ASHP national survey of pharmacy practice in hospital settings: Dispensing and

administration-2020

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**Purpose.** Results of the 2020 ASHP national survey of pharmacy practice in hospital settings are presented.

**Methods.** Pharmacy directors at 1,437 general and children's medical/surgical hospitals in the United States were surveyed using a mixed-mode method of contact by email and mail. Survey completion was online. IQVIA supplied data on hospital characteristics; the survey sample was drawn from the IMS hospital database.

**Results.** The response rate was 18.7%. Almost all hospitals (92.5%) have a method for pharmacists to review medication orders on demand. Most hospitals (74.5%) use automated dispensing cabinets (ADCs) as their primary method for drug distribution. A third of hospitals use barcodes to verify doses during dispensing in the pharmacy and to verify ingredients when intravenous medications are compounded. More than 80% scan barcodes when restocking ADCs. Sterile workflow management technology is used in 21.3% of hospitals. Almost three-quarters of hospitals outsource some sterile preparations. Pharmacists can independently prescribe in 21.1% of hospitals. Pharmacist practice in ambulatory clinics in 46.2% of health systems and provide telepharmacy services in 28.4% of health systems.

**Conclusion.** Pharmacists continue their responsibility in their traditional role in preparation and dispensing of medications. They have successfully employed technology to improve

safety and efficiency in performance of these duties and have employed emerging technologies to improve the safety, timeliness, and efficiency of the administration of drugs to patients. As pharmacists continue to expand their role to all aspects of medication use, new opportunities highlighted in ASHP's Practice Advancement Initiative 2030 have been identified.

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The ASHP national survey of pharmacy practice in hospital settings focuses on practices and technologies for managing and improving the medication-use system and the role that pharmacists play in this effort. The national surveys are organized according to 6 components of the medication-use system: prescribing, transcribing, dispensing, administration, monitoring, and patient education. Each year, the survey focuses on 2 components in the medication-use system. The 2020 survey cycle evaluated practices and technologies related to dispensing and administration. The most recent 3 surveys represent a composite picture of the ways hospitals and health systems are managing and improving the entire medication-use system and current roles of pharmacists in medication-use system management.

In assessing dispensing and administration practices, the 2020 survey was intended to describe and evaluate trends in the inpatient medication distribution system; the methods for medication preparation and dispensing; the use of technology in medication distribution, including the use of bar-coding technology in the preparation of compounded sterile preparations; sterile compounding environmental sampling program reporting; staff competency assessment to prepare compounded sterile preparations; and hazardous drug handling. It also addressed the quality assessment of medication administration processes, including the use of smart infusion pumps, the outsourcing of sterile preparation activities, the process of medication order review, baseline assessment of selected Practice Advancement Initiative (PAI) 2030 goals, and pharmacist-provided telehealth visits.

### Methods

An evaluation of dispensing and administration practices in US hospitals and health systems was conducted using methods similar to those used in past ASHP surveys.<sup>1-18</sup>

**Questionnaire development.** The 2020 questionnaire was developed using procedures suggested by Dillman.<sup>19</sup> Questions from previous surveys that pertained to topics of interest in this survey were evaluated for clarity and response. As with past surveys, data about hospital characteristics were available in the IMS hospital database.<sup>20</sup>

Survey sample. A sampling frame of 4,865 general and children's medical/surgical hospitals in the United States was constructed from the IMS database. Specialty, federal, and Veterans Health Administration hospitals were excluded from this sampling frame. Hospitals were stratified by size before sampling, and random samples of hospitals within these strata were taken to select the sample of 1,437 hospitals. We sampled 300 hospitals with fewer than 50 beds to compensate for historically lower response rates in hospitals of that size. We sampled all hospitals with 600 or more staffed beds (n = 137) to collect data from enough very large hospitals to provide reliable estimates. Two hundred hospitals were sampled in each of the other hospital size categories (Table 1).

**Data collection.** A mixed-mode survey method of contact by email and mail was used. Survey completion was done online using Qualtrics (Qualtrics, Provo, UT). Pharmacy directors in the sample were contacted up to a total of 8 times during the survey period. An announcement email was sent in August 2020. This first contact explained the survey and directed respondents to the online data collection site. At the beginning of September 2020, all hospital pharmacy directors were mailed an announcement letter that also explained the survey and directed respondents to the online data collection site. Email reminders were sent to all nonrespondents at the middle of September, end of September, and end of October 2020. All nonrespondents were mailed a postcard reminder at the middle of October 2020, and were mailed a reminder letter in the middle of November 2020. A final email reminder was sent to all nonrespondents on December 15, 2020. Data collection was closed on December 21, 2020.

**Data analysis.** Each hospital in the sample was assigned a unique identification number. This number allowed the survey response to be matched with the hospital characteristics in the IMS database. As with past surveys, data are presented by categories of staffed beds to more closely align with data from the American Hospital Association.<sup>21</sup> We used a design-based analysis.<sup>22</sup> This technique results in population estimates that are more accurate than reporting unweighted results.

Data were output from Qualtrics into an SPSS-readable file. All non-designed-based analyses were conducted using SPSS Version 27 (IBM Corporation, Armonk, NY). All designed-based analyses were conducted using StataSE Version 16 (StataCorp LLP, College Station, TX) using the set of survey commands. Weights were assigned to respondents to adjust their contribution to the population estimate. The weights were 32.80 for hospitals with fewer than 50 staffed beds, 21.03 for hospitals with 50 to 99 beds, 28.97 for hospitals with 100 to 199 beds, 19.77 for hospitals with 200 to 299 beds, 9.20 for hospitals with 300 to 399 beds, 7.76 for hospitals with 400 to 599 beds, and 4.15 for hospitals with 600 or more staffed beds. The strata were the categories for number of staffed beds, and the finite population correction was the total number of hospitals in the population (4,865).

Descriptive statistics were used extensively. Chi-square analysis and analysis of variance or regression was used to examine how responses differed as a function of hospital characteristics. The a priori level of significance was set at 0.05.

### Results

A total of 269 hospitals submitted usable data for analysis. The overall response was 18.7%.

**Hospital characteristics.** Table 1 shows the size, location, and ownership of the respondents' hospitals, the nonrespondents' hospitals, the surveyed hospitals, and the 4,865 general and children's medical/surgical hospitals. The characteristics of the surveyed hospitals are presented to highlight the complex sampling design employed in this survey. Respondents and nonrespondents were statistically different in regional location, and ownership status.

Medication order review. Overall, 92.5% of hospitals have a method for pharmacists to review and enter medication orders on demand (Table 2). Having the pharmacy department open and staffed 24 hours a day and 7 days a week is the most common method (42.8%), followed by after-hours medication order review and entry provided by a telepharmacy company (29.7%), use of an affiliated hospital with 24-hour services (15.0%), and having an employee pharmacist on call or at a remote location (5.1%) (Figure 1). In 7.5% of hospitals, orders are not reviewed by a pharmacist when the pharmacy department is closed. Smaller hospitals are more likely to not have the pharmacy review orders when the pharmacy department is closed (Table 2) and, when they do review orders, are more likely to use a telepharmacy company or an affiliated hospital to review medication orders. Larger hospitals are more likely to have a 24-hour pharmacy service. Regardless of the review method used, the percentage of hospitals where medication orders are not reviewed by a pharmacist has declined. The percentage of hospitals not reviewing orders after hours has declined annually since 2005, when we first surveyed order review, from 59.6% of US hospitals.<sup>3,5-10,12-15</sup>

Inpatient medication distribution technology. In 2020, 4.1% of general and children's medical/surgical hospitals used a robotic distribution system that automates the dispensing of unit dose inpatient medications in a centralized distribution system (Table 3). Most hospitals (74.5%) use automated dispensing cabinets (ADCs) as the primary method of maintenance dose distribution. Smaller hospitals are more likely than larger hospitals to use centralized manual unit doses systems for maintenance dose fulfillment, and larger hospital are more likely than smaller hospitals to use a robotics and ADCs for maintenance dose fulfillment,

The use of ADCs as the primary method of maintenance dose distribution increased since the 2002 survey,<sup>18</sup> rising from 22.3% that year to 37.8% in 2005,<sup>15</sup> 49.2% in 2008,<sup>12</sup> 62.5% in 2011,<sup>9</sup> and 70.2% in 2017<sup>3</sup> (Figure 2). There was a corresponding decrease in the use of centralized manual unit dose systems as the primary method of maintenance dose distribution since 2002, with 20.1% of hospitals using this method in 2020.

