

**Medicare Part D Data Linked with the Health Outcomes Survey: Association between Quality of Care using Prescription Drugs and Mortality as Outcomes among those Enrolled in the Medicare Advantage Program**

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## Table of Contents

Executive Summary	3
Introduction	5
Methods	8
Results	12
Discussion	14
Recommendations for CMS	18
Figure 1	19
Table 1	20
Table 2	21
Table 3	21
Appendix A	23
Appendix B	30
References	31

## Executive Summary

**Background.** On January 1, 2006, the Centers for Medicare and Medicaid Services implemented the Medicare Drug Benefit, or "Medicare Part D." Although reports point to positive effects of the Part D benefit, several questions remain unanswered about its overall effect on patient outcomes in the Medicare Advantage Program. One of the major unanswered questions involves the impact of the benefit of the Part D program on selected medications for those elderly with common chronic illnesses on their mortality.

**Objectives.** We evaluated the relationship between use of medications based upon nationally recognized clinical practice guidelines and health outcomes using mortality among Medicare Advantage (MA) patients enrolled in the Medicare Part D program.

**Methods.** We linked data from the Medicare Health Outcomes (HOS) Survey cohort 9 (April 2006 – May 2008) with the Medicare Part D prescription benefit files (January 1<sup>st</sup>, 2006 – December 31<sup>st</sup>, 2007) to calculate the medication based performance indicators for five high volume chronic conditions (diabetes, coronary artery disease (CAD), congestive heart failure (CHF), chronic obstructive lung disease (COPD)/asthma and depression). We used analysis of variance to examine the variations of the performance indicators across plans and correlation analyses to examine the associations of performance indicators and mortality at the patient and the plan levels. Mortality was operationalized as the ratio of observed/expected mortality rates with the expected mortality based upon propensity analysis with demographic and clinical covariates.

**Results.** Analysis of variance confirmed that the 203 MA plans differed significantly in their performance indicators ( $P < 0.001$ ). The smallest variation range was for diabetic patients

receiving from 52.2% to 100% of the indicated medications. The largest variation range was for patients with depression receiving antidepressants (7.4% to 66.7%). At the patient level, there were weak associations between the use of lipid-lowering medications and mortality among patients with diabetes ( $r = -0.02$ ,  $p = 0.005$ ) and CAD ( $r = -0.04$ ,  $p < 0.0001$ ). Beta-blocker use was also associated with a lower mortality among patients with CHF, albeit a weak correlation ( $r = -0.02$ ,  $p = 0.034$ ). At the plan level, the higher prevalence of angiotensin-converting enzyme (ACE) inhibitors/angiotensin receptor blockers (ARB) ( $r = -0.40$ ,  $p < 0.0001$ ) and beta-blockers ( $r = -0.27$ ,  $p < 0.0001$ ) were found to be significantly associated with a lower mortality among patients with CHF, while for lipid-lowering medications ( $r = -0.11$ ,  $p = 0.091$ ) it approached significance. Antidepressants showed a paradoxical correlation. They were associated with higher mortality in depressed patients at the patient level with a weak correlation ( $r = 0.02$ ,  $p = 0.007$ ), and a higher mortality at the plan level ( $r = 0.28$ ,  $p < 0.0001$ ).

**Conclusions.** While our results are not definitive, they reveal a complex relationship between outcome and different medication based process indicators. These findings support selective associations between mortality and process indicators that warrant further study using data such as the Healthcare Effectiveness Data and Information Set (HEDIS) and follow-up information that combines process measures and outcomes over a longer period of time.

## Introduction

By age 75, the average American has 2 or 3 chronic medical conditions.<sup>1</sup> The management of patients with chronic conditions can be challenging and costly.<sup>2</sup> The Medicare Modernization Act Prescription Drug Benefit plan (Part D) established a prescription drug benefit for all 43 million Medicare beneficiaries in the United States.<sup>3</sup> The prescription benefit has increased the proportion of Medicare enrollees with prescription drug coverage.<sup>4</sup> The Centers for Medicaid & Medicare Services (CMS) report that among seniors eligible for Medicare, Part D enrollees save on out-of-pocket expenses compared with those not enrolled in Part D.<sup>5</sup> Although reports point to positive effects of the Part D benefit, several questions remain unanswered about its overall effect on patient outcomes.<sup>6,7</sup> One of the major unanswered questions involves the impact of the benefit of the Part D program on selected medications for those elderly with common chronic illnesses on their mortality. The Medicare Health Outcomes Survey (HOS) provides a unique opportunity to examine the outcomes of Medicare Advantage (MA) patients enrolled in the Medicare Part D program.

There are now clearly defined therapeutic and preventive interventions that are known to affect health outcomes. Examples include the following: the use of HMG CoA reductase inhibitors for the treatment of hyperlipidemias that has reduced mortality by as much as 30% and the administration of angiotensin-converting enzyme (ACE) inhibitors in patients with heart failure and depressed left ventricular function that lowers the six month mortality by 40% and the 12-month by 31% ( $p=0.003$ ).<sup>8,9</sup> Aspirin for primary and secondary prevention of cardiovascular disease is another significant outpatient intervention to reduce mortality.<sup>10</sup> Inhaled

steroids have reduced the mortality of patient with chronic obstructive pulmonary disease by 29% (95% CI, 22% to 35%).<sup>11</sup> However, recent research indicates that some of the most debilitating, costly, and treatable illnesses are undertreated. In several large studies in the United States, less than half the patients with serious depression received effective medication or at least short-term psychotherapy.<sup>12</sup> Elderly patients with heart failure are less likely to receive evidence-based treatments and consequently have a worse prognosis.<sup>13</sup> They received ACE inhibitors, beta-blockers, and anticoagulants less frequently than younger patients did. The 1- year mortality rate was significantly higher in patients  $\geq 70$  years old (22% vs. 13.7%,  $P < 0.001$ ). Despite substantial evidence that antiplatelet therapy saves lives and reduces adverse events in patients with coronary artery disease (CAD), use of the most widely available and lowest cost antiplatelet agent, aspirin, continues to be disappointingly low.<sup>14</sup> At one teaching hospital, one third of patients with acute myocardial infarction who were eligible for treatment with beta-blocker drugs did not receive any of those drugs, contrary to the guidelines of the American College of Cardiology.<sup>15</sup> As a result, the mortality rate among these patients was 20 to 40 percent higher than that among patients given a beta-blocker. Few patients with diabetes are taking aspirin (44%), lipid-lowering (20%), or ACE inhibitors (39%) therapy.<sup>16</sup> Compared to those taking aspirin or ACE inhibitors, the odds of dying for those patients without ACE inhibitor therapy was 1.7 (95% CI, 1.2 - 3.9) and those with no aspirin therapy was 2.89 (95% CI, 1.9 - 4.3).

This study examined the relationship between use of medications based upon nationally recognized clinical practice guidelines (CPG) and health outcomes using mortality among MA patients enrolled in the Medicare Part D program. Linking process indicators and outcomes is critical to accurately estimating healthcare quality and quantifying its benefits. The specific

objectives of this study were (1) to evaluate the variation of nationally recommended use of medications for five high volume diagnoses (diabetes, CAD, congestive heart failure (CHF), chronic obstructive pulmonary disease (COPD) / asthma, and depression) across MA plans and (2) to examine the association of nationally recognized use of medications and mortality at the patient and plan levels.



