



The Effect of Hospital Isolation Precautions on Patient Outcomes and Cost of Care: A Multi-Site, Retrospective, Propensity Score-Matched Cohort Study

Kim Tran, BSc¹, Chaim Bell, MD PhD^{2,3}, Nathan Stall, MD⁴, George Tomlinson, PhD^{5,6}, Allison McGeer, MD^{7,8}, Andrew Morris, MD SM(Epi)^{9,10}, Michael Gardam, MSc MD^{10,11}, and Howard B. Abrams, MD^{1,9,12,13}

¹OpenLab, University Health Network, Toronto, ON, Canada; ²Institute of Health Policy, Management and Evaluation, University of Toronto, Toronto, ON, Canada; ³Department of Medicine, Division of General Internal Medicine, Mount Sinai Hospital, Toronto, ON, Canada; ⁴Core Internal Medicine, University of Toronto, Toronto, ON, Canada; ⁵Dalla Lana School of Public Health, University of Toronto, Toronto, ON, Canada; ⁶Department of Medicine, University Health Network, Toronto, ON, Canada; ⁷Departments of Laboratory Medicine and Pathobiology, Medicine and Dalla Lana School of Public Health, University of Toronto, Toronto, ON, Canada; ⁸Department of Microbiology, Division of Infection Control, Mount Sinai Hospital, Toronto, ON, Canada; ⁹Department of Medicine, University of Toronto, Toronto, ON, Canada; ¹⁰Department of Medicine, University of Toronto, Toronto, ON, Canada; ¹¹Infectious Diseases and Medicine, University of Toronto, Toronto, ON, Canada; ¹²Department of Medicine, Division of General Internal Medicine, University Health Network and Mount Sinai Hospital, Toronto, ON, Canada; ¹³, Toronto, ON, Canada; ¹⁴, Toronto, ON, Canada; ¹⁵, Toronto, ON, Canada; ¹⁵, Toronto, ON, Canada; ¹⁵, Toronto, ON, Canada; ¹⁶, Toronto, ON, Canada; ¹⁷, Toronto, ON, Canada; ¹⁸, Toronto, ON, Canada; ¹⁸, Tor

BACKGROUND: Isolation precautions have negative effects on patient safety, psychological well-being, and healthcare worker contact. However, it is not known whether isolation precautions affect certain hospital-related outcomes.

OBJECTIVE: To examine the effect of isolation precautions on hospital-related outcomes and cost of care.

DESIGN: Retrospective, propensity-score matched cohort study of inpatients admitted to general internal medicine (GIM) services at three academic hospitals in Toronto, Ontario, Canada between January 2010 and December 2012.

PARTICIPANTS: Adult (≥18 years of age) patients on isolation precautions for respiratory illnesses and methicillin-resistant *Staphylococcus aureus* (MRSA) were matched to controls based on propensity scores derived from nine covariates: age, sex, Resource Intensity Weight, number of hospital readmissions within 90 days, total length of stay for hospital admissions within 90 days, site of admission, month of isolation, year of isolation, and Case Mix Group.

MAIN MEASURES: Thirty-day readmission rates and emergency department visits, hospital length of stay, expected length of stay, adverse events, in-hospital mortality, patient complaints, and cost of care in Canadian doll ars (CAD).

KEY RESULTS: A total of 17,649 non-isolated patients were admitted to the participating hospitals during the study period. We identified 1506 patients isolated for respiratory illnesses and 745 patients isolated for MRSA. Compared to non-isolated individuals, those on isolation precautions for respiratory illnesses stayed 17 % longer (95 % CI: 9 %, 25 %), stayed 9 % longer than expected (95 % CI: 3 %, 15 %), and had 23 % higher cost of care (95 % CI: 14 %, 32 %). Patients isolated for MRSA had similar outcomes, but

they also had a 4.4 % higher (95 % CI: 1.4 %, 7.3 %) rate of readmission to hospital within 30 days.

CONCLUSIONS: Isolation precautions are associated with adverse effects which may result in poorer hospital outcomes. Balancing the benefits for the many with the harms to the few will be a future challenge.

KEY WORDS: Isolation; Hospital Outcomes.