Only 5.9% of hospitals do not have ADCs on patient care units. Of those hospitals with ADCs, 77.4% use individually secured lidded pockets as the predominant ADC configuration, and 22.6% use the original matrix drawer configuration that allows access to all medications stocked in a drawer (Table 4). The use of lidded pockets has increased over the last decade, from 51.5% of hospitals in 2008<sup>12</sup> to 61.9% in 2011,<sup>9</sup> 65.7% in 2014,<sup>6</sup> and 70.1% in 2017.<sup>3</sup>

Machine-readable coding in pharmacy. Robots, carousels, and, sometimes, manual unit dose pick stations use machine-readable coding to verify removal and replenishment of medications. Overall, 66.3% of hospitals use some form of machine-readable coding to verify doses during dispensing in the pharmacy (Table 5). The use of machine-readable coding in pharmacy departments has steadily increased over the past 12 survey years

(frequencies of use were 5.7% in 2002,<sup>18</sup> 11.5% in 2005,<sup>15</sup> 24.0% in 2008,<sup>12</sup> 33.9% in 2011,<sup>9</sup> 44.8% in 2014,<sup>6</sup> and 74.7% in 2017.<sup>3</sup> This practice differs by hospital size, with larger facilities using scanning during dispensing more than smaller facilities.

Furthermore, 81.4% of hospitals scan medication barcodes during restocking of ADCs; this differs by hospital size, with 100% of the largest hospitals (600 or more staffed beds) scanning barcodes while restocking ADCs, as compared with 90.2% of hospitals with 400 to 599 beds, 87.8% with 300 to 399 beds, 93.5% with 200 to 299 beds, 80.0% with 100 to 199 beds, 84.8% with 50 to 99 beds, and 72.2% with fewer than 50 beds.

The use of machine-readable coding during restocking of ADCs increased from 43.3% of hospitals in 2011,<sup>9</sup> 62.1% in 2014,<sup>6</sup> and 74.7% in 2017.<sup>3</sup>

**Sterile compounding technology.** Sterile preparation workflow management technology is used in 21.3% of hospitals (Table 6 and Figure 3). The use of workflow management technology for IV compounding differed significantly by hospital size, with larger hospitals more likely to have workflow management software compared to smaller hospitals. The use of this technology has increased from 6.5% in 2014<sup>6</sup> to 12.8% in 2017,<sup>3</sup> 16.4% in 2018,<sup>2</sup> and 19.8% in 2019.<sup>1</sup>

Barcode scanning to verify ingredients during the intravenous (IV) medication compounding process is used by 33.8% of hospitals (Table 6). Results in this area differed significantly by hospital size, with larger hospitals being more likely than smaller hospitals to use barcode scanning. The use of barcode scanning to verify ingredients has increased over the past 8 years, from 11.9% in 2011.<sup>1-3,6,9</sup>

The use of pictures or video of the compounding process was reported at 25.3% of hospital pharmacies (Table 6). The use of pictures or video has increased annually from 2017<sup>1-3</sup> and was significantly increased in 2020 vs 2019.<sup>1</sup>

Gravimetrics to verify dose, amount, and volume is used by 5.0% of hospital pharmacies (Table 6). Results in this area differed significantly by hospital size. The use of gravimetrics was stable over the last 3 years.<sup>1-3</sup>

Overall, 52.7% of hospitals do not use any technologies for compounding sterile preparations (Table 6). This differed significantly by hospital size, with the smallest hospitals most likely to not use any technologies for compounding sterile preparations. Hospital nonuse of any technology when compounding preparations has declined annually from 64.0% in 2017<sup>3</sup> to 59.9% in 2018<sup>2</sup> and 56.4% in 2019.<sup>1</sup>

**Sterile compounding automation.** Overall, 3.4% of hospitals are using a standalone robotic device in the pharmacy department for compounding flush solutions, syringe-based small-volume parenteral preparations, and minibags, excluding chemotherapy preparations (Table 7).

Robotic chemotherapy compounding devices are used in 1.6% of hospitals (Table 7). The use of a robot to compound sterile preparations differs significantly by hospital size, with larger hospitals more likely than smaller hospitals to have a robotic compounding device. However, adoption is currently limited, with 95.7% of hospitals not having an IV sterile compounding robot (Table 7).

Assessing staff competence to prepare compounded sterile products. Overall, 93.1% of hospitals use media-fill challenge testing, 92.5% use gloved fingertip sampling, 87.4% use direct periodic observation of garbing and gloving, 86.0% use direct periodic observation of sterile technique, 80.8% use direct periodic observation of cleaning and disinfecting, 68.6% use surface testing using contact plates, 59.0% use in-house testing materials (eg, *United States Pharmacopeia* chapter 797 [USP <797>] examination, calculations examination), 53.7% use commercial education and testing materials (eg, ASHP- endorsed materials [CriticalPoint, Gaithersburg, MD]), and 16.2% use end-product testing to assess staff members' sterile compounding competency (Table 8).

Over the last 3 years, hospitals have significantly increased the use of most of these competency assessment modalities.

Environmental sampling program and reporting. Overall, 83.6% of hospitals have an environmental sampling program that specifies sampling location, methods, frequency, action levels, and follow-up. About 61% report sterile compounding environmental sampling results through organizational quality reporting pathways. Use of these programs varies by hospital size, with more than 90% of the hospitals with 100 or more staffed beds having a detailed environmental sampling program, as compared with about three-fourths of hospitals with fewer than 100 beds. Likewise, about 70% or more of hospitals with 50 or more staffed beds report sampling results through quality reporting pathways, as compared with about 43% of hospitals with fewer than 50 beds (Table 9).

**Compliance with USP chapter 800.** Pharmacy directors were asked about their compliance with USP chapter 800 (USP <800>). Overall, 31.1% of respondents indicated that their hospitals were fully compliant with all sections of the chapter, 63.1% indicated they were not yet fully compliant but working on compliance, and 5.7% were not fully compliant and were not working on compliance (Table 10).

For those hospitals that were not yet compliant, 51.1% identified gaps in facilities/engineering controls; 50.8%, gaps in personnel training; 45.9%, gaps in hazardous drug lists, policies and procedures, and/or risk assessment; 35.1%, gaps in drug storage; 22.9%, gaps in garb/personal protective equipment; 17.6%, gaps in use of closed system drug-transfer devices; and 16.5%, gaps in use of a deactivating agent during cleaning (Table 10).

**Outsourced sterile compounding.** Overall, 72.5% of hospitals reported outsourcing non–patient-specific compounded sterile preparations (CSPs) from a registered 503B pharmacy (Table 11). The use of this practice varied significantly by hospital size; larger hospitals were more likely than smaller hospitals to outsource non–patient-specific CSPs.

Furthermore, 29.5% of hospitals reported outsourcing patient-specific CSPs from a registered 503A pharmacy (Table 11). The use of this practice varied significantly by hospital size, with the largest hospitals being more likely than smaller hospitals to outsource patient-specific CSPs.

Only 21.0% of hospitals did not outsource any CSPs from 503A or 503B pharmacies. The use of this practice varied significantly by hospital size; smaller hospitals were more likely than larger hospitals not to outsource CSPs. Overall, the outsourcing of patient-specific and non–patient-specific CSPs has increased since 2018.<sup>2</sup>

Hospital pharmacy directors were asked to describe their sterile compounding outsourcing strategy. Overall, 76.3% selectively outsource to facilitate management of drug shortages and/or preparation of parenteral nutrition, patient-controlled analgesia or epidural medications, or nonsterile-to-sterile preparations; 14.2% minimize outsourcing and prefer to compound all drugs internally and outsource only when we have no other option; and 9.5% maximize outsourcing by outsourcing as many preparations as possible.

Reasons for outsourcing CSPs include extended beyond-use (82.3%), purchasing repackaged/compounded medications for procedures (eg, anesthesia syringes) (58.7%), insufficient staff to meet compounding needs (34.7%), purchasing products on the Food and Drug Administration (FDA) drug shortages list (32.3%), complex formulations (eg, total parenteral nutrition) (31.3%), preparation of large batches for anticipatory use (30.6%), insufficient facilities to meet compounding needs (20.7%), nonsterile-to-sterile compounding (17.9%), patient clinical need (concentrations not commercially available; preservative-free formulations) (17.1%), and perceived higher-quality and/or safer products from outsourcing (15.8%).

The most common strategies used to evaluate external CSP providers were confirmation of state licensure (68.0%); availability of a group purchasing organization contract with the outsourcer (58.3%); establishing a contract with the pharmacy (legal review) (55.6%); commitment to receiving, at a minimum, quarterly quality assurance documentation (49.9%); evaluation of FDA-issued Form 483 reports and other FDA reports to state boards of pharmacy, if any (47.1%); rely on vetting at the corporate/health-system level (46.7%), receiving a signed attestation that the external provider follows *United States Pharmacopeia* chapter 797 requirements and current good manufacturing practices (40.6.9%), evaluation of state board of pharmacy inspections of the outsourcing pharmacy and corrective actions (35.0%), completing the previously available ASHP Foundation Outsourcing Sterile Products Preparation Vendor Assessment Tool (32.5%), and conducting a site-validation visit to the outsourcing pharmacy (23.5%).<sup>23,24</sup> Only 0.9% of hospitals indicated that no method was used to evaluate a CSP outsourcing vendor before purchasing from the vendor (Table 12).

