# CONNECTICUT **HEALTHCARE ASSOCIATED INFECTIONS (HAI) PREVENTION PLAN**

October 1

2015

In 2009 in response to the increasing concerns about the public health impact of healthcare-associated infections (HAIs), the US Department of Health and Human Services (HHS) developed an Action Plan template for all states to help prevent HAIs. In wake of the Ebola outbreak in West Africa in 2015 the plan was updated to include new Infection prevention activities for severe emerging and unusual infections. All states have been asked to update their plans by October 1, 2015. The HHS Action Plan includes recommendations for surveillance, research, communication, and metrics for measuring progress toward national goals. This Plan gives an historical overview of infection prevention actives achieved in Connecticut from 2009 forward and continuing into 2015-2018.

Submitted to the Centers for **Disease Control and Prevention** (CDC) Division of Healthcare **Quality Promotion (DHQP) Infection prevention Assessment** and Readiness (ICAR) Team

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#### Introduction

Healthcare Associated Infections (HAI) are infections that occur during, or as a consequence of, the provision of healthcare. HAIs are a significant medical and public health problem across the nation, and in Connecticut. Not only do HAIs cause suffering for patients and harm that is often preventable, but they also increase hospital lengths of stay and add considerably to healthcare costs.

In acute care hospitals alone in 2011, healthcare associated infections affected an estimated 648,000 patients with 721,800 health care—associated infections.<sup>1</sup> While more current data on hospital-based mortality from this 2011 study has not yet been published, a widely reported estimate from 2002 is that approximately 99,000 deaths occur annually associated with an HAI during an acute care hospitalization.<sup>2</sup> The estimated annual direct medical costs of HAI to hospitals in the United States ranged from \$28.4 to \$45 billion. The benefits of prevention range from \$5.7 to as much as \$31.5 billion.<sup>3</sup>

It should also be noted that much patient care is delivered outside of acute care hospitals, across a spectrum of intensity ranging from long term acute care, long term care, ambulatory facilities, to homecare. To account for the true costs of HAI, and to fully appreciate the human and economic costs and importance of this problem, costs in these other settings also need to be taken into account and research will need to be done to estimate these costs and impact on public health.

The CDC estimates that 2 million persons in the United States develop serious infections with antibiotic resistant bacteria each year, leading to 23,000 deaths. Nearly 250,000 patients are hospitalized and at least 14,000 people die each year due to *Clostridium difficile (C. difficile)* infections, which are facilitated by antibiotic use. This may lead to \$20 billion in excess direct healthcare costs, with additional costs to society as high as \$35 billion annually. <sup>4</sup>

The over or misuse of antibiotics is the single most important factor leading to antibiotic resistance. The development of antibiotic resistance began as soon as antibiotics became widely available in the 1940s, the development of antibiotic resistance has accelerated in recent decades. This coupled with a diminution of new antibiotics and other antimicrobial drugs in the developmental pipeline has created a public health crisis. Some strains of bacteria have developed that are resistant to nearly all or all available antibiotic, raising the specter of a return to the preantibiotic eras in which infections would frequently be untreatable. This has led the CDC to call on federal and state public health agencies to collaborate with healthcare providers and facilities to address this serious public health and economic threat through the fostering of antimicrobial stewardship programs in healthcare settings.

<sup>&</sup>lt;sup>1</sup> Magill S, Edwards J, et.al. Multistate Point-Prevalence Survey of Health Care—Associated Infections N Engl J Med 2014;370:1198-208.

<sup>&</sup>lt;sup>2</sup> Klevens RM, Edwards JR, et al. Estimating health care-associated infections and deaths in U.S. hospitals, 2002. *Public Health Reports* 2007 March-Apr; 122(2): 160-6. Scott <sup>3</sup> Scott, RD, The Direct Medical Costs of Healthcare-Associated Infections in U.S. Hospitals and the Benefits of Prevention, CDC, accessed on 9/8/15 at: http://www.cdc.gov/HAI/pdfs/hai/Scott CostPaper.pdf

<sup>4</sup> CDC, Antibiotic Resistance Threats in the United States, 2013, accessed on 9/8/15 at: http://www.cdc.gov/drugresistance/threat-report-2013/pdf/ar-threats-2013-508.pdf#page=1

In addition to addressing antibiotic and antimicrobial resistance, the scope of public health HAI programs has grown to address other HAI related issues beyond the initial focus on surveillance for and reporting of particular types of infections acquired during healthcare associated with medical devices or procedures in individual patients, such as central line associated bloodstream infections, catheter associated urinary tract infections, and surgical site infections. Such areas of expansion include patient exposure to contaminated medical products or equipment that may be associated with infection in patients, or which have not been associated with documented infections, but which have put groups of patients at risk. Another has been addressing outbreaks and clusters of HAIs to find underlying facilityspecific risks and rectify them. HAI programs have been asked to participate in the investigation of breaches of Infection prevention and injection safety, sometimes involving impaired healthcare providers, or intentional drug diversion. The impact of the Ebola outbreak in West Africa, and its impact on the United States, has highlighted an important role for HAI programs and their collaborators in the planning and preparedness for adequate Infection prevention while caring for serious emerging infectious diseases, such as the Viral Hemorrhagic Fevers (including Ebola), Middle Eastern Respiratory Virus (MERS) Co-V, pandemic influenza, and others. This has also highlighted the need for active HAI program assessment and technical assistance of the infection prevention infrastructure, training, and quality control in healthcare settings across the spectrum of healthcare. As the drive for quality and cost containment has come to include infections along with other sources of preventable harm, HAI programs have become involved in the collection and analysis of data that is used for the federal Centers for Medicare and Medicaid Services' (CMS) quality improvement and payment programs, which involves both data on HAIs and on vaccination status of healthcare staff.

There is a wide variety of stakeholders affected by HAIs or who have a role in preventing them. These include healthcare providers, patients, patient advocates, professional associations, facility associations, payors, and public health agencies. There is a critical need to collaborate among these groups to foster coordinated and effective action, and to find solutions that involve coordinated action among the stakeholders. One of the traditional activities of public health programs, such as HAI programs, has been to act as a convener of stakeholders to lead concerted coordinated action. Such collaboration is fostered by multidisciplinary groups and technical advisory groups to advise HAI programs on strategies and activities and to evaluate the progress in achieving program goals.

As the scope and complexity of the HAI programs grow, it is more important than ever to identify priorities to address the currently preventable fraction and not waste time and action, and avoid opportunity costs. This indicates the need to carefully plan. The detailed living plan, such as the one that follows, gives an historical record of what has been accomplished since 2009 when it was first published, and the continuing plans for 2015-2018 of the HAI program and its collaborators to guide their actions going forward. It can ensure coordination, foster evaluation of progress, incorporate changes, and find synergies. Further, it can ensure proactive rather than reactive efforts that will accomplish the goal of eliminating the preventable fraction of HAIs effectively and efficiently, without interfering with the other core functions of the provision of healthcare.

### **Template for State Healthcare-associated Infection Plan**

In response to the increasing concerns about the public health impact of healthcare-associated infections (HAIs), the US Department of Health and Human Services (HHS) has developed an Action Plan to help prevent Healthcare-associated Infections. The HHS Action Plan includes recommendations for surveillance, research, communication, and metrics for measuring progress toward national goals. Three overarching priorities have been identified:

- Progress toward 5-year national prevention targets (e.g., 50-70% reduction in bloodstream infections);
- Improve use and quality of the metrics and supporting systems needed to assess progress towards meeting the targets; and
- Prioritization and broad implementation of current evidence-based prevention recommendations

Background: The 2009 Omnibus bill required states who received Preventive Health and Health Services (PHHS) Block Grant funds to certify that they would submit a plan to reduce HAIs to the Secretary of Health and Human Services not later than January 1, 2010. In order to assist states in responding within the short timeline required by that language and to facilitate coordination with national HAI prevention efforts, the Centers for Disease Control and Prevention (CDC) created a template to assist state planning efforts.

This template helps to ensure progress toward national prevention targets as described in the HHS Action Plan. CDC is leading the implementation of recommendations on national prevention targets and metrics and states should tailor the plan to their state-specific needs.

Initial emphasis for HAI prevention focused on acute care, inpatient settings, and then expanded to outpatient settings. The public health model of population-based healthcare delivery places health departments in a unique and important role in this area, particularly given shifts in healthcare delivery from acute care settings to ambulatory and long term care settings. In non-hospital settings, infection control and oversight have been lacking which have resulted in outbreaks which can have a wide-ranging and substantial impact on affected communities. At the same time, trends toward mandatory reporting of HAIs from hospitals reflect increased demand for accountability from the public.

The State HAI Action Plan template targets the following areas:

- 1. Enhance HAI Program Infrastructure
- 2. Surveillance, Detection, Reporting, and Response
- 3. Prevention
- 4. Evaluation, Oversight, and Communication

## With new Ebola-related, infection control activities, the following two tables have been added to reflect those activities:

- 5. Infection Control Assessment and Response (Ebola-associated activity from FOA Supplement, CK14-1401PPHFSUPP15, Project A)
- 6. Targeted Healthcare Infection Prevention Programs (Ebola-associated activity from FOA Supplement, CK14-1401PPHFSUPP15, Project B)

#### **Framework and Funding for Prevention of HAIs**

CDC's framework for the prevention of HAIs builds on a coordinated effort of federal, state, and partner organizations and is based on a collaborative public health approach that includes surveillance, outbreak response, infection control, research, training, education, and systematic implementation of prevention practices. Legislation in support of HAI prevention provides a unique opportunity to strengthen existing state capacity for prevention efforts.

#### Template for developing HAI plan

The following template provides choices for enhancing state HAI prevention activities in the six areas identified above. For each section, please choose elements which best support current activities or planned activities. Current activities are those in which the state is presently engaged and includes activities that are scheduled to begin using currently available resources. Planned activities represent future directions the state would like to move in to meet currently unmet needs, contingent on available resources and competing priorities. A section for additional activities is included to accommodate plans beyond the principal categories.

#### 1. Enhance HAI program infrastructure

Successful HAI prevention requires close integration and collaboration with state and local infection prevention activities and systems. Consistency and compatibility of HAI data collected across facilities will allow for greater success in reaching state and national goals. Please select areas for development or enhancement of state HAI surveillance, prevention, and control efforts.

**Table 1:** State infrastructure planning for HAI surveillance, prevention, and control.

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
Complete	Planned	1. Establish statewide HAI prevention leadership through the formation of multidisciplinary group or state HAI advisory council  i. Collaborate with local and regional partners (e.g., state hospital associations, professional societies for infection control and healthcare epidemiology, academic organizations, laboratorians, and networks of acute care hospitals and long term care facilities).	Initiated 2006 and ongoing
$\boxtimes$	$\boxtimes$	In 2006, the Connecticut (CT) General Assembly passed Public Act 06-142, <i>An Act Concerning Hospital Acquired Infections</i> , now codified in state statue as CGS 19a 490 o. It created an 11-member Advisory Committee on Healthcare Associated Infections to advise the Department of Public Health (DPH) on the development, operation, and monitoring of a mandatory Healthcare Associated Infections (HAI) reporting system. The Committee included representation from consumers, the public, hospital infection preventionists, infectious disease physicians, the CT State Medical Society (CSMS), the CT Hospital Association (CHA), and DPH as outlined in Appendix 2. <sup>1</sup> The Advisory Committee has met quarterly since inception.  ii. NEW: Include hospital preparedness partners (e.g., hospital/healthcare coalitions funded through the ASPR	October 2015
		Hospital Preparedness Program). Additional representation from accrediting and/or licensing agency with surveyor authority is ideal.  Beginning in 2015, members of the current HAI Advisory Committee were asked to propose addition members to expand the committee in the wake of the Ebola outbreak. Proposed members are:  Pro Health (Group of private MDs)  Long Term Care Association  Long Term Care Infection Preventionists  CT Association of Ambulatory Surgical Centers	
		Long Term Acute Care Facilities (LTACs) Hospice School Based Clinics Urgent Care Centers Other Out Pt Clinics (MD owned) Nursing Homes	

<sup>&</sup>lt;sup>1</sup> CDC template footnote documentation begins with Appendix 1 (page 15). DPH footnote documentation of Appendix 2 begins on this page.

