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WZB117 (2-Fluoro-6-(m-hydroxybenzoyloxy) Phenyl m-Hydroxybenzoate) Inhibits GLUT1-mediated Sugar Transport by Binding Reversibly at the Exofacial Sugar Binding Site*

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WZB117 (2-fluoro-6-(m-hydroxybenzoyloxy) phenyl m-hydroxybenzoate) inhibits passive sugar transport in human erythrocytes and cancer cell lines and, by limiting glycolysis, inhibits tumor growth in mice. This study explores how WZB117 inhibits the erythrocyte sugar transporter glucose transport protein 1 (GLUT1) and examines the transporter isoform specificity of inhibition. WZB117 reversibly and competitively inhibits erythrocyte 3-O-methylglucose (3MG) uptake with \( K_{i(app)} = 6 \mu M \) but is a noncompetitive inhibitor of sugar exit. Cytochalasin B (CB) is a reversible, noncompetitive inhibitor of 3MG uptake with \( K_{i(app)} = 0.3 \mu M \) but is a competitive inhibitor of sugar exit indicating that WZB117 and CB bind at exofacial and endofacial sugar binding sites, respectively. WZB117 inhibition of GLUTs expressed in HEK293 cells follows the order of potency: insulin-regulated GLUT4 ≈ neuronal GLUT3 ≈ GLUT1. This may explain WZB117-induced murine lipodystrophy. Molecular docking suggests the following. 1) The WZB117 binding envelopes of exofacial GLUT1 and GLUT4 conformers differ significantly. 2) GLUT1 and GLUT4 exofacial conformers present multiple, adjacent glucose binding sites that overlap with WZB117 binding envelopes. 3) The GLUT1 exofacial conformer lacks a CB binding site. 4) The inward GLUT1 conformer presents overlapping endofacial WZB117, d-glucose, and CB binding envelopes. Interrogating the GLUT1 mechanism using WZB117 reveals that saturating WZB117 and CB stimulate erythrocyte 3MG uptake. Extracellular WZB117 does not affect CB binding to GLUT1, but intracellular WZB117 inhibits CB binding. These findings are incompatible with the alternating conformer carrier for glucose transport but are consistent with either a multisubunit, allosteric transporter, or a transporter in which each subunit presents multiple, interacting ligand binding sites.

Most cancer cells use anaerobic metabolism (glycolysis) to generate the ATP required for cellular processes and proliferation (1). This contrasts with normal, differentiated cells, which mostly use mitochondrial oxidative phosphorylation to sustain cellular function (2). The transition from aerobic to anaerobic metabolism is termed “the Warburg effect” and is driven by growth conditions, by mutations of proto-oncogenes and tumor-suppressor genes (2, 3). Vander Heiden et al. (1) have proposed that the metabolism of all proliferating cells (including cancer cells) is adapted to enhance nutrient uptake and nutrient incorporation into the biomass needed to produce a new cell. This would explain why the developing central nervous system is so susceptible to glucose deprivation resulting from glucose transport-deficiency at the blood-brain barrier (4) and why cancer cells are so sensitive to limited glucose availability (5).

This has prompted several groups to propose that suppressing anaerobic metabolism may offer an effective anti-cancer strategy. Three approaches have been used to limit glycolysis in cancer cells; that is glucose deprivation in cancer cells; that is glucose deprivation in vivo, in vitro, and in vivo use of glycolysis inhibitors (8), and cellular glucose transport inhibition in vitro and in vivo (8, 9). All three approaches cause cancer cell death.

Glucose transport in most mammalian cells is catalyzed by one or more members of the GLUT2 (SLC2A) family of passive glucose transporters (10). GLUT1 (the blood brain barrier/erythroid glucose transporter) and, in some instances, GLUT3 (the neuronal glucose transporter) or GLUT2 expression is significantly increased in proliferating cells (11).

Successful strategies for passive glucose transport inhibition have exploited a range of molecules that bind at or close to either the exofacial (e.g. maltose, ethyldiene glucose) or the endofacial sugar binding site (e.g. cytochalasin B; Refs. 12–14). A number of unrelated inhibitory molecules react with the GLUT1 purine binding site (e.g. ATP, AMP, caffeine; Ref. 15) or at other, less well defined sites (e.g. androgens, quercetin; Refs.

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16 and 17). These and related molecules have been suggested as potential scaffolds for designing inhibitors of glucose uptake in cancer cells or cancer stem cells (8, 18–20).

Using such an approach, Zhang et al. (21) identified a class of polyphenolic compounds with sugar transport inhibitory potency. Among these, WZB117 (2-fluoro-6-(m-hydroxybenzoxy)phenyl m-hydroxybenzoate) showed the highest affinity for sugar transport inhibition and inhibited tumor growth in a nude mouse model (9).

Although a promising strategy for cancer cell elimination, glucose transport inhibition in vivo may present several attendant complications. These include impaired insulin secretion, elevated blood glucose, diuresis, elevated glycation, impaired glucose transport across the blood-brain barrier, and reduced metabolic capacity in healthy cells that depends upon glycolysis for normal function. We, therefore, undertook a systematic analysis of WZB117 inhibition of glucose transport.