## Methods

### Study Population

The HOS Performance Measurement analysis is limited to beneficiaries who meet the following analytic criteria:

1. Were age sixty-five or older at the time of completing the baseline survey.
2. Had enough data to score baseline PCS or MCS scores using a previously validated Modified Regression Estimation (MRE) algorithm for missing data as well as adjustments for contextual issues such mode of administration (phone versus mail-out).<sup>17</sup>
3. Were members of a health plan at baseline that remained in HOS at follow-up.
4. Had a baseline survey disposition of M10 (mail, complete survey), M11 (mail, partial complete survey), M31 (mail, break-off), T10 (telephone, complete survey), T11 (telephone, partial complete survey), or T31 (telephone, break-off).

We used the Medicare HOS cohort 9 (April 2006 - May 2008) because on January 1, 2006, CMS implemented the Medicare Part D program. Beneficiaries were randomly sampled from each managed care plan participating in the Medicare Advantage Plans in 2006 for the cohort 9 baseline administration. There were 203 contract/market areas, and 1,000 beneficiaries were randomly sampled from each (if a contract/market area had fewer than 1,000 members, all beneficiaries were included). The total number of beneficiaries sampled was 188,515, including both the aged and disabled (see Figure 1). Among the 188,515 beneficiaries, 111,667 met all 4-study criteria. Of the 111,667 patients, 94,630 had Medicare Part D claims. Out of the 94,630 Medicare Part D patients, 7,148 (7.54%) died by the time of the two-years of follow-up.

## Study Measures

We used two study measures:

1) *Two-year mortality rates*: Building upon prior work,<sup>18</sup> we used mortality as the study outcome. Mortality is a measure that is particularly relevant to elderly patients and might reflect potentially poor quality of care.<sup>19</sup> We used the Medicare HOS mortality files to ascertain the vital status of the MA patients.

2) *Performance indicators*: We assessed how well plans followed nationally recognized guidelines for five high volume diagnoses: diabetes, coronary artery disease (CAD), congestive heart failure (CHF), chronic obstructive lung disease (COPD)/asthma and depression (table 1). These diagnoses were self-reported in the HOS survey. Within each of the five diagnoses, there were one to three drug interventions that were measured. The number of patients receiving the desired drug constituted the numerator. There were two performance indicators for the diagnosis of diabetes: percentage of patient with diabetes treated with ACE inhibitors or angiotensin receptor blockers (ARB) and lipid lowering medications. There were two indicators for the diagnosis of CAD: percentage of patients with CAD treated with beta-blockers and lipid lowering medications. There were three indicators for the diagnosis of CHF: percentage of patients with CHF treated with ACE inhibitors or ARBs, beta-blockers and lipid lowering medications. There was one indicator for the diagnoses of COPD/asthma and depression: percentage of patients with COPD/asthma treated with steroid inhalers and percentage of patients with depression treated with anti-depressants. We used the Medicare Part D prescription benefit file to calculate the performance indications. This file contains information that was collected between January 1<sup>st</sup>, 2006 and December 31<sup>st</sup>, 2007. We did not use National Drug Codes from

the U.S. Food and Drug Administration because this classification scheme has not been updated since 1976.<sup>20</sup> To identify the medications for each of the selected class groups, we used the Tarascon Pocket Pharmacopoeia 2009 Classic Shirt-Pocket Edition 23rd edition.

### **Analytic Plan**

First, we calculated the prevalence of the performance indicators for each plan (objective 1). Within each of the five diagnoses (diabetes, CAD, CHF, COPD / asthma and depression), we measured one to three drug interventions. The number of patients receiving the desired drug constituted the numerator. We used analysis of variance to examine the variations of the performance indicators across plans.

Second, we used correlation analyses at the patient and the plan levels to examine the association of performance indicators and mortality (objective 2). We expected negative correlations between the performance indicator prevalence estimates (proportion of those with an indicated medication given a specific diagnosis according to the accepted guidelines) and mortality because the higher the number of patients receiving the desired drug, the lower the mortality. For the analysis we used the ratio of observed/expected mortality rates as the study outcome. In calculating the expected mortality rates, we used 3 steps:

*Step 1:* we calculated propensity scores, defined as the probability of a patient on a drug or not given a set of known predictor variables, to build treatments groups that are similar with respect to outcome risk factors.<sup>21</sup> We applied a logistic regression model, in which the log of the “odds” of the event is modeled as a linear function of the predictor

variables. We used the following predictor variables: age, gender (male/female), race/ethnicity (whites/African-American/Hispanics/others), married (yes/no), less than high school education (yes/no), homeowner (yes/no), body mass index, baseline physical functioning, baseline mental functioning, activities of daily living (ADL), presence of medical/psychiatric conditions (diabetes, hypertension, stroke, COPD/asthma, cancer, arthritis of the hip, arthritis of the hand, low back pain, congestive heart disease, osteoporosis, sciatica, depression), current smoker (yes/no). We included only the significant predictor variables in the model in order to generate the propensity scores.

*Step 2:* We used the propensity scores to generate weights using the approach modeled after Horwitz and Thompson.<sup>22</sup> Assuming that there is no unmeasured confounding, we obtained unbiased estimates of the probability of being on treatment by comparing treatment groups within each of the levels of the propensity score. The weights were defined as the (Total number of patients on a drug/10)/ number of patients on a drug in one decile and the (Total number of patients not on a drug/10)/ number of patients not on a drug. Each individual was assigned with one of the two weights in each decile.

*Step 3:* We applied the weights from step 2 as a probability weight to a logistic model (probability weighted regression model) in order to generate the expected mortality rates. In this model, mortality was the dependent variable and being on a drug (yes/no) was the independent variable.

## Results

Our study focused on 94,630 MA patients with Medicare Part D claims (table 2). When compared with MA patients without Medicare Part D claims (N=17,037), MA patients with Medicare Part D claims were more likely to be female, non-White, non-married, with less than a high school education and with a lower income. The MA patients with Medicare Part D claims had higher prevalence of chronic conditions when compared to those without Medicare Part D claims, including diabetes (23.0% vs. 18.1%,  $p<0.0001$ ), hypertension (65.9% vs. 55.1%,  $p<0.0001$ ), coronary artery disease (15.7% vs. 14.6%,  $p=0.0003$ ), congestive heart failure (9.3% vs. 7.9%,  $p<0.0001$ ), COPD/asthma (14.4% vs. 11.8%,  $p<0.0001$ ) and depression (28.6% vs. 23.0%,  $p<0.0001$ ). The MA patients with Medicare Part D claims had lower physical health (39.0 (SD  $\pm$  12) vs. 40.8 (SD  $\pm$  11),  $p<0.0001$ ) and mental health (51.6 (SD  $\pm$  11) vs. 53.1 (SD  $\pm$  10),  $p<0.0001$ ) than those without Medicare Part D claims on the basis of the summary scores from the Veterans RAND 12 Item Health Survey (VR-12). However, there was no difference in the 2-year mortality rates between patients with or without Medicare Part D claims (7.55% vs. 7.54%, respectively).

The Medicare Part D file contained 5,943,875 prescriptions and 16,712 (11-digit) National Drug Codes. The mean number of prescriptions for patients for each drug code was 3.22 over the 2-year follow-up period with a minimum of 1 and a maximum of 38 prescriptions.

Analysis of variance confirmed that the 203 plans differed significantly in their performance indicators ( $P < 0.001$ ) (Appendix A). The indicator prevalence estimates for diabetic patients ranged from 52.2% to 100% for ACE inhibitors/ARBs and 33.3% to 90.0% for lipid lowering medications. For CAD patients estimates ranged from 42.4% to 100% for beta-blockers and 51.8% to 100% for lipid lowering medications. Estimates for CHF patients ranged from 38.5% to 86.7% for ACE inhibitors/ARBs, from 33.3% to 100% for lipid lowering medications and from 46.7% to 93.3% for beta-blockers. For COPD/asthma patients, the indicator for inhaled steroids ranged from 21.3% to 71.4%. Last, the indicator for antidepressants for patients with depression ranged from 7.4% to 66.7%.