**Smart infusion pumps.** Overall, 87.9% of hospitals use smart infusion pumps (Table 13). The use of smart infusion pumps varies by hospital size, with the largest hospitals being the most likely to have smart infusion pumps. The percentage of hospitals using smart infusion pumps has increased over the past 15 years from 32.2% in the 2005 survey.<sup>15</sup>

Overall, 13.4% of hospitals have a smart pump/EHR interface that autopopulates pumps with prescribed order and patient information from the EHR, eliminating the need to manually select the drug and infusion rate during setup; this varies by hospital size, with the largest hospitals being the most likely to have this functionality (Table 13). This has increased significantly since 2017 from 8.9%.<sup>3</sup>

Overall, 14.9% of hospitals have an interface through which smart-pump infusion use data autopopulate to the patient record in the EHR, with 85.1% of hospitals requiring the nurse to manually document infusion data into the EHR (Table 13).

The availability of interfaces between the smart pump and the EHR has increased significantly since 2017.<sup>3</sup>

**Medication administration quality metrics.** Overall, 64.1% of hospitals regularly review smart infusion pump data and quality metrics by a medication safety/quality committee. In addition, 87.5% of hospitals regularly review barcode medication administration (BCMA) data and quality metrics by a medication safety/quality committee.

**PAI 2030.** ASHP's Practice Advancement Initiative, formerly the Pharmacy Practice Model Initiative (PPMI), aspires to transform how pharmacists care for patients by empowering the pharmacy team to take responsibility for medication-use outcomes; to promote optimal, safe, and effective medication use; to expand pharmacist and technician roles; and to implement the latest technologies.<sup>25</sup> In this survey, we examined pharmacist's ability to independently prescribe medications, use of data analytics and technology to reduce the risk of adverse events or suboptimal outcomes in patients, the level of integration of pharmacy services across the continuum of care, advanced pharmacy technician roles, pharmacists practicing in clinic settings, and pharmacists providing ambulatory patient care via telehealth.

Independent prescribing. Overall, 21.1% of hospitals empower pharmacists to independently prescribe medications, including the selection, initiation, monitoring, and adjustment of medication therapy pursuant to a diagnosis of medical disease or condition

by a provider. This excludes therapeutic interchange, IV-to-oral switches, and renal dosing of antibiotics under policy or protocol.

*Data analytics.* Data analytics and technology can reduce the risk of adverse events or suboptimal outcomes in patients. Overall, 70.5% of hospitals use basic analytics from smart pumps, clinical decision support, and/or automation in dispensing and compounding; 2.6% use basic analytics and advanced analytics in the form of artificial intelligence, machine learning, and predictive analytics; and 26.9% do not use analytics to inform treatment decisions, for adverse event monitoring, or for outcomes monitoring (Table 14). The use of analytics varies by hospital size, with larger hospitals using some form of analytics to inform decision making more often than smaller hospitals.

Service level integration. The level of integration of pharmacy services across the continuum of care can drive effective patient care transitions and promote optimal patient outcomes. Pharmacy directors were asked to rate the level of integration in their hospital. Overall, 0.6% indicated seamless integration, 13.8% indicated pharmacy services were mostly integrated, 55.0% indicated some integration, and 30.6% indicated that pharmacy services were not at all integrated (Table 15). This varied by hospital size, with smaller hospitals reporting lower levels of integration than larger hospitals.

*Technician activities.* Pharmacy technicians perform important activities in the pharmacy department to support the pharmacy practice model. Many activities require advanced skill sets beyond typical activities such as packaging, medication delivery, restocking ADCs, unit dose cart fill, and sterile and nonsterile compounding. The most common advanced activities, by the percentage of survey respondents reporting them, included purchasing (94.7%), billing (75.6%), controlled substance system management (53.4%), management of 340B Drug Pricing Program activities (49.3%), information technology system management (44.3%), responsibility for USP chapter 795 (USP <795>), USP <797>, or USP <800> compliance (43.0%) (Table 16). Less common activities include regulatory compliance (39.6%), technician supervision of other technicians (37.4%), initiation of medication reconciliation (ie, obtaining a medication list) (32.0%), checking dispensing by other technicians (tech-check-tech) (30.6%), medication assistance program management (22.9%), facilitating transitions of care (14.6%), vaccine administration (5.0%), and assisting in outpatient patient care activities (intake and vital signs assessment) (2.5%).<sup>26</sup> Larger hospitals, in general, were more likely than smaller facilities to have technicians in various advanced roles. There were significant increases from the 2018 survey in the percentages of hospitals reporting that technicians were supervising other technicians, checking the dispensing of other technicians, and involved in medication assistant program management.<sup>26</sup>

*Outpatient clinics.* Overall, 46.2% of health systems had pharmacists practicing in primary or specialty care clinics (Table 17). The use of this practice varied by hospital size, with larger hospitals being more likely to have pharmacists practicing in clinic settings. The proportion of hospitals with pharmacists practicing in outpatient clinic settings has increased from 18.1% in 2010.<sup>10</sup> Within these settings, pharmacists most commonly practice in the areas of anticoagulation (25.8%), oncology (23.2%), general drug therapy management services (21.3%), diabetes (16.9%), family medicine (16.8%), cardiovascular disease (eg, dyslipidemia, hypertension, congestive heart failure) (15.8%), infectious disease (eg, human immunodeficiency virus disease, hepatitis) (11.6%), solid organ transplant (8.4%), pain and palliative care (6.5%), immunology (eg, gastroenterology, rheumatology, dermatology, neurology) (6.1%), and pharmacogenomics (3.3%). These figures varied by hospital size, with larger hospitals being more likely to have pharmacists practicing in all of

the aforementioned clinic types. The proportion of hospitals with pharmacists in all these areas has increased since 2018.<sup>2</sup>

*Pharmacist-provided telehealth.* Overall, 28.4% of hospitals have pharmacists provide ambulatory patient care via telehealth. This varies by hospital size, with larger hospitals being more likely to have pharmacists provide patient care via telehealth than smaller hospitals. For example, at 54.8% of the largest hospitals (those with 600 or more staffed beds) ambulatory pharmacists provide telehealth visits, as compared with 63.4% of hospitals with 400 to 599 beds, 47.5% with 300 to 399 beds, 33.3% with 200 to 299 beds, 29.4% with 100 to 199 beds, 25.0% with 50 to 99 beds, and 15.1% with fewer than 50 beds (uncorrected  $\chi^2$  = 25.2882, *df* = 6, design-based *F*(4.47, 1136.19) = 4.7554, *P* = 0.0005).

The primary tools used to provide telehealth services are the phone (70.9%), video chat (25.2%), EHR patient portal (3.3%), and email (0.6%). No hospitals reported using text messaging for pharmacist-provided telehealth visits.

Overall, 41.0% of hospitals report billing for pharmacist-provided telehealth services.
Discussion

Based on the ongoing results of the ASHP national hospital pharmacy survey over the past years, pharmacists can reflect on the progress that has been made in medication-use management and the important role that the profession of pharmacy has played in improving it. This is true in all phases of the medication-use process, including the cornerstone of pharmacy practice (drug preparation and dispensing) and is reflected in the results of the 2020 survey.

While pharmacists are potentially involved in all steps, a critical role is a review of the medication order before a dose is prepared and dispensed before administration to the patient. In the hospital setting, this has historically been a challenge given the remoteness of the site of medication ordering from the site where it is prepared and dispensed (ie, in the pharmacy). There is also a more urgent need for the drug in the hospital, and care is provided 24 hours per day. Early leaders in hospital pharmacy advocated and implemented a 24-hour pharmacy service, but adoption has been slow, particularly in smaller institutions. While the percentage of hospitals with this service is still below 50%, other methods have evolved, making it possible for more than 90% of hospitals to implement a method for pharmacists to prospectively review medication orders to detect and prevent errors before a dose is administered to the patient. Electronic access to health information and connectivity to decentralized ADCs have enabled this positive trend (Figure 1).

As length of stay has shortened and the acuity of care provided in the hospitals has increased, the need for a quick turnaround time between a treatment decision and drug administration is more important. Centralized unit dose drug dispensing systems were no longer responsive enough. Placing medications in patient care areas evolved as a way to make medications more readily available, and decentralized drug dispensing systems using ADCs have almost entirely replaced centralized unit dose drug distribution programs (Figure 2). This change carried with it the potential to bypass the pharmacist review of medication orders and the possibility of accessing drugs before such a review, potentially resulting in a less safe drug distribution system. Enhancements to ADCs are important, available, and necessary to ensure a safe decentralized drug distribution system. Examples of these enhancements include restricting access to medications until the pharmacist reviews the medication order and securing the bin in which individual medications are stored ("lidded pockets") so that drugs cannot be inadvertently obtained before the order review.

