Impact of a pharmacist-led diabetes management on outcomes, utilization, and cost

Daniel D Maeng  
Jove Graham  
Michael Bogart  
Jing Hao  
Eric A Wright  
Department of Epidemiology and Health Services Research, Geisinger, Danville, PA, USA

Purpose: Pharmacist-led medication therapy disease management (MTDM) has shown improvement in clinical outcomes in patients with certain chronic diseases. However, only limited data demonstrating the impact on health care utilization and cost of care are available. This study seeks to evaluate the impact of a pharmacist-led MTDM program on clinical surrogate outcomes, care utilization, and cost of care among patients with diabetes mellitus.

Methods: A retrospective cohort study was conducted by utilizing electronic health records and insurance claims data. Patients were identified between February 2011 and December 2014. Data were collected from Geisinger, a large integrated health care system located in Pennsylvania and southern New Jersey. A total of 5,500 patients with diabetes mellitus were identified; 2,750 were enrolled in MTDM and were 1-to-1 propensity score-matched to a comparison cohort not enrolled in a pharmacist-led MTDM program.

Results: There were no differences between groups in composite HbA1c, blood pressure, or low-density lipoprotein cholesterol goal attainment at 12 months (12% vs 12%, \( P = 0.53 \)). HbA1c goal was reached more frequently among patients without MTDM compared to those at 12 months (57% vs 51%, \( P < 0.0001 \)). There were no significant differences between the two cohorts in the attainment of blood pressure or low-density lipoprotein cholesterol goals at 12 months. MTDM was associated with reduced all-cause hospitalization rate (–19.6%; \( P = 0.02 \)) as well as increased primary care physician visits (18.5%; \( P < 0.001 \)) and lower average per-member-per-month medical cost (–13%, \( P = 0.027 \)).

Conclusion: Despite the lack of impact on the clinical surrogate outcomes, MTDM was associated with lower cost of care and fewer hospitalizations, possibly facilitated by increased monitoring (ie, higher primary care utilization).

Keywords: diabetes, pharmacist, medication therapy management, health outcomes, HbA1c, utilization, cost of care

Introduction

Over 30 million people in the United States have diabetes mellitus (DM).\(^1\) In addition to the morbidity and mortality associated with DM, the cost burden for an individual with DM is more than twice as high as it is for an individual without DM, while the total direct and indirect cost burden of DM in 2012 was estimated to be $245 billion.\(^2\) Although the clinical, humanistic, and economic burdens associated with DM are high, they can be improved through appropriate management of hyperglycemia and the often-associated comorbidities such as hypertension and hyperlipidemia. The potential impact of adequate treatment of all three of these risk factors has shown significant improvements in glycemic level, blood pressure, and lipid control; a >50%
control and cardiovascular risk in patients with diabetes. Despite evidence supporting the benefits of these risk factors in patients with DM, <10% of adults with DM in the United States have these factors adequately controlled. Among a multitude of other system-level initiatives, the American Diabetes Association recommend the expanding role of teams to intensify treatment. In addition to other team members, pharmacist-led management of diabetes is well-studied as an effective intervention to improve glycemic control and cardiovascular risk in patients with diabetes. However, universal adoption by health care systems has not occurred. One of the barriers for more widespread implementation is the lack of information on the relative impact of these services on health care utilization and cost of care. To improve the management of our patients with DM, Geisinger has developed and implemented a program of pharmacist-led management of DM, referred to as medication therapy disease management (MTDM). Initially piloted in 2008, the program embeds MTDM pharmacists within primary care provider (PCP) clinics where the pharmacists collaboratively manage a select population of patients with DM. This study assesses the impact of an MTDM program on the achievement of guideline-based disease targets, health care utilization, and cost.

