Structural selectivity of human SGLT inhibitors

Charles S. Hummel,1 Chuan Lu,1 Jie Liu,2 Chiari Ghezzi,1 Bruce A. Hirayama,1 Donald D. F. Loo,1 Vladimir Kepe,2 Jorge R. Barrio,2 and Ernest M. Wright1

Departments of 1Physiology and 2Molecular and Medical Pharmacology, David Geffen School of Medicine at University of California Los Angeles, Los Angeles, California

Submitted 1 September 2011; accepted in final form 20 September 2011

THE PHARMACEUTICAL INDUSTRY has targeted human renal Na+-d-glucose cotransporters (hSGLTs) to control blood glucose concentrations in patients with type 2 diabetes. A number of specific hSGLT inhibitors are in phase III clinical trials (4). Companies seek inhibitors with high specificity for hSGLT2 over hSGLT1 for three major reasons: first, hSGLT2 is believed to be responsible for the majority (>80%) of glucose reabsorption in the kidney under normal and hyperglycemic conditions; second, mutations in hSGLT2 cause familial renal glycosuria, a benign condition characterized by increased urinary glucose excretion (36, 37); and third, hSGLT1 inhibitors may cause diarrhea, mimicking the phenotype of individuals deficient in functional hSGLT1 protein (40).

Drug development has focused on modifying the chemical structure of the natural Na+-d-glucose cotransporter (SGLT) inhibitor phlorizin {1-[2-(β-d-glucopyranosyloxy)-4,6-dihydroxyphenyl]-3-(4-hydroxyphenyl)-1-propanone} to identify compounds with high potency and selectivity for hSGLT2 over hSGLT1. Early candidate drugs were O-glycosides like phlorizin, e.g., remogliflozin (10) and sergliflozin (15, 16, 21), but development was discontinued because they were rapidly hydrolyzed by glucosidases in vivo (17). This problem was overcome by removing the glycosidic linkage, e.g., C-aryl glycosides such as dapagliflozin (43). A number of studies have reported on structure-activity relationships in the development of C-aryl glycoside inhibitors with extensive modifications to both the sugar and chalcone moieties (11, 28, 35, 43).

This approach has led to promising candidate drugs, including dapagliflozin (28) and canagliflozin (30). Dapagliflozin increased urinary glucose excretion in control subjects and improved glucose homeostasis and insulin sensitivity in diabetic rats (12, 22). In diabetic human subjects, they improved glycemic control by lowering fasting and postprandial plasma glucose, all without major adverse events (2, 9, 23, 44). Despite promising clinical data, little is known about how these drugs work at a molecular level.

In our laboratory’s recent study of SGLTs expressed in human embryonic kidney 293T (HEK-293T) cells (14), we showed that differences in inhibitor dissociation rates accounted for differences in phlorizin inhibitory constant (Ki) for hSGLT2 and hSGLT1. Here we have extended these studies to dapagliflozin and several of its structural analogs. Our results provide insights into both dapagliflozin’s mechanism of action and its effects on renal glucose reabsorption.

MATERIALS AND METHODS

Cell Culture and Transfection

HEK-293T and COS-1 cells were maintained, passaged, and transfected with hSGLT2 or hSGLT1 cDNA, as described previously (14). HEK-293T is a highly transflectable derivative of the 293 cell line into which the temperature-sensitive gene for SV40 T-antigen was inserted.

Radioactive Tracer Determinations

[14C]-methyl-d-glucopyranoside (α-MDG) uptakes into COS-1 cells expressing hSGLT2 or hSGLT1 were as described (14). Cells were incubated in sodium buffer with 50 μM [14C]-MDG

Address for reprint requests and other correspondence: E. M. Wright, Dept. of Physiology, David Geffen School of Medicine at UCLA, Los Angeles, CA 90095-1751 (e-mail: ewright@mednet.ucla.edu). URL: http://140.142.237.182/

http://www.ajpcel1.org 0363-6143/12 Copyright © 2012 the American Physiological Society

C373
at 37°C for 1–40 min in the presence or absence of inhibitors: 100 μM phlorizin and 10 mM or 1 μM of each hSGLT2 inhibitor (dapagliflozin, fluoro-dapagliflozin, or galacto-dapagliflozin). Uptakes were expressed as picomoles per minute per microgram total protein, mean ± SE (Fig. 1). [14C]-α-MDG uptakes were linear for 15 min for hSGLT2 and 2 min for hSGLT1, and so the effects of inhibitors on initial rates of uptake were determined at 10 min for hSGLT2 and 1 min for hSGLT1.

