

# Trends in site of death and health care utilization at the end of life: a population-based cohort study

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## Abstract

**Background:** High rates of health care utilization at the end of life may be a marker of care that does not align with patient-stated preferences. We sought to describe trends in end-of-life care and factors associated with dying in hospital.

**Methods:** We conducted a population-level retrospective cohort study of adult decedents in Ontario between Apr. 1, 2004, and Mar. 31, 2015, using linked administrative data sets, including the Office of the Registrar General for Deaths database, the hospital Discharge Abstract Database, the National Ambulatory Care Reporting System and physicians' billing claims (Ontario Health Insurance Plan). The primary outcome was place of death. To determine health care utilization and health care costs during the 6 months before death, we also identified admissions to hospital and to the intensive care unit, emergency department visits, and receipt of mechanical ventilation and palliative care.

**Results:** In the last 6 months of life, 77.3% of 962 462 decedents presented to an emergency department, 68.4% were admitted to hospital, 19.4% were admitted to an intensive care unit, and 13.9% received mechanical ventilation. Forty-five percent of all deaths occurred in hospital, a proportion that declined marginally over time, whereas receipt of palliative care increased during terminal hospital admissions (from 14.0% in fiscal year 2004/05 to 29.3% in 2014/15,  $p < 0.001$ ) and in the last 6 months of life (from 28.1% in 2004/05 to 57.7% in 2014/15,  $p < 0.001$ ). The proportion of decedents who presented to the emergency department, were admitted to hospital or were admitted to the intensive care unit in the last 6 months of life did not change over 11 years. The mean total health care costs in the last 6 months of life were highest among those dying in hospital, with most costs attributable to inpatient medical care.

**Interpretation:** Health care utilization in the last 6 months of life was substantial and did not decrease over time. It is possible that increased capacity for palliative, hospice and home care at the end of life may help to better align health system resources with the preferences of most patients, a topic that should be explored in future studies.

High-quality end-of-life care is important for patients and families, and is an increasing focus for policy-makers and the research community. Although there is marked international variability in end-of-life care practices<sup>1-3</sup> among industrialized countries, including Canada, there are high rates of in-hospital death and use of resource-intensive services, including admission to an intensive care unit (ICU).<sup>1</sup> This resource-intensive care often does not align with patient preferences and accounts for a substantial proportion of health care expenditures.<sup>4-6</sup>

Earlier reports emphasized gaps in the provision of quality end-of-life care for dying Canadians and their families,<sup>7,8</sup> including marked variability in the provision of palliative care services.<sup>9</sup> Amid a rapidly aging Canadian population and the accompanying large increase in the prevalence of chronic diseases and frailty, whether and to what extent these gaps remain

**Competing interests:** James Downar has received speaker's fees from Boehringer-Ingelheim (Canada) and Novartis, as well as personal fees from the Ontario College of Family Physicians for participation on a steering committee for end-of-life care. Kenneth Rockwood is the founder and chief scientific officer of DGI Clinical, which works with various pharmaceutical companies to develop individualized outcome measurement and advanced data analytics. DGI Clinical is supported by the NRC's Industrial Research Assistance Program. Dr. Rockwood has asserted copyright in the Clinical Frailty Scale (through Dalhousie University's Industry Liaison Office), which is available free for use in education, research and not-for-profit health care. He has also received personal fees from various pharmaceutical companies for activities unrelated to this study. No other competing interests were declared.

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is largely unknown. Measuring trends and current health care practices at the end of life is important to better align patient preferences and health needs with health system resources and policies. However, estimates of temporal trends in the use of health care resources at the end of life are conflicting. For example, some studies suggest that rates of hospital death are declining,<sup>10,11</sup> whereas others do not.<sup>12,13</sup> In Canada, analyses between 1994 and 2004 found decreasing rates of hospital deaths;<sup>10</sup> however, whether this trend has continued or how it applies to other markers of end-of-life care is unclear.

We sought to describe contemporary temporal patterns in place of death, health care utilization and costs at the end of life and to identify factors associated with dying in a hospital in Ontario.

## Methods

### Design and settings

We conducted a population-based cohort study in Ontario, Canada's largest province (population 13.6 million, of whom about 16% are 65 years or older). Most physician and hospital services are universally insured for all residents of Ontario, through the province's single-payer, publicly funded health care system, with health services planning and delivery coordinated by the province's 14 regional Local Health Integration Networks (LHINs).

We identified individuals 18 years of age or older who died between Apr. 1, 2004, and Mar. 31, 2015 (fiscal years 2004 to 2014), using the Registered Persons Database, which contains vital statistics data on all residents of Ontario, including date of and age at death. We used the Office of the Registrar General for Deaths database to identify the cause of death. To describe health care utilization and costs before death, we linked information on decedents to population-based health administrative data sets. These data sets included the Canadian Institute for Health Information's Discharge Abstract Database, which captures data on all acute care hospital admissions in Ontario; the National Ambulatory Care Reporting System, which contains information on hospital-based ambulatory services, including emergency department visits, day surgery and outpatient clinic visits; the Continuing Care Reporting System, which includes demographic and clinical information on individuals receiving facility-based continuing care services; and the Home Care Database, which captures data on receipt of publicly funded home care services, including date and type of services. Information on physician services, including date and type of medical services provided and billing claims for inpatient and outpatient services, was obtained from the Ontario Health Insurance Plan (OHIP) database. These data sets were linked using unique encoded identifiers and were analyzed at ICES.

