groups will feed back on their progress at the monthly meetings to encourage action and momentum.

Measurement of improvement We have now completed departmental audits that are up to date and shared with the team. Our monthly meeting has a full agenda promoting and sharing best practice.

Effects of changes These changes have allowed the department to move to a more cohesive and coherent view of their work around quality. We have helped to standardise care, share good practice and maintain a quality service for the future. The staff wellbeing group has started a free fruit and lunchtime walking initiative. While the patient voice group have raised the profile of recording this in clinic letters and meetings.

Lessons learnt One of the greatest challenges was getting colleagues to prioritise “quality” work above their clinical case load. In a busy department it can be hard find time to implement change. The quality day provided a different environment away from clinical duties to allow people to think about what we could do better.

Message for others Identifying “quality” and the evidence required to show this is the first step. Forming a central easily accessed folder containing all the information along with an index or database to assess progress is the next. Using the CQC standards as a template for our files helped us to organise our thinking in line with national standards. Engaging colleagues and encouraging personal responsibility to support ongoing improvement is also essential.

**Abstracts**

**LOOKED AFTER CHILDREN AT RISK OF BLOOD-BORNE INFECTIONS: A QUALITY IMPROVEMENT AUDIT**

P Mikrou, G Cropp, V Sadavarte; Child Development Centre, University Hospital North Midlands, Stoke-on-Trent, UK

We assessed our service provision through an audit, basing our standards on the 2008 BAAF guidance. This was a retrospective audit from June 2013 to June 2014. In total 212 children attended the specialist LAC clinic. A risk assessment was carried out, based on information about parental health and lifestyle and the results of antenatal screening for Hepatitis B, Syphilis and HIV. Hepatitis C results were only available in high risk population (IV drug users). 37 children (17%) were identified as needing a BBI screen. Out of these, only 22 children (60%) were screened. 8 out of the 22 children (36%) had a complete screen (including Hepatitis B/C and HIV); with the remaining having a partial screen. 12 children had Hepatitis C positive mothers. Worryingly, only 8 of those 12 children (66%) had a BBI screen. There were no Hepatitis B or HIV positive mothers. Reasons for not having a BBI screen were difficulties in obtaining consent, failure to identify the children at risk or to get the extended information about parental lifestyle and screening results. BBI screen revealed 2 children positive for Hepatitis C antibodies and appropriate follow-up was arranged. No children had a BBI screen when that was not indicated.

We subsequently developed a protocol in the form of two flowcharts. These will be included in the LAC health assessment paperwork and aim to promote clarity and good clinical practice. As failure to obtain consent played an important hindering factor in getting our vulnerable population screened, we suggested, when possible, consent is taken at the time of consultation. Improved communication and information sharing between Health and Social Care is essential. Finally, team education is greatly important and will be reinforced by the integration of BBI risk assessment to the induction programme of the new community trainees.

The feedback has so far been very positive. We strongly believe it promotes good clinical practice. We plan to implement this in March 2015 and we aim to maintain and reinforce those changes by continuous monitoring and evaluation of our service.

Looked After Children are, sadly, a growing population in our society. Their health promotion and safeguarding is a responsibility of both Social Care and Health Authority. It is crucial that we, as health professionals, constantly strive to offer a high quality service, by enhancing clinical enquiries and audits, supporting changes to practice and implementing those for improved patient outcomes and experiences.

**IMPROVING RECORDING OF POSTNATAL WARD NEONATAL OBSERVATIONS**

K Parkinson, Y Lim, A Demetriou, J Ziprin. Department of Paediatrics and Neonatal Medicine, Imperial College Healthcare NHS Trust, London, UK

Context This audit was carried out on two postnatal wards at different sites within the same trust.

Problem Whilst working on the postnatal wards we observed that the observations that babies required (e.g. after meconium delivery) weren’t being carried out according to the frequency specified in the guidelines. Therefore causing a negative impact on patients due to the potential for missing a deterioration if observations were not carried out appropriately.

Assessment of problem and analysis of its causes We found on average 41% of the recommended frequency of observations were carried out on site 1 and 52% on site 2. More than 50% of the recommended observations were not done in the following categories; maternal GBS, phototherapy and hypoglycaemia in site 1 and in site 2; hypoglycaemia, and 33–35+6/40 gestation.

It was felt a possible reason why such a low percentage of observations were completed was due to lack of awareness of the frequency specified in the guidelines. Additionally at site 1 there is no set place to record observations whereas at site 2 separate charts are used. This may explain the higher percentage of recorded observations at site 2.