Retrospective Analysis of Long-Term Adherence to and Persistence with DPP-4 Inhibitors in US Adults with Type 2 Diabetes Mellitus

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Received: October 23, 2014 / Published online: December 12, 2014
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ABSTRACT

Introduction: Patients with type 2 diabetes mellitus (T2DM) must remain adherent and persistent on antidiabetic medications to optimize clinical benefits. This analysis compared adherence and persistence among adults initiating dipeptidyl peptidase-4 inhibitors (DPP-4is), sulfonylureas (SUs), and thiazolidinediones (TZDs) and between patients initiating saxagliptin or sitagliptin, two DPP-4is.

Methods: This retrospective cohort study utilized the US MarketScan® (Truven Health Analytics, Ann Arbor, MI, USA) Commercial and Medicare Supplemental health insurance claims databases. Adults aged ≥18 years with T2DM who initiated a DPP-4i, SU, or TZD from January 1, 2009 to January 31, 2012 were included. Patients must have been continuously enrolled for ≥1 year prior to and ≥1 year following initiation. Adherence was measured using proportion of days covered (PDC), with PDC ≥ 0.80 considered adherent. Persistence was measured as time to discontinuation, defined as last day with drug prior to a 60+ days gap in therapy. Multivariable logistic regression and Cox proportional hazards models compared the outcomes between cohorts, controlling for baseline differences.

Results: The sample included 238,372 patients (61,399 DPP-4i, 134,961 SU, 42,012 TZD). During 1-year follow-up, 47.3% of DPP-4i initiators, 41.2% of SU initiators, and 36.7% of TZD initiators were adherent. Adjusted odds of adherence were significantly greater among DPP-4i initiators than SU (adjusted odds ratio [AOR] = 1.678, P < 0.001) and TZD initiators (AOR = 1.605, P < 0.001). During 1-year follow-up, 55.0% of DPP-4i initiators, 47.8% of SU initiators, and 42.9% of TZD initiators did not discontinue therapy. Adjusted hazards of discontinuation were significantly greater for SU (adjusted hazard ratio [AHR] = 1.390,
and TZD initiators (AHR = 1.402, P < 0.001) compared with DPP-4i initiators. Saxagliptin initiators had significantly better adherence (AOR = 1.213, P < 0.001) compared with sitagliptin initiators, and sitagliptin initiators had significantly greater hazard of discontinuation (AHR = 1.159, P < 0.001). Results were similar over a 2-year follow-up.

Conclusions: US adults with T2DM who initiated DPP-4i therapy, particularly saxagliptin, had significantly better adherence and persistence compared with patients who initiated SUs or TZDs.

Keywords: Adherence; Dipeptidyl peptidase-4 inhibitors; Persistence; Type 2 diabetes mellitus

INTRODUCTION

In the US, an estimated 26 million people have diabetes, approximately 8% of the entire population [1]. Diabetic complications are an important medical and public health concern, as diabetes is associated with kidney failure, limb amputation, blindness, heart disease, and stroke [1]. Appropriate disease management, primarily maintenance of glycated hemoglobin (HbA1c) levels below 7% can decrease the risk of microvascular complications [2]. Current treatments for type 2 diabetes mellitus (T2DM) include lifestyle and dietary modifications, oral antidiabetic medications, non-insulin injectables, and insulin [2, 3]. For patients with mild hyperglycemia, lifestyle change and monotherapy with metformin or an acceptable alternative, such as a dipeptidyl peptidase-4 inhibitor (DPP-4i), are recommended [3]. Sulfonylureas (SUs) and thiazolidinediones (TZDs) can also be prescribed, although it has been suggested that the additional risks associated with these drug classes should be considered [3]. The American Diabetes Association, European Association for the Study of Diabetes, and the American Association of Clinical Endocrinologists recommend that patients who do not reach HbA1c targets should be prescribed combination therapy, followed by insulin initiation if necessary [2, 3].

To optimize the clinical benefits of antidiabetic pharmacotherapy, patients must remain adherent to and persistent with their prescribed medications. The importance of adherence to antidiabetic medications has been recognized by Medicare and appears as a Medicare Part D plan quality measure [4]. Increased adherence to medications is associated with a decrease in HbA1c [5–8], lower odds of mortality [7], fewer all-cause hospitalizations [7], and lower healthcare expenditures [9–12]. These improvements in outcomes may translate into substantial benefit to the healthcare system, averting a projected 699,000 emergency room visits and 341,000 hospitalizations per year, resulting in a savings of nearly $5 billion [13]. Despite the noted impact of good adherence, a systematic review found a wide range of adherence rates across retrospective studies, with rates ranging from 36% to 93% [14]. A second review found that when combining study results, only 58% of patients were considered adherent [15]. To date, the majority of interventions to improve adherence to diabetes medications have focused on education and one-on-one counseling [16].