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		Home Health Agencies	
		Private Insurance Company healthcare payors	
		Veterinarians	
		Dentists	
		Hospital Preparedness Coordinators	
		Connecticut Medical Society	Initiated 2008 and ongoing
		iii. NEW: Engage HAI advisory committee in potential roles and activities to improve antibiotic use in the state (antibiotic stewardship)	
		The Committee has also discussed developing and revising subcommittees:	
		1. Antimicrobial Stewardship	
		2. HAI Prevention	
		3. HAI Public Education	
		iv. NEW: Engage HAI advisory committee in activities to increase health department's access to data and subsequently use those data in prevention efforts	
		The HAI Advisory Committee made its initial recommendations in 2007 which launched the program in CT:  1. Participate in the CDC's National Healthcare Safety Network (NHSN) reporting system,	Initiated 2007 and ongoing
		2. Utilize one NHSN Patient Safety Module,	
		3. Provide education and training on CDC and NHSN enrollment, HAI definitions, data entry, and analysis,	
		4. Report outcome data on Central Line Associated Blood Stream Infections (CLABSIs) in one adult intensive care unit (ICU) per acute care hospital, and one pediatric ICU in acute care children's hospital(s) in the state,	
		5. Utilize first year data collection as a pilot to assure accuracy and completeness of reporting,	
		6. Plan validation testing of year two data and beyond,	
		7. Expand DPH technical staff for data analysis, education and training,	
		8. Use results to implement evidence-based prevention methods.	
		Beginning in 2008 CT acute care hospitals reported CLABSIs from one adult ICU per hospital and all pediatric ICUs via the NHSN data system.	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		In 2012 the federal Centers for Medicaid and Medicare Services (CMS) expanded HAI reporting requirements for healthcare institutions as a condition of receiving annual payments. As a result of these requirements the HAI Advisory Committee recommended that HAI reporting in CT followed the CMS reporting schedule. In January 2012, CMS required that acute care hospitals continue to report CLABSIs in adult, pediatric and neonatal ICUs and added catheter associated urinary tract infections (CAUTIs), and surgical site infections (SSI) in two operative sites. Long term care hospitals had to begin reporting CLABSIs and CAUTIs, and hemodialysis centers had to begin reporting dialysis events using NHSN definitions.	Initiated 2012 and ongoing
		Beginning in 2013 CMS required acute care hospitals in the state to report healthcare personnel (HCP) influenza vaccination rates for their employees, contract workers, students, licensed independent practitioners and volunteers. CMS schedule for HAI reporting and additional reporting criteria for reporting is outlined in Appendix 3.	Initiated 2013 and ongoing
		<ul> <li>v. Identify specific HAI prevention targets consistent with HHS priorities</li> <li>1. Central Line-associated Blood Stream Infections (CLABSI)</li> <li>2. Clostridium difficile Infections (CDI)</li> <li>3. Catheter-associated Urinary Tract Infections (CAUTI)</li> <li>4. Methicillin-resistant Staphylococcus aureus (MRSA) Infections</li> <li>5. Surgical Site Infections (SSI)</li> </ul>	2009 and Ongoing
		Other activities or descriptions:	
Complete		2. Establish an HAI surveillance prevention and control program i. Designate a State HAI Prevention Coordinator  In 2006, the Connecticut Legislature directed the DPH to create a state public health HAI program and to establish a state HAI Advisory Committee. In 2008, 3 FTE state-funded positions were created to oversee the HAI program. These positions included the state HAI coordinator, Richard Melchreit, a public health physician with 25 years of experience; a senior epidemiologist, Lauren Backman with nursing, microbiology and MHS degree and work experience as an infection preventionist; and an epidemiologist who provided data analysis skills and National Healthcare Safety Network (NHSN)	2006

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		technical assistance. In 2014, The HAI program experienced a permanent reduction in state funding that eliminated a key epidemiology position. This lost position makes it difficult to maintain the high quality HAI program standards now expected by CT healthcare facilities. The HAI program is part of the Infectious Disease (ID) Section under the direction of Matthew Cartter, MD, MPH, the State Epidemiologist. The ID Section also contains the Epidemiology and Emerging Infections (EIP) Program that works in partnership with HAI.	
Complete		<ul> <li>Develop dedicated, trained HAI staff with at least one FTE (or contracted equivalent) to oversee HAI activities areas (Integration, Collaboration, and Capacity Building; Reporting, Detection, Response, and Surveillance; Prevention; Evaluation, Oversight, Communication, and Infection Control)</li> </ul>	2008
		As above	
		Other activities or descriptions:	
		3. Integrate laboratory activities with HAI surveillance, prevention, and control efforts.	1995 and Ongoing
$\boxtimes$		i. Improve laboratory capacity to confirm emerging resistance in HAI pathogens and perform typing where	
Complete		appropriate (e.g., outbreak investigation support, HL7 messaging of laboratory results)	
		The ELC program was initiated in 1995 as one of the key activities under CDC's plan to address emerging infectious disease threats. Its purpose is to protect the public health and safety of the American people by enhancing the capacity of public health agencies to effectively detect, respond, prevent and control known and emerging (or re-emerging) infectious diseases. This is accomplished by providing financial and technical resources to:  1. Strengthen epidemiologic capacity;	
		2. Enhance laboratory capacity;	
		3. Improve information systems; and	
		4. Integrate epidemiology, laboratory, and information systems components of public health departments.	
		ELC Logic Model Overview  A logic model provides a graphic depiction of activities and expected outcomes. It helps ensure better alignment between what CDC asks DPH staff to do to build laboratory capacity, what DPH is expected to accomplish, and how DPH activities and achievement of outcomes will be measured. (see Figure 1)	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		Figure 1  Core Areas/Strategies  1. Strengthen Epidemiological Capacity Strategy 1a: Enhance Investigation response and reporting Strategy 1a: Improve surveillance to drive public health action Strategy 1a: Implement and evaluate public health ractice, and prevention and control strategies Strategy 1a: Coordinate and collaborate  2. Enhance Laboratory Capacity Strategy 2a: Sustain and enhance laboratory ciapnostic capacity Strategy 2a: Improve aboratory coordination and outreach/information Systems Strategy 3a: Enhance Health Information Systems Strategy 3a: Sustain and enhance integrated surveillance information systems  ELC Logic Model Overview  Mid-term Outcomes  Long-term/Pistal Outcomes  Improve better prepared to respond or lampore better prepared to respond and control health outcomes Improved surveillance  Better coordination and exchange Implementation  Improved surveillance  Improved surveillance Improved completeness and timeliness of reporting  More timely and efficient efforts: Data used to: Improve H response and control lead to: Improve public health practice and prevention of infectious diseases Improve dompleteness and timeliness of reporting  More timely and efficient efforts: Development and implementation of strong: Development and implementation of	
		Connecticut Laboratory Capacity Project Summaries  Project A: Epidemiology Capacity: Ensure health departments are equipped with staff, surveillance systems and tools to provide rapid response to infectious disease threats.  Project B: Laboratory Capacity: Develop well-equipped public health laboratories, with well-trained staff, that foster communication and appropriate integration between laboratory and epidemiology functions.  Project C: Health Information Systems — Electronic Lab Reporting Capacity: Develop and enhance health information systems infrastructure in public health agencies.  Project D: Laboratory Efficiencies Initiative: Develop and enhance PHL capacity to share testing, training and services acr states  Project F: Cross-Cutting Outbreak Capacity: Provide surge capacity for effective response to outbreak emergency.	oss

Check	Check	Items Planned for Implementation (or currently underway)	Target Dates for
Items Underway	Items Planned		Implementation
- Ciliaci III ay		Project G1: OutbreakNet and NORS: Outbreak Response and Surveillance Activities that support Connecticut statewide	
		epidemiology capacity to respond to foodborne disease outbreaks.	
		Project G2: FoodCORE: Supports Connecticut statewide epidemiology and laboratory capacity to identify foodborne	
		pathogens and monitor for clusters of disease.	
		<b>Project G4: PulseNet:</b> Supports laboratory capacity to identify foodborne pathogens and report them to PulseNet.	
		<b>Project G6: CaliciNet:</b> Capacity for Molecular Identification of Noroviruses supports laboratory capacity for molecular	
		detection of noroviruses.	
		Project G8: NARMS: Supports submission of isolates to the CDC for further testing.	
		<b>Project 11: Prevention Infrastructure:</b> Supports critical health program, epidemiology and infection control staff to support healthcare-associated infection activities.	
		Project 12: Antimicrobial Stewardship: Supports a contractor to promote best practices for AMS in Connecticut.	
		<b>Project I3: Clostridium difficile Infection (CDI) Prevention:</b> Supports a contractor to work with CT long term care facilities on prevention of CDI.	
		<b>Project I5: Data Validation:</b> Supports continued training of the DPH HAI data validation coordinator and additional staff to conduct the project as well as dialysis data validation	
		Project I6: Hemodialysis BSI: Supports a contractor to work to increase infection control capacity.	
		<b>Project J: West Nile Virus and other Arbovirals:</b> Develop and implement effective surveillance, prevention, and control of	
		arboviruses that occur. CT activities include support of a laboratory capacity at the DPH State Laboratory as well as at the	
		Connecticut Agricultural Experiment Station.	
		Project K: Lyme Disease: Assist state and local health departments to develop and implement effective surveillance for	
		diagnosis, prevention, and control of human infections of Lyme Disease supports epidemiology capacity to conduct surveillance for Lyme Disease in CT.	
		<b>Project M: Influenza:</b> Implement enhanced capacity for surveillance and diagnostic testing of respiratory viruses. Provide	
		laboratory and epidemiologic surge capacity necessary for response to a respiratory virus-related emergency.	
		Project M1: Influenza Surveillance and Diagnostic Testing: Supports epidemiology and laboratory capacity for influenza	
		testing, surveillance and analysis in CT.	
		Project M2: Influenza Outbreak Response: Supports additional epidemiology and laboratory capacity in Connecticut to	
		respond to an influenza pandemic.	
		Project N: Non-Influenza Respiratory Viruses: Strengthen laboratory capacity to identify non-influenza respiratory viruses	
		essential for case finding of non-influenza respiratory diseases.	
		Project N2: Non-Influenza Respiratory Diseases: Supports additional epidemiology and laboratory capacity during severe	
		non-influenza respiratory outbreaks.	
		<b>Project U: Tickborne (non-Lyme):</b> Epidemiology, laboratory and/or informatics support for projects designed to improve the	
		detection, investigation, reporting, and response to public health issues related to tickborne diseases.	2014 and

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
- Citaci ii ay		<b>Project W. Advanced Molecular Detection:</b> Supports laboratory capacity to implementing the ability to use advanced techniques to distinguish foodborne outbreak pathogens.	Ongoing
		DPH State Laboratory:  During 2014-2015 the DPH State Laboratory had two microbiologists fully dedicated to molecular subtyping activities including pulsed-field gel electrophoresis (PFGE). In August 2014, staff were reassigned to work on establishing protocols for whole genome sequencing of select enteric isolates  The Connecticut DPH has been able to develop in-house capacity to support informatics needs for electronic data exchange. This capacity includes staff from the Information Technology (IT) and the Infectious Disease (ID) Sections, in particular, the Epidemiology and Emerging Infections Programs Unit. The Informatics Specialist oversees the integrated surveillance system and electronic laboratory reporting activities and staff. The Informatics Unit is based in the Epidemiology and Emerging Infections Program. Connecticut's electronic disease surveillance system is a Consilience Software Maven-based application that supports the majority of reportable disease surveillance. DPH has completed the DPH "Electronic Laboratory Reporting (ELR) System" which is comprised of the PHIN MS secure message transport using the CDC's RnR hub, the PilotFish Technologies PilotFish integration engine and console, and an xml based vocabulary management system built on the BaseX library. The PilotFish integration engine and xml vocabulary management system together comprise the "Messaging Bus" that has been developed to process HL7 2.5.1 ELR messages that are conformant to the national ELR and Meaningful Use (MU) standards as well as non-HL7 messages. The DPH ELR System can be easily modified to support older version HL7 2.3.1 messages if needed.	
		By August 2015, DPH will be in production ELR processing with 4 hospital laboratories using the Cerner Health Sentry System, and the DPH State Laboratory using the ChemWare HORIZON LIMS, for HIV, Hepatitis A, B, and C results. Additional disease testing from these laboratories will be underway, including testing for blood lead. This work is being supported by ID program staff. DPH will have in place an on-boarding process for bringing new laboratories into ELR production. This process will be based on the "ELR Steps and Definitions" document published by the CSTE ELDR subcommittee, and includes a readiness assessment, sharing of documents and processes such as the Connecticut DPH ELR HL7 2.5.1 messaging guide (local guide), the DPH on-boarding protocols, test message validation, vocabulary mapping, acceptance testing and production processes, and data quality validation and monitoring.	
		Other activities or descriptions:	
		4. Improve coordination among government agencies or organizations that share responsibility for assuring or overseeing HAI surveillance, prevention, and control (e.g., State Survey agencies, Communicable Disease Control, state licensing boards)	1995 and Ongoing

Check Items	Check Items	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
Underway	Planned		•
		Emerging Infections Program:	
		Emerging Infections Program:  The ID Section also contains the Connecticut Epidemiology and Emerging Infections (EIP) Program. Since 1995 the mission of the Epidemiology and Emerging Infections Program is to prevent illness, disability, and death in Connecticut residents caused by infectious diseases. The Epidemiology and Emerging Infections Program accomplishes its mission by conducting surveillance for more than 30 infectious diseases of uplic health importance, investigating disease outbreaks, epidemiologic studies of emerging infectious diseases, training, and public education programs to develop, evaluate, and promote prevention and control strategies for infectious diseases. The EIP has sites in ten states, and works in partnership with medical practitioners, local and federal public health officials, other state agencies, medical and public health professional associations, infectious disease experts from academic and clinical practice, and public service organizations. (Figure 2 and 3)  Figure 2. The EIP Network has sites in 10 states:	
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Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		Figure 3. Connecticut Epidemiology and Emerging Infections (EIP) Program Structure:  Dept. of Public Health Laboratory  CT Dept. of Public Health Matthew Cartter, EIP P.I.  Departments	
		Leadership Group Cartter Heimer Meek Eaton Rabatsky-Ehr  ABCs Petit Hurd Phan FloodNet Heimer Response Cartter Response Cartter Heimer Niccolai Sosa Nelson Vasquez  Yale School of Public Health Robert Heimer, EIP Co-P.I.	
		Active Bacterial Core surveillance (ABCs): Active population-based laboratory surveillance for invasive bacterial disease. Pathogens included: groups A and B streptococcus, Haemophilus influenzae, Neisseria meningitidis, Streptococcus pneumoniae, and methicillin-resistant Staphylococcus aureus.  FoodNet: Active population-based laboratory surveillance to monitor the incidence of foodborne diseases. Surveillance is conducted for seven bacterial and two parasitic pathogens: E. coli O157:H7, Campylobacter, Listeria, Salmonella, Shigella, Yersinia, Vibrio, Cryptosporidium, and Cyclospora.  Influenza activities: Active population-based surveillance for laboratory confirmed influenza-related hospitalizations. EIP sites also conduct influenza vaccine effectiveness evaluations among groups for which ACIP recommends annual vaccination.  Healthcare Associated Infections-Community Interface (HAIC) projects: Active population-based surveillance for Clostridium difficile infection and other healthcare associated infections caused by pathogens such as MRSA, Candida, and multi-drug resistant gram-negative bacteria.  Surveillance efforts of these core EIP activities generate reliable estimates of the incidence of certain infections and provide the foundation for a variety of epidemiologic studies to explore risk factors, spectrum of disease, and prevention strategies.	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
Onderway	Tidilica	State Licensing Boards:	
		DPH Health Care Systems (HCS) Branch is the Connecticut survey agency for CMS, and has participated with the HAI Advisory Committee since its inception in 2006, this includes putting the state HAI program on the agenda of the state licensure boards (medical, nursing), convened by DPH, on a regular basis.	
		Quality Collaboratives:	
		Qualidigm, (the state quality improvement organization) is a national healthcare consulting and research company that provides innovative and scientifically-based solutions to transform and improve care delivery and patient outcomes. In 2014 Qualidigm was contracted to conduct two state collaboratives:	2006 and Ongoing
		The Antimicrobial Stewardship (AMS) prevention collaborative DPH contracted with Qualidigm, Inc. to establish an	
		antimicrobial stewardship prevention collaborative that performed an environmental scan, enrolled five "Communities of Care" which included acute care hospitals, and collecting six months of baseline MDRO data using the National Healthcare Safety Network (NHSN) LabID module as a benchmark.	2014 and Ongoing
		The Clostridium difficile (CDI) prevention collaborative worked with 33 participating long term care facilities to reduce CDI and multi-drug resistant infections using rapid cycle quality improvement techniques. The collaborative had a kick off meeting in April 2014 that discussed CDI pathophysiology, treatment and prevention. Participants also developed AIM statements and completed self-assessment tools utilizing APIC standards. Participants identified targeted areas for improvement such as hand hygiene, early diagnosis, rapid containment and environmental cleaning.	
		Other activities or descriptions:	
		5. Facilitate use of standards-based formats (e.g., Clinical Document Architecture, electronic messages) by healthcare facilities for purposes of electronic reporting of HAI data. Providing technical assistance or other incentives for implementations of standards-based reporting can help develop capacity for HAI surveillance and other types of public health surveillance, such as for conditions deemed reportable to state and local health agencies using electronic laboratory reporting (ELR). Facilitating use of standards-based solutions for external reporting also can strengthen relationships	2013 and Ongoing