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INTRODUCTION

Isolation precautions are used for patients hospitalized with known or suspected colonization or infection with pathogens transmitted in hospitals. These precautions may require a private room or placement behind drawn curtains in a multibed room, requiring staff and visitors to wear personal protective equipment, and restricting the movement of patients. The use of these precautions has been shown to control outbreaks (e.g. severe acute respiratory syndrome [SARS]) and, in some circumstances, the transmission of other pathogens. However, such precautions may also be associated with unintended adverse consequences.

Isolation precautions negatively affect several dimensions of patient care. Isolated patients have half as much contact with clinicians as do control patients, and are half as likely to be examined by attending physicians on rounds. 5–7 They are also twice as likely to experience adverse events during hospitalization, and eight times as likely to experience supportive care failure such as falls and pressure ulcers. 8,9 Studies have reported negative psychological effects of isolation including feelings of loneliness, stigmatization, and increased depression and anxiety. Finally, isolated patients experience lower rates of satisfaction with care. 8,19

Received August 28, 2015 Revised June 23, 2016 Accepted August 26, 2016 Published online October 17, 2016 Some criticisms of these conclusions have suggested that studies to date have not adequately adjusted for the severity of patients' underlying illness, and no studies have examined the trade-off between the known benefits of isolation precautions and their potential risks.

We hypothesize that even with more complete adjustment for patient characteristics and severity of underlying illness, there may be other, unrecognized effects on operational and patient outcomes compared to well-matched controls. We sought to examine whether isolation precautions increased 30-day readmissions and emergency department (ED) visits, hospital length of stay (LOS), expected length of stay (ELOS), adverse events, in-hospital mortality, patient complaints, and cost of care compared to propensity-matched controls.

METHODS

We conducted a multi-site, retrospective, propensity scorematched cohort study of patients consecutively admitted to general internal medicine (GIM) services using hospital administrative databases in Toronto, Canada. The study was approved by the institutional research ethics boards of the participating hospitals.

Setting

Patients admitted through the emergency department to GIM at three academic hospitals in Toronto, Canada, between January 2009 and December 2012 were included in the study. All three hospitals are in downtown Toronto, and are tertiary academic institutions affiliated with the University of Toronto. During the 2011 fiscal year, Toronto General (TG), Toronto Western (TW), and Mount Sinai (MS) hospitals admitted 3163 patients, 3250 patients, and 2881 patients, respectively, to their GIM services.

Infection Prevention and Control Practices

GIM patients are screened at admission for methicillinresistant *Staphylococcus aureus* (MRSA) colonization using nasal, rectal/groin, and wound swabs. At sites TG and TW, respiratory (droplet and contact) precautions are used for patients admitted with fever and respiratory symptoms; at MS, patients with respiratory or cardiac symptoms are tested for respiratory viruses on admission, and droplet and contact precautions used only for patients with influenza.

An infection control practitioner determines the requirement for isolation precautions. The decision to discontinue isolation is at the discretion of the infection control practitioner, based on improvement in symptoms, confirmatory tests or alternative diagnosis, and/or discussion with attending physician. Information on a patient's isolation status is updated in the patient record. Cases of infectious diseases are confirmed by infection control practitioners and are cross-referenced with the infection control database.

Design and Participants

Matched cohorts were selected from among all adults (>18 years of age) hospitalized on GIM between January 2009 and December 2012 and who had at least a 2-day LOS. The LOS requirement ensured that patients had adequate exposure to isolation precautions to potentially affect outcomes of care. Two groups of isolated patients were identified: (1) those placed on droplet precautions for respiratory illnesses and (2) those placed on contact precautions for MRSA. We report on these two groups separately, as patients with MRSA are placed on isolation for a longer period of time than patients with respiratory illnesses; as such, the effect of isolation may be more pronounced in the MRSA cohort. Patients in the isolated cohorts include those who had or were suspected of having an infectious disease prior to hospital admission as well as those with hospital-acquired infections; infections acquired during hospitalization represent only a small proportion of cases. The MRSA cohort included patients who were colonized or infected. Patients with no isolation precautions were eligible as controls.

