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2 The primary care and other health system use of home care patients: a retrospective cohort analysis

3 Authors

- 4 Aaron Jones, MSc¹
- 5 Susan E. Bronskill, PhD^{2,3}
- 6 Gina Agarwal, MBBS, PhD^{1,4}
- 7 Hsien Seow, PhD^{1,5}
- 8 David Feeny, PhD⁶
- 9 Andrew P. Costa, PhD^{1,7}

Author Affiliations

- 11 ¹Department of Health Research Methods, Evidence, and Impact, McMaster University,
- 12 Hamilton, Ontario, Canada.
- 13 ²ICES, Toronto, Ontario, Canada.
- ³Institute of Health Policy, Management & Evaluation, Dalla Lana School of Public Health, University of
- 15 Toronto
- ⁴Department of Family Medicine, McMaster University, Hamilton, Ontario, Canada.
- ⁵Department of Oncology, McMaster University, Hamilton, Ontario, Canada.
- 18 ⁶Department of Economics, McMaster University, Hamilton, Ontario, Canada.
- ⁷Department of Medicine, McMaster University, Hamilton, Ontario, Canada.

Corresponding Author

- 22 Aaron Jones, MSc
- 23 PhD Candidate
- 24 Department of Health Research Methods, Evidence, and Impact
- 25 McMaster University
- 26 1280 Main St. W., Hamilton, L8S 4K1
- e: jonesa13@mcmaster.ca
- 28 p: 416-912-9276

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33 Conflicts of interest

34 None to report.

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Data sharing statement

The dataset from this study is held securely in coded form at ICES. While data sharing agreements prohibit ICES from making the dataset publicly available, access may be granted to those who meet prespecified criteria for confidential access, available at www.ices.on.ca/DAS. The full dataset creation plan and underlying analytic code are available from the authors upon request, understanding that the computer programs may rely upon coding templates or macros that are unique to ICES and are therefore either inaccessible or may require modification.

Author contributions

Aaron Jones and Andrew Costa and conceived the study and developed the design in consultation with
the authors. Aaron Jones completed the analysis and drafted the initial manuscript. All authors
contributed to the interpretation of data, the critical revision of the manuscript for important
intellectual content, and approved the final version submitted for publication.

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<u>Abstract</u>

Background

- Robust and integrated primary care and home care are core components of effective chronic disease management in the community. However, primary care use by home care patients is not well studied.
- We examined the primary care and other health system use of a cohort of home care patients.

Methods

We conducted a population-based retrospective cohort study of publicly-funded home care patients in Ontario, Canada from October 2014 to September 2016. Primary outcomes were patterns of primary care physician visits including coordination with home care, home visits, and afterhours/weekend visits within six months of a home care assessment. Secondary outcomes included specialist physician visits, emergency department use, home care visits, and long-term care home placement. Multivariable models examined associations between patient characteristics and subsequent patterns of primary care use.

Results

Our cohort identified 226,054 home care patients with a median age of 81. Following assessment, home care patients visited primary care physicians at a rate of 0.78 visits per month. Physician-based home care coordination codes were billed for 3.9% of patients. Primary care home visits were received by 13.1% of patients, and 15.1% of patients utilized afterhours/weekend primary care.

Interpretation

Publicly-funded home care patients frequently visited a primary care physician but billing claims for coordination between primary care and home care were infrequent. Physician home visits were more

likely to be received by the oldest and most functionally impaired patients, suggesting that home visits are responsive to the needs of home care patients.

Background

Chronic disease management models frequently emphasize the importance of quality primary care for effective chronic disease management in the community. 1-3 Coordinated care can reduce depressive symptoms and improve the functional status of older adults with multimorbidity. 4 Home-based primary care has been shown to reduce emergency department (ED) visits and hospitalizations in homebound older adults. 5 Access to timely primary care and afterhours primary care could reduce ED visits. 6,7 However, research suggests that older adults with complex care needs frequently experience fragmentation of care and difficultly accessing primary care.

Home care patients are a population of complex community-dwelling older adults characterized by multiple chronic conditions, need for support in activities of daily living, and a high risk of adverse outcomes.^{9,10} Aging strategies have frequently called for robust and responsive primary care and home care to enable seniors to live well in the community as long as possible^{11,12}. However, the patterns of primary care use by home care patients have not been well studied.

The objective of this study was to describe the primary care physician use of publicly-funded home care patients in Ontario, including coordination between home care and primary care and advanced access to primary care such as home visits and afterhours/weekend care. We examined associations between patient characteristics and primary care use and also described the use of other health sectors to contextualize our findings.

Methods

Study Design and Data Sources

This study identified a population-based, retrospective cohort of adults in Ontario, Canada who received a comprehensive home care assessment and used multiple health administrative databases to identify patterns of health system use following the assessment. A description of all data sources can be found in Appendix 1. These datasets were linked using unique encoded identifiers and analyzed at ICES. This study was a granted an exemption of formal ethics review from the Hamilton Integrated Research Ethics Board.

