

Platelet Refractory Patient Transfusion Request

Clie	ent to Complete	9									
Hospital:			Phone #:	ne #: Fax			Acct #	:			
Red	uest Date:		Time: R	equested by	(Nam	e of Hospital Staff	i):				
Patient Last Name			First Name		Date	Medical Recor	d Number	ABO/Rh	Gender		
Clinical Status: Diagnosis:						Current Platelet Count: x 10³/uL					
М	ost Recent Pla		story: Active Bleeding:	□ No □ Y							
Date			atelet Product(s) Given			1-Hour Post-Transfusion Platelet Count 10 ³ uL					
L											
Ar	e any of the foll	owing factors believe	d to contribute to the plat	elet refracto	riness	? (check all that ap	oply):				
	Fever Infe	ction	herapy Splenomegal	ly DIC	Othe	er:					
Rec	uest is: ST/	AT Routine									
NO.	TE: If testing h	nas been performed	by another lab, please	send result	s with	this request.					
	Check here if yo	ou wish to consult with	n a Vitalant Field Medical	Director.							
		Testing and Produc	t Request Options		Sample Requirements						
NC	TE: Local opti	ions vary; consult y	our center for more info	ormation	NOTE: Gel separator tubes are NOT acceptable						
		Testing R	equests								
		ody screen (immune does not distinguish b	e adherence assay): etween – HLA* Class I a	nd HPA*	10 m	nL EDTA tube(s)					
		ody screen (ELISA): distinguishes betwee	n – HLA Class I and HPA	A antibodies	10 m	nL red top clot tub	e(s)				
	HLA Class I a	ntibody screen: Flo	w cytometry assay		10 m	nL red top clot tub	e(s)				
		ass I antibody identific	n: Flow cytometry assay cation assay when antibo		10 m	nL red top clot tub	e(s)				
		low-resolution gene	<u>otype</u>		20 m	nL EDTA or ACDA	tubes – U	nspun/unope	ened		
☐ Human platelet antigen genotype					10 mL EDTA tube(s) – Unspun/unopened						
		Product R	equests								
	incompatibilitie		erence assay): Detects HLA Class I and/or HPA a		10 m	nL EDTA tube(s)					
	SEARCH for H (No. of units I	ILA-compatible don needed:)	ors/products		No s	sample needed if t	esting was	done previo	usly		
	-		CMV-Serone No Other Needs:	_			I: Yes eed by Date				
**T	he term "HLA-d	compatible" refers to	n; HPA = Human donors/products selected avoidance of HLA-A and	based upon	: (1) th	ne matching of HL	A-A and -B		tween the		

Center Information:



Instructions

- Please contact the local Vitalant blood center to arrange sample pick up and/or shipping. Contact information can be found at: www.vitalant.org
- Fill out this request form as completely as possible. Attach copies of any related work previously performed at your facility or at other special testing labs.
- Label all samples with:
 - Full patient name,
 - Second unique patient identifier number, and
 - Date collected.
- Incorrectly or unlabeled specimens may be rejected.
- Please update Vitalant with any changes to the status of this request.

Important Information

- For patients who require ongoing platelet transfusion support and who have a history of defined HLA antibody specificities, a repeat antibody identification is recommended at least every 1-to-2 months.
- The use of platelet crossmatch should be reserved for patients:
 - Who require special platelet products more urgently than can be achieved through "HLA-centric" means (often while waiting for the HLA-related testing and donor/product search to be completed), and/or
 - Whose etiology for refractoriness is suspected to be due to HPA incompatibilities.
- Communicate with the on-duty Vitalant Medical Director (reachable through Hospital Services department) for assistance ordering testing, interpreting the results, and/or deciding what type(s) of special platelet products to order. The Medical Director may also help identify additional means by which to support refractory patients' platelet transfusion-related needs.

Vitalant to Complete

DIN/	Component Codes	Date/Time/EC	Unit Status					
	ABO/Rh	CMV Status	Center	WIP	Verified Date/EC			