

Concise Communication

Pathogens attributed to central-line-associated bloodstream infections in US acute-care hospitals during the first year of the coronavirus disease 2019 (COVID-19) pandemic

Lindsey M. Weiner-Lastinger MPH¹ , Kathryn Haass MPH¹, Cindy Gross BS^{1,2}, Denise Leaptrot MSA^{1,2} , Emily Wong MPH^{1,3}, Hsiu Wu MD¹ and Margaret A. Dudeck MPH¹

¹Centers for Disease Control and Prevention, Atlanta, Georgia, ²CACI, Atlanta, Georgia and ³Leidos, Atlanta, Georgia

Abstract

To assess potential changes in the pathogens attributed to central-line–associated bloodstream infections between 2019 and 2020, hospital data from the National Healthcare Safety Network were analyzed. Compared to 2019, increases in the proportions of pathogens identified as *Enterococcus faecalis* and coagulase-negative staphylococci were observed during 2020.

(Received 12 October 2021; accepted 6 January 2022; electronically published 8 February 2022)

During 2020, hospitals saw unprecedented increases in critically ill patients as coronavirus disease 2019 (COVID-19) spread across the United States. In response, hospitals were often required to modify their operations, services provided, and patient care practices. Several studies have documented an alarming increase in device utilization and healthcare-associated infections (HAIs) in the United States during 2020, particularly in intensive care units (ICUs)²⁻⁴; however, studies evaluating changes in HAI pathogens during the pandemic have been limited to a small number of facilities. To assess changes in the common pathogens reported from central-line-associated bloodstream infections (CLABSIs) at the national level between 2019 and 2020, we examined data reported to the National Healthcare Safety Network (NHSN) by acute-care hospitals.

Methods

For each CLABSI, hospitals are required to report 1–3 pathogens and select antimicrobial susceptibility results to NHSN.⁵ CLABSI pathogens identified in adult ICUs and wards in 2019 and 2020 were analyzed. "Wards" included all adult non–critical-care units such as step-down and mixed-acuity units, excluding inpatient rehabilitation units.

The 15 pathogens most frequently associated with CLABSIs in 2019 and 2020 were identified, and their frequencies and ranks within each location type were calculated. Pathogen distributions were also reviewed among the subset of hospitals that performed continuous HAI surveillance in both 2019 and 2020, with no

Author for correspondence: Lindsey M. Weiner-Lastinger, E-mail: Llastinger@cdc.

Cite this article: Weiner-Lastinger LM, et al. (2023). Pathogens attributed to centralline–associated bloodstream infections in US acute-care hospitals during the first year of the coronavirus disease 2019 (COVID-19) pandemic. Infection Control & Hospital Epidemiology, 44: 651–654, https://doi.org/10.1017/ice.2022.16 difference in results observed. Although COVID-19 patient status was an optional field for data entry in NHSN, the distribution of pathogens among COVID-19 ICU patients was assessed. Vancomycin resistance among *Enterococcus* (VRE) and methicillin resistance among *Staphylococcus aureus* (MRSA) were measured by calculating the percentage of tested pathogens that were resistant. A mid-P exact test result \leq .05 was used to identify significant differences.

Results

Most CLABSIs in 2019 (89.4%) and 2020 (89.1%) had a single pathogen identified, with no substantial change in the proportion of CLABSIs that were polymicrobial.

ICUs

In total, 7,675 ICU CLABSI pathogens were reported from 1,560 hospitals during 2019. The most common pathogens were *Candida* (29.3%), coagulase-negative staphylococci (CNS) (13.3%), and *Enterococcus faecium* (8.1%) (Table 1). In 2020, 12,635 pathogens were reported by 1,906 hospitals, and *Candida* (27.8%), CNS (18.2%), and *Enterococcus faecalis* (15.0%) were the 3 most frequently reported species.

