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Tipparaju, S.K., Muench, S.P., Mui, E.J. et al. (10 more authors) (2010) Identification and Development of Novel Inhibitors of Toxoplasma gondii Enoyl Reductase. *Journal of Medicinal Chemistry*, 53 (17). pp. 6287-6300. ISSN 0022-2623

<https://doi.org/10.1021/jm9017724>

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Published in final edited form as:

J Med Chem. 2010 September 9; 53(17): 6287–6300. doi:10.1021/jm9017724.

Identification and Development of Novel Inhibitors of *Toxoplasma gondii* Enoyl Reductase

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Abstract

Toxoplasmosis causes significant morbidity and mortality and yet available medicines are limited by toxicities and hypersensitivity. Since improved medicines are needed urgently, rational approaches were used to identify novel lead compounds effective against *Toxoplasma gondii* enoyl reductase (TgENR), a type II fatty acid synthase enzyme essential in parasites but not present in animals. Fifty-three compounds, including three classes that inhibit ENRs, were tested. Six compounds have anti-parasite MIC_{90s} ≤ 6 μM without toxicity to host cells, three compounds have IC_{90s} < 45 nM against recombinant TgENR and two protect mice. To further understand the mode of inhibition, the co-crystal structure of one of the most promising candidate compounds in complex with TgENR has been determined to 2.7 Å. The crystal structure reveals that the aliphatic side chain of compound **19** occupies, as predicted, space made available by replacement of a bulky hydrophobic residue in homologous bacterial ENRs by Ala in TgENR. This provides a paradigm, conceptual foundation, reagents, and lead compounds for future rational development and discovery of improved inhibitors of *T. gondii*.

Introduction

Toxoplasmosis causes substantial morbidity and mortality, especially in persons who are congenitally infected or immune-compromised and this parasite is the most frequent infectious cause of uveitis.^{1–4} *Toxoplasma gondii* (*T. gondii*, Tga) is acquired as a

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The PDB ID for TgENR co-crystallized with compound 19 is 3NJ8.

Supporting Information Available

Table S1 contains HPLC purity analysis data of the tested TgENR inhibitors.

sporozoite from oocysts formed in cats or bradyzoites from cysts in meat. In humans, this parasite has a simple life cycle consisting of two stages; tachyzoites and bradyzoites. The former are a rapidly growing, obligate intracellular forms of *T. gondii* present when parasites are first acquired in acute infections. *T. gondii* then develops into slowly growing, encysted, latent bradyzoites, sequestered within cysts inside cells, with a competent host immune response. When a cyst ruptures, stage transition from latent bradyzoites back to rapidly growing tachyzoites occurs, causing destruction of surrounding tissue.

Reasons for recrudescence of eye disease have not been completely defined but is a lifelong problem in individuals infected congenitally as well as some of those whose infection is acquired after birth.^{3,4} This is an especially pressing problem in Brazil, as 80% of the population is infected with particularly pathogenic parasite strains, with a high incidence beginning in childhood. In some regions of Brazil, 20% of these individuals and 50% of those over 50 years old have eye disease. In immunocompromised persons such as those with AIDS, disease due to recrudescence (especially in the brain) is frequent, occurring in 50% of those with AIDS whose HIV infection remains untreated. Life threatening toxoplasmosis occurs in those immunocompromised by malignancies, organ transplantations and autoimmune disease with associated treatments. Rarely, there is significant organ damage in those without known immune compromise. An epidemic of multivisceral, lethal disease caused by a hypervirulent strain of parasite was reported recently in Guyana, making this emerging infection potentially even more problematic with globalization of food supplies. This parasite can easily contaminate food supplies or the environment and is a potential bioterrorism pathogen. There have been several recent epidemics associated with contaminated water supplies.

Consequences of chronic infections present in ~30% of the population (~2 billion people) worldwide, throughout their lifetimes, are not thoroughly characterized. Recently, memory impairment was reported in healthy, young to middle aged professionals in association with this infection and presence of a susceptibility allele of a gene encoding an enzyme that degrades dopamine, catechol *O*-methyl transferase (COMT) (Yolken et al. 2009 Personal Communication). There is also a higher prevalence of antibody to *T. gondii* in those with cryptogenic epilepsy and schizophrenia, although cause and effect between *T. gondii* infection and these neurologic observations has not yet been proven.

There are only a few medicines that restrict growth of tachyzoites,¹⁻⁴ and use of these medicines is associated with significant incidences of hypersensitivity (up to 25%) and toxicity.⁵ No medicines eliminate encysted, latent bradyzoites. Better approaches to treat this disease are greatly needed including medicines that eliminate active parasites causing disease and means to eliminate latent parasites.

Recent work by our group,⁶⁻¹⁵ and a recent report by others,¹⁶ provide the foundation for the present work to develop a new class of medicines to better treat toxoplasmosis. Specifically, the prokaryotic-like type II fatty acid biosynthetic (fas) pathway in *T. gondii* is a validated molecular target in tachyzoites it is essential for parasite survival *in vitro* and *in vivo*.¹⁶ In particular, the enoyl reductase (ENR) enzyme, which catalyses the last reductive step of the type II fatty acid synthesis pathway, is present in all *T. gondii* life cycle stages except microgametes¹¹ and ENRs in other organisms have been shown to be the target for a wide range of potent inhibitors. Importantly, compounds which inhibit type II fatty acid synthesis (including triclosan and a number of newly designed and synthesized compounds) not only inhibit *T. gondii* tachyzoite growth but are effective against other apicomplexan

³Abbreviations: *T. gondii* and Tg, *Toxoplasma gondii*; ENR, enoyl reductase; COMT, catechol *O*-methyl transferase; HLM, Human liver microsomes.

parasites, such as the hepatic stage of *Plasmodia*^{7,16–20} which causes malaria. However, while triclosan is a potent inhibitor of the enzyme, its poor solubility makes it unsuitable as a potential therapeutic. This prompted the investigation into different triclosan scaffolds as potential therapeutic agents for toxoplasmosis, with improved activity, as well as physicochemical properties and toxicity profiles.

This report details activity of a panel of 53 compounds (Figure 1) based on previously identified ENR inhibitors, with two of these compounds (compounds **2** and **19**) showing low toxicity and activity in the low nM range. A promising adaptation to the triclosan scaffold is the addition of an *n*-propyl group at the 4-position, exploits an increase in space in the parasitic ENR binding pocket. In order to validate this, a co-crystal structure of one of the most promising inhibitors (compound **19**) in complex with TgENR, was determined to 2.7Å. This structure revealed that these inhibitors utilize the extra space within the binding pocket of the parasitic ENR family. The data presented herein provides insights into lead compounds which form the basis for future development of highly active inhibitors of ENR with potential for progression to medicines that effectively treat toxoplasmosis and related apicomplexan diseases.

Results

Synthesis of Inhibitors

Synthesis of compounds tested is shown in Figure 2 (Synthetic Schemes). The numbers, molecular weights, ClogPs and structures of compounds are shown in Figures 1 and 2. The requisite diphenyl ethers were synthesized either by nucleophilic aromatic substitution (Method A) or through Cu catalyzed coupling reactions (Method B). The choice of the method was dependent on the product desired and the choice of the starting phenol. In each case, the methoxy diarylether precursors so obtained were demethylated to phenols using boron tribromide (Scheme 1). The aryl boronic acid **20** was prepared by lithiation of 1-(3-chlorophenoxy)-2-methoxy-4-propylbenzene followed by quenching with trimethyl borate followed by acidic hydrolysis. The diarylether derivatives **2**, **4**, **6**, **8**, **9–12**, **14–19**, **21–24**, **29**, **31**,²¹ and the 2-pyridone derivatives **38–44** were previously published by us.²² Amides **7**, **25** and **27** were prepared from their corresponding aniline precursors using standard amide bond forming procedures (Scheme 2). Pivaloyl ester **3** was prepared from triclosan (Scheme 2). Aminopyridines **52** and **53** were synthesized from 3-(6-aminopyridin-3-yl)-acrylic acid²³ according to Scheme 3. Benzimidazolones **46–48** were prepared using standard procedures as depicted in Scheme 3. The HPLC and high resolution mass spectroscopy were employed to determine the purity of the tested compounds. Purity was between 96 and 100% as shown in Table S1.

Inhibition of *T. gondii* tachyzoites *in vitro*

A summary of ability of each compound to inhibit the parasite's growth by 50% (MIC₅₀) and 90% (MIC₉₀) and effect on growth and survival of nonconfluent cultures of the same fibroblasts host cells in the parasite inhibition assay, as an indication of toxicity, are in Figure 1.

Analysis of the 53 compounds identified 6 compounds that robustly inhibited parasites (~60–95% inhibition of parasite growth) with less than 20% toxicity to host cells with MIC₉₀s <6μM (Figure 1). An additional six compounds had modest inhibitory effect on parasites (~20% to 60% inhibition of parasite growth) and an additional three had ~20–40% inhibition of host cell uptake of tritiated thymidine (Figure 1). The next six compounds had less than 40% inhibitory effect on uptake of uracil by parasites or were very toxic to host cells precluding interpretation of parasite inhibition assays. The last compounds shown had

minimal inhibitory effect and/or harmed host cell growth substantially (Figure 1). These data showing IC₅₀ and IC₉₀ are in Figure 1. To illustrate these results in specific experiments with more detail, a representative experiment with compounds **25**, which was toxic for fibroblasts, and **19**, which was efficacious and not toxic are shown in Figure 3. Of the initial 53 compounds tested, compounds that displayed the greatest effect and least toxicity were **2**, **3**, **19**, and **39**.

Inhibition of TgENR activity

Compounds that inhibited the growth of *T. gondii* in culture (Figure 1, shaded regions) were initially tested for inhibition of TgENR enzymatic activity at three concentrations (0.2, 2, and 20 μM). Compounds which displayed significant inhibitory activity at 2 μM were assayed in triplicate at ten concentrations to determine IC₅₀ values as summarized in Figure 1 and shown in detail in Figure 3. This assay uses 20 nM TgENR, preventing the accurate measurement of IC₅₀ values below this concentration. Seven compounds (including triclosan) are listed as having IC₅₀ values below 20 nM. Compound **39** turned out to be a poor inhibitor of TgENR and hence appears to have an off target effect on parasites and requires further investigation. This result indicates that this compound may be of interest for further development but does not target ENR specifically.

