Forces and Dynamics of Glucose and Inhibitor Binding to Sodium Glucose Co-transporter SGLT1 Studied by Single Molecule Force Spectroscopy*

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Background: SGLT1 functions in intestinal glucose absorption and renal reabsorption.
Results: With increasing temperature, width of energy barrier and average life time increased for glucose binding to SGLT1 but decreased for phlorizin binding.
Conclusion: Sugar translocation and inhibitor binding involves several steps with different temperature sensitivity.
Significance: Force spectroscopy can be used to study dynamics and structure of membrane transporter.

Single molecule force spectroscopy was employed to investigate the dynamics of the sodium glucose co-transporter (SGLT1) upon substrate and inhibitor binding on the single molecule level. CHO cells stably expressing rbSGLT1 were probed by using atomic force microscopy tips carrying either thioglucose, 2'-aminoethyl β-D-glucopyranoside, or aminophlorizin. Poly(ethylene glycol) (PEG) chains of different length and varying end groups were used as tether. Experiments were performed at 10, 25 and 37 °C to address different conformational states of SGLT1. Unbinding forces between ligands and SGLT1 were recorded at different loading rates by changing the retraction velocity, yielding binding probability, width of energy barrier of the binding pocket, and the kinetic off rate constant of the binding reaction. With increasing temperature, width of energy barrier and average life time increased for the interaction of SGLT1 with thioglucose (coupled via acrylamide to a long PEG) but decreased for aminophlorizin binding. The latter indicates that in the membrane-bound SGLT1 the pathway to sugar translocation involves several steps with different temperature sensitivity. The latter suggests that also the aglucon binding sites for transport inhibitors have specific, temperature-sensitive conformations.

Glucose uptake into cells is essential for supply of energy in almost every organism from invertebrate to mammalian. The sugar can enter cells either by passive diffusion via facilitating transporters of the GLUT family or by secondary active transport via sodium-glucose co-transporters (SGLTs)3 using the electrochemical Na+/K-ATPase as the driving force (1–3). SGLT1 is a member of the large solute carrier family (SCL5) present in the apical membrane of the epithelial cells of the small intestine and the kidney and participating in intestinal glucose absorption and renal glucose reabsorption in many species (1). Mutations of SGLT1 can cause severe malfunctions such as glucose-galactose-malabsorption, a serious disease in newborn children in which they may die due to diarrhea and dehydration (4). Moreover, sodium-glucose co-transporters are therapeutic targets to treat hyperglycemia in type 2 diabetes (5, 6). Hence, a detailed elucidation of the structure and function of SGLT1 is required.

To this end various methods have been used such as kinetic studies (7), electrophysiology methods (8), tryptophan scanning studies (9, 10), mutagenesis studies (11, 12), x-ray crystallography (13), and plasmon resonance spectroscopy (14) to name a few. Crystallographic data are available for Vibrio para-hemolyticus sodium/galactose symporter (vSGLT) (13) in the sodium- and galactose-bound state. Overall, a group of seven central helices contributes side-chain interactions for ligand selectivity. These are stabilized by seven supporting helices. The model proposed recently by Sala-Rabanal et al. (15) integrates the kinetic and structural data available to date into a six-step alternating access model.

Our group has successfully used atomic force microscopy (AFM) and single molecule recognition force spectroscopy (16–18) to probe the transporter in its natural environment embedded in the plasma membrane of living cells under near-physiological conditions (19). The extracellular location and

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3 The abbreviations used are: SGLT1, sodium glucose co-transporter I; AFM, atomic force microscopy; thio-glc, 1-thio-β-D-glucose; AcS-, acetyltio-; hex-glc, AcS-hexyl-glucose; hex-phil, AcS-aminophlorizin; NHS, N-hydroxysuccinimide; N, newtons; AA-PEG5000, acrylamide-PEG-linker; BIC, Bayesian information criterion; mal-PEG1300, maleimide-PEG1300-NHS.

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accessibility of three extramembranous loops (loop 6–7, loop 8–9, and loop 13–14) was identified. They form a vestibule for the entry of the sugar into the translocation pathway and contain the first of several sugar recognition sites. This vestibule is accessible to the sugar only in the presence of sodium (20, 21). Phlorizin acts as a competitive inhibitor of SGLT1 with an apparent \( K_i \) of 1 \( \mu \)M (22). The phlorizin carrier complex represents a dead end conformation of the transporter in which it is locked into a condensed, rigid conformation unable to mediate translocation (23, 24). Phlorizin consists of a pyranose ring (sugar residue) and two aromatic rings joined by an alkyl spacer (the aglucon moiety, phloretin) (22). It is supposed that phlorizin binds via a two-step mechanism to the sugar translocation site and an aglucon binding site of the transporter (8, 25). One of the extracellular loops, loop 13–14, was found to provide an additional aglucon binding site. Alkyl-glucosides, such as hexyl-glucoside, also inhibit glucose transport competitively (26). The sites of interaction between the aglucon of the inhibitors and loop 13–14 differ and overlap only partly (10).