Study Cohort

All publicly-funded home care patients in Ontario who are receiving on-going care are periodically assessed with the Resident Assessment Instrument for Home Care (RAI-HC), a comprehensive clinical assessment¹³. We selected all RAI-HC assessments of adult (19+), home care patients completed in Ontario between October 1, 2014 and September 30, 2016. If an individual was assessed more than once during the period, their most recent assessment was selected. The assessment date was considered the index date for follow-up. Patients receiving palliative home care at baseline were excluded from the cohort as their health utilization patterns and outcomes vary greatly from other home care patients.

Baseline Characteristics

Patient characteristics were identified from the baseline assessment and included demographic, health and functional characteristics, frailty¹⁰, and health-related quality of life¹⁴. We identified patients with three key conditions known to be primary drivers of home care: congestive health failure, chronic pulmonary obstructive disease (COPD), and dementia. Similar to other studies we also classified each patient's primary care enrollment models at baseline. The three main model types are: a) Family

Health Teams, which are team-based, interprofessional primary health care organizations funded primarily through capitation payments; b) Other blended capitation models that are funded similarly but lack the explicit interprofessional approach; and c) Enhanced fee-for-service models which are funded primarily through billing claims. A few rural and specialty models were grouped together in an "Other" category and patients not rostered with a physician were considered a distinct category.

Primary care use among home care patients

We linked the index assessment records to other health administrative databases to identify health service use within six months (182 days) of the assessment date. A six-month follow-up was chosen as it aligns with the standard RAI-HC assessment interval and at least three-quarters of patients can be expected to experience a meaningful clinical change within six months¹⁹. Primary care physician (PCP) visits were defined as office, home, or phone-based services provided by a general practice/family practice physician or community medicine physician with a maximum of one visit per patient per physician per day. We identified primary care coordination with home care using billing codes specific to PCP supervision of home care or participation of a PCP in a case conference concerning a home care patient. For measures of advanced access, we identified PCP visits to a patient's home and PCP visits that occurred afterhours or on a weekend or holiday. Details of the calculation of each physician measure can be found in Appendix 2.

Other health system use among home care patients

To contextualize the primary care use, we also measured other health sector use of patients and their transitions between care settings. Specialist physician visits were defined similarly to primary care and included all physicians other than general practice/family practice, community medicine, and pediatrics. Home care use was measured as hours of personal support and number of home nursing visits. Other measures included: unplanned ED visits, unplanned acute hospital admissions, long-term care home

admission, and death. We also tracked the care setting of the patient (community, hospital, long-term care, dead) across the six-month follow-up period and calculated the total number of transitions in care settings.

Descriptive analysis

We reported the proportions of patients with any PCP visit, PCP coordination with home care, PCP home visit, and PCP afterhours/weekend visit, and the rate of PCP visits per month. We also reported the proportion of patients with any specialist physician visit, the average number of specialties seen and the rate of specialist physician visits per month. For home care we reported the proportions of patients who received or were authorized at baseline for personal support and home nursing as well as the rate of visits/hours per month among patients with the service. Other measures reported included the proportions of patients with an unscheduled ED visit, acute hospital admission, and long-term care home admission, the rate of ED visits per month, the average number of transitions of care settings, and the proportion of patients who died in the follow-up window.

All rates were based on the number of days during follow-up that the patient spent in the community, i.e. not dead, in a long-term care home, or in hospital. Home care use rates were additionally restricted to exclude days after home care services were discharged when applicable. Monthly rates were produced by multiplying the daily rate by 30. Additionally, the proportion of patients who received a PCP home visit was reported by functional impairment strata and we stratified the proportion of patients with PCP coordination with home care by Ontario's 14 health regions to explore potential variation in the rates based on regional initiatives to promote coordination. All descriptive measures were reported both for the entire cohort as well as the important subpopulations with congestive heart failure, COPD, and dementia.

Multivariable analysis

We fit multivariable regression models to examine associations between patient characteristics and the primary care use measures. The rate of PCP visits was fit with a quasi-poisson generalized linear model²⁰ with an offset term for days spent in the community. The proportions of PCP coordination with home care, PCP home visit, and a PCP afterhours/weekend visit were separately fit with logistic regression models. Each model included the following independent variables: sex, age, region, rurality, patient enrollment model type, home care services received or authorized at baseline, functional impairment, cognitive impairment, mood symptoms, comorbid conditions, and number of concurrent medications. Results were reported as rate ratios or odds ratios with 95% confidence intervals. All analyses were performed using SAS 9.4.

Results

Our cohort identified 226,054 adult home care patients with an assessment between October 1, 2014 and September 30, 2016. The median age of patients in the cohort was 81 years and just under two-thirds (63%) were female (Table 1). Over 40% of patients needed at least limited assistance with personal hygiene, locomotion, eating, or toileting, and over 60% had at least a mild cognitive impairment. Around 13% of patients had a diagnosis of congestive heart failure at baseline, 20% had COPD, and 25% had a diagnosis of dementia. Roughly 30% of patients were enrolled in each of the three broad types of primary care patient enrollment models at baseline. Transitions between care settings across the follow-up period were common (Figure 1). At the end of six months, 71% of the patients were still living in the community, 13% were in long-term care homes, 4% were in hospitals, and 13% had died.