Check	Check	Items Planned for Implementation (or currently underway)	Target Dates for
Items	Items		Implementation
Underway	Planned		
		between healthcare facilities and regional nodes of healthcare information, such as Regional Health Information	
		Organizations. (RHIOs) and Health Information Exchanges (HIEs). These relationships, in turn, can yield broader benefits	
		for public health by consolidating electronic reporting through regional nodes.	
		Before the expansion of HAI reporting in Connecticut in 2012, hospitals were reluctant to use the CDA of HLA massaging in NHSN as it was a burden to set up and required much data entry for very narrow disease reporting. Since disease reporting has expanded, there was a greater interest in using the CDA beginning in 2013. The HAI program has distributed guidance and has consulted with hospital IT departments and IPs to facilitate development of this capacity. Currently 12 hospitals are using CDA.	
		Other activities or descriptions:	

#### 2. Surveillance, Detection, Reporting, and Response

Timely and accurate monitoring remains necessary to gauge progress towards HAI elimination. Public health surveillance has been defined as the ongoing, systematic collection, analysis, and interpretation of data essential to the planning, implementation, and evaluation of public health practice, and timely dissemination to those responsible for prevention and control.<sup>2</sup> Increased participation in systems such as the National Healthcare Safety Network (NHSN) has been demonstrated to promote HAI reduction. This, combined with improvements to simplify and enhance data collection, and improve dissemination of results to healthcare providers and the public are essential steps toward increasing HAI prevention capacity.

The HHS Action Plan identifies targets and metrics for five categories of HAIs and identified Ventilator-associated Pneumonia as an HAI under development for metrics and targets (Appendix 1):

- 6. Central Line-associated Blood Stream Infections (CLABSI)
- 7. Clostridium difficile Infections (CDI)
- 8. Catheter-associated Urinary Tract Infections (CAUTI)
- 9. Methicillin-resistant Staphylococcus aureus (MRSA) Infections
- 10. Surgical Site Infections (SSI)
- 11. Ventilator-associated Pneumonia (VAP)

State capacity for investigating and responding to outbreaks and emerging infections among patients and healthcare providers is central to HAI prevention. Investigation of outbreaks helps identify preventable causes of infections including issues with the improper use or handling of medical devices; contamination of medical products; and unsafe clinical practices.

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<sup>&</sup>lt;sup>2</sup> Thacker SB, Berkelman RL. Public health surveillance in the United States. Epidemiol Rev 1988;10:164-90.

 Table 2: State planning for surveillance, detection, reporting, and response for HAIs

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		1. Improve HAI outbreak detection and investigation	1995 and ongoing
		<ul> <li>i. Work with partners including CSTE, CDC, state legislatures, and providers across the healthcare continuum to improve outbreak reporting to state health departments</li> </ul>	
		ii. Establish protocols and provide training for health department staff to investigate outbreaks, clusters, or unusual cases of HAIs.	
		iii. Develop mechanisms to protect facility/provider/patient identity when investigating incidents	
		and potential outbreaks during the initial evaluation phase, where possible, to promote reporting of outbreaks	
		iv. Improve overall use of surveillance data to identify and prevent HAI outbreaks or transmission	
		in HC settings (e.g., hepatitis B, hepatitis C, multi-drug resistant organisms (MDRO), and other reportable HAIs)	
		Other activities or descriptions:	
		As above:	1995 and ongoing
		i. Connecticut has a reportable disease list that is developed by the State Epidemiologist and	
		authorized by the legislature annually. The list is capable of identifying diseases of interest, and can indicate an outbreak by the numbers reported.	
		ii. DPH received federal funding and can hire and train additional HAI staff. As part of their duties	
		these staff will assist in developing training for HAI outbreak response and HAI prevention.	
		iii. Connecticut has a stringent public health confidentiality law, 19a-25, which protects the identity	
		of patients, providers, and facilities during any investigation of infectious diseases (whether in healthcare facilities or community settings).	
		iv. The Epidemiology and Emerging Infections Program conducts surveillance for more than 30	
		infectious diseases of public health importance, investigating disease outbreaks, epidemiologic	
		studies of emerging infectious diseases, training, and public education programs to develop,	
		evaluate, and promote prevention and control strategies for infectious diseases. This program	
		works in partnership with medical practitioners, local and federal public health officials, other	
		state agencies, medical and public health professional associations, infectious disease experts	
		from academic and clinical practice, and public service organizations.	
		v. A sole source contract was executed between IPRO Network of New England and DPH in early  December 2014. The project director, Shelli Eason, a nurse with extensive project management	December 2014 and ongoing

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		and evaluation experience was hired. On March 10 forty-eight nurse trainers from 44 centers (100%) attended the opening day-long conference to inform these key personnel on the benefit of using NHSN to guide training and to evaluate the degree facilities are implementing best practices, policy changes, training, and evaluation with feedback. Information on NHSN included how NHSN data relates to ESRD CMS coverage; the CDC collaborative project; the interventions that led to tangible change in centers; the importance of NHSN data for QI review and intervention; the implications of NHSN data being shared on the CT DPH public website; and how surveys provide opportunities to verify patient safety in facilities. The training also covered infection control practices, including commonly seen infections. Recommended infection control practices were reviewed, including CDC tools and guidance for infection control in the dialysis setting. In year 2, DPH will renew its contract with IPRO. The contract requires that CDC recommendations, tools, and materials are the primary source for all assessment, training and technical assistance. IPRO will continue monthly conference calls of its Advisory Committee, made up of representatives from Large Dialysis Organizations, independent centers, hospitals, DPH HAI program staff, Mass DPH, Qualidigm (the Connecticut QIO), and infection preventionists vi. The new ELC 2015 Ebola funding will add nurse consultant/IPs and data analysis staff to support the development of a state infection control (IC) assessment program. The staff will provide increased data collection, management, analysis, reporting, training and technical assistance that	2015
		<ul> <li>will lead to identifying and mitigating gaps in infection control practices and outbreak reporting.</li> <li>2. Enhance laboratory capacity for state and local detection and response to new and emerging HAI issues.</li> <li>CRE: <ul> <li>Managers at all 18 clinical laboratories were trained on the Connecticut CRE surveillance protocol and case definition in 2014. Connecticut has been receiving reports from 18 clinical laboratories and has entered the data into a database. Laboratory audits have not yet been completed to validate the data. In 2014, 117 isolates were reported and resistance profiles were received on 68% of these. All had appropriate data on clinical site (all were active infections as we are not receiving reports on tests for colonization, e.g., stool samples). Clinical follow-up is not routine yet. We have received reports of one novel resistance mechanism isolate, confirmed by CDC.</li> </ul> </li> <li>Other activities or descriptions:</li> </ul>	2014 and ongoing
		3. Improve communication of HAI outbreaks and infection control breaches	See EIP Program

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		<ul> <li>i. Develop standard reporting criteria including, number, size, and type of HAI outbreak for health departments and CDC</li> <li>ii. Establish mechanisms or protocols for exchanging information about outbreaks or breaches among state and local governmental partners (e.g., State Survey agencies, Communicable Disease Control, state licensing boards)</li> <li>The new ELC 2015 Ebola funding will add nurse consultant/IPs and data analysis staff to support the development of a state infection control (IC) assessment program. The staff will provide increased data collection, management, analysis, reporting, training and technical assistance that will lead to identifying and mitigating gaps in infection control practices and outbreak reporting.</li> </ul>	above, page 10-12  2015 and ongoing
		4. Identify at least 2 priority prevention targets for surveillance in support of the HHS HAI Action Plan:  I. Central line-associated bloodstream infections (CLABSI)  II. Catheter—associated urinary tract infections (CAUTI)  III. Surgical site infections (SSI)  IV. Clostridium difficile infections (CDI)  V. Methicillin-resistant Staphylococcus aureus (MRSA) infections  VI. Ventilator-associated Pneumonia (VAP)	Completed I.2009 II.2012 III.2012 IV.2013 V.2013 VI. Not yet mandated by CMS

Check	Check	Items Planned for Implementation (or currently underway)	Target Dates for
Items	Items		Implementation
<b>Jnderway</b>	Planned	Con manufacture	
		See results below	
		The following tables provide data on all mandated HAI reporting requirements to CMS via NHSN.	
		Infections are reported using the Standard Infection Ratio (SIR).	
		Figure 4. What does the standardized infection ratio number mean?	
		If the SIR is more than 1:	
		There was an increase in the number of infections reported in the nation compared to the national	
		baseline.	
		A high SIR usually reflects a need for stronger HAI prevention efforts. Other factors may also play a	
		role in a high SIR, such as intense data validation activities that lead to the discovery and reporting	
		of more infections than in previous years.	
		Man.	
		If the SIR is 1:	
		There were about the same number of infections reported in the nation compared to the national	
		baseline.	
		·	
		If the SIR is less than 1:	
		There was a decrease in the number of infections reported in the nation compared to the national	
		baseline. Usually, a low SIR reflects the results of robust HAI prevention strategies.	
		baseline. Osaany, a low six reflects the results of robust firm prevention strategies.	

Check Items Underway	Check Items Planned	Items Planne	ed for Implem	entation (or c	urrently underv	vay)			Target Dates for Implementation
			2009 and ongoing						
		Acute Care I	CUs		Data Jan 2009 t				
		Summary Year	CLABSI Count	Number Expected	Catheter Line Days	CT SIR	SIR 95% Confidence Intervals	Interpretation	
		2009	109	120	56707	0.905	0.747, 1.087	4	
		2010	84	118	54726	0.713	0.572, 0.878	4	
		2011	71	128	59237	0.554	0.436, 0.695	4	
		2012	138	225	105191	0.613	0.517, 0.721	14	
		2013	99	212	99477	0.466	0.381, 0.565	1	
		2014	84	200	94607	0.420	0.337, 0.518		
		their composi	and 2014 Co te SIR from 0. rget. Howeve	nnecticut acut 90 to 0.42 for	CLABSIs. By 201	13 Connectio	e units (ICUs) were ut had reached the ximately 57,000 to	2013 national	

Check Items Underway	Check Items Planned	Items Plann	ed for Imp	lementation	or current	ly under	way)			Target Dates for Implementation	
Onaciway	1 Idillica			Table 2. C	T Acute Car	e Hospit	al Pediatri	ic ICU CLABSI			
					Rep	orts to I	NHSN			2009 and ongoing	
					-		009-2014				
		Pediatric IC CLABSI	Pediatric ICUs								
		Summary Year	CLABSI Count	Number Expected	Catheter Line Days	CT SIR	SIR p-value	SIR 95% Confidence Intervals	Interpretation		
		2009	11	12	4137	0.886	0.4155	0.442, 1.586	4	1	
		2010	9	11	3793	0.791	0.3008	0.362, 1.501		1	
		2011 10 12 3835 0.869 0.4012 0.417, 1.598									
		2012	6	10	3416	0.585	0.1153	0.215, 1.274	-		
		2013	5	10	3367	0.495	0.0908	0.181, 1.097	4		
		2014	5	10	3484	0.478	0.0736	0.175-1.060		1	
		Summary of Between 200 from 0.89 to	9 and 2014		t pediatric h	nospitals	ICUs were	able to reduce thei	r composite SIR		

Check	Check	Items Plan	ned for Im	plementat	ion (or cur	rently un	ns Planned for Implementation (or currently underway)										
Items	Items									Implementation							
derway	Planned																
				Table 3	. CT Acute	Care Hos	pital Neo	natal ICU CLABSI									
						Reports	to NHSN			2012 and ongoing							
					Sum	mary Dat	ta 2012-20	014									
		NICU								7							
		CLABSI															
		Summary Year	CLABSI Count	Number Expected	Catheter Line Days	CT SIR	SIR p-value	SIR 95%Confidence Intervals	Interpretation								
		2012	12	30	12816	0.4	0.0002	0.207, 0.699	16								
		2013	12	30		0.4	0.0003	0.219, 0.688	1								
			.4 Connect	icut neona			_	o.160, 0.656	national baseline.								
		Summary o	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								
		<b>Summary o</b> In 2012-201	f Results 2 .4 Connect	<b>012-2014:</b> icut neona	tal ICUs CLA	ABSI SIRs	were signi	ficantly below the i	national baseline.								

