

# Mind the Gap:

## Pre-hospital blood and the blood banker

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# Disclaimer

The views and opinions expressed in this presentation are those of the author and do not reflect official policy or the position of NHS Blood and Transplant or the Defence Medical Services.

## Aim of presentation

Pre-hospital transfusion may impact on the blood banker. Examples include:

- Legal and regulatory requirements

- Demand and stock management

- Impact on continuing transfusion care

The aim of this presentation is to explore some of the challenges and potential solutions

# The military influence: Damage Control Resuscitation (DCR)

Aims of DCR  
include:

Minimise blood  
loss

Maximise tissue  
oxygenation

Optimise  
outcome



*Transfusion is an integrated part  
of Damage Control Resuscitation  
Transfusion may begin in the  
pre-hospital space*



## Potential challenges

Sufficient 'universal' blood components

Secure cold chain – sufficient proof of compliance to return unused blood to hospital stock

Traceability – confirm fate and ability to re-call for up to 30 years

Haemovigilance and serious event reporting

Confirmation of ABO and RhD groups

**Compliance with 'national' regulators**

Providing further evidence - RCT

# Mind the gap



- **Safe blood**
- **Safe transfusion**
- **Safe transfer**
- **Safe systems**

# Safe Blood

# Hierarchy of formal blood supply: (EU model)



Blood is normally collected, tested and processed by a **blood establishment**

Supply to ambulance/pre-hospital groups trusts may be:

- Direct from blood service

- Indirect via **blood bank**

Pre-hospital group may be a **facility** or extension of the blood bank (if formal arrangements SLA/MOU in place)

# Meeting the demand for Universal Blood Components

Group O RED CELLS  
(RCC) can be used for all  
ABO groups

Group Rhesus D neg red  
cells prioritised for females

Group AB PLASMA used for  
all ABO groups (the RhD  
group does not matter)

Consider 'Universal plasma'  
or Group A Plasma with low  
titre anti-B



**Group O RhD neg Blood and  
Group AB plasma are in short  
supply globally**

**Pre-hospital transfusion may  
have an impact on local and  
national blood supplies**

# Blood component storage and shelf-life

Component	Storage	Shelf-life	Extended shelf-life
Red cells	4°C $\pm$ 2	35 days	42
FFP	-25°C	3 years	
Thawed FFP	4°C $\pm$ 2	24 hours	5 days (if dry thaw)
Platelets	22°C $\pm$ 2 + agitation	<b>7 days</b>	

# The working environment



*Simulate during staff training and validation of equipment and systems*

## External:

Temperature

Dust

Wind

Cold

Wet

## Internal:

Dark

Noisy

Vibration

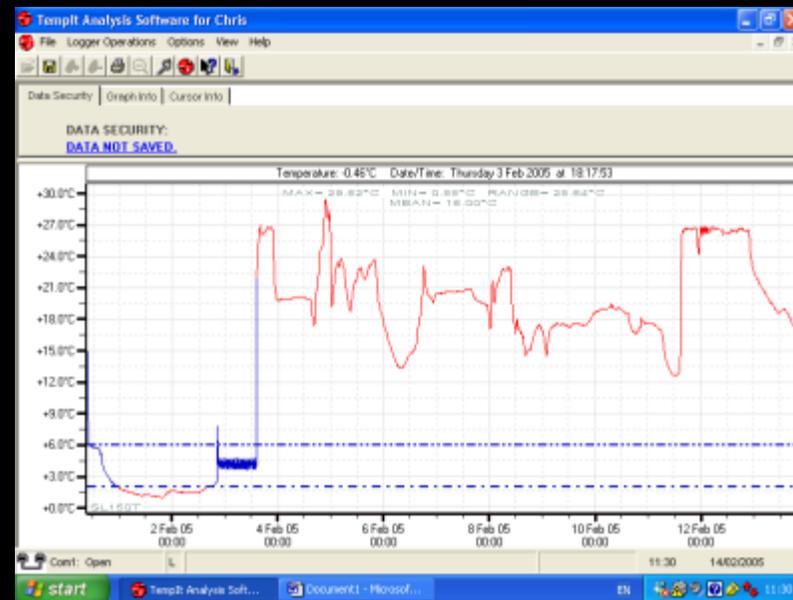
Constrained

Dangerous

# Validate delivery systems



# Monitoring the cold chain



*Retain records for internal and external audit*

# Traceability: An example of a 3 part label

F Med 682  
(Rev 01/06)

**MOD CROSSMATCH LABEL**

ISSUED FOR

Donation No:.....  
Hospital/Trauma No:.....  
Service No:.....  
Surname:.....  
Forename:.....  
DOB:.....  
Blood Group:.....

