

PT ID																				
SURNAME															D.O.B.....					
OTHER NAMES															SEX					
ADDRESS.....															MARITAL STATUS					
															REL.....					

Reaction Type	Signs and Symptoms	Clinical Action for ALL Transfusion Reactions.
Mild Allergic	Localised urticaria, pruritus, rash, wheeze	1. STOP the Transfusion Immediately. 2. Check Vital Signs – Call for Assistance if required. 3. Notify RMO.
Severe Allergic	Flushing, wheezing, hypotension anaphylaxis	4. Maintain IV Access – withdraw 5-10mL prior to flushing with normal saline.
Febrile - (Non Haemolytic)	Unexpected fever (>1° C) which may be accompanied by chills and rigors	5. Recheck patient ID and product ID at the patient's bedside. 6. Notify Pathology for guidance.
Acute Haemolytic ABO Incompatibility	Rigors, fever, flank or IV site pain, tachycardia, dyspnoea, hypotension, unexplained bleeding, oliguria, haemoglobinuria	7. Collect the following: * 1 x 6ml EDTA tube (Pink) * 1 x set of blood cultures * 1 x serum sample (Gold top) * 1 x urine sample.
Septic Reaction	Fever, chills, rigors, nausea, vomiting, hypotension	8. Return all Blood Bags with giving sets to Pathology.
Transfusion-Related Acute Lung Injury (TRALI)	Dyspnoea, respiratory failure, non cardiogenic pulmonary oedema, chills, fever	9. Document Event. <input type="checkbox"/> Progress Notes <input type="checkbox"/> EIMS <input type="checkbox"/> Blood Product & Reaction Investigation Form

Transfusion Safety
Transfusions must commence within 30 minutes once product collected from controlled storage and must be completed within 4 hours from removal from controlled storage. Red Cells must be administered over a minimum of 2 hours, unless an emergency.
Patients receiving blood components or blood products must be directly observed for the first 15 minutes.

Refusal to permit transfusion of blood components or blood products:

1) Refusal to permit
I, hereby **withhold** my consent to and forbid the administration of blood or its derivatives under any circumstances to during this stay in hospital.
Relationship to patient if not self.....

The benefits and risks of this transfusion have been explained to me and I accept that this decision may have risks of permanent damage or possible death.

I have had the chance to ask questions and have had all of my questions answered.

I accept full responsibility for the risk and potential consequences of my decision.

The reason for my refusal is

I refuse the transfusion of Blood Components/Blood Products
Patient/guardian Date.....
Patient/guardian signature

2) Confirmation of refusal
I Dr..... have explained to the potential consequences which may flow from the refusal to allow blood or its derivatives to be administered. It is my understanding that this explanation has been understood.
Medical Officer Date.....
Medical Officer signature

PT ID																				
SURNAME															D.O.B.....					
OTHER NAMES															SEX					
ADDRESS.....															MARITAL STATUS					
															REL.....					

Medical Officer to complete section 1, 2, or 3 for all episodes of transfusion.

CONSENT	Section 1 I Dr..... Medical Officer have discussed with (patient/guardian) the recommendation of a transfusion of blood components/blood products for Name of person if not self Please initial where applicable <input type="checkbox"/> We have discussed the benefits and the foreseeable risks of the proposed transfusion <input type="checkbox"/> We have discussed the alternatives to the proposed transfusion <input type="checkbox"/> The person giving consent has been given the brochure 'Blood Transfusions – Have all your questions been answered' I have had the chance to ask questions and have had all of my questions answered. I consent to the transfusion of Blood Components/Blood Products Patient/guardian Date Patient/Responsible Person signature Medical Officer obtaining consent Medical Officer Date Medical Officer signature
	Section 2 Transfusion dependent patients: Patients requiring regular transfusions for the same indication/condition, the consent for transfusion is valid for 12 months. Patient's Indication/Condition Date annual consent completed If a transfusion is required for an indication/condition other than that described on their annual consent form, a new consent form must be completed for each episode of transfusion (e.g. an acute admission) Name Signature..... Designation..... Date Medical Officer/RN print name Medical Officer/RN signature <input type="checkbox"/> This consent is to remain valid for the next 12 months unless the indications for transfusion change (see Section 2).
	Section 3 Life Threatening Situation (Emergency Transfusion): Medical Officer to Complete This episode of transfusion is due to an emergency situation. The patient/responsible person is incapable of agreement to the proposed transfusion. Completion of documented consent by the patient/responsible person for transfusion was unobtainable. Medical Officer Signature Date Medical Officer print name Medical Officer signature

Medical Officer to complete:

Red cells: Patient <input type="checkbox"/> unstable <input type="checkbox"/> transfusion dependant Blood Loss <input type="checkbox"/> rapid <input type="checkbox"/> ongoing <input type="checkbox"/> anticipated Symptoms <input type="checkbox"/> dyspnoea <input type="checkbox"/> fatigue <input type="checkbox"/> dizziness <input type="checkbox"/> angina <input type="checkbox"/> Impaired Reserve <input type="checkbox"/> cardiac <input type="checkbox"/> pulmonary <input type="checkbox"/> bone marrow dysfunction <input type="checkbox"/> cerebrovascular Altered O ₂ Requirements <input type="checkbox"/> tissue hypoxia <input type="checkbox"/> sepsis <input type="checkbox"/> anaesthesia	Current Hb.....g/L; Ferritin
Fresh Frozen Plasma <input type="checkbox"/> Haemorrhage with INR > 1.5 <input type="checkbox"/> Haemorrhage – patient over anti-coagulated <input type="checkbox"/> Pre-surgery with INR > 1.5 <input type="checkbox"/>	Cryoprecipitate <input type="checkbox"/> Haemorrhage with Fibrinogen deficiency (<1.0g/L) <input type="checkbox"/> Acute DIC <input type="checkbox"/>
Platelets <input type="checkbox"/> Thrombocytopenia <input type="checkbox"/> Bone marrow failure – plt count <10x10 ⁹ L or <20x10 ⁹ L with risk factors <input type="checkbox"/> Bleeding <input type="checkbox"/> Invasive surgical procedure <input type="checkbox"/> Current Platelet Count	Biostate <input type="checkbox"/> FVIII or <input type="checkbox"/> vWF Albumex <input type="checkbox"/> Prothrombinex <input type="checkbox"/> with bleeding due to factor II, IX and or X deficiency <input type="checkbox"/> Warfarin reversal <input type="checkbox"/> Current INR
<input type="checkbox"/> Massive Transfusion <input type="checkbox"/> Human Anti-D Immunoglobulin Refer to appropriate form	IVIG <input type="checkbox"/> Immunology <input type="checkbox"/> Haematology <input type="checkbox"/> Neurology <input type="checkbox"/>