Several factors associated with adherence to antidiabetic medications have been identified in the literature. Younger age, female gender, residence in the Southern region of the US, and high cost-sharing have been associated with lower adherence in multiple analyses [10, 17, 18]. Additional factors include depression,
polytherapy, and twice-daily regimens [14]. Analyses have also compared patients initiating specific non-insulin antidiabetic drugs and drug classes in terms of adherence and persistence, including metformin, SUs, glucagon-like peptide-1 receptor agonists (GLP-1RAs), TZDs, and DPP-4is [10, 17, 19, 20].

To our knowledge, no US-based analyses comparing adherence to and persistence with DPP-4is as a medication class with other classes of antidiabetic medications have been reported. Therefore, the objective of this analysis was to compare adherence and persistence among patients initiating DPP-4is, SUs, and TZDs over 1- and 2-year follow-up periods. Additional objectives were to compare adherence and persistence between patients initiating saxagliptin and sitagliptin and to describe treatment patterns following initiation.

METHODS

Data Source

This retrospective, observational cohort study used administrative claims data from the Truven Health MarketScan® (Truven Health Analytics, Ann Arbor, MI, USA) Commercial and Medicare Supplemental databases. The databases contain inpatient medical, outpatient medical, and outpatient prescription administrative claims data and enrollment records of individuals with employer-sponsored primary or Medicare supplemental insurance. The Commercial and Medicare Supplemental Databases have been utilized in numerous published analyses [21]. The data were previously collected, de-identified, and compliant with the Health Insurance Portability and Accountability Act privacy regulations. Therefore, Institutional Review Board approval and written informed consent were not required for this study.

Patient Selection

Patients with at least one outpatient pharmacy claim with a National Drug Code (NDC) for a DPP-4i, SU, or TZD from January 1, 2009 to January 31, 2012, were selected. A rolling selection method was used, whereby each drug claim was evaluated on the criteria described below and the earliest claim meeting all criteria was selected as the index date and the medication class (DPP-4i, SU, or TZD) was considered the index drug class. Patients must have had at least 28 days of continuous supply of the index drug class and have been at least 18 years old on the index date. All patients were required to have at least 365 days of continuous enrollment with medical and pharmacy benefits prior to the index date (pre-period) and at least 365 days of enrollment following the index date (1-year follow-up period). A subset of patients was required to have 730 days of post-index enrollment (2-year follow-up period). To capture initiators only, patients were required to have no claims for the index drug class in the pre-period. Lastly, patients were required to have at least one non-diagnostic medical claim with a T2DM diagnosis (International Classification of Diseases, Ninth Revision, Clinical Modification [ICD-9-CM] 250.x0, 250.x2) during the study period. Patients with medical claims with diagnoses of type 1 diabetes (ICD-9-CM 250.x1, 250.x3) or gestational diabetes (ICD-9-CM 648.8x) or with multiple index drug classes on index date were excluded from analysis.

Outcome Variables

The primary outcomes were adherence to and persistence with the index drug class. Both
measures were calculated using the service date and days’ supply fields existing on outpatient pharmacy claims for DPP-4is, SUs, and TZDs identified by NDC codes. Adherence was measured as proportion of days covered (PDC), calculated by taking the number of days a patient had the index drug class on hand during the 1- or 2-year follow-up based on the days’ supply field on pharmacy claims divided by follow-up time (365 or 730 days). Days’ supply for early refills was appended to the end of days’ supply of the previous prescription. Patients with a PDC \( \geq 0.80 \) were considered adherent. Persistence was defined as the time from index date to last day with index drug class on hand prior to a gap of \( \geq 60 \) consecutive days without index drug class [22]. A cut-point of 60 days was utilized as a conservative definition of discontinuation, as patients with gaps of up to 60 days were considered persistent. Switching within drug class was allowed for the drug class comparison. When comparing adherence and persistence outcomes between patients initiating saxagliptin and sitagliptin, PDC and time to discontinuation were calculated at the drug level, rather than the drug class level.

Patients were classified into the following mutually exclusive groups based on first event after index date during the 1-year follow-up period: remained on index drug class with no augmentation, augmentation with additional drug class, discontinuation of index drug class and switch to a new drug class, discontinuation of study drug class and continuation on other medication classes with no switch, and discontinuation of all antidiabetic medications. An augmentation was defined as the addition of a medication class not part of the initial regimen that overlapped with the index drug class for \( \geq 30 \) days. A switch was defined as the discontinuation of index drug class and the addition of medication class not in the initial regimen prior to discontinuation with overlap \( \leq 30 \) days or following discontinuation. Discontinuation was measured at the drug class level for all cohort comparisons.

**Independent Variables**

The primary independent variable of interest was index drug class: DPP-4i, SU, or TZD. When comparing within the DPP-4i medication class, the primary predictor was index drug: saxagliptin or sitagliptin. A third DPP-4i, linagliptin, was not compared within the DPP-4i–specific analysis because few linagliptin initiators had 1 year of follow-up, and no linagliptin initiators had 2 years of follow-up in the claims data. A fourth DPP-4i, alogliptin, was not available in the US during the patient selection period.

