HEALTHCARE-ASSOCIATED INFECTIONS PROGRAM

Central Line Associated Bloodstream Infection Surveillance

Last updated 2019

Basics of Infection Prevention Healthcare-Associated Infections Program Center for Health Care Quality California Department of Public Health



Objectives

- Review CLABSI surveillance definitions
- Discuss importance of accurate data collection
- Demonstrate how to report CLABSI events summary data in NHSN
- Discuss NHSN data analysis and feedback to staff



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CLABSI Surveillance for Prevention

- 1. Perform surveillance for CLABSI using NHSN standardized definitions and methods
- 2. Compare SIR or rate over time to assess prevention progress
- 3. Monitor CLABSI incidence over time using the standardized infection ratio (SIR) metric

(See Introduction to NHSN slides)



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CLABSI Surveillance Key Terms

- Lab confirmed bloodstream infection (LCBI)
 - Blood culture positive for a pathogen
- Commensal
 - Organism not usually considered pathogenic
 - Include (but not limited to)
 - Diphtheroids
 - Propionibacterium spp.
 - coagulase-negative staphylococci
 - viridans group streptococci
 - Aerococcus spp.
 - Micrococcus spp.

See NHSN Patient Safety Manual: Chapter 4, pp 4-10, NHSN organism list https://www.cdc.gov/nhsn/pdfs/pscmanual/4psc_clabscurrent.pdf



CLABSI Surveillance

- For BSI to be considered a CLABSI, a **central line** must be
 - In place for >2 days on the date of the event (date device placed = day one)

AND

- <u>Still in place on day of event -or- in place on the day prior</u> to the event
- The CLABSI event date is defined as the day the <u>first</u> element used to meet the surveillance definition occurs within the seven-day window period

NHSN Patient Safety Module: Chapter 4



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CLABSI Surveillance Definition

LCBI 1 Patient of any age has a recognized pathogen cultured from one or more blood cultures and Organism cultured from blood is not related to an infection at

another site

LCBI 2*

 Patient of any age
 has common skin commensals cultured from 2 or more blood cultures drawn on separate occasions

and

has at least one of the
 following signs or symptoms
 Fever (>38°C), chills, or
 hypotension

and

 Signs and symptoms and (+) lab results are not related to an infection at another site

*All criteria occur within 7 day infection window period

LCBI 3*

- Patient of ≤ 1 year of age
- has common skin commensals cultured from 2 or more blood cultures drawn on separate

occasions

and

has at least one of the following signs or symptoms Fever (>38°C), hypothermia (<36°C core),

apea, or bradycardia and

Signs and symptoms and (+) lab results are not related to an infection at another site



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Mucosal Barrier Injury (MCBI) BSI

- More specific BSI definition for oncology patients
- BSI resulting when intestinal organisms from compromised intestinal wall mix into the bloodstream
- Occurs in post allogeneic hematopoietic transplant or severely neutropenic patients
- MCBI SIR is calculated separately from CLABSI SIR



CLABSI Infection Criteria- Acute Care Hospitals

Diagnostic Test for Possible CLABSI Localized Sign or Symptoms for Possible CLABSI (ONLY used with 2 blood commensals)

- Positive blood culture with a pathogen OR-
- 2 positive blood cultures with common commensals

- Fever
- Chills
- Hypotension



CLABSI due to Common Commensal Organisms

- Two blood cultures have been collected on the <u>same or</u> <u>consecutive days</u>
 - One positive culture may be due to poor skin prep prior to lab draw (skin contaminant)
 - Two matching positive cultures of the same commensal, meeting criteria, are considered a true pathogen

Example: Blood cultures positive for common commensal organism (e.g., S. epi) collected on Mon-Tues meets LCBI 2; cultures collected on Mon-Wed are too far apart



CLABSI Infection Window Period

- Defined as the 7-days during which all site-specific infection criteria must be met
- Includes the <u>day the first positive blood culture</u> was obtained, <u>3 calendar days before</u> and <u>3 calendar days after</u>

