



Patient Blood Management, Consent for Transfusion and Patient Information

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BBT (Better Blood Transfusion)

Department of Health Health Service Circular (HSC)

- HSC 1998/224: Better Blood Transfusion HTCs; Transfusion guidelines and protocols; SHOT participation
- HSC 2002/009: Better Blood Transfusion [2] Appropriate Use of Blood HTTs; BSMS participation; lab. accreditation; traceability; national guidelines for appropriate use
- HSC 2007/001: Better Blood Transfusion [3] Safe and Appropriate Use of Blood



BBT3

All NHS Trusts providing blood transfusion should [have]:

- Functioning HTT & HTC
- Action plan for compliance with BSQR
- Meet NPSA competency assessment
- Policies for blood transfusion & refusal; patient identification and technologies to support this
- Participate in BSMS & NCA
- Laboratory: CPA accreditation, out of hours covers, & GMP
- Report to SHOT & SABRE
- Regular audit of transfusion practice

BBT3

..... and should [have]:

- Guidance for the use of red cells, FFP & platelets [indications]
- Establish protocols to empower laboratory staff to ensure appropriate clinical information is provided with requests
- Pre-operative assessment/ anaemia management of surgical patients
- Develop blood conservation strategies: cell salvage, point-of-care testing, pharmacological agents
- Procedure for prescription & administration of Anti-D, and training / assessment for the prevention & management of HDN
- Procedure for identification & management of maternal anaemia
- Increase patient & public involvement, ensuring patients are informed o transfusion & alternatives.



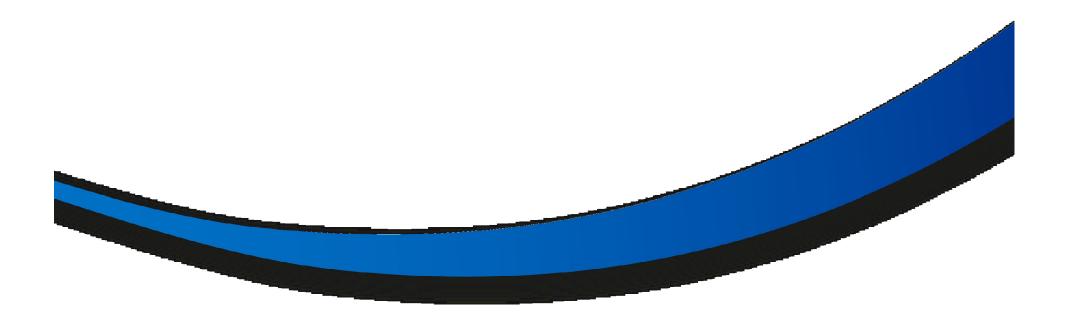
BBT3

NHSBT & NBTC required to support HTT/Cs

• Trusts required to ensure representatives from HTC can attend regional & national committees.



Patient Blood Management





Patient Blood Management

Sub heading

Text

•Evidence-based

Multidisciplinary approach

•Optimising the care of patients who might need a blood transfusion.

•Puts the patient at the heart of decisions

•Ensure that they receive the best treatment

•Avoid the inappropriate use of blood and blood components.



Why does it matter?

- Improves care
- Reduces inappropriate transfusion
- Ensures availability where there are no transfusion alternatives available



PBM – Key Messages

- Patient at the heart of decision making
- **Blood** conserve patient's own blood, avoid transfusion where appropriate
- Management organise and co-ordinate

Further information can be found at:

http://www.transfusionguidelines.org.uk

under 'National Blood Transfusion Committee'

PBM

- Represents an international initiative in best practice for transfusion medicine; NHSBT is working with the Department of Health and the National Blood Transfusion Committee to support NHS Trusts to manage their blood use effectively.
- Evidence shows that there is inappropriate use that can be reduced and that the current trend of use is not sustainable.

