

Central-Line–Associated Bloodstream Infections in Québec Intensive Care Units: Results from the Provincial Healthcare-Associated Infections Surveillance Program (SPIN)

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BACKGROUND. Following implementation of bundled practices in 2009 in Quebec and Canadian intensive care units (ICUs), we describe CLABSI epidemiology during the last 8 years in the province of Québec (Canada) and compare rates with Canadian and American benchmarks.

METHODS. CLABSI incidence rates (IRs) and central venous catheter utilization ratios (CVCURs) by year and ICU type were calculated using 2007–2014 data from the Surveillance Provinciale des Infections Nosocomiales (SPIN) program. Using American and Canadian surveillance data, we compared SPIN IRs to rates in other jurisdictions using standardized incidence ratios (SIRs).

RESULTS. In total, 1,355 lab-confirmed CLABSIs over 911,205 central venous catheter days (CVC days) were recorded. The overall pooled incidence rate (IR) was 1.49 cases per 1,000 CVC days. IRs for adult teaching ICUs, nonteaching ICUs, neonatal ICUs (NICUs), and pediatric ICUs (PICUs) were 1.04, 0.91, 4.20, and 2.15 cases per 1,000 CVC days, respectively. Using fixed SPIN 2007–2009 benchmarks, CLABSI rates had decreased significantly in all ICUs except for PICUs by 2014. Rates declined by 55% in adult teaching ICUs, 52% in adult nonteaching ICUs, and 38% in NICUs. Using dynamic American and Canadian CLABSI rates as benchmarks, SPIN adult teaching ICU rates were significantly lower and adult nonteaching ICUs had lower or comparable rates, whereas NICU and PICU rates were higher.

CONCLUSION. Québec ICU CLABSI surveillance shows declining CLABSI rates in adult ICUs. The absence of a decrease in CLABSI rate in NICUs and PICUs highlights the need for continued surveillance and analysis of factors contributing to higher rates in these populations.

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Central-line–associated bloodstream infection (CLABSI) is associated with serious morbidity and mortality in intensive care units (ICUs) and is one of the costliest hospital-acquired infections (HAIs).^{1,2} In the province of Québec, the perceived public health importance of HAI due to a large *C. difficile* outbreak led to the development of provincial surveillance programs in 2003: Surveillance Provinciale des Infections Nosocomiales (SPIN) under the Institut National de Santé Publique du Québec (INSPQ).³ Currently, all ICUs in Québec with ≥ 10 beds are required to report CLABSIs year-round to SPIN, giving the program the advantage of having representative population surveillance.⁴ SPIN objectives include acquiring data to track epidemiology, incidence, and causative pathogens, as well as providing benchmark incidence rates. Importantly, the program's continuous surveillance throughout the year enables both intra- and interfacility benchmarking of CLABSI rates and central line use.

Canadian and American surveillance have revealed an overall decline in CLABSI since 2006.^{5,6} Québec CLABSI rates also reflected this downward trend from 2003 to 2009.⁷ These declines coincided with the implementation of several important programs and guideline updates such as the Centers for Disease Control and Prevention (CDC) revised intravascular catheter-related infection prevention guidelines⁸ and the Canadian Patient Safety Institute's program, Safer Healthcare Now! The Safer Healthcare Now! program was created in 2009 to support the implementation of evidence-based bundles of central-line insertion and maintenance and has been effective in decreasing CLABSI rates.⁹ On a regional level, a survey of ICUs in Québec revealed that most ICUs implemented bundled practices; however, practices such as performing regular audits were less optimal in most adult ICUs.¹⁰ A recent study of American pediatric ICUs (PICUs) showed similar bundle use and compliance practices.¹¹ Due to changing practices and overall

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decreasing CLABSI rates within the last decade, as seen in several national and regional surveillance programs including Québec, we aimed to determine the effect of changing practices and culture of CLABSI prevention efforts in Québec during the last 8 years, as well as to ascertain how SPIN rates compared with other populations to guide future prevention efforts. Our specific aims were (1) to describe CLABSI rates in Québec during the surveillance period, (2) to determine whether any significant rate trends existed, especially after newer guideline publications, and (3) to benchmark Quebec rates dynamically with annual SPIN, Canadian, and American surveillance CLABSI rates.

METHODS

SPIN Surveillance Network

SPIN is a year-round active and prospective CLABSI surveillance program, mandatory for all ICUs with ≥ 10 beds in the province of Québec since 2007, with individual-level data. ICUs with < 10 beds voluntarily submit data. Retrospective analysis of the program's reporting validity during the study period showed excellent results when compared with other regional surveillance networks, having a sensitivity and specificity of 88% and 92%, respectively.¹² By 2014, 70 ICUs from 51 different hospitals participated in the program (969 beds), comprising 33 nonteaching adult ICUs, 24 adult teaching ICUs, 8 neonatal ICUs (NICUs), and 5 PICUs. Of these, 57 ICUs (851 beds) participated in all 8 years of surveillance (Table 1) and were used in rate descriptions. All ICUs were included in benchmarking for standardized incidence ratio (SIR) analyses, regardless of full or partial participation because subgroup analyses demonstrated similar incidence. A previously published surveillance report of SPIN CLABSI rates included 2 years that overlap the present study (2007–2008 and 2008–2009);⁷ nevertheless, because mandatory SPIN CLABSI surveillance began in 2007, we included data from 2007 onward for optimal validity.

Definitions and CLABSI Identification

Central venous catheters (CVCs) were defined as intravenous catheters that end in a vessel in proximity to the heart, eg, the subclavian, internal jugular, or femoral vein. In accordance with NHSN and CNISP practices, peripherally inserted catheters, total implanted catheters, and umbilical catheters were also considered CVCs. SPIN has been following the National Healthcare Safety Network (NHSN) definition of CLABSI since April 1, 2010.¹³ SPIN CLABSI cases from 2007 to 2010 were retrospectively recomputed to reflect the new definition. NHSN 2006–2008 data reports already reflected this new definition, whereas CNISP CLABSI reports adopted the change as of April 1, 2010.^{6,14}

Data Collection and Surveillance

Patients with CVC in the ICU were followed 48 h after CVC removal or discharge from the ICU. Infection control

practitioners prospectively identified positive blood cultures in ICU patients, confirmed CVC placement and timing, and performed the chart review for criteria fulfillment. Data on CLABSIs that occurred between April 1, 2007, and March 31, 2015, were extracted in June 2015. The present study is a retrospective longitudinal cohort analysis that was approved by the INSPQ and did not require institutional board review because it was a secondary analysis of collected data.

