

BMJ Open Access to palliative care by disease trajectory: a population-based cohort of Ontario decedents

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To cite: Seow H, O'Leary E, Perez R, *et al.* Access to palliative care by disease trajectory: a population-based cohort of Ontario decedents. *BMJ Open* 2018;**8**:e021147. doi:10.1136/bmjopen-2017-021147

► Prepublication history and additional material for this paper are available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2017-021147>).

Received 13 December 2017
Revised 16 February 2018
Accepted 19 February 2018



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ABSTRACT

Objectives To examine access to palliative care between different disease trajectories and compare to other geographic areas.

Design A retrospective population-based decedent cohort study using linked administrative data.

Setting Ontario, Canada.

Participants Ontario decedents between 1 April 2010 and 31 December 2012. Patients were categorised into disease trajectories: terminal illness (eg, cancer), organ failure (eg, chronic heart failure), frailty (eg, dementia), sudden death or other.

Interventions Receipt of palliative care services from institutional and community settings, derived from a validated list of palliative care codes from multiple administrative databases.

Outcome measures Receiving any palliative care services in the last year of life (yes/no), intensity (total days) and time of initiation of palliative care, in hospital and community sectors. Multivariable analysis examined the association between disease trajectory and the receipt of palliative care in the last year of life.

Results We identified 235 159 decedents in Ontario. In the last year of life, 88% of terminal illness, 44% of organ failure and 32% of frailty decedents accessed at least one palliative care service. Most care was provided during an inpatient hospitalisation. Terminal illness decedents received twice as many palliative care days (mean of 49 days) compared with organ failure and frailty decedents. Patients with terminal illness initiated palliative care median of 107 days before death compared with median of 19 days among those using the US Medicare hospice benefit.

Conclusions Terminal illness decedents are more likely to receive any palliative care, with increased intensity and earlier before death than organ failure or frailty decedents. These data serve as a useful comparison for other countries with similar and different healthcare systems and eligibility criteria.

INTRODUCTION

With the population ageing and living longer with more comorbidities, health systems are focused on providing quality end-of-life care through improved palliative care services.^{1 2} Earlier availability of palliative care to patients with terminal illness has been shown to improve quality of life, reduce late-life health

Strengths and limitations of this study

- This study examines palliative care access and time to initiation across a comprehensive list of health-care services by disease trajectory.
- This is a large population-based study, within a universal health system, where patients have access to both institutional and community palliative care.
- This work provides a measure of access and time to initiation to palliative care for patients in all trajectories and can be compared with other countries.
- Using administrative health data to capture use of palliative care is limited by undercoding of palliative care delivered, particularly in the community setting.
- We are unable to account for the quality of care, privately obtained care or patients' end-of-life care preferences and how those differ between trajectories.

services use and even extend survival.^{3–5} However, palliative care is often not delivered or initiated until very late in the dying trajectory. Research shows that dying occurs in three main trajectories: (1) terminal illness, typical of cancer (high function followed by acute decline); (2) organ failure, typical of heart and lung disease (medium–high function, intermittent acute exacerbations and partial recovery) and (3) frailty, typical of dementia (low function and prolonged gradual dwindling).^{6–8}

Evidence shows that palliative care is more often provided to patients with cancer versus those without cancer^{9–14} because of the 'predictability' of decline^{8 15 16} and the history of hospice care for patients with cancer. This 'predictability' can sometimes be formalised into health policy, such as in the US Medicare Hospice Benefit, which requires a doctor's certification that death is expected within 6 months and that the patient forego any hospital or curative care. Whereas in other countries, like the UK, Australia and Canada, the eligibility criteria for palliative care does not require either condition. Given the growing body of literature of the benefits of

early palliative care in non-cancer diagnoses,^{17–20} there is a dearth of research describing how access to palliative care, particularly time to initiation before death and intensity and type of service use, differs by disease trajectory, and how that may be influenced by health system and various criteria to access palliative care at a population level.

