A Pooled Analysis of the Efficacy and Safety of Saxagliptin as Monotherapy in Patients with Type 2 Diabetes

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Abstract

Objective: Saxagliptin is a once-daily, orally administered dipeptidyl peptidase-4 inhibitor, approved for the treatment of type 2 diabetes mellitus (T2DM). In 4 double-blind, placebo-controlled, phase 3 trials, saxagliptin 2.5 or 5 mg significantly reduced glycated hemoglobin (HbA1c) from baseline at 24 weeks. A pooled analysis of these clinical trials was conducted to assess the therapeutic profile of saxagliptin monotherapy.

Methods: A post hoc pooled analysis of 4 saxagliptin monotherapy trials in patients with T2DM was conducted to determine the consistency of treatment effects, assessed as change from baseline in HbA1c at week 24, in patient subgroups stratified by race, sex, age, and baseline HbA1c. Secondary end points included change from baseline at week 24 in fasting plasma glucose and proportion of patients achieving HbA1c <7%. Safety assessments included adverse event reports, laboratory test results, and vital sign measurements.

Results: At week 24, saxagliptin 2.5 mg, saxagliptin 5 mg, and placebo decreased HbA1c from baseline by −0.66%, −0.64%, and −0.13% respectively (P<0.0001 for each saxagliptin dose vs placebo). The treatment effects of saxagliptin 2.5 mg on HbA1c were consistent across subgroups, and no treatment interactions were observed, with the exception of baseline HbA1c (P=0.003). The overall occurrence of adverse events was similar among groups (66.0% saxagliptin 2.5 mg; 53.0% saxagliptin 5 mg; 45.3% placebo). The incidence of hypoglycemic events was low and comparable among groups, and no cases of confirmed hypoglycemia were reported.

Conclusions: Consistent with findings from individual studies, this pooled analysis showed saxagliptin to significantly improve glycemic measures compared with placebo. Outcomes confirmed saxagliptin treatment was associated with a weight-neutral effect and a low risk for hypoglycemia. Based on the overall findings, saxagliptin has a favorable benefit:risk profile and may be considered an alternative first-line therapy for patients with T2DM in whom metformin is contraindicated or not tolerated.

Keywords: Type 2 diabetes mellitus; Saxagliptin; DPP-4 inhibitor; Glycated hemoglobin; Hypoglycemia; Metformin; Glycemic control

Introduction

Diabetes mellitus is a highly prevalent, chronic condition affecting more than 29 million people in the United States [1]. Diabetes affects numerous organ systems, and is associated with serious complications, including heart disease, stroke, kidney failure and lower-limb amputations [1].

In adults, type 2 diabetes mellitus (T2DM), which is characterized by insulin resistance with a progressive insulin secretion defect and β-cell failure, accounts for more than 90% of all diagnosed cases [1-3]. The natural history of T2DM has been well described, and it is now understood that multiple pathophysiologic defects contribute to hyperglycemia [3].

There is a well-established correlation between hyperglycemia, as measured by elevated glycated hemoglobin (HbA1c) levels, and an increased risk for microvascular and macrovascular complications [4,5]. Therefore, attaining good glycemic control may help prevent or delay long-term complications of T2DM, especially microvascular disease [1,6]. In an observational study, a reduction in the risk of T2DM complications was observed for every 1% decrease in HbA1c. The lowest risk for disease-related complications was observed in patients with HbA1c levels below 6.0% [4].

Recent guidelines from the American Diabetes Association and European Association for the Study of Diabetes therefore recommend a therapeutic target of <7% HbA1c for most adults with T2DM and a more stringent target of <6.5% if this can be achieved without significant hypoglycemia or other treatment-related adverse effects [5]. However, less stringent HbA1c goals may be more appropriate for patients with a limited life expectancy, history of severe hypoglycemia, and extensive comorbid conditions [5]. It should be emphasized that glycemic targets should be individualized and tailored to the patient’s preferences and needs [5].