Statistical Analysis

Pooled CLABSI incidence rates (IRs; cases per 1,000 CVC days), CVC utilization ratios (CVCURs, an indicator of CVC usage), and SIRs were calculated by ICU type (adult teaching or nonteaching, pediatric or neonatal) and by surveillance year. Incidence rate by each reporting period (1 calendar year comprises 13 4-week intervals) was examined for seasonal trends. The surveillance year begins April 1, which acts as day 1 of reporting period 1. Henceforth, calendar years written singly as “2007” refers to the start of surveillance year, which spans from April 1, 2007, to March 31, 2008. ICUs were defined as “teaching” if associated with medical training and research programs, and “nonteaching” otherwise. NICUs and PICUs are all associated with teaching hospitals. Poisson confidence intervals for rates and SIRs were used to compare CLABSI rates. Statistical calculations were performed using Stata version 14 (StataCorp; College Station, Texas).

SIRs use indirect standardization to compare rates between 2 populations.¹⁵ SIRs were obtained by dividing the observed number of CLABSI cases by the expected number of cases. Expected rates were taken from a reference population and were multiplied by the observed number of CVC days to generate the expected numbers of cases. A SIR of 1 denotes no difference between the observed and expected number of CLABSIs; a SIR < 1 denotes a rate less than expected, and a SIR > 1 denotes a rate higher than expected. The 95% SIR confidence intervals (CIs) were derived using upper and lower 95% CI limits of CLABSI IRs to calculate the corresponding number of expected cases.

To examine intraregional CLABSI rate trends over time, we used pooled SPIN rates from April 2007 to March 2010 as the benchmark because several important prevention guidelines and initiatives were published in 2009 (eg, the Safer Healthcare Now! program in Canada and the World Health Organization's launch of the Save Lives: Clean Your Hands initiative for hand hygiene).¹⁶ Using these pooled rates as benchmarks allowed us to measure the impact of these initiatives over time. To determine whether sustained rate trends existed, dynamic SPIN benchmarks were also used: SIRs for a particular year were calculated using pooled SPIN rates from the preceding 3 years for a given ICU type (eg, the 2010 adult teaching ICU SIR used pooled CLABSI rates of 2007 to 2009 in adult teaching ICUs as the benchmark to calculate expected rates).¹⁷

TABLE 1. Total ICU Units, CLABSI Cases, CVC Days, Pooled Means (95% CI), CVCUR by Year, and ICU Type

	Surveillance Year ^a							
	2007	2008	2009	2010	2011	2012	2013	2014
Adult Teaching ICUs								
No. of units (no. total participants)	24 (383)	24 (383)	24 (399)	23 (399)	23 (399)	23 (399)	23 (399)	23 (399)
No. of fully participating ICUs (no. ICU beds)	18 (383)							
Total cases, full participants (total cases, all participants)	90	107	85 (81)	78	59 (57)	63 (62)	64 (62)	44 (43)
Total CVC days (CVC days, full participants)	67,992 (67,992)	67,402 (67,402)	69,835 (68,483)	72,491 (70,928)	71,397 (69,805)	72,250 (70,431)	71,867 (70,038)	71,698 (70,132)
IR, full participants (95% CI)	1.32 (1.06–1.63)	1.59 (1.30–1.92)	1.18 (0.94–1.47)	1.10 (0.87–1.37)	0.82 (0.62–1.06)	0.88 (0.67–1.13)	0.89 (0.68–1.13)	0.61 (0.44–0.83)
CVCUR, full participants	0.65	0.64	0.62	0.64	0.61	0.61	0.59	0.59
Pooled IR, all years (95% CI)	1.04 (0.96–1.13)							
Adult Nonteaching ICUs								
No. of units (no. of beds)	22 (248)	22 (250)	24 (270)	26 (286)	27 (292)	29 (308)	31 (319)	32 (332)
No. of fully participating ICUs (no. of ICU beds)	21 (242)							
Total cases (cases, full participants)	30 (29)	20 (20)	40 (23)	29 (20)	32 (23)	38 (28)	24 (16)	21 (12)
Total CVC days (CVC days, full participants)	21,272 (20,758)	21,743 (21,553)	27,051 (23,072)	30,095 (23,208)	32,426 (24,862)	34,140 (24,912)	35,549 (23,808)	34,377 (24,912)
IR for all participants (95% CI)	1.40 (0.94–2.01)	0.93 (0.57–1.43)	1.00 (0.63–1.50)	0.86 (0.53–1.33)	0.93 (0.59–1.39)	1.12 (0.75–1.62)	0.67 (0.38–1.09)	0.48 (0.25–0.84)
CVCUR for complete participants only	0.33	0.35	0.37	0.36	0.39	0.39	0.38	0.38
Pooled IR all years	0.91 (0.78, 1.06)							
NICUs								
No. of units (no. of beds)	7 (172)	7 (172)	7 (172)	7 (172)	7 (172)	7 (172)	8 (184)	8 (184)
No. of fully participating ICUs (no. of ICU beds)	7 (172)							
Total cases (cases, full participants)	40	35	53	74	101	80	71	40
CVC days (total CVC days, full participants)	11,129	11,585	12,762	14,793	16,939	15,100	17,454 (17,452)	17,898 (17,895)
IR overall (95% CI)	3.59 (2.49–4.79)	3.02 (2.10–4.20)	4.15 (3.11–5.43)	5.00 (3.93–6.28)	5.96 (4.86–7.25)	5.30 (4.14–6.52)	4.07 (3.18–5.13)	2.23 (1.60–3.04)
CVCUR	0.15	0.15	0.17	0.19	0.22	0.19	0.23	0.24
Pooled IR, all years (95% CI)	4.20 (3.84–4.59)							
PICUs								
No. of units (no. of beds)	5 (54)	5 (54)	5 (54)	5 (54)	5 (54)	5 (54)	5 (54)	5 (54)
CLABSI cases	12	10	12	15	12	19	15	15
Total CVC days	5,375	5,629	6,194	6,531	6,643	6,730	6,855	7,283
IR (95% CI)	2.23 (1.15–3.89)	1.78 (0.85–3.27)	1.94 (1.00–3.38)	2.30 (1.29–3.79)	1.81 (0.93–3.20)	2.82 (1.70–4.41)	2.19 (1.22–3.61)	2.06 (1.15–3.40)
CVCUR	0.54	0.55	0.59	0.60	0.56	0.51	0.60	0.59
Pooled IR, all years (95% CI)	2.15 (1.76–2.59)							

