

# Epidemiology and Surveillance

Last Updated 2019

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



# Objectives

- Discuss basic principles of epidemiology and how they apply to healthcare-associated infection (HAI) surveillance
- Review recommended surveillance practices
- Describe surveillance outcome and process measures for infection prevention

# Epidemiology

- Definition: Study of disease factors affecting populations  
Clinical care: focus on the individual  
vs  
Epidemiology: focus on the group
- Healthcare epidemiology answers questions such as:
  - What factors contribute to increased HAI rates?
  - What populations are at higher risk for developing HAI?
  - How have HAI changed over time?
- Assessment of trends over time

# Infection Prevention and Healthcare Epidemiology

- Goal is HAI prevention
- Discipline professional societies
  - Association for Professionals in Infection Control and Epidemiology (APIC)
  - Society for Healthcare Epidemiology of America (SHEA)
  - Infectious Diseases Society of America (IDSA)
- Epidemiologic research and surveillance underlie HAI prevention
  - Use data for action!

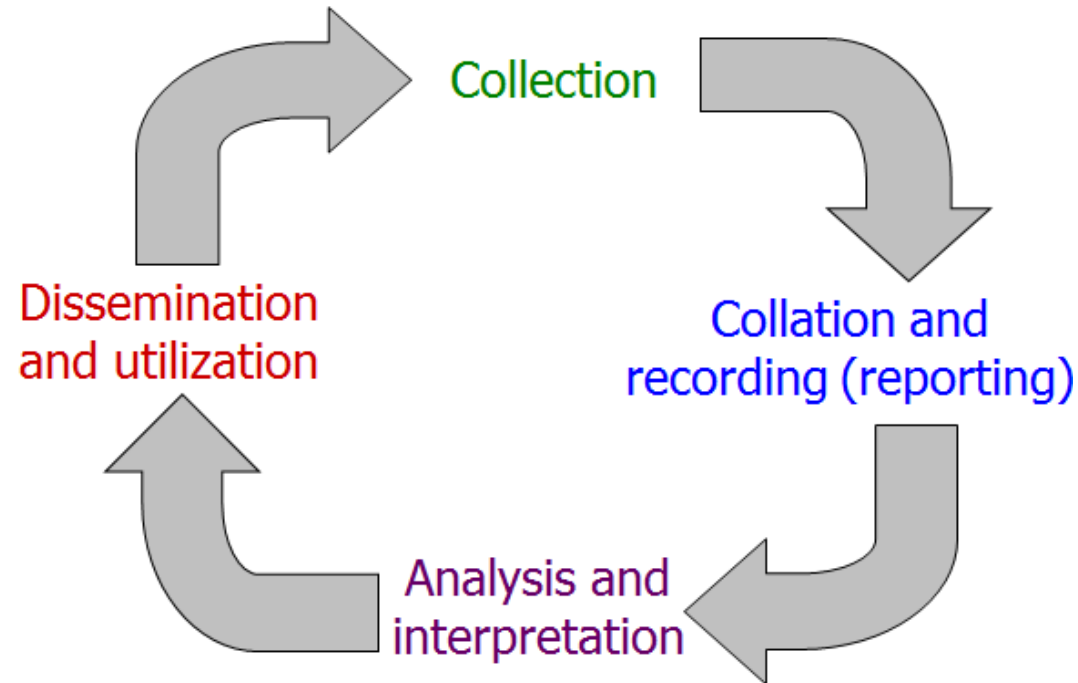
# Epidemiologic Surveillance

- The ongoing, systematic collection, recording, analysis, interpretation, and dissemination of data
- Reflects rate of disease onset or current health/disease status of a community or population (e.g., healthcare patients)
- Aims to identify risk factors for disease
- Used for public health action to reduce morbidity and mortality, and to improve health

# Surveillance

A surveillance system is an information loop that starts and ends with communication and action

## Flow of Surveillance Data



# Key Tenets of HAI Surveillance

- A written plan serves as the foundation
  - What HAI am I tracking? Why?
  - How will data be used?
  - Where are opportunities to prevent HAI in my facility?
- The intensity of surveillance efforts need to be maintained over time
- Stay consistent over time; always apply same surveillance definitions

# Outcome Measure Examples

- CLABSI, CDI, and SSI Standardized Infection Ratio (SIR)
- MRSA and VRE BSI rate per 10,000 patient days



## Process Measure Examples

- CAUTI prevention: percent urinary catheters with appropriate indication
- CLABSI prevention: percent adherence to CLIP bundle (all or none)
- CDI prevention: thoroughness of environmental cleaning
- HAI prevention: percent adherence to hand hygiene

# Measuring Infections

## Incidence

Number of persons in a population who develop a disease or condition within a specified period of time

Measure of **new** infections

## Prevalence

Proportion of persons in a population who have a disease or condition at a given point in time

Measure of infections that **are present**

# Incidence

Incidence measures the frequency of **disease onset** (i.e., rate).  
Answers: 'What is the risk of X occurring?'

$$\text{Incidence} = \frac{\text{\# of new cases during a specified time period}}{\text{size of a specific population}}$$

Example:

$$\frac{5 \text{ SSI}}{97 \text{ Kidney Surgeries}} = 0.05 \text{ new infections per 97 kidney surgeries, During the time period of Jan-Dec 2017}$$

# Prevalence

Prevalence measures disease status in a population at a particular time. Answers: 'How common is X?'

Prevalence =  $\frac{\text{\# of existing cases during a specified time period}}{\text{size of a specific population}}$

Examples:

$$\frac{160 \text{ employees vaccinated}}{200 \text{ employees total}} = 0.8 \times 100 = \mathbf{80\%}$$

$$\frac{2 \text{ patients colonized with MRSA}}{10 \text{ patients admitted on same day}} = 0.2 \times 100 = \mathbf{20\%}$$

# Incidence Density Rate

Incidence density accounts for variation in the time each person is at risk for an event

Incidence density rate =

$$\frac{\text{\# of new cases during a specified time period}}{\text{person-time at risk}}$$

Example:

$$\frac{\text{\# hospital onset CDI}}{\text{\# of patient days}}$$

# HAI Surveillance Definitions

- Case definition (surveillance definition)
  - Clinical and laboratory characteristics that a patient must have to be counted as an event or case for surveillance purposes
  - **Time, place, & person** (e.g., age, sex)
  - Universal case reporting
  - A surveillance system in which all cases of a disease are to be reported

# Laboratory-based surveillance

A surveillance method in which the reports of cases come from clinical laboratory data only (forgoing case review/symptomatology)

# Applying Surveillance Definitions

- Always refer to written definitions to ensure accuracy of applying case definitions
  - Use standardized, published, validated definitions where available
- For accurate and valid comparisons, use the same definitions
  - If definitions change, the comparability of rates over time will be compromised

*“Align criteria and definitions and decrease subjectivity while maintaining epidemiologic standardization and clinical relevance.”*  
*(NHSN Patient Safety Module, Chapter 2, 2019)*



# Clinical vs Surveillance Definitions

- Clinical
  - Patient centered
  - Used for therapeutic decisions
- Surveillance
  - Population based
  - Applied exactly the same way each time

## Collect Surveillance Data

- Include IP, clinical staff, and others share the responsibility
- Limit collection to only what is needed
- Be involved in efforts when creating or revising the electronic health records to enable HAI data collection

# Prospective Surveillance

- Initiated when patient is still under the care
- Advantages
  - Ability to capture information in real time
  - Can interview caregivers
  - Can gather findings not recorded in patient record
  - Easier to demonstrate temporality (e.g., before and after observations) and therefore make causal inferences

# Retrospective Surveillance

- Closed record review after patient has been discharged
- Advantages:
  - Allows for comprehensive review of sequential events
  - Efficient
- Disadvantage:
  - Does not allow for prompt intervention
  - Important/relevant information may be missing
- Administrative (billing, coding) data alone **cannot accurately identify HAI**
  - May be useful for identifying **possible HAI**

# Numerator Data

- Numerator = number of instances of the “event” being measured
- Examples:
  - HAI identified through **active** surveillance: CLABSI, CAUTI, SSI, VAP
  - HAIs identified by **laboratory** finding alone: CDI, MRSA BSI, VRE BSI
  - Care **practices, processes**, observations: CLIP, hand hygiene compliance
- Record point in time or time period

# Denominator Data

- Denominator = number of patients or procedures being followed, the population size, or person-time at risk (patient or line days)
- Examples:
  - Procedures
  - Patient days
  - Patient visits

## Calculate and Analyze Infection Rates

Calculate rates and ratios by denominator type

- Total population at risk, or time at risk
- Used to calculate raw rate or incidence density rate:

Examples:

$$\frac{5 \text{ SSI}}{300 \text{ cardiac procedures}} \times 100 = 1.67$$

$$\frac{2 \text{ CLABSI}}{1500 \text{ line days}} \times 1000 = 1.33$$

$$\frac{218 \text{ patient days with central line}}{360 \text{ total patient days}} = 0.61$$

# Risk Factor Data

- Factors that increase a patient's risk for HAI include
  - Patient characteristics and co-morbidities
  - Facility characteristics
  - Unit / ward type
  - Community disease prevalence
  - Invasive device use and duration
  - Surgical procedure type, duration, approach, and other circumstances
- Data collection includes risk factor data necessary for risk adjustment



# Apply Risk Adjustment Methodology

- **CLABSI and CAUTI:** Infection risk takes into account patient location
- **SSI:** Probability of infection calculated for each surgical patient; varies by surgery
- **CDI & MDRO (LabID):** Infection type risk accounts for facility characteristics, disease burden (community prevalence), and testing method (for CDI)

# Standardized Infection Ratio (SIR)

- Summary measure used to track HAI incidence
- Allows for tracking over time
- Compares the actual number of HAI reported to what would be predicted using 2015 baseline data
- Adjusted for risk factors significantly associated with HAI

# Calculating Standardized Infection Ratio (SIR)

- Standardized infection ratio

$$\text{SIR} = \frac{\text{Observed HAI}}{\text{Predicted HAI}}$$

Example:

Hospital A has 4 MRSA BSI over 23,500 patient days. National data predicted 2.5 MRSA BSI.

$$\text{SIR} = \frac{4}{2.5} = 1.6$$

# NHSN: A Guide to the SIR

- How to interpret SIR
- How SIR is calculated
- Risk adjustment factors for specific HAI

