Efficacy and safety of sitagliptin added to metformin and insulin compared with voglibose in patients with newly diagnosed type 2 diabetes

Chunhong Shi, Ru Zhang, Ran Bai, Dan Liu, Yongbo Wang, Xueyang Zhang, Hao Wang, Jianling Du

Department of Endocrinology, The First Affiliated Hospital of Dalian Medical University, Dalian, PR China.

OBJECTIVE: To assess the efficacy and safety of sitagliptin compared with voglibose added to combined metformin and insulin in patients with newly diagnosed type 2 diabetes (T2DM).

METHODS: In this 12-week prospective, randomized, parallel trial, 70 newly diagnosed T2DM patients with glycosylated hemoglobin (HbA1c) ≥9% and/or fasting plasma glucose (FPG) ≥11.1 mmol/L were randomized (1:1) to receive sitagliptin 100 mg per day + metformin + insulin glargine or voglibose 0.2 mg three times daily + metformin + insulin glargine. Change in HbA1c at week 12 was the primary endpoint.

RESULTS: The mean baseline HbA1c was 11.0% in the patients. The changes in HbA1c from baseline were -6.00% in the sitagliptin group and -3.58% in the voglibose group, and the between-group difference was -2.42% (95% CI -1.91 to -2.93, p=0.02). The differences in FPG and homeostatic model assessment of β-cell function (HOMA-β) and the change in body weight between groups from baseline were -2.95 mmol/L (p=0.04), 43.91 (p=0.01) and -2.23 kg (p=0.01), respectively. One patient (2.9%) in the sitagliptin group and three patients (8.6%) in the voglibose group exhibited hypoglycemia.

CONCLUSIONS: Sitagliptin added to combined metformin and insulin therapy showed greater efficacy and good safety regarding hypoglycemia in patients with newly diagnosed T2DM compared with voglibose.

KEYWORDS: Diabetes Mellitus, Type 2; Sitagliptin Phosphate; Voglibose; Combined Therapy; Efficacy; Safety.
newly diagnosed T2DM and confirmed that after two weeks of treatment, sitagliptin had a stronger effect on decreasing mean blood glucose, fasting blood glucose and glucose fluctuation (13).

At present, no studies have been carried out on the long-term efficacy of adding sitagliptin compared with voglibose to combined metformin and insulin therapy for the treatment of newly diagnosed T2DM patients experiencing high glucose toxicity. In the present study, we added sitagliptin or voglibose to combined metformin and insulin therapy for treating newly diagnosed T2DM patients with HbA1c > 9.0% and/or FPG ≥ 11.1 mmol/L. Twelve weeks later, the efficacy and safety of these two treatments were compared and analyzed.

### MATERIALS AND METHODS

#### Design and subjects

This study used a randomized, prospective, parallel design. A total of 83 newly diagnosed T2DM patients from the First Affiliated Hospital of Dalian Medical University were screened. Patients were diagnosed with T2DM within the past year according to the 2013 American Diabetes Association (ADA) criteria, age from 18 to 65 years and fasting FPG ≥ 11.1 mmol/L and/or HbA1c > 9%. These patients had never taken oral hypoglycemic agents or received insulin treatment prior to their participation in the present trial.

The exclusion criteria were as follows: the presence of acute complications of diabetes, such as diabetic ketoacidosis or hyperosmolar hyperglycemic syndrome; severe cardiovascular or cerebrovascular events within the past 6 months; kidney damage (estimated glomerular filtration rate less than 60 ml/min/1.73 m²), or liver damage (alanine aminotransferase or aspartate aminotransferase 2.5 times more than the normal upper limit); the presence of a tumor, severe infection, or stress; a history of acute pancreatitis; or a history of gastrointestinal surgery.

The protocol was approved by the ethics committee of the First Affiliated Hospital of Dalian Medical University (Ethics References No: YJ-KY-FB-2015-02) and performed in accordance with the Declaration of Helsinki and good clinical practice guidelines. All patients provided written informed consent and then received screening.

#### Randomization and masking

Using a computer-generated random number sequence, the patients were randomly and evenly divided into groups receiving sitagliptin 100 mg per day + metformin + insulin glargine or voglibose 0.2 mg three times daily + metformin + insulin glargine. The random numbers were stored in a closed opaque envelope. The researchers and patients were aware of the grouping and the drug interventions, but the investigators in charge of data collection and analysis were masked to the grouping and drug interventions.

