Short- and medium-term efficacy of sodium glucose cotransporter 2 (SGLT-2) inhibitors for the treatment of type 1 diabetes: systematic review and meta-analysis

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Abstract

Introduction: Sodium glucose cotransporter 2 (SGLT2) inhibitors are insulin-independent and glucose-dependent anti-hyperglycaemic drugs that have shown potential as an adjuvant therapy to insulin for the treatment of type 1 diabetes mellitus (T1DM). The purpose of this meta-analysis is to systematically collect available data from randomised trials to determine SGLT-2 inhibitor efficacy in terms of glycaemic control, body mass index, and renal protection when compared with placebo.

Material and methods: Cochrane Library, MEDLINE, and EMBASE databases were searched for randomised controlled trials and meta-analyses (without language restrictions) conducted from January 2010 to September 2019.

Results: Seventeen randomised controlled trials with 7325 participants were included. Sodium glucose cotransporter 2 therapy significantly reduced the level of glycated haemoglobin (HbA1c) (by 0.37%), body weight (by 2.88 kg), and estimated glomerular filtration (eGFR) (by 0.67 mL/min/1.73 m²) when compared with placebo (all outcomes, p < 0.00001). Subgroup analysis by HbA1c levels showed significant differences between six and 12 months of treatment (p < 0.1). The magnitude of the HbA1c lowering effect waned with longer duration of treatment after six months (up to 12 months). Subgroup analysis by body weight showed significant differences between 1 and 3–4 months of treatment (p < 0.1). Weight loss plateaued after 3–4 months of treatment; subsequently, the weight remained relatively stable until 12 months. Subgroup analysis by eGFR showed significant differences between six and 12 months of treatment (p < 0.1). The magnitude of the eGFR lowering effect increased with longer duration of treatment after six months (up to 12 months).

Conclusions: Sodium glucose cotransporter 2 inhibitors show significant therapeutic effects when compared with placebo. Although changes in HbA1c, body weight, and eGFR vary during treatment, the therapeutic effects of SGLT-2 inhibitors measured by these three outcomes can last up to 12 months. More long-term, randomised trials and extended studies are needed to determine the long-term effects of SGLT2 inhibitors as adjuvant therapy for T1DM patients. (Endokrynol Pol 2020; 71 (4): 325–333)

Key words: sodium-glucose co-transporter 2; therapeutic effects; type 1 diabetes; meta-analysis; systematic review

Introduction

Type 1 diabetes mellitus (T1DM) is an autoimmune disease characterised by chronic hyperglycaemia due to an absolute deficiency of endogenous insulin and progressive metabolic alterations that increase the risk of cardiovascular diseases [1]. Type 1 diabetes affects about 22 million adults and 400,000 children, accounting for less than 5% of the total diabetes cases in the world [2]. Insulin replacement is still the main therapy for type 1 diabetes. Compared with conventional therapy, initial intensive therapy for diabetes has been shown to result in a moderate reduction in all-cause mortality [3]. However, intensive insulin therapy to achieve strict blood sugar control is usually associated with hypoglycaemia, and insulin cannot solve other important abnormalities associated with type 1 diabetes, such as glucagon imbalance and increased gastric emptying rate (resulting in large and rapid fluctuations of postprandial blood glucose levels) [4]. In addition, insulin therapy itself is associated with weight gain and related hypertension [5]. Therefore, there is a need for adjuvant therapy that can improve key parameters continuously (such as glycated haemoglobin (HbA1c), weight, and length of time in which the correct blood glucose range is maintained) and which can reduce the hypoglycaemic risk.