Demographic, clinical, and cost and utilization characteristics were measured to describe the study sample and to control for potential confounding in multivariable models. Demographic measures included: sex, age, geographic region, urbanicity, insurance plan type, primary payer, presence of capitated services, and year of index date. Clinical characteristics were measured during the pre-period and included use of study drugs other than index drug class, and metformin and insulin use based on pharmacy claims, overall health captured by the Deyo Charlson Comorbidity Index (CCI) [23] and the number of unique ICD-9-CM codes appearing on a patient’s medical claims, diagnosis of macrovascular disease (acute myocardial infarction, other ischemic heart disease, congestive heart failure, cerebrovascular accident, or peripheral vascular disease) and microvascular disease (diabetic peripheral
neuropathy, diabetic retinopathy, leg and foot amputation, or nephropathy), and diagnosis or procedure indicative of renal impairment. Pregnancy during the follow-up period was captured, as it may affect diabetes treatment. Cost and utilization covariates captured during the pre-period were evidence of an endocrinologist visit and cardiologist visit, total healthcare expenditures, and diabetes prescription expenditures.

Several characteristics of the index drug prescription claim were also captured: the days’ supply, mail-order status, fixed-dose combination with metformin, cost-sharing index, which was the average patient out-of-pocket cost for a 30-day supply, calculated annually for each study drug for each insurance plan in the database, and index regimen [24]. A patient’s initial regimen was defined as the index drug class plus any concomitant use of additional antidiabetic medication classes (alpha-glucosidase inhibitors, amylin analogs, dopamine receptor agonists, GLP-1RAs, insulin, meglitinides, and metformin). Claims with service dates occurring 60 days prior to the index date to 45 days after were evaluated. A medication class was considered part of the initial regimen if (a) the patient received a prescription for the drug class within 60 days before index date and received a second prescription for that drug class within 45 days after index date or (b) the patient received a prescription for the drug within 60 days before index date or 45 days after index date and had ≥30 days’ overlap between index drug class and additional drug class between index date and index date + 45 days. This definition was purposely conservative so as not to misclassify switches as part of a patient’s initial regimen. Patients with no additional medication classes meeting this definition were considered to be on monotherapy.

Analysis

In descriptive analyses, pairwise comparisons using t tests and Chi-squared tests were conducted comparing SU initiators and TZD initiators with DPP-4i initiators and saxagliptin initiators with sitagliptin initiators. Multivariable logistic regression models were fit to compare the odds of being adherent, and multivariable Cox proportional hazards models were fit to compare time to discontinuation (persistence) between index drug classes and index drugs. To determine which covariates were included in the multivariable models, descriptive statistics and statistical tests were evaluated to identify characteristics that differed between cohorts and also may be associated with adherence or persistence (i.e., were potential confounders). Models controlled for the following variables: age, sex, presence of capitated services, payer, region, urbanicity, plan type, indicator for fixed-dose metformin index drug, indicator for index drug filled via mail order, index drug regimen (monotherapy, index drug plus additional non-insulin antidiabetic drugs, index drug plus insulin), baseline total expenditures and diabetes prescription expenditures, index diabetes medication class cost-sharing, baseline endocrinologist and cardiologist visits, baseline renal impairment, baseline macrovascular and microvascular disease, pregnancy during follow-up, baseline number of unique three-digit ICD-9-CM diagnoses and Charlson Comorbidity Index. Sensitivity analyses of monotherapy and non-mail-order patients were also conducted, as patients initiating monotherapy have the simplest treatment profile and those with an initial prescription filled via mail order may have inflated persistence due to larger days’ supply on the index claim and automatic refill. P values
were compared with an a priori alpha 0.05 to determine statistical significance. Analyses were conducted with SAS version 9.3 (SAS Institute Inc. Cary, NC, USA).

RESULTS

Patient Selection

Patient attrition is presented in Fig. 1. There were 1,788,908 patients identified from the databases with at least one prescription claim for a DPP-4i, SU, or TZD from January 1, 2009 to January 31, 2012. After applying additional inclusion and exclusion criteria, the final sample consisted of 238,372 patients; 61,399 patients initiated a DPP-4i, 134,961 initiated an SU, and 42,012 initiated a TZD. Within DPP-4i initiators, 11,219 patients initiated saxagliptin and 49,400 initiated sitagliptin. A total of 134,444 patients had 2 years of follow-up data.

Descriptive Analysis

Demographic, clinical, and treatment characteristics are presented in Tables 1 and 2. The average age of patients in the sample was 56.7 years (standard deviation [SD] 12.2), and 54.3% were men. The majority of patients were insured through a commercial plan (77.7%). Approximately, one-fifth (20.3%) had evidence of macrovascular disease in the pre-period. Metformin use was common in the pre-period (55.0%), whereas insulin use was less common (6.7%). There were significant differences in baseline characteristics among the three drug class cohorts. DPP-4i initiators were younger on average than SU initiators but older than TZD initiators, and a greater proportion were women than both SU and TZD initiators. The proportion of patients using metformin in the pre-period was higher among DPP-4i initiators, and a greater proportion of patients initiated a study drug that was part of a fixed-dose combination with metformin. A mail-order index prescription was more common among DPP-4i initiators, whereas initiating monotherapy was less common. Significant differences were also found between the saxagliptin and sitagliptin initiators. On average, saxagliptin initiators were younger and more commonly insured through commercial plans. The prevalence of renal impairment, macrovascular disease, and microvascular disease was lower among saxagliptin initiators. More saxagliptin initiators initiated monotherapy compared with sitagliptin initiators.

