The durability of sitagliptin in elderly patients with type 2 diabetes

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Aim: To evaluate the durability of sitagliptin and to assess changes in clinical chronic complications following sitagliptin monotherapy for 48 months in elderly patients with type 2 diabetes mellitus (T2DM).

Subjects and methods: We enrolled 76 drug-naive patients (40 women and 36 men; mean age: 71.3±11.7 years) with T2DM who received 25–100 mg of sitagliptin therapy from an outpatient clinic. The observational period for each patient was >48 months, beginning at the time sitagliptin therapy was initiated. The following were measured or performed at the beginning of each year: body mass index; serum total cholesterol, low-density lipoprotein, high-density lipoprotein; triglyceride levels; creatinine (Cr) levels; urine albumin and urine Cr; nonmydriatic fundusgraphy; and semiquantified neuropathy. The fasting plasma glucose and glycated hemoglobin (HbA₁c) was measured every 3–6 months.

Results: The change in HbA₁c was significantly reduced after 6 months of therapy (7.1%±0.8% to 6.3%±0.2%). No changes in fasting plasma glucose, Cr, serum total cholesterol, triglyceride, low-density lipoprotein, high-density lipoprotein, body mass index, and microvascular complications were apparent. Using repeated measures to test the sequential changes in HbA₁c from month 6 to month 48, the test of within-subjects effect was not significant (P=0.34).

Conclusion: Sitagliptin has a durable effect and stabilizes microvascular complication progression in elderly patients. This study can provide useful data for clinicians and health care professionals using sitagliptin monotherapy in the treatment of elderly patients with T2DM.

Keywords: type 2 diabetes mellitus, sitagliptin, oral antidiabetic drugs, dipeptidyl peptidase IV inhibitor

Introduction

Type 2 diabetes mellitus (T2DM) is a progressive disease due to the gradual dysfunction of pancreatic β-cells.1 A single oral antidiabetic drug (OAD) for blood glucose control in patients with T2DM has a 50% failure rate after 3 years and a 75% failure rate after 9 years.1 In clinical practice, the ideal OAD should have the following effects: efficacy; no weight gain; no risk of hypoglycemia; no risk of cardiovascular events; and preserving pancreatic β-cells. In the elderly, the risks of renal insufficiency and hypoglycemia are higher than in younger patients with T2DM. There are limited data regarding the use of pharmacologic agents in elderly patients with T2DM. According to the recommendation of American Diabetic Association and European Association for the Study of Diabetes, dipeptidyl peptidase IV (DPP-4) inhibitors have an intermediate effect on patients with T2DM, a lower risk of hypoglycemia, a lower risk of weight gain, and fewer side effects than other OADs and insulin.2 No increase in cardiovascular events has been associated with DPP-4 inhibitors based on two recent large clinical trials.3,4
The mechanism of action of DPP-4 inhibitors involves augmentation of the incretin axis by increasing endogenous glucagon-like peptide-1 (GLP-1) and glucose-dependent insulinotropic peptide active form concentrations, both of which result in the stimulation of glucagon-dependent insulin and the inhibition of glucagon; thus, a reduction in blood glucose follows.\textsuperscript{3–7} When compared with other established OADs in several randomized controlled trials,\textsuperscript{8–12} sitagliptin demonstrated noninferiority in efficacy endpoints. In addition to metformin, sulfonylurea (SU) is another popular OAD based on a Taiwanese study.\textsuperscript{13} SU, which directly stimulates pancreatic \(\beta\)-cells for insulin secretion, leads to earlier \(\beta\)-cell exhaustion. The thiazolidinediones can improve insulin resistance and durability in glycemic control, but they may lead to an increased risk of cardiovascular events and bladder cancer.\textsuperscript{14–16} Heart function is less in the elderly than in young adults; the risk for cancer is also high in the elderly. Metformin-induced lactic acidosis is also a concern. Sitagliptin was the first DPP-4 inhibitor used to control blood glucose in Taiwan. A previous meta-analysis confirmed the safety of sitagliptin in elderly patients with T2DM.\textsuperscript{17} According to a previous study, sitagliptin decreases \(\beta\)-cell apoptosis and is associated with a decrease of \(\beta\)-cell function deterioration.\textsuperscript{18} In human studies, however, the durability of sitagliptin is seldom mentioned. In all studies involving sitagliptin as monotherapy, the duration of treatment was \(\leq 1\) year.\textsuperscript{19} In the current study, we observed the blood glucose changes after monotherapy sitagliptin was used for 48 months in elderly patients with T2DM; we also evaluated the durability of sitagliptin and chronic complications.

