Toxoplasma gondii is a globally prevalent parasitic protist. It is well-known for its ability to infect almost all nucleated vertebrate cells, which is reflected by its unique metabolic architecture. Its fast-growing tachyzoite stage catabolizes glucose via glycolysis to yield L-lactate as a major by-product that must be exported from the cell to prevent toxicity; the underlying mechanism remains to be elucidated, however. Herein, we report three formate–nitrite transporter (FNT)–type monocarboxylate/proton symporters located in the plasma membrane of the T. gondii tachyzoite stage. We observed that all three proteins transport both L-lactate and formate in a pH-dependent manner and are inhibited by 2-hydroxy-chromanones (a class of small synthetic molecules). We also show that these compounds pharmacologically inhibit T. gondii growth. Using a chemical biology approach, we identified the critical residues in the substrate-selectivity region of the parasite transporters that determine differential specificity and sensitivity toward both substrates and inhibitors. Our findings further indicate that substrate specificity in FNT family proteins from T. gondii has evolved such that a functional repurposing of prokaryotic-type transporters helps fulfill a critical metabolic role in a clinically important parasitic protist. In summary, we have identified and characterized the lactate transporters of T. gondii and have shown that compounds blocking the FNTs in this parasite can inhibit its growth, suggesting that these transporters could have utility as potential drug targets.
This work focuses on the biochemical characterization, pharmacological inhibition, and mechanism of FNT proteins to demonstrate their physiological importance during the lytic cycle of *T. gondii* in mammalian cells.

**Results**

*T. gondii* harbors three FNTs on its surface

We started by searching the annotated genome of *T. gondii* for genes encoding putative monocarboxylate transporters. This search did not reveal the presence of MCT-type transporters (22); instead, we identified three FNT-encoding genes (ToxoDB IDs: TGGT1_209800, TGGT1_292110, and TGGT1_229170). The first one, TGGT1_209800, located on the chromosome Ib, appeared intron-less; we refer to the gene product of 412 amino acids (aa) as *TgFNT1* (44.3 kDa) herein. The second, TGGT1_292110, with six exons on the chromosome IX, encodes for *TgFNT2* with 463 aa (50.8 kDa), and the third TGGT1_229170, with five exons present on the chromosome VIII, translates into *TgFNT3* with 501 aa (53.3 kDa). The three *TgFNT* proteins display typical signature motifs, *e.g.* in the L2 and L5 loop regions preceding the two half-helices, and a six-transmembrane topology with both termini facing intracellularly shared by all FNT family members (Fig. 1A) (23). Quite notably, however, the *TgFNTs* harbor extended C termini that are about 115 to 225 aa longer than the previously reported FNTs. *TgFNT2* also contains an additional stretch of 70 aa at the N terminus. The sequences of the *TgFNT1–3* termini appear to be nonhomologous, whereas the protein cores, *i.e.*, the transmembrane domains plus connecting loops, exhibit an overall 61–66% sequence identity and 75–78% similarity with each other. Finally, we cloned and verified their ORF sequences using the mRNA isolated from tachyzoites of *T. gondii*.

Next, we generated full-length ectopic expression constructs of *TgFNT1–3* carrying a hemagglutinin epitope (HA) at the respective C terminus for immunofluorescent localization in tachyzoites. The constructs were transfected into fresh extracellular parasites followed by drug selection to make stable transgenic lines expressing *TgFNT1–3* under the control of *Tg* GRA1 elements targeted at the UPRT locus. Immunofluorescence studies revealed that all *TgFNTs* were expressed at the surface of intracellular tachyzoites (Fig. 1B). Immunostaining of *TgFNT1–3* using an anti-HA antibody indicated a peripheral transgenic lines expressing *TgFNT-HA* proteins. The ensuing strains expressed *TgFNT1–3* under the control of *Tg* GRA1 elements targeted at the UPRT locus. Immunofluorescence studies revealed that all *TgFNTs* were expressed at the surface of intracellular tachyzoites (Fig. 1B). Immunostaining of *TgFNT1–3* using an anti-HA antibody indicated a peripheral expression of all transporters, as judged by their colocalization with a *bona fide* marker of the inner membrane complex (*Tg* Gap45; Fig. S1A), located just underneath the parasite’s plasma lemma (24). To determine the exact site of protein location, we treated extracellular parasites with bacterial \(\alpha\)-toxin to split the plasma membrane from the inner membrane complex and performed co-staining of *TgFNT1–3* with a plasma membrane marker (*Tg* Sag1), and with *Tg* Gap45 (Fig. S1A). The HA signal overlapped clearly with *Tg* Sag1, but not with *Tg* Gap45, confirming the occurrence of all three *TgFNT-HA* constructs at the parasite surface.