Infection Window Period:	3 days before first positive diagnostic test Mar 7 Mar 8 Mar 9		FIRST POSITIVE DIAGNOSTIC TEST		after first p agnostic te	
Example:	Mar 7 Mar 8 Mar 9		Mar 10	Mar 11	Mar 12	Mar 13



CLABSI Event Date

- The <u>date of event</u> is the date the first element is used to meet the definition for the first time
- May or may not be the positive blood culture date



CLABSI Location Attribution

- A CLABSI is attributed to the location of the patient on the day of event
 - Defined as the date that the <u>first</u> element used to meet the LCBI criterion occurred
- If the date of event for a CLABSI is the day of transfer or discharge, or the next day, the infection is attributed to the transferring location
- Attribute CLABSI to correct location for accurate SIR calculations. Each location has different risk adjustments in NHSN



CLABSI Cannot Re-Occur in the Same Patient within a 14-Day Timeframe

- The date of the CLABSI event is considered day 1
- A new CLABSI is not reported until 14 days have elapsed
- If a new pathogen is identified in the blood within the 14 day timeframe, it should be <u>added</u> to the CLABSI already reported
 - Refer to the CLABSI protocol for more details



Secondary BSI Attribution

- The period in which a positive blood culture must be collected to be considered a secondary BSI to a primary site of infection
 - Includes the 7-day infection window combined with the 14-day repeat infection timeframe, or 14-17 days depending on the date of the event
 - A positive blood culture collected outside this 14-17 date range cannot be considered a secondary BSI to the primary infection
- A primary BSI (CLABSI) cannot have a secondary BSI



Secondary BSI Attribution -2

- A secondary BSI may be attributed to a primary site of infection if one of the following is true:
 - 1. The blood culture pathogen matches an organism also cultured in the primary infection site

OR

- 2. A positive blood culture is an element used to meet the primary site infection
- See the Secondary BSI Guide (Table B1) of the CLABSI protocol for more details



Secondary BSI Attribution -3

- NHSN Infections that include a positive blood culture as an element in the primary site definition:
 - Bone-Osteomyelitis
 - Burn
 - Disc space infection
 - Endocarditis
 - GI tract infection
 - Intra-abdominal infection

- Meningitis
- Other infection-reproductive tract
- Pneumonia
- Spinal abscess
- Omphalitis
- Urinary System Infection

• Joint

NHSN Patient Safety Module: Chapter 4, Secondary BSI Guide, pp 4-27, Table B1 <u>https://www.cdc.gov/nhsn/pdfs/pscmanual/pcsmanual_current.pdf</u>



Pathogen Assignment

- If a new blood pathogen is identified within the 14-day repeat infection timeframe, it should be added to the already reported CLABSI as an additional pathogen
- Do not report it as a new CLABSI
- Pathogens excluded from specific infection definitions (e.g. yeast for UTI and PNEU) are also excluded from being considered secondary bloodstream infections
 - Example: Yeast in the blood and urine would be reported as a CLABSI, as yeast is excluded from the UTI definition
- Refer to the NHSN protocol for more details on pathogen assignment and secondary BSI



Pathogens Associated with CLABSI

•	Coagulase-negative Staphylococci	16%
•	Staphylococcus aureus	13%
•	Klebsiella (pneumoniae/oxytoca)	8%
•	Enterococcus faecalis	8%
•	Enterococcus faecium	7%
•	Candida albicans	6%
•	Escherichia coli	5%
•	Candida spp	5%

NHSN Antimicrobial Resistance Report: Distribution of all Pathogens Reported by HAI Type, Appendix to Table 4, 2011-2014

https://www.cdc.gov/nhsn/xls/reportdatatables/2014-appendix-pathogens.xlsx



How do I Apply the CLABSI Surveillance Definitions?

Let's look at some





NHSN HAI and POA Worksheet Generator

Based on the information you provided: Admit Date: Tue Jan 01 2019 The event is: HAI Date of Event: Fie Jan 04 2019 Infection Window Period: Tue Jan 01 2019 - Mon Jan 07 2019 Repeat Infection Timeframe (RIT): Fri Jan 04 2019 - Thu Jan 17 2019 Event Type: BSI

Start Over...