### Baseline characteristics

Decedents' demographic characteristics, including age, sex, income, location of residence and cause of death, were measured at the time of death. Income (ascribed as neighbourhood income quintile) and urban or rural residence were

determined by linking each decedent's residential postal code at the time of death to the Statistics Canada 2006 census. Cause of death was determined from the Office of the Registrar General for Deaths database and classified according to the *International Statistical Classification of Diseases and Related Health Problems, 10th revision* (ICD-10). A primary chronic condition was defined according to a 9-level variable indicating the presence of cancer, chronic obstructive pulmonary disease (COPD), coronary artery disease, congestive heart failure, peripheral vascular disease, severe liver disease, diabetes with end organ failure, renal failure or dementia.<sup>14</sup> Decedents were assigned to one of these conditions on the basis of ICD-10 diagnostic codes during the last hospitalization, which were available for more than 90% of decedents. When multiple chronic conditions were identified, the following hierarchy was applied: most responsible diagnosis, then comorbid diagnoses, then minor diagnoses.

For each LHIN, we determined the number of acute care beds per 100 000 population and calculated the median bed capacity for the province over the study period. We classified LHINs falling below and above the provincial median for acute care beds as having "low acute care bed capacity" and "high acute care bed capacity," respectively, and categorized decedents according to the acute care bed capacity of their LHIN.

### Outcomes

Our primary outcome was place of death, classified as a hospital, the patient's home, a long-term care facility or another location. Deaths in hospital were further classified according to whether they occurred in an ICU or another location. Secondary outcomes were the factors associated with dying in hospital, the use of health services during the 6 months before death (including acute care hospitalizations and emergency department admissions, receipt of mechanical ventilation and palliative care, and number of physician visits) and health care costs.

We classified place of death using information contained in the vital statistics (Registered Persons Database), hospital (Discharge Abstract Database) and emergency department (National Ambulatory Care Reporting System) databases. For decedents who were admitted to hospital within the 6 months before death, we identified the number of admissions to an ICU using special care unit codes in the Discharge Abstract Database.<sup>15,16</sup> We also measured the total number of inpatient and ICU days for these individuals. We identified receipt of palliative care using a comprehensive set of diagnostic and billing codes across multiple providers and health care settings, including information contained in the home care (Home Care Database), hospital (Discharge Abstract Database), emergency department (National Ambulatory Care Reporting System), long-term care and complex continuing care (Continuing Care Reporting System) and physician billing (OHIP) databases, consistent with prior studies.<sup>17,18</sup>

We estimated health care costs (from the perspective of the health care payer, the Ontario Ministry of Health and Long-term Care) from the health administrative data sets using an established approach for person-level costing.<sup>19,20</sup> This algorithm summarizes costs attributed to several health sectors,

for example, inpatient acute care, emergency department visits, complex continuing care and physician services (for details, see Appendix 1, available at [www.cmajopen.ca/content/7/2/E306/suppl/DC1](http://www.cmajopen.ca/content/7/2/E306/suppl/DC1)). Briefly, we derived costs for institutional care (e.g., hospital and rehabilitation admissions) by applying resource intensity weights for the specific care sector to the cost per weighted case determined from amounts paid by the Ontario Ministry of Health and Long-term Care. Costs for health services associated with specific fees (e.g., physician visits, diagnostic tests, medications) were based directly on the fee paid per use of these services.<sup>19</sup> All costs were adjusted to 2014 Canadian dollars.

### Statistical analyses

We used descriptive statistics to characterize the location of death and the use of health services for all decedents and according to groups defined by age, chronic condition and year of death. We summarized categorical and continuous variables as proportions and means (standard deviations [SDs]) or medians (interquartile ranges [IQRs]), respectively. Because of our large sample size and the potential for small differences to be statistically significant, we do not report *p* values for group comparisons. We compared the characteristics of individuals who died in hospital and out of hospital using standardized differences, where a difference less than 0.1 is commonly used to indicate negligible differences between groups.<sup>21</sup> To summarize differences in health care resource use between groups and over time (fiscal year 2004 v. fiscal year 2014), we calculated absolute differences (with 95% confidence intervals [CIs]). We used the Cochran–Armitage trend test to examine changes over time in the location of death. We used multilevel logistic regression, accounting for the clustering of decedents within LHINs, to identify factors associated with dying in hospital. We considered putative factors on the basis of their clinical relevance and availability in the data set. At the patient level, these factors were age, sex, Charlson comorbidity score, rural residence, income, year and cause of death. The LHIN acute care bed capacity was included as a region-level factor. All analyses were conducted using SAS Enterprise Guide 7.1 (SAS Institute).

### Ethics approval

This study was approved by the institutional review board at Sunnybrook Health Sciences Centre, Toronto, Ontario.

