



Trends in *Clostridioides difficile* infection rates in Canadian hospitals during the coronavirus disease 2019 (COVID-19) pandemic

Kelly B. Choi MSc¹ , Tim Du MSc² , Anada Silva MSc¹ , George R. Golding PhD², Linda Pelude MSc¹, Robyn Mitchell MSc¹, Wallis Rudnick PhD¹ , Romeo Hizon BSc², Ghada N Al-Rawahi MD³ , Blanda Chow MSc⁴, Ian Davis MD⁵, Gerald A. Evans MD⁶ , Charles Frenette MD⁷, Jennie Johnstone MD⁸ , Pamela Kibsey MD⁹, Kevin C. Katz MD¹⁰, Joanne M. Langley MD^{11,12} , Bonita E. Lee MD¹³ , Yves Longtin MD¹⁴ , Dominik Mertz MD¹⁵ , Jessica Minion MD¹⁶ , Michelle Science MD¹⁷, Jocelyn A. Srigley MD¹⁸, Paula Stagg RNMN¹⁹, Kathryn N. Suh MD²⁰, Nisha Thampi MD²¹ , Alice Wong MD²² , Jeannette L. Comeau MD¹¹ , Susy S. Hota MD²³ and for the Canadian Nosocomial Infection Surveillance Program (CNISP)

¹Public Health Agency of Canada, Ottawa, Ontario, Canada, ²National Microbiology Laboratory, Winnipeg, Manitoba, Canada, ³British Columbia Children's Hospital, Vancouver, British Columbia, Canada, ⁴Alberta Health Services, Calgary, Alberta, Canada, ⁵Queen Elizabeth II Health Sciences Centre, Halifax, Nova Scotia, Canada, ⁶Kingston Health Sciences Centre, Kingston, Ontario, Canada, ⁷McGill University Health Centre, Montréal, Quebec, Canada, ⁸Sinai Health, Toronto, Ontario, Canada, ⁹Royal Jubilee Hospital, Victoria, British Columbia, Canada, ¹⁰North York General Hospital, Toronto, Ontario, Canada, ¹¹Dalhousie University, Halifax, Nova Scotia, Canada, ¹²IWK Health Centre, Halifax, Nova Scotia, Canada, ¹³Stollery Children's Hospital, Edmonton, Alberta, Canada, ¹⁴Jewish General Hospital, Montréal, Quebec, Canada, ¹⁵Hamilton Health Sciences, Hamilton, Ontario, Canada, ¹⁶Regina General Hospital, Regina, Saskatchewan, Canada, ¹⁷The Hospital for Sick Children, Toronto, Ontario, Canada, ¹⁸BC Children's & Women's Hospitals, Vancouver, British Columbia, Canada, ¹⁹Western Memorial Regional Hospital, Corner Brook, Newfoundland, Canada, ²⁰The Ottawa Hospital, Ottawa, Ontario, Canada, ²¹Children's Hospital of Eastern Ontario, Ottawa, Ontario, Canada, ²²Royal University Hospital, Saskatoon, Saskatchewan, Canada and ²³University Health Network, Toronto, Ontario, Canada

Abstract

The coronavirus disease 2019 (COVID-19) pandemic has placed significant burden on healthcare systems. We compared *Clostridioides difficile* infection (CDI) epidemiology before and during the pandemic across 71 hospitals participating in the Canadian Nosocomial Infection Surveillance Program. Using an interrupted time series analysis, we showed that CDI rates significantly increased during the COVID-19 pandemic.

(Received 25 May 2022; accepted 3 August 2022)

Changes in clinical practice during the coronavirus disease 2019 (COVID-19) pandemic, including the potential increased use of broad-spectrum antibiotics, may have exacerbated the burden of *Clostridioides difficile* infection (CDI).¹ Single-center studies have reported increased,² decreased,^{3,4} or unchanged CDI rates.^{5,6} Furthermore, the impact of the COVID-19 pandemic on rates of CDI in Canada has not yet been described. Using national surveillance data, we compared CDI rates, patient characteristics, and clinical outcomes before and during the COVID-19 pandemic in Canada.

Methods

Data collection and participating hospitals

Using standardized case definitions, we included patients with CDI identified from hospitals participating in the Canadian

Nosocomial Infection Surveillance Program (CNISP) between January 1, 2015, and June 30, 2021. Healthcare-associated (HA) and community-associated (CA) CDI rates were classified as previously described.⁷ Beginning January 2021, COVID-19 diagnosis status was collected for all CDI patients.