#### Procedures

All patients received diabetes health education, diabetes diets were provided by the nutrition department, and the patients were instructed to comply with a routine exercise plan.

Patients in the sitagliptin group took sitagliptin (100 mg, MSD, Hangzhou) 15 minutes before breakfast; patients in the voglibose group took voglibose (0.2 mg, Takeda, Japan) three times daily with meals. Metformin hydrochloride (0.5 g, SASS, Shanghai) was also orally administered three times daily with meals.

Subcutaneous injection of insulin glargine (Sanofi, France) was administered prior to bedtime. Insulin glargine was used at a dosage of 0.2 U·kg⁻¹·d⁻¹ initially. The amount of insulin glargine was adjusted based on the results of a finger-prick test. For patients who had fasting blood glucose (FBG) ≥ 7 mmol/L for two consecutive days, insulin glargine was increased by 2 U. For patients who had FBG < 5 mmol/L or showed hypoglycemia without obvious cause, the insulin glargine dose was decreased by 2 U, as described previously (14). Administration of insulin glargine was discontinued when the dose was ≤ 10 U/d.

Every two weeks, telephone follow-up was conducted to record the current amount of insulin intake, fasting and postprandial blood glucose levels, hypoglycemic events and fingertip blood glucose levels at that time. At baseline and three months, the HbA1c level was examined, a 75-g oral glucose tolerance test was conducted, and FPG, two-hour postprandial plasma glucose (2-h PPG), fasting C peptide (FCP) and 2-h C peptide (CP) were measured. In addition, body weight was measured.

C peptide test: After fasting overnight, patients consumed a mixture of 150 mL of warm water and 150 mL of 50% glucose solution over a five-minute period. Venous blood was drawn before and two hours after consumption of the mixture. Serum FCP and 2-h CP levels were examined by electrochemiluminescence. Morning FBG and 2-h PPG levels after three meals were monitored using a finger-prick test (ACCU-CHEK Performa, Roche, Germany) according to the instructions (15). FPG, liver function, kidney function, and blood lipids were measured by an automatic biochemical analyzer (HITACHI 7600-210, Japan), and HbA1c was determined using HPLC (VARIANT-II, Bio-Rad, CA, USA). Homoeostatic model assessment of insulin resistance (HOMA-IR) and β function (HOMA-β) were calculated according to well-established methods (16).

Symptomatic hypoglycemia was defined as the occurrence of dizziness, palpitations and other symptoms of sympathetic nerve stimulation; biochemical hypoglycemia was defined as a finger-prick blood glucose < 4.4 mmol/L.

#### Outcomes

The change in HbA1c from baseline to week 12 was the primary endpoint. The changes in FPG, 2-h PPG, HOMA-IR, HOMA-β, and body weight at week 12 compared with baseline, and duration and dosage of insulin application were the secondary endpoints.

#### Statistical methods

The standard deviation was assumed to be 0.6%, as described previously (9). If the difference in HbA1c between the sitagliptin group and voglibose group was 0.5%, 33 patients were required in each group (66 patients in total) to reach 90% power, permitting a 10% dropout rate; thus, a total of 70 patients were included in the randomization.

The analysis of outcomes was conducted in accordance with the principle of intention to treat.

Data that accorded with a normal distribution are expressed as the mean ± standard deviation (X ± s), intergroup differences were compared by an independent samples t-test, and intragroup differences were compared by a paired sample t-test. All p-values were two-sided, and p < 0.05 indicates a statistically significant difference.
RESULTS

Baseline information
From March 2015 to February 2016, 83 patients were screened, and 70 were enrolled and randomized into two groups. Six patients did not complete the 12-week follow-up (Figure 1). No statistically significant differences in initial characteristics were seen between the two groups (Table 1).

Primary endpoint
At week 12, the initial changes in HbA1c were -6.00% in the sitagliptin group and -3.58% in the voglibose group. The difference between the groups was -2.42% (95% CI -1.91 to -2.93, p=0.02) (Figure 2).

Secondary endpoints
After treatment, the decrease in FPG in the sitagliptin group (-8.71 ± 2.99 mmol/L) was significantly greater than that in the voglibose group (-5.76 ± 4.04 mmol/L), the difference was -2.95 mmol/L (p=0.04), although the decrease in 2-h PPG between the groups was not significantly different (p=0.16) (Figure 3).

After treatment, the increase in HOMA-β in the sitagliptin group (71.39 ± 55.53) was significantly greater than that in the voglibose group (27.48 ± 25.14), with a difference of 43.91 (p=0.01). The decrease in HOMA-IR between the groups was not significantly different (p=0.75) (Figure 4).

