### CHRISTUS SPOHN HEALTH SYSTEM **POLICY and PROCEDURE MANUAL**

## TITLE: Laboratory Critical Results, Notification, and Documentation-Procedure

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Originator: Laboratory

Approved By: \*See last page for Laboratory Medical Director / Administrator approvals

(Original with signature archived in Document Control)

#### **PURPOSE:**

To establish standardized critical values for laboratory test procedures which require notification of Physician/LIP (Licensed Independent Practitioner) and/or other appropriate licensed caregiver\*, e.g. RN, Respiratory Therapist. Critical values have been determined with the use of pertinent Clinical Laboratory Science journals and in consultation with Administrative Laboratory Directors, Lead Technologists, and Laboratory Medical Directors and other Medical Staff as indicated.

\*Only licensed caregivers are authorized to receive critical test results1

### **DEFINITION:**

Critical Values: those values that fall outside of the predetermined high and low critical limits and which therefore require immediate notification of the Physician/LIP and/or other authorized licensed caregiver and/or Infection Control for certain Microbiology values.

### **SECTION 1.01 Procedure for Notification and Documentation**

#### 1. Notification:

- a. A list of critical values requiring notification is attached to this document for reference.
- b. If a patient testing value is obtained that falls into the critical value range, the result will be called to the ordering Physician/LIP and/or other appropriate licensed caregiver immediately.
- c. A "read-back" of the result by the person to whom the result is given is required for successful completion of notification.
- d. For Inpatients: The ordering Physician/LIP or appropriate licensed caregiver is notified of the critical value(s).
- e. External Laboratory: Test results that have been referred to the laboratory from outside, intra-System Laboratory hospital clients, nursing homes, and/or skilled nursing facilities will be called to the appropriate Physician/LIP/other authorized caregiver of the patient.
- f. Other: For discharged, clinic or other outpatients not mentioned above, the ordering Physician/LIP, Physician/LIP's nurse, or other authorized party shall be notified of critical values in accordance with hospital policy H-143-P1, "CSHS Critical Values Policy," Procedures #6 and #7, excerpted below for convenience:
  - 6. For a discharged patient the critical values are called directly to the ordering physician or on call physician within one hour.
  - 7. All Critical diagnostic test results received by the patient's nurse will be reporting according to the following sequential notification system:
    - a. First call to MD/LIP #1 (ordering or on call)
    - b. If no response after 15 minutes, call MD/LIP #1 again
    - c. After 30 minutes, escalate notification by contacting the ED Physician and subsequently document the variance on the ordering physician.

g. Patient values determined by the Laboratory testing personnel to exhibit significant trends affecting patient care will be called to the Medical Director, Physician/LIP and/or licensed caregiver as defined in H-143-P<sup>1</sup> and accordingly herein.

#### 2. Documentation:

- a. Documentation of notification for critical results will be entered into the Laboratory computer system and the report will include:
  - i. The patient's full name (first and last name) with two patient identifiers of the patient for whom critical result(s) are being called, e.g. the patient's DOB, and/or the patient's hospital account number
  - ii Date of call
  - iii. Time of call
  - iv. Name (first and last name) and title of person to whom the result(s) are given
- b. Documentation of successful "read-backs" will be entered into the Laboratory computer system with the critical value result. *In order to be successful, the read-back <u>must include at least two patient identifiers</u> the patient's full name (first and last name), <i>and* the patient's DOB, and/or the patient's hospital account number. *A patient's room number is NOT a valid patient identifier.*
- c. Should all attempts for proper notification fail, successful read-backs can include sufficient documentation of faxed results to the outpatients'/discharged patients' Physician/LIPs via a fax transmission confirmation report.

### Reviews:

09/11, 03/13, 12/13, 04/15, 06/16, 07/16, 03/17

#### Reference:

<sup>1</sup> CHRISTUS Spohn Health System Policies and Procedures Manual, "Administrative Policies and Procedures, Section H – Patient Care, H-143-P, 'CSHS Critical Values Policy'" procedure steps 6 and 7. Available at: CSHS CHRISTUS Connect Intranet:

http://christusconnect.echristus.net/regions/spn/OSandGuidelines/Pages/Spohn-PoliciesAndProceduresPage.aspx.