Co-crystallization and structure solution of TgENR in complex with NAD⁺ and compound **19**

In order to gain insights into the mode of binding for compound **19** which is very active against the ENR enzyme (IC₅₀ < 20 nM; Figure 3 bottom panel) and the parasite in tissue culture, structural studies were conducted. TgENR was co-crystallized in the presence of NAD⁺ and compound **19** with the subsequent crystals diffracting to beyond 2.7 Å (Data are in Table 1). The refined structure showed clear and continuous density for the bound inhibitor allowing for its unambiguous placement within the structure (Figure 5a). The superposition of TgENR in complex with compound **19** and triclosan shows that the mode of binding for both inhibitors is similar with the aromatic rings of both compounds adopting the same position, with ring A in both cases forming stacking interactions with the NAD⁺ co-factor. The additional bulk of the alkyl substituent of compound **19**, when compared to triclosan, occupies the space made available by replacement of a bulky hydrophobic residue common to the bacterial enzyme family (Met206 in *E. coli*) by alanine (Ala247 in TgENR) which is conserved in the apicomplexan family (Figure 5c). Comparison of the triclosan and compound **19** TgENR complex also reveals that there is little change in the position of the mainchain atoms with an overall Cα r.m.s.d value of 0.3 Å (Figure 5b). However, the side chain of Phe 243 does move significantly by ~1.5 Å about its Cβ atom to accommodate the extra bulk of compound **19** compared to triclosan, without significant change to its mainchain position (Figure 5b, c). As such elaborating the nature of the substituent on the phenoxy ring of the candidate compound exploits the increase in space which is a conserved feature of the apicomplexan ENR family, without perturbing the overall fold of the binding pocket.

Inhibition of *T. gondii* tachyzoites *in vivo*

Effect of Compounds **2** and **19** on *T. gondii in vivo* are in Figure 4. Representative experiment of two replicate studies, indicate that compounds **2** and **19** can reduce parasite burden in mice (*P* < 0.05).

Cytochrome P450 inhibition and human liver microsomal stability

2 and **19** were tested for *in vitro* inhibition of recombinant CYP450 isozymes 2C9, 2D6 and 3A4. At 10 μM, **2** and **19** showed strong inhibition of CYP2C9 (both >85%) and relatively

low inhibition of CYP2D6 and CYP3A4 (both <22%). Further, metabolic stabilities in pooled human liver microsomes (HLM) indicated that they were moderately metabolized in HLM, and remaining parent drugs after incubation for 60 minutes were 51.23% and 45.14%, respectively. Intrinsic clearances, metabolic rates, of **2** and **19** in HLM were slower than the control drug verapamil.

Discussion

Triclosan is a broad spectrum antibacterial agent incorporated into a wide range of consumer products.^{24–26} It inhibits the ENR (FabI) enzyme in a number of microorganisms like *E. coli*,^{27,28} *P. aeruginosa*²⁹ and *S. aureus*.³⁰ Significant variation in the degree of inhibition of different pathogens by triclosan has been observed^{28–30} suggesting that it should be possible to modify the compound to improve its efficacy against a particular organism through systematic structure-based as well as ligand-based drug design methods to improve properties such as potency, solubility and toxicity.

The crystal structure of *T. gondii* and *P. falciparum* ENR in complex with triclosan shows the binding geometry of triclosan to be similar to that seen with other organisms. Thus the phenoxy 'ring A' (Figure 1 and 2) of triclosan and the nicotinamide ring of NAD⁺ engage in π -stacking interactions, while the phenolic hydroxyl group on ring A (Figure 1) is involved in a hydrogen bond with a conserved Tyr (OH), and the ether linkage has two hydrogen bonds, one to the hydroxyl group of Tyr, and the other to the ribose unit of the NAD⁺. Encouraged by our earlier results on triclosan activity against TgENR,⁷ we tested a number of newly synthesized triclosan derivatives in addition to those available to us from our previous work.^{21,22} In addition, we designed and tested compounds based on diazaborine and aminopyridine inhibitor families shown to inhibit the enzyme with the goal of attaining improved inhibitors.

Several of the triclosan derivatives identified were active against parasites at low micromolar levels with sub-micromolar inhibitory activity toward TgENR (Figures 1 and 3). In general, presence of a strong electron withdrawing group on the B ring contributed to improved activity while the presence of an electron donating group reduced the activities of these compounds (compare compounds **2** vs **1**, **8** vs **10**, **4** vs **10**. Freundlich and co-workers showed that compound **2** has an IC₅₀ of 0.2 μ M against *P. falciparum* ENR.³¹ This compound is highly active against TgENR as well, which is not surprising as the active sites of these two enzymes share significant homology. Unlike the observation of Sivaraman *et al.*,^{32,33} in the case of TgENR, deletion of chlorine atoms on the B ring of triclosan (as in compound **4**) does not improve activity toward both TgENR and the parasite, suggesting that the presence of a more lipophilic group would favor the improvement in activity. However, modification of the B ring to a biphenyl to increase its hydrophobicity did not improve activity (compound **6**). This is not surprising since this group faces the NAD⁺ co-factor and would cause a large degree of steric hindrance accounting for the large increase in IC₅₀ value (>2000 nM). When the B ring of triclosan is replaced by a hetero-aromatic ring resulted in compounds **17** and **18** with compound **17**, the pyridine analog of compound **4** showing some activity. However, compound **18**, the corresponding pyrazine analog had no activity. Since compound **18** is smaller than triclosan this lack of activity is unlikely to be caused simply through steric hindrance within the binding pocket highlighting the importance of presentation of an appropriate electronic distribution in this ring to allow for optimal interaction with the active site. Moreover, the pyrazine may be attracted towards the NAD⁺ cofactor, destabilizing both the rings A and B, driven in part by the formation of a hydrogen bond between the backbone oxygen of NAD⁺ and nitrogen of compound **18**. Further studies are required to test this hypothesis. Although of the 11 compounds reflecting changes focused on the B ring (**2**, **4**, **8–10**, **14–18**) only compound **2** displays a comparable

efficacy to triclosan. These observations, taken together open up additional avenues for further development of a new class of heteroaryl analogs of triclosan. However, caution must be taken to not disturb the important interaction the A ring makes with the NAD⁺ cofactor through changes in the B ring.

The phenolic hydroxyl group on ring A is considered to be critical to triclosan's activity, as it is involved in hydrogen bonding to a conserved Tyr residue in the active site and forming packing interactions with the NAD⁺ cofactor. This has been observed by us in our earlier work,¹⁰ and is also evident by the inactivity of compound **5**. The significant change in potency between compounds **4** (IC₅₀ >20<200 nM) and **5** (IC₅₀ >20000 nM), can only be attributed to a change of the OH group to NH₂. In light of this observation, it is interesting to note that **3** shows improved activity on the parasites over triclosan itself. Since the pivalate group would produce severe steric clashes within the binding pocket (as highlighted by the increased IC₅₀ value) we suspect that this heightened activity may be due to the enhanced ability of this pro-drug form to enter parasites, whereby the ester is hydrolyzed. Similarly, while the antiparasitic activities of compounds **4** and **13** were comparable, there is significant difference in their TgENR activity indicating that a tertiary butyl group is indeed too bulky at this position. The differences in the antiparasitic activity and TgENR inhibitory activity observed in some of the analogs is not surprising and could be attributed to off target effects and problems with delivery, particularly with delivery of an inhibitory compound into the apicoplast that has 4 surrounding membranes and in which ENR resides. We intend to investigate this finding further.

Diazaborines are known to inhibit ENR through covalent bond formation between the 2'-hydroxyl of the nicotinamide ribose unit of the NAD⁺ molecule and the boron atom of the drug.³⁴ Consequently, we believed that it would be valuable to incorporate this type of functionality into our triclosan based ENR inhibitors, and thus we prepared the boronic acid analog **20**. This compound also proved to be a very effective inhibitor of TgENR, although further studies are needed to determine whether a covalent bond has been formed with the nicotinamide ribose. Other attempts to vary the *meta* position of the B ring with H-bond donors and acceptors did not give encouraging results (compounds **29**, **31**, and **33**).

Although the indole aminopyridine compound **50**²³ was very effective against *S. aureus* and *E. coli* ENR, it shows only moderate activity toward TgENR. We synthesized several aminopyridine mimics of this compound, namely **50**, **52** and **53**, in order to determine whether this series had any special merits that might warrant further investigation. Unfortunately, none of these showed any promising activity, this may in part be due to the steric hindrance which can be brought about by Phe243 (TgENR numbering) in the binding pocket as previously reported.¹⁵

When the A ring was modified by replacing the chlorine atom at the *para* position by an hydrophilic OH group, the activity decreased (compounds **11** and **12**). These changes on ring A may be brought about by the disturbance of the stacking interactions with the NAD⁺ cofactor which are critical for inhibitor binding. Furthermore, this inactivity could also be influenced by the lower lipophilicity of these compounds rather than the electron-donor effect of the hydroxyl group. To test this hypothesis, we attempted to append a long hydrophobic alkyl chain at the *para* position. This can be accommodated since the bacterial and plant ENR family contain a bulky hydrophobic residue (Met, Leu or Ile) close to the ring A of triclosan, whereas the ENR of apicomplexan species such as *T. gondii* and *Plasmodium* have a fully conserved alanine residue. This sequence change produces an increase in space at the base of the binding pocket and reduces the van der Waals packing interactions with the triclosan based inhibitors in comparison to other members of the ENR family.¹⁵ When we synthesized a number of triclosan derivatives bearing an *n*-propyl group

at the 4-position which were predicted to be optimal length to fit within the binding site, these inhibitors showed, in general, better or similar activities than the triclosan scaffold (compounds **19–24**), with no detrimental effect of this extension to their IC₅₀ values with derivatives that bear electron withdrawing groups on the B ring (compounds **19, 21–24**) producing among the most active compounds identified to date.

