Trypanosoma brucei Bloodstream Forms Depend upon Uptake of myo-Insitol for Golgi Complex Phosphatidylinositol Synthesis and Normal Cell Growth

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myo-Insitol is a building block for all inositol-containing phospholipids in eukaryotes. It can be synthesized de novo from glucose-6-phosphate in the cytosol and endoplasmic reticulum. Alternatively, it can be taken up from the environment via Na+- or H+-linked myo-insitol transporters. While Na+-coupled myo-insitol transporters are found exclusively in the plasma membrane, H+-linked myo-insitol transporters are detected in intracellular organelles. In Trypanosoma brucei, the causative agent of human African sleeping sickness, myo-insitol metabolism is compartmentalized. De novo-synthesized myo-insitol is used for glycosylphosphatidylinositol production in the endoplasmic reticulum, whereas the myo-insitol taken up from the environment is used for bulk phosphatidylinositol synthesis in the Golgi complex. We now provide evidence that the Golgi complex-localized T. brucei H+-linked myo-insitol transporter (TbHMIT) is essential in bloodstream-form T. brucei. Downregulation of TbHMIT expression by RNA interference blocked phosphatidylinositol production and inhibited growth of parasites in culture. Characterization of the transporter in a heterologous expression system demonstrated a remarkable selectivity of TbHMIT for myo-insitol. It tolerates only a single modification on the inositol ring, such as the removal of a hydroxyl group or the inversion of stereochemistry at a single hydroxyl group relative to myo-insitol.

In all eukaryotes, myo-insitol is the precursor for all inositol-containing phospholipids, including phosphatidylinositol (PI), phosphatidylmyo-inositol (poly)phosphates, inositol phosphorylceramide (IPC), and glycosylphosphatidylinositol (GPI). In mammalian cells, it is taken up from the environment via sodium/myo-insitol cotransporters (SMITs) or proton-linked myo-insitol transport (T. brucei H+-linked myo-insitol transporter [TbHMIT]). Human SMIT1 and SMIT2 belong to the sodium/glucose transporter family, SGLT1/SLC5, whose members in general mediate uptake of sugars and osmolytes in the gastrointestinal tract and the kidney (1). They are localized in the plasma membrane and, besides myo-insitol, also transport xylose and glucose (2, 3). In contrast, the human HMIT belongs to the sugar/polysaccharide transport facilitator family, GLUT/SLC2A (4). While most members of this family are also located in the plasma membrane and regulate sugar homeostasis within the body, members of the family of subclass III transporters, to which HMIT belongs, are typically localized intracellularly (5, 6). Interestingly, GLUT12/SLC2A12 and HMIT/SLC2A13 have been found to colocalize with Golgi complex markers (7, 8). Although the substrate specificities of the subclass III GLUT/SLC2A transporters have been studied in model systems, their physiological roles have not been firmly established (5, 6). Notably, HMIT completely lacks sugar transport activity but instead transports myo-insitol with a Km of approximately 100 μM in a Xenopus oocyte expression system (8).

As an alternative to uptake, myo-insitol can be synthesized de novo in a reaction sequence that is conserved from bacteria to mammals, using glucose-6-phosphate as an initial substrate (9). Endogenously produced as well as imported myo-insitol can then be used for inositol phospholipid synthesis in a process that is generally believed to occur in the endoplasmic reticulum (ER). In yeast, plants, protozoa, and mammals, PI synthesize has been localized to the ER using cell fractionation and immunolocalization studies (10–13). Reconstitution experiments involving purified PI synthase from Saccharomyces cerevisiae showed that it incorporates asymmetrically into model vesicles, suggesting that its active site may face the cytosolic side of the ER in yeast (10). However, the in vivo topology of the active site of PI synthase has not been determined experimentally.