Figure 2 presents treatment patterns following index drug class initiation. Among all study cohorts, augmentation was the least common event, although augmentation was more common among DPP-4i initiators. A significantly larger proportion of DPP-4i initiators remained on index drug class compared with SU and TZD initiators. Among DPP-4i initiators, the proportion of patients remaining on DPP-4is was significantly larger among the saxagliptin initiators. Overall, 20.6% of the study sample discontinued all antidiabetic medications for >60 days without first switching or augmenting. Discontinuing all antidiabetic medications was least common among DPP-4i initiators because, as noted above, remaining on initial regimen or augmenting was more common. In the DPP-4i-specific comparison, discontinuing all antidiabetic medications was less common among saxagliptin initiators. Within the first year, over 40% of SU and TZD initiators stopped refilling their index drug class, either discontinuing all antidiabetic medications or
continuing on the other medications in their initial regimen.

Unadjusted adherence and persistence results are presented in Table 3. Over the 1-year follow-up period, a significantly greater proportion of DPP-4i initiators were adherent to index drug class than SU and TZD initiators. This relationship was consistent over the 2-year follow-up period and in the monotherapy and the non-mail-order sub-samples. Significantly more DPP-4i initiators were persistent with index drug class during the 1-year follow-up period than SU initiators and TZD initiators. As with adherence, this was true over the 2-year

Fig. 1 Patient attrition. DPP-4i Dipeptidyl peptidase-4 inhibitor, ICD-9-CM International Classification of Diseases, Ninth Revision, Clinical Modification, SU Sulfonylurea, TZD Thiazolidinedione
| Table 1 Patient demographic characteristics |
|---------------------------------------------|
|                                      | Full study population | DPP-4i initiators | SU initiators | TZD initiators | DPP-4i initiators: Saxagliptin | DPP-4i initiators: Sitagliptin |
|                                      | N = 238,372            | N = 61,399         | N = 134,961   | N = 42,012     | N = 11,219                    | N = 49,400                    |
| Age, years (mean, SD)                | 56.7/12.2              | 56.4/11.7          | 57.2/12.6     | 55.6/11.6     | 55.6/11.3                    | 56.5/11.8                    |
| Sex (N, %)                           |                          |                    |              |              |                              |                              |
| Men                                  | 129,456/54.3%          | 31,407/51.2%       | 74,303/55.1% | 23,746/56.5% | 5,779/51.5%                 | 25,208/51.0%                |
| Women                                | 108,916/45.7%          | 29,992/48.8%       | 60,658/44.9% | 18,266/43.5% | 5,440/48.5%                 | 24,192/49.0%                |
| Region (N, %)                        |                          |                    |              |              |                              |                              |
| Northeast                            | 31,785/13.3%           | 10,129/16.5%       | 16,768/12.4% | 4,888/11.6%  | 1,727/15.4%                 | 8,297/16.8%                 |
| North Central                        | 61,753/25.9%           | 14,334/23.3%       | 37,056/27.5% | 10,363/24.7% | 2,428/21.6%                 | 11,781/23.8%                |
| South                                | 100,688/42.2%          | 27,670/45.1%       | 54,397/40.3% | 18,621/44.3% | 5,825/51.9%                 | 21,405/43.3%                |
| West                                 | 41,702/17.5%           | 8,623/14.0%        | 25,416/18.8% | 7,663/18.2%  | 1,135/10.1%                 | 7,388/15.0%                 |
| Unknown                              | 2,444/1.0%             | 643/1.0%           | 1,324/1.0%   | 477/1.1%      | 104/0.9%                    | 529/1.1%                    |
| Population density (N, %)            |                          |                    |              |              |                              |                              |
| Urban                                | 194,717/81.7%          | 50,939/83.0%       | 109,486/81.1%| 34,292/81.6% | 9,223/82.2%                 | 41,094/83.2%                |
| Rural                                | 41,367/17.4%           | 9,847/16.0%        | 24,260/18.0% | 7,260/17.3%  | 1,900/16.9%                 | 7,799/15.8%                 |
| Unknown                              | 2,288/1.0%             | 613/1.0%           | 1,215/0.9%   | 460/1.1%      | 96/0.9%                     | 507/1.0%                    |
| Insurance plan type (N, %)           |                          |                    |              |              |                              |                              |
| Comprehensive                        | 29,943/12.6%           | 7,699/12.5%        | 18,045/13.4% | 4,199/10.0%  | 1,384/12.3%                 | 6,256/12.7%                 |
| EPO                                  | 2,741/1.1%             | 784/1.3%           | 1,418/1.1%   | 539/1.3%      | 159/1.4%                    | 613/1.2%                    |
| HMO                                  | 41,690/17.5%           | 8,733/14.2%        | 26,238/19.4% | 6,719/16.0%  | 1,349/12.0%                 | 7,297/14.8%                 |
| POS                                  | 19,565/8.2%            | 5,607/9.1%         | 10,626/7.9%  | 3,332/7.9%    | 1,177/10.5%                 | 4,351/8.8%                  |
| PPO                                  | 120,966/50.7%          | 31,827/51.8%       | 65,674/48.7% | 23,465/55.9% | 5,652/50.4%                 | 25,772/52.2%                |
| POS with capitation                  | 962/0.4%               | 326/0.5%           | 475/0.4%     | 161/0.4%      | 53/0.5%                     | 267/0.5%                    |
### Table 1 continued