To further investigate the mode of binding of the most promising inhibitor which contains an *n*-propyl group at the 4 position, TgENR was co-crystallized in the presence of compound **19** and NAD⁺. The structure reveals that compound **19** binds in very similar mode to triclosan (Figure 5c).^{35–45} A structural comparison of the complex with that of the *E. coli* ENR/NAD⁺/triclosan enzyme reveals that the additional bulk of the alkyl substituent of compound **19** compared to triclosan does occupy the space made available by the replacement of a bulky hydrophobic residue of the *E. coli* enzyme Met by alanine in TgENR within the substrate binding pocket. Moreover, the structure shows that *n*-propyl is not the most optimal group as there is still some additional space at the base of the pocket which may be further utilized through further modifications. The only significant difference in side chain positions between the compound **19** and triclosan complex is a small change in the position of Phe243 to accommodate the larger bulk of compound **19** (Figure 5c). Presence of an *n*-propyl group on the 4-position clearly does not adversely affect the activity of these inhibitors (compare compounds **2** and **22**), hence this side chain could be further modified to modulate both physicochemical properties and activity of these compounds. Moreover, despite triclosan and compound **19** showing similar solubility, as shown by their ClogP, ALOGpS, and tPSA values (Table 2), there is more scope for improvement through modifications of compound **19**, (in particular the additional *n*-propyl group) when compared to triclosan. Since the ability to bind the triclosan analogues with the extension on the *n*-propyl group is only achieved within the parasitic ENR it may be possible to develop an inhibitor which is specific for the parasitic and not bacterial family. Although the addition of the *n*-propyl side chain maximizes packing interactions within the active site, these interactions are not fully exploited and could explain why the *n*-propyl group did not enhance activity. The additional *n*-propyl group does however produce a compound which is more tailored towards the apicomplexan and not bacterial ENR family. Moreover, further development of this inhibitor family is feasible particularly with the crystal structure showing there is still additional space at the base of the inhibitor binding pocket which could be further modified to improve binding.

By maintaining the A ring of these compounds which is involved in packing interactions with the NAD⁺, and changing the B ring, it is clear that subtle variations in the MIC₉₀ value can be achieved. Since these changes are small it may be that they affect the packing of the B ring within the active site or change its electrostatic properties without destabilizing the interaction with the A ring and NAD⁺ such that the mode of binding will be very similar to triclosan. Among these, inhibitor **19**, bearing a 4-cyano group on the B ring seemed to be most promising with the least toxicity to host cells. Interestingly, compound **24** with a 4-amino functionality on the B ring also showed reasonable activity albeit lower than compound **19**. This lower activity is caused through the removal of the chlorine and replacement of CN with NH₂ on ring B, this again affects the properties of the B ring whilst allowing the A ring to stack onto the NAD⁺ cofactor such that the IC₅₀ value is still comparable with triclosan. From a medicinal chemistry perspective, triclosan with two chlorine atoms at the 2' and 4' positions on the ring B presents little opportunity to incorporate any polar substitutions in order to reduce the lipophilicity of the drug. The rationale behind introduction of functionalities such as CN or NO₂ on ring B is to create an opportunity to modify these groups in future rounds of ligand optimization to improve the physicochemical properties of these compounds. However, increased electron density on the

A ring (a difference between compound **19** vs. **30**, and **32**) which enhances ability of compound **19** to π -stack with the nicotinamide ring of NAD⁺ did not improve activity.

As a final proof of concept for the promising activity of compounds **2** and **19**, they were tested in an animal model for *Toxoplasma* infection with both protecting mice against parasite burden *in vivo* with an efficacy approximately 5 times higher than triclosan itself (Figure 4). In addition, a further study into the metabolism of these derivatives was performed. Both compounds **2** and **19** were tested for *in vitro* inhibition of recombinant CYP450 isozymes 2C9, 2D6 and 3A4. At 10 μ M, **2** and **19** showed strong inhibition of CYP2C9 (both >85%) and relatively low inhibition of CYP2D6 and CYP3A4 (both <22%). Further metabolic stabilities in pooled HLM indicated that they were moderately metabolized in HLM, and remaining parent drugs after incubation for 60 minutes were **51.23%** and **45.14%**, respectively. Intrinsic clearances and metabolic rates of **2** and **19** in HLM were slower than the control drug verapamil. Interestingly, when certain of these compounds recently were used to test their efficacy against *Bacillus anthracis* some were noted to be effective.^{21,22} The scavenging of fatty acids which abrogates the lethal effect of the elimination of Type 2 FAS with *Staphylococcus aureus* in mice is not seen with *T. gondii*.⁴⁶ Triclosan was noted to be safe in mice in earlier studies.^{10,47}

Conclusions

In all a series of fifty-three compounds were tested for their activity against *T. gondii*, with six compounds having anti-parasite MIC_{90s} $\leq 6\mu$ M without toxicity to host cells. Three compounds had IC_{90s} <45nM against recombinant TgENR and of the 44 triclosan analogs tested, 6 showed better activity than triclosan. Moreover, those with the best toxicological profiles were tested in mice (compounds **2** and **19**), and were found to be several times more active than triclosan itself in protecting mice. To investigate if, as predicted, the larger bulk of a subset of these compounds exploits the increase in space in the parasitic enzymes, a co-crystal structure of compound **19** in complex with TgENR, and NAD⁺ was solved. This structure clearly revealed that the compound's aliphatic side chain does, as predicted, occupy space made available in the parasitic ENR family and reveals that further modifications may be possible. Moreover compound **19** with its adaptations on both the A and B ring is more amenable to chemical modifications which could improve both potency and solubility.

This work provides a foundation for discovery of medicines to treat toxoplasmosis by targeting ENR and for a program which includes a structure based program to develop new medicines against ENR. This work has produced new insights into the efficacy of a range of inhibitor scaffolds. The co-crystal structures proved useful in understanding structure activity relationships. In particular the co-crystal structure of compound **19** has revealed that a subset of these compounds exploit the increase in space within the parasitic ENR family which these inhibitors target. This strategy can now be utilized in future medicinal chemistry efforts to develop compounds that inhibit TgENR with increased avidity along with modifications which will increase bioavailability.

Experimental Section

Chemistry

Synthesis of Inhibitors—¹H NMR and ¹³C NMR spectra were recorded on Bruker spectrometer with TMS as an internal standard. Standard abbreviation indicating multiplicity was used as follows: s = singlet, d = doublet, t = triplet, q = quadruplet, quin = quintuplet, m = multiplet and br = broad. HRMS experiment was performed on Q-TOF-2TM (Micromass). The progress of all reactions was monitored by TLC on precoated *silica gel*

plates (Merck *Silica Gel* 60 F254). Preparative TLC was performed with Analtech 1000–mm *silica gel* GF plates. Column chromatography was performed using Merck *silica gel* (40–60 mesh).

Compounds triclosan, **5**, **36**, **49**, **54** and **37** were commercially available and have been purchased. Compounds **50** and **51** were synthesized according to the reported procedures.²³ The new diarylether analogs of triclosan were synthesized by Schemes 1 and 2.²¹ The synthesis of aminopyridine compounds **53** and **52**, as well as benzimidazole-2-one derivatives **46**, **47**, and **48** were carried out according to the Scheme 3. All the compounds not illustrated in the synthetic schemes were reported earlier.^{21,22} Descriptions of synthetic experimental details and analytical data of compounds in Figure 1 is as follows:

General Methods

Method A—The aryl halide (1 mmol), phenol (1 mmol), and K₂CO₃ or Cs₂CO₃ (2–4 mmol) in DMSO (1.5 mL) were heated to 100 °C under nitrogen until the reaction was shown to be complete by TLC (8–12 h). After cooling to rt, the reaction mixture was diluted with ethyl acetate, and washed with 5% aqueous NaOH solution. The aqueous layer was further extracted with ethyl acetate, and the combined organic layers were washed with brine. The organic layer was dried over Na₂SO₄, and concentrated under vacuum to give the crude product, which was subsequently purified by flash chromatography on *silica gel*.

Method B—To the phenol (1 mmol) dissolved in DMF (1.75 mL) was added KO^tBu (1.1 mmol) in one portion and the mixture was heated at 45 °C under mild vacuum. After 2 h, the reaction mixture was cooled to rt. Aryl halide (1 mmol) and (CuOTf)₂·Ph (0.05 mmol) were added and the mixture was heated to reflux at 145 °C for 16–20 h. After cooling to rt, the reaction mixture was diluted with ethyl acetate, filtered over Celite, and extracted with ethyl acetate. The extract was washed with 5% aqueous NaOH solution. The aqueous layer was then extracted with ethyl acetate, and the combined organic layers were washed with brine. The organic layer was dried over Na₂SO₄, and concentrated under vacuum to give the crude product, which was subsequently purified by flash chromatography on *silica gel*.

Method C—In each case, the methoxy diarylether derivatives obtained by the above methods were converted to the corresponding phenols using boron tribromide according to the following procedure: A solution of boron tribromide (2–8 mmol of 1.0 M solution in dichloromethane) was added to a solution of the corresponding methoxy diphenylether analog (1 mmol) in dry dichloromethane (4 mL) maintained at –78 °C under nitrogen. The reaction mixture was stirred at the same temperature (1 h), and then at rt (3–8 h) while monitoring by TLC. Following completion, the reaction was quenched with methanol at –78 °C and concentrated under vacuum. The concentrate was further dissolved in ethyl acetate, washed with 10% aqueous sodium bicarbonate solution, and the organic layer was separated and washed with water and then with brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel*.

2-(3-Chlorophenoxy)-5-propylphenol (33)—To the 2-methoxy-4-propyl-phenol (1 g, 6.0 mmol) dissolved in DMF (11 mL) was added KO^tBu (0.81 g, 7.2 mmol) in one portion and the mixture was heated at 45 °C under vacuum. After 2 h, the reaction mixture was cooled to room temperature. 1-chloro-3-iodobenzene (1.72 g, 7.2 mmol), and (CuOTf)₂·Ph (0.31 g, 0.6 mmol) were added, and the mixture was heated to reflux at 145 °C for 18 h (Method B above). Usual workup and purification by flash chromatography (SiO₂, 10% EtOAc/Hexanes) gave analytically pure **1-(3-chlorophenoxy)-2-methoxy-4-propylbenzene** as a colorless oil (62%). ¹H NMR (400 MHz, CDCl₃): δ = 1.00 (t, *J* = 7.3 Hz, 3H), 1.70

(quin, $J = 7.5$ Hz, 2H), 2.62 (t, $J = 7.5$ Hz, 2H), 3.83 (s, 3H), 6.79 (dd, $J = 8.0, 1.7$ Hz, 1H), 6.85–6.83 (m, 2H), 6.91 (t, $J = 2.1$ Hz, 1H), 6.95 (d, $J = 8.0$ Hz, 1H), 7.00 (dd, $J = 7.9, 1.0$ Hz, 1H), 7.21 ppm (t, $J = 8.1$ Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 13.5, 24.3, 37.6, 55.5, 112.8, 114.3, 116.3, 120.6, 121.4, 121.7, 129.8, 134.4, 140.3, 141.2, 151.0, 159.0$ ppm.

