

APPENDIX A

Patient Details (or pre-printed label)

## **Consent Form for the Refusal of Blood Transfusion**

Patient Surna	me/ family	nan	ne										
Date Of Birth	1				Male		Fema	le					
CHI:						<b>-</b>	J						
This part to be completed by a Registered Medical Practitioner													
Type Of Operation Investigation or Treatment:													
A patient 'ag	reement to	o in	vestigation o	r treatm	ent conse	nt form' has	Yes		No				
been complet	ted, if no, please give reason why above												
I acknowledge that this limited consent will not be over-ridden unless revoked or modified, this should be													
recorded in w	nung.												
I am the	patient			parent			guardia	ın					
I agree	• To	To what is proposed, which has been explained to me by the doctor named on this											
(subject to	form												
the	To the use of the type of anaesthetic that I have been told about												
exclusions	<ul> <li>To the use of non-blood volume expanders; pharmaceuticals that control haemorrhage and/ or stimulate the production of red cells.</li> </ul>												
below)					-								
I have told the doctor		ınat	I am one of t	ne Jeno	van's with	esses with fir	m religi	ous con	viction	S			
(tick as	I	I am refusing blood for personal reasons											
appropriate)	That the procedure might not be done by the doctor who has been treating me so far.												
understand													
understand	That my express refusal of allogeneic blood or primary blood components, as indicated on page two, will be regarded as absolute and will NOT be over- ridden in												
					_								
	any circumstances by a purported consent of a relative or other person or body. Such refusal will be regarded as remaining in force even though I may be unconscious and/												
	or affected by medication/ stroke, or other condition rendering me incapable of												
	expressing my wishes and consent to treatment options, and the doctors(s) treating												
	me consider that SUCH REFUSAL MAY BE LIFE THREATENING.												
	That any	prod	cedure in addi	tion to tl	he investig	ation or treati	ment de	escribed	on this	form, but			
	with the exclusion of the transfusion of allogenic blood or primary blood components, will only												
	be carried out if necessary and in my best interests and can be justified for medical reasons.												
	That details of my treatment, and any consequences resulting, will not be disclosed to any												
	source without my express consent or that of my instructed agent(s) unless required by												
	law.												



Patient Details (or pre-printed label)											
Patient Surname/ family name											
Date Of Birth	Male		Female								
CHI:	1										
Please indicate your requirements by ticking appropriate boxes -:											
	Acce	pt	Refuse								
Primary Blood Components											
Red Blood cells											
Fresh Frozen Plasma (FFP, plasma)											
Platelets											
White cells (Granulocytes)											
Products containing a minor blood fraction											
Cryoprecipitate											
Albumin											
Intravenous immunoglobulin											
Anti-D immunoglobulin											
Other immunoglobulins e.g. tetanus											
Procedures involving my own blood											
Cell salvage											
Acute normovolaemic haemodilution											
Renal Dialysis											
Plasmapheresis											
Blood radio-labelling											
Recombinant products – not blood sourced											
rFVIIa (Novoseven)											
Erythropoietin											
Others e.g. FVIII											
Other Components/Procedures (please specify)											
	•										
Patient		, ,									
I confirm that I have indicated above my wishes. I accept detailed above.	or retuse the l	olood comp	oonents & proce	edures as							
Signature: Print name:		Date:									
Doctor											
Signature: Print name:		Date:									



## NOTES OF GUIDANCE

## Medical Practitioner



The Doctor completing this form should be suitably qualified and experienced to explain the consequences of the patient's refusal for blood or blood products in relation to the patient's condition and proposed treatment. This should ideally be the consultant in charge or the practitioner who is operating or supervising the treatment.

Information must be given in a way the patient can understand with important risks and possible alternatives explained. He/she should be given sufficient time to consider the issues and have the opportunity to ask questions.

Where the treatment requires an anaesthetic, (especially where blood loss is likely to be an issue), it is advisable to discuss this with the consultant anaesthetist involved in the case at the earliest opportunity to ensure optimal care.

In all adult cases, advanced directives carry the weight of law in conscious and unconscious patients.

The patient can withdraw this declaration to refuse blood or blood products at any time following completion of this form. If this occurs it is recommended that the patient's wishes are witnessed by another Health Professional and documented clearly in the medical notes.

Where the patient is under 16 years it is advisable to refer to the relevant sections of the Refusal of Blood Transfusion Policy – suitable for Jehovah Witnesses (part 3) and Policy on Consent to Healthcare Assessment, Care & Treatment (part 8) and seek any further advice as may considered appropriate in the circumstances.

Please refer to the Refusal of Blood Transfusion Policy for further information and guidance.