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ABSTRACT

Fifty years of evolution in infection prevention and control programs have involved significant accomplishments related to clinical practices, methodologies, and technology. However, regulatory mandates, and resource and research limitations, coupled with emerging infection threats such as the COVID-19 pandemic, present considerable challenges for infection preventionists. This article provides guidance and recommendations in 14 key areas. These interventions should be considered for implementation by United States health care facilities in the near future.

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2022 marks the 50th anniversary of the founding of the Association for Professionals in Infection Control and Epidemiology (APIC), an organization devoted to advancing the science and practice of infection prevention and control. Over these years, the position and department title has transitioned from “Infection Control” to “Infection Prevention and Control” in order to better reflect the fundamental goals. In addition, infection prevention (IP) programs in health care facilities nation-wide have enhanced the culture of safety through modifications in each organization’s systems of care by assessing efficacy, and revising, standardizing, and monitoring clinical and ancillary practices. Coordinated efforts at reducing health care-associated infections (HAIs) have been determined to be effective to varying degrees when IP programs are provided with adequate resources and supported by the implementation of evidence-based strategies. Multifaceted HAI prevention programs have been proven to be cost-effective, a finding of vital importance in the present landscape of health care reimbursement and therefore, in the overall financial health of the institution. IP efforts have been facilitated by the application of such fundamental tools as core components, competency models, and implementation science which assist infection preventionists (IPs) to bridge gaps.
between organizational barriers to change and successful outcomes.

Although marked with many significant triumphs over our first 50 years, IP challenges posed in present-day health care settings have created new questions and compelled IPs to re-assess our approaches to reduce infection risk and to control known and emerging pathogens. This article, authored by experienced IPs, epidemiologists, and other content experts, provides guidance for addressing critical issues currently affecting IP programs with an emphasis on the implementation of innovative, cost-effective, and evidence-based interventions, engaging health care leaders and experts in clinical care in proven prevention measures, holding staff accountable, and adopting high-reliability principles.

**KEY IP CHALLENGES AND RECOMMENDATIONS FOR CHANGE:**

**IP Program Standardization**

The creation of reliable design of processes is a methodology that has been used in businesses including the airline industry and health care to produce better outcomes by reducing defects such as the ineffective use of time. According to the science of reliable design, the establishment of such a template IP program would support replication of best practices, avoidance of errors, and would ultimately optimize processes. The Infrastructure Report and the APIC IP Competency Model provide a starting point for the creation of this reliable design, by recommending standard priority areas of focus for IP professionals. However, neither provides scientific evidence to recommend a standard percentage of time dedicated to each priority area, in addition to recommendations regarding other critical components of an IP program including staffing levels, (interns, surveillance assistance, IP lead, manager, director, etc.) reporting structure and physician participation.

Currently, how IPs allocate their time each day varies widely among health care facilities and across the continuum of care, driven in part by regulations, the priorities of the IPs’ manager, and by the strengths and interests of the IP professional. The 2015 APIC Mega Report provides a listing of the average percent of time spent on key IP focus areas but does not address the variability from hospital to hospital.

Although reporting structure is key to the success of an IP program, there is currently wide variation regarding hierarchical reporting with some programs reporting to Quality departments, some to Nursing, some to patient safety, and a few directly to the C Suite or executive level. Physician partnership in IP programs have been proven to be important for optimal outcomes, although at present there is no national certification process to support standardization of the role and training of the physician in IP programs. And finally, IP department staffing is also lacking standardization. A recent peer-reviewed study concludes that the actual IP staffing level in US hospitals is anywhere between 31% and 66% above the current outdated benchmark of 0.5–1.0 IP per 100 occupied beds. Research is needed in order to define the ideal for each of these essential program components, which would together provide a reliable design for a best practice IP program.

Also, the pandemic has shown us the importance and need for more robust IP in different care settings, for example, nursing homes, behavioral health centers, and long-term acute care (LTAC) centers. In addition, ambulatory and outpatient IP is an ever important area requiring additional IP presence that is also lacking and frequently on the lowest priority for the average IP assigned to the acute care setting. The best practice IP program must, therefore, address IP across the continuum of care, spanning from outpatient to acute, to alternative care settings.

**Surveillance**

The National Healthcare Safety Network (NHSN) has served as the backbone of HAI surveillance with nearly 25,000 participating medical facilities. Standardized surveillance definitions and their application became all-encompassing to a point that for many IPs it defined their profession. As years passed, surveillance definitions became complicated, inconsistent from year to year, and subject to interpretative bias. In 2015, a systematic review was published highlighting this variability with 1 significant-conclusion, surveillance definitions need to be revised. NHSN definitions have been scrutinized over the years due to criteria that are perceived to not better reflect the clinical representation. Therefore, new partnerships between NHSN and the IP community, and other stakeholders will be important to evolve to future measures that are easily captured and more reflective of the quality of clinical care and processes. In addition, surveillance metrics became tied to public reporting measures, federal incentives, and penalties, as well as private insurer quality metrics that drove reimbursement. This phenomenon has increased pressure on IP programs to ensure prospective surveillance that is rooted in accuracy and speed.

Given the scope and complexity of surveillance activities, it has been estimated that 45% of IP time is consumed by this activity. According to Hebdon, most IP programs have not addressed this unsustainable time requirement by adopting an electronic solution which is also known as a data mining system. This has a significant impact on performance improvement activities and other IP program-related functions, particularly in a low-resourced department. Surveillance accuracy was also determined to be poor and according to a 2017 study respondents to case studies over 6 years showed 62.5% accuracy with a range of 16%-87%. Manual surveillance has proven itself to have inaccuracies and unsustainable time commitment which negatively impact IP programs exponentially.

Automated surveillance software has recently been addressing a need for accuracy and time efficiency. Multiple vendors are now competing to provide medical facilities with agile and user-friendly tools to achieve surveillance goals. However, caution must be taken when determining the appropriate surveillance system for IP programs. A systematic review published in 2020, concluded that estimated benefits of automated surveillance were still premature and that less than 20% of the studies were able to cite any efficiencies gained. These findings are directly tied to multiple factors that IP must consider. Many systems offer a wide array of solutions and integration within facility electronic health records (EHR). However, the ability to detect specific EHR information will render automated surveillance a success or a failure. The main factors that IP must consider include implementation, maintenance, and training costs, prerequisites for integrating communication between the surveillance system and the multitude of modules in an EHR, reporting and install capabilities, malleability of the software to tailor its focus and streamline data points and resourcing of ongoing maintenance. It is critical for current and future IPs to have a robust knowledge of electronic solutions and integrate the right platform taking these variables into consideration.

**HAND HYGIENE**

Hand hygiene (HH) compliance among health care providers (HCP) continues to be a significant challenge for many organizations. HH noncompliance occurring in health care institutions are multifactorial with the main causes reported in a Joint Commission assisted 8-hospital study including ineffective placement of dispensers or sinks, HH data compliance data not collected or reported accurately, lack of coaching, an issue not included as part of safety culture, ineffective education, and HCP distractions. IP programs struggle to
meet the demands HH noncompliance places on their organization through HH is undisputed as the number 1 method for reducing HAIs.