A mixture of above intermediate **1-(3-chlorophenoxy)-2-methoxy-4-propylbenzene** (0.22 g, 0.79 mmol) and BBr_3 (1M, 2.5 mL, 2.5 mmol) were subjected to the general demethylation procedure outlined in method C above. The crude product was purified by flash chromatography (SiO_2 , 10% EtOAc/Hexanes) to give analytically pure **33** as a colorless oil (62%). ^1H NMR (400 MHz, CDCl_3): $\delta = 0.99$ (t, $J = 7.3$ Hz, 3H), 1.67 (quin, $J = 7.5$ Hz, 2H), 2.58 (t, $J = 7.5$ Hz, 2H), 5.43 (br s, 1H), 6.72 (d, $J = 8.1$ Hz, 2H), 6.86 (d, $J = 8.2$ Hz, 2H), 6.92–6.89 (m, 2H), 7.02 (s, 1H), 7.09 (d, $J = 8.0$ Hz, 1H), 7.26 ppm (t, $J = 8.1$ Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 13.5, 24.1, 37.2, 115.1, 116.2, 117.3, 119.2, 120.5, 122.9, 130.2, 134.8, 139.9, 140.3, 146.9, 157.8$ ppm. HRMS (ESI-positive): calcd for $\text{C}_{15}\text{H}_{15}\text{ClO}_2$ ($[\text{M}-\text{H}]^+$): 261.06878, found: 261.0693.

3-Chloro-4-(2-hydroxy-6-methoxy-4-propylphenoxy)benzotrile (30)—2,6-Dimethoxy-4-propylphenol (1.0 g, 5.1 mmol), 3-chloro-4-fluoro-benzotrile (0.79 g, 5.1 mmol) and K_2CO_3 (1.41 g, 10.2 mmol) in DMSO (8 mL) were heated to 100 °C under nitrogen for 12 h (Method A above). Usual workup followed by purification of the crude reaction mixture by column chromatography (SiO_2 , 10% EtOAc/Hexanes) afforded the compound **3-chloro-4-(2,6-dimethoxy-4-propylphenoxy)benzotrile** (1.57 g, 93%). ^1H NMR (400 MHz, CDCl_3): $\delta = 1.00$ (t, $J = 7.3$ Hz, 3H), 1.70 (quin, $J = 7.5$ Hz, 2H), 2.62 (t, $J = 7.5$ Hz, 2H), 3.78 (s, 6H), 6.49 (s, 2H), 6.62 (d, $J = 8.6$ Hz, 1H), 7.38 (dd, $J = 8.6, 1.8$ Hz, 1H), 7.73 ppm (d, $J = 1.8$ Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 13.4, 24.2, 38.2, 55.8, 105.0, 114.5, 117.6, 122.8, 128.3, 131.5, 133.4, 141.4, 151.9, 157.6$ ppm. MS (ESI) m/z : 354.1 $[\text{M} + \text{Na}]^+$

A mixture of above intermediate **3-chloro-4-(2,6-dimethoxy-4-propylphenoxy)benzotrile** (0.29 g, 0.86 mmol) and BBr_3 (1M, 3.4 mL, 3.4 mmol) were subjected to the general demethylation procedure outlined in method C above. The reaction resulted in two products (**30**, 20% yield) and (**26**, 60% yield) that were separated by flash chromatography (SiO_2 , 15% EtOAc/Hexanes) to give analytically pure samples as white solids.

3-Chloro-4-(2-hydroxy-6-methoxy-4-propylphenoxy)benzotrile (30)— ^1H NMR (400 MHz, CDCl_3): $\delta = 0.99$ (t, $J = 7.3$ Hz, 3H), 1.68 (quin, $J = 7.5$ Hz, 2H), 2.57 (t, $J = 7.5$ Hz, 2H), 3.73 (s, 3H), 5.67 (s, 1H), 6.40 (s, 1H), 6.54 (s, 1H), 6.73 (d, $J = 8.6$ Hz, 1H), 7.40 (d, $J = 8.6$ Hz, 1H), 7.70 ppm (s, 1H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 13.4, 23.9, 37.8, 55.6, 104.4, 105.8, 108.9, 115.0, 117.2, 123.0, 126.8, 131.6, 133.5, 141.8, 148.3, 151.2, 156.8$ ppm. HRMS (ESI-positive): calcd for $\text{C}_{17}\text{H}_{16}\text{ClNO}_3$ ($[\text{M}-\text{H}]^+$): 316.0746, found: 316.0755.

3-Chloro-4-(2,6-dihydroxy-4-propylphenoxy)benzotrile (26)— ^1H NMR (400 MHz, CDCl_3): $\delta = 0.98$ (t, $J = 7.3$ Hz, 3H), 1.66 (quin, $J = 7.5$ Hz, 2H), 2.53 (t, $J = 7.5$ Hz, 2H), 5.04 (s, 2H), 6.47 (s, 2H), 6.85 (d, $J = 8.6$ Hz, 1H), 7.40 (d, $J = 8.6$ Hz, 1H), 7.77 ppm (d, $J = 1.9$ Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 13.4, 23.7, 37.4, 106.7, 108.8, 115.1, 117.0, 123.2, 126.0, 131.9, 133.8, 142.3, 147.7, 156.1$ ppm. HRMS (ESI-positive): calcd for $\text{C}_{16}\text{H}_{14}\text{ClO}_3$ ($[\text{M}-\text{H}]^+$): 302.05894, found: 302.0606.

4-(2,6-Dihydroxy-4-propylphenoxy)benzotrile (32)—2,6-Dimethoxy-4-propylphenol (0.63 g, 3.2 mmol), 4-fluoro-benzotrile (0.39 g, 3.2 mmol) and Cs_2CO_3 (2.1 g, 6.4 mmol) in DMSO (9 mL) were reacted according to the Method A above. Usual

workup followed by purification of the crude reaction mixture by column chromatography (SiO₂, 10% EtOAc/Hexanes) afforded the compound **4-(2,6-dimethoxy-4-propylphenoxy)benzotrile** (0.81 g, 86%). A mixture of this intermediate compound (0.20 g, 0.67 mmol) and BBr₃ (1M, 3.4 mL, 3.4 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO₂, 10% EtOAc/Hexanes) to give analytically pure **32** as a colorless oil (74%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.90 (t, *J* = 7.2 Hz, 3H), 1.55 (quin, *J* = 7.2 Hz, 2H), 2.40 (t, *J* = 7.2 Hz, 2H), 6.27 (s, 2H), 6.91 (d, *J* = 8.8 Hz, 2H), 7.73 (d, *J* = 8.8 Hz, 2H), 9.41 ppm (s, 2H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 14.1, 24.2, 37.6, 103.7, 107.9, 116.3, 119.6, 127.0, 134.4, 140.5, 150.7, 162.3 ppm. HRMS (ESI-positive): calcd for C₁₆H₁₅NO₃ ([M-H]⁺): 268.09792, found: 268.0988.

3-Chloro-4-(2,6-dihydroxy-4-propylphenoxy)benzoic acid (34)—A mixture of the nitrile **3-chloro-4-(2,6-dimethoxy-4-propylphenoxy)benzotrile** (0.60 g, 1.8 mmol) whose synthesis is described in the preparation of **30** above and 25% aqueous NaOH (0.9 mL) in ethanol (5 mL) was refluxed with stirring (20 h). After cooling, the reaction mixture was acidified with dilute hydrochloric acid. The precipitate was filtered, dried, and purified by column chromatography (SiO₂, 6% MeOH/CHCl₃) to yield the **3-chloro-4-(2,6-dimethoxy-4-propylphenoxy)benzoic acid** (0.51 g, 80%) as a white solid. ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.94 (t, *J* = 7.2 Hz, 3H), 1.66 (quin, *J* = 7.6 Hz, 2H), 2.58 (t, *J* = 7.6 Hz, 2H), 3.38 (br s, 1H), 3.71 (s, 6H), 6.53 (d, *J* = 8.8 Hz, 1H), 6.67 (s, 2H), 7.76 (d, *J* = 8.8 Hz, 1H), 7.97 ppm (s, 1H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 14.2, 24.5, 38.2, 56.4, 105.9, 114.1, 121.0, 125.4, 128.3, 130.3, 131.6, 141.7, 152.3, 157.4, 166.3 ppm. MS (ESI) *m/z*: 372.1 [M + Na]⁺ A mixture of this intermediate (0.40 g, 1.14 mmol) and BBr₃ (1M, 6.8 mL, 6.8 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO₂, 5% MeOH/Chloroform) to give analytically pure **34** as a white solid (76%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.91 (t, *J* = 7.2 Hz, 3H), 1.56 (quin, *J* = 7.2 Hz, 2H), 2.41 (t, *J* = 7.6 Hz, 2H), 6.28 (s, 2H), 6.61 (d, *J* = 8.4 Hz, 1H), 7.77 (dd, *J* = 8.6, 1.6 Hz, 1H), 7.95 (d, *J* = 2.0 Hz, 1H), 9.45 ppm (br s, 1H); ¹³C NMR (100 MHz, CDCl₃): δ = 14.1, 24.2, 37.6, 107.9, 114.6, 121.3, 125.0, 127.2, 130.1, 131.4, 140.6, 150.6, 157.8, 166.4 ppm. HRMS (ESI-positive): calcd for C₁₆H₁₅ClO₅ ([M-H]⁺): 321.0535, found: 321.0545.