Statistical analysis

We calculated HA-CDI rates per 10,000 patient days and CA-CDI rates per 1,000 patient admissions. We used interrupted time series (ITS) analysis to assess changes in CDI rates between the prepandemic period (62 monthly periods from January 1, 2015, to February 29, 2020) and the COVID-19 pandemic period (16 periods from March 1, 2020 to June 30, 2021) by modeling monthly CDI rates, accounting for time trends and seasonality. Models were fitted using a generalized linear mixed model with Poisson distribution and log-link function with first-order autocorrelation. The outcome was monthly CDI counts. Models included an offset term (ie, the natural log of either patient days or patient admissions). Changes over time for HA- and CA-CDI rates by month were displayed graphically. Predictor variables were pandemic status (pre pandemic = 0 and pandemic = 1), West (British Columbia,

© The Author(s), 2022. Published by Cambridge University Press on behalf of The Society for Healthcare Epidemiology of America.

Author for correspondence: Dr. Susy S. Hota, MSc, MD, FRCPC, Infection Prevention and Control, University Health Network, 8th floor Room 102, 200 Elizabeth Street, Toronto, Ontario, Canada, M5G 2C4. E-mail: susy.hota@uhn.ca

Cite this article: Choi KB, et al. (2022). Trends in Clostridioides difficile infection rates in Canadian hospitals during the coronavirus disease 2019 (COVID-19) pandemic. Infection Control & Hospital Epidemiology, https://doi.org/10.1017/ice.2022.210



Fig. 1. Monthly HA-CDI and CA-CDI rates between January 2015 and June 2021. (a) Actual and predicted HA-CDI rates per 10,000 patient days. (b) Actual and predicted CA-CDI rates per 1,000 patient admissions. Note. Red dotted line represents the declaration of the pandemic. Quarterly denominators, proportional to the number of days in each month.

Alberta, Saskatchewan and Manitoba, Central; Ontario and Quebec) or East (Nova Scotia, New Brunswick, Prince Edward Island, Newfoundland and Labrador, and Nunavut) region, month and time (78 monthly periods from January 2015 to June 2021). Results were reported as incidence rate ratios (IRRs) comparing prepandemic and COVID-19 pandemic periods with 95% confidence intervals (CIs).

The χ^2 test was used for categorical variables and the Student t test or Wilcoxon rank-sum test was used for continuous variables to compare characteristics and clinical outcomes among CDI patients by COVID-19 status and pandemic period. All statistical analyses were conducted using SAS version 9.4 software (SAS Institute, Cary, NC) with a significance level of 0.05.

Results

We summarized CDI rates from 71 hospitals (38 adult hospitals, 24 mixed hospitals, and 9 pediatric standalone hospitals) between January 1, 2015, and June 30, 2021. All 71 hospitals participated in HA-CDI surveillance, and 60 (85%) participated in CA-CDI surveillance. Crude HA-CDI rates per 10,000 patient days increased by 4.5%, but this increase was not significant (P = .0896). Crude CA-CDI rates per 1,000 patient admissions increased significantly by 21.8% from the prepandemic period (January 1, 2019, to February 29, 2020) to the pandemic period (March 1, 2020, to June 30, 2021) (Supplementary Table 1).

Figure 1 displays the results of the ITS analyses of HA-CDI and CA-CDI rates across the entire surveillance period (January 1,

2015, to June 30, 2021). Before the COVID-19 pandemic, both HA-CDI and CA-CDI rates decreased over time. We modeled the monthly CDI rates and adjusting for region and month; the results indicated that pandemic status was a significant predictor for both the HA-CDI model (P = .0007) and the CA-CDI model (P < .0001). We detected a statistically significant rate increase of 11% of HA-CDI (IRR, 1.11; 95% CI, 1.04–1.18) and 25% of CA-CDI (IRR, 1.25; 95% CI, 1.14–1.37) during the pandemic period.

We did not detect significant differences in age, sex, temperature, leukocyte counts, albumin levels or severe outcomes (death, admission to the ICU or colectomy) between the prepandemic cohort (n = 23,315, January 1, 2015, to February 29, 2020) and the pandemic cohort (n = 6,214) (data not shown).