## Results

### Patterns and temporal trends in place of death and use of palliative care

There were 962 462 decedents in Ontario between fiscal year 2004 and fiscal year 2014. The mean age at death was 76.3 (SD 14.8) years, and 50.1% of decedents were women (Table 1). Overall, 45.0% of decedents died in hospital. The proportion who died in hospital declined only slightly over time (45.9% in fiscal year 2004 and 43.2% in fiscal year 2014; absolute difference 2.7%, 95% CI 2.3% to 3.2%; test for trend *p* < 0.001; Figure 1), with a corresponding increase

in deaths at home (37.9% and 43.2%, respectively; absolute difference 5.3%, 95% CI 4.9% to 5.8%; test for trend *p* < 0.001; Figure 1). Patterns in place of death varied according to the primary chronic condition, with those who had liver disease (73.3%), peripheral vascular disease (67.5%), congestive heart failure (61.1%), COPD (60.8%), coronary artery disease (57.7%) and cancer (54.0%) being more likely to die in hospital than those who had dementia (34.2%) (absolute differences ranging from 19.8% [95% CI 19.4% to 20.2%] to 39.1% [95% CI 38.3% to 39.8%]; see Figure A1 in Appendix 2, available at [www.cmajopen.ca/content/7/2/E306/suppl/DC1](http://www.cmajopen.ca/content/7/2/E306/suppl/DC1)).

The proportion of decedents who received palliative care during the terminal hospital admission more than doubled, from 14.0% in fiscal year 2004 to 29.3% in fiscal year 2014 (absolute difference 15.3%, 95% CI 14.9% to 15.7%; Figure 1). Decedents with cancer had the highest rates of palliative care during the terminal hospitalization throughout the study (37.9% in fiscal year 2004 to 44.0% in fiscal year 2014; absolute difference 6.1%, 95% CI 5.2% to 7.1%), but increased uptake of palliative care was observed for patients with all conditions, including those with dementia (7.9% in fiscal year 2004 to 23.7% in fiscal year 2014; absolute difference 15.8%, 95% CI 14.7% to 16.8%), coronary artery disease (7.6% in fiscal year 2004 to 25.6% in fiscal year 2014; absolute difference 18.0%, 95% CI 16.4% to 19.6%) and congestive heart failure (10.3% in fiscal year 2004 to 35.3% in fiscal year 2014; absolute difference 25.0%, 95% CI 24.0% to 26.2%) (see Figure A1 in Appendix 2). When the analysis was restricted to decedents who died in hospital, receipt of palliative care during the terminal hospital admission increased significantly, from 30.5% in fiscal year 2004 to 67.9% in fiscal year 2014 (absolute difference 37.4%, 95% CI 36.8% to 38.1%). Similar increases were observed among patients who had chronic conditions, with decedents who had cancer having among the highest rates of palliative care during the terminal hospital admission (see Figure A2 in Appendix 2).

For the adjusted analysis, it was necessary to recut the data in the course of responding to requests from the reviewers and editors; by the time of this re-analysis, there had been updates to the project data sources and an additional 21 decedents were included. In this adjusted analysis, men and older decedents were more likely to die in hospital than out of hospital (Table 2), with the association between age and death in hospital attenuating among those 85 years of age and older. In addition, the odds of dying in hospital were higher for those residing in more rural areas (odds ratio [OR] 1.12, 95% CI 1.11 to 1.13) and neighbourhoods with the lowest (compared with the highest) income quintile (OR 1.13, 95% CI 1.12 to 1.14). There was no difference in the odds of in-hospital death among decedents living in LHINs with high and low acute care bed capacity (Table 2).

### Decedents' use of health care resources in last 6 months of life

The majority of decedents visited the emergency department (77.3%) and saw 10 or more physicians (52.6%) during the last

**Table 1 (part 1 of 2): Demographic characteristics of Ontario decedents, Apr. 1, 2004, to Mar. 31, 2015 (fiscal year 2004 to fiscal year 2014)**

Characteristic	Location of death; % of decedents*			Standardized difference‡
	All locations n = 962 462	In hospital n = 433 074	Out of hospital n = 529 388†	
Age at death, yr, mean ± SD	76.3 ± 14.8	75.8 ± 13.5	76.6 ± 15.8	0.06
Age category at death, yr				
18–49	5.8	4.6	6.8	0.10
50–54	3.4	3.3	3.5	0.01
55–59	4.7	4.7	4.6	0.01
60–64	6.0	6.4	5.6	0.03
65–69	7.4	8.3	6.6	0.07
70–74	9.5	11.0	8.3	0.09
75–79	12.8	14.7	11.4	0.10
80–84	16.6	17.9	15.6	0.06
≥ 85	33.7	28.9	37.6	0.18
Sex, female	50.1	48.0	51.9	0.08
Charlson Comorbidity Index score				
0	14.1	6.5	20.4	0.42
1	13.9	14.3	13.5	0.02
≥ 2	61.7	79.2	47.3	0.70
Missing	10.3	0	18.7	NA
Income quintile§				
1 (lowest)	23.5	23.8	23.3	0.02
2	21.1	21.9	20.4	0.04
3	19.1	19.2	19.0	0
4	18.5	18.2	18.8	0.02
5 (highest)	17.8	16.9	18.5	0.04
Rural residence§	14.8	15.3	14.5	0.02
Chronic conditions				
Cancer	20.3	24.4	17.0	0.18
Congestive heart failure	11.5	15.6	8.1	0.23
Dementia	9.0	6.9	10.8	0.14
COPD	5.1	6.9	3.6	0.15
Coronary artery disease	4.7	6.1	3.6	0.11
Renal failure	3.0	4.1	2.1	0.12
Severe liver disease	1.5	2.5	0.7	0.14
Peripheral vascular disease	1.0	1.5	0.6	0.09
Diabetes with end-organ damage	0.1	0.1	0.1	0
None	43.8	32.1	53.3	0.44
Cause of death (top 20)				
Ischemic heart disease	13.6	9.9	16.7	0.20
Cancer				
Lung	6.2	7.1	5.5	0.06
Colorectal	3.0	3.0	2.9	0.01
Hematologic	2.5	3.5	1.6	0.12
Breast	1.8	1.8	1.8	0