Check Items	Check Items	Items Plan	ned for Im	plementat	ion (or cur	rently und	derway)			Target Dates for Implementation
Underway	Planned									implementation
				Ta	able 4. CT	Acute Care	e Hospital	ICU CAUTI		2012 and ongoing
						ts to NHSI	-			
					Sum	mary Data	a 2012 - 2	014		
		Acute								
		Care ICU								
		CAUTIS Summary CAUTI Number Catheter CT SIR SIR SIR%95 Interpretation								
		Summary Year	CAUTI Count	Number Expected	Catheter Days		SIR p-value	Confidence Intervals	Interpretation	
		2012	510 27	272	120232	1.872	0.000	1.715, 2.040	1	
		2013	423	254	112819	1.663	0.000	1.510, 1.828	19	
		2014	396	235	104955	1.683	0.000	1.523, 1.855	1	
		Summary of Since 2012 hospitals h measures,								

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		Figure 5. Surgical Site Infections Definitions of three SIR Models  There are three SSI SIR models available from NHSN, each briefly described in the table below.  All SSI SIR  Includes Superficial, Deep & Organ/Space SSIs	
		Model  Superficial & Deep incisional SSIs limited to primary incisional SSIs only Includes SSIs identified on admission, readmission & via post-discharge surveillance	
		Includes only Deep incisional primary SSIs & Organ/Space SSIs      Includes only SSIs identified on Admission/Readmission to facility where procedure was performed     Includes only inpatient procedures     Used for the HAI Progress Report, published annually by CDC	
		<ul> <li>Includes only in-plan, inpatient COLO and HYST procedures in adult patients (i.e., ≥ 18 years of age)</li> <li>Includes only deep incisional primary SSIs and organ/space SSIs with an event date within 30 days of the procedure</li> <li>Uses only age and ASA to determine risk</li> <li>Used only for CMS IPPS reporting and for public reporting on Hospital Compare</li> </ul>	

Check Items Underway	Check Items Planned	Items Pla	nned for Im <sub>l</sub>	olementatio	n (or curre	ently unde	rway)				Target Dates for Implementation
Sincerway	Tiumeu	ALL SSI Co Includes superficia	OLO al, deep	Table 5. CT /	lodel by Pi	•	-				2012 and ongoing
		Summary Year	Procedure Code	Procedure Count	All SSI Model Infection Count	All SSI Model Number Expected	All SSI Model SIR	SIR p-value	All SSI Model 95% Confidence Interval	Interpretation	
		2012	COLO	3612	324	201	1.611	0.0000	1.443, 1.794	140	
		2013	COLO	3652	309	202	1.526	0.0000	1.363, 1.703	19	
		2014	COLO	3746	311	212	1.470	0.0000	1.313, 1.640	1	
		Overall rate	of Results 20 es for "All SS enchmarks fo	I SIR Model		s have rem	ained ur	nchanged a	and statistically a	above the	

Check Items Underway	Check Items Planned	Items Plan	ned for Impl	ementation	(or current	ly underwa	y)				Target Dates for Implementation	
			2012 and ongoing									
		ALL SSI HY Includes superficial and organ	, deep									
		Summary Year	Procedure Code	Procedure Count	All SSI Model Infection Count	All SSI Model Number Expected	All SSI Model SIR	SIR p- value	All SSI Model 95% Confidence Interval	Interpretation		
		2012	Abd. HYST	3955	96	74	1.29	0.02	1.051, 1.569	1		
		2013	Abd. HYST	4040	105	76	1.38	0.00	1.136, 1.666	19		
		2014	Abd. HYST	3979	91	75	1.20	0.08	0.976, 1.473	-1		
		Overall rate							ged and statis	cically above the		

Check Items	Check Items Planned	Items Pla	nned for Im	nplementat	ion (or currently	underway)					Target Dates for Implementation
Underway	Planneu			Table 7. Cī	FAcute Care Hosp For CMS IPF Summary I	S: Colon Su	ırgery	y SSI Mod	del*		2012 and ongoing
		-	30 day SSIs d Organ Spa								
		Summary Year	Procedure	Procedure count	Number of infections(comple x 30 day)	Number of expected infections	CT SIR	SIR p-value	SIR 95% Confidenc e Intervals	Interpretation	
		2012	COLO	3591	141	109	1.295	0.004	1.094, 1.522	14	
		2013	COLO.	3639	141	111	1.270	0.007	1.073, 1.493	19	
		2014	COLO.	3714	171	115	1.489	0.000	1.278, 1.725	19	
		Pt ≥ 18 yrs Incisional S Complex 3 website Summary	Secondary S 0 Day SSI M of Results 2	ent date with SIs, and on lodel is use	thin 30 days of proly includes those different	procedures ng and for	/SSIs wit public rep	h primary porting o	v closure te n Hospital (	chniques.	

Check Items	Check Items	Items Pla	nned for Im	plementati	ion (or currently	underway)	)				Target Dates for Implementation	
Underway	Planned											
		Complex	. 30 day SSIs	For C	Acute Care Hos MS IPPS: Abdom Summary	inal Hyster	ectomy	-	odel*		2012 and ongoing	
		·	Complex 30 day SSIs  Deep and Organ Space									
		Abd. Hys	•									
		Summary Year	Procedure	Procedure count	Number of infections (complex 30 day)	Number of expected infections	CT SIR	SIR p-value	SIR 95% Confidence Intervals	Interpretation		
		2012	Abd. Hyst.	3950	47	33	1.417	0.023	1.053, 1.868	1		
		2013	Abd. Hyst.	4039	59	34	1.713	0.000	1.316, 2.194	1		
		2014	Abd. Hyst.	3976	37	35	1.058	0.716	0.756, 1.443	-1		
		Pt ≥ 18 yrs Incisional Complex 3 website. Summary These rat	Secondary S 30 Day SSI M of Results 2	ent date wit SIs, and onl lodel is used 1012-2014: ained unch	hin 30 days of pr y includes those I for CMS reporti anged and statist	procedures ing and for	s/SSIs wi <sup>.</sup> public re	th primar porting o	ry closure te on Hospital (	chniques. Compare		

Check Items	Check Items	Items Pla	Target Dates for Implementation							
Inderway	Planned	nned								
			2013 and ongoing							
					MRSA Bloc	2013-2014		,		
					IVINSA BIOC	ou racwiu	eliv LabiL	,		
		Hospital	Number	Number	Number of pt.	SIR	SIR	SIR 95%	Interpretation	
		Onset	of HO	predicted HO	days		p-value			
		(HO) MRSA	MRSA	MRSA				Confidence Intervals		
		2013	110	148	1,993,507	0.744	0.0014	0.614, 0.893	1	
		2013	110	1-10	1,333,307	0.744	0.0014	0.014, 0.033		
		2014	87	135	1,969,098	0.645	0.0000	0.520, 0.792		
		Definition:								
			llin-resistant Sta	phylococcus aureus						
					ita relevant to MRSA I	olood LabID rep	orting AND sep	arately for outpatient em	ergency department, and 24-	
		hour observatio	n location(s).							
		Healthcare Facil	ity-Onset (HO): I	abID Event collected	d >3 days after admiss	ion to the facili	ty (i.e., on or af	ter day 4).		
		MDRO Bloodstr	eam Infection In	cidence Rate = Numl	per of all unique blood	d source LabID I	Events per patie	nt per month identified >	3 days after admission to the	
						verall facility-v	vide inpatient=F	acWideIN) / Number of p	atient admissions to the	
		location or facili	ity x 100 (will be	removed from NHSN	l analysis in July					
		Summary	of Results	2013-2014:						
		•			te care hospita	als were a	ble to mai	ntain a statically	significant reduction	
					red to other h			•		
		Hoopital	acquirea i	viito, t compai	ed to other m	oopitalo i c	por till 8 to			

Check Items Underway	Check Items Planned	Items Pla	Items Planned for Implementation (or currently underway)							Target Dates for Implementation
			2013 and ongoing							
		Hospital Onset (HO) C. diff.	Number of HO C. diff.	Number predicted HO C. diff.	Number of pt. days	SIR	SIR p-value	SIR 95%  Confidence Intervals	Interpretation	
		2013	1475	1442	1,829,590	1.023	0.3942	0.972, 1.076		
		2014	1515	1406	1,828,577	1.077	0.0044	1.024, 1.132	19	
		month in Overall Fa Summary In 2014 Co	the facility cility of Results onnecticut	/ Number of Inpatient re 2013-2014:	patient days for porting) nospital rates o	or the faci	lity x 10,00	00 (this calculation	OI LabID Events per on is only accurate for tically worse than	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		Figure 6. CT Acute Care Hospital Healthcare Personnel Influenza Vaccination Compliance by Type Influenza Season 2013-2014	2013 and ongoing
		30 96.7 87.8 87.6 Total HCP (including contract workers) 60 Licensed Independent Practitioners (LIPs) Employees  Summary of Results During the 2013-2014 influenza season, 86% (range 67%-100%) of all HCP in acute care hospitals were vaccinated. Slight improvement from the 2012-2013 influenza season, 83% HCP vaccinated (range 30%-99%).	

Check	Check	Items Pla	nned for	Implemen	tation (or	currently	underway)	)		Target Dates for
Items Jnderway	Items Planned									Implementation
Jiidei Way	· iaiiica		2014 and ongoing							
					_		CH PPS-Ov	'ACH) CLABSI Reports erall		
		LTACH CLABSI								
		Summary Year	CLABSI Count	Number Expected	Catheter Line Days	CT SIR	SIR p-value	SIR 95%Confidence Intervals	Interpretation	
		2014	12	18	17111	0.673	0.1565	0.364, 1.143	-10	
		•	<b>of Result</b> : ABSIs in (		were com	parable to	national ba	aseline.		
		•	ABSIs in (	CT LATACs	ng Term Ac	cute Care		「ACH) CAUTI Reports	to NHSN	2014 and ongoing
		•	ABSIs in (	CT LATACs	ng Term Ac	cute Care	Hospital (L	FACH) CAUTI Reports erall	to NHSN	2014 and ongoing
		•	ABSIs in (	CT LATACs	ng Term Ac	cute Care	Hospital (L <sup>1</sup> CH PPS-Ov	FACH) CAUTI Reports erall	to NHSN	2014 and ongoing
		In 2014 CL	ABSIs in (	CT LATACs	ng Term Ac	cute Care	Hospital (L <sup>1</sup> CH PPS-Ov	FACH) CAUTI Reports erall	to NHSN  Interpretation	2014 and ongoing
		LTACH CAUTI Summary	ABSIs in (	12. CT Lor	ng Term Ad	cute Care   for CMS L1 Summa	Hospital (L <sup>T</sup> CH PPS-Ov ry Data 202	FACH) CAUTI Reports rerall L4 SIR 95%Confidence		2014 and ongoing
		LTACH CAUTI Summary Year	Table  CAUTI Count  46	Number Expected	ng Term Ad	cute Care for CMS LT Summa CT SIR	Hospital (LTCH PPS-Ov ry Data 20: SIR p-value	FACH) CAUTI Reports rerall 14 SIR 95%Confidence Intervals		2014 and ongoing
		LTACH CAUTI Summary Year 2014 Summary	Table  CAUTI Count  46  of Result:	Number Expected 32	Catheter Days	cute Care for CMS LT Summa  CT SIR  1.438	Hospital (L'CH PPS-Ov ry Data 20: SIR p-value 0.0191	FACH) CAUTI Reports rerall 14 SIR 95%Confidence Intervals		2014 and ongoing
		LTACH CAUTI Summary Year 2014 Summary	Table  CAUTI Count  46  of Result:	Number Expected 32	Catheter Days	cute Care for CMS LT Summa  CT SIR  1.438	Hospital (L'CH PPS-Ov ry Data 20: SIR p-value 0.0191	SIR 95%Confidence Intervals		2014 and ongoing

Check	Check	Items Pla	Target Dates for													
Items	Items															Implementation
nderway	Planned															
			2014 and ongoin													
					A	dult and	Pediatric \	Wards								
					(	CAUTI Re	ports to N	HSN								
						for CMS	S PPS-Over	all								
						Summa	ry Data 20	14								
		IRF														
		CAUTI														
		Summary	CAUTI	Number	Catheter	CT SIR	SIR	SIR 95%Confidence	Interpretation							
		Year	Count	Expected	Days		p-value	Intervals								
		2014	11	9	2571	1.253	0.4445	0.659,2.170	4							
									=							
		Summary In 2014 CA			comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								
					comparabl	le to natio	onal baseli	ne.								

