

## Risk factors for surgical site infection following cesarean delivery: a retrospective cohort study

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### Abstract

**Background:** The rate of cesarean delivery is increasing in North America. Surgical site infection following this operation can make it difficult to recover, care for a baby and return home. We aimed to determine the incidence of surgical site infection to 30 days following cesarean delivery, associated risk factors and whether risk factors differed for pre-discharge versus post-discharge infection.

**Methods:** We identified a retrospective cohort in Nova Scotia by linking the provincial perinatal database to hospital admissions and physician billings databases to follow women for 30 days after they had given birth by cesarean delivery between Jan. 1, 1997 and Dec. 31, 2012. Logistic regression with generalized estimating equations was used to determine risk factors for infection.

**Results:** A total of 25 123 women had 33 991 cesarean deliveries over the study period. Of the 25 123, 923 had surgical site infections, giving an incidence rate of 2.7% (95% CI 2.54%–2.89%); the incidence decreased over time. Risk factors for infection (adjusted odds ratios  $\geq 1.5$ ) were prepregnancy weight 87.0 kg or more, gaining 30.0 kg or more during pregnancy, chorioamnionitis, maternal blood transfusion, anticoagulation therapy, alcohol or drug abuse, second stage of labour before surgery, delivery in 1997–2000 and delivery in a hospital performing 130–1249 cesarean deliveries annually. Women who gave birth earlier in the study period, those who gave birth in a hospital with 130–949 cesarean deliveries per year and those with more than 1 fetus were at a significantly higher risk for surgical site infection before discharge; women who smoked were at significantly higher risk for surgical site infection after discharge.

**Interpretation:** Most risk factors are known before delivery, and some are potentially modifiable. Although the incidence of surgical site infection decreased over time, targeted clinical and infection prevention and control interventions could further reduce the burden of illness associated with this health-care-related infection.

The proportion of infants born by cesarean delivery has been increasing in North America. In Canada, the cesarean delivery rate was 20% in 1988<sup>1</sup> and rose to 26% in 2012.<sup>2</sup> Surgical site infection following this procedure can adversely influence the postpartum period for the women and her newborn, making it difficult to recover, care for a baby and return home. Although Canada has no national surveillance for surgical site infection following cesarean delivery, the US National Healthcare Safety Network follows this infection and reported a mean rate of 0.16% in 2014.<sup>3</sup> An Australian study of 81 health care facilities from 2002 to 2013 showed a rate of 2.05%, which decreased over the study period.<sup>4</sup> Some hospitals conduct surveillance on these infections up until discharge, and others also conduct postdischarge surveillance. The rate of surgical site infection until hospital discharge following cesarean delivery varies, with reported values ranging from 0.16% to 3.2%.<sup>3,5,6</sup> Although rates are higher if postdischarge surveillance is conducted, this activity is resource intensive and therefore is not routinely performed.

Risk factors for surgical site infection following cesarean delivery have been identified in the literature. Most studies

have shown a significant association between obesity and risk for surgical site infection,<sup>7,8</sup> and a dose-response relation between body mass index and risk for infection has been noted.<sup>5</sup> Findings for other risk factors, such as hypertension and diabetes, have been less consistent.<sup>5,9,10</sup> Both appropriately timed antibiotic prophylaxis<sup>11</sup> and chlorhexidine antiseptic skin preparation<sup>12</sup> have been shown to decrease the infection risk. Few studies report infection rates to 30 days post partum. There is little or no evidence regarding whether risk factors

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differ based on time of infection presentation, and the influence of hospital size, anticoagulation therapy and smoking on infection risk.

In this retrospective population-based cohort study, we followed women from the perinatal period to 30 days post partum to determine 1) the incidence of surgical site infection including before and after hospital discharge, 2) risk factors associated with infection and 3) whether risk factors differed with time of presentation of infection.

## Methods

### Setting

This study took place in Nova Scotia (population roughly 950 000), which in 2011 had 8860 births and a cesarean delivery rate of 26.6%.<sup>13</sup> About half of deliveries are performed at the regional and tertiary care centre, IWK Health Centre, based in the capital city (Halifax).

### Participants

Women with a Nova Scotia health card who gave birth via cesarean delivery in the province between Jan. 1, 1997 and Dec. 31, 2012 were eligible. Before 1997, International Classification of Diseases, 9th revision (ICD-9) codes were not captured with decimals in the physician billings database and, therefore, could not be used to distinguish surgical site infections from other types of infection. When this study was begun, administrative health databases were complete up until 2012. In Nova Scotia, everyone is eligible for a health card if they can provide proof of citizenship or immigration. Women were excluded if they delivered a baby weighing less than 500 g or at less than 20 weeks' gestational age.