Programs designed to improve HH compliance involve leadership support and extensive system integration. A strategy advocated by the World Health Organization (WHO) is comprised of 5 elements which are all touted as essential and complementary: System Change (Build it), Education (Teach it), Monitoring and Feedback (Check it), Communication (Sell it), and Culture Change (Live it).24

Within these elements are issues that require further research. For example, although direct observation is the gold monitoring standard there remains a lack of standardization in the methodology, with direct or overt HH observations rates reported as much higher than indirect or covert observations.25 One study that determined baseline HH average compliance rates of 47.5% across 8 hospitals reported that observers were initially unable to determine the causes of non-compliance simply by visually observing practice due to behavioral influences.23

Observation methods have their own inherent problems as evidenced by the well-documented Hawthorne effect. Education of the observer can vary however a standardized method of “train-the-trainer” has been demonstrated as an effective and efficient method for sharing knowledge. This method along with simulation was used in a multicountry 3-day course based on the WHO multimodal HH improvement strategy resulting in a statistically significant improvement in participant knowledge.26 Furthermore, sustainability was achieved after evaluation 2 years later. Though proven effective, simulation in a lab environment is not always available and not widely used. Regardless of the improvement methodology used or the strength of individual components, multimodal strategies have proven to be the most successful.27-29

Recent interest has been given to supplemental methods for monitoring HH compliance. To address this issue, an automated hand hygiene surveillance system (AHHSS) has been developed. A recent study examining the impact of an AHHSS installed on 4 patient units in 1 hospital, reported unit-specific baseline quarterly averages of 66%-95% by direct observation with postintervention of 77%-90%.30 It is important to note improvements in HH compliance rates were not observed on all units in this study using the AHHSS, while those units that did increase compliance rates improved after implementation of additional interventions, including the Toyota Kata performance improvement methodology. The methodology attempts to change behavior using a defined step-by-step process. Reductions in non-Clostridioides difficile HAIs occurred but were not statistically significant.

A recent review of 73 studies using AHHSS indicated that such systems “…face issues of accuracy, data integration, privacy and confidentiality, usability, and associated costs as well as infrastructure improvements.”31 Such limitations, including lack of standardization, reduce accuracy, thus circumventing the intended process. Data validation and reporting can be problematic when the system isn’t supported internally by executives, therefore, placing an additional burden on IP responsibilities. Improvements in automated systems include integration with cloud technology, therefore, decreasing the need for facility information technology (IT) department support.32

Additional influences in monitoring are in part due to regulatory and nonprofit groups. In 2019, The Leapfrog Group, a nonprofit consumer “watchdog” organization, added a new HH standard to their hospital and ambulatory surgery center (ASC) surveys.33 Adherence to the standard by participating hospitals is measured by compliance in 4 of 5 domains, which include training and education, infrastructure, culture, and adherence to a defined number of observations which may be assisted by the use of an electronic monitoring system. The consideration for implementation of this monitoring methodology should take into account various issues including the impact on an organization’s finances, training capabilities, and staffing levels as the program requires meeting specific standards such as a minimum number of observations covering all units, shifts, and days of the week, feedback on HH compliance data to individuals who have contact with patients or with items to be used by patients while conducting a direct overview of observers to ensure that the process is uniform and consistent. Hospitals that elect not to participate in Leapfrog, should realize that publicly reported information will instead be used in their published reports.

In conclusion, recommendations to address future HH challenges include the need to conduct further studies addressing the influence of human behavior and system culture on HH practices, new approaches in training and education, investigation of the most effective monitoring strategies, as well as determining the best processes for enhancing communication of monitoring results, and the linking of said results to health care personnel performance reviews.

ENVIRONMENTAL CONTAMINATION

The sources and routes of pathogen transmission in health care settings have been well researched.34 The most significant source is the patient who is colonized or infected and may shed organisms from body sites, bedding, gowns, and contaminate nearby environmental surfaces and portable equipment used in their care. The frequency of pathogen transfer from patients and their environment to HCP hands, gloves, and gowns have been demonstrated to be 33%, 30%, and 10%, respectively. Also concerning is the finding that the hands of HCPs are just as likely to be contaminated by touching an environmental surface as by direct contact with a patient.35

In acute care settings, the patient environment is “…defined as the area inside the curtain, including equipment, medical devices, furniture, telephone, personal belongings, and the bathroom.”36 Multiple studies indicate that the patient environment plays an important role in the transmission of many pathogens of concern in health care including methicillin-resistant Staphylococcus aureus (MRSA), vancomycin-resistant Enterococcus (VRE), gram-negative organisms such as Pseudomonas aeruginosa, and C difficile.37-39 These and many other pathogens have been known to persist on environmental surfaces from hours to days, and in the case of spore-forming organisms, months.30 Studies have demonstrated that when a patient is colonized or infected with organisms including MRSA, VRE, and Acinetobacter, the risk of acquisition of these organisms by a newly admitted patient to the same room is increased.41 In a large study of 10,289 HAIs occurring over 7 years in 4 hospitals,42 the risk of acquiring an HAI was nearly 6-fold when a prior bed occupant was colonized or infected with a pathogen.

Evidence suggests that adequate cleaning of patient rooms is often insufficient. In 2 large studies43,44 (23 and 16 hospitals, respectively) researchers used a fluorescent solution applied on room surfaces located in a patient’s immediate environment and a hand-held ultraviolet light device to assess the adequacy of cleaning. The studies concluded that only 49% and 57.1% of the surfaces were adequately cleaned.45

Several important points should be considered when determining how best to improve patient room cleaning and disinfection in a health care facility. First, studies examining the efficacy of education of environmental services (EVS) staff, key patient safety persons, have demonstrated minimal improvement, with 5%-30% of surfaces remaining potentially contaminated.45 Secondly, the effect of improved environmental cleaning and disinfection on patient acquisition of pathogens, appears to be variable. One review of the literature reported a reduction in the rates of MRSA, VRE, and C difficile infection (CDI) from 0% to 49% after improvement in cleaning practices, with 1 study demonstrating an 83% reduction in VRE bacteremia.44 Thirdly, monitoring of cleaning practices, which include
traditional visual inspection, microbiological sampling, and nonmicrobiological testing such as fluorescent markers used as surrogates for residual contamination and quantification of adenosine triphosphate (ATP) levels to determine persistence of organic material, lack universal acceptable standard levels of residual contamination.46 Fourth, concentration solely on cleaning “high-touch” surfaces often prevents thorough and complete room decontamination.57

Hospitals are investigating automated supplemental environmental disinfection processes with increasing frequency in response to failures associated with human factors such as inadequate or overlooked manual cleaning of objects, lack of proper supervision and monitoring, lack of resources, and low levels of hand hygiene.

Automated supplemental disinfection technology can currently be categorized as mobile devices that are used for terminal room disinfection and technologies that provide continuous surface disinfection. The principle mobile technologies represented in peer-reviewed studies incorporate the automated emission of chemical vapors, aerosols or ultraviolet (UV), and hand-operated electrostatic sprayers. The most comprehensive clinical trial to assess a UV-C “no-touch” technology is the Benefits of Enhanced Terminal Room Disinfection (BETR-D) study, the results of which were published in 2017.48 This multihospital study concluded that the addition of a UV-C device to the standard manual process of terminal cleaning and disinfection decreased patient acquisition of a target organism by approximately 10% to 30%, suggesting that the environment is responsible for a significant portion of antibiotic-resistant organism (ARO) acquisition.49 Factors such as organic load, pathogen type, intensity, surface types, distance of the surface from the UV device, placement of the device in the room, exposure time, room size and configuration, and air movements, all contribute to the relative efficacy.49

Studies using hydrogen peroxide (HP) decontaminating systems report 86%–100% reduction in MRSA, VRE, and multidrug-resistant gram-negative bacteria on contaminated surfaces in hospital rooms,50,51 with reported reductions in 10 published studies,52 although an analysis of 5 studies using HP did not find a statistical reduction in CDI rates.53 Perhaps the most significant limitation in the use of HP robot technology is the total procedural time required (approximately 1.5–8.0 hours) due, in part, to room setup which requires occlusion of doorways and vents. Additionally, all these automated supplemental systems cannot be used in the presence of persons in the room and require dedicated staff to perform the function.