Back...

Admit date: 1/1/2019

www.cdc.gov/nhsn

Hospital Day/Date	First Diagnostic Test	Infection Window Period (*)	Date of Event	Repeat Infection Timefran (*)	ne		
1 1/1/2019 - Admit Date			•				
2 1/2/2019			•				
3 1/3/2019			•				
4 1/4/2019	√	v	- HAI				
5 1/5/2019			•				
6 1/6/2019			·				
7 1/7/2019			· ·				
8 1/8/2019							
9 1/9/2019	Enter 3 dr	ata points into Workshee	et Generator:				
10 1/10/2019		•					
11 1/11/2019		of Admission					
12 1/12/2019	Date	of first diagnostic test					
13 1/13/2019		Event type					
14 1/14/2019							
15 1/15/2019	 NHSN Ger 	 NHSN Generates a worksheet for you to enter additional data 					
16 1/16/2019	• You must	determine if the HAI def	finition is me	t			
17 1/17/2019		1	r				

Print Friendly Window...

Generate Table...

BSI Event Date

Hospital Day/Date	First Diagnostic Test	Infection Window Period	Date of Event	Repeat Infection Timeframe			
nospital Day/ Date		(*)	Date of Event		(*)		
12/30/2018			-				
12/31/2018							
1 1/1/2019 - Admit Date			. /	Autom	atically		
2 1/2/2019	√	BC + Staph aureus	- POA	populates H			
3 1/3/2019				or POA	on date		
4 1/4/2019				of event			
5 1/5/2019				016	vent		
6 1/6/2019							
7 1/7/2019							
8 1/8/2019	BSI: POA						
9 1/9/2019	Date of Ev	ent: date of the first o	liagnosti	c test			
10 1/10/2019	Pathogen:						
11 1/11/2019	Fathogen	staph A	I				
12 1/12/2019			-				
13 1/13/2019			-				
14 1/14/2019							
15 1/15/2019			•				