After 12 weeks, weight loss in the sitagliptin group (-3.77 ± 1.88 kg) was significantly greater than that in the voglibose group (-1.54 ± 1.05 kg), with a difference of -2.23 kg (p=0.01) (Figure 5).

The duration of insulin glargine use in the sitagliptin group (32 ± 15 d) was significantly shorter than that in the voglibose group (50 ± 27 d), with a difference of 18 d (p=0.04). The amount of insulin glargine administered was not significantly different between the groups (sitagliptin group 0.28 ± 0.08 U·kg⁻¹·d⁻¹ vs. voglibose group 0.31 ± 0.05 U·kg⁻¹·d⁻¹) (p=0.55).

Table 1 - Baseline characteristics of the S Group and V Group (± S).

|               | S Group (n=35) | V Group (n=35) |
|---------------|---------------|---------------|
| Sex (male/female) | 20/14         | 18/16         |
| Age (years)    | 41.2 ± 10.0   | 43.1 ± 12.0   |
| Weight (kg)    | 76.15 ± 13.57 | 76.72 ± 9.36  |
| WC (cm)        | 92.99 ± 11.38 | 94.19 ± 8.29  |
| BMI (kg/m²)    | 26.23 ± 3.45  | 26.86 ± 2.57  |
| SBP (mmHg)     | 130.26 ± 8.35 | 131.33 ± 12.61|
| DBP (mmHg)     | 82.75 ± 5.50  | 82.26 ± 8.47  |
| FPG (mmol/L)   | 14.99 ± 2.73  | 14.50 ± 3.11  |
| HbA1c (%)      | 11.9 ± 1.5    | 10.6 ± 1.1    |
| TC (mmol/L)    | 5.54 ± 1.36   | 5.16 ± 1.04   |
| TG (mmol/L)    | 2.17 ± 1.54   | 1.73 ± 0.88   |
| HDL-C (mmol/L) | 1.72 ± 1.02   | 1.34 ± 0.26   |
| LDL-C (mmol/L) | 3.32 ± 0.98   | 3.26 ± 0.74   |

S Group: Sitagliptin, V Group: Voglibose. WC: waist circumference, BMI: body mass index, SBP: systolic blood pressure, DBP: diastolic blood pressure, FPG: fasting plasma glucose, HbA1c: glycated hemoglobin A1c, TC: total cholesterol, TG: triglyceride, HDL-C: high-density lipoprotein cholesterol, LDL-C: low-density lipoprotein cholesterol.
Safety endpoint
One patient in the sitagliptin group (2.9%) and three patients in the voglibose group (8.6%) exhibited symptomatic hypoglycemia. No incidents of biochemical hypoglycemia or nocturnal hypoglycemia occurred in patients in either group.

DISCUSSION
For newly diagnosed T2DM patients with high blood glucose levels, intensive insulin therapy can quickly relieve high glucose toxicity and improve islet cell function. In this study, sitagliptin and voglibose added to insulin glargine combined with metformin were compared in a 12-week treatment regimen of a group of T2DM patients with high blood glucose levels. The results showed that the decreases in HbA1c and FPG in the sitagliptin group were significantly higher than those in the voglibose group, and the decrease in 2-h PPG between the groups did not differ. This result might have occurred because DPP-4 inhibitors reduce the degradation of glucagon-like peptides in the body, thereby increasing glucose-dependent insulin secretion, inhibiting glucagon secretion, and ultimately lowering fasting and postprandial blood glucose levels (17). Alpha glucosidase inhibitors (AGIs) reduce postprandial blood glucose primarily by reducing intestinal carbohydrate absorption. Therefore, sitagliptin is superior to voglibose for lowering fasting blood glucose levels when either is combined with insulin and metformin.

Figure 2 - Comparison of HbA1c decline in the Sitagliptin and Voglibose groups after treatment.

Figure 3 - Comparison of FPG and 2-h PPG decline in the Sitagliptin and Voglibose groups after treatment.

Figure 4 - Comparison of HOMA-β and HOMA-IR change from baseline between the Sitagliptin and Voglibose groups after treatment.