This study focuses on patients in Ontario, Canada, who can access palliative care services in community and institutional settings without foregoing curative treatment through its universal insured hospital and physician system.²¹ Criteria for palliative care referral in the hospital is at the physician's discretion; whereas in the community, they often use the 'surprise question' of not being surprised if the patient died within a year,²² combined with performance status decline.²³ In short, eligibility in Ontario is not formally standardised, which is unlike the standardised criteria of the Gold Standards Framework, which is widespread in the UK. Ontario is the largest province in Canada and has the highest number of deaths.²⁴ Previous studies have shown that half of patients in Ontario received at least one palliative care service in their last year of life,²⁵ though they did not examine variations by disease trajectory. This study examines how disease trajectory is associated with access to palliative care services in multiple settings, including time of initiation before death and intensity and type of service use. We also compare our data to other geographic areas, namely the USA, UK and Western Australia. Our hypothesis is that compared with the USA, Ontarians will initiate palliative care services earlier, across all disease trajectories and compared with UK and Western Australia, access will be similar across all disease trajectories.

METHODS

We conducted a retrospective cohort study of Ontario decedents who died between 1 April 2010 and 31 December 2012. We used linked administrative health databases, held at the Institute for Clinical Evaluative Sciences (ICES), to identify palliative care services used across multiple health sectors in the 12 months before death. We used a previously derived comprehensive list of palliative care billing codes to capture palliative care services provided by physicians, nurses and personal support workers in multiple sectors from multiple administrative databases.^{25 26} The databases included: Physician claims database, which captured palliative care services billed by physicians in both community and hospital settings; Home Care Database and the interRAI databases captured publicly funded home care services, such as nursing or personal support care, with palliative care intent; Discharge Abstract Database and the National Ambulatory Care Reporting System captured hospitalisations and emergency department (ED) visits, respectively, where palliative care was the main reason for admission or consulted; Continuing Care Reporting System captured palliative care provided in long-term care and complex continuing care settings. We also linked with the Vital Statistics database for date of death,

sex, age and postal code and Statistics Canada Census data for income quintile and rurality via postal codes.²⁷

We further categorised decedents by the major trajectories of functional decline at end of life, defined by main cause of death as per prior research,^{7 8 28} which have also been validated in Canada.^{29 30} Using International Classification of Diseases 10th Revision codes from the death certificate as defined previously,²⁹ decedents were classified into these trajectories: terminal illness (eg, cancer), organ failure (eg, chronic heart failure), frailty (eg, Alzheimers), sudden death (eg, accident) and other (see online supplementary appendix 1 for main causes of death).

Outcomes of interest

The primary outcome of interest was whether a decedent received palliative care at least once in the last 12 months of life. We further categorised palliative care services delivered in 'any institutional care' setting (ie, hospital inpatient, complex continuing care (analogous to subacute care), long-term care and ED) and 'any community care' settings (ie, outpatient care, home care and home-based physician billing). If both a home care and a physician home visit occur on the same day, they count as a separate home care day and separate physician home visit in subcategory analysis. However, both care events count as a single community care day in 'any community care' so as not to double count for community care that happen on the same day and count more care days than calendar days. The same definition applies to 'any institutional care'. In an acute hospital setting, palliative care days were counted for the entire duration of stay when the most responsible diagnosis for the hospital stay was palliative, palliative medicine was a service provider or a palliative service was provided. For all remaining palliative acute hospital encounters only a single day of the hospitalisation was counted (eg, patient had a postadmission palliative diagnosis). In the community-based settings of care, a palliative care day must have a record of a palliative care service in billing codes; we did not assume that care following the initiation of a palliative care code had a palliative intent in the community settings.

We also examined timing to initiation of palliative care, defined as first instance of any palliative care service captured in the last year of life. If a decedent had the first palliative care service outside of the window, initiation was represented as 365 days. We also examined intensity of palliative care by totalling the number of days palliative care was delivered, categorised by service type.