In further assays, we assessed the orientation of *TgFNT1–3* in the plasma membrane by immunostaining detergent-permeabilized and nonpermeabilized parasites (Fig. S1B). The presence of HA staining only in permeabilized parasites confirmed that the C termini of *TgFNT1–3* do face the cytosol, as predicted above (Fig. 1A). We also examined the endogenous expression profiles of these proteins based on the MS datasets in ToxoDB. Interestingly, only *TgFNT1* and *TgFNT2* are expressed at the protein level, whereas *TgFNT3* is not detectable in tachyzoites of the type I strain (*i.e.*, RH strain used herein). The latter is however readily detectable in the cyst-forming type II strain (ME49 strain). These results together demonstrate the occurrence of three distinct FNTs located in the plasma membrane of *T. gondii*, of which only *TgFNT1* and *TgFNT2* are naturally expressed during the lytic cycle of the type I strain.

*TgFNTs* transport \(L\)-lactate as well as formate in a pH-dependent manner

To determine substrate specificity and kinetics of *TgFNT* transport, we carried out biophysical import assays using radio-
labeled formate and lactate in an established Saccharomyces cerevisiae strain (W303–1A jen1/H9004 ady2/H9004) lacking endogenous monocarboxylate transporters (7, 8, 25). Transport was monitored over a short time scale (second to minute) to prohibit any metabolic and phenotypic effects. Under these conditions, uptake of radioisotope is directly linked to specific substrate transport via the recombinantly expressed TgFNT. Others and we have shown before that FNT transport is bidirectional (6, 8).

Initially, we expressed full-length ORFs of TgFNT1–3. It turned out however that the resulting proteins were highly unstable in the yeast system, appearing as multiple fragments in immunoblots (Fig. 2A). Consequently, our import assays with 14C-radioabeled formate or l-lactate as substrates yielded erratic results. Earlier studies on the structure and transport mechanism of the FNTs suggested that despite a probably flexible N-terminal domain, the core protein assumes a rather rigid, channel-like conformation (25). Hence, we presumed that protein-stabilizing deletions in TgFNTs might be tolerated without affecting their basic transport function and substrate selectivity. Indeed, shortening of the termini to the lengths indicated in Fig. 1 resulted in reasonably stable expression of all three transporters in yeast (Fig. 2B), which enabled a detailed functional characterization.

The truncated TgFNT1–3 proteins were functional transporting formate and L-lactate. The three transporters produced similar substrate uptake curves for formate and L-lactate in 1 mM inward-directed gradients at pH 6.8 (Fig. 2C), which is comparable in magnitude to the FNT from P. falciparum (8). The initial import rates of all TgFNTs, as determined by exponential fittings, indicated nearly equal preference for the formate molecule (45 Da) and for the twice as large L-lactate (89 Da) (Fig. 2D). In the steady state of transport, lactate accumulated at 50–60% higher levels in the yeast cells than formate. Earlier, we established that the transmembrane proton gradient is a driving force for monocarboxylate transport via FNT-type proteins (25). Accordingly, increasing acidity of the external buffer increases import rates of monocarboxylates. Hence, we monitored lactate import by TgFNT1–3 at a lowered pH of 4.8, i.e. by two logarithmic units, which resulted in about 4-fold higher transport rates (Fig. 3, A and B). In summary, our data reveal the encoding of three functional FNT-type monocarboxylate/H+ symporters in T. gondii.
nesulphonate (pCMBS) to the TgFNT assays (Fig. 4B). As a control, we tested PfFNT, which is insensitive to pCMBS because of absence of cysteines in the transport path (8). Indeed, pCMBS fully inhibited TgFNT2 and TgFNT3, whereas TgFNT1 remained entirely functional (Fig. 4B). The data imply that, unlike TgFNT1, the filter regions in TgFNT2 and TgFNT3 expose pCMBS-accessible cysteines.

