Results Operative incised abdominal wall displayed profound allodynia which was reduced by ropivacaine with low dose ketamine combination in the 4 hours following incision. Blood samples these patients showed enhanced levels of 3 cytokines: IL-1β, IL-6, tumor necrosis factor alpha (TNFα). Ropivacaine with low dose ketamine administration reduced levels. First group lower cytokines levels over second group (mean ± SD, IL-1β - 4.4 ± 2.2 vs. 14.2 ± 2.4 pg/mg protein; IL-6 - 204.8 ± 80.0 vs. 441.2 ± 90.4 pg/mg protein; TNFα - 14.4 ± 4.6 vs. 58.2 ± 7.2 pg/mg) (p<0.001).

Conclusion Ropivacaine with low dose ketamine administration reduces cytokine expression. These studies suggest that Ropivacaine with low dose ketamine combination may alter the inflammatory reaction.

1619 IDENTIFICATION OF NOXIOUS EVENTS FOR NEWBORN INFANTS WITH A NEURAL NETWORK
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Background Recognition of pain experienced by immature and/or critically ill newborns in the Neonatal Intensive Care Unit remains a challenge despite the use of objective scoring systems that depend on physiological and behavioural parameters. We consider there is a need to identify pain using only physiological data streams.

Methods Data were collected from three preterm male, gestational age 27.25±0.95 weeks (mean±SD), birth weight 941.25±189.31 grams. Heart rate (HR), mean arterial pressure (MAP), respiratory rate (RR), blood oxygen saturation (SpO2) were considered for the NN Input Vector. NN’s output were set to ‘1’ for noxious stimuli pattern (NSP) defined in 100%. Event ‘routine care’ coincided with the NSP defined in 100%. Event ‘breakthrough analgesia’ identified by NN with the artifact nociceptive event.

Results Noxious events were captured in previous study and integrated with real-time physiological data streams. In this study we correlated the nociceptive event identified by NN with the artifact nociceptive event. These studies suggest that Ropivacaine with low dose ketamine combination may alter the inflammatory reaction.

Introduction Although children with painful sickle cell crises (PSCC) frequently present to the Emergency Department (ED), pain in sickle cell disease is often under-recognised, under-treated and treatment may be delayed. We aimed to evaluate pain assessment and management in children presenting to the ED with PSCC.

Methods A 12-month prospective descriptive study of acute pain management of PSCC at an urban tertiary paediatric ED. Pain was assessed by the triage nurse or physician using a validated age appropriate pain scale (Faces, Legs, Activity, Cry, Consolability (FLACC) Scale; Manchester Pain Ruler).

Results There were 96 presentations in 66 patients with PSCC (Table 1). Nineteen (19.7%) patients received no pre-hospital analgesia.

Abstract 1619 Table 1

<table>
<thead>
<tr>
<th></th>
<th>Entire Cohort (n=56)</th>
<th>Severe pain Cohort* (n=30)</th>
<th>Moderate Cohort* (n=30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Triage Pain Score</td>
<td>7/10 (IQR 5–8)</td>
<td>8/10 (IQR 7–10)</td>
<td></td>
</tr>
<tr>
<td>Pain Score at 60 minutes</td>
<td>5/10 (IQR 2.25–6)</td>
<td>7/10 (IQR 5–8)</td>
<td></td>
</tr>
<tr>
<td>Cases in line with PED analgesia guidelines (%)</td>
<td>45%</td>
<td>50%</td>
<td>95%</td>
</tr>
<tr>
<td>Median time for opioid breakthrough/analgesia (min)</td>
<td>87 minutes</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*severe pain defined as ≥ 7/10 and moderate pain as 3–6 on age-appropriate pain scale

Conclusion PSCC pain is under-treated, under-monitored and adequate treatment of pain is delayed in our ED. Patients with severe pain appear at highest risk for treatment guideline violation. This is predominantly related to lack of opiate administration. An educational intervention, with/without the inclusion of an easily administered, fast-onset and short-acting opiate e.g. intranasal fentanyl, may decrease the time from ED arrival to effective pain relief.

1620 EMERGENCY ANALGESIA ADMINISTRATION IN CHILDREN: RETROSPECTIVE ANALYSIS AND RECOMMENDATIONS
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Introduction Emergency analgesia administration in children is inadequate and guideline is insufficient. We aimed to analyse our department’s paediatric pain management to inform and recommend necessary alterations to current practice.

Methods 800 children (0–16 years old) presenting with painful conditions to Queen Elizabeth Hospital Emergency Department within a 40-month period (01/01/2008–28/02/2012) were randomly identified from a prospective audit database and allocated into four groups according to pain scores (no, mild, moderate and severe pain; 200 children in each group). Analgesia types and differential diagnoses were recorded.