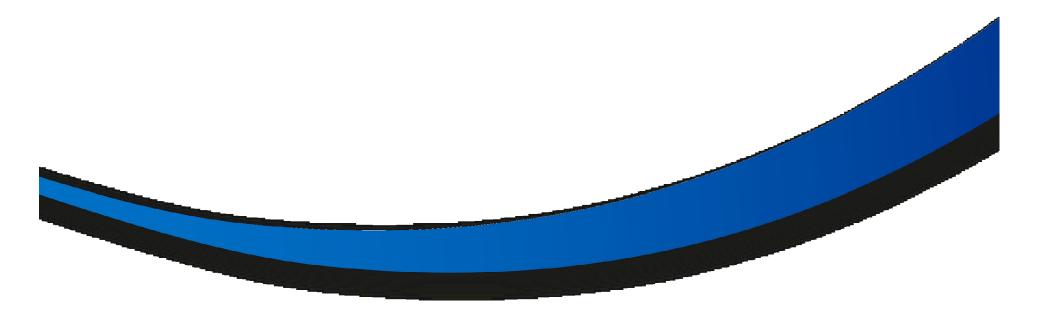


PBM Key Elements

- Use of performance measures (KPIs) in transfusion
- Consent for transfusion
- Cell salvage, IV iron, and use of alternatives
- Management of anaemia
- Clinical audit/tracking of blood use
- NICE guidelines



Consent for Blood Transfusion





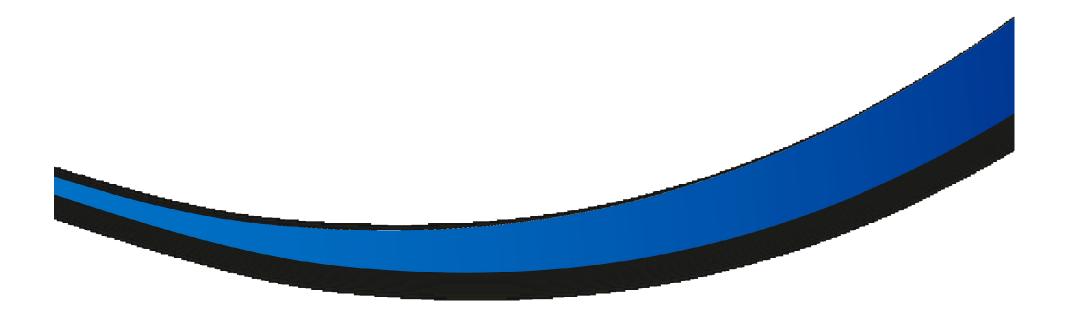
Background

In March 2010 SaBTO initiated a public consultation on patient consent for blood transfusion



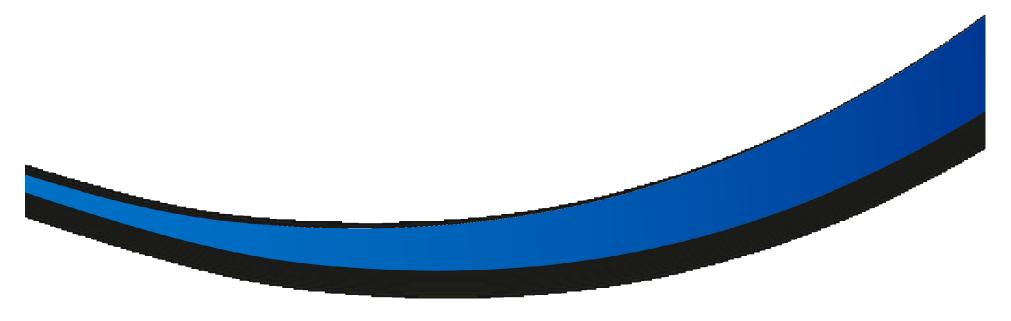
Why?

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- Public Health: Recipients may not be aware that they have received blood and then go on to donate.



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- Patient Choice: Many patients may not wish to receive a blood transfusion and / or may wish to know what the alternatives are.
- Public Health: Recipients may not be aware that they have received blood and then go on to donate.
- General legal and ethical principle: Valid consent should be obtained from a patient before they are treated.