Interestingly, recent reports in protozoan parasites indicated that the Golgi complex represents an additional site for inositol phospholipid synthesis. Direct evidence for the presence of PI synthase in the Golgi complex was obtained from immunolocalization studies in Trypanosoma brucei bloodstream forms (13), showing that the enzyme has a dual localization in the ER and Golgi complex. In support of the idea of the Golgi complex being the site of bulk PI synthesis in trypanosomes (13), a subsequent study revealed that T. brucei procyclin forms express a plasma membrane- and Golgi complex-localized proton-linked myo-insitol transporter, TbHMIT (14). Downregulation of TbHMIT in-
hibited bulk PI formation but had no effect on GPI synthesis, demonstrating that PI synthesis in T. brucei is compartmentalized, with the Golgi complex representing the site of synthesis of bulk membrane PI utilizing exogenous myo-inositol (14) and the ER being the site of PI synthesis for GPI production utilizing de novo-synthesized myo-inositol (15). Transporter-mediated myo-inositol uptake has also been characterized in other protozoan parasites, including Leishmania donovani (16–19) and Trypanosoma cruzi (20, 21). These parasites all cause devastating human diseases, including leishmaniasis, Chagas disease, and human African sleeping sickness. Membrane transporters are of particular importance for these pathogens to acquire essential nutrients from their respective hosts and offer attractive targets for rational drug design and/or the delivery of cytotoxic substrate analogues. The reported dependence of T. brucei proyclic forms in culture on the presence of exogenous myo-inositol (13, 14) validates HMIT as a potential drug target.

In this report, we extend our previous analysis of myo-inositol uptake in T. brucei proyclic forms to the pharmacologically more relevant bloodstream form. We demonstrate that expression of TbHMIT is essential for normal growth of T. brucei bloodstream parasites in culture and that it is involved in myo-inositol transport and PI formation within the Golgi complex. In addition, we have tested a series of myo-inositol stereoisomers and structural analogs and related compounds to characterize the substrate specificity of TbHMIT.

MATERIALS AND METHODS

Unless otherwise stated, all reagents were of analytical grade and purchased from Merck (Darmstadt, Germany), Sigma-Aldrich (Buchs, Switzerland), or ICN Biomedicals (Tägerig, Switzerland). Antibiotics and fetal bovine serum (FBS) were obtained from Invitrogen (Basel, Switzerland). Primers and sequencing services were from Microsynth AG (Balgach, Switzerland). Restriction enzymes were purchased from Thermo Scientific (Wohlen, Switzerland). myo-[2-3H(N)]inositol (15 to 20 Ci/mmol) (myo-[3H]inositol) and [3H]ethanolamine (40 to 60 Ci/mmol) were from American Radiolabeled Chemicals (St. Louis, MO, USA), and dCTP-[α-3P] (3000 Ci/mmol) was from PerkinElmer Life Sciences (Schwerzenbach, Switzerland).

Trypanosomes and culture conditions. Bloodstream-form T. brucei parasites derived from line MiTat 1.2, coexpressing T7 RNA polymerase and a tetracycline repressor (22), were cultured at 37°C with 5% CO2 in RPMI 1640 containing 5% heat-inactivated FBS. Restriction enzymes were purchased from Thermo Scientific (Darmstadt, Germany) and Sigma-Aldrich (Buchs, Switzerland) and quantified using the Rita Star software provided by the manufacturer. For analysis of GPI precursors, bloodstream-form trypanosomes were labeled for 16 h with trace amounts of [3H]ethanolamine and transferred to GeneScreen Plus nylon membranes (PerkinElmer Life Sciences). 32P-labeled probes were made by random priming of the same PCR products as were used as insertions in the stem-loop vector using a Prime-a-Gene labeling system (Promega). Hybridization was performed overnight at 60°C in hybridization buffer containing 7% (wt/vol) SDS, 1% (wt/vol) bovine serum albumin, 0.9 mM EDTA, and 0.5 M Na2HPO4 (pH 7.2), and the membrane was analyzed by autoradiography using BioMax mass spectrometry (MS) film and a Trans-Blot-HE intensifying screen. rRNA was visualized on the same formyldehydro-agarose gel by ethidium bromide staining to control for equal loading.

myo-Inositol uptake assays. T. brucei bloodstream cells (1 × 107 cells) at the mid-log phase were harvested by centrifugation at 1,250 × g for 10 min and suspended in phosphate-buffered saline (PBS; 135 mM NaCl, 1.3 mM KCl, 3.2 mM Na2HPO4, 0.5 mM KH2PO4, pH 7.4) at 27°C. Uptake of myo-[3H]inositol was measured by adding 50 nM myo-[3H]inositol to cells at 37°C. At various time points, uptake of the label was terminated by pelleting 1.5 × 106 parasites by centrifugation (1,500 × g, 5 min, 4°C) and washing three times in ice-cold PBS. After resuspension of the pellet in 100 μL PBS, radioactivity was measured by scintillation counting using a Packard Tri-Carb 2100TR liquid scintillation analyzer (PerkinElmer Life Sciences). Aliquots of the parasite suspensions before centrifugation were used to determine the total amount of radioactivity in the assay.