Electrophysiological Experiments

Whole cell patch-clamp recordings were performed on HEK-293T cells 2 days posttransfection (14). The extracellular solution contained (in mM) 150 NaCl, 1 CaCl₂, 1 MgCl₂, 10 HEPES, pH 7.4 (“Na+ buffer”), or 150 choline chloride, 1 CaCl₂, 1 MgCl₂, pH 7.4 (“choline+” buffer). For experiments at 37°C, mannitol (100 mM) was added to the external solution to reduce noise and increase the stability of the whole cell recordings. The pipette (internal) solution contained (in mM) 145 CsCl, 5 NaCl, 11 EGTA, and 10 HEPES. Membrane potential was held at −60 mV. For hSGLT2, all experiments were performed at 37°C, and for hSGLT1, at 37 or 22°C.

Inhibitor Kinetics

Our general approach to the interaction of inhibitors with the SGLT's followed that used previously by Oulianova et al. (32) to investigate phlorizin binding to SGLTs in rabbit renal brush-border membranes but here we measure binding to specific human SGLTs using electrophysiological assays. Inhibition of steady-state, glucose-induced hSGLT current was measured as a function of external inhibitor concentration to determine the inhibition constant \( K_i \). As described previously, when the glucose concentration is equal to the sugar half-maximal inhibition constant \( K_{i,50} \), the concentration of the inhibitor producing 50% inhibition (IC50) is twice the \( K_i \) (14, 38). The hSGLT2 d-glucose \( K_{i,50} \) was 5 mM at 37°C (14) and for hSGLT1 it was 2 mM at 37°C and 0.5 mM at 22°C.

The sequence of each experiment was as follows. Current was induced by the \( K_{i,50} \) concentration of d-glucose. The solution was changed to d-glucose plus inhibitor. Once a new steady-state current was achieved, the cell was washed with d-glucose and Na+-free (choline+”) buffer for ≥3 min to remove the inhibitor. This protocol was repeated at multiple inhibitor concentrations to estimate the \( K_i \).

Some of the hSGLT1 galacto-dapagliflozin \( K_i \) determinations were performed using the two-electrode voltage clamp on Xenopus laevis oocytes expressing hSGLT1 to minimize the quantity of the inhibitor needed to complete the experiments. Oocyte isolation, preparation, injections, and electrophysiological methods were performed as described previously (24, 25).

Inhibitor off rates \( (k_{off}) \) were determined by recording the time course of recovery of the d-glucose-coupled current after removal of an inhibitor [such as phlorizin (Pz)], assuming pseudo-first-order binding and dissociation kinetics (14):

\[
\frac{k_{on}[Pz]}{[hSGLT]} = \frac{k_{off}}{Pz-hSGLT}
\]

The d-glucose-coupled current recovery followed an exponential time course with time constant \( 1/k_{off} \), and half-time for recovery was related to the off rate \( (t_{1/2, off}) = \ln(2)/k_{off} \). The on rate \( (k_{on}) \) was calculated from the empirically measured \( K_i \) values and \( k_{off} \) values \( (K_i = k_{off}/k_{on}) \). The \( k_{off} \) for hSGLT1 at 37°C was within the half-time of the solution change (5 s), and so the temperature was lowered to 22°C, and the SF-77B Fast-step solution changer (Warner Instruments, Hamden, CT) was utilized to wash out inhibitors (solution change half-time of ≤20 ms).

Synthesis of Inhibitors

Dapagliflozin \( [(1S)-1S,1S,4S,4aR,5S,9aS,11bR,11bH]-3-(4-chlorophenyl)iminophenyl]-d-glucopyranoside \) was synthesized by a reported procedure (28). Galacto-dapagliflozin was synthesized starting from dapagliflozin by adequate sugar protection and selective configuration conversion at C4’, and this was followed by fluorination to give fluoro-dapagliflozin. The phenol-aglycone of dapagliflozin [4-chloro-3-(4-ethoxybenzyl)phenol] was obtained by hydrolysis of the corresponding bromo-aglycone, prepared from commercial 4-bromo-2-chlorobenzoic acid (28). The synthesized products were characterized by Electrospray ionization mass spectrometry, 1H-NMR, 19F-NMR, and 13C-NMR, and their anomic configuration at C1’ and the epimeric configuration at C4’ were confirmed by advanced two-dimensional NMR techniques and X-ray crystallography data. [The syntheses and spectroscopic characterization will be published elsewhere.] Figure 1 shows the structures of the substrate and inhibitors used in this study.

