structured approach to the history and examinations of a child with possible tic disorder and when investigations are indicated. Management is discussed under two categories of (a) Simple tics and (b) Tics with co-morbidities. Advice is given on who will benefit from psychology input and what to do when co-morbidities are noted. Indications for paediatric neurology and CAHMS referral are discussed. The Appendix includes community referral forms, the Yale tic questionnaire and some links to information for carers.

**Conclusion**
The authors are hopeful this guideline has fulfilled the aims outlined. Consideration will be given to seeking feedback from users when it is formally adopted. Paediatricians from other Trusts may also find much of the guidance helpful though caution would be needed with the referral pathways which are different in different Trusts.

**REFERENCES**

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**P149 IS VIRTUAL THE FUTURE OF OPD?**

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**Introduction**
The General Paediatric Outpatients Department at Our Lady’s Children’s Hospital Crumlin has a greater than two year waiting list for new routine appointments. Long waiting time is considered as one of the most concerning matters in the majority of health care organisations.

A wait time of no more than 6 months for routine appointments is proposed. A virtual clinic was run by three General paediatricians from November 2017 to October 2018, with the aim to reassess and prioritise the ‘long-waiters’ on the waiting list. A Virtual clinic is a contact between the clinical team and the patient, without a face to face meeting, to plan clinical care with the use of telecommunication.

**Methods**
Parents of the longest waiting children were called by the paediatricians, after going through their referral letters. If parents were not contactable, their GP’s were contacted. Progression of the symptoms, persistent concerns, any reviews in the meantime were recorded. Subsequently patients were either removed from the list with parental consent or remained on the list with or without prioritisation.

**Results**
A total of 65 patients were reviewed via virtual clinics. 43 patients were subsequently removed from the list with parental consent. In 15 (23%) referring reason was resolved, 4 (6%) were seen by other hospitals and 2 (3%) were seen privately, while 7 (11%) were reviewed by other specialties. 22 patients remained on the waiting list. Among these 65, 41 (63%) parents were contactable, GPs were contacted in 9 (14%) cases while no one was contactable in 15 (23%) cases.

**Conclusion**
Virtual clinics can play an important role in dealing with the long waiting lists in paediatric population.

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**P150 PRIME (PRETERM INFANTS NEED MILK EARLY): A QUALITY IMPROVEMENT INITIATIVE IN A TERTIARY NEONATAL UNIT**

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**Aim**
Maternal milk (MM) protects against necrotizing enterocolitis and sepsis. PRIME is a multi-disciplinary initiative to improve the early provision of MM for preterm infants and enhance outcomes. Our aim was to increase the number of high-risk infants receiving MM in the first day of life in our tertiary neonatal unit.

**Method**
We retrospectively reviewed time to first MM for infants born <32 weeks gestational age (GA) or with a birth weight (BW) <1500 g in 2016. We conducted a cross-sectional survey to evaluate the knowledge and attitudes of staff towards breast milk for preterm infants. Deficits in background knowledge and training informed a teaching programme. Education involved training sessions, development of guidelines, distribution of posters, and presentation at meetings. The first 5 infants per month born <32 weeks GA or BW <1500 g were included in the post-intervention analysis. The effect of the interventions were evaluated using a before and after study design. Time to first MM was our key performance indicator.

**Results**
We reviewed 123 infants born in 2016 [Median (IQR) GA 29 (26, 31) weeks, BW 1140 (820, 1410) g]. Many infants didn’t start feeds in 1st24 hours as MM was not available; median (IQR) time to 1st MM 35 (17, 53) hours, 34% of infants received MM in the 1st24 hours of life. Prospective data was collected from 25 infants, born May – October 2018, following interventions [Median (IQR) GA 30 (26, 31) weeks, BW 997 (890, 1560) g]. The median (IQR) time to 1st MM was 17 (8, 25) hours and 76% of infants received MM in the 1st24 hours.

**Conclusion**
The initial results of this hospital-wide QI initiative are promising. The median time to first MM has halved in the 5 months since this initiative commenced. Further PDSA cycles are indicated to ensure ongoing improvement.

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**P151 ASSESSING THE USE OF THE ISBAR3 HANDOVER FRAMEWORK IN AN IRISH PAEDIATRIC HOSPITAL: A MEDICAL STUDENT PERSPECTIVE**

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**Intro**
Medical staff are routinely educated on the ISBAR3 framework for clinical handover. This tool is implemented to encourage safer and more effective task transfer during clinical emergencies, referral to other specialties, transfer of hospital and wards.
**Abstracts**

**P152 IMPLEMENTATION OF A BILIRUBIN MONITORING DOCUMENT IN THE NEONATAL ICU AND POSTNATAL WARD SETTING AT UNIVERSITY MATERNITY HOSPITAL LIMERICK**

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**Introduction** Transcutaneous (TcB) and serum bilirubin (SBR) levels are often monitored during the neonatal period in order to decide if a newborn requires phototherapy. Other details, however, are required to assess the infant’s risk factors and threshold of treatment\(^1\). Gestational age, ABO/Rh incompatibility and Direct Coombs test (DCT) are important amongst other risk factors\(^2\) to guide our management.

**Aim** To improve clarity and safety when recording TcB and SBR results by creating a one-page document encompassing all necessary pieces of information.

**Method** Our trial bilirubin monitoring document consisted of 3 sections. The first section is a standard Phototherapy nomogram\(^1\) followed by a section for details such as name, chart number, gestational age, date of birth, time of birth, DCT status and other risk factors. The third section consists of a table with columns for date, time of blood sample, age in hours, TcB, SBR, risk line, risk Zone\(^2\) and plan. Moreover, we included 8 rows for continual monitoring. Three months after implementation and education of this trial document, a survey was carried out amongst midwives and neonatal nurses in UMHL. Various questions were asked regarding the clarity and safety of the new document when compared with the old way of documenting. Feedback and comments were collected and analysed.

**Results** 24 individuals responded to the survey. Of these, 58% found the old way of documenting ‘unclear’ whereas 20% found it ‘very unclear’. 21% found the new document ‘clear’ and the remaining 79% found it ‘very clear’. 71% found the new document ‘easy to use’ and the remaining 29% found it ‘very easy to use’. 33% found the new document to be ‘safer’ and the remaining 67% found it ‘much safer’ to use when compared with the old way of documenting. Suggestions received included creating a column for the doctor’s signature next to the plan and also to include a line for the mother’s blood group in the second section.

**Discussion** Prior to implementation of this new document, all of these details were scattered throughout the chart making it difficult for another doctor to ascertain the information when it comes to interpreting a new bilirubin result for a particular newborn. Survey response was in favour of using our new document, due to its clarity, ease of use, enhanced safety and continuity of care. We plan to consider adding the suggested changes and finalising this document as a quality improvement initiative.