NOTE. ICU, intensive care unit; CI, confidence interval; IR, incidence rate; CLABSI, central-line-associated bloodstream infection; CVC, central venous catheter; CVCUR, central venous catheter ratios (CVCUR); PICU, pediatric ICU.

^aThe surveillance year begins April 1, which acts as day 1 of reporting period 1. Calendar years are written singly, eg, 2007 refers to the period from April 1, 2007, to March 31, 2008.

To compare SPIN rates with American and Canadian ICU CLABSI rates, we obtained published CLABSI rates from available CNISP and NHSN reports during 2007–2014. CNISP surveillance data were extracted from published reports for 2006, 2009, 2010, and 2011,⁶ and NHSN data were extracted for years 2006–2008 (pooled rates)¹⁴ and for subsequent yearly reports from 2009 to 2013.^{5,18–21} CDC/NHSN and consumer groups release ongoing reports publishing SIRs using NHSN 2006–2008 CLABSI benchmarks; therefore, we included this benchmark to be consistent with ongoing publications. However, to better explore whether SPIN rates were similar to NHSN rates over this period in the context of practice changes affecting both healthcare populations, dynamic SIRs using NHSN rates from the preceding 3 years were also used as benchmarks for the examined year. CLABSI rates for NHSN medical and/or surgical ICUs described as “major teaching” were used to obtain expected rates for SPIN adult teaching ICU SIR derivations; ICUs classified as “all other” were considered nonteaching adult ICUs. NICUs were not compared because they used birth weights in their reporting; this information was not collected in SPIN during the entire study period. However, as of April 2013, birth-weight-specific CLABSI rates were being reported.

Due to gaps in published reports between CNISP and SPIN during this period, the most recently available CNISP rates were used as benchmarks for any corresponding SPIN year. CNISP 2006 rates served as benchmarks for SPIN surveillance years 2007–2009 inclusive; pooled CNISP 2009–2010 rates were used to benchmark SPIN years 2010 and 2011; and CNISP 2011 rates were used to benchmark SPIN surveillance years 2012–2014, inclusive. Because the vast majority of CNISP hospitals are tertiary hospitals with academic affiliations, adult nonteaching ICUs were excluded from CNISP SIR derivations.

RESULTS

CVCURs and Pooled IRs

Total participating ICUs, CLABSI cases, CVC days, CVCUR, and pooled IR by year and ICU type are shown in Table 1. Over the surveillance period, ICU participation increased from 56 to 67 facilities, 11 of which were nonteaching adult ICUs. A total of 1,428 laboratory-confirmed CLABSIs and 970,498 CVC days were recorded, and the overall pooled mean rate was 1.47 (95% CI, 1.40–1.55) cases per 1,000 CVC days. Restricting analysis to ICUs that participated for the entire surveillance period, the overall incidence remained at 1.49 (95% CI, 1.41–1.57). Incidence rates and CVCURs by ICU type and year are shown in Table 1 and illustrated in Figure 1. Figure 2 shows rates by reporting period: no significant evidence of seasonality was observed in rates for each ICU type. CVCURs for adult teaching and nonteaching, pediatric, and neonatal ICUs, which participated for the entire 8 years were 0.62, 0.37, 0.57, and 0.20, respectively (Table 1).

SIRs Against SPIN 2007–2009 and NHSN 2006–2008 Benchmarks

SIRs with fixed SPIN 2007–2009 and NHSN 2006–2008 benchmarks were calculated to study rate changes over time before and after important guideline and program launches in 2009–2010. Table 2 presents SIRs for each ICU type and by year: adult teaching ICUs showed a significant rate decline over the period, with 2014 SIRs of 0.45 (95% CI, 0.33–60) and 0.26 (95% CI, 0.19–0.36) using SPIN and NHSN benchmarks, respectively. Adult nonteaching ICUs also decreased, with a SPIN SIR of 0.48 (95% CI, 0.30–0.73) and an NHSN SIR of 0.39 (95% CI, 0.24–0.59) in 2014. PICUs did not show a significant rate change with either benchmark. Neonatal ICU rates varied; a significant rate increase was observed in SIRs for years 2011 and 2012, followed by a significant rate decrease in 2014, with an SIR of 0.62, (95% CI, 0.44–0.84).

SIRs Using Dynamic Benchmarks

Dynamic SIRs using SPIN, CNISP, and NHSN benchmarks by ICU type are shown in Table 3. For adult teaching ICUs using SPIN benchmarks, rates for most years were similar to the preceding year's rates, except for 2014, which showed a statistically significant decline with an SIR of 0.71 (95% CI, 0.52–0.96). With CNISP benchmarks, SPIN adult teaching ICUs had lower rates compared with the most recent CNISP rates published in 2007, 2009–2011, and 2014. Dynamic NHSN benchmarks yielded significantly lower SIRs for adult teaching ICUs for all years. In adult nonteaching ICUs, SIRs in 2013 and 2014 using SPIN benchmarks showed significantly lower rates compared with preceding years, and SIRs using NHSN benchmarks were significantly lower in 2010 and 2014.

