## SHOT & Lessons Learned from Major Haemorrhage

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### **SHOT Aims**

### Improving patient safety by

- Raising standards of hospital transfusion practice
- Informing policy with UK Blood Services
- Aiding production of clinical guidelines
- Educating users on transfusion hazards and their prevention

## Haemovigilance in the UK

#### **MHRA**

### SHOT

Medicines & Healthcare Products Regulatory Agency

**Serious Hazards of Transfusion** 

- Competent Authority' for the BSQR 2005
  - QMS in blood establishments and hospital blood banks.
- Competent Authority for the Medicines Act 1968
- Competent Authority for the Medical Devices Regulations 2008
- STATUTORY reporting

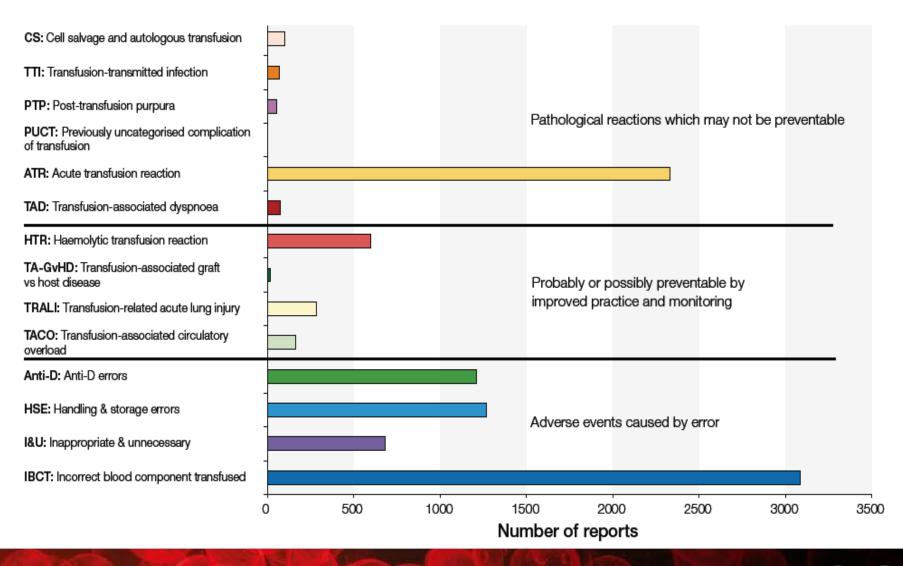
- Confidential enquiry
- Serious adverse reactions/events AND near misses all of which occur in BOTH a laboratory and CLINICAL environment.
- PROFESSIONALLY MANDATED reporting

## SHOT Reports 2011

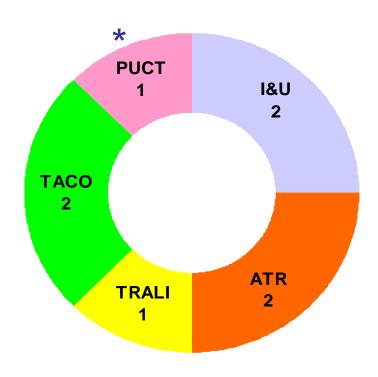
- 3435 reports made to the scheme
  - 2768 analysed
    - 204 in an inappropriate category, moved
  - -667 withdrawn (20%)
- + 270 reports made in 2010, but only completed in 2011
- 3038 cases including 'near miss' and 'right blood right patient'



## Cumulative Reports 1996-2011



# Deaths where transfusion was causal or contributory 2011 (n=8)



\*
The PUCT incident was a case of NEC in a baby that is not categorically linked to the transfusion, but is an association that could not be ignored

**I&U** – Inappropriate & unnecessary transfusion

**ATR** – Acute transfusion reaction

**TRALI** – Transfusion related acute lung injury

**TACO** – Transfusion associated circulatory overload

**PUCT** – Previously uncharacterised complication of transfusion

## Critical points in the transfusion process

- Decision to transfuse
- Prescription/request
- Sampling for pre-transfusion testing
- Laboratory testing
- Collection of blood from storage site
- Bedside administration