Among 1,576 CDI patients identified in 2021, data regarding COVID-19 status were reported for 1,244 (89%). Of those, 64 (5%) were coinfected with Severe Acute Respiratory Syndrome Coronavirus-2 (SARS-CoV-2). A significantly higher proportion of CDI patients with COVID-19 had HA-CDI (78%) compared to those without COVID-19 (66%; P = .0366). Both groups were similar in age, sex, and severe outcomes (Supplementary Table 2). Time to CDI positive test from admission, measured in days, was longer for COVID-19 patients (10.5 days) than non-COVID-19 patients (4.0 days; P = .0004).

Discussion

This study is the first Canadian multicenter research to assess the impact of COVID-19 on CDI. We have described CDI rate trends,

patient characteristics, and clinical outcomes before and during the COVID-19 pandemic for 71 hospitals across Canada.

Our previous surveillance data showed a steady decline in CDI rates before the pandemic.^{7,8} However, during the pandemic, we observed an increase in HA- and CA-CDI rates, whereas other jurisdictions reported varying changes in healthcare-associated infections.9 CDI and COVID-19 coinfection was uncommon (5%); however, COVID-19 patients were more likely to be associated with HA-CDI acquisition than non-COVID-19 patients. The increased HA-CDI rates in COVID-19 patients could be attributable to increased use of broad-spectrum antibiotics for clinical management of COVID-19.7,10 Changes in IPAC practices due to pandemic-related surge pressures may have also contributed to the observed increase. Additionally, COVID-19 disproportionately affected the elderly population in Canada during the early phases of the pandemic.^{11,12} The elderly are at higher risk of respiratory infections leading to empiric antibiotic treatment, resulting in increased risk for acquiring CDI.¹³ Furthermore, COVID-19 outbreaks in Canadian long-term care facilities (LTCFs) resulted in a surge of residents being admitted to hospitals at the beginning of the pandemic.¹⁴ LTCF residents are at higher risk of acquiring CDI, further contributing to increased HA-CDI rates.

Our results are similar to those of previous studies showing a slight increase in HA-CDI rates during the pandemic^{5,6}; however, neither prior study demonstrated statistical significance or controlled for time. Although our unadjusted rates were not significantly different when comparing prepandemic and pandemic periods, the ITS analysis showed a significant HA-CDI rate increase of 11% after adjusting for region and month.

Other studies have reported decreases in CDI rates during the COVID-19 pandemic. Single-center studies by Ponce-Alonso et al³ and Hazel et al⁴ reported significant reductions in CDI rates, citing enhanced infection control measures as a probable contributor with factors such as antimicrobial consumption remaining constant.

CA-CDI rates increased significantly from the prepandemic to pandemic periods. The increased CA-CDI rates among CNISP hospitals may be attributable to reduced accessibility and/or avoidance of healthcare facilities during the pandemic. Those seeking medical care were often assessed through virtual visits or emergency departments, which are suboptimal settings for determining antibiotic appropriateness, resulting in excessive or broad-spectrum antibiotic usage. Additionally, it has been reported that COVID-19 can affect the normal intestinal flora similar to treatment with antibiotics, causing a predisposition to CDI acquisition.¹⁵

Our findings show an increase in CDI rates during the COVID-19 pandemic. Considering the significant strain that COVID-19 has placed on the Canadian healthcare system, these data further our understanding of CDI epidemiology and assist in our ability to manage current and future pandemics.

Our study had several limitations. Data analysis was limited to 16 months of the pandemic; hence, longer-term impacts could not be assessed. Comparisons of prepandemic and pandemic rates are preliminary and should be interpreted with caution. Infection control policies and pandemic restrictions were dynamic throughout the pandemic period and may have impacted on monthly rates differentially. Additionally, our study did not assess changes in screening practices, antibiotic usage, or associations between CDI rates and COVID-19 rates across Canada or within the participating hospital sites. Such studies would have been beneficial to furthering our understanding of the impact of COVID-19 on CDI rates. Future studies should aim to elucidate factors that may have influenced changes in CDI rates during the pandemic.

Supplementary material. To view supplementary material for this article, please visit https://doi.org/10.1017/ice.2022.210

Acknowledgments. We gratefully acknowledge the contribution of the physicians, epidemiologists, infection control practitioners, and laboratory staff at each participating hospital. We thank the staff at Public Health Agency of Canada in the Centre for Communicable Diseases and Infection Control, Ottawa, Ontario: Cecilia McClellan, Joelle Cayen, and Jessica Bartoszko. We also thank Sean Ahmed of the the National Microbiology Laboratory, Winnipeg, Manitoba. CNISP is a collaborative effort between sentinel hospitals across Canada, the Public Health Agency of Canada's Centre for Communicable Diseases and Infection Control (CCDIC), the National Microbiology Laboratory (NML), and the Association of Medical Microbiology (AMMI) Canada.