Statistical analysis

Descriptive mean and median statistics describe the usage patterns of decedents as well as the trajectory of care in the last year of life. Multivariate logistic regression was used to predict the likelihood of any use of palliative care. A negative binomial regression was used to predict the number of days of palliative care that a decedent would receive in the last year of life. Covariates included in the models include: sex, age, income quintile, rurality^{27 31} and number

of chronic conditions. The number of chronic conditions is derived using a combination of validated Institute for Clinical Evaluative Sciences (ICES) algorithms that use prior hospital and physician claims records to identify the disease and hospital and physician claims records in the prior 2 years before death. Ethics approval for this study was received from the Ottawa Hospital Research Institute Ethics Board in Ottawa, Canada.

RESULTS

During the study period, we identified 235 159 decedents, who used a total of 4 497 685 days of palliative care services in the last year of life (mean 19.1 days per decedent). Our

cohort was categorised into end-of-life trajectories: 32% as terminal illness, 31% organ failure, 29% frailty, 5% other and 3% as sudden death (table 1). Decedent characteristics were similar across all the trajectories, with the exception of frailty which had more older women and sudden death which had younger decedents with fewer comorbidities. Men and women were equally represented, and 80% were aged 65 years or older. Seventy-nine per cent of the cohort had three or more comorbidities, where hypertension was the most prevalent, followed by osteoarthritis, cancer, diabetes and congestive heart failure. Remaining results will focus on the three major disease trajectories: terminal illness, organ failure and frailty.

Table 1 Cohort demographics by end-of-life disease trajectory

	Terminal illness		Organ failure		Frailty		Other		Sudden death		Overall	
	N	%	N	%	N	%	N	%	N	%	N	%
Total cohort*	75657	32	72363	31	67513	29	11784	5	7842	3	235159	100
Sex												
Male	39125	52	34371	48	30703	45	5295	45	4987	64	114481	49
Female	36532	48	37992	53	36810	55	6489	55	2855	36	120678	51
Age												
<19	172	<1	691	1	47	<1	827	7	435	6	217	1
19–44	1886	2	1601	2	479	1	332	3	2636	34	6934	3
45–54	5454	7	3247	4	1738	3	442	4	1547	20	1242	5
55–64	12311	16	6631	9	4193	6	730	6	1090	14	24955	11
65–74	18042	24	10885	15	7472	11	1229	10	676	9	38304	16
75–84	22790	30	21447	30	18990	28	2959	25	780	10	66966	28
85–94	13730	18	23514	32	27641	41	4257	36	592	8	69734	30
95+	1272	2	4347	6	6953	10	1008	9	86	1	1366	6
Income†												
Lowest	16014	21	17288	24	15637	23	2545	22	2008	26	53492	23
Low	15931	21	15344	21	13634	20	2317	20	1626	21	48852	21
Middle	14698	19	13727	19	13059	19	2086	18	1474	19	45044	19
High	14621	19	13074	18	12884	19	2063	18	1358	17	44000	19
Highest	13996	19	12136	17	11850	18	1967	17	1258	16	41207	18
Rurality†												
Urban	64302	85	61171	85	57853	86	9752	83	6564	84	199642	85
Rural	1123	15	1074	15	9558	14	1286	11	1211	15	34027	14
No. of Chronic Diseases												
0	348	<1	2049	3	1649	2	1166	10	1791	23	7003	3
1	6496	9	3732	5	3674	5	672	6	1891	24	16465	7
2	11388	15	6463	9	7144	11	1150	10	1358	17	27503	12
3	14846	20	9543	13	9710	14	1559	13	1022	13	36680	16
4	14238	19	11296	16	11059	16	1815	15	674	9	39082	17
5	11260	15	11772	16	10730	16	1740	15	457	6	35959	15
6+	17081	23	27508	38	23547	35	3682	31	649	8	72467	31

*Percentages of 'total cohort' row represent the proportion of the whole cohort. All other percentages in each descriptive category are representative of the proportion of patients in each category under each trajectory and are not summative across a whole row.

†Does not equal 100%: a small number of records are missing this information.