Table 1: Baseline characteristics of adult home care patients, Ontario, September 2014 to October 2016

<Insert Table 1 here >

Figure 1: Transitions between care settings, adult home care patients, Ontario, September 2014 to

196 October 2016

<Insert Figure 1 here>

Primary care use among home care patients

The overall PCP visit rate during the follow-up period was 0.78 visits per month (Table 2), with 84% of patients having at least one PCP visit. Billing claims related to PCP coordination with home care occurred in only 3.9% of patients. PCP home visits were received by 13.1% of patients and 15.1% of patients used after hours/weekend primary care. Patients with congestive health failure and COPD had somewhat higher primary care use than patients than the overall population. Among the most impaired patients, just over one-quarter (27.5%) received a PCP home visit (Table 3). Stratifying PCP coordination with home care by health regions revealed significant regional variation in the use of the codes, particularly in the differential between family health teams and other patient enrollment models (Appendix 3).

- **Table 2:** Primary care physician and other health system use, adult home care patients, Ontario,
- 209 September 2014 to October 2016
- 210 <Insert Table 2 here>
- **Table 3**: Proportion of patients with a PCP home visit by functional impairment strata
- 212 <Insert Table 3 here>

Other health system use among home care patients

Around three-quarter of patients received personal support while just over a third received home nursing. Nearly half of patients visited the ED and over a quarter had an unplanned hospital admission across the six-month follow-up. Patients with congestive heart failure or COPD had higher rates of ED

visits, hospitalizations, and death. Patients with dementia had similar ED, and hospital utilization as the overall population, but were significantly more likely to be admitted to a long-term care home.

Multivariable analysis

Increasing age, enrollment in an enhanced fee-for-service model, home care nursing, and having 9 or more prescription medications at baseline was associated with greater primary care use (Table 3). PCP coordination with home care was more common when a patient was enrolled with a family health team, receive or was authorized at baseline for home care nursing, or resided in southern Ontario. Age, severe functional impairment, and home care nursing were strongly associated with an increased likelihood of receiving a PCP home visit. Finally, enrollment in an enhanced fee-for-service model was associated with higher likelihood of an afterhours/weekend primary care visit, while living in eastern or northern Ontario or in a rural location was negatively associated.

Table 4: Multivariable regression models of PCP use

<Insert Table 4 here>

Interpretation

Publicly-funded home care patients with continuing care needs frequently utilized primary care and other health services within six months of assessment. Nearly all of the patients visited a primary care physician at least once, however just under 4% had a billing for home care coordination, 13% had a primary care physician home visit, and 15% used primary care afterhours or a weekend or holiday. Coordination codes were more common in interprofessional primary care practices while afterhours care was more prevalent in non-capitated practices. Physician home visits were more likely to be received by the oldest and most functionally impaired patients.

Comparing primary care use between studies can be imprecise due to differences in methodology and time frames, but a study with similar primary care definitions to ours reported a PCP visit rate of 0.52 visits per month among all older adults in Ontario²¹. As home care patients in our study had a PCP visit rate of 0.78 per month, this suggests home care patients had around 50% higher primary care use than a general older adult population. Looking at other health sectors, comparisons with the same study suggest home care patients had twice the specialist physician use and 3.5 times the ED visits of an older adult population.

The billing codes specific to PCP coordination with home care were rarely used. This could suggest low levels of coordination possibly due to difficulties in communication and lack of integration between home care and primary care²². However, it may also be the result of the coordination codes not being billed due to lack of awareness of the specific codes or a sense that the codes are not worth the effort of claiming. Family health teams were considerably more likely than other enrollment models to bill the coordination codes, which could be due to their interprofessional orientation, or that regional planning organizations specifically target family health teams in initiatives to promote coordination. For example, in some regions home care coordinators have been aligned with specific family health teams and may work out of the same office²³. There was significant variation in the overall frequency of coordination billing between different health planning regions, as well as in the difference between family health teams and other models, suggesting that regional initiatives to promote coordination influenced the rate of billing.

Older patients with severe functional impairments were much more likely to receive a primary care home visit than those without impairments, which suggests physician home visits are responsive to the functional needs of patients. Around one quarter of patients (27.5%) with a severe functional impairment received a PCP home visit, which is slightly higher than the 22% of a palliative patients in

Ontario reported to have received a physician home visit in the last six months of life²⁴. Even among patients with the lowest levels of functional impairment however, almost 10% received a home visit. Historically part of general practice, physician house calls in Canada have become more frequent recently after falling in previous decades partially due to changes in how primary care was funded^{25–28}. Finally, home care patients enrolled in non-capitated models were considerably more likely than those enrolled in capitation-based models to use afterhours or weekend primary care. This effect has been previously noted, including in a study done shortly after the implementation of the first capitation models in Ontario.²⁹ All enrollment models are mandated to make a minimum amount of afterhours care available but differences how these models are funded could be contributing to fewer afterhours visits in the capitated models.

