

# UNIVERSITY OF SASKATCHEWAN

# **The Impact of Continuity of Care on Medication Adherence**

Affiliations: 1. Department of Pharmacy, College of Pharmacy and Nutrition, University of Saskatchewan, Saskatoon, Canada; 2. Department of Community Health Sciences, Rady Faculty of Health Sciences, University of Manitoba, Winnipeg, Manitoba, Canada; 3. Population Public and Indigenous Health, Alberta Health Services, Calgary, Alberta, Canada.

# **Background:**

Objectives: Continuity of care (COC) is considered an important determinant of medication adherence based on measures such as the usual provider continuity index (UPCI) that are derived exclusively from physician visit claims. This study aimed to: a) determine if high UPCI values predict physicians who deliver different clinical services; and b) compare UPCI with an integrated COC measure in a multivariable model of patients receiving statin medications.

### **Methods:**

This was a retrospective cohort study of new statin users between 2012 and 2017 in Saskatchewan, Canada. We calculated sensitivity/specificity of a high UPCI value for predicting physicians who were prescribers of statins and/or providers of complete medical examinations. Next, we used logistic regression models to test two measures of COC (high UPCI value or an integrated COC measure) on the outcome of optimal statin adherence (proportion of days covered  $\geq 80\%$ ). The DeLong test was used to compare predictive performance of the two models.

# **Results:**

Among 55,144 new statin users, a high UPCI was neither a sensitive or specific marker of physicians who prescribed statins or performed a complete medical exam. The integrated COC measure had a stronger association with optimal adherence (adjusted odds ratio [OR] =1.56, 95% confidence interval [CI] 1.50 to 1.63) than UPCI (adjusted OR =1.23, 95% CI 1.19 to 1.28), and improved predictive performance of the adherence model.

# **Conclusion:**

The number of physician visits alone appears to be insufficient to represent COC. An integrated measure improves predictive performance for optimal medication adherence in patients initiating statins.



#### **Figure 1. Cohort flow chart**

Number of individuals eligible to receive health insurance in Saskatchewan Receiving statin medications between January 1st, 2012 and December 31st, 2017 179,923 Decease 158,861 Receiv Study cohort (new users of statin medications) 58,549 Admittee 57,761 Staying 57,420 Pregn 57,240 Not visi Number of individuals in the final cohort 55,144

<sup>a</sup>Index date = the earliest date receiving a statin medication between January 1st, 2012 and December 31st, 2017;  $^{b}GP = general practitioner.$ 

### Table 1. Baseline characteristics<sup>a</sup> of the final cohort.

	All	Patients grouped by UPCI <sup>b</sup>		Patients grouped by integrated COC <sup>c</sup>	
		High(>=0.82)	Low(<0.82)	Yes	No
	n=55,144	n=27,859	n=27,285	n=15,579	n=39,565
Median age (Q25, Q75)	59.0 (51.0, 67.0)	59.0 (52.0, 68.0)	58.0 (50.0, 67.0)	59.0 (51.0, 67.0)	59.0 (51.0, 67.0)
Females (n, %)	24,385 (44.2%)	11,635 (41.8%)	12,750 (46.7%)	6,840 (43.9%)	17,545 (44.3%)
Patients with one or more hospitalizations for acute care (n, %)	12,528 (22.7%)	6,203 (22.3%)	6,325 (23.2%)	2,626 (16.9%)	9,902 (25.0%)
Median number of visits to GPs <sup>e</sup> (Q25, Q75 <sup>d</sup> )	6.0 (3.0, 9.0)	5.0 (3.0, 9.0)	6.0 (3.0, 10.0)	6.0 (3.0, 9.0)	5.0 (3.0, 9.0)
Median number of visits to specialists (Q25, Q75 <sup>d</sup> )	2.0 (0.0, 6.0)	2.0 (0.0, 6.0)	2.0 (0.0, 6.0)	2.0 (0.0, 5.0)	2.0 (0.0, 7.0)
Patients with one or more visits to emergency department (n, %)	11,450 (20.8%)	5,519 (19.8%)	5,931 (21.7%)	2,739 (17.6%)	8,711 (22.0%)
Patients by income level (n, %)					
1 (lowest)	10,339 (18.7%)	4,787 (17.2%)	5,552 (20.3%)	2,675 (17.2%)	7,664 (19.4%)
2	10,207 (18.5%)	5,058 (18.2%)	5,149 (18.9%)	2,761 (17.7%)	7,446 (18.8%)
3	10,093 (18.3%)	5,182 (18.6%)	4,911 (18.0%)	2,942 (18.9%)	7,151 (18.1%)
4	11,289 (20.5%)	5,897 (21.2%)	5,392 (19.8%)	3,251 (20.9%)	8,038 (20.3%)
5 (highest)	10,268 (18.6%)	5,456 (19.6%)	4,812 (17.6%)	3,052 (19.6%)	7,216 (18.2%)
missing	2,948 (5.3%)	1,479 (5.3%)	1,469 (5.4%)	898 (5.8%)	2,050 (5.2%)
Patients by residence location (n, %)					
Rural	17,811 (32.3%)	8,666 (31.1%)	9,145 (33.5%)	4,364 (28.0%)	13,447 (34.0%)
Urban	37,333 (67.7%)	19,193 (68.9%)	18,140 (66.5%)	11,215 (72.0%)	26,118 (66.0%)

<sup>a</sup> Median age, number of females, residence (rural/urban), and patient income level were measured on the index date; Number of patients with one or more hospitalizations, median visits to GPs/specialists, patients with one or more visits to emergency departments were measured within one year prior to the index date; <sup>b</sup>UPCI = Usual Provider Continuity index; <sup>c</sup>COC = continuity of care;  $^{d}Q25 = 25\%$  percentile, Q75 = 75% percentile;  $^{e}GP =$  general practitioners.