Palliative care access

Among the full cohort, 54% received at least one palliative care service in the last year of life. Palliative care from an institutional and community setting was mainly delivered by hospital inpatient services (46% of overall cohort) and community outpatient services (25%), respectively. Palliative care physician home visits were delivered to 6% of the overall decedent cohort. However, there was wide variation in use of palliative care across end-of-life trajectories (table 2). Across all settings, 88% of those in the terminal illness trajectory received palliative care compared with 44% of the organ failure trajectory and 32% in the frailty trajectory. Within particular settings, the terminal illness trajectory had nearly twice as many decedents receiving palliative care services in the hospital inpatient setting (76%) than the other trajectories. Many terminal illness decedents received outpatient palliative care (53%) and end-of-life home care services (47%), which was four and eight times more, respectively, than in the other two trajectories. Palliative care physician home visits were delivered to 15% of terminal illness decedents compared with 3% of organ failure decedents and 2% of frailty decedents.

Intensity of palliative care

Among users of palliative care in any setting, terminal illness has the highest mean number of palliative care days,

ranging from 17 in an institution and 32 in the community, compared with 12 and 11 for organ failure and 11 and 10 for frailty trajectories. In all trajectories, about half of all palliative care days used occurred in the last 2 months of life, with a twofold increase in the last month of life. For example, decedents in the terminal illness trajectory averaged eight palliative care days in the second to last month before death, which increased to 13 days in the final month of life.

Initiation of palliative care

Decedents in the terminal illness trajectory had palliative care initiated a median of 107 days before death, more than four times earlier than organ failure (median 22 days) and frailty (median 24 days). In terms of intensity, the terminal illness trajectory had palliative care on 37% of days after initiation versus 25% and 23% in organ failure and frailty decedents (table 3).

Multivariable analyses of odds of using any palliative care services

When examining the odds of using any palliative care services in the last year of life, decedents with a terminal illness trajectory have an OR of 17.0 (OR 95% CI 17.03 to 17.09) when compared with those with a frailty trajectory controlling for sex, age, income quintile, rurality and

Table 2 Use (≥ 1 encounters) of palliative care by end-of-life trajectory and sector in the last year of life

Sector and setting of palliative care	End-of-life trajectory			
	Terminal illness (N=75 657)	Organ failure (N=72 363)	Frailty (N=67 513)	Overall* (N=235 159)
Any palliative care in any setting (%)	88.0	44.4	32.4	53.6
Palliative care in an institutional care setting (%)				
Any institutional care†	76.4	39.9	26.1	46.5
Hospital inpatient	75.6	39.4	25.2	45.9
Complex continuing care	6.0	1.4	1.1	2.7
Long-term care	0.4	0.4	0.9	0.5
Emergency room	0.2	<0.1	<0.1	0.1
Palliative care in a community care setting (%)				
Any community care†	68.6	17.2	15.1	32.4
Outpatient	52.7	12.4	11.9	24.8
Home care	46.8	6.0	3.4	18.0
Physician home visits	14.8	2.5	1.9	6.2
Among users of palliative care				
Mean days of institutional care	16.54	12.02	10.71	14.10
Mean days of community care	32.08	10.74	9.68	21.59
Initiation and intensity				
Median number of days before death to palliative care initiation (IQR)	107 (33, 246)	22 (6, 124)	24 (6, 132)	59 (13, 200)
Proportion of days following initiation in which palliative care was recorded (IQR)	37% (0.18–0.67)	25% (0.1–0.7)	23% (0.1–0.64)	33% (0.14–0.67)

*Overall includes the sudden death (3%) and other (5%) trajectories which account for 8% of the total cohort. These are not individually shown here.

†Multiple services received on the same calendar day are counted as a single unit of 'Any community care' or 'Any institutional care'. This avoids double counting palliative care in a single day and prevents decedents from having more service days than total days.

