The Effect of Liraglutide on Epicardial Adipose Tissue in Type 2 Diabetes

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Abstract

To study the effect of liraglutide on the thickness of epicardial adipose tissue (EAT) in type 2 diabetes mellitus (T2DM) patients with abdominal obesity. Methods. Abdominal obesity T2DM patients with poor glycemic control were collected and treated with liraglutide. Results. After 3 months of treatment with liraglutide, glycosylated hemoglobin (HbA1c) decreased from 9.81 ± 1.46% to 6.94 ± 1.29% (95%CI = 2.14 – 3.59, p < 0.001). The weight decreased from 91.67 ± 16.29 kg to 87.29 ± 16.43 kg (95%CI = 2.97 – 5.79, p < 0.001). Waist circumference before treatment was 103.69 ± 9.14 cm, and after treatment was 96.42 ± 8.42 cm (95%CI = 5.04 – 9.50, p < 0.001). Total cholesterol (TC), triglyceride (TG), and low-density lipoprotein cholesterol (LDL-C) were significantly lower than those before treatment. TC decreased from 5.34 ± 1.05 mmol/L to 4.86 ± 0.97 mmol/L (95%CI = 0.15 – 0.82, p < 0.001). TG was 1.89 (1.48-3.17) and then to 1.92 ± 0.69 (p = 0.03). LDL-C decreased from 3.39 ± 0.84 mmol/L to 3.01 ± 0.74 mmol/L (95%CI = 0.17 – 0.59, p = 0.001). HDL-C increased by 1.7% after treatment, with no significant difference (p = 0.062). More importantly, the thickness of EAT decreased from 5.0 (5.0-7.0) mm to 3.95 ± 1.43 mm (p < 0.001) after liraglutide administered for 3 months. Conclusion. Liraglutide significantly reduces EAT thickness in T2DM with abdominal obesity, which provides theoretical support for the cardiovascular benefits of liraglutide.

1. Introduction

Type 2 diabetes mellitus (T2DM) is a metabolic syndrome with disorders of glucose, protein, and fat caused by deficiency of insulin secretion and/or insulin function. It is often accompanied by obesity. Obesity, especially visceral fat accumulation, can lead to islet dysfunction, insulin resistance (IR), prediabetes, T2DM, and inflammatory reaction and increase the incidence of coronary heart disease, hypertension, and other events [1, 2]. Quantification and reduction of visceral fat represent effective methods to identify high-risk individuals and reduce their cardio-metabolic risk of T2DM patients [3]. Epicardial adipose tissue (EAT), as visceral fat, has attracted much attention due to its close relationship with myocardium and coronary artery [4]. The increase of EAT may represent a chronic inflammatory injury and is related to coronary artery disease [5], and T2DM patients had significantly thicker EAT [6]. Liraglutide is a glucagon-like peptide-1 (GLP-1) analog which can activate GLP-1 receptor (GLP-1R). In T2DM patients, studies suggested that GLP-1 receptor agonist (GLP-1 RA) could redistribute adipose tissue deposits and reduce visceral fat [7, 8]. Additionally, liraglutide can reduce the risk of cardiovascular events. Our present study was to explore whether, in addition to its effects on blood glucose and blood lipids, liraglutide reduces cardiovascular events by acting on EAT in T2DM with abdominal obesity.
2. Design and Methods