We found that WZB117 interacts with GLUT1 at the exofacial sugar binding site and thus acts as a reversible, competitive inhibitor of net glucose uptake and exchange glucose transport but as a noncompetitive inhibitor of sugar efflux from cells. WZB117 inhibits the insulin-sensitive glucose transport (GLUT4) with greater potency than its inhibition of either GLUT1 or GLUT3.

Results

Sugar transport in human erythrocytes is catalyzed by GLUT1 (22). Transport theory states that ligands binding reversibly at the exofacial sugar binding site act as competitive inhibitors of sugar uptake and as noncompetitive inhibitors of exit. Conversely, ligands binding reversibly at the endofacial sugar binding site act as noncompetitive inhibitors of sugar uptake and as competitive inhibitors of exit (23–25).

To understand whether WZB117 inhibits GLUT1 by binding at the exofacial or endofacial sugar binding site, we examined its effects on three modes of erythrocyte sugar transport: zero-trans 3MG uptake (uptake into cells lacking sugar), zero-trans 3MG exit (efflux from cells into medium lacking sugar), and equilibrium exchange 3MG uptake (unidirectional uptake of 3MG in cells where \[ [3\text{MG}]_i = [3\text{MG}]_o \].

WZB117 inhibits zero-trans 3MG (0.1 mM) uptake by erythrocytes in a dose-dependent manner with \( K_{(\text{app})} = 6.2 \pm 1.6 \mu M \) (Fig. 1). CB (a known GLUT1 endofacial site ligand) inhibits 0.1 mM 3MG zero-trans uptake with \( K_{(\text{app})} = 0.3 \pm 0.1 \mu M \) (see Fig. 4).

We assessed the reversibility of inhibition in two ways. The first method involved exposing RBCs to buffer containing or lacking WZB117 (3 \( \mu M \)) or CB (0.5 \( \mu M \)) for 10 min, then removing the inhibitors using a centrifugation/wash cycle and incubating the cells in inhibitor-free buffer for a further 15 min. We then measured 0.1 mM 3MG uptake in these cells. WZB117 (3 \( \mu M \)) or CB (0.5 \( \mu M \)) inhibit sugar uptake by 50% (Fig. 2). Removing the inhibitor before sugar uptake determinations restored sugar uptake to levels of 75% or greater of uninhibited controls (Fig. 2).

WZB117 Is a Competitive Inhibitor of 3MG Uptake by Human Erythrocytes—The second method of assessing reversibility of inhibition evaluates the type of sugar transport inhibition produced by WZB117. Competitive inhibitors act by binding reversibly either to the active site of an enzyme or to a site whose occupancy occludes occupancy of the active site and vice versa (26).

Preincubation with WZB117 (7 \( \mu M \)) before initiation of transport measurements increased \( K_{(\text{app})} \) for zero-trans 3MG uptake from 1.0 \( \pm \) 0.2 mM to 2.8 \( \pm \) 0.4 mM, but \( V_{\max} \) for 3MG uptake (0.39 \( \pm \) 0.03 mmol/liter of cell water/min) was unaffected (Fig. 3). This result indicated that WZB117 is a competitive inhibitor of human erythrocyte sugar uptake. If WZB117 bound irreversibly to the active site, preincubation with the
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FIGURE 3. Effects of WZB117 and CB on Michaelis-Menten kinetics of zero-trans 3MG uptake (A), zero-trans 3MG exit (B and C), and equilibrium exchange 3MG uptake (D). Results are shown for control cells (○), WZB117-treated cells (●), and for CB-treated cells (△). Each data point is the mean ± S.E. of three duplicate measurements. A, ordinate: 3MG uptake in mmol/liter of cell water/min. Abscissa: [3MG] in mM. Curves drawn through the points were computed by nonlinear regression assuming Michaelis-Menten uptake kinetics (Equation 2) and have the following results: control cells (○), $V_{max} = 0.59 ± 0.03$ mmol/liter of cell water/min, $K_{m(app)} = 1.05 ± 0.16$ mM, $R^2 = 0.992$, standard error of regression = 0.018 mmol/liter of cell water/min; WZB117 treatment (●): $V_{max} = 0.68 ± 0.04$ mmol/liter of cell water/min, $K_{m(app)} = 2.77 ± 0.35$ mM, $R^2 = 0.997$, standard error of regression = 0.018 mmol/liter of cell water/min; CB treatment (△): $V_{max} = 0.29 ± 0.05$ mmol/liter of cell water/min, $K_{m(app)} = 1.67 ± 0.63$ mM, $R^2 = 0.966$, standard error of regression = 0.010 mmol/liter of cell water/min. B and C, ordinate: [3MG] in mmol/liter of cell water. Abscissa: time in minutes. Curves drawn through the points were computed by nonlinear regression and numerical integration assuming Michaelis-Menten exit kinetics and have the following results: Control cells (○): $V_{max} = 2$ mmol/liter of cell water/min, $K_{m(app)} = 15$ mM; WZB117 treatment (●): $V_{max} = 0.8$ mmol/liter of cell water/min, $K_{m(app)} = 15$ mM; CB treatment (△): $V_{max} = 2$ mmol/liter of cell water/min, $K_{m(app)} = 47$ mM. D, Hanes-Woolf plot of equilibrium exchange transport. Ordinate: [3MG]/rate of equilibrium exchange 3MG uptake in minutes. Abscissa: intracellular and extracellular [3MG] in mM. Lines drawn through the points were computed by nonlinear regression assuming that each line is described by Equation 3. The results are: control cells (○): $V_{max} = 13.39 ± 0.79$ mmol/liter of cell water/min, $K_{m(app)} = 40.25 ± 3.57$ mM, $R^2 = 0.990$, standard error of regression = 0.17 min; WZB117 treatment (●): $V_{max} = 13.36 ± 2.02$ mmol/liter of cell water/min, $K_{m(app)} = 82.25 ± 15.30$ mM, $R^2 = 0.956$, standard error of regression = 0.43 min; CB treatment (△): $V_{max} = 8.87 ± 0.33$ mmol/liter of cell water/min, $K_{m(app)} = 49.85 ± 2.55$ mM, $R^2 = 0.996$, standard error of regression = 0.16 min.