Sodium glucose cotransporter 2 (SGLT2) inhibitors are a new type of widely studied therapeutic adjuvants that can improve blood glucose control without relying on insulin. Sodium glucose cotransporter 2 inhibitors work by blocking glucose reabsorption in the renal proximal tubules, increasing urinary glucose excretion and heat and weight loss [6]. The safety and efficacy of SGLT-2 inhibitors for the treatment of type 2 diabetes have been confirmed in many studies [7–10]. On the
other hand, many studies of combined treatment of type 1 diabetes with SGLT-2 inhibitors and insulin have shown that SGLT-2 inhibitors effectively control blood sugar levels, reduce insulin demand, and promote weight loss. However, potential side effects, such as diabetic ketoacidosis and infection, have attracted widespread scrutiny [11–14]. So far, the U.S. Food and Drug Administration (FDA) and the European Medicines Agency (EMA) have approved four SGLT2 inhibitors (canagliflozin, dapagliflozin, empagliflozin, and ertugliflozin). In addition, the double SGLT1/SGLT2 inhibitor sotagliflozin has been approved by the EMA, and in Japan another three SGLT2 inhibitors (ipragliflozin, luseogliflozin, tofogliflozin) have been approved [28].

Previous meta-analyses [11–14] have confirmed the effectiveness of SGLT-2 inhibitors as adjuvant drugs. However, these reports did not analyse the effects of these drugs at different time points in detail. In addition, the duration of most registered trials is less than or equal to 24 weeks, but the therapeutic effects of drugs may differ when the observation period is prolonged. Therefore, we conducted a meta-analysis to determine the magnitude and persistence of the therapeutic effects of SGLT-2 inhibitors as auxiliary drugs when compared with placebo, focusing mainly on three outcomes: blood glucose control, body mass index, and renal protection.

Material and methods

Data sources and search
This study followed the preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 guidelines [15]. We searched the MEDLINE, Embase, Cochrane Library (January 2010–October 30, 2019), and clinical trial.gov databases, as well as online reports of major diabetes conferences, for randomised controlled trials comparing the efficacy of SGLT-2 versus placebo in adult type 1 diabetic patients. We also checked the reference lists of the original studies and reviewed articles and meta-analyses downloaded electronically, to identify other qualifying reports. There were no language restrictions. The search terms were: “Dapagliflozin OR Empagliflozin OR Sotagliflozin OR Canagliflozin OR SGLT2 OR Sodium-Glucose Cotransporter 2 Inhibitor OR SGLT2 Inhibitor OR ipragliflozin OR Sodium Glucose transporter 2 Inhibitors OR SGLT2 Inhibitors” AND “Diabetes Mellitus, Type 2 [Mesh] OR Diabetes Mellitus, Type 1 OR Type 1 Diabetes OR Diabetes Mellitus, Ketosis-Prone OR Diabetes, Autoimmune OR Diabetes Mellitus, Juvenile-Onset OR Juvenile-Onset Diabetes OR Diabetes Mellitus, Insulin-Dependent OR IDDMM OR Diabetes Mellitus, Insulin-Dependent, 1 OR Brittle Diabetes Mellitus OR Diabetes Mellitus, Sudden-Onset”.

Study selection
Eligible randomised trials had to meet the following criteria:
— they compared the efficacy and safety of SGLT2 inhibitors with placebo;
— they were randomised controlled trials written in English or non-English languages
— they compared SGLT2 inhibitors with placebo as adjunctive therapy to insulin for T1DM in patients aged 18 years or above, regardless of participant gender or race.

Other therapeutic, not double-blind, non-randomised studies, or those that involved non-human species, were excluded. Studies involving children and observational studies were also excluded.

Outcome measures
The main outcome measures of this analysis were: glycated haemoglobin (HbA1c) levels, body weight, and estimated glomerular filtration rate (eGFR) after 1, 3–4, 6, and 12 months of treatment, compared with placebo.