**Critical Laboratory Results Requiring Notification** 

SECTION 1,02 CHEMISTRY CRITICAL VALUES				
Assay (Serum, unless otherwise specified)	Units of Measure	Gritical Low	Critical High	
Calcium	mg/dL	6.6	13.0	
From 0 Days				
Ionized Calcium	mmol/L	0.82	1.55	
From 0 Days				
Chloride	mmol/L	74	125	
From 0 Days				
CO2	mmol/L	10	40	
From 0 Days				
Glucose Neonatal From 0 Days – 11 Months	mg/dL	35	200	
Glucose From 1 Year	mg/dL	50	485	
CSF Glucose	mg/dL	35	435	
From 0 - 4 Days	9 =	• •		
Magnesium Neonatal	mg/dL	1.1	5.0	
•	mg/ac	1.1		
From 0 - 4 Days **Magnesium	mg/dL	1.2	5.0	
	mg/uL	1,4	**OB 7.0	
From 5 Days	m O o m II ca	250	325	
Osmolality, Serum	mOsm/kg	200	320	
From 0 Days				
Phosphorous Neonatal	mg/dL	1.2	9.6	
From 0 Days – 1 Year				
Phosphorous	mg/dL	1.2	8.9	
From 2 Years				
Potassium Neonatal	mmol/L	2.5	7.9	
From 0 - 7 Days	<u></u>			
Potassium	mmol/L	2.8	6.2	
From 8 Days – 12 Years				
Potassium	mmol/L	3.0	6.0	
From 13 Years				
Sodium	mmol/L	120	160	
From 0 Days				
Troponin-I*	ng/mL	none	0.5	
From 0 Days			0.10	
Troponin-T* From 0 Days	ng/ml	none	U. IU	

<sup>\*</sup>Troponin notification will only be performed upon initial critical value and with a subsequent spike during the same admission.

<sup>\*\*</sup>For OB Patients on magnesium therapy only: Critical magnesium levels of >=7.0 mg/dL will be called to RN and documented per protocol. A standardized canned text result comment citing this critical values policy revision will be added in these circumstances for OB patients on magnesium therapy only.

Assay (Serum, unless otherwise specified)	Units of Measure	Critical Low	Critical High (Toxic Level)
Acetaminophen	mcg/mL	none	Greater than or equal to 50
Amikacin Random	mcg/mL	none	Greater than 8.0
Amikacin Trough	mcg/mL	none	Greater than 8.0
Amikacin Peak	mcg/mL	none	Greater than or equal to 35.0
Carbamazepine	mcg/mL	none	Greater than or equal to 20.0
Digoxin	mg/mL	none	Greater than 2.0
Ethyl Alcohol	mg/dL	none	Greater than or equal to 400
Gentamicin Random	mcg/mL	none	Greater than 2.0
Gentamicin Trough	mcg/mL	none	Greater than 2.0
Gentamicin Peak	mcg/mL	none	Greater than 10.0
Lithium	mmol/L	none	Greater than or equal to 1.2
Phenobarbital	mcg/mL	none	Greater than 40.0
Random From 0			
Phenobarbital Random From 12	mcg/mL	none	Greater than or equal to 55.0
	maalm)	nana	Greater than 40.0
Phenobarbital Trough From 0 Days	mcg/mL	none	Greater than 40.0
Phenobarbital	mcg/mL	none	Greater than or equal to 55.0
Trough From 12			<b>1</b>
Phenytoin	mcg/mL	none	Greater than or equal to 30.0
Salicylate	mg/dL	none	Greater than or equal to 30.0
Theophylline	mcg/mL	none	Greater than 20.0
Trough From 0 Days			Cracter than 2E 0
Theophylline	mcg/mL	none	Greater than 25.0
Trough From 12 Theophylline	mcg/mL	none	Greater than 20.0
Random From 0	inog/inc	110110	J. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5.
Theophylline	mcg/mL	none	Greater than 25.0
Random From 12			
Tobramycin Random	mcg/mL	none	Greater than 2.0
Tobramycin Trough	mcg/mL	none	Greater than 2.0
Tobramycin Peak	mcg/mL	none	Greater than 10.0
Valproic Acid	mcg/mL	none	Greater than or equal to150.0
Vancomycin Random	mcg/mL	none	Greater than 20.0
Vancomycin Trough	mcg/mL	none	Greater than 20.0
Vancomycin Peak	mcg/mL	none	Greater than or equal to 50.0
SECTION 1,04 TRANSFUSION SER	VICES		
Test		Units of	Measure Critical Low Critical Hi
Neonatal DAT	Reactions grade	· · · · · · · · · · · · · · · · · · ·	N/A Positive
(From 0 days)	negative to 4+ p		