3-(2-Hydroxy-4-propylphenoxy)phenyl boronic acid (20)—1-(3-chlorophenoxy)-2-methoxy-4-propylbenzene (0.83 g, 3.0 mmol), whose synthesis is described in the preparation of **33**, was dissolved in 20 mL of anhydrous THF and treated with n-BuLi (1.6 M in hexanes, 3.8 mL, 6 mmol) at -78 °C for 2.5 h under Ar. Trimethyl borate (3.1 g, 30 mmol) was added drop wise to this solution at -78 °C and the reaction mixture was stirred for 3 h. The reaction was quenched by careful addition of 14 mL of 2N HCl at -78 °C and stirred at room temperature for 18 h. The solvent was evaporated and the concentrate was further dissolved in ethyl acetate, washed with 10% aqueous sodium bicarbonate solution, and the organic layer was separated and washed with water and brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (20% EtOAc/Hexanes) to result in **3-(2-methoxy-4-propylphenoxy)phenyl boronic acid** (0.65 g, 76%) as a grey solid. ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.91 (t, *J* = 7.3 Hz, 3H), 1.62 (quin, *J* = 7.4 Hz, 2H), 2.56 (t, *J* = 7.4 Hz, 2H), 3.35 (s, 3H), 6.36 (d, *J* = 8.2 Hz, 1H), 6.77 (d, *J* = 8.0 Hz, 1H), 6.93 (d, *J* = 8.0 Hz, 1H), 7.00–6.98 (m, 2H), 7.16 (t, *J* = 8.0 Hz, 1H), 8.38 ppm (s, 2H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 13.7, 24.2, 37.2, 55.7, 112.6, 120.8, 121.7, 121.8, 128.5, 130.1, 135.7, 139.9, 142.0, 151.2, 160.6 ppm. A mixture of this boronic acid intermediate (0.34 g, 1.2 mmol) and BBr₃ (1M, 6.0 mL, 6.0 mmol) were subjected to the general demethylation

procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO₂, 2% MeOH/chloroform) to give analytically pure **20** as a grey solid (72%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.89 (t, *J* = 7.1 Hz, 3H), 1.57 (quin, *J* = 7.2 Hz, 2H), 2.48 (t, *J* = 7.1 Hz, 2H), 6.65 (d, *J* = 7.4 Hz, 1H), 6.82–6.78 (m, 3H), 6.91 (d, *J* = 8.0 Hz, 1H), 7.03 (d, *J* = 7.6 Hz, 1H), 7.29 ppm (t, *J* = 8.1 Hz, 1H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 13.7, 24.0, 36.9, 114.5, 115.6, 117.3, 119.7, 121.5, 122.2, 130.9, 133.7, 139.4, 140.4, 149.1, 159.4 ppm.

4-(2,6-Dihydroxy-4-propylphenoxy)benzamide (28)—A mixture of the (2,6-dimethoxy-4-propylphenoxy)benzamide (0.60 g, 1.8 mmol) whose synthesis is described in the preparation of **32** above, H₂O₂ (38%, 1.5 mL, 12.4 mmol) and 3N NaOH (0.3 mL) in ethanol (10 mL) was refluxed with stirring (20 h). After cooling, the reaction mixture was acidified with dilute hydrochloric acid. The precipitate was filtered, dried to obtain **4-(2,6-dimethoxy-4-propylphenoxy)benzamide** as a white solid. A mixture of this intermediate compound (0.15 g, 0.48 mmol) and BBr₃ (1M, 3.4 mL, 3.4 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO₂, 10% MeOH/chloroform). The product was recrystallized from MeOH/chloroform to give analytically pure **28** as white prisms (74%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.91 (t, *J* = 7.2 Hz, 3H), 1.55 (quin, *J* = 7.2 Hz, 2H), 2.40 (t, *J* = 7.2 Hz, 2H), 6.26 (s, 2H), 6.78 (d, *J* = 8.4 Hz, 2H), 7.18 (br s, 1H), 7.80–7.78 (m, 3H), 9.28 ppm (s, 2H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 14.2, 24.3, 37.6, 107.9, 114.6, 127.5, 129.5, 140.0, 150.9, 161.1, 167.9 ppm.

5-Propyl-2-(4-(2H-tetrazol-5-yl)phenoxy)-3-hydroxyphenol (35)—A mixture of the **4-(2,6-dimethoxy-4-propylphenoxy)benzamide** (0.18 g, 0.6 mmol) (from above), SiCl₄ (0.20 g, 1.2 mmol) and NaN₃ (0.15 g, 2.3 mmol) was refluxed at 90 °C in CH₃CN (0.5 mL) for 6 h under Ar. After cooling, the reaction mixture was diluted with EtOAc, washed with saturated sodium bicarbonate solution, the organic layer was separated and washed with water followed by brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (8% MeOH/chloroform) to result in **5-(4-(2,6-dimethoxy-4-propylphenoxy)phenyl)-2H-tetrazole** (0.18 g, 97%) as a grey solid. ¹H NMR (400 MHz, CDCl₃): δ = 0.96 (t, *J* = 7.2 Hz, 3H), 1.66 (quin, *J* = 7.4 Hz, 2H), 2.57 (t, *J* = 7.6 Hz, 2H), 3.72 (s, 6H), 6.45 (s, 2H), 6.88 (d, *J* = 8.1 Hz, 2H), 8.01 (d, *J* = 8.2 Hz, 2H), 12.56 ppm (br s, 1H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 13.9, 24.2, 37.9, 55.9, 105.5, 114.9, 121.1, 128.2, 128.5, 140.6, 152.4 ppm. A mixture of this intermediate compound (0.17 g, 0.50 mmol) and BBr₃ (1M, 4.0 mL, 4.0 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO₂, 10% MeOH/chloroform) to give analytically pure **35** as off-white solid (78%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 0.91 (t, *J* = 7.2 Hz, 3H), 1.56 (quin, *J* = 7.2 Hz, 2H), 2.40 (t, *J* = 7.6 Hz, 2H), 6.28 (s, 2H), 6.97 (d, *J* = 8.0 Hz, 2H), 7.94 (d, *J* = 8.4 Hz, 2H), 9.34 ppm (s, 2H); ¹³C NMR (100 MHz, (CD₃)₂SO): δ = 14.2, 24.3, 37.6, 107.9, 116.1, 127.4, 128.9, 140.1, 150.9, 161.0 ppm. HRMS (ESI-positive): calcd for C₁₆H₁₆N₄O₃ ([M+H]⁺): 313.1295, found: 313.1296.

Hexa-2,4-dienoic acid (4-(4-chloro-2-hydroxyphenoxy)-phenyl)amide (25)—A mixture of **4-(4-chloro-2-methoxyphenoxy)phenylamine**²¹ (0.19 g, 0.8 mmol), EDAC·HCl (0.15 g, 0.8 mmol) and HOBt (0.1 g, 0.8 mmol) in 8 mL DMF was stirred for 10 minutes at rt under Ar. To this, a mixture of sorbic acid (0.09 g, 0.8 mmol) and triethylamine (0.08 g, 0.8 mmol) in 2 mL of DMF was added at rt and the reaction mixture was stirred at rt. After 16 h, the reaction mixture was diluted with EtOAc, washed with saturated sodium bicarbonate solution, the organic layer was separated and washed with water followed by

brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na_2SO_4 , concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (3% MeOH/chloroform) to result in **hexa-2,4-dienoic acid (4-(4-chloro-2-methoxyphenoxy)-phenyl)amide** (0.18 g, 67%) as a grey solid. $^1\text{H NMR}$ (300 MHz, CDCl_3): δ = 1.87 (d, J = 7.1 Hz, 3H), 3.85 (s, 3H), 5.89 (d, J = 19.7 Hz, 1H), 6.20 (dd, J = 19.7, 6.5 Hz, 2H), 7.03–6.89 (m, 5H), 7.42–7.32 (m, 2H), 7.53–7.50 ppm (m, 1H). A mixture of this intermediate compound (0.18 g, 0.50 mmol) and BBr_3 (1M, 1.5 mL, 1.5 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO_2 , 2% MeOH/chloroform) to give analytically pure **25** as a brown gel (76%). $^1\text{H NMR}$ (300 MHz, CDCl_3): δ = 1.88 (d, J = 6.9 Hz, 3H), 6.18–5.74 (m, 2H), 6.22–6.16 (m, 2H), 6.80–6.77 (m, 2H), 7.07–6.98 (m, 3H), 7.54 ppm (d, J = 11.2 Hz, 2H); $^{13}\text{C NMR}$ (100 MHz, CD_3OD): δ = 17.2, 116.7, 117.1, 117.2, 119.3, 121.0, 121.4, 121.5, 121.6, 129.0, 129.7, 137.9, 141.7, 149.7, 154.0 ppm. HRMS (ESI-positive): calcd for $\text{C}_{18}\text{H}_{16}\text{ClNO}_3$ ($[\text{M}+\text{H}]^+$): 330.0891, found: 330.0883.