There is a wide array of medications available to treat T2DM [7]. Aside from insulin, most medications are expected to reduce HbA1c levels by 0.5% to 2.0% when administered as monotherapy [8]. A reduction in HbA1c of 0.5% is generally considered to be a clinically meaningful and therapeutic response [9,10]. Metformin, a biguanide, is the preferred initial pharmacologic therapy for T2DM because it has a long-standing evidence base of clinical efficacy and no associated risk for weight gain or hypoglycemia [2,11]. However, metformin is not

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appropriate for all patients. It is contraindicated in patients with renal insufficiency and is commonly associated with gastrointestinal (GI) side effects, particularly in the early stages of treatment [12].

Dipeptidyl peptidase-4 (DPP-4) inhibitors are a newer class of antihyperglycemic therapy that targets the incretin system. The incretin hormones, glucagon-like peptide 1 (GLP-1) and glucose-dependent insulinotropic peptide (GIP), which are released from the intestinal tract, facilitate postprandial insulin secretion from the pancreas and account for approximately 50% of the postprandial insulin response [13]. GLP-1 has been shown to have multiple physiologic benefits, including stimulating glucose-dependent insulin secretion, decreasing glucagon secretion, and slowing gastric emptying [13,14]. Native GLP-1 and GIP, however, are degraded within minutes by the DPP-4 enzyme [15]. Research has shown DPP-4 expression and activity to be higher in patients with T2DM compared with healthy controls [16]. By slowing the degradation of GLP-1 and GIP, DPP-4 inhibitors thus sustain the incretin effects of increasing postprandial insulin secretion and suppressing glucagon production [13]. Preclinical data have also suggested that DPP-4 inhibitors may limit ß-cell apoptosis and stimulate ß-cell proliferation [17]. Current guidelines support the addition of DPP-4 inhibitors for patients experiencing suboptimal glycemic control with metformin alone [2] or as a first-line alternative if metformin monotherapy cannot be tolerated [5,11].

Saxagliptin is a once-daily (QD), orally administered, potent, competitive, and reversible DPP-4 inhibitor [18,19]. Saxagliptin is currently approved as an adjunct to diet and exercise to improve glycemic control in adults with T2DM [20,21]. As appropriate, saxagliptin may also be used in combination with metformin, sulfonylureas (SUs), thiazolidinediones, insulin, and metformin plus an SU [5,11].

In 4 individual phase 3 trials, saxagliptin monotherapy (2.5 or 5 mg QD) significantly reduced HbA1c from baseline at 24 weeks compared with placebo [10,22-24]. Consistent with the characteristics of a DPP-4 inhibitor [5,11], saxagliptin was associated with a low risk of hypoglycemia and with weight neutrality [10,22-24]. A pooled analysis of these 4 phase 3 saxagliptin monotherapy trials was performed to better characterize the therapeutic profile of saxagliptin monotherapy for the treatment of T2DM.

Methods

This was a post hoc pooled analysis of all of the phase 3, controlled, saxagliptin monotherapy trials in patients with T2DM (clinical trial registration: NCT00918879, NCT00316082, NCT00698932, NCT00121641). Each of the 4 studies had a similar study design and a primary end point of change from baseline in HbA1c at week 24. Individual study methodology has been previously published [10,22-24]. Briefly, the studies included patients with T2DM who were ≥ 18 years of age and treatment-naïve, who had early-stage disease (HbA1c, 7%-10%), and who might otherwise seek first-line treatment with metformin [10,22-24]. All patients who met certain criteria for poor glycemic control were allowed rescue medication with metformin. In each study, patients with significant cardiovascular (CV) disease, or significant or unstable renal disease were excluded. Of note, 2 of the 4 studies included only the saxagliptin 5-mg dose and were conducted in Asian populations [10,23].

All studies were performed in accordance with guidelines set forth by the International Conference on Harmonization and Declaration of Helsinki. Study protocol and amendments were approved by the individual review board/independent ethics committee for each participating site. All patients provided informed written consent.