### Design and data sources

We identified the retrospective cohort using the Nova Scotia Atlee Perinatal Database (NSAPD), administered by the Reproductive Care Program of Nova Scotia. It contains information on all pregnancies and deliveries in the province, which is collected by health care professionals using standardized forms and patient charts. The database is regularly analyzed for accuracy and reliability. From the database, we obtained relevant information on pregnancy, labour, delivery and the postpartum period, including surgical site infection identified before discharge.

### Outcome

The outcome was surgical site infection following cesarean delivery within 30 days of surgery. The NSAPD was linked with the Canadian Institute for Health Information's Discharge Abstract Database (to capture the delivery admission or readmission) and with the Nova Scotia Medical Services Insurance Physician Billings Database (to capture insured service encounters rendered by a physician) to identify women with a diagnostic code indicative of surgical site infection. The Discharge Abstract Database used the ICD-9 until 2000 and the Canadian enhancement of the 10th revision (ICD-10-CA) from 2001 onward; the Physicians Billings Database used

ICD-9 codes. Diagnostic codes indicative of surgical site infection were those for wound infection or endometritis according to any of the coding systems: the NSAPD, ICD-9 (614.3, 674.30, 674.32, 674.34, 998.51, 998.59) or ICD-10-CA (N71.0, N71.9, N73.0, O86.0xx). We categorized infections according to whether the infection was diagnosed during the hospital stay for delivery of the infant or after discharge. The timing of pre-discharge surgical site infections could not be defined as only admission and discharge dates are recorded in the Discharge Abstract Database. For infections identified after discharge, we estimated the number of days from surgery to infection identification based on the date of either hospital readmission or physician encounter.

### Potential risk factors

We examined several potential risk factors for surgical site infection based on a literature review and the variables available in the NSAPD (Table 1). This database has information on smoking at various times (i.e., at first prenatal visit, 20 weeks' gestation and delivery), from which we determined whether the woman smoked. Height was not recorded in the NSAPD until 2003; therefore, we approximated standard body mass index categories<sup>14</sup> using weight cut-points determined by means of receiver operator characteristic curve analysis: less than 53.0 kg = underweight, 53.0–66.9 kg = normal, 67.0–76.9 kg = overweight, 77.0–86.9 kg = obese I, 87.0–97.9 kg = obese II and 98.0 kg or more = obese III. In supplementary analyses of deliveries from 2003 on, we examined body mass index (< 18.5 kg/m<sup>2</sup>, 18.5–24.9 kg/m<sup>2</sup>, 25.0–29.9 kg/m<sup>2</sup>, 30.0–39.9 kg/m<sup>2</sup> and ≥ 40.0 kg/m<sup>2</sup>) for comparison. Hypertension, diabetes and depression variables were all combinations of both diagnostic codes and codes indicating that the woman was taking medication for these medical conditions from the NSAPD.

### Statistical analysis

The unit of analysis was cesarean delivery rather than women. We used descriptive statistics to estimate the incidence of pre-discharge, postdischarge and total surgical site infection to 30 days with exact confidence intervals (CIs). We used  $\chi^2$  tests to determine which risk factors were significantly associated with surgical site infection. We entered risk factors associated with infection at  $p < 0.10$  into a multiple logistic regression model, from which we removed risk factors that were not independently associated with surgical site infection using backward stepwise selection. Only risk factors that were associated with infection at  $p < 0.05$  were retained. From the logistic models, we estimated adjusted odds ratios (ORs) with 95% CIs. We examined risk factors for pre- and postdischarge surgical site infection using multinomial logistic regression. Generalized estimating equations were used for all logistic models to account for potential correlation between women who had more than 1 cesarean delivery over the study period. If 5% or more of data was missing for a risk factor, we created a "missing" category. Because a missing-value category would have low numbers for risk factors with less than 5% of data missing, we excluded deliveries with missing

**Table 1: Potential risk factors for surgical site infection that were examined, to determine inclusion in multivariable model in univariate analyses**