Geisinger is a physician-led integrated health care system serving ~3 million residents across central and northeastern Pennsylvania and southern New Jersey. As of 2017, it consists of 12 hospitals, two research centers, two skilled nursing facilities, a substance abuse treatment center, and health insurance through the Geisinger Health Plan (GHP). It also includes 83 primary and specialty clinic sites, 44 of which are community-based primary care clinics. Approximately 40% of Geisinger patients have insurance coverage through GHP.

Medication therapy and disease management protocol
For DM, the defining characteristics are so unambiguous and the evidence base for successful treatment is so extensive to allow for rules-based, precision management that can be skillfully carried out by non-physician health care professionals, such as appropriately trained pharmacists. The distinguishing features of Geisinger’s pharmacist-led MTDM program relative to the prevailing DM standard of care include embedding of a MTDM pharmacist within each PCP clinic site and autonomy of the pharmacist in performing the MTDM functions.

Patients diagnosed with DM may be referred to the MTDM program at any time. Patients are referred by their physician who completes a pharmacist MTDM referral electronically in Geisinger’s electronic health records (EHRs) system. The decision for referral to MTDM is made by the physician based on his or her clinical judgment; no predetermined set of criteria are used to determine the patients’ eligibility for MTDM. At the initial visit, the pharmacist extensively interviews and educates the patient and verifies information with the EHR. The pharmacist is authorized to manage prescriptions for all the DM-related conditions. If a new prescription is needed, the MTDM pharmacist ensures that a written, e-prescribed, or telephoned prescription is generated by following collaborative practice guidelines. The pharmacist schedules subsequent MTDM appointments or laboratory testing as needed, independent of the referring physician. Every attempt is made to coordinate these appointments with preexisting appointments to make it convenient for the patient.

Methods
This study was funded by GlaxoSmithKline and approved by Geisinger’s Institutional Review Board. Geisinger’s EHR and GHP claims databases were queried to obtain the retrospective data covering a 6-year period from January 1, 2009, to December 31, 2014. To ensure patient confidentiality, patient information, such as patient name, address, and contact information, was removed from the final data sets. As noted above, the MTDM program was initially piloted in 2008 with one pharmacist in a single primary care site, and it was subsequently expanded after February 2011 to include 24 primary care sites as of 2015. For the purposes of this study, those patients who have enrolled in that initial pilot phase of the MTDM program were excluded, because it was deemed that the early pilot MTDM program had not been fully developed and mature. As such, the MTDM intervention group in this study included only those patients who had been enrolled in the MTDM program between February 2011 and December 2014. The non-MTDM comparison group was defined as those meeting the inclusion and exclusion criteria as described below but were not referred to the MTDM at any point during the same period. Pre-MTDM claims data from 2009 to 2010 were used to account for any preexisting differences between the intervention and the comparison groups.

The study population was defined as patients who were aged ≥18 years during the study period, who had a primary or secondary diagnosis for DM defined as an ICD, ninth Revision, Clinical Modification (ICD-9-CM) diagnosis code.
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of 250.xx; received health care services from a Geisinger primary care clinic; and had ≥1 month of both medical and prescription drug coverage through GHP. Patients were excluded if they were pregnant (ICD-9-CM 650.xx) or had gestational diabetes (ICD-9-CM 648.8); if they had any MTDM encounter prior to February 2011 (ie, exposed to MTDM during the pilot phase); or if they had previously requested not to be included in any research studies. Patients with <6 months of EHR data prior to their index date (defined as the date of first visit with a MTDM pharmacist) or <12 months of EHR data after index date were excluded to ensure adequate baseline and follow-up observation. Patients with only one MTDM encounter were also excluded, as a single encounter was attributable to a lack of engagement within the MTDM. Figure 1 shows the illustration of the observation schema.