# YAO SZ<sup>1</sup>, Lix L<sup>2</sup>, Teare G<sup>3</sup>, Charity E<sup>1</sup>, Blackburn DF<sup>1</sup>

Not receiving statin 1,190,736
Tissing birth/sex information or age $< 18$ on the index date <sup>a</sup> <b>87</b>
d or not continuously receiving health insurance within fiver years prior to, or one year on and after the index date <sup>a</sup> 21,062
ing statin medications within five years prior to the index date <sup>a</sup> 100,312
d to long term care facilities within five years prior to, or one year on and after the index date <sup>a</sup> <b>788 (1.3% of the study cohort)</b>
g in an out-of-province facility for acute care within one year on and after the index date <sup>a</sup> 341 (0.6% of the study cohort)
ant within one year prior to, or one year on and after the index date <sup>a</sup> 180 (0.3% of the study cohort)
iting a GP <sup>b</sup> within one year on and after the index date <sup>a</sup> , or none of the visited physicians can be identified <b>2,096 (3.6% of the study cohort)</b>

#### Table 2. Measures of accuracy using UPCI<sup>a</sup> to predict USP<sup>b</sup>, CMEP<sup>c</sup>, and integrated COC<sup>d</sup> status.



<sup>a</sup>UPCI = usual provider continuity index; <sup>b</sup>USP=usual statin prescriber; <sup>c</sup>CMEP = complete medical examination provider; <sup>d</sup>COC = continuity of care; <sup>e</sup>CI = confidence interval; <sup>f</sup>PPV=positive predictive value; <sup>g</sup>NPV=negative predictive value.

#### Table 3: Odds ratios (OR<sup>a</sup>) and 95% confidence intervals (95% CI<sup>b</sup>) for the association of measures of COC<sup>c</sup> with optimal adherence (PDC<sup>d</sup> >= 80%)

	Unadjusted model OR <sup>a</sup> (95%CI <sup>b</sup> )	Adjusted model <sup>g</sup> OR <sup>a</sup> (95%CI <sup>b</sup> )
Integrated COC <sup>e</sup>	1.45 (1.40, 1.51)	1.56 (1.50, 1.63)
Among patients with high UPCI <sup>f</sup>		1.48 (1.40, 1.56)
Among patients with low UPCI <sup>f</sup>		1.60 (1.51, 1.70)
UPCI <sup>f</sup>	1.28 (1.24, 1.32)	1.23 (1.19, 1.28)
Patients presenting integrated COC <sup>c</sup>		1.13 (1.06, 1.21)
Patients not presenting integrated COC <sup>c</sup>		1.22 (1.17, 1.27)

<sup>a</sup>OR = odds ratio; <sup>b</sup>CI = confidence interval; <sup>c</sup>COC = continuity of care; <sup>d</sup>PDC = proportion of days covered; <sup>e</sup>Integrated COC = having a single physician identified as the usual care provider, the usual statin prescriber, and the complete medical examination provider; <sup>f</sup>UPCI = usual provider continuity index; <sup>g</sup>Covariates in the adjusted model included 1) age, sex, residence (rural/urban), and income level (i.e., the neighborhood median household income quintile lowest=1, highest=5) on the index date; 2) the following measured within 365 days prior to the index date: number of hospitalizations, number of out-patient visits (to GPs and to specialists, respectively), number of emergency department visits, Charlson comorbidity score, number of distinct prescription medications (by drug identification numbers), and percentage of prescription medication cost paid by government health insurance; and 3) a list of chronic conditions identified between January 1<sup>st</sup>, 1996, and the index date, including osteoporosis, rheumatoid arthritis, hypertension, stroke, ischemic heart disease, acute myocardial infarction, heart failure, multiple sclerosis, Parkinson's disease, Alzheimer's disease and dementia, epilepsy, asthma, chronic obstructive pulmonary disease, diabetes, mood and anxiety diseases, schizophrenia, schizophrenia, and cancer.

Acknowledgement: "The authors acknowledge the Health Quality Couns for use of de-identified data provided by the Saskatchewan Ministry of Health and eHealth Saskatchewan. The interpretation and conclusions contained herein do not necessarily represent those of the Government of Saskatchewan, the Saskatchewan Ministry of Health, or eHealth Saskatchewan.

**Disclosures:** David Blackburn is the Chair in Patient Adherence to Drug Therapy within the College of Pharmacy and Nutrition, University of Saskatchewan. This position was created through unrestricted financial support from AstraZeneca Canada, Merck Canada, Pfizer Canada, and the Province of Saskatchewan's Ministry of Health. None of the sponsors were involved in developing this study or writing the manuscript. Shenzhen Yao, Lisa Lix, Gary Teare, and Charity Evans declare no conflicts.



Sensitivity	Specificity	<b>PPV<sup>f</sup> (95%CI<sup>e</sup>)</b>	NPV <sup>g</sup>	Карра
(95%CI <sup>e</sup> )	(95%CI <sup>e</sup> )		(95%CI <sup>e</sup> )	(95%СІ <sup>е</sup> )
0.55	0.61 (0.60, 0.62)	0.78	0.35	0.13 (0.13, 0.14)
0.55 (0.54, 0.56)	0.52 (0.51, 0.52)	0.39 (0.39, 0.40)	0.67 (0.66, 0.68)	0.06 (0.05, 0.07)
0.58	0.53	0.33	0.76	0.09 (0.08, 0.09)
(0.58, 0.59)	(0.52, 0.53)	(0.32, 0.33)	(0.76, 0.77)	