Category	Factors not associated with infection ( $p \geq 0.10$ )	Factors associated with infection ( $p < 0.10$ )
Institution-related	–	No. of cesarean deliveries in delivery hospital (< 130, 130–949, 950–1249, $\geq 1250$ )
Area-level	Maternal residence (urban, rural)	Region of maternal residence (A, B, C, D) Neighbourhood-level income quintile (1, 2, $\geq 3$ )
Maternal demographic characteristics	–	Maternal age (< 25 yr, 25–34 yr, $\geq 35$ yr) Prepregnancy weight (< 53.0 kg, 53.0–66.9 kg, 67.0–76.9 kg, 77.0–86.9 kg, 87.0–97.9 kg, $\geq 98.0$ kg, missing) Marital status (married or common-law, other, missing) Smoking during pregnancy (no, yes) Alcohol or drug abuse during pregnancy (no, yes)
Maternal medical conditions	Maternal influenza immunization in pregnancy (no, yes)	Nonobstetric preexisting health condition affecting pregnancy (no, yes) Hypertension (no, preexisting, gestational or unspecified, preeclampsia) Diabetes (no, preexisting, gestational) Anemia during pregnancy (no, yes) Depression (no, yes) Anticoagulation therapy during pregnancy (no, yes)
Pregnancy history	Surgical site infection after previous cesarean delivery (no, yes)	Parity (primiparous, multiparous) Mode of delivery in last pregnancy (none, vaginal, cesarean) Previous cesarean delivery (0, 1, $\geq 2$ )
Pregnancy characteristics	–	Weight gain during pregnancy (< 10.0 kg, 10.0–19.9 kg, 20.0–29.9 kg, $\geq 30.0$ kg, missing) Chorioamnionitis (no, yes) Diagnostic and/or therapeutic procedure(s) performed in woman (no, yes) Steroid use $\geq 48$ h before delivery for fetal lung maturity (no, yes)
Labour		Cervical dilation at last examination before cesarean delivery (0 cm, 1–3 cm, 4–10 cm) Time between rupture of membranes to delivery ( $\leq 1$ h, 2–11 h, $\geq 12$ h) Stage of labour before cesarean delivery (none, first, second) Spontaneous rupture of membranes (yes, no)
Delivery	Season of delivery (December–February, March–May, June–August, October–November) Type of cesarean delivery (low segment transverse, other) Instrumentation use (no, forceps/vacuum) General anesthesia during labour and/or delivery (no, yes) Other procedure(s) performed during cesarean delivery (no, yes)	Length of antepartum stay (< 24 h, 24–49 h, $\geq 50$ h) Year of delivery (1997–2000, 2001–2004, 2005–2008, 2009–2012) Delivery on weekend (no, yes) Primary indication for cesarean delivery (breech presentation, dystocia, fetal distress, other, previous cesarean delivery) Regional anesthesia during labour and/or delivery (no, yes) Antibiotic therapy during labour and delivery (yes, no) Maternal blood transfusion (no, yes)
Fetal and neonatal	Presentation at delivery (vertex, other, missing) No. of fetuses (1, $\geq 2$ ) Diagnostic and/or therapeutic procedure(s) performed in fetus (no, yes)	Infant birth weight (< 2500 g, 2500–3999 g, $\geq 4000$ g) Apgar score at 5 min (< 7, $\geq 7$ ) Gestational age (< 37 wk, 37 wk to 39 wk + 6 d, $\geq 40$ wk) Breastfeeding at discharge (yes, no)

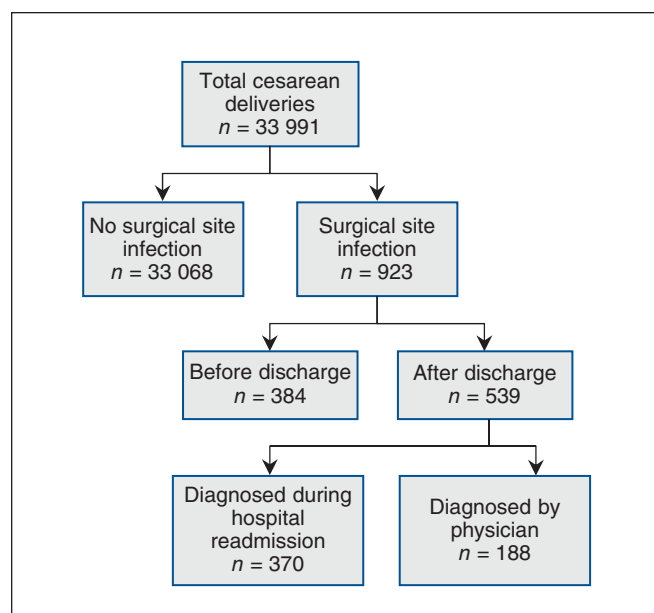
values for these risk factors from regression analyses. We did not use multiple imputation for our primary analyses owing to the large size of our data set. Estimates from analyses using 5 data sets with missing values imputed with chained equations were very similar to those shown herein. We conducted all analyses using Stata/SE 13 (StataCorp LP).

### Ethics approval

This study was approved by the Reproductive Care Program of Nova Scotia's Joint Data Access Committee, Health Data Nova Scotia and the IWK Health Centre's Research Ethics Board.