A large increase in the proportion of ICU CLABSI pathogens identified as CNS and *E. faecalis* were noted in 2020 compared to 2019. The increase in absolute number of *E. faecalis* CLABSIs was widespread; 388 hospitals reported at least 1 *E. faecalis* ICU CLABSI pathogen in 2019, compared to 848 hospitals in 2020 (data not shown). The reporting of *E. faecalis* varied by month in 2020, with the proportion of pathogens identified as *E. faecalis* ranging from 8%–9% (January–March) to 17%–18% (November and December). Little variation by month was observed in 2019, when

© The Author(s), 2022. Published by Cambridge University Press on behalf of The Society for Healthcare Epidemiology of America. This is an Open Access article, distributed under the terms of the Creative Commons Attribution licence (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted re-use, distribution and reproduction, provided the original article is properly cited.



Table 1. Frequency and Distribution of the 15 CLABSI Pathogens Most Frequently Reported to the NHSN from Adult ICUs in 2019 and 2020

		2019			2020		
Pathogen	No.	%	Rank	No.	%	Rank	
All <i>Candida</i> spp	2,246	29.3	1	3,513	27.8	1	
Coagulase-negative staphylococci	1,023	13.3	2	2,296	18.2	2	
Enterococcus faecium	622	8.1	3	787	6.2	5	
Staphylococcus aureus	613	8.0	4	882	7.0	4	
Enterococcus faecalis	590	7.7	5	1,895	15.0	3	
Selected <i>Klebsiella</i> spp ^{a,b}	432	5.6	6	541	4.3	6	
Escherichia coli	306	4.0	7	333	2.6	8	
Enterobacter spp ^b	263	3.4	8	228	1.8	9	
Pseudomonas aeruginosa	235	3.1	9	340	2.7	7	
Serratia spp	179	2.3	10	196	1.6	11	
Other <i>Enterococcus</i> spp ^c	135	1.8	11	205	1.6	10	
Acinetobacter spp	101	1.3	12	120	0.9	13	
Bacteroides spp	85	1.1	13	102	0.8	14	
Yeast, not specified	81	1.1	14	154	1.2	12	
Proteus spp	74	1.0	15	89	0.7	16	
Viridans group streptococci	55	0.7	16	93	0.7	15	
Other pathogen	635	8.3		861	6.8		
Total	7,675	100.0		12,635	100.0		

Note. CLABSI, central-line-associated bloodstream infection; NHSN, National Healthcare Safety Network; ICU, intensive care unit.

the proportion of *E. faecalis* pathogens ranged from 6% to 9% for almost all months in the year.

In 2019, 5.9% of tested *E. faecalis* were resistant to vancomycin (VRE); the resistance percentage was significantly lower in 2020, at 3.0% (Table A1).

Wards

In total, 1,821 hospitals reported 14,508 CLABSI pathogens from wards in 2019, of which *Candida* (12.1%), *S. aureus* (11.8%), and *Escherichia coli* (11.5%) were the most frequently reported (Table 2). In 2020, 1,848 hospitals reported 13,943 pathogens, and CNS replaced *E. coli* to become the third most common pathogen (11.1%). Increases were observed in the proportion of pathogens in 2020 that were CNS and *E. faecalis* compared to 2019. The percentage of *E. faecalis* that were resistant to vancomycin was significantly lower in 2020 than 2019 (5.2% vs 7.6%) (Table A1).

CLABSIs in COVID-19 ICU patients

Data on COVID-19 patient status were available for 4,232 (33.5%) of ICU CLABSI pathogens, with 2,787 (65.9% of those with data) occurring in patients with confirmed or suspected COVID-19 (Table A2). *Candida* (28.9%), *E. faecalis* (21.1%), and CNS (19.7%) were the 3 most frequently reported CLABSI pathogens among ICU patients with COVID-19.

Table 2. Frequency and Distribution of the 15 CLABSI Pathogens Most Frequently Reported Reported to NHSN from Adult Wards^a in 2019 and 2020