Data extraction and risk of bias assessment
Y.X. Huang and Z.J. Jiang performed data extraction and risk of bias assessment, respectively, and independently extracted the data in duplicate according to the Cochrane Intervention System Evaluation Manual, using a pre-designed data collection form. The extracted data included the following: the first author, year of publication, country, study population size, research and design, baseline characteristics of participants, and details of the experimental and control interventions (including dose and follow-up times). Any disputes were settled by consensus. The quality of the randomised controlled trials was assessed by means of the Cochrane bias risk tool. The bias risk map and bias summary risk for this study are shown in the Supplementary Data Figure S1 and S2. [16].

Data synthesis and analysis
The analysis was conducted using RevMan5.3 and Stata V.14.0 software, following the Cochrane Manual for Systematic Evaluation of Interventions [16], and was reported according to PRISMA Guidelines [17] (Preferred Reporting Item for Systematic Evaluation and Meta-analysis). For continuous variables, we extracted the average difference between the intervention and control groups, and the corresponding standard deviation from baseline. When the standard deviation was not reported, it was obtained from the standard error, as described in the Cochrane manual. When only the difference between the average values of the two groups was reported, the standard deviation was also calculated according to the Cochrane Manual [16]. Heterogeneity was evaluated based on I² statistics. We used regression analysis to explore the reasons for results showing large heterogeneity [18]. Publication bias was visually assessed by drawing funnel plots and by performing Egger weighted regression tests [19]. We conservatively used the prior random effects model, assuming there were significant differences in therapeutic effects between studies.

Results

Literature retrieval and study selection
A database search led to the identification of 2711 eligible research records, 298 of which were excluded due to duplication. After screening the titles and abstracts of the remaining 2413 studies, a total of 2386 studies were excluded based on the inclusion and exclusion criteria: 24 were duplicates, 1756 involved type 2 diabetes, 134 were animal experiments, 441 had no original data, and 31 were unrelated. Finally, a total of 27 full articles were considered for further analysis. Because 10 of these were duplicates, a total of 17 studies evaluating the efficacy of SGLT2 combined with insulin therapy in patients with type 1 diabetes were included. The articles written by Dandona 2017 and Dandona 2018 were about the same study but differed in the follow-up duration.
(24 and 52 weeks, respectively). The entire process is summarised in Figure 1.

**Research characteristics**
All the studies included in this meta-analysis were double-blind, randomised controlled trials and were published as full-length articles from 2014 to 2018. The follow-up time in individual studies ranged from 1 week to 52 weeks. Baseline characteristics were well balanced in the individual studies. A placebo was administered in the control arm in all trials (Tab. 1).

**Search results and quality assessment**

**Glycated haemoglobin (HbA1c)**
The meta-analysis revealed that SGLT2 therapy significantly reduced HbA1c levels by 0.37% (weighted mean difference [WMD]: –0.37%; 95% CI: –0.51, –0.25; I² = 91%; p < 0.00001), the 6-months subgroup (WMD: –0.40%; 95% CI: –0.44, –0.36; I² = 99%; p < 0.00001), and the 12-months subgroup (WMD: –0.30%; 95% CI: –0.35, –0.26; I² = 98%; p < 0.00001). Moreover, the 1-month subgroup showed no significant difference with the 3–4-months subgroup. Similarly, the 3–4-months subgroup showed no significant difference with the 6-months subgroup. However, there was a significant difference between the 6-months subgroup and the 12-months subgroup (p < 0.1). The magnitude of the HbA1c lowering effect waned with longer duration of treatment after six months (up to 12 months) (Fig. 2).

**Body weight**
The meta-analysis revealed that SGLT2 therapy significantly reduced body weight by 2.88 kg (WMD: –2.88; 95% CI: –3.11, –2.65; I² = 99%; p < 0.00001). The subgroup analysis indicated that all subgroups showed statistically significant differences with placebo, including the 1-month subgroup (WMD: –1.77; 95% CI: –1.93, –1.61; I² = 0%; p < 0.00001), the 3–4-months
Table 1. Research characteristics

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<th>First author</th>
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subgroup (WMD: –2.83; 95% CI: –3.90, –1.76; F = 86%; p < 0.00001), the 6-months subgroup (WMD: –2.87; 95% CI: –3.19, –2.55; F = 100%; p < 0.00001), and the 12-months subgroup (WMD: –3.21; 95% CI: –3.86, –2.55; F = 100%; p < 0.00001). Moreover, the 1-month subgroup showed a significant difference with the 3–4-months subgroup (p < 0.1), but the 3–4-months subgroup showed no significant difference with the 6-months subgroup. Similarly, there was no significant difference between the 6-months subgroup and the 12-months subgroup.

