

Allocation
Label
on unit

ZZTESTSQEPIC, LABEICURV DOB 08/09/1970
MR# 61431898 AUX ID TT1236
Pt ABO/RH AB-Positive RBC AS L
W0358 17 147983 EXP 01/31/2018 2359
UNIT: AB-Posit XM: Electronic BY: 9070


W0358 17 147983 SM


8400

BloodSource, Inc.
Mather, CA 95655
FDA Registration Number 3007101484
US License Number 1776

AB

Properly identify intended Recipient
See Circular of Information for Indications,
contraindications, cautions and methods of infusion.
This product may transmit infectious agents.
Rx Only

Rh POSITIVE

VOLUNTEER DONOR


E0336V00

 Expiration
Date
0180312359

RED BLOOD CELLS
ADENINE-SALINE (A8-1) ADDED
LEUKOCYTES REDUCED
From 500 mL CPD Whole Blood
Store at 1 to 6 C

31 JAN 2018

 Neg for Zika Virus
by Investigational NAT

Prt Date: 01/16/2018

TRANSFUSION SERVICE USE

Name: ZZTESTSQEPIC, LBEICURV
 MR# 61431898 DOB 08/09/1970 Acc No: T10002905
 ABO/Rh: AB-Positive AUXIL ID: TT1236 Loc: RVDIXA Ord Phy: GO,ORSON DY

UN# W0358 17 147983 RBC AS Leukoreduced DIV: 00
 ABO/Rh: AB-Positive EXP: 01/31/2018 2359 VOL: 325
 XM: Electronically Compat 9070

BBLR

ISSUE

ISSUED & INSPECTED BY _____ DATE _____ TIME _____
 Signature

PERSON ACCEPTING UNIT _____
 Signature

TRANSFUSIONIST

TWO AUTHORIZED PERSONNEL MUST VERIFY EACH UNIT IMMEDIATELY PRIOR TO TRANSFUSION AT THE BEDSIDE

• I CONFIRMED RECIPIENT'S IDENTITY BY ID WRISTBAND	• I CONFIRMED RECIPIENT'S IDENTITY BY ID WRISTBAND
• RECIPIENT IDENTITY CONFIRMED ON THIS FORM AND LABEL ON UNIT TO BE GIVEN	• RECIPIENT IDENTITY CONFIRMED ON THIS FORM AND LABEL ON UNIT TO BE GIVEN
• DONOR NUMBER ON UNIT AND THIS FORM ARE THE SAME	• DONOR NUMBER ON UNIT AND THIS FORM ARE THE SAME
• ANY SPECIAL REQUIREMENTS ARE CORRECT	• ANY SPECIAL REQUIREMENTS ARE CORRECT

 VERIFIED AND TRANSFUSED BY: TRANSFUSION START DATE/TIME VERIFIED BY:

RECORD OF TRANSFUSION	TIME	TEMP	PULSE	B/P	ABNORMAL REACTION		SIGNATURES
	PRE-TRANSFUSION (Within 30 min. of Start)				/		
15 MIN. AFTER START				/	NO	YES	
END OF TRANSFUSION VOLUME TRANSFUSED _____				/	NO	YES	
COMMENTS: <input type="checkbox"/> Blood warmer used							

INSTRUC-TIONS

If a reaction is suspected, immediately stop transfusion and notify Transfusion Service. Refer to Transfusion Reaction in Blood Product Transfusion Policy in PolicyStat.

RECORD OF SUSPECTED TRANSFUSION REACTION

• Does patient ID armband, donor unit label, crossmatch label and *Transfusion Record* all match? Yes No → Notify Transfusion Service STAT
 • Was any medication added to blood? No Yes. If Yes, what? _____
 • List IV solutions running in same line: _____

REACTION SYMPTOMS:	TIME	REACTION SYMPTOMS:	TIME
<input type="checkbox"/> CHILLS, WITH / WITHOUT RIGORS		<input type="checkbox"/> URTICARIA / HIVES	
<input type="checkbox"/> DYSPNEA / WHEEZING / COUGHING / SOB / CYANOSIS		<input type="checkbox"/> DARK OR BLOODY URINE	
<input type="checkbox"/> HYPOTENSION (SHOCK)		<input type="checkbox"/> DECREASED URINE OUTPUT	
<input type="checkbox"/> HYPERTENSION		<input type="checkbox"/> RASH	
<input type="checkbox"/> PAIN & LOCATION		<input type="checkbox"/> FLUSHING	
<input type="checkbox"/> ANXIETY		<input type="checkbox"/> PRURITUS	
<input type="checkbox"/> TEMP ≥38°C (100.4°F) & RISE OF ≥1°C (2°F) FROM PRE-TRANSFUSION		<input type="checkbox"/> NAUSEA / VOMITING	
<input type="checkbox"/> FROTHY EXUDATE IN ETT		<input type="checkbox"/> GENERALIZED / ABNORMAL BLEEDING	

NAME OF LAB PERSONNEL NOTIFIED: _____ NAME OF PROVIDER NOTIFIED: _____ REPORTED BY: _____
 Name Date/Time Name Date/Time Name Date/Time