

APHERESIS PRODUCT TAG

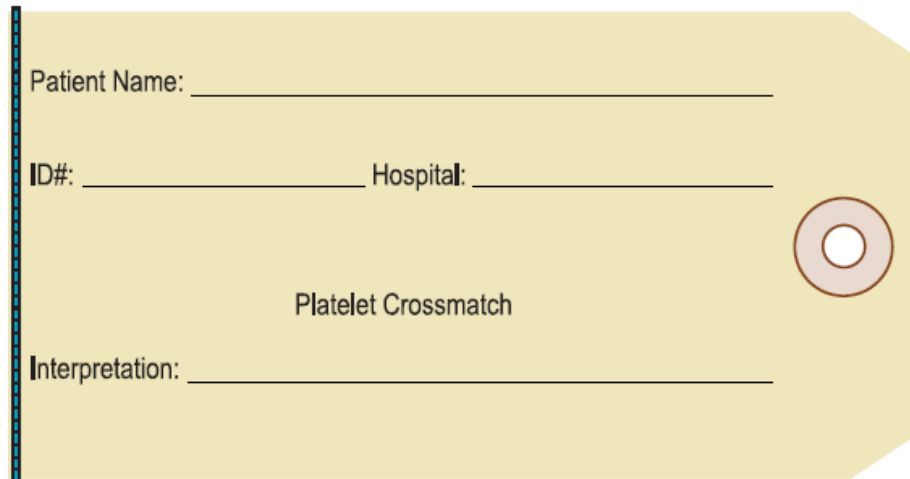
Color is Manila

FRONT



The front view of the tag is a light beige rectangular label with a rounded right side. On the left edge, there is a vertical blue and white dashed line. The word "APHERESIS" is printed vertically in bold black letters. Below it, the text "Unit Number" is followed by a horizontal line for writing. On the right side, there is a circular hole with a red outline. To the left of the hole, the text "Carter BloodCare" is printed vertically. To the right of the hole, the text "APL100", "Version: 04", and "Effective Date: 06/24/2014" is printed vertically.

BACK



The back view of the tag is a light beige rectangular label with a rounded right side. On the left edge, there is a vertical blue and white dashed line. The text "Patient Name:" is followed by a horizontal line for writing. Below that, "ID#: _____ Hospital: _____" is followed by two horizontal lines for writing. In the center, the text "Platelet Crossmatch" is printed. At the bottom, "Interpretation:" is followed by a horizontal line for writing. On the right side, there is a circular hole with a red outline.

APHERESIS PRODUCT TAG

Color is Manila

FRONT


APHERESIS

Unit Number _____

**PLATELET
CROSSMATCHED
IRRADIATION REQUIRED**

APL 100
Version: 04
Effective Date: 06/24/2014

Carter BloodCare




BACK

Patient Name: _____

ID#: _____ Hospital: _____

Platelet Crossmatch

Interpretation: _____



CARTER BLOODCARE
2205 HIGHWAY 121 BEDFORD, TX 76021

Facility: Bedford Central Office Client: CBC - Carter Blood Care

Unit #: W035208656565-* Exp. Date/Time: 9/2/08 23:59

Unit Group/Type: O NEG

Component: E4545 AS-3 RED BLOOD CELLS LEUKOCYTES REDUCED

Patient: DOE,JANE

BBID: 102

Hospital #: 1022

Case #: 082108DOJA

Patient Group/Rh: O NEG

Doctor: Physician,Staff

Crossmatch: COMP

Crossmatch Expires: 8/24/08 23:59

By: 6542 8/21/08 13:31

Unit Antigens:

Unit Attributes: Leuko-reduced

The Patient's Name, Hospital Number, Product Type, Unit Number, and additional applicable identifiers were verified in patient's presence prior to transfusion according to institutional policies.