The main outcome variables of interest were the percent of patients meeting three predefined targets concurrently and percent of patients meeting each individual target at 12 months following the initial MTDM visit with a pharmacist. Those three targets were: glycemic control of HbA1c <8; blood pressure of systolic pressure <130 mmHg and diastolic pressure <80 mmHg; and low-density lipoprotein cholesterol (LDL-C) levels of <100 or <70 mg/dL for patients with coronary heart disease or chronic kidney disease. Because this was an observational study, some patients did not have visits scheduled at exactly 12 months of follow-up; in these cases, the most recent measures prior to 12 months were carried forward.

For health care utilization and cost of care, GHP claims data were used to capture the per-member-per-month (PMPM) rates of all-cause acute inpatient admissions, emergency department (ED) visits, physician office visits, and the average PMPM total cost of care. Cost of care was defined as the allowed amounts, defined as the sum of GHP’s payments to the health care providers plus patients’ out-of-pocket costs in the forms of deductibles, copays, and coinsurance. All-cause costs were reported to avoid any potential bias that may result from inaccurate identification of diabetes-related care utilization. Total cost was divided into medical and prescription drug costs. Also, the analysis of utilization and cost was limited to the subsample who had prescription drug coverage through GHP to ensure that all the patients in the analysis had comparable access to prescription drugs. Within this subsample, the 1-to-1 matched balanced cohort was maintained by dropping from the subsample of those in the MTDM intervention group whose matched comparison did not have GHP prescription drug coverage.

Statistical analysis

We used a propensity scoring model to estimate the probability of MTDM enrollment and match the MTDM and control patients with similar scores at a 1:1 ratio. For MTDM patients, propensity was calculated from the first MTDM visit (i.e. index date). For control patients, since there was no enrollment date, propensity scores were calculated for every encounter they had during the period, with each encounter serving as a potential index date. Propensity was calculated as a function of the following patient characteristics at index date: sex; Caucasian race; whether the index

Figure 1 Study observation schema.

Abbreviation: MTDM, medication therapy disease management.
within a medical home environment. The impact of MTDM was then estimated as the difference between the regression-adjusted “observed” and “expected” values. “Observed” values represent the regression-adjusted mean estimates for the MTDM cohort, while “expected” values represent the regression-adjusted estimates for the same patients with the MTDM group indicators set to zero, that is, using counterfactual data from the non-MTDM cohort to estimate the outcomes of the MTDM cohort if those patients never enrolled in the MTDM program. Bootstrap standard errors with 100 replications were obtained to calculate the corresponding 95% confidence intervals.

**Results**

The final matched sample was 5,500 patients with diabetes (Figure 2). Approximately 22% (615/2,750) of MTDM cases had between two and five MTDM visits, and over half (1378/2750) had ≥15 MTDM visits, up to a maximum of 146 MTM visits. After the propensity score match, patient characteristics were well balanced (Table 1). Furthermore, prevalence of comorbid conditions was also similar after the matching (shown in Table S1).
Patient follow-up ranged from 365 to 1679 days, with a median of 935 days. As noted above, not all patients had scheduled follow-up visits exactly at or near 365 days. Therefore, a sensitivity analysis was conducted in which patients whose outcome values were carried forward >6 months were excluded from the sample (Table 2).

The percent of patients achieving all three targets simultaneously were similar between the MTDM and the non-MTDM cohorts (MTDM, 318 [12%]; non-MTDM, 333 [12%]; \(P=0.53\)), even after excluding patients whose latest outcome values were taken before 6 months following the index date (MTDM, 228 [13%]; non-MTDM, 233 [14%]; \(P=0.19\)) (Table 2). Both groups had reductions noted in HbA1c from baseline, but the change in HbA1c was more modest in the MTDM group than the comparison group (–0.5% vs –0.7%, \(P<0.0001\)), resulting in 51% of MTDM patients and 57% of comparison patients at goal (\(P<0.0001\)). The MTDM cohort also showed a smaller decline in DBP from baseline than the non-MTDM comparison group.
cohort (−0.9 to −1.8 mmHg, \(P=0.003\)); however, DBP goal attainment remained similar between groups. No additional differences between the cohorts were noted.