Another promising supplemental disinfection technology is the electrostatic sprayer which can be used to apply any sporidical disinfectant directly on equipment and other environmental surfaces in various health care areas. A recent study using this type of device demonstrated a >6 log reduction against C difficile and a SARS-CoV-2 surrogate.54

Understanding the operating parameters of mobile environmental disinfecting technology has led science to the next evolutionary level in decontamination concepts. Research engineers have designed continuous disinfection technologies incorporated in room designs that emit either visible light at specific wavelengths or produce disinfecting emissions at levels that are not toxic to humans. “Blue light” in the spectrum of 400-470 nm has demonstrated a consistent 1- to 2-log reduction of surface bacteria.55 A novel photocatalytic conversion device inserted into the ducts of an HVAC system, uses a multi-wavelength ultraviolet light to illuminate target surfaces. The device is comprised of a honeycomb matrix treated with a photocatalytic coating of titanium oxide (TiO2) and other reactive metals. The device converts H2O and O2 in the air into hydroxyl radicals and HP which work to reduce bioburden on surfaces. In a published ICU-based study, a 95% reduction in average microbial burden, 81% reduction in the prevalence of MRSA, and a 54% reduction were reported in HAIs over a 4-month period.56 Another study in 5 medical units using HVAC-installed devices that emit dry HP on a continuous basis demonstrated an overall 96.5% microbial reduction for all combined surfaces.57 Regardless of the available technology, IPs need to acquire evidence-based information on the device’s effectiveness in reducing contaminating microorganisms on health care environmental surfaces.

ANTIBIOTIC-RESISTANT ORGANISMS

Antibiotic-resistant organisms (AROs) are associated with 4.95 million deaths worldwide.58 On December 7, 2021, the Centers for Disease Control and Prevention (CDC) announced that it has awarded $22 million to nearly 30 organizations around the world to combat antimicrobial resistance and other health care threats through the establishment of 2 new networks—the Global Action in Healthcare Network (GAHIN) and the Global AR Laboratory and Response Network (Global AR Lab & Response Network). ARO control is an international priority that requires all health care facilities and agencies to assume responsibility to prevent transmission. Such initiatives are driven by the fact that infections with AROs are associated with increased lengths of stay, costs, and, more importantly, mortality.59

IPs in coordination with other key health care personnel should review core information regarding AROs in order to determine facility policy on such issues as isolation, appropriate therapy, and antibiotic stewardship; ensure accurate microbiology test results using the latest Clinical and Laboratory Standards Institute (CLSI) determination of minimum inhibitory concentration (MIC) antibiotic breakpoints60; prioritize pathogens using WHO document that categorizes (Critical, High, Medium) resistant bacteria based on treatment options and potential for spread; for example, carbapenem-resistant A baumannii, Paeruginosa, Enterobacteriaceae61; or CDC’s phenotype definitions62 and Antibiotic Resistance Threats report.63 In addition, AROs are detailed in the NHSN MultiDrug-Resistant Organism & C difficile Infection module which addresses AROs associated with HAIs.64 “Enterobacteriales” was adopted as a new taxonomy and scientific order in 2020. The microbiologic family of Enterobacteriales includes Carbapenem-resistant Enterobacteriaceae (CRE) organisms. CRE bacteria pose a serious public health threat and are associated with high mortality due in part to limited antibiotic therapy. The principal CRE organisms include Escherichia coli, Klebsiella pneumoniae, and Enterobacter spp.65

Organisms with newly identified antibiotic-resistant mechanisms continue to be identified throughout the world. Access to electronic communication across the continuum of health, for example, ambulatory to acute to long term care, by key health care personnel on emerging AROs is, therefore vital. IPs should assist in developing a procedural and administrative policy for transferring adult and pediatric patients identified with AROs, utilizing the CDC Inter-Facility Transfer guidelines for communication between hospitals, including out of state/territory facilities.57

One recent report identified an emerging “superbug” that was cultured from a patient in a Florida long-term care facility. The event was associated with a P aeruginosa isolate that was producing VIM (Verona integron-encoded metallo-beta-lactamate), an enzyme that makes it resistant to a class of antibiotics normally highly effective against antibiotic-resistant bacteria. This was Florida’s first reported case, classified as VIM-CRPA, and immediate containment was necessary.58

It is critical to identify the specific pathogen in serious, life-threatening infections, especially for situations that are likely to require prolonged therapy. Due to the inherent time associated with microbiologic culturing methods, clinicians often use a standard empiric
antimicrobial therapy approach that often is composed of a regimen of 2 or more antibiotics administered at a time. Adverse outcomes associated with overuse of antimicrobials includes emergence of AROs and *Clostridioides difficile*. One of the top concerns for antibiotic stewardship endeavors is the delayed receipt of results of organism identification (ID) and antimicrobial susceptibility test (AST). Generally, such testing can take a microbiology laboratory from 48 to 72 hours, and in some circumstances a longer period due to the growth characteristics of the particular organism. Institutions should investigate the use of rapid diagnostic technology that reduces both ID (<1 hour) and AST (< 8 hours). Advancements in rapid diagnostics have shortened the time to results from days to hours and have had positive effects when paired with a transition from the use of broad-spectrum antimicrobial regimens to pathogen-specific therapy on clinical outcomes and on efforts to combat antimicrobial resistance. Further advances, including the provision of point-of-care testing, along with increased funding and government initiatives, will require review in order to further close the gap between current culturing methods and rapid diagnostic results.

**DECOLONIZING PATIENTS**

Colonization with health-care-associated pathogens such as *S. aureus*, enterococci, and Gram-negative organisms, is associated with increased risk of infection. Decolonization is an evidence-based intervention whose goal is to reduce or eliminate the bacterial bioburden in order to reduce the risk of infection. Most decolonization interventions consist of a nasal product plus bathing with chlorhexidine gluconate (CHG). This section will focus on decolonization to reduce HAIs in the intensive care unit (ICU), noncritical care, and surgery.

In 2013, 3 randomly selected cluster trials on the topic of CHG bathing among ICU patients were published. One cluster-crossover study reported that daily 2% CHG cloth bathing in the ICU resulted in a 23% reduction of vancomycin-resistant *Enterococcus* (VRE) and methicillin-resistant *S. aureus* (MRSA) acquisition and a 28% reduction in bloodstream infections (BSIs). In another study of 2014 by Milstone et al found a significant association between 2% CHG cloth bathing and a decline in BSIs compared with standard bathing. Another trial, called the REDUCE MRSA study, used a cluster-randomly selected methodology in 43 hospitals (including 74 adult ICUs) to evaluate 3 MRSA prevention interventions: the first cluster implemented MRSA screening and isolation, the second cluster included screening, isolation, and decolonization of MRSA carriers with CHG bathing and nasal mupirocin (ie, targeted decolonization), and the ICUs in the third cluster did not screen any patients but instead, all patients were decolonized with CHG cloth bathing and nasal mupirocin (ie, universal decolonization). Universal decolonization was found to be associated with the greatest decrease in all-cause BSIs (44%; *P < .001) and rates of MRSA clinical cultures (37%; *P < .01). In a secondary analysis, CHG bathing was also shown to reduce blood culture contamination by 45% (*P < .02), and to be the most cost-effective. In 2014, the Compendium on Strategies to Prevent Central Line–Associated Bloodstream Infections (CLABSI) in Acute Care Hospitals recommended bathing for ICU patients over 2 months of age with a CHG preparation daily (quality of evidence: I).

Although the patient-specific risk is highest in ICUs, an equal number of CLABSI occur in hospital units outside of the ICU. Questions remain about the use of ICU-proven strategies in noncritical care units. The ABATE Infection (active bathing to eliminate infection) trial, a 2-arm cluster randomly selected trial is the first and only large-scale cluster-randomly selected trial designed to evaluate whether universal CHG bathing for all patients plus targeted mupirocin for MRSA carriers in noncritical-care units reduces multidrug-resistant organisms and all-cause BSIs. In the initial analysis there was no difference in MRSA and VRE clinical cultures or BSIs, but the posthoc analysis of the trial found that non-ICU patients with vascular access devices (central lines or midlines) had a significant 37% reduction in MRSA and VRE and a significant 32% reduction in all-cause BSI. Based on this study, many facilities implemented CHG bathing with or without mupirocin in patients in noncritical care units with vascular access devices. In 2019 the CDC recommended CHG plus an intranasal antibiotic/antiseptic product in patients with a central line or midline as a supplemental strategy in noncritical care.