Serial No: 025203

Donation No:..... Product:.....  
The above listed unit was transfused into the following  
Hospital/Trauma No:.....  
**Insert into Patients Records**  
Surname:..... Forename:.....  
On: / / at : hrs  
Checked by:..... Signature:.....

Serial No: 025203

Donation No:..... Product:.....  
The above listed unit was transfused into the following  
Hospital/Trauma No:.....  
**Return to Blood Bank**  
Surname:..... Forename:.....  
On: / / at : hrs  
Checked by:..... Signature:.....

Serial No: 025203

**Return this section to the Blood Bank**

Grey - Leave on bag

Blue – put into patient record

Pink – return to blood bank

# Safe Transfusion

# Serious Hazards



*'Risk is relative'*

Context

Commitment to patient safety

Consider requirement to capture both blood donor and patients events

Challenge of recognition

Challenge of global reporting

# Risks at reception

Baseline blood sample  
Switch from universal to  
group specific  
Switch from a manual to  
digital system  
Completion of  
transfusion records

*There must be a safe  
system to link Pre-  
hospital and Hospital  
records and ID*

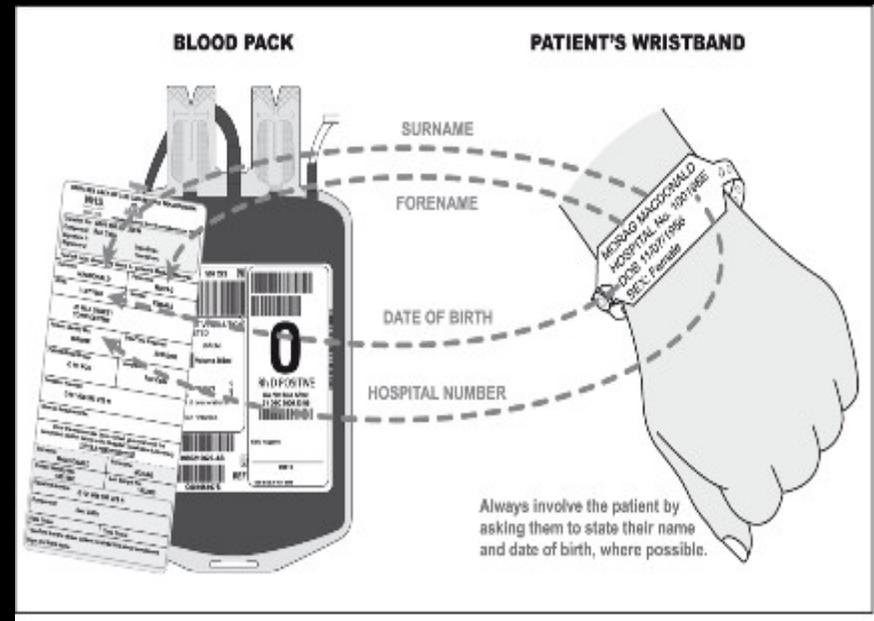


*The biggest risk in transfusion is  
giving ABO incompatible blood  
due to human error. Risks  
increase during the transfer from  
pre-hospital to hospital care*

# Changing identification

Identification is required in hospital for:

- ❑ Blood sampling
- ❑ Collecting blood components
- ❑ Administration of blood



*There should be compatibility between local emergency ID systems and receiving blood bank Information systems*