### Results

During the period 1997–2012, 25 123 women (with 33 991 cesarean deliveries) were identified from the NSAPD and could be linked to hospital and physician billings data (> 96% linkage). A total of 923 surgical site infections followed the 33 991 deliveries within 30 days, giving an incidence rate of 2.7% (95% CI 2.54%–2.89%) (Figure 1). Fifteen women had more than 1 surgical site infection during the study period. The incidence of surgical site infections decreased from 5.2% in 1997 to 2.0% in 2012 (Figure 2). The incidence of infection was higher after discharge (1.6% [95% CI 1.45%–1.72%]) than before discharge (1.1% [95% CI 1.02%–1.24%]) ( $p < 0.001$ ). Of the 539 postdischarge surgical site infections, 370 (68.6%) were identified during hospital readmission. Of the 539 women who presented with surgical site infection after discharge, 469 (87.0%) did so within 2 weeks of their surgery. The mean length of postpartum stay decreased from 3.86 (standard deviation 1.53) days in 1997–2000 to 3.23 (standard deviation 1.04) days in 2009–2012 ( $p < 0.001$ ). The



**Figure 1:** Participant flow diagram. “Diagnosed during hospital readmission” and “diagnosed by physician” do not add to 539 as 19 women were coded as having a surgical site infection during a hospital readmission and a physician office visit.

rate of administration of antibiotic therapy for prophylaxis or treatment during labour and delivery increased from 46.9% in 1997–2000 to 73.6% in 2009–2012; we were unable to distinguish between these indications from the data available. We were able to determine that, from 2003 to 2012, the indication was group B *Streptococcus* infection in 972 (8.2%) of the 11 854 women who received antibiotics.

Table 2 shows the risk factors independently associated with the odds of developing surgical site infection within 30 days following cesarean delivery at  $p < 0.05$  (see Appendix 1, available at [www.cmajopen.ca/content/5/3/E546/suppl/DC1](http://www.cmajopen.ca/content/5/3/E546/suppl/DC1), for univariate analysis). Women who gave birth in 1997–2000 had a higher risk of infection (adjusted OR 2.31, 95% CI 1.88–2.84) than those who gave birth in 2009–2012. Compared to women who were primiparous, those who were multiparous had a lower risk of surgical site infection (adjusted OR 0.81, 95% CI 0.69–0.95). A dose–response relation was observed between prepregnancy weight and infection, with women who weighed 98.0 kg or more having more than 3 times the odds of infection as women who weighed 53.0–66.9 kg. In supplementary analyses conducted for 2003–2012, body mass index was similarly positively associated with infection (data not shown).

The results of the multinomial logistic regression analysis comparing the risk factors for pre- and postdischarge infection are given in Table 3 (see Appendix 2, available at [www.cmajopen.ca/content/5/3/E546/suppl/DC1](http://www.cmajopen.ca/content/5/3/E546/suppl/DC1), for univariate associations). Women who smoked during pregnancy had significantly higher odds of infection presenting after discharge (adjusted OR 1.39, 95% CI 1.13–1.70) than before discharge (adjusted OR 0.94, 95% CI 0.73–1.21). Deliveries earlier in the study period, of more than 1 fetus and in a hospital performing 130–949 cesarean deliveries annually had significantly higher odds of surgical site infection presenting before discharge than after discharge.

### Interpretation

In this 16-year study of nearly 34 000 births by cesarean delivery, the incidence of surgical site infection to 30 days post partum was 2.7%. More infections presented after discharge than before discharge (58.4% v. 41.6%), as noted by other investigators.<sup>15</sup> Infection rates decreased over time. Our analysis identified some risk factors that are modifiable or known before delivery, such as prepregnancy weight and anticoagulation therapy, and some risk factors were associated differently with pre- versus postdischarge surgical site infection.

The incidence of surgical site infection of 2.7% observed in the present study is comparable to the lower end of the infection rates reported in the literature, which range from 0.16%<sup>3</sup> to 9.6%.<sup>5</sup> Rates reported from infection prevention and control programs are fairly consistent and are at the lower end of the range.<sup>3,4</sup> In a study from Nova Scotia, the authors reported an incidence of surgical site infection following cesarean delivery of 1.5% for 1988–2002 but followed women only to discharge.<sup>16</sup> We do not know why Nova Scotia rates are at the lower end of the reported range. Possible contributing factors include universal access to

health care, introduction of standardized antibiotic administration before cesarean delivery at some sites and the salutary effect of a provincial reproductive care program that sets quality standards and provides education to all obstetric health care providers. Variation in reported infection rates can often be attributed to different populations, time periods and definitions of surgical site infection, secular trends and improvements in health care.<sup>17</sup> For example, Wloch and colleagues<sup>5</sup> found a considerably higher infection rate than we did, possibly owing to higher prevalence of comorbidities such as diabetes and active follow-up. The decrease in our rate over a 16-year period, from 5.2% to 2.0%, is similar to the absolute decrease of 2.05% reported in an Australian study<sup>4</sup> and to trends in health-care-associated infections seen in the United States.<sup>3</sup> Possible reasons for these temporal trends include quality-improvement interventions such as surgical checklists, use of 2% chlorhexidine skin antiseptics, standing orders for systemic antimicrobial prophylaxis with adjusted dosages for obesity and increased administration of appropriate prophylactic antibiotics following published guidelines by the Centers for Disease Control and Prevention in 1999,<sup>18</sup> the American College of Obstetricians and Gynecologists in 2003<sup>19</sup> and the Canadian Society of Obstetricians and Gynaecologists in 2010.<sup>11</sup> Reported rates could also erroneously decrease if only pre-discharge infections are surveilled, as the length of postpartum stay has decreased over time.