Table 3 Predictive models for the use of palliative care

Exposure	Use of palliative care (Yes/No)*		Number of palliative care days†	
	OR (95% CI)		Incidence rate ratio (95% CI)	
Trajectory				
Terminal illness	17.06	(17.03 to 17.09)	6.94	(6.91 to 6.97)
Organ failure	1.70	(1.68 to 1.72)	1.56	(1.54 to 1.58)
Frailty	Ref.		Ref.	
Other	1.60	(1.56 to 1.64)	0.97	(0.93 to 1.01)
Sudden death	0.35	(0.27 to 0.43)	0.22	(0.16 to 0.28)
Sex				
Female	1.06	(1.04 to 1.08)	1.08	(1.06 to 1.10)
Male	Ref.		Ref.	
Age				
<19	0.72	(0.64 to 0.80)	0.78	(0.72 to 0.84)
19–45	0.89	(0.84 to 0.94)	0.98	(0.93 to 1.03)
45–54	Ref.		Ref.	
55–64	1.08	(1.04 to 1.12)	0.97	(0.93 to 1.01)
65–74	1.17	(1.13 to 1.21)	0.95	(0.92 to 0.98)
75–84	1.16	(1.12 to 1.20)	0.90	(0.86 to 0.94)
85–94	1.00	(0.84 to 1.16)	1.67	(1.55 to 1.79)
≥95	1.10	(1.05 to 1.15)	0.91	(0.86 to 0.96)
Income quintiles				
Q1	Ref.		Ref.	
Q2	1.05	(1.02 to 1.08)	1.09	(1.06 to 1.12)
Q3	1.01	(0.98 to 1.04)	1.08	(1.05 to 1.11)
Q4	1.07	(1.04 to 1.10)	1.10	(1.07 to 1.13)
Q5	1.09	(1.06 to 1.12)	1.19	(1.16 to 1.22)
Rurality				
Rural	Ref.		Ref.	
Urban	1.28	(1.25 to 1.31)	1.23	(1.2 to 1.26)
No. of comorbidities				
0	Ref.		Ref.	
1	3.27	(3.18 to 3.36)	2.82	(2.75 to 2.89)
2	3.74	(3.65 to 3.83)	3.13	(3.06 to 3.20)
3	4.12	(4.03 to 4.21)	3.43	(3.36 to 3.50)
4	4.53	(4.44 to 4.62)	3.69	(3.62 to 3.76)
5	4.75	(4.66 to 4.84)	3.97	(3.90 to 4.04)
≥6	5.40	(5.31 to 5.49)	4.83	(4.76 to 4.90)

*Multivariable logistic regression was used to determine OR.

†Negative binomial regression was used to determine incidence rate ratio.

number of comorbidities (table 3). Decedents in the organ failure trajectory are nearly twice (OR 1.7, 95% CI 1.68 to 1.72) as likely to use any palliative care compared with frailty trajectory.

Multivariable analyses of number of palliative care days received

Negative binomial regression analysis shows that decedents in the terminal illness trajectory receive seven times

more days of palliative care (incidence rate ratio: 6.94, 95% CI 6.91 to 6.97) in the last year of life than decedents with a frailty trajectory. Increasing comorbidity was associated with higher number of days of palliative care received in the last year of life.

Comparison to palliative care access in other countries

In our cohort, among those who received any palliative care services, 55% died from terminal illness, 27%

from organ failure and 18% from frailty illness trajectories. Whereas among those who received the Medicare Hospice Benefit in the USA, 27% had cancer, 17% had dementia and 30% had cardiac, circulatory or respiratory failure³² (table 4). Data from Western Australia shows 69% of patients with cancer and 14% of patients without had access to specialist palliative care services³³ (compared with 88% of cancer and 39% non-cancer in Ontario, Canada). In UK, among palliative care inpatient admissions, 88% had cancer.³⁴

Length of stay also varies by country. In Ontario, UK and Western Australia, patients with cancer had longer median lengths of stays (range 37–107 days) than other disease trajectories (range 6–43 days).^{33 34} However in the USA, the trend is the opposite, with patients with dementia having the longest median lengths of stay (56 days) and patients with cancer have the shortest (19 days).³²

DISCUSSION

Our population-based analysis of decedents in Ontario, Canada shows that while nearly half of decedents receive at least one palliative care service, there are large disparities based on dying trajectory. Eighty-eight per cent of those dying in the terminal illness trajectory (predominantly cancer deaths) received palliative care services compared with organ failure (44%) or frailty trajectories (32%). The terminal illness group also received twice as many palliative care services and four times earlier than the other two trajectories. In our universal insured hospital and physician system that does not require patients to forego curative treatment to receive palliative care, the median time from first palliative care service to death is 107 days for terminal illness, 22 days for organ failure and 24 days for frailty trajectories.