Energy-Minimized Inhibitor Structures

Representative energy-minimized in silico chemical structures of phlorizin, dapagliflozin, fluoro-dapagliflozin, and galacto-dapagliflozin were generated using HyperChem version 6.0 (Hypercube, Gainesville, FL) (see also Ref. 13).

Statistical Analysis

Data fitting and t-tests for significance were performed using either Sigma Plot 10.0 (Systat Software, San Jose, CA) or Excel (Microsoft, Redmond, WA).

Reagents

All reagents were purchased from Fisher Scientific or Sigma and were of the highest purity available. [14C]α-MDG was purchased from Moravek Biochemicals (Brea, CA).

Fig. 1. Chemical structures of the Na+-d-glucose cotransporter (SGLT) substrate and inhibitors used in this study. α-MDG, [14C]α-methyl-d-glucopyranoside.
RESULTS

Biochemical

To provide a direct comparison with previous work on SGLT inhibitors, we first measured the effects on quasi-steady-state (40 min) \( ^{14} \text{C}\)l-\( \alpha \)-MDG uptakes into COS-1 cells expressing hSGLT1 and hSGLT2. Dapagliflozin at 10 \( \mu \)M blocked 90 ± 1% of phlorizin-sensitive sugar transport by hSGLT2, whereas fluoro-dapagliflozin and galacto-dapagliflozin blocked 80 ± 1 and 30 ± 3%, respectively (Fig. 2A). In parallel experiments with hSGLT1, we found that 1 \( \mu \)M dapagliflozin blocked 50 ± 3% of the phlorizin-sensitive \( \alpha \)-MDG uptake, whereas fluoro-dapagliflozin and galacto-dapagliflozin blocked 40 ± 3 and 20 ± 6%, respectively (Fig. 2B). The results with dapagliflozin and phlorizin are comparable with the published data (12). Next we measured the effect of the inhibitors on the initial rates of \( \alpha \)-MDG transport, and IC\(_{50}\) values for dapagliflozin and phlorizin were 6 ± 1 and 65 ± 10 \( \mu \)M for hSGLT2 and 400–800 and 400 ± 100 \( \mu \)M for hSGLT1, respectively. For the initial rate measurements, the potency of dapagliflozin for hSGLT2 was ~100-fold greater than for hSGLT1, as opposed to 1,200-fold difference reported for steady-state uptakes (12, 28).

Role of sugar moeity. To assess the contribution of the sugar moiety to inhibitor potency, we measured the effect of phloretin (the aglycone of phlorizin) and dapagliflozin-aglycone [4-chloro-3-(4-ethoxybenzyl)phenol] on the 40-min 50 \( \mu \)M \( \alpha \)-MDG uptakes: 250 \( \mu \)M phloretin inhibited hSGLT1 and hSGLT2 transport by 70 ± 5 and 90 ± 8%, respectively, which is consistent with reported IC\(_{50}\) values [140 and 25 \( \mu \)M (33)], and 300 \( \mu \)M dapagliflozin-aglycone inhibited hSGLT1 and hSGLT2 transport by 25 and 60%, respectively (Table 1). Additional experiments showed dapagliflozin-aglycone IC\(_{50}\) values of ~1,000 \( \mu \)M for hSGLT1 and 200 \( \mu \)M for hSGLT2 (Lu C, Hummel CS, and Wright EM, unpublished observations). These results demonstrate that removing glucose from the phlorizin and dapagliflozin molecules reduces their inhibitory potency by more than three orders of magnitude against both hSGLT isoforms.

Biophysical

\( K_i \) values. To investigate further the kinetics and specificity of drug interactions with hSGLT2 and hSGLT1, we turned to electrophysiological methods and examined the inhibition of the sugar-coupled Na\(^{+}\) currents, as previously developed for phlorizin (14). We first determined the \( K_i \) values from blockade of steady-state currents generated by glucose at its half-maximal concentration (\( K_{0.5} \)). For hSGLT2, dapagliflozin and fluoro-dapagliflozin had the same \( K_i \) values, ~6 \( \mu \)M, whereas for galacto-dapagliflozin it was 25 \( \mu \)M. These values are comparable to the \( K_i \) of 11 \( \mu \)M for phlorizin reported previously (Table 2). For hSGLT1, the dapagliflozin \( K_i \) was 360 ± 20 \( \mu \)M compared with 140 \( \mu \)M for phlorizin. These electrophysiological results confirm the higher selectivity of dapagliflozin as an inhibitor for hSGLT2, \( K_i \) 6 vs. 360 \( \mu \)M, and this 60-fold selectivity is consistent with our biochemical studies (see above). Again, this selectivity is 10-fold lower than the 1,000-fold ratio reported previously for the inhibition of the steady-state (2 h) accumulation of \( \alpha \)-MDG in transfected cells (12, 28).