There were significant correlations between the performance indicators and the observed/expected mortality ratios. At the patient level, there were weak associations, albeit in the hypothesized direction, between lipid-lowering medications and a lower mortality among patients with diabetes ( $r = -0.02$ ,  $p = 0.005$ ) and CAD ( $r = -0.04$ ,  $p < 0.0001$ ). Beta-blockers were also associated with a lower mortality among patients with CHF ( $r = -0.02$ ,  $p = 0.034$ ). At the plan level for those patients with CHF, ACE inhibitors/ARBs ( $r = -0.40$ ,  $p < 0.0001$ ) and beta-blockers ( $r = -0.27$ ,  $p < 0.0001$ ) were found to be significantly associated with a lower mortality. Lipid-lowering medications ( $r = -0.11$ ,  $p = 0.091$ ) were associated with lower mortality but not significant. Antidepressant medications showed a paradoxical correlation. They were associated with a higher mortality in depressed patients at the patient ( $r = 0.02$ ,  $p = 0.007$ ) and at the plan level ( $r = 0.28$ ,  $p < 0.0001$ ).

## Discussion

This study showed a wide variation of nationally recommended use of drugs across MA plans. There were significant correlations between selected use of medications and mortality. We were able to identify performance indicators for conditions such CHF, which were significantly associated with better outcomes (lower mortality) at the plan level. The Medicare Part D prescription benefit file can supplement the quality of medical care information obtained from process of care measures. The fact that we found a few but important correlations between process indicators and mortality highlight the additional contribution of mortality for quality assessments.

Studies have shown that the Medicare Part D has led to increases in drug use and decreases in expenses for older adults.<sup>23</sup> However, large variations in the drug utilization across plans are confirmed by our results. Such variation may in part be a function of the cost and drugs covered by plan. Exact coverage and costs are different for each Medicare drug plan, but all plans must provide at least a standard level of coverage set by Medicare. The actual drug plan costs vary depending on the prescriptions used, whether the drugs are on the plan's formulary, and whether the patient qualifies for extra help paying the Part D costs. Further, studies have found recent increases in premiums.<sup>24</sup> Policymakers continue to question government's role in areas such as negotiating prices directly with pharmaceutical manufacturers and limiting the number of plans offered. The substantial variation in drug utilization reported in this study may be influenced by the complex differences found in the plan design, enrollment, and the impact of the program on beneficiaries.

Adherence to pharmacotherapy for heart failure is poor among older adults in part because of high prescription drug costs. Studies have found that Medicare Part D was associated with improved access to medications and adherence to pharmacotherapy in older adults with heart failure.<sup>25</sup> Those previously lacking drug coverage were more likely to fill prescriptions for an ACE/ARB plus a beta-blocker after Part D (adjusted odds ratio = 1.73, 95% CI 1.42-2.10,  $P < 0.0001$ ) and more likely to be adherent to such pharmacotherapy (adjusted odds ratios = 2.95, 95% CI 1.85-4.69,  $P < 0.0001$ ) relative to the comparison group. This report highlights a few important correlations between process indicators for CHF (ACE inhibitors/ARBs, lipid lowering medications and beta blockers) and mortality noting the additional contribution of these process indicators for quality assessments.

We also found a paradoxical relationship between mortality and the use of antidepressants at the patient and plan levels. Depression imposes important burdens on the elderly. Despite this, rates of initiation of and adherence to recommended pharmacotherapy are frequently low in this population.<sup>26</sup> Among a large Medicare cohort of fee-for-service beneficiaries with diabetes, comorbid depression was associated with an increase in all-cause mortality over a two-year period corroborating our findings in this study.<sup>27</sup> Future research will be required to determine whether the increase in mortality associated with depression is due to potential behavioral mediators (i.e., smoking, poor adherence to diet).



Though our efforts to link mortality with process indicators yielded mixed results, this is not unique to our experience. Studies in which both process and outcomes were evaluated found that they were not always consistent with each other.<sup>28</sup> Studies have found weak relationships between processes of care and overall mortality.<sup>29</sup> Similar results were found in studies of specific conditions. Bradley et al, for example, found that only a few acute myocardial infarction process measures (beta-blocker at admission, aspirin at admission, and smoking cessation) correlated with hospitals' mortality rates.<sup>30</sup>

There are several limitations of this study. First, we found that 15.2% of the HOS patients did not have Part D Medicare claims. We can formulate several possibilities regarding these patients: they are not enrolled in Part D period or they are enrolled but not using meds at all or they have other prescription coverage (e.g., they have a company's retirement insurance covering it). Second, the Medicare Part D claim file does not contain drug class assignment. We compared our drug class algorithm with the HEDIS 2008 National Drug Coding class assignment. We found that both approaches produced similar results (Appendix B). Third, the Medicare Part D data included medication information from January 2006 thru December 2007 (24 months), while HOS cohort 9 data represented a time frame from April 2006 thru May 2008. The HOS data did not overlap with the Medicare data for the first three months in 2006 and the Medicare data did not overlap with the HOS data for the last 4 months in 2008. This lack of overlap provides more potential error in the attempt to establish baseline medication profiles in 2006 with the other clinical baseline measures that include morbidities and physical and mental functioning (VR-12). Future analysis would be improved with better overlap of the Medicare data with the HOS data including baseline measures of physical and mental functional status

using the VR-12. Fourth, we examined mortality over a 2 year period. The efficacious impacts of medications according to the clinical guidelines may take a longer period of time in chronic conditions such as diabetes and CAD. For example, the progression and regression of CAD may take at least 3 years or more to occur.<sup>31</sup> Future studies should examine the impact of process indicators on improved mortality using longer follow-up periods. They can also consider mortality analysis that includes time to event (mortality) as there is greater follow-up time using mortality curves (e.g. Cox Proportional Hazard Models). The time limited nature of the data for this study limited the analysis to expected mortality. As the Medicare Advantage Program has a longer experience it is hoped that more data will become available that spans a greater time frame. Fifth, those patients for whom the drug is contraindicated were not excluded from the denominator; e.g. a CHF patient who is allergic to ACE/ARB. A more accurate count of patients receiving the desired drug can be made if patient allergies are listed in the Medicare Part D file. Sixth, we did not examine the doughnut hole and the effect of changes on clinical outcomes. This issue continues to raise equity in access concerns among elderly patients.<sup>32</sup>

In summary, our results are not definitive, and reveal a complex relationship between outcome and different process indicators. Nevertheless, there is support for the relationship between mortality and processes of care that warrants further investigation.