Study end points and assessments

Pooled assessments were performed for saxagliptin 2.5 mg, saxagliptin 5 mg, and placebo groups. The focus of this analysis was the primary end point of change from baseline in HbA1c at week 24. Subgroup analyses were performed to assess the consistency of treatment effects. Patients were stratified by race (white, Asian, black, other), sex, age (<65 or ≥ 65 years), and baseline HbA1c (<8%, 8%-<9%, or ≥ 9%). Secondary end points included in this analysis were change from baseline in fasting plasma glucose (FPG) at week 24, proportion of patients achieving a therapeutic glycemic response (defined as HbA1c <7%), and change from baseline in postprandial glucose (PPG) area under the curve (AUC), and 2-hour PPG. For all efficacy evaluations, data were pooled for the saxagliptin 2.5-mg groups, saxagliptin 5-mg morning (AM) groups, and placebo groups. Data for patients who were titrated from saxagliptin 2.5 mg to 5 mg, received saxagliptin 10 mg, or received saxagliptin 5 mg in the evening (PM) were not included. All pooled efficacy analyses were performed on data collected before patients received rescue medication.

Safety and tolerability assessments were based on adverse event (AE) reports, laboratory test results, vital sign measurements, electrocardiogram, and physical examinations. Treatment-emergent AEs were defined as any new, untoward medical occurrence or worsening of a preexisting medical condition, regardless of relationship to the study treatment. A serious AE (SAE) was defined as any untoward medical condition that was fatal, life threatening, resulted or prolonged inpatient hospitalization, caused significant disability, resulted in a congenital anomaly or birth defect, or was considered an important medical event.

Safety events of special interest included those that were of potential importance to antihyperglycemic agents, were relevant to the mechanism of action of DPP-4 inhibitors or the safety profile of other DPP-4 inhibitors, or were selected based on preclinical and clinical findings from the saxagliptin clinical development program. AEs of special interest included hypoglycemia, infections (including opportunistic), GI-related AEs, CV events, hypersensitivity reactions, lymphopenia, thrombocytopenia, pancreatitis, skin disorders, and bone fractures. A confirmed hypoglycemic event was characterized as having symptoms of hypoglycemia and fingerstick glucose ≤ 50 mg/dL.

For the safety analysis, data for the saxagliptin 2.5- to 5-mg titration group were pooled with the 2.5-mg groups, and data from the saxagliptin 5-mg PM group were pooled with 5-mg AM groups; both groups were compared with the pooled placebo group. Data from the saxagliptin 10-mg group were not included.

Statistical analyses

The primary and secondary efficacy end points, including treatment by subgroup interactions, were evaluated using an analysis of covariance model that used last observation carried forward to account for missing data. Supportive efficacy analyses using repeated measures were also conducted.

Results

A total of 1452 randomized patients (saxagliptin 2.5 mg, n=247; saxagliptin 5 mg, n=645; placebo, n=560) were included in this analysis. Of these patients, 73% completed 24 weeks of treatment.
in the saxagliptin 2.5-mg group as did 84% in the saxagliptin 5-mg group and 81% in the placebo group. In each treatment group, early discontinuation before 24 weeks was primarily due to perceived lack of efficacy with study medication or withdrawal of consent.

With the exception of race, patient demographics and disease characteristics were generally well balanced across treatment groups (Table 1). The mean age was 52 years, and 52% of patients were men. The mean weight was 77.4 kg. Patients generally had recent-onset T2DM (mean duration at baseline, 1.5 years) and early-stage disease (mean baseline HbA1c, 8.1%; mean baseline FPG, 164 mg/dL).

**Efficacy analyses**

At week 24, saxagliptin 2.5 mg, saxagliptin 5 mg, and placebo decreased HbA1c from baseline by −0.66%, −0.64%, and −0.13%, respectively (P<0.0001 for each saxagliptin dose vs placebo; Figure 1).

The treatment effects of saxagliptin on HbA1c were consistent across patient subgroups, and no treatment interactions were observed in subgroups categorized by race, sex, or age. However, a significant interaction was observed for baseline HbA1c (P=0.003; Figure 2).

Changes from baseline in FPG at week 24 were statistically significant with each saxagliptin dose compared with placebo (P<0.0001; Figure 3). Saxagliptin 5 mg also significantly increased the proportion of patients who achieved a therapeutic glycemic response (HbA1c <7%) at week 24 compared with placebo (P<0.0001), although only a numerical increase was observed with saxagliptin 2.5 mg (Figure 4). Changes from baseline in PPG AUC and 2-hour PPG at week 24 were statistically significant with each saxagliptin dose group versus placebo (P<0.001; Figure 5). Supportive analyses using repeated measures were consistent with the primary analysis of change from baseline in HbA1c and FPG at week 24 and further demonstrated consistent treatment effects on HbA1c across patient subgroups (Supplementary Appendix).