Hexa-2,4-dienoic acid (4-(2,4-dichlorophenoxy)-3-hydroxyphenyl)amide (27)—

A mixture of 2,4-dichlorophenol (1.0 g, 6.3 mmol), KOtBu (0.83 g, 7.5 mmol), 2-iodo-5-nitroanisole (1.9 g, 6.9 mmol), and $(\text{CuOTf})_2 \text{PhCH}_3$ (0.16 g, 0.3 mmol) in DMF (11 mL) were reacted according to the Method B above. Usual workup and purification by flash chromatography (SiO_2 , 5% EtOAc/Hexanes) gave analytically pure **4-(2,4-dichlorophenoxy)-3-methoxynitrobenzene** as a brown oil (62%). $^1\text{H NMR}$ (400 MHz, $(\text{CD}_3)_2\text{SO}$): δ = 4.14 (s, 3H), 6.36 (d, J = 4.8 Hz, 1H), 7.13 (d, J = 5.2 Hz, 1H), 7.46 (dd, J = 8.8, 2.4 Hz, 1H), 7.61 (dd, J = 4.2, 2.4 Hz, 1H), 7.69 (d, J = 2.4 Hz, 1H), 8.09 (d, J = 8.4 Hz, 2H); $^{13}\text{C NMR}$ (100 MHz, $(\text{CD}_3)_2\text{SO}$): δ = 57.5, 105.9, 108.7, 117.3, 117.6, 118.1, 122.2, 125.7, 129.5, 129.7, 130.7, 140.2, 150.2 ppm. A suspension of the above compound (2.0 g, 6.5 mmol), and 10% Pd/C (0.32 g) in EtOH (25 mL) was stirred under hydrogen atmosphere at 40 PSI at room temperature for 16 h. The catalyst was removed by filtration through a pad of Celite, and the residue was thoroughly washed with EtOAc. The solvent was evaporated and purification by flash chromatography (SiO_2 , 20% EtOAc/Hexanes) gave the compound **4-(2,4-dichlorophenoxy)-3-methoxyaniline** (0.7 g, 38%). $^1\text{H NMR}$ (400 MHz, CDCl_3): δ = 3.48 (br s, 2H), 3.69 (s, 3H), 6.27 (dd, J = 4.2, 2.4 Hz, 1H), 6.36 (d, J = 2.4 Hz, 1H), 6.61 (d, J = 8.8 Hz, 1H), 6.85 (d, J = 8.4 Hz, 1H), 7.06 (dd, J = 4.4, 2.3 Hz, 1H), 7.42 ppm (d, J = 2.4 Hz, 1H); $^{13}\text{C NMR}$ (100 MHz, CDCl_3): δ = 55.4, 100.2, 106.7, 116.2, 122.4, 122.9, 126.1, 127.1, 129.5, 144.7, 151.7, 153.2 ppm. A mixture of this aniline intermediate (0.15 g, 0.5 mmol), EDAC-HCl (0.10 g, 0.5 mmol) and HOBt (0.07 g, 0.5 mmol) in 6 mL DMF was stirred for 10 minutes at rt under Ar. To this, a mixture of sorbic acid (0.06 g, 0.5 mmol) and triethylamine (0.08 g, 0.8 mmol) in 2 mL of DMF was added at rt and the reaction mixture was stirred at rt. After 16 h, the reaction mixture was diluted with EtOAc, washed with saturated sodium bicarbonate solution, the organic layer was separated and washed with water followed by brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na_2SO_4 , concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (3% MeOH/chloroform) to result in **hexa-2,4-dienoic acid (4-(2,4-dichlorophenoxy)-3-methoxyphenyl)amide** (0.18 g, 93%) as brown liquid. $^1\text{H NMR}$ (400 MHz, $(\text{CD}_3)_2\text{SO}$): δ = 1.84 (d, J = 6.2 Hz, 3H), 3.34 (s, 3H), 6.35–6.09 (m, 3H), 6.64 (d, J = 8.6 Hz, 1H), 7.04 (d, J = 8.6 Hz, 1H), 7.28–7.15 (m, 3H), 7.65 (d, J = 8.6 Hz, 2H), 10.15 ppm (s, 1H); $^{13}\text{C NMR}$ (100 MHz, $(\text{CD}_3)_2\text{SO}$): δ = 18.8, 56.0, 105.1, 112.0, 117.5, 122.2, 122.9, 123.2, 126.4, 128.7, 130.1, 130.3, 138.3, 138.5, 141.4, 151.2, 153.2, 164.4 ppm. MS (ESI) m/z : 402.1 $[\text{M} + \text{H}]^+$ A mixture of this intermediate compound (0.12 g, 0.33 mmol) and BBr_3 (1M, 1.6 mL, 1.6 mmol) were subjected to the general demethylation procedure outlined in Method C above. The crude product was purified by flash chromatography (SiO_2 , 3% MeOH/chloroform) to give analytically pure **27** as a brown oil (76%). $^1\text{H NMR}$ (400 MHz, $(\text{CD}_3)_2\text{SO}$): δ = 1.83

(d, $J = 6.2$ Hz, 3H), 6.33–6.09 (m, 2H), 6.64 (d, $J = 8.6$ Hz, 1H), 6.96 (d, $J = 8.6$ Hz, 1H), 7.06 (d, $J = 8.4$ Hz, 1H), 7.19–7.13 (m, 1H), 7.28 (d, $J = 8.6$ Hz, 1H), 7.55 (s, 1H), 7.65 (s, 1H), 10.02 ppm (s, 1H); ^{13}C NMR (100 MHz, $(\text{CD}_3)_2\text{SO}$): $\delta = 18.8, 108.7, 111.0, 117.5, 122.4, 122.8, 123.4, 126.1, 128.6, 129.9, 130.3, 137.4, 137.9, 138.2, 141.2, 149.4, 153.3, 164.2$ ppm.

5-Chloro-2-phenoxy-N-pivaloyl aniline (13)—A suspension of commercially available 5-chloro-2-phenoxyaniline (0.13 g, 0.61 mmol) and Et_3N (0.15 g, 1.5 mmol) in 2 mL methylene chloride at 0°C was treated with pivaloyl chloride (0.07 g, 0.61 mmol), and the reaction mixture was stirred for 3h. The reaction mixture was diluted with ethyl acetate, extracted and the combined organic layers were washed with brine. Purification by flash chromatography (SiO_2 , 5% EtOAc/Hexanes) gave analytically product. ^1H NMR (400 MHz, CD_3OD): $\delta = 1.14$ (s, 9H), 6.97 (t, $J = 8.5$ Hz, 3H), 7.12 (t, $J = 7.4$ Hz, 3H), 7.18 (dd, $J = 4.4, 2.4$ Hz, 3H), 7.35 ppm (t, $J = 8.2$ Hz, 3H); ^{13}C NMR (100 MHz, CD_3OD): $\delta = 25.7, 38.8, 99.6, 116.5, 120.3, 122.9, 123.6, 124.8, 128.4, 129.3, 130.4, 145.9, 156.4, 177.7$ ppm. HRMS (ESI-positive): calcd for $\text{C}_{17}\text{H}_{18}\text{ClNO}_2$ ($[\text{M}+\text{H}]^+$): 304.1099, found: 304.1098.

2-Amino-N-(5-chloro-2-phenoxyphenyl)nicotinamide (7)—A mixture of 2-aminonicotinic acid (0.27 g, 1.9 mmol), EDAC·HCl (0.38 g, 1.9 mmol) and HOBt (0.26 g, 1.9 mmol) in 15 mL DMF was stirred for 10 minutes at rt under Ar. To this, a mixture of commercially available 5-chloro-2-phenoxyaniline (0.43 g, 1.9 mmol) and triethylamine (0.2 g, 1.9 mmol) in 5 mL of DMF was added at rt and the reaction mixture was stirred at rt. After 16 h, DMF was evaporated, the reaction mixture was diluted with EtOAc, washed with saturated sodium bicarbonate solution. The organic layer was separated and washed with water followed by brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na_2SO_4 , concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (1% MeOH/chloroform) to result in **7**. ^1H NMR (400 MHz, CDCl_3): $\delta = 6.46$ (br s, 2H), 6.63–6.59 (m, 1H), 6.83 (d, $J = 8.7$ Hz, 1H), 7.08–7.02 (m, 3H), 7.21 (t, $J = 7.4$ Hz, 1H), 7.41 (t, $J = 8.2$ Hz, 2H), 7.59–7.57 (m, 1H), 8.21–8.13 (m, 1H), 8.40 (br s, 1H), 8.61 ppm (d, $J = 2.2$ Hz, 1H); ^{13}C NMR (100 MHz, $(\text{CD}_3)_2\text{SO}$): $\delta = 109.7, 111.8, 118.6, 121.1, 124.0, 126.4, 126.5, 127.5, 130.4, 131.0, 137.7$ ppm. HRMS (ESI-positive): calcd for $\text{C}_{18}\text{H}_{14}\text{ClN}_3\text{O}_2$ ($[\text{M}+\text{H}]^+$): 340.0847, found: 340.0851.

5-chloro-2-(2,4-dichlorophenoxy)phenyl pivaloate (3)—To a mixture of **1** (0.26 g, 0.9 mmol) and DMAP (0.02 g, 0.2 mmol) in 8 mL of methylene chloride at 0°C was added Et_3N (0.10 g, 1.0 mmol) and stirred for 10 minutes. To this, pivaloyl chloride (0.12 g, 1.0 mmol) was added at 0°C and the reaction mixture was stirred for 1h. The reaction mixture was stirred for an additional 2 h at rt. After evaporation of methylene chloride, the reaction mixture was diluted with ethyl acetate, washed with water followed by brine. The combined organic layers were extracted by ethyl acetate, dried over Na_2SO_4 and concentrated under vacuum. Purification by flash chromatography (SiO_2 , 3% EtOAc/Hexanes) gave analytically product **3** (85%). ^1H NMR (400 MHz, CDCl_3): $\delta = 1.25$ (s, 9H), 6.81 (d, $J = 8.8$ Hz, 1H), 7.21–7.18 (m, 3H), 7.46 ppm (d, $J = 2.4$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3): $\delta = 26.5, 38.7, 118.7, 120.5, 124.1, 124.8, 126.4, 127.6, 128.5, 129.3, 129.9, 142.2, 145.6, 150.9, 175.6$ ppm.

3-(6-Aminopyridin-3-yl)-N-(4-methoxyphenyl)acrylamide (52)—A mixture of 3-(6-aminopyridin-3-yl)-acrylic acid²³ (0.12 g, 0.7 mmol), EDAC·HCl (0.13 g, 0.7 mmol) and HOBt (0.10 g, 0.7 mmol) in 15 mL DMF was stirred for 10 minutes at rt under Ar. To this, a mixture of *p*-anisidine (0.09 g, 0.7 mmol) and triethylamine (0.07 g, 0.7 mmol) in 5 mL of DMF was added at rt and the reaction mixture was stirred at rt. After 16 h, DMF was

evaporated, the reaction mixture was diluted with EtOAc, washed with saturated sodium bicarbonate solution. The organic layer was separated and washed with water followed by brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum, and the crude product was purified by flash chromatography on *silica gel* (5% MeOH/chloroform) to result in **52**. ¹H NMR (400 MHz, CD₃OD): δ = 3.79 (s, 3H), 6.56 (d, *J* = 15.6 Hz, 1H), 6.62 (d, *J* = 15.6 Hz, 1H), 6.90 (d, *J* = 9.0 Hz, 1H), 7.50 (s, 1H), 7.55 (d, *J* = 9.1 Hz, 2H), 7.75 (dd, *J* = 4.4, 2.2 Hz, 1H), 8.08 ppm (d, *J* = 1.8 Hz, 1H); ¹³C NMR (100 MHz, CD₃OD): δ = 54.1, 108.6, 113.2, 115.7, 119.7, 121.0, 131.3, 135.1, 137.6, 148.0, 156.1, 159.9, 164.9 ppm. HRMS (ESI-positive): calcd for C₁₅H₁₅N₃O₂ ([M+H]⁺): 270.1237, found: 270.1239.