In the present work AFM was employed to further characterize the molecular interaction between SGLT1 and \( \beta \)-glucose and inhibitors with regard to their dynamics and forces. Molecular interaction between receptors and ligands is controlled by a complex array of intermolecular forces that can be characterized by their free energy landscape. AFM can be used to directly quantify the range and magnitude of the interaction forces between proteins and other molecules (28, 29). Dynamic aspects of bond rupture, e.g. dissociation rate constants, commonly used to describe the affinity between a ligand and a protein, and width of energy barrier, interpreted as the distance of the energy barrier from the energy minimum along the direction of the applied force, can be obtained by varying the loading rate of the force appliance. This provides insights into the molecular dynamics and the energy landscape for substrate/inhibitor-transporter complexes. Location of energy barriers and nature of interaction forces have been studied extensively for proteins by investigating their properties at different temperatures (30). We used a similar approach as it has been shown that sodium-dependent glucose transport is strongly temperature-dependent (11), ceasing below the transition temperature of the membrane lipids in vitro (31). In contrast, sodium-dependent, glucose-inhibitable binding of phlorizin is still demonstrable at temperatures close to 0 °C (4). Therefore, studies were performed at 10, 25, and 37 °C to investigate further the properties of the glucose translocation pathway and the inhibitor binding sites.

**EXPERIMENTAL PROCEDURES**

**AFM Tip Functionalization**—1-Thio-\( \beta \)-D-glucose (thio-glc) (Sigma), 2′-aminoethyl \( \beta \)-D-glucopyranoside, and 3-amino-phlorizin were coupled to the AFM tip using a well established three-step protocol. Therefore, 2′-aminoethyl \( \beta \)-D-glucopyranoside was synthesized as described (32) and \( N \)-acyilated with \( N \)-succinimidyl 6-(acetyltio)hexanoate as previously described for “tris-nitritoltriacetic acid” (33) yielding the product AcS-hex-glc. Immediately before coupling to the AFM tip, the acetyl group was removed with hydroxylamine (34) to obtain a free thiol, abbreviated as hex-glc (Fig. 1A). 3-Aminophlorizin was synthesized as described (35) except that sodium dithionate was used for reduction of 3-nitrophlorizin instead of catalytic hydrogenation (36). 3-Aminophlorizin was also extended with 6-(acetyltio)hexanoic acid, yielding a conjugate termed AcS-hex-phil. Immediately before coupling to the AFM tip, the acetyl group was removed with hydroxylamine (34) to obtain a free thiol. The molecule was abbreviated as hex-phil (Fig. 1A). Two different polyethylene glycol (PEG) spacers were used to couple the glucose derivatives to the tip. Maleimide-PEG\(_{1300}\)-NHS was purchased from Polypure (Oslo, Norway). Acrylamide-PEG\(_{5000}\)-NHS was synthesized by reacting NH\(_2\)-PEG\(_{5000}\)-COOH hydrochloride (Nektar Therapeutics, Huntsville, Alabama) with acrylic acid chloride and activating the terminal COOH group as \( N \)-hydroxysuccinimide (NHS) ester (37). Covalent coupling of the glucose derivatives was performed as follows: (i) amino functionalization with aminopropyltriethoxysilane in the gas phase (as described in detail in mal-linker testing and by Riener et al. (39) and Ebner et al. (38), (ii) reaction of the amino function with the NHS ester function of the cross-linker (39), and (iii) attachment of the ligands to the thiol-reactive groups on the free end of the cross-linker (Fig. 1C) (40). In particular, thio-glc (Fig. 1A) was coupled to tip-PEG\(_{5000}\)-acrylamide (Fig. 1B) by incubating these tips in 100 \( \mu \)M of buffer A (150 mM NaH\(_2\)PO\(_4\)-H\(_2\)O, 1 mM EDTA-Na\(_2\), pH 7.5, adjusted with NaOH) containing 100 mM thio-glc at room temperature for 4 h (19). 20 \( \mu \)M of 100 mM Tris(2-carboxyethyl)phosphine hydrochloride were added into the incubation solution to activate the thiol group of thio-glc. Tips were washed 3 times in buffer A (100 mM NaCl, 50 mM NaH\(_2\)PO\(_4\)-H\(_2\)O, 1 mM EDTA-Na\(_2\), pH 7.5, adjusted with NaOH) and stored in buffer A at 4 °C for at least 3 weeks. For coupling of 2′-aminooxyethyl \( \beta \)-D-glucopyranoside (hex-glc; Fig. 1A) to tip-PEG\(_{1300}\)-mal (Fig. 1B), 50 \( \mu \)L of AcS-hex-glc (4 mM) in buffer A were mixed with 5 \( \mu \)L of hydroxylamine reagent (500 mM hydroxylamine hydrochloride, 25 mM EDTA, titrated to pH 7.5 with Na\(_2\)CO\(_3\)) to remove the acetyl group. The AFM tip with the thiol-reactive end group was incubated in this solution for 4 h at room temperature. Tips were washed in buffer A and stored in buffer A at 4 °C for at least 3 weeks. For coupling of hex-phil (Fig. 1A) to tip-PEG\(_{1300}\)-mal (Fig. 1B), the same procedure was used as described for hex-glc.

**Cell Culture**—RB-SGLT1-expressing G6D3 cells, a CHO cell line stably transfected with rabbit SGLT1 generated by Lin et al. (41) were grown in 25-cm\(^2\) flasks (BD Falcon™ tissue culture flask, VWR, Vienna, Austria) under 5% CO\(_2\) at 37 °C. This cell line was cultured in Dulbecco’s modified Eagle’s medium containing high glucose (25 mM) (PAN-Biotech, Aidenberg, Germany) supplemented with 5% fetal calf serum (PA, Linz, Austria), 1 mM sodium pyruvate (Invitrogen), 2 mM l-glutamine (Invitrogen), minimal essential medium (Invitrogen), and 25 \( \mu \)M \( \beta \)-mercaptoethanol (Invitrogen). Culture medium contained 400 \( \mu \)g/mL sulfate G418 Geneticin (MedPro, Vienna, Austria) to maintain selection of transfected cells. Culture medium was renewed 3 times per week, and the cells were subcultured at 80% confluence. Cell passages below 20 were used.