Surgical site infections (SSI) remain one of the most common and expensive HAIs, with *S. aureus* among the most frequent etiologic pathogens. Studies confirm *S. aureus* carriage increases the risk of *S. aureus* SSIs. There is strong evidence that nasal and skin decolonization prior to cardiac and orthopedic surgery is effective in reducing SSIs caused by MSSA or MRSA. A recent study implemented a bundled intervention in 20 hospitals to prevent complex *S. aureus* SSIs after cardiac surgery and hip and knee arthroplasty. The bundle included preoperative screening for MRSA and MSSA nasal colonization, CHG bathing for all patients, nasal mupirocin decolonization for *S. aureus* carriers, and both vancomycin and cefazolin perioperative antibiotic prophylaxis for MRSA carriers. The mean rate of complex *S. aureus* SSIs significantly decreased by 42% from 36% infections per 10,000 operations during the baseline period to 21 infections per 10,000 operations during the intervention period (rate ratio 0.58; 95% CI, 0.37−0.92).

Moving forward, challenges related to decolonization include expansion of these strategies for additional surgical patient populations, as well as nonsurgical invasive procedures. For example, interventional radiology patients receiving implants such as cardiac implantable electronic devices (CIED) have been associated with significant numbers of postimplant infections, often caused by *S. aureus*. For patients known to be colonized with MRSA or MSSA prior to a CIED procedure, a multigroup British Working Party guideline recommends the use of nasal and topical antimicrobial agents preprocedure in order to suppress carriage. Although mupirocin remains the best-studied nasal agent, there is some evidence on the use of safe and effective alternative agents such as povidone-iodine (PI), as well as photodynamic therapy (PDT). Although PI may be a useful alternative decolonizing agent for the prevention of *S. aureus* infections, additional clinical data are required to further confirm the effectiveness of PI in preventing *S. aureus* infections. There is only 1 alcohol-based study that was conducted at a single-center using a quasi-experimental design with historical controls. Lastly, photodynamic therapy is another promising approach for topical MRSA decolonization, but larger clinical trials are needed to evaluate different nasal decolonization protocols (including determining the optimal sensitizer) using clinically significant infection as the outcome.

In addition, there is a need to determine if the widespread use of CHG-based products promotes reduced CHG activity. Testing for CHG susceptibility is currently not standardized and the clinical impact of reduced chlorhexidine susceptibility in bacteria is unknown and not yet well-defined.

**DECREASING CONTACT PRECAUTIONS**

The recommended use of contact precautions (CPs) in acute care as a proposed strategy for preventing ARO transmission has been a mainstay for half a century with successes documented particularly when the organism is emerging rather than endemic. However, recommended duration of CPs as a strategy for prevention in hospitals remains unsettled. Determining when CPs may be discontinued and thus safely reduce risk of transmission is influenced by 3 key factors. First, involves the organisms themselves: which antibiotic resistant bacteria trigger an isolatable condition, and whether the patient...
is deemed colonized or infected with the particular ARO. Second, involves what type of isolation is applied. In nonhospital settings such as ambulatory clinics, dialysis centers, and home care settings recommendations for handling patients with AROs include use of standard precautions, while skilled nursing facilities deploy enhanced barrier precautions (use of Personal Protective Equipment [PPE], but without room restriction) for high contact care activities and CP when wounds or body fluids are unable to be contained or when ongoing transmission is suspected or confirmed. The third factor involves whether a health care facility implements advances in health care strategies to reduce transmission risk or bioburden. Single patient rooms, alcohol-based hand sanitizers, hand hygiene monitoring protocols, supplemental environmental disinfection technologies (eg, ultraviolet/continuously active disinfection), dedicated/disposable medical equipment/devices, and patient decolonization with CHG bathing, nares decolonization and/or antiseptic oral rinse are some examples of interventions that may reduce the risk of transmission. Currently, these strategies are neither uniformly nor equitably adopted across (or even within) health care facilities.

There has been a growing body of evidence indicating the development of potential unintended consequences (patient dissatisfaction, safety, and worsening noninfectious outcomes) with the use of CPs for AROs in a 1-size-fits-all approach. A contemporary, popular approach has been to reserve CP for patients infected (vs colonized) with MRSA or VRE presumably under the hypothesis that colonization may correlate with a reduced organism density and therefore, reduced risk of transmission, the evidence in support of this specific hypothesis is not very strong. Meanwhile, there have been many reports of organizations suspending CPs without an increase in MRSA or VRE infection rates, often in the context of the aforementioned health care advances. A recent SHEA expert guidance document regarding the duration of CPs acknowledges that “no universally recommended approach exists for making decisions regarding CP duration or discontinuation for any epidemiologically significant organism”. The same document recommends that organizations develop ARO management policies including the use and duration of CPs in the context of their risks, priorities, and resources and that these policies be organism-specific, evaluated, and modifiable as/when/if the epidemiology changes. What is lacking and needed is greater consensus on ARO transmission risk reduction strategies in the context of utilizing CPs judiciously for resource conservation and patient safety.

As mentioned, the list of AROs has expanded over the last 50 years and few would argue that further diversification is unexpected or that all AROs are equally transmissible. While suspending CPs for extended spectrum beta-lactamase (ESBL) Gram-negative organisms without an increase in ESBL infections has been reported, this has not been as commonly reported with carbapenem resistant Enterobacteriaceae (CRE) though modeling in subacute settings suggest decolonization and screening for clearance may yield better results and no evidence to date with the recently emerged Candida auris. Per the HCIPAC meeting minutes of August 19, 2021, pertaining to the Isolation Precautions Guideline Workgroup and forthcoming efforts to provide further guidance, “Most of this will focus on the framing of transmission processes as opposed to the gritty isolation practices. Airborne and contact isolation practices should not change.” This suggests that the way forward is a re-evaluation of the transmission modalities of AROs at the organism level (eg, MRSA vs CRE) and hopefully in the context of the aforementioned, disproportionally deployed advances in practice and technology. Decision support tools have been successfully utilized for infection prevention initiatives for years and are well suited for this challenge. Such tools could be developed and be evidence-based, algorithmic in nature and incorporate organism prevalence, health care setting and available control strategies including, but not exclusive to contact precautions.

**Diagnostic Stewardship**

Diagnostic Stewardship (DS) involves the ordering, collection, and timely reporting of diagnostic tests to improve treatment of infectious conditions. Suboptimal culturing practices can adversely impact patient safety and the quality of care in a wide variety of ways including: reporting of asymptomatic bacteriuria misidentified as urinary tract infection (UTI) and blood culture contamination mistaken for bloodstream infection resulting in a wrong diagnosis, increasing the numbers of AROs and *C. difficile* as a result of overuse of antimicrobials, adverse reactions, and unwarranted financial expenditures stemming from inappropriate specimen testing and drug treatment, as well as the potential for over-reporting of HAIs such as catheter-associated urinary tract infections (CAUTIs). Recent reviews and surveys examining urine and blood culturing practices have underscored this issue and have identified potential underlying causes.

In order for health care organizations to optimize clinical and epidemiological outcomes related to urine and blood culturing, the CDC has recently identified key needs that will be essential to address, including the need for drivers to incentivize or require health care facilities to invest in DS, the need for meaningful measures development, and perhaps most important, the need for leadership support for implementation. In addition, system-wide management programs must be directed by standardized practice guidelines, jointly written by IPs, microbiologists, and other key stakeholders, providing guidance in the preanalytic phase, that is, pathways for selecting the appropriate test according to patient’s syndrome, methods for obtaining optimal collection of clinical specimens, and interpreting microbiology results. In addition, efforts must be focused on education of nurses to understand the “why” behind proper culturing techniques, and of physicians on appropriate ordering and interpretation of culture results. Additionally, in order to ensure the integration of all areas of infection management (diagnosis, treatment, and prevention) a unified stewardship strategy is recommended such as reflected in the AID (antimicrobial, IP, and DS) model. The basis of this model is the understanding that outcomes improve when providers understand and cooperate in implementing a system that promotes the appropriate test, the right interpretation of a test result, selection of the appropriate antimicrobial, and administration at the right time. DS targets all relevant patient populations, for example, ensuring that established protocols capture both catheterized and noncatheterized patients relative to urine culture collection. DS also targets all relevant patient settings, for example, in specimen-intensive areas such as Emergency Departments and focuses on prevention of overutilization of blood and urine cultures. This can be accomplished in part by reviewing and revising order sets in electronic medical records (EMR) to reflect essential modifications such as urine culture reflex testing, a practice that reduces unwarranted urine cultures while continuing to provide accurate clinical findings, and integration of electronic medical records to improve compliance with ordering protocols. An important aspect of DS is ongoing evaluation of novel devices, such as those designed to reduce blood culture contamination.