Signature: _____ Signature: _____

Transfusion Record Date: _____

Time From _____ AM/PM To _____ AM/PM

Temperature - Pre _____ Post _____

Amount Transfused _____ Transfusion Reaction: YES NO

If reaction occurs - Stop Transfusion, Initiate Report

Rev. 12/15/2004

CARTER BLOODCARE
2205 HIGHWAY 121 BEDFORD, TX 76021

Facility: Bedford Central Office

Client: CBC - Carter Blood Care

Unit #: W035208565656-4

Exp. Date/Time: 8/23/08 23:59

Unit Group/Type: O NEG

Component: E2827 PLATELETS LEUKOCYTE REDUCED

Patient: DOE,JANE

BBID: _____

Hospital #: 1022

Case #: 082108DOJA

Patient Group/Rh: O NEG

Doctor: Physician,Staff

Crossmatch Not Required

By: _____

Unit Attributes: Irradiated

Leuko-reduced

The Patient's Name, Hospital Number, Product Type, Unit Number, and additional applicable identifiers were verified in patient's presence prior to transfusion according to institutional policies.

Signature: _____

Signature: _____

Transfusion Record Date: _____

Time From _____ AM/PM To _____ AM/PM

Temperature - Pre _____ Post _____

Amount Transfused _____ Transfusion Reaction: YES NO

If reaction occurs - Stop Transfusion, Initiate Report

Rev. 12/15/2004

Phone: 817-412-5740 Fax: 817-412-5749

SAMPLE TYPE AND APPROPRIATE SAMPLE LABELING

Required Sample(s) 5 - 15 mls (EDTA) - (NO Serum Separator) Samples were collected using a validated electronic ID system? Yes No

Patient Name (Last, First): _____ Patient ID: _____ Ordering Physician: _____	Sample(s) Collected Date/Time/By: _____ Requesting Facility: _____ Blood Bank ID (if app.): _____	Order Status (Circle One) <p style="text-align: center; font-weight: bold;">STAT ASAP ROUTINE</p> To Be Delivered by Date/Time: _____
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Date of Birth: _____ Gender: (Mark One) <input type="checkbox"/> M <input type="checkbox"/> F <input type="checkbox"/> Other: _____ Diagnosis: _____ Blood Bank ID Stickers Included (circle one): Yes No N/A	FOR CBC USE ONLY: Issued to the Distribution Department												
Transfusion History Transfused within last 3 months NO YES UNKNOWN If yes, date of last red cell transfusion: _____	<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">DATE</th> <th style="width: 10%;">TIME</th> <th style="width: 15%;">Clerical Check</th> <th style="width: 15%;">Visual Inspection</th> <th style="width: 10%;">Tech 1</th> <th style="width: 10%;">Tech 2</th> </tr> </thead> <tbody> <tr> <td> </td> <td> </td> <td> </td> <td> </td> <td> </td> <td> </td> </tr> </tbody> </table>	DATE	TIME	Clerical Check	Visual Inspection	Tech 1	Tech 2						
DATE	TIME	Clerical Check	Visual Inspection	Tech 1	Tech 2								

Pregnancy History Number of Pregnancies: _____ Pregnant Now? (circle one) Yes No RhIG (circle one) NO YES If yes, date of RhIG administration: _____ Date Due: _____	Special Instructions (Circle applicable) Irradiated Sickle Cell Negative CMV negative
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TESTING REQUESTED (check applicable) _____ Type & Screen and Crossmatch _____ Type & Screen Only _____ Blood Type (ABO/RH) _____ RHIG Evaluation (RH immunoglobulin) _____ Additional Crossmatched Units (specify number) _____ Other (specify) _____	PRODUCT REQUESTED (indicate number needed) _____ LRBC(s) If applicable (circle one) Autologous Directed _____ APHERESIS PLATELET(s) _____ FFP(s) _____ Cryo(s) _____ Pack each unit separately	Other (specify) _____ Infusion Sets (indicate number needed) / Misc. _____ Y type filter (red cells) _____ Component filter (plasma, platelets, cryo) _____ Blood Bank Armbands (10 per box; specify number of boxes) _____ Forms _____ Other: _____
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Pretransfusion Criteria (Check applicable)*

*For compliance with regulatory agencies this section must be completed for any blood component requested.