		2019			2020		
Pathogen	No.	%	Rank	No.	%	Rank	
All <i>Candida</i> spp.	1,758	12.1	1	1,822	13.1	1	
Staphylococcus aureus	1,710	11.8	2	1,681	12.1	2	
Escherichia coli	1,666	11.5	3	1,502	10.8	4	
Coagulase-negative staphylococci	1,453	10.0	4	1,544	11.1	3	
Selected <i>Klebsiella</i> spp ^{b,c}	1,300	9.0	5	1,187	8.5	5	
Enterococcus faecalis	957	6.6	6	1,081	7.8	6	
Enterococcus faecium	880	6.1	7	809	5.8	7	
Viridans group streptococci	707	4.9	8	647	4.6	8	
Pseudomonas aeruginosa	691	4.8	9	573	4.1	9	
Enterobacter spp ^c	566	3.9	10	416	3.0	10	
Serratia spp.	225	1.6	11	199	1.4	13	
Other <i>Enterococcus</i> spp ^d	213	1.5	12	218	1.6	11	
Bacteroides spp	166	1.1	13	216	1.5	12	
Stenotrophomonas maltophilia	146	1.0	14	127	0.9	15	
Proteus spp	137	0.9	15	130	0.9	14	
Other pathogen	1,933	13.3		1,791	12.8		
Total	14,508	100.0		13,943	100		

Note. CLABSI, central-line–associated bloodstream infection; NHSN, National Healthcare Safety Network.

Discussion

This paper describes the CLABSI pathogens commonly isolated during the first year of the COVID-19 pandemic, using data from almost all US hospitals.⁶ Our results showed that the common pathogens among COVID-19 ICU patients at a national level, particularly *E. faecalis* and CNS, were consistent with results from local studies.^{7–10}

Even though the stark increase in *E. faecalis* pathogens reported in 2020 was unexpected, an increase in *Enterococcus* BSIs in 2020, compared to 2018–2019, was also observed in a hospital in northern Italy.⁷ These results, along with substantially higher proportions of *E. faecalis* identified in November and December 2020, during which a large number of COVID-19 hospitalizations occurred in the United States, ¹¹ suggest that COVID-19 patients and/or patients hospitalized during times of heightened COVID-19 burden may be particularly susceptible to CLABSIs caused by *E. faecalis*. The reasons for this are unclear, but several local studies from the United States and Italy identified *Enterococcus* as a common BSI pathogen among COVID-19 patients.^{7–10}

In addition to host factors, changes in the amount and overall pattern of antibiotic use for hospitalized patients in 2020 could have contributed to a rise in *E. faecalis*. Giacobbe et al⁸ reported

^aIncludes *K. oxytoca* and *K. pneumoniae*. For 2020, this group also includes *K. aerogenes*. ^b*K. aerogenes* (formerly known as *Enterobacter aerogenes*) is classed in the *Enterobacter* spp group in 2019 and the *Klebsiella* spp group in 2020.

^cThe group 'other *Enterococcus* spp' combines enterococci identified to the species level, excluding *E. faecium* and *E. faecalis*, and enterococci for which the species was not reported.

^aIncludes all non-critical-care unit types, including specialty care areas, step-down units, and mixed-acuity units.

^bIncludes *K. oxytoca* and *K. pneumoniae*. For 2020, this group also includes *K. aerogenes*. ^c*K. aerogenes* (formerly known as *Enterobacter aerogenes*) is classed in the *Enterobacter* spp. group in 2019 and the *Klebsiella* spp. group in 2020.

^dThe group 'other *Enterococcus* spp' combines enterococci identified to the species level, excluding *E. faecium* and *E. faecalis*, and enterococci for which the species was not reported.

that almost all COVID-19 patients in their 1,200-bed hospital were treated with a cephalosporin, and an increase in antibiotic use, especially ceftriaxone, was observed in 2 large US hospital cohorts during 2020. The additional antibiotic use in 2020, or other antibiotic effects, may have contributed to changes in the selection pressure for pathogens in hospitals that favored the growth of *E. faecalis*. Interestingly, increases in *E. faecium* were not observed during 2020. Additional research is needed to understand the mechanism behind the increases in *E. faecalis* and to more fully explore the impact of the COVID-19 pandemic on pathogens and antimicrobial resistance patterns in hospitals.

CNS was the second most reported pathogen for ICU CLABSIs in 2019 and 2020, with a marked increase in 2020. Due to the surge of case load and relative scarce healthcare resources early in the pandemic, inadequate adherence to aseptic blood culture collection technique may have resulted in some increases in CNS isolates. However, the NHSN CLABSI definition includes stipulations to reduce the impact of contamination by offering separate criteria for common commensals and known pathogens; thus, the increase in CNS CLABSIs during 2020 is unlikely to have been caused by contamination alone.