|                  | Full study population | DPP-4i initiators: Saxagliptin | DPP-4i initiators: Sitagliptin |
|------------------|-----------------------|-------------------------------|-------------------------------|
|                  | $N = 238,372$         | $N = 61,399$                  | $N = 42,012$                  |
|                  | $N/\text{Mean} \quad \%/\text{SD}$ | $N/\text{Mean} \quad \%/\text{SD}$ | $N/\text{Mean} \quad \%/\text{SD}$ |
| CDHP/HDHP        | 10,412 4.4%           | 2,943 4.8%                    | 1,691 4.0%                    |
| Unknown          | 12,093 5.1%           | 3,480 5.7%                    | 1,906 4.5%                    |

**Presence of capitation ($N, \%$)**

- **Capitated**
  - 26,321 11.0%
  - 4,804 7.8%
  - 17,106 12.7%
  - 4,411 10.5%
  - 621 5.5%
  - 4,146 8.4%
- **Non-capitated**
  - 212,051 89.0%
  - 56,595 92.2%
  - 117,855 87.3%
  - 37,601 89.5%
  - 10,598 94.5%
  - 45,254 91.6%

**Primary payer ($N, \%$)**

- **Commercial**
  - 185,281 77.7%
  - 48,999 79.8%
  - 102,055 75.6%
  - 34,227 81.5%
  - 9,278 82.7%
  - 39,060 79.1%
- **Medicare**
  - 53,091 22.3%
  - 12,400 20.2%
  - 32,906 24.4%
  - 7,785 18.5%
  - 1,941 17.3%
  - 10,340 20.9%

*P* values for all comparisons between DPP-4i initiators and SU initiators and between DPP-4i initiators and TZD initiators are <0.0001. *P* values for all comparisons between saxagliptin and sitagliptin initiators are <0.0001 except for sex ($P = 0.356$) and population density ($P = 0.004$).

CDHP: Consumer-driven health plan, DPP-4i: Dipeptidyl peptidase-4 inhibitor, EPO: Exclusive provider organization, HDHP: High-deductible health plan, HMO: Health maintenance organization, POS: Point-of-service plan, PPO: Preferred provider organization, SD: Standard deviation, SU: Sulfonylurea, TZD: Thiazolidinedione.
| Table 2 Patient clinical and treatment characteristics |
|-----------------------------------------------|
|                                      | Full study population | DPP-4i initiators | SU initiators | TZD initiators | DPP-4i initiators: Saxagliptin | DPP-4i initiators: Sitagliptin |
|                                      | N = 238,372 | N = 61,399 | N = 134,961 | N = 42,012 | N = 11,219 | N = 49,400 |
| Pre-period clinical characteristics     |                |                |                |                |                |                |
| Deyo CCI* (mean, SD)                    | 1.5 1.4       | 1.6 1.4       | 1.5 1.5       | 1.4 1.3       | 1.5 1.3       | 1.6 1.4       |
| Number of unique 3-digit ICD-9-CM      | 10.0 8.0      | 10.6 8.0      | 10.0 8.1      | 9.3 7.2       | 10.3 7.6      | 10.6 8.0      |
| Comorbid conditions (N, %)              |                |                |                |                |                |                |
| Renal impairment                       | 16,913 7.1%   | 4,071 6.6%    | 10,308 7.6%   | 2,534 6.0%    | 602 5.4%      | 3,366 6.8%    |
| Macrovascular disease                  | 48,408 20.3%  | 12,719 20.7%  | 28,509 21.1%  | 7,180 17.1%   | 2,066 18.4%   | 10,480 21.2%  |
| Microvascular disease                  | 28,491 12.0%  | 7,701 12.5%   | 15,916 11.8%  | 4,874 11.6%   | 1,230 11.0%   | 6,333 12.8%   |
| Pregnancy during follow-up             | 659 0.3%      | 140 0.2%      | 410 0.3%      | 109 0.3%      | 23 0.2%       | 115 0.2%      |
| Medication use (N, %)                  |                |                |                |                |                |                |
| Metformin                              | 131,008 55.0% | 37,143 60.5%  | 72,020 53.4%  | 21,845 52.0%  | 7,171 63.9%   | 29,488 59.7%  |
| Insulin                                | 15,861 6.7%   | 4,675 7.6%    | 7,417 5.5%    | 3,769 9.0%    | 759 6.8%      | 3,842 7.8%    |
| Pre-period utilization and expenditures |                |                |                |                |                |                |
| Visit to endocrinologist (N, %)        | 14,997 6.3%   | 5,903 9.6%    | 6,455 4.8%    | 2,639 6.3%    | 925 8.2%      | 4,855 9.8%    |
| Visit to cardiologist (N, %)            | 53,003 22.2%  | 15,163 24.7%  | 29,891 22.1%  | 7,949 18.9%   | 2,554 22.8%   | 12,401 25.1%  |
| Total healthcare expenditures (mean, SD) | $11,741 $29,070 | $12,100 $28,645 | $12,205 $30,918 | $9,726 $22,803 | $10,385 $22,394 | $12,427 $29,699 |
| Diabetes medication expenditures (mean, SD) | $255 $789 | $339 $920 | $197 $657 | $317 $940 | $312 $905 | $344 $920 |
Table 2  continued