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\(^4\) W. Frasch and K. R. H. Kinne, unpublished data.
for all experiments. For AFM investigations, the cells were seeded on 22-mm
diameter coverglasses from subconfluence to monolayers, and the experiments were performed within 1–5
days after seeding the coverglasses.

Cell Viability—At all temperatures, G6D3 cells stably expressing rbSGLT1 (rabbit isoform of SGLT1) appeared to be vital during AFM investigations. Topographical and deflection images of the cells (data not shown), acquired in contact mode using a magnetically driven dynamic force microscope (Pico-SPM II, Agilent Technologies, Chandler, AZ) and non-functionalized cantilevers of 0.01–0.03 N/m nominal spring constant, showed a firm attachment on glass slides as the area surrounding the nucleus region revealed to be flat. Typical cells were ~30–70 µm in diameter with a prominent bright zone representing the nucleus with a height of 2–4 µm. The pronounced flatness of the cells and the presence of characteristic morphological features such as the cytoskeleton implied that the cells were alive and vital; this allowed for stable measurements over several hours. Moreover, the cells were very suitable for force spectroscopy measurements due to their elongated spatial dimensions and their low height.

In pilot transport and binding experiments it was found that, as expected, sodium-dependent α-methyl-D-glucose uptake at 10 °C was only 5% that of the uptake observed at 37 °C, whereas a slightly increased phlorizin binding was observed.

AFM Force Microscopy/Spectroscopy—All AFM measurements were carried out in Kreb’s- Ringer-HEPES buffer (120 mM NaCl, 4.7 mM KCl, 2.2 mM CaCl₂, 1.2 mM MgCl₂, and 10 mM HEPES; the pH was adjusted to 7.4 with Tris-base) using a magnetically driven dynamic force microscope (PicoSPM II, Agilent Technologies, Tempe, AZ). Topographical and deflection images of the cells were acquired in contact mode AFM using non-functionalized cantilevers (Veeco Instruments, Mannheim, Germany) of 0.01–0.03 N/m nominal spring constant in order to record the viability of the cells. For detection of glucose/phlorizin-SGLT1 recognition, force-distance cycles were performed at controlled temperature conditions, i.e. 10 °C, 25 °C, and 37 °C, using ligand-functionalized cantilevers (Veeco Instruments) with 0.01–0.03 N/m nominal spring constant in the conventionally contact force spectroscopy mode. In particular, the cells grown on a plastic slide were mounted on an Agilent Peltier-100 sample compartment spanning a temperature range from 0 to 40 °C. A DC power supply device (McVoice) was connected with the Peltier-100 to sustain a constant temperature by applying a current. The temperature was controlled by permanently monitoring the resistance of the Peltier element using a digital multimeter (SkyTronic). A gravity-fed water cooling system was installed to decrease the minimum sample temperature, reduce the power requirement, and prevent overheating of the device when the sample is cooled. The flow rate of the water through the Peltier element was adjusted to 25 ml/min. In addition, the scanner housing was floated with argon gas when the temperature was set to 10 °C to prevent moisture on the glass of the “AFM nose.”

Force-distance cycles were recorded with the assistance of a CCD camera for positioning the cantilever above a single cell.

At a fixed lateral position an AFM tip carrying a glucose or phlorizin molecule approached (trace) the cell surface and was subsequently retracted (retrace) (see Fig. 2B). During this cycle the cantilever deflection (Δx), which can directly be converted into a force (f) according to Hook’s law (f = kΔx, k being the cantilever spring constant), was continuously monitored and plotted versus tip-surface separation (i.e. distance). Upon approach no binding of the cantilever occurred. As the cantilever touched the surface it deflected upward due to a repulsive tip-sample interaction. Subsequent tip-surface retraction led to relaxation of the cantilever bending until the repulsive force dropped to zero. If a ligand on the tip had bound to a surface protein, an attractive force developed upon retraction, causing the cantilever to bend downward. At a particular unbinding force (f_u) the ligand was detached from the surface-molecule, and the cantilever jumped back to zero deflection.

The interaction efficiency of thio-glucose on AA-PEG₅₀₀₀, hex-gluc, and hex-phl on maleimide-PEG₃₀₀₀-NHS (mal-PEG₃₀₀₀) was probed at 10 °C, 25 °C, and 37 °C by recording up to 1000 force-distance cycles with a z-range of 1000 N/m/s and a sweep frequency of 1 Hz (amounting to a sweep rate of 2000 nm s⁻¹) on a single cell. Within these 1000 force-distance cycles the position of the cantilever where it touched the cell surface was varied several times. Binding probabilities (the chance for observing an unbinding event within one force-distance cycle) were derived for each linker-molecule-compound at a certain temperature and expressed as the mean value ± S.E. The specificity of the recognition events between SGLT1 and glucose derivatives at each temperature was proven by adding 0.5 mM phlorizin to the buffer solution and thus blocking the substrate interaction sites of SGLT1. The binding probability was acquired for initial conditions (cells in Krebs-Ringer-HEPES buffer without phlorizin), phlorizin-containing conditions, and washout conditions. Binding probabilities are given as the mean values ± S.E.