**Table 1 (part 2 of 2): Demographic characteristics of Ontario decedents, Apr. 1, 2004, to Mar. 31, 2015 (fiscal year 2004 to fiscal year 2014)**

Characteristic	Location of death; % of decedents*			Standardized difference‡
	All locations <i>n</i> = 962 462	In hospital <i>n</i> = 433 074	Out of hospital <i>n</i> = 529 388†	
Prostate	1.4	1.2	1.5	0.02
Pancreas	1.4	1.4	1.4	0
Dementia and Alzheimer disease	5.9	2.5	8.7	0.27
Cerebrovascular disease	5.1	6.0	4.3	0.08
Chronic lower respiratory disease	3.5	4.1	3.0	0.06
Diabetes mellitus	2.8	2.2	3.4	0.07
Influenza and pneumonia	2.2	3.0	1.5	0.10
Diseases of urinary system	2.0	2.8	1.3	0.11
Falls	1.6	2.4	1.0	0.11
Heart failure and complications	1.6	1.7	1.5	0.02
Cirrhosis and other liver disease	1.2	1.9	0.7	0.11
Hypertensive disease	1.0	0.6	1.4	0.08
Intentional self-harm	1.0	0.2	1.6	0.16
Cardiac arrhythmias	1.0	1.2	0.8	0.04
Sepsis	0.9	1.6	0.3	0.13

Note: COPD = chronic obstructive pulmonary disease, NA = not applicable, SD = standard deviation.  
 \*Except where indicated otherwise.  
 †Out-of-hospital deaths occurred in the home (*n* = 389 250), in long-term care facilities (*n* = 77 280) and in other locations (*n* = 62 858).  
 ‡Standardized differences for comparisons by location of death, where a difference less than 0.1 is commonly deemed to indicate negligible differences between groups.  
 §Information on income and location of residence was missing for 0.5% (5213) and 0.1% (617) of decedents, respectively.

6 months of life (Table 3). In the same period, 68.4% were admitted to hospital at least once (median number of hospitalizations 1, IQR 0 to 2), about 1 in every 5 decedents was admitted to an ICU, and 13.9% received mechanical ventilation. Decedents spent a median of 6 (IQR 0 to 19) days in hospital during the last 6 months of life. Over the 11-year study period, there was no substantial change in the proportion of decedents who were admitted to hospital or the ICU or who used emergency department services in the last 6 months of life. There was, however, an increasing trend in the proportion of decedents seeing 10 or more physicians and receiving mechanical ventilation in the 6 months before death (Table 3).

Rates of health service use during the last 6 months of life were higher among decedents who died in hospital than among those who died in other locations, including a higher proportion of admissions to the ICU (36.3% v. 5.6%; absolute difference 30.7%, 95% CI 30.6% to 30.9%) and a higher proportion receiving mechanical ventilation (27.7% v. 2.6%; absolute difference 25.1%, 95% CI 25.0% to 25.2%). Among individuals who died in hospital, rates of use of these high-intensity resources (ICU and mechanical ventilation) were lower among those who received palliative care within the last 6 months of life (Table 4). Compared with decedents who had a chronic organ failure-related condition (e.g., COPD, congestive heart failure), those who had cancer or dementia were less likely to be admitted to the ICU and less likely to receive mechanical ventilation (see Table A1 in Appendix 3, available at [www.cmajopen](http://www.cmajopen).

[ca/content/7/2/E306/suppl/DC1](http://cmajopen.ca/content/7/2/E306/suppl/DC1)). About half of decedents received palliative care services during the last 6 months of life, with use of these services increasing significantly over the study period (from 28.1% in fiscal year 2004 to 57.7% in fiscal year 2014; absolute difference 29.6%, 95% CI 29.2% to 30.1%) (Figure 2). Across all years, receipt of palliative care during the last 6 months of life was highest among decedents dying in a long-term care facility, ranging from 61.9% in fiscal year 2004 to 88.3% in fiscal year 2014 (absolute difference 26.4%, 95% CI 25.1% to 27.8%) (Figure 2).