inhibitor would most likely have reduced $V_{max}$ for transport in addition to increasing $K_{m(app)}$.

Conversely, CB (0.7 μM) reduces $V_{max}$ for 3MG uptake (0.29 ± 0.05 mmol/liter of cell water/min) but has no effect on $K_{m(app)}$ (Fig. 3A). This result confirms that CB is a noncompetitive inhibitor of uptake and is expected for an inhibitor which binds reversibly to the endofacial active site (25).

WZB117 Is a Noncompetitive Inhibitor of Net Sugar Exit—Because WZB117 competes for 3MG binding at or close to the exofacial sugar entry site, we hypothesized that WZB117 would act as a noncompetitive inhibitor of sugar exit (25). To this end, we performed sugar exit experiments where we measured the time course of zero-trans sugar exit from pre-loaded red cells into sugar-free external medium. The time course of exit was analyzed by simulating sugar exit by numerical integration using Berkeley Madonna software assuming Michaelis-Menten exit kinetics then adjusting simulated Michaelis-Menten parameters using the Levenberg-Marquardt algorithm until the fits between simulated and experimental exit data were optimal. Our analysis shows that WZB117 (0.7 μM) reduces $V_{max}$ for 3MG exit from 2 mmol/liter of cell water/min to 0.8 mmol/liter of cell water/min, whereas $K_{m(app)}$ for exit (15 mM) was unaffected (Fig. 3B). Consistent with published data (13), CB (0.7 μM) was competitive for sugar exit, increasing $K_{m(app)}$ to 47 mM with no effect on $V_{max}$ (Fig. 3C). WZB117 Is a Competitive Inhibitor of Equilibrium Exchange Sugar Transport—Equilibrium exchange 3MG transport in human red blood cells ([3MG]$_i$ = [3MG]$_o$) was competitively
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inhibited by WZB117. WZB117 (7 μM) increased \( K_{m(app)} \) for 3MG exchange transport from 40.25 ± 3.57 mM to 82.25 ± 15.30 mM but had no effect on \( V_{max} \) (13.39 ± 0.79 mmol/liter of cell water/min; Fig. 3D). CB (0.7 μM), however, was a mixed-type inhibitor of equilibrium exchange 3MG transport reducing \( V_{max} \) (8.87 ± 0.33 mmol/liter of cell water/min) and increasing \( K_{m(app)} \) (49.85 ± 2.56 mM; Fig. 3D).

Subsaturating Levels of CB and WZB117 Stimulate Sugar Uptake in Human Erythrocytes—Previous studies have shown that subsaturating concentrations of CB or exofacial inhibitors of GLUT1 produce a modest stimulation of erythrocyte sugar uptake (27, 28). Subsaturating levels of CB (0.1 μM) and WZB117 (0.1 μM) stimulate zero-trans 3MG uptake in human erythrocytes by up to 25% above untreated cells (Fig. 4). This effect is significant (\( p = 0.0075 \)) at 10 and 50 nm CB and at 100 nm WZB117 (\( p < 0.001 \)). CB and WZB117 may stimulate transport by reducing \( K_{m(app)} \) for sugar uptake (Table 1), but the data are not conclusive.