DS interventions should also be applied to *C. difficile* testing to ensure more appropriate testing, facilitate prompt isolation and optimize clinical treatment. DS interventions related to *C. difficile* testing should include: provision of education of nurses on the appropriate documentation of patient bowel movements and use of laxatives, implementation of *C. difficile* order sets and EMR best practice alerts to assist providers in avoiding testing when patients do not meet current recommendations (recent laxative use, < 3 unformed stools in 24 hours); optimizing testing strategies to distinguish between toxigenic and nontoxigenic strains of *C. difficile*. Specifically, the nucleic acid amplification test (NAAT) is a widely adopted laboratory diagnostic tool for *C. difficile*, due to the high sensitivity and rapid
and long-term care following discharge, increased direct care costs, day readmission rates, a greater need for intensive care management and a longer term care stay. Therefore, best practice strategies should incorporate a 2-step reflex testing algorithm that utilizes advanced laboratory screening methodologies.

**HAI SURVEILLANCE AND PREVENTION: BSI**

Several important interventions are needed to achieve improvements in surveillance and prevention of BSI including expansion of the definition, improved documentation of clinical findings to prevent missed events and over-diagnosis, identification of provider knowledge gaps followed by education to support the introduction of an enhanced central line maintenance bundle and evaluation of novel technologies including advanced antimicrobial IV dressings that may extend the effective antimicrobial period over the entire recommended duration of the dressing. Future endeavors should include in vivo studies that test the effectiveness of advanced dressings in reducing BSIs.

Device-associated BSIs account for significant numbers of HAIs in United States health care facilities. However, the full extent of the problem is unknown because surveillance and federal reporting is currently mandated only for CLABSI events. Published studies indicate that considerable numbers of BSIs occur as a complication after placement of arterial, hemodialysis, midline, and peripheral intravenous catheters (PIV). This evidence supports a need to implement a more comprehensive prevention strategy that addresses all types of intravenous catheters, which could be captured using a concept known as Hospital Onset Bacteremia (HOB). This expanded prevention strategy should extend to settings outside of hospitals, such as home infusion, where caregiver practices are a notable infection factor. It should also reflect recent learnings from the COVID-19 pandemic based on analysis of the causes for the increase in BSI during this period.

**HAI SURVEILLANCE AND PREVENTION: NONVENTILATOR HOSPITAL ACQUIRED PNEUMONIA**

Nonventilator hospital acquired pneumonia (NV-HAP) is the most common hospital acquired infection in the US affecting 1% of hospitalized patients with a crude mortality rate of 15%–30%. NV-HAP is associated with a significantly longer length of stay, high 30-day readmission rates, a greater need for intensive care management and long-term care following discharge, increased direct care costs, and perhaps most importantly, it is a frequent source of sepsis.

As noted in a September 2021 Joint Commission Safety Briefing. NV-HAP is a substantial threat to patient safety and a large driver of health care cost, yet it is not formally tracked or reported to the National Database of Nursing Quality Indicators or the CDC NHSN nor does it impact the Centers for Medicare and Medicaid Services pay-for-reporting or performance programs. NV-HAP surveillance is not currently standardized, and risk stratification is not reliable, therefore, consideration should be given to application of universal prevention guidelines throughout every patient’s hospital or long-term care stay. The most promising evidence-based prevention measures include primary source control (eg, oral care), early and frequent ambulation, head of a bed elevation, and aspiration precautions. Additional randomly selected controlled trials evaluating the impact of individual interventions (rather than a “bundle”) are warranted. However, it is equally important to help health care facilities improve quality and safety now and develop practical prevention strategies to translate existing evidence into practice to improve the health and safety of patients.

NV-HAP prevention recommendations include engaging facility leadership and interdisciplinary team members in discussions about the importance of NV-HAP surveillance and prevention, considering provision of adequate resources, and facilitating implementation of evidence-based interventions, nursing documentation templates on NV-HAP prevention that capture process measures and an electronic tracking system to monitor NV-HAP. Other measures include mentoring influential nurses and nursing assistants in direct care positions to serve in leadership roles (including those who are not in official management positions) to champion prevention efforts within the daily workflow, raising the profile of oral care as a new standard and an essential infection prevention practice rather than an optional comfort measure, and supporting a safety culture by facilitating regular reminders and communication, particularly in areas with high staff turnover. Lastly, maintaining staff education and competencies in fundamental nursing care (eg, oral care, mobility), creating an efficient process to stock needed supplies and equipment at the bedside, empowering patients to ask for assistance with oral care, feeding, and mobility when they need it and to report when they completed their care independently so it can be documented, and establishing the prevention plan as a standard of practice facility wide.

**HAI SURVEILLANCE AND PREVENTION: VENTILATOR-ASSOCIATED EVENTS**

For many years IP programs conducted surveillance for ventilator-associated pneumonia (VAP) using an NHSN definition that included several clinical components that were often subjective and difficult to interpret. In 2013, the VAP definition changed to a tiered system represented by ventilator associated events (VAE) subcategorized by objective criteria for infection-related ventilator associated condition and then more specifically by possible and probable VAP. The VAE definition eliminates subjectivity by using clearly defined criteria often contained in medical records and therefore, facilitates the automated collection of data. Additionally, VAE definitions broadened the focus of surveillance from pneumonia alone to the syndrome of nosocomial complications in ventilated patients.

However, the current tiered VAE definition lacks sensitivity, potentially resulting in underreporting of true events. An initial goal of the VAE definition was to identify all causes of respiratory deterioration as well as broadening the safety surveillance for patients on a ventilator including pneumonia, atelectasis, fluid overload and acute respiratory distress syndrome. A meta-analysis by Fan et al determined the pooled sensitivity of VAE for traditionally defined VAP was 42%. Additionally, the pooled positive predictive value of VAE for traditionallydefined VAP was 23%. Another study by Zosa et al identified that the VAE definition grossly underestimates the clinical diagnosis of VAP and reports less than a third of the patients treated for VAP. The fact that VAE surveillance misses many traditionally defined VAPs would appear to undermine its claim to detect clinically important respiratory complications of care. Enhancing surveillance and intervention protocols are supported by the severe consequences of such infections: VAPs have a significant attributable mortality (4.6%) along with increases in morbidity, hospital length of stay (LOS) and cost of care.

When the VAE definition was implemented, its purpose was not to be a method to specifically identify VAP but rather to broaden safety surveillance for a variety of ventilator-associated events. Updated definitions will enhance consistency, accuracy, and reproducibility of surveillance information. A collaborative effort among the IP community and NHSN with the aim to enhance a balance between epidemiological and clinical occurrences based on expert review should increase infection sensitivity, leading to broader and more effective prevention efforts, as well as raising event agreement among respiratory therapists, intensivists, and IPs.
HAI SURVEILLANCE AND PREVENTION: URINARY TRACT INFECTIONS

UTIs represent a common diagnosis for patients in the ambulatory, acute, and long-term care settings. Contrary to the evaluation of patients for bacteremia where blood is normally sterile, colonization of the urine increases with age and is considered physiologic. Asymptomatic bacteriuria (AB) is prevalent with older age, diabetes, impaired voiding, and urinary catheterization. A widespread practice is to obtain a urine specimen for analysis for both infectious and noninfectious conditions. The urinalysis, commonly used to test for noninfectious conditions, includes metabolic, inflammatory, and renal components. Some elements indicating bacteriuria (e.g., nitrites), and others reflecting inflammation (e.g., leukocyte esterase) are often mistakenly equated with infection triggering the order of a urine culture. Almost half of patients admitted to hospitals have a urinalysis performed and a quarter will have a urine culture done.

