Dapagliflozin: a Sodium Glucose Cotransporter 2 Inhibitor in Development for Type 2 Diabetes

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ABSTRACT

Type 2 diabetes mellitus (T2DM) is a growing worldwide epidemic. Patients face lifelong therapy to control hyperglycemia and prevent the associated complications. There are many medications, with varying mechanisms, available for the treatment of T2DM, but almost all target the declining insulin sensitivity and secretion that are associated with disease progression. Medications with such insulin-dependent mechanisms of action often lose efficacy over time, and there is increasing interest in the development of new antidiabetes medications that are not dependent upon insulin. One such approach is through the inhibition of renal glucose reuptake. Dapagliflozin, the first of a class of selective sodium glucose cotransporter 2 inhibitors, reduces renal glucose reabsorption and is currently under development for the treatment of T2DM. Here, we review the literature relating to the preclinical and clinical development of dapagliflozin.

Keywords: dapagliflozin; diabetes; SGLT2 inhibitor

INTRODUCTION

More than 285 million people worldwide have diabetes and this number is estimated to climb to 438 million by 2030.1 Type 2 diabetes mellitus (T2DM) is the most common form, accounting for approximately 85% of all cases.2

T2DM is a progressive disorder characterized by hyperglycemia, peripheral insulin resistance (in most cases), and a decrease in insulin secretion.3,4 Development of T2DM is a lengthy multistage process and changes in insulin resistance, insulin secretion, and plasma glucose can be present many years before a diagnosis of T2DM is made.5 Increasing insulin resistance (reduced sensitivity of tissues to insulin) generally occurs as the first in this sequence of events and is followed by a compensatory increase in insulin secretion by the pancreatic beta-cells, thereby maintaining close to normal plasma glucose levels.4,6 In those individuals who ultimately develop T2DM a combination of beta-cell dysfunction and a reduction in beta-cell mass culminates in decreasing insulin secretion and associated hyperglycemia. By the time
T2DM is diagnosed 50% of the beta-cell function may already be lost. In addition, excessive and inappropriate glucagon secretion is commonly observed in T2DM, which might exacerbate hyperglycemia and complicate management of the disorder.

As a result of the deleterious effects of chronic hyperglycemia the majority of individuals with T2DM will experience microvascular and/or macrovascular complications. These include diabetic retinopathy, nephropathy, neuropathy, peripheral arterial disease, stroke, and coronary artery disease. The ultimate goal of any pharmacological intervention in T2DM is to limit microvascular and potentially macrovascular complications by maintaining plasma glucose levels within a relatively normal range. This link between glycemic control and vascular complications was highlighted in the UK Prospective Diabetes Study (UKPDS). Glycated hemoglobin A1c (HbA1c) levels were used as a determinant of glycemic control, and for every 1% reduction in HbA1c it was estimated that there would be an associated 37% risk reduction for microvascular complications and a 21% reduction in diabetes-related deaths.

CURRENTLY AVAILABLE TREATMENTS

There is a well-established association between weight gain and T2DM, and the majority of T2DM patients are overweight or obese. While in some countries treatment of T2DM often begins with the introduction of medications in addition to lifestyle modification, in others lifestyle changes are the first course of treatment implemented and are aimed at restoring glycemic control through diet modification and the introduction of a regular exercise program. However, most patients will be unable to maintain normoglycemia for any length of time without pharmacological intervention. Furthermore, due to the progressive nature of T2DM, an intensification of this initial treatment is often required to maintain glycemic control. This can involve the escalation of medication dosage and/or the inclusion of additional antidiabetes medications into the ongoing treatment plan. A wide choice of antidiabetes medications are available, the majority of which target the increasing insulin resistance or decreasing insulin secretion and are listed below.

