Austin Laboratory AUSTIN HEALTH (APA) 145 Studley Road HEALTH | Pathology Heidelberg 3084 A/Prof K.Ireland-Jenkin, Dr M.Leroi, Dr Q. Lam. Dr C. Hogan

Reason patient cannot sign

Signature_



REQUEST FOR BLOOD / BLOOD PRODUCTS

RIGHT BLOOD RIGHT PATIENT RIGHT REASON

Date:

Time:

Patient Details ∪.	R. No		Hosp		Ward		Lab numb	per (lab use only)	A/3815
		DATE/TIME REQUIRED:		IT or Bleeding	Within 2 ho	ours 🗌	NOT Urgent		
Postcode: Tel No		Tests Requested: Group & Screen is valid for 3 days							
MEDICARE or DVA (Repat) Number Clinical Notes / Indication for Transfusion Specify indications below									
Transfusion History (must be completed for extended expiry) Yes No Unknown		Specify indicatio Irradiated CMV Negative	ns below				Se		
			Surname:(print).	octor (Must	Sign & Date)	No:		Pager No: Initials:	
Patient status at the time of service or when the specimen was collected Private patient in a private hospital or approved day care facility Public patient in a recognized hospital Outpatient of a recognized hospital Outpatient of a recognized hospital MEDICARE ASSIGNMENT FORM Section 20A of the Health Insurance Act 1973 (not required for inpatients) I assign my rights to benefits to the approved practitioner who will render the requested pathology service(s).			Person Drawing Blood Specimen (must sign and date) I(PRINT NAME) certify that the blood specimen(s) accompanying this request was drawn from the patient named above and I established the identity of this Patient by direct inquiry and/or inspection of wrist band, and immediately upon the blood being drawn I labelled the specimen(s) and signed the tube(s), including date and time.						

SIGNED

Date:__/_/

SPECIAL REQUIREMENTS / ATTRIBUTES FOR CELLULAR BLOOD COMPONENTS (red cells and platelets)

In general, most patients do not require cellular blood components (red cells or platelets) with special attributes. If you are not sure whether your patient requires special blood components, please contact the **Austin Haematology Laboratory Registrar on 9496 5981,** or if afterhours contact the **on-call Haematology Registrar via Austin Switchboard on 9496 5000.** The responsibility for the appropriate ordering of blood products lies with the requesting clinician.

Please refer to the National Blood Authority website: http://www.blood.gov.au for Patient Blood Management Guidelines on the appropriate use of blood components; or refer to the Australian Red Cross Blood Service website: http://www.transfusion.com.au for detailed explanation and recommendations regarding the use of special blood components.

In an emergency situation and / or if there is significant bleeding, a life saving transfusion should not be withheld if the suggested special component is not readily available.

PATIENT POPULATION	CMV NEGATIVE BLOOD COMPONENTS INDICATIONS	IRRADIATED BLOOD COMPONENTS INDICATIONS			
Haematology patients	Haematology patients will NOT require CMV negative blood components regardless of their CMV serology results	Allogeneic and autologous haematopoietic stem cell transplant Autograft: min 6m post transplant Allograft: min 12m post transplant; long term if active GVHD Haematological malignancy and receiving/completed chemotherapy within the last 12 months			
Oncology patients	Only if specifically requested by the treating consultant	Aplastic anaemia on immunosuppression Malignancy and receiving long term or high dose steroid therapy Alemtuzumab (note also used for some neurology patients)			
Solid organ transplant	Required if both the recipient and donor of the transplanted organ are CMV negative	Not required			
Pregnant women	Require CMV negative blood components regardless of their CMV status In an emergency, If CMV negative components are not available, CMV-unknown components may be given as universal leucodepletion confers a high level of safety in preventing CMV transmission	Not required			
Neonatal and Interuterine Transfusions	Required for all	 Low birth weight <1500g until 40 weeks (corrected gestational age) Intrauterine transfusion and subsequent exchange transfusions Prior intrauterine transfusion until 6 months corrected gestational age 			
Congenital cellular immunodeficiency (Excludes HIV/AIDS)	Not required	Required for all patients			
Patients receiving nucleoside analogues	Not required	Required for all patients			

^{*}Extended expiry for Group and Hold can be requested in a pre-operative setting, for surgical patients, who have not received any blood transfusion, and who have not been pregnant, in the preceding three months.

PRIVACY NOTE: The information provided will be used to assess any Medicare benefit payable for the service rendered and to facilitate the proper administration of government health programs, and may be used to update enrolment records. Its collection is authorised by provisions of the Health Insurance Act 1973. The information may be disclosed to the Department of Health and Ageing or to a person in the medical practice associated with this claim, or as authorised/required by law.