**Subjects and methods**

**Subjects**

There were 76 patients with T2DM (40 women and 36 men; mean age: 71.3±11.7 years) who were first diagnosed and had participated in a diabetic share–care education program for \(\geq 6\) months. All of the patients were \(\geq 65\) years of age and had never received medical treatment before enrollment in the diabetes education program and study. The patients received a diabetic diet and share–care education every 3 months since T2DM was diagnosed, and sitagliptin monotherapy was prescribed. The inclusion criteria limited the study subjects to outpatients with T2DM from our Metabolism and Endocrinology Clinic who were treated with sitagliptin for diabetic control. The exclusion criteria included the following: type 1 diabetes mellitus; treatment with other antidiabetic agents; diabetes secondary to other causes, such as surgery, pharmaceutical products, malnutrition, and infections; and participation in a clinical trial or other clinical study during the index period. The case identification period or index period was defined as the time between January 1, 2009 and January 30, 2010. The observational period for each patient was \(>48\) months, beginning at the time that sitagliptin monotherapy was started. The dose of sitagliptin was based on the initial estimated glomerular filtration rate (eGFR), as follows: 100 mg/day for patients with an eGFR \(\geq 50\) mL/minute/1.73 m\(^2\); 50 mg/day for patients with an eGFR \(\geq 25\) mL/minute/1.73 m\(^2\) but \(<50\) mL/minute/1.73 m\(^2\) 50 mg/day; and 25 mg/day for patients with an eGFR \(<25\) mL/minute/1.73 m\(^2\).

**Methods**

Among this continuous sitagliptin user cohort, the following clinical and laboratory parameters were measured at baseline and every year thereafter: any self-reported hypoglycemia symptoms; body mass index; nonmydriatic fundusgraphy; semiquantification of neuropathy; serum levels of total cholesterol (TC), low-density lipoprotein (LDL), high-density lipoprotein (HDL), triglycerides (TG), and creatinine (Cr); and urine albumin and urine Cr. The blood lipid profile, including TC, LDL, HDL, and TG, were checked after fasting for \(>10\) hours. The comorbidity (dyslipidemia and hypertension) and chronic complications of T2DM at baseline and before the last follow-up evaluation were collected and analyzed. The glycated hemoglobin (HbA\(_{1c}\)) and fasting plasma glucose (FPG) levels were measured every 3–6 months in each patient. Protective sensation was assessed using a 5.07 Semmes–Weinstein (SW) nylon filament test (10 g monofilament) and vibration perception using 128 Hz Rydel Seiffer forks. Two observers confirmed all measurements. Neuropathy was evaluated based on the vibration perception threshold of the great toe and internal malleolus bilaterally. The patients were asked to indicate when the vibration sensation was lost. The vibration was determined on a 9-point grading scale (0/8–8/8) of a tuning fork. Patients were diagnosed with neuropathy if the vibration perception threshold was \(\leq 4\) on an 8-point scale.\textsuperscript{19} The vibration perception threshold yielded a sensitivity of 86% and a specificity of 76%.\textsuperscript{20}

Two examiners conducted the SW monofilament test when patients were unable to observe their feet. The 10 g SW monofilament was used on ten sites of the foot, as follows: dorsal surface of the foot between the base of the first and second toes; the first, third, and fifth toes; the first, third,
and fifth metatarsal heads; the medial and lateral midfoot; and the heel. The SW monofilament was pressed on the test site with enough pressure to bend the monofilament for 1 second. Patients were asked to answer “Yes” or “No” when they felt or did not feel the touch of the monofilament, respectively. If a patient did not feel the filament at >4 sites, neuropathy was diagnosed. The sensitivity and specificity at ten sites with a 10 g SW monofilament was 93.1% and 100%, respectively.21

