467. Healthcare Workers Perceptions Regarding the Use of an Electronic Hand Hygiene Monitoring System at a VA Hospital
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Background. A cornerstone of healthcare-associated infection prevention is hand hygiene which has resulted in regulatory requirements to monitor hand hygiene compliance. Direct observation is the gold standard for hand hygiene compliance monitoring, but has several drawbacks. Electronic monitoring systems have begun to replace direct observation with several potential advantages, including larger sample size and more timely feedback. End user acceptance and adoption is a critical step to evidence-based practice implementation. To evaluate potential barriers and facilitators to adoption, we conducted a qualitative evaluation of nursing perceptions following a trial of an electronic hand hygiene compliance monitoring system.

Methods. We conducted four focus groups of 21 nursing staff on a medical/surgical inpatient unit at a tertiary care VA hospital. Nursing staff consisted of Registered Nurses, Nursing Assistants, and Health Technicians; of which there were 19 females and 2 males. Groups were audio recorded and tapes transcribed. Content analysis of transcriptions was undertaken to identify codes, categories, and themes.

Results. Themes identified as facilitators included: (1) unit champion; (2) electronic monitoring (vs. human observation); and (3) timely feedback. Themes identified as barriers included: (1) concern with data accuracy; (2) feasibility of frequent electronic observation (vs. human observation); and (3) timely feedback. Themes identified as facilitators included: (1) unit champion; (2) electronic monitoring (vs. human observation); and (3) timely feedback. Themes identified as barriers included: (1) concern with data accuracy; (2) feasibility of frequent electronic observation (vs. human observation); and (3) timely feedback.

Conclusion. Nurses staff perceived electronic monitoring improved hand hygiene compliance. Staff verbalized negative perceptions with hand hygiene compliance monitoring but preferred electronic monitoring vs. human monitoring. Most barriers discussed revolved around the need to understand how the electronic monitoring system works and need to believe the data are accurate. Implementation of this innovative technology will require extensive planning to address staff knowledge and understanding to ensure staff acceptance and adoption.

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468. The Efficacy of Alcohol Based Wipes, Gel, Foam, and Spray Compared With Liquid Soap in Eliminating Transient Hand Bacteria
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Background. Hand hygiene is a proven method of preventing the spread of pathogens and reducing healthcare-associated infections. Studies have shown that up to 50% of healthcare professionals’ (HCPs) hands were contaminated with the same pathogen as a patient with a confirmed multidrug-resistant organism, such as MRSA or VRE. This study examined the interaction of the soap with skin.

Methods. Twenty-five healthy adults were randomly chosen to participate in one of the five hand hygiene tests. Before implementing hand hygiene, moistened sterile swabs were used to rub the fingers, thumbs, and palms of both hands. The volunteers then performed one of the hand hygiene methods following WHO recommendations for hand washing and hand rubs. Wipes were used by applying a pulling motion on fingers and thumbs followed by rubbing the palms. The swabs were agitated for 15 seconds in a petri dish and plated onto Petri dishes for incubation of 48 hours at 37°C.

Results. The percent reduction in transient hand bacteria using aerobic colony counts were enumerated and calculated as follows: 90% for wipes, 82% for liquid soap, 80% for gel, 72% for foam, and 71% for spray. The wipes eliminated hand bacteria significantly better than the liquid soap (P = 0.0247) while the gel (P = 0.7239) and foam (P = 0.0661) showed no significance. Lastly, the soap performed significantly better than the spray (P = 0.0182).

Conclusion. This study demonstrated that alcohol-based wipes performed better at removing transient bacteria from the hands than liquid soap and water. This result potentially provides another method for HCPs in reducing the risk of infection for their next patient and decreasing the likelihood of transmitting an infectious agent via hands.

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469. Microbial Removal Efficacy of a Novel Nonantimicrobial Hand Soap
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Background. The CDC Hand Hygiene Guidelines recommend washing hands with soap when hands are visibly soiled. Pending changes to the United States healthcare antibiotic regulations are decreasing the availability of antimicrobial soap active ingredients making it important to understand key performance differences across soap types. The purpose of this study was to investigate the germ removal properties of a novel, nonantimicrobial soap exhibiting improved interfacial tension properties, a measure of the interaction of the soap with skin.

Methods. The novel nonantimicrobial soap was compared with a control nonantimicrobial soap. In study 1, the soaps were tested according to ASTM E2575 to determine reduction of Serratia marcescens after one use where 5 mL of soap was applied to dry hands, lathered and rinsed 30s (N = 12). Studies 2 and 3 compared the control soap and the same novel nonantimicrobial soap on both healthy skin and hand skin bacteria. In study 2, the control soap achieved a 2.26 log reduction compared with a 1.70 log reduction for the control soap (P < 0.0001). In studies 2 and 3, the nonantimicrobial soap achieved log reductions that were 0.34 (P = 0.0236) and 0.53 (P < 0.005) greater than the control soap, respectively.

Conclusion. This study indicates that a nonantimicrobial soap can achieve a high level of microbe removal (>99%) on skin. Additionally, product formulation appears to impact the microbial removal properties of nonantimicrobial soap on both healthy and irritated human skin. Therefore, this novel soap may be a good option in a high-frequency hygiene activity such as healthcare hand hygiene.

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466. Use of Administrative Data to Characterize Clostridium difficile Infections (CDI) Reported by California Hospitals to the California Department of Public Health (CDPH) via the National Healthcare Safety Network (NHSN): 2014–2015
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Background. In 2014–2015, CDI accounted for more than half of all healthcare-associated infections (HAIs) reported by California hospitals. The CDPH HAIP Program used an administrative dataset from the California Office of Statewide Health Planning and Development (OSHPD) to identify admission source (e.g., home, skilled nursing facility), length of stay, payer category, and outcome (e.g., death) of patients with CDI reported by California hospitals via NHSN.

Methods. We merged NHSN CDI events with OSHPD hospital discharge data for the period January 1, 2014, to December 31, 2015. NHSN classifies CDI cases as community onset (CO) if the CDI test specimen was collected during the first three hospital days and hospital onset (HO) if collected on day 4 or later. We used OSHPD discharge datasets (ICD-9-CM: 00845 and ICD-10-CM: A047 codes). We matched NHSN CDI cases with OSHPD hospital discharge records by hospital, admission date, and date of birth.

Results. Hospitals reported 58,841 NHSN inpatient incident and recurrent CDI events in 2014–2015. We matched 42,172 (71.7%) NHSN CDI records with an OSHPD hospital discharge record; 60.5% of matched cases were CO-CDI and 39.5% were HO-CDI. Sources of admission included home (78.2%; CO: 81.0% and HO: 74.0%), skilled nursing/intermediate care facility (10.7%; CO: 10.9% and HO: 10.4%), acute care hospital (6.0%; CO: 3.2% and HO: 10.4%), and residential care facility (1.7%; CO: 2.0% and HO: 1.4%). Payers included Medicare (61.8%), Medi-Cal (18.7%), and private insurance (16.8%). The median length of stay for CO cases was 5 days (interquartile range [IQR]: 3–9), and for HO cases, 15 days (IQR: 9–25); 8.7% (CO: 7.1% and HO: 11.2%) of patients with CDI died during hospitalization.

Conclusion. Our analysis demonstrates use of an administrative dataset to supplement NHSN HAI data. Patients with CDI were predominantly admitted from home and had prolonged hospitalizations and substantial in-hospital mortality. We are evaluating use of these data to identify hospital admissions at various time intervals before