Check Items Underway	Check Items Planned	Items Planne	Target Dates for Implementation						
Officer way	riamieu	Hemo.	Table 14	Rate	sis Access-Relate es per 100 Patie Report to NH cess (fistula, gra ımmary Data 20	nt Months ISN ft, catheter)	m Infection (BS	ı)	2013 and ongoing
		Access- related BSI							
		Summary year	Access Type	Total # BSI	Total # (count) of Patient Months	BSI rate /100 patient- months	NHSN pooled mean BSI rate/100 patient months	Interpretation	
		2013	All	279	36374	0.767	1.27	16	
		2014	All	316	37127	0.851	1.27		
		-	ficant reduc		onnecticut hemo elated bloodstre	•			
		5. Adopt nat	tional stand	ards for data an	d technology to	track HAIs (e	.g., NHSN).		Completed
$\boxtimes$			elop metric	-	ogress towards	national goal	s (align with tar	geted state	2009
		_			nts for prevention	n targets			
	_	Other activitie							

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		6. Develop state surveillance training competencies         i. Conduct local training for appropriate use of surveillance systems (e.g., NHSN) including facility and group enrollment, data collection, management, and analysis Since 2008, the DPH HAI program has conducted annual NHSN training programs. As of 2015, training has continued for Acute Care Hospitals, Long Term Acute Care Hospitals, Inpatient Rehabilitation Facility staff, and ambulatory surgical centers on the NHSN in preparation for new Centers for Medicaid and Medicare Services (CMS) reporting requirement from those facilities.	Initiated 2007 and ongoing as new NHSN modules are added
		Other activities or descriptions	
		7. Develop tailored reports of data analyses for state or region prepared by state personnel Since 2008 DPH HAI Program is mandated, based on state statute, to provide an annual report on program activities. The report includes data analysis of each hospital in the state.	Initiated 2008
		8. Validate data entered into HAI surveillance (e.g., through healthcare records review, parallel database comparison) to measure accuracy and reliability of HAI data collection  Data validation – DPH completed CLABSI validation process for CLABSI numerators and denominators in 2010 and 2014 respectively	Completed and Published 2010, 2014 Ongoing for each new NHSN module
		<ol> <li>Develop a validation plan</li> <li>Pilot test validation methods in a sample of healthcare facilities</li> <li>Modify validation plan and methods in accordance with findings from pilot project</li> <li>Implement validation plan and methods in all healthcare facilities participating in HAI surveillance</li> <li>Analyze and report validation findings</li> <li>Use validation findings to provide operational guidance for healthcare facilities that targets</li> </ol>	
		any data shortcomings detected  Other activities or descriptions:  Publications:  1. Backman LA, Melchreit R, Rodriguez R. Validation of the surveillance and reporting of central line-associated bloodstream infection data to a state health department. Am J Infect Control.	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		<ul> <li>2010 Dec;38(10):832-8. doi: 10.1016/j.ajic.2010.05.016.</li> <li>2. Backman LA, Nobert G, Melchreit R, Fekieta R, Dembry LM. Validation of the surveillance and reporting of central line-associated bloodstream infection denominator data. Am J Infect Control. 2014 Jan;42(1):28-33. doi: 10.1016/j.ajic.2013.06.014. Epub 2013 Oct 29.</li> </ul>	
		<ul> <li>9. Develop preparedness plans for improved response to HAI         <ul> <li>i. Define processes and tiered response criteria to handle increased reports of serious infection control breaches (e.g., syringe reuse), suspect cases/clusters, and outbreaks</li> </ul> </li> <li>Surveillance         <ul> <li>i. The state HAI Advisory Committee recommended that public HAI reporting in Connecticut continues to mirror the scope of CMS reporting. The Reportable Conditions Committee concurred and therefore recommended that CLABSI and CAUTI from ACH and LTACs, CAUTI from IRFs, and the MRSA and c. difficile LabID from acute care hospitals be reported along with colon and hysterectomy SSIs. Hemodialysis centers also report NHSN "dialysis events."</li></ul></li></ul>	Activities from 2009 - 2015

Check	Check	Items Pla	anned for Implementation (or currently underway)	Target Dates for
Items	Items			Implementation
Underway	Planned		Control of the contro	
			included acute care hospitals into the collaborative, and collected six months of baseline	
			MDRO and CDI data using the National Healthcare Safety Network (NHSN) LabID module as a	
			benchmark.	
		ii.	DPH received a Public Health Foundation Future of Public Health Award to incorporate	
			quality Improvement methods into public health and medical practice. The DPH led a	
			Clostridium difficile prevention collaborative with 25 long term care facilities.	
		iii.	DPH supported the End Stage Renal Disease Network of New England in its development of	
			the first NHSN dialysis validation protocol in the nation.	
		Training		
		j.	In 2011, Federal stimulus act (ARRA)-funded APIC training in infection control (EPI 101 and	
			201).	
		Policy and	d Leadership	
		i.	The Multi-Disciplinary HAI Advisory Committee focused on policy changes affecting	
			healthcare associated infections, education campaigns such as promoting hand hygiene; and	
			the evaluation of hospital resources associated with healthcare associated infection	
			prevention.	
		ii.	The Technical Advisory Group (TAG) provided technical advice to the DPH Healthcare	
			Associated Infections Program for HAI surveillance including advice on medical care,	
			epidemiology, statistics, infectious diseases etc.	
		iii.	Public Reporting Group is a committee charged with making recommendations to the	
			Connecticut Department of Health on the public reporting of healthcare-associated	
			infections.	
		iv.	The Education Sub-Committee provides education and training about healthcare associated	
			infections & prevention of healthcare associated infections to applicable persons and	
			healthcare disciplines.	
		V.	Connecticut Leadership for Quality Healthcare Coalition is a multi-disciplinary and multi-	
			stakeholder advisory committee established by Qualidigm as a component of its CMS 10 <sup>th</sup>	
			Scope of Work. HAIs are one of the focus areas for this group, and the DPH HAI Coordinator	
			is one of several DPH staff participating on the group with Commissioner Mullen in 2010.	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		<ul> <li>Outbreaks         <ol> <li>In 2012, Connecticut was heavily involved in the surveillance and response to the Multistate                 Fungal Meningitis Outbreak as one of the 27 states receiving shipment of the contaminated lots                 of methyl prednisolone from the New England Compounding Center. We have collaborated with                 the state Department of Consumer Protection (which regulates pharmacies in Connecticut) on                 communications with both the medical and pharmacy communities about patient notification                 and monitoring.</li> </ol> </li> </ul>	
		Other activities or descriptions:  With the award of the 2015 Ebola Grant the DPH will enhance outbreak response and reporting	Beginning June 2015 and ongoing
		<ul> <li>DPH HAI Program will: <ol> <li>Identify 2012-2015 NHSN HAI rates and HAI outbreaks by individual acute care hospitals (ACH);</li> <li>Identify 2012-2015 outbreak reports to DPH from Reportable Disease databases;</li> <li>Query individual ACH facility for facility specific outbreak identification and response activities;</li> <li>Correlate facility specific NHSN rates with facility specific outbreak/prevention activities;</li> <li>Request participation and attend IC Committee meetings in each of the 29 ACH,s beginning with ACHs with HAI rates above baseline;</li> <li>Present NHSN data, identified HAI reporting gaps, outbreak/prevention activities (individual and statewide), and examples of successes, for discussion and problem solving at IC meetings</li> <li>Offer education, consultation, technical assistance, staff trainings, collaboration with state prevention partners, and updates at future IC meetings;</li> <li>Prepare progress reports summarizing DPH HAI program activities for state HAI Advisory group.</li> </ol> </li> </ul>	
		10. Collaborate with professional licensing organizations to identify and investigate complaints related to provider infection control practice in non-hospital settings and set standards for continuing education and training	July 2015 and ongoing
		DPH is working in collaboration with CT state licensing agency by reviewing all ACH licensing agency visitations (either as state or CMS reviewers) for infection control violations or recommendations. Nurse	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		consultants will be categorizing infection control gaps, and work with stakeholders on improvements.	
		Other activities or descriptions:	
		11. Adopt integration and interoperability standards for HAI information systems and data sources	Complete
		<ul> <li>i. Improve overall use of surveillance data to identify and prevent HAI outbreaks or transmission in HC settings (e.g., hepatitis B, hepatitis C, multi-drug resistant organisms (MDRO), and other reportable HAIs) across the spectrum of inpatient and outpatient healthcare settings</li> <li>See Emerging Infections Program description of projects, pg. 12</li> </ul>	
		<ul> <li>ii. Promote definitional alignment and data element standardization needed to link HAI data across the nation.</li> <li>NHSN standard definitions</li> </ul>	
		Other activities or descriptions:	
$\boxtimes$		<ul> <li>12. Enhance electronic reporting and information technology for healthcare facilities to reduce reporting burden and increase timeliness, efficiency, comprehensiveness, and reliability of the data         <ul> <li>i. Report HAI data to the public</li> </ul> </li> <li>NHSN data is reported to CMS and healthcare facilities throughout the state. It is reportable to DPH based on state statute. Reports are also available to the public on the DPH website, and Medicare</li> </ul>	Complete
		Hospital Compare website	
		Other activities or descriptions:	
		<ul><li>13. Make available risk-adjusted HAI data that enable state agencies to make comparisons between hospitals.</li><li>Using NHSN SIR for comparison between hospitals</li></ul>	Complete
		Other activities or descriptions:	
		14. Enhance surveillance and detection of HAIs in nonhospital settings	Initiated 2012 and ongoing
		CT DPH surveillance is aligned with CMS reporting requirements. CMS schedule for HAI reporting and additional reporting criteria for reporting is outlined in Appendix 3.	

#### 3. Prevention

State implementation of HHS Healthcare Infection Control Practices Advisory Committee (HICPAC) recommendations is a critical step toward the elimination of HAIs. CDC and HICPAC have developed evidence-based HAI prevention guidelines cited in the HHS Action Plan for implementation. These guidelines are translated into practice and implemented by multiple groups in hospital settings for the prevention of HAIs. CDC guidelines have also served as the basis for the Centers for Medicare and Medicaid Services (CMS) Surgical Care Improvement Project. These evidence-based recommendations have also been incorporated into Joint Commission standards for accreditation of U.S. hospitals and have been endorsed by the National Quality Forum. Please select areas for development or enhancement of state HAI prevention efforts.

Table 3: State planning for HAI prevention activities

Check	Check	Items Planned for Implementation (or currently underway)	Target Dates for
Items	Items		Implementation
Underway	Planned		
		1. Implement HICPAC recommendations	Initiated 2009 and
			ongoing
		i. Develop strategies for implementation of HICPAC recommendations for at least 2	
		prevention targets specified by the state multidisciplinary group.	
		Other activities or descriptions:	
		The HICPAC recommendations for CLABSIs include use of a set of prevention practices that need to	
		be used in concert to be effective in preventing HAIs, commonly referred to as "bundles." The	
		implementation of these bundles is a strategy promoted by DPH. The Comprehensive Unit Based	
		Safety Program developed at Johns Hopkins was introduced in CT in 2009 with 17 intensive care	
		unit (ICU) teams participating from 14 hospitals. The Connecticut hospitals that have participated in	
		the project have committed their ICU teams to work collaboratively to prevent CLABSIs by	
		standardizing processes related to the insertion, maintenance, and removal of central-lines, and	
		measurably improving the culture of safety in the ICU. In its final year (2011) of the project, teams	
		continued to spread their successful interventions hospital-wide and attended a final session to	
		celebrate their collective achievement.	
		In the fall of 2011 CHA expanded the Stop BSI project to encompass the Stop CAUTI project, a	
		national initiative aimed at reducing catheter-associated urinary tract infections (CAUTI). The goal	
		of the project is to reduce CAUTIs by 25 percent through the implementation of best practices for	
		the appropriate placement, continuance, and timely removal of urinary tract catheters.	
		2. Establish prevention working group under the state HAI advisory council to coordinate state	Initiated 2009 and
		HAI collaboratives	ongoing
		i. Assemble expertise to consult, advise, and coach inpatient healthcare facilities involved	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		in HAI prevention collaboratives	
		Other activities or descriptions: The DPH works in partnership with the Connecticut Hospital Association (CHA) and Qualidigm. Together they collaborate with healthcare providers to facilitate sharing local and national best practices, tools and resources, and strategies for implementing prevention initiatives and garnering leadership support. The DPH has either hosted or participated in a number of seminars on infection prevention and approaches for promoting quality improvement. DPH Commissioner Mullen has been regularly communicating with hospital Chief Executive Officers through circular letters and memos about HAI reporting initiatives.	
		All hospitals licensed by the DPH have a hospital-wide program for the prevention, control, and investigation of infectious diseases. Nurses, physicians, medical technologists, and other professionals who have acquired special training in infection control or epidemiology manage these programs. Hospitals collaborate via the CHA Infection Prevention Coordinators Conference and the DPH HAI Advisory Committee.	
		The efforts of these infection prevention and control programs have resulted in the development of several national HAI prevention programs to help eliminate HAIs. One of these is the Comprehensive Unit-based Safety Program (CUSP) described above, and now managed by Health Research and Educational Trust (HRET). This program uses carefully crafted quality improvement and workplace culture change methods to achieve the goal of consistently and sustainably by incorporating proven best practices to prevent CLABSIs and CAUTIs.	
$\boxtimes$		<ul> <li>3. Establish HAI collaboratives with at least 10 hospitals (this may require a multi-state or regional collaborative in low population density regions)</li> <li>i. Identify staff trained in project coordination, infection control, and collaborative</li> </ul>	Initiated 2009 and ongoing
		coordination  ii. Develop a communication strategy to facilitate peer-to-peer learning and sharing of best practices	
		iii. Establish and adhere to feedback from standardized outcome data to track progress  Other activities or descriptions:  As above  Additionally, Qualidigm, a national healthcare consulting and research company that provides	2014

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		<ul> <li>innovative and scientifically-based solutions to transform and improve care delivery and patient outcomes, was contracted with the state to conduct two collaboratives through July 2014.</li> <li>i. The Antimicrobial Stewardship (AMS) prevention collaborative was used to assist 11 participating communities to implement antimicrobial stewardship programs based on their community specific needs.</li> <li>ii. The Clostridium Difficile (CDI) prevention collaborative worked with 33 participating long term care facilities to reduce CDI and multi-drug resistant infections using rapid cycle quality improvement techniques. The collaborative had a kick off meeting in April 2014 that discussed CDI pathophysiology, treatment and prevention. Participants also developed AIM statements and completed self-assessment tools utilizing APIC standards. Participants identified targeted areas for improvement such as hand hygiene, early diagnosis, rapid containment and environmental cleaning</li> </ul>	
		<ul> <li>Develop state HAI prevention training competencies</li> <li>i. Consider establishing requirements for education and training of healthcare professionals in HAI prevention (e.g., certification requirements, public education campaigns, and targeted provider education) or work with healthcare partners to establish best practices for training and certification</li> </ul>	Initiated 2009 and ongoing
		<ul> <li>Other activities or descriptions:</li> <li>DPH received a Public Health Foundation Future of Public Health Award to incorporate quality Improvement methods into public health and medical practice. The DPH led a Clostridium difficile prevention collaborative with 25 long term care facilities</li> <li>DPH supported the End Stage Renal Disease Network of New England in its development of the first NHSN dialysis validation protocol in the nation.</li> </ul>	2011