Limitations

Our study has a number of strengths, including having a large, population-based sample and ability to measure health service use across multiple sectors. There are also some important limitations. Our measure of coordination between primary care and home care relies on three specific billing codes. The degree to which this measure underrepresents the true level of coordination cannot be ascertained by this study and would require qualitative, primary data collection. Also, findings around Ontario-specific primary care models or billing codes may have less generalizability.

Conclusion

We found that home care patients with continuing care needs in Ontario, Canada frequently visited a primary care physician. Physician billing claims specific to coordination between primary care and home care were rarely utilized but may underrepresent the true level of coordination. Physician home visits

- were more likely to be received the oldest and most functionally impaired patients, suggesting
 responsiveness to patient needs. These findings provide important data on the primary care use of
 home care patients and can inform future research on how patterns of primary care and home care can
 influence the health outcomes of home care patients.
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Table 1: Baseline characteristics of adult home care patients in Ontario, September 2014 to October 2016

Demographics Age, yrs (Median (Q1-Q3)) 81 (71-88) Sex, female 83978 (62.9) Lived Alone 110137 (48.7) Health Activities of Daily Living (ADL) impairment¹ Independent/Supervision 127725 (56.5) Limited/Extensive 72220 (32.0) Maximal/ Dependent 26109 (11.6) Cognitive impairment² 11081 (53.6) Intact / Borderline intact 85613 (37.9) Mild / Moderate 121081 (53.6) Severe 19360 (8.6) Mood symptoms³ 108918 (48.2) Some symptoms 59684 (26.4) Daily symptoms 59684 (26.4) Bladder incontinence 94535 (41.8) Fall in last 90 days 91962 (40.7) Five or more concurrent medications 189760 (83.9) Congestive heart failure 29875 (13.2) COPD 44209 (19.6) Dementia 58413 (25.8) Frailty Index 80ust (0-0.19) Robust (0-0.19) 46043 (20.4) Pre-frail (0.2-0.29) 68562 (30.3) </th <th>Patient Characteristics</th> <th>No. (%) of patients n=226,054</th>	Patient Characteristics	No. (%) of patients n=226,054
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Acitivites of Daily Living (ADL) impairment¹ Independent/Supervision 127725 (56.5) Limited/Extensive 72220 (32.0) Maximal/ Dependent 26109 (11.6) Cognitive impairment² Intact / Borderline intact 85613 (37.9) Mild / Moderate 121081 (53.6) Severe 19360 (8.6) Mood symptoms³ No symptoms 108918 (48.2) Some symptoms 59684 (26.4) Daily symptoms 57452 (25.4) Bladder incontinence 94535 (41.8) Fall in last 90 days 91962 (40.7) Five or more concurrent medications 189760 (83.9) Congestive heart failure 29875 (13.2) COPD 44209 (19.6) Dementia 58413 (25.8) Frailty Index Robust (0-0.19) 46043 (20.4) Pre-frail (0.2-0.29) 68562 (30.3) Frail (>= 0.3) 111449 (49.3) Health-related quality of life (Median (Q1-Q3)) ⁴ 0.19 (-0.01-0.42) Patient Enrollment Model Type Enhanced fee-for-service 73150 (28.7) Family health team 75031 (32.4) Other capitation 64908 (33.2) Other	Lived Alone	110137 (48.7)
Independent/Supervision Limited/Extensive Naximal/ Dependent Cognitive impairment² Intact / Borderline intact Mild / Moderate Severe 19360 (8.6) Mood symptoms³ No symptoms Some symptoms Daily symptoms Saldder incontinence Fall in last 90 days Five or more concurrent medications COPD 44209 (19.6) Dementia Robust (0-0.19) Pre-frail (0.2-0.29) Frail (>= 0.3) Patient Enrollment Model Type Enhanced fee-for-service Family health team Other capitation Cognitive impairment² 26109 (11.6) Congestive heart failure COPD 46408 (33.2) Congestive heart failure COPD 46043 (20.4) Pre-frail (0.2-0.29) Frail (>= 0.3) Congestive heart failure COPD Congestive heart failure C	Health	
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Cognitive impairment2	Limited/Extensive	72220 (32.0)
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Fall in last 90 days Five or more concurrent medications Congestive heart failure COPD COPD A4209 (19.6) Dementia 58413 (25.8) Frailty Index Robust (0-0.19) Pre-frail (0.2-0.29) Frail (>= 0.3) Health-related quality of life (Median (Q1-Q3)) ⁴ Patient Enrollment Model Type Enhanced fee-for-service Family health team Other capitation Other 91962 (40.7) 189760 (83.9) 29875 (13.2) 44209 (19.6) 58413 (25.8) Frailty Index 46043 (20.4) 68562 (30.3) 111449 (49.3) 0.19 (-0.01-0.42)		
Congestive heart failure COPD 44209 (19.6) Dementia 58413 (25.8) Frailty Index Robust (0-0.19) Pre-frail (0.2-0.29) Frail (>= 0.3) Health-related quality of life (Median (Q1-Q3)) ⁴ Patient Enrollment Model Type Enhanced fee-for-service Family health team 75031 (32.4) Other capitation 64908 (33.2) Other	Fall in last 90 days	
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Dementia 58413 (25.8) Frailty Index 46043 (20.4) Robust (0-0.19) 46043 (20.4) Pre-frail (0.2-0.29) 68562 (30.3) Frail (>= 0.3) 111449 (49.3) Health-related quality of life (Median (Q1-Q3)) ⁴ 0.19 (-0.01-0.42) Patient Enrollment Model Type Enhanced fee-for-service 73150 (28.7) Family health team 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Congestive heart failure	29875 (13.2)
Frailty Index Robust (0-0.19)	COPD	44209 (19.6)
Frailty Index Robust (0-0.19)	Dementia	58413 (25.8)
Pre-frail (0.2-0.29) 68562 (30.3) Frail (>= 0.3) 111449 (49.3) Health-related quality of life (Median (Q1-Q3)) ⁴ 0.19 (-0.01-0.42) Patient Enrollment Model Type 73150 (28.7) Enhanced fee-for-service 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Frailty Index	
Frail (>= 0.3)	Robust (0-0.19)	46043 (20.4)
Health-related quality of life (Median $(Q1-Q3)$) ⁴ 0.19 (-0.01-0.42) Patient Enrollment Model Type Enhanced fee-for-service 73150 (28.7) Family health team 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Pre-frail (0.2-0.29)	68562 (30.3)
Patient Enrollment Model Type Enhanced fee-for-service 73150 (28.7) Family health team 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Frail (>= 0.3)	111449 (49.3)
Enhanced fee-for-service 73150 (28.7) Family health team 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Health-related quality of life (Median (Q1-Q3)) ⁴	0.19 (-0.01-0.42)
Family health team 75031 (32.4) Other capitation 64908 (33.2) Other 8403 (2.0)	Patient Enrollment Model Type	
Other capitation 64908 (33.2) Other 8403 (2.0)	Enhanced fee-for-service	73150 (28.7)
Other 8403 (2.0)	Family health team	75031 (32.4)
,	Other capitation	64908 (33.2)
Not enrolled 4562 (3.7)	Other	8403 (2.0)
100011011011	Not enrolled	4562 (3.7)