2.1. Participant Identification. T2DM patients with abdominal obesity who had poor blood glucose control in Endocrinology Department of Jinpu District, the First Affiliated Hospital of Dalian Medical University from November 2016 to February 2018 were consecutively collected. Participants’ inclusion criteria were as follows: (1) aged between 18 and 70 years, (2) body mass index (BMI) > 25 kg/m² and waist circumference ≥ 90 cm in males and ≥ 85 cm in females, (3) glycosylated hemoglobin (HbA1c) was between 7 and 11%, and (4) the basic drugs were metformin and/or α-glucosidase inhibitors and/or insulin at the time of enrollment and met the liraglutide indications. Exclusion criteria are as follows: (1) inflammatory bowel disease and diabetic gastroparesis; (2) pregnant or breastfeeding women; (3) moderate to severe liver dysfunction (Child’s-Pugh score); (4) acute or chronic renal insufficiency, glomerular filtration rate (GFR) < 60 ml/(min × 1.73 m²); (5) history of thyroid tumor or pancreatitis; (6) diabetes with acute complications or with tumor; (7) electronic implants such as cardiac pacemaker, insulin pump, and magnetic metal; and (8) uncontrollable involuntary movement, claustrophobia patients, and so on who are not suitable for MRI examination. With the approval of the medical ethics committee, all participants were informed of the possible side effects of liraglutide and provided informed consent.

2.2. Drugs and Dosage. Liraglutide (Victoza, manufactured by Novo Nordisk, Denmark) was injected subcutaneously once a day for 3 months, starting from 0.6 mg/day and increasing to 1.2 mg/day in 3-5 days if there was no adverse. The curative effect of liraglutide was evaluated by monitoring fasting plasma glucose (FPG) and 2-hour blood glucose after three meals every day and before bedtime and early morning if necessary. For patients with poor hypoglycemic effect, the dose was further increased to 1.8 mg/day.

2.3. Biochemical Blood Tests. The patients kept a light diet for three days and fasting water for 12 hours before the blood tests. Blood samples were drawn in the next morning. FPG, HbA1c, total cholesterol (TC), triglyceride (TG), low-density lipoprotein cholesterol (LDL-C), and high-density lipoprotein cholesterol (HDL-C) were tested ahead of and after 3 months liraglutide being administered by automatic biochemical analyzer (Hitachi 7600, Japan). The 2-hour postprandial blood glucose (2hPBG) was measured by Accu-Chek advantage electronic sensing portable blood glucose meter (Roche, Germany).

2.4. EAT Thickness Measurement. EAT thickness was measured before and after 3-month liraglutide treatment by 1.5T cardiac MRI (platform HDxt; General Electric Medical Systems, Waukesha, WI, USA). Scanning parameters were TR 3.6 ms, TE1.6 ms, turning angle 50°, bandwidth 125 kHz, FOV 350 mm × 350 mm, matrix 192 × 224, layer thickness 10 mm, and layer spacing 0. EAT thickness was measured at the right ventricle late diastolic phase with four-chamber echocardiography by two well-experienced radiologists. All cardiac MRI scans were performed by the same radiographer.

2.5. Statistical Analysis. The SPSS 19.0 software was used for statistical analysis. Normal distributed data were presented as the mean ± standard deviation (M ± SD). Abnormally distributed data were presented as the median (interquartile range). Paired t-test and Wilcoxon test were used to compare the differences of data before and after liraglutide treatment. It is considered statistically significant when the p value was less than 0.05.

3. Results

3.1. Subject Enrollment. The basic demographic characteristics of 21 out of 27 subjects who completed the trial are shown in Table 1. After the injection of liraglutide, the patients had varying but tolerable degrees of nausea and loss of appetite. For most of the patients, the above symptoms gradually disappeared without drug intervention after 1-3 days. No severe hypoglycemia was observed during the whole trial, while among the 27 subjects, 5 were withdrawn from the group because of intolerable gastrointestinal reaction, and 1 was withdrawn because of skin allergy at the injection site. The average dose of liraglutide in 3 months was 1.2 mg/day.

3.2. FPG, 2hPBG, and HbA1c Decreased. After the patients received 3 months of liraglutide, compared with that before liraglutide, FPG decreased from 10.31 ± 2.41 mmol/L to 7.17 ± 1.28 mmol/L (95%CI = 2.07 – 4.20, p < 0.001), and 2hPBG decreased from 12.68 ± 3.33 mmol/L to 9.12 ± 1.37 mmol/L (95%CI = 2.05 – 5.07, p < 0.001), shown in Figure 1(a). What is more, HbA1c decreased from 9.81 ± 1.46% to 6.94 ± 1.29% (95%CI = 2.14 – 3.59, p < 0.001), shown in Figure 1(b). FPG, 2hPBG, and HbA1c were dropped by 30.5%, 28.1%, and 29.3% on average.