Metabolic labeling of trypanosomes, lipid extraction, and TLC. Metabolic labeling of trypanosomes was performed as described before (27). Briefly, myo-[3H]inositol was added to bloodstream-form or procyclic trypanosomes at mid-log phase, and incubation was continued for 16 h. Cells were harvested by centrifugation at 1,750 × g for 10 min and washed with ice-cold Tris-buffered saline (10 mM Tris, 144 mM NaCl, pH 7.4) to remove unincorporated label, and bulk phospholipids were extracted twice with 10 ml chloroform:methanol (CM; 2:1 [vol/vol]). CM fractions were pooled, dried under nitrogen, and resuspended in a small volume of CM. Aliquots were treated with 6 μl PI-specific phospholipase C from Bacillus cereus (Thermo Scientific, Wohlen, Switzerland) for 60 min, as described elsewhere (28). Lipids were analyzed by thin-layer chromatography (TLC) on Silica Gel 60 plates (Merck) using solvent system 1, composed of chloroform:methanol:acetic acid:water (25:15:4:2 [vol/vol/vol/vol]). Appropriate lipid standards were run alongside the samples to be analyzed. Radioactivity was detected by scanning the air-dried plate with a radiisotope detector (Berthold Technologies, Regensdorf, Switzerland) and quantified using the RiaStar software provided by the manufacturer. For analysis of GPI precursors, bloodstream-form trypanosomes were labeled for 16 h with trace amounts of [3H]ethanolamine (28). After the cells were harvested and bulk lipids were extracted as described above, GPI lipids were extracted from the pellet using chloroform:methanol:water (10:10:3 [vol/vol/vol]) and partitioned between water
and butan-1-ol. [3H]-labeled GPI lipids in the butan-1-ol-rich upper phase were analyzed by TLC using solvent system 2, composed of chloroform:methanol:water (10:10:3 [vol/vol/vol]) (28). Radioactivity was detected as described above.

**Mass spectrometry and inositol analysis.** Total lipids for mass spectrometry analysis were extracted using a modification of the Bligh and Dyer method (30). Briefly, T. brucei bloodstream forms were collected at the mid-log phase, washed with PBS, resuspended in 100 µl of fresh PBS, and transferred to a glass tube. Chloroform:methanol (1:2 [vol/vol]) (375 ml) was added and subjected to vigorous vortex mixing for 10 to 15 min. The sample was made biphasic by the addition of 125 ml chloroform and 125 ml water, subjected to vortex mixing again, and centrifuged at 1,000 × g at room temperature (RT) for 5 min. The lower phase was dried under nitrogen and stored at 4°C. Total lipid extracts were dissolved in 15 ml of chloroform:methanol (1:2 [vol/vol])–15 ml of acetone:isopropanol:water (6:7:2 [vol/vol/vol]) and analyzed with a Abscetix 4000 QTrap, a triple-quadrupole mass spectrometer equipped with a nanoelectrospray source. Samples were delivered using a Nanomate interface in direct infusion mode (–125 nM). Lipid extracts were analyzed in both positive- and negative-ion modes using a capillary voltage of 1.25 kV. Tandem mass spectrometry (MS/MS) scanning (daughter, precursor, and neutral loss scans) was performed using nitrogen as the collision gas with collision energies of between 35 and 90 V. Each spectrum encompassed at least 50 repetitive scans. MS/MS spectra were obtained with collision energies as follows: 35 to 45 V, phosphatidylcholine-sphingomyelin (PC/SM) in positive-ion mode, parent-ion scanning of m/z 184: 35 to 55 V, PI in negative-ion mode, parent-ion scanning of m/z 241; 35 to 65 V, phosphatidyethanolamine (PE) in negative-ion mode, parent-ion scanning of m/z 196; 20 to 35 V, phosphatidylycerine (PS) in negative-ion mode, neutral loss scanning of m/z 87; and 40 to 90 V. MS/MS daughter ion scanning was performed with collision energies of between 35 and 90 V. Assignment of phospholipid species is based upon a combination of survey, daughter, precursor and neutral loss scans, as well previous assignments (31). The identity of phospholipid peaks was verified using the LIPID Metabolites And Pathways Strategy (LIPID MAPS) Nature Lipidomics Gateway (www.lipidmaps.org).