**TABLE 1**

<table>
<thead>
<tr>
<th>Experimental condition</th>
<th>( V_{max} ) (mmol/liter cell water/min)</th>
<th>95% Confidence interval</th>
<th>( K_{m(app)} ) (μM)</th>
<th>95% Confidence interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.63 ± 0.01</td>
<td>0.59–0.68</td>
<td>2.26 ± 0.12</td>
<td>1.92–2.68</td>
</tr>
<tr>
<td>WZB117 0.7 μM</td>
<td>0.55 ± 0.04</td>
<td>0.46–0.69</td>
<td>1.74 ± 0.26</td>
<td>1.08–2.91</td>
</tr>
<tr>
<td>CB 0.025 μM</td>
<td>0.60 ± 0.02</td>
<td>0.55–0.67</td>
<td>1.73 ± 0.13</td>
<td>1.36–2.23</td>
</tr>
</tbody>
</table>

**Effect of WZB117 on CB Binding**—CB is membrane-permeant and rapidly crosses the cell membrane to interact with the GLUT1 endofacial sugar binding site (23, 25). The simple carrier hypothesis for facilitated diffusion predicts that CB and WZB117 binding to GLUT1 are mutually exclusive (23–25). Accordingly, CB should competitively inhibit WZB117 inhibition of and binding to GLUT1. The IC\(_{50}\) for CB inhibition of 3MG uptake (0.35 μM) is not materially affected when WZB117 (5 μM) is present (Fig. 4).

WZB117 is without effect on \(^{[3]H}\)CB binding to human RBCs but served as a low affinity inhibitor of CB binding to red cell membranes depleted of peripheral membrane proteins (Fig. 5; \( K_{i(app)} = 156 ± 24 \mu M \)). RBC membranes depleted of peripheral proteins are unsealed, and both membrane surfaces are accessible to buffer (29). Collectively, these results indicate: 1) CB binding to the GLUT1 endofacial sugar binding site is insensitive to WZB117 binding at the exofacial sugar binding site; 2) CB and WZB117 compete for binding at the GLUT1 endofacial sugar binding site; 3) RBCs are impermeable to WZB117 because inhibition of binding by a membrane-permeant WZB117 would have been evident in measurements with intact RBCs. In addition, these findings are consistent with the actions of D-glucose on CB binding to RBCs; extracellular D-glucose is without effect on binding, whereas intracellular D-glucose competitively inhibits binding (30).

**Isoform Specificity of Transport Inhibition**—The GLUT (SLC2A) family of passive glucose transporters can show distinct, isoform-specific affinities for inhibitors. For example,
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The insulin-sensitive glucose transporter of fat and muscle (GLUT4) is inhibited by HIV protease inhibitors, but GLUT1 is not (31), whereas GLUT1, GLUT3, and GLUT4 are inhibited by CB and the intestinal fructose transporters GLUT5 and GLUT7 are not (32). We, therefore, stably transfected HEK293 cells with GLUT1, GLUT3, or GLUT4 and measured the concentration dependence of WZB117 inhibition of 2-deoxy-D-glucose (2DG, 0.1 mM) uptake at 37 °C. WZB117 inhibits GLUT1- and GLUT3-mediated 2DG uptake with $K_{i(app)} = 10 \mu M$ (Fig. 6A) but is a more potent inhibitor of GLUT4-mediated uptake ($K_{i(app)} = 0.2 \mu M$; Fig. 6A).

Heterologous expression of human GLUT1 or GLUT4 (Fig. 6E) significantly reduced endogenous GLUT3 mRNA (Fig. 6B) and protein (Fig. 6, C and D) levels in HEK293 cells as judged by qPCR and Western blotting analyses. This explains why high affinity, WZB117 inhibition of transport in GLUT4-transfected cells reduced transport more effectively than observed in untransfected cells (Fig. 6A). We hypothesize that endogenous GLUT3 expression is reduced in GLUT4-transfected cells as a compensatory response to increased cellular uptake.

**Molecular Docking Studies**—We undertook molecular docking studies of $\alpha$-glucose, WZB117, and CB binding to a homology model of the exofacial (e2) open conformation of GLUT1 (hm-GLUT1-e2). We used the ligand-depleted chain A of the maltose-bound human GLUT3 (PDB code 4ZWC) structure (33) as the template to generate the homology model of the GLUT1-e2 (open) state. GLUT1 and GLUT3 share 65% sequence identity and 88% sequence similarity (10).

$\beta$-d-Glucose docks at three high affinity sites in hm-GLUT1-e2 (Fig. 7A; Fig. 8, a–f). We term these sites peripheral, intermediate, and core based upon their location within the exofacial, interstitium-exposed cavity of the GLUT1-e2 conformer. Eighteen $\beta$-d-Glc binding configurations were obtained at the core site. The affinity of binding ($\Delta G$) ranged from $-6$ to $-4$ kcal/mol. Of these 18 possibilities, only 4 match the known stereospecificity of sugar inhibition of GLUT1-mediated L-sorbose entry in human erythrocytes (34). We illustrate one such complex in Fig. 8, e and f. $\beta$-d-Glc binding at this site is coordinated by hydrogen bonds to Gln161, Gln282, Asn317, and Asn318, and Glu380. Two additional $\beta$-d-Glc binding sites were detected in the exofacial cavity. $\beta$-d-Glc binding at the peripheral site (Fig. 8, a and b) and at the intermediate site (Fig. 8, c and d) presents fewer configurations, and in both instances C1 and its hydroxyl are solvent-exposed, whereas C6 and its hydroxyl are not. $\beta$-d-Glc binding at the intermediate site is coordinated by hydrogen bonds to Thr35, Asn34, Val182, Ser173, and Asn131. This orientation of sugar is incompatible with the stereochemistry of L-sorbose uptake inhibition described by Barnett et al. (34).

