Central Line Associated Bloodstream Infection Surveillance

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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



CLABSI Surveillance for Prevention

- 1. Perform surveillance for CLABSI using NHSN standardized definitions and methods
- 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)



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

- Still in place on day of event -or- in place on the day prior 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

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 \leq 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

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

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

Localized Sign or Symptom s for Possible CLABSI (ONLY used with 2 blood 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>



CLABSI Infection Window Period

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



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
 - Gl tract infection
 - Intra-abdominal infection
 - Joint

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

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

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





CLABSI Event Date

Date the first element used to meet the definition for the first time

		ı		
HOSPITAL	INFECTION		HOSPITAL	INFECTION
DAY	WINDOW PERIOD		DAY	WINDOW PERIOD
				Central Line
1			1	inserted
	Blood Culture			
2	+Staph A		2	
3			3	Fever 38.8C
				Blood Culture +
4			4	Staph epi
				Blood Culture +
5			5	Staph epi
6			6	
7			7	
8			8	
9			9	
10			10	
11			11	
12			12	
13			13	
14			14	
15			15	
16			16	
17			17	
18			18	
	BSI-POA			CLABSI-HAI
	Date of Event =2			Date of Event =3
	Pathogen= Staph A			Pathogen= Staph
				epi

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Infection

Window Period

RIT

1

2

13

14

Primary and Secondary BSI Examples
nfection Window Period 1st positive diagnostic test, 3 ays before and 3 days after)
epeat Infection Timeframe 4 days. Date of event = day 1
econdary BSI Attribution

R ıy 1

Period (Infection window Period + RIT)

HEALTHCARE-ASSOCIATED INFI							
Hosp Day	BSI	RIT	Infection Window Period				
1							
2							
3		1	Dysuria				
4		2	Urine culture >100,00cuf/ml E. faecalis				
5		3					
6		4					
7		5					
8		6					
9		7					
10		8					
11		9	Blood Culture E. faecalis/Yeast				
12		10					
13		11					
14		12					
15		13					
16		14					
17			UTI & Secondary				
18			BSI DOE=3				
19							
20			Pathogen:				
21			E.faecalis				

22

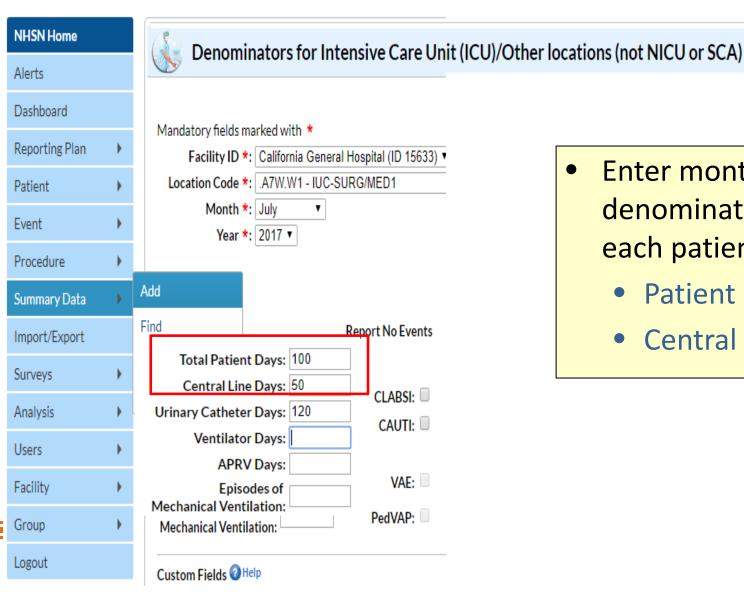
23

24

Blood culture

E. faecalis/Yeast

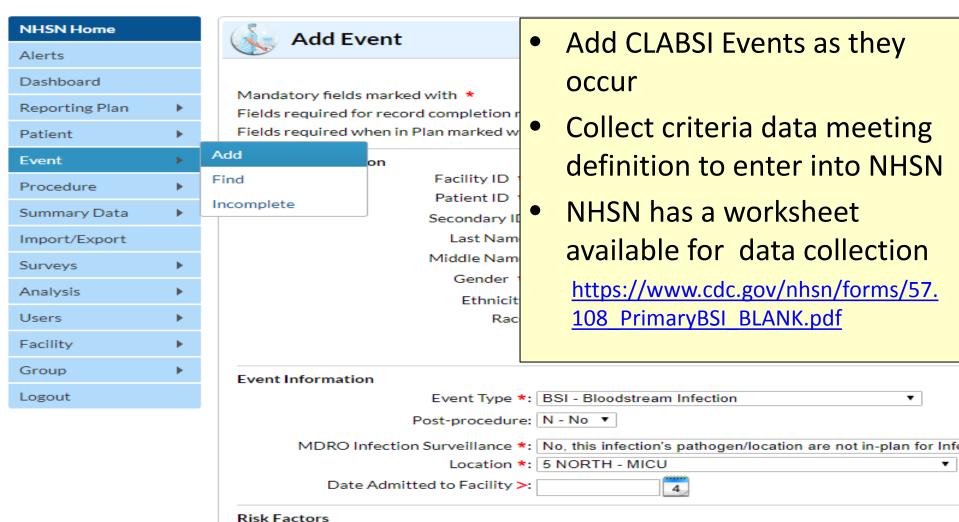
Add Monthly CLABSI Summary Data to NHSN



- **Enter monthly** denominator data for each patient location
 - Patient days
 - Central line days