Figure 5 - Comparison of weight change between the Sitagliptin and Voglibose groups after treatment.
Since sitagliptin was superior to voglibose for lowering fasting blood glucose levels, even though no significantly different effects were seen on postprandial blood glucose levels between the groups, sitagliptin was also more effective than voglibose in reducing HbA1c. A similar conclusion was reported in a previous study. Compared with AGIs, the efficacy of sitagliptin in patients with poorly controlled T2DM compared with that of stable glimepiride alone (mean HbA1c 7.7%) was assessed after 12 weeks. Notably, HbA1c was reduced by -0.44% in the sitagliptin group (p < 0.001) (18). A meta-analysis confirmed that changes in HbA1c (weighted mean deviation -0.30%; 95%CI -0.47 to -0.13%, p < 0.001) and FPG levels (weighted mean deviation -0.50 mmol/L; 95% CI -0.89 to -0.11 mmol/L, p=0.01) were significantly greater in the DPP-4 inhibitor treatment group compared with that in the AGI group (19).

The present study found that HOMA-β in the sitagliptin group increased significantly after 12 weeks, suggesting a significant improvement in islet function. In a double-blind, crossover clinical trial, Tremblay et al. (20) found that T2DM patients receiving sitagliptin therapy displayed significantly improved islet function compared with patients receiving a placebo. In the present study, the voglibose treatment group also showed improved islet cell function and reduced insulin resistance, consistent with the results reported by Do et al. (21). The increase in HOMA-β in the sitagliptin group was significantly greater than that in the voglibose group. This finding suggests that sitagliptin more effectively protected and improved islet cell function, possibly because DPP-4 inhibitors inhibit the degradation of incretin, which has an indirect role in the inhibition of pancreatic β-cell apoptosis, and promote the proliferation of β cells (22). Voglibose relieves only high glucose toxicity to pancreatic β cells; therefore, compared with the voglibose group, the sitagliptin group showed better islet β cell protection.

In patients with newly diagnosed T2DM, glucotoxicity and lipotoxicity could be effectively reduced by early insulin therapy to alleviate insulin resistance and protect islet function (23). After short-term insulin therapy, patients with highly elevated blood glucose levels can gradually reduce or discontinue insulin treatment. In the present study, when the insulin glargine dose was reduced to less than 10 U/d and blood glucose was still well controlled, administration of insulin glargine was discontinued. The duration of insulin glargine use in the sitagliptin group (mean 32 d) was significantly shorter than that in the voglibose group (mean 50 d). This result might be due to a better effect of sitagliptin on lowering blood glucose and improving islet function; therefore, less insulin glargine was required to control blood glucose. The fact that there was no difference between the two groups in the average insulin dose per day per kilogram of body weight indicates that the shorter period of insulin use required in the sitagliptin group compared with that in the voglibose group was not due to the insulin dose.

Weight loss can improve insulin sensitivity; therefore, weight loss is essential for obese T2DM patients. The present study showed significant weight loss in both groups after treatment. Compared with placebo, DPP-4 inhibitors (24) and AGIs (25) can reduce body weight, consistent with the conclusions of our study. In addition, the sitagliptin group displayed significantly more weight loss than the voglibose group. A prospective, randomized, multicenter study treated T2DM patients with sitagliptin (50 mg/d) or voglibose (0.6 mg/d) for 24 weeks, and weight loss with sitagliptin treatment was significantly higher than that with voglibose treatment (-1.3 ± 3.2 kg vs. 0.4 ± 2.8 kg) (26), consistent with the results of the present study.

Hypoglycemia increases cardiovascular risk and is a barrier to blood glucose control in patients with diabetes. Our study showed that the sitagliptin group had one occurrence and that the voglibose group had three occurrences of symptomatic hypoglycemia. No incidents of biochemical hypoglycemia or nocturnal hypoglycemia occurred in either group. Both treatments therefore resulted in a low incidence of hypoglycemia and showed excellent safety. In our previous study, we compared the incidence of hypoglycemia in newly diagnosed T2DM patients receiving combination treatment with continuous subcutaneous insulin injection and DPP-4 inhibitors or AGIs; two weeks of dynamic blood glucose monitoring showed that both therapies have excellent safety (12).

The limitations of our study are that the number of included patients was relatively small and the follow-up time was relatively short. The results therefore require verification in large-scale, multicenter clinical trials with more patients and longer follow-up times.

In conclusion, compared with voglibose, sitagliptin added to combined insulin and metformin therapy can achieve a significantly better effect on lowering HbA1c and FPG and shows excellent safety for newly diagnosed T2DM patients with highly elevated blood glucose.

ACKNOWLEDGMENTS

We thank all investigators and patients who participated in the study.