This study had several limitations. All data from adult locations were analyzed, including data from pediatric patients housed in adult locations at the time of their infection. The CMS granted a reporting exception for the first half of 2020, leading some hospitals to pause HAI reporting to the NHSN. Any underestimation in the number of pathogens during 2020 is assumed to be minimal due to the high volume of reporting that continued throughout the year. It was optional for hospitals to report patient COVID-19 status to the NHSN. Given the limited responses available, we acknowledge that the pathogen distribution among COVID-19 ICU patients is not representative of all COVID-19 ICU patients who experienced an HAI.

Compared to the pre–COVID-19 period, we identified national increases during 2020 in the proportion of CLABSIs caused by *E. faecalis* and CNS. Infection prevention professionals are encouraged to review the common pathogens and antimicrobial resistance patterns in their hospitals and jurisdictions to identify opportunities to strengthen HAI prevention and antimicrobial stewardship efforts.

Supplementary material. To view supplementary material for this article, please visit https://doi.org/10.1017/ice.2022.16

Acknowledgments. The findings and conclusions in this article are those of the authors and do not necessarily represent the official position of the US Centers for Disease Control and Prevention.

Financial support. No financial support was provided relevant to this article.

Conflicts of interest. All authors report no conflicts of interest relevant to this article.

References

- Rebmann T, Alvino RT, Holdsworth JE. Availability and crisis standards of care for personal protective equipment during fall 2020 of the COVID-19 pandemic: a national study by the APIC COVID-19 task force. *Am J Infect Control* 2021;49:657–662.
- Patel PR, Weiner-Lastinger LM, Dudeck MA, et al. Impact of COVID-19 pandemic on central-line–associated bloodstream infections during the early months of 2020, National Healthcare Safety Network. Infect Control Hosp Epidemiol 2021. doi: 10.1017/ice.2021.108.
- Fakih MG, Bufalino A, Sturm L, et al. Coronavirus disease 2019 (COVID-19) pandemic, central-line-associated bloodstream infection (CLABSI), and catheter-associated urinary tract infection (CAUTI): The urgent need to refocus on hardwiring prevention efforts. *Infect Control Hosp Epidemiol* 2022;43:26–31.
- 4. Weiner-Lastinger L, Pattabiraman V, Konnor RY, et al. The impact of coronavirus disease 2019 (COVID-19) on healthcare-associated infections in 2020: A summary of data reported to the National Healthcare Safety Network. *Infect Control Hosp Epidemiol* 2021;1–14.
- National Healthcare Safety Network. Patient safety component manual. Centers for Disease Control and Prevention website. https://www.cdc.gov/nhsn/pdfs/pscmanual/pcsmanual_current.pdf. Updated January 2021. Accessed September 30, 2021.
- Hospital-acquired condition reduction program. Centers for Medicare and Medicaid Services website. https://www.cms.gov/Medicare/Medicare-Feefor-Service-Payment/AcuteInpatientPPS/HAC-Reduction-Program. Updated December 2021. Accessed December 16, 2021.
- Bonazzetti C, Morena V, Giacomelli A, et al. Unexpectedly high frequency
 of enterococcal bloodstream infections in coronavirus disease 2019 patients
 admitted to an Italian ICU: an observational study. Crit Care Med 2021;
 49:e31–e40.
- Giacobbe DR, Battaglini D, Ball L, et al. Bloodstream infections in critically ill patients with COVID-19. Eur J Clin Invest 2020;50:e13319.
- Cataldo MA, Tetaj N, Selleri M, et al. Incidence of bacterial and fungal bloodstream infections in COVID-19 patients in intensive care: an alarming "collateral effect." J Glob Antimicrob Resist 2020;23:290–291.
- DeVoe C, Segal M, Wang L, et al. Increased rates of secondary bacterial infections, including Enterococcus bacteremia, in patients hospitalized with coronavirus disease 2019 (COVID-19). Infect Control Hosp Epidemiol 2021. doi: 10.1017/jce.2021.391.
- COVID Data Tracker. Centers for Disease Control Prevention website. https://covid.cdc.gov/covid-data-tracker/#hospitalizations. Accessed June 15, 2021.
- 12. The intersection of antibiotic resistance (AR), antibiotic use (AU), and COVID-19. Department of Health and Human Services website. https://www.hhs.gov/sites/default/files/antibiotic-resistance-antibiotic-use-covid-19-paccarb.pdf. Published February 10, 2021. Accessed June 28, 2021.
- LeRose J, Sandhu A, Polistico J, et al. The impact of COVID-19 response on central-line-associated bloodstream infections and blood culture contamination rates at a tertiary-care center in greater Detroit area. *Infect Control Hosp Epidemiol* 2021;42:997–1000.
- Sapiano MR, Dudeck MA, Soe M, et al. Impact of coronavirus disease 2019 (COVID-19) on US hospitals and patients, April–July 2020. Infect Control Hosp Epidemiol 2022;43:32–39.
- Wu H, Soe MM, Konnor R, et al. Hospital capacities and shortages of healthcare resources among US hospitals during COVID-19 pandemic, National Healthcare Safety Network, March 27–July 14, 2020. Infect Control Hosp Epidemiol 2021. doi: 10.1017/ice.2021.280.