Nonmydriatic retinal photography was performed using a CR6-45NM nonmydriatic retinal camera (Canon Inc., Tokyo, Japan) by two examiners. Retinopathy was diagnosed according to the American Academy of Ophthalmology Retina/Vitreous Panel (Preferred Practice Patterns Committee) and confirmed by one ophthalmologist.22 The effectiveness of the nonmydriatic retinal camera in the diagnosis of retinal disease has a sensitivity of 92% and a specificity of 96%.23 Nephropathy was defined as an increase in urine albumin/urea Cr >30 mg/dL in the first morning urine.24 The abbreviated Modification of Diet in Renal Disease (MDRD) Study Group equation was used to calculate the eGFR (mL/minute/1.73 m²), as follows:

\[
186.3 \times (\text{serum creatinine}^{-1.154}) \times (\text{age}^{-0.203}) \times 0.742 \text{ (if female)}.
\]

Comorbidities, including hypertension and dyslipidemia, were also recorded. Patients who received an antihypertensive drug for >1 week, except a diuretic for edema, were recorded as having hypertension. Dyslipidemia was defined based on a patient ever receiving statin therapy. Hypoglycemia was defined as a self-monitoring blood glucose level <70 mg/dL or self-reported hypoglycemia. This study was conducted according to the guidelines of the Declaration of Helsinki, and the research protocol was approved by the Ethics Committee of the Chang Gung Memorial Hospital.

Assays

The concentrations of FPG, TC, LDL, HDL, TG, and Cr were measured from serum using a commercial enzyme-linked immunosorbent assay kit and performed with an autoanalyzer (Hitachi 7250; Hitachi Ltd., Tokyo, Japan). Urine albumin concentrations were determined by immunonephelometry (Dade Behring Marburg GmbH, Marburg, Germany). The HbA\textsubscript{1c} level was measured by high-pressure liquid chromatography (Bio-Rad Laboratories Inc., Hercules, CA, USA).

Statistical analysis

The variables are summarized as the mean ± standard deviation. Differences in baseline clinical and biochemical characteristics were tested using a paired t-test. The repeated measures of a general linear model were used to analyze the sequential changes in HbA\textsubscript{1c}. A P-value <0.05 was considered statistically significant. All statistical operations were performed using the Statistical Package for Social Science program (SPSS for Windows, version 11.5; IBM Corporation, Armonk, NY, USA).

Results

The average duration of sitagliptin therapy for the 76 patients in the current study was 52.4±4.1 months. All of the patients were prescribed sitagliptin monotherapy for >48 months. Two patients used sitagliptin for >52 months. Seventeen, 21, and 38 patients received 25 mg/day, 50 mg/day, and 100 mg/day of sitagliptin, respectively. The change in dose of sitagliptin during the 48-month observation period was 69.4±24.3 mg to 67.3±23.6 mg (P=0.36). No hypoglycemic episodes were reported.