A close geometrical and chemical fit within the binding pocket has to be accomplished as a prerequisite for molecular recognition; the underlying specific interactions are a distinct class of highly complementary non-covalent bonds composed of several weak interactions such as electrostatic, polar, van der Waals, hydrophobic, and hydrogen bonds. The binding energy E_p, given by the free energy difference between the bound and the free state, is the common parameter to describe the strength of a bond. Ligand-receptor binding is generally a reversible reaction. Thus, the average lifetime of a ligand-receptor bond, τ(0), is given by the inverse of the kinetic off rate k_off. Therefore, ligands will dissociate from receptors without any force applied to the bond at times larger than τ(0). In contrast, if molecules are pulled faster than τ(0) the bond will resist and require a force for detachment. The activation energy for the dissociation is the difference in energy between the initial state and the transition state of the highest energy to which the system must be raised before dissociation can occur. In force spectroscopy experiments, the dynamics of pulling on specific receptor-ligand bonds is varied, which leads to detailed structural and kinetic information of the bond breakage. Moreover, energy landscapes of receptor-ligand bonds can be detectable for certain time scales in force-induced unbinding; especially, force
Single Molecule Force Spectroscopy of SGLT1

spectroscopy appears to be a proper tool to discover hidden activation barriers and explore the energy landscape in greater detail.

To study ligand/SGLT1 unbinding forces and kinetic and structural parameters of the binding pocket, the tip approach-and-retract velocities were varied in the range from 1 μm/0.2 s to 1 μm/2 s. Thus, the loading rate, which was determined from the product of the effective spring constant (cantilever spring and bound molecule) times retraction velocity, was varied in the range from ~1.5 × 10^4 to ~2 × 10^4 pN/s. The measurements were carried out at 10 °C, 25 °C, and 37 °C. Several individual measurements were performed for each linker-molecule compound (thio-gluc on AA-PEG5000, hex-gluc on to mal-PEG1300 and hex-phil on mal-PEG1300) using cantilevers of 0.01 and 0.02 N/m nominal spring constant. For each tip-ligand system, 1000–2000 force-distance cycles were recorded at each loading rate. Single unbinding events were collected, and the most probable unbinding force (f_p) was plotted against the log-arithmetic of the loading rate (r) to gain the width (x_p) of prominent energy barriers along the dissociation pathway and the dissociation rate constant (k_off). In particular, a maximum likelihood approach was employed as a statistical estimation technique to fit a statistical model to the obtained data, thus providing estimations for model’s parameters. Here, the consequent unbinding force was plotted against the particular loading rate for every single detected unbinding event. The data points of each recognition event were merged in one dynamic force spectra plot to obtain dynamic aspects of a bond. As an analytical model to estimate for the maximum likelihood of x_p and k_off the Evans model was employed. The Bayesian information criterion (BIC) was intended to choose the best model from among several competing/equal models. The BIC was designed to select the best model that describes the data adequately without using too many parameters. The formulation for the BIC is given by,

\[
\text{BIC} = -2(l(\psi|y) - l(\psi^*|y)) + 2\log(n) \quad \text{(Eq. 1)}
\]

where \(l(\psi|y)\) is the log-likelihood of the model under consideration, \(l(\psi^*|y)\) is the log-likelihood of the most likely model in the subset of models considered, \(p\) is the number of parameters fit in the model, and \(n\) is the number of observations. Thus, the model with the smallest BIC value is chosen as the best model (41). Student’s t test was performed to compare the gained \(k_{off}\) and \(x_p\) values and account for significant differences.

The spring constant of the cantilevers was determined by using the thermal noise method (42, 43). The deflection sensitivity of the photodetector was determined from the slope of the force-distance plot, taken on the hard bare surface of a glass coverslip. Force-distance cycles were analyzed using Matlab Version 7.1 (MathWorks Inc., Natick, NA) where empirical force distributions of the rupture forces of the unbinding event (probability density function (pdf)) were calculated as described previously (44).

RESULTS

Design of Various Substrate and Inhibitor Derivatives for Tethering to AFM Tips—To investigate SGLT1 interactions at the single-molecule level by AFM, three different compounds were coupled to AFM tips, thio-gluc, as an example for the substrate glucose, hex-gluc, as representative of alkyl-glucosides, and 3-aminophlorizin (aminophil) (Fig. 1A; see “Experimental Procedures”). The molecules were tethered via heterobifunctional PEG linkers varying in length (N) and end-group (TRG) having the general structure TRG-PEG_N-NHS (see Fig. 1, B and C), “TRG” denoting a thiol-reactive group (maleimide or acrylamide) and “NHS” indicating a reactive NHS ester. In detail, thiglucose was linked to the tip via a long (~30–40 nm) acrylamide-PEG-linker (AA-PEG5000) using a well established three-step protocol (see Fig. 1C). The latter has one of the thinnest and least bulky functional PEG-chain end groups. Amino ethyl β-n-glucopyranoside as well as 3-aminophlorizin was coupled to the tip using a short (~8 nm) mal-PEG1300.

Single Molecule Recognition of SGLT1—A scheme of single-molecule recognition force detection of a glucose derivative-SGLT1 complex on living G6D3 cells by using AFM is illustrated in Fig. 2A. In this so-called force-distance mode, the deflection angle of the cantilever is measured as a function of the vertical position of the cantilever. A typical force curve (retraction) for a single molecule glucose-SGLT1 binding event on the surface of a G6D3 cell with a ligand-conjugated tip is shown in Fig. 2B. The specificity of the interaction between all tip-tethered glucose derivatives and SGLT1 on the cell membrane could be confirmed by masking the substrate binding site of SGLT1 with free phlorizin (see the inset in Fig. 2B and Fig. 3). The number of recognition events dropped significantly, to ~20% of the unblocked conditions. The same low binding probability was observed when parental CHO cells not expressing SGLT1 were investigated (data not shown).

