present during hospitalisation. Twenty five questionnaires have been collected (response rate 53%). Among patients there were 18 infants and toddlers, 3 children and 4 adolescents. Mean length of stay in the PICU was 11 days, 60% of admissions were unplanned. Questionnaires were completed mainly by mothers (84%). Not satisfactory opinions have been given mainly for understandable information on examinations and tests (12%) and on possibility to stay close to the child during intensive procedures (16%). All of parents declared that the team worked efficiently and the team showed respect for the patients but only 72% of parents responded that during stay in the PICU the staff regularly asked for parent’s experiences.

Conclusions The EMPATHIC-30 empowers parents to provide feedback on their experiences in paediatric intensive care and may facilitate health care professionals to improve quality of care. Following a single centre experience the EMPATHIC 30 Poland study should be continued as a national project.

**PO-0275** ASSESSMENT AND COMPARISON OF A LAB-SCORE AND A CLINICAL PREDICTION MODEL FOR DETECTING SERIOUS BACTERIAL INFECTIONS IN FEBRILE YOUNG CHILDREN

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**Background and aims** C-reactive protein and Procalcitonin have been lately the most researched biomarkers in identifying serious bacterial infections (SBI) in febrile children. The Lab-score (2008) includes CRP, PCT and urinalysis for detecting SBI and the Clinical Prediction Model (CPM) (2013) combines clinical variables with CRP value for detecting pneumonia and other SBI separately. We aimed to assess and compare the value of the Lab-score and the CPM in identifying febrile children at risk for SBI in the Emergency Department (ED).

**Method** This survey is part of a prospective observational study aimed to identify children with fever without source at risk for SBI. Patients were recruited from Tîrgu Mures Emergency Clinical County Hospital, Romania. SBI diagnosis was based on urine, blood and CSF cultures and chest radiographs. For children included, aged 1 to 36 months, the Lab-score and the CPM were calculated. Positive and negative likelihood ratios and post test probabilities were calculated for each test.

**Results** From 134 children, SBI was diagnosed in 31 (23.13%): 11 pneumonia and 20 other SBI, mostly urinary tract infections. Positive and negative likelihood ratios for Lab-score (≥3), CPM-Pneumonia (≥10%) and CPM-Other SBI (≥10%) were 7,25/0,25, 22/0,65 and 5,23/0,50 and the post test probabilities were 69%, 66% and 48% for the same cut-off values.

**Conclusions** Both the Lab-score and CPM-Pneumonia are valuable tools in detecting SBI in febrile young children. CPM-Other SBI showed less performance than Lab-score and CPM-Pneumonia, possibly due to the lack of urinalysis value in CPM-Other SBI, which are mostly UTI.

**PO-0276** FEVERISH CHILDREN IN A DGH IN NORTHERN IRELAND – WHAT ARE WE DOING?

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**Aims** We undertook this audit to review the management of febrile children in our emergency departments (ED) compared to The College of Emergency Medicine (CEM) standards.

**Methods** The data was collected using a tool designed by CEM. Entry criteria: under 5 years old and temp >38°C on arrival.

**Results** Total number of patients was 50. The assessed risk profile for this population (using NICE guidelines) were 24 low risk, 14 intermediate risk, 11 high risk and 1 we were unable to risk stratify from the clinical notes. Nine children were pre-scribed antibiotics (5 low risk, 2 intermediate and 2 high risk).

Of the 11 patients who were high risk, 7 had a clear source of infection. Of the 4 who had no source identified, one had bloods and urine performed in ED but these were not recorded in the notes. 2 had bloods performed on the paediatric ward.

For the 14 patients in the intermediate risk, 8 had a source of infection, 5 had no obvious source identified and one was not clearly documented. No patients without a source were prescribed antibiotics. No documentation was recorded about discharge advice.

18 patients (36%) did not have a blood pressure (BP) or a capillary refill time (CRT) documented in the notes and 10 patients (20%) did not have their GCS or AVPU recorded.

**Conclusions** There are areas that require review. Improvements must be made to ensure a full set of observations are recorded, emphasising the importance of BP/CRT as well as GCS/AVPU.

**PO-0277** REVIEW OF THE MANAGEMENT AND OUTCOME FOR PATIENTS TREATED FOR WHEEZE IN A TERTIARY PAEDIATRIC EMERGENCY DEPARTMENT (PED)

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**Background and aims** To review the management of patients who were treated for wheeze in the PED and compare them to those in the British Thoracic Society (BTS) Guidelines.

**Methods** Reviewed PED files of those patients over 2 who were treated for wheeze in PED in March.

**Results** Number of patients 46.

93% received bronchodilators while 7% were given Prednisolone only. 65% were given maximum nebulised bronchodilator therapy (x3 sets). 89% received steroids and 2% received IV medication. 41% received antibiotics in ED and 50% had a chest x-ray performed.