We also measured hSGLT1 \( K_i \) values at 22°C (Table 3). For dapagliflozin and phlorizin, the \( K_i \) values were quite similar at 22°C to those at 37°C, 390 and 80 \( \mu \)M vs. 360 and 140 \( \mu \)M (Table 3), indicating a relatively low temperature dependence of inhibitor binding. This is consistent with the minimal temperature dependence of the hSGLT1 apparent affinities (\( K_{0.5} \)) at 22°C (28).

Table 1. Inhibition of \( ^{14} \text{C}\)\( \alpha \)-methyl-\( \beta \)-glucopyranoside uptake by aglycones

<table>
<thead>
<tr>
<th>Compound</th>
<th>Concentration, ( \mu )M</th>
<th>hSGLT1, %inhibition</th>
<th>hSGLT2, %inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phloretin</td>
<td>250</td>
<td>70 ± 5</td>
<td>90 ± 8</td>
</tr>
<tr>
<td>Dapagliflozin-aglycone</td>
<td>300</td>
<td>25 ± 7</td>
<td>60 ± 3</td>
</tr>
</tbody>
</table>

Values are means ± SE for \( n \geq 3 \) wells. hSGLT1 and hSGLT2, human Na\(^{+}\)-\( \beta \)-glucose cotransporters 1 and 2, respectively. The “% inhibition” is determined by the ratio of inhibition of the test compound (phloretin, dapagliflozin-aglycone) to that of 100 \( \mu \)M phlorizin (100% inhibition). The phloretin inhibition values were consistent with IC\(_{50}\) values reported in the literature [140 \( \mu \)M for hSGLT1 and 25 \( \mu \)M for hSGLT2 (33)].
for d-glucose transport over the same 15°C temperature range, 0.5 (22°C) to 1.8 mM (37°C) (14). On the other hand, the temperature dependence of the maximal rate of transport is high, ≈25 kcal/mol (14).

The affinity of hSGLT1 for fluoro-dapagliflozin (Kᵢ = 330 nM) at 22°C was similar to the parent compound, but it was 70-fold lower for galacto-dapagliflozin (Kᵢ = 25,000 nM) (Table 3). Inhibitor constants were higher for hSGLT1 relative to hSGLT2 (Tables 2 and 3): 7.2-fold for phlorizin, and 62- to 65-fold for dapagliflozin and fluoro-dapagliflozin, (390 and 330 vs. 6 nM), and 1,000-fold for galacto-dapagliflozin (25,000 vs. 24 nM).

Rates of inhibitor association and dissociation. To interpret differences in Kᵢ values among the different inhibitors and between the two hSGLT isoforms, we investigated the rates of association and dissociation. The dissociation rates of the inhibitors were determined by measuring the time course of the association and dissociation. The dissociation rates, i.e., kᵩᵢ values for each inhibitor are summarized in Table 3. The kᵩᵢ were comparable for hSGLT1 and hSGLT2, ~1 × 10⁶ M/s, except that for galacto-dapagliflozin that was 0.01 × 10⁶ M/s. The kᵩᵢ of 2- to 100-fold were faster for hSGLT1 than hSGLT2.

Phlorizin/dapagliflozin exchange. We evaluated whether dapagliflozin bound to hSGLT1 or hSGLT2 can be exchanged with phlorizin (Fig. 5). As shown for hSGLT2 in Fig. 5A, the 100 mM d-glucose current was completely blocked by 200 nM dapagliflozin. When dapagliflozin in the buffer was replaced with 1,000 nM phlorizin, the inhibition of Na⁺/glucose current was maintained, but, on washing out phlorizin, the transporter remained blocked for >5 min. As the phlorizin block of hSGLT2 is normally rapidly reversed (t½,Off = 25 s), these data suggest that phlorizin did not exchange with dapagliflozin bound to hSGLT2. In contrast, phlorizin readily exchanged with dapagliflozin bound to hSGLT1 (Fig. 5B). After inhibition of hSGLT1 with 750 nM dapagliflozin, replacement with 160 nM phlorizin maintained the inhibition. With phlorizin washout, the Na⁺/glucose current returned with a t½,Off of 9 s, identical to the time course for phlorizin alone (Table 3).