Weight loss plateaued after 3–4 months of treatment, but thereafter the weight remained relatively stable and was maintained until 12 months (Fig. 3).

Estimated glomerular filtration rate (eGFR)
The meta-analysis revealed that SGLT2 therapy significantly reduced the eGFR by 0.67 mL/min/1.73 m² (WMD: –0.67; 95% CI: –0.71, –0.63; F = 100%; p < 0.00001). The subgroup analysis indicated that both the 6-months subgroup (WMD: –0.51; 95% CI: –0.55, –0.46; F = 100%; p < 0.00001) and the 12-months subgroup showed statistically significant differences with placebo (WMD: –1.32; 95% CI: –1.42, –1.23; F = 72%; p < 0.00001). Moreover, there was a significant difference between the 6-months subgroup and the 12-months subgroup (p < 0.1). Hence, the eGFR lowering effect increased with longer duration of treatment after six months (up to 12 months) (Fig. 4).

**Evaluated parameters in different time-points**
Figure 5 shows the reduction of HbA₁c (black line) compared with the control group in different follow-up time (1 month, 3–4 months, 6 months, 12 months); it also shows the reduction of body weight (red line) compared with the control group in different follow-up time (1 month, 3–4 months, 6 months, 12 months), and that the reduction of eGFR (blue line) compared with the control group in different follow-up time (6 months, 12 months). The data are represented by the table included in the figure. The analysis was performed with log-transformed values.
by analysis of covariance and corrected for the baseline values. In both panels, means and 95% confidence intervals are shown.

**Risk of bias in the study and meta-regression**

We used the Cochrane Risk Bias Assessment Tool included in Revman 5.3 to assess bias in the 17 randomised
trials and constructed a bias risk map and a bias risk summary map. We identified three studies with one high risk of bias, six studies with one or two unknown risks of bias, and eight high-quality studies. We conducted meta-regression on the outcome indicators showing large heterogeneity to explore the causes of the heterogeneity, and finally only the HbA1c results showed statistical significance. Our meta-regression analysis detected a linear correlation between HbA1c levels and follow-up time (regression coefficient 1.003; p = 0.013) (Supplementary File — Figure S3).

Discussion

Short and medium-term efficacy of sodium glucose cotransporter 2 inhibitors are a new class of antidiabetic drugs, which work independently of insulin. They have been shown to improve blood sugar control, body weight, and blood pressure in T2DM [9]. Previous studies [12, 13] also confirmed that SGLT2 inhibitors are effective when used to treat T1DM. Because adjuvant drug therapy for type 1 diabetes is usually long-term, it is crucial to determine the magnitude and persistence of the therapeutic effects of these agents. In this review, we summarise the data of 17 clinical trials conducted in patients with type 1 diabetes. When the results of all the placebo-controlled trials were combined, we found that although changes in HbA1c, body weight, and eGFR differed during the treatment period, the therapeutic effects of the SGLT-2 inhibitors, based on these three outcomes, could last up to 12 months. At present, there are no large-scale experiments longer than 12 months to determine the duration of the therapeutic effects of SGLT2 inhibitors in patients with type 1 diabetes.