Data Sources

Variables including patient demographics, clinical characteristics, and outcomes of interest were obtained from two linked hospital administrative databases: the Discharge Abstract Database (DAD) and Enterprise Data Warehouse (EDW). The DAD is an administrative database containing demographic, administrative, and clinical data for hospital discharges that are provided by participating hospitals to the Canadian Institute of Health Information (CIHI).²⁰ The EDW is the data repository for clinical and corporate information at TG and TW hospitals. The costs associated with isolated patients and matched controls, in Canadian dollars (CAD), were obtained through the Ministry of Health and Long-Term Care's Ontario Case Costing Initiative (OCCI). OCCI provides participating hospitals with a standardized methodology for the collection of case costing data, allowing for comparisons between hospitals. 21-24 Only direct costs were considered. Physician costs are not paid from hospital budgets and not included in OCCI case costing.

Study Outcomes

We collected data on 30-day readmission and ED visits, hospital LOS, ELOS, adverse events, in-hospital mortality, patient complaints, and cost of care. Thirty-day readmission and ED visits were defined as a hospital readmission or ED visit to the same hospital within 30 days of discharge. Hospital LOS was defined as the number of days spent in the hospital from admission until the patient was discharged, transferred to another facility, or died. ELOS is a variable created by CIHI that predicts expected length of hospitalization, adjusting for the reason for hospitalization, comorbidities, age, complications, and inpatient Resource Intensity Weights (RIW); the ELOS values are derived from patient LOS data of all Canadian acute care hospitals within the current 2 years.²⁵ We

captured the following adverse events through hospital incident reports: falls, infection control incidents, medication-related incidents, equipment-related incidents, treatment-related incidents, laboratory-related incidents, and operative or invasive incidents. In-hospital mortality was defined as any death occurring during a patient's hospital admission. Patient complaints include any formal complaint made to the hospital patient relations office.

Propensity Score Development

Propensity score modeling was used to match isolated and non-isolated patients.^{26–28} The propensity score for the likelihood of being placed on isolation precautions was developed using a multivariable logistic regression model. A total of nine covariates were included in the model: age, sex, RIW, number of hospital readmissions within 90 days, total LOS for hospital admissions within 90 days, site of admission, month of isolation, year of isolation, and Case Mix Group (CMG). Two-way interactions were included between the first six covariates, and CMG had a factor for each of the 52 most common groups, which accounted for 75 % of observations. These covariates were all selected as important predictors of isolation for infection control. Controls for each isolated case were sampled with replacement by nearest-neighbor matching using a caliper width of 0.1 standard deviations of the logit of the propensity score.^{29–31} With this approach, isolated patients are randomly ordered, and the first isolated patient is selected and matched to the control patient with the closest propensity score that is within the caliper. If multiple controls are almost equally close to the case, they are all selected and re-weighted to reflect the number of matches. This approach estimates what is called the average treatment effect in the treated (ATT); here, ATT is the effect of isolation on patients who are characteristic of those actually isolated.³² Two other effects that could have been estimated, the average treatment effect and the average treatment effect in controls, are of little interest, as they would estimate the effect of isolation in patient groups where isolation is much less likely. Covariate balance between cohorts was assessed graphically and by the standardized difference; meaningful imbalance is indicated by an absolute standardized difference of 0.10.33

Statistical Analysis

Propensity score development, matching, and the analyses below were repeated for the two separate isolated patient groups. We calculated means and percentages for the covariates in the matched and unmatched cohorts. Summaries and comparisons of outcomes were one of three types. Binary outcomes (mortality, readmissions, ED visits, and adverse events) were summarized as percentages and compared with the risk difference. Our continuous outcomes (those related to cost and LOS) were all right-skewed and were summarized by geometric means and compared by the ratio of means. Complaints and adverse events were summarized as rates (number per 100 patient-days) and compared with the difference in

rates. All estimates of effects were accompanied by 95 % confidence intervals and p values. Model diagnostics were performed to ensure that all model assumptions were valid. Two-tailed p values of 0.05 or less were considered statistically significant. All analyses were conducted using R software (Vienna, Austria), and matching and estimation of the effects of isolation used the Match package.³⁴

Sensitivity Analysis

To validate results, we also took three other approaches to matching and comparing isolated patients with controls. The first two, stratification and 1:1 nearest-neighbor matching with no ties, used the same main propensity score as in the main analyses. First, stratification involved grouping all isolated patients and all controls into ten strata based on the same propensity score used in our main analysis. Second, we used nearest-neighbor matching without replacement and used only the nearest control to each case to obtain 1:1 matches. The third approach used the package 'twang' in R to estimate the propensity score with generalized boosted regression trees (Table 1).35 To obtain the ATT, isolated patients were given a weight of 1, and all control patients were given a weight of ps/(1-ps), where ps is the patient's propensity score. The effects of isolation were estimated in weighted regression models. We examined the consistency of results across methods, but present full results only for the nearest-neighbor matching within a 0.1 caliper width. Differences between the approaches were also noted.