DISCUSSION

Clinical interest in SGLTs has intensified with the recent development of new oral inhibitors for treatment of type 2 diabetes. These drugs (several of them in late phase III clinical trials) are designed to lower blood glucose concentration by preventing renal glucose reabsorption by specifically targeting hSGLT2 (4). Despite this success, relatively little is known

<Table 2. Binding properties of phlorizin and dapagliflozin derivatives with hSGLT2 (37°C)

<table>
<thead>
<tr>
<th>Compound</th>
<th>n</th>
<th>Kᵢ, nM</th>
<th>t½,On, s</th>
<th>k½,Off, s⁻¹</th>
<th>kᵩᵢ, M/s (calculated)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phlorizin</td>
<td>3</td>
<td>11 ± 0.9</td>
<td>24 ± 3</td>
<td>0.030 ± 0.003</td>
<td>2.7 × 10⁶</td>
</tr>
<tr>
<td>Dapagliflozin</td>
<td>5</td>
<td>6.0 ± 0.6</td>
<td>&gt;300</td>
<td>&lt;0.002</td>
<td>&lt;3.3 × 10⁶</td>
</tr>
<tr>
<td>Fluoro-dapagliflozin</td>
<td>10</td>
<td>5.3 ± 0.8</td>
<td>170 ± 20</td>
<td>0.0048 ± 0.0006</td>
<td>0.91 × 10⁶</td>
</tr>
<tr>
<td>Galacto-dapagliflozin</td>
<td>2</td>
<td>24 ± 6</td>
<td>21 ± 3</td>
<td>0.030 ± 0.007</td>
<td>1.3 × 10⁶</td>
</tr>
</tbody>
</table>

Values are means ± SE; n, no. of cells. Kᵢ, inhibitory constant; t½,On, half-time off rate; k½,Off, inhibitor off rate; kᵩᵢ, inhibitor on rate. Phlorizin parameters are from Ref. 14.

<Table 3. Binding properties of phlorizin and dapagliflozin derivatives with hSGLT1 (22°C)

<table>
<thead>
<tr>
<th>Compound</th>
<th>n</th>
<th>Kᵢ, nM</th>
<th>t½,On, s</th>
<th>kᵩᵢ, M/s (calculated)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phlorizin (37°C)</td>
<td>&gt;3</td>
<td>140 ± 15</td>
<td>4 ± 0.5</td>
<td>0.19 ± 0.02</td>
</tr>
<tr>
<td>Phlorizin</td>
<td>4</td>
<td>80 ± 10</td>
<td>9.4 ± 6</td>
<td>0.07 ± 0.01</td>
</tr>
<tr>
<td>Dapagliflozin</td>
<td>6</td>
<td>390 ± 25</td>
<td>1.8 ± 0.1</td>
<td>0.40 ± 0.05</td>
</tr>
<tr>
<td>Fluoro-dapagliflozin</td>
<td>7</td>
<td>330 ± 30</td>
<td>1.2 ± 0.1</td>
<td>0.61 ± 0.03</td>
</tr>
<tr>
<td>Galacto-dapagliflozin</td>
<td>4</td>
<td>25,000 ± 6,000</td>
<td>0.68 ± 0.08</td>
<td>1.1 ± 0.1</td>
</tr>
</tbody>
</table>

Values are means ± SE; n, no. of cells (galacto-dapagliflozin: n = 3, determined in X. laevis oocytes expressing bSGLT1). Phlorizin parameters at 37°C are from Ref. 14.
about the mechanism of action of these compounds or the basis of selectivity for hSGLT2 over hSGLT1. We have set out to probe the mechanism of one such inhibitor, dapagliflozin (28), which provides a reference point for comparison with other structurally related hSGLT2 inhibitors currently under development by examining the kinetics of interaction with hSGLT1 and hSGLT2 (14, 25, 45). Specifically, we compared and contrasted dapagliflozin, galacto-dapagliflozin, fluoro-dapagliflozin, and phlorizin (Fig. 6) inhibition kinetics of hSGLT2 and hSGLT1, via determinations of $K_i$, $k_{On}$, and $k_{Off}$ (Figs. 3 and 4, Tables 2 and 3).