As shown in Table 1, the change in HbA\textsubscript{1c} from baseline was significant after 6 months and 48 months of treatment (7.1%±0.8%, 6.3%±0.2%, and 6.4%±0.4%, respectively; P<0.01). Changes in baseline renal function were not apparent after 48 months of treatment (Cr, 1.36±0.73 mg/dL and 1.32±0.67 mg/dL, respectively [P=0.20]; eGFR, 53.4±24.3 mL/minute/1.73 m² and 57.6±23.0 mL/minute/1.73 m², respectively; P=0.36). Body mass index was stable after sitagliptin monotherapy (26.5±4.5 kg/m² and 26.1±4.2 kg/m², respectively; P=0.45). There were no apparent changes in FPG (123±20.1 mg/dL and 112±24.4 mg/dL, respectively; P=0.06), TG (159±94.2 mg/dL and 161±111.4 mg/dL, respectively; P=0.72), TC (176.5±40.5 mg/dL and 178.8±38.7 mg/dL, respectively; P=0.61), HDL (47.1±12.6 mg/dL and 46.8±12.1 mg/dL, respectively; P=0.48), and LDL (99.1±35.1 mg/dL and 100.1±31.6 mg/dL, respectively; P=0.28). The prevalence of patients with hypertension and dyslipidemia did not appear to increase from the time of enrollment until the study had ended (22 versus 29, P=0.07; 30 versus 33, P=0.83, respectively). Patients received angiotensin receptor blockers or angiotensin-converting enzyme inhibitor therapy for hypertension; statin use for dyslipidemia did not increase (22 versus 23; 29 versus 30; all P=0.32). Chronic microvascular complications (retinopathy [1 versus 1] [the number
of patient with retinopathy before and after 48 months], nephropathy [19 versus 20, P=0.40], and neuropathy [24 versus 25; P=0.34]) were not significantly increased during the 48-month follow-up in our share–care program (Table 1). There were no hospital admissions for cardiovascular events during the observation period in the study group.

In the general linear model, we used repeated measures to test the sequential changes in HbA	extsubscript{1c} from month 6 to month 48. The test of within-subjects effect was not significant (P=0.34; Figure 1). We also used sex as a between-subjects factor; there were no apparent differences between the sexes (P=0.98).

The patients who received different doses of sitagliptin had apparent reductions in HbA	extsubscript{1c} from baseline during the first 6 months of treatment (25 mg, 7.3±1.3% and 6.0±0.8%, respectively; 50 mg, 7.1±0.8% and 6.3±0.6%, respectively; and 100 mg, 7.1±0.8% and 6.3±0.6%, respectively; all P<0.001); the HbA	extsubscript{1c} levels remained stable in patients receiving different doses of sitagliptin between month 6 and month 48 (Figure 2). There were no statistically significant differences in HbA	extsubscript{1c} concentrations between sitagliptin regimens (P=0.23 [tests of the between-subjects effect]).

**Discussion**

We studied 76 drug-naïve patients who received sitagliptin monotherapy for >48 months. The HbA	extsubscript{1c} level decreased significantly after 6 months of sitagliptin therapy, but the HbA	extsubscript{1c} levels fluctuated mildly; the HbA	extsubscript{1c} was well controlled for >48 months. Although the sample size was small, the durability of sitagliptin was demonstrated and there were no apparent changes in body weight, renal function, and the lipid profile. No hypoglycemic episodes were reported during monotherapy with sitagliptin. Before beginning sitagliptin therapy, all patients had already received lifestyle modification counseling, including diet education, for >6 months. Although neither a diet diary nor a daily activity record was maintained, lifestyle modification counseling clearly had a meaningful effect on the patients. In a review article, the author concluded that based on nonglycemic effects, such as the risk of hypoglycemia, weight gain, and durability, DPP-4 inhibitors may be considered an alternative to SUs, which directly stimulate pancreatic β-cells to secrete insulin. Of note, in the A Diabetes Outcome Progression Trial (ADOPT), the cumulative incidence of monotherapy failure at 5 years was 15% with rosiglitazone, 21% with metformin, and 34% with glyburide. Indeed, the increasing risk of cardiovascular events and bladder cancer may be of greater concern than the durability of thiazolidinediones.