We identified several risk factors for surgical site infection after cesarean delivery, including higher prepregnancy weight and weight gain during pregnancy. Like Wloch and colleagues,<sup>5</sup> we observed a dose-response relation between prepregnancy weight and infection, with weighing 98.0 kg or

more (approximating obese class III) being associated with more than 3 times the odds of surgical site infection compared to weighing 53.0–66.9 kg (approximating normal weight). Prepregnancy obesity has been a significant and strong independent risk factor for surgical site infection in most studies<sup>7,8</sup> and, given its high prevalence (estimated at 35% in our cohort), is therefore important. Other risk factors identified in the present study that are theoretically modifiable included smoking, alcohol or drug abuse, gaining 30.0 kg or more during pregnancy and no antibiotic therapy. Antibiotic prophylaxis is an established protective factor for surgical site infection,<sup>20</sup> whereas smoking has not been observed consistently to be a risk factor.<sup>21</sup> Our findings regarding alcohol or drug abuse and weight gain as risk factors can be used to refine or to initiate and evaluate targeted clinical and infection prevention and control interventions to decrease the infection rate and lower this burden of illness following cesarean delivery on patients and the health care system.

Some factors associated with surgical site infection before discharge in our study may be linked to an increased potential for contamination during surgery (e.g., not receiving antibiotic therapy) and longer surgery duration (e.g., obesity and multiple fetuses). In contrast, risk factors associated with post-discharge infection may be indicative of increased potential for wound contamination (e.g., smoking and low socioeconomic status) and delayed wound healing or wound separation (e.g., obesity). These findings suggest that the effect of some individual risk factors is influenced by the length of time since surgery. A previous study of various operations (excluding cesarean delivery) also showed a significant difference between risk factors for surgical site infection before and after discharge, including age and duration of surgery.<sup>22</sup> More

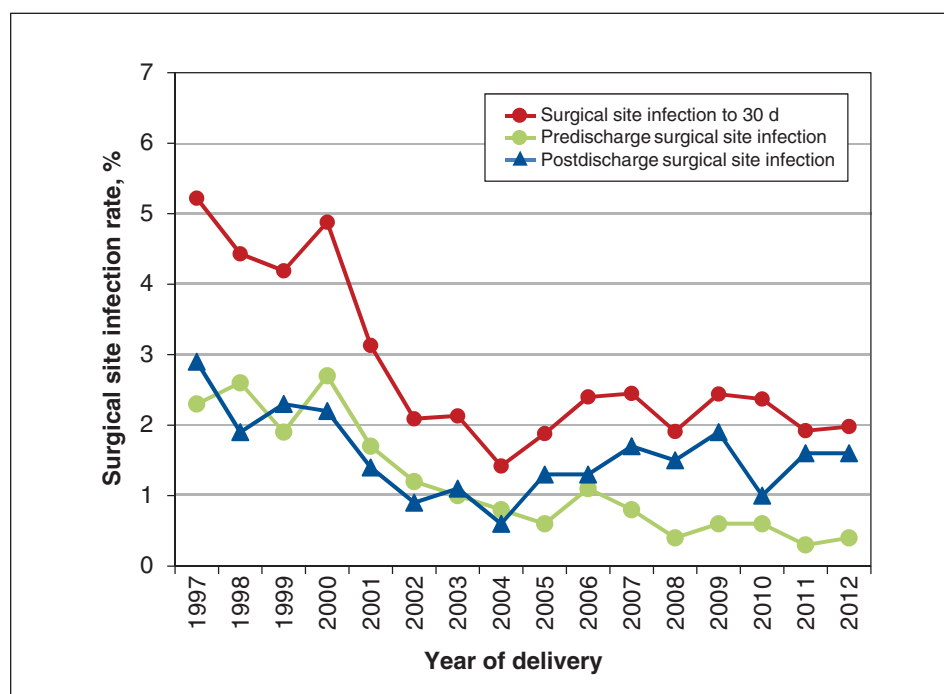


Figure 2: Rate of surgical site infection by year.