We chose to measure inhibition kinetics under initial rate conditions rather than under steady-state conditions used by others (i.e., we measured the kinetics of $\alpha$-MDG uptake in the absence of intracellular sugar). We have confirmed that dapagliflozin is a selective hSGLT2 inhibitor, and that this selectivity over hSGLT1 is largely due to differences in drug-SGLT dissociation rates: the $k_{Off}$ for hSGLT2 is more than 100-fold slower than that for hSGLT1. Under the conditions in our experiments, the dapagliflozin $K_i$ was 6 nM for hSGLT2 and 360 nM for hSGLT1. By comparison, the phlorizin $K_i$ was 10 nM for hSGLT2 and 140 nM for hSGLT1. Based on 2-h $\alpha$-MDG uptakes in transfected Chinese hamster ovary cells, IC$_{50}$ values for dapagliflozin and phlorizin were 1 and 36 nM for hSGLT2 and 1,390 and 330 nM for hSGLT1 (12, 28). The greatest discrepancy between the previous and our present results is that we observed a marked selectivity for hSGLT2 over hSGLT1, which was not apparent in previous studies.

Fig. 3. Time course of inhibitor dissociation in hSGLT2 (−60 mV, 37°C). Continuous records of sugar current (100 mM d-glucose) show reduction in steady-state transport [represented by glucose-coupled current ($I_{glucose}$)] upon application of inhibitor [half-time on rate ($t_{1/2,On}$), not shown]. Upon removal of inhibitor, the time course of d-glucose current recovery was monitored [half-time off rate ($t_{1/2,Off}$)]. A: PZ inhibited current with a $t_{1/2,On}$ of 9 s, and $I_{glucose}$ recovery occurred with $t_{1/2,Off}$ = 30 s. B: DAPA had a $t_{1/2,On}$ of 25 s, but it showed no recovery of current, even after 5 min, and thus $t_{1/2,Off}$ >> 300 s. C: F-DAPA’s $t_{1/2,On}$ was 18 s, and it showed a slow recovery of $I_{glucose}$, $t_{1/2,Off}$ = 200 s. D: G-DAPA association occurred with $t_{1/2,On}$ = 10 s, and its dissociation was rapid, $t_{1/2,Off}$ = 19 s.
work is the selectivity of dapagliflozin for hSGLT2 over hSGLT1, 1,400-fold, vs. the 60-fold ratio reported here. Such a pronounced difference is likely due to contrasting experimental approaches between the measurement based on initial rates in our biophysical assays compared with the earlier reported steady-state assays (12, 28). In our hands, the dapagliflozin IC₅₀ estimated from initial rates of [¹⁴C]H₂⁵-glucitol uptake was 5 nM, whereas, under steady-state conditions, this value was determined to be 1 nM.

Multiple lines of evidence support the conclusion that the inhibitors interact with both hSGLT1 and hSGLT2 at the glucose binding site, as first suggested by Alvarado (1). First, phlorizin binding is competitive with glucose (33). Second, phlorizin and glucose bind to SGLTs only in the presence of Na⁺ (14, 45). Third, changes in the sugar moiety changes the potency of the inhibitors: the galacto-dapagliflozin Kᵢ was higher than dapagliflozin for both hSGLT2 and hSGLT1 (Tables 2 and 3), and the binding of the aglycones was three orders of magnitude lower than those of the parent drugs (e.g., for hSGLT2, phloretin and phlorizin Kᵢ values were 25 μM and 11 nM; dapagliflozin-aglycone and dapagliflozin Kᵢ values were 200 μM and 5 nM, Tables 1–3). Fourth, modification of the glucose ring produced predictable changes in the inhibitor Kᵢ: the 2-deoxy- and 3-deoxy-sugar derivatives were very poor inhibitors of hSGLT2 (IC₅₀ ≥ 2,000 nM) (35), also in agreement with the observation that these deoxy sugars are poor substrates for hSGLT2 transport (Lu C et al., unpublished observations). In contrast, substitution of -F for -OH at C4 in both glucose and dapagliflozin (fluoro-dapaglifozin, Fig. 6) did not alter the kinetics of transport or inhibition of hSGLT1 or hSGLT2 (Tables 2 and 3 (42); Lu et al., unpublished observations). Parenthetically, these results also suggest that [¹⁸F]fluoro-dapagliflozin (τ₁/₂ of the positron-emitter fluorine-18; τ₁/₂ = 110 min) may also be a useful molecular imaging probe to map and quantify the distribution of SGLT2 in both animal and human subjects using positron emission tomography (3).
What accounts for the inhibitor selectivity of hSGLT2 over hSGLT1-phlorizin = 10:1, dapagliflozin = 60:1? Our data offer a kinetic explanation: basically, dissociation rates for inhibitors were orders of magnitude slower for hSGLT2 than for hSGLT1, e.g., the $k_{\text{off}}$ for fluoro-dapagliflozin at 37°C was $0.005 \text{ s}^{-1}$ for hSGLT2 vs. $0.6 \text{ s}^{-1}$ for hSGLT1. The reciprocal of the off rate ($1/k_{\text{on}}$) is the mean occupancy time in the inhibitor-bound conformation. Thus dapagliflozin is bound to hSGLT2 for periods $>500 \text{ s}$ compared with 30 s for phlorizin. In contrast, dapagliflozin is bound to hSGLT1 for 2.5 s compared with 14 s for phlorizin. The tighter binding of dapagliflozin to hSGLT2 over hSGLT1 is consistent with the inability of phlorizin to exchange with dapagliflozin bound to hSGLT2, unlike the rapid exchange with hSGLT1 (see Fig. 5). The calculated $k_{\text{on}}$ were comparable for hSGLT1 and SGLT2, $1–100 \times 10^4 \text{ M/s}$ (Tables 2 and 3), except that for the galacto-dapagliflozin $k_{\text{on}}$ for hSGLT1 is only 1% of that for the other inhibitors.