SECTION 1.05 HEMATOLOGY CRITICAL VALUES				
Test	Units of Measure	Critical Low	Critical High	
WBC*	x10 <sup>3</sup> /μL	Less than 2.0	Greater than 35	
Neonatal WBC (0-7 days)	x10 <sup>3</sup> /μL	Less than 2.0	Greater than 50	
Hemoglobin	g/dl	Less than 7.0	N/A	
eonatal Hemoglobin	g/dl	Less than 9.5	Greater than 25	
Hematocrit	%	Less than 20.0	N/A	
Neonatal Hematocrit	%	Less than 29.0	Greater than 75	
Platelets*	x10 <sup>3</sup> /μL	Less than 50	Greater than 1,000	
PT (non-Coumadin)	Seconds	N/A	Greater than 25	
PT (Coumadin) **	INR	N/A	Greater than or equal to 4.5	
PTT (non-heparin)	Seconds	N/A	Greater than 68	
PTT (heparin) **	Seconds	N/A	Greater than 120	
Fibrinogen	mg/dL	Less than 100	N/A	
CSF WBC	WBC/cumm	N/A	Greater than 10	

<sup>\*</sup> For Oncology patients, WBC and platelet count critical values are called upon admission only; rechecks

are performed on all subsequent results.

\*\*PT and PTT critical values for patients <u>not</u> on anticoagulants will not be flagged in the Laboratory LIS but will be called upon admission only. This applies to all CHRISTUS Spohn Hospital – Corpus Christi facilities.

SECTION 1,06 MICROBIOLOGY CRITICAL VALUES
Positive Blood Culture (non-contaminated)
Positive CSF Culture or Gram Stain
Positive CSF Cryptococcal Antigen
Positive Sterile-Site Body Fluid Culture or Gram Stain
Positive Sterile-Site Tissue Culture or Gram Stain
Positive AFB Smear or Culture*
Positive Mycobacteria tuberculosis identification*
Positive Ocular Culture with:
Bacillus species
Neisseria gonorrhoeae
Pseudomonas aeruginosa
Staphylococcus aureus
Acanthamoeba species
Positive Stool Culture Pathogens (Salmonella, Shigella, Campylobacter, Shiga Toxin producer, E.cole 0157.H7)*
Positive Herpes Simplex Virus from Newborns (up to one year of age)

<sup>\*</sup>Also requires notification to Infection Control Department

The following Microbiology results ("notifiable conditions") are required by state and/or federal law to be reported to the appropriate authorized party. The Microbiology Department calls the following conditions upon the first occurrence of each patient admission, directly to the nursing unit for in-house patients only. The Infection Control Department is also notified of the same conditions via the HIS Infection Control Report. Infection Control personnel notify state authorities with communicable disease reports as required by state law.

### SECTION 1.06,a, MICROBIOLOGY NOTIFIABLE CONDITIONS (ALERTS, NOT CRITICAL VALUES)

- ♦ Methicillin Resistant Staphylococcus aureus (MRSA)
- Vancomycin Resistant Enterococcus (VRE)
- Extended Spectrum Beta Lactamase producing Enterobacteriaceae (ESBL)
- ♦ Carbapenem Resistant Enterobacteriaceae (CRE)
- Multidrug Resistant Acinetobacter
- ♦ Multidrug Resistant Pseudomonas aeruginosa
- ♦ Pan-Resistant Organisms
- Toxigenic Clostridium difficile
- ♦ Select Agents and other highly pathogenic organisms (Bacillus anthracis, Brucella spp., Burkholderia mallei, Burkholderia pseudomallei, Coccidiodes immitis, Francisella tularensis, Yersinia pestis, and C. diphtheriae, Salmonella typhi, Vibrio cholera)\*

# Section 1.07 Anatomic Pathology Critical Procedure

Frozen Section: A frozen section is considered a critical test/procedure and the Pathologist reports results immediately to the requesting Physician/LIP.

Approved:	CSHCC – Memorial	
	Joe Lewis M.D Laboratory Medical Director	05/08/17 Date
	George C. Bost	05/08/17
	Laboratory Administrator	Date
	CSHCC – Shoreline	05/004/3
	Joe Lewis M.D. Laboratory Medical Director	<u>05/08/17</u> Date
	George C. Bost Laboratory Administrator	05/08/17 Date
	CSHCC - South	
	Joe Lewis M.D. Laboratory Medical Director	<u>05/08/17</u> Date
	George C. Bost Laboratory Administrator	05/08/17 Date
	CSH – Alice	
	Randall Simonsen M.D. Laboratory Medical Director	<u>05/11/17</u> Date
	Barbara A. Herro Laboratory Administrator	<u>05/11/17</u> Date
	CSH Beeville	
	Randall Simonsen M.D. Laboratory Medical Director	05/11/17 Date
	Barbara A. Herro Laboratory Administrator	<u>05/11/17</u> Date
	CSH – Kleberg	
	Randall Simonsen M.D. Laboratory Medical Director	05/11/17 Date
	Barbara A. Herro	05/11/17 Date: