

## Can we use automated analyses of electronic health record data to improve surveillance for healthcare-associated infections?

Seminar on Application of Artificial Intelligence (AI) on Infectious Diseases and Infection Control  
November 13, 2024

Michael Klompas MD, MPH, FIDSA, FSHEA  
Professor, Harvard Medical School  
Hospital Epidemiologist, Brigham and Women's Hospital

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## Disclosures

- **Grant funding**
  - Centers for Disease Control and Prevention
  - Massachusetts Department of Public Health
  - Agency for Healthcare Research and Quality
- **Royalties**
  - UpToDate

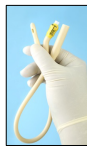
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## Current State

We focus the bulk of our effort on:



### CAUTI



Catheter-associated urinary tract infection

### CLABSI



Central-line associated bloodstream infection

### SSI (HYST & COLO)



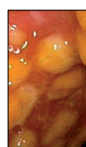
Surgical site infection (hysterectomy & colectomy)

### HO-MRSA Bacteremia



Hospital-onset MRSA bacteremia

### HO-CDI



Hospital-onset C. difficile infection

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## An Enduring Legacy of SENIC

- Landmark study designed by CDC in 1974 to describe US infection control programs, measure nosocomial infection rates, and determine if implementing infection surveillance and control programs lead to lower HAI rates
- Resounding success
- Set the framework for infection control for the next 50 years

American Journal of Epidemiology  
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**THE SENIC PROJECT**  
STUDY ON THE EFFICACY OF NOSOCOMIAL INFECTION CONTROL (SENIC PROJECT)

SUMMARY OF STUDY DESIGN

ROBERT W. HALEY, DANA QUADRI, HOWARD E. FREEDMAN, JOHN V. BENNETT, AND THE CDC SENIC PLANNING COMMITTEE

Haley, R. W. (SENIC Project, CDC, Atlanta, GA 30333), D. Quadri, H. E. Freedman, J. V. Bennett and the CDC SENIC Planning Committee. Study on the efficacy of nosocomial infection control (SENIC Project). Summary of study design. *Am J Epidemiol* 111:472-485, 1980.

With the emergence of nosocomial infections as a serious problem among US hospitals, the Center for Disease Control undertook in 1974 a nationwide study to evaluate approaches to infection control. The three-phase project, now known as the Study on the Efficacy of Nosocomial Infection Control, or SENIC Project, was designed with three primary objectives: (1) to determine whether and, if so, to what degree the implementation of infection surveillance and control programs (ISCPs) has lowered the rate of nosocomial infection; (2) to describe the current status of ISCPs and infection rates; and (3) to demonstrate the relationship among characteristics of hospitals and patients, components of ISCPs, and changes in the infection rate. With data collection completed in a nationally representative sample of hospitals, analysis is underway to identify approaches to infection control that are most effective in the least cost to hospitals and to point out additional specific questions to be answered by future research.

**cross infection; hospital infections; nosocomial infections; health surveys; medical records; epidemiologic methods**

Received for publication September 11, 1979, and in final form January 15, 1980.  
Reprints: Robert W. Haley, Center for Disease Control, International Epidemiology and Health Statistics Branch, 1600 Clifton Road, NE, Atlanta, GA 30333.  
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Presented in part at the 10th Annual Meeting of the American Epidemiological Society, Denver, CO, December 1979, and at the 1980 National Conference on Antimicrobial Agents and Chemotherapy, Atlanta, GA, October 1980.

The design of the study and design, the general project methodology, and the data collection have been outlined in the SENIC Project Design Manual, Bureau of Epidemiology, CDC. The method of selecting the study sites and the methodology with the Bureau of Training, CDC. Other organizations participating in varying phases of the project include the American Hospital Association, School of Public Health, University of Minnesota, School of Public Health, University of North Carolina at Chapel Hill, and the University of Texas at Dallas.

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Haley, *Am J Epi* 1980;111:472-485

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**AMERICAN**  
**Journal of Epidemiology**

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JOURNAL OF THE AMERICAN SOCIETY OF EPIDEMIOLOGISTS  
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VOL. 121,	FEBRUARY 1985	NO. 2
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**Original Contributions**

**THE NATIONWIDE NOSOCOMIAL INFECTION RATE:  
A NEW METHOD FOR VITAL STATISTICS**

ROBERT W. HALEY,<sup>1</sup> DAVID H. CULVER,<sup>2</sup> JAMES W. WHITE,<sup>2</sup> W. MEREDITH MORRIS<sup>3</sup> AND  
T. CLAYTON THOMPSON<sup>4</sup>

<sup>1</sup>Haley, R. W. (CDC) Project, CDC, Atlanta, GA 30333; <sup>2</sup>Culver, D. H., White, J. W., Morris, W. M. & T. C. Thompson. The nationwide nosocomial infection rate: a new method for vital statistics. *Am J Epidemiol* 1985;121:159-167.

<sup>3</sup>From a random sample of patients and hospitals and extrapolation ratios derived from the last available sources of data, the authors estimate that the nationwide nosocomial infection rate among the 1,643 acute-care US hospitals in 1975-1976 was 1.7 nosocomial infections per 100 admissions and that over 2 million nosocomial infections occurred annually. Nosocomial urinary tract infections constituted 47% of the infections, surgical wound infections 24%, nosocomial pneumonia 19%, nosocomial bacteremia 15%, and nosocomial infections at all other sites 19%. If the estimates are scaled for the existence of the original method, the true rates may be as low as one-half, and the number of nosocomial infections is varying further. However, it may be a good approximation if the authors' assumptions are reasonable. This study also presents preliminary estimates and calls for timely and accurate vital statistics on the problem.

<sup>4</sup>cross infection; health policy; health services research; health surveys; hospitals; sampling studies; vital statistics

The infectious complications suffered by hospitalized patients (nosocomial infections) constitute a serious public health problem. There may be no more successful means of preventing them than better surveillance. In 1981, and in 1982, the Centers for Disease Control (CDC), Division of Field Epidemiology, conducted the National Nosocomial Infection Study (NNIS). This study was designed to provide information on the magnitude of the problem, to identify risk factors, and to evaluate control measures. The study was conducted in two phases. The first phase was a descriptive study of the problem in 1981. The second phase was an analytic study of the problem in 1982. The results of the first phase are reported here. The results of the second phase will be reported in a subsequent issue of the journal.

Received for publication November 4, 1983, and in revised form March 1, 1984.  
Address reprint requests to Dr. Haley, Center for Disease Control, Division of Field Epidemiology, 1600 Clifton Road, NE, Atlanta, GA 30333.  
This work was supported by the National Institutes of Health, Contract Grant No. CA 30333, and the National Institute for Environmental Health Sciences, Contract Grant No. ES 00001.  
Reprints: Haley, Robert W. (CDC) Project, CDC, Atlanta, GA 30333.  
Copyright © 1986 by The American Society of Epidemiologists and Biostatisticians. All rights reserved. Printed in the United States of America. 0950-2688/85/0002-0159\$01.00/0.

# An Enduring Legacy of SENIC

- Used detailed chart reviews to estimate an HAI rate of 5.7 events per 100 admissions
  - UTI 42%
  - SSI 24%
  - HAP 10%
  - Bacteremia 5%
  - Other sites 19%
  
- Documented that robust infection control programs were associated with substantial reductions in HAIs
  
- Parallel data demonstrated that devices, procedures, and intensive care accounted for the majority of nosocomial infections

Haley, Am J Epidemiol 1985;121:159-167  
Stamm, Ann Intern Med 1978;89:764-769

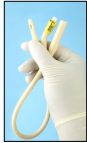




[illegible]

## Is this still the right approach for the current era?

- Are we capturing the right events (completeness, morbidity)?
- Are we conducting surveillance in an efficient and objective fashion?
- Is our approach commensurate with current data sources and documentation practices (i.e. EHRs)?

