Pre-Transfusion Blood Sampling: Predicting Future Performance from Simulated Practice

Marc Feng, Simon Smith, Helen Hynes, Medical Education Unit, UCC



Rationale

• This study reviews the type and frequency of labelling errors, occurring when blood is ordered for Pre-transfusion sampling in a simulated teaching environment and compares it to reported data from real-world clinical settings.

Background

- The challenges to healthcare in reducing avoidable medical error have been well documented since the publication in 2000 of the Institute of Medicine's report 'To Err is Human' (1).
- National and international data shows that errors in pre-transfusion sampling are common (2,3) and may result in fatal errors (3)
- Mislabelled samples are up to 40 times more likely to contain blood from the wrong patient. (4)

 Anonymized data was collected from standard teaching sessions delivered in Final Medical year: the Procedural Skills Laboratory and the Simulated Ward.

- In the Simulated Ward students were asked by a nurse to take blood for Group and Hold from a manikin arm attached to a role player who was "experiencing a miscarriage".
- The students are told that this is a repeat sample, as the sample sent on admission was rejected due to incomplete data.



 In the Procedures Laboratory students were given full clinical / demographic details for a simulated case of post-partum haemorrhage and asked to take a sample for Group and Crossmatch of 4 units of blood from a manikin arm.



The students are given the following details:

- Pt's name, address, DOB, MRN, Ward, Consultant
- Clinical details:
 - \odot Group and Cross Match 4 Units
 - \odot Post partum haemorrhage
 - $\odot\,G^2P^1$
 - \circ Blood Group O –ve
 - \circ 1 miscarriage 2 years ago
 - \odot Had anti D after miscarriage
 - $\odot\,\text{No}$ known antibodies
 - \circ No previous transfusions

 Students were instructed to prepare the bottles and forms for the Blood Bank. These were examined by 2 researchers and coded for omissions and errors. A judgement was made on whether the samples would be processed based on current Pre-Transfusion sampling guidelines.⁵

• 223 samples were collected in total

Source	
Procedures Laboratory	207
Simulated Ward	16
Total	223

Errors	Procedures Laboratory	Simulated Ward
Incomplete or Missing Information on sample tube / request form	39	5
Mismatched Information between sample tube and request form	9	0
No Signature on sample tube and / or request form	17	2
Sample tube was Unlabelled	1	1
Illegible data on sample tube	25	0
Missing Clinical Data	66	3
No Reason Given for Request	32	4

Errors	Procedures Laboratory	Simulated Ward
Incomplete or Missing Information on sample tube / request form	39	5
Mismatched Information between sample tube and request form	9	0
No Signature on sample tube and / or request form	17	2
Sample tube was Unlabelled	1	1
Illegible data on sample tube	25	0
Missing Clinical Data	66	3
No Reason Given for Request	32	4

	Procedures Lab	Simulated Ward
Would be processed	145	11
Would not be processed	62 (31%)	5 (30%)
Total	207	16

- Error types in this study corresponds to the same reporting categories as the results from a National Survey in 2011
- Overall, no significant difference between the 2 types of teaching sessions

- Pre-transfusion Sampling in Ireland: Results of a National Survey 2011
- 71,314 pre-transfusion sample received in June, July and August 2011 from 41 hospitals in Ireland
- To evaluate sampling practice in Ireland
- On average, 4% of pre-transfusion samples rejected (up to 10.8% in some hospitals)

NHO REPORT 2010/2011



Reason sample rejected (n = 2922 samples rejected)	No of Samples
Incomplete or missing information from sample tube and/or request form	1181
Mismatched information between sample tube and request form	437
No signature on sample tube and/or request form	248
Addressograph label used on sample tube	194
Sample tube was unlabelled	62
Illegible data on sample tube	6
Other	746

- Similar findings with 2011 study and our data with exception of sample size
- Most common errors in both studies were due to incomplete and mismatched information
- According to the SHOT report, number of near-misses continue to increase
- "Wrong Blood in Tube" (WBIT) errors have the potential to result in ABOincompatible transfusions
- Majority of WBIT errors due to misidentification of patient and labelling sample away from bedside
- Highlights importance of strict adherence to pre-transfusion sampling guidelines

- Multidisciplinary training needed starting from medical school
- Majority of students agree that simulated practice is a great learning opportunity (6)
- More frequent simulated assessments on pre-transfusion sampling may be needed to further reduce sample rejection rate
- Use of 2-person dependent check (challenge and response) may provide additional safety for mislabeled samples
- Implementation of zero-tolerance approach
- Further studies should focus on why such errors occur in the clinical setting

References

- Kohn LT, Corrigan JM, Donaldson MS. To err is human: building a safer health system. Washington, DC: National Academies Press, 1999. https://www.iom.edu/Reports/1999/To-Err-is-Human-Building-A-Safer-Health- System.aspx
- 2. Cronin et al, National Haemovigilance Office Report, 2011, pages 24-28
- 3. SHOT annual report 2016, https://www.shotuk.org/shot-reports/
- 4. Lumadue et al. 1997 Adherence to a strict labelling policy decreases the incidence of erroneous blood grouping of blood specimens. Transfusion 37(11-12):1169-72
- Guidelines for the administration of blood and blood components. Irish Blood Transfusion Board. https://www.giveblood.ie/Clinical-

Services/Haemovigilance/Publications/Guidelines for the Administration of Blood and Blood Components.pdf

6. Hogg G, Pirie E, Ker J. The use of simulated learning to promote safe blood transfusion practice. Nurse Education in Practice. 2006;6(4):214-223.