As noted above, the analysis of utilization and cost was limited to the subsample of patients who had GHP prescription drug coverage. This subsample consisted of 2,058 patients in each of the MTDM and non-MTDM groups, or ~75% of the propensity score-matched sample. Patients’ exposure to MTDM was associated with a reduction in the acute inpatient admission (270 vs 335 per 1,000 members per year or 19.6% reduction; \(P=0.02\)) as well as an increase in the PCP visit rate (5,555 vs 4,687 per 1,000 members per year or 18.5% increase; \(P<0.001\)), following the index date (Tables 3 and 4). MTDM was also associated with a statistically significant total medical cost savings ($1,061 vs $1,230 PMPM or 13.7% reduction; \(P=0.027\)) (Table 5). No significant impact on prescription drug costs was observed.

### Table 1 Baseline characteristics of the propensity-matched cohorts

| Patient characteristics | Non-MTDM (n=2,750) | MTDM (n=2,750) | \(|d|^*\) |
|-------------------------|---------------------|----------------|--------|
| Males, N (%)            | 1,419 (52)          | 1,383 (50)     | 0.03   |
| Age in years, mean (SD) | 59 (13)             | 59 (13)        | 0.00   |
| Age in years by category, N (%) |                  |                |        |
| 18–24                   | 1 (<1)              | 0 (0)          | 0.03   |
| 25–34                   | 94 (3)              | 99 (4)         | 0.01   |
| 35–44                   | 268 (10)            | 266 (10)       | 0.00   |
| 45–54                   | 614 (22)            | 612 (22)       | 0.00   |
| 55–64                   | 830 (30)            | 807 (29)       | 0.02   |
| 65–74                   | 645 (23)            | 659 (24)       | 0.01   |
| 75+                     | 308 (11)            | 307 (11)       | 0.00   |
| Race, N (%)             |                     |                |        |
| American Indian         | 4 (<1)              | 7 (<1)         | 0.02   |
| Asian                   | 12 (<1)             | 17 (<1)        | 0.03   |
| African-American        | 80 (3)              | 78 (3)         | 0.00   |
| Native Hawaiian/Pacific Islander |   |              |        |
| Unknown                 | 10 (<1)             | 6 (<1)         | 0.03   |
| White/Caucasian         | 2,640 (96)          | 2,640 (96)     | 0.00   |
| Weight in lbs, mean (SD)| 224 (59)            | 223 (55)       | 0.02   |
| BMI, mean (SD)          | 35.4 (8.4)          | 35.4 (7.8)     | 0.00   |
| Systolic BP in mmHg, mean (SD) | 129 (17)    | 129 (16)       | 0.00   |
| Diastolic BP in mmHg, mean (SD) | 74 (10)     | 73 (10)        | 0.10   |
| HbA1c in mg/dL, mean (SD) | 8.7 (1.8)      | 8.8 (1.8)      | 0.06   |
| LDL-C, mean (SD)        | 96 (36)             | 96 (38)        | 0.00   |
| Medication use in baseline 12 months, N (%) | | | |
| Antihypertensive         | 2,481 (90)          | 2,507 (91)     | 0.03   |
| Antihyperlipidemic       | 2,316 (84)          | 2,362 (86)     | 0.05   |
| Antidiabetic             | 1,106 (40)          | 1,385 (50)     | 0.20   |
| Insulin                  | 1,707 (62)          | 2,071 (75)     | 0.29   |
| Any oral antidiabetic    |                     |                |        |
| Quan-Charlson Comorbidity Index, mean (SD) | 1.38 (1.46)    | 1.48 (1.53)     | 0.07   |
| Quan-Charlson Index by category, N (%) | | | |
| 0                       | 896 (33)            | 884 (32)       | 0.01   |
| 1                       | 866 (31)            | 758 (28)       | 0.09   |
| 2                       | 489 (18)            | 516 (19)       | 0.03   |
| 3                       | 247 (9)             | 306 (11)       | 0.07   |
| 4                       | 136 (5)             | 148 (5)        | 0.02   |
| 5                       | 71 (3)              | 94 (3)         | 0.05   |
| 6                       | 25 (1)              | 21 (1)         | 0.02   |
| 7+ (max 12)             | 23 (<1)             | 23 (1)         | 0.01   |
| Patients with GHP coverage at time of index date, N (%) | 1,560 (57) | 1,560 (57) | 0.00 |