3.3. Weight, Waist Circumference, and BMI Decreased. The patients’ average body weight was 91.67 ± 16.29 kg before liraglutide treatment and 87.29 ± 16.43 kg after treatment, dropped by 4.38 kg on average (95%CI = 2.97 – 5.79, p < 0.001). BMI decreased on average by 1.49 kg/m² from 30.97 ± 4.04 kg/m², and after treatment was 29.48 ± 4.08 kg/m² (95%CI = 1.02 – 1.98, p < 0.001). Waist circumference decreased by 7.27 cm on average from 103.69 ± 9.14 cm to 96.42 ± 8.42 cm (95%CI = 5.04 – 9.50, p < 0.001). Results are shown in Figure 2.

3.4. Effect of Liraglutide on Blood Lipid Level. TC decreased from 5.34 ± 1.05 mmol/L to 4.86 ± 0.97 mmol/L (95%CI = 0.15 – 0.82, p = 0.001), TG decreased from 1.89 (1.48-3.17) and then to 1.92 ± 0.69 (p = 0.03), and LDL-C decreased from 3.39 ± 0.84 mmol/L to 3.01 ± 0.74 mmol/L (95%CI = 0.17 – 0.59, p = 0.001) (Figure 3). HDL-C increased by 1.7% on average from 0.96 (0.89-1.45) mmol/L to 1.19 ± 0.25 mmol/L, but the difference was not statistically significant (p = 0.062).

3.5. EAT Thickness Decreased. The EAT thickness was 5.0 (5.0-7.0) mm to 3.95 ± 1.43 mm after treatment measured by MRI, with an average decrease of 1.62 mm. The changes suggested that liraglutide can significantly reduce EAT thickness (p < 0.001, Figure 4).
Liraglutide, as a GLP-1 analog, has 97% homology structurally with endogenous human GLP-1 [9]. GLP-1 receptor (GLP-1R) mainly exists in fat, liver, gastrointestinal tract, kidney, cardiovascular, and central nervous system [10]. Liraglutide activates GLP-1R to promote insulin secretion of pancreatic β cells, reduce body weight, promote vasodilation, and reduce the level of atherosclerotic risk factors in endothelial cells [11–13]. Our results showed that FPG, 2hPBG, and HbA1c of T2DM patients with abdominal obesity were significantly decreased after liraglutide treatment for 3 months, and the body weight, waist circumference, BMI, TG, TC, and LDL-C levels were also significantly decreased. Similar to the results of the present study, liraglutide can effectively reduce Hba1c [14] and body weight [9]. Aoki and his colleagues found that the non-HDL-C and calculated TC were decreased significantly after administrating of liraglutide for 3 months [15]. In addition, in T2DM patients receiving standard care, liraglutide can reduce the risk of cardiovascular events compared with the placebo group [16], but the specific reason is yet not fully clear.

Epicardial adipose tissue (EAT), characterized by brown adipose tissue, is closely connected with the myocardium and coronary artery without fascia barrier. Under physiological conditions, EAT can mitigate the impact, absorb fatty acids, and protect myocardium and coronary arteries from cold injury [17]. It can balance the secretion of proinflammatory and anti-inflammatory factors, regulate the cardiovascular system, and play an important role in obesity-related inflammation and atherosclerosis [18, 19]. Scholars have also confirmed that EAT is positively correlated with obesity, impaired fasting blood glucose regulation, insulin resistance, metabolic syndrome, hypertension, and coronary heart disease through adipocyte hypertrophy and hyperplasia [20, 21]. T2DM patients had significantly thicker EAT associated with cardiac systolic dysfunction [6]. What is more, EAT volume in patients with coronary heart disease with abdominal obesity is significantly higher than that in patients without coronary heart disease [22]. Therefore, for T2DM patients with abdominal obesity, we should not only effectively control blood glucose levels but also monitor EAT levels.

