







Dr Jeremy McComish 230814HH Prof James McCluskey 0170567B

NATA Accreditation No: 18808 ASHI Accreditation No: 12-9-AU-05-1

Request for HLA-HPA Compatible Platelets - Clinical Information and Investigation Request Form

Instructions for completing this form. The form can be completed electronically prior to signing:

- 1. Complete appropriate sections on Page 1, 2 and 3.
- 2. Collect the appropriate sample tubes as specified on Page 4
- 3. Send samples with completed copy of Pages 1, 2 and 3 only.

<u>Note:</u> Referring clinician to complete all parts of this form when requesting HLA or HPA compatible platelets for a specific patient.

Testing Laboratory							
Please send samples to							
Contact details	5						
Phone			Emai	I			
Patient details							
Last Name			First I	Name			
Gender			MRN	/UR		DOB	
Referring Clini	Referring Clinician details						
Name							
Signature			Phon	e			
Address			Emai	I			
Tick if a hard copy report is required							
Referring Labo	oratory name						
Phone			Emai	l			
Name of perso	n completing the for	m (if different froi	n abov	/e)			
Name							
Phone			Emai	l			
Sample collect	tion						
Collector's name			Date collec	& time of tion			
Collector's signature							
Specimen type (s) include	EDTA	ACD	Serur	n (clot)			
Please attach sample label/barcode				Patient's signa	ature & Date		









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Patient details									
Last Name				First N	lame				
Gender			MRN/	JR	DOB		В		
Clinical Diagnosis and History									
Clinical Inform	ation (Tick	as required)		Clinic Inforn as req	nation (Tick	Clinical History (Add any other relevant details)			
Sepsis				Anti	ibody				
Fever					apy				
Bleeding				e.g. Rituximab					
Coagulopath History of tra	-				ecent IVIG				
Antibiotics (/					usion,				
					vide date.				
					evious gnancy				
	_				yes, how				
Splenomega	-	Add dataila)		ma	any)				
Recent Che	motherapy (/	400 details)							
HAPLO BMT Only	HLA Typin	lg: Yes	No		lf Yes, date t	ested:			
HPC Transplant Date				a mul	nt blanted with tiparous e donor?	Ye	es No	Unsure	
Туре				Blood (Dono	l Group or)				
Deficit	А	В	0		AB F	RhD Ne	gative F	RhD Positive	
Patient blood group	Patient blood group <u>IMPORTANT NOTE:</u> please include a validated/authorised blood group report for the patient when submitting this request.					or the patient			
Platelet Increments – At least 2 post transfusion Increments <u>MUST</u> be provided to determine refractoriness.									
Data of	transfused platelets		Blood grou	o of			Post	count	
Date of Transfusion			platelets transfused		Pre-count		10- 60 mins	24h	
*	Apheresi	s Pooled	*				*		
*	Apheresi	s Pooled	*				*		
Current Platelet	t Count						·	·	









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Patient details							
Last Name			First Name				
Gender			MNR/UR	[ООВ		
Current WCC			Red blood cell detected?	antibodies	Yes, Specify		
Neutrophil Count					No		
Request detail	S						
Platelets	HLA HPA		Is the patient	Yes	No		
compatible for			pregnant?	platelets will on currently pregna during labour ar available. All pla	body non-reactive y be considered for ant recipients (but not nd delivery) and only if atelets are leucodepleted ly prevents transfusion CMV infection		
	ABO compatible platelets will be provided where possible.						
	If ABO compatible platelets are not available, low titre anti A/B components will be provided.						
	Are RhD negative platel	et required? (for	r female recipient	of child bearing p	otential)		
Pland group	Yes No)					
Blood group product	NB: If patient is receiving ABO mismatched bone marrow transplant please select acceptable blood group/s						
	ABO 1 st Choice		RhD 1 st Choice				
	ABO 2 nd Choice		RhD 2 nd Choice				
	ABO 3 rd Choice				1		
Required testing on patient	HLA Typing Yes No	HLA Ab Testi Yes	ng HP No	A Typing Yes No	HPA Ab Testing Yes No		
Note: A Lifeblood Transfusion Nurse will discuss the transfusion requirements if HLA Compatible Platelet support is indicated based on test results.							

Select State contact details to return completed forms or for any urgent request.

State	
Fax	
Phone	
Email	





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Platelet Transfusion Refractoriness (PTR) Sample Collection Guideline

Investigation request and samples		Special instructions and indicative turnaround time	Storage and transport instructions	
Platelet Transfusion Refractoriness (PTR)		EDTA from pre-transfusion collection is acceptable	Store and transport at either room temperature or 4°C.	
	8 mL EDTA or ACD and 12 mL serum (clot)	Note: Laboratory turnaround time is 1 – 3 working days.		