For the inositol analysis, bloodstream forms were collected and lipids were extracted as described above. An internal standard of D6-myoinositol was added to samples prior to hydrolysis by strong acid (6 M HCl, 110°C), derivatization with tetramethylsilane (TMS), and analysis by gas chromatography-mass spectrometry, as published elsewhere (32). myo-Inositol was quantified, and the means and standard deviations of the results of three separate analyses were determined.

**Microscopy.** For immunolocalization of HA-tagged TbHMIT, trypanosomes were cultured in the presence of tetracycline for 24 h to induce protein expression and processed as described previously (14). HA-tagged proteins were detected using monoclonal mouse anti-HA antibody (Covance, Munich, Germany) at a dilution of 1:250 in PBS for 1 h at room temperature. The Golgi complex was visualized by incubating fixed parasites for 1 h at room temperature with rabbit anti-TbGRASP antibody (kindly provided by G. Warren, University of Vienna) (used at a dilution of 1:1,000). Subsequently, the slides were washed three times with PBS and incubated with the corresponding secondary antibodies, Alexa Fluor 594 goat anti-mouse IgG and Alexa Fluor 488 goat anti-rabbit IgG (Invitrogen), at dilutions of 1:1,000 and 1:500, respectively, for 1 h at room temperature. Slides were washed three times with PBS and mounted using Vectashield containing 4',6-diamidino-2-phenylindol (DAPI; Vector Laboratories, Burlingame, CA, USA). Fluorescence microscopy was performed on a Leica AF6000 microscope (Leica Microsystems, Heerbrugg, Switzerland), using the software provided by the manufacturer.