N-(2-Aminobenzyl)-3-(6-aminopyridin-3-yl)acrylamide (53)—A mixture of 3-(6-aminopyridin-3-yl)-acrylic acid²³ (0.13 g, 0.8 mmol), EDAC·HCl (0.15 g, 0.8 mmol), HOBT (0.10 g, 0.8 mmol), 2-aminobenzylamine (0.09 g, 0.8 mmol) and triethylamine (0.08 g, 0.8 mmol) in 15 mL DMF were reacted according to the procedure described above for **52**. ¹H NMR (400 MHz, CD₃OD): δ = 4.42 (s, 2H), 6.41 (d, *J* = 15.6 Hz, 1H), 6.62 (d, *J* = 15.7 Hz, 1H), 6.60 (d, *J* = 8.8 Hz, 1H), 6.68 (t, *J* = 7.3 Hz, 1H), 6.75 (d, *J* = 7.9 Hz, 1H), 7.06 (t, *J* = 7.5 Hz, 1H), 7.11 (d, *J* = 7.4 Hz, 1H), 7.45 (d, *J* = 15.7 Hz, 1H), 7.72 (d, *J* = 8.6 Hz, 1H), 8.04 ppm (s, 1H); ¹³C NMR (100 MHz, CD₃OD): δ = 39.4, 108.7, 115.4, 116.1, 117.2, 119.7, 122.2, 127.9, 129.3, 135.2, 137.2, 145.1, 147.7, 159.8, 167.1 ppm. HRMS (ESI-positive): calcd for C₁₅H₁₆N₄O ([M+H]⁺): 269.1397, found: 269.1402.

1-(4-Nitrophenyl)-1,3-dihydrobenzimidazol-2-one (46)—1,3-Dihydrobenzimidazol-2-one (0.50 g, 3.7 mmol), 1-fluoro-4-nitrophenol (0.53 g, 3.7 mmol) and K₂CO₃ (1.00 g, 7.5 mmol) in DMSO (8 mL) were heated to 100 °C under nitrogen for 12 h (Method A above). The reaction mixture was cooled to rt and filtered. The yellow solid was repeatedly washed with water (3 × 5 mL), EtOAc (3 × 2 mL), ether (3 × 2 mL) and finally with hexanes (3 × 2 mL). The solid was dried in vacuum to obtain analytically pure **46** (0.70 g, 74%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 6.92 (s, 2H), 7.26 (br s, 1H), 7.34 (br s, 1H), 8.00 (d, *J* = 8.5 Hz, 2H), 8.48 ppm (d, *J* = 8.7 Hz, 1H).

1-(4-Nitrobenzoyl)-1,3-dihydrobenzimidazol-2-one (48)—To a mixture of 1,3-dihydrobenzimidazol-2-one (0.51 g, 3.8 mmol) in 9 mL of anhydrous pyridine at 0 °C was added 4-nitrobenzoyl chloride (1.00 g, 5.6 mmol) portion wise and the reaction mixture was stirred for 15 minutes. The contents were then refluxed at 80 °C for 4 h. The reaction mixture solidified when cooled to rt, to which, 10 mL water and 20 mL of EtOAc were added and stirred for 10 minutes. The organic layer was separated and washed with 1N HCl followed by water and brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum to result in **48** (1.50 g, 98%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 6.91 (s, 2H), 8.16 (d, *J* = 8.5 Hz, 2H), 8.31 (d, *J* = 8.6 Hz, 1H), 10.59 ppm (s, 1H); ¹³C NMR (100 MHz, CDCl₃): δ = 108.9, 120.8, 124.1, 130.1, 131.1, 136.9, 150.0, 150.4, 155.7, 166.3 ppm.

1-(4-Nitrobenzyl)-1,3-dihydrobenzimidazol-2-one (45)—1,3-Dihydrobenzimidazol-2-one (0.20 g, 1.5 mmol), 4-nitrobenzyl bromide (NOTE: strong lachrymator) (0.32 g, 1.5 mmol) and K₂CO₃ (0.31 g, 2.2 mmol) in MeOH (6 mL) were refluxed at 100 °C under nitrogen for 4 h. The reaction mixture was cooled to rt and MeOH was evaporated, 10 mL water and 20 mL of EtOAc were added and stirred for 10 minutes. The organic layer was separated and washed with water followed by brine. The aqueous layer was extracted twice with ethyl acetate. The combined organic layers were then dried over Na₂SO₄, concentrated under vacuum and the crude product was purified by flash chromatography on *silica gel* (15% EtOAc/Hexanes) to result in **45**. ¹H NMR (400 MHz,

(CD₃)₂SO): δ = 4.98 (s, 2H), 6.98 (d, J = 8.5 Hz, 2H), 7.15 (d, J = 8.5 Hz, 2H), 8.28 (d, J = 8.6 Hz, 2H), 11.38 ppm (s, 1H); ¹³C NMR (100 MHz, CDCl₃): δ = 43.8, 108.4, 108.9, 109.5, 120.8, 121.1, 121.7, 122.0, 154.7 ppm. HRMS (ESI-positive): calcd for C₁₄H₁₁N₃O₃ ([M+H]⁺): 270.0873, found: 270.0875.

1-(4-Nitrobenzenesulfonyl)-1,3-dihydrobenzimidazol-2-one (47)—To a mixture of 1,3-dihydrobenzimidazol-2-one (0.50 g, 3.7 mmol) in 8 mL of methylene chloride and 2 mL of DMF at 0 °C was added Et₃N (0.42 g, 4.1 mmol) and stirred for 10 minutes. To this, 4-nitrobenzenesulfonyl chloride (0.83 g, 3.73 mmol) was added at 0 °C and the reaction mixture was stirred for 15 minutes. The reaction mixture was stirred for an additional 2 h at rt. The reaction mixture was quenched by adding crushed ice. The precipitated yellow solid was repeatedly washed with water (3 × 5 mL), EtOAc (3 × 2 mL), ether (3 × 2 mL) and finally with hexanes (3 × 2 mL). The solid was dried in vacuum to obtain analytically pure **46** (0.70 g, 74%). ¹H NMR (400 MHz, (CD₃)₂SO): δ = 6.91 (s, 2H), 7.85 (d, J = 8.4 Hz, 2H), 8.20 (d, J = 8.4 Hz, 2H), 10.56 ppm (s, 1H); ¹³C NMR (100 MHz, CDCl₃): δ = 108.6, 120.5, 123.4, 127.0, 129.7, 155.3 ppm.

Biology

Testing of inhibitors *in vitro* against *T. gondii* Tachyzoites—Four-day old confluent cultures of human foreskin fibroblasts (HFF) were infected with 10³ tachyzoites of RH strain of *T. gondii* and cultured for 1 hour to allow parasite invasion, inhibitors added and cells cultured for 3 days, supplemented with ³H uracil and incubated for a further day, whereupon uracil incorporation into cells and thus parasite growth were assessed by liquid scintillation counting.⁴⁷ Lack of toxicity for mammalian host cells was demonstrated first by visual inspection of monolayers and in separate ³H thymidine incorporation assays using non-confluent cell monolayers.

Cloning, Sequencing, Overexpression, and Purification of TgENR—These were performed as described previously.^{14,48}

TgENR inhibitor assay—Activity of TgENR was assayed by monitoring consumption of NADH ($\epsilon_{340}=6220\text{M}^{-1}\text{cm}^{-1}$) with a scanning spectrophotometer. ENR enzymes catalyze the conversion of trans-2-acyl-ACP to acyl-ACP, however, they are typically assayed with a surrogate substrate, trans-2-butyryl-Coenzyme A (crotonyl-CoA).¹⁸ We used a reaction mixture which would contain 20nM TgENR, 100mM Na/K phosphate buffer pH7.5, 150mM NaCl, and 100 μ M NADH (Sigma) when diluted to a final volume of 50 μ L. Compounds dissolved in DMSO were added to this mixture (2% final concentration of DMSO) and reactions were initiated at 25°C by addition of 100 μ M crotonyl-CoA (Sigma). This assay had also previously been used to monitor TgENR inhibition by an octaarginine tagged triclosan compound over a 24-hour period. Hydrolysis of the octaarginine tag released triclosan, leading to decreasing IC₅₀ values (11nM at 24 hours) over the time course of the experiment¹⁰. Nonlinear regression analysis was performed using GraphPad Prism software

Co-crystallization and structure determination of TgENR in complex with NAD⁺ and compound 19. Preparation of protein crystals—Crystals of TgENR in complex with NAD⁺ and compound **19** were grown using the hanging drop vapor diffusion technique at 290K and by screening around the conditions which had previously been shown to be appropriate for crystallizing the TgENR/NAD⁺/triclosan complex (0.1M Tris-HCL pH9.0 and 6% PEG 8,000).¹⁴ In order to co-crystallize the TgENR with compound **19** the poor solubility was overcome by dissolving the inhibitor in DMSO and then diluting with the appropriate buffer for co-crystallization experiments.

Data collection and image processing—Crystals of both the TgENR/NAD⁺/compound **19** complex were flash frozen in 20% glycerol and data were collected at 100K to 2.7Å at the Daresbury SRS. The data were subsequently processed using the DENZO/SCALEPACK package (Table 1).⁴⁹ The structure of the complex was determined by molecular replacement using the program Amore and the TgENR/NAD⁺/triclosan structure as a search model from which the triclosan and NAD⁺ coordinates were omitted. A clear solution was obtained in Amore and the model was subjected to rigid-body refinement and was subsequently rebuilt and refined in an iterative process using REFMAC5⁵⁰ and WinCoot. Although omitted in the search model, clear and continuous density could be seen for both the NAD⁺ cofactor and compound **19** allowing for their unambiguous fitting. Data collection and model refinement statistics can be seen in Table 1.

Testing of inhibitors *in vivo*—Compounds effective in enzyme assays and then *in vitro* (IC₅₀ <10µM) were tested in an acute parenteral infection.¹⁰ CD1 mice were infected intraperitoneally with 2000 *T. gondii* (RH strain, Type 1). This inoculation dose is capable of killing one hundred percent of mice by day 10 post-infection. Mice were administered inhibitors intraperitoneally at 10mg/kg doses. Protection was measured by determining parasite burdens in the peritoneal cavity by microscopy at day 5 post-infection and survival up to 5 days post infection.