Our hypotheses were incorrect. While our Canadian data demonstrated terminal illness (predominantly cancer) patients received palliative care much earlier before death than in the USA, patients without cancer in Ontario were identified closer to death than in the USA. Importantly, the type of palliative care services offered, the training of providers and the organisation of the delivery system are not equivalent between countries or within Canada.³⁵ Nonetheless, comparing similar statistics between geographic areas can generate hypotheses on how different eligibility criteria and health systems may explain differences in results. For instance, the in-home visiting hospice insured services offered in the USA includes extensive teams of specialist physicians and nurses and interprofessional providers, which is more comprehensive and coordinated than the services offered across Ontario, Canada.³⁶ Indeed, our results show the vast majority of palliative care services were delivered in hospital inpatient units, not the home as in the USA. Yet the requirement to forego curative treatment to receive hospice care in the USA may be

a factor in its relatively late initiation for patients with cancer, particularly with advancements in cancer treatment. Conversely, the comprehensive home-based focus of the US hospice insured benefit may explain the higher proportion of patients without cancer using it and for longer compared with Ontario, Canada which does not have widespread access to home-based fully insured palliative care teams.

Our data are also interesting compared with UK (universal health system) and Western Australia (mix of public and private health systems), which also have no requirements for an expected death certification or to forego curative treatments. Despite this similarity in eligibility, access to palliative care, use by disease trajectories, initiation before death and intensity and type of service use differ. The physician ratio is lower in Ontario, Canada than the other countries. The UK and USA have more physician specialists (75%) to generalists (25%) (all specialties) compared with Western Australia and Ontario, which is half-half.³⁷ The availability of human resources and their training likely affects palliative care access and the delivery model (ie, specialist or generalist driven). For instance, in Ontario, one study showed that there were only 276 of 9732 family physicians, where palliative care services comprised more than 10% of their billings (40% of the cohort billed no palliative care at all).³⁸ Indeed receipt of physician home-based visits for palliative care was very low across all disease trajectories in our data, which may be related to inadequate billing fees for home visits.³⁹ The limited availability of palliative care physician specialists may explain preferential access to patients with terminal illness, who may traditionally be easier to identify as needing palliative care. Considering the growing body of evidence of efficacious palliative care interventions for non-cancer diseases,^{17–20 40} the marked disparities in access to patients without cancer ought to be a policy priority and will likely require overcoming the stigma of imminent death and medical failure as well as education on the benefits of early integration.^{41 42}

Limitations of using administrative health data to capture the use of palliative care include the potential undercoding of palliative care delivered, particularly in the community and long-term care.²⁵ In the community, despite financial incentives to use specialised billing codes for palliative care, physicians may provide care reflecting palliative intent or elements of a palliative approach but not bill as such. This may include discussions about coping, basic symptom management and so on. In long-term care, palliative care billing codes are uncommon, rather, monthly management codes and subsequent visit codes are used.^{25 43 44} There are potential issues with reliability and validity when using cause of death data to group decedents into disease trajectories, particularly with the non-terminal illness trajectories. For example, not all stroke recovery follow the trajectory pattern of organ failure. We cannot describe the quality of care or include services provided by volunteers, family members