NICU SIRs showed significantly higher rates with SPIN benchmarks in 2010 and 2011, having SIRs of 1.39 (95% CI, 1.09–1.74) and 1.44 (95% CI, 1.17–1.75), respectively. Likewise, using CNISP benchmarks, NICUs SIRs for 2012 (1.82; 95% CI, 1.42–2.24) and 2013 (1.40; 95% CI, 1.09–1.76) were also significantly higher. PICUs demonstrated no significant differences in dynamic SIRs using SPIN data, but they did yield significantly higher SIRs when using NHSN benchmarks in 2011 (SIR 1.92; 95% CI, 1.16–3.00) and CNISP data in 2012 (SIR 2.12; 95% CI, 1.28–3.31).

DISCUSSION

From 2007 to 2014, the overall rate of 1.49 cases per 1,000 CVC days for all ICU types was comparable to CLABSI rates in other developed countries after bundle intervention, such as Germany (1.64 cases per 1,000 CVC days in 2008–2010),²² and Victoria, Australia (1.26 cases per 1,000 CVC days in 2009–2013).²³ Neither a seasonality effect nor a “July effect” on rates due to influx of new residents in hospitals was identified.²⁴

Importantly, Québec adult teaching and nonteaching ICUs showed lower and decreasing CLABSI rates over the

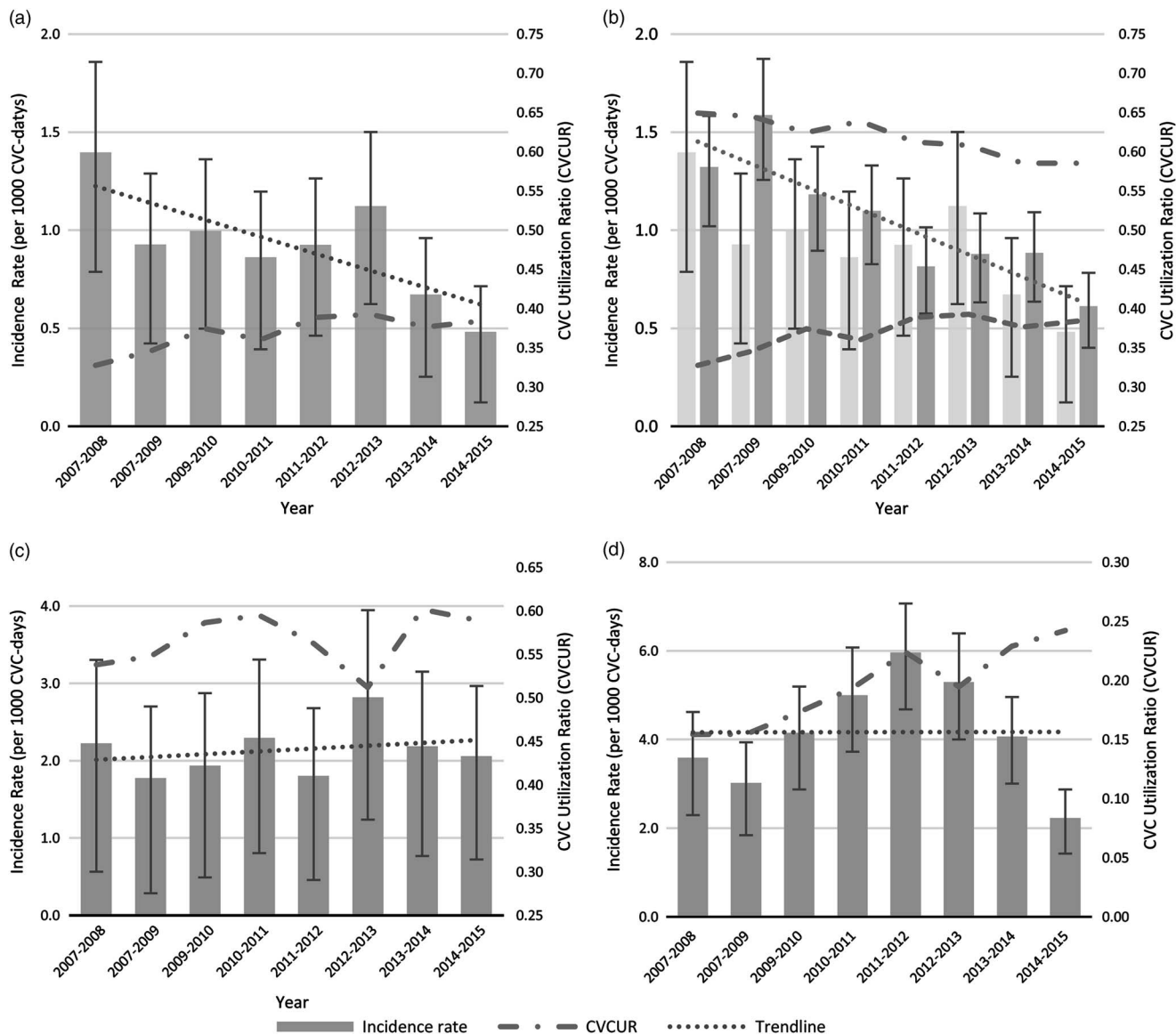


FIGURE 1. Incidence Rate (IR) expressed as CLABSI cases per 1,000 CVC days, with 95% Poisson confidence interval bars, and central venous catheter utilization Ratios (CVCURs) by year for (a) Adult nonteaching ICUs, (b) Adult teaching ICUs, (c) PICUs, and (d) NICUs.

surveillance period. Later, adult teaching and nonteaching ICU rates demonstrated statistically significant declines when using SPIN 2007–2009 benchmarks, decreasing by 55% (95% CI, 40%–67%) for adult teaching ICUs, and by 52% (95% CI, 27%–70%) for adult nonteaching ICUs in 2014. Using dynamic benchmarks to examine significant year-to-year changes, SPIN adult ICUs also had lower rates compared with NHSN and CNISP benchmarks for most years. SIR was not statistically significant when using dynamic SPIN benchmarks, perhaps to a lack of power because SPIN is a smaller network. Post-hoc power calculation showed that power was <80%, ranging from 5% to 51% for most ICU types and most years. Dynamic adult nonteaching ICU SIRs with NHSN

referents were more comparable to SPIN rates overall, with significantly lower SIRs in 2008 and 2010.

Adult ICU rate reduction may be attributed to several factors. In 2009, a national campaign from the Canadian Patient Safety Institute implemented guidelines on the use of evidence-based bundles in hospitals. Furthermore, greater HAI awareness from updated CDC intravascular catheter guidelines in 2011 and WHO hand hygiene recommendations in 2009 may have contributed to decreasing rates^{8,16}; this effect was seen in a multicenter time series study in Germany.²² SPIN rates for adult teaching ICUs were comparable to CNISP benchmarks, suggesting Québec CLABSI interventions paralleled that of national efforts. Recent results from 1 Québec

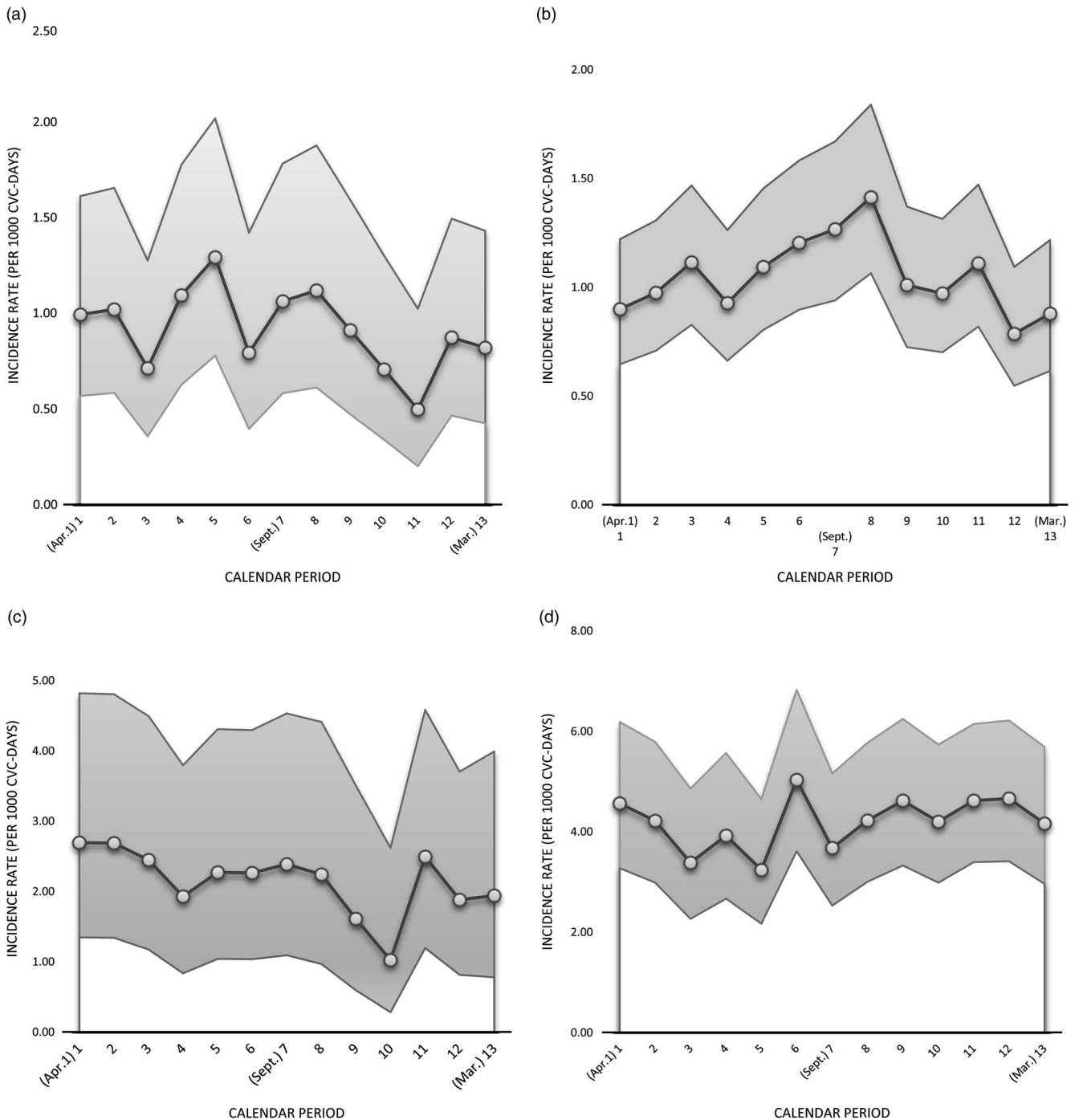


FIGURE 2. Incidence by each calendar period for each ICU type by year for (a) Adult nonteaching ICUs, (b) Adult teaching ICUs, (c) PICUs, and (d) NICUs. Period are calendrical, with April 1 (start of yearly reporting period) corresponding to period 4 in graphs. Shading represents 95% CI.

academic center with 7 ICUs demonstrated continual decreases during the last 8 years of stepwise prevention.²⁵ Moreover, surveillance in itself has been shown to decrease rates of device-associated infections, which may in part explain decreasing rates prior to guideline changes.²⁶

Unlike adult ICUs, Québec NICU and PICU rates did not show the same downward trend. In PICUs, no significant rate changes were observed using either fixed or dynamic SPIN-derived SIRs, although smaller sample size should be noted. PICU rates at outset of SPIN surveillance were comparable

TABLE 2. Standardized Incidence Rate (SIR) by Intensive Care Unit (ICU) Type Using Baseline NHSN 2006–2008 and SPIN 2007–2009 Rates as Benchmarks for All SPIN Surveillance Years from 2007 to 2014