Financial support. This work was supported by the Public Health Agency of Canada.

Conflict of Interest. Y.L. has received research grants from Becton Dickinson and Merck, unrelated to this work. S.S.H. received research funding from Finch Therapeutics Group for participating as a site principal investigator for a clinical trial. All authors report no conflicts of interest relevant to this article.

References

- Abelenda-Alonso G, Padullés A, Rombauts A, et al. Antibiotic prescription during the COVID-19 pandemic: a biphasic pattern. *Infect Control Hosp Epidemiol* 2020;41:1371–1372.
- Lewandowski K, Rosołowski M, Kaniewska M, et al. Clostridioides difficile infection in coronavirus disease 2019 (COVID-19): an underestimated problem? *Polish Arch Intern Med* 2021;131:121–127.
- Ponce-Alonso M, Sáez De La Fuente J, Rincón-Carlavilla A, et al. Impact of the coronavirus disease 2019 (COVID-19) pandemic on nosocomial *Clostridioides difficile* infection. *Infect Control Hosp Epidemiol* 2021;42: 406–410.
- Hazel K, Skally M, Glynn E, et al. The other "C": hospital-acquired Clostridioides difficile infection during the COVID-19 pandemic. Infect Control Hosp Epidemiol 2022;43:540–541.
- Luo Y, Grinspan LT, Fu Y, et al. Hospital-onset Clostridioides difficile infections during the COVID-19 pandemic. Infect Control Hosp Epidemiol 2021;42:1165–1166.
- Yadlapati S, Jarrett SA, Lo KB, Sweet J, Judge TA. Examining the rate of *Clostridioides* (formerly *Clostridium*) difficile infection pre– and post– COVID-19 pandemic: an institutional review. *Cureus* 2021;13:e20397.
- Healthcare-associated infections and antimicrobial resistance in Canadian acute-care hospitals, 2014–2018. Canada Commun Dis Rep 2020;46:99–112.
- Du T, Choi KB, Silva A, et al. Characterization of healthcare-associated and community-associated *Clostridioides difficile* infections among adults, Canada, 2015–2019. Emerg Infect Dis 2022;28:1128–1136.
- Weiner-Lastinger LM, Abner S, Edwards JR, et al. Antimicrobial-resistant pathogens associated with adult healthcare-associated infections: summary of data reported to the National Healthcare Safety Network, 2015–2017. *Infect Control Hosp Epidemiol* 2020;41:1–18.
- Langford BJ, So M, Raybardhan S, *et al.* Bacterial coinfection and secondary infection in patients with COVID-19: a living rapid review and meta-analysis. *Clin Microbiol Infect* 2020;26:1622–1629.
- 11. Flint AJ, Bingham KS, Iaboni A. Effect of COVID-19 on the mental health care of older people in Canada. *Int Psychogeriatrics* 2020;32:1.
- Hsu AT, Lane NE, Sinha SK, et al. Report: understanding the impact of COVID-19 on residents of Canada's long-term care homes—ongoing challenges and policy responses. LTC COVID website. https://ltccovid.org/wpcontent/uploads/2020/06/LTCcovid-country-reports_Canada_June-4-2020.pdf. Published June 4, 2020. Accessed July 28, 2022.
- van Heijl I, Schweitzer VA, Zhang L, van der Linden PD, van Werkhoven CH, Postma DF. Inappropriate use of antimicrobials for lower respiratory

tract infections in elderly patients: patient- and community-related implications and possible interventions. *Drugs Aging* 2018;35:389–398.

14. Mitchell R, Choi KB, Pelude L, Rudnick W, Thampi N, Taylor G. Patients in hospital with laboratory-confirmed COVID-19 in a network of Canadian

acute-care hospitals, March 1 to August 31, 2020: a descriptive analysis. *CMAJ Open* 2021;9:E149–E156.

 Leonardi I, Gao IH, Lin W-Y, *et al.* Mucosal fungi promote gut barrier function and social behavior via type 17 immunity. *Cell* 2022;185:1–16.