Safety in the medication-use system has also been improved by acknowledging the limits of human performance and adopting technologies that are more reliable. These

technologies include the use of robotic technologies and machine-readable coding for verification and documentation. The application of some of these technologies is more feasible in the hospital and healthcare setting than in other settings. For example, robots used to support a centralized drug distribution system, while likely more accurate, are expensive and difficult to cost-justify in all but the largest of hospitals or multihospital systems with centralized operations. This is reflected in the survey data showing that only 4.1% of hospitals make use of robotic technology to support a centralized unit dose drug distribution system, and only 3.4% use standalone robotic devices to support a centralized sterile compounding program. In contrast, barcode verification and documentation have gained more widespread use to support CSPs, stocking of automated drug storage devices, and drug administration.

The safety of CSPs has been an important issue due to highly publicized events of patient harm and an increased interest in employee safety. In response, increasingly stringent practice standards and regulatory oversight have emerged. The response by pharmacists has been primarily driven by the USP standards, namely USP <797> and USP <800>. As indicated by the national survey results, many changes in pharmacy practice have taken decades (Figures 1 and 2), but those prompted by enforceable standards of practice evolve more quickly. This is reflected by changes in practices for compounding sterile preparations driven by USP <797> and changes in practices for handling of hazardous drugs driven by USP <800>. Survey results show significant improvement in these practices during a short (3-year) time period; they also explain the increase in outsourcing the preparation of sterile compounded products, particularly complex preparations, for which compliance with practice standards is more challenging.

In addition to the compounding of sterile preparations, there are heightened risks associated with the administration of sterile medications. These risks have resulted in the almost universal use of smart pumps that provide alerts when an infusion is programmed incorrectly. An additional patient safety feature that is emerging is the linking of the smart pump to electronic health information systems to import prescribing information into the pump and export drug administration information out of the pump to the healthcare record. This capability prevents problem-prone transcription of information to healthcare records and the device and eliminates errors of omission in drug administration. While the use of smart pumps for IV drug administration is almost universal, the availability of such an electronic interface is not; however, it is increasing. Another use of smart pumps is as a measurement instrument to record quality information based on pump programming errors and alert overrides. This is a feature that is being used more often now, with almost twothirds of hospitals using such information as a quality metric and doing so through a hospital quality or patient safety committee.

While pharmacists still focus attention on their traditional roles in drug preparation and dispensing, they have been turning their attention to improving other steps in the medication-use process. This is the focus of PAI 2030, which is intended to provide pharmacists with the tools and guidance they need to continue to lead and shape the profession. PAI 2030 consists of 59 recommendations on providing optimal, safe, and effective medication use to provide aspirational guidance as a roadmap to pharmacy practice advancement. The 2020 ASHP national hospital pharmacy survey included several questions to provide baseline data on some of the PAI 2030 recommendations. These data reveal both challenges and opportunities. For example, only 21.1% of hospitals permit pharmacists to independently prescribe. More than a quarter of hospitals do not use analytics to inform treatment decisions. Less than 1% seamlessly integrate pharmacy services, and almost a third do not integrate these services at all. Most hospitals make use of pharmacy technicians for traditional activities, but less than 40% use them for advanced roles such as supervising other technicians, medication reconciliation, medication assistance program coordination, and facilitating transitions of care.

Part of transitions of care programs include ambulatory care services provided in outpatient clinics. Almost 50% of health systems surveyed had pharmacists practicing in the ambulatory care setting; this is a dramatic increase from less than 20% reported in 2010. An alternative model for providing service to patients when they are out of the hospital is telehealth, and pharmacists can provide services to patients using this platform. Almost 30% of hospitals have such a program, and it is likely that this will continue to increase based on changes in care delivery in response to the coronavirus disease 2019 epidemic.

### Conclusion

The annual ASHP national survey of pharmacy practice is an important tool to assess the current state of practices related to medication use and the role that pharmacists play in monitoring, managing, and improving it. The results of the 2020 survey show that pharmacists continue their responsibility in their traditional role in preparation and dispensing of medications, and they have successfully employed technology to improve safety and efficiency in these duties. Moreover, they have employed emerging technologies to improve the safety, timeliness, and efficiency of the administration of drugs to patients. As pharmacists continue to expand their role to all aspects of medication use, new opportunities highlighted in the ASHP PAI 2030 have been identified. Tracking progress towards these new roles will be an important role for the ASHP national survey.

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Figure 1. 24-hour review of medication orders by pharmacists.

Figure 2. Evolution of drug distribution systems (1975-2020). ADC indicates automated dispensing cabinet.

Figure 3. Growth in implementation of sterile product preparation technologies. Numbers above bars indicate percentage of surveyed hospitals reporting use or nonuse of technology.

# **Key Points**

- Decentralized drug distribution systems now predominate in US hospitals.
- Machine-readable coding is now widely used throughout the medication-use process.
- The availability of electronic health information has enabled remote models of care.

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			ci	8				
Table 1. Size, Location, and Ownership	of Respondents' Ho	ospitals <sup>ª</sup>	9					
	Respo	ndents	Nonres	oondents	Surv	eyed	Рори	lation
Characteristic	n	Row %	 n	 Row %	n	 Row %	n	Col %
All hospitals	269	18.7	1,168	81.3	1,437	28.9	4,865	100
Staffed beds								
<50	55	18.3	245	81.7	300	16.6	1,804	36.3
50–99	33	16.5	167	83.5	200	28.8	694	14.0
100–199	35	17.5	165	82.5	200	19.7	1,014	20.4
200–299	31	15.5	169	84.5	200	32.6	613	12.3
300–399	41	20.5	159	79.5	200	53.1	377	7.6
400–599	42	21.0	158	79.0	200	61.3	326	6.6
≥600	32	23.4	105	76.6	137	100.0	137	2.8
Region <sup>b</sup>								
West	55	22.2	193	77.8	248	25.8	963	19.4
Midwest	97	23.2	322	76.8	419	28.8	1,453	29.3
South	74	14.0	453	86.0	527	27.9	1,888	38.0
Northeast	43	17.7	200	82.3	243	36.8	661	13.3
Ownership <sup>c</sup>								
For-profit	29	13.7	182	86.3	211	27.7	763	15.4
Nonprofit	240	19.6	986	80.4	1,226	29.2	4,202	84.6
a								

<sup>a</sup>From the IQVIA hospital database.

 ${}^{b}\chi^{2} = 14.453, df = 3, P = 0.002.$ 

 $^{c}\chi^{2} = 4.024, df = 1, P = 0.045.$ 

Table 2. Medication Order Review and Entry  Afterhours medication order review and entry provided by									
	ò	Hospital open 24/7 and orders reviewed in-house in real time	National or regional company	Affiliated hospital with 24-hour pharmacy services	Employee pharmacist (on-call or remote location)	Orders not reviewed afterhours			
Characteristic	n	%	%	%	%	%			
Staffed beds	2								
<50	53	1.9	50.9	20.8	9.4	17.0			
50–99	32	28.1	40.6	18.8	3.1	9.4			
100–199	34	61.8	20.6	14.7	2.9	0.0			
200–299	30	76.7	6.7	13.3	3.3	0.0			
300–399	40	92.5	2.5	2.5	2.5	0.0			
400–599	40	95.0	5.0	0.0	0.0	0.0			
≥600	31	100.0	0.0	0.0	0.0	0.0			
All hospitals—2020	260	42.8 <sup>°</sup>	29.7 <sup>a,b</sup>	15.0 <sup>a</sup>	5.1 <sup>ª</sup>	7.5 <sup>ª</sup>			
All hospitals—2017 <sup>3</sup>	692	43.0	20.9	19.8	5.4	11.1			

			G			
All hospitals—2015 <sup>5</sup>	299	41.6	16.6	16.8	11.2	13.8
All hospitals—2014 <sup>6</sup>	425	40.2	20.5	14.5	3.4	21.4
All hospitals—2013 <sup>7</sup>	411	38.4	13.4	17.5	4.7	25.9
All hospitals—2012 <sup>8</sup>	480	37.0	15.9	12.9	2.3	32.0
All hospitals—2011 <sup>9</sup>	562	38.7	11.1	11.7	1.9	36.7
All hospitals—2010 <sup>10</sup>	565	34.5	8.8	9.8	3.6	43.4
All hospitals-2008 <sup>12</sup>	527	35.9	4.9	6.2	2.2	50.9
All hospitals—2007 <sup>13</sup>	531	33.8	5.0	6.5	3.1	51.6
All hospitals—2006 <sup>14</sup>	457	32.3	2.5	7.5	3.6	54.0
All hospitals—2005 <sup>15</sup>	510	30.1	3.1	5.3	1.9	59.6

<sup>a</sup>Uncorrected  $\chi^2$  = 148.4344, df = 24, design-based F(15.41, 3899.19) = 7.1497, P < 0.0001.