Binding Behavior of Thioglucose, Alkyl-glucose, and Amino-phlorizin Tethered AFM Cantilevers at Different Temperatures—It was found that SGLT1 could specifically be detected at 10 °C, 25 °C, and 37 °C when living cells were probed using cantilevers carrying the substrate or inhibitors. The observed binding probabilities (probability of finding an unbinding event in force-distance cycles) of all tips to SGLT1 from several experiments ranged from 9 to 27% but were not statistically different.

Comparing the binding probability of thioglucose on AA-PEG5000 at different temperatures, the binding probability was low at 10 °C (8.7 ± 2.28%) and increased significantly for 25 °C (14.62 ± 3.69%) and further at 37 °C (26.72 ± 4.56%) (Fig. 4A). It is noteworthy that at 37 °C thioglucose on AA-PEG5000 shows the highest binding probabilities of all derivatives tested. The interaction of hexyl-glucose on mal-PEG1300 with the cells was not altered significantly by changing the temperature, as the binding probability ranged from 11.3 ± 1.16 to 15.6 ± 2.03% for all temperatures (Fig. 4B). Similarly, the recognition events for aminophlorizin on mal-PEG1300 showed only small changes: a slightly increased binding at 37 °C (19.38 ± 3.28%) compared with 10 °C (16.9 ± 1.55%) and lesser binding probability at 25 °C (11.13 ± 1.00%) (Fig. 4C).

Kinetic and Structural Parameters of the Binding Pockets—Experiments were performed at various tip-retraction velocities, leading to a force spectrum in dependence on the loading rate (pulling velocity times effective spring constant). Unbinding forces for thioglucose on AA-PEG5000 hexyl-glucose on
mal-PEG1300, and aminophlorizin on mal-PEG1300 were deter-
mixed for retraction speeds in the range between $1.5 \times 10^3$ to
$2 \times 10^4$ pN/s. All collected force-distance cycles were evalu-
ated by selecting every single unbinding event and generating
the probability density function of the force, effective spring
constant, and length. The individual forces of each unbinding
event were plotted against the logarithm of the loading rate and
thus merged into one particular force spectrum (45). The
results obtained for thioglucose on AA-PEG5000 probed at dif-
ferent temperatures are shown in Fig. 5. As expected (46, 47),
the forces increased with increasing loading rates. Subse-
quently, data were analyzed using a maximum likelihood
approach yielding a comprehensible estimation for the width of
the energy barrier, $x$, and the dissociation rate constant, $k_{off}$.
Herein, a cloud of data points was gained to which a maximum
likelihood fit using the Evans model as the analytical model was

FIGURE 1. Linkage of substrates and inhibitors of SGLT1 to the AFM tip via heterobifunctional PEG linker by a three-step protocol. Thio-glc, hex-glc, and AcS-aminophlorizin (amino-phi) (A) were coupled to cross-linkers varying in lengths having a maleimide end group (~8 nm) or an acrylamide-end group (~30–40 nm) (B). C, amino groups were generated on the silicon nitride AFM cantilever tip using aminopropyltriethoxysilane (step 1) to attach the flexible distensible PEG-linker covalently via its NHS end to the tip (step 2). The ligand was linked to the free thio-reactive end of the PEG chain (step 3). Thioglucose was coupled to the acrylamide-linker, hexyl-glucose, and aminophlorizin were bound to the maleimide-linker. Circles in B and C indicate the thio-reactive group (TRG). In C the dotted line circles the ligand, which is coupled to the PEG linker.

FIGURE 2. Single molecule recognition of SGLT1 on the surface of living cells. A, schematic representation of a force-distance cycle, where the cantilever approached the cell surface and subsequently retracted (solid line). The deflection angle of the cantilever remained zero (1) until the tip contacted the sample surface from where the cantilever was bent upward (2). Upon retraction a force signal with a distinct shape occurred when an interaction between the ligand and a receptor occurred (3). The cantilever was increasingly bent downward until the bond broke at a defined unbinding force $f_u$ required to break a single ligand-receptor interaction (4). B, force curves show specific interactions between thioglucose coupled via an acrylamide-linker and SGLT1 upon tip-surface retraction. The specific recognition disappeared in the presence of phlorizin (inset).
applied to estimate $x_B$ and $k_{off}$ (Fig. 5; also see “Experimental Procedures”). From these, $x_B$ and $k_{off}$ of the SGLT1-thioglucose bond were calculated. As illustrated in Fig. 5, for instance, the width of the energy barrier, $x_B$, and dissociation rate constant, $k_{off}$, at 37 °C were 6.24 ± 0.08 Å and 0.26 ± 0.02 s$^{-1}$, respectively (Fig. 5C).