**Table 2 (part 1 of 2): Risk factors for surgical site infection following cesarean delivery to 30 days post partum**

Risk factor	% of total*	No. of deliveries		Adjusted OR (95% CI)
		No infection	Infection	
<b>Institution-related</b>				
No. of cesarean deliveries per hospital per year				
< 130	25.1	8165	158	Reference
130–949	21.7	6910	282	2.20 (1.79–2.71)
950–1249	27.7	8861	317	1.85 (1.50–2.29)
≥ 1250	25.5	8305	154	1.39 (1.09–1.79)
<b>Area-level</b>				
Quintile of neighbourhood-level income				
1 (lowest)	20.1	6410	262	1.40 (1.19–1.65)
2	18.1	5824	172	1.09 (0.91–1.30)
3–5	61.8	20 007	477	Reference
<b>Maternal demographic characteristics</b>				
Age, yr				
< 25	18.6	5929	225	1.13 (0.94–1.36)
25–34	61.5	19 880	511	Reference
≥ 35	19.9	6432	175	1.21 (1.01–1.45)
Prepregnancy weight, kg				
< 53.0	8.0	2603	61	1.05 (0.78–1.41)
53.0–66.9	29.3	9519	194	Reference
67.0–76.9	16.8	5429	135	1.28 (1.02–1.61)
77.0–86.9	12.1	3888	108	1.43 (1.12–1.82)
87.0–97.9	7.8	2488	85	1.86 (1.42–2.43)
≥ 98.0	9.5	2961	186	3.63 (2.90–4.54)
Missing	16.6	5353	142	1.52 (1.12–2.06)
Marital status				
Married/common-law	72.7	23 511	578	Reference
Single/divorced/separated/ widowed	22.3	7116	285	1.34 (1.13–1.58)
Missing	5.0	1614	48	1.14 (0.83–1.58)
Alcohol or drug abuse				
No	98.6	31 809	885	Reference
Yes	1.4	432	26	1.71 (1.12–2.61)
<b>Maternal medical conditions</b>				
Nonobstetric preexisting health conditions affecting pregnancy				
No	80.9	26 160	676	Reference
Yes	19.1	6081	235	1.28 (1.09–1.51)
Diabetes				
No	93.5	30 181	809	Reference
Preexisting	0.6	196	11	1.62 (0.86–3.04)
Gestational	5.9	1864	91	1.49 (1.17–1.88)

**Table 2 (part 2 of 2): Risk factors for surgical site infection following cesarean delivery to 30 days post partum**

Risk factor	% of total*	No. of deliveries		Adjusted OR (95% CI)
		No infection	Infection	
<b>Depression during pregnancy</b>				
No	95.6	30 826	855	Reference
Yes	4.4	1415	56	1.40 (1.05–1.87)
<b>Anticoagulation therapy during pregnancy</b>				
No	99.1	31 965	888	Reference
Yes	0.9	276	23	2.58 (1.64–4.05)
<b>Pregnancy history</b>				
<b>Parity</b>				
Primiparous	49.4	15 837	539	Reference
Multiparous	50.6	16 404	372	0.81 (0.69–0.95)
<b>Pregnancy characteristics</b>				
<b>Weight gain during pregnancy, kg</b>				
< 10.0	16.7	5372	155	0.81 (0.66–0.99)
10.0–19.9	41.8	13 522	350	Reference
20.0–29.9	13.6	4352	142	1.26 (1.03–1.54)
≥ 30.0	1.9	601	37	2.14 (1.49–3.06)
Missing	26.0	8394	227	1.10 (0.86–1.41)
<b>Chorioamnionitis</b>				
No	97.8	31 569	856	Reference
Yes	2.2	672	55	2.78 (2.07–3.75)
<b>Labour</b>				
<b>Stage of labour before cesarean delivery</b>				
None	48.6	15 793	332	Reference
First	31.9	10 235	350	1.20 (1.01–1.43)
Second	19.4	6213	229	1.54 (1.27–1.88)
<b>Delivery</b>				
<b>Year of delivery</b>				
1997–2000	22.0	6947	342	2.31 (1.88–2.84)
2001–2004	26.5	8607	191	1.14 (0.91–1.41)
2005–2008	27.0	8742	193	1.02 (0.83–1.25)
2009–2012	24.5	7945	185	Reference
<b>Antibiotic therapy during labour and delivery</b>				
Yes	62.9	20 355	506	Reference
No	37.1	11 886	405	1.30 (1.12–1.51)
<b>Maternal blood transfusion</b>				
No	99.0	31 932	885	Reference
Yes	1.0	309	26	2.70 (1.77–4.12)
<b>Fetal and neonatal</b>				
<b>Gestational age</b>				
< 37 wk	9.8	3125	120	1.36 (1.10–1.69)
37 wk to 39 wk + 6 d	57.1	18 475	443	Reference
≥ 40 wk	33.1	10 641	348	1.09 (0.93–1.28)

Note: CI = confidence interval, OR = odds ratio.  
 \*A total of 33 152 cesarean deliveries were included in the analysis; 839 deliveries were excluded because of missing values for neighbourhood-level income ( $n = 724$ ), stage of labour ( $n = 1$ ) and/or gestational age ( $n = 118$ ).

