patient's tissue during scope insertion. The results were a relief to the patients and families. **Conclusions:** It is prudent to investigate residual foreign tissue in a medical device that is being used on patients with mucosal breaches. Molecular pathology involving human identity testing is a very useful tool in the investigation of these types of events.

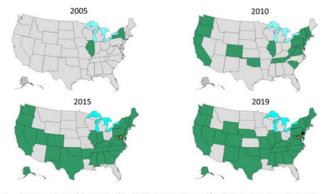
Funding: None Disclosures: None Doi:10.1017/ice.2020.1024

Presentation Type:

Poster Presentation

Shifting Landscape of Healthcare-Associated Infection and Antimicrobial Resistant Infection Reporting Policy, 2005-2019 Jeremy Goodman, Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, CDC; Samuel Clasp, Population Health and Healthcare Office, Office of the Associate Director for Policy and Strategy, CDC; Arjun Srinivasan, Centers for Disease Control and Prevention; Elizabeth Mothershed, Centers for Disease Control and Prevention; Seth Kroop, Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, CDC; Lyn Nguyen, Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, CDC; Tara Holiday, Centers for Disease Control and Prevention

Background: Healthcare-associated infections (HAIs) are a serious threat to patient safety; they account for substantial morbidity, mortality, and healthcare costs. Healthcare practices, such as inappropriate use of antimicrobials, can also amplify the problem of antimicrobial resistance. Data collected to target HAI prevention and antimicrobial stewardship efforts and measure progress are an important resource for assuring transparency and accountability in healthcare, tracking adverse outcomes, investigating healthcare practices that may spread or protect against disease, detecting and responding to the spread of resistant pathogens, preventing infections, and saving lives. Methods: We discuss 3 healthcareassociated infection and antimicrobial Resistant infection (HAI-AR) reporting types: NHSN HAI-AR reporting, reportable diseases, and nationally notifiable diseases. HAI-AR reporting requirements outline facilities and data to report to NHSN and the health department to comply with state laws. Reportable diseases are those that facilities, providers, and laboratories are required to report to the health department. Nationally notifiable diseases are those reported by health departments to the CDC for nationwide surveillance and analysis as determined by Council of State and Territorial Epidemiologists (CSTE) and the CDC. Data presented are based on state and federal policy; NHSN data are based on CDC reporting statistics. Results: Since the 2005 launch of the CDC NHSN and publication of federal advisory committee HAI reporting guidance, most states have established policies stipulating healthcare facilities in their jurisdiction report HAIs and resistant infections to the NHSN to gain access to those data, increasing from 2 states in 2005, to 18 in 2010, and to 36 states, Washington, DC, and Philadelphia in 2019. Reporting policies and NHSN participation expanded greatly following the 2011 inception of CMS HAI quality reporting requirements, with several states aligning state requirements with CMS reporting. States listing carbapenem-resistant Enterobacteriaceae (CRE) as a reportable disease increased from 7 in 2013 to 41 states and the District of Columbia in 2019. Vancomycin-intermediate and



States with Healthcare Facility NHSN Reporting and/or Data Access Requirements

Fig. 1.

NHSN Extension of Coverage

Facilities Reporting to NHSN and CMS Reporting Requirement Start Dates



Fig. 2.

vancomycin-resistant Staphylococcus aureus (VISA/VRSA) was added as a nationally notifiable disease in 2004, carbapenemaseproducing CRE (CP-CRE) was added in 2018, and Candida auris clinical infections were added in 2019. The CDC and most jurisdictions with HAI reporting mandates issue public reports based on aggregate state data and/or facility-level data. States may also alert healthcare providers and health departments of emerging threats and to assist in notifying patients of potential exposure. Conclusions: Through efforts by health departments, facilities, patient advocates, partners, the CDC, and other federal agencies, HAI-AR reporting has steadily increased. Although reporting laws and data uses vary between jurisdictions, data provided serves as valuable tools to inform prevention.

Funding: None **Disclosures:** None Doi:10.1017/ice.2020.1025

Presentation Type:

Poster Presentation

Site Visits Reveal Common Gaps in Instrument Reprocessing and Sterilization at Philadelphia Dental Clinics

Tiina Peritz, Philadelphia Department of Public Health; Susy Rettig; Susan Coffin, Children's Hospital of Philadelphia

Background: Most dental clinics lack resources and oversight related to infection prevention and control (IPC) practices. Few dental clinics undergo inspections by regulatory authorities unless the state licensing authorities receive a specific complaint. Many states, including Pennsylvania, do not have continuing IPC education requirements for dental providers. In 2018-2019, the Philadelphia Department of Public Health (PDPH) received and responded to multiple complaints and concerns related to IPC practices at dental clinics. Complaints were investigated in collaboration with the Pennsylvania Department of State (PADOS). Methods: Unannounced site visits were conducted at 7 Philadelphia dental clinics from December 2018 through September 2019 as part of the public health responses. Clinic evaluations and observations by PDPH certified infection preventionists focused on (1) IPC policies and procedures, (2) staff IPC training, (3) hand hygiene, (4) personal protective equipment, (5) instrument reprocessing and sterilization, (6) injection safety, and (7) environmental cleaning and disinfection. The CDC and the Organization of Safety, Antisepsis and Prevention (OSAP) checklists were adapted for this purpose. Results: Most dental practices we visited were small, unaffiliated, owner-operated clinics. The most common gaps we identified were associated with instrument reprocessing and sterilization practices, including inadequate separation between clean and dirty work areas, limited space and availability of sinks, inappropriate use of glutaraldehyde products for instrument cleaning (n = 3, 43%), extended reuse of cleaning brushes (n = 5, 71%), sterilization or storage of sterilized instruments without appropriate packaging (n = 2, 29%), lack of spore testing or reviewing results (n = 2, 29%), and lack of documentation of sterilizer run cycles and maintenance (n = 7, 100%). Additionally, most clinics did not have well-developed IPC policies and procedures, and staff IPC trainings were neither documented nor conducted annually. Alcohol-based hand sanitizer was often not available at the point of use. Conclusions: In Philadelphia, dental clinics often lacked IPC support and oversight. Lapses across multiple key IPC domains were common. These findings suggest that public health may have a role in providing IPC support to unaffiliated dental clinics. Licensing entities can also serve a role in improving IPC practices by more widely mandating continuing IPC education as part of the dental license renewal process.

