



April 2024

Blood Bank Request Form / Transfusion Record Form Changes

Effective 6th May 2024, Pathlab will be moving to using the nationalised standard pre transfusion Blood Bank Request Form and Transfusion Record Form. These forms replace the local forms currently used in Rotorua and Taupo Hospitals. This standardisation will make it easier for clinical staff as they move between hospitals.

This has become urgent due to NZBS adopting the ISBT numbering system for blood components. From 09/06/24 a 14-digit unit number will be introduced for blood components. **There is no requirement for this number to be transcribed onto the Transfusion Record Form and doing so will only introduce errors leading to unnecessary delays.**

This will allow familiarisation for clinical staff and laboratory staff before the ISBT component numbers are introduced.

The new forms can be ordered via your current process.

Additionally, the consent form for prescription of blood components or products to issue blood still **MUST** be completed by the clinical staff who will hold the responsibility for these steps to be completed. However, **NZBS and the rest of Pathlab do NOT require the sighting of the consent forms and prescriptions** so these forms are no longer required by the laboratory for blood to be issued.

Mark Howard
Service Lead, Transfusion

Dr Gustavo Faulhaber
Haematologist

CLINICAL UPDATE

Please ensure all members of your institution
receive a copy of this clinical update.

Patient:			
Family Name		Place of Surgery / Transfusion	
Given Names		Date of Surgery / Transfusion	
NHI	Date of Birth	Gender	
If patient is a neonate please provide mother's details:		Family Name	NHI
		Given Names	Date of Birth
Diagnosis / Indication for transfusion		Consultant	
History affects validity of the blood sample for the provision of red cells for transfusion – please see over			
You must complete all 3 questions – please tick: Transfused in the last 3 months? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown <input type="radio"/> NA			
Pregnant in the last 3 months? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown <input type="radio"/> NA			
RhD Ig given in the last 3 months? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown			
Blood Bank Tests: If the request is urgent please phone the Blood Bank			
<input type="radio"/> Group & Antibody Screen <input type="radio"/> Cord Group & DAT <input type="radio"/> Direct Antiglobulin Test <input type="radio"/> Other <input type="radio"/> 1st Antenatal Group & Screen <input type="radio"/> Subsequent Antenatal Group & Screen <input type="radio"/> Antenatal Antibody Titre			
Component Required:		Fractionated Product Required:	
Red Cells	Adult	Number	Date and Time Required
	Neonatal		
Platelets	Adult	RhD Immunoglobulin	Dose <input type="radio"/> 625 IU <input type="radio"/> 250 IU
	Neonatal	Prothrombinex-VF	
FFP	Adult	IVIg* <small>*Requires prior approval</small>	
	Neonatal	Albumex 4	
Cryoprecipitate		Albumex 20	
Other Components		Hep. B Immunoglobulin	
		Other	
Requester			
Signature	Print Name (if not the labeller)	Contact No. / Pager	Date
MANDATORY DECLARATION by Person Collecting the Sample – Failure to complete may result in sample rejection			
I certify that I collected the sample(s) accompanying this request from the patient named above			
I confirmed the identity of this patient by direct enquiry and / or inspection of their wristband			
I labelled the sample(s) by hand immediately after collection at the bedside and in the presence of the patient			
Collector			
Signature Mandatory	Print Name (if not in the box above)	Contact No. / Pager	Time
			Date

		Patient Label Name: _____ NHI: _____ DOB: _____ Address: _____																																											
Use for collection of Blood products from Blood Bank																																													
STOP Transfusing incompatible blood can cause death. If you have ANY concerns with checking or transfusing blood products, please contact the Transfusion Laboratory																																													
<ul style="list-style-type: none"> Checks must be performed at the bedside Both staff must check independently The transfuser must be a certified blood product nurse, midwife, doctor or anaesthetic technician who will be responsible for monitoring the transfusion The checker may be any of the above or an anaesthetic technician or enrolled nurse 		<ul style="list-style-type: none"> Both must confirm patient identify (wrist label, swing label and prescription) are all the same patient The right product for the right person Prior to transfusion check that <ul style="list-style-type: none"> The written consent for transfusion is current T, P, R and BP have been recorded on the observation chart within 60mins before transfusion starts 																																											
Product(s) required from blood bank:																																													
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BLOOD PRODUCTS ADMINISTRATION RECORD

CLINICAL UPDATE

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