<sup>b</sup>95% confidence interval did not include previous-year point estimate.

Table 3. Primary Method of Maintenance Dose Distribution

xer		Decentralized Automated (eg, Automated Dispensing Cabinets)	Centralized Manual (eg, Unit Dose)	Centralized Automated (eg Robot)	Decentralized Manual (eg, Satellite)
Characteristic	n	%	%	%	%
Staffed beds					
<50	55	63.6	34.5	1.8	0.0
50–99	33	78.8	21.2	0.0	0.0
100–199	35	91.4	5.7	0.0	2.9
200–299	31	77.4	12.9	6.5	3.2
300–399	41	65.9	12.2	19.5	2.4
400–599	42	76.2	14.3	9.5	0.0
≥600	32	78.1	0.0	18.8	3.1
All hospitals—2020	269	<b>74.5</b> <sup>a</sup>	20.1 <sup>ª</sup>	4.1 <sup>a,b</sup>	1.3 <sup>ª</sup>

All hospitals—2017 <sup>3</sup>	693	70.2	20.7	8.4	0.7		
All hospitals—2011 <sup>9</sup>	554	62.5	28.9	8.4	0.2		
All hospitals—2008 <sup>12</sup>	523	49.2	43.1	7.6	0.1		
All hospitals—2005 <sup>15</sup>	510	37.8	52.5	7.8	1.9		
All hospitals-2002 <sup>18</sup>	504	22.3	69.7	7.1	1.6		

<sup>a</sup>Uncorrected  $\chi^2$  = 50.1677, *df* = 18, design-based *F*(10.49, 2748.98) = 4.1149, *P* < 0.0001.

<sup>b</sup>95% confidence interval did not include previous-year point estimate.

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Table 4. Predominant Configuration of Automated Dispensing Cabin	nets	Predominate o	configuration
Characteristic	n	Nurse selects from multiple drugs in matrix drawer %	Only requested medication is available to nurse (eg, in lidded pockets) %
Staffed beds			
<50	48	29.2	70.8
50-99	30	23.3	76.7
100–199	35	25.7	74.3
200–299	31	12.9	87.1
300–399	41	12.2	87.8
400–599	41	14.6	85.4
≥600	32	9.4	90.6
All hospitals—2020	258	22.6 <sup>ª</sup>	77.4 <sup>ª</sup>
All hospitals—2017 <sup>3</sup>	668	29.9	70.1

All hospitals—2014 <sup>6</sup>	413	32.3	65.7				
All hospitals—2011 <sup>9</sup>	517	38.1	61.9				
All hospitals—2008 <sup>12</sup>	452	48.5	51.5				

L not include , <sup>a</sup>95% confidence interval did not include previous-year point estimate.

Table 5. Use of Machine-Readable Coding	Durin	g Dispensing	2	During Restocking of ADCs
Characteristic	n	%	n	%
Staffed beds				
<50	54	50.0	54	72.2
50–99	32	71.9	33	84.8
100–199	35	68.6	35	80.0
200–299	31	77.4	31	93.5
300–399	41	82.9	41	87.8
400–599	41	85.4	41	90.2
≥600	32	93.8	32	100.0
All hospitals—2020	266	66.3ª	267	81.4 <sup>b,c</sup>
All hospitals—2017 <sup>3</sup>	693	61.9	692	74.7
All hospitals—2014 <sup>6</sup>	417	44.8	407	62.1
All hospitals—2013 <sup>7</sup>	411	54.2	NS	NS
All hospitals—2012 <sup>8</sup>	479	47.3	NS	NS

All hospitals—2011 <sup>9</sup>	561	33.9	517	43.3		
All hospitals—2008 <sup>12</sup>	526	24.0	NS	NS		
All hospitals—2007 <sup>13</sup>	531	18.4	NS	NS		
All hospitals—2005 <sup>15</sup>	510	11.5	NS	NS		
All hospitals—2004 <sup>16</sup>	492	9.2	NS	NS		
All hospitals—2002 <sup>18</sup>	511	5.7	NS	NS		

Abbreviations: ADC, automated dispensing cabinet; NS, not surveyed.

<sup>a</sup>Uncorrected  $\chi^2$  = 21.7525, df = 6, design-based F(4.40, 1138.93) = 4.2696, P = 0.0013.

<sup>b</sup>Uncorrected  $\chi^2$  = 12.0858, *df* = 6, design-based *F*(4.43, 1152.83) = 2.3735, *P* = 0.0443.

<sup>c</sup>95% confidence interval did not include previous-year point estimate.

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Table 6. Technology Used During Sterile Product Preparation No IV workflow												
		Barcode scanning performed to verify ingredients <sup>a</sup>	Pictures or video of compounding process obtained	IV workflow management software used <sup>b</sup>	Gravimetrics used to verify dose/amount/volume	technologies used for sterile product preparation activities						
Characteristic	n	%	%	%	%	%						
Staffed beds	.0											
<50	55	14.5	16.4	7.3	0.0	78.2						
50–99	33	24.2	21.2	15.2	0.0	60.6						
100–199	35	45.7	31.4	34.3	14.3	34.3						
200–299	30	46.7	30.0	23.3	0.0	40.0						
300-399	41	58.5	34.1	31.7	12.2	24.4						
400–599	41	58.5	41.5	48.8	9.8	19.5						
≥600	32	65.6	34.4	40.6	18.8	21.9						
All hospitals—2020	267	33.8 <sup>c</sup>	25.3 <sup>d</sup>	21.3 <sup>e</sup>	5.0 <sup>f</sup>	52.7 <sup>g</sup>						
All hospitals—2019 <sup>1</sup>	503	35.7	19.5	19.8	4.4	56.4						
All hospitals—2018 <sup>2</sup>	760	31.6	14.1	16.4	5.4	59.9						
All hospitals—2017 <sup>3</sup>	691	26.9	12.5	12.8	5.5	64.0						

NS All hospitals-2014<sup>6</sup> 425 19.5 All hospitals-2011<sup>9</sup> 560 11.9 Abbreviations: IV, intravenous; NS, not surveyed. <sup>a</sup>Examples include Epic Dispense Prep, or internally developed system. <sup>b</sup>Examples include Baxter DoseEdge, MedKeeper, PharmacyKeeper. <sup>c</sup>Uncorrected  $\chi^2$  = 37.1929, *df* = 6, design-based *F*(4.46, 1158.41) = 7.0542, *P* < 0.0001. <sup>d</sup>95% confidence interval did not include previous-year point estimate. <sup>e</sup>Uncorrected  $\chi^2$  = 28.5893, df = 6, design-based F(4.48, 1165.81) = 5.3561, P = 0.0001. <sup>f</sup>Uncorrected  $\chi^2$  = 24.6760, *df* = 6, design-based *F*(2.39, 621.21) = 9.8617, *P* < 0.0001. <sup>g</sup>Uncorrected  $\chi^2$  = 52.8652, *df* = 6, design-based *F*(4.41, 1147.69) = 10.1855, *P* < 0.0001. <sup>h</sup>Not surveyed.

NS NS

NS

NS

6.5

NS

Table 7. Robotics Used During Sterile Product Preparation											
		ompounding device used nazardous products <sup>a</sup>		mpounding device used for hemotherapy <sup>b,c</sup>	IV sterile com	pounding robotics not used					
Characteristic	n	%	n	%	n	%					
Staffed beds	55	1.8	25	0.0	55	98.2					
50-99	33	3.0	18	5.6	33	93.9					
100-199	35	2.9	27	0.0	35	97.1					
200–299	30	0.0	25	0.0	30	100.0					
300–399	41	2.4	37	2.7	41	95.1					
400–599	41	19.5	36	0.0	41	80.5					
≥600	32	9.4	32	15.6	32	78.1					
All hospitals—2020	267	3.4 <sup>d</sup>	200	1.6 <sup>e</sup>	267	95.7 <sup>f</sup>					
All hospitals—2017 <sup>3</sup>	691	2.3	453	0.9	NS	NS					

All hospitals—2014 <sup>6</sup>	425	2.9	347	0.3	NS	NS					
All hospitals—2011 <sup>9</sup>	560	2.5	478	0.1	NS	NS					
All hospitals-2008 <sup>12</sup>	526	1.1	NS	NS	NS	NS					
Abbreviation: IV, intravenou	IS.										

<sup>a</sup>Examples include ARXIUM RIVA, Baxter IntelliFill i.v., Omnicell i.v. Station, Grifols KIRO SP.

<sup>b</sup>Examples include Omnicell i.v. Station Onco, Loccioni Apoteca Chemo, ARXIUM RIVA, Equashield Pro, Grifols KIRO Oncology.

<sup>c</sup>For those hospitals that prepare chemotherapy.

<sup>d</sup>Uncorrected  $\chi^2$  = 16.4906, *df* = 6, design-based *F*(4.15, 1078.73) = 3.1469, *P* = 0.0127.