**Substrate specificity of TbHMIT.** *Xenopus laevis* oocytes were prepared, injected with Tb927.11.5350 cRNA, and depolished as described previously (14). Electrophysiological experiments were performed as described before (14). myo-Inositol, epimeric myo-inositol, diso-inositol, protosterol, chiro-inositol, 1,2-inositol, 1,2-inositol, 1,4-inositol, 1,2-inositol, 1,2-inositol, 1,4-inositol, 1,4-inositol, 1,4-inositol, 1,4-inositol, 1,4-inositol, 1,4-inositol.
FIG 1 Essentiality of TbHMIT in T. brucei bloodstream forms. (A) Growth of T. brucei bloodstream-form RNAi parasites. Expression of TbHMIT in T. brucei bloodstream forms was downregulated by tetracycline (Tet)-inducible RNAi, and growth of trypanosomes was monitored for 5 days. Data points represent cumulative numbers of RNAi cells incubated in the absence (filled symbols) or presence (open symbols) of tetracycline and correspond to mean values ± standard deviations of the results of three experiments performed using clone A3. The inset shows a Northern blot analysis of total RNA extracted from trypanosomes after 2 days of incubation in the absence (−) or presence (+) of tetracycline and probed with [32P]-labeled oligonucleotides used as insertions for the respective stem-loop vectors (top panel); rRNA was stained with ethidium bromide and used as loading control (bottom panel). Two other RNAi clones showed similar growth defects and reductions in TbHMIT mRNA levels upon treatment with tetracycline. (B) \( \text{myo-}\text{inositol} \) uptake by T. brucei bloodstream forms. After 2 days of incubation in the absence (filled symbols) or presence (open symbols) of tetracycline, RNAi parasites (clone A3) were washed and subsequently incubated with trace amounts of \( \text{myo-}\text{[3H]inositol} \) (50 nM final concentration). After the indicated times, parasites were washed, and the amount of radioactivity in the cell pellet was measured. Uptake of \( \text{myo-}\text{[3H]inositol} \) at each time point was calculated and plotted as a function of incubation time. Data points represent mean values ± standard deviations of the results of triplicate determinations from three independent experiments. (C) Analysis of \( \text{myo-}\text{[3H]inositol} \)-labeled lipids. T. brucei bloodstream-form TbHMIT RNAi cells (clone A3) were incubated in the absence or presence of tetracycline for 3 days. During the last 16 h of incubation, parasites (2 \( \times \) 10\(^6\) cells) were labeled with 25 \( \mu \)Ci of \( \text{myo-}\text{[3H]inositol} \). \( \text{[3H]}-\text{labeled lipids} \) were extracted from parasites incubated in the absence (top panel) or presence (middle panel) of tetracycline to downregulate TbHMIT, separated by one-dimensional TLC using solvent system 1, and visualized by scanning the plate (14). Extracts from cell equivalents were applied. The bottom panel represents extracts from T. brucei procyclic forms run on the same TLC plate for comparison, indicating the migration of \( \text{[3H]inositol phosphorylceramide} \) ([3H]IPC) and \( \text{[3H]phosphatidylinositol} \) ([3H]PI). The vertical lines indicate the site of sample application (left line) and the migration of the solvent front (right line), respectively. (D) Analysis of \( \text{[3H]ethanolamine-labeled GPI lipids} \). Trypanosomes were cultured in the absence (top panel) or presence (middle panel) of tetracycline for 3 days to down regulate TbHMIT and were then labeled during 16 h with \( \text{[3H]ethanolamine} \). \( \text{[3H]GPI lipids} \) were extracted and analyzed by TLC using solvent system 2. The major DPI lipids, designated P2 and P3, were identified based on published R\(_f\) values (28) and the migration of \( \text{[3H]}\Pi \), i.e., the major DPI precursor lipid in T. brucei procyclic forms, run on the same plate (bottom panel). The migration of residual \( \text{[3H]DE} \) is also indicated. The left and right vertical lines indicate the site of sample application and the migration of the solvent front, respectively.
phospholipid profile of wild-type parasites (Fig. 2A; see reference 31 for comparison). These include the major plasmalogen (alk-kenyl acyl) phosphatidylethanolamine (PE) species at 727 m/z and PI molecular species at 865, 887, and 913 m/z, corresponding to 36:0, 38:3, and 40:4 PI. Upon induction of the RNAi cells for 48 h with tetracycline (Fig. 2B), the intensity of all PI molecular species clearly diminished compared to that seen with uninduced cells, while the plasmalogen PE was still the dominant species (compare Fig. 2A to panel B). To confirm the decrease in the level of the PI molecular species whose levels increased or decreased after RNAi performed against TbHMIT compared to noninduced cells (A).

In contrast, in extracts from parasites after 48 h of tetracycline induction, the intensities of these fragments were drastically decreased (see Fig. S2B). In addition, the amounts of inositol-containing phospholipids were quantified by measuring the myo-inositol contents in lipid extracts from control and induced cells and normalizing to cell numbers. The results show that RNAi cells after downregulation of TbHMIT only had 79.5% ± 2.5% myo-inositol-containing lipids compared to noninduced cells (100% ± 4% [mean values ± standard deviations of the results from three independent experiments]).

In the induced cells, apart from the reduction in PI species, the intensities of two phospholipid species at 762 and 795 m/z had increased (Fig. 2B). These two species were subjected to fragment-
tation (see Fig. S3A and B in the supplemental material, respectively) and identified as PS 34:0 and phosphatidylglycerol (PG) 38:5, respectively. The only other obvious change in the phospholipids was observed in positive-ion mode (see Fig. S4 in the supplemental material), which shows the choline-phosphate-containing species of phosphatidylcholine (PC) and sphingomyelin (SM). The level of the species at 794 m/z, representing PC 40:4, was clearly decreased in cells after TbHMIT RNAi compared to that in uninduced cells (compare Fig. S4A with panel B). The cells were obviously trying to compensate for a lack of PI, but the reasons for these specific changes, other than maintenance of the correct membrane fluidity for normal cellular functions, are unknown.

**Localization of TbHMIT in *T. brucei* bloodstream forms.** Immunofluorescence microscopy revealed that ectopically expressed HA-tagged TbHMIT in *T. brucei* bloodstream forms localized to a distinct intracellular structure located between the nucleus and the kinetoplast (Fig. 3). The signal completely colocalized with TbGRASP (Fig. 3), a marker for the Golgi complex in *T. brucei* (34). These findings are in line with a previous report showing that TbHMIT localizes to the Golgi complex in *T. brucei* procyclic forms (14).