- Metformin is generally recommended as the first line of treatment in T2DM. It decreases hepatic glucose production, improves glucose clearance through an improvement of hepatic insulin sensitivity, decreases fatty acid oxidation, and increases glucagon-like peptide 1 (GLP-1; an endogenous peptide secreted from the gastrointestinal tract and a potent stimulator of glucose-dependent insulin release) levels.
- Sulfonylureas, such as glimepiride and glipizide, inhibit pancreatic beta-cell K<sub>ATP</sub> channels and enhance insulin secretion.
- Thiazolidinediones (TZDs), such as rosiglitazone and pioglitazone, are peroxisome proliferator-activated receptor-gamma agonists. They increase the sensitivity of muscle, fat, and liver to endogenous and exogenous insulin indirectly reducing hepatic glucose production by altering adipose tissue lipid metabolism.
- Meglitinides, such as repaglinide and nateglinide, also bind to the beta-cell K<sub>ATP</sub> channel, albeit at a different site, and stimulate insulin secretion.
- GLP-1 mimetics, including exenatide and liraglutide, bind to GLP-1 receptors at several sites including pancreatic beta-cells. They potentiate meal-related glucose-dependent insulin secretion and glucagon suppression.
and delay gastric emptying resulting in reduced postprandial hepatic glucose production and enhanced peripheral glucose uptake.19

- Dipeptidyl peptidase-4 (DPP-4) inhibitors, such as sitagliptin, vildagliptin, and saxagliptin, prevent the degradation of endogenous GLP-1, thereby prolonging its insulinotropic activity.20,21

- Amylin mimetics (eg, pramlintide) are synthetic analogs of the beta-cell hormone amylin.22 They act by slowing down the movement of food through the intestine and the absorption of glucose from the intestine, reducing postprandial glucose levels. Amylin mimetics also inhibit postprandial glucagon production.16

- The alpha-glucosidase inhibitors, such as acarbose, are one of the few classes of antidiabetes agents that do not have an insulin-dependent mechanism of action. They act by reducing the breakdown of oligosaccharides to monosaccharides in the proximal small intestine, thereby lowering postprandial glucose levels.16

- Insulin treatment provides glycemic control through direct stimulation of the insulin receptor.16

With the continual decline in insulin secretion and sensitivity that occurs as T2DM progresses, medications that rely upon those mechanisms for their activity frequently lose efficacy and, despite the availability of several different classes of antidiabetic agents, up to 60% of T2DM patients still do not achieve their target glycemic goals.23 There is a need, therefore, for orally active antidiabetes medications that act via insulin-independent mechanisms. One such approach currently under clinical investigation is through inhibition of renal glucose reabsorption and the consequent enhancement of urinary glucose excretion.

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**RENAL GLUCOSE REABSORPTION AND SODIUM GLUCOSE COTRANSPORTER 2 (SGLT2) INHIBITORS**

The role of the kidneys in maintaining normoglycemia, through the filtration and reabsorption of glucose as well as gluconeogenesis, is well established. Every day 180 L of plasma are filtered through the kidneys and, in normoglycemic individuals, this translates to approximately 180 g of glucose.24,25 Under normal conditions the ability of the kidneys to reabsorb glucose from the glomerular filtrate is extremely effective, with less than 0.5 g/day of this filtered glucose ultimately appearing in the urine. Under periods of hyperglycemia the amount of filtered glucose reabsorbed increases in proportion to the plasma glucose concentration until the resorptive capacity of the tubules is exceeded, at which point the excess glucose is excreted in urine (glucosuria).26

Glucose reabsorption in the renal tubules is accomplished by way of SGLTs that move glucose into the renal epithelial cells. The majority of the glucose is reabsorbed from the glomerular filtrate by SGLT2.24 SGLT2 is a high-capacity, low-affinity transporter predominantly expressed in the kidney where it is exclusively found in the brush border membrane of the S1 segment of the proximal tubule.25,27,28 The remainder of the glucose is reabsorbed from the filtrate in the distal S3 segment of the renal proximal tubule by the high-affinity, low-capacity glucose transporter sodium glucose cotransporter 1, SGLT1 (Figure 1A).29,30 However, while SGLT2 is predominantly expressed in the kidney, SGLT1 is also highly expressed in the small intestine, where it is involved in the transport of glucose across the brush border membrane.30 In the renal tubule an electrochemical gradient generated by the Na⁺/K⁺ ATPase located in the
basolateral membrane drives the movement of sodium ions across the luminal membrane and provides the driving force for glucose cotransport (Figure 1B).²⁴