Adult Teaching ICUs			NICUs		
Year ^a	Benchmark Used		Year ^a	Benchmark Used	
	SPIN 2007–2009	NHSN 2006–2008		SPIN 2007–2009	NHSN 2006–2008
2007	0.97 (0.78–1.19)	0.57 (0.46–0.70)	2007	1.00 (0.69–1.33)	N/A
2008	1.16 (0.95–1.40)	0.68 (0.56–0.83)	2008	0.84 (0.58–1.16)	N/A
2009	0.89 (0.71–1.10)	0.52 (0.42–0.65)	2009	1.15 (0.86–1.50)	N/A
2010	0.79 (0.62–0.98)	0.46 (0.37–0.58)	2010	1.39 (1.09–1.74)	N/A
2011	0.60 (0.46–0.78)	0.36 (0.27–0.58)	2011	1.65 (1.35–2.01)	N/A
2012	0.64 (0.49–0.81)	0.38 (0.29–0.48)	2012	1.47 (1.15–1.81)	N/A
2013	0.65 (0.50–0.83)	0.38 (0.30–0.49)	2013	1.13 (0.88–1.42)	N/A
2014	0.45 (0.33–0.60)	0.26 (0.19–0.36)	2014	0.62 (0.44–0.84)	N/A

Adult Nonteaching ICUs			PICUs		
Year ^a	Benchmark Used		Year ^a	Benchmark Used	
	SPIN 2007–2009	NHSN 2006–2008		SPIN 2007–2009	NHSN 2006–2008
2007	1.10 (0.74–1.57)	0.90 (0.61–1.28)	2007	1.12 (0.58–1.96)	0.95 (0.49–1.66)
2008	0.72 (0.44–1.11)	0.59 (0.36–0.90)	2008	0.90 (0.43–1.65)	0.76 (0.36–1.39)
2009	1.16 (0.83–1.57)	0.94 (0.67–1.28)	2009	0.98 (0.51–1.71)	0.82 (0.43–1.44)
2010	0.75 (0.50–1.08)	0.61 (0.41–0.88)	2010	1.16 (0.65–1.91)	0.98 (0.55–1.61)
2011	0.77 (0.53–1.09)	0.63 (0.43–0.89)	2011	0.91 (0.47–1.59)	0.77 (0.40–1.34)
2012	0.87 (0.58–1.15)	0.71 (0.47–0.94)	2012	1.43 (0.86–2.23)	1.20 (0.72–1.88)
2013	0.53 (0.34–0.78)	0.43 (0.28–0.64)	2013	1.11 (0.62–1.82)	0.93 (0.52–1.54)
2014	0.48 (0.30–0.73)	0.39 (0.24–0.59)	2014	1.04 (0.58–1.72)	0.88 (0.49–1.45)

NOTE. NHSN, National Healthcare Safety Network; SPIN, Surveillance Provinciale des Infections Nosocomiales; NICU, neonatal ICU; PICU, pediatric ICU; N/A, not available.

^aThe surveillance year begins April 1, which acts as day 1 of reporting period 1. Calendar years are written singly, eg, 2007 refers to the period from April 1, 2007, to March 31, 2008.

with NHSN rates, and lower than CNISP rates. SPIN PICU rates remained constant over time with no decrease in rates during the period, while NHSN and CNISP PICU rates decreased more than SPIN PICU rates.

When compared with SPIN 2007–2009 benchmarks, SPIN NICUs had statistically significant rate increases from 2007 to 2011, peaking at 5.96 (95% CI, 4.86–7.25) cases per 1,000 CVC days in 2011, corresponding to an SIR of 1.65 (1.35, 2.01). Subsequently, rates and SIR declined, resulting in a statistically significant SIR decreases of 48% (95% CI, 16%–56%). Prior to 2012, NICU SPIN SIRs for most years were significantly lower using CNISP benchmarks; however, CNISP-derived SIRs became significantly higher in 2012–2013. Similarly, dynamic SPIN-derived SIRs were also significantly higher in 2010–2012.

Higher NICU and PICU rates may have several explanations. First, evidence for insertion and maintenance bundles in these populations are less robust than in adults. Several studies show that children have longer central catheter dwell times, emphasizing greater importance on maintenance bundle adherence.^{27,28} Consequently, there is greater heterogeneity in bundle element types for children than for bundles designed for adults.^{29,30} Successful strategies described include incorporating

elements based on facility-specific challenges, involving parents in prevention efforts, and holding regular meetings with stakeholders to discuss outcomes and directions.^{31–33} The rising NICU and PICU rates during in Québec around 2012 may also be due to outbreaks leading to persisting local CLABSI endemics. For example, between 2010 and 2013, 46% of all CLABSI NICU cases and 52% of all PICU CLABSIs originated in 1 facility, compared to 30% and 37%, respectively, in that facility for all other years. HAI rates also greatly differ across NICUs in Canada and may be explained by regional strains, difference in case mix, and clinical practices.³⁴ Québec has 4 large academic centers, a distinguishing feature offering unique challenges. Following a combination of molecular and epidemiological characterization of what led to the rate increase and subsequent decline in PICUs and NICUs, sharing of knowledge and strategies regularly among the 4 centers will be important for future prevention efforts.