# Recognition and management of Acute Transfusion Reactions

Focus on 'C'ABC and managing physiology

Consider risks

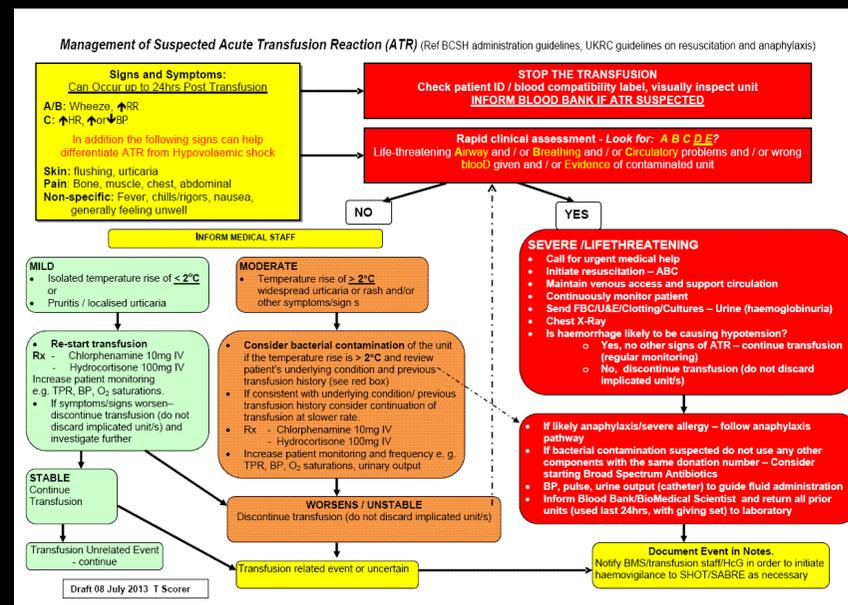
? Group O blood – then ABO unlikely

Group specific – return to group O

Previous transfusion – unknown – haemolytic reaction possible treat later

Use of plasma – manage allergic response

Continue resuscitation



Management of ATRs

CGOs 2014

# Challenge of recognition

Reactions are rarely reported in the critically ill

Some reactions may occur late (after transfer) and are may not be recognised as transfusion reactions.

**Novel problems** may present e.g. mechanical haemolysis and the pre-hospital treatment should be considered during investigation

# Mechanical haemolysis



Non immune haemolysis  
in the rapidly transfused

Causes could include:

Narrowed subclavian  
lines (as reported in  
renal dialysis)

Administration system  
i.e. Rapid infusers  
and connectors

Blood forcefully given  
via IO devices

Cases must be reported to  
national haemovigilance  
systems

# Safe transfer

# Reducing the risk at Patient transfer



**Pre-hospital/Previous transfusion (and blood group if known) should be included in notification and handover**

# Transfusion support at receiving hospital

## Continue providing blood components

Establish (or confirm) patient blood group (ABO and RhD)

Move to group specific blood components when blood group confirmed

Screen and identify atypical antibodies and advise

Follow-up of untested blood, transfusion reactions or serious/adverse events

Complete traceability and transfusion record

*Complete research records*

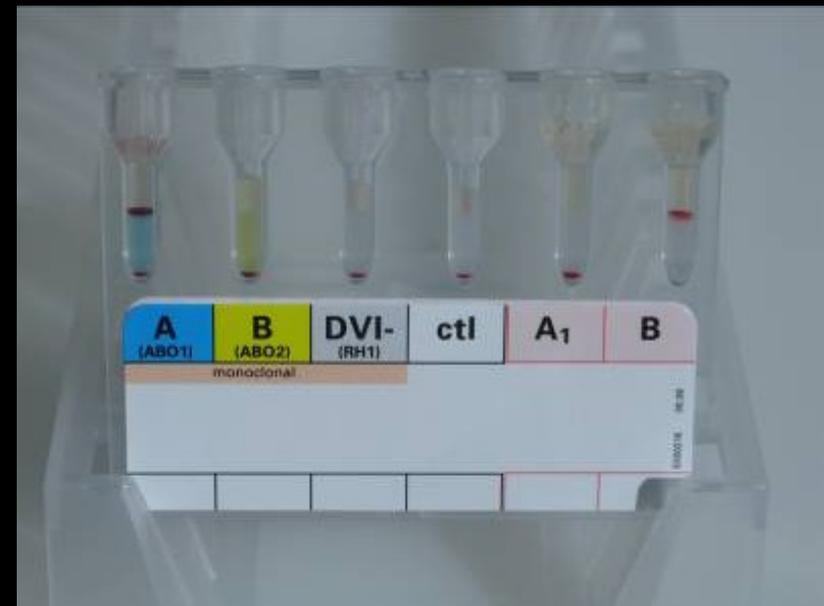
Post exposure consent to transfusion

# Pre-hospital transfusion: Changing ABO groups

The use of large amounts of ABO non-identical blood may 'obscure' patient blood groups

