

Quest Diagnostics At Shady Grove Medical Center and Washington Adventist Hospital

BLOOD BANK STAFF MEETING

MINUTES

(03/03/2015)

PRESENT

3.3.2015 @ 0640-0705 (SGMC) STEPHANIE CODINA, MARY-DALE ABELLANO, HOJAT GOUDARZI, DIPTI PATEL,

RONALD ROJAS, ANNE RIENKS, NAMRATA SHRESTHA, HAMERE TADESSE

3.3.2015 @ 1435-1450 (SGMC) STEPHANIE CODINA, SARAH DELINGER, HABIBA, LAKO

3.4.2015 @ 0635-0700 (WAH) STEPHANIE CODINA, VANESSA ROBINSON, SHAKIMAH RODNEY

3.4.2015 @ 1620-1640 (WAH) STEPHANIE CODINA, TARA APPELBAUM

DISTRIBUTION: BLOOD BANK STAFF MEMBERS

MEETING COMMENCED

Item	Discussion	Action	Follow-up
Minutes			
FDA Inspection	The FDA recently inspected the SGMC blood bank. We had a clean inspection with only a few recommendations. • We will be changing the PI/Variance form a bit to accommodate recommendations (new copy attached). Some boxes were changed to only require an answer when applicable. Other boxes were removed completey. • Beginning in April, the SGMC form will be blue and WAH will be yellow.	None	None
Audits	Nursing seems to be doing a better job with the audits. No major issues were brought up.	None	None
Pneumatic Tube	We are currently validating the pneumatic tube for use in blood product transport at SGMC. When implemented, we will use the same process that WAH is using whereby the person who removes the blood from the tube station will sign and return the form. We need to validate cryo. Instructions and forms are at the issue desk. Please work on the validation if you have the opportunity to thaw cryo.	None	None
OL Monitor	SGMC will be getting a second OL monitor soon. We will set one up for "unreceived" and the other for "received" samples to hopefully keep samples on one screen.	None	None
Printers	We are getting rid of the pink tag. The new printers should arrive any day (they were shipped last week). Once they arrive, we will need to validate before going live. More instructions once they are installed. 1. An example of the new tag is attached. 2. We will have a new form for downtime blood administration (see attached).	None	None

Item	Discussion	Action	Follow-up
Outside Results	We recently had a hospital call to "demand" that we fax the results of an ARC workup to them. Per our SOP, we will give verbal results. If the lab wants faxed results, they must go through HIS/medical records and send a patient release. They also have the option of sending a patient release to ARC for the results. We will not fax.	None	None
Competencies	The BB1 competency is out. BB2 and BB4 will be out soon. Please provide your thoughts about the Immucor competencies.	None	None
New T&S Requirement	 The non-RBC SOPs are being updated to reflect changes to the T&S requirements: Inpatients = entire hospitalization. You can tell it's the same hospitalization by the FIN number. Outpatients = 1 year This will go live as soon as all SOPs are out. Note: OIC will be using a card system to maintain the BB armband for patients. See attached. 	None	None
Sick Calls	Reminder, per SOP, you must call both the department AND the supervisor when calling off from work.	None	None
Open Forum	WAH stated they are not getting the transfuse orders for red cell exchange procedures in a timely manner. Stephanie to follow up.	None	None
Meeting adjourned			
Next meeting week of April 13, 2015			

