

Versiti Provider Signature:

Cellular Therapy Product Collection Request

Instructions: The requesting provider or designee is responsible for completing all fields of this form. The Versiti provider is responsible for reviewing the order. Please scan or fax the completed form using state specific contact information below with the Autologous Donor Suitability Determination or the Allogeneic Donor Eligibility Determination and supporting documents.

State	Address	Email		Pnone	Fax	
Wisconsin	638 N. 18 th St, Milwaukee, WI, 53233	WI-CellCollection@versiti.org		414-937-6154	414-933-6833	
Michigan	1036 Fuller Ave NE, Grand Rapids, MI 49503	MI-CellCollection@versiti.org		616-233-8569	616-233-8671	
Indiana 3450 N. Meridian St, Indianapolis, IN 46208 IN-Ce		IN-CellC	Collection@versiti.org	010-233-8309	010-233-8071	
				Tentative Collection Start Date:		
AUTOLOGOUS DONOR/ALLOGENEIC RECIPIENT INFORMATION						
Apply Hospital Label or complete: Diagnosis:						
Donor/Recipient Name:			Sex: M F ABO/Rh:			
Date of Birth:			Height: inches cm			
Medical Record #:			Weight: kg			
ALLOGENEIC DONOR INFORMATION (DO NOT COMPLETE FOR AUTOLOGOUS DONORS)						
Apply Hospital Label or complete:			Sex: M F ABO/Rh:			
Donor Name:			Height: i	nches cm w	eight: kg	
Date of Birth:					•	
Medical Record #:			Has donor or family been made aware of the			
			availability of a don	or advocate?	YES NO	
PRODUCT TYPE						
HPC, Apheresis: Target Dose: x 10 ⁶ CD34/kg MNC, Apheresis						
COMMERCIAL OR CLINICAL PROTOCOL, IF APPLICABLE						
Commercial: List Company/Product Name:						
Clinical/Research: List Sponsor/Protocol Name:						
DONOR INFORMATION AND RECORDS						
All DONORS: Are there communication barriers or issues that pertain to the safety of the collection procedure?						
YES* NO *If yes, describe:						
ALL DONORS: Vein Assessment Performed by:				Peripheral veins acceptable		
Central Venous Catheter (CVC) Ultrasound Guided Peripheral Access (schedule back-up CVC appt.)						
ALL DONORS: Is there a signed consent on file? YES NO						
Female Donors Only: Date Pregnancy Test was completed:			N/A- Not indicated			
HPC Donors Only: Date Hemoglobinopathy assessment was completed:						
HPC Donors Only: Start Date of planned mobilization regimen:						
PROCESSING ORDER, IF APPLICABLE						
Tentative Infusion Date: Process for Fresh Infusion Process for cryopreservation and storage						
Allogeneic Products Only: Perform CD3 counts. Prepare product for Donor Lymphocyte Infusion if counts are sufficient.						
Plasma Depletion or Volume Reduction per Versiti policy.						
AUTHORIZATION SIGNATURES						
Form Completed by: Ordering Provider:						
Ordering Provider Signature:		Date:				

 Document No:
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 Version:
 1
 Page:
 1 of 1

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