Obtaining a urine culture without an evaluation of the patient’s clinical condition and the likelihood of symptomatic urinary tract infection leads to misdiagnosis, inappropriate antimicrobial use, and increase the risk for antimicrobial resistance. In addition, variation in testing occurs based on setting. In the ambulatory arena, often a UTI is diagnosed based on clinical grounds, and urine culture may not be obtained for the first episode of cystitis. The elderly, especially those that reside in long-term care facilities, have a high prevalence for AB, and are prone for misdiagnosis of UTI when clinical parameters are not included in the decision making to test. On the other hand, the clinically defined catheter-associated UTI (CAUTI) is reached by excluding other diagnoses.

The main challenge in addressing UTI relies on the importance of incorporating clinical signs and symptoms as cardinal factors, with laboratory support, to reach the diagnosis. DS starts with a focus on the preanalytic component which is heavily dependent on clinician knowledge and competencies. Understanding the pretest probability before obtaining a urinalysis or culture, and incorporating the patient’s symptoms in the decision to order the test will reduce the likelihood of detecting patients with AB. Specimen contamination is minimized by obtaining a clean midstream sample for those noncatheterized and a fresh specimen from sampling port in those catheterized. The analytic phase is where the laboratory can actively support unnecessary testing by limiting urine cultures to those ordered for urinary symptoms. The absence of pyuria on a urinalysis is predictive of not having a symptomatic UTI. Different algorithms have been implemented in hospitals to reduce the unnecessary use of urine cultures based on the quantitative evaluation of pyuria on a urinalysis. However, the cutoff for pyuria to adequately exclude asymptomatic bacteriuria is still not clear, and it is uncertain if using the urinalysis to decide whether a culture is warranted discriminates adequately between those with or without a clinical infection. Reflex urine cultures should only be performed for patients with suspected symptomatic UTI, and not only based on the urinalysis results. Exclusions include pregnancy and urologic procedures associated with mucosal trauma. In the postanalytic phase, framing the culture results and the corresponding antimicrobial susceptibilities in context will help enhance the clinician’s decision making.

DS is a prerequisite to achieving successful antimicrobial stewardship outcomes. We need to promote standardized processes to improve the diagnostic testing of patients with suspected UTI. First, clinicians need to eliminate the old habits of “panculturing” to a more reflective behavior where testing the urine is based on suspicion of infection. Second, the use of “urinalysis with reflex to culture” requires more research to standardize the parameters that would trigger a culture. Third, future process measures to evaluate the variation in utilization of urinalysis and urine culture may help understand the under- and over-use of diagnostic tools. Fourth, in the hospital setting, more inclusive metrics such as the standardized utilization ratio (SUR) are useful tools to evaluate both infectious and noninfectious device risk. From an outcome measure perspective, the current surveillance definition, especially for CAUTIs, suffers from a low positive predictive value for true instances of disease, leading to skepticism from clinicians. More objective, electronically captured data, such as combining patient-level catheter-associated bacteriuria and antimicrobial use may help better understand the stewardship and infection prevention problem. Finally, in the ambulatory or long-term settings, evaluating the incidence of UTI and associated antimicrobial treatment of populations at risk for misdiagnosis (e.g., diabetics, elderly) may provide valuable information on potential future areas of focus.

HAI SURVEILLANCE AND PREVENTION: SURGICAL SITE INFECTIONS

One well studied approach to improving surgical patient outcomes is the use of care bundles. The care bundle concept was first introduced into the surgical arena by the Institute for Healthcare Improvement (IHI) in 2006 as a strategy to reduce the risk of SSI and was based upon 4 core measures: (1) timely and appropriate antimicrobial prophylaxis, (2) appropriate hair removal, (3) normothermia, and (4) glycemic control (focusing at that time on vascular and cardiac patients). While each of these measures had a relevant evidence base, together they were not sufficiently robust to lead to a sustainable improvement in patient outcome across the surgical spectrum.

However, over time with the addition of key elements, the surgical care bundle (SCB) has emerged as an accepted method of packaging the best, evidence-based measures into routine care for all surgical patients to prevent SSIs. Most of the selective evidence-based measures listed below have been identified by systematic review and meta-analyses, providing moderate to high clinical evidence (1A) and recommended in national, international and societal guidelines. Currently, there is no consensus as to the maximal number of measures within an evidence-based surgical care bundle. However, consensus suggests that individual care bundle components should attempt to address the myriad of patient risk factors present at the time of surgery.

One bundle measure supported by several evidence-based studies in selective surgical disciplines is a standardized preadmission shower/cleansing strategy using CHG to reduce the microbial burden on the surface of the skin at the time of surgery. This should be viewed as adjunctive to perioperative skin antisepsis protocols (i.e., skin prep) which is a separate SSI prevention care bundle measure. The 2021 AORN Guideline for Preoperative Patient Skin Antisepsis states the following: (1) follow manufacturer instructions for use (IFU) and safety sheet for prepping, handling, safety and storage; (2) establish a standardized skin prep protocol which includes a selected skin marking strategy; (3) confirm the surgical site and isolated contaminated sites prior to prepping with a barrier drape; (4) apply surgical skin prep agent using sterile technique and sterile supplies, starting at incision site and move outwards; (5) implement patient and staff safety measures such as minimizing the risk of fire, especially with alcohol-based products; (6) at end of procedures, assess patient skin for injury after removing skin antiseptic agent; (7) document prep in the EMR. The 2013 Antimicrobial Prophylaxis Guidelines were developed jointly by the American Society of Health-System Pharmacists (ASHP), the Infectious Diseases Society of America (IDSA), the Surgical Infection Society (SIS), and the Society for Healthcare Epidemiology of America (SHEA). For the first time, this document addresses weight-based considerations to insure maximal tissue concentrations within the wound bed during the intraoperative period.
Another bundle measure shown to reduce the incidence of SSIs, especially in high-risk surgery involving implants such as knee and hip arthroplasty is the screening of patients for both MSSA and MRSA nasal colonization. While mupirocin has been for many years the standard agent for nasal decolonization, studies using nasal povidone iodine (5% or 10%) or a novel alcohol-based (70%) nasal antiseptic have been documented to be effective in reducing staphylococcal nasal colonization but further clinical investigations are warranted.151 In addition, perioperative supplemental oxygen (80%) or hyperoxia is a recognized bundle measure, as it increases tissue oxygen tension, which may lead to an increase in oxidative killing of surgical pathogens and a reduction in SSIs. Current national, societal and international guidelines are supportive of perioperative hyperoxia to reduce SSIs in colorectal surgery patients.152 The SSI prevention care bundle element related to the patient’s core temperature recommends monitoring throughout the surgical procedure and with maintenance of normothermia preoperatively, intraoperatively, and postoperatively. Hypothermia is defined as a core temperature below 36 °C and patients should be prewarmed a minimum of 30 minutes prior to the induction of anesthesia.153 The presence of diabetic hyperglycemia at the time of surgery is a significant risk factor across the spectrum of surgical disciplines, so glycemic control is yet another evidence-based SSI prevention care bundle element. Hemoglobin A1C represents a more accurate indicator of glycemic control than blood glucose. Preoperative and inpatient diabetes management improves glycemic control on the day of surgery and postoperatively, decreasing the incidence of hypoglycemia which leads to improved clinical outcome.154 A systematic literature review (SLR) and meta-analysis identified 15 randomly selected controlled trials, producing a risk ratio of 0.67, 95% CI 0.54-0.84 (P ≤ 0.001), demonstrating that use of antimicrobial coated/impregnated sutures were effective in demonstrating a statistically significant, lower risk of SSI in clean, clean-contaminated and contaminated surgical procedures.155 Two recent studies have documented that use of an antimicrobial suture for wound closure as part of an evidence-based surgical care bundle can provide a significant fiscal benefit to the hospital and third party-payer, suggesting that antimicrobial sutures should be considered for both superficial and deep layer closure after all surgical operations.156,157