It had the following Objectives:

- Identify the preferred option for recording consent
- Explore the potential operational impact of implementing a standardised form of consent for transfusion
- Confirm what type of information patients should receive



Identified key issues in transfusion practice

- Not always given information on risks, benefits, and alternatives to transfusion, or the right to refuse transfusion
- Not always made aware that they have had a transfusion
- Those unaware that they have received a transfusion may go on to donate blood when they should not
- There is inconsistent practice across the UK

Summary of Action Plan



14 recommendations / 3 broad categories:

Clinical practice:

What should be done / hospital policy Recommendations 1-6

Governance: Review of clinical practice Recommendations 7 -10

Education:

To help support clinical practice Recommendations 11-14



1.

Valid consent for blood transfusion should be obtained and documented in the patient's clinical record by the healthcare professional



2.

There should be a standardised information resource for clinicians indicating the key issues to be discussed by the healthcare professional when obtaining valid consent from a patient for a blood transfusion

Standardised Information Resource Blood and Transplant

http://www.transfusionguidelines.org.uk/index.asp?Publication=BBT&Section=22&pagei d=7691



GUIDANCE FOR CLINICAL STAFF

TO SUPPORT PATIENT CONSENT FOR BLOOD TRANSFUSION

Patient may require Blood / Blood Component Transfusion

Patients receiving a blood transfusion (red cells, platelets or plasma) whether for a medical or surginal cause should be informed of the indication for the transfusion including risks, benefits and alternatives. A record of this discussion should be documented in the patient's diring a records.

Ideally the decision to transfuse should be made with the patient or parent/carer in advance of any planned transfusion.

In the emergency setting, the information will need to be given retrospectively.

Prospective Information

Retrospective Information

Valid consent* should be obtained prior to any planned transfusion and documented in the patient's dinical record. Patients treated in emergency setting where it was not possible to obtain valid consent pre-transfusion

Patients who were told pre-procedure (e.g. previsito consent entails the provision of information on risks, benefits and attentives available tetrice axing the patients dure consent. This descend have to include a signature from the patient.
Patients who were told pre-procedure (e.g. preoperatively) that they might require a transfusion then need to be informed whether they did/did not receive a transfusion.

Key issues to be discussed when obtaining valid consent

- 1. The following information should be discussed:
 - o Type of blood / blood component
 - o Indication for transfusion
 - o Benefits of the transfusion
 - o Risks of transfusion
 - Possible alternatives to transfusion
 - o How the transfusion is administered and the importance of correct patient identification
 - Inform patient that following a blood transfusion they can no longer be a blood donor.
- 2. Provide written information.
- 3. Check if patient needs time to consider or requires further information.
- Document the discussion in the patient's dinical records.

At discharge

- 1. If patient has had a transfusion, ensure that they have been informed.
- Record information about the transfusion in the discharge summary, also stating that the patient has been informed.

Resources

This guidance applies to the transfusion of all blood components (red cells, white cells, platelets, fresh frozen plasma & cryoprecipitate) and should be used by healthcare organisations to strengthen the consent processes already in place.

Specific guidance should also be used to ensure that alternatives have been considered for blood and blood components e.g. pre-operative iron therapy, intra-operative cell salvage where appropriate for avoidance of red cell transfusion and prothrombin complex concentrate in place of FFP for warfanin reversal.

Adverse events

Clinical learns involved with the prescribing and administration of blood and components must be aware of adverse events that can be associated with transfusion including prompt recognition and management (<u>www.shotuk.orc</u>). These include:

incorrect Blood Component Transfused (IBCT)	Inappropriate, Unnecessary, Under/Delayed Transfusion (landU)	
Acufe and Haemolytic Transfusion Reactions (ATR and HTR)	Transfusion-Transmitted infection (TTI)	
Transfusion-Associated Circulatory Overload (TACO)	Transitusion Associated Acute Lung Injury (TRALI)	
Transfusion-Associated Dysphoea (TAD)	Transfusion Associated Graft-versus-Host Disease (TA-GvHD)	

Post Transfusion Purpura (PTP)

Clinicians should refer to the HPA website (www.hoa.org.uk) to get the latest information on the risks of transmissible infections. Current guidance from the HPA states that the risk of getting hepatitis from a blood transfusion in the UK is currently (January 2011) about 1 in 670,000 for hepatitis B and 1 in 83 million for hepatitis C. The chance of getting HIV (Human Immunodeficiency Virus) infection is about 1 in 5 million or HTLV (Human T-Lymphotropic Virus) infection is about 1 in 18 million.