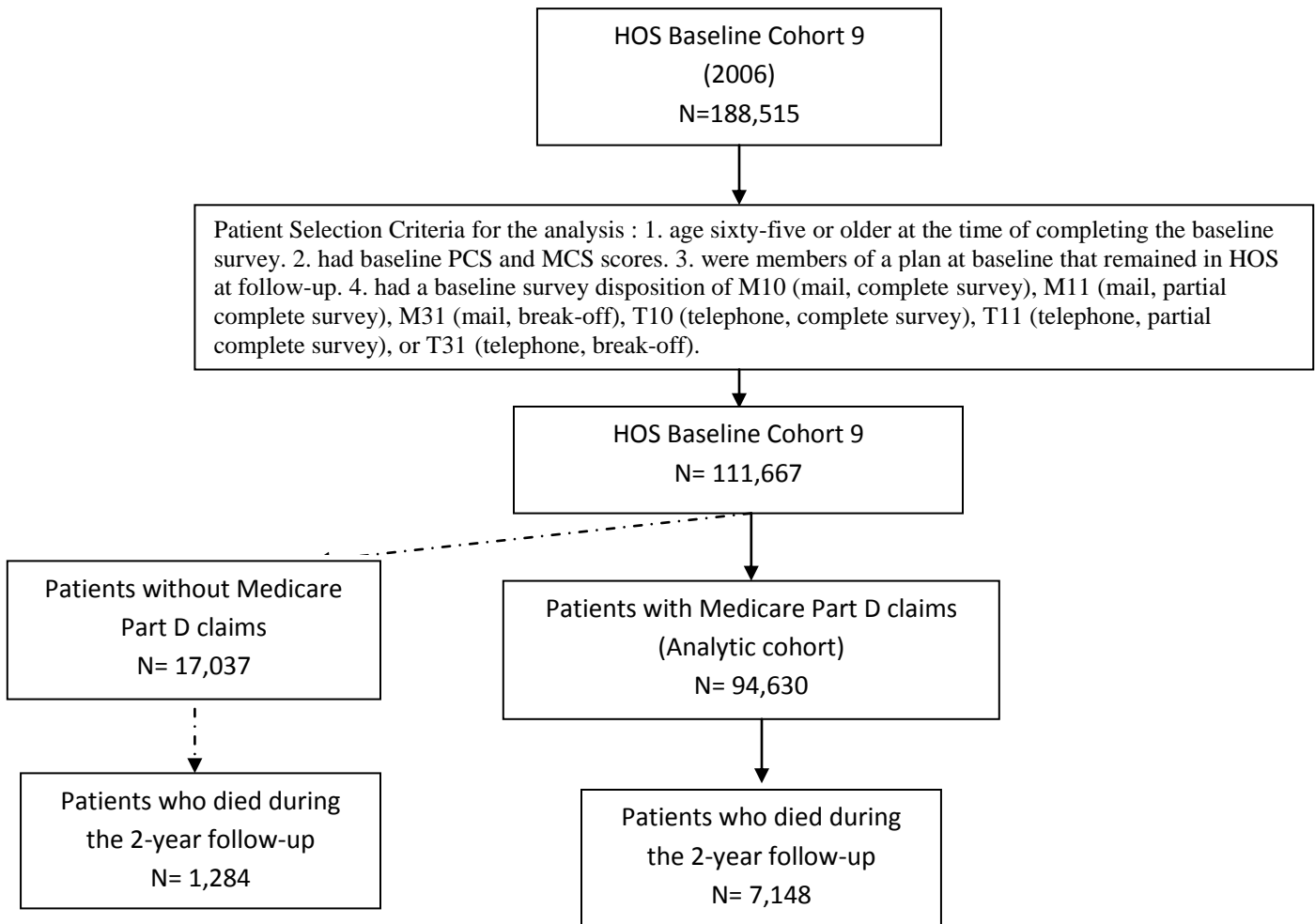
## **Recommendations for CMS**

With the growing number of MA plans, CMS is looking to refine their methods of measuring and improving the performance of the MA plans. Our findings highlight the additional contribution of mortality for quality assessments.

The fact that we found a few but important correlations between process indicators and mortality might indicate that the Medicare Part D claim file was sufficiently robust to find such associations. However, this database can be improved by including: 1) drug class assignments, 2) Part D enrollees without claims, 3) age, sex and ICD-9-DM diagnoses and 4) over the counter medications such as aspirin.

Health care organizations are increasing their scrutiny of the care provided to Medicare enrollees with chronic diseases. Medicare Part D can supplement the quality of medical care information obtained from process of care measures such as the Healthcare Effectiveness Data and Information Set (HEDIS). We look forward to conducting future studies with richer information that tracks the use of medications as indicators of quality of care coupled with HEDIS measures with lengthier follow-up to examine the effect on the survival of Medicare Advantage recipients.

**Figure 1. Medicare Health Outcomes Survey (HOS) Analytic Cohort 9**



**Table 1. Performance Indicators**

<b>Condition</b>	<b>Performance Indicator</b>	<b>Clinical Practice Guideline/Reference</b>
Diabetes	ACE inhibitors/ARB	ADA Standards of Care 2005/DQIP
	Lipid lowering Medication	ADA Standards of Care 2005/DQIP
Coronary Artery Disease	Beta-blockers	ACC/AHA
	Lipid Lowering Medication	ACC/AHA
Congestive Heart Failure	On ACEI/ARB	ACC/AHA
	Beta-blocker	ACC/AHA
	Lipid Lowering Medication	ACC/AHA
COPD	On steroid inhaler	ATS
Depression	On antidepressant <sup>33</sup>	APA

ACC/AHA= American College of Cardiology/American heart Association; ACCP= American College of Clinical Pharmacy; ACE= angiotensin converting enzyme inhibitor; ADA= American Diabetes Association; ARB= angiotensin receptor blocker; DQIP= Diabetes Quality Improvement Project; ATS=American Thoracic Society; APA=American Psychiatric Association

**Table2. Patient Characteristics of Medicare Advantage (MA) Patients with and without Medicare Part D claims from the Medicare Health Outcomes Survey cohort 9**

	<b>MA patients with Medicare Part D claims (n=94,630)</b>	<b>MA patients without Medicare Part D claims (n=17,037)</b>	<b>p-value</b>
Age	75.4 (+6)	75.1 (+6)	NS
Female	60.9%	49.7%	<.0001
White	85.4%	90.1%	<.0001
African American	9.1%	7.2%	
Hispanics	1.9%	0.6%	
Others	3.7%	2.1%	
Married	54.4%	60.0%	<.0001
Less than high school education	27.8%	24.6%	<.0001
Income			<.0001
Less than \$5,000	3.8%	2.6%	
\$5,000-\$9,999	8.5%	6.0%	
\$10,000-\$19,999	26.4%	24.9%	
\$20,000-29,999	11.4%	21.2%	
\$30,000-39,999	19.3%	13.7%	
\$40,000-49,999	7.0%	8.2%	
\$50,000-79,999	7.6%	8.1%	
\$80,000-99,999	1.9%	2.0%	
\$100,000 or more	2.4%	1.9%	
Don't know	11.7%	11.2%	
Medicaid eligibility	7.3%	2.5%	<.0001
Diabetes	23.0%	18.1%	<.0001
Hypertension	65.9%	55.1%	<.0001
Coronary Artery Disease	15.7%	14.6%	0.0003
Congestive Heart Failure	9.3%	7.9%	<.0001
COPD/asthma	14.4%	11.8%	<.0001
Depression	28.5%	23.0%	<.0001
Baseline Physical Health (Mean $\pm$ SD)*	39.0 ( $\pm$ 12)	40.8 ( $\pm$ 11)	<.0001
Baseline Mental Health (Mean $\pm$ SD)*	51.6 ( $\pm$ 11)	53.1 ( $\pm$ 10)	<.0001
Mortality rate, 2-year	7.55%	7.54%	NS

\*Physical and mental summary scores standardized to a U.S. population with a norm of 50.0. Higher scores denote better health.