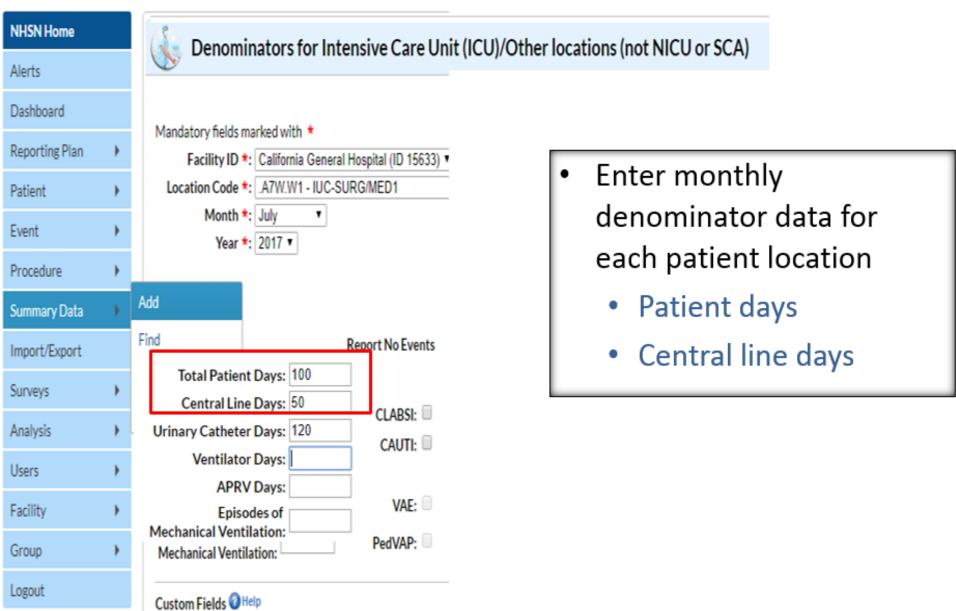
CLABSI Event Date

Hospital Day/Date	First Diagnostic Test	Infection Window Period (*)	Date of Event		Repeat Infection Timeframe (*)	Secondary BSI Attribution Period (*)	d	
1 1/1/2019 - Admit Date		Central line inserted						
2 1/2/2019								
3 1/3/2019		Fever 38.8	· HAI					
4 1/4/2019	√	🖉 BC + Staph epi						
5 1/5/2019		BC + Staph epi						
6 1/6/2019			·	Remember:				
7 1/7/2019			·					
8 1/8/2019			The <u>date of event</u> is the					
9 1/9/2019			date the first element is					
10 1/10/2019			·	used	to meet the d	efinition		
11 1/11/2019				for t	he first time			
12 1/12/2019				_				
13 1/13/2019								
14 1/14/2019								
15 1/15/2019			-					
16 1/16/2019			-					

Primary and Secondary Examples

Hospital Day/Date	First Diagnostic Test	Infection Window Period (*)		Repeat Infection Timeframe (*)	Secondary BSI Attribution Period (*)	
1 1/1/2019 - Admit Date			•			
2 1/2/2019						
3 1/3/2019		 Dysuria 	· HAI			
4 1/4/2019	√	Urine culture >100,000cfu/ml E.				
5 1/5/2019						
6 1/6/2019			•			
7 1/7/2019						
8 1/8/2019			-			
9 1/9/2019			•			
10 1/10/2019			•			
11 1/11/2019		Blood culture E. faecalis/Yeast			Blood Culture E. faecalis/Yeast	
12 1/12/2019						
13 1/13/2019		& Secondary BSI		Primary	BSI	
14 1/14/2019	DOE= 1/3/19			DOE = 1/	/11/19	
15 1/15/2019	Pathogen: E. faecalis			Pathogen: Yeast		
16 1/16/2019	ratin			Pathogen: reast		

Add Monthly CLABSI Summary Data to NHSN

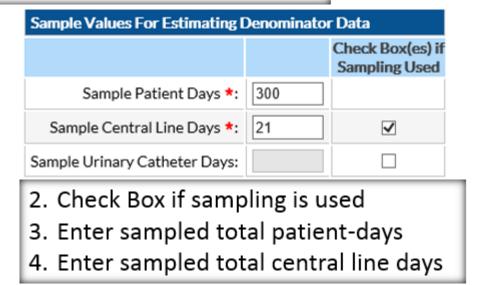


Optional: Denominator Data Sampling

How to sample: Count the number of the location patient days and the number of central lines on a designated day each week. Not on Saturday or Sunday. Add those numbers for the month and enter here.

1. Enter Monthly patient days for this location based on daily collection

Denominator Data						
	V	Report No Events				
Total Patient Days:	450					
Central Line Days:	32	CLABSI:				
Urinary Catheter Days:		CAUTI:				
Ventilator Days:		VAE: PedVAP:				
5. NHSN will estimate the central line days for the month						



Note: Sampling may not be used for NICU or specialty care areas/oncology

Add CLABSI Event to NHSN

NHSN Home	🗼 Add Event
Alerts	 Add CLABSI Events as they occur
Dashboard	· · · ·
Reporting Plan	Fields required for record complete Collect criteria data meeting
Patient 🕨	Fields required when in Plan mate definition to enter into NHSN
Event •	Add aerinition to enter into NHSN
Procedure 🕨	Find Facili NHSN has a worksheet available
Summary Data 🔹 🕨	Incomplete
Import/Export	Las for data collection
Surveys 🕨	https://www.cdc.gov/nhsn/forms/57.108_
Analysis	E PrimaryBSI_BLANK.pdf
Users 🕨	
Facility •	
Group	Event Information
Logout	Event Type *: BSI - Bloodstream Infection
	Post-procedure: N - No 🔻
	MDRO Infection Surveillance *: No, this infection's pathogen/location are not in-plan for Infe
	Location *: 5 NORTH - MICU
	Date Admitted to Facility >:
	Risk Factors
	Central line *: Y - Yes 🔻
	Any hemodialysis catheter present: Y - Yes 🔻
	Location of Device Insertion: ED - EMERGENCY DEPARTMENT (ED)
	Date of Device Insertion:
	Event Details
	Specific Event ➤: LCBI - Laboratory confirmed bloodstream infection ▼