**Note:** \(|d|\) denotes standardized difference in mean or percentages and is not confounded by sample size as \(P\)-values are.

**Abbreviations:** LDL-C, low-density lipoprotein cholesterol; MTDM, medication therapy disease management; GHP, Geisinger Health Plan.
Our findings are consistent with a previous study conducted in another integrated health system setting, in which authors considered patients in a pharmacist-led diabetes management program within a patient-centered medical home versus patients not enrolled in the program. They reported similar

**Discussion**

Patients in a pharmacist-led MTDM program experienced lower rates of inpatient admissions and lower medical costs. Glycemic control among patients in the MTDM program has improved although to a lesser extent than matched controls.
observations of no significant differences in clinical outcomes but a significantly higher estimated rate of ambulatory care visits and lower rate of hospitalizations.

Other studies have shown positive associations between pharmacist-led management program and the clinical outcomes considered in this study.13–16 A meta-analysis of 15 randomized controlled trials also noted significant improvements in cardiovascular risk factors related to pharmacist-managed care.17 However, there are clear differences between these previous studies and the current study including differences in settings (eg, Veterans Affairs and community health center vs integrated health system), study design (eg, randomized clinical trial vs observational study), lower individual study sample sizes, and the differences in the design and maturity of the disease management programs. In our study, the MTDM program resulted in significant reduction in high-cost medical utilization leading to significant per-member per-month savings despite moderate improvements in HbA1c and less improvement than the comparator. Future research should seek to explore the mechanisms explaining the optimal medication management for DM patients.

Despite these explanations and comparisons with other studies, interpreting our seemingly contradictory findings is challenging. Our initial hypothesis, based on the collection of evidence to date, has been that pharmacist management would improve glycemic control, which would in-turn drive lower utilization and cost. Yet, despite a smaller reduction in HbA1c in the MTDM group, we have observed a decrease in inpatient admissions and a reduction in medical costs. While evidence suggests that improving glycemic control is associated with improved microvascular and possible macrovascular outcomes, evidence on acute health care utilization is not clear. In addition, there is evidence to suggest that the tight glycemic control increases the risk of harm via hypoglycemia18 without a benefit on most micro- or macrovascular outcomes.19

It is possible that based on the MTDM treatment protocol, pharmacists have tailored the regimens to the patients in ways that avoided hypoglycemia. They may have also avoided over-basalization (eg, hyper-focus on morning blood glucose at goal while disregarding glucose readings throughout the rest of day) and chose drugs with inherently better health/safety profile (eg, medication with less weight gain, lower cardiovascular and heart failure risk, and b-cell sparing medications). The MTDM treatment protocol includes early initiation and titration of metformin along with the initiation of newer evidence-based antidiabetic agents (eg, glucagon-like peptide-1 (GLP-1) agonists) for glycemic control over sulfonylureas. In a separate analysis, we have noted a 14% lower odds of using sulfonylureas in MTDM-managed patients versus comparison patients (OR 0.86, 95% CI 0.79, 0.94, P=0.002), higher odds of being placed on GLP-1 agonists (OR 1.90, 95% CI 1.45, 2.49, P<0.0001), biguanides (OR 1.12, 95% CI 1.02, 1.23, P=0.02), meglitinide analogs (OR 2.97, 95% CI 2.31, 3.81, P<0.0001), and insulin (OR 1.11, 95% CI 1.01, 1.21, P=0.02). These complementary mechanisms may support our non-congruent findings and may also support a reevaluation of HbA1c goal attainment as a metric for success in this type of program.