- 1 ADL Hierarchy Scale: Includes personal hygiene, locomotion, eating and toileting
- 2 Cognitive Performance Scale
- 3 Depression Rating Scale
- 4 HUI3 scores are based on a scale where 1 represents perfect health and 0 represents dead The range of the values is -0.36 to 1 with scores less than 0 representing states worse than dead

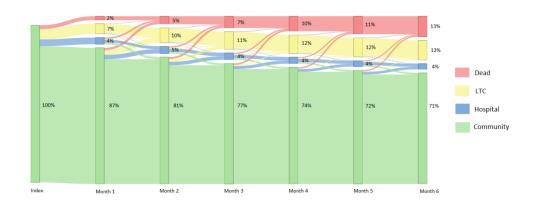


Figure 1: Transitions between care settings, adult home care patients, Ontario, September 2014 to October 2016

Table 2: Primary care physician and other health system use, adult home care patients, Ontario, September 20

		Ch	ronic Conditio	ns
Hardela Hill and a Reserve	All	Heart Failure	COPD	Dementia
Health Utilization Measure	n=226,054	n=29,875	n=44,209	n=58,413
Primary Care				
Any PCP visit (%)	84.4%	85.7%	85.7%	82.0%
Rate of PCP visits per month	0.78	0.94	0.86	0.79
Any PCP home care coordination (%)	3.9%	5.1%	4.7%	3.5%
Any PCP home visit (%)	13.1%	16.3%	13.5%	14.8%
Any PCP afterhours/weekend visit (%)	15.1%	15.2%	15.2%	13.2%
Other Health Sectors				
Any specialist visit (%)	68.1%	70.8%	70.4%	55.5%
Rate of specialist visits per month	0.60	0.70	0.64	0.42
Average count of specialties seen	1.50	1.64	1.62	1.00
Any personal support (%)	71.8%	77.7%	73.8%	78.5%
Any home nursing (%)	35.7%	46.3%	41.5%	24.5%
Rate of personal support hours per month	22.3	23.4	20.4	29.3
Rate of home nursing visits per month	6.8	6.3	6.1	5.1
Any ED visit (%)	46.9%	58.4%	55.0%	46.7%
Any acute care hospitalization (%)	27.2%	40.2%	34.3%	27.7%
LTCH admission (%)	15.7%	15.5%	13.4%	34.1%
Death (%)	13.2%	23.2%	17.3%	14.7%
Rate of ED visits per month	0.17	0.25	0.23	0.15
Transitions of care settings (%) ¹				
0	60.4%	50.0%	56.3%	47.5%
1-2	28.4%	33.7%	29.4%	40.7%
3+	11.2%	16.4%	14.4%	11.8%

¹ Includes transfers between any of the following: community, acute hospital, rehab hospital, mental health hospital, continuing care hospital, or long-term care home