RESULTS

A total of 17,649 non-isolated patients were admitted to the participating hospital sites during the study period. We identified 1506 patients isolated for respiratory illnesses and 745 patients isolated for MRSA. Of the MRSA cohort, 20–25 % of cases represented clinical infection, whereas the remainder represented colonization (verbal report, authors MG and AM). The mean age of patients was 67 years, and 50 % were women. The propensity-matched cohorts had similar baseline characteristics (Table 1).

Among the respiratory isolation cohort, 207 (13.7 %) were readmitted to the hospital, and 164 (10.9 %) visited the ED within 30 days after discharge. Their mean cost for hospitalization was CAD \$7254, and their mean length of stay was 8.5 days. Most patients did not experience an adverse event or file a formal complaint with the hospital. Nearly all (99.7 %, n = 1502) patients isolated for respiratory illness were successfully matched to a non-isolated control (Table 2).

Among the cohort with MRSA, 141 (18.9 %) were readmitted to the hospital, and 85 (11.4 %) visited the ED within 30 days of discharge. Their mean cost for hospitalization was CAD \$12,129, with a mean length of stay of 12.8 days. Overall, 14.4 % of patients experienced an adverse event, and 1.1 patients per 100 admissions filed a formal complaint with the hospital. A total of 737 cases (98.9 %) had a matched control.

Table 1 Characteristics of Hospitalized Individuals by Group (pre-match and post-match)

Variable	Before matching				After matching						
	Control (n = 17,649)	MRSA (n = 745)		Respiratory illness (n = 1506)		MRSA (n = 737) 8 cases dropped		Respiratory illness (n = 1502) 4 cases dropped			
		Value	Std. diff.	Value	Std. diff.	Isolated	Control	Std. diff.	Isolated	Control	Std. diff.
Age (years)	67.9	69.0	0.08	71.7	0.29	69.0	69.0	0.00	71.7	70.6	0.06
Male (%)	50	58	0.22	53	0.10	57	58	0.00	53	55	-0.03
Number of admissions in	0.24	0.38	0.33	0.27	0.08	0.38	0.39	-0.01	0.27	0.28	-0.01
previous 90 days											
Length of stay in previous 90 days (days)	2.12	4.38	0.36	2.95	0.13	3.88	3.4	0.04	2.95	2.94	0.00
Resource Intensity Weight	1.52	2.87	0.76	1.84	0.21	2.3	2.08	0.07	1.8	1.84	-0.01
CMG196 – heart failure without cardiac catheter (%)	6	7	0.06	5	-0.04	7	7	0.01	5	5	0.00
CMG138 – viral/unspecified pneumonia (%)	3	6	0.28	25	1.47	7	7	-0.02	25	24	0.04
CMG139 – chronic obstructive pulmonary disease (%)	3	6	0.29	15	0.92	6	7	-0.02	15	16	-0.03
CMG254 – gastrointestinal hemorrhage (%)	3	2	-0.10	0	-0.25	2	2	0.00	0	0	N/A
CMG487 – lower urinary tract infection (%)	4	7	0.24	3	-0.02	7	7	-0.02	3	3	0.01

MRSA methicillin-resistant Staphylococcus aureus, CMG Case Mix Group

Outcomes for Patients Isolated for Respiratory Illness

Individuals isolated for respiratory illnesses were not significantly more likely to experience a 30-day readmission or ED visit, death, or adverse event during hospitalization or to file a formal complaint compared to non-isolated individuals (Table 3). However, those on isolation precautions for respiratory illnesses stayed 17 % longer (LOS, 8.5 days vs. 7.6 days; 95 % CI: 1.09, 1.12), stayed 9 % longer than expected, (LOS/ELOS, 1.08 vs. 1.01; 95 % CI: 1.03, 1.15), and had 23 % higher cost of care (direct cost, CAD \$7194 vs. CAD \$6294; 95 % CI: 1.14, 1.32) than non-isolated individuals. These findings are consistent across all approaches.