APPENDIX

Table A1. The Percentage of CLABSI Pathogens Resistant (%R) to Vancomycin (VRE) or Methicillin (MRSA) in 2019 and 2020, by Location

		2019			2020		
Location, Pathogen	No. Tested	% Tested	% R ^a	No. Tested	% Tested	% R ^a	<i>P</i> Value
Adult ICUs							
Enterococcus faecalis	525	90.0	5.9	1,681	88.7	3.0	.004
E. faecium	574	92.3	81.5	713	90.6	77.7	.091
Staphylococcus aureus	531	86.6	47.8	758	85.9	47.6	.941
Adult Wards ^b							
E. faecalis	854	89.2	7.6	939	86.9	5.2	.039
E. faecium	811	92.2	72.5	713	88.1	70.3	.336
S. aureus	1,498	87.6	46.2	1,432	85.2	48.3	.249

Note. ICU, intensive care unit; CLABSI, central-line–associated bloodstream infection; VRE, vancomycin-resistant *Enterococcus*; MRSA, methicillin-resistant *Staphylococcus aureus*.
^aPercent resistance (%R) is measured for VRE or MRSA, as appropriate. VRE is defined as *Enterococcus* resistant to vancomycin. MRSA is defined as *S. aureus* resistant to methicillin, oxacillin, or cefoxitin.

Table A2. Frequency and Distribution of the CLABSI Pathogens Most Frequently Reported Among Adult ICU Patients With Confirmed or Suspected COVID-19

		•	
Pathogen	No.	%	Rank
All <i>Candida</i> spp.	806	28.9	1
Enterococcus faecalis	589	21.1	2
Coagulase-negative staphylococci	550	19.7	3
Staphylococcus aureus	186	6.7	4
Enterococcus faecium	148	5.3	5
Selected Klebsiella sppa	77	2.8	6
Pseudomonas aeruginosa	60	2.2	7
Escherichia coli	48	1.7	8
Yeast, not specified	42	1.5	9
Other <i>Enterococcus</i> spp ^b	37	1.3	10
Serratia spp	29	1.0	11
Acinetobacter spp	20	0.7	12
Enterobacter spp	19	0.7	13
Viridans group streptococci	16	0.6	14
Stenotrophomonas maltophilia	14	0.5	15
Other pathogen	146	5.2	
Total	2,787	100.0	

Note. ICU, intensive care unit; CLABSI, central-line–associated bloodstream infection; COVID-19, coronavirus disease 2019.

 $^{^{\}rm b}$ lncludes all non-critical-care unit types, including specialty care areas, step-down units, and mixed-acuity units.

^aIncludes K. oxytoca, K. pneumoniae, and K. aerogenes.

^bThe group 'other *Enterococcus* spp' combines enterococci identified to the species level, excluding *E. faecium* and *E. faecalis*, and enterococci for which the species was not reported.