RED BLOOD CELLS (RBCs)	PLATELETS (PLTs)	FRESH FROZEN PLASMA (FFP) / CRYO
_____ Current Hgb or HCT* (Indicate All of the Following that Apply) _____ Pre-Surgery (Pre-op) _____ Hemoglobin ≤ 8 g/dl or Hematocrit ≤ 24% _____ Symptoms of Anemia _____ Active bleeding/Acute blood loss _____ Other (specify) _____	_____ 10 ³ cells / ul (Current Platelet Count*) (Indicate All of the Following that Apply) _____ Platelet count of 20,000 / ul or less _____ Platelet count of < 50,000 / ul, bleeding or planned surgery in 24 hours _____ Platelet dysfunction and bleeding / surgery planned _____ Platelet count < 50,000 after blood loss _____ Other (specify) _____	_____ Active bleeding _____ Coagulation Deficiency _____ INR > 1.5 _____ Other (specify) _____

UNCROSSMATCHED OR INCOMPATIBLE PRODUCT RELEASE

UNCROSSMATCHED

This is an acute emergency. The current status of the patient's condition dictates that these units are needed with sufficient urgency to waive the performance of the compatibility testing by Carter BloodCare prior to shipment or administration. The benefits of the product(s) being transfused outweigh the risk(s) involved.

Please sign and fax to the Reference and Transfusion Lab at Carter BloodCare - Fax #817-412-5749

Physician's Signature: _____		Date: _____	
Physician's Name Printed: _____		Facility: _____	
RN may sign as instructed by physician: _____ (If signed, needs physician signature within 24 hours)			
Unit Number	Product code	ABO/Rh	Expiration Date

R&T Tech: _____ Date: _____

Patient Name: _____
Patient Account Number: _____
Patient ABO/Rh (if known): _____

INCOMPATIBLE

The current status of the patient's condition dictates that these units are needed and I acknowledge that the product(s) listed below is/are incompatible with this patient. The benefits of the product(s) being transfused outweigh the risk(s) involved. I understand that close monitoring of this patient for evidence of hemolysis or other transfusion reaction should occur throughout the transfusion period.

Please sign and fax to the Reference and Transfusion Lab at Carter BloodCare - Fax #817-412-5749

Physician's Signature: _____		Date: _____	
Physician's Name Printed: _____		Facility: _____	
RN may sign as instructed by physician: _____ (If signed, needs physician signature within 24 hours)			
Unit Number	Product	ABO/Rh	Expiration Date

Reason for Incompatible Blood Release

- Reactivity due to auto antibody
- Antibody to a high frequency antigen
- Reactivity to a drug antibody

R&T Tech: _____ Date: _____

Patient Name: _____
Patient Account Number: _____

UNTESTED PRODUCT RELEASE

Patient Name

Identification Number

Facility

ABO/RH (if known)

Reason for Product Release:

Checked Test Procedures Not Performed

Collected from a Donor known to be Negative on
_____ for Checked Test Procedures

Date

_____ Anti-HIV 1/2

_____ Anti-HTLV I/II

_____ NAT - HIV1/HCV/HBV

_____ HBsAg

_____ STS (Syphilis)

_____ NAT - WNV

_____ Anti-HBc

_____ IAT

_____ NAT - ZIKA

_____ Anti-HCV

_____ CMV

_____ Bacterial Detection (Platelets)

_____ Anti-T.cruzi (Chagas') or Previously Tested

UNIT NUMBER

PRODUCT CODE

PRODUCT DESCRIPTION

ABO/RH

Form Completed by: _____ Date: _____

Current conditions dictate that these units are needed with sufficient urgency to waive the performance of the above tests by Carter BloodCare prior to shipment or administration.

Requesting Physician or Medical Director Signature _____ Date: _____

HLA MATCHED

Patient _____

ID No. _____

Facility _____

Unit No. _____

HLA MATCHED GRADE _____

IRRADIATION REQUIRED BEFORE INFUSION

RTL 422.01

Patient Name: _____

ID#: _____ Hospital: _____

Platelet Crossmatch

Interpretation: _____

APHERESIS

Unit Number _____

PLATELET CROSSMATCHED

APL100
Version: 03
Effective Date: 11/11/2008

Carter BloodCare

RTL207.01A Confirmed Antigen Typing (green label)

_____ has been antigen tested and found positive or negative for the following:

C _____	K _____	Jka _____	Lea _____	P1 _____
c _____	k _____	Jkb _____	Leb _____	_____
E _____	Fya _____	S _____	M _____	_____
e _____	Fyb _____	s _____	N _____	_____