<sup>e</sup>Uncorrected  $\chi^2$  = 15.4968, *df* = 6, design-based *F*(2.04, 393.37) = 7.5051, *P* = 0.0006.

<sup>h</sup>Uncorrected  $\chi^2$  = 18.5507, *df* = 6, design-based *F*(4.13, 1073.12) = 3.5569, *P* = 0.0063.

<sup>g</sup>Not surveyed.

 Table 8. Methods Used to Assess Staff Competence to Prepare Compounded Sterile Preparations

		Media Fill Testing	Gloved Fingertip Testing	Direct Periodic Observation of Garbing and Gloving	Direct Periodic Observation of Sterile Technique	Direct Periodic Observation of Cleaning and Disinfecting	Surface Testing Using Contact Plates to Assess Cleaning Effectiveness	In-house Testing Materials (eg, USP <797> Exam, Calculations Exam)	Commercial Education and Testing Materials (eg, ASHP, CriticalPoint)	End-Product Testing
Characteristic	n	%	%	%	%	%	%	%	%	%
Staffed beds	5									
<50	51	82.4	82.4	78.4	80.4	72.5	56.9	45.1	39.2	5.9
50–99	33	100.0	100.0	84.8	87.9	84.8	63.6	72.7	51.5	21.2
100–199	35	97.1	94.3	94.3	88.6	85.7	71.4	57.1	60.0	14.3
200–299	30	100.0	100.0	100.0	90.0	90.0	80.0	60.0	73.3	23.3
300–399	41	97.6	97.6	92.7	87.8	80.5	85.4	70.7	58.5	26.8
400–599	41	100.0	100.0	85.4	92.7	82.9	85.4	82.9	63.4	31.7
≥600	32	100.0	100.0	93.8	87.5	81.3	81.3	81.3	75.0	34.4
All hospitals—2020	263	93.1 <sup>ª</sup>	92.5 <sup>b,c</sup>	87.4 <sup>c,d</sup>	86.0 <sup>c</sup>	80.8 <sup>c</sup>	68.6 <sup>c,e</sup>	59.0 <sup>f</sup>	53.7 <sup>c,g</sup>	16.2 <sup>h</sup>

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All hospitals—2017 <sup>3</sup>	687	89.0	85.6	75.8	76.8	74.5	75.7	57.2	43.4	18.8
All hospitals—2014 <sup>6</sup>	419	85.1	66.9	71.9	NS	71.3	71.5	NS	NS	NS
Abbreviation: NS, not surveyed; USP <797>, United States Pharmacopeia chapter 797.										
<sup>a</sup> Uncorrected $\chi^2$ = 25.45	16 <i>, df</i> = 6, d	esign-based F	2.52, 644.16) =	= <b>7</b> .1164, P = 0.00	03.					
<sup>b</sup> Uncorrected $\chi^2$ = 22.22	.65 <i>, df</i> = 6, d	esign-based F	2.51, 643.63) =	= 6.2449 <i>, P</i> = 0.00	08.					
<sup>c</sup> 95% confidence interva	al did not inc	lude previous	-year point est	imate.						
<sup>d</sup> Uncorrected $\chi^2$ = 14.82	.04, <i>df</i> = 6, d	esign-based F	(4.18, 1070.64)	= 3.0537, P = 0.0	147.					
<sup>e</sup> Uncorrected $\chi^2$ = 13.97	17, <i>df</i> = 6, d	esign-based F	(4.41, 1128.59)	= 2.7033, P = 0.0	248.					
<sup>f</sup> Uncorrected $\chi^2$ = 17.19	51, <i>df</i> = 6, de	esign-based F(	4.42, 1131.34)	= 3.3056, <i>P</i> = 0.0	081.					
<sup>g</sup> Uncorrected $\chi^2$ = 15.93	94, <i>df</i> = 6, d	esign-based F	4.45, 1139.03)	= 3.0360, P = 0.0	132.					
<sup>h</sup> Uncorrected $\chi^2 = 15.88$	24, <i>df</i> = 6, d	esign-based F	(4.47, 1144.41)	= 2.8978, P = 0.0	169.					

Table 9. Environmental Sampling Program and Report	Environmental san	ppling program (location, methods, ction levels, follow-up) used		pounding environmental sampling h organization's quality reporting pathways
Characteristic	n	%	n	%
Staffed beds				
<50	54	72.2	54	42.6
50-99	33	78.8	33	72.7
100–199	35	91.4	35	68.6
200–299	30	93.3	30	70.0
300-399	41	92.7	41	75.6
400–599	41	97.6	41	75.6
≥600	32	96.9	32	68.8
All hospitals—2020	266	83.6 <sup>a,b</sup>	266	61.0 <sup>b,c</sup>
All hospitals—2017 <sup>3</sup>	687	71.0	688	44.3

<sup>a</sup>Uncorrected  $\chi^2$  = 19.0081, *df* = 6, design-based *F*(4.38, 1133.24) = 3.7873, *P* = 0.0034.

<sup>b</sup>95% confidence interval did not include previous-year point estimate.

<sup>c</sup>Uncorrected  $\chi^2$  = 21.8388, *df* = 6, design-based *F*(4.45, 1152.02) = 4.2317, *P* = 0.0014.

Table 10. USP <800> Compliance and Gaps <sup>a</sup>	
Characteristic	%
Compliance ( <i>n</i> = 261)	
Fully compliant with all sections	31.1
Not yet fully compliant but working on compliance	63.1
Not fully compliant and not working on compliance	5.7
Gaps <sup>a</sup> ( <i>n</i> = 169)	
Facilities/engineering controls	51.5
Personnel training	50.8
Hazardous drug list/policies and procedures/assessment of risk	45.9
Drug storage	35.1
Garb/PPE	22.9
Closed system drug-transfer devices	17.6
Use of deactivating agent during cleaning	16.5
Abbreviation: PPE, personal protective equipment; USP <800>, United States Pharmaceopeia chapter 800.	

<sup>a</sup>For those hospital not currently fully compliant with USP <800>.

Table 11. Outsourcing of Compounded		Patient-specific CSPs prepared by 503A pharmacy	Non-patient-specific CSPs prepared by 503B pharmacy	CSPs not outsourced
Characteristic	n	%	%	%
Staffed beds				
<50	54	20.4	53.7	37.0
50–99	32	21.9	65.6	21.9
100–199	34	32.4	85.3	14.7
200–299	30	40.0	86.7	6.7
300–399	40	42.5	92.5	5.0
400–599	41	46.3	95.1	2.4
≥600	31	45.2	90.3	3.2
All hospitals—2020	262	29.5	72.5 <sup>ª</sup>	21.0 <sup>b</sup>
All hospitals—2018 <sup>2</sup>	761	27.0	68.6	24.5

Abbreviation: CSP, compounded sterile preparation. <sup>a</sup>Uncorrected  $\chi^2$  = 35.0178, *df* = 6, design-based *F*(4.38, 1116.77) = 6.9486, *P* < 0.0001.

<sup>b</sup>Uncorrected  $\chi^2$  = 28.1128, *df* = 6, design-based *F*(4.36, 1111.33) = 5.6465, *P* = 0.0001.

Table 12. Strategies to Evaluate External Compounded Sterile Preparation Outsourcers

		Confirmation of licensure in state	Availability of GPO contract with 503B outsourcer	Establish a contract with the pharmacy (legal review)	Receive (at a minimum) quarterly quality assurance documentation	Evaluation of any FDA-issued Form 483 reports and FDA reports to state boards of pharmacy	Vetting completed at health- system and/or corporate level	Signed attestation that outsourcer follows USP <797> and current good manufacturing practices	pharmacy inspections of outsourcing pharmacy and corrective actions	Outsourcing Sterile Products Preparation Vendor Assessment Tool	Complete a site validation visit to the outsourcing pharmacy	No method used
Characteristic	n	%	%	%	%	%	%	%	%	%	%	%
Staffed beds	<u>8</u>											
<50	33	51.5	45.5	51.5	21.2	27.3	36.4	45.5	24.2	27.3	6.1	3.0
50–99	25	80.0	52.0	60.0	44.0	44.0	44.0	44.0	32.0	24.0	16.0	0.0
100–199	28	71.4	67.9	46.4	60.7	50.0	46.4	28.6	39.3	28.6	28.6	0.0
200–299	27	70.4	59.3	55.6	66.7	63.0	48.1	48.1	40.7	33.3	29.6	0.0
300–399	37	70.3	73.0	67.6	67.6	56.8	59.5	40.5	35.1	48.6	40.5	0.0
400–599	40	82.5	72.5	67.5	72.5	60.0	65.0	37.5	47.5	52.5	40.0	0.0

\*

≥600	29	86.2	55.2	69.0	75.9	86.2	62.1	37.9	58.6	41.4	58.6	0.0
All hospitals—2020	219	68.0 <sup>ª</sup>	58.3	55.6	49.9 <sup>b</sup>	47.1 <sup>c</sup>	46.7	40.6	35.1	32.5	23.5 <sup>d</sup>	0.9
All hospitals-2018 <sup>2</sup>	583	70.7	NS	59.9	51.8	48.3	NS	42.9	36.5	29.0	30.4	1.5

Abbreviations: FDA, Food and Drug Administration; GPO, group purchasing organization. <sup>a</sup>Uncorrected  $\chi^2$  = 13.2462, *df* = 6, design-based *F*(4.44, 942.14) = 2.3111, *P* = 0.0494.