Increasing urinary glucose excretion through an inhibition of glucose reabsorption represents an attractive method of maintaining blood glucose control without the accompanying risk of hypoglycemia seen with those antidiabetes medications that increase insulin secretion. In addition, the caloric loss associated with the excreted glucose may be anticipated to cause weight loss.

The concept of normalizing glucose levels through an increase in urinary glucose excretion is not a new one. The antihyperglycemic properties of the glucosuric agent phlorizin, an SGLT inhibitor derived from apple tree bark, have been known for many years. However, clinical use of phlorizin was not feasible due to nonselectivity. In addition, phlorizin had limited oral bioavailability due to the degradation of an O-glucoside linkage by gastrointestinal beta-glucosidases.³¹,³²

In the quest for a more attractive clinical candidate a number of specific inhibitors of SGLT2 have been developed. Several are undergoing late phase clinical testing for T2DM, eg dapagliflozin, canagliflozin, ASP1941, LX4211, and BI10773.³³ Two further SGLT2 inhibitors that displayed promising initial results, sergliflozin and remogliflozin etabonate, were discontinued for a variety of reasons, including nonselectivity, unfavorable pharmaceutical properties, or development of replacement SGLT2 compounds.³⁴-³⁶

Dapagliflozin is furthest along in development and is currently in phase 3 trials. For the remainder of this article we will review the preclinical and clinical data available for dapagliflozin.

DAPAGLIFLOZIN

Preclinical Studies

In preclinical studies dapagliflozin exhibited potent inhibition of human SGLT2 with an EC₅₀ of 1.1 nM and a 1200-fold selectivity for human SGLT2 over human SGLT1, and contained a beta-glucosidase-resistant C-glucoside in place of the O-glucoside linkage, permitting
oral administration. In both normal and experimentally diabetic rats dapagliflozin induced significant renal glucose excretion. Normal rats exhibited an improved glucose tolerance profile with a single dose of dapagliflozin and this was associated with reductions in glucose excursions following oral glucose tolerance testing (OGTT). In two different rat models of diabetes (streptozotocin and Zucker diabetic fatty [ZDF] rats) hyperglycemia was reduced after administration of a single oral dose of dapagliflozin and was observed within 6 hours of dosing. Reductions in both fasting and postprandial glucose levels were maintained in ZDF rats over 2 weeks with once-daily dosing with dapagliflozin. The promising efficacy, tolerability, and overall favorable absorption, distribution, metabolism, and excretion profile of dapagliflozin led to its clinical evaluation in healthy and T2DM subjects.

Clinical Studies

Pharmacokinetics

Single-ascending (SAD) and multiple-ascending (MAD; 2-week duration) dose studies were performed in healthy and T2DM subjects to evaluate the pharmacokinetic and pharmacodynamic profile of dapagliflozin (2.5-500 mg SAD; 2.5-100 mg/day MAD). After oral administration absorption of dapagliflozin was rapid in both healthy and T2DM participants (Tmax of approximately 1-2 hours). It demonstrated a half-life of approximately 16 to 17 hours in both populations. Dapagliflozin is highly protein bound and renal excretion was minimal throughout the 2-week studies in both populations. Dapagliflozin is primarily metabolized via uridine diphosphate glucuronosyltransferase 1-9 (UGT1A9) to form the inactive glucuronidated metabolite, dapagliflozin 3-O-glucuronide (BMS-801576), which is excreted in the urine. Total exposure to dapagliflozin was proportional to dose and similar on day 1 and day 14 in both healthy and T2DM subjects. After 14 days dapagliflozin accumulation was minimal and not different between the healthy and T2DM subjects (accumulation index: 1.20-1.30 and 1.23-1.33, respectively). Thus, the pharmacokinetic profile of dapagliflozin was consistent with a once-daily administration protocol.

