increasing interest in administration by continuous infusion in adult studies, but paediatric data is lacking. A continuous infusion, if given via an elastomeric pump could also facilitate earlier discharge from hospital.

**Methods**

Children with an oncology/haematology diagnosis admitted to Alder Hey Children’s Hospital with febrile neutropenia, normally treated with Piperacillin/Tazobactam via elastomeric device were considered eligible for the study. Patients received 24–36 hours of intermittent dosing before continuous dosing commenced via elastomeric pump. We analysed the data from 5 patients with double lumen central lines as a pilot phase to determine if expanded recruitment to include patients with single lumen lines was possible.

**Results**

Five patients were recruited, four of which had the continuous infusion. The mean Cmax following intermittent dosing of piperacillin and tazobactam from the lumen used for drug administration were 189.7 mg/L, 95% CI [71.7 - 307.9] and 18.5 mg/L, 95% CI [10.7, 26.3] respectively. The mean Cmax following intermittent dosing for the ‘empty’ lumen were 160.5 mg/L, 95% CI [121.6 - 199.4] for piperacillin and 13.7 mg/L, 95% CI [11.2 - 16.3] for tazobactam. The largest difference seen was on patient 002 with a concentration of piperacillin almost double that seen in the ‘empty’ lumen (429 mg/L vs 233 mg/L).

**Conclusion**

The lumen used for drug administration has enough residual drug to influence the results, so expanding recruitment to include single lumen lines is not going to be undertaken. The study has been amended to recognise this, and recruitment will continue with double lumen central lines only.

**Disclosure(s)**

Nothing to disclose

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**P70**

**IDENTIFYING AND MANAGING PROBLEMATIC POLYPHARMACY IN CHILDREN AND YOUNG PEOPLE?**

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**Background**

Polypharmacy may be necessary in a patient with complex disease or multiple illnesses. Problematic polypharmacy (PP) is defined as the prescribing of multiple medications inappropriately, or where the intended benefit of the medication is not realised. Identification and management of PP already occurs in adult medicine, with evidence based guidelines to manage potential PP. No specific deprescribing guideline or tools to guide management were identified. Paediatric clinical pharmacology is well placed to create and implement such guidelines.

**Disclosure(s)**

Nothing to disclose

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**P71**

**POPULATION PHARMACOKINETICS AND PHARMACOGENETICS OF CAFFEINE IN CHINESE PREMATURE INFANTS WITH APNOEA OF PREMATURITY**

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**Background**

Caffeine is commonly regarded as the treatment of choice for neonatal apnoea. However, limited data on the developmental pharmacokinetics and pharmacogenetics were available in Chinese premature infants. The objective of this study was to develop a population pharmacokinetic model of caffeine after intravenous administration to Chinese neonates with apnoea of prematurity (AOP) and evaluate the impact of developmental pharmacogenetics of CYP1A2.

**Methods**

Sparse pharmacokinetic samples were collected from AOP newborns receiving caffeine citrate at a loading dose of 20 mg/kg/d and maintenance dose of 5–10 mg/kg/d. Population pharmacokinetic-pharmacogenetic analysis of caffeine was performed using NONMEM. Eight CYP1A2 polymorphisms were genotyped.

**Results**

A total of 99 newborns with a mean (SD) postmenstrual age of 32.0 (2.16) (range 22.3 - 38.0) weeks were included in the present study. Pharmacokinetic data fitted an one-compartment model with first-order absorption and elimination. Current weight, postmenstrual age and serum creatinine concentration were significant covariates influencing caffeine clearance. None of tested CYP1A2 polymorphisms had significant impact on caffeine pharmacokinetics.

**Conclusion**

The population pharmacokinetics-pharmacogenetics of caffeine was evaluated in Chinese AOP premature infants. This developmental pharmacokinetic model can be helpful to individualize caffeine therapy.

**Disclosure(s)**

Nothing to disclose