Williams-Herman et al analyzed >10,000 subjects in clinical trials and concluded that sitagliptin was well tolerated for as long as 2 years. The longest published study of sitagliptin monotherapy was up to 104 weeks and has been reported

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**Table 1** The clinical and biochemical characteristics of patients before and after 48 months of sitagliptin therapy

<table>
<thead>
<tr>
<th></th>
<th>Before sitagliptin use</th>
<th>After sitagliptin use (48 months)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>71.3±11.7</td>
<td>36/40</td>
<td></td>
</tr>
<tr>
<td>Sex (male/female)</td>
<td>36/40</td>
<td>67.3±23.6</td>
<td>0.36</td>
</tr>
<tr>
<td>Dose (mg)</td>
<td>69.4±24.4</td>
<td>112±24.4</td>
<td>0.06</td>
</tr>
<tr>
<td>FPG (mg/dL)</td>
<td>123±20.1</td>
<td>1.3±0.73</td>
<td>0.20</td>
</tr>
<tr>
<td>Creatinine (mg/dL)</td>
<td>53.4±24.3</td>
<td>57.6±23.0</td>
<td>0.36</td>
</tr>
<tr>
<td>eGFR (mL/minute/1.73 m²)</td>
<td>7.1±0.8</td>
<td>6±0.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>26.5±4.5</td>
<td>26.1±4.2</td>
<td>0.45</td>
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<tr>
<td>TG (mg/dL)</td>
<td>159.2±94.2</td>
<td>161.1±111.4</td>
<td>0.72</td>
</tr>
<tr>
<td>TC (mg/dL)</td>
<td>176.5±40.5</td>
<td>178.8±38.7</td>
<td>0.61</td>
</tr>
<tr>
<td>HDL (mg/dL)</td>
<td>47.1±12.6</td>
<td>46.8±12.2</td>
<td>0.48</td>
</tr>
<tr>
<td>LDL (mg/dL)</td>
<td>99.1±35.1</td>
<td>100.1±31.6</td>
<td>0.28</td>
</tr>
<tr>
<td><strong>Comorbidities (number)</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
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<td>29</td>
<td>0.07</td>
</tr>
<tr>
<td>ARB or ACEI</td>
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<td>23</td>
<td>0.32</td>
</tr>
<tr>
<td>Dyslipidemia</td>
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<td>33</td>
<td>0.83</td>
</tr>
<tr>
<td>Statin</td>
<td>29</td>
<td>30</td>
<td>0.32</td>
</tr>
<tr>
<td>Retinopathy</td>
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<td>1</td>
<td></td>
</tr>
<tr>
<td>Neuropathy</td>
<td>24</td>
<td>25</td>
<td>0.34</td>
</tr>
<tr>
<td>Nephropathy</td>
<td>19</td>
<td>20</td>
<td>0.40</td>
</tr>
</tbody>
</table>

**Abbreviations:** FPG, fasting plasma glucose; eGFR, estimated glomerular filtration rate; HbA	extsubscript{1c}, glycated hemoglobin; BMI, body mass index; TG, triglycerides; TC, total cholesterol; HDL, high-density lipoprotein; LDL, low-density lipoprotein; ARB, angiotensin receptor blocker; ACEI, angiotensin-converting enzyme inhibitor.
by Williams-Herman et al\textsuperscript{28} (American Diabetes Association 2009, abstract 540P) in which the HbA\textsubscript{1c} decreased from 8.5\% to 7.4\% in 50 patients. In a review article involving published randomized, double-blind studies in patients with inadequately controlled T2DM (HbA\textsubscript{1c} =6.5\%–10\%), the studies were only 12–24 weeks in length.\textsuperscript{28} The reductions in baseline HbA\textsubscript{1c} ranged from 0.48\%–0.70\% among patients randomized to receive sitagliptin (all $P$, 0.001). In the present study, the enrolled patients did not have higher HbA\textsubscript{1c} concentrations (6.3\%–7.9\%) when compared to the 2-year study of Williams-Herman et al\textsuperscript{27} because we followed a previous study involving sitagliptin monotherapy, in which the HbA\textsubscript{1c} concentration was lowered by 0.6\%–0.7\% and the patients had a target HbA\textsubscript{1c} of 6.5\%–7\%.\textsuperscript{28} In the initial 6 months of therapy, the HbA\textsubscript{1c} concentration was reduced by approximately 0.8\% in the current study, followed by a 0.1\% increase from 6 months to 48 months of sitagliptin therapy. We also showed that a different dose of sitagliptin had the same HbA\textsubscript{1c}-lowering effect, and that there was no statistically significant difference in the HbA\textsubscript{1c} concentration between sitagliptin regimens. The patients enrolled in our study were all elderly and at higher risk for developing ischemic heart disease, congestive heart failure, and age-related nephropathy. Hypoglycemia may be a sequential problem or factor precipitating these complications. According to a previous study, sitagliptin has a lower risk of hypoglycemia and a better safety profile in elderly patients.\textsuperscript{29} Generally, metformin, GLP-1 analogs/DPP-4 inhibitors, and long-acting insulin are preferred over other treatment options (SU, glinides, and glitazones) in older patients.\textsuperscript{29} The cardiovascular safety of DPP-4 inhibitors has also been reported in two recent large clinical trials.\textsuperscript{3,4} In an open-label, randomized study involving elderly patients in Taiwan, Chien et al\textsuperscript{31} reported that 24 weeks of sitagliptin produced a reduction