## Rapid Response Report

NPSA/2010/RRR017

From reporting to learning

21 October 2010

The transfusion of blood and blood components in an emergency

For action by Trusts by April 2011



## Key points from NPSA RRR

- Local protocols with a trigger phrase
- Dedicated communicator / coordinator
- Early / easy release of components by lab
- Adequate support (eg porters / transport)
- All cases reviewed by Hospital Transfusion
   Committee and delays or problems investigated locally / reported externally

#### MAKE A PHONE CALL

to the **Blood Bank** or on-call **Haematology BMS** 

#### Tell them...

- Situation
- Patient Details

Name

Sex

ID number

• What blood component is required, how much and how soon.

#### TAKE SAMPLES

Label them properly and ensure they get to the lab.

- Blood Bank sample
- FBC
- Coagulation screen



## Overview of process (military)

- Alert Major Haemorrhage Team
- Inform OR, ICU, Radiology & Laboratory
- Transfer patient to relevant location



#### **Assess Bleeding**

Bleeding approaches **500ml** and is continuing

Bleeding approaches **1000ml** and is continuing

Bleeding approaches **2000ml** and is continuing, and/or suspected amniotic fluid embolus

Bleeding continues after >12 units red cells, and appropriate administration of at least 2 doses platelets and/or 2 litres FFP / 10 bags cryo in <12 hours and surgical measures have failed.

#### **Contact Blood Bank**

FBC, PT, APTT, Fib. Crossmatch 4 units

#### **Transfuse Red Cells**

If cross match blood is not available, use **O Neg** emergency stock if appropriate (no more than 2), or **group-specific** units if patient is known.

If bleeding continues, order **group-compatible** units (6 at a time)

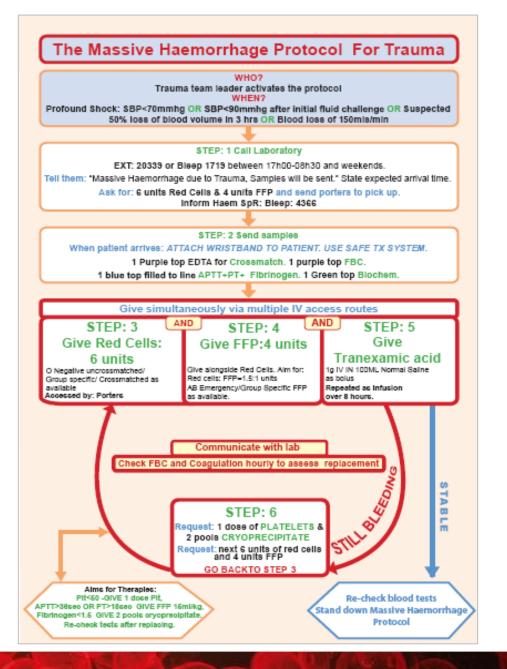
# Contact Haematologist. Transfuse platelets and /or FFP and/or cryoprecipitate to maintain platelet count

maintain platelet count >50 PT<30 sec Fibrinogen >1.5g/l

- ONE ATD of platelets
- ONE LITRE of FFP
- TWO packs of cryoprecipitate

**Discuss** possible use of rFVIIa with Haematologist Initial dose of 90ug/kg, followed by a second dose 2 hours later if blood loss continues.





# Failure to monitor transfusion requirements during a GI haemorrhage

- An elderly patient was admitted to the MAU with a haematemesis and an initial Hb of 10.6 g/dL.
- She had further episodes of vomiting blood.
- Five units of red cells were transfused before a repeat Hb was performed, which was 20.4 g/dL.
- The patient was recognised to have circulatory overload and died shortly thereafter.

# Failure to monitor transfusion requirements during a GI haemorrhage

- An elderly patient with a severe GI bleed had repeat Hbs of 6.1 and 6.4 g/dL.
- Six units of red cells were transfused prior to rechecking the Hb, which was 17.1 g/dL.
- The patient developed circulatory overload and required venesecting 2 units.

