Evaluation of Association Between Diabetes-Related Quality Measure Achievement and Diabetes Complications in a Medicare Advantage Population

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Background

- Individuals with Type 1 diabetes mellitus (T1DM) or Type 2 diabetes mellitus (T2DM) may be at risk for a range of complications including retinopathy, nephropathy, limb amputation, and the development of cardiovascular complications.
- Centers for Medicare and Medicaid Services (CMS) rates Medicare Part C and D plans on several • quality measures for the purpose of educating consumers on the quality of Medicare plans in the marketplace including increasing the transparency of health plan performance and comparison.¹
- Specific quality measures for Medicare Part C and D focus on indicators that track quality of medical ٠ care and medication management, in diabetic patients.^{1,2}
- It remains unclear if attaining these CMS diabetes quality indicators improves outcomes.

Objective

Assess the achievement of eight diabetes-related quality measures, at the patient level, and examine whether achievement was associated with fewer complications.

Methods

- Study Design: Retrospective cohort study
- **Data Source:** Humana administrative database, which contains integrated medical, pharmacy, and lab-related claims, and member-level achievement indicators for Medicare Part C quality measures. Medicare Part D measures were provided to Humana by Accumen, LLC, a third party vendor.
- **Patient Selection:** ٠
 - Medicare Advantage plan with Prescription Drug coverage (MAPD) patients, aged 18 to 75 years as of December 31, 2011, continuously enrolled from January 1, 2010 to December 31, 2011, with a diagnosis of diabetes during the pre-index period, defined by at least one of the following:
 - ≥1 prescription claims for insulin or oral hypoglycemics/antihyperglycemics.
 - ≥2 two face-to-face encounters, in an outpatient setting or nonacute inpatient setting, on different dates of service, with a diagnosis of diabetes (250.x).
 - ≥1 face-to-face encounter in an acute inpatient or ED setting, with a diagnosis of diabetes.
 - Exclusions included:
 - Diagnosis of gestational diabetes (648.8) or pregnancy (630.xx-679.xx or v22.x-v24.x) at any position during the study period.
 - Diagnosis of both T1DM (250.x1 or 250.x3) and T2DM (all other 250.x ICD-9 CM codes) AND the presence of ≥ 1 prescription claims for oral hypoglycemics/antihyperglycemics.
 - Long-term care facility stay >30 days at any point during the study period.
 - Quality Measure-Specific Exclusion Criteria:
 - Each quality measure has its own CMS-defined exclusion criteria¹ and sample size.

- LDL-C Screening: Clinically desirable
 - measure achieved if an LDL-C test was performed during the measurement year, as identified by claim/encounter or automated laboratory data.
- LDL-C Control <100 mg/dL: Clinically desirable -
 - measure achieved if the most recent LDL-C level is <100 mg/dL.
- Medical Attention for Nephropathy: Clinically desirable -
 - measure achieved if a nephropathy screening test or evidence of nephropathy, as documented through administrative data, is present during the measurement year.
- Medication Adherence to Oral Diabetes Medications: Clinically desirable
 - measure achieved if proportion of days covered (PDC) was greater than or equal to 80% across biguanides, sulfonylureas, and thiazolidinediones, or combinations thereof.
- Diabetes Medication Dosing (DMD): Not clinically desirable
 - measure achieved if patient dispensed a dose higher than the daily recommended dose for the following diabetes treatment therapeutic categories of oral hypoglycemic: biguanides, sulfonylureas, and thiazolidinediones (TZDs).
 - DMD quality measure was analyzed separately across the three classes.
 - Due to data availability, 1 of 62 MAPD was not available for this measure.
- Diabetes Treatment (DT): Not clinically desirable
 - measure achieved if patient had a day's supply of renin angiotensin system (RAS) antagonist less than the largest days' supply of a different antihypertensive agent.
 - Due to data availability, 38 of the 62 MAPD plans were available for this measure.
- **Outcome: New or Worsening Diabetic Complications** •
 - Diabetes Complications Severity Index (DCSI) score has been shown to predict adverse outcomes including hospitalization and mortality based on number and severity of complications associated with diabetes.³
 - Patients were assessed for an increase in diabetes-related complications that occurred during the measurement year.
 - Baseline DCSI was scored for the pre-index period (2010).
 - Measurement year DCSI was scored across both the pre-index (2010) AND the post-index (measurement year; 2011) periods.
 - A member with a measurement year DCSI greater than the baseline DCSI was categorized as having "New or Worsening Complications."
- **Statistical Analyses**
 - Unadjusted association between guality measure achievement and new or worsening diabetes complications was assessed using χ^2 tests or the Fisher exact test, as appropriate.
- Baseline Characteristics: Pre-index demographic and clinical characteristics were assessed.
- Quality Measures: Eight DM-related quality measures¹ studied and clinical desirability of measure achievement listed below:
 - HbA1c Poor Control >9%: Not clinically desirable
 - measure achieved if the most recent HbA1c level is >9.0%, missing, or was not done during the measurement year.
- Eye Exam: Clinically desirable
 - measure achieved if ≥1 of the following exams is completed by an eye care professional: a retinal or dilated eye exam in the measurement year, OR a negative retinal or dilated eye exam (negative for retinopathy) in the year prior to the measurement year.
- Analyses were conducted for all Part D measures (Adherence, DMD, DT) and for two of the Part C Measures (HbA1c Poor Control >9% and LDL-C Screening).
- Adjusted impact of quality measure achievement on new or worsening diabetes complications was assessed by a stepwise logistic regression model, adjusted for all preindex (baseline) demographic characteristics and the baseline DCSI score.
 - Models prepared for all Part D measures (Adherence, DMD, DT) and for two of the Part C Measures (HbA1c Poor Control >9% and LDL-C Screening).
 - Odds Ratios and 95% confidence intervals (CI) reported.
- A priori alpha level for all inferential analyses was set at 0.05, and all statistical tests were two-tailed.