# Difficulties with our current surveillance targets

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<b>CAUTI</b>	<b>CLABSI</b>	<b>SSI (HYST &amp; COLO)</b>	<b>HO-MRSA Bacteremia</b>	<b>HO-CDI</b>
				
Catheter-associated urinary tract infection	Central-line associated bloodstream infection	Surgical site infection (hysterectomy & colectomy)	Hospital-onset MRSA bacteremia	Hospital-onset <i>C. difficile</i> infection

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**Complicated**

**Labor Intensive**

**Subjective**

**Non-Specific**

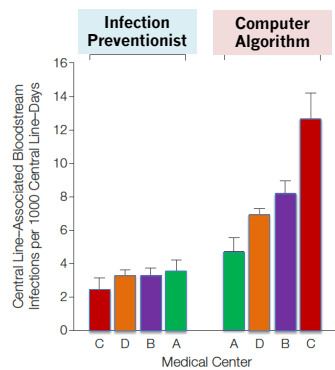
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### It's Hard!

- Is this patient's fever due to infection or some other cause?
- Is this bacteremia due to abdominal infection or line-infection?
- Is this coag-negative Staph a contaminant or a true pathogen?
- Is this positive *C.diff* test colonization or infection?
- Was this present on admission or was it hospital-acquired?
- Does the surgeon's prescription for antibiotics for "erythema" count as an attending physician's diagnosis of infection?
- Is cholecystitis after colectomy a surgical site infection?
- etc.

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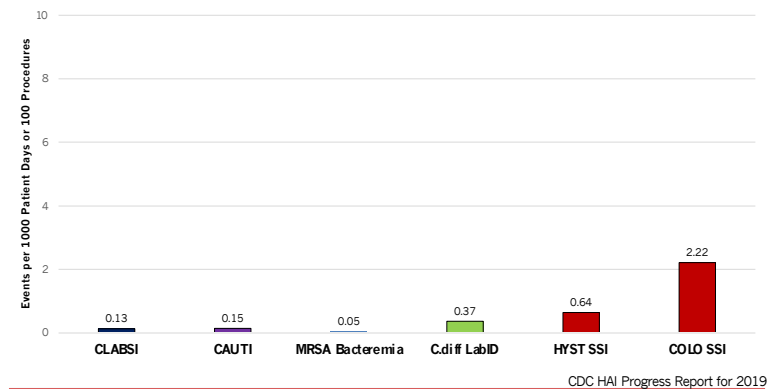
### Manual versus Automated Surveillance on CLABSI Rank



Lin, JAMA 2010;304:2035-41

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### They Are Rare!

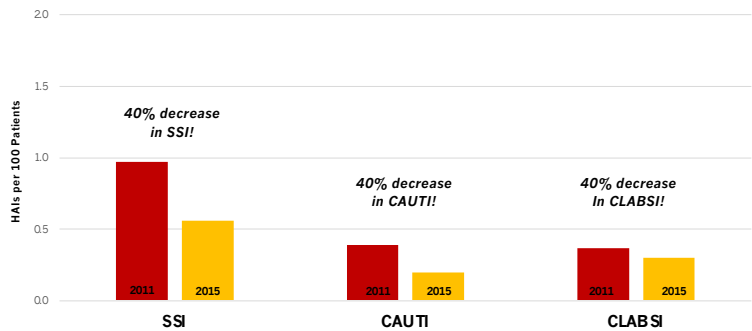


CDC HAI Progress Report for 2019

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## And Becoming Rarer

CDC point-prevalence surveys for HAI in 2011 (183 hospitals) vs 2015 (199 hospitals)



Magill, N Engl J Med 2018;379:1732-1744

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## They Miss the Most Common HAI

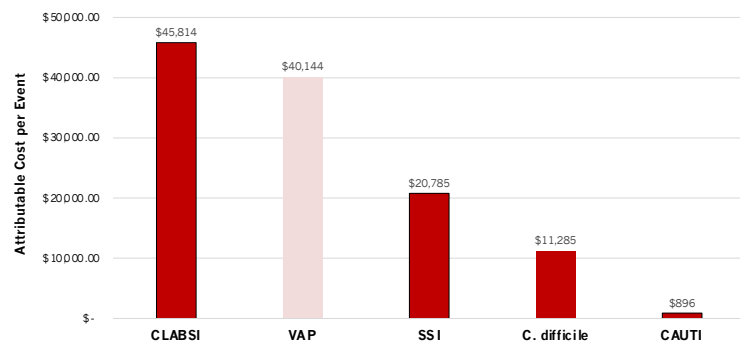
CDC one day point-prevalence survey of HAI in 199 hospitals in 10 U.S. states

1. **Hospital-acquired pneumonia**
2. GI infections including *C. difficile*
3. Surgical site infection
4. Primary bloodstream infections
5. Urinary tract infections

Magill, N Engl J Med 2018;379:1732-1744

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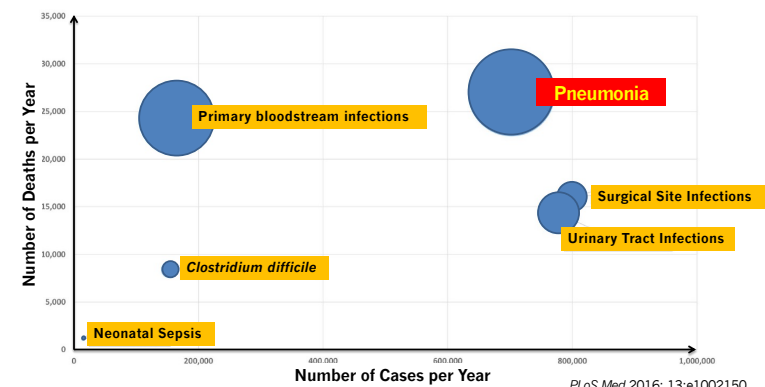
## They vary widely in their impact



Zimlichman, JAMA Internal Med 2013;173:2039-46

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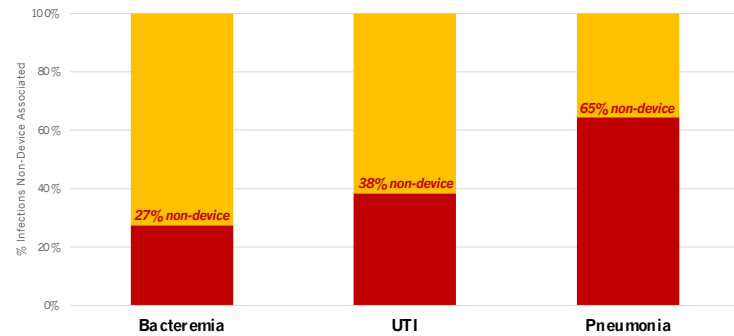
## Burden & Mortality of HAIs in Europe



PLoS Med 2016; 13:e1002150

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## They miss non-device-associated infections



Magill, N Engl J Med 2018;379:1732-1744

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## What's the alternative?

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CDC 24/7: Saving Lives. Protecting People™

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[A-Z Index](#)

### National Healthcare Safety Network (NHSN)

CDC > NHSN Home > Patient Safety Component > Calculators & Worksheets

**NHSN Home**

- NHSN Login
- About NHSN
- Enroll Facility Here
- CMS Requirements
- Change NHSN Facility Admin
- Resources by Facility
- Patient Safety Component**
  - Annual Surveys, Locations & Monthly Reporting Plans
  - Analysis Resources
  - Antimicrobial Use & Resistance

#### MDRO & CDI LabID Event Calculator Version 2.0

Welcome to Version 2.0 of the MDRO & CDI LabID Event Calculator. Version 2.0 operates based upon the currently posted LabID Event protocols in the NHSN Multidrug-Resistant Organism (MDRO) & *Clostridioides difficile* Infection (CDI) Module. The calculator is a web-based tool that is designed to help users learn how to accurately apply the MDRO & CDI LabID Event algorithms and assist users in making the correct MDRO & CDI LabID Event determinations.

Please note that the MDRO & CDI LabID Event Calculator does not ask users to enter any patient identifiers (other than dates of specimen collection, which can be changed as needed). The MDRO & CDI LabID Event Calculator does not save, store, or report any data that is entered. Likewise, LabID Event determination data are NOT reported to the NHSN application, and users will not be able to export data entered into the Calculator. Therefore, events that are determined by the Calculator to be LabID Events will need to be entered into the NHSN application either manually or via CDA.

If you have questions or suggestions about the Calculator, please feel free to send them to the NHSN mailbox: [nhsn@cdc.gov](mailto:nhsn@cdc.gov).

**MDRO & CDI LabID Event Calculator**  
Version 2.0  
(must have javascript enabled)

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**CDC** Special Point of View Article

## Developing a New, National Approach to Surveillance for Ventilator-Associated Events\*

Shelley S. Magill, MD, PhD<sup>1</sup>; Michael Klompas, MD, MPH<sup>2,3,4</sup>; Robert Balk, MD<sup>5,6</sup>; Suzanne M. Burns, RN, ACNP, MSN, RRT<sup>6,7</sup>; Clifford S. Deutschman, MS, MD<sup>6,8</sup>; Daniel Diekema, MD<sup>9,10</sup>; Scott Fridkin, MD<sup>1</sup>; Linda Greene, RN, MPS<sup>11,12</sup>; Alice Guh, MD, MPH<sup>1</sup>; David Gutterman, MD<sup>6,13</sup>; Beth Hammer, RN, MSN, ANP-BC<sup>6,14</sup>; David Henderson, MD<sup>15</sup>; Dean Hess, PhD, RRT<sup>16,17,18</sup>; Nicholas S. Hill, MD<sup>6,19</sup>; Teresa Horan, MPH<sup>1</sup>; Marin Kollef, MD<sup>6,20</sup>; Mitchell Levy, MD<sup>6,21</sup>; Edward Septimus, MD<sup>22,23</sup>; Carole VanAntwerpen, RN, BSN<sup>24,25</sup>; Don Wright, MD, MPH<sup>26</sup>; Pamela Lipsett, MD, MHPE<sup>6,27</sup>

*Critical Care Medicine* 2013;41:2467-2475

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**CDC** Centers for Disease Control and Prevention  
CDC 24/7: Saving Lives. Protecting People<sup>SM</sup>

**nhsn.cdc.gov/VAECalculator/vaecalc.html**

National Healthcare Safety Network (NHSN)

CDC > NHSN > Materials for Enrolled Facilities

NHSN Ventilator-Associated Event (VAE) Calculator Ver. 10.0

Start Over Calculate VAE Explain... Goto PVP

Day	Date	Min PEEP (cmH <sub>2</sub> O)	Min FiO <sub>2</sub> (%)	VAE	T+30 <sup>WBC</sup> or T+38 <sup>WBC</sup>	WBC	CEP/EPD	Choose a Drug	Choose a Drug	QAD
3	10/3/2024	8	50							
4	10/4/2024	5	40							
5	10/5/2024	5	40							
6	10/6/2024	5	40							
7	10/7/2024	8	50	1 VAE						1 yes
8	10/8/2024	10	60							1 yes
9	10/9/2024	10	60							1 yes
10	10/10/2024	8	50							1 yes
11	10/11/2024									1 yes
12	10/12/2024									1 yes
13	10/13/2024									1 yes

Legend: 1- VAE Window 2- VAE Date 3- Qualifying Antimicrobial Day (QAD)

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**Brief report**  
Assessment of an automated surveillance system for detection of initial ventilator-associated events  
Dooshanveer Nuckchady MD<sup>1</sup>, Michael G. Heckman MS<sup>2</sup>, Nancy N. Diehl Tara Creech RN<sup>3</sup>, Darlene Carey RN, MSN<sup>4</sup>, Robert Domnick BS<sup>5</sup>, Walter C. Hellinger MD<sup>6,7</sup>

**Major Article**  
Development and validation of an automated ventilator-associated event electronic surveillance system: A report of a successful implementation  
Courtney Hebert MD, MS<sup>1,2,\*</sup>, Jennifer Flaherty RN, MPH, CIC<sup>3</sup>, Justin Smyer MLS (ASCP)<sup>4</sup>, MPH, CIC<sup>5</sup>, Jing Ding PhD<sup>6</sup>, Angino MD<sup>7</sup>

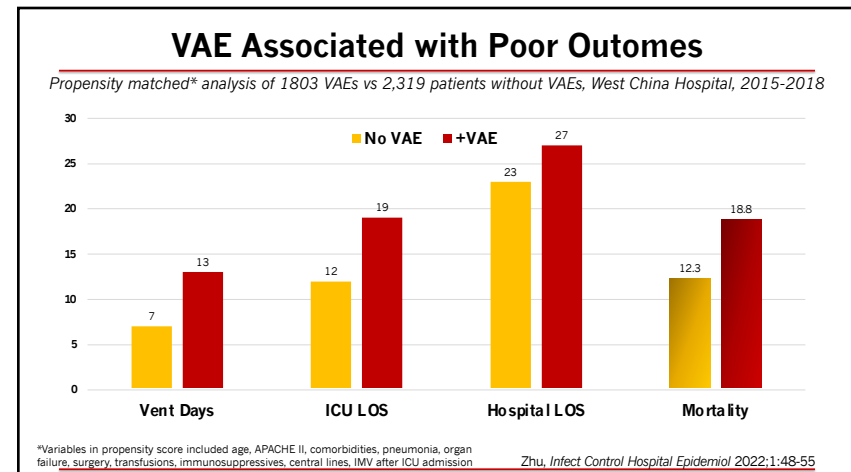
**Electronic Implementation of a Novel Surveillance Paradigm for Ventilator-associated Events**  
Feasibility and Validation  
Peter M. C. Klein Klouwenberg<sup>1,2,3,\*</sup>, Maaike S. M. van Mourik<sup>4,\*</sup>, David S. Y. Ong<sup>1,2,3</sup>, Jannette M. Marcus J. Schultz<sup>1</sup>, Olaf L. Cremer<sup>2</sup>, and Marc J. M. Bonten<sup>1,2,3</sup>, on behalf of the MARIS

**Building and Validating a Computerized Algorithm for Surveillance of Ventilator-Associated Events**  
Joseph Elsworth, BSH<sup>1</sup>, Naga Huda MD<sup>2</sup>, Anupama Nedukumar MD, MPH<sup>3</sup>, Thomas Chavaler, BSN, CIC<sup>4</sup>, Sima, MPH, CIC<sup>5</sup>, Sorab Dhar, MD<sup>6</sup>, Mary E. Robinson, BSH<sup>2</sup>, Keith S. Kaye, MD, MPH

**Development, Implementation and Use of Electronic Surveillance for Ventilator-Associated Events (VAE) in Adults**  
Erving Resetar, MIM, PMP<sup>1,3</sup>, Kathleen M. McMullen, MPH, CIC<sup>2</sup>, Anthony J. Rus MPH<sup>1</sup>, Joshua A. Doherty, BS<sup>3</sup>, Kathleen A. Gase, MPH, CIC<sup>2</sup>, Keith F. Woeltje, MD,

**An automated retrospective VAE-surveillance tool for future quality improvement studies**  
Oliver Wolfers<sup>1,2,3</sup>, Martin Faltys<sup>4</sup>, Janos Thomann<sup>5</sup>, Stephan M. Jakob<sup>6</sup>, Jonas Marschall<sup>7</sup>, Tobias M. Merz<sup>8,9</sup>, and Rami Sommerstein<sup>1,10</sup>

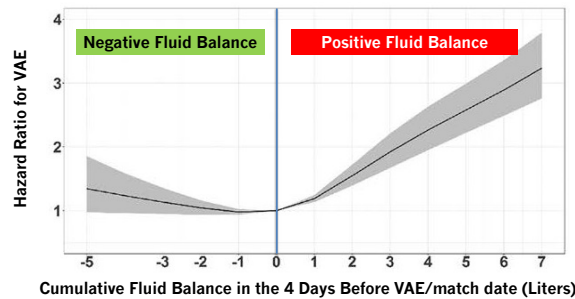
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## Strong Association between Fluid Balance and VAEs

Cumulative fluid balance amongst 1,528 VAE patients matched to 3,038 non-VAE patients on basis of age, time to VAE, and time from ICU admission until initiation of mechanical ventilation, West China Hospital, 2015-2018. Adjusted for demographics, ICU type, comorbidities, ICU diagnosis, APACHE II, meds, procedures, and others.

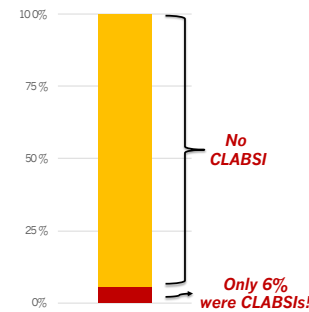


Wang, *Critical Care Medicine* 2022;50:307-316

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## Hospital Onset Bacteremia (HOB)

### Hospital-Onset Bacteremia in 80 ICUs



- Prospective surveillance for hospital-onset bacteremia in 80 ICUs in 16 hospitals from 2012-2013
  - 11,280 episodes of hospital-onset bacteremia
  - Only 663 were classified as CLABSIs (6%)
- HOB allowed for much more discrimination between ICUs versus using CLABSI rates
  - 75% of ICUs' CLABSI SIRs included 1 versus 25% of ICUs' projected HOB SIRs

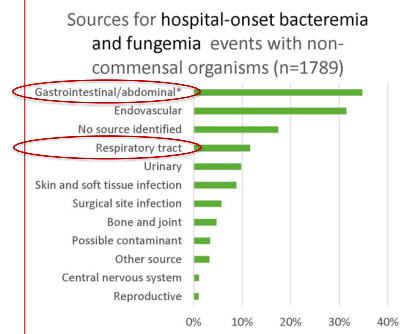
Rock, *JCHE* 2016;37:143-148  
Dante, *JCHE* 2019;40:358-361

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## Characteristics of Hospital-Onset Bacteremia Patients

Cross-sectional analysis of 2109 hospital-onset bacteremia, 13 US hospitals, 2016-2019

- Median age: 56
  - ICU: 32%
  - Active malignancy: 22%
  - Any device present: 92%
- Median hospital length-of-stay
  - 24 days
- Hospital mortality: 23%
- 40% of events not captured through existing surveillance definitions

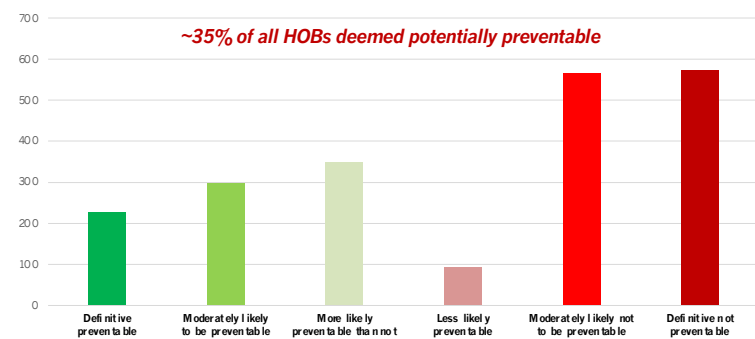


Leekha, *BMJ Quality Safety* 2024;33:487-498

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## Estimated Preventability of Hospital Onset Bacteremia Events

Cross-sectional analysis of 2109 hospital-onset bacteremia, 13 US hospitals, 2016-2019



Leekha, *BMJ Quality Safety* 2024;33:487-498

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[illegible]

## Non-Ventilator HAP

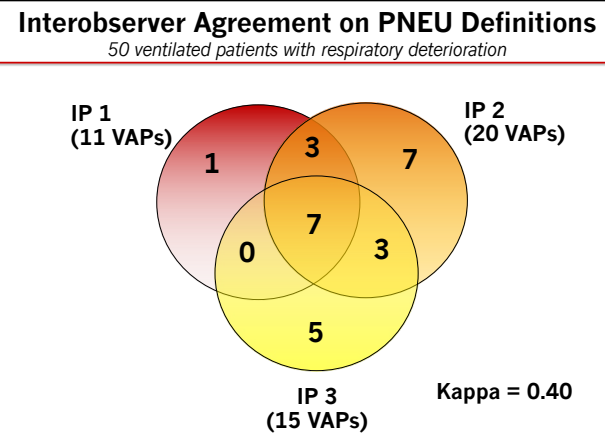
- o Pilot surveillance strategy using EHR data including vital signs, oxygen devices, oxygen flow rates, lab tests, antibiotics...
- o Flags patients on hospital day  $\geq 3$  with:
  - o Sustained deterioration in oxygenation, *and*
  - o Abnormal temperature or WBC count, *and*
  - o Order for chest imaging, *and*
  - o  $\geq 3$  days of new antibiotics
- o Applied to 2 years of data from 4 hospitals (489,519 admissions)
  - o 0.6 events per 100 hospitalizations
  - o 28% mortality
  - o Moderate correlation with clinical dx and PNEU

Ji, *JAMA Network Open* 2019;2(10):e1913674  
Ramirez Battle, *ICHE* 2020;41:219-221



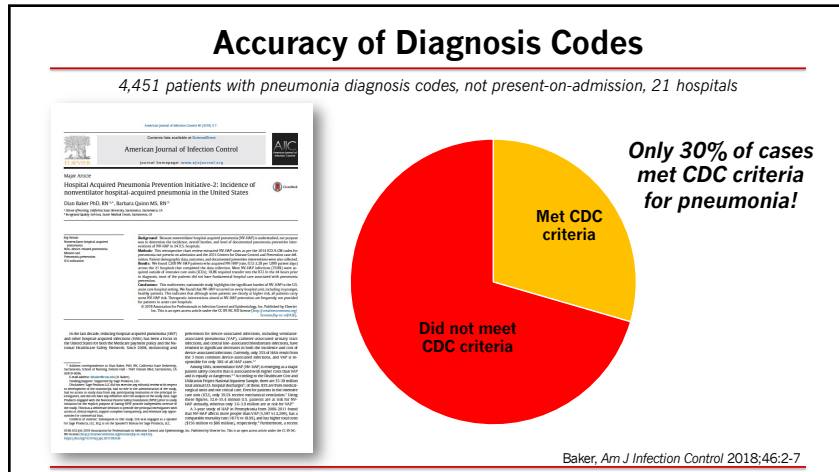
## The Challenge of NV-HAP Diagnosis & Surveillance

<p><u><b>Imaging</b></u></p> <p>New or progressive and persistent</p> <p>Infiltrate or Consolidation or Cavitation</p>	<p><u><b>Systemic Signs</b></u></p> <p>Fever <math>&gt;38.0^{\circ}\text{C}</math></p> <p>WBC <math>&lt;4\text{K}</math> or <math>&gt;12\text{K}</math></p> <p>Altered mental status</p>	<p><u><b>Pulmonary Signs</b></u></p> <p>New onset purulent sputum, change in character of sputum, increased secretions</p> <p>New onset or worsening cough, dyspnea, or tachypnea</p> <p>Rales of bronchial breath sounds</p> <p>Worsening gas exchange, increased oxygen requirement</p>
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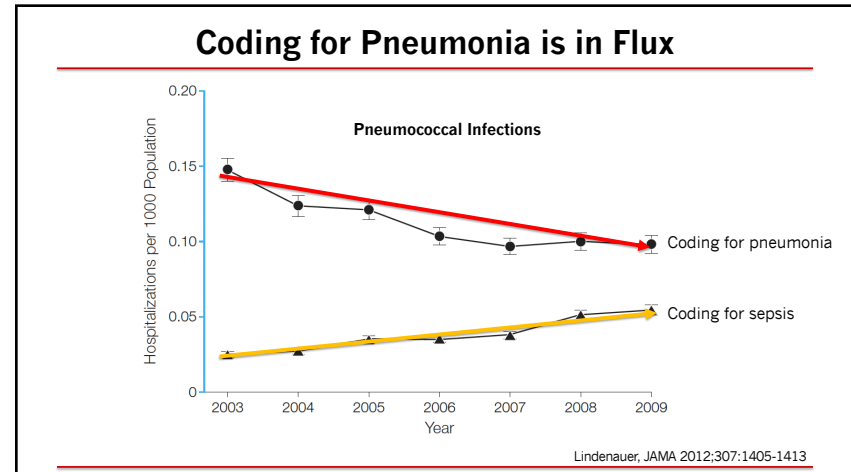


Am J Infect Control 2010;38:237





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**JAMA Network | Open.**

Original Investigation | Infectious Diseases

## Development and Assessment of Objective Surveillance Definitions for Nonventilator Hospital-Acquired Pneumonia

Wenjing Ji, PhD; Caroline McKenna, MPH; Aileen Ochoa, MPH; Haiyan Ramirez Battle, MD; Jessica Young, PhD; Zilu Zhang, MS; Chanu Rhee, MD, MPH; Roger Clark, DO; Erica S. Shenoy, MD, PhD; David Hooper, MD; Michael Klompas, MD, MPH; for the CDC Prevention Epicenters Program

**Abstract**

**IMPORTANCE** Hospital-acquired pneumonia is the most common health care-associated infection in the United States. Most cases occur in nonventilated patients, but many hospitals track hospital-acquired pneumonia only in ventilated patients because of the complexity and subjectivity of conducting surveillance for large numbers of nonventilated patients.