Assuming that pCMBS might be too voluminous to approach Cys-46 in TgFNT1, we switched to a smaller cysteine-reactive compound, i.e. methyl methanethiosulphonate (MMTS), which transfers a thiomethyl moiety to cysteines forming a disulfide bridge (Fig. 4C). PfFNT with a cysteine-less Ψ/K filter was again included as a negative control. The observed slightly reduced transport rates of PfFNT in the presence of MMTS are probably because of competition with the substrate, because MMTS is similar in size as formate and L-lactate, and was present at the same concentration. Notably however, MMTS treatment fully inhibited formate and lactate transport of TgFNT1, showing that the cysteine modification can indeed block the selectivity filter (Fig. 4C). Remarkably, the effect of MMTS on TgFNT2 and TgFNT3 was rather distinguished: both excluded L-lactate yet allowed the formate to pass (Fig. 4C). Apparently, the modification by MMTS led to an intermediate filter diameter, which then exhibited the selectivity profile similar to the prokaryotic FNT proteins. Collectively, these experiments not only underline the mechanism of FNT functioning, but also reveal a divergent evolution of TgFNT1–3, which likely serves special purposes during the natural life cycle of T. gondii.

**Druglike inhibitors can block L-lactate transport by TgFNT1–3**

Our earlier study has described druglike, high-affinity inhibitors of PfFNT (26). By selection of a PfFNT G107S resistance mutant, we located the drug-binding site to the intracellular substrate-entry region of PfFNT. In this study, we chose the original screening hit compound from the *malaria box*, MMV007839, along with four other compounds with the same pharmacophore but different lipophilicity from our own collection for testing on TgFNT1–3 (Fig. 5A). BH-296 lacks a phenolic hydroxyl moiety maintaining the molecule in the linear, active form, whereas BH-317, BH-326, and BH-388 can undergo the same repeated conversion between the active vinylogous acid form and a cyclic hemiketal form, as we showed before for MMV007839 (26). All compounds ablated L-lactate transport by TgFNT1–3 (Fig. 5B and Fig. S2). Consistent with their deviant substrate selectivity filter, TgFNT1 required double-digit micromolar concentrations for half-maximal inhibition, whereas TgFNT2 and TgFNT3 were more susceptible yielding IC_{50} values even in the nanomolar range (Table 1).

To test whether the binding site of the inhibitors to TgFNT1–3 corresponds to PfFNT, we introduced single mutations in TgFNT1 (G93S), TgFNT2 (G166S), and TgFNT3 (G93S), mimicking the resistance mutation observed in the PfFNT protein. The altered proteins were expressed in yeast at the same levels as the corresponding nonmutated variants (Fig. 2B), albeit the transport rates for lactate were reduced by about half (Fig. 6A). In each case, the glycine-to-serine mutation shifted the IC_{50} value of MMV007839 toward a higher value, i.e. 4× in TgFNT1 G93S (108.0 ± 8.1 μM), 46× in TgFNT2 G166S...
Inhibition of TgFNT1–3 impairs the lytic cycle

To determine the anti-parasitic effect of the TgFNT1–3 inhibitors, we set up plaque assays, which recapitulate the successive lytic cycles of tachyzoites, and thus the overall growth fitness of the parasite in cultures (27). To begin with, we analyzed concentrations of 1 μM and 10 μM for all compounds (Fig. 7A); higher concentrations were not tolerated by the host cells. The parasite growth appeared normal at 1 μM, but was differentially inhibited at 10 μM. As shown in Fig. 7B, BH-296 and BH-388 resulting in the smallest average plaque size, were most potent among all. Parasite growth was inhibited by about 35% in the presence of BH-296 and BH-388, whereas BH-312 and BH-326 exerted a modest 10% inhibition.

