the mother and her offspring. Consolidated Standards of Reporting Trials (CONSORT) reporting criteria apply to these trials and should include safety data on the offspring. The safety of maternal treatments is a key issue for health care professionals and parents. Diabetes, human immunodeficiency virus (HIV) infection and hypertension are among the most frequent chronic diseases worldwide in women of child bearing potential. The objective of this systematic review was to analyse offspring data reported in clinical trials conducted in pregnant women receiving chronic drug treatment for one of these conditions.

Methods Pubmed and Embase (01/01/1997–31/12/2017) were searched for drug trials in pregnant women with diabetes, HIV infection or hypertension. Titles and abstracts were screened, followed by a full text review of eligible articles. Inclusion criteria were interventional clinical trials in pregnant women treated with chronic medication and full text in English. Trial characteristics, maternal and offspring data were extracted. Data was summarised by disease and study. Twelve key items were considered for the offspring. The protocol was registered on PROSPERO (CRD42017057024).

Results Overall, 196 articles reporting 132 clinical trials (diabetes n=55; HIV n=59; hypertension n=18) were included. The number of births were frequently not reported (diabetes 40%; HIV 24%; hypertension 56%). Congenital malformations were infrequently reported with sufficient detail (diabetes 27%; HIV 34%; hypertension 6%). Similar observations were made for other key items (e.g. foetal losses, neonatal deaths, birth weight corrected for gestational age).

Conclusions Underreporting of key data for the offspring was frequent in publications of clinical trials in pregnant women with diabetes, HIV infection or hypertension making the assessment of the benefit-risk ratio of treatment options during pregnancy difficult.

Disclosure(s) Nothing to disclose

P11 NEONATAL AND PAEDIATRIC PROTOCOL DEVELOPMENT AND DRUG SAFETY: POINTS TO CONSIDER FOR RISK MANAGEMENT AND SAFETY DATA COLLECTION

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Background Protocol development for neonatal or paediatric clinical trials needs to take into account the age group specifics of the study population (e.g. pharmacokinetics, reference values for laboratory data and vital signs). Drug safety and risk management for neonatal/paediatric trials require an understanding of how these change throughout childhood. We were interested in reviewing and summarising the literature to identify publications which provide researchers with practical information of how the neonatal/paediatric drug safety profile informs age group specific safety data collection and risk management in the protocol.

Methods Pubmed, Embase and regulatory authority (RA) websites were searched for publications up to 31/12/2018 for children (0–18 years). In addition, the bibliography of included publications was reviewed to identify additional publications.

Results RA websites provided general and disease specific guidance on neonatal/paediatric clinical trials with sections relating to drug safety. No publication was identified describing the practicalities of how the neonatal/paediatric drug safety profile can be included throughout the various sections of a clinical trial protocol. The existing literature was summarised providing an overview of how the neonatal/paediatric drug safety profile supports the development of the various protocol sections. For example laboratory values in the exclusion criteria and safety monitoring sections need to be adjusted for age. Vital sign and psychomotor assessment should be done at least at baseline, trial completion and follow-up. Monitoring of