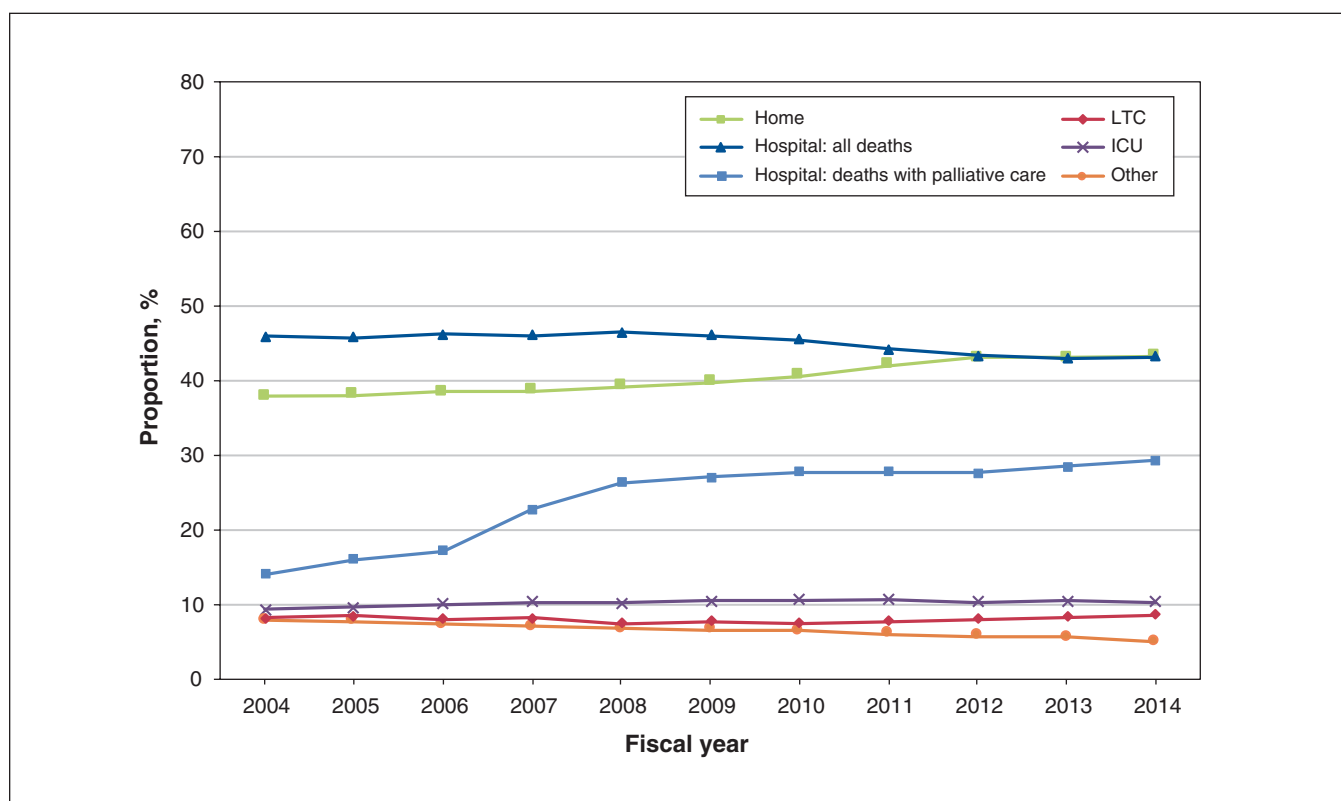
### Health care costs in the last 6 months of life

Median total health care costs in the last 6 months of life were higher among decedents who died in hospital (\$32 291, IQR \$18 245 to \$55 308) than among those who died out of hospital (\$22 895, IQR \$7636 to \$36 931) (see Table A2 in Appendix 3). Overall, most non-patient-borne health care costs were attributed to acute care inpatient costs, which ranged from a median of \$0 (IQR \$0 to \$11 060) among decedents who died in settings other than a hospital to \$16 284 (IQR \$7290 to \$33 216) among those dying in hospital.

### Interpretation

In this population-based study, we found that nearly half of all deaths occurred in hospital, with one-quarter of these deaths occurring in the ICU. We also found substantial health care





**Figure 1:** Temporal trends in location of death for Ontario, from Apr. 1, 2004, to Mar. 31, 2015 (fiscal year 2004 to fiscal year 2014). Note: ICU = intensive care unit, LTC = long-term care.

utilization among decedents in the 6 months before death. For example, 3 of every 4 decedents were admitted to a hospital or emergency department. Between fiscal years 2004 and 2014, deaths in hospital decreased, with concomitant increases in the proportions of terminal hospitalizations involving palliative care and deaths at home; however, other markers of acute care and use of hospital-based health care resources did not change.

Like others,<sup>1-3,22,23</sup> we found considerable use of acute care health resources at or near the time of death. Overall, these findings suggest a continued reliance on hospitals to provide end-of-life care and underscore the gaps in options for palliative, hospice or home-based care at the end of life within the Canadian health care system.<sup>7-9,24</sup> A lack of outpatient structures and processes for delivering end-of-life services, including comprehensive palliative care services, may therefore help to explain the observed utilization patterns, especially in more rural and remote areas. Further research is needed to establish whether, and which models of, palliative care service are effective in reducing the use of acute care health resources by individuals who are dying.

In our study, we observed temporal trends in markers of end-of-life care practices, notably decreasing rates of in-hospital deaths and increasing receipt of palliative care. From 1980 to 1997, more than 66% of Ontarians died in hospital, with relatively stable rates over that period.<sup>25</sup> Although methodological differences partly explain the substantial drop in deaths occurring in hospital between that earlier report and the findings of the current study, the change (from 66% to 45%) may also

have been related to substantial health care restructuring efforts in Ontario, which resulted in the closure of thousands of hospital beds and the creation of new long-term care beds.<sup>26</sup> Nonetheless, that almost half of decedents received some form of palliative care in the last 6 months of life is encouraging and generally similar to recent observations using comparable methods.<sup>17,18</sup> We extend these earlier findings by providing evidence of an increasing trend in the receipt of palliative care at the population level. This trend is important, as our finding likely reflects increasing recognition of the benefits of palliative care for improving care at the end of life and the growth of palliative care services in the last decade.<sup>27-31</sup> It is also plausible that shifting awareness regarding palliative care services among individuals who are dying and their caregivers might have influenced end-of-life care decisions. Future studies aimed at understanding the drivers of the observed increase will be informative to efforts to improve care at the end of life.