We also measured the replication rates in the presence of these chemicals using a strain that expresses the yellow fluorescent protein (YFP) in its cytosol (28). Because the YFP signal correlates with the replication rate of tachyzoites, it provides a rapid and accurate measurement of the parasite growth (Fig. 7C). Similar to the plaque assays, BH-296 and BH-388 were most effective against the parasite (Fig. 7C; Table 1), whereas the other three compounds were less active, as judged by a much higher IC50 of the growth inhibition (Fig. S3). Irrespective of their distinguishable IC50 values in the transport assays, the IC50 values of growth inhibition fall into a similar range. To conclude, these results suggest that FNT inhibitors can block the asexual reproduction of T. gondii, which resonates with our biochemical assays.

Discussion

Plasmodium and Toxoplasma are long known to be voracious consumers of glucose during their acute infection phase

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**Figure 5. Inhibition of TgFNT1–3 L-lactate transport by small molecules in yeast.** A, the chemical structures of the compounds varying in their aromatic substituent. MMV007839, BH-317, BH-326, and BH-388 undergo repeated conversions between a cyclic hemiketal transport form and a linear vinylogous acid active form. BH-296 lacks a phenolic hydroxyl moiety (arrow) maintaining the molecule in the linear form. B, dose-dependent inhibition of TgFNT1–3 by MMV007839 (pH 6.8, 1 mM inward gradient). Three biological assays were done, each with triplicate data points. Error bars denote S.E.

**Table 1**

| Inhibitors | TgFNT1 (yeast) | TgFNT2 (yeast) | TgFNT3 (yeast) | Parasite growth |
|------------|---------------|---------------|---------------|----------------|
| MMV007839  | 28.8 ± 10.5   | 1.94 ± 1.05   | 0.55 ± 0.08   | 47.1 ± 33.6    |
| BH-296     | 56.0 ± 3.3    | 6.14 ± 4.59   | 3.00 ± 1.54   | 9.8 ± 1.8      |
| BH-317     | 13.0 ± 2.4    | 0.92 ± 0.27   | 0.65 ± 0.08   | 15.8 ± 3.1     |
| BH-326     | 22.3 ± 2.4    | 0.57 ± 0.13   | 0.59 ± 0.08   | 23.4 ± 8.2     |
| BH-388     | 24.0 ± 3.0    | 4.78 ± 0.84   | 2.59 ± 0.08   | 5.8 ± 2.4      |

(88.8 ± 13.4 μM), and 149× in TgFNT3 G93S (82.0 ± 7.9 μM) (Fig. 6B). Yet again, TgFNT1 was quantitatively less susceptible to a mutation-driven change in the IC50 value.
Glucose is catabolized primarily via glycolysis, resulting in a substantial production of \( \gamma \)-lactate as a key by-product. Both parasites release the excess of \( \gamma \)-lactate to the external milieu to sustain their proliferation. Previously, we have characterized PfFNT, which is the sole \( \gamma \)-lactate transporter expressed in \( P. \) falciparum. Herein, we show the presence of multiple FNT isoforms to perform the same task in tachyzoites of \( T. \) gondii. A functional link between glycolysis and FNT proteins is corroborated by concordant transcript profiles of \( TgFNT1–3 \), the glucose transporter, and the enzymes of glycolysis during the lytic cycle of tachyzoites (Fig. S4). An apparent redundancy in \( \gamma \)-lactate transport is also in accordance with the exceptional metabolic plasticity observed in the central carbon metabolism of tachyzoites (16, 18, 19). FNTs discriminate between different monocarboxylate substrates by size selection in the \( \Phi/K \) filter region (7). In view of an almost identical filter layout of \( TgFNT1–3 \) the question arises why \( T. \) gondii expresses three isoforms. Besides mere redundancy, it is plausible that they serve different parasite stages during the natural life cycle. Glucose import is critical for \( P. \) falciparum species (17, 18), whereas \( T. \) gondii tachyzoites can survive without it, albeit with a modest growth defect (13, 16). In accord with \( PfFNT \), inhibition of the singular hexose transporter in \( P. \) falciparum (\( PfHT \)) kills the parasite (17, 29), validating both \( PfHT \) and \( PfFNT \) as antimalarial targets. It is indeed surprising to see a lethal phenotype in tachyzoites upon inhibition of \( TgFNT \)s in the micromolar range of these compounds. Notwithstanding putative off-target effects, the data suggest a critical role of lactate transport, which remains to be confirmed by gene knockout studies.