Funding: None
Disclosures: None
Doi:10.1017/ice.2020.1026

Presentation Type:

Poster Presentation

Standardized Antimicrobial Administration Ratio (SAAR) Clinical Outcomes Assessment in a Large Community Healthcare System

Hayley Burgess, HCA Healthcare; <u>Mandelin Cooper, HCA Healthcare</u>; Laurel Goldin, HCA Healthcare; Kenneth Sands, Hospital Corporation of America

Background: Research on the association between the standardized antimicrobial administration ratio (SAAR) and clinical outcomes is lacking. Objective: We compared SAAR and patient outcomes in 97 acute-care facilities affiliated with a large health-care system. Methods: Facilities were classified using the broadspectrum hospital-onset (BSHO) SAAR for medical, surgical, and medical-surgical wards as low, moderate, or high antimicrobial use: low use SAAR, <0.8; moderate use SAAR, 0.95−1.05; and high-use SAAR, >1.2. Data were included from patients aged ≥18 years who were discharged between the first quarter of 2018 and the second quarter of 2019, had nonmissing matching criteria,

BMI between 10 and 90, and at least 1 BSHO medication administered in a medical, surgical, or medical-surgical ward. Patients were matched for gender, age group, BMI category, year and quarter of discharge, ICU stay, and diagnosis-related group (DRG). Eligible drugs included all routes for cefepime, ceftazidime, doripenem, imipenem/cilastatin, meropenem, and piperacillin/tazobactam and IV only for amikacin, aztreonam, gentamicin, and tobramycin. Outcomes were evaluated in a pairwise manner using t tests or y^2 tests. **Results:** Each of the 3 study groups consisted of 6,327 patients, 51% of whom were men; average age, 63 years; 70% of whom were obese or overweight, and 19% of whom had an ICU stay. The most common DRG code was infectious and parasitic diseases (57%) followed by digestive system (9%), respiratory system (7%), and kidney and urinary tract (6%). High antibiotic use was associated with longer length of stay and a higher estimated cost per visit. Low antibiotic use was associated with higher rate of mortality and a lower rate of readmissions compared to moderate use. The low-usage group did not exhibit a statistically significant difference in mortality, readmissions, or rate of C. difficile compared to the high-usage group. Conclusions: The optimal antibiotic utilization group varied among outcomes. Further evaluation of outcomes is needed for the SAAR to understand the ranges and the relationship between the measure and clinical outcomes. Funding: None

Disclosures: None
Doi:10.1017/ice.2020.1027

Presentation Type:

Poster Presentation

Staphylococcus spp Resistance to Chlorhexidine: Is There Any Impact Related to the Routine Use for Hand Hygiene?

Icaro Boszczowski, Hospital das Clínicas University of Sao Paulo; William Kazumassa Minami, Laboratory of Medical Microbiology, University of São Paulo; Marcia Baraldi, Hospital Alemão Oswaldo Cruz; Ana Paula Marchi, Laboratory of Medical Microbiology, University of São Paulo; Sânia Alves dos Santos, Laboratory of Medical Microbiology, University of São Paulo cristiane schmitt, School of Nursing, University of São Paulo; Amanda Luiz Pires Maciel, Hospital Alemão Oswaldo Cruz; Maria Eduarda Rufino Zani, Laboratory of Medical Microbiology, University of São Paulo; Letícia Muniz Souza, Laboratory of Medical Microbiology, University of São Paulo; Silvia Figueiredo Costa, São Paulo University

Background: Although guidelines recommend the use of chlorhexidine gluconate (CHG) for hand hygiene (HH), the impact of its routine use on antimicrobial resistance is not clear. **Objective:** To analyze the impact on the CHG susceptibility among isolates obtained from hands of HCW during its routine use for HH. Methods: We conducted a crossover study at 4 medical-surgical wards of a tertiary-care hospital in São Paulo, Brazil. In 2 units (intervention group), we established routine use of CHG for HH. For the other 2 units (control group), regular soap was provided. The availability of alcohol formulation for HH was not changed during the study. Every 4 months we swapped the units, ie, those using CHG changed for regular soap and vice versa. At baseline, we cultured the hands of HCWs. Only nursing staff hands were investigated. For hand culturing, HCWs placed their hands inside a sterile bag containing a solution of phosphate-buffered saline, Tween 80, and sodium thiosulfate. After the solution incubated overnight,