What does dapagliflozin binding reveal about the inhibitor binding sites in hSGLTs? First, as discussed above, it is probable that the sugar moiety of these inhibitors binds in the glucose binding site. The fact that the inhibitor binding constants are $10^3$- to $10^6$-fold lower for glucose indicates that the structural and chemical properties of the aglycone are compatible with a specific binding site, probably in the entry vestibule to the glucose binding site. Alvarado (1) was among the first to postulate the synergistic effect of glucose and the aglycone in phlorizin binding. Second, the $K_i$ for phlorizin is 10-fold lower for hSGLT2 than hSGLT1, indicating a difference between hSGLT1 and hSGLT2 in this location and that the interaction between the chalone and the protein is stronger in hSGLT2. Third, changing the sugar moiety of dapagliflozin from glucose to galactose reduced affinity of both hSGLT1 and hSGLT2, but less for hSGLT2, suggesting that the position of the sugar in the binding site is not rigid, and so the position of the aglycone is shifted. In this case, the change affects binding less in hSGLT2 and may be an indication that the hSGLT2 binding site is more constrained than that of hSGLT1, perhaps explaining the low rate of galacto-dapagliflozin binding to hSGLT1 (Table 3).

At this time, it is difficult to pinpoint structural differences between the inhibitor sites in hSGLT1 and hSGLT2 based on the crystal structure of Vibrio parahaemolyticus SGLT1 (vSGLT1) (8), despite the general validity of hSGLT structural models (45). There is 32% amino acid identity (60% similarity) between vSGLT and hSGLT1, and all of the gating and coordinating residues are conserved between vSGLT1, hSGLT1, and hSGLT2. It is possible to dock the inhibitors to the occluded sugar binding site in the bacterial and human SGLTs, but, given the flexibility of the aglycones (Fig. 6), it is not yet possible to draw meaningful conclusions about the differences in inhibitor binding sites between hSGLT1 and hSGLT2 based on existing evidence. The successful determination of the crystal structures of inhibitors bound to the SGLTs would permit a more accurate interpretation of this differential binding.

---

**Fig. 5.** PZ exchange with DAPA bound to SGLTs. $p_i$/glucose (100 mM for hSGLT2 in A; 0.5 mM for hSGLT1 in B) was blocked by DAPA. The extracellular (bath) solution was subsequently replaced with PZ: 1 μM in hSGLT2 (A) and 160 nM in hSGLT1 (B). Next, solution was replaced with the original α-glucose buffer. A: for hSGLT2, there was no measurable dissociation time course, suggesting PZ did not exchange with DAPA binding to hSGLT2. B: the hSGLT1 time course of current recovery ($t_{1/2,\text{off}} \approx 9 \text{ s}$) was identical to that observed for PZ alone (Fig. 3A), suggesting that PZ can be exchanged for DAPA.