The Golgi complex localization of TbHMIT in *T. brucei* bloodstream forms is noteworthy. It is believed that PI synthesis occurs on the cytosolic side of the ER (10, 35). It should be noted, however, that in several studies in plants (11), yeast (36), protozoa (13), and mammals (12), PI synthase was found to localize not only in the ER but also in close proximity to or in the Golgi complex. As has been demonstrated in *T. brucei* (13, 14), PI synthesis in the ER and PI synthesis in Golgi complex may serve different purposes. PI production in the ER is required for GPI synthesis, while PI produced in the Golgi complex provides bulk PI for membrane formation. Based on the localization of TbHMIT in the Golgi complex, these results suggest that the last step in PI synthesis, i.e., the attachment of myo-inositol to cytidine diphosphate-diacylglycerol (CDP-DAG), may occur in the lumen of the Golgi complex. Interestingly, a recent report showed that *T. brucei* CDP-DAG synthase localizes to the ER and the Golgi complex (37), which would be consistent with PI synthesis taking place in the lumen of the ER and the Golgi complex. In addition, results of studies using membrane topology prediction programs TMHMM (38) and Phobius (39) indicate that the active site of *T. brucei* PI synthase is on the luminal side of the ER. On the basis of these observations, we propose the following model for compartmentalization of PI synthesis in *T. brucei* (Fig. 4). In procyclic forms, myo-inositol is taken up from the environment via plasma membrane-localized TbHMIT; in bloodstream forms, myo-inositol

![FIG 3](image_url) Localization of TbHMIT in *T. brucei* bloodstream forms. Trypanosomes cultured in the presence of tetracycline for 24 h to induce expression of HA-tagged TbHMIT were washed, allowed to settle onto microscope slides, fixed with paraformaldehyde, and permeabilized with Triton X-100. TbHMIT was detected using anti-HA antibody (first panel), whereas the Golgi complex was stained with anti-GRASP antibody (second panel). The third panel shows an overlay of panels A and B, with DNA stained with DAPI. The corresponding differential interference contrast (DIC) micrograph is shown in the fourth panel.

![FIG 4](image_url) Schematic of compartmentalized myo-inositol metabolism in *T. brucei*. For details, see main text. glc-6-P, glucose-6-phosphate; GPI, glycosylphosphatidylinositol; ins, inositol; ins-3-P, inositol-3-phosphate; IPC, inositolphosphoryl ceramide; PI, phosphatidylinositol; TbHMIT, *T. brucei* H+-coupled myo-inositol transporter; GlcNAc-PI, N-acetylglucosaminyl phosphatidylinositol; GlcN-PI, glucosaminyl phosphatidylinositol; TbPIS, *T. brucei* PI synthase; TbSLS1, *T. brucei* sphingolipid synthase 1; TbINO1, *T. brucei* 1-β-myoinositol-3-phosphate synthase; TbIMPase, *T. brucei* inositol monophosphatase.
uptake likely occurs via a different transporter. Cytosolic myo-inositol is then transported into the Golgi complex via TbHMIT, where it is used for bulk PI and, subsequently, IPC synthesis. In contrast, de novo synthesis of myo-inositol starts with the cytosolic conversion of glucose-6-phosphate to inositol-3-phosphate, which in turn is transported into the ER via an unknown transporter. After dephosphorylation, newly synthesized myo-inositol is used for PI formation by PI synthase. Subsequently, PI is translocated from the luminal to the cytosolic face of the ER, where it is used to initiate GPI synthesis. Recently, a similar mechanism has also been proposed to occur in the intraerythrocytic stage of the malaria parasite, Plasmodium falciparum (40). In addition, it is worth mentioning that the two PI synthase isoforms described in Arabidopsis thaliana show different substrate specificities (11), suggesting that two pools of PI may exist and that these may enter alternative routes of metabolism. Furthermore, a Golgi complex localization of PI synthase (11, 12, 36) and HMIT (8) has also been documented in other cells. Thus, we propose that PI formation and metabolism may be similarly compartmentalized in other eukaryotes as well.