Results

Table 1. Baseline Demographic and Clinical Characteristics by Quality Measure(s)

Characteristic	General Population (n=164,238)	Part C Measure* Qualifiers (n=159,454)	Adherence Measure Qualifiers (n=95,978)	DMD – Biguanides Measure Qualifiers (n=83,759)	DMD – Sulfonylureas Measure Qualifiers (n=52,541)	DMD – Thiazolidinediones Measure Qualifiers (n=14,323)	Diabetes Treatment Measure Qualifiers (n=4,464)
Age, years , mean (SD)	68.0 (±6.4)	67.9 (±6.5)	67.9 (±6.3)	67.7 (±6.5)	68.2 (±6.1)	67.9 (±6.4)	67.7 (±6.3)
Age Category							
18-29	44 (0.03%)	45 (0.03%)	23 (0.02%)	23 (0.03%)	<10 (0.01%)	<10 (0.03%)	<10 (0.0%)
30-39	578 (0.4%)	580 (0.4%)	303 (0.3%)	307 (0.4%)	108 (0.2%)	44 (0.3%)	15 (0.3%)
40-49	3,274 (2.0%)	3,399 (2.1%)	1,892 (1.9%)	1,769 (2.1%)	881 (1.7%)	282 (1.9%)	77 (1.7%)
50-59	13,336 (8.1%)	13,266 (8.3%)	8,006 (8.3%)	7,320 (8.7%)	4,108 (7.8%)	1,215 (8.5%)	418 (9.4%)
60-69	62,541 (38.1%)	60,476 (37.9%)	37,222 (38.8%)	32,974 (39.4%)	20,170 (38.4%)	5,444 (38.0%)	1,837 (41.2%)
70-79	84,465 (51.4%)	81,688 (51.2%)	48,532 (50.6%)	41,366 (49.4%)	27,270 (51.9%)	7,334 (51.2%)	2,117 (47.4%)
Gender							
Female	82,447 (50.2%)	80,148 (50.3%)	48,020 (50.0%)	42,432 (50.6%)	24,725 (47.1%)	6,474 (45.2%)	2,274 (50.9%)
Male	81,791 (49.8%)	79,306 (49.7%)	47,958 (50.0%)	41,327 (49.4%)	27,816 (52.9%)	7,849 (54.8%)	2,190 (49.1%)
Race/Ethnicity							
White	131,073 (79.8%)	125,554 (78.7%)	75,960 (79.1%)	66,582 (79.5%)	40,936 (77.9%)	11,220 (78.3%)	3,710 (83.1%)
Black	25,357 (15.4%)	25,715 (16.1%)	14,985 (15.6%)	12,678 (15.1%)	8,753 (16.7%)	2,220 (15.5%)	367 (8.2%)
Hispanic	2,677 (1.6%)	3,099 (1.9%)	1,721 (1.8%)	1,553 (1.9%)	975 (1.9%)	320 (2.2%)	72 (1.6%)
Other/Unknown	5,131 (3.1%)	5,086 (3.2%)	3,312 (3.5%)	2,946 (3.5%)	1,877 (3.6%)	563 (3.9%)	315 (7.1%)
Geographic Region							
Midwest	44,954 (27.4%)	42,381 (26.6%)	21,931 (22.8%)	19,234 (22.9%)	12,053 (22.9%)	3,509 (24.5%)	619 (13.9%)
Northeast	2,925 (1.8%)	2,711 (1.7%)	1,803 (1.9%)	1,591 (1.9%)	998 (1.9%)	275 (1.9%)	489 (10.9%)
South	104,073 (63.4%)	102,770 (64.4%)	64,109 (66.8%)	55,581 (66.4%)	35,487 (67.5%)	9,276 (64.7%)	1,917 (42.9%)
West	12,286 (7.5%)	11,592 (7.3%)	8,135 (8.5%)	7,353 (8.8%)	4,003 (7.6%)	1,263 (8.8%)	1,439 (32.2%)
Population Density							
Urban	100,672 (61.3%)	98,931 (62.0%)	58,306 (60.7%)	50,946 (60.8%)	32,095 (61.1%)	8,500 (59.3%)	2,838 (63.6%)