014 to October 2016



Table 3: Proportion of patients with a PCP home visit by functional impairment strata

ADL Impairment	PCP home visit (%)
Independent/Supervision	9.3
Limited/Extensive	14.5
Maximal/ Dependent	27.5



Table 4: Multivariable regression models of PCP use

Variable		PCP visits			ome care
Variable		Rate Ratio	95% CI	Odds Ratio	95% CI
Sex	Female vs. Male	0.97	(0.95, 0.98)	1.01	(0.96, 1.06)
	19-59	(ref)	(ref)	(ref)	(ref)
	60-69	1.02	(0.98, 1.05)	0.94	(0.86, 1.03)
Age	70-79	1.08	(1.05, 1.12)	1.01	(0.93, 1.10)
	80-89	1.15	(1.11, 1.18)	1.02	(0.95, 1.11)
	90+	1.20	(1.16, 1.25)	1.02	(0.93, 1.12)
	Central	(ref)	(ref)	(ref)	(ref)
Region ¹	East	0.96	(0.94, 0.99)	0.95	(0.89, 1.01)
region	North	0.84	(0.81, 0.87)	1.11	(1.02, 1.20)
	South	1.07	(1.05, 1.09)	1.41	(1.34, 1.48)
Rurality ²	Rural vs. Urban	0.96	(0.94, 0.99)	1.06	(1.00, 1.13)
	EFS	(ref)	(ref)	(ref)	(ref)
Dationt on allocant	FHT	0.77	(0.76, 0.79)	2.73	(2.57, 2.90)
Patient enrollment	Other Capitation	0.80	(0.78, 0.82)	1.24	(1.16, 1.32)
model type ³	Other	0.66	(0.62, 0.71)	0.87	(0.70, 1.07)
	Not enrolled	0.79	(0.75, 0.83)	0.66	(0.55, 0.79)
	Personal support	0.98	(0.96, 1.00)	1.18	(1.12, 1.25)
Home care services	Nursing	1.34	(1.31, 1.36)	3.21	(3.07, 3.36)
	Independent/Supervision	(ref)	(ref)	(ref)	(ref)
Function	Mild/Moderate impairment	1.01	(0.99, 1.03)	1.06	(1.01, 1.12)
	Severe impairment	1.05	(1.02, 1.08)	1.06	(0.98, 1.14)
Comition	Independent/Supervision	(ref)	(ref)	(ref)	(ref)
Cognition	Mild/Moderate impairment	0.98	(0.97, 1.00)	0.94	(0.90, 0.99)
	Severe impairment	0.98	(0.94, 1.02)	1.00	(0.91, 1.09)
	No symptoms	(ref)	(ref)	(ref)	(ref)
Mood	Some symptoms	1.10	(1.08, 1.13)	1.12	(1.06, 1.18)
	Daily symptoms	1.22	(1.19, 1.24)	1.17	(1.11, 1.24)
	CHF	1.12	(1.09, 1.14)	1.08	(1.01, 1.14)
Chronic conditions	COPD	1.07	(1.05, 1.09)	1.06	(1.04, 1.12)
	Dementia	1.04	(1.02, 1.07)	0.99	(0.93, 1.05)
NI	0-4	(ref)	(ref)	(ref)	(ref)
Number of	5-8	1.17	(1.14, 1.21)	1.03	(0.95, 1.11)
medications	9+	1.38	(1.34, 1.42)	1.15	(1.07, 1.23)

^{1:} Region is defined by the first letter of a postal code: P - North, K - East, M,N - Central, L - South

2: Rural is defined as having a postal code with a Rurality Index of Ontario 2008 score >= 40

3: EFS - Enhanced fee-for-service, FHT - Family Health Team

PCP I	nome visit		fterhours/ kend visit
Odds Ratio	95% CI	Odds Ratio	95% CI
1.05	(1.02, 1.08)	1.08	(1.04, 1.1)
(ref)	(ref)	(ref)	(ref)
1.34	(1.25, 1.44)	0.91	(0.86, 0.95)
1.80	(1.69, 1.92)	0.95	(0.91, 1.00)
2.63	(2.47, 2.79)	0.96	(0.92, 1.00)
3.89	(3.65, 4.14)	0.96	(0.91, 1.01)
(ref)	(ref)	(ref)	(ref)
0.82	(0.79, 0.85)	0.55	(0.53, 0.57)
0.61	(0.58, 0.65)	0.50	(0.47, 0.53)
1.00	(0.97, 1.03)	1.10	(1.07, 1.13)
0.88	(0.84, 0.92)	0.63	(0.60, 0.65)
(ref)	(ref)	(ref)	(ref)
0.78	(0.76, 0.81)	0.40	(0.39, 0.41)
0.81	(0.79, 0.84)	0.64	(0.63, 0.66)
1.03	(0.93, 1.14)	0.32	(0.28, 0.37)
1.03	(0.96, 1.10)	0.44	(0.41, 0.47)
1.46	(1.41, 1.51)	1.12	(1.09, 1.15)
2.08	(2.02, 2.13)	1.22	(1.19, 1.26)
(ref)	(ref)	(ref)	(ref)
1.39	(1.35, 1.43)	1.00	(0.97, 1.03)
2.69	(2.59, 2.80)	1.27	(1.22, 1.32)
(ref)	(ref)	(ref)	(ref)
1.07	(1.03, 1.10)	0.90	(0.88, 0.92)
1.08	(1.03, 1.13)	0.90	(0.86, 0.95)
(ref)	(ref)	(ref)	(ref)
1.02	(0.99, 1.06)	1.05	(1.02, 1.08)
1.10	(1.07, 1.14)	1.12	(1.09, 1.16)
1.03	(0.99, 1.07)	0.97	(0.93, 1.00)
1.08	(1.04, 1.12)	1.05	(1.02, 1.08)
0.91	(0.88, 0.94)	0.80	(0.78, 0.83)
(ref)	(ref)	(ref)	(ref)
1.08	(1.03, 1.13)	1.13	(1.09, 1.17)
1.20	(1.15, 1.25)	1.22	(1.18, 1.27)