WZB117 docking studies indicate that WZB117 can assume three positions in the exofacial cavity of hm-GLUT1-e2. When binding at its highest affinity site, WZB117 spanned the core.
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and intermediate β-D-Glc binding sites (Fig. 7b and Fig. 8, g and h). Phenol ring 1 occupied the intermediate glucose binding site and was coordinated by hydrogen bonds to GLUT1 residues Val169 and Asn415. Phenol ring 2 occupied the core glucose binding site and was coordinated by hydrogen bonds to GLUT1 residues Asn317 and Glu380 (Fig. 8, g and h). The two lower affinity WZB117 binding sites comprise one in which WZB117 docks to and spans the peripheral and core D-Glc binding sites and a second where WZB117 interacted with the peripheral and intermediate D-Glc binding sites (not shown). CB did not dock at any high affinity sites in hm-GLUT1-e2 but did dock to the endofacial, GLUT1-e1 conformer (GLUT1-e1; Fig. 7d). The e1 GLUT1 conformation presented overlapping endofacial WZB117 and D-glucose sites that partially overlapped with the benzene ring of CB (Fig. 7, c and d).

Discussion

WZB117, a prototypic anticancer drug, inhibited human erythrocyte sugar transport by acting as a competitive inhibitor...
of sugar uptake and as a noncompetitive inhibitor of sugar exit. This suggests that WZB117 and extracellular glucose compete for binding to the same site on the erythrocyte sugar transporter. Molecular docking and ligand binding studies support this hypothesis by showing that extracellular D-glucose and WZB117 binding sites comprise overlapping amino side chains distinct from those involved in binding CB, an inhibitor that acts at the endofacial sugar binding site.

### Sidedness of Inhibition

Several studies have demonstrated (regardless of the assumed carrier model) that reversible transport inhibitors binding at the exofacial sugar binding site of the passive glucose transporters serve as competitive inhibitors of net sugar uptake and equilibrium exchange transport and as noncompetitive inhibitors of net sugar exit (23–25). Conversely, reversible transport inhibitors binding at the endofacial sugar binding site are competitive inhibitors of net sugar exit and equilibrium exchange transport and are noncompetitive inhibitors of net sugar uptake (23–25). WZB117, therefore, fits the profile of a reversible inhibitor binding at the exofacial sugar binding site. CB behaves as a reversible inhibitor that binds at the endofacial sugar binding site, although the mixed-type inhibition of equilibrium exchange sugar transport and our molecular docking studies suggest that CB binding may involve more than simple interaction with the endofacial sugar binding site.

### Determinants of Affinity and Isoform Specificity

Although several permutations of D-glucose and WZB117 binding sites were identified in GLUT1 docking studies, the highest affinity binding sites are consistent with the transport and ligand binding studies presented here and with previous studies of ligand binding to GLUT1 (12, 25, 30, 33, 35–37). Three exofacial D-glucose docking sites are present in the interstitium exposed exofacial cavity of hm-GLUT1-e2, peripheral, intermediate, and core. The stereochemistry of sugar binding at the peripheral and intermediate sites is incompatible with the findings of previous studies describing the stereochemistry of sugar transport inhibition by D-glucose analogs (34), whereas several configurations of sugar binding at the core site are compatible with the earlier studies. We, therefore, propose that the core D-glucose binding site may be the site at which hexose binding subsequently gives rise to sugar translocation.

WZB117 occupies three envelopes in GLUT1-e2. Envelope 1 comprises peripheral and core Glc binding sites, envelope 2 comprises intermediate and core Glc binding sites, and envelope 3 comprises peripheral and intermediate Glc sites. A majority (93%) of computed WZB117/GLUT1 complexes correspond to WZB117 binding within envelope 2. The remaining 7% are equally distributed between envelopes 1 and 3. WZB117 geometry is fully extended (linear) in envelope 1 but is “L” shaped in envelopes 2 and 3. Although each envelope would leave one available Glc binding site available for extracellular hexose, it is easy to see how Glc and WZB117 would compete for binding to GLUT1.

CB does not dock to hm-GLUT1-e2 but can be docked with high affinity to GLUT1-e1. D-Glucose and WZB117 also dock to GLUT1-e1 but bind at overlapping (and presumably mutually exclusive) sites deeper within the e1 cavity. These sites do overlap with the CB binding site, and the observed competitions between sugar, intracellular WZB117, and CB for binding to GLUT1 suggest that mutually exclusive binding could result from steric hindrance.

GLUT1 and GLUT3 are inhibited by WZB117 with $K_{i(app)}$ of $\sim 5–10 \mu M$. GLUT4, the insulin-sensitive glucose transporter of fat and muscle cells (38–40), is inhibited by WZB117 with a $30–100$-fold greater potency.

