

CONSENT FOR USE OF BLOOD PRODUCTS

Affix patient labe	<u>l here</u>		
I	(Medical O	Officer/Midwife giving information to patient) have	ve
given and explained information in	relation to the adminis	stration of blood components/blood products	to
	(Person rece	eiving the information).	
This information included: The purpose of giving blood comp The type of blood, blood compone The risks associated with their use Available alternatives to the use of	nts or blood products to be	e used;	
answered them appropriately and to the	ne best of my ability.	ns and where questions have been asked I have	ve
Signature (Medical Officer)	Designation	Date	
	PATIENT CON	ISENT	
	-	stand about the explanation or ask before signing this form.	
	administration of blood co	person giving consent) have been provided wi omponents/blood products. I have been given the actorily answered.	
		_ (type of blood/blood product to be used). I als may be found necessary during the use of these	
I give this consent for myself/for		who is my	

NOTE: This consent is for the total number of blood products administered for the ongoing management of a particular disease, disorder or pregnancy.

Author: Dhana Gounder Authoriser: Sarah Morley Approver: Fidelma Murphy

Signature (Person giving consent)

Effective Date: 21/04/2022 Page 1 of 1
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Refer to document(s): 111M001, 101M036

Date