Appendix 1: Database

ICES Databases

Discharge Abstract
Database(DAD)

Ontario Health Insurance Policy Claims (OHIP)

National Ambulatory Care Reporting System (NACRS)

Home care
Database(HCD)

Resident Assessment Instrument (RAI)-Home care source (OACCAC)

Ontario Mental Health Reporting System (OMHRS)

National Rehabilitation Reporting System(NRS)

Continuing Care
Reporting
System(CCRS)

The Corporate Provider Database (CPDB)

Registered Persons Database(RPDB)



es used in the study

Description

The DAD is compiled by the Canadian Institute for Health Information and contains administrative, clinical (diagnoses and procedures/interventions), demographic, and administrative information for all admissions to acute care hospitals, rehab, chronic, and day surgery institutions in Ontario. At ICES, consecutive DAD records are linked together to form 'episodes of care' among the hospitals to which patients have been transferred after their initial admission.

The OHIP claims database contains information on inpatient and outpatient services provided to Ontario residents eligible for the province's publicly funded health insurance system by fee-for-service health care practitioners (primarily physicians) and "shadow billings" for those paid through non-fee-for-service payment plans. The main data elements include patient and physician identifiers (encrypted), code for service provided, date of service, associated diagnosis, and fee paid.

The NACRS is compiled by the Canadian Institute for Health Information and contains administrative, clinical (diagnoses and procedures), demographic, and administrative information for all patient visits made to hospital- and community-based ambulatory care centres (emergency departments, day surgery units, hemodialysis units, and cancer care clinics). At ICES, NACRS records are linked with other data sources (DAD, OMHRS) to identify transitions to other care settings, such as inpatient acute care or psychiatric care. The HCD is a clinical client centric database that captures all services that are provided by or coordinated by Community Care Access Centres (CCACs). The data elements captured include information on: client, intake, assessment, admission & discharge, diagnosis and surgical procedure, and care delivery. ICES receives home care data from the Ontario Ministry of Health and Long-Term Care (MOHLTC). The primary purpose of the information collected through the HCD is to aid in planning and better clinical insight into clients who encounter service through CCACs.

The RAIHC database is managed by the Community Care Access Centres (CCACs) and is a standardized clinical assessment to all long-stay home care clients in Ontario defined as clients receiving ongoing support for at least 60 consecutive days. Data collected include comprehensive clinical, functional and resource utilization information that are used to inform client needs. When used over time, it provides the basis for an outcome-based assessment of the person's response to care or services.

The OMHRS is compiled by the Canadian Institute for Health Information and contains administrative, clinical (diagnoses and procedures), demographic, and administrative information for all admissions to adult designated inpatient mental health beds. This includes beds in general hospitals, provincial psychiatric facilities, and specialty psychiatric facilities. Clinical assessment data is ascertained using the Resident Assessment Instrument for Mental Health (RAI-MH), but different amounts of information are collected using this instrument depending on the length of stay in the mental health bed. Multiple assessments may occur The NRS is compiled by the Canadian Institute for Health Information and contains client data collected from participating adult inpatient rehabilitation facilities and programs across Canada. Main data elements contain socio-demographic information, administrative data (e.g. referral, admission and discharge), health characteristics, activities and participation (e.g. ADL, communication, social interaction), and interventions. The CCRS database is compiled by the Canadian Institute for Health Information and contains demographic, clinical, functional, and resource utilization information for individuals receiving facility-based continuing care (also known as extended, auxiliary, or complex chronic care) in Ontario hospitals and residential care providing 24 hour nursing services (i.e. nursing home). Clinical assessment data (on the physical, functional, cognitive, and social domains of health) is ascertained using the Resident Assessment Instrument Minimum Data Set (RAI-MDS) version 2.0 which is administered by trained healthcare professionals.

The Corporate Provider Database (CPDB) contains information on all physician and some non-physician (such as chiropractors, physiotherapists, and optometrist) providers funded by the Ministry, either through OHIP or other funding arrangements. The data includes demographic, eligibility, specialty, practice location, (encrypted) provider billing number, limited demographic information (year of birth, gender, year of graduation, specialty, and location of practice).