The pH-dependence of \( TgFNT1–3 \) reported herein is comparable with \( PfFNT \) (8), suggesting that both parasites employ the same mechanism to transport lactate. The underlying “dielectric slide” mechanism is based on the electrostatic attraction of the deprotonated substrate anion into the FNT vestibule and subsequent neutralization by protonation allowing for passage through two hydrophobic constriction sites in the protein (25, 30). Transport of the comparatively large \( \gamma \)-lactate molecule via \( TgFNT1–3 \) is facilitated by a substrate selectivity filter, which is wider than that of bacterial formate–nitrite transporters. The \( \Phi/K \) selectivity filter with its size exclusion principle and the usual FNT fold are highly reminiscent of the aquaporin (AQP) water and solute channels (31, 32). Six transmembrane helices in conjunction with two half-helices fold into one transduction unit (Fig. 1A), which oligomerizes to form a homomeric FNT pentamer. AQPs with their distinct primary sequences form homotetramers, yet the FNT and AQP protomers can be superimposed with only small deviations (23). The access path from the extracellular side to the selectivity filter appears deeper in \( TgFNT1 \) compared with \( pCMBS \)-sensitive AQPs that carry a respective cysteine residue (33). On the other hand, it is narrower compared with \( TgFNT2 \) and \( TgFNT3 \), because the organomercurial did not reach the cysteine at the filter site of \( TgFNT1 \), leaving its functionality unaffected (Fig. 4B). In support of this notion, our structure models of \( TgFNT2 \) and \( TgFNT3 \) show a 20% larger section area in the \( \Phi/K \) region than \( TgFNT1 \) (Fig. 4A). Cysteine-modification by MMTS provides direct evidence for the difference in width

![Figure 6](image-url)  
**Figure 6.** Functionality and inhibition of \( TgFNT \) glycine-to-serine mutants in yeast. \( A \), a transport of \( \gamma \)-lactate by the \( TgFNT \) mutants (pH 6.8, 1 mM inward gradient) compared with the nonmutated proteins (dashed lines). \( B \), glycine-to-serine mutations in \( TgFNT1–3 \) shift the MMV07839 inhibition curves to higher concentrations (dashed lines represent the positions of curves with nonmutated \( TgFNTs \) from Fig. S5A). The experiments were carried out three times, each in triplicates. Error bars denote S.E.