Of those discharged (n = 32), 75% were discharged within 4 h, 31% patients were discharged within 1 h of last bronchodilator and a further 41% within 2 h.

Of those treated with maximum nebulised bronchodilators (n = 30), 43% were admitted, 23% discharged within 1 h post last bronchodilator and a further 27% within 2 h.

8% re-attended within 48 h -4% due to elevated temperature, 2% were uncertain with inhaler technique and 2% due to increase in symptoms.

No patient had documented evidence of written asthma management plan given or to attend primary care for review within next 48 hrs.

**Conclusions** The low re-attendance rate is supportive that those attending received good clinical care and in a timely manner, with 75% of patients being discharged within the 4 h target. However, education is required to ensure patients stay 3-4 h post last bronchodilator and the need for documented discharge.
MANAGEMENT OF PAIN IN ACUTE PRESENTATIONS TO A TERTIARY PEDIATRIC EMERGENCY DEPARTMENT (PED)

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Background and aims To review pain assessment and management in our PED.

Methods We reviewed the films for patients who were coded as soft tissue injury, fracture or burn over a 6 day period in June 2013 and compared this to standards set by College of Emergency Medicine (CEM) in the UK.

Results Number of patients = 67
98% did not have a pain score recorded from triage. No recorded pain score from any medical personnel.

49% received analgesia with 82% receiving paracetamol alone and 15% receiving oramorph. Of those receiving analgesia, 70% did so within 20 min of arrival and 85% within first hour.

There was no documented re-assessment of pain scores although 6% of patients did receive further analgesia.

Conclusion The results hi-lighted a need for re-education of nursing and medical staff on the benefits of pain scores. Coupled with this re-education there will be a review of the current PED filmsy with a greater emphasis on pain scales, pain scores and prompts to re-score.

SKIN CONDUCTANCE CHANGES DIFFER BETWEEN PAINFUL STIMULI AND GENERAL HYPOXIA DIFFERENT FROM PERIPHERAL OXYGEN SATURATION

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Background and aims Peripheral oxygen saturation (SpO2) decreases during general hypoxia and painful events in preterm infants. Skin conductance responses per sec (SCR/sec) increase during painful procedures. The purpose of this observational study was to examine if SCR/sec can help to diagnose if SpO2 is due to general hypoxia or painful events.

Methods Ten infants, diagnosed as ventilator unstable with birth weight 1248 (± 710) grams, gestational age 30 (± 5) weeks and at postnatal day 3 (± 1) were observed for 1 h when venous blood sampling was performed. SpO2, SCR/sec, heart rate (HR), and respiratory rate (RR) were recorded each 2nd minute and during the painful and hypoxic events (defined as SpO2 lower than 80%). The variables were studied during the painful and hypoxic events as well as in situations without events. Non-parametric tests for within individual variables (Friedman’s Anova) were used.

Results There were statistical differences for physiological measures during the no-painful- and hypoxic events; mean (SD): SCR/sec does not increase during SpO2 and may be used to differ between pain and general hypoxia in preterm infants.

WITHDRAWN

ETHICAL AND CLINICAL IMPLICATIONS RELATED TO EMERGENCE PHENOMENA IN PEDIATRIC KETAMINE SEDATION

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Background and aims Ketamine is an efficient, economical dissociative sedative used during painful procedures as an alternative to general anaesthesia. Its physiological risk profile has been thoroughly researched; however, less well understood are the psychological events (‘emergence phenomena’) associated with its use. In adult practice, ketamine use is declining given the traumatic effects of emergence phenomena on patients, family, and staff. In paediatric practice, however, ketamine use is increasing despite a limited understanding of the occurrence of paediatric emergence phenomena and it’s associated potential for non-physiological harm. This study analyses the ethical implications of paediatric ketamine sedation by exploring healthcare practitioners’ (HCPs) accounts of paediatric emergence phenomena.

Methods HCPs (doctors, nurses, paramedics, and a play specialist) were interviewed about their experiences with emergence phenomena during paediatric ketamine sedation. Interviews were then analysed using a descriptive exploratory approach underpinned by hermeneutic narrative methods to identify common themes.

Results A focus on physiological risk has resulted in a trial-and-error approach to paediatric ketamine sedation practice. Understandings of non-physiological or psychological risk and the potential negative longitudinal outcomes in paediatric populations remain vague. “Dream seeding” (guided imagery) is widely yet inconsistently used as an anxiolytic for reducing negative psychotropic events, despite only anecdotal evidence of its effectiveness.

Conclusions Although ketamine sedation can help protect children from the pain of treatment and HCPs from emotional distress of delivering that treatment, there is potential for non-physiological harm. The practice of dream seeding should be investigated as a tool to mitigate adverse psychotropic events.