Another potential mechanism is increased collaboration with PCPs. The MTDM pharmacists are co-located in clinics with PCPs. This increased access could encourage greater patient–clinician interactions on an on-going basis that would not otherwise be possible in the traditional DM disease management model. The significant increase in the PCP visit rates among the MTDM patient cohort may reflect this. To the extent that increased PCP visits imply greater opportunity for surveillance, early detection, and prevention of potentially acute events, the increased PCP visit rate may be indicative of the underlying mechanism through which cost reductions and lower hospitalizations can be achieved.

### Table 5 MTDM impact on cost of care

| MTDM exposure      | Total medical ($ per-member-per-month) | Prescription drug ($ per-member-per-month) |
|--------------------|----------------------------------------|--------------------------------------------|
|                    | Observed | Expected | Difference (95% CI) | % Difference (P-value) | Observed | Expected | Difference (95% CI) | % Difference (P-value) |
| Pre-MTDM            | 763      | 763      | –                   | –                  | 193      | 193      | –                   | –                  |
| Post-MTDM: 1–12 months | 1,044   | 1,254   | –210 (–382, –38)    | –16.7 (0.016)      | 332      | 329      | 4                   | 0.9                |
| Post-MTDM: >12 months | 1,081  | 1,198   | –118 (–302, 67)     | –9.8 (0.213)       | 366      | 358      | 7                   | 2.2                |
| Post-MTDM: all months | 1,061  | 1,230   | –169 (–319, –19)   | –13.7 (0.027)      | 348      | 342      | 6                   | 1.8                |

Abbreviation: MTDM, medication therapy disease management.
However, further studies are necessary to reveal the more precise mechanism. Although we have mitigated the bias in our study design via propensity-score matching, which resulted in very close agreement on observed baseline characteristics between groups, we recognize that unmeasured confounding variables could also have influenced our unexpected findings. Our study observed and compared patients managed by an MTDM program and those not managed by an MTDM program after program implementation, and there may have been other dissimilar factors between cohorts that we did not measure or balance on. One example is baseline medication use, which in a post hoc analysis we noted had some differences between cohorts that could indicate differences in acuity or treatment (ie, the MTDM group had higher baseline use of insulin, sulfonylureas, metformin, meglitinides, and DPP4 inhibitors). We acknowledge that baseline differences like these, observed or unobserved, may have contributed to our results.

There are several additional potential limitations. The non-MTDM comparison group has been drawn from DM patients who were likely eligible for the MTDM program enrollment but were not referred to it by physicians for unknown reasons, implying a potential selection bias. We have attempted to mitigate this potential bias via an extensive propensity score matching algorithm. Additionally, if physicians were selecting higher-risk or more-difficult-to-manage patients into MTDM, our results are expected to underestimate the true effect. Another limitation is the unknown generalizability of the MTDM program beyond Geisinger. Although the MTDM program was designed to be scalable and generalizable, future studies are needed to examine the feasibility of similar MTDM programs elsewhere.

Finally, because patients were in an observational study and not adhering to a strict visit schedule, some patients were assigned as their end points the clinical values that were taken close to, rather than exactly at, the 12-month point. We attempted to assess the impact of this limitation by performing a sensitivity analysis as shown in Table 2, and the results were virtually identical. The patients considered “at goal” in this study, however, might not have remained so throughout the study period or vice versa. As such, future research could use longitudinal models to capture temporal dynamics between the clinical measures and the patients’ exposure to the MTDM program.

**Conclusion**

Despite the lack of impact on clinical surrogate outcomes, the results suggest that MTDM was associated with an overall lower cost of care and fewer hospitalizations. This finding is consistent with the expectation that pharmacist-led MTDM directly impact utilization and cost of care among a diabetic cohort and should be considered as an important member of a multidisciplinary team in the management of diabetes.