Table 4 Comparison of PC access and initiation across geographic areas

	Ontario	UK	USA	Western Australia
Criteria to access PC	<ul style="list-style-type: none"> ▶ 94 000 deaths in Ontario 2014/2015 ▶ Universal insured hospital and physician system ▶ No restrictions on curative along with PC ▶ No written document required to initiate PC, though often the 'surprise question' of expected death of 1 year to 6 months is used to initiate care⁴⁷ ▶ Provided by general practitioners, specialists and home-care providers 	<ul style="list-style-type: none"> ▶ 548 000 deaths 2015 ▶ Primary care delivered heavily by general practitioners and primary care trusts ▶ Universal health insurance ▶ Patients may be terminal (expected to die within 12 months, have a life-limiting illness or chronic condition with a trajectory that has a sharp functional decline or extensive acute episodes, or require extended care) ▶ Can mix palliative and curative care³⁴ 	<ul style="list-style-type: none"> ▶ 2.6 M deaths in 2015 ▶ Hospice benefit includes visiting interprofessional providers in home, residential hospices, hospitals, long-term care, etc ▶ Available to Medicare patients ▶ Must have signed physician note stating expected death within 6 months ▶ Must waive access to curative treatments to access hospice benefit³² 	<ul style="list-style-type: none"> ▶ 23852 deaths in Western Australia in 2009/2010 ▶ Mix of private and government service providers ▶ Use 'normative need' to assess access to PC specialists³³
Physician ratio	<ul style="list-style-type: none"> ▶ 2.2 physicians/1000 ppl (2015) ▶ 47%/53%: generalists/specialists³⁷ 	<ul style="list-style-type: none"> ▶ 2.8 physicians/1000 ppl (2015) ▶ 29%/71%: generalists/specialists³⁷ 	<ul style="list-style-type: none"> ▶ 2.5 physicians/1000 ppl (2011) ▶ 12%/88%: generalists/specialists³⁷ 	<ul style="list-style-type: none"> ▶ 3.5 physicians/1000 ppl (2015) ▶ 45%/47%: generalists/specialists (8%: medical doctors not further defined)³⁷
Per cent that get any service	<ul style="list-style-type: none"> ▶ 54% of decedents between 2010 and 2012 received at least PC services (from billing claims) in any setting.(table 2) 	<ul style="list-style-type: none"> ▶ 74% of people who are in need of PC receive either specialist or generalist services ▶ 18% of non-malignant access to PC was for chronic respiratory illness, 11% for heart failure³⁴ 	<ul style="list-style-type: none"> ▶ 46% of Medicare (>65 years old) decedents received ≥1 day of hospice care (via the Medicare hospice benefit) in 2015³² 	<ul style="list-style-type: none"> ▶ 46% of decedents received any PC³³
Cancer and non-cancer access	<ul style="list-style-type: none"> ▶ 88% of terminal illness, 44% of organ failure and 32% of frailty decedents (or 39% non-cancer) received any PC services (table 2) ▶ Among those receiving any PC services, 55% died from terminal illness, 27% from organ failure and 18% from frailty illness trajectories 	<ul style="list-style-type: none"> ▶ 88% of PC inpatients have cancer diagnosis ▶ 20% of inpatient referrals are for non-cancer³⁴ 	<ul style="list-style-type: none"> ▶ Among those who received the hospice benefit, the principal diagnoses were: 27% cancer, 19% cardiac, 17% dementia and 11% respiratory³² 	<ul style="list-style-type: none"> ▶ 69% of patients with cancer had access to specialist care ▶ 14% of patients without cancer had access to specialists³³
Average Length of stay in PC	<ul style="list-style-type: none"> ▶ Median days of initiation of service to death: ▶ Terminal illness 107 days ▶ Organ failure 22 days ▶ Frailty 24 days (table 2) 	<ul style="list-style-type: none"> ▶ Median days on service in one large study in one region (Leeds, UK): ▶ 37 days for cancer ▶ 16 days for non-cancer⁴⁸ 	<ul style="list-style-type: none"> ▶ Mean/(median) days on service ▶ Cancer: 47/(19) days ▶ cardiac: 76/(28) days ▶ dementia: 105/(56) days ▶ respiratory 69/(19) days ▶ stroke 77/(20) days³² 	<ul style="list-style-type: none"> ▶ Median number of days receiving specialist PC was 30 (cancer), 8 (COPD) and 5 (Alzheimers and heart failure)³³ ▶ Median days PC initiated before death: 62 (cancer), 6 (Alzheimers) and 43 (COPD)³³
Location of service (community, home, hospital)	<ul style="list-style-type: none"> ▶ 68% of cancer decedents have PC in a community setting ▶ 76% in an acute care setting ▶ <1% of PC for any trajectory was received in an LTC facility (table 2) 	<ul style="list-style-type: none"> ▶ ~20% of LTC residents were seen by a PC specialist nurse, 96% were seen by a GP ▶ Poor access to hospitals. Only 21% of hospitals provide face-to-face PC 24/7 ▶ 27% of hospital outpatient PC and 17% of community PC provided to non-malignant disease³⁴ 	<ul style="list-style-type: none"> ▶ Home 56.0% ▶ Nursing facility 41.3% ▶ Hospice inpatient facility 1.3% ▶ Acute care hospital 0.5% ▶ Other 0.9%³² 	<ul style="list-style-type: none"> ▶ Organ failure patients (eg, liver failure) tended to receive care in hospital over community settings ▶ Motor neuron and cancer decedents had increased access to community services³³