The monitoring of EAT included abdominal circumference, echocardiography, multislice computed tomography (MCT), and magnetic resonance imaging (MRI). Among them, abdominal circumference is the cheapest and most easily obtained visceral fat marker, but it lacks sensitivity and specificity. Echocardiography is noninvasive, reliable, easy to repeat, and relatively accurate, but it cannot obtain a linear measurement value in a single position. Because of its high accuracy, low variability, and high repeatability, MRI results can be used as the reference standard for quantification of EAT and it is considered as the gold standard for measuring visceral fat [23–25]. There are few studies on the measurement of EAT by cardiac MRI. By using cardiac MRI measurements, we found that after 3 months of liraglutide treatment, the thickness of EAT decreased significantly along with the decrease of waist circumference. Similar to the results of this study, Bouchi et al. found that liraglutide can reduce visceral fat and improve the quality of life of diabetic patients [8].

4. Discussion

Liraglutide, as a GLP-1 analog, has 97% homology structurally with endogenous human GLP-1 [9]. GLP-1 receptor (GLP-1R) mainly exists in fat, liver, gastrointestinal tract, kidney, cardiovascular, and central nervous system [10]. Liraglutide activates GLP-1R to promote insulin secretion of pancreatic β cells, reduce body weight, promote vasodilation, and reduce the level of atherosclerotic risk factors in endothelial cells [11–13]. Our results showed that FPG, 2hPBG, and HbA1c of T2DM patients with abdominal obesity were significantly decreased after liraglutide treatment for 3 months, and the body weight, waist circumference, BMI, TG, TC, and LDL-C levels were also significantly decreased. Similar to the results of the present study, liraglutide can effectively reduce Hba1c [14] and body weight [9]. Aoki and his colleagues found that the non-HDL-C and calculated TC were decreased significantly after administrating of liraglutide for 3 months [15]. In addition, in T2DM patients receiving standard care, liraglutide can reduce the risk of cardiovascular events compared with the placebo group [16], but the specific reason is yet not fully clear.

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Table 1: Demographic characteristics of T2DM patients.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>T2DM patients</th>
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<tbody>
<tr>
<td>Number</td>
<td>21</td>
</tr>
<tr>
<td>Gender (female/male)</td>
<td>6/15</td>
</tr>
<tr>
<td>Age (mean ± SD, years)</td>
<td>42.86 ± 11.15</td>
</tr>
<tr>
<td>Height (mean ± SD, cm)</td>
<td>171.62 ± 8.49</td>
</tr>
<tr>
<td>Weight (mean ± SD, kg)</td>
<td>91.67 ± 16.30</td>
</tr>
<tr>
<td>BMI (mean ± SD, kg/m²)</td>
<td>30.97 ± 4.04</td>
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considered to be that it reverses the whitening of brown fat in EAT by stimulating GLP-1R after administration of liraglutide, thus reducing the thickness of EAT and protecting cardiovascular. Diabetes patients showed obvious early and progressive myocardial lipid accumulation. Whether or not liraglutide and metformin have the same alleviating effect on diabetic cardiomyopathy need further investigation.

Scholars also found that liraglutide can alleviate endotoxemia-related microvascular thrombosis through GLP-1 receptor signal-mediated immune regulation without affecting blood glucose or HbA1c level, to prevent systemic inflammation, vascular dysfunction, and end organ injury, which has potential clinical significance for the treatment of sepsis [34]. GLP-1R activation reduces vascular inflammation by selectively acting on endothelial cells rather than myeloid cells to reduce the cardiovascular complications of arterial hypertension [35]. Liraglutide could regulate sepsis-induced endothelial dysfunction by reducing vascular inflammation and oxidative stress [36]. Moreover, in the early stage of thrombosis, a specific anti-thrombotic therapy may inhibit the specific target pathway of coagulation cascade and change the thrombosis process to improve the clinical outcome of patients [37], while it is still
The authors declare that they have no conflicts of interest.

Conflicts of Interest

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References