## Transfusion of unnecessary components and with inappropriate doses

- A patient was bleeding after a sub-total colectomy and a request was made for 2 doses of platelets and 2 units of FFP.
- The patient had a normal platelet count (245 x 109/L) and a normal INR of 1.2 but the doctor did not check these results.
- The BMS did not telephone these results to the doctor or contact the consultant haematologist in order to challenge the inappropriate decision.

# Lack of knowledge around major haemorrhage protocol (1)

- A middle-aged male was admitted to A&E with a massive haematemesis and received 2 litres of colloid.
- No Incident Communication Coordinator had been identified in A&E and the transfusion laboratory had not been alerted to activate the major haemorrhage protocol
- The clinical staff in A&E were unaware of how to access the 2 emergency O RhD negative units, which were then issued by the lab

contd...

# Lack of knowledge around major haemorrhage protocol (2)

- A further 2 units of red cells were then requested and issued as group specific.
- The clinicians also requested FFP and cryoprecipitate but were refused on the basis that a coagulation screen should have been interpreted by a haematologist prior to issue.
- The patient subsequently arrested and died, having received 10.5 L of colloid and 4 units of red cells.

# Delay in obtaining units following major haemorrhage protocol being initiated

- A child involved in a road traffic accident (RTA) was found to be asystolic at the scene and cardiopulmonary resuscitation (CPR) was commenced.
- The ambulance staff had alerted A&E to major blood loss and had requested blood to be available there on arrival.
- The major haemorrhage protocol, however, required a unique patient number to be allocated prior to issuing emergency O RhD negative units from the transfusion laboratory and there was a delay following the patient's arrival in A&E for any red cells to be made available.

# Failure to replace blood volume after post partum haemorrhage

- A woman in her mid-thirties had a ventouse-assisted vaginal delivery for fetal distress at term, complicated by massive haemorrhage.
- The major haemorrhage protocol was activated, six units of blood were delivered within 5 minutes and one was started immediately.
- She was transferred from the delivery room to theatre and the bleeding was controlled within 30 min.
- The blood loss was unclear with losses recorded in both the delivery suite and theatre. A second unit was commenced.
- About 2 hours later, she suffered cardiac arrest from which she could not be resuscitated despite transfusion of 12 units of blood and 3 units of Fresh Frozen Plasma (FFP).
- The coroner confirmed cause of death to be cerebral hypoxia secondary to haemorrhage.

# Late request for blood to cover surgery leads to inappropriate use of emergency O RhD negative blood.

- An elderly lady was admitted on the morning of surgery for major abdominal surgery and a sample was sent for grouping with request for a crossmatch.
- She was taken to theatre without waiting for results.
- The antibody screen was positive the BMS phoned theatre, but surgery was already underway.
- Four units of O RhD negative emergency blood and 4 units of FFP were transfused.
- The antibody was anti-E and fortunately the O RhD negative units used were compatible.

## Delay in patient transfusion during AAA surgery caused by a BMS error and IT malfunction

- A 75 year old man was bleeding in theatre during repair of AAA. The massive haemorrhage protocol was activated, and 6 units of group-specific blood were issued to the theatre refrigerator using the electronic blood tracking system.
- The units were retrospectively cross matched and results added to the Laboratory Information Management System which sent a message to the theatre refrigerator to quarantine the units, possibly because the system had received two conflicting messages about the units.
- Theatre staff were denied access to the refrigerator and nobody knew how to proceed
- Eventually the refrigerator was unlocked remotely and the blood obtained after a 25 minute delay.