Stephanie (Codina
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Recorder



BB Variance Form

GEC	
□ SGMC	
WAH	

Occu	urrence Date://		Occurrence Location:		<u>. </u>	
Patie	ent name:			1	MR#	
Accession #: (L Name) Test Code:			(F Name)			
	Description of Variance (specify the in Patient Identification Error				Issuing	
	Incorrect patient drawn by lab		Not labeled with blood bank label		Issued unit did not meet transfusion requirement	
П	Incorrect patient drawn by nursing	ТП	Not labeled		Wrong product issued	
	Patient not banded with blood bank armband		Double labeled (label beneath BB label)		Autologous or directed products not issued when available	
	Armband information incomplete or incorrect		Misspelled name and/or incorrect MRN		Product returned outside of acceptable temperature range	
	Blood bank armband removed		Label missing one or more required pieces	201		
	Patient's Current ABO/Rh does not match patient's historical ABO/Rh		of information (name, MRN, collector's initials, time or date of collection)		ABO confirmation not completed prior to product issue	
	Admitting Department error in patient identification or MRN merge		Duplicate specimen		Procedural Error	
	Wrong blood in tube		Specimen collected on patient at incorrect		Failure to follow SOP	
	Application of the Control of the Co		time (TS drawn with current TS on file,		Unclear SOP	
	Ordering	1_	Fetal Screen collected prior to delivery)		No procedure available	
ļ U	Test or blood product ordered on wrong patient (computer entry)		Collected in wrong tube type		Tech misunderstood SOP	
	Blood product requested on incorrect		Missing specimen (i.e. cord blood)			
	patient (transfusion request)		Maintanana (Famour)		Supplier Error	
	Incorrect BB number on request form		Maintenance/Temperature/QC Temp out of range; no corrective		Mfg recall (excludes blood products) Incorrect or unfilled order	
	Testing		documentation		incorrect or unfilled order	
	Corrected report due to erroneous result		Temp not recorded		Quality Concern / Complaint	
	Clerical (computer data entry error)		QC not documented		Tech Quality Concern (Document on back)	
	Blood Product		QC failure, no corrective action	Ħ	Customer Complaint (Document on back)	
	Incorrect label	_	20 Amino, no concentro abito.	2; —	Customer Complaint (Bootiment on back)	
	Extended expiration date					
	Wasted Product	1	Comments (use space on back of this for	m)		
			Date:		RQI # FDA #	
			•	_		
Notif	ied:	(dat	e/time)	_ by: .	(Tech Code)	
1						
B. S	Supervisor Action and Recommendati	on: (<i>da</i>	cument all follow-up actions taken on reverse	e) (Tra	No lab involvement (No lab involvement (
C. I	Level of Severity No patient impact	Mino	r impact		The last involvement (v)	
D. F	Follow-Up: Hospital Incident Report #		Date:	_		
E. S	Signatures (Sign/Initial and date)					
Super	visor: Medical Director:	:	QA Specialist:		Admin. Director:	



BB Variance Form

□ GEC
□ SGMC
□ WAH

Laboratory RQI (Reportable Quality Issues)
Any FDA reportable event
Any revised result for Blood Bank testing including but not limited to ABO group, Rh type, atypical antibody screen/identification, DAT, RBC antigen typing
Any Significant Procedural Delay resulting in known or potential impact to patient care (treatment or discharge) including: Inability to provide timely blood products during an emergent event
Any Significant Specimen Collection issue (by laboratory staff) causing physical or psychological harm to the patient
Any issue or event judged by a Pathologist or the CLIA Laboratory Director to have known or potential impact on current patient care
A variation or deviation from the local Hospital Policy and Procedure that has known or potential impact on current patient care.
For RQI or FDA reportable - Notify a Supervisor immediately and document on the front of the form
Use these lines for additional information or to document Tech Quality Concerns:
Supervisor Action and Recommendation:

Label Output:

The output label will be as shown in the following graphic. There will no barcodes on this label.

Patient Name	Donor ID/ Unit Number
Medical Record Number	Expiration Date/Time
Blood Bank Armband Number	Volume
Date of Birth	
ABO/Rh	ABO/Rh
Crossmatch Results	Unit Attributes

There will be field headings on the label next to the data fields that are stripped from the incoming data stream. Based on the sample data on the previous page, the label will look like the following.