![Figure 7](image-url)  
**Figure 7.** \( TgFNT \) inhibitors can impair the tachyzoite growth. \( A \), overall growth fitness of tachyzoites, as judged by plaque assays. Plaques form after parasite-inflicted lysis in the host cell monolayer during the lytic cycles of tachyzoites. The plaque size correlates with the parasite fitness. Parasites were treated with the carrier solvent DMSO (–) or with \( TgFNT \) inhibitors at 1 mM and 10 \( \mu \)M. \( B \), plaque size determination by image analysis at 1 \( \mu \)M (open bars) and 10 \( \mu \)M (closed bars) inhibitor concentration. Asterisks label significant differences (p < 0.05; Student’s t test); \( C \), effect of \( TgFNT \) inhibitors (100 \( \mu \)M, 10 \( \mu \)M, and 1 \( \mu \)M) on the parasite replication as determined by fluorescence intensity of YFP-expressing tachyzoites. The compound color code is indicated above the graphs. Pyrithymamine (1 \( \mu \)M in all plots) and DMSO-treated parasites (–) served as the positive and negative controls, respectively. The results show three independent assays, each done in triplicates (± S.E.).
of the selectivity filters in TgFNT1 and TgFNT2/3 proteins. Whereas the former was blocked upon addition of a small thiomethyl moiety, the latter two tolerated the chemical modification, albeit with the restriction of smaller transport substrates.

Similarly, our inhibitor compounds, which block TgFNT1–3 at the intracellular entrance of the transport path, were less efficient in accessing and binding TgFNT1 than TgFNT2 or TgFNT3. As a result, the IC_{50} values turned out to be considerably higher for TgFNT1. IC_{50} values obtained in the in vitro cultures of T. gondii tachyzoites were in a similar range with TgFNT1 expressed in yeast, suggesting it to be the major target of inhibitors used here. However, it should be noted that the compounds must enter the parasite cytosol to reach the TgFNT-binding site. This requires the inhibitors to pass the plasma membranes of the host cell and parasites by diffusion or active transport. Further, the compounds must exhibit a certain metabolic stability inside the cells to maintain their chemical structure for binding to the TgFNTs. Our prior observations in the Plasmodium system show that even small structural differences of the inhibitor compounds, e.g., the presence or absence of a methoxy moiety, can have major effects on their potency in the in vitro parasite culture despite an equal activity in the yeast system (26). Therefore, phenotypic drug screenings should stand a better chance to identify compounds with good efficiency against living parasites than assays with isolated target proteins.

In summary, this study has identified and characterized the previously unknown lactate transporters of T. gondii. Moreover, we showed that blockers of TgFNTs can inhibit the parasite growth, rendering them potential novel drug targets. The work advocates further studies on the comparative metabolic importance of three TgFNT isoforms.

**Experimental procedures**

**DNA sequences and protein analysis**

DNA and protein sequences were derived from ToxoDB and the NCBI databases. Protein structure data were derived from RCSB data bank. Sequence alignments and logos were set using TeXshade and TeXtopo (34, 35). TgFNT structure models were generated with SWISS-MODEL (36) using S. typhimurium FocA (PDB ID: 3Q7K) (37) as a template. Visualization of protein structures was done using the PyMOL Molecular Graphics System, Schrödinger, LLC.

**Expression of TgFNT1–3 in yeast**

The ORF sequences of TgFNT1 (TGGT1_209800), TgFNT2 (TGGT1_292110), and TgFNT3 (TGGT1_229170) were codon-optimized, synthesized (GenScript), and cloned into the yeast expression vector pDR196 using SpeI and SalI (46). The truncated TgFNT versions were generated by PCR using the primers (Life Technologies) indicated in Table S1 and cloned into the WT constructs. Point mutations were introduced by site-directed mutagenesis using the QuikChange protocol (Stratagene) (primers with the respective nucleotide exchanges are shown in Table S1). All constructs were sequenced for accuracy prior to further use. Transformation of W303-1A (MATa, can1-100, ade2-loc, his3-11-15, leu2-3,-112, trp1-1-1, ura3-1, jen1::kanMX4, ade2::hphMX4) yeast cells, kindly provided by M. Casal (38), was done using the lithium acetate/single stranded carrier DNA/PEG procedure (39).Transformed cells were grown at 30 °C in selective media (S.D.) containing adenine, histidine, leucine, tryptophan, and 2% (w/v) glucose in the absence of uracil.

