residual chlorine); 63.1% used specific tests for Legionella; and 35.6% performed heterothetic plate counts (HPCs). We analyzed new, 2018 hospital survey data to assess further progress toward meeting CMS requirements for WMPs. Methods: We analyzed 2018 NHSN Annual Hospital Survey responses for facilities that reported on WMPs in 2017. Responses included information regarding risk assessments for Legionella and other waterborne pathogens as well as details regarding WMP teams and water-monitoring practices. WMP team members were categorized as administrative (hospital administrator, compliance officer, risk or quality management), epidemiology or infection control (epidemiologist or infection preventer, other clinical), or environmental or facilities (consultant, facility manager or engineer, equipment or chemical supplier, maintenance). Statistical significance was assessed using the McNemar test, where appropriate.

Results: Of hospitals reporting on WMPs in 2017, 4,087 of 4,929 (83%) responded again in 2018. The proportion of facilities that reported having a WMP increased from 3,258 of 4,087 (79.7%) in 2017 to 3,647 of 4,087 (89.2%) in 2018 (P < .0001). Of the 3,647 hospitals that reported having a WMP in 2018, 95.9% had conducted a risk assessment for waterborne pathogens; 67.3% of these facilities had most recently done so within 1 year of the survey. WMP teams had representation from environmental or facilities staff at 98.8% of hospitals, epidemiology or infection control staff at 89.8% of hospitals, and administrative staff at 71.7% of hospitals. Of facilities with WMPs in 2018, 90.5% reported regular monitoring of water temperature, 72.2% disinfectant, 67.4% tests for Legionella, and 48.8% HPCs. Conclusions: More hospitals reported having a WMP in 2018 than in 2017. However, ~1 in 10 respondents lacked a WMP. Differences in water monitoring practices across facilities potentially reflect a lack of standardization in how WMPs are implemented. Some hospital WMPs do not incorporate routine monitoring of water temperature and disinfectant, which is a basic practice. CDC continues to develop tools, resources, and training to support facility WMP teams in meeting CMS requirements and protecting patients from water-associated pathogens.

Funding: None
Disclosures: None

Doi:10.1017/ice.2020.577

Presentation Type: Late Breaker Poster
Communications and Screening for 2019 Novel Coronavirus at a Tertiary-Care Medical Center
 Lorinda Sheeler, University of Iowa Hospitals & Clinics; Mary Kukla, University of Iowa Hospitals and Clinics; Oluchi Abosi, University of Iowa Hospitals & Clinics; Holly Meacham, University of Iowa Hospital and Clinics; Stephanie Holley, University of Iowa Hospital & Clinics; Jorge Salinas, University of Iowa Hospitals and Clinics

Background: In December of 2019, the World Health Organization reported a novel coronavirus (severe acute respiratory coronavirus virus 2 [SARS-CoV-2]) causing severe respiratory illness originating in Wuhan, China. Since then, an increasing number of cases and the confirmation of human-to-human transmission has led to the need to develop a communication campaign at our institution. We describe the impact of the communication campaign on the number of calls received and describe patterns of calls during the early stages of our response to this emerging infection. Methods: The University of Iowa Hospitals & Clinics is an 811-bed academic medical center with >200 outpatient clinics. In response to the coronavirus disease 2019 (COVID-19) outbreak, we launched a communications campaign on January 17, 2020. Initial communications included email updates to staff and a dedicated COVID-19 webpage with up-to-date information. Subsequently, we developed an electronic screening tool to guide a risk assessment during patient check-in. The screening tool identifies travel to China in the past 14 days and the presence of symptoms defined as fever >37.7°C plus cough or difficulty breathing. The screening tool was activated on January 24, 2020. In addition, university staff contacted each student whose primary residence record included Hubei Province, China. Students were provided with medical contact information, signs and symptoms to monitor for, and a thermometer. Results: During the first 5 days of the campaign, 3 calls were related to COVID-19. The number of calls increased to 18 in the 5 days following the implementation of the electronic screening tool. Of the 21 calls received to date, 8 calls (38%) were generated due to the electronic travel screen, 4 calls (19%) were due to a positive coronavirus result in a multiplex respiratory panel, 4 calls (19%) were related to provider assessment only (without an electronic screening trigger), and 2 calls (10%) sought additional information following the viewing of the web-based communication campaign. Moreover, 3 calls (14%) were for people without travel history but with respiratory symptoms and contact with a person with recent travel to China. Among those reporting symptoms after travel to China, mean time since arrival to the United States was 2.7 days (range, 0–11 days). Conclusion: The COVID-19 outbreak is evolving, and providing up to date information is challenging. Implementing an electronic screening tool helped providers assess patients and direct questions to infection prevention professionals. Analyzing the types of calls received helped tailor messaging to frontline staff.