Receiving hospital may have to use 'universal components'

**Transfusion history important to determine blood group and risk**



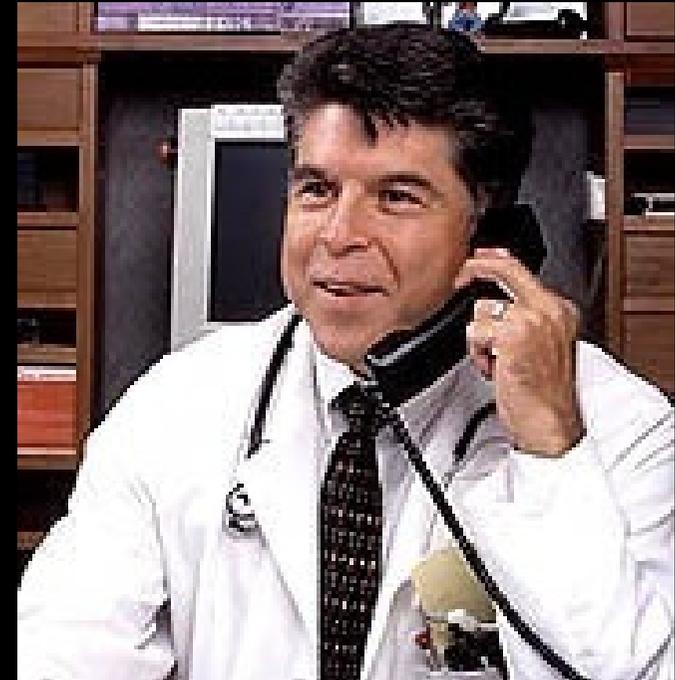
*Group O cells in a group A RhD neg patient*

# Communication errors from bench to bedside

822 outgoing  
telephone calls from  
laboratories

29 errors were detected  
(3.5%)

*“A significant proportion  
had the potential to  
endanger the patient”*



# Improving communication Applying human factors

A small number of simple interventions would reduce errors, e.g.:

- ✧ Structured messages
- ✧ 'Read back'
- ✧ Protocols for speaking up
- ✧ Better focussed, rather than more extensive, checking procedures



# Safe systems

## Blood facility compliance

An organisation which receives blood from a hospital blood bank for transfusion purposes is defined as a **facility**.

The 3 key tasks covered by a facility blood compliance report are

- Cold chain management

- Reporting serious adverse events

- Traceability records

Evidence provided via documentation within a quality system

# Documentation e.g. Training



All staff involved in the transfusion process should be trained.

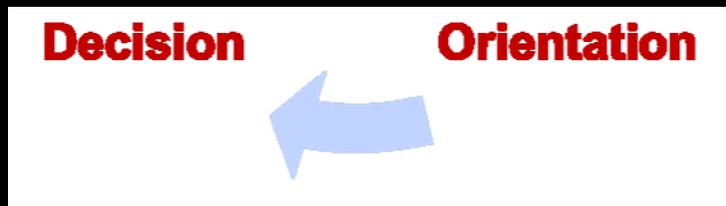
Documents required include:

- Policy

- Roles and responsibilities

- Group and individual Training records

# Clinical governance systems



*Early, local investigation of transfusion reactions and safety events*

## Pre-hospital representative

Support implementation of local policy and practice

Promote best transfusion practice through training, education and audit

Review usage and wastage of blood and advise on stock holdings

Review traceability statistics and investigate non-compliance

**Support investigation and management of transfusion related events**

# Good blood in bad places

## Providing continuity of transfusion care



**Allcock, E.C., Woolley, T., Doughty, H., Midwinter, M., Mahoney, P.F., Mackenzie, I. (2011) The Clinical Outcome of UK Military Personnel who received a Massive Transfusion in Afghanistan during 2009. *Journal of the Royal Army Medical Corps*, 157, 365-369.**

# Together – We 'Mind the gap'