**Table3. Correlations between Performance Indicators and Mortality**

<b>Condition</b>	<b>Performance Indicator</b>	<b>At the patient</b>	<b>At the plan level</b>
Diabetes	ACE inhibitor/ARB	-0.01 (p=0.190) (n=17,608)	-0.07 (p=0.294) (n=203)
	Lipid lowering medication	<b>-0.02</b> <b>(p=0.005)</b> <b>(n=17,653)</b>	-0.09 (p=0.199) (n=203)
CAD	Lipid lowering medication	<b>-0.04</b> <b>(p&lt;.0001)</b> <b>(n=12,155)</b>	0.06 (p=0.352) (n=203)
	Beta blockers	0.01 (p=0.401) (n=11,937)	-0.07 (p=0.257) (n=203)
CHF	ACE/ARB	<b>-0.001</b> <b>(p=0.006)</b> <b>(n=7,157)</b>	<b>-0.40</b> <b>(&lt;.0001)</b> <b>(n=203)</b>
	Lipid lowering medications	<b>-0.061</b> <b>(p&lt;.0001)</b> <b>(n=6,936)</b>	<b>-0.11</b> <b>(p=0.091)</b> <b>(n=203)</b>
	Beta blockers	<b>-0.02</b> <b>(p=0.034)</b> <b>(n=6,835)</b>	<b>-0.27</b> <b>(p&lt;.0001)</b> <b>(n=203)</b>
COPD	Inhaled steroids	-0.01 (p=0.350) (n=10,024)	0.03 (p=0.608) (n=203)
Depression	Antidepressants	<b>0.02</b> <b>(p=0.007)</b> <b>(n=19,991)</b>	<b>0.28</b> <b>(p&lt;0.0001)</b> <b>(n=203)</b>

Appendix A: Variation of Performance Indicators by Medicare Advantage Plans

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
<b>Overall</b>	<b>70.35</b>	<b>69.05</b>	<b>72.26</b>	<b>66.94</b>	<b>68.03</b>	<b>62.38</b>	<b>68.30</b>	<b>43.32</b>	<b>39.09</b>
H0150	71.32	68.38	79.17	72.22	67.24	56.90	65.52	44.16	38.33
H0151	65.83	64.17	58.11	66.22	68.09	59.57	57.45	31.65	34.21
H0154	76.69	67.67	62.82	71.79	74.55	56.36	69.09	39.19	51.74
H0302	76.52	68.70	70.79	69.66	54.55	52.27	68.18	46.39	41.72
H0303	71.03	67.29	60.81	66.22	76.09	45.65	67.39	40.00	45.97
H0307	73.00	65.00	78.95	67.11	74.00	72.00	74.00	47.62	38.67
H0316	74.62	66.15	70.83	53.13	57.45	48.94	57.45	41.18	42.46
H0317	66.67	33.33	66.67	66.67		100.00			54.55
H0351	72.00	68.00	59.09	62.50	67.92	43.40	60.38	38.96	38.13
H0354	74.19	68.82	67.65	67.65	66.67	69.23	79.49	47.22	42.48
H0502	83.64	73.64	69.74	75.00	84.00	52.00	66.00	57.14	53.29
H0504	75.44	67.54	73.86	65.91	70.18	70.18	71.93	52.24	37.80
H0523	68.32	55.45	70.89	63.29	71.43	59.18	61.22	27.71	33.33
H0524	77.39	88.70	78.46	76.92	69.23	69.23	73.08	71.43	46.97
H0532	70.48	69.52	80.77	70.51	77.50	67.50	70.00	48.60	48.61
H0543	69.47	68.42	78.13	65.63	75.61	58.54	70.73	49.18	35.04
H0544	82.47	77.27	85.25	73.77	68.09	70.21	68.09	44.12	34.97
H0545	66.67	71.32	71.26	66.67	57.14	55.36	60.71	38.36	33.72
H0562	72.73	77.27	77.94	64.71	76.32	57.89	50.00	47.69	31.34
H0564	69.31	60.40	75.44	70.18	70.59	67.65	79.41	30.67	31.58
H0571	64.24	70.30	72.34	46.81	77.78	66.67	77.78	31.25	13.50
H0602	75.36	66.67	68.85	55.74	66.67	44.44	55.56	43.86	45.83
H0609	65.05	60.19	71.21	65.15	66.67	58.82	54.90	49.40	37.70
H0630	71.72	73.74	83.08	76.92	78.95	76.32	78.95	49.43	48.91
H0657	56.34	50.70	76.00	52.00	56.67	46.67	46.67	42.00	44.44



	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H0752	66.67	69.70	79.37	66.67	51.52	63.64	75.76	44.44	32.28
H0755	64.91	74.56	70.73	70.73	51.22	60.98	70.73	44.29	34.75
H1013	80.58	74.76	82.35	72.55	81.82	72.73	63.64	58.54	29.09
H1016	65.00	70.00	75.58	61.63	78.72	78.72	70.21	42.86	38.89
H1019	67.90	76.54	74.00	58.00	56.52	65.22	65.22	52.08	33.09
H1026	73.08	75.64	62.30	62.30	85.71	71.43	90.48	36.36	40.00
H1032	67.24	69.83	68.49	68.49	72.09	58.14	74.42	39.39	39.74
H1034	73.91	78.99	80.00	79.05	74.58	72.88	76.27	47.17	45.03
H1035	76.32	74.56	73.81	61.90	58.18	67.27	58.18	27.85	37.41
H1036	69.44	75.00	73.91	60.87	80.56	72.22	66.67	38.18	37.69
H1045	75.00	75.00	79.49	84.62	75.00	85.71	92.86	32.14	35.11
H1076	70.24	66.67	82.61	76.09	80.77	73.08	69.23	31.91	41.91
H1080	67.62	57.14	71.43	65.71	64.00	48.00	76.00	35.48	34.29
H1099	72.88	74.58	70.45	64.77	60.00	65.00	52.50	46.58	40.97
H1170	73.28	75.86	86.57	82.09	67.44	72.09	74.42	44.29	37.17
H1230	78.57	81.82	80.65	72.58	82.86	68.57	74.29	62.50	24.64
H1251	71.43	68.91	69.57	50.00	80.00	64.00	60.00	50.00	27.35
H1349	72.83	59.78	64.00	60.00	65.67	53.73	62.69	42.31	47.29
H1350	67.74	61.29	80.36	55.36	67.57	62.16	62.16	45.83	48.51
H1406	72.50	71.67	73.68	73.68	78.85	69.23	76.92	41.38	23.38
H1415	79.02	65.03	55.56	83.33	74.36	56.41	66.67	45.45	17.81
H1463	79.38	68.04	77.14	68.57	66.04	60.38	69.81	41.89	45.30
H1468	68.85	68.03	69.23	64.84	62.12	62.12	69.70	32.95	36.13
H1472	75.00	75.00	100.00	50.00				50.00	57.14
H1553	68.75	75.89	76.67	71.11	72.50	57.50	70.00	35.29	47.10
H1555	72.03	61.86	72.94	69.41	68.42	50.88	68.42	35.37	51.03
H1558	72.88	61.02	67.27	74.55	73.81	57.14	73.81	44.23	49.54
H1651	68.33	70.83	75.00	82.81	58.93	58.93	69.64	50.00	42.06
H1716	71.43	66.07	84.62	63.46	83.87	74.19	74.19	41.82	51.47
H1806	64.29	60.71	66.67	66.67	58.33	33.33	58.33	58.33	55.00