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**Disclosure**

The authors report no conflicts of interest in this work.

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### Table S1 Baseline comorbidities/medical history of the two cohorts

| Comorbidity                        | Non-MTDM (n=2,750) | MTDM (n=2,750) | |d|* |
|------------------------------------|--------------------|----------------|---|---|
| Acute myocardial infarction        | 125 (5%)           | 127 (5%)       | 0.00 |
| Unstable angina                    | 61 (2%)            | 68 (2%)        | 0.02 |
| Stable angina                      | 112 (4%)           | 127 (5%)       | 0.03 |
| Coronary heart disease             | 586 (21%)          | 642 (23%)      | 0.05 |
| Ischemic stroke                    | 48 (2%)            | 51 (2%)        | 0.01 |
| Peripheral artery disease          | 202 (7%)           | 222 (8%)       | 0.03 |
| Type 1 diabetes                    | 269 (10%)          | 257 (9%)       | 0.01 |
| Type 2 diabetes                    | 2683 (98%)         | 2706 (98%)     | 0.06 |
| Hypertension                       | 2226 (81%)         | 2226 (81%)     | 0.00 |
| Hyperlipidemia                     | 2391 (87%)         | 2391 (87%)     | 0.00 |
| AIDS                               | 3 (<1%)            | 2 (<1%)        | 0.01 |
| Congestive heart failure           | 255 (9%)           | 282 (10%)      | 0.03 |
| Chronic obstructive pulmonary disease | 920 (33%)        | 920 (33%)      | 0.00 |
| Dementia                           | 15 (<1%)           | 7 (<1%)        | 0.05 |
| Hemiplegia or paraplegia           | 12 (<1%)           | 8 (<1%)        | 0.02 |
| Leukemia                           | 6 (<1%)            | 7 (<1%)        | 0.01 |
| Lymphoma                           | 5 (<1%)            | 6 (<1%)        | 0.01 |
| Any malignancy                     | 218 (8%)           | 242 (9%)       | 0.03 |
| Mild liver disease                 | 137 (5%)           | 186 (7%)       | 0.08 |
| Moderate to severe liver disease   | 8 (<1%)            | 9 (<1%)        | 0.01 |
| Peptic ulcer disease               | 57 (2%)            | 58 (2%)        | 0.00 |
| Rheumatic disease                  | 101 (4%)           | 83 (3%)        | 0.04 |
| Renal disease                      | 488 (18%)          | 523 (19%)      | 0.03 |
| Valvular disease                   | 249 (9%)           | 308 (11%)      | 0.07 |
| Bipolar disorder                   | 36 (1%)            | 56 (2%)        | 0.06 |
| Depressive disorder                | 695 (25%)          | 775 (28%)      | 0.07 |
| Schizophrenia                      | 21 (1%)            | 15 (1%)        | 0.03 |

Note: *|d| denotes standardized difference in means or percentages.

Abbreviation: MTDM, medication therapy disease management.