The obtained forces and the corresponding loading rates of all investigated cantilevers at a distinct temperature, i.e. 10 °C, 25 °C, or 37 °C, were merged into one dynamic force spectra plot (Fig. 6). Thus, hundreds of particular unbinding events were taken into account individually making estimates for the structural and kinetic parameter more precise. The values obtained for dissociation rate and the width of the energy barriers are listed in Table 1. Thioglucose on AA-PEG5000 revealed the lowest dissociation rate constant, $k_{off}$ (0.26 ± 0.02 s$^{-1}$), at 37 °C indicating that the lifetime of the SGLT1-thioglucose complex is longest at this temperature compared with 25 °C (0.83 ± 0.07 s$^{-1}$) and 10 °C (0.6 ± 0.05 s$^{-1}$). These values were significantly different, with $p < 0.001$. In contrast, hexyl-glucose bound to mal-PEG$_{1300}$ dissociated from the transporter in a quite similar time scale at each temperature ($k_{off} = 0.41 ± 0.03$ s$^{-1}$ at 10 °C, 0.50 ± 0.07 s$^{-1}$ at 25 °C and 0.4 ± 0.07 s$^{-1}$ at 37 °C). However, aminophlorizin on mal-PEG$_{1300}$ showed an inversed dissociation behavior compared with thioglucose, as the lifetime of the aminophlorizin-SGLT1 complex is shortened with increasing temperatures. Thus, aminophlorizin was the molecule that dissociated significantly faster at 37 °C ($k_{off} = 1.00 ± 0.06$ s$^{-1}$) than at 10 °C ($k_{off} = 0.34 ± 0.03$ s$^{-1}$).

Moreover, the single molecular binding energy width for glucose/inhibitor/cotransporter interactions could be extracted from the plot of unbinding force versus the loading rate (see “Experimental Procedures”). The width of the energy barrier, $x_B$, can thereby be taken as a measure for the location of the maximum energy barrier within the binding pocket. The obtained values for the $x_B$ of thioglucose on AA-PEG$_{5000}$ increased with increasing temperatures ($x_B = 3.96 ± 0.08$ Å at 10 °C, 4.99 ± 0.1 Å at 25 °C, and 6.24 ± 0.08 Å at 37 °C). $x_B$ for thioglucose on AA-PEG$_{5000}$ at 37 °C was largest among all linker compounds investigated. The length scales of the binding pocket for hexyl-glucose on mal-PEG$_{1300}$ probed at different temperatures were also altered within a $p$ value < 0.01, showing the highest value at 10 °C ($x_B = 3.48 ± 0.05$ Å) and the lowest at 37 °C ($x_B = 2.75 ± 0.1$ Å). Similar values for $x_B$ were observed for aminophlorizin on mal-PEG$_{1300}$ at 10 °C ($x_B = 3.32 ± 0.06$ Å) and 37 °C ($x_B = 2.93 ± 0.04$ Å).

**DISCUSSION**

In the present study SGLT1 was mapped on the membrane of living G6D3 cells stably expressing the transporter by AFM cantilevers being loaded with 1-thio-β-D-glucose, modified amino-hexyl-glucose, and aminophlorizin. The binding behavior, e.g. binding probability, interaction forces, etc. of particular ligands was ascertained at different temperatures by performing single molecule recognition force spectroscopy experiments. Moreover, new structural and kinetic parameters of the

**FIGURE 3. Specific interaction of glucose and inhibitors at 37 °C.** Data of one typical experiment are presented for each cantilever, probed in different conditions: initial condition (first column), block condition (second column), and washout condition (third column). Values are the mean ± S.E. for $n = 2000$–3000, where 2–3 different cells were probed at each condition. The specificity of the recognition events could be demonstrated by a significantly reduced binding probability in the presence of phlorizin when compared with control levels (initial condition) with $p < 0.005$. After a careful washout procedure the interaction increased again. **FIGURE 4. Binding probabilities of glucose, hexyl-glucose, and aminophlorizin at different temperatures.** The interaction between thioglucose on AA-PEG$_{5000}$ (A), hexyl-glucose on mal-PEG$_{1300}$ (B), and aminophlorizin (C) on mal-PEG$_{1300}$ was studied at 10 °C, 25 °C, and 37 °C using the same cantilever and the same cells. Values are the mean ± S.E. of three to five different cantilevers and two different cells for one cantilever. On each cell up to 1000 force-distance cycles were recorded.
interactions of SGLT1 with glucose and transport inhibitors could be gained.

**New SGLT1 Sensors**—In the current study new coupling chemistry for tethering thioglucose, hexyl-glucose, and aminophlorizin to the AFM tip was developed. Previously, AFM studies on SGLT1 had been performed mainly with a vinylsulfone linker between the PEG chain and thioglucose or a modified 3′-aminophlorizin (19, 20, 48). By using acrylamide as the thio-reactive group to attach thioglucose to PEG5000, the size (and charge) of the thioester linkage is reduced compared with vinylsulfone-PEG800. In addition, the PEG chain is extended, providing the least bulky and most flexible sugar probe used until now (Fig. 1B).

In AcS-hexyl-glucose, terminally coupled to the thio-reactive group of the PEG chain, a long stretch of an alkyl chain is exposed, allowing additional interactions of this entity with the transporter similar to those occurring with free hexyl-glucose (26). In addition, phlorizin was tethered to the AFM tip via a C6 chain to an amino group at the C3 atom of its aromatic ring B; this provides more mobility to phlorizin on the cantilever tip but leaves the major binding sites of its aglucon moiety, the 4′-OH and 6′-OH of the adjacent aromatic ring A, freely accessible to SGLT1 (49). Replacement of vinylsulfone by maleimide accelerates the coupling reaction.