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H1849	70.07	68.61	68.27	64.42	66.67	59.65	70.18	34.18	43.11
H1903	67.11	67.11	62.67	65.33	77.36	56.60	77.36	25.00	31.40
H2108	76.27	63.56	72.13	68.85	73.21	69.64	75.00	58.21	29.70
H2150	83.97	86.54	83.51	72.16	84.62	79.49	79.49	66.67	39.73
H2204	70.77	70.00	76.19	75.00	61.90	59.52	73.81	41.77	44.07
H2206	61.79	73.98	82.50	73.75	58.33	80.56	80.56	54.02	44.20
H2256	66.10	71.19	78.48	73.42	72.09	72.09	74.42	52.44	49.65
H2261	63.85	71.54	76.60	67.02	51.92	65.38	69.23	49.44	38.82
H2312	74.04	74.04	73.49	78.31	74.14	63.79	84.48	55.06	33.11
H2354	72.22	50.00	51.85	70.37	45.00	45.00	65.00	35.00	32.26
H2450	78.90	79.82	77.45	79.41	74.55	67.27	76.36	43.48	50.34
H2459	67.68	63.64	79.17	73.61	56.67	50.00	73.33	38.46	45.80
H2461	78.45	72.41	73.08	73.08	73.33	55.56	73.33	46.43	46.79
H2462	70.34	73.73	81.00	78.00	65.08	68.25	82.54	47.76	51.72
H2610	64.34	75.19	75.00	61.96	71.43	61.90	66.67	35.62	42.33
H2611	100.00	85.71	100.00	100.00	60.00	60.00	80.00	71.43	66.67
H2649	68.60	70.25	67.50	62.50	60.66	55.74	65.57	41.38	36.36
H2654	64.84	60.94	60.98	62.20	65.91	52.27	65.91	36.14	39.60
H2663	64.80	64.00	68.89	74.44	67.92	60.38	75.47	40.96	45.08
H2667	68.04	62.89	62.07	65.52	63.33	53.33	56.67	36.67	53.44
H2672	65.63	62.50	67.47	63.86	64.81	57.41	62.96	40.00	50.00
H2802	65.04	60.16	63.83	71.28	61.22	65.31	77.55	40.66	38.13
H2803	63.64	68.60	72.58	72.58	70.59	61.76	70.59	29.85	42.96
H2931	75.44	64.04	65.14	66.06	71.88	64.06	78.13	30.51	31.25
H2949	67.35	66.33	76.56	64.06	57.78	62.22	66.67	43.84	35.56
H2960	75.38	65.38	82.56	54.65	74.51	60.78	60.78	49.49	41.42
H2961	65.52	58.62	65.15	57.58	76.00	58.00	60.00	35.35	31.36
H3107	61.86	64.41	73.77	75.41	65.79	73.68	78.95	43.14	24.63
H3112	70.00	90.00	76.92	84.62	81.82	81.82	81.82	52.63	19.23
H3152	57.98	60.50	57.14	64.94	67.57	62.16	78.38	58.67	35.09

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H3154	68.81	61.47	63.24	70.59	61.54	61.54	56.41	37.70	41.22
H3156	72.41	82.76	72.73	77.27	52.17	60.87	86.96	60.00	40.98
H3164	78.99	74.79	86.00	74.00	82.98	70.21	59.57	50.88	24.28
H3204	66.00	69.00	72.60	53.42	58.82	62.75	58.82	40.91	50.63
H3251	68.38	66.91	73.61	63.89	68.18	52.27	59.09	58.82	46.81
H3305	66.04	66.04	85.37	73.17	86.67	66.67	93.33	50.00	39.47
H3307	77.59	63.79	73.85	72.31	61.90	76.19	85.71	39.02	29.36
H3312	64.04	64.04	74.42	72.09	48.57	65.71	74.29	55.56	29.13
H3327	57.14	71.43	90.00	70.00	66.67	75.00	83.33	66.67	33.33
H3328	60.00	86.67	77.78	88.89	85.71	71.43	71.43	50.00	29.41
H3330	79.69	73.44	75.00	83.33	77.42	58.06	74.19	55.00	23.31
H3335	57.78	62.22	66.20	60.56	57.58	45.45	66.67	49.32	41.60
H3336	89.61	79.22	86.67	60.00	85.71	76.19	76.19	45.16	11.83
H3351	74.19	72.58	84.21	55.26	42.31	69.23	65.38	42.31	40.00
H3359	88.43	79.34	82.00	66.00	82.93	73.17	73.17	61.02	15.71
H3361	71.60	66.67	58.54	58.54	66.67	51.52	57.58	34.21	27.66
H3362	68.69	72.73	76.56	70.31	66.67	53.33	73.33	39.29	41.18
H3366	69.07	73.20	67.12	60.27	59.46	72.97	70.27	48.48	34.93
H3370	71.55	75.86	79.52	69.88	75.76	63.64	75.76	50.82	29.50
H3379	70.00	63.33	63.16	65.79	86.67	80.00	66.67	46.15	36.36
H3384	57.14	67.86	70.21	63.83	52.17	56.52	69.57	43.48	40.74
H3387	73.27	57.43	58.70	69.57	60.00	57.50	72.50	48.72	7.38
H3388	56.76	60.81	79.63	79.63	66.67	66.67	70.83	47.17	48.86
H3449	69.09	67.27	77.65	70.59	73.81	69.05	78.57	44.26	47.37
H3456	74.13	69.93	66.67	70.67	69.05	66.67	76.19	39.73	34.97
H3503	74.07	51.85	72.88	79.66	75.68	54.05	72.97	42.42	42.22
H3619	72.45	77.55	78.57	60.71	71.11	66.67	62.22	43.28	49.21
H3653	76.56	64.06	62.64	75.82	73.21	62.50	71.43	43.68	36.97
H3655	58.52	61.48	72.41	66.67	53.19	74.47	68.09	33.72	39.60
H3657	78.49	65.59	74.32	75.68	71.05	60.53	73.68	43.14	28.18

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H3659	70.63	60.32	65.48	61.90	73.91	54.35	73.91	37.14	32.14
H3660	69.23	68.38	72.90	60.75	60.87	50.00	58.70	41.94	45.27
H3664	66.67	67.41	62.00	69.00	58.33	50.00	63.89	47.62	47.75
H3668	68.84	67.39	69.44	67.59	80.00	71.67	66.67	39.80	44.63
H3672	63.03	63.03	61.40	56.14	67.57	56.76	54.05	48.39	36.36
H3706	54.76	64.29	71.79	73.08	56.16	68.49	63.01	35.58	40.63
H3749	61.61	64.29	62.50	54.69	67.19	59.38	68.75	46.34	41.18
H3755	76.47	58.82	64.47	68.42	70.59	50.98	70.59	27.85	42.01
H3805	82.02	77.53	75.00	72.22	76.56	67.19	68.75	40.28	38.46
H3851	70.19	72.12	76.25	60.00	63.83	51.06	61.70	50.00	45.38
H3856	75.51	72.45	73.02	68.25	58.54	48.78	65.85	34.38	54.40
H3864	65.43	67.90	80.00	52.31	81.25	59.38	59.38	55.38	44.83
H3907	61.22	72.45	65.85	62.20	65.71	68.57	74.29	38.71	43.70
H3909	67.96	76.70	75.34	73.97	81.08	67.57	72.97	40.43	47.96
H3916	68.07	68.91	80.00	61.25	67.74	58.06	61.29	47.89	41.59
H3931	70.19	75.00	79.07	73.26	76.19	73.81	83.33	50.00	34.85
H3949	69.68	65.16	68.92	66.22	69.57	62.32	66.67	40.63	27.81
H3952	64.18	73.13	78.57	80.36	71.43	71.43	75.00	35.71	36.17
H3954	62.75	64.71	71.25	73.75	46.81	59.57	82.98	43.64	37.90
H3957	66.67	60.12	72.81	71.93	61.40	71.93	66.67	43.21	42.08
H3959	70.16	61.29	69.90	70.87	61.11	59.26	68.52	48.89	32.57
H3962	66.67	65.97	79.05	62.86	65.57	67.21	62.30	51.85	39.07
H3964	73.83	71.03	77.50	72.50	78.38	81.08	67.57	49.23	26.58
H4003	75.00	63.33	61.90	52.38	70.69	58.62	65.52	23.64	18.56
H4004	70.65	66.17	61.59	42.38	75.00	67.24	49.14	32.53	32.50
H4005	68.83	72.08	65.31	46.94	70.24	63.10	50.00	36.07	30.77
H4006	70.45	69.32	54.93	50.70	71.05	63.16	68.42	32.08	24.34
H4102	69.11	60.16	68.75	77.08	70.00	68.00	86.00	33.96	42.70
H4152	64.29	62.24	68.42	66.32	85.00	62.50	75.00	46.75	44.44
H4406	73.79	66.21	65.57	62.30	74.51	56.86	47.06	29.73	38.29