## Obstetric major haemorrhage with delay in transfusion caused by a fire alarm.

- A 40 year old woman was undergoing elective caesarean section and started to bleed excessively. At the same time, the fire alarm sounded.
- The obstetrician and theatre staff were aware of the alarm, but management of the bleeding continued.
- Urgent bloods were sent to haematology via the tube system and the laboratory was telephoned to alert them to the need for urgent analysis and a need for blood products.
- However, there was no answer so an assumption made that the laboratory had been evacuated.
- The general manager (outside the building with evacuated staff) was contacted and located haematology staff who were cleared to return to the laboratory.
- Blood samples were analysed and major haemorrhage pack was issued

### Misidentification during multiple trauma

- Multiple RTA involving 5 victims
- 2-year old female allocated 'Unknown Female 1'
- Child's mother allocated 'Unknown Female 2'
- Theatre nurse volunteered to help ED teams
- Nurse administered blood for 'Unknown Female 2' to the child, as she interpreted the '2' as the age of the patient
- The child died of her injuries soon afterwards



## Lack of correct final identity check leads to a Haemolytic Transfusion Reaction

- A patient with a haematemesis needed an urgent blood transfusion.
- The patient's wristband was contaminated with blood and could not be read, so the electronic bedside checking system was not used.
- The compatibility form filed in the patient's notes, which belonged to another patient, was used to provide the identifiers for collecting the blood.
- O Pos patient was transfused with >50 mL of A RhD positive red cells before the error was noticed.
- The patient was admitted to ITU with intravascular haemolysis and renal impairment.

## Communication failure results in inappropriate transport of red cell units

- Telephone request from ED for 4 units of blood which was subsequently issued to the laboratory fridge.
- 2 hours later, the ED called asking if blood was ready, as the patient was being transferred
- The ward were told that it would take 10–15
  minutes to package the blood and issue the
  appropriate documentation, but the BMS found that
  it had already been removed
- The receiving hospital contacted the lab to inform them that 1 unit had arrived with the patient and had been transported in an ASDA bag.

# Unlabelled products transfused to wrong patient in error

- Platelets for a patient on ITU were delivered to the ED by taxi from BTS
- The ED had also requested platelets for a different patient.
- ED took delivery of the platelets, assumed they were for their patient, and transfused them despite there being no documentation or label with any patient details.

# Blood gas analyser Hb used as trigger for emergency transfusion

- An Hb of 5.0 g/dL was obtained from an ED blood gas machine on a middle-aged female patient who was asymptomatic and not actively bleeding.
- Two units of O RhD negative red cells were issued as an emergency
- One unit of O neg had already been transfused when the laboratory result became available which was 8.9 g/dL.
- The second unit of O neg was wasted due to inappropriate storage (on the patient's bed)



## Overestimation of blood loss from acute GI bleed

- A patient was admitted to the ED with a GI bleed. Hb on admission was 12.1 g/dl.
- 2 units of emergency blood were given, followed by 6 units of crossmatched blood over the next 12 hours.
- The FBC was not re-checked until all 8 units had been transfused, and the Hb was 18.5 g/dl

# **Excessive transfusion follows**misinterpretation of verbal instructions

- A 48 year old male patient in ED with a GI bleed.
- Five units of blood ordered and a verbal order for 2 units was given by the doctor, who then wrote them up on a prescription chart.
- Staff nurse asked the doctor if he wanted the blood given through the rapid infuser, and he confirmed that "all the blood can go though this".
- All five units were transfused instead of the intended 2 units.

### In brief....

- Delay in provision of blood to an off-site obstetric haemorrhage as the courier had not been trained.
- Adult with Hb of 4.7 g/dl in ED. Nurse was sent for the emergency O Neg, but returned with a paedipack, which was administered after checking at the bedside by two doctors.
- Had to resort to using O Negs during an obstetric haemorrhage as SpR had labelled crossmatch sample with another patient's details.



## **Key Learning Points**

- Continue to sample, monitor and observe during a transfusion
- Appropriate assessment and prescription
- Good patient ID at all stages
- Communication with laboratory
- Early release of components by laboratory
- Clear instructions / easy to follow protocol
- Do your own job well





### Thanks to

- The SHOT Team
  - Vicky, Julie, Chris, Hema, Debbi & Alison
- Paula Bolton-Maggs, SHOT Medical Director
- You for listening

www.shotuk.org