**OBJECTIVE** To propose and assess potentially objective, efficient, and reproducible surveillance definitions for nonventilator hospital-acquired pneumonia (NV-HAP) using routine clinical data stored in electronic health record systems.

**DESIGN, SETTING, AND PARTICIPANTS** This cohort study was conducted in 2 tertiary referral and 3 community hospitals in Massachusetts between May 1, 2005, and July 1, 2008. All nonventilated

**Key Points**

**Question** Is it possible to conduct operational surveillance using the clinical data routinely recorded in electronic health records to identify nonventilated adults with hospital-acquired pneumonia?

**Findings** In this cohort study of 310 651 patients with 489 519 admissions, an electronic surveillance definition based on worsening oxygenation, at least 3 days of new antibiotics, fever or leukocytosis, and performance of chest imaging

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### NV-HAP Electronic Surveillance Definition

**Worsening Oxygenation**

*Drop in SpO2 to <95% on room air, initiation of supplemental oxygen, or escalation of supplemental oxygen*

**Fever or Leukocytosis**

*WBC <4 or >12 or Temp <36° or >38° C*

**Performance of Chest Imaging**

*Chest X-ray or CT Scan*

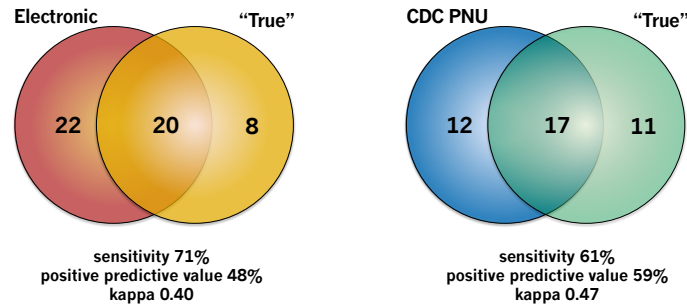
**≥3 days of New Antibiotics**

*New = not given in previous 2 days; can be different antibiotics for each day so long as each one is new*

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## Correlation with “True” Pneumonia per Reviewer

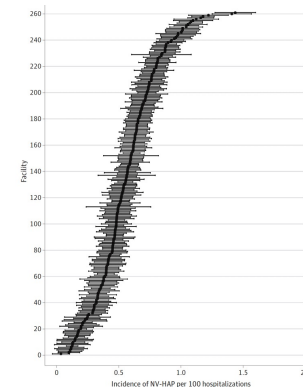
Chart review exercise to assess correlation between the electronic NV-HAP surveillance definition vs “truth” (N=120)



Ramirez Batlle, *ICHE* 2020;41:219-221

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## NV-HAP Incidence and Characteristics



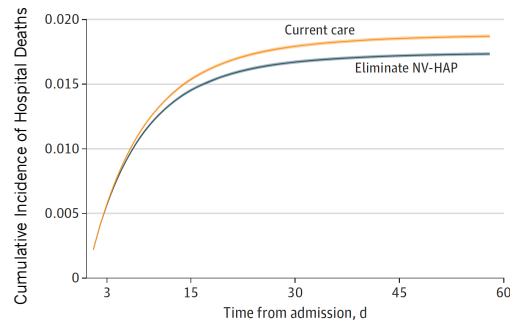
- Electronic NV-HAP criteria applied to 284 US hospitals, 2015-2020
- Median 0.55 NV-HAP events per 100 admissions
- 75% of cases in non-ICU settings
- Crude mortality 22.4%
- Median length-of-stay 16 days (IQR 11-26)
- 250 charts reviewed for accuracy: pneumonia confirmed in 81%

Jones, *JAMA Network Open* 2023;6(5):e2314185

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## What if we could eliminate NV-HAP?

Analysis of 6.02 million admissions to 284 US hospitals. Modeled impact of eliminating NV-HAP accounting for hospital factors and patients' demographics, comorbidities, service, daily vital signs, & daily laboratory test results



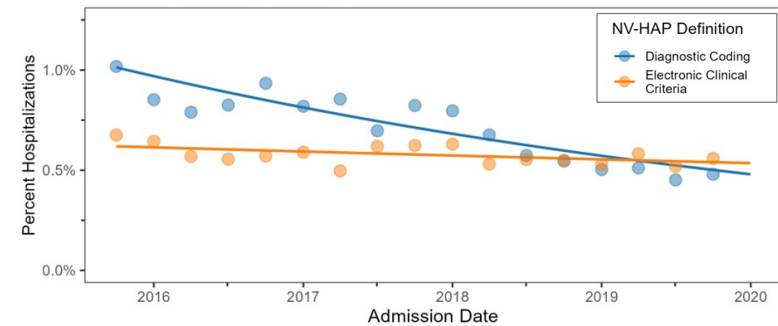
**Eliminating NV-HAP would reduce overall hospital mortality by ~7.3%**

Jones, *JAMA Network Open* 2023;6(5):e2314185

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## NV-HAP Trends with Codes vs Electronic Clinical Criteria

Trends in NV-HAP rates detected with diagnosis codes vs electronic clinical criteria, 17 VA hospitals, 2015-2019

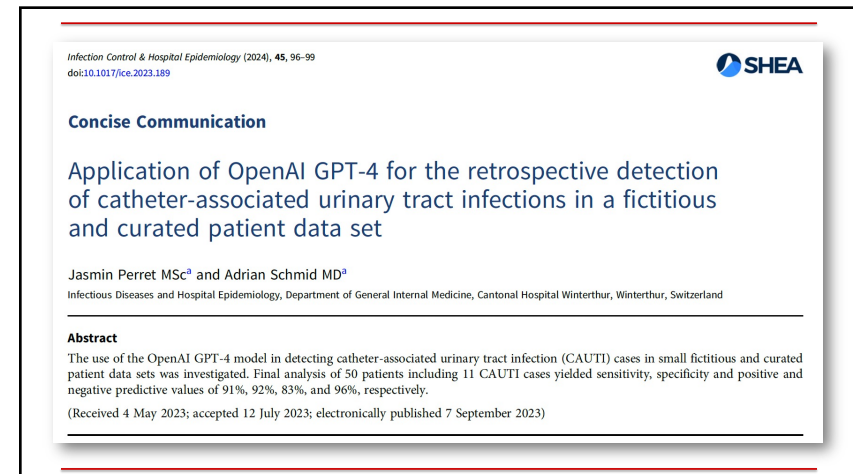


Jones 2024, unpublished data

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### Large Language Models for CAUTI Surveillance

- Researchers in Switzerland created two fictitious sets of patients to feed to GPT-4. Case descriptions brief due to token limits (max 8,192)
- Dataset 1 (structured data only, 79 patients)
  - GPT identified 6 CAUTIs (sensitivity 80%, positive predictive value 67%)
    - 4 confirmed
    - 2 false positives (one had 3 species in the urine, one did not meet all criteria in infection window)
    - 1 false negative (catheter removed day before infection)
  - Performance improved with training on the same patients
- Dataset 2 (structured data + symptoms, 50 patients)
  - GPT identified 12 CAUTIs
    - 10 confirmed
    - 2 false positives (catheter only present one day, not all infection criteria in infection window)
    - 1 false negative (catheter removed the day before infection)

Perret, ICHE 2024;45:96-99

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### Performance of a large language model for identifying central line-associated bloodstream infections (CLABSI) using real clinical notes

Guillermo Rodriguez-Nava MD<sup>1</sup>, Goar Egoryan MD<sup>2</sup>, Katherine E. Goodman PhD, JD<sup>3,4</sup>, Daniel J. Morgan MD, MS<sup>3,5</sup> and Jorge L. Salinas MD<sup>1</sup>