Our finding that individuals dying in hospital had higher costs of care in the last 6 months of life and that about 50% of these costs were attributable to inpatient care align with previous data from Ontario.<sup>32</sup> Although we do not propose that all such admissions can or should be avoided, given the challenges of predicting death,<sup>33</sup> these data suggest that there may be opportunities to improve end-of-life care that is better aligned with patients' wishes. In particular, the observed higher risk of dying in hospital among individuals from rural residences and lower-income neighbourhoods may indicate

**Table 2 (part 1 of 2): Multilevel model of factors associated with dying in hospital versus all other locations\***

Factor	OR (95% CI)	
	Unadjusted†	Adjusted
<b>Patient-related</b>		
Age category, yr		
18–49	1.0 (Ref.)	1.0 (Ref.)
50–54	1.40 (1.36 to 1.44)	1.34 (1.30 to 1.38)
55–59	1.54 (1.50 to 1.58)	1.47 (1.43 to 1.51)
60–64	1.70 (1.66 to 1.74)	1.63 (1.59 to 1.67)
65–69	1.88 (1.84 to 1.92)	1.80 (1.76 to 1.84)
70–74	1.98 (1.94 to 2.03)	1.91 (1.86 to 1.95)
75–79	1.92 (1.88 to 1.96)	1.89 (1.85 to 1.93)
80–84	1.71 (1.68 to 1.74)	1.74 (1.70 to 1.78)
≥ 85	1.15 (1.13 to 1.17)	1.22 (1.20 to 1.25)
Sex		
Men	1.0 (Ref.)	1.0 (Ref.)
Women	0.86 (0.85 to 0.86)	0.88 (0.87 to 0.89)
Income quintile		
1 (lowest)	1.0 (Ref.)	1.0 (Ref.)
2	1.04 (1.03 to 1.05)	1.02 (1.01 to 1.04)
3	0.96 (0.95 to 0.98)	0.96 (0.94 to 0.97)
4	0.93 (0.92 to 0.94)	0.92 (0.91 to 0.93)
5 (highest)	0.89 (0.88 to 0.90)	0.88 (0.88 to 0.89)
Rural residence		
No	1.0 (Ref.)	1.0 (Ref.)
Yes	1.13 (1.11 to 1.14)	1.12 (1.11 to 1.13)
Cause of death		
Ischemic heart disease	1.0 (Ref.)	1.0 (Ref.)
Cancer		
Lung	2.16 (2.12 to 2.21)	2.00 (1.96 to 2.04)
Colorectal	1.76 (1.72 to 1.81)	1.71 (1.66 to 1.75)
Hematologic	3.59 (3.48 to 3.69)	3.51 (3.41 to 3.61)
Breast	1.65 (1.60 to 1.71)	1.79 (1.73 to 1.85)
Prostate	1.43 (1.38 to 1.48)	1.31 (1.26 to 1.36)
Pancreas	1.66 (1.60 to 1.72)	1.60 (1.54 to 1.66)
Dementia and Alzheimer disease	0.48 (0.47 to 0.50)	0.55 (0.54 to 0.56)
Cerebrovascular disease	2.35 (2.30 to 2.40)	2.48 (2.43 to 2.53)
Chronic lower respiratory disease	2.31 (2.25 to 2.37)	2.28 (2.22 to 2.34)
Diabetes mellitus	1.08 (1.05 to 1.11)	1.06 (1.03 to 1.09)
Influenza, pneumonia	3.35 (3.25 to 3.45)	3.72 (3.61 to 3.83)
Diseases of urinary system	3.60 (3.49 to 3.72)	3.84 (3.72 to 3.97)
Falls	4.21 (4.07 to 4.37)	4.67 (4.50 to 4.84)

**Table 2 (part 2 of 2): Multilevel model of factors associated with dying in hospital versus all other locations\***

Factor	OR (95% CI)	
	Unadjusted†	Adjusted
Heart failure	1.92 (1.85 to 1.98)	2.09 (2.02 to 2.17)
Cirrhosis and other liver diseases	4.65 (4.47 to 4.85)	4.64 (4.45 to 4.83)
Hypertensive disease	0.75 (0.72 to 0.79)	0.82 (0.79 to 0.86)
Intentional self-harm	0.16 (0.15 to 0.18)	0.19 (0.18 to 0.21)
Cardiac arrhythmias	2.66 (2.55 to 2.78)	2.94 (2.82 to 3.07)
Sepsis	8.90 (8.42 to 9.41)	9.29 (8.78 to 9.82)
Year‡	0.86 (0.85 to 0.88)	0.81 (0.79 to 0.82)
<b>LHIN-related</b>		
Acute care bed capacity§		
Low	1.0 (Ref.)	1.0 (Ref.)
High	1.00 (0.97 to 1.03)	1.02 (1.00 to 1.05)

Note: CI = confidence interval, LHIN = Local Health Integration Network, OR = odds ratio, Ref. = reference category.  
 \*The sample size of 962 483 for this analysis reflects multilevel models performed on Oct. 25, 2018.  
 †Accounts for clustering by LHIN.  
 ‡ORs reflect per 10-year unit change.  
 §Defined using the median bed capacity per 100 000 population for the province over the study period. LHINs falling below and above the provincial median were classified as having “low acute care bed capacity” and “high acute care bed capacity,” respectively.