Strengths and Limitations

A major strength of the study is the complete population-level surveillance of SPIN ICUs, which includes a mix of different hospitals (both teaching and nonteaching) and ICU types in

TABLE 3. Standardized Incidence Rate (SIR) Using Dynamic SPIN, NHSN, or CNISP Benchmarks of CLABSI Rates of Either the Preceding 3 Years or the Most Recent Rate, by Intensive Care Unit (ICU) Type

Adult Teaching ICUs				NICUs			
Year ^a	SIR			Year ^a	SIR		
Benchmark	SPIN ^b	CNISP	NHSN	Benchmark	SPIN ^b	CNISP	NHSN
2007	N/A	0.74 (0.60, 0.91)	N/A	2007	N/A	0.69 (0.48–0.91)	N/A
2008	N/A	0.89 (0.73–1.08)	N/A	2008	N/A	0.58 (0.40–0.80)	N/A
2009	N/A	0.68 (0.55–0.85)	0.52 (0.42–0.65)	2009	N/A	0.79 (0.59–1.04)	N/A
2010	0.79 (0.62–0.98)	0.60 (0.48–0.75)	0.46 (0.37–0.58)	2010	1.39 (1.09–1.74)	0.72 (0.54–0.95)	N/A
2011	0.64 (0.49–0.83)	0.62 (0.47–0.80)	0.64 (0.49–0.82)	2011	1.44 (1.17–1.75)	0.62 (0.47–0.80)	N/A
2012	0.84 (0.62–1.07)	0.93 (0.71–1.19)	0.58 (0.45–0.74)	2012	1.03 (0.81–1.27)	1.82 (1.42, 2.24)	N/A
2013	0.96 (0.74–1.22)	0.95 (0.73–1.21)	0.67 (0.52–0.86)	2013	1.04 (0.81–1.31)	1.40 (1.09–1.76)	N/A
2014	0.71 (0.52–0.96)	0.65 (0.47–0.88)	0.59 (0.36–0.66)	2014	0.44 (0.31–0.60)	0.77 (0.55–1.05)	N/A

Nonteaching Adult ICUs				PICUs			
Year ^a	SIR			Year ^a	SIR		
Benchmark	SPIN ^b	CNISP	NHSN	Benchmark	SPIN ^b	CNISP	NHSN
2007	N/A	N/A	N/A	2007	N/A	0.74 (0.60–0.91)	N/A
2008	N/A	N/A	N/A	2008	N/A	0.89 (0.73–1.08)	N/A
2009	N/A	N/A	0.94 (0.67–1.28)	2009	N/A	0.68 (0.55–0.85)	0.82 (0.43–1.44)
2010	0.75 (0.50–1.08)	N/A	0.61 (0.41–0.88)	2010	1.16 (0.65–1.91)	0.60 (0.48–0.75)	0.98 (0.55–1.61)
2011	0.87 (0.60–1.23)	N/A	1.15 (0.76–1.52)	2011	0.89 (0.46–1.56)	0.93 (0.48–1.62)	1.92 (1.16–3.00)
2012	0.99 (0.65–1.30)	N/A	0.99 (0.65–1.30)	2012	1.40 (0.84–2.18)	2.12 (1.28–3.31)	1.47 (0.89–2.30)
2013	0.66 (0.42–0.98)	N/A	0.68 (0.43–1.00)	2013	0.95 (0.53–1.56)	1.65 (0.92–2.71)	1.33 (0.75–2.20)
2014	0.66 (0.41–1.02)	N/A	0.64 (0.40–0.98)	2014	0.91 (0.51–1.50)	1.55 (0.87–2.55)	1.43 (0.80–2.36)

NOTE. NHSN, National Healthcare Safety Network; SPIN, Surveillance Provinciale des Infections Nosocomiales; CNISP, Canadian Nosocomial Infection Surveillance Program; NICU, neonatal ICU; pediatric ICU; N/A, not available.

^aThe surveillance year begins April 1, which acts as day 1 of reporting period 1. Calendar years are written singly, eg, 2007 refers to the period from April 1, 2007, to March 31, 2008.

^bSPIN dynamic benchmarks for each year were calculated using the incidence rates for preceding 3 years of surveillance; dynamic SIRs for SPIN surveillance years 2007–2009 were thus not calculated because it is the benchmark.

Québec, which lead to accurate CLABSI benchmarking. This surveillance program has been validated in the past and has been shown to be accurate,⁷ resulting in greater accuracy in intraregional rate comparisons. That said, as always when comparing rates and generalizability between different networks, differences in surveillance methods and infection control practice should be kept in mind. Nevertheless, here, both incidence rates and SIRs illustrate that CLABSI rates are declining in Québec adult ICUs during 2007 to 2014.

Our study demonstrates that CLABSI rates in adult teaching ICUs in Québec were significantly lower than CNISP and NHSN rates and that rates continued to decline throughout the surveillance period. SPIN adult nonteaching ICUs rates also decreased, at a pace more comparable to NHSN nonteaching adult ICUs. On the contrary, IRs in SPIN NICUs and PICUs increased from 2011 to 2013, unlike other American and Canadian facilities, which saw a continual decline in rates. Future efforts should be directed at delineating and understanding causes of persistently higher rates in the NICU and PICU and at identifying strategies to further decrease these rates.

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REFERENCES

- Sydnor ERM, Perl TM. Hospital epidemiology and infection control in acute-care settings. *Clin Microbiol Rev* 2011;24:141–173.
- Zimlichman E, Henderson D, Tamir O, et al. Health care-associated infections: a meta-analysis of costs and financial impact on the US health care system. *JAMA Intern Med* 2013;173:2039–2046.
- Fontela PS, Platt RW, Rocher I, et al. Surveillance Provinciale des Infections Nosocomiales (SPIN) Program: implementation of a