AUTHOR CONTRIBUTIONS

Shi C and Zhang R conducted the statistical analysis and wrote the manuscript. Liu D and Wang Y collected the data. Zhang X, Wang H and Du J participated in the design of the study and edited the manuscript. Bai R designed and directed the entire study, and revised the manuscript.

REFERENCES

1. Xu Y, Wang L, He J, Bi Y, Li M, Wang T, et al. Prevalence and control of diabetes in Chinese adults. JAMA. 2013;310(9):948-59, https://doi.org/10.1001/jama.2013.168118.
2. Weng J, Lu J, Jia W, Zou D, Zha D, Zhou Z, et al. Type 2 diabetes prevention and treatment guidelines (2013 Version). Chin J Diabetes Mellitus. 2014;6:447-98.
3. Bretenot MF, Iberl M, Shimomura K, Zhang Q, Adriaenssens AE, Proks P, et al. Reversible changes in pancreatic islet structure and function produced by elevated blood glucose. Nat Commun. 2014;5:4639, https://doi.org/10.1038/ncomms5639.
4. Inzucchi SE, Bergenstal RM, Buse JB, Diamant M, Ferrannini E, Nauck M, et al. Management of hyperglycemia in type 2 diabetes, 2015: a patient-centered approach: update to a position statement of the American Diabetes Association and the European Association for the Study of Diabetes. Diabetes Care. 2015;38(1):140-9, https://doi.org/10.2337/dc14-2441.
5. Gao HW, Xie C, Wang HN, Lin YJ, Hong TP. Beneficial metabolic effects of nateglinide versus acarbose in patients with newly-diagnosed type 2 diabetes. Acta Pharmacol Sin. 2007;28(4):534-9, https://doi.org/10.1111/j.1745-7254.2007.00534.x.
6. Standl E, Schnell O. Alpha-glucosidase inhibitors 2012 - cardiovascular considerations and trial evaluation. Diab Vasc Dis Res. 2012;9(3):163-9, https://doi.org/10.1177/1479164112441524.
7. Tasyurek HM, Altunbas HA, Balci MK, Sanlioglu S. Incretins: their physiology and application in the treatment of diabetes mellitus. Diabetes Metab Res Rev. 2014;30(5):354-71, https://doi.org/10.1002/dmrr.2501.
8. Barzilai N, Guo H, Mahoney EM, Caporossi S, Golm GT, Landng RB, et al. Efficacy and tolerability of sitagliptin monotherapy in elderly patients with type 2 diabetes: a randomized, double-blind, placebo-controlled trial. Curr Med Res Opin. 2011;27(5):1049-58, https://doi.org/10.1185/03007995.2011.568059.
Sitagliptin has greater efficacy

Shi C et al.

9. Iwamoto Y, Tajima N, Kadowaki T, Nonaka K, Taniguchi T, Nishii M, et al. Efficacy and safety of sitagliptin monotherapy compared with voglibose in Japanese patients with type 2 diabetes: a randomized, double-blind trial. Diabetes Obes Metab. 2010;12(7):633-22, https://doi.org/10.1111/j.1463-1326.2010.01197.x.

10. Yokoh H, Kobayashi K, Sato Y, Takemoto M, Uchida D, Kanatsuka A, et al. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor sitagliptin compared with alpha-glucosidase inhibitor in Japanese patients with type 2 diabetes inadequately controlled on metformin or pioglitazone alone (Study for an Ultimate Combination Therapy to Control Diabetes with Sitagliptin-1): A multicenter, randomized, open-label, non-inferiority trial. J Diabetes Investig. 2015;6(2):182-91, https://doi.org/10.1111/jdi.12282.

11. Sato S, Saitoh Y, Kou K, Meguro S, Tanaka M, Irie J, et al. Efficacy and safety of sitagliptin added to insulin in Japanese patients with type 2 diabetes: the EDIT randomized trial. PLoS One. 2015;10(3):e0121988, https://doi.org/10.1371/journal.pone.0121988.

12. Takai M, Ishikawa M, Maeda H, Kanamori A, Kubota A, Amemiya H, et al. Safety and efficacy of adding sitagliptin to insulin in patients with type 2 diabetes: the ASSIST-K study. Diabetes Res Clin Pract. 2014;103(3):e30-e3, https://doi.org/10.1016/j.diabres.2013.12.045.

13. Shi CH, Wang L, Bai R, Wang VB, Liu D, Zhang XY, et al. Comparison of therapeutic effects between sitagliptin and voglibose both combined with sensor-augmented insulin pump in newly diagnosed type 2 diabetes. Zhonghua Yi Xue Za Zhi. 2016;96(32):2554-8, https://doi.org/10.3760/cma.j.issn.0376-2491.2016.32.008.