At week 24, minor decreases in weight (mean [95% CI]) from baseline were observed in the saxagliptin 2.5-mg group (−0.9 [−1.3, −0.5] kg), saxagliptin 5-mg group (−0.4 [−0.6, −0.2] kg), and placebo group (−1.3 [−1.5, −1.0] kg).

**Safety and tolerability**

Based on the pooled safety data, mean exposure was 20.6 weeks in the saxagliptin 2.5-mg group, 21.8 weeks in the saxagliptin 5-mg group, and 20.9 weeks in the placebo group. The majority of patients (74%) were exposed to study medication for ≥23.7 weeks, consistent with the primary analysis of change from baseline in HbA1c and FPG at week 24 and further demonstrated consistent treatment effects on HbA1c across patient subgroups (Supplementary Appendix).

The overall proportion of patients reporting AEs was similar across treatment groups, and no dose-response was observed (saxagliptin 2.5 mg, 66.0%; saxagliptin 5 mg, 53.0%; placebo, 45.3%; Table 2). The most common AEs (≥3% of patients) occurring in the saxagliptin dose groups were upper respiratory tract infection, urinary tract infection, and nasopharyngitis (Table 2). With the exception of hypoglycemia (discussed below), arthralgia was the only AE with an incidence of ≥

![Adjusted Mean (95% CI) change from baseline in HbA1c at week 24. Number of patients by treatment group and HbA1c outcomes. Data collected before rescue treatment. Change from baseline in HbA1c was evaluated using an ANCOVA model with last observation carried forward to account for missing data. At week 24, saxagliptin 2.5 mg and saxagliptin 5 mg significantly decreased HbA1c from baseline compared with placebo (P<0.0001 for each saxagliptin dose). ANCOVA=analysis of covariance; HbA1c=glycated hemoglobin. Number of randomized patients with baseline and week 24 values.](image-url)

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The occurrence of SAEs and AEs leading to discontinuation was low (<5% and <3%, respectively, in any treatment group; Table 2). SAEs related to cardiac disorders were reported for 0.8% of patients in each saxagliptin group and for 0.2% of patients in the placebo group. In each case, the patient had either preexisting CV disease or multiple CV risk factors, and each event was considered not related or unlikely related to study treatment. Overall, no specific type of SAE or AE leading to discontinuation prevailed (Supplementary Appendix).

There were two deaths reported during the 24-week treatment period, 1 due to pneumococcal sepsis in the saxagliptin 2.5-mg group and 1 due to myocardial infarction (MI) in the saxagliptin 5-mg group. Neither death was considered related to study treatment. Two additional deaths were reported after the 24-week treatment period, 1 due to cerebral hemorrhage in the placebo group and 1 due to pancreatic and hepatic cancer in the saxagliptin 2.5-mg group (approximately 16 months after treatment discontinuation). The incidence of reported hypoglycemic events was low and similar between the saxagliptin 2.5- and 5-mg groups (4.0% and 3.0%, respectively, vs 1.6% with placebo; Table 2). However, no hypoglycemic event was considered serious, 2% in the saxagliptin 5-mg group with a difference of >1% compared with placebo (2.6% vs 1.3%, respectively).

Figure 2: Adjusted mean (95% CI) change from baseline in HbA1c at w24: subgroup analysis. Subgroup analyses were performed to assess the consistency of treatment effects on change from baseline in HbA1c. Patients were stratified by race (white, Asian, black, other) sex, age (<65 or ≥ 65 years), and baseline HbA1c (<8%, 8%–<9%, or ≥9%). Only a few patients identified as “other” in the subgroup for race (saxagliptin 2.5 mg, n=3; saxagliptin 5 mg, n=4; placebo, n=7). The adjusted mean change (95% CI) from baseline in HbA1c in these patients was −1.96 (−3.06, −0.86) in the saxagliptin 2.5 mg group, −1.78 (−2.74, −0.82) in the saxagliptin 5 mg group, and −0.06 (−0.80, 0.68) in the placebo group at week 24. No significant treatment interaction was observed, with the exception of baseline HbA1c (P=0.003). HbA1c=glycated hemoglobin. Number of randomized patients with baseline and week 24 values.