With all three newly synthesized sensors a specific recognition of SGLT1 can occur, as the major binding sites of glucose are freely accessible. Thus, a proper fitting of the sugar moieties into the binding pocket of SGLT1, mainly formed by transmembrane helices 10–13, can be achieved (20, 50–53). The specificity of interaction of the probes was demonstrated first of
all by the fact that a significant number of unbinding events was only observed in CHO cells transfected with the sglt1 gene. In addition, for all probes the number of unbinding events was almost identical, and low concentrations of phlorizin reversibly reduced the binding.

Effect of Temperature on SGLT1 Properties—In the current study SGLT1 was investigated in living cells incorporated into the plasma membrane. Temperature changes can affect such a system in various ways. With regard to the membrane, phospholipids undergo phase transitions and change their fluidity. Previous studies have shown that SGLT1 activity changes in parallel to the changes in fluidity when it is incorporated into liposomes composed of phospholipids with different transition points (54). This strong dependence is probably due to the fact that ~40% of the Trps responsible for hSGLT1 fluorescence are in close contact to the membrane phospholipids (23).

The SGLT1 protein is postulated to undergo a series of conformational changes while transporting glucose from the extracellular side of the cell membrane into the cytoplasm in an alternating accessibility model. Starting with an outward-facing configuration, sodium binding opens up access to the sugar translocation pathway. Thereafter, glucose is bound and occluded in the translocation site. This is followed by a conformational change from an outward-facing occluded state to an inward-facing occluded state. Upon opening of the inward gate, the Na\(^+\) ions and sugar are released into the cell interior. The transport cycle is completed by the transition from the inward-facing ligand-free state to the outward-facing ligand-free state (15). This transport cycle involves rearrangement of intramolecular binding sites as well as changes in the orientation of transmembrane helices. By changing the temperature, the conformational equilibrium to the various states might be altered.

FIGURE 6. Dynamic force spectra of the different SGLT1 ligand complexes at different loading rates. The most probable unbinding forces were plotted against the logarithm of the loading rate. Comparisons of force spectra with thioglucose on AA-PEG\(_{5000}\), hexyl-glucose on mal-PEG\(_{1300}\), and aminophlorizin on mal-PEG\(_{1300}\) at 10 °C (A), 25 °C (B), and 37 °C (C) are shown.

TABLE 1
Kinetic property of glucose, hexyl-glucose, and aminophlorizin binding to rbSGLT1
The values were obtained from the experiments depicted in Fig. 6. \(k_{\text{off}}\) of thioglucose on AA-PEG\(_{5000}\), at 37 °C compared with \(k_{\text{off}}\) of thioglucose on AA-PEG\(_{5000}\), at 10 °C and 25 °C and \(k_{\text{off}}\) of aminophlorizin on mal-PEG\(_{1300}\), at 37 °C when compared with \(k_{\text{off}}\) of aminophlorizin on mal-PEG\(_{1300}\), at 10 °C and 25 °C were significantly different. \(x_\text{gg}\) of thioglucose on AA-PEG\(_{5000}\), at 37 °C when compared with \(x_\text{gg}\) of the other ligands was significantly different.

| SGLT1 sensors         | Width of the energy barrier, \(x_{\text{gg}}\) | Dissociation rate constant, \(k_{\text{off}}\) |
|-----------------------|---------------------------------------------|---------------------------------------------|
|                       | 10 °C | 25 °C | 37 °C | 10 °C | 25 °C | 37 °C |
| Thio-glc on AA-PEG\(_{5000}\) | 3.96 ± 0.08 | 4.99 ± 0.1 | 6.24 ± 0.08 | 0.60 ± 0.05 | 0.83 ± 0.07 | 0.26 ± 0.02 |
| hex-glc on mal-PEG\(_{1300}\) | 3.48 ± 0.05 | 3.18 ± 0.1 | 2.75 ± 0.1 | 0.41 ± 0.03 | 0.50 ± 0.07 | 0.40 ± 0.07 |
| Amino-phl on mal-PEG\(_{1300}\) | 3.23 ± 0.06 | 4.28 ± 0.1 | 2.93 ± 0.04 | 0.34 ± 0.03 | 0.54 ± 0.07 | 1.00 ± 0.06 |

\(^a_p < 0.005.\hspace{1cm} ^b_p < 0.001.\)
Thermal fluctuations of proteins are in general important for biological activity (55); thus the temperature changes in this work may modulate the protein flexibility, and folding and orientation of substructures within the membrane can be affected or accessibility to the sugar binding residues. Finally the interaction forces between ligands and proteins (hydrogen bonding and Coulomb and van der Waal forces) are temperature-dependent (56).

At 10 °C, which is below the transition point of the membrane lipids, SGLT1 is probably constrained in its structural movements and trapped in non-transporting conformational states. Interestingly, even at this temperature, SGLT1 was able to interact specifically with all ligands, suggesting that sodium and sugars can be bound by the transporter. Evidence for this interaction can also be derived from phlorizin binding studies where a sodium-dependent, glucose inhibitible binding was found.4,5 This implies that the opening of the vestibule and the initial hydrophilic sugar binding cavity is rather temperature-independent as it occurs at the surface of the transporter. Similar conclusions can be drawn from tryptophan fluorescence studies with purified hSGLT1 (57). With an increase of the temperature to 25 °C, all binding probabilities increase similar to the transport activity and turnover rate; at a higher turnover rate probably more SGLT1s in the outward open conformation face the membrane surface per time unit; thus, more binding events can occur. At 37 °C, another increase in binding probability is observed for thioglucose on AA-PEG5000 but not for hexyl-glucose or aminophlorizin on mal-PEG1300. This result might suggest that at this temperature the thigolucose probe recognized SGLT1 in more than one conformation. Further evidence for this assumption is presented below.