**EMERGING PATHOGENS**

The National Institute of Allergy and Infectious Diseases defines emerging infectious diseases “as infectious diseases that have newly appeared in a population or have existed but are rapidly increasing in incidence or geographic range”.158 Although these emerging pathogens may be well studied in laboratory settings, by definition, when they emerge in populations that have not had prior exposure or experiences with such diseases, the public, as well as IP programs are woefully unprepared to address the public health implications and, in turn, the socio-economic ramifications of being so ill-prepared. IP programs find themselves all too often being reactionary rather than proactive in the light of emerging pathogens. Two very different but recent examples can be seen with Ebola and COVID-19. First identified in 1976, the CDC describes 33 outbreaks in 19 countries of the Ebola virus hemorrhagic fever (including the 3 species that cause disease in humans: Zaire, Sudan, and Bundibugyo; and 1 species that does not cause disease in humans, Reston) prior to the Ebola outbreak of 2014.159 Although this pathogen was known for 38 years, when Ebola was first diagnosed in a patient admitted to a hospital in Texas, United States (US), it was evident that infection prevention programs were not prepared to properly control potential transmission. Transmitted through infected body fluid, PPE took on a new meaning for front-line staff, including the importance of selecting the correct PPE, as well as conducting extensive training to learn the proper process of donning and doffing PPE to reduce the risk of blood or body fluid contamination. In addition, special arrangements in terms of the patient care environment were needed to address the patient’s isolation room, temporary “mini-labs” where testing can be run by specially trained staff, 1-way traffic flow to prevent accidental contamination, nearby emergency showers for staff in the event of exposure, and special waste management precautions. More recently, in December 2019, when COVID-19 first appeared in California and rapidly spread across the United States, it quickly became apparent that the US public health and health care infrastructure were not designed to provide rapid identification of infected individuals, control transmission among the vulnerable population, and manage and treat large surges of patients in our health care facilities. Emerging pathogen events have underscored the need for IP programs to emerge from positions of remaining reactive to becoming proactive by developing robust systems capable of properly handling future pandemic diseases.

As described by Abouleish, a lack of information, conflicting information, unfamiliarity with the disease, along with human perception all work together to formulate a perceived risk level of disease transmission.160 This helps explain, in part, acceptance, or lack thereof, of infection prevention measures by the public. In the context of the latest pandemic, universal infection prevention and control approaches to reduce disease transmission in the health care setting focus on: (1) early identification and appropriate isolation of patients suspected of having the disease through symptom screening or testing; (2) universal source control through mandatory masking to contain respiratory droplets; (3) quarantine after exposure; (4) the use of appropriate personal protective equipment; and (5) appropriate environmental decontamination.161,162

Airborne infection isolation rooms (AIIR), formerly known as negative pressure isolation rooms, are designed to protect other patients and health care team members from diseases transmitted via the airborne route. Most facilities are designed with few AIIR in comparison to standard rooms, and as a result, these were rapidly allocated during the surges of COVID-19.163 Soon, infection prevention programs found themselves working creatively to develop geographic cohorts (entire units made up of patients with a common infectious disease) or find ways to create more AIIRs, for example, by using portable HEPA scrubbers or changing the ventilation dynamics of a particular unit or floor. Through the course of the pandemic, research has found that the large majority of COVID-19 virus is spread via respiratory droplets that drop precipitously within 6 feet of the infected, not only putting unprotected individuals at risk within that 6-foot space, but also able to contaminate the environment near infected individuals.164 However, the relationship of finding virus particles in the environment and the role these particles play in the ability to cause disease requires further investigation.

Aerosol generating procedures (AGP) have gained considerable attention since the pandemic began. Yet there remains poor consensus and conflicting evidence on which specific procedures should be considered AGPs, and to what extent these procedures can or do, pose the greatest risk in COVID-19 transmission.162 The answer undoubtedly lies within a spectrum of factors that contribute to determining acceptable or nonacceptable risk. As an example, nebulizers, which by design aerosolize medicine for patient inhalation, have not been fully researched to determine their association with disease spread given such elements as variation in disease state of the host and ventilation/air quality of the surrounding environment. Optimal air quality includes having minimum fresh air exchanges—adequate ventilation is considered to be 60 liters/s/patient (AGPs not performed)162 which may not always be possible, especially given a surge situation. Manufacturers of technologies that use HEPA or Ultra-HEPA filtration, ultraviolet-C radiation, and various combinations of other technology promise to rid the air of impurities and kill
airborne infectious agents, often with a paucity of concrete data in practical applications. Although scientific evidence conducted in laboratory settings has supported the effectiveness of these technologies, there is a need for new research and validation that these technologies delivered in real-world settings can verify these claims.

An additional important issue concerns the interplay between emerging pathogens and PPE. As the COVID-19 pandemic progressed rapidly through the first year, the extremely high demand for, and subsequent lack of, PPE forced many infection prevention programs to develop policies and practices related to extended use and re-use of PPE. To meet the need, infection preventionists were forced to reevaluate disposable PPE and develop various methodologies of extending their use or reprocessing single-use equipment so it could safely be used again. Most notably, reprocessing filtering facepiece respirators (FFR) became exceedingly important. A systematic literature review identified 14 methods for decontaminating FFR, with 4 predominant methods surfacing: ultraviolet germicidal irradiation, moist heat, microwave-generated steam, and hydrogen peroxide vapor. In addition, cautionary tales of using the various technologies that decontaminate respirators may indeed alter their performance. Further research is needed to ensure that decontamination and reuse of PPE can be done effectively and efficiently without compromising safety. Alternatively, identifying existing PPE or developing new PPE that can be worn safely and effectively for extended periods of time (such as powered air-purifying respirators) should be further investigated.

Despite attempts at understanding how to handle IP issues associated with COVID-19, the financial ramifications incurred by health care institutions due to the pandemic may represent the greatest ongoing challenge. The impact incurred by rising operating costs and decreasing revenue may very well translate into reduced budgets for programs such as infection prevention. This may present itself with cuts to IP staffing, infrastructure resources, product limitations, or pauses in innovative technologies.

**CONCLUSION**

Fifty years of IP evolution has brought us to a crossroads in which we face increasingly diverse and complex issues. This article addresses many of those issues, providing insight on how to more effectively improve IP programs, standardize metrics, and better control potential HAI events in the future. (Table 1 summarizes recommendations for change in 14 topics of concern.) These advances must be accomplished with the understanding of the importance of a structure for infection prevention nationally that spans across the continuum of care from acute to skilled nursing to ambulatory to postacute settings and which is resilient to mammoth events such as pandemics. Regardless of which strategies are considered, IP successes will depend largely on strong leadership support. A recent analysis of management methods identified 3 practices as important facilitators in the prevention of HAI: First involves engagement of executive staff. Establishment of IP goals by executive leadership emphasizes an organizational priority among managers and frontline staff and enables open communication with persons who are empowered to make change. Second addresses information sharing. Establishment of an organization-wide system to relay, display, and discuss relevant infection data with frontline staff is an important activity. Third involves management coaching. The coaching activities identified as most needed involve providing staff with feedback on how to perform clinical care processes correctly and re-educating staff on best practices for IP. The future success of IP programs will therefore, lay in identifying and implementing cutting edge program modifications and best practices while supported by targeted executive actions.