Outcomes for Patients Isolated for MRSA

Patients isolated for MRSA were 4.4 % more likely than non-isolated individuals to be readmitted within 30 days after discharge (rate, 19.0 % vs. 14.7 %; 95 % CI: 1.4 %, 7.3 %; Table 4). There were no significant differences in 30-day ED visits, formal complaints, or inpatient mortality rates between the cohorts.

Similar to patients with respiratory illness, patients isolated for MRSA stayed 30 % longer (LOS, 11.9 days vs. 9.1 days; 95 % CI: 1.22, 1.39), stayed 13 % longer than expected, (LOS/ELOS, 1.3 vs. 1.2; 95 % CI: 1.07, 1.20), and had 43 % higher cost of care (direct cost, CAD \$11,009 vs. CAD \$7670; 95 %

Table 2 Descriptive Statistics for Outcome Variables Comparing Isolated to Non-Isolated Cohorts (pre-match)

Outcome	Control	MRSA		Respiratory illness	
		Isolated	p value	Isolated	p value
Inpatient mortality, %	5.7	8.2	0.0048	6.9	0.0531
30-day readmission, %	12.9	18.9	< 0.0001	13.7	0.4297
30-day ED visit, %	12.8	11.4	0.2978	10.9	0.0379
Readmission or ED visit, %	18.4	22.6	0.0049	17.3	0.3260
ELOS (days)	7.3	10.2	< 0.0001	8.3	< 0.0001
LOS (days)	7.6	12.8	< 0.0001	8.5	< 0.0001
LOS/ELOS	1.2	1.4	0.0001	1.1	0.0086
Cost, CAD	6146	12129	< 0.0001	7254	< 0.0001
Cost per day, CAD	852	946	< 0.0001	855.7	0.0025
Adverse events, n/100					
Fall	5.6	12.1	< 0.0001	4.2	0.0231
Medication incident	1.5	2.1	0.1741	2.1	0.0676
Laboratory incident	0.4	0.3	0.5622	1.0	0.0035
Treatment incident	0.4	0.5	0.5731	0.6	0.2720
Infection control incident	0.1	1.9	< 0.0001	1.0	< 0.0001
Safety incident	0.3	0.3	0.8529	0.3	0.8628
Equipment incident	0.1	0.7	0.0010	0.5	0.0012
Any adverse event	6.8	14.4	< 0.0001	9.2	0.0012
Patient complaints, $n/100$	1.0	1.1	0.9113	2.7	< 0.0001

Table 3 Respiratory Isolation Cohort - Outcome Variables Comparing Isolated to Non-Isolated Cohorts (post-match)

Outcome	Isolated	Control	Effect	95 % CI	p value
Inpatient mortality*, %	6.9	8.5	-1.6	-3.9, 0.6	0.1578
30-day readmission*, %	13.7	15.7	-2.0	-5.0, 1.1	0.2088
30-day ED visit*, %	10.9	11.2	-0.2	-2.9, 2.5	0.8664
Readmission or ED visit*, %	17.4	18.5	-1.1	-4.5, 2.2	0.5005
ELOS [†] (days)	8.3	7.8	1.07	1.03, 1.12	0.0019
LOS [†] (days)	8.5	7.6	1.17	1.09, 1.25	< 0.0001
LOS/ELOS	1.08	1.01	1.09	1.03, 1.15	0.0046
Cost [†] , CAD	7194	6294	1.23	1.14, 1.32	< 0.0001
Cost per day [†] , CAD	855	826	1.05	1.02, 1.08	0.0030
Adverse events [‡] , n/100					
Fall	4.2	5.1	-0.9	-3.2, 1.3	0.4236
Medication incident	2.1	1.6	0.5	-0.7, 1.7	0.4122
Laboratory incident	0.9	0.6	0.3	-0.7, 1.2	0.5824
Treatment incident	0.6	0.7	-0.1	-0.8, 0.6	0.7554
Infection control incident	1.0	0.6	0.4	-0.4, 1.2	0.2831
Safety incident	0.3	0.1	0.3	-0.6, 1.2	0.5418
Equipment incident	0.4	0.2	0.2	-0.4, 0.8	0.4399
Any adverse event	9.1	8.9	0.2	-2.9, 3.4	0.8830
Patient complaints [‡] , n/100	2.6	1.0	1.6	-0.2, 3.4	0.0839

^{*}Reported as percentages; effect is the difference in percentage between the isolated and non-isolated cohorts

CI: 1.33, 1.54) compared to matched controls. These findings were consistent across all statistical approaches.