<sup>1</sup>Division of Infectious Diseases & Geographic Medicine, Department of Medicine, Stanford University School of Medicine, Stanford, CA, USA, <sup>2</sup>Division of Oncology, Department of Medicine, Stanford University School of Medicine, Stanford, CA, USA, <sup>3</sup>Department of Epidemiology and Public Health, University of Maryland School of Medicine, Baltimore, MD, USA, <sup>4</sup>University of Maryland Institute for Health Computing, Bethesda, MD, USA and <sup>5</sup>VA Maryland Healthcare System, Baltimore, MD, USA

**Abstract**  
We evaluated one of the first secure large language models approved for protected health information, for identifying central line-associated bloodstream infections (CLABSIs) using real clinical notes. Despite no pretraining, the model demonstrated rapid assessment and high sensitivity for CLABSI identification. Performance would improve with access to more patient data.  
(Received 7 June 2024; accepted 30 August 2024)

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## Large language models for CLABSI surveillance

- Pilot study in Stanford of CLABSI surveillance using GPT 4.0
  - 40 patients already reviewed by IPs for possible CLABSI
    - 20 with CLABSI, 20 without CLABSI
  - Provided GPT with just the index positive blood culture result & the last two progress notes in the infection window period
  - GPT detected 16/20 CLABSIs and 7/20 non-CLABSIs
    - Sensitivity 80%, Specificity 35%
  - Investigators attributed 11/17 incorrect classifications to missing information elsewhere in the chart (i.e. not in the last 2 notes)
    - e.g. BSI present on admission, another source for BSI, additional clinical signs, etc.
  - Providing this additional information to GPT improved performance
    - Sensitivity 90%, Specificity 75%
  - Residual errors due to misclassification of commensals and mucosal barrier injury organisms as pathogens, misclassification of CLABSI organism as commensal, misattribution of BSI to pancreatitis.

Rodriguez-Nava, *ICHE* 2024; ePub: 10.1017/ice.2024.164

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NEJM AI 2024;1(11)  
DOI: 10.1056/AIcs2400420

### CASE STUDY

## Large Language Models for More Efficient Reporting of Hospital Quality Measures

Aaron Boussina , Ph.D.,<sup>1</sup> Rishivardhan Krishnamoorthy , M.S.,<sup>1</sup> Kimberly Quintero , R.N., M.S.,<sup>2</sup> Shreyansh Joshi , Gabriel Wardi , M.D.,<sup>1,3,4</sup> Hayden Pour , M.S.,<sup>1</sup> Nicholas Hilbert , R.N., M.S.N.,<sup>1</sup> Atul Malhotra , M.D.,<sup>1</sup> Michael Hegar , M.D.,<sup>1</sup> Amy M. Sitapati , M.D.,<sup>1</sup> Chad VanDerBerg , M.P.H.,<sup>2</sup> Karandeep Singh , M.D., M.M.Sc.,<sup>1</sup> Christopher A. Longhurst , M.D., M.S.,<sup>1</sup> and Shamim Nemat , Ph.D.<sup>1</sup>

Received: April 25, 2024; Revised: August 9, 2025; Accepted: August 16, 2024; Published: October 21, 2024

### Abstract

Hospital quality measures are a vital component of a learning health system, yet they can be costly to report, statistically underpowered, and inconsistent due to poor interrater reliability. Large language models (LLMs) have recently demonstrated impressive performance on health care-related tasks and offer a promising way to provide accurate abstraction of complete charts at scale. To evaluate this approach, we deployed an LLM-based system that ingests Fast Healthcare Interoperability Resources data and outputs a completed Severe Sepsis and Septic Shock Management Bundle (SEP-1) abstraction. We tested the system on a sample of 100 manual SEP-1 abstractions that University of California San Diego Health reported to the Centers for Medicare & Medicaid Services in 2022. The LLM system achieved agreement with manual abstractors on the measure category assignment in 90 of the abstractions (90%;  $\kappa=0.82$ ; 95% confidence interval, 0.71 to 0.92). Expert review of the 10 discordant cases identified four that were mistakes introduced by manual abstraction. This pilot study suggests that LLMs using interoperable electronic health record data may perform accurate abstractions for complex quality measures. (Funded by the National Institute of Allergy and Infectious Diseases [1R42AI177108-1] and others.)

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## SEP-1 Mandatory Sepsis Bundle

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## SEP-1 Mandatory Sepsis Bundle

3 Hours

Measure lactate

Draw blood cultures

Administer broad spectrum antibiotics

Shock? 30cc/kg IV crystalloids

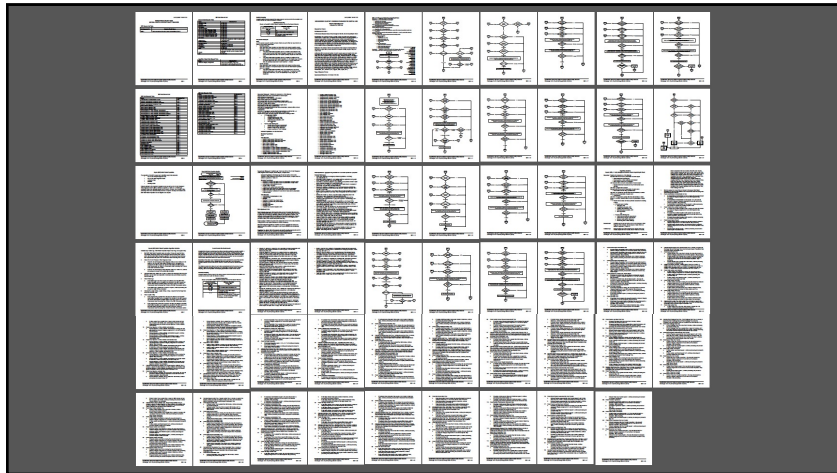
6 Hours

Repeat lactate if initial level was elevated

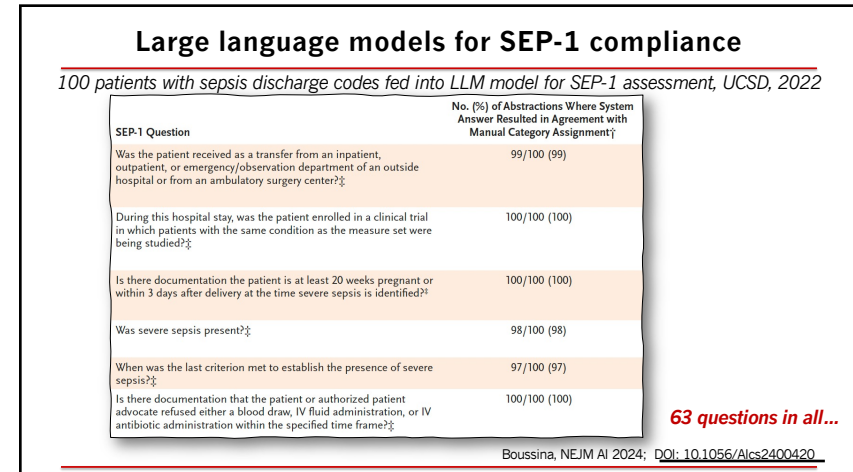
Administer vasopressors if still hypotensive