**Western blotting**

A 100 ml yeast liquid culture in selective media (S.D.) was prepared and cells were collected at an A_{600} of 1 (1000 g, 5 min, 4 °C), washed with 25 ml water and 10 ml extraction buffer (5 mM EDTA, 25 mM Tris, pH 7.5), pelleted, and resuspended in 0.5 ml of extraction buffer. Cells were disrupted by adding 0.5 g acid-washed glass beads (diameter 0.5 mm; Sigma-Aldrich) followed by 10 cycles of vortexing (30 s each) and intermittent cooling on ice for 1 min. Cell lysates were cleared (1000 × g, 5 min, 4 °C) and the membrane fraction was collected at 10,000 × g for 5 min at 4 °C. The pellet was resuspended in 0.1 ml of 100 mM phosphate buffer, pH 8.0, with 50 mM NaCl. Proteins were separated by SDS-PAGE and blotted on PVDF membranes (Hybond-P, Amersham Bioscience). Detection was done with a monoclonal mouse anti-hemagglutinin antibody (Roche) and a horseradish peroxidase–conjugated secondary antibody (Jackson ImmunoResearch Laboratories) using the ECL Plus system (GE Healthcare) for chemiluminescence (Lumi-Imager F1, Roche).

**TgFNT1–3 transport assays in yeast**

Assays were done as described earlier (8). Briefly, a yeast liquid culture in selective media (S.D.) was harvested at an A_{600} of 1, and the cells were resuspended in 50 mM HEPES/Tris buffer (pH 6.8), or in 50 mM citric acid/Tris buffer (pH 4.8). The yeast cell suspensions were adjusted to an A_{600} of 50 (±10%) and kept on ice. The transport assays were started by adding 20 μl of the substrate solution to 80 μl of yeast suspension to yield a final concentration of 1 mM substrate and 0.04 μCi radiolabeled [1-14C] substrate (formate or L-lactate). Assays were done at 18 °C. Substrate uptake was stopped at specified time points by abrupt dilution with 1 ml ice-cold water and rapid transfer onto a vacuum filtration unit fitted with a GF/C filter membrane (Whatman). After washing with 7 ml ice-cold water, the filter membranes were transferred into scintillation vials filled with 3 ml of scintillation fluid (Quicksafe A, Zinsser Analytic). Vials were analyzed using a liquid scintillation counter (Packard Tri-Carb, PerkinElmer Life Sciences). The transport rates were calculated from single-exponential fittings. Unless specified otherwise, all assays were performed with three independent biological replicates. The background subtraction was done using the nonexpressing parental yeast cells. The given S.E. error margins can be converted into standard deviation by multiplication with the square root of 3, i.e. 1.732.

**Chemical modification and inhibition of TgFNTs in yeast**

80 μl of yeast cell suspension were incubated with 100 μM pCMBS or 1 mM MMTS for 30 min at 18 °C. The transport assays were stopped at 2 min, i.e. in the initial linear uptake phase. For determination of IC_{50} values, 1 μl of DMSO with or
without test compound at different concentrations was added to 80 μl of the cell suspension 15–20 min prior to the assay solution. Uptake was stopped in the initial linear phase, i.e. at 2 min for truncated WT TgFNT1–3, and at 4 min for the Gly to Ser mutants, respectively. Three independent assays, each in triplicate was done.

Parasite culture

Tachyzoites of *T. gondii* (RH-hxgprt− and RH-hxgprt−–YFP strain) were cultured in confluent monolayers of primary human foreskin fibroblast (HFF) cells, essentially as described before (28, 40). Briefly, the host cells were maintained in DMEM supplemented with fetal bovine serum (10%), glutamine (2 mM), sodium pyruvate (1 mM), penicillin (100 units ml−1), streptomycin (100 μg ml−1), and 1 × nonessential amino acids (minimum Eagle’s medium). Cultures were incubated at 37 °C in a humidified incubator with 5% CO2. For regular parasite cultures, HFF cells were infected with a multiplicity of infection of 3, and tachyzoites were passaged every second day into fresh host cells.