The sugar moiety of inhibitors appears to interact with the glucose binding site of hSGLT1 and hSGLT2, but selectivity of inhibitor binding is synergistic and depends on the chemistry of both the sugar moiety and the aglycone, indicating the dual-specific binding elements. This is clearly evident for the studies of galacto-dapagliflozin, where the results were unanticipated when considering selectivity of hexose transport alone: for hSGLT1, both glucose and galactose are transported with the same kinetics ($K_{0.5}$ and $V_{\text{max}}$), whereas galactose is a poor substrate for hSGLT2 transport ($K_{0.5} > 100 \text{ mM}$) (14). In contrast, the $K_i$ for galacto-dapagliflozin compared with dapagliflozin was 25 vs. 6 nM for hSGLT2 and 25,000 vs. 350 nM for hSGLT1.
Clinical Significance

In control human subjects, oral dapagliflozin inhibited up to 50% of the renal glucose reabsorption by the kidney (19, 22). The maximum glucose excretion, ≈60 g/24 h, occurred with 50-mg oral dapagliflozin, and, over this time, the plasma concentration of the drug rose to 4 μM at 1.5 h and decayed to 0.25 μM at 24 h. Ninety percent of dapagliflozin was found to be bound to serum proteins, and only 1% of the injected dose was excreted in the urine (see also Refs. 20, 31). Most of the oral dose appeared in plasma as an inactive glucuronidated metabolite, dapagliflozin-3-O-glucuronide, and this was excreted in the urine. These data, therefore, suggest that the free (unmodified and unbound) drug concentration in plasma and the glomerular filtrate, in the 24 h following a 250-mg dose, ranges from as high as 400 to as low as 25 nM. This is significantly higher than the dapagliflozin $K_i$ for hSGLT2 (5 nM), and so it would be expected that glucose excretion due to hSGLT2 inhibition would be close to the filtered glucose load, if hSGLT2 were responsible for 90% of glucose reabsorption.

What accounts for the fact that the selective hSGLT2 inhibitors only produce a 50% block of renal glucose reabsorption, whereas phlorizin produces complete blockage (5)? One possibility is that hSGLT1 accounts for a larger fraction of glucose reabsorption than previously recognized. Three recent studies in transgenic mice support this possibility: homozygous SGLT2 knockout (SGLT2−/−) mice retained up to 40% of renal α-glucose reabsorptive capacity (18, 27, 41). Given the above discussion of the pharmacokinetic data (for a 250-mg maximal dose), we estimate that the mean free dapagliflozin concentration in the glomerular filtrate is well below the hSGLT1 $K_i$ (100 nM).

Another important question is why only traces of dapagliflozin are found in the urine. Since the major metabolite, the 3-O-glucuronide, is excreted, it is likely that free dapagliflozin in plasma is also passed into the glomerular filtrate. If dapagliflozin inhibits by binding to the luminal SGLTs, once those binding sites are saturated, any additional dapagliflozin in the glomerular filtrate should be passed through the tubule and be excreted. Since this does not happen, and only a trace of dapagliflozin is found in urine, it suggests that there is a mechanism for dapagliflozin absorption somewhere in the renal tubule, possibly by one or both of the SGLTs, as transport of β-D-glucosides by SGLT1 is well documented (6, 26). Another explanation for dapagliflozin’s low in vivo potency and low urinary excretion is that it may only block hSGLT2 by gaining access to the apical membrane of tubular cells across the basolateral membrane from plasma. This possibility would depart from known phlorizin binding mechanisms to SGLT1; e.g., phlorizin does not inhibit intestinal absorption from the blood side (29) and does not inhibit SGLT1 from the cytosolic side of the plasma membrane (7, 34). These alternative possibilities are currently being investigated.

Aside from dapagliflozin, there currently is a paucity of published data on other hSGLT2 inhibitors. In only one study has the major functional differences between dapagliflozin and canagliflozin been reported: whereas the potency and specificity of canagliflozin in vitro are very similar to dapagliflozin, hSGLT2 $K_i = 2$ nM, hSGLT1 $K_i = 1,000$ nM (30), signifi-
cantly higher oral doses of canagliflozin than dapagliflozin are required to produce equivalent effects on renal glucose excretion (39). Kinetic specificity for hSGLT2 vs. SGLT1 alone cannot explain this discrepancy at this time, and so other pharmacokinetic factors must be in play.

In summary, the high affinity of dapagliflozin for hSGLT2 is the result of its tight binding, reflected by its surprisingly slow dissociation from the transporter, and this is the biological basis for the difference in dapagliflozin affinity between hSGLT1 and hSGLT2. Inhibitor affinity is the result of a synergistic relationship between binding sites for sugar and the aglycone, with alterations in the sugar resulting in surprising differences in selectivity. The sugar moiety has an important role in determining inhibitor specificity and is likely essential for positioning of the aglycone for interaction with residues in the sugar pathway, and the aglycone also influences the inhibitor interaction in ways we have yet to determine.

REFERENCES


Downloaded from http://ajpcell.physiology.org/ by 10.220.33.3 on October 20, 2017

AJP-Cell Physiol • doi:10.1152/ajpcell.00328.2011 • www.ajpcell.org