|                      | Full study population | DPP-4i initiators | SU initiators | TZD initiators | DPP-4i initiators: Saxagliptin | DPP-4i initiators: Sitagliptin |
|----------------------|-----------------------|-------------------|---------------|---------------|--------------------------------|--------------------------------|
|                      | N = 238,372           | N = 61,399        | N = 134,961   | N = 42,012    | N = 11,219                      | N = 49,400                      |
| Diabetes medication cost-sharing index for 30-day supply (mean, SD) | $14 $13               | $26 $14           | $5 $3        | $25 $13       | $28 $14                        | $25 $14                        |
| Characteristics of index drug |                      |                   |               |               |                                |                                |
| Index prescription was fixed dose with metformin (N, %) | 42,825 18.0%       | 21,435 34.9%     | 11,240 8.3%  | 10,150 24.2%  | 2,871 25.6%                    | 18,564 37.6%                   |
| Index prescription was mail-order (N, %) | 32,146 13.5%       | 11,551 18.8%     | 13,458 10.0% | 7,137 17.0%   | 1,660 14.8%                    | 9,786 19.8%                    |
| Drug regimen at index (N, %) |                      |                   |               |               |                                |                                |
| Monotherapy | 88,382 37.1%       | 19,097 31.1%     | 55,007 40.8% | 14,278 34.0%  | 3,980 35.5%                    | 14,719 29.8%                   |
| Study drug plus 1 other NIAD | 128,763 54.0%     | 36,194 58.9%     | 70,515 52.2% | 22,054 52.5%  | 6,386 56.9%                    | 29,502 59.7%                   |
| Study drug plus 2+ other NIAD | 7,730 3.2%        | 2,479 4.0%       | 2,997 2.2%   | 2,254 5.4%    | 303 2.7%                       | 2,159 4.4%                     |
| Study drug plus insulin and 1+ other NIAD | 7,595 3.2%        | 2,229 3.6%       | 3,422 2.5%  | 1,944 4.6%    | 325 2.9%                       | 1,882 3.8%                     |
| Study drug plus insulin only | 5,902 2.5%        | 1,400 2.3%       | 3,020 2.2%  | 1,482 3.5%    | 225 2.0%                       | 1,138 2.3%                     |

*A higher value on the Deyo-CCI measure indicates a greater number of comorbidities and greater risk of mortality. P values for all comparisons between DPP-4i and SU initiators are <0.0001 except for baseline macrovascular disease (P = 0.039), pregnancy during follow-up (P = 0.003), and baseline total healthcare expenditures (P = 0.477). P values for all comparisons between DPP-4i and TZD initiators are <0.0001 except for pregnancy during follow-up (P = 0.311). P values for all comparisons between saxagliptin and sitagliptin initiators are <0.0001, except for pregnancy during follow-up (P = 0.577) and baseline diabetes medication expenditures (P = 0.001).

CCI Charlson Comorbidity Index, DPP-4i Dipeptidyl peptidase-4 inhibitor, ICD-9-CM International Classification of Diseases, Ninth Revision, Clinical Modification, NIAD Non-insulin antidiabetic drug, SD Standard deviation, SU Sulfonylurea, TZD Thiazolidinedione
follow-up period and in both sub-samples. Among DPP-4i initiators, patients initiating saxagliptin had better unadjusted persistence to index drug than patients initiating sitagliptin overall and in sub-samples. There were significant differences in adherence between the two cohorts in all samples except monotherapy patients. Descriptively, a smaller proportion of linagliptin patients were adherent (43.5%) compared with saxagliptin patients (49.1%) and sitagliptin patients (46.1%) and a smaller proportion of linagliptin patients were persistent (50.9% vs. 56.6% for saxagliptin and 53.6% for sitagliptin) during the first year of follow-up. No statistical comparisons were conducted due to small sample size.