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H4454	78.57	74.29	69.57	63.77	78.57	69.05	78.57	38.36	47.62
H4456	73.11	63.87	75.27	63.44	73.47	59.18	73.47	38.82	46.15
H4461	68.14	76.11	77.78	60.61	79.07	69.77	67.44	37.66	45.34
H4506	72.88	71.19	65.79	63.16	78.05	63.41	60.98	43.42	34.94
H4510	75.63	73.11	73.61	62.50	58.33	56.25	62.50	48.78	32.37
H4513	73.11	62.18	62.50	60.94	63.64	63.64	70.45	32.73	35.92
H4520	73.56	65.52	82.00	62.00	58.62	68.97	62.07	42.37	47.12
H4564	80.23	77.91	73.42	69.62	66.67	64.71	72.55	48.21	52.00
H4590	71.43	70.63	74.63	67.16	70.37	81.48	55.56	47.95	34.13
H4604	63.20	64.80	74.68	43.04	66.04	67.92	49.06	40.74	46.20
H5005	68.75	71.88	76.27	57.63	62.00	60.00	56.00	43.08	43.61
H5050	78.43	72.55	82.76	75.86	62.50	45.83	79.17	50.00	50.00
H5102	67.54	60.53	58.10	68.57	70.89	49.37	72.15	21.33	43.57
H5151	63.89	60.42	66.67	72.50	64.21	56.84	67.37	30.49	40.68
H5211	65.79	68.42	80.00	80.00	57.14	66.67	76.19	52.38	44.19
H5215	52.24	59.70	65.22	60.87	70.83	62.50	58.33	46.94	27.85
H5253	69.44	69.44	65.57	59.02	61.54	58.97	66.67	29.33	23.73
H5254	65.48	60.71	68.25	66.67	62.50	53.13	68.75	41.82	41.25
H5256	63.16	56.14	60.47	72.09	64.86	45.95	59.46	32.50	51.28
H5262	76.79	75.00	70.59	86.27	81.25	65.63	90.63	51.11	47.56
H5264	59.49	58.23	76.92	58.97	53.66	58.54	58.54	50.00	58.33
H5402	71.55	75.86	71.62	55.41	60.00	56.67	63.33	30.77	43.36
H5404	77.05	77.05	75.00	54.55	60.47	67.44	58.14	44.78	32.03
H5407	86.96	80.43	71.43	66.67	64.29	71.43	57.14	37.50	32.69
H5411	66.67	84.06	80.56	69.44	66.67	66.67	70.83	66.67	38.78
H5415	71.43	80.36	75.00	61.36	56.52	65.22	60.87	34.38	37.31
H5416	76.12	65.67	64.71	50.00	57.14	50.00	64.29	33.33	46.15
H5500	69.05	64.29	83.33	73.81	71.43	71.43	61.90	34.15	50.00
H5506	71.15	67.31	69.23	66.67	43.48	52.17	60.87	50.00	56.00
H5507	70.59	62.50	71.03	70.09	73.33	53.33	61.67	50.54	47.89

	Diabetes (N=21,059)		CAD (N=14,176)		CHF (N=8,464)			COPD (13,126)	Depression (N=26,096)
	ACE inhibitor/ARB	Lipid-lowering medications	Lipid-lowering medications	Beta-Blockers	ACE inhibitor/ARB	Lipid-lowering medications	Beta-Blockers	Inhaled Steroids	Antidepressants
H5508	70.54	70.54	78.95	54.74	62.75	70.59	64.71	37.33	43.75
H5509	71.08	69.88	66.97	57.80	64.10	66.67	76.92	38.54	44.00
H5510	68.21	75.50	84.92	76.19	70.69	75.86	74.14	46.91	44.14
H5512	71.19	84.75	70.59	68.63	70.00	65.00	80.00	51.85	47.44
H5515	76.47	67.65	73.58	66.04	78.79	57.58	72.73	54.17	41.05
H5516	61.94	75.37	80.85	61.70	63.64	70.45	79.55	43.82	43.04
H5517	61.62	58.92	59.03	60.42	62.79	50.00	67.44	25.00	41.67
H5521	69.85	72.06	83.53	71.76	71.05	68.42	76.32	44.05	45.52
H5524	75.68	67.57	70.91	63.64	58.62	62.07	62.07	48.72	52.11
H5525	65.63	68.75	69.62	74.68	71.93	56.14	77.19	35.00	46.27
H5526	62.50	75.00	74.63	70.15	51.85	62.96	62.96	42.86	33.78
H5527	69.23	65.38	60.00	64.00	63.33	63.33	66.67	37.93	46.84
H5528	61.11	62.04	66.22	64.86	48.00	56.00	52.00	55.10	18.75
H5529	61.46	70.83	75.95	63.29	73.53	79.41	70.59	40.28	50.00
H5530	92.00	84.00	72.22	55.56	66.67	55.56	66.67	63.64	57.14
H5531	74.07	66.67	70.83	58.33	42.86	42.86	57.14	68.18	39.53
H5532	67.26	66.37	75.00	67.31	52.00	64.00	66.00	45.83	49.28
H5533	62.34	66.23	80.36	71.43	38.46	53.85	80.77	44.68	57.75
H6360	83.84	82.83	80.56	72.22	71.11	77.78	80.00	40.35	33.33
H9001	61.68	79.44	74.12	72.94	68.57	65.71	74.29	38.81	43.94
H9003	75.28	80.90	75.00	79.41	66.67	66.67	63.64	44.87	52.63
H9005	61.80	66.29	71.43	77.14	71.19	54.24	69.49	53.03	50.00
H9011	70.51	70.51	86.67	76.67	85.71	71.43	85.71	51.61	28.21
H9016	74.19	74.19	65.67	59.70	69.44	66.67	61.11	44.83	28.68
H9047	67.50	72.50	82.67	73.33	79.17	64.58	66.67	53.52	40.85
H9101	76.35	75.68	76.06	83.10	71.88	68.75	78.13	56.67	31.64
H9103	82.91	84.18	83.33	79.17	72.55	70.59	77.45	63.72	60.92
H9104	76.05	69.46	74.67	62.67	73.33	53.33	63.33	48.24	41.71

ACEI= angiotensin converting enzyme inhibitor; ARB= angiotensin receptor blocker

Appendix B. Comparison between our drug class classification and the HEDIS 2008 National Drug Coding class assignment

<b>Condition</b>	<b>Indicator</b>	<b>Our drug class classification</b>	<b>HEDIS 2008 National Drug Coding class assignment</b>
Diabetes (N=21,059)	ACEIARB	14,814 (70.35%)	14,803 (70.29%)
CAD (N=14,176)	ACEIARB	8,731 (61.59%)	8,726 (61.55%)
	BetaBlockers	9,490 (66.94%)	9,416 (66.42%)
CHF (N=8,464)	ACEIARB	5,758 (68.03%)	5,750 (67.93%)
	BetaBlockers	5,781 (68.3%)	5,776 (68.24%)
COPD (N=13,126)	InhaledCort	5,686 (43.32%)	4,857 (37%)
Depr (N=26,096)	Antidepressants	10,200 (39.09%)	10,187 (39.04%)