Funding: None
Disclosures: None

Doi:10.1017/ice.2020.578

Presentation Type: Late Breaker Poster
Effectiveness of Annual Flu Vaccination Until Week Four of the 2019–2020 Season
Pablo Chico-Sánchez, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante; Juan Gabriel Mora-Muriel, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Paula Gras-Valenti, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Natali J. Jimenez-Sepulveda, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Pablo Chico-Sánchez, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Isel Gomez-Sotero, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Natividad Algado-Sellés, Alicante Institute for Health and Biomedical Research (ISABIAL), Epidemiology Unit, Preventive Medicine Department, General University Hospital of Alicante, Alicante, Spain; Ana Esclapez-Martinez, Hospital General de Alicante; Sandra Canovas-Javega, Alicante Institute for Health and
Background: Annual flu vaccination is the most effective way to prevent the disease and its complications. Vaccine effectiveness (EV) varies from season to season, requiring annual re-evaluation. The objective of this study was to estimate the preliminary effectiveness of the influenza vaccine until epidemiological week 4 of the 2019–2020 season, in patients admitted to a tertiary-level hospital. Method: We conducted a case-control study at University General Hospital, Alicante, Spain, during the 2019–2020 season. We included all patients hospitalized with influenza confirmed by laboratory test (ie, PCR positive for influenza) during the period between epidemiological week 40 of 2019 and epidemiological week 4 of 2020. These were considered cases, and those with clinical suspicion of influenza and negative RT-PCR were considered controls. Vaccination coverage was calculated in cases and in controls, determining the odds ratio. We calculated the vaccine effectiveness (VE) and its 95% confidence interval using the following formula: $VE = (1 - \text{odds ratio}) \times 100$.

Result: We included 545 patients: 61 cases and 484 controls. The overall EV for influenza cases prevention was 40.7% (95% CI, -17.1 to 70.1), and for those >1 year of age, the overall EV was 56.9% (95% CI, 13.9–78.5).

Conclusion: The 2019–2020 Influenza vaccine was effective in preventing influenza cases in patients admitted up to week 4 of the 2019–2020 season. These results are preliminary and may vary; they should be re-evaluated at the end of the season.

Funding: None
Disclosures: None
Doi:10.1017/ice.2020.579

Presentation Type: Late Breaker Poster

Low Carriage Rates of Multidrug-Resistant Organisms in Prospective Stool Donors in a Large Fecal Microbiota Transplantation (FMT) Stool Bank

Amanda Zaman, OpenBiome; Taha Qazi, OpenBiome; Pooja Pai, OpenBiome; Tricia Peters, OpenBiome; Susie Nicolaysen, OpenBiome

Background: Fecal microbiota transplantation (FMT) has emerged as standard of care for Clostridioides difficile not responsive to antibiotic therapy. Rigorous screening of healthy donors is critical to patient safety. As part of routine donor evaluation for FMT, multidrug-resistant organism (MDRO) screening is performed to assess the presence of extended-spectrum β-lactamase–producing organisms (ESBLs), vancomycin-resistant enterococci (VRE), carbapenem-resistant Enterobacteriaceae (CRE), and methicillin-resistant Staphylococcus aureus (MRSA). Carriage rates of these organisms in a healthy, low-risk population are largely unknown. We report MDRO carriage rates among individuals screened for a stool donation program at a large-scale FMT stool bank.

Methods: Individuals were screened at a non-profit stool bank (OpenBiome, Cambridge, MA). Potential donors underwent in-person clinical assessment, including MDRO risk factors (eg, travel, occupation, healthcare exposure). If they met the clinical assessment criteria, laboratory testing, including MDROs, was performed. Once enrolled in the donor program, donors underwent repeated clinical and laboratory screening at 60-day intervals, with intermittent health checks throughout the donation period. Stool samples provided at 60-day intervals were screened for MDROS (ie, ESBL, CRE, VRE), and nasal swabs for MRSA were tested using culture-based methods. All stool samples tested for MDROS from prospective and enrolled donors were included.

Results: Between February 2017 and July 2019, 247 individuals were screened for MDROS. Overall, 11 samples (0.04%) tested positive for ESBL, MRSA, or VRE. No CRE carriers were identified. Also, 2 individuals tested positive twice for ESBL, resulting in 13 of 1,688 (0.77%) positive screens. International travel in the previous 12 months was reported by 6 of 11 MDRO carriers. Occupations typically associated with MDROs were not observed in carriers. Most of the MDRO-positive donors were students; however, students make up the majority of the stool donor cohort.

Conclusions: This study is the first to report background MDRO carriage rates in a population of otherwise healthy, low-risk donors.