Our data using exogenously added myo-[3H]inositol showed that, after depletion of TbHMIT by RNAi treatment, uptake of myo-[3H]inositol into bloodstream-form parasites still occurred, albeit at a clearly reduced level (Fig. 1B), and yet the formation of [3H]PI was blocked (Fig. 1C). Together with the observed localization of TbHMIT in the Golgi complex, these data suggest that myo-[3H]inositol is taken up in T. brucei bloodstream forms via a different transporter, located in the plasma membrane, and subsequently remains metabolically inactive because of the lack of the Golgi complex-localized TbHMIT, preventing entry of (cytosolic) myo-[3H]inositol into the Golgi complex for [3H]PI production. These results differ slightly from our previous findings in T. brucei procyclic forms (14), where treatment with RNAi against TbHMIT blocked not only [3H]PI formation but also myo-[3H]inositol uptake into parasites. Interestingly, in procyclic forms, TbHMIT is not only present in the Golgi complex but can also be detected in the plasma membrane, mediating myo-[3H]inositol uptake into the cell. We are currently addressing these differences between the bloodstream and procyclic forms with further experiments.

Collectively, these results demonstrate that expression of TbHMIT is essential for normal growth of T. brucei bloodstream forms in culture and that it is involved in myo-inositol transport into the Golgi complex for PI formation within the lumen of the Golgi complex.

**Substrate specificity of TbHMIT.** Perfusion of TbHMIT-expressing Xenopus laevis oocytes with 200 μM myo-inositol resulted in inward currents of 20 ± 1 nA (mean value ± standard deviation, using 13 oocytes from two independent batches). The same concentration of myo-inositol did not elicit any currents in water-injected control oocytes. These results are consistent with a previous report (14). To identify other potential substrates, oocytes expressing TbHMIT were exposed to a series of commercially available compounds that are structurally related to myo-inositol (Fig. 5). The currents elicited by these compounds (each applied at a concentration of 200 μM) were compared to those obtained with myo-inositol (Fig. 6). Interestingly, we found that two quercitol isomers, epimeric and vibo-quercitol, elicited currents comparable to those induced by myo-inositol (76% ± 9% and 83% ± 3%, respectively, of the myo-inositol current). In contrast, <5% of the current obtained with myo-inositol was elicited by another quercitol isomer, proto-quercitol (Fig. 6). Quercitols comprise a group of 6-C-containing polyols which, compared to the group of inositol isomers, lack one hydroxyl group (Fig. 5) and, in the case of proto-quercitol, contain a C1 epimerization. In addition, low currents (15% to 25% of the myo-inositol current) were also obtained with scyllo-inositol, epi-inositol, 1L-chiro-inositol, and phytic acid (myo-inositol-1,2,3,4,5,6-hexakisphosphate). In contrast, the uracils induced by myco-inositol, allo-inositol, 1L-chiro-inositol, 1L-epi-2-inososide (2L-2,3,4,6/5-pentahydroxyhexanone), 1-quebrachitol (2-O-methyl-1-chiro-inositol), d-pinitol (3-O-methyl-d-chiro-inositol), N-00601 [(1R,4S)-6-methoxyhexahexanone-1,2,3,4,5-pentol], N-05350 [(1R,3S)-6-methoxyhexahexanone-1,2,3,4,5-pentol], and d-inositol-3-phosphate represented <5% of the control myo-inositol current (Fig. 6).

These results, together with our previous findings showing that TbHMIT lacks transport activity for C5 (xylitol) and other C6 (mannitol and sorbitol) polyols and for sugars (glucose and mannose) (14), provide an insight into the selectivity of the transporter in terms of functional groups and stereoselectivity. Among the different analogs tested, deleting a single hydroxyl group at the 1 position [as in (−)-vibo-quercitol] or the 6 position [as in (+)-...
6-methoxycyclohexane-1,2,3,4,5-pentol. Inversion of stereochemistry at a single hydroxyl group relative to myo-inositol ring only, such as the removal of a hydroxyl group at position 1 or position 6 or the inversion of stereochemistry at a single hydroxyl group relative to myo-inositol, but no additional modifications. Interestingly, TbHMIT (14) and its orthologs from T. cruzi (20) and Leishmania parasites (18, 41) show no transport activity for monosaccharides. This is in marked contrast to all other members of the GLUT/SLC2A family, including the intracellularly located members of subclass III, which transport many different monosaccharides (1, 5, 6). In addition, HMIT’s transport specificity is also different from that of the SMITs, which transport both myo-inositol and monosaccharides (2, 3).

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