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		<ul> <li>Federal stimulus act (ARRA)-funded APIC trainings in infection control (EPI 101 and 201) in 2011.</li> <li>CT DPH also developed public service ads and announcements, winning a 2012 Silver Award in the Integrated Communications – Not-For-Profit category, the campaign was designed to increase Connecticut residents' awareness of statewide prevention efforts for reducing healthcare associated infections (HAIs) and what Connecticut healthcare patients, visitors, workers and providers could do to aid in these efforts. Through a series of television ads, city transit bus posters, and prevention materials displayed in healthcare facilities around Connecticut, the infection prevention messages were communicated to Connecticut residents.</li> <li>Steps that Connecticut residents can do to prevent healthcare associated infections, include;</li> <li>Good Hand Hygiene: Wash your hands</li> <li>Good Respiratory Etiquette: Sneeze in your Sleeve</li> <li>Know Your Vaccination Status Be up-to-date on immunizations for both childhood diseases and adult diseases: Get your Flu Shot</li> <li>Know the proper use of antibiotics: Finish Your Antibiotics</li> <li>Pre-Surgery Preparedness: Make My Day - Follow your pre-surgical instructions.</li> <li>Speak-Up: Be your own advocate or bring a family member or friend with you to all healthcare appointments. Bring a friend with you to your doctor's appointment.</li> </ul>	
		5. Implement strategies for compliance to promote adherence to HICPAC recommendations	Initiated 2009 and ongoing
		<ul> <li>Consider developing statutory or regulatory standards for healthcare infection control and prevention or work with healthcare partners to establish best practices to ensure adherence</li> </ul>	
		<ul> <li>ii. Coordinate/liaise with regulation and oversight activities such as inpatient or outpatient facility licensing/accrediting bodies and professional licensing organizations to prevent HAIs</li> </ul>	

Check Items Underway	Check Items Planned	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
		iii. Improve regulatory oversight of hospitals, enhance surveyor training and tools, and add sources and uses of infection control data iv. Consider expanding regulation and oversight activities to currently unregulated settings	
		where healthcare is delivered and work with healthcare partners to establish best practices to ensure adherence	
		Other activities or descriptions:	
		The DPH created a statewide health improvement plan that provides a blueprint for action to address some of Connecticut's most challenging health issues. The plan helps to promote	
		Healthy Connecticut 2020, the DPH plan for improving the health of Connecticut residents by the end of 2020. The plan provides a common framework for organizations to use in leveraging	
		resources, engaging partners, identifying their own priorities, and strategies for collective action. The DPH will work with the HAI Advisory Committee and other partners and	
		stakeholders on the objectives in the plan. Some of the objectives include:	
		<ul> <li>i. Enhancing the State's public reporting infrastructure for healthcare associated infections</li> <li>ii. Reducing the rate of catheter-associated urinary tract infections in long term care facilities</li> <li>iii. Reducing the rate of <i>Clostridium difficile</i> infections in long term care facilities</li> </ul>	
		<ul><li>iv. Reducing the rate of central line-associated bloodstream infections in hemodialysis facilities</li><li>v. Reducing the number of surgical site infections in ambulatory surgical centers</li></ul>	
		6. Enhance prevention infrastructure by increasing joint collaboratives with at least 20 hospitals (i.e. this may require a multi-state or regional collaborative in low population density regions)  As above	Initiated 2009 and ongoing
		Other activities or descriptions:	
$\boxtimes$		7. Establish collaborative(s) to prevent HAIs in nonhospital settings (e.g., long term care, dialysis)  As above	Initiated 2009 and ongoing
		Other activities or descriptions:	

#### 4. Evaluation and Communication

Program evaluation is an essential organizational practice in public health. Continuous evaluation and communication of findings integrates science as a basis for decision-making and action for the prevention of HAIs. Evaluation and communication allows for learning and ongoing improvement. Routine, practical evaluations can inform strategies for the prevention and control of HAIs. Please select areas for development or enhancement of state HAI prevention efforts.

Table 4: State HAI communication and evaluation planning

Check Items	Check	Items Planned for Implementation (or currently underway)	Target Dates for
Underway	Items		Implementation
	Planned		
		1. Conduct needs assessment and/or evaluation of the state HAI program to learn	July-August 2015
		how to increase impact	
		i. Establish evaluation activity to measure progress toward targets and	
		An assessment survey sent to all ACH IPs in spring 2015; survey results analyzed July 2015.	
		ii. Establish systems for refining approaches based on data gathered	
		Data from ACH IP survey will be presented to the Multidisciplinary Group and members	
		will work with DPH HAI staff on writing of the relevant section in our new state HAI plan.	
		Other activities or descriptions (not required):	
		2. Develop and implement a communication plan about the state's HAI program	Initiated 2008 and
		and about progress to meet public and private stakeholders needs	ongoing
		<ul> <li>Disseminate state priorities for HAI prevention to healthcare organizations, professional provider organizations, governmental agencies, non-profit public health organizations, and the public</li> </ul>	
		Transparency and Communication with Public and Private Stakeholders: Connecticut's DPH has alway been open and transparent regarding information and statistics on outbreaks of disease and infection control and prevention in the state. The DPH provides a variety of on-line publications reporting statistics, outbreaks, and preventive measures for both healthcare agencies and the public, including the	

	<ul> <li>following:         <ol> <li>The Connecticut Epidemiologist Newsletter, this publication has been produced by the Connecticut Department of Public Health, since 1982. It is an important tool for the Division of Infectious Diseases to supply current disease information to primary care physicians, infection preventionists, other health professionals, and the public. It includes information on outbreaks of infectious disease and other epidemiologically significant diseases in the state. It is available on the DPH website at: <a href="http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=388262">http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=388262</a></li> </ol> </li> <li>ii. The DPH HAI website also contains information on types of HAIs, multi-drug resistant bacteria, infection prevention, and links to CDC's website: <a href="http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=417318">http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=417318</a></li> <li>iii. Reportable disease statistics can be found at: <a href="http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=388390&amp;dphNav_GID=1601">http://www.ct.gov/dph/cwp/view.asp?a=3136&amp;q=388390&amp;dphNav_GID=1601</a></li> <li>iv. Connecticut's NHSN reported infection rates can be found at: <a href="http://www.cdc.gov/hai/pdfs/stateplans/factsheets/ct.pdf">http://www.cdc.gov/hai/pdfs/stateplans/factsheets/ct.pdf</a></li> </ul>	
	Other activities or descriptions:	
	<ul> <li>3. Provide consumers access to useful healthcare quality measures</li> <li>i. Disseminate HAI data to the public</li> <li>See above.</li> <li>Information can also be found on Medicare's Hospital Compare website:</li> <li><a href="http://www.medicare.gov/hospitalcompare/search.html">http://www.medicare.gov/hospitalcompare/search.html</a></li> </ul>	Initiated 1982 and ongoing  2012
<u> </u>	Other activities or descriptions:	
	4. Guide patient safety initiatives  i. Identify priorities and provide input to partners to help guide patient safety initiatives and research aimed at reducing HAIs  Every year since 2008, the DPH HAI program has conducted data validation audits followed by HAI training programs for the purpose of reviewing validation results, and reviewing NHSN protocols and surveillance definitions based on common errors,	2008 and ongoing
	knowledge gaps, and misinterpretations identified from the validation audits. These trainings have included many statewide HAI prevention partners such as: CT HAI Advisory Committee members; Qualidgm; the CT Hospital Association (CHA); the CT Emerging Infections Program (EIP) at Yale University; the CT Infectious Disease Society;	

APIC-New England; the CT Infection Prevention Group; the Infection Control Nurses of CT; and the IPRO End State Renal Disease (ESDR) Network of New England. Additionally, these trainings have enhanced communication with the IPs, resulting in DPH acting as a resource and advocate for consistent, objective and independent application of NHSN definitions. A review of the 2012 validation results and intense NHSN surveillance definition review was presented to the Acute care hospitals (ACH), Long term acute care hospitals (LTACs), and Inpatient Rehabilitation Facilities (IRFs) staff on March 27 and April 10, 2014. In addition to sharing the validation audit results with each individual hospital through telephone calls and a formal "CT DPH Audit Discrepancy Report", DPH has discussed this data with the CDC NHSN team through publication of the data, presentation of the data with the NHSN team which included the audit challenges and lessons learned, and participation as a member of the DHQP CAUTI ad-hoc expert panel.	
Other activities or descriptions:	

### Healthcare Infection Control and Response (Ebola-associated activities)

The techniques and practice on which infection control protocols are based form the backbone of infectious disease containment for pathogens that are otherwise amplified and accelerated in healthcare settings. Investments in a more robust infection control infrastructure will prevent many HAIs transmitted to, and among, patients and health care workers.

**Table 5: Infection Control Assessment and Response** 

Check Items Underway	Check Items	Items Planned for Implementation (or currently underway)	Target Dates for Implementation
	Planned		
		Create an inventory of all healthcare settings in state. List must include at least one infection control point of contact at the facility	October 1, 2015
$\boxtimes$		Since 2008, DPH has maintained a list of healthcare facilities reporting to NHSN. The list is updated annually. As of July 1, 2015, 131 CT facilities report to NHSN. In 2015 DPH is in the process of re- designing an Excel spreadsheet for each category of healthcare facility with name, address, # of beds, type of regulatory/licensure, infection control point of contact, etc. see example, Appendix 4	
		2. Identify current regulatory/licensing oversight authorities for each healthcare facility and explore ways to expand oversight As above. Working with Wendy Furniss, RNC, MS, Branch Chief, Healthcare Quality and Safety Branch, CT DPH to identify oversight authority for each healthcare facility.	
		Other activities or descriptions:	
		<ul> <li>Assess readiness of Ebola-designated facilities within the state         <ol> <li>Use CDC readiness assessment tool and determine gaps in infection control</li> <li>Address gaps (mitigate gaps)</li> <li>Conduct follow-up assessments</li> </ol> </li> <li>There are no designated Ebola treatment centers in CT. CDC came to CT in November 2014 to assess 2 hospitals for Ebola readiness as Assessment Facilities. A third evaluation will be conducted in a third hospital on 8/25/15. From these evaluations gaps will be</li> </ul>	October 1, 2015

identified, mitigated, and followed-up assessments will be conducted.	
Other activities or descriptions:  4. Assess outbreak reporting and response in healthcare facilities i. Use standard assessment tool and determine gaps in outbreak reporting and response ii. Address gaps (mitigate gaps) iii. Track HAI outbreak response and outcome  DPH HAI Program will:  • Identify 2012-2015 NHSN HAI rates and HAI outbreaks by individual acute care hospitals (ACH);  • Identify 2012-2015 outbreak reports to DPH from Reportable Disease databases;  • Query individual ACH facility for facility specific outbreak identification and response activities;  • Correlate facility specific NHSN rates with facility specific outbreak/prevention activities;  • Request participation and attend IC Committee meetings in each of the 29 ACHs beginning with ACHs with HAI rates above baseline;  • Present NHSN data, identified HAI reporting gaps, outbreak/prevention activities (individual and statewide), and examples of successes, for discussion and problem solving at IC meeting;  • Offer education, consultation, technical assistance, staff trainings, collaboration with state prevention partners, and updates at future IC meetings;  • Prepare progress reports summarizing DPH HAI program activities for state HAI Advisory group	October 1, 2015 and ongoing

**Table 6: Targeted Healthcare Infection Prevention Programs** 

Check Items Underway	Check Items Planned	Target Dates for Implementation	
Underway		<ol> <li>Expand infection control assessments</li> <li>Expand assessments to other additional facilities and other healthcare settings and determine gaps in infection control</li> <li>Address gaps (mitigate gaps)</li> <li>Conduct follow-up assessments</li> <li>CT has 131 healthcare facilities reporting data to CMS/NHSN: 29 acute care hospitals (ACH), 6 Long Term Acute Care (LTAC), 8 Inpatient Rehabilitation facilities (IRFs), 44 End Stage Renal Dialysis facilities (ESRD) and 44 Ambulatory Surgical Facilities (ASCs).</li> <li>Data activities will expand to the other NHSN healthcare facilities once ongoing ACH reports are established.</li> <li>By September 1, 2015, DPH will prepare and conduct a hands-on, interactive NHSN data training with attendance from all 29 ACHs, 6 LTAC, and 8 IRFs. Since reporting modules are different for the CT ESRDs and ASCs, separate trainings will be conducted. Written guidance on data analysis and preparation of HAI data reports (monthly, quarterly, annual, trends over time) by patient care unit and pathogens will be provided. One-on-one or other additional trainings will be scheduled as needed.</li> <li>In May 2016, DPH will conduct an annual training on 2016 NHSN Surveillance Definitions. The six state QIO and CT Hospital Association will partner with this training. This training will be attended by all ACH, LTAC, IRF IPs.</li> <li>By May 1, 2016, in partnership with the six-state New England QIO and CT hospital association, DPH HAI will organize and present a state conference on HAI Prevention.</li> </ol>	October 1, 2015 and ongoing