disparities in access to end-of-life care services, including availability of home, palliative and hospice care. Further, although our data are unable to provide insights regarding the appropriateness of care, the observation that a large proportion of decedents received care from 10 or more physicians within the last 6 months of life raises concerns about the potential for fragmented care. Care continuity may be important not only in terms of the quality of care provided to these individuals, but inadequacies may also adversely affect health care costs.<sup>34</sup>

Our findings suggest important differences in care depending upon predominant terminal clinical diagnoses. For example, there were higher rates of ICU admission, receipt of mechanical ventilation in the period before death and hospital-based deaths among those with organ failure (e.g., congestive heart failure, COPD, coronary artery disease, liver failure, renal failure) than among those with cancer or dementia. This finding has implications for planning health services, supporting individuals dying from chronic non-cancer conditions and potentially supporting subspecialist clinicians treating those conditions. Having greater options for end-of-life and palliative care in the community setting or in hospice may increase quality-adjusted life-days, and may reduce both hospitalization rates and overall costs.<sup>22,35–37</sup> This is particularly relevant for individuals dying with non-cancer conditions, for whom options for palliative care within the Canadian setting are limited.<sup>38</sup>

**Table 3: Utilization of health care services in the last 6 months of life among all decedents**

Fiscal year	Health care service utilized; % of patients*						
	Hospital admission	Hospital days, median (IQR)	ICU admission	ICU days,† mean ± SD	ED visits	Saw ≥ 10 physicians	Mechanical ventilation
All	68.4	6 (0 to 19)	19.4	1.7 ± 6.5	77.3	52.6	13.9
2004	68.7	6 (0 to 20)	19.1	1.5 ± 6.1	76.4	44.8	12.3
2005	68.8	6 (0 to 20)	19.2	1.6 ± 6.5	77.1	45.9	12.6
2006	68.2	6 (0 to 20)	19.1	1.6 ± 6.2	77.1	48.7	13.2
2007	68.4	6 (0 to 20)	19.3	1.6 ± 6.3	77.2	51.3	13.6
2008	68.2	6 (0 to 20)	18.6	1.6 ± 6.4	77.0	52.2	13.5
2009	68.6	6 (0 to 19)	19.5	1.7 ± 6.9	77.2	53.8	14.3
2010	68.5	6 (0 to 19)	19.2	1.7 ± 6.4	77.6	54.8	14.3
2011	68.5	6 (0 to 19)	19.5	1.8 ± 6.9	77.7	55.7	14.3
2012	67.9	6 (0 to 19)	19.7	1.7 ± 6.6	77.2	55.1	14.5
2013	68.5	6 (0 to 19)	20.1	1.8 ± 6.8	77.7	57.1	15.1
2014	68.6	6 (0 to 18)	19.9	1.8 ± 6.7	77.9	57.5	15.1
Absolute difference, % (95% CI)‡	0 (−0.5 to 0.4)	NA	0.7 (0.4 to 1.1)	0.3 (0.2 to 0.4)	1.5 (1.1 to 1.9)	12.6 (12.2 to 13.1)	2.8 (2.5 to 3.1)

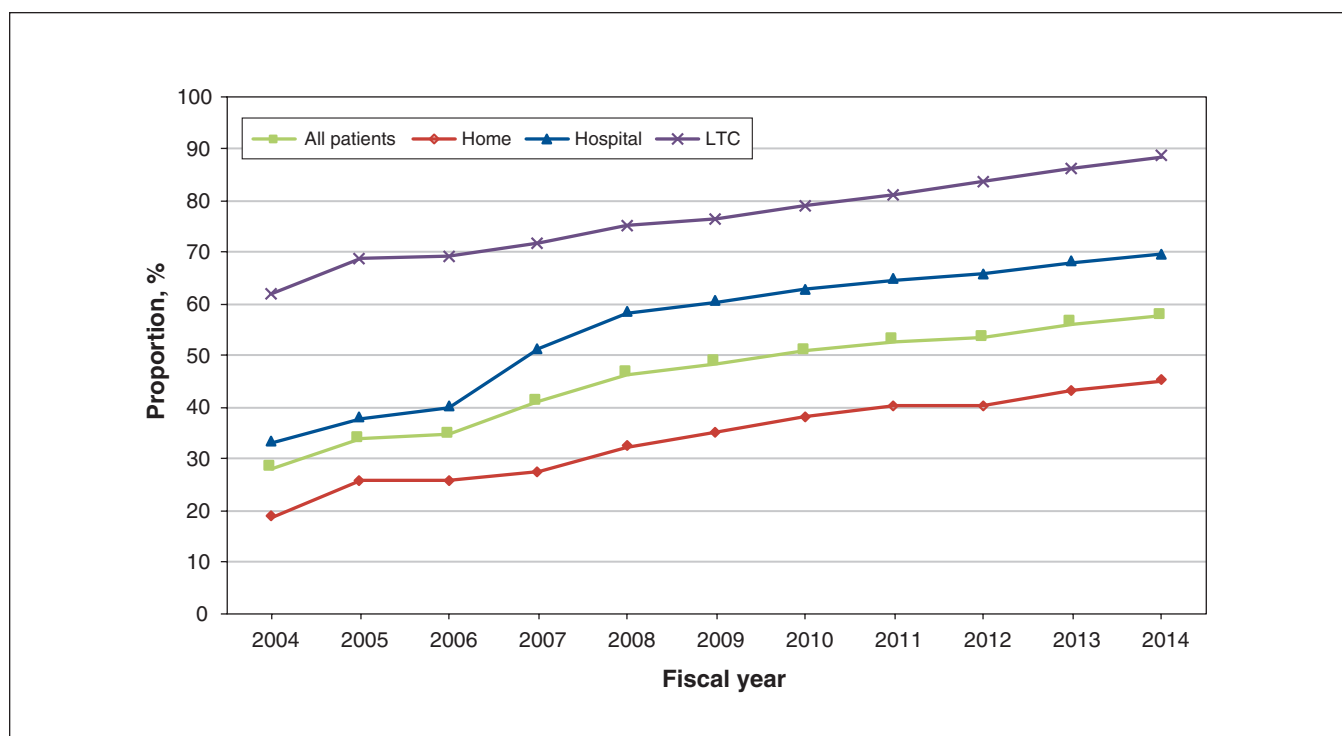
Note: CI = confidence interval, ED = emergency department, ICU = intensive care unit. IQR = interquartile range, NA = not applicable, SD = standard deviation.  
 \*Except where indicated otherwise.  
 †For all years, the median number of ICU days was 0.  
 ‡The absolute difference, calculated by subtraction, compares utilization of each health care service between fiscal year 2004 and fiscal year 2014.

