Recognising and reporting acute transfusion reactions: Experience from STIR

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www.health.vic.gov.au/bloodmatters









Serious Transfusion Incident Reporting (STIR)

Blood Matters:

- Clinical transfusion practice improvement program
- Victoria and Tasmania
- Partnership between DoH and Blood Service
- Clinical governance
- Transfusion nurses
- Graduate certificate transfusion practice
- Haemovigilance











Serious Transfusion Incident Reporting (STIR)

- Pilot 2006
- Multi-state participation
- Voluntary reporting
- Fresh components
- Categories based on SHOT
- National HV data dictionary
- Near misses
- Links with state sentinel event program
- Report to national HV program







Aims of STIR

- Measure and monitor serious transfusion incidents, including near misses, relating to administration and handling of fresh components and pretransfusion samples
- Derive recommendations for better, safer transfusion practice and disseminate these to health services, state and federal governments and the Australian Red Cross Blood Service











STIR

- Two-stage on-line reporting:
 - Initial notification
 - Detailed case investigation form
- MS Word fillable form, MS Access database
- De-identified for patient, staff, institution after report
- Review by HTC prior to submission

Review by multidisciplinary expert group: final classification, severity, imputability













ATRs

- Acute haemolytic
- FNHTR
- Allergic/anaphylaxis
- Bacterial infection
- TACO
- TRALI

Occurring during or up to 24h following transfusion

(Transfusion-associated dyspnoea)





Initial notifications 2006-2012

•	984	reported	events
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- 973 patients (54% F)
- Mean age 49y
- 190 events in <18y
- Initially reported as
 - suspected: 396
 - confirmed: 592
- 542 events (55%) a/w RBCs

Reaction	N	%
ATR	499	50
WBIT	239	24
Near miss	110	11 42%
IBCT	66	7
DTR	27	3
Bacterial*	18	2
TRALI*	29	3
PTP*	1	<1

^{*} Suspected, not confirmed



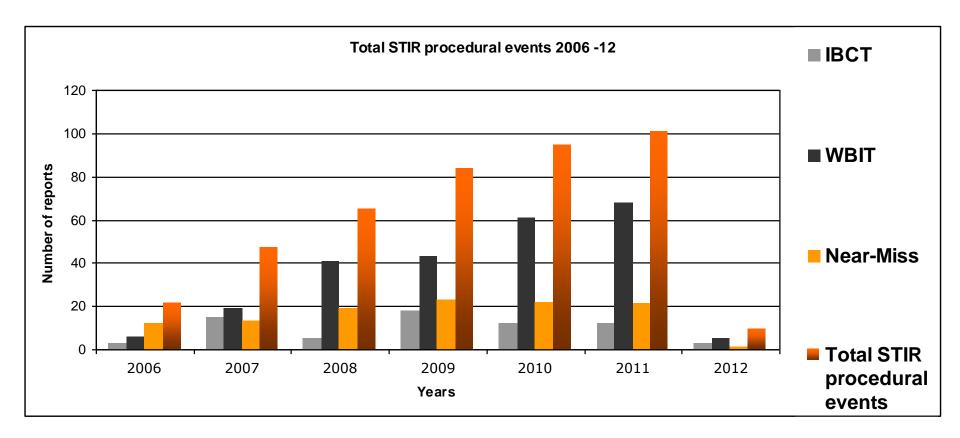


ATRs 2006-2012

- 60% occurred in general ward
- Fever most common reported symptom, often with chills and/or rigors
- Sometimes with dyspnoea, hypertension, tachycardia, nausea, vomiting
- Average 68 min from start tx to identification symptoms
- 9 sentinel events, mostly ABO inc tx
- No deaths
- Many cases ultimately called FNHTR
- 192 allergic reactions (37% all ATRs)
- Severity allergic reactions:
 - mild to moderate (56%)
 - other severe allergic requiring adrenaline (23%), anaphylaxis (21%)







(To March 2012)







Case 1: IBCT: storage and patient ID

- Anaesthetised patient emergency surgery for spinal cord compression
- Incorrect unit and paperwork retrieved from theatre fridge
- Checked unit against paperwork match
- Not checked against patient wristband
- Blood group of red cell unit: B RhD positive
- Blood group of patient: O RhD positive
- Unclear how many mL administered
- Admitted to ICU post-op and made a full recovery



"Yee-ouch! That's gotta hurt."









Case 2: IBCT: ID and labelling

- Sample tube unlabelled and left behind when ED patient removed from resuscitation bay
- Deteriorating (new) patient moved from ED bay to resuscitation bay
- Tube labelled with the new patient's label
- Request for blood sent to lab
- Lab issued 4 units group A RBC
- 2 units transfused in ED and 2 issued for transfer
- Patient found to be group O at receiving hospital















Case 2 cont'd: IBCT

- Lack of compliance with patient ID and specimen labelling protocols
- Lab queried Hb discrepancy AND and unsigned collector's declaration person but still accepted and processed sample
- Lab "was urged to continue" due to deterioration of patient
- Provision of group A blood when could have issued emergency group O unXM











Case 3: IBCT: transcription error

- Preterm infant emergency transfusion in NICU
- Prescribed red cells for Hb 93g/L
- Pathology results transposed
 - Platelets 93, Hb 130
- Post-tx Hb not documented
- Clinical consequences unknown











Case 4: Blood group discrepancy/patient ID

- Historical blood group different to current specimen
- Nursing staff collecting specimen correct patient ID procedure



Patient using sister's name, DOB and Medicare card











Our experience

- ATRs common but often difficult to classify (data, definitions)
- Many mild but accepted anyway to encourage participation
- Improvements in data completeness with e-forms reduce delays/re-work and assist with assessing cases
- Hospitals generally now review before submission data quality
- Role of TN and HTC
- Imputability and severity assignments still challenging esp WBIT and near misses
- Value of central expert review group
- Role of human factors in procedural errors/ATR
- Patient education and participation





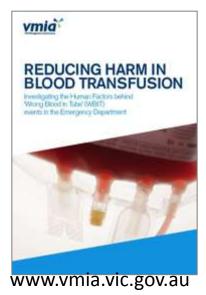






Taking action

- Share data with reporting hospitals
- Share experience: TN network, ANZTP SIG (www.anzsbt.org.au)
- Feedback to hospitals on sentinel events and investigations
- Recommendations to DH on policy and practice issues
- Advice/instruction to hospital CEOs, quality managers
- Educational activities for diverse range of staff
- Provide data to national HV system
- Research
- Design and test interventions



How do we monitor hospital transfusion practice using an end-to-end electronic transfusion management system? Murphy MF et al, Transfusion 2012





www.health.vic.gov.au/bloodmatters

- Blood Matters program
- STIR definition guide
- 2006-07 & 2008-09 reports







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