NHSN CLABSI Analysis Reports

NHSN Home		
Alerts		Add Event Add CLABSI Events as they occur
Dashboard		· · · · ·
Reporting Plan	•	Fields required for record complete Collect criteria data meeting
Patient	•	Fields required when in Plan may
Event	•	Add definition to enter into NHSN
Procedure	•	Find Facili Patie • NHSN has a worksheet available
Summary Data	•	Incomplete
Import/Export		for data collection
Surveys	•	https://www.cdc.gov/nhsn/forms/57.108_
Analysis	•	E PrimaryBSI_BLANK.pdf
Users	•	
Facility	•	
Group	•	Event Information
Logout		Event Type *: BSI - Bloodstream Infection
		Post-procedure: N - No 🔻
		MDRO Infection Surveillance *: No, this infection's pathogen/location are not in-plan for Infe
		Location *: 5 NORTH - MICU
		Date Admitted to Facility >:
		Risk Factors Central line *: Y - Yes V
		Any hemodialysis catheter present: Y - Yes V
		Location of Device Insertion: ED - EMERGENCY DEPARTMENT (ED)
		Date of Device Insertion:
		Event Details
		Specific Event >: LCBI - Laboratory confirmed bloodstream infection ▼

NHSN TAP Report - CLABSI

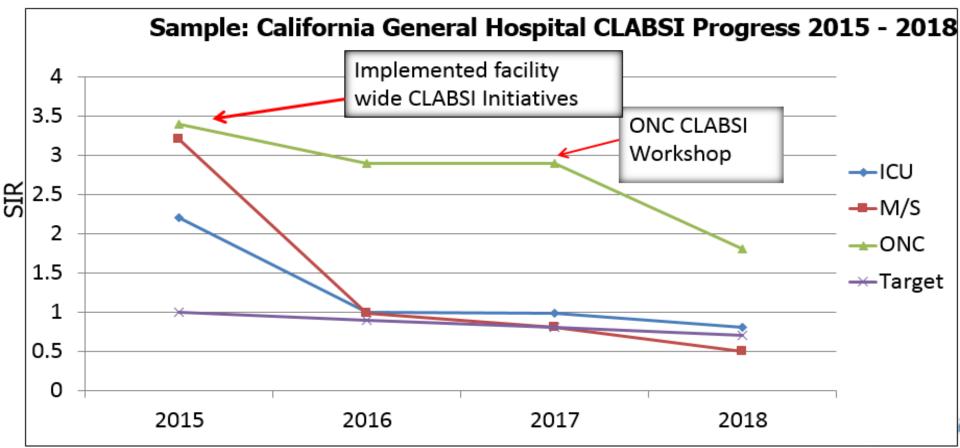
	-								
Facility	Location				Central				SIR
CAD	Rank	Location	CDC Location	Events	Line Days	DUR %	CAD	SIR	Test
20.52	1	1 West	IN:ACUTE:WARD:M	14	2269	49	13.10	7.81	
	2	2 West	IN:ACUTE:WARD:M	4	1349	42	3.40	3.34	
	3	SICU	IN:ACUTE:CC:S	3	1062	9	2.58	-	
	4	5 West	IN:ACUTE:WARD:M	2	983	9	1.61		

- Identifies the number of infections that needed to prevented to reach targeted goal (CAD)
 - Lists results high-to-low by location
 - Assists in deciding where to focus infection prevention resources



Measure CLABSI Prevention Progress

- Feedback results to your staff and leadership
- Changes in CLABSI incidence should be visible over time
- In the example, we can see ONC needed additional interventions



CLABSI Surveillance Summary

- Consistent use of standard surveillance methods and CLABSI definitions are essential for accurate case finding
- Capturing complete and accurate data is necessary for precise CLABSI SIR calculation
- Perform surveillance and feedback CLABSI SIR with adherence monitoring results to all units and leadership



Questions?

For more information, please contact any HAI Program Liaison IP Team member

> Or email <u>HAIProgram@cdph.ca.gov</u>