<sup>b</sup>Uncorrected  $\chi^2$  = 35.3053, *df* = 6, design-based *F*(4.43, 940.21) = 6.0518, *P* < 0.0001.

<sup>c</sup>Uncorrected  $\chi^2$  = 19.9237, *df* = 6, design-based *F*(4.44, 940.91) = 3.4240, *P* = 0.0065.

<sup>d</sup>Uncorrected  $\chi^2$  = 24.0832, *df* = 6, design-based *F*(4.48, 949.40) = 4.0106, *P* = 0.0021.

<sup>e</sup>Not surveyed.

Table 13. Use of Smart Infusion Pumps	Smart infus	ion pumps	prescribed information f need to man	mps autopopulate l order and patient from EHR, eliminating ually select drug and rate during setup		Nurses manually document infusion data into health record/EHR	Infusion data autopopulate from smart pump to EHR
Characteristic	n	%	n	%	n	%	%
Staffed beds	3						
<50	54	79.6	43	2.3	43	95.3	4.7
50–99	32	84.4	27	14.8	27	85.2	14.8
100–199	34	91.2	31	16.1	31	83.9	16.1
200–299	30	96.7	29	24.1	29	72.4	27.6
300–399	40	100.0	40	25.0	40	72.5	27.5
400–599	41	95.1	39	15.4	39	82.1	17.9
≥600	31	100.0	31	22.6	31	80.6	19.4
All hospitals—2020	262	87.9 <sup>ª</sup>	240	13.4 <sup>b,c</sup>	240	85.1 <sup>c,d</sup>	14.9 <sup>c,d</sup>
All hospitals—2017 <sup>3</sup>	683	88.1	599	8.9	600	90.8	9.2
All hospitals—2014 <sup>6</sup>	426	80.5	368	8.7	364	90.5	9.5
All hospitals—2013 <sup>7</sup>	413	80.8	355	5.9	355	93.1	6.9

All hospitals—2012 <sup>8</sup>	480	77.0		NS	NS	NS	NS			
All hospitals-2011 <sup>9</sup>	561	67.9	NS	NS	NS	NS	NS			
All hospitals-2010 <sup>10</sup>	563	65.0	NS	NS	NS	NS	NS			
All hospitals—2009 <sup>11</sup>	550	56.2	NS	NS	NS	NS	NS			
All hospitals—2008 <sup>12</sup>	525	59.1	NS	NS	NS	NS	NS			
All hospitals—2007 <sup>13</sup>	531	41.1	NS	NS	NS	NS	NS			
All hospitals—2006 <sup>14</sup>	460	37.0	NS	NS	NS	NS	NS			
All hospitals—2005 <sup>15</sup>	510	32.2	NS	NS	NS	NS	NS			

Abbreviations: EHR, electronic health record; NS, not surveyed.

<sup>a</sup>Uncorrected  $\chi^2$  = 13.9717, *df* = 6, design-based *F*(4.24, 1080.41) = 2.7369, *P* = 0.0251.

<sup>b</sup>Uncorrected  $\chi^2$  = 15.0431, *df* = 6, design-based *F*(4.44, 1034.26) = 2.6254, *P* = 0.0282.

<sup>c</sup>95% confidence interval did not include previous-year point estimate.

<sup>d</sup>Uncorrected  $\chi^2$  = 13.5979, *df* = 6, design-based *F*(4.45, 1036.11) = 2.3883, *P* = 0.0430.

Table 14. Use of Data Analytics and Technology to Reduce Risk of Adverse Events or Suboptimal Outcomes

	12	Basic analytics from smart pumps, clinical decision support, and/or automation used in dispensing and compounding	Basic and advanced analytics in the form of artificial intelligence, machine learning, and predictive analytics used	Analytics not used
Characteristic	n	%	%	%
Staffed beds	0			
<50	54	50.0	0.0	50.0
50-99	32	81.3	3.1	15.6
100–199	34	70.6	5.9	23.5
200–299	30	93.3	0.0	6.7
300–399	40	90.0	5.0	5.0
400–599	41	87.8	4.9	7.3
≥600	31	90.3	9.7	0.0
All hospitals—2020	262	70.5 <sup>°</sup>	2.6 <sup>ª</sup>	26.9 <sup>ª</sup>

<sup>a</sup>Uncorrected  $\chi^2$  = 52.3918, *df* = 12, design-based *F*(7.89, 2010.70) = 5.9818, *P* < 0.0001.

Table 15. Level of Pharmacy Services Integration to Optimize Patient Care Transitions

		Not At All Integrated	Some Integration	Mostly Integrated	Seamless Integration
Characteristic	n	%	%	%	%
Staffed beds					
<50	53	43.4	49.1	7.5	0.0
50–99	32	31.3	53.1	12.5	3.1
100–199	34	29.4	50.0	20.6	0.0
200–299	30	13.3	70.0	16.7	0.0
300–399	40	20.0	65.0	15.0	0.0
400–599	41	17.1	63.4	17.1	2.4
≥600	31	6.5	64.5	29.0	0.0
All hospitals—2020	261	30.6 <sup>a</sup>	55.0 <sup>ª</sup>	13.8 <sup>ª</sup>	0.6 <sup>a</sup>

All -

<sup>a</sup>Uncorrected  $\chi^2$  = 26.4182, *df* = 18, design-based *F*(10.58, 2686.54) = 2.1971, *P* = 0.0136.

Table 16. Advanced Pharmacy Technicians Activities

	×	Purchasing	Billing	Controlled Substance System Management	340B Program Management	Information Technology System Management	Responsible for USP <795>, USP <797>, or USP <800> Compliance	Regulatory Compliance	Technician Supervising Other Technicians	Initiation of Medication Reconciliation (Obtaining List)	Checking Dispensing by Other Technicians (Tech-Check-Tech)	Medication Assistance Program Management	Facilitating Transitions of Care	Vaccine Administration	יווסטשיט די אושטיווי די אינאפאר ווו אנוכנא Clinic Intake, Vital Signs Assessmemt)
Characteristic	n	%	%	%	%	%	%	%	%	%	%	%	%	%	%
Staffed beds															
<50	50	90.0	78.0	56.0	50.0	26.0	30.0	24.0	26.0	14.0	28.0	12.0	2.0	8.0	2.0
50–99	32	90.6	81.3	50.0	43.8	40.6	50.0	53.1	25.0	28.1	34.4	15.6	18.8	3.1	0.0
100–199	33	100.0	69.7	57.6	48.5	45.5	60.6	54.5	45.5	39.4	36.4	30.3	15.2	6.1	0.0
200–299	29	100.0	86.2	37.9	41.4	65.5	41.4	27.6	48.3	44.8	27.6	31.0	13.8	0.0	3.4
300–399	40	97.5	60.0	42.5	55.0	62.5	35.0	42.5	47.5	57.5	27.5	27.5	30.0	0.0	7.5
400–599	40	97.5	75.0	65.0	65.0	72.5	50.0	60.0	47.5	47.5	25.0	40.0	40.0	5.0	7.5
≥600	31	96.8	58.1	80.6	58.1	71.0	54.8	51.6	83.9	58.1	35.5	51.6	45.2	6.5	9.7
All hospitals—2020	255	94.7 <sup>ª</sup>	75.6	53.5	49.4	44.3 <sup>b</sup>	43.0 <sup>c</sup>	39.6 <sup>d</sup>	37.4 <sup>e,f</sup>	32.0 <sup>g</sup>	30.6 <sup>f</sup>	22.9 <sup>f,h</sup>	14.6 <sup>i</sup>	5.0	2.5

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All hospitals—2018 <sup>26</sup>	681	95.1	74.8	54.2	NS	44.9	NS	NS	29.8	26.3	17.9	13.8	12.6	NS	NS
All hospitals—2016 <sup>4</sup>	370	96.2	73.9	60.7	NS	40.7	NS	NS	34.5	30.1	17.6	9.3	9.6	NS	NS
All hospitals—2014 <sup>6</sup>	424	95.2	81.3	61.4	NS	37.8	NS	NS	28.1	18.0	17.8	11.3	8.4	NS	NS
All hospitals-2013 <sup>7</sup>	410	93.6	76.2	54.8	NS	38.4	NS	NS	29.5	14.3	16.4	12.4	7.7	NS	NS
All hospitals—2012 <sup>8</sup>	475	93.8	80.0	61.2	NS	39.5	NS	NS	26.8	11.4	17.0	10.6	NS	NS	NS
All hospitals – 2011 <sup>9</sup>	547	94.0	79.0	56.0	NS	41.8	NS	NS	28.9	11.5	15.1	8.2	NS	NS	NS
All hospitals—2008 <sup>12</sup>	523	91.5	84.1	47.2	NS	36.8	NS	NS	32.3	NS	15.7	NS	NS	NS	NS

Abbreviations: NS, not surveyed; USP <795>/<797>/<800>, United States Pharmacopeia chapter 795/797/800.