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure1}
\caption{Change in the HbA\textsubscript{1c} levels from baseline at 6-month intervals.}
\end{figure}

\textit{Abbreviation}: HbA\textsubscript{1c}, glycated hemoglobin.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure2}
\caption{Change in HbA\textsubscript{1c} levels as a function of sitagliptin dose.}
\textbf{Notes}: The patients who received different doses of sitagliptin had apparent reductions in HbA\textsubscript{1c} from baseline during the first 6 months of treatment. The HbA\textsubscript{1c} levels remained stable in patients receiving different doses of sitagliptin between month 6 and month 48.
\textit{Abbreviation}: HbA\textsubscript{1c}, glycated hemoglobin.
\end{figure}
in the HbA1c concentration of 1.3% from a baseline average HbA1c of 9.5% compared with the control group. The elderly patients are more likely to have age-related kidney problems, which may require an adjustment in the dose for patients receiving sitagliptin. In the current study, the average eGFR was <60 mL/minute/1.73 m² and the dose of sitagliptin was <70 mg.

Despite only 4 years of observation, chronic microvascular complications (retinopathy, nephropathy, and neuropathy) were evaluated every year, and were not increased in the current study. The changes in antihypertensive and statin treatments were also not apparent before or after 4 years of sitagliptin therapy. In an animal study, DPP-4 inhibitors in combination with angiotensin II receptor blocker treatment significantly reduced urinary albumin excretion and oxidative stress in diabetic endothelial nitric oxide synthase knockout mice and offered a new therapeutic approach for patients with diabetic nephropathy. In a large-scale clinical trial involving saxagliptin and cardiovascular outcomes in patients with T2DM, a decrease in microalbuminuria was reported. Functional GLP-1R expressed in retinal pigment epithelium cells also revealed the potential direct beneficial effects of GLP-1 treatment or an increase in GLP-1 via a DPP-4 inhibitor against diabetic retinopathy. A long-acting DPP-4 inhibitor (vildagliptin analog), when used in streptozotocin-induced diabetic peripheral neuropathy, can improve nerve function by preventing, blocking, or counteracting the progression of peripheral neuropathy in experimental diabetic rats. Therefore, sitagliptin may decrease the progression of microvascular complications.

There were several limitations to this study. First, 4 years of observations is inadequate to assess chronic complications; further long-term evaluation is indicated. Second, because of the age of the patients, age-related nephropathy cannot be ruled out. Although only one patient developed retinopathy, we cannot diagnose “diabetic nephropathy”. We used “nephropathy” to present the changes in kidney complications. Third, the sample size was small, so a further large-scale study may be needed.

Conclusion

This study has revealed the endurance of sitagliptin in elderly patients with T2DM. Glycemic control improved significantly, even in patients with a lower baseline HbA1c concentration, and the progression of microvascular complications stabilized, with sustained effects as long as 4 years. Long-term observation is indicated for these patients. This study can provide useful data for clinicians and health care professionals using sitagliptin monotherapy in the treatment of elderly patients with T2DM.

Disclosure

The authors report no conflicts of interest in this work.

References