Other activities or descriptions:	
<ul> <li>Increase infection control competency and practice in all healthcare settings through training</li> <li>i. Incorporate general infection control knowledge and practice assessments of competency into state licensing board requirements, credentialing, and continuing education requirements for clinical care providers (e.g., medical license, admitting privileges) and/or licensing/accreditation requirements for healthcare facilities.</li> </ul>	October 1, 2015 and ongoing
CT is in the process of restructuring the HAI Advisory Committee and adding more organizations to the multidisciplinary team. Prospective additional members are:  Pro Health (Group of private MDs)  Long Term Care Association  CT Association of Ambulatory Surgical Centers  CT Association of Ambulatory Surgical Centers  CT Hospice  School Based Clinics  Urgent Care Centers  Other Out Pt Clinics (MD owned)  Nursing Homes  Home Health Agencies  Insurance Payers (Aetna)  Veterinarians  Dentists  Hospital Preparedness Coordinators  Connecticut Medical Society  By working closely with these additional stakeholders greater opportunities for improvement can be identified, more focused training can be developed and issues of regulation/licensure/privileges can be addressed.	

		ii. Develop a sustainable training program based on CDC guidance and technical assistance to perform training, prioritizing on-site train-the-trainer programs in
		key domains of infection control, including the incorporation of hands-on
		evaluations and competency assessments of best practices and system to
		monitor ongoing compliance and competency.
		<ul> <li>The Yale School of Public Health, through its Office of Public Health Practice</li> </ul>
		(OPHP), will develop and deliver a train-the-trainer style continuing education
$\boxtimes$	$\boxtimes$	(CE) program on basic infection control (IC) to frontline staff in selected priority
		outpatient healthcare settings: federally qualified health centers (FQHCs); school
		based health centers (SCHCs); physician offices and urgent care clinics. The
		project will consist of three phases: a needs assessment phase, a program
		development phase and a program delivery phase.
		<ul> <li>In the needs assessment phase, OPHP will reach out to representatives of the</li> </ul>
		outpatient clinics above to gather information on 1) current IC CE training being
		provided to staff (content and delivery method); 2) ideal length of training
		program; 3) training audiences/staff composition; 4) potential trainers/ IC staff;
		5) perceived barriers to consistently implementing IC protocols in the work
		setting and 5) possible relationships between this IC training program and other
		agency goals (performance management, licensure, etc.). A summary report of
		findings will be written by OPHP.
		Based on the needs assessment results, OPHP will lead the development of the
		IC curricula and the train-the-trainer program.
		OPHP will contract with an IC subject matter expert (SME) to design and develop
		the curricula and a basic learning assessment and evaluation.
		OPHP will contract with a consultant with expertise in the design and  development of weaking as trains the training areas as a develop training.
		development of workplace train-the-trainer programs to develop training
		program materials and a training schedule. The curricula will be reviewed by
		representatives of the outpatient settings and approved by CT DPH.
		OPHP will partner with representatives of the outpatient clinics, and other
		community groups as needed, to identify appropriate trainers to be trained to
		deliver the curricula either as "master trainers" and/or as trainers in the
		workplace.

		<ul> <li>OPHP will coordinate the train-the-trainer program implementation in partnership with representatives of the outpatient clinics, the SME and the consultant.</li> <li>OPHP will arrange for the training sessions for trainers, including the training venues, instructors and program materials. In partnership with representatives of the outpatient clinics,</li> <li>OPHP will assist trainers in planning program delivery, developing their delivery schedules and administering the assessment and evaluation.</li> <li>As is practicable, OPHP will work with the trainers/worksites to track the delivery of the training and results of the assessment and evaluation, such as sampling.</li> </ul>	
		Other activities or descriptions: Infection Prevention in Hemodialysis – Nurse "Train the Trainer" Session	March 2015
		IPRO and the Connecticut Dept. of Public Health (Healthcare Associated Infection Division) have collaborated to create a program that trains one RN from each licensed hemodialysis facility in CT to obtain the skills needs to train the staff in their respective facilities, using the Centers of Disease Control and Prevention guidance and materials.	
		<ul> <li>3. Enhance surveillance capacity to improve situational awareness, describe emerging threats, and target onsite assessments to implement prevention programs         <ol> <li>Build capacity to analyze data reported by facilities in a defined region to allow for a comprehensive assessment of potential healthcare-associated infection threats, and communicate results with healthcare facilities.</li> <li>Work with CDC to guide analytic direction and identify facilities for prioritized assessments/response</li> <li>Improve outbreak reporting capacity by developing an infrastructure that includes clear definitions of infectious threats of epidemiologic importance</li> </ol> </li> </ul>	October 1, 2015 and ongoing
		that are communicated to facilities  iv. Implement a response plan to address potential emerging threats identified by using enhanced surveillance	

<ul> <li>Other activities or descriptions:         <ul> <li>Since 9/11/2001 CT acute care hospital have had to report state-wide syndromic surveillance based on unscheduled hospital admissions (HASS). The system's objectives are to monitor for outbreaks caused by Category A biologic agents and evaluate limits in space and time of identified outbreaks. Twenty nine acute-care hospitals are required to report their previous day's unscheduled admissions for 11 syndromes: pneumonia, hemoptysis, respiratory distress, acute neurologic illness, nontraumatic paralysis, sepsis and nontraumatic shock, fever with rash, fever of unknown cause, acute gastrointestinal illness, possible cutaneous anthrax, and suspected illness clusters.</li> <li>Ebola Response: There are no designated Ebola treatment centers in CT. CDC came to CT in spring 2015 to assess 2 hospitals for Ebola readiness as Assessment Facilities. A third evaluation will be conducted in a third hospital on 8/25/15. From these evaluations gaps will be identified.</li> <li>DPH in in the process of assessing all current outbreak reporting systems (HASS, reportable diseases, etc.) to develop a more inclusive and comprehensive outbreak reporting plan. DPH is currently reviewing other state plans for guidance.</li> <li>The HAI Advisory Committee is analyzing NHSN data through TAP reports</li> </ul> </li> </ul>	2001 and ongoing

#### Appendix 1

The HHS Action plan identifies metrics and 5-year national prevention targets. These metrics and prevention targets were developed by representatives from various federal agencies, the Healthcare Infection Control Practices Advisory Committee (HICPAC), professional and scientific organizations, researchers, and other stakeholders. The group of experts was charged with identifying potential targets and metrics for six categories of healthcare-associated infections:

- Central Line-associated Bloodstream Infections (CLABSI)
- Clostridium difficile Infections (CDI)
- Catheter-associated Urinary Tract Infections (CAUTI)
- Methicillin-resistant Staphylococcus aureus (MRSA) Infections
- Surgical Site Infections (SSI)
- Ventilator-associated Pneumonia (VAP)

Following the development of draft metrics as part of the HHS Action Plan in January 2009, HHS solicited comments from stakeholders for review.

#### Stakeholder feedback and revisions to the original draft Metrics

Comments on the initial draft metrics published as part of the HHS Action Plan in January 2009 were reviewed and incorporated into revised metrics. While comments ranged from high level strategic observations to technical measurement details, commenters encouraged established baselines, both at the national and local level, use of standardized definitions and methods, engagement with the National Quality Forum, raised concerns regarding the use of a national targets for payment or accreditation purposes and of the validity of proposed measures, and would like to have both a target rate and a percent reduction for all metrics. Furthermore, commenters emphasized the need for flexibility in the metrics, to accommodate advances in electronic reporting and information technology and for advances in prevention of HAIs, in particular ventilator-associated pneumonia.

To address comments received on the Action Plan Metrics and Targets, proposed metrics have been updated to include source of metric data, baselines, and which agency would coordinate the measure. To respond to the requests for percentage reduction in HAIs in addition to HAI rates, a new type of metric, the standardized infection ratio (SIR), is being proposed. Below is a detailed technical description of the SIR.

Below is a table of the revised metrics described in the HHS Action plan. Please select items or add additional items for state planning efforts.

Metric Number and Label	Original HAI Elimination Metric	HAI Comparison Metric	Measurement System	National Baseline Established (State Baselines Established)	National 5-Year Prevention Target	Coordinator of Measurement System	Is the metric NQF endorsed?
1. CLABSI 1	CLABSIs per 1000 device days by ICU and other locations	CLABSI SIR	CDC NHSN Device- Associated Module	2006-2008 (proposed 2009, in consultation with states)	least 50% from baseline or to	CDC	Yes*
2. CLIP 1 (formerly CLABSI 4)	Central line bundle compliance	CLIP Adherence percentage	CDC NHSN CLIP in Device- Associated Module	2009 (proposed 2009, in consultation with states)	100% adherence with central line bundle	CDC	Yes <sup>†</sup>
3a. C diff 1	Case rate per patient days; administrative/disc harge data for ICD-9 CM coded Clostridium difficile Infections	Hospitalizations with <i>C. difficile</i> per 1000 patient discharges	Hospital discharge data	(proposed 2008, in consultation	At least 30% reduction in hospitalizations with <i>C. difficile</i> per 1000 patient discharges	AHRQ	No
3b. C diff 2 (new)		C. difficile SIR	CDC NHSN MDRO/CDAD Module LabID <sup>‡</sup>		Reduce the facility-wide healthcare facility-onset <i>C. difficile</i> LabID event SIR by at least 30% from baseline or to zero	CDC	No
4. CAUTI 2	# of symptomatic UTI per 1,000 urinary catheter days	CAUTI SIR	CDC NHSN Device- Associated Module	2009 for ICUs and other locations 2009 for other hospital units (proposed 2009, in consultation	Reduce the CAUTI SIR by at least 25% from baseline or to zero in ICU and other locations	CDC	Yes*

Metric Number and Label	Original HAI Elimination Metric	HAI Comparison Metric	Measurement System	National Baseline Established (State Baselines Established)	National 5-Year Prevention Target	Coordinator of Measurement System	Is the metric NQF endorsed?
	Incidence rate (number per 100,000 persons) of invasive MRSA infections	MRSA Incidence rate	CDC EIP/ABCs	(for non-EIP states, MRSA	At least a 50% reduction in incidence of healthcareassociated invasive MRSA infections	CDC	No
5b. MRSA 2 (new)		MRSA bacteremia SIR	CDC NHSN MDRO/CDAD Module LabID <sup>‡</sup>		Reduce the facility-wide healthcare facility-onset MRSA bacteremia LabID event SIR by at least 25% from baseline or to zero	CDC	No
	Deep incision and organ space infection rates using NHSN definitions (SCIP procedures)	SSI SIR	CDC NHSN Procedure- Associated Module	(proposed 2009, in consultation	Reduce the admission and readmission SSI <sup>§</sup> SIR by at least 25% from baseline or to zero	CDC	Yes <sup>¶</sup>
(formerly SSI 2)	Adherence to SCIP/NQF infection process measures	SCIP Adherence percentage	CMS SCIP	,	At least 95% adherence to process measures to prevent surgical site infections	CMS	Yes

<sup>\*</sup> NHSN SIR metric is derived from NQF-endorsed metric data

<sup>&</sup>lt;sup>†</sup> NHSN does not collect information on daily review of line necessity, which is part of the NQF

<sup>&</sup>lt;sup>‡</sup> LabID, events reported through laboratory detection methods that produce proxy measures for infection surveillance

<sup>§</sup> Inclusion of SSI events detected on admission and readmission reduces potential bias introduced by variability in post-discharge surveillance efforts

<sup>&</sup>lt;sup>¶</sup> The NQF-endorsed metric includes deep wound and organ space SSIs only which are included the target.

#### Understanding the Relationship between HAI Rate and SIR Comparison Metrics

The Original HAI Elimination Metrics listed above are very useful for performing evaluations. Several of these metrics are based on the science employed in the NHSN. For example, metric #1 (CLABSI 1) for CLABSI events measures the number of CLABSI events per 1000 device (central line) days by ICU and other locations. While national aggregate CLABSI data are published in the annual NHSN Reports these rates must be stratified by types of locations to be risk-adjusted. This scientifically sound risk-adjustment strategy creates a practical challenge to summarizing this information nationally, regionally or even for an individual healthcare facility. For instance, when comparing CLABSI rates, there may be quite a number of different types of locations for which a CLABSI rate could be reported. Given CLABSI rates among 15 different types of locations, one may observe many different combinations of patterns of temporal changes. This raises the need for a way to combine CLABSI rate data across location types.

A standardized infection ratio (SIR) is identical in concept to a standardized mortality ratio and can be used as an indirect standardization method for summarizing HAI experience across any number of stratified groups of data. To illustrate the method for calculating an SIR and understand how it could be used as an HAI comparison metric, the following example data are displayed below:

Risk Group Stratifier		Observed CLABSI Rate	s	NHSN CLABSI Rates for 2008 (Standard Population)			
Location Type	#CLABSI	#Central line-days	CLABSI rate <sup>*</sup>	#CLABSI	#Central line-days	CLABSI rate*	
ICU	170	100,000	1.7	1200	600,000	2.0	
WARD	58	58 58,000		600	400,000	1.5	
$SIR = \frac{\text{observed}}{\text{expected}} = \frac{170 + 58}{100000 \times \left(\frac{2}{1000}\right) + 58,000 \times \left(\frac{1.5}{1000}\right)} = \frac{228}{200 + 87} = \frac{228}{287} = 0.79 \qquad 95\%CI = (0.628,0.989)$							

\*defined as the number of CLABSIs per 1000 central line-days

In the table above, there are two strata to illustrate risk-adjustment by location type for which national data exist from NHSN. The SIR calculation is based on dividing the total number of observed CLABSI events by an "expected" number using the CLABSI rates from the standard population. This "expected" number is calculated by multiplying the national CLABSI rate from the standard population by the observed number

of central line-days for each stratum which can also be understood as a prediction or projection. If the observed data represented a follow-up period such as 2009 one would state that an SIR of 0.79 implies that there was a 21% reduction in CLABSIs overall for the nation, region or facility.

The SIR concept and calculation is completely based on the underlying CLABSI rate data that exist across a potentially large group of strata. Thus, the SIR provides a single metric for performing comparisons rather than attempting to perform multiple comparisons across many strata which makes the task cumbersome. Given the underlying CLABSI rate data, one retains the option to perform comparisons within a particular set of strata where observed rates may differ significantly from the standard populations. These types of more detailed comparisons could be very useful and necessary for identifying areas for more focused prevention efforts.