DISCUSSION

The results of our study show that in comparison to non-isolated patients, patients isolated for respiratory illnesses and MRSA had longer lengths of hospital stay, stayed in the hospital longer than expected, and had higher costs. Differences also existed between

the respiratory illness and MRSA cohorts. Patients isolated for MRSA were more likely to be readmitted to the hospital within 30 days after discharge, which was not found for patients isolated for respiratory illnesses. Neither isolated cohort was more likely to experience a 30-day ED visit or death or to file a formal complaint compared to matched controls.

Evidence is lacking as to whether respiratory precautions reduce nosocomial spread of respiratory disease; therefore, it is difficult to assess benefits versus adverse impacts of this practice.

Table 4 MRSA Isolation Cohort - Outcome Variables Comparing Isolated to Non-Isolated Cohorts (post-match)

Outcome	Isolated	Control	Effect	95 % CI	p value
Inpatient mortality*, %	8.0	7.0	1.0	-1.0, 3.1	0.3276
30-day readmission*, %	19.0	14.7	4.4	1.4, 7.3	0.0041
30-day ED visit*, %	11.4	11.7	-0.3	-2.6, 2.1	0.8228
Readmission or ED visit*, %	22.7	19.3	3.4	0.2, 6.6	0.0366
ELOS [†] (days)	9.8	8.4	1.15	1.09, 1.20	< 0.0001
LOS [†] (days)	11.9	9.1	1.30	1.22, 1.39	< 0.0001
LOS/ELOS	1.3	1.2	1.13	1.07, 1.20	< 0.0001
Cost [†] , CAD	11009	7670	1.43	1.33, 1.54	< 0.0001
Cost per day [†] , CAD	938	868	1.10	1.06, 1.13	< 0.0001
Adverse events ‡ , $n/100$					
Fall	10.3	8.0	2.3	-1.5, 6.1	0.2309
Medication incident	2.2	2.4	-0.3	-1.7, 1.2	0.7274
Laboratory incident	0.27	0.33	-0.1	-0.4, 0.3	0.7834
Treatment incident	0.4	0.5	-0.1	-0.6, 0.4	0.7620
Infection control incident	1.8	0.11	1.7	0.7, 2.6	0.0007
Safety incident	0.1	0.3	-0.2	-0.5, 0.1	0.2006
Equipment incident	0.5	0.2	0.3	-0.4, 1	0.3629
Any adverse event	12.4	10.7	1.7	-2.3, 5.7	0.4100
Patient complaints [‡] , $n/100$	1.09	1.25	-0.2	-1.3, 1	0.7864

^{*}Reported as percentages; effect is the difference in percentage between the isolated and non-isolated cohorts

[†] Reported as geometric means; effect is the ratio of means between the isolated and non-isolated cohorts

[‡] Reported as rates per 100 admissions; effect is the absolute differences in the rates per 100 patients between isolated and non-isolated cohorts ED emergency department, ELOS expected length of stay, LOS length of stay

[†] Reported as geometric means; effect is the ratio of means between the isolated and non-isolated cohorts

[‡] Reported as rates per 100 admissions; effect is the absolute difference in the rates per 100 patients between isolated and non-isolated cohorts MRSA methicillin-resistant Staphylococcus aureus, ED emergency department, ELOS expected length of stay, LOS length of stay

For MRSA, the reproductive ratio, R(A), for a single admission of an MRSA patient not on precautions is about 0.67. The reproductive ratio for MRSA rises above 1 only when readmissions are included, since patients can be colonized for years. Precautions reduce the R(A) by >90 %. 36,37 Thus, each patient in precautions from admission, as opposed to nonisolated, would result—from that admission—in about 0.67 × 0.35 = 0.23, i.e. just under one quarter of one infection. If more than one admission is counted, the number of prevented infections increases. In addition, if started early enough, the intervention prevents not only the first generation of transmission, but also later ones, hence further increasing the benefits. 36,37