Figure 3: Adjusted mean (95% CI) change from baseline in FPG (mg/dL) at week 24. Number of patients by treatment group and FPG outcomes. Data collected before receiving rescue medication. Change from baseline in FPG was evaluated using ANCOVA model with last observation carried forward to account for missing data. At week 24, saxagliptin 2.5 mg and 5 mg significantly reduced FPG from baseline compared with placebo (P<0.0001 for each saxagliptin dose). ANCOVA=analysis of covariance; FPG=fasting plasma glucose. Number of randomized patients with baseline and week 24 values.

Figure 4: Proportion of patients achieving a therapeutic glycemic response (HbA1c <7%) at week 24. Proportion of patients achieving a therapeutic glycemic response. Data collected before receiving rescue medication. The proportion of patients achieving HbA1c was evaluated using ANCOVA model with last observation carried forward to account for missing data. Compared with placebo, saxagliptin 5 mg significantly increased the proportion of patients who achieved HbA1c <7% at week 24 (P<0.0001) and saxagliptin 2.5 mg was associated with a numerical increase. ANCOVA=analysis of covariance; HbA1c=glycated hemoglobin; NS=not significant. Number of randomized patients with week 24 values.
led to treatment discontinuation, or required medical management or third-party assistance. There were no cases of confirmed hypoglycemia.

There was no increased risk for infection with saxagliptin 5 mg compared with placebo (22.1% vs 18.1%, respectively), although a higher occurrence of infection was observed with saxagliptin 2.5 mg (30.4%). There was no also notable association between saxagliptin and GI tolerability issues (Table 2 and Supplementary Appendix). Diarrhea was the most commonly reported GI-related AE; however, the overall incidence was low (<5% in each saxagliptin group). Two serious GI-related AEs, abdominal pain and intestinal obstruction, were reported in 1 patient each in the saxagliptin 5-mg group. Two GI-related AEs leading to treatment discontinuation were also reported in the saxagliptin 5-mg group: 1 patient each reporting dry mouth and gastric disorder.

Hypersensitivity reactions occurred infrequently (<3% in any treatment group), and no events were considered serious or led to treatment discontinuation. The incidence of individual AEs related to cardiac disorders was low: less than 1% in any treatment group. CV SAEs reported in the saxagliptin 2.5-mg, saxagliptin 5-mg, and placebo groups included unstable angina (0.4%, 0.3%, and 0, respectively), atrial fibrillation (0.4%, 0.2%, and 0), MI (0, 0.2%, and 0), cerebrovascular accident (0, 0.3%, and 0), supraventricular tachycardia (0, 0.2%, and 0), and coronary artery disease (0, 0, and 0.2%).

All other AEs of special interest, including opportunistic infections, lymphopenia, thrombocytopenia, pancreatitis, skin disorders, and bone fracture, occurred in <1% of patients in any treatment group. Subgroup analyses by sex, age, and race showed no difference in the type or incidence of AEs across patients. Overall, there was no notable difference in the incidence of marked laboratory abnormalities among treatment groups.

**Discussion**

Findings from this pooled analysis of 4 phase 3 clinical trials provide greater insight on the efficacy, safety, and tolerability of saxagliptin in Type 2 Diabetes.
patients with T2DM. Consistent with the individual studies [10,22-24], saxagliptin 2.5 and 5 mg significantly reduced HbA1c from baseline at week 24 compared with placebo.

Saxagliptin produced a clinically meaningful and therapeutic response [9,10], as evidenced by changes from baseline in HbA1c that exceeded those achieved with placebo by ≥ 0.5%. Secondary efficacy outcomes supplement these findings and further demonstrate the clinical benefit of saxagliptin on glycemic measures.