## THE NHSN STANDARDIZED INFECTION RATIO (SIR)

*A Guide to the SIR*

*Updated July 2017. Please see [Page 2](#).*



NHSN: A Guide to the SIR

<https://www.cdc.gov/nhsn/pdfs/ps-analysis-resources/nhsn-sir-guide.pdf>

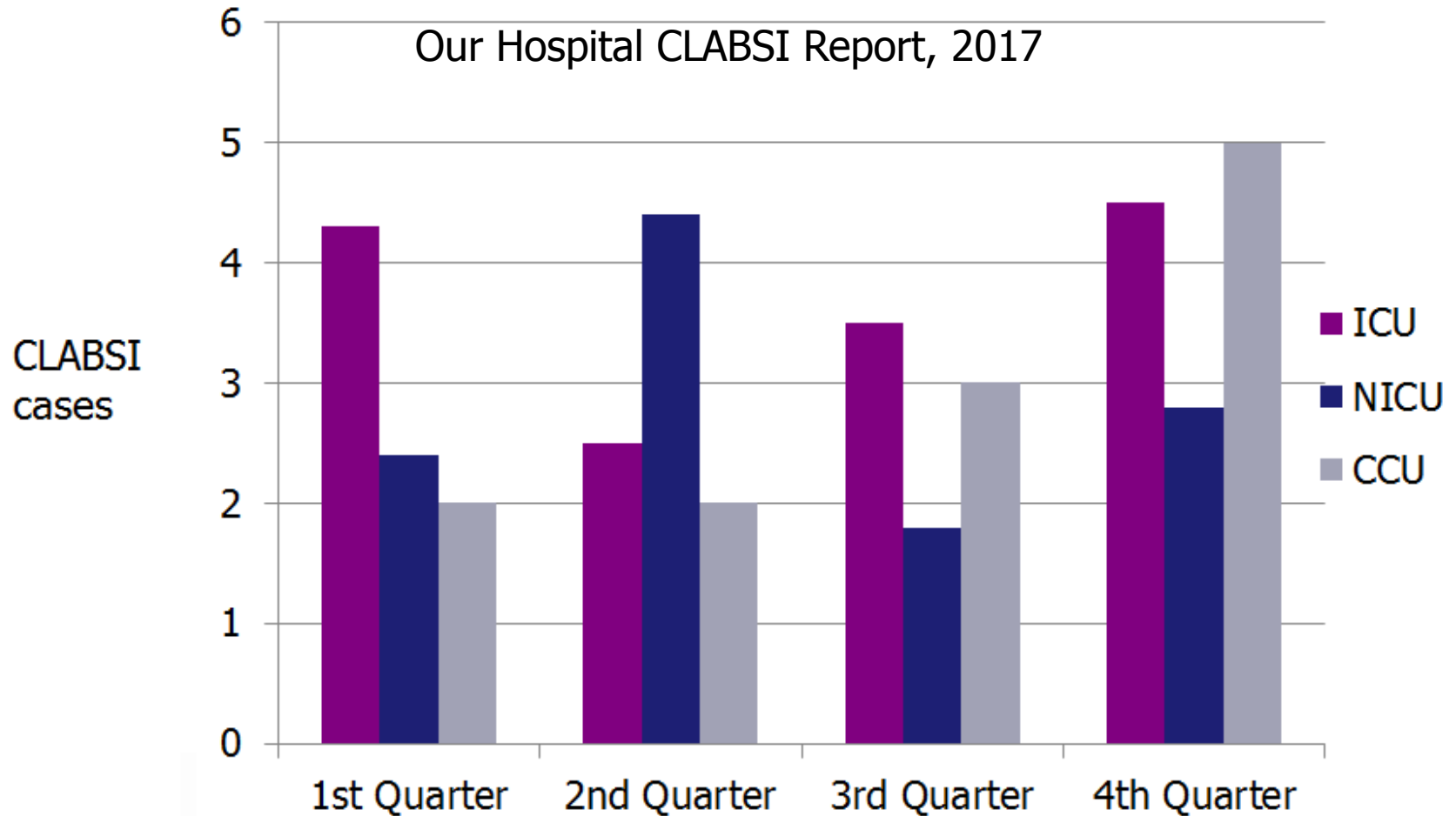
# Report and Use Surveillance Data

*“The demonstrable power of surveillance is in sharing findings with those who need to know and who can act on the findings to improve patient safety.”*