| IP Program Standardization | • Based on outcomes based scientific research (eg, HAI rates, process compliance, patient satisfaction), establish a standard template for IP programs to support the replication of best practices, avoid errors, and optimize processes. |
| --- | --- |
| Surveillance | • Conduct research focusing on determining IPs time allocation taking into account variability between health care facilities. |
| Hand hygiene | • Conduct research into IP program reporting structure. |
| Environmental Contamination | • Establish a certification process for physicians in IP programs. |
| Antibiotic Resistant Organisms | • Conduct research to determine ideal IP staffing levels based on essential program components. |
| Surveillance | • Establish a collaborative association between NHSN and the IP community to evaluate HAI definitions in order to increase accuracy and reflect quality of clinical care and processes. |
| Hand hygiene | • Conduct research into methods and training regimens to improve accuracy when conducting manual surveillance. |
| Environmental Contamination | • Establish a collaborative association between vendors of automated surveillance software and the IP community to evaluate the standardization and improvement of HAI accuracy across all available platforms. |
| Antibiotic Resistant Organisms | • Review available national and international programs addressing behavior modification of health care personnel for improving hand hygiene. |
| Surveillance | • Review published studies on hand hygiene improvement strategies that include enhancements in education, monitoring, infrastructure, and culture. |
| Hand hygiene | • Consider the use of automated hand hygiene systems designed to assist in the verification of compliance while providing the ability to track compliance history. |
| Environmental Contamination | • Review studies addressing limitations in environmental cleaning and conduct a gap analysis to determine which factors need to be addressed. |
| Antibiotic Resistant Organisms | • Implement new strategies based on societal guidelines including those addressing the education of EVS staff to increase cleaning and disinfection. |
| Surveillance | • Establish facility-specific acceptable levels of cleaning and disinfection. |
| Hand hygiene | • Consider the use of supplemental disinfection technologies taking into consideration. such factors as cost, staffing needs, time allotments, and effectiveness of disinfection process. |
| Environmental Contamination | • Using key national and regional information, establish facility-specific listing of AROs integrating such information into the EMR in order to expedite isolation, therapy strategies, and antimicrobial stewardship program review. |
| Antibiotic Resistant Organisms | • Ensure that the facility receives state and local public health organizations’ timely information on emerging AROs. |

(continued)
• Consider the use of rapid diagnostic technology for AROs; such technologies reduce both
  identification and antibiotic susceptibility testing, thereby expediting isolation protocols and
  narrowing antibiotic therapies.

Decolonizing Patients
• Consider the use of a universal decolonization strategy for ICU patients (strategy includes use of
  CHG bathing in conjunction with a nasal antibiotic/antiseptic regimen).
• Consider the use of a decolonization strategy for non-ICU patients with vascular access
  devices (strategy includes use of CHG bathing with or without the use of a nasal
  antibiotic/antiseptic regimen).
• Consider the use of a decolonization strategy for patients undergoing cardiac and
  orthopedic surgery.
• Consider the use of a decolonization strategy for patients receiving CIED implants;
  Conduct further research on decolonization strategies for patients receiving other
  types of implants.

Decreasing Contact Precautions
• Discontinuing CPs requires health care facilities to assess which AROs are isolatable and
  under what conditions (colonized or infected), type of isolation initiated (standard
  precautions, enhanced barrier precautions, etc.), and whether supplemental strategies are
  used that may reduce the risk of transmission.
• Health care facilities should establish protocols for discontinuing CPs based on ARO
  transmission risk, organizational priorities, and resources.

Diagnostic Stewardship
• In order for health care facilities to invest in DS, there will be a need for the establishment
  of drivers to incentivize implementation, the development of meaningful measures,
  and directed leadership support.
• Establish system-wide standardized practice guidelines, jointly written by IPs, microbiologists,
  and other key stakeholders, providing guidance in the pre-analytic phase, that is, pathways
  for selecting the appropriate test according to patient's syndrome, methods for obtaining
  optimal collection of clinical specimens, and interpreting microbiology results.
• Implement educational efforts on education of nurses to understand the "why" behind
  proper culturing techniques, and of physicians on appropriate ordering and interpretation
  of test results.
• In order to ensure the integration of all areas of infection management (diagnosis, treatment,
  and prevention), establish a unified stewardship strategy such as reflected in the AID
  (antimicrobial, IP, and DS) model. The basis of this model is the understanding that outcomes
  improve when providers understand and cooperate in implementing a system that promotes
  the appropriate test, the right interpretation of a test result, selection of the appropriate
  antimicrobial, and administration at the right time.
• Revise order sets in EMRs to reflect essential modifications in testing, to include urine
  and blood culture ordering and management protocols.
• Consider evaluating novel methods to reduce blood culture contamination.
• DS interventions related to C difficile testing should include provision of education to
  clinicians on what constitutes clinically significant diarrhea, the appropriate documentation
  of patient bowel movements and use of laxatives, implementation of C difficile order sets
  and EMR best practice alerts to assist providers in order to optimize testing and enhance
  the identification of patients with active disease, and not colonization.

HAI Surveillance and Prevention: Bloodstream Infections
• Establish collaborative association between NHSN and the IP community to evaluate
  BSI definitions in order to increase accuracy and reflect quality of clinical care and processes.
• Conduct a gap analysis and knowledge assessment to determine educational and process
  needs, followed by institution of enhanced maintenance practices.
• Consider evaluating novel technologies that extend the effective antimicrobial period over
  the entire recommended duration of an IV dressing.
• Expand NHSN surveillance utilizing the model Hospital Onset Bacteremia, an initiative that
  captures BSI events not only related to central lines but to all types of intravascular catheters;
  such a strategy may be considered for application to other health care settings such as
  extended care facilities.

HAI Surveillance and Prevention: Nonventilator
Hospital Acquired Pneumonia
• Establish a collaborative association between NHSN and the IP community to review
  published information on the occurrence of NV-HAP events, draft definitions, and to consider
  surveillance trials.
• Consider establishing universal prevention measures including oral care, early and frequent
  ambulation, head of bed elevation, and aspiration precautions.

HAI Surveillance and Prevention: Ventilator Associated Events
• Establish a collaborative association between NHSN and the IP community to evaluate
  VAE definitions in order to increase accuracy and reflect quality of clinical care and processes.
• Establish a collaborative association between NHSN and the IP community to facilitate the
  transition to institute ventilator-associated pneumonia as a requirement for national reporting,
  a measure that would incentivize prevention initiatives.

HAI Surveillance and Prevention: Urinary Tract Infections
• Establish a collaborative association between NHSN and the IP community to evaluate
  UTI definitions in order to increase accuracy and reflect quality of clinical care and processes.
• Establish comprehensive education programs for clinicians emphasizing such important
  preanalytic issues as understanding pretest probabilities before obtaining a urinalysis
  or culture and incorporating the patient's symptoms in the decision process regardless
  of the clinical setting.
• Establish standardized methods of urine collection for both catheterized and
  noncatheterized patients.
• Establish laboratory processes for carefully reviewing urinalysis findings prior to
  processing a urine specimen for culture.
• Conduct research to better understand the urinalysis parameters that should
  trigger a urine culture.
• Conduct research to better understand the incidence of UTI and associated antimicrobial
  treatment in populations at risk for misdiagnosis (eg, ambulatory or long-term settings).
• Consider establishment of an advanced, evidence-based surgical care bundle, with new
  measures to include preadmission CHG shower/cleansing, weight-based antimicrobial

(continued)
Emerging pathogens

prophylaxis, a nasal decolonization strategy, perioperative supplemental oxygen, maintenance of normothermia, glycemic control, and use of antimicrobial sutures.

- Conduct research addressing the appropriate control of patients with emerging diseases to include methodologies to improve early identification, surge management, isolation including use, necessity, and alternate technologies for AIRRs, transportation of patients, selection, proper use, and reprocessing issues related to such PPE items as facemask respirators, employee exposure management, waste management, internal and external communication enhancement, initiation and duration of quarantine, effective environmental decontamination, and identification of AGPs that pose greatest risk of organism transmission.

**AGP, Aerosol generating procedures; AIRR, Airborne infection isolation rooms; ARO, antibiotic-resistant organism; BSI, Bloodstream Infections; CHG, chlorhexidine gluconate; CIED, cardiac implantable electronic devices; CP, contact precautions; DS, Diagnostic Stewardship; EDR, electronic medical records; EVS, environmental services staff; HA, Healthcare associated infections; IP, infection prevention; NV-HAP, Nonventilator hospital acquired pneumonia; NHSN, The National Healthcare Safety Network; PPE, Personal Protective Equipment; UTI, Urinary tract infection; VAE, ventilator associated events.**

**STATEMENT**

The views of the authors may not be representative of the corresponding institutions.

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