Suburban	42,885 (26.1%)	40,963 (25.7%)		21,900 (26.2%)		3,882 (27.1%)	1,127 (25.3%)
Rural	19,550 (11.9%)	18,484 (11.6%)	11,847 (12.3%)	10,313 (12.3%)	6,521 (12.4%)	1,842 (12.9%)	458 (10.3%)
Unknown	1,131 (0.7%)	1,076 (0.7%)	687 (0.7%)	600 (0.7%)	385 (0.7%)	99 (0.7%)	41 (0.9%)
Low Income Subsidy							
Yes	30,788 (18.8%)	34,513 (21.6%)	20,632 (21.5%)	17,595 (21.0%)	11,124 (21.2%)	4,020 (28.1%)	994 (22.3%)
Dual Eligibility Status		, <u>,</u> ,					
Yes	3,688 (2.3%)	4,532 (2.8%)	2,509 (2.6%)	2,177 (2.6%)	1,344 (2.56%)	534 (3.7%)	113 (2.5%)
Diabetes Type							
Type 1	5,194 (3.2%)	4,923 (3.1%)	310 (0.3%)	317 (0.4%)	242 (0.5%)	68 (0.5%)	12 (0.3%)
Type 2	153,736 (93.6%)	152,663 (95.7%)		81,332 (97.1%)		14,086 (98.3%)	4,321 (96.8%)
Unknown	5,308 (3.2%)	1,868 (1.2%)	2,221 (2.3%)	2,110 (2.5%)	612 (1.2%)	169 (1.2%)	131 (2.9%)
Comorbidity Measures	, <u>, , , , , , , , , , , , , , , , , , </u>						
DCCI Score, mean (SD)	2.2 (±1.9)	2.3 (±1.9)	2.2 (±1.8)	2.0 (±1.7)	2.4 (±1.9)	2.1 (±1.7)	1.9 (±1.6)
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*Part C measures include: HbA1c Poor Control >9%, Eye Exam, LDL-C Screening, LDL-C Control <100 mg/dL, Medical Attention for Nephropathy; Sample sizes depending on the quality measure's CMS-defined exclusion criteria and patient-level data availability; SD, Standard Deviation; DCCI, Deyo-**Charlson Comorbidity Index**

Table 2. Unadjusted Associations of New or Worsening **Diabetes Complications and Quality Measure Achievement**

Quality Measure	Quality Measure Achieved?	Definition	% of Patients Experiencing New or Worsening Diabetes Complications	p-value	
HbA1c Poor Control >9%	Yes	HbA1c Poor Control >9%	39.7%	<0.0001	
	No	HbA1c Control ≤9%*	36.0%		
LDL-C Screening	Yes	At least 1 LDL Screen in Year*	38.3%	<0.0001	
	No	No LDL Screens in Year	31.9%		
Adherence	Yes	PDC ≥ 80%*	37.8%	0.0003	
	No	PDC < 80%	36.4%		
	Yes	Dosing EXCEEDS Recommended	41.2%	0.0296	
DMD - Biguanides	No	Within Recommended Dosing*	37.1%		
DMD - Sulfonylureas	Yes	Dosing EXCEEDS Recommended	43.6%	0.1354	
	No	Within Recommended Dosing*	39.9%		
DMD - Thiazolidinediones	Yes	Dosing EXCEEDS Recommended	44.0%	0.5425	
	No	Within Recommended Dosing*	38.1%		
Diabetes	Yes	Suboptimal Antihypertensive Therapy	42.3%	<0.0001	
Treatment	No	RAS Antagonist Overlap 100%*	34.1%		