Pharmacodynamics

Renal glucose excretion was enhanced by dapagliflozin in healthy and T2DM individuals in a dose-dependent fashion and reached a plateau at the 20 mg/day dose. Following 2 weeks of daily dapagliflozin dosing, cumulative amounts of urinary glucose ranged from 20 to 55 g/day in healthy subjects and from 37 to 70 g/day in T2DM patients. In two separate 12-week trials once-daily dapagliflozin was administered to T2DM participants who were either treatment naive (dapagliflozin 2.5-50 mg once daily) or had ongoing insulin therapy with insulin sensitizers (dapagliflozin 10-20 mg once daily). A similar

Figure 2. Total cumulative amount of glucose excreted in the urine over 24 hours from type 2 diabetes mellitus patients following 14 days of treatment with either dapagliflozin or placebo. A significant increase in urinary glucose excretion was seen at all doses plateauing at the 25 mg once-daily dose. Reproduced with permission.
increase in urinary glucose excretion (up to approximately 85 g/day) was observed at the conclusion of the 12-week treatment period in both populations.\textsuperscript{40,41}

\textbf{Clinical Efficacy in T2DM Patients}

Healthy subjects administered dapagliflozin (2.5-100 mg once daily) for up to 2 weeks exhibited no change in glycemic parameters.\textsuperscript{38} However, T2DM patients administered dapagliflozin (5-100 mg once daily) over the same time period exhibited significant dose-dependent reductions in fasting serum glucose at day 13. In addition, improvements in OGTT were also observed on day 13 with all doses of dapagliflozin in the T2DM individuals.\textsuperscript{39}

In 12-week studies of once-daily dapagliflozin significant reductions in glucose excursions were seen in T2DM patients following OGTT, as well as dose-proportional reductions in fasting plasma glucose. Significant improvements in HbA\textsubscript{1c} levels were also seen with dapagliflozin in T2DM subjects over the 12-week dosing period.\textsuperscript{40,41} A $-0.55\%$ to $-0.90\%$ change in HbA\textsubscript{1c} was seen following 12 weeks of dapagliflozin treatment (2.5-50 mg once daily) versus $-0.18\%$ with placebo in treatment-naïve T2DM patients (baseline mean HbA\textsubscript{1c} 7.6\% to 8\%). Furthermore, significantly more patients achieved their target glycemic goal of $<7\%$ HbA\textsubscript{1c} with 50 mg once daily dapagliflozin (59\%) compared with placebo (32\%) following the 12-week treatment period.\textsuperscript{40} In T2DM patients currently receiving insulin plus an insulin sensitizer, changes of $-0.61\%$ to $-0.69\%$ in HbA\textsubscript{1c} (baseline mean 8.3\% to 8.5\%) were observed compared with a mean increase of 0.09\% in the placebo group (Figure 3). Around 63\% of those subjects achieved a $\geq0.5\%$ reduction at week 12 in HbA\textsubscript{1c} levels with dapagliflozin (10 and 20 mg once daily) versus 16\% in the placebo group.\textsuperscript{41} This reduction in HbA\textsubscript{1c} in patients inadequately controlled on insulin has been shown to be maintained following 24 and 48 weeks of dapagliflozin treatment.\textsuperscript{42,43} A significant reduction in HbA\textsubscript{1c} of $-0.75\%$ to $-0.90\%$ (mean baseline HbA\textsubscript{1c} 8.5\%) was observed at 24 weeks with dapagliflozin compared with placebo ($-0.3\%$).\textsuperscript{42} At 48 weeks a change of $-0.74\%$ to $-0.94\%$ with HbA\textsubscript{1c} (baseline mean approximately 8.5\%) was recorded with dapagliflozin (2.5-10 mg once daily) compared with $-0.43\%$ in the placebo group.\textsuperscript{43} Dapagliflozin also reduced the rate of insulin uptitration and study discontinuation as a result of a lack of glycemic control following 48 weeks of treatment in patients with T2DM.\textsuperscript{44} In comparison, the change in mean HbA\textsubscript{1c} from baseline mean (8.41\%) seen with the GLP-1 mimetic exenatide in T2DM patients inadequately controlled on insulin was $-1.71\%$ with exenatide versus $-1.00\%$ with placebo.
over 30 weeks.\textsuperscript{45} Owing to differences between the trial designs it is not possible to draw direct comparisons between the trials, although it is notable that while the reduction in HbA\textsubscript{1c} with exenatide was numerically greater than observed with dapagliflozin the reduction in HbA\textsubscript{1c} in the placebo group was also larger than that seen in the dapagliflozin trial.