Multivariable Analysis

Figure 3 presents adherence results. In adjusted logistic regression models, patients initiating DPP-4is had significantly greater odds of being adherent than patients initiating an SU or TZD. During the 1-year follow-up period, the adjusted odds of being adherent among DPP-4i initiators was over 1.6 times the odds of being adherent among SU initiators (adjusted odds ratio [AOR] = 1.678, 95% confidence interval [CI] = 1.631, 1.727) and TZD initiators (AOR = 1.605, 95% CI = 1.563, 1.647). Among DPP-4i initiators, patients initiating saxagliptin had significantly greater odds of being adherent than patients initiating sitagliptin (AOR = 1.213, 95% CI = 1.161, 1.266). Results for the 2-year follow-up period and for subsets of monotherapy and non-mail-order patients were of similar direction, magnitude, and significance.

Figures 4 and 5 present adjusted curves for time to discontinuation. In adjusted Cox proportional hazards models, the adjusted hazard of discontinuation was approximately 40% higher among SU initiators (adjusted hazard ratio [AHR] = 1.390, 95% CI = 1.363, 1.418) and TZD initiators (AHR = 1.402, 95% CI = 1.377, 1.427) compared with DPP-4i initiators during 1-year follow-up. Over the same time period, patients who initiated sitagliptin were significantly more likely to discontinue than those who initiated...
Table 3  Unadjusted adherence and persistence

|                     | Full study population | DPP-4i initiators | SU initiators | TZD initiators | DPP-4i initiators: Saxagliptin | DPP-4i initiators: Sitagliptin |
|---------------------|-----------------------|-------------------|---------------|----------------|--------------------------------|--------------------------------|
|                     | N                     | %                 | N             | %             | N                             | %                              |
| 1-year follow-up    |                       |                   |               |               |                                |                                |
| Adherent patients   | 100,092               | 42.0              | 29,070        | 47.3          | 55,597                        | 41.2                           |
| Persistent patients | 116,283               | 48.8              | 33,765        | 55.0          | 64,503                        | 47.8                           |
| 2-year follow-up    |                       |                   |               |               |                                |                                |
| Adherent patients   | 46,505                | 34.6              | 12,578        | 40.5          | 26,157                        | 34.6                           |
| Persistent patients | 45,123                | 33.6              | 12,410        | 39.9          | 25,318                        | 33.5                           |
| Monotherapy patients—1-year follow-up | N = 88,382 | N = 19,097 | N = 38,407 | N = 14,278 | N = 3,980 | N = 14,719 |
| Adherent patients   | 35,616                | 40.3              | 8,878         | 46.5          | 21,658                        | 39.4                           |
| Persistent patients | 40,768                | 46.1              | 10,050        | 52.6          | 24,926                        | 45.3                           |
| Non-mail-order patients—1-year follow-up | N = 206,226 | N = 49,848 | N = 121,503 | N = 34,875 | N = 9,559 | N = 39,614 |
| Adherent patients   | 81,596                | 39.6              | 21,966        | 44.1          | 47,725                        | 39.3                           |
| Persistent patients | 96,981                | 47.0              | 26,304        | 52.8          | 56,368                        | 46.4                           |

Patients were considered to be adherent if the proportion of days covered by index drug class/index drug over the follow-up period was greater than or equal to 0.80. Patients were considered to be persistent if they did not discontinue (i.e., have a gap of >60 days) index drug class/index drug during the follow-up period. P values for all comparisons between DPP-4i and SU initiators, DPP-4i and TZD initiators were <0.001. All P values for saxagliptin and sitagliptin initiator comparison were <0.05, with the exception of 1-year adherence for monotherapy patients (P = 0.184).

DPP-4i Dipeptidyl peptidase-4 inhibitor, SU Sulfonylurea, TZD Thiazolidinedione
saxagliptin (AHR = 1.159, 95% CI = 1.122, 1.195). As with adherence, these associations held over the 2-year follow-up period and within the two patient subgroups.

**DISCUSSION**

In this US administrative claims-based analysis of adults with T2DM, patients who initiated a DPP-4i had significantly better adherence to and persistence with index drug class compared with patients who initiated an SU or TZD. Among DPP-4i initiators, saxagliptin patients had significantly better adherence and persistence than sitagliptin patients. These associations were observed over a 1-year follow-up period, a 2-year follow-up period, and within sub-samples of patients initiating monotherapy and filling index prescription through a non-mail-order method.

Patients may discontinue their oral antidiabetic medication following the onset of adverse events particular to their medication,
such as hypoglycemia while on an SU [25, 26]. Additionally, clinicians may discontinue a medication when the patient develops a comorbidity that places the patient at greater risk for an adverse event with a particular medication, for example, a patient at increasing risk for fractures on a TZD [27].