However, isolation precautions may negatively impact several dimensions of patient care. Our study adds to knowledge in this area by noting that isolation precautions are associated with certain negative hospital-related outcomes. Previous studies have reported negative psychological effects, more adverse events, and less healthcare provider contact for isolated patients. ^{6–11,13,16–18}This may lead to a disproportionate amount of 'hospitalization-induced disability', i.e. the impact of the hospitalization itself, with subsequent de-conditioning, over and above the severity of the illness, resulting in loss of function, prolonging hospital stay, and possibly leading to an increased 30-day readmission rate.

Others have reported higher rates of adverse events, especially supportive care failures such as falls, for isolated versus non-isolated patients. Report 10 Our study does not fully support these findings, as we found no statistically significant difference in fall rates; it is possible that our propensity score-matched design better adjusted for severity of illness than studies that found a significant difference in these rates. Our work contributes novel information on the effect of isolation for respiratory illnesses and MRSA on various hospital-related outcomes, specifically 30-day readmission rates and ED visits, in-hospital mortality, LOS, ELOS, cost of care, adverse events, and patient complaints.

Our work has limitations. Although we studied three separate hospitals from two different health systems, they are all University of Toronto teaching hospitals located in downtown Toronto, Canada. Hence, this may reduce the generalizability of the results.

When comparing isolated and non-isolated patients, caution must be used in attributing differences in outcomes and cost of care to the conditions of isolation rather than illness severity. We attempted to mitigate this by using a propensity scorematched design. Additionally, the literature suggests that MRSA infections lead to worse outcomes, higher costs, and longer lengths of stay than those observed for methicillinsusceptible infections—thus, some of the negative outcomes here may be due to the underlying infection and/or an inherent but unmeasured difference in patient status, and not the isolation practice itself. Another limitation noted in 'Methods' is that the practice of isolation for infection control differs

between sites. At sites 1 and 2, respiratory (droplet and contact) precautions are used for patients admitted with fever and respiratory symptoms; at site 3, patients with respiratory or fever with cardiac symptoms are tested for respiratory viruses on admission, whereas droplet and contact precautions are used only for patients with influenza. Additionally, the sites differ in their requirement for hospital staff to wear masks when caring for patients colonized or infected with MRSA. Finally, we acknowledge that data for 30-day hospital readmissions are limited to the hospital the patient was originally discharged from. However, in our region, two-thirds of readmissions are to the same site as the initial hospitalization. Moreover, the potential for missing readmission data should be equally distributed across both isolated and non-isolated patients.

Our results have some important implications. Isolation precautions are seen as important public health and infection control measures. Despite this expected benefit, we found that these practices are associated with unintended consequences. It is possible that infection control procedures for hospitalized patients contribute to poorer outcomes that they experience, and hence more attention needs to be paid to mitigating the risks associated with isolation.

The finding of higher costs for isolated patients is important. Many healthcare institutions are remunerated through activity-based funding, which incorporates patient volume and case mix. However, isolated patients are staying longer than expected based on this methodology, and thus institutions may not be sufficiently compensated to care for them if this characteristic is not considered. Institutional remuneration schemes may require re-evaluation so that hospitals are properly compensated for isolated patients, who consist of a significant minority of the patient population on GIM wards.

CONCLUSION

Isolation precautions for infection control purposes are considered an important public health intervention. However, isolation may adversely affect the isolated individual and, in our study, was associated with longer LOS, higher costs, and increased hospital readmissions at 30 days for patients with MRSA. Balancing the benefits for the many with the harms to the few is a future challenge for hospital isolation practices.

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or integrity of any part of the work are appropriately investigated and resolved

Corresponding Author: Howard B. Abrams, MD; , TGH EN14-218, 200 Elizabeth Street, Toronto, ON M5G 2C4, Canada (e-mail: Howard.abrams@uhn.ca).

Compliance with Ethical Standards:

Conflict of Interest: The authors declare that they do not have a conflict of interest.

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