Given the baseline demographics and characteristics included in the individual studies, pooling allowed a comprehensive evaluation of saxagliptin in a diverse study population. The treatment effect of saxagliptin was consistent across patient subgroups categorized by race, sex, and age. The only notable treatment interaction was observed in patients with higher HbA1c at baseline (≥ 9%). However, a greater treatment effect in patients with higher baseline HbA1c levels is consistent with previous observations with oral antidiabetic drugs [25].

Pooled safety data were consistent with the known profile of saxagliptin and DPP-4 inhibitors [5,11], and no new or unexpected safety issues emerged. Although metformin is widely accepted as the preferred first-line pharmacologic treatment in T2DM [5,11], lack of tolerability and contraindications may bar its use in some patients. For example, nearly 90% of patients starting metformin therapy report GI symptoms (eg, diarrhea, heartburn, nausea), the presence of which is associated with decreased quality of life and adherence [26]. This may be especially problematic for patients with diabetes who have preexisting GI complications caused by abnormal intestinal motility secondary to diabetic autonomic neuropathy [27]. Metformin is contraindicated in patients with renal dysfunction or disease (serum creatinine levels ≥ 1.4 in women and ≥ 1.5 in men) [12]. Because increased age is associated with reduced renal function, this could limit the use of metformin in some elderly patients [12]. In contrast, saxagliptin is not associated with GI tolerability issues [10,22-24], and the incidence of AEs and hypoglycemic events with saxagliptin 2.5 mg (the recommended dose for patients with moderate or severe renal impairment or end-stage renal disease) was similar to that of placebo in patients with renal impairment [28]. Therefore, saxagliptin could be considered an alternative treatment in patients for whom metformin is not tolerated or contraindicated.

Additional findings from the saxagliptin clinical development program supplement those reported here and provide greater insight on treatment effects in a patient population commonly burdened with serious comorbidities. For example, the Saxagliptin Assessment of Vascular Outcomes Recorded in patients with diabetes mellitus (SAVOR) trial included 16,492 patients with T2DM and established CV disease or multiple CV risk factors, many of whom also had mild-to-moderate renal impairment [29]. Findings showed there was no difference in the occurrence of the primary composite end point of CV death, myocardial infarction, or stroke between saxagliptin (7.3%) and placebo groups (7.2%; P=0.99 for superiority, P<0.001 for noninferiority) in patients with CV disease or risk factors. The occurrence of the secondary end point (primary composite end point plus hospitalization for heart failure, coronary revascularization, or unstable angina) also did not differ significantly between saxagliptin and placebo groups (12.8% and 12.4%, respectively; P=0.66 for superiority, P=0.001 for noninferiority). Analysis of the individual components of the secondary end point showed a higher incidence of hospitalization for heart failure in patients treated with saxagliptin compared with placebo (3.5% vs 2.8%; P=0.007). However, this finding was unexpected, and the statistical analysis of this secondary component was not adjusted for multiplicity, which could have resulted in a false-positive result, and thus warrants further study [29]. It may be noted that deaths due to heart failure were similar for saxagliptin and placebo (0.5% each). Supplementary findings also showed HbA1c was significantly lower with saxagliptin versus placebo at 1 year, 2 years, and end of treatment (P<0.001 for all) [30]. Additional ongoing trials of DPP-4 inhibitors in large populations of patients with T2DM and diverse medical histories will help further establish the CV risk profile of these treatments [31-33].

Although this analysis was somewhat limited by the inherent differences among the 4 studies (eg, differences in doses studied, patient ethnicities), these differences do not preclude the value of pooling these data. Numerical comparison of the relative efficacy from the pooled analyses for the 2.5-mg saxagliptin dose should be interpreted with caution because this dose was only examined in 2 of the 4 studies. However, no treatment-by-study interaction was observed for any of the doses in the pooled analysis, and saxagliptin 2.5 mg produced clinically relevant and statistically significant reductions in HbA1c that were similar to those noted with the saxagliptin 5-mg dose.

Conclusions

This pooled analysis provides a comprehensive assessment of the therapeutic profile of saxagliptin monotherapy and further establishes its efficacy and tolerability in patients with T2DM. Based on these overall findings, in addition to the findings from the SAVOR trial, saxagliptin is associated with a generally favorable benefit:risk profile, and may be considered an alternative first-line therapy for patients with T2DM in whom metformin is contraindicated or not tolerated.

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