Expression of TgFNT1–3 in tachyzoites

The full-length ORFs of TgFNT1–3 were cloned into pGRA1-LIPKO vector digested with Nsi I/Pac I sites. The forward primer for TgFNT1 contained Sbf I (compatible with Nsi I in the vector), whereas the forward primers of TgFNT2–3 were engineered with Nsi I sites (Table S1). The reverse primers for all ORFs contained Pac I sites. This vector allowed stable genomic insertion of the single-copy TgFNT1–3 expression cassettes at the uracil phosphoribosyltransferase locus via negative selection with 5-fluoro-oxyuridine (FuDR) (41). To make transgenic strains, fresh extracellular tachyzoites (~10^7) were harvested and suspended in cytomix buffer (120 mM KCl, 0.15 mM CaCl2, 10 mM K2HPO4/KH2PO4, 25 mM HEPES, 2 mM EGTA, 5 mM MgCl2, 5 mM GSH, 5 mM ATP) and 10 μg plasmid, immediately followed by electroporation (BTX instruments). Transfected tachyzoites were used to infect confluent host cells and drug-selected with 5-fluoro-oxyuridine before subjecting them to immunolocalization assays.

Indirect immunofluorescence assays

HFF cells seeded on glass coverslips were infected with freshly isolated tachyzoites for 24 h, fixed with 4% paraformaldehyde for 10 min, and then neutralized with 0.1 M glycine in PBS for 5 min. Samples were permeabilized by 0.1% Triton X-100 in PBS for 20 min, blocked with 2% BSA in PBS/0.1% Triton X-100 for 20 min. Subsequently, they were stained with primary antibodies (mouse anti-HA, 1:10,000; rabbit anti-Tg-Gap45, 1:10,000) for 1 h and then with respective secondary antibodies (Alexa Fluor 488 and Alexa Fluor 594, 1:3000) for 45 min in the dark. Samples were mounted into a mix of Fluoromount-G and DAPI, dried at 4 °C overnight and examined by fluorescence microscopy (Apotome, Zeiss). To resolve the C-terminal topology of TgFNT1–3, extracellular parasites were stained with a mouse α-HA antibody (1:3000), suspended either in 3% BSA/PBS solution for pre-permeabilization staining, or in 2% BSA/0.2% Triton X-100/PBS for post-permeabilization, as shown in Fig. S1. The inner membrane complex was separated from the plasma membrane by treating extracellular parasites with α-toxin from *Clostridium septicum* (20 nM, 2 h) (List Biological Laboratories) followed by fixation on BSA-coated (0.01%) coverslips. In both cases, the standard secondary antibody staining procedure was performed afterward, as described above.

Parasite growth assays

Plaque assays were done in 6-well plates seeded with HFF cells (42). Inhibitors were dissolved in DMSO, and then mixed with 2 ml of culture media before diluting them to desired amounts into each well. The carrier solvent DMSO was also included as a control alongside in all assays. Host cell monolayers were infected with 150 parasites, and parasitized cells were cultured unperturbed for 7 days (37 °C, 5% CO2). Samples were fixed with ice-cold methanol for 10 min and then stained with crystal violet dye for 15 min. The plaque sizes were measured using the ImageJ program (National Institutes of Health). To set up YFP-based growth assays, HFF monolayers in 96-well plates were infected with 7 × 10^3 tachyzoites of the RH-hxgprt−–YFP strain. After 4 h, the inoculation medium was replaced by colorless medium (free of phenol red) containing 1, 10, or 100 μM of the indicated compounds. The YFP fluorescence intensity was recorded every 24 h for up to 120 h (excitation, 485 nm; emission, 528 nm; area scan mode). Parasites treated with 1 μM pyrimethamine or with DMSO served as the positive and negative controls, respectively.

**Author contributions**—H. E., B. R., N. G., and E. B. methodology; H. E. and B. R. writing—review and editing; N. G. and E. B. conceptualization; N. G. and E. B. writing—original draft; E. B. formal analysis; E. B. supervision; E. B. funding acquisition; E. B. project administration.

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