**Table 4: Utilization of health care services in the last 6 months of life among individuals who died in hospital (n = 433 074)**

Fiscal year	Health care service utilized; % of patients receiving ("yes") and not receiving ("no") palliative care									
	Hospital admission		ED visits		Saw ≥ 10 physicians		ICU admission		Mechanical ventilation	
	Yes	No	Yes	No	Yes	No	Yes	No	Yes	No
All	99.8	99.5	96.2	96.0	76.9	66.7	26.6	48.6	18.9	38.9
2004	99.9	99.9	93.2	95.6	69.3	58.4	16.5	44.5	8.2	31.9
2005	99.9	99.8	94.0	95.9	69.9	59.0	18.5	45.9	10.2	33.8
2006	99.9	99.8	94.8	96.1	72.9	64.5	19.7	45.6	11.2	35.5
2007	99.9	99.6	95.4	96.1	75.2	67.5	24.0	47.4	16.0	37.5
2008	99.8	99.4	95.9	95.9	75.8	68.0	25.0	46.8	18.1	37.5
2009	99.8	99.2	96.1	95.6	77.0	69.1	27.0	49.7	19.6	41.1
2010	99.8	99.1	96.6	95.7	77.7	70.7	27.1	50.6	19.9	42.7
2011	99.8	99.3	96.8	96.3	78.7	72.0	28.6	52.1	20.8	43.6
2012	99.8	99.3	97.2	96.2	78.9	71.6	30.2	52.5	21.9	44.7
2013	99.8	99.2	97.2	96.1	80.5	73.2	31.4	54.6	23.6	46.9
2014	99.8	99.2	97.3	96.4	80.6	73.3	31.5	53.8	24.2	46.5
Absolute difference, % (95% CI)*	0.3 (0.3 to 0.3)		0.2 (0.1 to 0.3)		10.2 (10.0 to 10.5)		22.0 (21.7 to 22.3)		20.1 (19.8 to 20.3)	

Note: CI = confidence interval, ED = emergency department, ICU = intensive care unit.  
 \*The absolute difference, calculated by subtraction, compares utilization of each health care service between fiscal year 2004 and fiscal year 2014 for individuals who did ("yes") and did not ("no") receive palliative care in the last 6 months of life.





**Figure 2:** Temporal trends in proportion of patients receiving palliative care in the last 6 months of life, overall and by location of death, from Apr. 1, 2004, to Mar. 31, 2015 (fiscal year 2004 to fiscal year 2014). Note: LTC = long-term care.

## Limitations

Our population-based observational study describes associations but cannot explain the causes for high rates of health care utilization near the end of life. In particular, we lacked information on several factors previously shown to be associated with the use of health services, for example, patient and family preferences, provider decision-making, and differential availability of hospice and palliative care services across various patient populations and geographic locations.<sup>22,39–41</sup> These factors may have important influences on decisions with respect to location of death. Furthermore, we have reported costs from the perspective of the health care system payer; therefore, we have not considered all societal costs, including out-of-pocket expenses and informal care costs. These latter costs may vary by chronic condition<sup>42</sup> and can be a substantial burden on patients and informal care providers.<sup>42,43</sup> We used primarily physician billing codes to identify receipt of palliative care, so we likely underestimated the reported rates, particularly for care delivered in non-acute care settings and by non-physicians.<sup>17</sup> It is also plausible that changes in coding practices over time might have affected our reported estimates. However, given the relatively short duration of our study and the lack of sharp spikes in year-to-year trends, we believe that such changes would have minimal effects on our results and the conclusions drawn.

## Conclusion

In this population-based study, we found decreasing rates of deaths in hospital and a corresponding increase in the provision of palliative care to dying Canadians. However, other

markers of end-of-life care did not change over time, with about three-quarters of decedents being admitted to hospital and about 1 in 5 being admitted to the ICU during the last 6 months of life. Rates of hospitalization before death were higher for those living in more rural and economically disadvantaged neighbourhoods. It is possible that increased capacity for palliative, hospice and home care at the end of life may help to better align health system resources with the preferences of most patients, a topic that should be explored in future studies.

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