Table S2 Codes used for identification of comorbid conditions

| Disease or procedure | ICD-9 CM or CPT Codes |
|----------------------|-----------------------|
| Acute myocardial infarction | 410.00, 410.01, 410.02, 410.10, 410.11, 410.12, 410.20, 410.21, 410.22, 410.30, 410.31, 410.32, 410.40, 410.41, 410.42, 410.50, 410.51, 410.52, 410.60, 410.61, 410.62, 410.70, 410.71, 410.72, 410.80, 410.81, 410.90, 410.91, 410.92 |
| Unstable angina       | 411.1, 411.81         |
| Stable angina         | 413.0, 413.1, 413.9   |
| Chronic obstructive pulmonary disease | 416.8, 416.9, 490.0, 491.0, 491.1, 491.20-490.22, 491.8, 491.9, 492.0, 492.0, 492.0-493.02, 493.00-493.12, 493.20-493.22, 493.81, 493.82, 493.9, 493.90-493.92, 494, 494.0, 494.1, 495.0, 495.1, 495.7-495.9, 496, 500-505, 506.4, 508.1, 508.8 |
| Coronary heart disease | 414, 414.0, 414.00-414.07, 414.1, 414.10-414.12, 414.19, 414.2, 414.3, 414.8, 414.9 |
| Ischemic stroke        | 433.01, 433.11, 433.21, 433.31, 433.81, 433.91, 434.01, 434.11, 434.91 |
| Peripheral artery disease | 00.55, 00.61, 00.63, 00.64, 38.13, 38.18, 39.50, 39.72, 39.74, 39.90, 333.00, 433.10, 433.20, 433.30, 433.80, 433.90, 441.3, 441.4, 443.9, 445.0, 445.01, 445.02, 35301, 34800-34805, 35081-35103, 35450-35459, 35470-35475, 35480-35485, 35490-35495, 35501-35571, 35583-35587, 35601-35671, 37205-37208, 37215-37216, 37220-37235, 93668 |
| Type 1 diabetes        | 250.01, 250.03, 250.11, 250.13, 250.21, 250.23, 250.31, 250.33, 250.41, 250.43, 250.51, 250.53, 250.61, 250.63, 250.71, 250.73, 250.81, 250.83, 250.91, 250.93 |
| Type 2 diabetes        | 250.00, 250.02, 250.10, 250.12, 250.20, 250.22, 250.30, 250.32, 250.40, 250.42, 250.50, 250.52, 250.60, 250.62, 250.70, 250.72, 250.80, 250.82, 250.90, 250.92 |
| Hypertension           | 401.0, 401.1, 401.9, 402.0, 402.02, 402.10, 402.11, 402.90, 402.91, 403.00, 403.01, 403.10, 403.11, 403.9, 403.90, 403.91, 404.00, 404.01, 404.03, 404.10, 404.11, 404.90, 404.91, 405.01, 405.09, 405.11, 405.19, 405.91, 405.99 |

(Continued)
### Table S2 (Continued)

| Disease or procedure          | ICD-9 CM or CPT Codes                                                                 |
|-------------------------------|---------------------------------------------------------------------------------------|
| Hyperlipidemia                | 272.0, 272.1, 272.2, 272.4                                                           |
| Valvular disease              | V42.2, V43.3, 35.01-35.14, 35.20-35.28, 35.96, 35.97, 35.99, 424.0, 424.1, 424.2, 424.3, 427.31, 424.90, 424.91, 424.99, 746.00, 746.01, 746.02, 746.09, 746.1-746.7 |
| Bipolar disorder              | 296.4, 296.41-296.44, 296.46, 296.5, 296.51, 296.52, 296.54, 296.55, 296.6, 296.61-296.64, 296.7, 296.8, 296.82, 296.89, 296.9, 296.99 |
| Depression                    | 296.2, 296.21-296.26, 296.3, 296.31-296.36, 300.4, 311                              |
| Schizophrenia                 | 295.00, 295.01, 295.20, 295.22, 295.30, 295.32, 295.34, 295.40, 295.44, 295.52, 295.54, 295.60, 295.62, 295.64, 295.70, 295.72, 295.74, 295.75, 295.80-295.82, 295.90, 295.92 |
| Coronary artery bypass graft  | 36.10-36.17, 36.19, 36.20, 33510-33519, 33521-33523, 33533-33536                   |
| Coronary revascularization    | 00.66, 36.0, 36.03, 36.04, 36.06, 36.07, 36.09, 36.1, 36.10, 36.11, 36.12, 36.13, 36.14, 36.15, 36.16, 36.17, 36.19, 36.2, 36.3, 36.31, 36.32, 36.33, 36.34, 36.39, 92980, 92981, 92982, 92984, 92995, 92996, 33510-33536 (except 33530), G0290, G0291, S2205-S2209 |