*Clinically-desirable outcome

Table 3. Adjusted* Odds of New or Worsening Diabetes **Complications by Quality Measure Achievement**

Quality Measure	Quality Measure Achieved?	Definition	Odds of New or Worsening Diabetes Complications OR (95% CI)	p-value	
HbA1c	Yes	HbA1c Poor Control >9%	1.12 (1.10 - 1.15)	-0.0001	
Poor Control >9%	No	HbA1c Control ≤9%**	reference	<0.0001	

Figure 1. Quality Measure Achievement Rates



+ = Higher achievement rate is desirable; + = Lower achievement rate is desired.

Results Summary

- Cohort size ranged from 4,464 to 159,454, depending on the guality measure and patient-level data availability (Table 1).
- Most patients (>80%) achieved LDL-C screening, nephropathy, and adherence standards (Figure 1).
- <1% of patients exceeded dosing standards for biguanides, sulfonylureas, and TZDs (Figure 1).
- Eye screening and use of appropriate anti-hypertensive treatments had low achievement levels (50.3% and 54.2%, respectively; Figure 1).
- A majority (61%) of patients achieved HbA1c<9% while 29% achieved LDL-C control <100mg/dl (Figure 1).
- Statistically significant univariate associations were identified for new or worsening diabetes complications and achievement of: HbA1c Poor Control >9%, LDL-C Screening, Adherence, DMD-Biguanides, Diabetes Treatment (Table 2).
- Logistic regression estimates showed that the odds of developing new or worsening diabetes complications (Table 3) statistically significantly increased with:
 - Failure to reduce HbA1c below 9% [(OR, 1.12 (95% CI, 1.10-1.15); p<0.0001)].
 - Failure to use RAS Antagonist anti-hypertensive treatment [(OR, 1.40, (95% CI, 1.24-1.59); p<0.0001].
- Counter-intuitively, the odds of developing new or worsening diabetes complications (Table 3):
 - Statistically significantly increased in patients with at least 1 LDL-screen [(OR, 1.32 (95% CI, 1.28-1.36); p<0.0001)].
 - Statistically significantly decreased in patients with a PDC < 80% [(OR, 0.94 (95% CI, 0.91-0.97); p=0.0005)]

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Yes At least 1 LDL Screen in Year** 1.32 (1.28 - 1.36) LDL-C Screening <0.0001 No LDL Screens in Year reference No PDC ≥ 80%** Yes reference 0.0005 Adherence No PDC < 80% 0.94 (0.91 - 0.97) **Dosing EXCEEDS Recommended** 1.15 (0.99 - 1.35) Yes 0.0733 **DMD** - Biguanides No Within Recommended Dosing** reference **Dosing EXCEEDS Recommended** 1.15 (0.94 - 1.40) Yes DMD -0.1826 Sulfonylureas No Within Recommended Dosing** reference **Dosing EXCEEDS Recommended** Yes 1.37 (0.61 - 3.07) DMD · 0.4466 Thiazolidinediones No Within Recommended Dosing** reference Suboptimal Antihypertensive Yes 1.40 (1.24 - 1.59) Diabetes Therapy <0.0001 Treatment

*Adjusted for all pre-index (baseline) demographic characteristics and the baseline DCSI score; **Clinicallydesirable outcome; OR, odds ratio; CI, confidence interval.

RAS Antagonist Overlap 100%**

Conclusion

The assessment of a 1-year measurement period suggests that attainment of several CMS diabetes quality measures may be associated with lower new or worsening complication risk. Follow-up longitudinal studies may provide clarity on the long-term impact of achieving quality measures.

No

Limitations

Comprehensive

Health Insights

A Humana.Company

- This study utilized data from Humana MAPD health plans only and may not be generalizable.
- Due to limitations of data provided by Humana's quality data third party vendor, a number of plans were excluded due to incomplete data.
- Causal inference cannot be directly determined as relationships between quality measure achievement and outcomes were based on statistical associations and temporal relationships.

References

reference

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