<sup>a</sup>Uncorrected  $\chi^2$  = 10.4406, *df* = 6, design-based *F*(3.30, 819.28) = 2.7777, *P* = 0.0352.

<sup>b</sup>Uncorrected  $\chi^2$  = 28.4510, *df* = 6, design-based *F*(4.44, 1101.48) = 5.3684, *P* = 0.0002.

<sup>c</sup>Uncorrected  $\chi^2$  = 14.8736, *df* = 6, design-based *F*(4.46, 1104.90) = 2.8014, *P* = 0.0203.

<sup>d</sup>Uncorrected  $\chi^2$  = 22.2263, *df* = 6, design-based *F*(4.46, 1106.22) = 4.1733, *P* = 0.0015.

<sup>e</sup>Uncorrected  $\chi^2$  = 18.7811, *df* = 6, design-based *F*(4.45, 1103.95) = 3.5401, *P* = 0.0051.

<sup>f</sup>95% confidence interval did not include previous-year point estimate.

<sup>g</sup>Uncorrected  $\chi^2$  = 27.5050, *df* = 6, design-based *F*(4.46, 1105.62) = 5.1041, *P* = 0.0002.

<sup>h</sup>Uncorrected  $\chi^2$  = 16.4352, *df* = 6, design-based *F*(4.48, 1111.38) = 3.0477, *P* = 0.0127.

<sup>i</sup>Uncorrected  $\chi^2$  = 30.1397, *df* = 6, design-based *F*(4.50, 1115.26) = 5.3812, *P* = 0.0001.

Table 17. Outpatient Cli	nic Operatic	ons	Types of clinics pharmacists participate in													
ç	2,2	Pharmacist work in ambulatory or primary care clinics		Anti-Coagulation	Oncology	Medication Therapy Management Services	Diabetes	Family Medicine	Cardiovascular Disease (eg, Lipids, Hypertension, CHF)	Infectious Disease (eg, HIV, Hepatitis)	Solid Organ Transplant	Pain & Palliative Care	เททพนกอเอยูy เeg, นเ, Rheumatology, Dermatology, Neurology)	Pharmacogenomics		
Characteristic	n	%	n	%	%	%	%	%	%	%	%	%	%	%		
Staffed beds																
<50	53	30.2	53	15.1	5.7	11.3	9.4	7.5	5.7	0.0	0.0	0.0	0.0	0.0		
50–99	32	40.6	32	31.3	21.9	18.8	12.5	18.8	12.5	15.6	3.1	3.1	3.1	3.1		
100–199	34	50.0	34	23.5	26.5	23.5	14.7	17.6	11.8	11.8	8.8	8.8	8.8	5.9		
200–299	30	56.7	30	26.7	30.0	26.7	23.3	26.7	20.0	16.7	6.7	13.3	3.3	3.3		

						.C								
300–399	40	60.0	40	32.5	42.5	27.5	22.5	15.0	25.0	15.0	15.0	5.0	10.0	2.5
400–599	41	80.5	41	61.0	58.5	48.8	41.5	31.7	51.2	39.0	34.1	22.0	29.3	7.3
≥600	31	87.1	31	48.4	67.7	41.9	48.4	45.2	64.5	45.2	67.7	29.0	25.8	19.4
All hospitals—2020	261	46.2 <sup>a,b</sup>	261	25.8 <sup>c</sup>	23.2 <sup>d</sup>	21.3 <sup>e</sup>	16.9 <sup>f</sup>	16.8 <sup>g</sup>	15.8 <sup>h</sup>	11.6 <sup>i</sup>	8.4 <sup>j</sup>	6.5 <sup>k</sup>	6.1 <sup>1</sup>	3.3 <sup>m</sup>
All hospitals—2018 <sup>2</sup>	744	32.9	743	21.5	16.3	16.6	13.4	13.5	10.8	8.6	7.1	4.5	4.2	1.9
All hospitals—2016 <sup>4</sup>	391	29.0	391	19.1	13.3	12.1	8.4	9.2	7.6	NS	NS	3.6	NS	NS
All hospitals—2015 <sup>5</sup>	312	25.1	312	13.3	15.2	8.5	8.6	7.0	5.1	NS	NS	2.2	NS	NS
All hospitals—2014 <sup>6</sup>	425	31.4	425	16.8	14.9	9.6	7.3	6.7	5.4	NS	NS	3.0	NS	NS
All hospitals—2013 <sup>7</sup>	412	27.1	411	16.6	14.1	10.5	9.0	6.3	5.3	NS	NS	2.6	NS	NS
All hospitals—2010 <sup>10</sup>	560	18.1	556	11.0	9.7	6.2	4.6	3.1	1.1	NS	NS	2.6	NS	NS
All hospitals—2008 <sup>12</sup>	526	17.1	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS	NS
All hospitals—2006 <sup>14</sup>	460	19.2	460	10.7	8.1	3.9	5.1	2.3	2.5	NS	NS	2.3	NS	NS

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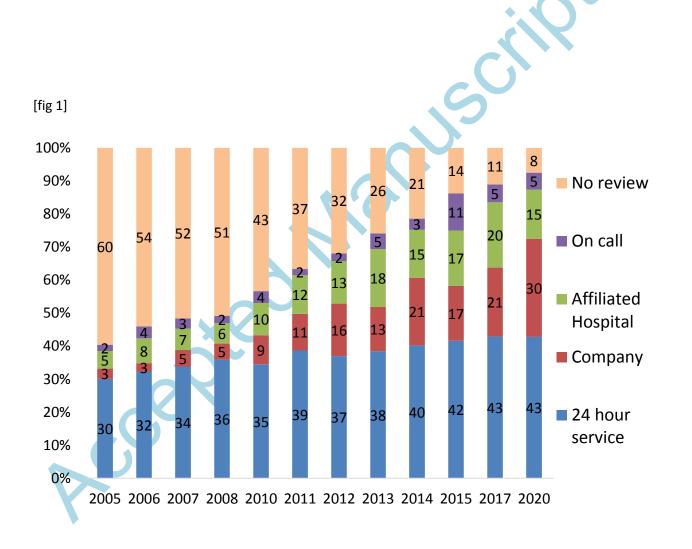
Abbreviations: CHF, congestive heart failure; GI, gastrointestinal; NS, not surveyed.

<sup>a</sup>Uncorrected  $\chi^2$  = 26.4703, *df* = 6, design-based *F*(4.42, 1121.74) = 5.0932, *P* = 0.0003.

<sup>b</sup>95% confidence interval did not include previous-year point estimate.

<sup>c</sup>Uncorrected  $\chi^2$  = 19.9265, *df* = 6, design-based *F*(4.47, 1136.42) = 3.7451, *P* = 0.0034.

<sup>d</sup>Uncorrected  $\chi^2 = 41.7524$ , df = 6, design-based F(4.46, 1133.86) = 7.7074, P < 0.0001. <sup>e</sup>Uncorrected  $\chi^2 = 16.5195$ , df = 6, design-based F(4.49, 1139.83) = 3.0894, P = 0.0117. <sup>f</sup>Uncorrected  $\chi^2 = 18.3748$ , df = 6, design-based F(4.52, 1147.28) = 3.4107, P = 0.0063. <sup>g</sup>Uncorrected  $\chi^2 = 15.1228$ , df = 6, design-based F(4.47, 1134.87) = 2.8141, P = 0.0197. <sup>h</sup>Uncorrected  $\chi^2 = 39.1714$ , df = 6, design-based F(4.52, 1148.18) = 7.1606, P < 0.0001. <sup>i</sup>Uncorrected  $\chi^2 = 34.4970$ , df = 6, design-based F(4.36, 1107.68) = 7.5306, P < 0.0001. <sup>j</sup>Uncorrected  $\chi^2 = 59.4503$ , df = 6, design-based F(4.36, 1108.24) = 12.7897, P < 0.0001. <sup>k</sup>Uncorrected  $\chi^2 = 22.9836$ , df = 6, design-based F(4.37, 1110.70) = 4.9526, P = 0.0004. <sup>i</sup>Uncorrected  $\chi^2 = 11.1926$ , df = 6, design-based F(4.38, 1111.89) = 2.3864, P = 0.0440.



scill

[fig 2]