- mandatory surveillance program for central line-associated bloodstream infections. *Am J Infect Control* 2011;39:329–335.
4. Fontela PS, Quach C, Buckeridge D, Pai M, Platt RW. Surveillance length and validity of benchmarks for central line-associated bloodstream infection incidence rates in intensive care units. *PloS One* 2012;7:e36582.
 5. Dudeck MA, Edwards JR, Allen-Bridson K, et al. National Healthcare Safety Network report, data summary for 2013, device-associated Module. *Am J Infect Control* 2015;43:206–221.
 6. Public Health Agency of C. *Central Venous Catheter-Associated Bloodstream Infections in Intensive Care Units in Canadian Acute-Care Hospitals: Surveillance Report January 1, 2006 to December 31, 2006 and January 1, 2009 to December 31, 2011*. Public Health Agency of Canada; 2014. 55.
 7. Fontela PS, Platt RW, Rocher I, et al. Epidemiology of central line-associated bloodstream infections in Quebec intensive care units: a 6-year review. *Am J Infect Control* 2012;40:221–226.
 8. O'Grady NP, Alexander M, Burns LA, et al. Guidelines for the prevention of intravascular catheter-related infections. *Clin Infect Dis* 2011;52:e162–e193.
 9. The Joint Commission. *Preventing Central Line-Associated Bloodstream Infections: A Global Challenge, a Global Perspective*. Oak Brook, IL: Joint Commission Resources; 2012/05// 2012.
 10. Gonzales M, Rocher I, Fortin E, et al. A survey of preventive measures used and their impact on central line-associated bloodstream infections (CLABSI) in intensive care units (SPIN-BACC). *BMC Infect Dis* 2013;13:562.
 11. Edwards JD, Herzig CT, Liu H, et al. Central line-associated blood stream infections in pediatric intensive care units: longitudinal trends and compliance with bundle strategies. *Am J Infect Control* 2015;43:489–493.
 12. Fontela PS, Rocher I, Platt RW, et al. Evaluation of the reporting validity of central line-associated bloodstream infection data to a provincial surveillance program. *Infect Control Hosp Epidemiol* 2013;34:217–219.
 13. Center of Disease Control and Prevention/National Health Surveillance Network. Bloodstream Infection Event (Central Line-Associated Bloodstream Infection and Non-central line-associated Bloodstream Infection) 2015. http://www.cdc.gov/nhsn/pdfs/pscmanual/4psc_clabscurrent.pdf
 14. Edwards JR, Peterson KD, Mu Y, et al. National Healthcare Safety Network (NHSN) report: data summary for 2006 through 2008, issued December 2009. *Am J Infect Control* 2009;37:783–805.
 15. Gustafson TL. Practical risk-adjusted quality control charts for infection control. *Am J Infect Control* 2000;28:406–414.
 16. Pittet D, Allegranzi B, Boyce J. The World Health Organization Guidelines on hand hygiene in health care and their consensus recommendations. *Infect Control Hosp Epidemiol* 2009;30:611–622.
 17. Saman DM, Kavanagh KT, Abusalem SK. Redefining the standardized infection ratio to aid in consumer value purchasing. *J Patient Saf* 2013;9:55–58.
 18. Dudeck MA, Horan TC, Peterson KD, et al. National Healthcare Safety Network report, data summary for 2011, device-associated module. *Am J Infect Control* 2013;41:286–300.
 19. Dudeck MA, Horan TC, Peterson KD, et al. National Healthcare Safety Network (NHSN) Report, data summary for 2010, device-associated module. *Am J Infect Control* 2011;39(10):798–816.
 20. Dudeck MA, Horan TC, Peterson KD, et al. National Healthcare Safety Network (NHSN) report, data summary for 2009, device-associated module. *Am J Infect Control* 2011;39(5):349–367.
 21. Dudeck MA, Weiner LM, Allen-Bridson K, et al. National Healthcare Safety Network (NHSN) report, data summary for 2012, device-associated module. *Am J Infect Control* 2013;41(12):1148–1166.
 22. Hansen S, Schwab F, Schneider S, Sohr D, Gastmeier P, Geffers C. Time-series analysis to observe the impact of a centrally organized educational intervention on the prevention of central-line-associated bloodstream infections in 32 German intensive care units. *J Hosp Infect* 2014;87:220–226.
 23. Worth LJ, Spelman T, Bull AL, Brett JA, Richards MJ. Central line-associated bloodstream infections in Australian intensive care units: time trends in infection rates, etiology, and antimicrobial resistance using a comprehensive Victorian surveillance program, 2009–2013. *Am J Infect Control* 2015;43:848–852.
 24. Young JQ, Ranji SR, Wachter RM, Lee CM, Niehaus B, Auerbach AD. “July effect”: impact of the academic year-end changeover on patient outcomes: a systematic review. *Ann Intern Med* 2011;155:309–315.
 25. Paquet F, Frenette C, Patterson C, Decary A. Impact of an eight-year program to reduce central line-associated blood stream infection (CLABSI) in a University Teaching Hospital. *Open Forum Infectious Diseases* 2015;2:S117.
 26. Gastmeier P, Schwab F, Sohr D, Behnke M, Geffers C. Reproducibility of the surveillance effect to decrease nosocomial infection rates. *Infect Control Hosp Epidemiol* 2009;30:993–999.
 27. Miller MR, Griswold M, Harris JM, 2nd, et al. Decreasing PICU catheter-associated bloodstream infections: NACHRI's quality transformation efforts. *Pediatrics* 2010;125:206–213.
 28. Milstone AM, Reich NG, Advani S, et al. Catheter dwell time and CLABSIs in neonates with PICCs: a multicenter cohort study. *Pediatrics* 2013;132:e1609–e1615.
 29. Huskins WC. Quality improvement interventions to prevent healthcare-associated infections in neonates and children. *Curr Opin Pediatr* 2012;24:103–112.
 30. Smulders CA, van Gestel JP, Bos AP. Are central line bundles and ventilator bundles effective in critically ill neonates and children? *Intensive Care Med* 2013;39:1352–1358.
 31. Fisher D, Cochran KM, Provost LP, et al. Reducing central line-associated bloodstream infections in North Carolina NICUs. *Pediatrics* 2013;132:e1664–e1671.
 32. McMullan C, Propper G, Schuhmacher C, et al. A multi-disciplinary approach to reduce central line-associated bloodstream infections. *Joint Comm J Qual Patient Safety* 2013;39:61–69.
 33. Neill S, Haithcock S, Smith PB, et al. Sustained reduction in bloodstream infections in infants at a large tertiary care neonatal intensive care unit. *Adv Neonat Care* 2016;16:52–59.
 34. Aziz K, McMillan DD, Andrews W, et al. Variations in rates of nosocomial infection among Canadian neonatal intensive care units may be practice-related. *BMC Pediatr* 2005;5:22.