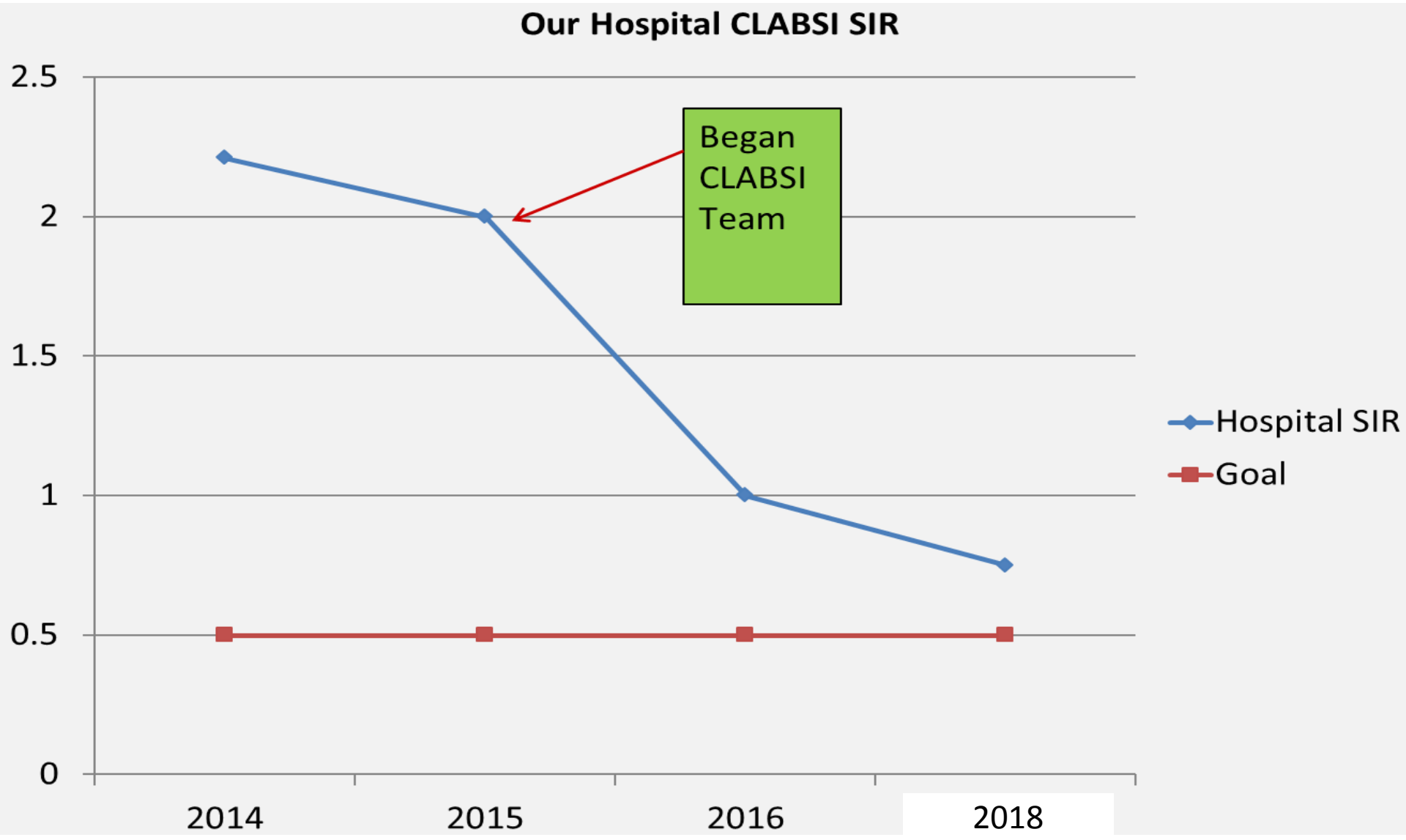
AJIC Am J Infect Control, 35:427-40, 2007

- Plan for distribution of findings
- Report to health care providers most able to impact patient care
- Report in a manner to stimulate process improvement
- Use visual displays of data (e.g., charts, graphs, tables)

# Sample Bar Charts



## Sample Line Graphs and Histograms - 3



## Summary

- The IP must understand the basic principles of epidemiology and apply them to HAI surveillance
- Accurate and consistent data collection, recording, analysis, interpretation, and communication of findings is an essential part of the infection prevention and surveillance plan
- Surveillance of process measures helps focus prevention activities to improve outcomes



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- Ebbing Lautenbach, K. F. Woeltje, and P.N. Malani., Practical Healthcare Epidemiology, 3<sup>rd</sup> Edition, 2010.
- Horan, T.C., Andrus, M., and Dudeck, M.A. CDC/NHSN surveillance definition of health care-associated infection and criteria for specific types of infections in the acute care setting. *Am J Infection Control* 36: 309-332, 2008.
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## Questions?

For more information,  
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