When administered for 24 weeks as a monotherapy to treatment-naïve T2DM patients dapagliflozin (2.5-10 mg once daily) produced dose-dependent reductions in HbA\textsubscript{1c} levels from a baseline mean of 7.8% to 8.0% that were statistically significant with the 5 mg and 10 mg doses (–0.77% and –0.89%, respectively) versus –0.23% with placebo.\textsuperscript{46} In patients with ongoing metformin therapy there was a –0.67% to –0.84% change in HbA\textsubscript{1c} from a baseline mean of 7.92% to 8.17% with dapagliflozin (2.5-10 mg once daily) at 24 weeks as compared with –0.30% in the placebo group.\textsuperscript{47} Furthermore, in a 24-week study evaluating the effectiveness of dapagliflozin in patients with T2DM inadequately controlled on the sulfonylurea glibenclamide, significant reductions in HbA\textsubscript{1c} from the baseline mean (8.07% to 8.15%) of –0.58% to –0.82% with dapagliflozin (2.5-10 mg once daily) were observed compared with –0.13% with placebo.\textsuperscript{48}

In the longest reported trial with dapagliflozin so far (52 weeks) reductions in HbA\textsubscript{1c} with dapagliflozin from baseline mean (approximately 7.72%) were reported in patients poorly controlled with metformin. The adjusted mean change from baseline was –0.52% with dapagliflozin (≤10 mg once daily) compared with a similar reduction of –0.52% with the sulfonylurea glipizide after 52 weeks of treatment.\textsuperscript{49}

Total body weight loss occurred in all groups in the 12-week study in treatment-naïve T2DM patients. The reduction in body weight was continuous across the dosing period and ranged from –2.5% to –3.4% at 12 weeks and –1.2% in placebo although this did not reach statistical significance.\textsuperscript{40} After 12 weeks the change in body weight with dapagliflozin in T2DM patients taking insulin and insulin sensitizers was –4.3 to –4.5 kg in the dapagliflozin treatment groups versus –1.9 kg with placebo (Figure 4).\textsuperscript{41} Both 24-week and 48-week studies were consistent with this effect of dapagliflozin on weight loss (24 weeks: –1.67 kg with 10 mg vs. 0.02 kg placebo, \(P\leq0.0001\); 48 weeks: –1.79 kg with 10 mg vs. –0.18 kg).\textsuperscript{42,43} In treatment-naïve T2DM patients, decreases of up to 3.3 kg were observed after 24 weeks of treatment with dapagliflozin and decreases of 2.2 kg were evident in the placebo group, although this was not reported as being significant. In metformin-treated T2DM patients, reductions of up to 3.0 kg in body