Add CLABSI Event to NHSN



Central line *: Y - Yes ▼

Location of Device Insertion: ED - EMERGENCY DEPARTMENT (ED)

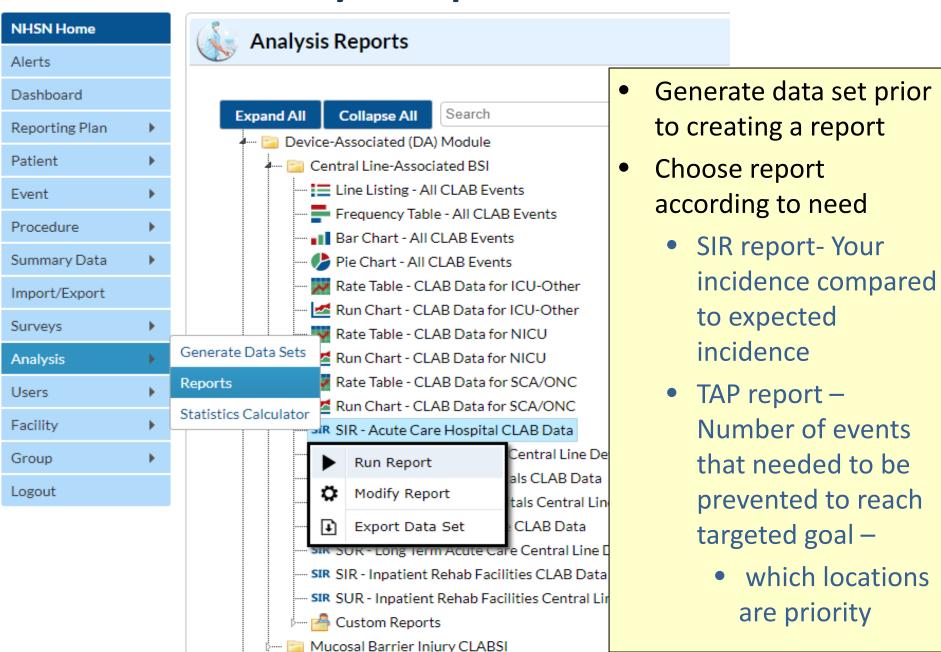
Specific Event >: LCBI - Laboratory confirmed bloodstream infection ▼

Any hemodialysis catheter present: Y - Yes ▼

Date of Device Insertion:

Event Details

NHSN CLABSI Analysis Reports



NHSN TAP Report - CLABSI

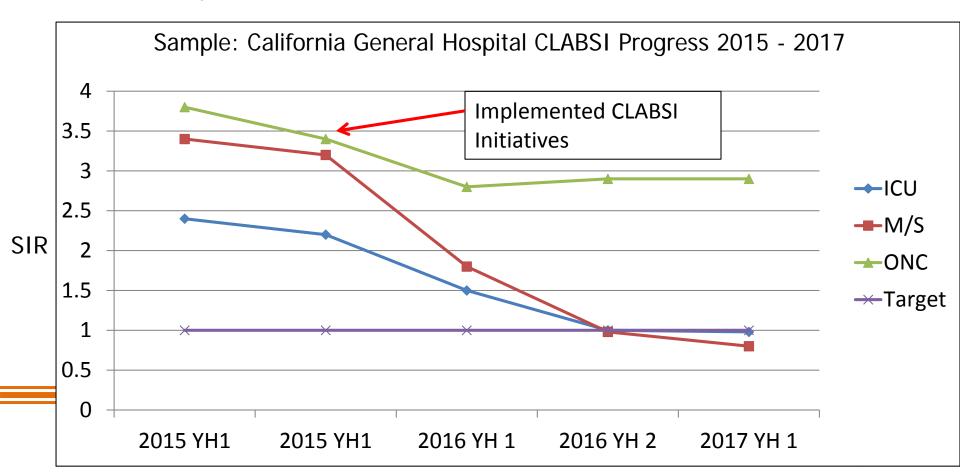
	_								
Facility	Location				Central				SIR
CAD	Rank	Location	CDC Location	Events	Line Days	DUR %	CAD	SIR	Test
20.52	1	1 West	IN:ACUTE:WARD:M	14	2269	49	13.10	7.81	
	2	2 West	IN:ACUTE:WARD:M	4	1349	42	3.40	3.34	
	3	SICU	IN:ACUTE:CC:S	3	1062	9	2.58		
	4	5 West	IN:ACUTE:WARD:M	2	983	9	1.61		

- 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 needs some 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 HAIProgram@cdph.ca.gov