COPD, chronic obstructive pulmonary disease; GP, general practitioner; LTC, long-term care; PC, palliative care; ppl, people.

or private care that is not recorded in the health administrative databases. We also do not have an administrative database for hospice services and cannot account for care provided in a residential hospice. However, only 1%–3% of deaths occur in a residential hospice, and the majority of hospice care occurs after initiation of palliative home care services—which is included in our study.

In conclusion, our study quantifies a large disparity in access to palliative care for those dying from organ failure and frailty trajectories. Decedents with a terminal illness trajectory, exemplified by a cancer diagnosis, are significantly more likely to receive palliative care services than the other dying trajectories; they receive more services (intensity) both in hospital and community, and these services are initiated earlier in the dying trajectory. All trajectories could benefit from increased access to palliative home care services and physician home visits. This data will be useful to compare to in the future since a national palliative care framework was an identified need⁴⁵ and has recently been passed into law.⁴⁶ These data also serve as a useful comparison for other countries with similar and different health-care systems and eligibility criteria to explore palliative care access across disease trajectories.

Contributors HS and PT conceptualised the study and RP performed the data abstraction and analysis. All authors contributed to interpretation and critical analysis of results. EO and HS drafted the first manuscript and all authors were responsible for contributing to the critical content and review of the manuscript. All authors act as guarantors and affirm that the article is an honest, accurate and transparent account of the study being reported; that no important aspects of the study have been omitted and that any discrepancies from this study as planned have been explained. All authors take responsibility for the integrity of the data and the accuracy of the data analysis.

Funding The author(s) disclosed receipt of the following financial support for the research, authorship and/or publication of this article. This research was supported by a research grant from the Ontario MOHLTC to the Health System Performance Research Network (HSPRN). This study was also supported by the ICES, which is funded by an annual grant from the Ontario MOHLTC. The opinions, results and conclusions reported in this article are those of the authors and are independent from the funding sources. No endorsement by MOHLTC is intended or should be inferred.

Disclaimer The funders had no influence in study design; in the collection, analysis and interpretation of data; in the writing of the report and in the decision to submit the article for publication.

Competing interests None declared.

Patient consent Not required.

Ethics approval This study has been approved by the research ethics board at the ICES, at Sunnybrook Health Sciences Centre in Toronto, ON, Canada and by the research ethics board at the Ottawa Hospital Research Institute at Ottawa, ON Canada.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement Using encrypted health card numbers as unique identifiers, records of healthcare use and costs were linked across various administrative databases. No written consent was obtained; all data were encrypted using health card numbers as unique identifiers. Thus, all records used were deidentified and anonymised. All data were housed and analyses at ICES, a prescribed entity for the purposes of section 45 of Ontario's Personal Health Information Privacy Act.

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