![Figure 4. Changes from baseline in mean body weight over 12 weeks of treatment with dapagliflozin (DAPA) or placebo (PLA) in subjects with ongoing insulin (INS) therapy. A trend towards continuous and greater weight loss was seen over 12 weeks with 10 and 20 mg once-daily dapagliflozin versus placebo. LOCF=last observation carried forward. Copyright 2009 American Diabetes Association. From Diabetes Care, Vol. 32, 2009; 1656-1662. Reprinted with permission from The American Diabetes Association.](image-url)
weight were seen with dapagliflozin (2.5-10 mg once daily) as compared with 0.9 kg in the placebo group following 24 weeks of treatment, with similar reductions following 52 weeks of treatment.46,47,49

**Tolerability and Side Effects**

In T2DM patients adverse events observed in the dapagliflozin treatment and placebo groups were similar in frequency and were generally mild in nature whether for patients who were treatment naïve or for those receiving ongoing metformin or insulin plus insulin sensitizer therapy.40-47 Few cases of hypoglycemia were observed and these were generally mild, self-limiting, and occurred with a similar frequency in the placebo group; none were severe.40-47

A mild diuretic effect of dapagliflozin was observed at week 12 in the treatment-naive patients (an increase in urinary output of up to 470 mL/day above baseline compared with a 112 mL/day decrease in placebo). In those T2DM patients taking insulin plus oral insulin sensitizers there was an increase in urine output of up to 444 mL/day above baseline compared with 255 mL/day with placebo (although the difference was not reported as being significant).40,41 Consistent with this observation, dapagliflozin appears to be associated with a mild reduction in mean blood pressure with no evidence of orthostatic hypotension.40,41

The long-term effects of dapagliflozin on renal function are as yet unknown, as are the effects of dapagliflozin when used in cases of compromised renal function or when coadministered with certain antihypertensive drug classes that may affect renal physiology. However, no clinically relevant changes in glomerular filtration rate have been reported with dapagliflozin treatment.39-41 Owing to the inhibition of the cotransportation of sodium and glucose, dapagliflozin administration may result in an increase in urinary sodium. Acute transient increases in urine sodium have been observed with dapagliflozin treatment.39 Median changes from baseline in urinary sodium excretion were +34.7, +40.2, and +48.0 mEq for the 5, 25, and 100 mg dapagliflozin treatment groups, and −15.1 mEq for the placebo group over the initial 24 hours. With continuing daily administration this increase in sodium excretion appeared to normalize, with median changes from baseline at day 13 of +1.8, +8.9, and −5.7 mEq for the 5, 25, and 100 mg dapagliflozin dose groups and +16.4 mEq for placebo. Despite the reported transient increase in urinary sodium excretion there is no evidence to suggest that this is reflected in any changes in serum sodium levels.39

Although dapagliflozin has been associated with increases in urine volume, serum magnesium, serum phosphate, and serum uric acid these changes were still within normal physiological ranges and there was little evidence to suggest that dapagliflozin would produce clinically significant changes in electrolyte balance.

Because it was hypothesized that increasing urinary glucose may promote microbial growth, potential urinary tract infections (UTIs) and genital infections were carefully monitored. While the frequency of reported UTIs was similar between the dapagliflozin and placebo groups over the 12-week time period, symptoms suggestive of genital infections (in particular vulvocandidiasis) were reported more frequently in the groups administered dapagliflozin.40,41

In the 24-week studies, signs and symptoms suggestive of genital infection were reported in up to 13% of T2DM patients who were either treatment naïve or who were inadequately controlled with metformin as compared with 1.3% and 5% in the respective placebo groups.46,47 No increased rate of discontinuation occurred as a result of these signs and symptoms suggestive of UTIs or genital infections.39,41-43,46-48 It is not
known at this time how these observations in controlled trials will translate into infection rates in everyday clinical practice until more widespread experience is gained over a longer time frame.