Although the risk of getting variant Creutzfeldt-Jakob Disease (vCJD) from a blood transfusion is probably very low with a single blood transfusion, the risk of any infection will increase with each additional blood component.

Long-term transfusion-dependent patients

Long-term transfusion-dependent patients will need modified consent. This should where possible include an initial discussion at the start of a transfusion regime with a regular update including appropriate information regarding the benefits and risks of transfusion and specific relevant issues e.g. iron overload, risk of allo-immunisation including haemolysis risks (red cells) and platelet refractoriness (HLA antiboties), infective risks and other transfusion reactions.

Other information

Where needed, patients should be provided with contact details of key specialists for further discussion around blood transfusion issues relevant to their specific clinical diagnosis e.g. hospital transfusion practitioner, local haematologist or other clinician such as anaesthetist, surgeon or obstetrician. The Trust website can be used to provide information for patients about consent and safe blood transfusion.

Useful websites	www.transfusionouidelines.org.uk.	www.blood.co.uk
www.nhs.uk/conditions/blood-transfusion	www.nhshealthguailty.org	www.shotuk.org
www.hpa.org.uk	www.bcshquidelines.co.uk	www.sign.ac.uk/guidelines/
Patient information leaflets are available from : www.hosoital.blood.co.uk		



5.

A UK Comparative Audit of consent for transfusion should be carried out, facilitated by the National Comparative Audit for Blood Transfusion

NCA 2014 Audit of Patient Information and Consent





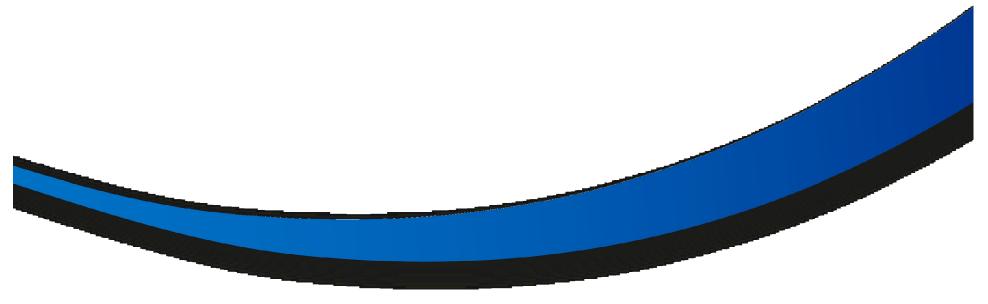
6.

There should be a standardised source of information for patients who may receive a transfusion in the UK



Patient Information Leaflets

• There is a list of key points that should be included in all adult patient information leaflets.



Key issues

- Discuss indications, benefits, risks and alternatives
- How transfusion is administered and importance of positive patient identification
- Provide the patient with written information
- Document the discussion in the patient's clinical records
- Ensure patient knows that can no longer donate

Local Resources

- Transfusion Practitioner
- Transfusion Laboratory Staff
- Hospital Transfusion Committee
- YOUR Trust Transfusion policy
- The Patient Blood Management Team
- Regional Transfusion Committee

Web based Resources

•www.transfusionguidelines.org
•ww3.access-24.co.uk
•http://hospital.blood.co.uk
•www.blood.co.uk
•www.learnbloodtransfusion.org.uk



Patient information can also be found at NHS Choices at: *http://www.nhs.uk* and via Facebook, Twitter and You Tube





- The majority of blood transfusions go to elective patients where there is time to discuss transfusion and the alternatives available
- It is very important that you find time to give your patient a leaflet and discuss with them the risks and benefits of receiving a blood transfusion



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