The GLUT4-e2 conformer also presents peripheral, intermediate, and core Glc binding sites. However, the presence of Ile$^{42}$ in GLUT4 transmembrane helix 1 (Thr$^{30}$ in GLUT1 H1) leaves only linear envelope 1 (peripheral and core Glc sites) accessible to WZB117 (Fig. 7f). Molecular docking, therefore, suggests that WZB117 preferentially occupies envelope 2 in GLUT1-e2 and envelope 1 in GLUT4-e2. $K_{i(app)}$ for sugar transport and $K_{i(app)}$ for transport inhibition are products of multiple steps in the catalytic cycle regardless of the presumed mechanism of transport (25, 41). It is not surprising, therefore, that WZB117 occupancy of different binding envelopes in two transporters characterized by very different transport kinetics (42) would result in significantly different $K_{i(app)}$ for sugar transport inhibition.

### Implications for Mechanism of Passive Sugar Transport

Four observations in the present study impact our understanding of glucose transport mechanism as follows. 1) As is the case for extracellular D-glucose (30), extracellular WZB117 does not affect CB binding to GLUT1. Hence, the transporter’s exofacial sugar/WZB117 binding site and the endofacial CB binding site are not mutually exclusive. If CB binds only to the GLUT1-e1 conformer, which presents an endofacial sugar binding site, this behavior refutes the simple (alternating conformer) carrier model for sugar transport which predicts that inhibitor binding to the exofacial sugar binding site should competitively inhibit CB binding (43–45). 2) Extracellular WZB117 does not increase $K_{i(app)}$ for CB noncompetitive inhibition of sugar uptake. Again, if CB binds only to the endofacial e1 conformer of GLUT1, this observation refutes the simple (alternating conformer) carrier model for sugar transport (43–45). 3) Like extracellular maltose and intracellular CB (27, 28), subsaturating extracellular WZB117 modestly stimulates sugar uptake. Transport stimulation by exofacial and endofacial transport inhibitors has been ascribed to cooperative interactions between co-exis- tant exofacial and endofacial sugar binding sites present in the tetrameric GLUT1 transporter complex (27, 28). The transporter is proposed to comprise four subunits (GLUT1 proteins) in which each subunit behaves as an alternating conformer carrier but where intersubunit cooperativity results in two subunits presenting exofacial (e2) orientations and 2 subunits presenting endofacial (e1) conformations at any instant (46, 47). At subsaturating WZB117, there is a greater probability that only one of the two available GLUT1-e2 conformers is occupied by WZB117, and its occupancy is proposed to allosterically activate sugar uptake via the remaining, inhibitor-free e2 subunit (28). As WZB117 is further increased, the probability that inhibitor occupies both e2 subunits increases and leads to sugar uptake inhibition. An alternative hypothesis is that exofacial ligand-induced allostery is an intrasubunit phenomenon in which occupancy of the exofacial cavity by poly- phenolic WZB117 increases GLUT1 affinity for, or transport of, the transported sugar through the same GLUT1 subunit (48). 4)
Molecular docking studies reveal three exofacial D-glucose binding sites in the hm-GLUT1-e2 cavity, peripheral, intermediate, and core. This is consistent with previous observations that GLUT1 binds ≈2 mol of transported sugar per mol GLUT1 (49). Each envelope would leave one Glc binding site available for substrate. The results of this docking analysis, therefore, do not permit resolution of the mechanism of exofacial allostery (i.e. inter- or intrasubunit interactions).

Implications for Use as an Anticancer Agent—Although a promising strategy for cancer cell elimination, glucose transport inhibition using WZB117 could perturb organismal carbohydrate homeostasis in several important ways. Blood-brain and blood-tissue barrier GLUT1-dependent glucose transport will be depressed, perturbing peripheral and central neuro-metabolic function (50). Human, GLUT1-dependent, pancreatic beta cell insulin secretion in response to hyperglycemia will be diminished (51), and insulin-dependent, GLUT4-mediated skeletal muscle and adipose glucose transport will be prone to high affinity WZB117 inhibition thereby further compounding hyperglycemia. The net effects will include acute hyperglycemia and perturbations of neuro-metabolic function and insulin-dependent lipogenesis. These predictions are consistent with the reported hyperglycemia and lipodystrophy in mice after WZB117 treatment (9) and suggest that WZB117-treatment could induce metabolic syndrome/insulin resistance in individuals undergoing therapy.

The current study demonstrates that WZB117 inhibits facilitative sugar transport by competing with sugars for occupancy of the exofacial substrate binding site of the transporter. GLUT1, GLUT3, and GLUT4 are sensitive to transport inhibition, suggesting that sugar transport across the blood brain barrier in CNS astrocytes and neurons and in insulin-sensitive tissues is susceptible to inhibition (52). Most cancer cells rely on anaerobic glycolysis to generate the ATP and substrates required for cellular processes and proliferation (1). This contrasts with normal, differentiated cells, which mostly use mitochondrial oxidative phosphorylation to sustain cellular function (2). This could explain why sugar transport inhibition is a successful anti-cancer strategy in mice (9). However, human CNS development and function are especially dependent on glucose transport and metabolism (50), and the consequences of inhibition of glucose transport into and within the CNS could be catastrophic. We suggest, therefore, that a combination therapy of ketogenic diet (such as is used to manage GLUT1 deficiency; Refs. 4 and 53–55) plus sugar transport inhibition might be more successful.