Dynamic force spectroscopy measurements track the dynamic behavior of SGLT1 upon substrate and inhibitor binding. Here, structural and kinetic parameters of their SGLT1 bond were gained by varying the cantilever retraction speed over several orders of magnitude. In particular, when probed at 10 °C, thioglucose was detached faster from SGLT1 than at 25 °C and 37 °C. Thermodynamically, the opposite would be expected for simple binding reactions. The long-lasting average lifetime of the SGLT1-thioglucose-AA-PEG5000 bond at 37 °C might be explained by additional conformation of SGLT1 in which thioglucose on the long AA-PEG5000-linker is able to penetrate deeper into the translocation pathway of the transporter, beyond the initial sugar binding site. The very slim reactive end group of the PEG linker could facilitate this entry. This assumption is supported by the previously reported increased unbinding force when thioglucose is coupled to the AFM tip via an acrylamide linker as compared with the bulkier vinylsulphone linker (58). The highest value for the width of the binding pocket observed for thioglucose on AA-PEG5000, at 37 °C (see Table 1) provides further evidence. The latter can be directly related to the number of protein residues contributing to binding sites (40). Hirayama et al. (60, 2007) reported that 3 residues between transmembrane helices 10 and 13 play a direct role in sugar binding and translocation from the external environment, 3 more face the binding site, and 8 further residues, although not contributing to the binding of sugar directly, affect sugar recognition. Thus, it might be assumed that at 37 °C some of the primarily not accessible binding sites further down in the translocation pathway are reached by thioglucose on AA-PEG1300 (21). A sequential multistep mechanism for sugar recognition and translocation has also been proposed based on prior AFM investigations (20).

Thioglucose on AA-PEG5000 and aminophlorizin on mal-PEG1300 showed an inverse behavior in $k_{off}$ when compared at 10 °C and 37 °C, and thioglucose on AA-PEG5000 was bound longest to the transporter at 37 °C among all probed derivatives and temperatures, whereas aminophlorizin yielded the highest dissociation rate constant (shortest time) at 37 °C. The findings for phlorizin dissociation are in good agreement with previous studies (3) as inhibitor association and dissociation were reported to be enhanced at 37 °C (59–61). This observation suggests that the glucose moiety of phlorizin is not capable of penetrating as deep into the translocation pathway as thioglucose on AA-PEG5000 due to steric hindrance by the aromatic aglucon. This assumption is supported by the lower width of the energy barrier. PEG is an ideal force transducer (62), and the force spectrum measured with dynamic force spectroscopy is largely independent on length of the handle (45, 63) through which the ligands are attached to the AFM tips. It thus seems improbable that the observed effects are due to the difference in length of the linker PEG chains of the two probes, ~170 Å for phlorizin and ~700 Å for thioglucose. Moreover, the hydrophilic cavity of the transporter, which leads to the translocation, is only 7 Å deep (57).

One has to postulate, in addition, that the aglucon binding site of SGLT1 on the membrane surface is altered at this temperature. Previous studies by our groups have shown that, in rbSGLT1, loops 13–14 facing the extracellular space is critically involved in the binding of the aglucon (64). In vitro studies suggest that the loop has a substructure of two small helices and interacts in several regions with membrane phospholipids (10, 65). These structural requirements for a tight interaction with the aglucon might be partly lost due to the increased lipid fluidity, and thereby overall binding affinity is reduced.

Hexyl-glucose, as a model for the interaction between SGLT1 and alkyl-glucosides, shows binding probabilities quite similar to those obtained for the aminophlorizin-tagged probe; with regard to $k_{off}$ and width of the energy barrier it lies in between thioglucose on AA-PEG5000 and aminophlorizin. Thus, as expected, the sugar moiety and the alkyl chain are involved in the binding to SGLT1; however, the relative role of the aglucon-receptor interactions seems to be less important for the overall binding reaction. This is also reflected in the 10× lower affinity to the carrier in transport studies (26).

In conclusion, a binding schematic for the substrate and transport inhibitors at different temperatures might be hypothesized; (i) glucose, phlorizin, and hexyl-glucose at all temperatures interact with SGLT1, the latter two via both sugar and aglucon binding sites located on the membrane surface, (ii) with increasing temperature the sugar translocation pathway opens, and additional binding sites for glucose become available and accessible, and (iii) the increase in temperature perturbs the conformation of the aglucon binding sites required for optimum interaction with transport inhibitors. The distinct temperature dependence observed in the current study sup-
ports the existence of multiple steps in the outward-facing translocation pathway of SGLT1, from the initial sugar binding followed by stereo-selection to occlusion of the sugar. It also corroborates the assumption of a two-site two-step mechanism for the binding of transport inhibitors. To describe the dynamical behavior of SGLT1 upon substrate and inhibitor binding more precisely, e.g. determining distinct binding sites or account for distinct conformations, single-molecule force spectroscopy experiments combined with mutagenesis studies would be required in which specific residues involved in sugar binding and translocation are exchanged. Furthermore, it has to be determined whether the changes induced by the different temperatures reflect conformations occurring in the normal translocation cycle.

Finally, these studies demonstrate that AFM force spectroscopy is a useful method for exploring the functional dynamics of membrane transporters and paves the way for investigating the influence of transport-inhibiting substances potentially applicable as therapeutics on the single molecule level under near-native conditions.

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