Potential explanations for the observed association between initiation of DPP-4is and better adherence and persistence include a better tolerability profile compared with SUs, which are associated with weight gain and hypoglycemia, and TZDs, which are associated with weight gain and bone fracture [3].
Differences noted between saxagliptin and sitagliptin may be due in part to a one-step dose adjustment for patients with renal impairment for saxagliptin, compared with a two-step dose adjustment for sitagliptin [28, 29]. However, more research utilizing additional types of data sources is needed to specifically test these hypotheses for the differences in adherence and persistence between drug classes and DPP-4i-specific drugs. An analysis conducted by Rathmann et al. [20] comparing DPP-4is and SUs reported results consistent with the findings presented here. Utilizing data from general practices in Germany, researchers found that the risk of discontinuation was higher for DPP-4i initiators

**Fig. 5** Adjusted curves for time to discontinuation by index drug for 1- and 2-year follow-up and within subsamples of monotherapy and non-mail-order patients for 1-year follow-up. **a** Adjusted curves and hazard ratios for discontinuation of index drug over 1-year follow-up period. **b** Adjusted curves and hazard ratios for discontinuation of index drug over 2-year follow-up period. **c** Adjusted curves and hazard ratios for discontinuation of index drug over 1-year follow-up period among monotherapy patients. **d** Adjusted curves and hazard ratios for discontinuation of index drug over 1-year follow-up period among non-mail-order patients. Models controlled for age, sex, presence of capitated services, payer, region, urbanicity, plan type, indicator for fixed-dose metformin index drug, indicator for index drug filled via mail order in monotherapy samples, index drug regimen (monotherapy, index drug plus additional non-insulin antidiabetic drugs, index drug plus insulin) in non-mail-order samples, baseline total expenditures and diabetes prescription expenditures, index diabetes medication class cost-sharing, baseline endocrinologist and cardiologist visits, baseline renal impairment, baseline macrovascular and microvascular disease, pregnancy during follow-up, baseline number of unique 3-digit ICD-9 diagnoses and Charlson Comorbidity Index. **AHR** Adjusted hazard ratio, **CI** Confidence interval, **DPP-4i** Dipeptidyl peptidase-4 inhibitor.
in the first 3 months and then fell below that of SU initiators for the remainder of follow-up, a maximum of 2 years [20]. Using US health insurance claims, Curkendall et al. [10] evaluated adherence and persistence over 1-year follow-up between patients initiating saxagliptin and patients initiating an SU, TZD, or GLP-1RA. These investigators found that patients initiating saxagliptin had greater odds of being adherent and lower risk of discontinuation than patients initiating medications within the other three drug classes [10]. No analyses over a 2-year follow-up or within monotherapy or non-mail-order subgroups were conducted [10]. Beyond adherence and persistence, Rathmann et al. [20] found that compared with SUs, initiating a DPP-4i was associated with lower risk of hypoglycemia and newly diagnosed macrovascular outcomes, including coronary heart disease, myocardial infarction, and stroke. This study has limitations that should be recognized. Administrative claims data are not collected for research purposes, and the diagnostic coding on administrative claims is recorded by healthcare providers to support reimbursement. Consequently, misclassification of certain study variables, comorbid conditions in particular, may have occurred. Similarly, unmeasured confounding may exist, as clinical variables such as HbA1c level and duration of disease, socioeconomic characteristics such as income and education, and other factors such as provider characteristics were not available for inclusion in the claims databases. As this was a retrospective, observational analysis, causal inferences should be drawn with caution. Calculations of adherence and persistence were based on the service date and days’ supply from outpatient prescription drug claims and assume that patients took the medication as directed. It cannot be determined whether patients skipped doses or discontinued the medication prior to the end of the current days’ supply. Any prescriptions obtained without generating a health insurance claim were not captured in this analysis. The reason for discontinuation (e.g., adverse events, failure to achieve treatment goals, etc.) is not available in claims databases. Lastly, the MarketScan Commercial and Medicare Supplemental databases contain data only on patients covered through employers, private commercial insurance, or employer-sponsored supplemental Medicare. Thus, findings from the study may not be generalizable to the whole US Medicare population, Medicaid population, or the uninsured.

CONCLUSIONS

Patients initiating DPP-4is were more adherent to and persistent with the index drug class than patients initiating SUs or TZDs. These findings were robust to sensitivity analyses. Similarly, patients initiating saxagliptin were more likely to be adherent and persistent than patients initiating sitagliptin. These associations between DPP-4is and adherence and persistence support the use of DPP-4i medications, specifically saxagliptin, in management of T2DM, as adherence and persistence are important to experiencing optimal clinical outcomes.

ACKNOWLEDGMENTS

The authors would like to thank Jerry Kagan of Truven Health Analytics for programming and data analyses. Funding for this analysis was provided by Bristol-Myers Squibb and AstraZeneca. AstraZeneca funded the article processing charges associated with this publication. Parts of this manuscript were
presented previously at the Academy of Managed Care Pharmacy 26th Annual Meeting & Expo in Tampa, Florida, and the International Society for Pharmacoeconomics and Outcomes Research 19th Annual International Meeting in Montreal, Canada. All named authors meet the ICMJE criteria for authorship for this manuscript, take responsibility for the integrity of the work as a whole, and have given final approval to the version to be published.

Conflict of interest. Ms. Farr, Dr. Smith, and Mr. Johnston are employees of Truven Health Analytics, which was providing funding to conduct this study. Dr. Curkendall was an employee of Truven Health Analytics at the time the study was conducted. Drs. Sheehan and Kalsekar are employees of AstraZeneca, which funded this analysis.

Compliance with ethics guidelines. The analyses presented in this article are based on previously collected, de-identified data and do not contain any new studies with human subjects performed by any of the authors.

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