Improving the Haemoglobinopathy Screening Programme for High-Risk Patients in a Tertiary Maternity Hospital in Dublin, a Pilot Quality Improvement Initiative

Introduction
A more coordinated haemoglobinopathy screening is desired across European countries. A non-funded pilot screening programme was implemented in six Irish maternity units in 2005. It aims to identify patients with haemoglobinopathies such as sickle cell disease and thalassemia, by screening patients from high-risk ethnic origins.

As screening involves identifying disease carriers, ensuring informed consent is vital. Since the introduction of an electronic healthcare record in the Rotunda hospital in November 2017, patients no longer sign consent forms before blood samples are taken. Attendance rates at haematology clinics are poor, it is proposed that this is due to lack of education about screening and lack of early contact with patients from high-risk ethnic origins.

Methods
A standardised admissions document was created and introduced using the Plan Do Study Act (PDSA) Cycle. The document included all essential components of the paediatric history. Six essential criteria were identified for inclusion in the initial paediatric admission documentation. These included: Medication reconciliation, Social work involvement, Smoking in the household, Immunisation history, Birth History and Developmental History. The admission document was intended to be used by all junior paediatric and GP trainees in the clinical decisions unit for the admission of all patients under general paediatrics.

Results
Significant improvements were seen across all domains after the introduction of the admission document. A marked improvement in medication documentation was noted post-intervention. Complete documentation of medications rose from 45% pre-intervention, to 75% post intervention. Immunisation history documentation improved from 55% to 87.5%. Birth history documentation improved from 60% pre-intervention to 87.5% post intervention. The most significant improvements were observed in documentation of Social work involvement (15% pre-intervention to 82.5% post-intervention), Developmental history (22.5% pre-intervention to 95% post-intervention) and Household smoking history documentation (22.5% pre-intervention to 85% post-intervention).

Conclusion
The introduction of a standardised admissions document is a simple tool by which to improve care and standardise documentation of all paediatric admissions. Having a specific admissions document is shown to improve patient safety by reducing medication omissions by implementing medication reconciliation in the admissions document. A standardised document allows for a focused paediatric history encompassing all essential domains to be obtained for every child on their initial admission to the department, thereby reducing omissions of important medical information.

References

Abstracts
P155 Improving patient safety in a tertiary centre through the introduction of a standardised admission document for paediatric patients

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Background
There was a lack of uniformity in the initial medical admission documentation for patients admitted under general paediatrics at the Royal Hospital for Children, Glasgow. There was no standardised medical admission document used for admitting patients. Thus significant variability was seen in terms of quality of initial medical admission documentation and crucial elements of the paediatric history were frequently omitted. A method of improving and standardising care through improving documentation was sought.

Objective
To enhance patient safety through improved documentation of admissions under general paediatrics. A standardised admissions document was introduced with the aim of reducing medication omissions, improving medication reconciliation and standardising documentation, to include all crucial elements of the paediatric history in one document.

Methods
A standardised admissions document was created and introduced using the Plan Do Study Act (PDSA) Cycle. The document included all essential components of the paediatric history. Six essential criteria were identified for inclusion in the initial paediatric admission documentation. These included: Medication reconciliation, Social work involvement, Smoking in the household, Immunisation history, Birth History and Developmental History. The admission document was intended to be used by all junior paediatric and GP trainees in the clinical decisions unit for the admission of all patients under general paediatrics.

The notes of 40 patients pre-intervention and 40 patients post-intervention were reviewed. The notes were reviewed for documentation of the 6 essential criteria as defined above.

Results
Significant improvements were seen across all domains after the introduction of the admission document. A marked improvement in medication documentation was noted post-intervention. Complete documentation of medications rose from 45% pre-intervention, to 75% post intervention. Immunisation history documentation improved from 55% to 87.5%. Birth history documentation improved from 60% pre-intervention to 87.5% post intervention. The most significant improvements were observed in documentation of Social work involvement (15% pre-intervention to 82.5% post-intervention), Developmental history (22.5% pre-intervention to 95% post-intervention) and Household smoking history documentation (22.5% pre-intervention to 85% post-intervention).

Conclusion
The introduction of a standardised admissions document is a simple tool by which to improve care and standardise documentation of all paediatric admissions. Having a specific admissions document is shown to improve patient safety by reducing medication omissions by implementing medication reconciliation in the admissions document. A standardised document allows for a focused paediatric history encompassing all essential domains to be obtained for every child on their initial admission to the department, thereby reducing omissions of important medical information.