Carter BloodCare

RTL207.01C Molecular Matched Antigen Typing

**Molecular Matched
Antigen Typing**

DIN: _____

Donor has been antigen tested using a Molecular Genotyping Assay and predicted negative for the following:

_____ Neg.	_____ Neg.
_____ Neg.	_____ Neg.
_____ Neg.	_____ Neg.
_____ Neg.	_____ Neg.
_____	_____
_____	_____

RTL207.01C
Version: 02
Carter BloodCare Effective Date: 01/20/2020

_____ has been antigen tested and found positive or negative for the following:

C _____ K _____ Jka _____ Lea _____ P1 _____

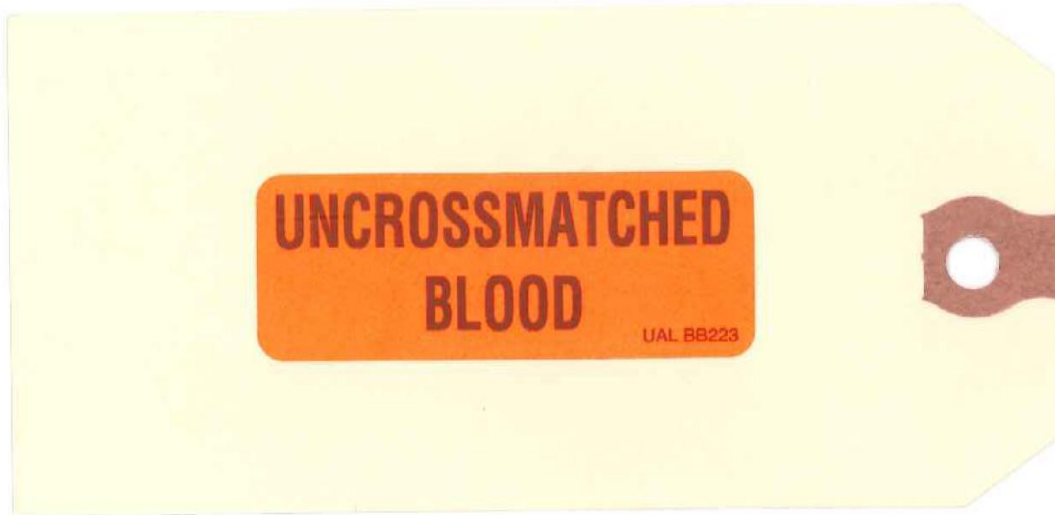
c _____ k _____ Jkb _____ Leb _____ _____

E _____ Fya _____ S _____ M _____ _____

e _____ Fyb _____ s _____ N _____ _____

Carter BloodCare

RTL214.01
Emergency Release Uncrossmatched Blood Label



**EMERGENCY RELEASE TIE TAG – “COLLECTED FROM A DONOR KNOWN TO BE
NEGATIVE ON”**

EXAMPLE: Collected from a donor known to be negative on _____ for

Collected from a donor known to be
negative on _____ for

Anti-HIV-1 / 2	STS
HBsAg	IAT
Anti-HBc	CMV
Anti-HCV	NAT HIV-1/ HCV/ HBV
Anti-HTLV-I/II	NAT WNV
Anti-T. cruzi (Chagas')	Negative or Previously Tested

RTL214.03A
Version: 07
Effective Date: 07/21/2021

EMERGENCY RELEASE TIE TAG – “TESTING NOT PERFORMED”**EXAMPLE: Testing Not Performed****TESTING NOT PERFORMED**

Anti-HIV-1 / 2	IAT
HBsAg	CMV
Anti-HBc	NAT HIV-1/ HCV/ HBV
Anti-HCV	NAT WNV
Anti-HTLV-I/II	STS
Bacterial Detection (Platelets)	
Anti-T. cruzi (Chagas')	Crossmatch, if applicable

RTL214.03B
Version: 07
Effective Date: 07/21/2021

HLA MATCHED APHERESIS PRODUCT TAG

HLA MATCHED

Patient _____

ID No. _____

Facility _____

Unit No. _____

HLA MATCHED GRADE _____

IRRADIATION REQUIRED BEFORE INFUSION RTL 422.01