**Table 3 (part 1 of 2): Risk factors for surgical site infection before and after discharge following cesarean delivery**

Risk factor	No infection, no. of deliveries*	Infection before discharge		Infection after discharge	
		No. of deliveries*	Adjusted OR (95% CI)	No. of deliveries*	Adjusted OR (95% CI)
<b>Institution-related</b>					
No. of cesarean deliveries per hospital per year					
< 130	8070	51	Reference	104	Reference
130–949	6893	125	3.16 (2.26–4.44)	157	1.76 (1.34–2.30)†
950–1249	8787	142	2.46 (1.73–3.51)	170	1.58 (1.20–2.08)
≥ 1250	8223	59	1.76 (1.16–2.67)	95	1.26 (0.92–1.73)
<b>Area-level</b>					
Quintile of neighbourhood-level income					
1 (lowest)	6349	97	1.22 (0.95–1.57)	162	1.54 (1.25–1.90)
2	5784	70	1.03 (0.78–1.36)	102	1.15 (0.91–1.46)
3–5	19 840	210	Reference	262	Reference
<b>Maternal demographic characteristics</b>					
Age, yr					
< 25	5876	82	1.08 (0.81–1.43)	142	1.18 (0.93–1.49)
25–34	19 719	211	Reference	295	Reference
≥ 35	6378	84	1.41 (1.08–1.83)	89	1.08 (0.84–1.38)
Prepregnancy weight, kg					
< 53.0	2593	27	1.04 (0.67–1.60)	34	1.08 (0.72–1.60)
53.0–66.9	9463	92	Reference	102	Reference
67.0–76.9	5410	51	1.02 (0.72–1.45)	82	1.46 (1.09–1.97)
77.0–86.9	3868	46	1.29 (0.90–1.85)	62	1.53 (1.11–2.12)
87.0–97.9	2480	37	1.70 (1.15–2.51)	48	1.97 (1.38–2.81)
≥ 98.0	2954	71	3.07 (2.21–4.25)	114	4.06 (3.02–5.45)
Missing	5205	53	1.06 (0.68–1.67)	84	1.95 (1.28–2.97)
Marital status					
Married/common-law	23 323	250	Reference	323	Reference
Single/divorced/separated/widowed	7044	111	1.34 (1.03–1.75)	172	1.28 (1.03–1.58)
Missing	1606	16	1.26 (0.73–2.18)	32	1.07 (0.72–1.60)
Smoking during pregnancy					
No	25 151	282	Reference	361	Reference
Yes	6822	95	0.94 (0.73–1.21)	165	1.39 (1.13–1.70)†
<b>Maternal medical conditions</b>					
Nonobstetric preexisting health conditions affecting pregnancy					
No	25 941	278	Reference	393	Reference
Yes	6032	99	1.37 (1.08–1.75)	133	1.22 (0.99–1.50)
Diabetes					
No	29 923	335	Reference	467	Reference
Preexisting	194	6	2.29 (0.96–5.48)	5	1.28 (0.52–3.16)
Gestational	1856	36	1.50 (1.05–2.15)	54	1.47 (1.09–1.98)
Depression during pregnancy					
No	30 567	357	Reference	490	Reference
Yes	1406	20	1.44 (0.91–2.29)	36	1.45 (1.01–2.08)
Anticoagulation therapy during pregnancy					
No	31 700	365	Reference	515	Reference
Yes	273	12	3.94 (2.02–7.70)	11	1.93 (1.03–3.63)



**Table 3 (part 2 of 2): Risk factors for surgical site infection before and after discharge following cesarean delivery**