Experimental Procedures

Reagents—([3H]Cytochalasin B and 3-O-[3H]methylglucose ([3H]3MG) were purchased from American Radiolabeled Chemicals (St. Louis, MO). Unlabeled 3MG, CB, and phloretin were purchased from Sigma. WZB117 was purchased from Molecular Targeting Technologies (St. Louis, MO). Unlabeled 3MG, CB, and phloretin (100 μM) sample buffer contained 0.125 M Tris-HCl, pH 6.8, 4% SDS, 20% glycerol, and 50 mM DTT. Transfer buffer was composed of 12 mM Tris base, 96 mM glycine, 20% methanol.

Cells—De-identified whole human blood was purchased from Biological Specialty Corporation (Colmar, PA). HEK293 cells were maintained in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum, 100 units/ml penicillin, and 100 μg/ml streptomycin in a 37 °C humidified 5% CO2 incubator. Resealed, human red cell ghosts were made as described in Sage and Carruthers (56), and erythrocyte membranes were depleted of peripheral membrane proteins and resuspended in KCl medium at 4 mg protein/ml as described in Carruthers (57).

Heterologous Expression of GLUTs—GLUT1, GLUT3, and GLUT4 heterologous expression in HEK293 cells was as described previously (42, 58). Both GLUT1 and GLUT3 contain a myc-epitope in exofacial loop 1 (42). GLUT1, GLUT3, GLUT4, and NaKATPase expression in whole cell lysates was analyzed by Western blotting as previously described (42, 58).

Erythrocyte Sugar Transport Measurements—All human red cell sugar transport experiments were performed at 4 °C as previously described (15, 59, 60).

Zero-trans Uptake—Five volumes (100 μl) of 3MG uptake medium ± WZB117 or CB were added to 1 volume (20 μl) of sugar-depleted, 50% hematocrit (Ht) red cells, and sugar uptake allowed to proceed for 0.5–1 min at 4 °C. Uptake was stopped by adding 50 volumes (1 ml) of ice-cold stop solution containing 100 mM phloretin and 10 μM CB. Cells were washed in stop solution and lysed in 3% perchloric acid, and centrifugation-clarified lysates were assayed for radioactivity in duplicates using liquid scintillation counting.

Zero-trans Exit—Sugar-depleted packed RBCs were loaded with 10 mM 3MG by incubating with 20 mM 3MG (containing 10 μCi [3H]3MG/10 ml of cold 3MG for 1 h at 37 °C). One volume (0.5 ml) of sugar-loaded RBCs was added to 50 volumes of KCl medium ± CB or WZB117 on a shaker with magnetic stirrer. Aliquots (0.5 ml) of the suspension were withdrawn at the indicated time intervals, washed in ice-cold stop solution, lysed, and assayed for remaining radioactivity.

Equilibrium Exchange Transport—Sugar-depleted RBCs were preloaded with 2.5–50 mM 3MG at 25 °C for 30 min. These cells were collected by centrifugation, incubated in 10 volumes of 3MG ± WZB117 or CB for 10 min at 4 °C, centrifuged again, and resuspended to 50% Ht in the appropriate 3MG and inhibitor medium. 3MG uptake medium (100 μl) ± WZB117 or CB was added to 20 μl of sugar-loaded cells (50% Ht), and the experiment was allowed to proceed for 0.5–1 min at 4 °C. Uptake was arrested by adding 50 volumes (1 ml) of ice-cold stop solution. Cells were centrifuged, and the pellet was washed in 1 ml of stop solution, collected by centrifugation, lysed in 3% perchloric acid, clarified, and assayed for radioactivity in duplicates using liquid scintillation counting.

HEK293 Cell Sugar Transport Measurements—All HEK293 cell sugar transport experiments were performed at 37 °C using 0.1 mM 2-deoxy-D-glucose as previously described (42, 58).
**WZB117 Binds at the GLUT1 Exofacial Sugar Binding Site**

**CB Binding**—Equilibrium CB binding to red cells and to red cell membranes depleted of peripheral membrane proteins was performed as previously described (30). Briefly, 1 volume of sugar-depleted red cells (50% Ht) or red cell membranes (4 mg of membrane protein/ml) was suspended in one volume of ice-cold wash buffer 150 mM KCl, 5 mM HEPES, pH 7.4, containing 0.5 mM [3H]CB, 10 μM cytochalasin D, and varying concentrations of WZB117 (1–60 μM) for 15 min at 4 °C with constant end-over-end rotation. For total CB, 2 × 10 μl of the cell suspension were lysed in 100 μl of 3% perchloric acid and clarified by centrifugation, and the associated radioactivity was assayed by liquid scintillation counting. For free CB, the cell suspension was centrifuged at 10,000 × g for 30 s, and 10 μl of clarified supernatant were assayed in duplicate by scintillation counting. Bound CB was obtained as: total CB - free CB.