 Name: SUNQUEST,APPLE
 Donor ID: W2004 14 100929 00

 MRN: 159753
 Exp Date: 10/31/2012 2359

 BB#: 1235689
 Volume: 335 ml

 DOB: 01/10/1953
 ABO/Rh.: O-NEG

 E Neg Kell Neg HgB S neg
 CMVN++

	el here or complete th	<u> </u>	Patient Information		
			Name		
			Medical record number		
			Blood bank number		
			Date of birth		
			ABO/Rh	****	
			Crossmatch Results	ompatible □N	I/A ☐Least Incompatible
Unit Information			I L		
Donor ID	Expiration date/ti	me Al	BO/Rh	Attributes	
Date/Time of issue:	<u> </u>	Issued to:		Visual ins	pection:
We certify that before	starting the blood tra	nefusion we have	checked the following		ceptable Not acceptable
 □ 4. Compare the patien match exactly. □ 5. Compare the blood of the following of the donor matches exactly. □ 7. Compare the expiration and has not been expiration of the provider order of the following of the provider order of the following of the	t's last and first name and bank number on the patie identification number (unition date (and time if appliceeded. ed attributes for the blood ative, HLA-matched).	I medical record numbers blood bank armbatt number) on the blood picable) on the blood picable, verify that the blood picable is the blood picable.	per on the hospital armbal and and the patient/unit lated product label and the patient product label and the patient label and the patient label and the patient label and the patient label product contains	nd and patient/und and patient/unit label. ent/unit label.	Type of blood product: red cells plasma platelets cryoprecipitate
			le af also at		☐ whole blood
Time	No ☐ Yes If yes, stop	HR	Resp	Temp	Initials
Pre-txn			iveah	Temp	initials
	* ***				
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15 min					
15 min 60 min					
15 min 60 min 2 hrs					
15 min 60 min 2 hrs 3 hrs Txn Stop					





Downtime Blood Administration

520-300 (2/15)

Fax both sides of this form to blood bank. DOWNTIME BLOOD TRANSFUSION REACTION REPORT

Procedure:

- 1. Clamp off blood immediately; disconnect blood tubing, all extension tubing and saline. Keep vein open using new saline and
- electronic medical record. Document below. 2. Check for agreement of all identifying names and numbers on patient armbands, blood product label, patientumit label, and
- 3. Notify attending physician immediately, or house physician if warranted to manage patient symptoms.
- 4. Phone Blood Bank. A phlebotomist will be sent to draw post-reaction specimen(s), if indicated.
- 5. Monitor vital signs every 15 min x 3, then every 30 min x 2, then hourly x 2, continue as needed. Record on frequent monitoring
- 6. Return blood container and transfusion tubing to the Blood Bank immediately. Fax a copy of this form to Blood Bank or record and document pertinent information on nursing flowsheet.
- 7. Maintain hourly I & O until further orders are obtained. photocopy both sides of this form and send to Blood Bank immediately.

	gnature:	!S		Signature:
			(Specific)	Other comments
☐ Bleeding	☐ Heat/Pain at IV Site	☐ Rash	EinunidolgomeH []	Сота
Petechiae □	Tudine Output □	eoibnust 🗌	Dyspnea	☐ Fever
Tenderness	səviH □	Back Pain	П Неадасће	Chills
elosuM 🗀	Pruritus 🗆	gniĭimoV □	□ Chest Pain	□ ↑ B\b
muiriləQ 🗌	gnidaul∃ 🗀	EesusN 🗌	☐ Shucobe	əsin9 ↑ 🗌
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		7w	(approx.)	Volume Transfused
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	When?			History of previous re
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——————————————————————————————————————	When?		or abortion?	History of pregnancy
			ior to transfusion:	Clinical Diagnosis pr
	oduct label and the patient/unit label. Ensure blood product contains the ordered attributed.		xpiration date (and time if applica en exceeded.	and has not bee of the brovider o
iber matches exactly.	d and the patient/unit label. Ensure the num product label and the patient/unit label. En	s plood bank armband	ood bank number on the patient' pnor identification number (unit r	match exactly. 3. Compare the blo
sure they match exactly. abel. Ensure they	n the blood bank and hospital armbands. En: r on the hospital armband and patient/unit l	o record number o nedical record numbe	am bns əmsn first name and mə İsən İsət and nəmə sənə n	T. Compare the part of the par

Patient Identification

Downtime Blood Mashington Adventist Hospital

Adventist HealthCare



250-300 Administration

Blood Bank Identification Card

I ha

Blood Bank

ive witnessed the drawing and labeling of my blood sample for the blood bank.	Number
Patient Signature:	
or	7
*Responsible Party:	
[photo ID is required; attach a copy to this form]	

*Note: Whenever possible, the cards should be signed by the patient. If the patient is unable to sign, please have the responsible family member or companion sign the cards. This "responsible" person should be the patient the patient the day of transfusion to confirm the blood bank identification information.

Patient: I have identified my signature above and my name on the blood bank which corresponds to the blood bank identification number on this card. Nurse: I have identified the patient per hospital policy.

Date	Time	Patient Signature	Nurse
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Blood Bank Identification **CHART CARD** 9400-103 (2/15)

Patient Identification

Date	Time	Patient Signature	Nurse
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Blood Bank Identification CHART CARD 9400-103 Patient Identification