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- <sup>1</sup> Centers for Disease Control and Prevention. Comorbid conditions by age, sex, and race. United States, 1981-2001. Available at: <http://209.217.72.34/aging/TableViewer/tableView.aspx?ReportId=185>. Accessed September 7, 2008.
- <sup>2</sup> Gandara E, Moniz TT, Dolan ML, Melia C, Dudley J, Smith A, Kachalia A. Improving adherence to treatment guidelines: a blueprint. *Crit Pathw Cardiol*. 2009;8:139-45.
- <sup>3</sup> Doherty RB. Assessing the new Medicare prescription drug law. *Ann Intern Med*. 2004;141:391-5.
- <sup>4</sup> Kaiser Family Foundation. Medicare Prescription Drug Coverage among Medicare Beneficiaries. Publication 7453. Washington, DC: Kaiser Family Foundation; 2006.
- <sup>5</sup> Bach PB, McClellan MB. The first months of the prescription-drug benefit—a CMS update. *N Engl J Med*. 2006;354:2312-4.
- <sup>6</sup> Kaiser Family Foundation. The Medicare Rx Law: Estimates of Medicare Beneficiaries Out-of-Pocket Drug Spending in 2006. Modeling the impact of the MMA. Publication 7201. Washington, DC: Kaiser Family Foundation; 2006.
- <sup>7</sup> Lichtenberg FR, Sun SX. The impact of Medicare Part D on prescription drug use by the elderly. *Health Aff (Millwood)*. 2007;26:1735-44.
- <sup>8</sup> Hobbs FD. Management of heart failure: evidence versus practice. Does current prescribing provide optimal treatment for heart failure patients? *Br J Gen Pract*. 2000;50:735-42.
- <sup>9</sup> Shepherd J, Cobbe SM, Ford I, et al. Prevention of coronary heart disease with pravastatin in men with hypercholesterolemia. West of Scotland Coronary Prevention Study Group. *N Engl J Med*. 1995;333:1301-7.
- <sup>10</sup> Hennekens CH, Dyken ML, Fuster V. Aspirin as a therapeutic agent in cardiovascular disease: a statement for healthcare professionals from the American Heart Association. *Circulation*. 1997;96:2751-3.
- <sup>11</sup> Sin DD, Tu JV. Inhaled corticosteroids and the risk of mortality and readmission in elderly patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2001 Aug 15;164(4):580-4.
- <sup>12</sup> Dwight-Johnson M, Lagomasino IT, Simpson GM. Underuse of evidence-based pharmacotherapies for affective disorders. *Psychiatr Serv*. 2003;54:1076-8.
- <sup>13</sup> Pulignano G, Del Sindaco D, Tavazzi L, et al. Clinical features and outcomes of elderly



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outpatients with heart failure followed up in hospital cardiology units: data from a large nationwide cardiology database (IN-CHF Registry). *Am Heart J.* 2002;143(1):45-55.

<sup>14</sup> Califf RM, DeLong ER, Ostbye T, et al. Underuse of aspirin in a referral population with documented coronary artery disease. *Am J Cardiol.* 2002;89:653-61.

<sup>15</sup> Horwitz RI, Viscoli CM, Clemens JD, Sadock RT. Developing improved observational methods for evaluating therapeutic effectiveness. *Am J Med* 1990;89:630-638.

<sup>16</sup> Wahid ST, Baines LA, Savopoulos L, et al. Longitudinal analysis of blood pressure, lipid, and glycemic control in diabetic patients with nephropathy attending a hospital outpatient clinic and their relationship to renal function, mortality, and cardiovascular morbidity. *Diabetes Care.* 2001;24:789-90.

<sup>17</sup> Imputing the Physical and Mental Summary Scores (PCS and MCS) for the MOS SF-36 and the Veterans SF-36 Health Survey in the presence of Missing Data.  
[http://www.hosonline.org/surveys/hos/download/HOS\\_Veterans\\_36\\_Imputation.pdf](http://www.hosonline.org/surveys/hos/download/HOS_Veterans_36_Imputation.pdf). Accessed July 30, 2007.

<sup>18</sup> Selim AJ, Kazis LE, Rogers W, Qian S, Rothendler JA, Lee A, Ren XS, Haffer SC, Mardon R, Miller D, Spiro A 3rd, Selim BJ, Fincke BG.. Risk-adjusted mortality as an indicator of outcomes: comparison of the Medicare Advantage Program with the Veterans' Health Administration. *Med Care.* 2006;44:359-65.

<sup>19</sup> Reason J. Human error: models and management. *BMJ.* 2000;320: 768–770.

<sup>20</sup> [National Drug Code Directory.](http://www.fda.gov/Drugs/InformationOnDrugs/ucm142438.htm)  
<http://www.fda.gov/Drugs/InformationOnDrugs/ucm142438.htm>. Accessed July 24, 2010.

<sup>21</sup> Rosenbaum PR, Rubin DB: The central role of the propensity score in observational studies for causal effects. *Biometrika.* 1983,70:41-57.

<sup>22</sup> Horvitz DG, Thompson DJ. A generalization of sampling without replacement from a finite universe. *J of Am Stat Assoc.* 1952;47:663-685.

<sup>23</sup> Yin W, Basu A, Zhang JX, Rabbani A, Meltzer DO, Alexander GC. The effect of the Medicare Part D prescription benefit on drug utilization and expenditures. *Ann Intern Med.* 2008;148:169-77.

<sup>24</sup> Hoadley J, Simon K. Medicare Part D turns four: trends in plan design, enrollment, and the impact of the program on beneficiaries. *Adv Health Econ Health Serv Res.* 2010;22:123-47.

<sup>25</sup> Donohue JM, Zhang Y, Lave JR, Gellad WF, Men A, Perera S, Hanlon JT. The Medicare drug benefit (Part D) and treatment of heart failure in older adults. *Am Heart J.* 2010;160:159-65.

- 
- <sup>26</sup> Wiley-Exley E, Wang PS, Patrick AR, Dormuth C, Maclure M, Avorn J, Canning CF, Schneeweiss S. Impact of drug cost sharing on service use and adverse clinical outcomes in elderly receiving antidepressants. *J Ment Health Policy Econ.* 2010;13:37-44.
- <sup>27</sup> Katon W, Fan MY, Unützer J, Taylor J, Pincus H, Schoenbaum M. Depression and diabetes: a potentially lethal combination. *J Gen Intern Med.* 2008;23:1571-5.
- <sup>28</sup> Kupersmith J. Quality of care in teaching hospitals: a literature review. *Acad Med.* 2005;80:458-66.
- <sup>29</sup> Dubois RW, Rogers WH, Moxley III JH, Draper D, Brook RH. Hospital inpatient mortality: is it a predictor of quality? *N Engl J Med.* 1987;317:1674-80.
- <sup>30</sup> Bradley EH, Herrin J, Elbel B, McNamara RL, et al. Hospital quality for acute myocardial infarction: correlation among process measures and relationship with short-term mortality. *JAMA.* 2006;296:72-8.
- <sup>31</sup> Lichtlen PR, Nikutta P, Jost S, Deckers J, Wiese B, Rafflenbeul W. Anatomical progression of coronary artery disease in humans as seen by prospective, repeated, quantitated coronary angiography. Relation to clinical events and risk factors. The INTACT Study Group. *Circulation.* 1992;86:828-38.
- <sup>32</sup> Kanavos P, Gemmill-Toyama M. Prescription drug coverage among elderly and disabled Americans: can Medicare-Part D reduce inequities in access? *Int J Health Care Finance Econ.* 2010 Mar 6.
- <sup>33</sup> Charbonneau A, Rosen AK, Owen RR, Spiro A, Ash AS, Miller DR, Kazis L, Kader B, Cunningham F, Berlowitz DR. Monitoring depression care: in search of an accurate quality indicator. *Med care.* 2004;42:522-531.