The National 5-year prevention target for metric #1 could be implemented using the concept of an SIR equal to 0.25 as the goal. That is, an SIR value based on the observed CLABSI rate data at the 5-year mark could be calculated using NHSN CLABSI rate data stratified by location type as the baseline to assess whether the 75% reduction goal was met. There are statistical methods that allow for calculation of confidence intervals, hypothesis testing and graphical presentation using this HAI summary comparison metric called the SIR.

The SIR concept and calculation can be applied equitably to other HAI metrics list above. This is especially true for HAI metrics for which national data are available and reasonably precise using a measurement system such as the NHSN. The SIR calculation methods differ in the risk group stratification only. To better understand metric #6 (SSI 1) see the following example data and SIR calculation:

Risk Grou	p Stratifiers		Observed SSI Rates		NHSN SSI Rates for 2008 (Standard Population)		
Procedure Code	Risk Index Category	#SSI <sup>†</sup>	#procedures	SSI rate <sup>*</sup>	#SSI <sup>†</sup>	#procedures	SSI rate <sup>*</sup>
CBGB	1 315		12,600	2.5	2100	70,000	3.0
CBGB	2,3	210	7000	3.0	1000	20,000	5.0
HPRO	1	111	7400	1.5	1020	60,000	1.7
SIR = -	$\frac{\text{observed}}{\text{expected}} = \frac{12600}{12600}$	(30)	$0 \times \left(\frac{5.0}{100}\right) + 7400 \left(\frac{1.7}{100}\right)$	$=\frac{636}{378+350+125.8}$	$=\frac{636}{853.8}=0.74$	95%CI = (0.649	,0.851)

This example uses SSI rate data stratified by procedure and risk index category. Nevertheless, an SIR can be calculated using the same calculation process as for CLABSI data except using different risk group stratifiers for these example data. The SIR for this set of observed data is 0.74 which indicates there's a 26% reduction in the number of SSI events based on the baseline NHSN SSI rates as representing the standard population. Once again, these data can reflect the national picture at the 5-year mark and the SIR can serve as metric that summarizes the SSI experience into a single comparison.

There are clear advantages to reporting and comparing a single number for prevention assessment. However, since the SIR calculations are based on standard HAI rates among individual risk groups there is the ability to perform more detailed comparisons within any individual risk group should the need arise. Furthermore, the process for determining the best risk-adjustment for any HAI rate data is flexible and always based on more detailed risk factor analyses that provide ample scientific rigor supporting any SIR calculations. The extent to which any HAI rate data can be risk-adjusted is obviously related to the detail and volume of data that exist in a given measurement system.

In addition to the simplicity of the SIR concept and the advantages listed above, it's important to note another benefit of using an SIR comparison metric for HAI data. If there was need at any level of aggregation (national, regional, facility-wide, etc.) to combine the SIR values across mutually-exclusive data one could do so. The below table demonstrates how the example data from the previous two metric settings could be summarized.

		Observed HAI	S		Expected HA	Als
HAI Metric	#CLABSI	#SSI <sup>†</sup>	#Combined HAI	#CLABSI	#SSI <sup>†</sup>	#Combined HAI
CLABSI 1	228			287		
SSI 1		636			853.8	
Combined HAI			228 + 636 = 864			287+853.8 = 1140.8
	SIR	=	$\frac{6+636}{+853.8} = \frac{864}{1140.8} = 0$	).76 95%CI =	(0.673,0.849)	

<sup>†</sup>SSI (surgical site infection)

<sup>&</sup>lt;sup>†</sup>SSI, surgical site infection

defined as the number of deep incision or organ space SSIs per 100 procedures

# 2014 CT Healthcare Associated Infection Advisory Committee Members Public Act No. 06-142

## AN ACT CONCERNING HOSPITAL ACQUIRED INFECTIONS

There is established a Committee on Healthcare Associated Infections, which shall consist of the commissioner or the commissioner's designee, and the following members appointed by the commissioner:

Two members representing the Connecticut Hospital Association;

Two members from organizations representing health care consumers;

Two members who are either hospital-based infectious disease specialists or epidemiologists with demonstrated knowledge and competence in infectious disease related issues;

One representative of the Connecticut State Medical Society;

One representative of a labor organization representing hospital based nurses; and

Two public members.

Healthcare Associated Infections Advisory Committee Composition: June 2014

# Healthcare Associated Infections Advisory Committee Composition: June 2014 Commissioner or Commissioner's designee:

1. Wendy Furniss, RNC, MS, Branch Chief, Healthcare Quality and Safety Branch, CT DPH, Hartford, CT

#### Two Representatives from the Connecticut Hospital Association

- 1. Allison L. Hong, MD, Interim VP, Quality and Patient Safety, CT Hospital Association, Wallingford, CT
- 2. Carl Schiessel, CT Hospital Association, Wallingford, CT

#### Two Representatives from Organizations Representing Health Care Consumers

- 1. Valerie Wyzykowski, Office of Healthcare Advocate, State of CT, Hartford, CT
- 2. Jean Rexford, Exec. Dir., CT Center for Patient Safety, Hartford, CT

#### Two Representatives from Hospital-based Infectious Disease Specialist or Epidemiologist

- 1. Louise Dembry, MD, Hospital Epidemiologist, Yale-New Haven Hospital, New Haven, CT
- 2. Brenda Grant, RN, MPH, CIC, CHES, Manager, Infection Prevention Stamford Hospital, Stamford, CT

#### One Representative from Connecticut State Medical Society

1. Jack Ross, MD, Chief, Infectious Diseases & Epidemiology, Hartford Hospital, Hartford, CT

#### One Representative from Labor organization Representing Hospital-based Nurses

1. Dale Cunningham, American Federation of Teachers, Rocky Hill, CT

#### One Member from the public

- 1. Lynne Garner, PhD, President and Trustee, The Donaghue Medical Research Foundation, West Hartford, CT
- 2. Raymond S. Andrews, retired, West Hartford, CT

#### **Total Voting Membership: 11 members**

# **APPENDIX 3**

# Healthcare Facility HAI Reporting Requirements to CMS via NHSN--Current or Proposed Requirements

CMS Reporting Program	HAI Event	Reporting Specifications	Reporting Start Date		
	CLABSI	Adult, Pediatric, and Neonatal ICUs	January 2011		
	CAUTI	Adult and Pediatric ICUs	January 2012		
	SSI: COLO	Inpatient COLO Procedures	January 2012		
	SSI: HYST	Inpatient HYST Procedures	January 2012		
Hospital	MRSA Bacteremia LabID Event	FacWidelN	January 2013		
Inpatient Quality Reporting (IQR)	C. difficile LabID Event	FacWidelN	January 2013		
Program	Healthcare Personnel Influenza Vaccination	All Inpatient Healthcare Personnel	January 2013		
	Medicare Beneficiary Number	All Medicare Patients Reported into NHSN	July 2014		
	CLABSI	Adult & Pediatric Medical, Surgical, & Medical/Surgical Wards	January 2015		
	CAUTI	Adult & Pediatric Medical, Surgical, & Medical/Surgical Wards	January 2015		
Hospital Outpatient Quality Reporting (OQR) Program	Healthcare Personnel Influenza Vaccination	All Outpatient Healthcare Personnel	October 2014		
ESRD Quality Incentive Program (QIP)	Dialysis Event (includes Positive blood culture, I.V. antimicrobial start, and signs of vascular access infection)	Outpatient Hemodialysis Facilities	January 2012		
	Healthcare Personnel Influenza Vaccination	All Healthcare Personnel	October 2015		
	CLABSI	Adult & Pediatric LTAC ICUs & Wards	October 2012		
Long Term Care	CAUTI	Adult & Pediatric LTAC ICUs & Wards	October 2012		
Hospital* Quality Reporting (LTCHQR) Program	Healthcare Personnel Influenza Vaccination	All Inpatient Healthcare Personnel	October 2014		
	MRSA Bacteremia LabID Event	FacWidelN	January 2015		
	C. difficile LabID Event	FacWidelN	January 2015		
	VAE	Adult LTAC ICUs & Wards	January 2016		
Inpatient Rehabilitation Facility Quality Reporting (IRFQR) Program	CAUTI	Adult & Pediatric IRF Wards	October 2012		
	Healthcare Personnel Influenza Vaccination	All Inpatient Healthcare Personnel	October 2014		
	MRSA Bacteremia LabID Event	FacWidelN	January 2015		
	C. difficile LabID Event	FacWidelN	January 2015		
* Long Term Care Hospitals are called Long Term Acute Care Hospitals in NHSN					

CMS Reporting Program	HAI Event	Reporting Specifications	Reporting Start Date	
Ambulatory Surgery Centers Quality Reporting (ASCQR) Program	Healthcare Personnel Influenza Vaccination	All Healthcare Personnel	October 2014	
PPS-Exempt Cancer Hospital Quality Reporting (PCHQR) Program	CLABSI	All Bedded Inpatient Locations	January 2013	
	CAUTI	All Bedded Inpatient Locations	January 2013	
	SSI: COLO	Inpatient COLO Procedures	January 2014	
	SSI: HYST	Inpatient HYST Procedures	January 2014	
Inpatient Psychiatric Facility Quality Reporting (IPFQR) Program  Healthcare Personnel Influenza Vaccination		All Inpatient Healthcare Personnel	October 2015	

# **APPENDIX 4**

# Sample Excel Spreadsheet to Be Expanded To Include Infection Control Point of Contact, Regulatory/Licensure Authority, Etc.

	CHILDREN'S GE	NERAL HOSPITAL (1)				
Facility Name <sup>a</sup>	Provider Name	Facility Address	City	Zip code	Licensed Beds	Licensed Bassinets
Connecticut Children's Medical Center	Connecticut Children's Medical Center	282 Washington Street	Hartford	06106	115	72
Center		HOSPITALS (27)				
Facility Name <sup>a</sup>	Provider Name	Facility Address	City	Zip code	Licensed Beds	Licensed Bassinets
Bridgeport Hospital	Bridgeport Hospital	267 Grant Street	Bridgeport	06610	373	10
Bristol Hospital, Inc.	Bristol Hospital, Inc., The	41 Brewster Road	Bristol	06010	134	20
Charlotte Hungerford Hospital, The	Charlotte Hungerford Hospital, The	540 Litchfield Street	Torrington	06790	109	13
Danbury Hospital, The b	Danbury Hospital, The	24 Hospital Avenue	Danbury	06810	430	26
Day Kimball Hospital	Day Kimball Healthcare, Inc.	320 Pomfret Street	Putnam	06260	104	18
Greenwich Hospital	Greenwich Hospital	5 Perryridge Road	Greenwich	06830	174	32
Griffin Hospital	Griffin Hospital, The	130 Division Street	Derby	06418	160	20
Hartford Hospital	Hartford Hospital	80 Seymour Street and	Hartford	06106	819	48
πατιτοτά πουριτάτ	,	200 Retreat Avenue	Haitiblu	00100	013	40
Hospital of Central Connecticut, The	Hospital of Central Connecticut at New Britain General and Bradley Memorial, The	100 Grand Street	New Britain	06050	414	32
John Dempsey Hospital	University or Connecticut Hearth	263 Farmington Avenue	Farmington	06030	224	10
Johnson Memorial Hospital	Johnson Memorial Hospital, Inc.	201 Chestnut Hill Road	Stafford	06076	92	9
Lawrence and Memorial Hospital	Lawrence and Memorial Hospital,	365 Montauk Avenue	New London	06320	280	28
Manchester Memorial Hospital	Manchester Memoriai Hospitai,	71 Haynes Street	Manchester	06040	249	34
Middlesex Hospital	Middlesex Hospital	28 Crescent Street	Middletown	06457	275	22
MidState Medical Center	MidState Medical Center	435 Lewis Avenue	Meriden	06451	144	12
			Milford	06460	106	12
Milford Hospital, Inc.	Milford Hospital, Inc., The	300 Seaside Avenue				
Norwalk Hospital	Norwalk Hospital Association, The ROCKVIITE General Hospital, Inc.,	34 Maple Street	Norwalk	06856	328	38
Rockville General Hospital	Tho	31 Union Street	Vernon	06066	102	16
Saint Francis Hospital and Medical Center	Saint Francis Hospital and Medical Center	114 Woodland Street and 500 Blue Hills Avenue	Hartford	06105	617	65
Saint Mary's Hospital, Inc.	Saint Mary's Hospital, Inc.	56 Franklin Street	Waterbury	06706	347	32
Sharon Hospital	Essent Heartncare of Connecticut,	50 Hospital Hill Road	Sharon	06069	78	16
St. Vincent's Medical Center	St. Vincent's Medical Center	2800 Main Street	Bridgeport	06606	473	47
Stamford Hospital, The	Stamford Hospital, The	30 Shelburne Road	Stamford	06904	305	25
Waterbury Hospital	Waterbury Hospital, The	64 Robbins Street	Waterbury	06708	357	36
Facility Name <sup>a</sup>	Provider Name	Facility Address	City	Zip code	Licensed Beds	Licensed Bassinets
William W. Backus Hospital, The	William W. Backus Hospital, The	326 Washington Street	Norwich	06360	213	20
Windham Community Memorial Hospital and Hatch Hospital	Windham Community Memorial Hospital, Inc.	112 Mansfield Avenue	Windham	06226	130	14
Corporation Yale-New Haven Hospital, Inc.	Yale-New Haven Hospital, Inc.	20 York Street	New Haven	06510	1407	134
	· ' ' ' '	20 TOTA SHEEL	INC W Havell	00310	1407	134
Total of 8,444 Beds and 789 Bassinets,						
<b>Total of 8,559 Beds and 861 Bassinets,</b> Source: DPH licensure files and e-licensure						
the license	34143435 4301 0010001 2014					
	tellite campus of Danbury Hospital ef ospital license.	fective October 1, 2014 and	Danbury Hosp	oital became	licensed for 4	30 beds and