The RPDB provides basic demographic information (age, sex, location of residence, date of birth, and date of death for deceased individuals) for those issued an Ontario health insurance number. The RPDB also indicates the time periods for which an individual was eligible to receive publicly funded health insurance benefits and the best known postal code for each registrant on July 1st of each year.



Appendix 2: Definitions of physician utilization measures

Measure	Database	Field	Values
PCP Visit	OHIP	Spec	00 - FP/GP
			05 - Comm
	OHIP	Feecode	Any feecod
			G202
			G212
			G372
			G373
			G365
			G538
			G539
			G590
			G591
PCP coordination with	OHIP	Spec	00 - FP/GP
home care		•	05 - Comm
	OHIP	Feecode	K071
			K072
			K124
PCP home visit	OHIP	Spec	00 - FP/GP
			05 - Comm
	OHIP	Feecode	A900
			A901
			B960
			B961
			B962
			B963
			B964
			B966
			B990
			B992
			B993
			B994
			B996
			вээо В997
202 6: 1 / 1	OLUB	Cooo	B998
PCP afterhours/weekend	OHIP	Spec	00 - FP/GP
visit	01115		05 - Comm
	OHIP	Feecode	Q012
			Q016
			Q017
			B962
			B963
			B964
			B994
			B993
			B996

	OHIP: Location	Any feecod
Physician specialist visits	OHIP: Spec	Not 00 - FP,
		A888
		A996
		A998
		A994
		A964
		A963
		A962

Note: Visits were limited to 1 per patient per physician per day Note: Visits were excluded if patient was in LTCH at time of billing

unity Medicine
le with location of Office, Home, or Phone

unity Medicine

unity Medicine

unity Medicine

/GP, 05 - Community Medicine, or 26 - Pediatrics le with location of Office, Home, or Phone

g

Appendix 3. Proportion of patients with a PCP coordination with home care k

	PCP home care	coordination (%
LHIN	FHT	non-FHT
Erie St. Clair	3.7	3.7
South West	6.6	6.2
Waterloo Wellington	5.3	1.5
Hamilton Niagara Haldimand Brant	12.5	3.2
Central West	23.1	6.2
Mississauga Halton	1.7	1.7
Toronto Central	3.0	2.3
Central	3.5	2.4
Central East	3.0	1.7
South East	12.6	3.7
Champlain	2.4	1.1
North Simcoe Muskoka	4.8	2.6
North East	4.0	4.2
North West	9.0	1.2



STROBE Statement—Checklist of items that should be included in reports of *cohort studies*

	Item No	Recommendation	Line #
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	2
		(b) Provide in the abstract an informative and balanced summary of what was done	61-83
		and what was found	_
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	86-97
Objectives	3	State specific objectives, including any prespecified hypotheses	98-102
Methods			
Study design	4	Present key elements of study design early in the paper	
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment,	_ 112-115
		exposure, follow-up, and data collection	133
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of	_ 112-11
		participants. Describe methods of follow-up	_
		(b) For matched studies, give matching criteria and number of exposed and	NA
		unexposed	_
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect	120-15
		modifiers. Give diagnostic criteria, if applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of methods of	App. 1
measurement		assessment (measurement). Describe comparability of assessment methods if there is	
		more than one group	
Bias	9	Describe any efforts to address potential sources of bias	_ 162,17
Study size	10	Explain how the study size was arrived at	114-1
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable,	153-18
Statistical methods	12	describe which groupings were chosen and why	_ 173-18
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding (b) Describe any methods used to examine subgroups and interactions	_ 173-16 169
		(c) Explain how missing data were addressed	NA
		(d) If applicable, explain how loss to follow-up was addressed	162
		(e) Describe any sensitivity analyses	NA
		(e) Describe any sensitivity analyses	- '''
Results	124		-
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study,	183
		completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	NA
		(c) Consider use of a flow diagram	- NA
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and	_ 1 7 \ 183-19
Descriptive data	1.	information on exposures and potential confounders	100-19
		(b) Indicate number of participants with missing data for each variable of interest	NA
		(c) Summarise follow-up time (eg, average and total amount)	- Fig 1, 19
Outcome data	15*	Report numbers of outcome events or summary measures over time	
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and	Tbl 2, 4 199-22
		their precision (eg, 95% confidence interval). Make clear which confounders were	199-22
		adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were categorized	Tbl4
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a	- Tbl3, 2
		meaningful time period	-, -

Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and	205
		sensitivity analyses	_
Discussion			
Key results	18	Summarise key results with reference to study objectives	231-237
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or	273-278
		imprecision. Discuss both direction and magnitude of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,	238-270
		multiplicity of analyses, results from similar studies, and other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	278
Other information			<u></u>
Funding	22	Give the source of funding and the role of the funders for the present study and, if	30
		applicable, for the original study on which the present article is based	

^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at http://www.strobe-statement.org.