14. Yki-Jarvinen H, Juurinen L, Alvarsson M, Bystedt T, Caldwell I, Davies M, et al. Initiate insulin by Aggressive Titration and Education (INITIATE): a randomized study to compare initiation of insulin combination therapy in type 2 diabetic patients individually and in groups. Diabetes Care. 2007;30(6):1364-9, https://doi.org/10.2337/dc06-1357.

15. Chinese Diabetes Society. Clinical application guide of blood glucose monitoring in China (Edition 2011). Zhonghua Yi Xue Za Zhi. 2011;91(10):656-64.

16. Li X, Zhou ZG, Qi HY, Chen XY, Huang G. Replacement of insulin by fasting C-peptide in modified homeostasis model assessment to evaluate insulin resistance and islet beta cell function. Zhong Nan Da Xue Xue Bao Yi Xue Ban. 2004;29(4):149-23.

17. Bae EJ. DPP-4 inhibitors in diabetic complications: role of DPP-4 beyond glucose control. Arch Pharm Res. 2016;39(8):1114-28, https://doi.org/10.1007/s12272-016-0681-x.

18. Kobayashi K, Yokoh H, Sato Y, Takemoto M, Uchida D, Kanatsuka A, et al. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor sitagliptin compared with alpha-glucosidase inhibitor in Japanese patients with type 2 diabetes inadequately controlled on sulfonylurea alone (SUCCESS-2): a multicenter, randomized, open-label, non-inferiority trial. Diabetes Obes Metab. 2014;16(8):761-5, https://doi.org/10.1111/dob.12264.

19. Cai X, Yang W, Zhou L, Zhang S, Han X, Ji L. Comparisons of the efficacy of glucose control, lipid profile, and beta-cell function between DPP-4 inhibitors and AGI treatment in type 2 diabetes patients: a meta-analysis. Endocrine. 2015;50(3):590-7, https://doi.org/10.1007/s12020-015-0653-3.

20. Tremblay AJ, Lamarche B, Deacon CF, Weinsnagel SJ, Couture P. Effect of sitagliptin therapy on postprandial lipoprotein levels in patients with type 2 diabetes. Diabetes Obes Metab. 2011;13(4):366-73, https://doi.org/10.1111/j.1463-1326.2011.01362.x.

21. Do HJ, Jin T, Chung JH, Hwang JW, Shin MJ. Voglibose administration regulates body weight and energy intake in high fat-induced obese mice. Biochem Biophys Res Commun. 2014;443(3):1110-7, https://doi.org/10.1016/j.bbrc.2013.12.120.

22. Mundil D, Cameron-Vendrig A, Husain M. GLP-1 receptor agonists: a clinical perspective on cardiovascular effects. Diab Vasc Dis Res. 2012;9(2):95-108, https://doi.org/10.1177/1479164112441526.

23. Weng J, Li Y, Xu W, Shi L, Zhang Q, Zhu D, et al. Effect of intensive insulin therapy on beta-cell function and glycaemic control in patients with newly diagnosed type 2 diabetes: a multicentre randomised parallel-group trial. Lancet. 2008;371(9626):1753-60, https://doi.org/10.1016/s0140-6736(08)60762-x.

24. Karasik A, Aschner P, Katzeff H, Davies MJ, Stein PP. Sitagliptin, a DPP-4 inhibitor for the treatment of patients with type 2 diabetes: a review of recent clinical trials. Curr Med Res Opin. 2008;24(2):489-96, https://doi.org/10.1185/030079908X261069.

25. van de Laar FA, Lucassen PL, Akkermans RP, van de Lisdonk EH, Rutten GE, van Weel C. Alpha-glucosidase inhibitors for patients with type 2 diabetes: results from a Cochrane systematic review and meta-analysis. Diabetes Care. 2005;28(1):154-63, https://doi.org/10.2337/diacare.28.1.154.

26. Oe H, Nakamura K, Kihara H, Shimada K, Fukuda S, Takagi T, et al. Efficacy and safety of adding sitagliptin to insulin in patients with type 2 diabetes inadequately controlled on sulfonylurea alone (SUCCESS-2): a multicenter, randomized, open-label, non-inferiority trial. Diabetes Obes Metab. 2014;16(8):761-5, https://doi.org/10.1111/dob.12264.