Additional data are necessary to fully elucidate the long-term effects of glucose excretion induced by dapagliflozin. It is noteworthy, however, that people with a condition known as familial renal glucosuria (FRG), arising primarily from a mutation of the gene encoding for the SGLT2 protein (SLC5A2), live normal healthy lives, with no adverse events, despite the consistently high levels of urinary glucose. Experience with FRG is based on a very limited number of cases, and caution should be exercised when making comparisons with the general diabetic population. In these patients glucosuria can range from <1 to >150 g/1.73 m² per day. While FRG patients generally have few clinical complications polyuria and enuresis were observed in a single case study and were followed later by a slight delay in growth and pubertal maturation. Other reported complications with FRG include episodes of dehydration and ketosis as well as urinary tract infections and natriuresis.

However, unlike FRG, T2DM is associated with impaired immune responses, and patients with T2DM may have side effects relating to elevated urinary glucose levels. Furthermore, while glucose-induced diuresis does not appear to lead to dehydration in the majority of cases of FRG, the mild diuretic properties of dapagliflozin may require careful monitoring to ensure adequate fluid intake and warrants further investigation. One episode of renal failure did occur during treatment with 10 mg dapagliflozin in a patient concomitantly being treated with insulin and several antihypertensive medications (enalapril, carvedilol, and furosemide). The patient discontinued dapagliflozin as a result of dehydration and prerenal azotemia. The condition was resolved with oral rehydration and the withholding of enalapril and furosemide.

SUMMARY AND THERAPEUTIC APPLICATIONS

T2DM is a progressive disease with a significant impact on patients, society, and healthcare systems. The aim of treatment is to achieve near normal glycemia in order to prevent the development or slow the progression of microvascular and possibly macrovascular complications. Currently available antidiabetes treatments have many limitations, including hypoglycemia (sulfonylureas, meglitinides, and insulin), weight gain (sulfonylureas, meglitinides, TZDs, and insulin), heart failure (TZDs), gastrointestinal side effects (alpha-glucosidase inhibitors, metformin), and the need for multiple injections and/or self-monitoring (insulin, amylin mimetics, and GLP-1 analogs). Furthermore, due to the progressive nature of the disease, most patients will ultimately require multiple antidiabetes treatments to achieve glycemic targets. Hence, there is a need for new antidiabetes treatments that produce a sustainable impact on glycemic control with low risk for hypoglycemia and weight loss, and minimal need for self-monitoring. SGLT2 inhibitors are a novel class of antidiabetes therapy that are taken orally, result in improvements in glycemic parameters with a low risk for hypoglycemia, and are associated with weight loss. They have the potential to harness what has long been considered a manifestation of diabetes, glucosuria, and turn it into a therapeutic strategy. However, like other newer antidiabetes treatments, this class lacks long-term (including cardiovascular) safety data.
With such a plethora of options to treat patients with T2DM, the decision to use a particular drug, or combination of drugs, in a particular patient should be individualized based on the patient’s specific risk-benefit balance and not solely upon a drug’s perceived ability to lower HbA1c. Metformin (in addition to lifestyle changes) remains a well-established first-line treatment for patients with T2DM, due to good long-term safety data, lack of hypoglycemia or weight gain, and evidence for cardiovascular protection. However, as metformin does not halt the progression of the disease, patients with T2DM are likely to need additional antidiabetic medications administered alone or in combination with metformin. Should SGLT2 inhibitors, such as dapagliflozin, prove to have an acceptable safety profile they may have the potential to be administered alone or in combination with metformin or insulin. The lack of long-term (including cardiovascular) safety data and other outcome data may limit their use initially to specific/defined low-risk patient groups.

CONCLUSION

In conclusion, dapagliflozin has the potential to be a useful addition to currently available antidiabetes treatments as it lowers fasting and postprandial glucose levels, improves glycemic control, and causes weight loss with a low risk of hypoglycemia. However, data regarding long-term safety including urinary tract infection, genital infection, and cardiovascular safety are needed, and its place in the algorithm of T2DM management is still to be determined.

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