Glycosylated haemoglobin (A1c) is an effective surrogate marker for evaluating the success of long-term diabetes treatment. According to the U.S. Food and Drug Administration (FDA), the primary endpoint to determine the efficacy of drugs that lower blood glucose levels should be A1c reduction [37]. HbA1c has been proven to be highly valuable for patient management, it is an important measure of population health, and is still an effective indicator of glycation as a complicating risk factor [38]. Controlling glycosylated haemoglobin levels reduces the side effects of insulin therapy in T1DM patients. Based on our meta-analysis, SGLT2 inhibitors show a relatively stable HbA1c lowering effect during the first six months of treatment, but then it wanes after six months of treatment (up to 12 months).

Compared with short-term treatment, long-term treatment with SGLT2 inhibitors can lead to continuous weight loss. These findings are important. Although significant progress has been made in insulin preparation, delivery systems, and blood glucose monitoring, only one third of patients achieve the desired blood glucose target, and many patients become overweight.

Figure 5. Evaluated parameters in different time-points. HbA1c — glycated haemoglobin; eGFR — estimated glomerular filtration rate.
or obese [39–41]. Previous evidence shows that moderate weight loss is associated with a beneficial reduction of cardiovascular risk factors [42, 43]. Losing weight may also help patients get psychological support from treatment plans. Based on our meta-analysis, weight loss plateaued after 3–4 months of treatment with SGLT2 inhibitors; thereafter, the weight was relatively stable up to 12 months.

In the early stages of diabetes, the glomerular filtration rate can increase, and this is one of the mechanisms of diabetic nephropathy. The early effects of SGLT2 inhibitors on glomerular ultrafiltration and urinary protein excretion are an indication of their potential renal protective effects [11]. Our study showed that the eGFR lowering effect increased after six months of treatment (up to 12 months). Hence, SGLT2 inhibitors as adjuvant drugs in T1DM also have long-term renal protective effects.

EI Masri et al. [13] carried out two subgroup analyses of dose and treatment duration, but only included four randomised controlled trials and only analysed the effects of dose and treatment duration on HbA1c and body weight, two outcome indicators. Xu et al. [12] performed a meta-analysis, together with three gradient dose sub-analyses, of SGLT2 inhibitors in T1DM, and pointed out that more sub-analyses were needed, such as long- and short-term treatment analyses. In addition to previous studies [12], four new studies [24, 30, 31, 34] have been added to the current meta-analysis, including the experimental results of the new drug ipragliflozin [30, 31]. Hence, this is the most comprehensive and reliable meta-analysis so far, including 17 trials. We performed subgroup analysis based on treatment duration at four time points and analysed changes in HbA1c, body weight, and eGFR over time. The extensive data collected in this study allows us to compare SGLT2 inhibitors and placebo at different time points, revealing the effects of SGLT-2 inhibitors over time. We also explored heterogeneity and identified the heterogeneous sources of glycated haemoglobin. Unfortunately, we were unable to successfully explore the heterogeneous source of other parameters. We studied the overall effects of SGLT2 inhibitors as a class of drugs, and in the process, we may have neglected to consider the influence of different drug types. The therapeutic effects may also be influenced by the characteristics of the participants in clinical trials and may differ for different molecules. In addition, most of this research is industry-funded, which may lead to further bias. We may also have missed unpublished reports. Many other reasons can explain the observed heterogeneity, and these viewpoints deserve further attention in specially designed meta-analyses.

Conclusion
This review provides information about the therapeutic effects of SGLT2 inhibitors in T1DM. More long-term randomised trials are needed to determine the long-term effects of SGLT2 inhibitors as adjuvants in the treatment of T1DM patients.

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Availability of data and materials
The analysed data sets generated during the study are available in the published article.

Authors’ contributions
Y.H. and Z.J. contributed equally; Y.H. conceptualised and designed the study; Z.J. and Y.H. conducted the database search, study selection, and data extraction process; Y.W. conducted quality assessment of screened studies; Z.J. and Y.H. wrote the manuscript; Y.W. approved the final version of this manuscript and takes responsibility for its contents.

Ethical approval and consent to participate
Not applicable.

Patient consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

References