**Quantitative RT-PCR**—Expression levels of hGLUT3 mRNA were measured by qPCR using the QuantiTect SYBR Green PCR kit (Qiagen). Reactions were performed according to the manufacturer’s protocol. Total RNA was isolated from HEK293 cells stably overexpressing hGLUT1 or hGLUT4 using the RNasey kit and QiShredder (Qiagen). cDNA was synthesized using the QuantiTect Reverse Transcriptase kit (Qiagen). qPCR reactions contained 100 ng of the indicated cDNA and 0.3 μM each human GLUT3-specific qPCR primers (G3, 5′-qF-AGCTCTCTGGGATCATGTCGTGTG and G3 3′qR-ATGGTGCCATAGATGGCTCTTGA, where qF denotes qPCR forward and qR denotes qPCR reverse). Samples were run in triplicate on a Research PTC-200 Peltier Thermal Cycler with a Chromo4 real time PCR detector running Opticon system. No restraints were used during the docking.

**Data Analysis**—Linear and nonlinear regression analysis of data sets and statistical tests were accomplished using GraphPad Prism (Version 7.0a; La Jolla, CA).

Michelis-Menten inhibition of sugar transport is assumed to be described by

\[
v = v_0 - \frac{v_o[I]}{K_{(i)app} + [I]} \quad \text{(Eq. 1)}
\]

uptake, where \(v_o\) is \(v\) measured in the absence of inhibitor \(I\), \([I]\) is the concentration of inhibitor and \(K_{(i)app}\) is that \([I]\) producing 50% inhibition of uptake.

Michelis-Menten transport is assumed to be described by

\[
v = \frac{V_{max}\[3MG]}{K_{m(app)} + [3MG]} \quad \text{(Eq. 2)}
\]

uptake, where \(V_{max}\) is the maximum rate of 3MG transport, \([3MG]\) is the concentration of 3MG, and \(K_{m(app)}\) is that \([3MG]\) where the rate of uptake is \(V_{max}/2\).

Hanes-Woolf analysis of transport assumes Michelis-Menten kinetics, and when data are plotted as \([3MG]/v\) versus \([3MG]\), the results fall on a straight line described by

\[
\frac{[3MG]}{v} = \frac{[3MG]}{V_{max}} + \frac{K_{m(app)}}{V_{max}} \quad \text{(Eq. 3)}
\]

where \(V_{max}\) is the maximum rate of 3MG transport, \([3MG]\) is the concentration of 3MG, and \(K_{m(app)}\) is \([3MG]\), where the rate of uptake is \(V_{max}/2\).

Transport stimulation followed by inhibition by inhibitors was approximated first by normalizing all uptake to \(v_o\) and then using the following model.

\[
v = 1 + \frac{[I] \Delta_1}{K_1 + [I]} - 1 \frac{[I] \Delta_1}{K_2 + [I]} \quad \text{(Eq. 4)}
\]

**Homology Modeling**—The homology models of the e2 (open) state of GLUT1 and GLUT4 were generated using the malsequence-bound human GLUT3 (PDB code 4ZWC) structure (33). GLUT1 and GLUT3 have 65% sequence identity and 88% sequence similarity (10). Ligands were removed, and chain A was used as the template for each modeled sequence. Sequence alignments were generated using ClustalX. Homology models were built using Modeler-9.9 and analyzed using PROCHECK. The GLUT1-e1 structure (PDB code 4PYP; Ref. 61) was used directly.

**Stochastic Docking**—The GLUT1 crystal structure was obtained from the protein data bank using PDB code 4PYP. The structures for β-D-glucose and CB were obtained from ZINC. The WZB117 structure was generated using the three-dimensional structure generator Corina from Molecular Networks GmbH. Docking was performed using the Schrodinger software suite. The protein structure was preprocessed with the Protein Preparation Wizard, bond orders were assigned, hydrogen were added, and the H-bond network was optimized. The system was energy-minimized using the OPLS 2005 force field. Ligand structures were prepared with the LigPrep module, and the pKaPr of the ligands was calculated using the Epik module. Computational docking was performed with the GLIDE module in standard precision (SP) mode and default values for grid generation. Cavities for docking were calculated using the CastP server, and the grid was centered on the residues forming the cavity. No restraints were used during the docking.

**Author Contributions**—O. O. conducted most of the experiments, analyzed the results, and wrote most of the paper. K. L. undertook the molecular docking studies. A. S. assisted with the quantitative analysis of data. J. D. conducted the experiments on the isoform specificity of WZB117 inhibition of transport. A. C. conceived the idea for the project and co-wrote the paper with O. O.

**References**

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WZB117 Binds at the GLUT1 Exofacial Sugar Binding Site


WZB117 (2-Fluoro-6-(m-hydroxybenzoyloxy) Phenyl m-Hydroxybenzoate) Inhibits GLUT1-mediated Sugar Transport by Binding Reversibly at the Exofacial Sugar Binding Site

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