Risk factor	No infection, no. of deliveries*	Infection before discharge		Infection after discharge	
		No. of deliveries*	Adjusted OR (95% CI)	No. of deliveries*	Adjusted OR (95% CI)
<b>Pregnancy history</b>					
Parity					
Primiparous	15 723	208	Reference	329	Reference
Multiparous	16 250	169	0.87 (0.69–1.08)	197	0.75 (0.61–0.93)
<b>Pregnancy characteristics</b>					
Weight gain during pregnancy, kg					
< 10.0	5353	62	0.83 (0.61–1.12)	93	0.80 (0.62–1.05)
10.0–19.9	13 468	140	Reference	208	Reference
20.0–29.9	4341	61	1.40 (1.02–1.90)	80	1.14 (0.87–1.48)
≥ 30.0	599	18	2.98 (1.79–4.96)	19	1.68 (1.02–2.75)
Missing	8212	96	1.43 (1.01–2.02)	126	0.92 (0.65–1.29)
Chorioamnionitis					
No	31 307	352	Reference	496	Reference
Yes	666	25	3.30 (2.11–5.16)	30	2.47 (1.67–3.63)
Diagnostic and/or therapeutic procedure(s) performed in woman					
No	31 049	365	Reference	500	Reference
Yes	924	12	0.88 (0.49–1.57)	26	1.52 (1.01–2.29)
<b>Labour</b>					
Stage of labour before cesarean delivery					
None	15 645	155	Reference	172	Reference
First	10 159	132	1.00 (0.77–1.29)	217	1.40 (1.11–1.77)
Second	6169	90	1.35 (1.02–1.79)	137	1.66 (1.28–2.15)
<b>Delivery</b>					
Year of delivery					
1997–2000	6891	172	5.69 (4.01–8.07)	163	1.30 (1.01–1.68)†
2001–2004	8532	102	2.91 (2.00–4.25)	88	0.63 (0.48–0.84)†
2005–2008	8670	63	1.52 (1.03–2.25)	130	0.88 (0.69–1.12)†
2009–2012	7880	40	Reference	145	Reference
Antibiotic therapy during labour and delivery					
Yes	20 185	187	Reference	316	Reference
No	11 788	190	1.48 (1.18–1.84)	210	1.17 (0.97–1.41)
Maternal blood transfusion					
No	31 675	367	Reference	511	Reference
Yes	298	10	2.80 (1.43–5.48)	15	2.66 (1.53–4.62)
<b>Fetal or neonatal</b>					
No. of fetuses					
1	30 939	350	Reference	515	Reference
≥ 2	1034	27	1.78 (1.13–2.80)	11	0.64 (0.34–1.20)†
Gestational age					
< 37 wk	3057	62	1.42 (1.02–1.96)	54	1.21 (0.88–1.66)
37 wk to 39 wk + 6 d	18331	191	Reference	248	Reference
≥ 40 wk	10585	124	0.98 (0.77–1.24)	224	1.19 (0.98–1.46)

Note: CI = confidence interval, OR = odds ratio.

\*A total of 32 876 cesarean deliveries were included in the analysis; 1115 deliveries were excluded because of missing values for neighbourhood-level income ( $n = 724$ ), smoking ( $n = 283$ ), stage of labour ( $n = 1$ ) and/or gestational age ( $n = 118$ ).

† $p < 0.05$  for association between risk factor and timing of infection (before v. after discharge).

studies are necessary to see whether these findings are observed in other populations and to separate the influence of location (hospital v. home) from that of time since surgery.

### Limitations

As demand for quality improvement in health care increases, administrative claims databases have been used in recent years for surveillance of health-care-associated infections. In 2 systematic reviews of the diagnostic accuracy of administrative claims databases for this purpose, highly variable accuracy was found.<sup>23,24</sup> One of the reviews<sup>23</sup> included surgical site infection following cesarean delivery and identified 3 validation studies.<sup>15,25,26</sup> Leth and colleagues<sup>26</sup> found sensitivities of infection diagnosed in hospital and after discharge of 77.1% and 68.9%, respectively, and specificities of 99.5% and 98.2%, respectively. A Canadian study showed that the sensitivity for detection of surgical site infection after cesarean delivery was 16.7% when using hospital data but 77.3% when including emergency department and physician claims databases; those authors concluded that the low sensitivity and positive predictive value make claims databases inadequate for use as quality indicators.<sup>25</sup> A third, small study showed that infection after cesarean delivery identified through such data sources was confirmed in 40% of 204 cases, with some criteria for surgical site infection met in an additional 27% of cases.<sup>15</sup> Since we used administrative databases to identify infections, there is likely some risk of misclassification. However, the comparability of our infection rates to those reported to the National Healthcare Safety Network<sup>3</sup> and to our institution's infection prevention control program, serving 50% of the population of Nova Scotia (data not shown), is reassuring. Although hospital-based surveillance conducted by infection prevention and control programs remains the gold standard for detection of health-care-associated infection, it is resource intensive and time consuming. Such programs often do not have the resources to detect postdischarge infections and therefore will miss most of these events. Use of administrative databases, as in our study, can detect trends over time, facilitate multisite surveillance when surveillance systems are not integrated across administrative health regions and allow linking of other data sets, such as those with risk factors or longer-term health outcomes.

Data on some risk factors (e.g., maternal height) were unavailable for the entire study period, and some risk factors had a high proportion of missing values. Although our study covered a 16-year period, the data are historical, as we followed patients only until 2012. Finally, the NSAPD does not contain certain surgical information (e.g., length of surgery), so we did not include such details as potential risk factors for surgical site infection.

### Conclusion

Use of administrative databases can identify postdischarge infection associated with health care that is not detected by hospital-based surveillance. The incidence of surgical site infection following cesarean delivery in Nova Scotia decreased over time. Obesity and weight gain during preg-

nancy were key risk factors for infection. Our findings are generalizable to populations with health care systems and demographic and clinical factors similar to those in Nova Scotia. Knowing which risk factors are associated with pre-versus postdischarge infection can assist clinicians in identifying women with specific risk factor profiles who may be at risk for surgical site infection. This offers an opportunity to develop systematic approaches to eliminate or reduce risk factors through patient, public and health care provider education, and system approaches such as timely administration of appropriate antibiotics.

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