Reassess volume status & tissue perfusion

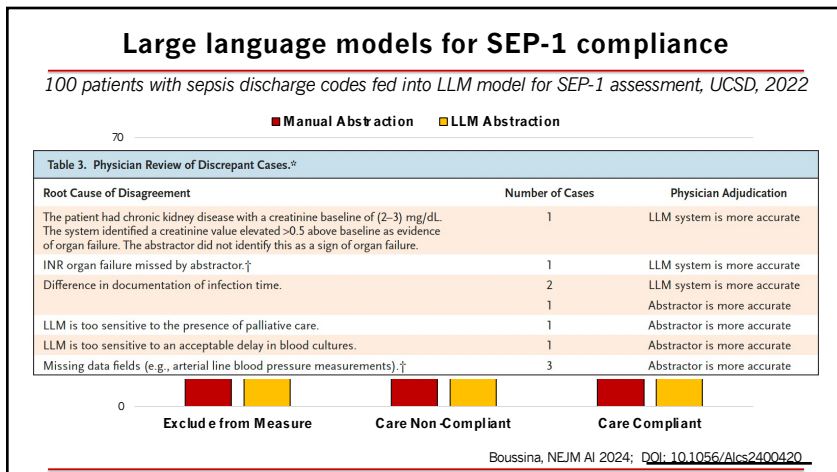
48



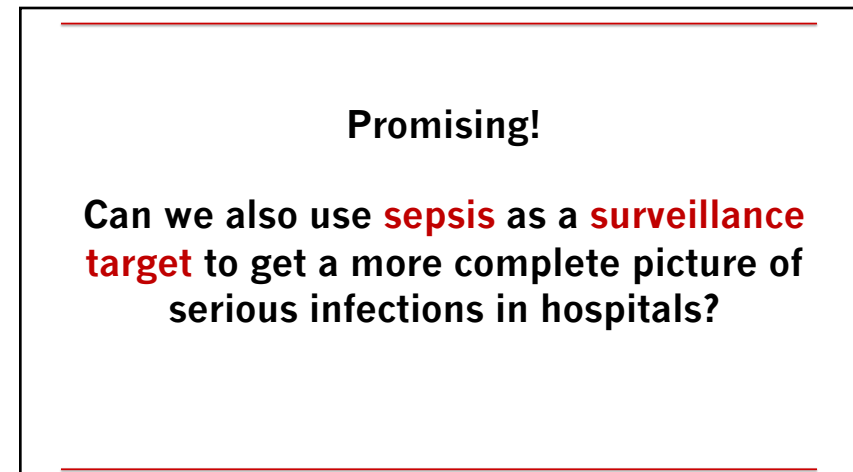
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## GET AHEAD OF SEPSIS

KNOW THE RISKS. SPOT THE SIGNS. ACT FAST.



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## Sepsis Surveillance Methods Using Diagnosis Codes

### Angus Method:

**1286** infection codes, **13** organ dysfunction codes

### Martin Method:

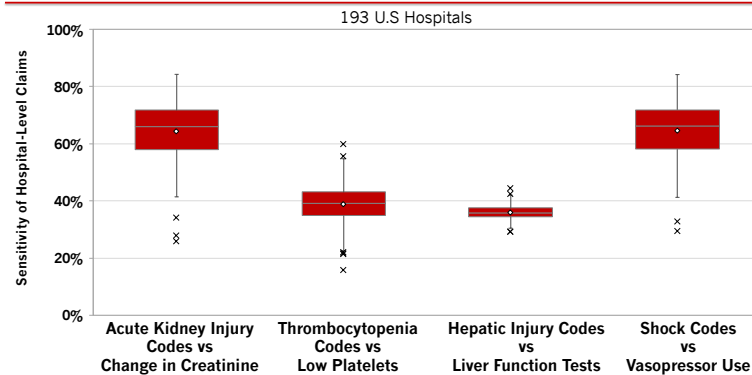
**6** infection codes, **13** organ dysfunction codes

### Dombrovskiy Method:

**18** infection codes, **22** organ dysfunction codes

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## Sensitivity of Diagnosis Codes vs Clinical Criteria

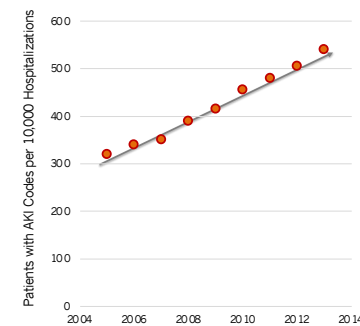


Rhee, *Crit Care Med* 2019;47:493-500

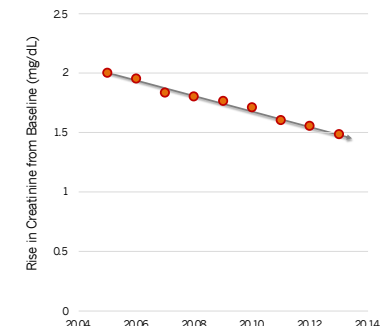
55

## Changes in Coding Thresholds

### Count of Patients with Acute Kidney Injury Codes



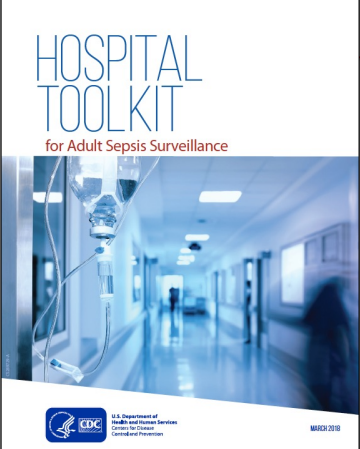
### Mean Change in Creatinine



Rhee et al, *Critical Care* 2015;19:338

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**HOSPITAL TOOLKIT**  
for Adult Sepsis Surveillance

U.S. Department of Health and Human Services  
Centers for Disease Control and Prevention

MARCH 2018

### CDC to the rescue!

- CDC created an EHR-based surveillance definition for sepsis called “Adult Sepsis Events”
- Uses detailed EHR-data to identify patients with suspected infection + organ dysfunction
- Implementation toolkit available that describes the required data elements, how to organize them, and provides analytic code for event detection

[cdc.gov/sepsis/clinicaltools](http://cdc.gov/sepsis/clinicaltools)

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### Clinical Criteria for Sepsis

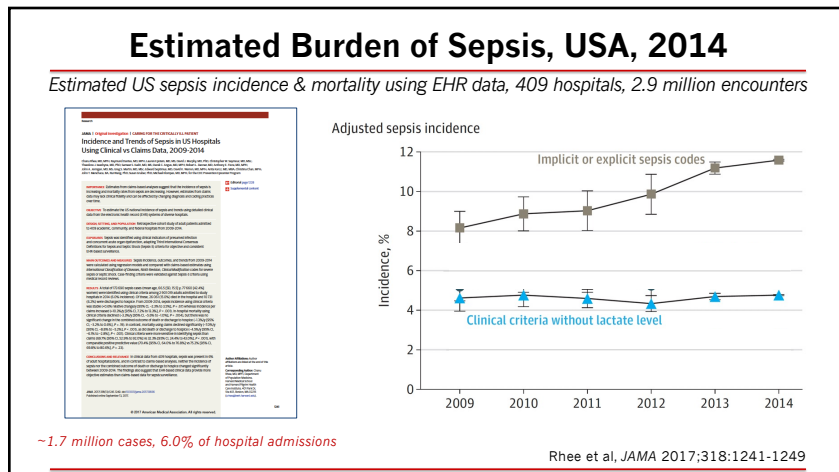
<b>Suspected Infection</b>	<b>Order for blood cultures</b> <i>and</i> <b>At least four days of new antibiotics</b>
<b>Concurrent Organ Dysfunction</b>	<b>Vasopressors</b> <i>or</i> <b>Initiation of mechanical ventilation</b> <i>or</i> <b>Doubling in creatinine (exclude dialysis pts)</b> <i>or</i> <b>Rise in bilirubin to <math>\geq 2.0</math>mg/dL</b> <i>or</i> <b>Fall in platelets to <math>\leq 100</math> cells/<math>\mu</math>L</b> <i>or</i> <b>Serum lactate <math>\geq 2.0</math> mmol/L</b>

**If all criteria are met on hospital day  $\geq 3$  than classify as**

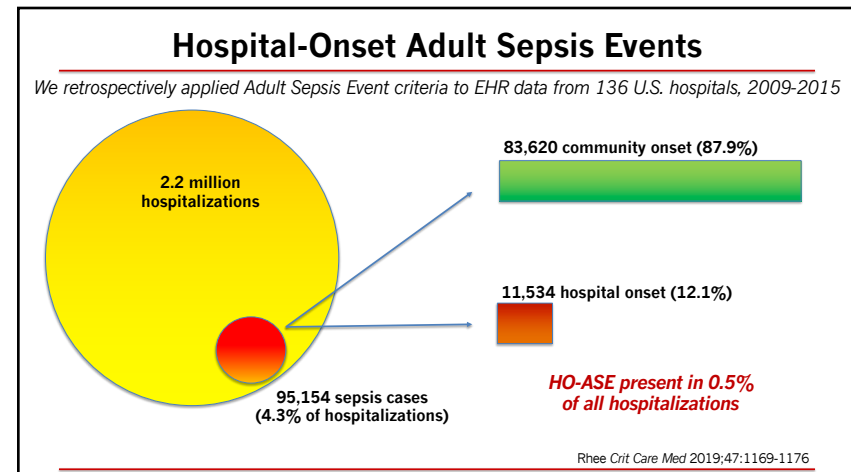
**Hospital-Onset Adult Sepsis Event (HO-ASE)**

Rhee, JAMA 2017;318:1241-1249

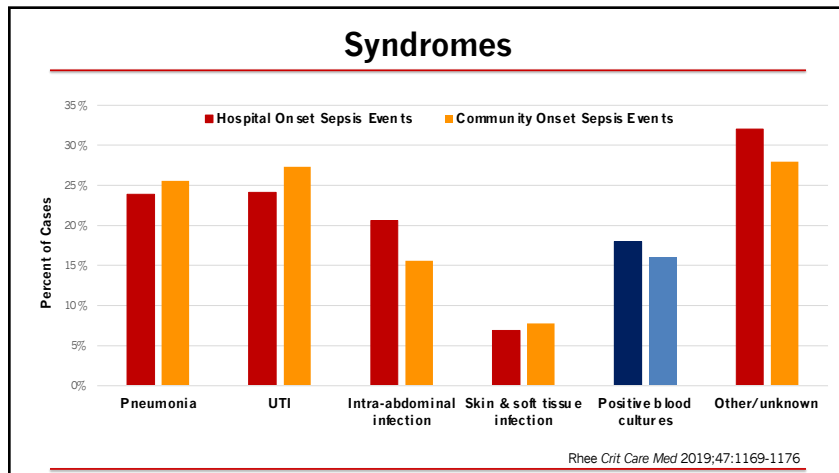
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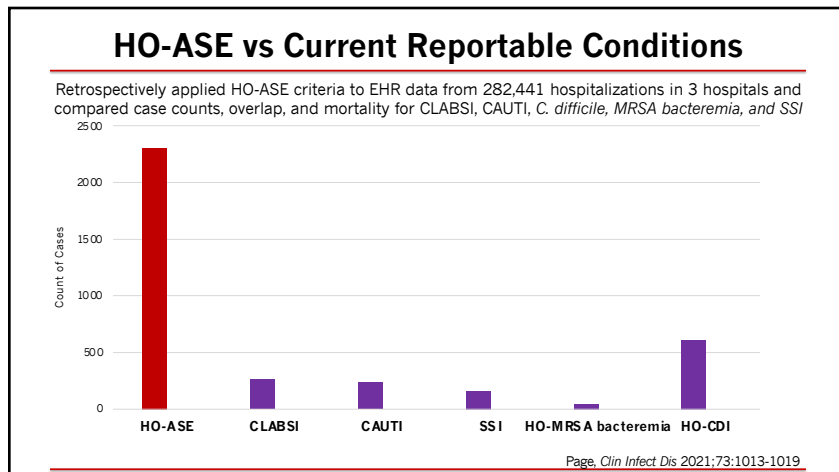
60



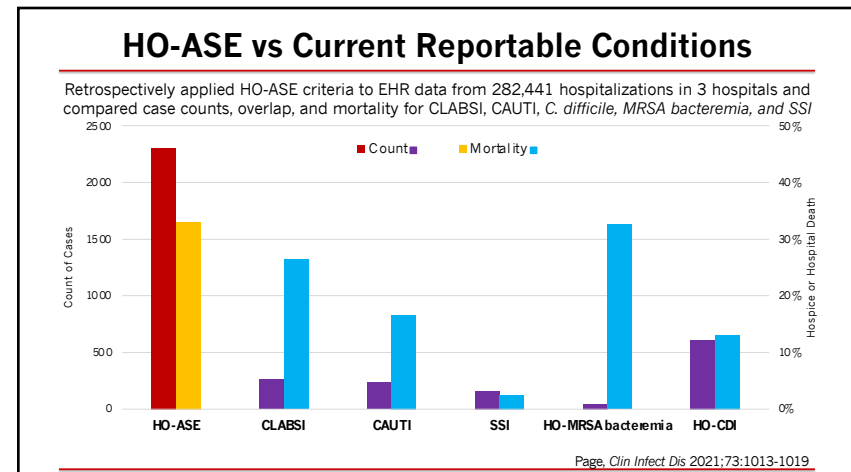
61

## How does Hospital-Onset Adult Sepsis Event surveillance compare to current surveillance targets?

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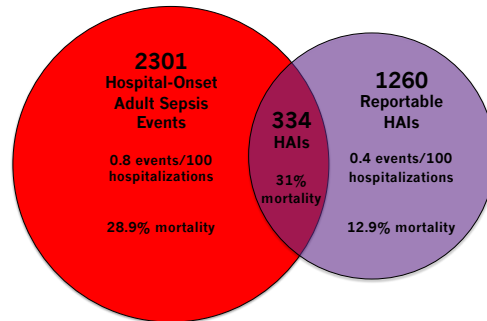


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## HO-ASE vs Current Reportable Conditions

Retrospectively applied HO-ASE criteria to EHR data from 282,441 hospitalizations in 3 hospitals and compared case counts, overlap, and mortality for CLABSI, CAUTI, *C. difficile*, MRSA bacteremia, and SSI



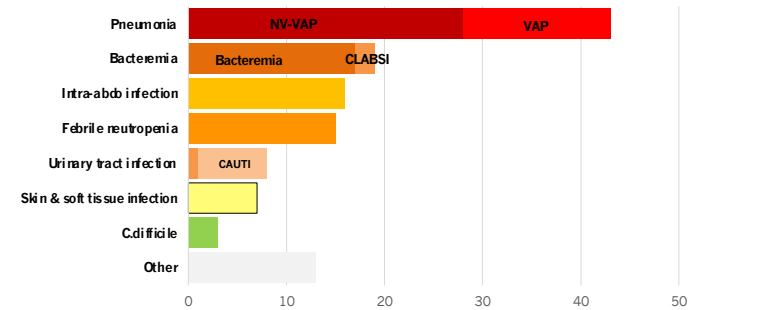
*HO-ASE identified a large number of high mortality hospital-onset infections missed by current reportables*

Page, Clin Infect Dis 2021;73:1013-1019

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## Distribution of Infections Identified by HO-ASE

Retrospectively applied HO-ASE criteria to EHR data from 282,441 hospitalizations in 3 hospitals and compared case counts, overlap, and mortality for CLABSI, CAUTI, *C. difficile*, MRSA bacteremia, and SSI

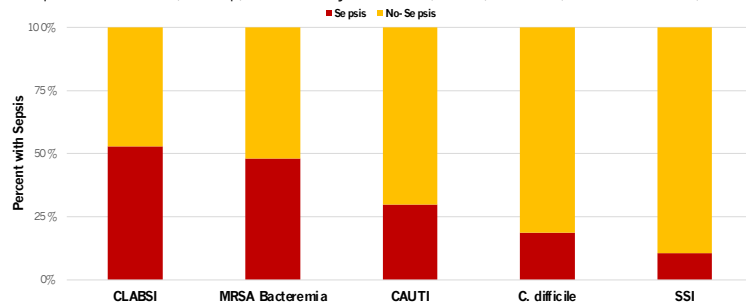


Page, Clin Infect Dis 2021;73:1013-1019

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## Proportion of Reportables that Qualified as Sepsis

Retrospectively applied HO-ASE criteria to EHR data from 282,441 hospitalizations in 3 hospitals and compared case counts, overlap, and mortality for CLABSI, CAUTI, *C. difficile*, MRSA bacteremia, and SSI



Page, Clin Infect Dis 2021;73:1013-1019

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## Summary

- **Our current surveillance strategy was developed before electronic clinical data became ubiquitous**
  - Focuses on relatively few infections, uses clinically nuanced definitions, requires manual chart review
  - Mandatory reporting requirements have intensified our limited focus
  - Our current approach is labor intensive yet subjective – undermines benchmarking
  - We are missing many high stakes hospital-acquired infections
- **Promising new surveillance strategies emerging (HOB, NV-HAP, HO-ASE)**
  - Use detailed electronic health record data rather than discharge codes or chart reviews
  - Can be efficiently applied to large populations
  - Makes surveillance objective and reproducible
  - Increases the number and breadth of nosocomial infections under surveillance (ICU and non-ICU, device and non-device)

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