ADMET assays—Cyp450 inhibition and human liver microsomal stability assays were performed as is standard.

Statistical Analyses—Sample size and number of experiments. There were 6 replicate samples per group for *in vitro* experiments and 6 mice for each experimental group. All experiments were performed with sufficient sample sizes to have an 80% power to detect differences at the 5% level of significance. Groups included untreated or controls. Results were compared using students T test, Chi square analysis or Fisher's exact test.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

Supported by NIAID U01 AI082180-01. We gratefully acknowledge support of this work by gifts from the Finley John Gubbins Samuel Special Needs Trust, R Blackfoot, R Thewind, A Akfortseven, S Gemma, S Jackson, AK Bump, S. Pettican, the Rooney Aldens, the Dominique Cornwell and Peter Mann Family Foundation, the Morel, Rosenstein, Kapnick and Kiewit families, Intervet/Schering Plough, Toxoplasmosis Research Institute, and The Research to Prevent Blindness Foundation. This work was also supported by the Johns Hopkins Malaria Research Institute and the Bloomberg Family Foundation (JLZ and STP). We thank J. McCammon and Dr. Marco Pieroni for their assistance in preparation of this manuscript.

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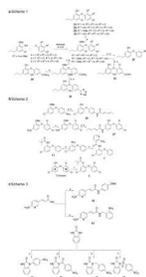
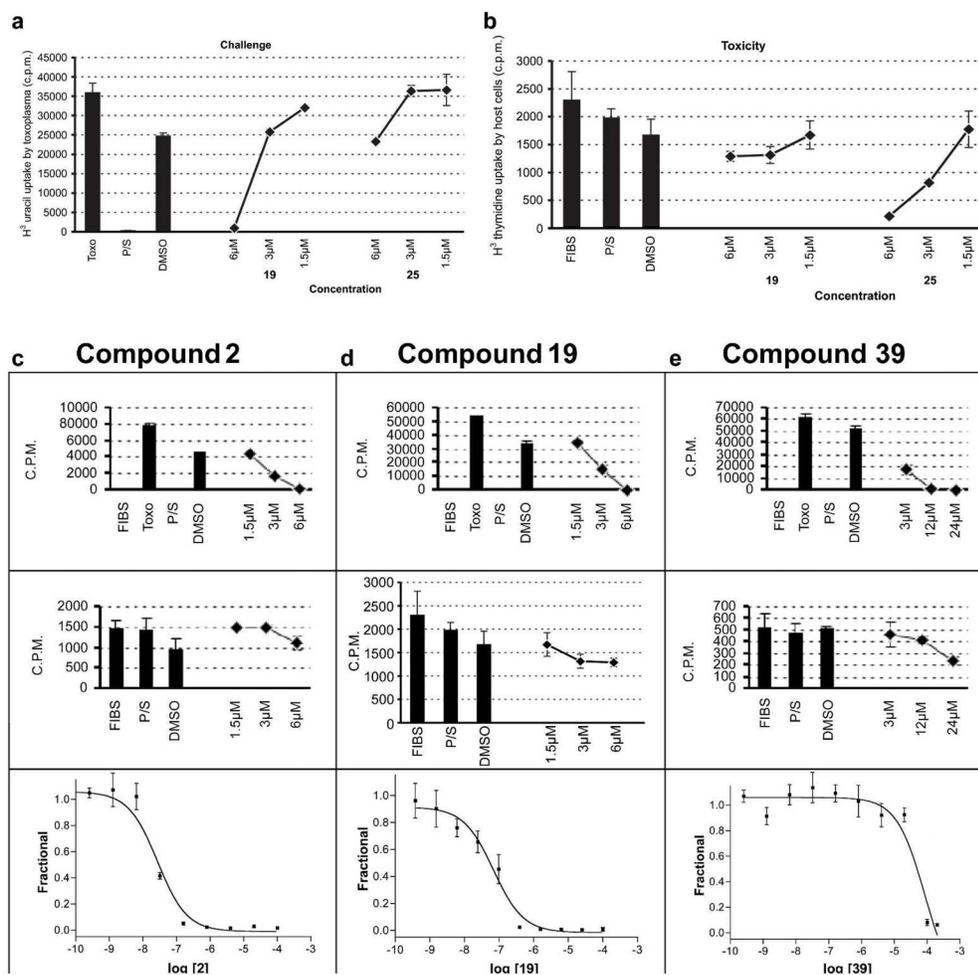


Figure 2.

(a) Synthesis of compound in 3 schemes: **Scheme 1.** Synthesis of compounds: When X = F and R² = Cl, R³ = H, R⁴ = CN; or when X = F and R² = R³ = H, R⁴ = CN, Method A: K₂CO₃ or Cs₂CO₃, DMSO, 100 °C, 8–12 h. When X = I and R² = R⁴ = H, R³ = Cl, Method B: KOtBu, DMF, (CuOTf)₂-PhH, 140 °C, 16–20 h. a) Excess BBr₃, CH₂Cl₂, -78 °C to rt, 2–6 h; b) n-BuLi, B(OMe)₃, 2N HCl, -78 °C to rt, 18 h; c) 35% H₂O₂, 3N NaOH, EtOH, 30 °C, 18 h; d) 25% NaOH, EtOH, reflux, 20 h; e) NaN₃, SiCl₄, CH₃CN, 90 °C, 6 h; **(b)** **Scheme 2.** Synthesis of compounds: a) Sorbic acid, EDAC·HCl, HOBt, Et₃N, DMF, rt, 16 h; b) Excess BBr₃, CH₂Cl₂, -78 °C to rt, 6 h; c) Pd/C, H₂, 40 PSI, TEA, EtOH, rt, 16 h; d) Pivaloyl chloride, Et₃N, THF, rt, 4 h; e) 2-Aminonicotinic acid, EDAC·HCl, HOBt, Et₃N, DMF, rt, 16 h; f) Pivaloyl chloride, Et₃N, DMAP, CH₂Cl₂, 0 °C to rt, 6 h; **(c) Scheme 3.** Synthesis of compounds: a) p-Anisidine, EDAC·HCl, HOBt, Et₃N, DMF, rt, 20 h; b) 2-Aminobenzylamine, EDAC·HCl, HOBt, Et₃N, DMF, rt, 16 h; c) 4-Fluoronitrobenzene, K₂CO₃, DMSO, 100 °C, 16 h; d) 4-Nitrobenzoyl chloride, pyridine, 0–80 °C, 4 h; e) 4-Nitrobenzyl bromide, K₂CO₃, MeOH, 100 °C, 4 h; f) 4-Nitrobenzenesulfonyl chloride, Et₃N, DMF, 0 °C–rt, 4 h.

**Figure 3.**

(a,b) Representative experiments showing effects of compounds **25** and **19** on parasite uptake of tritiated uracil and on growth of nonconfluent host cell fibroblasts. (a) Inhibition of *T. gondii* uptake of tritiated uracil by compound **19** and lack of inhibition by compound **25**. (b) Lack of toxicity to fibroblasts (uptake of ^3H Thymidine by nonconfluent HFF) by compound **19**. Toxicity to HFF from compound **25**. (c,d,e) Representative experiments showing inhibition of *T. gondii* tachyzoites (top panels), lack of inhibition of fibroblast growth (middle panels) and inhibition of ENR enzyme activity (bottom panels) by compounds **2**, **19**, and **39**. Error bars in panels A–E show the standard deviation between triplicate measurements. Compound **2** IC_{50} = 45nM and **19** = 20nM, and **39** less active (low micro molar), possibly off target.

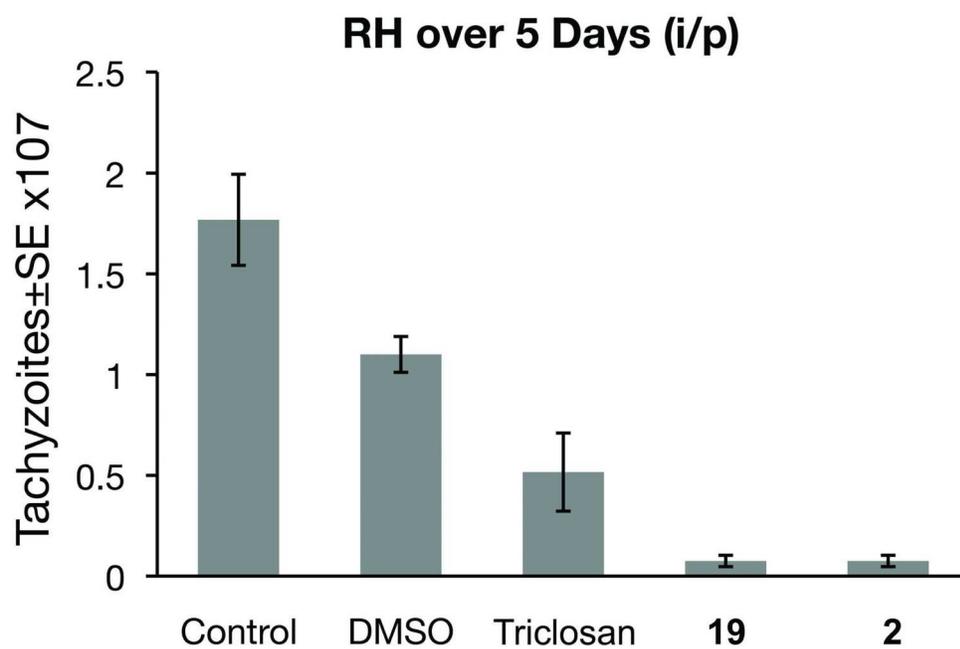


Figure 4. Reduction of parasite burden by **2** and **19** in mice.

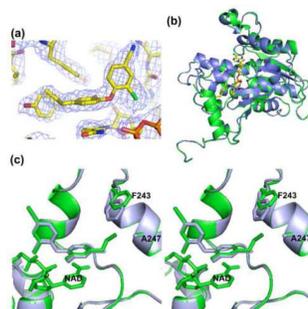


Figure 5. (a) $2F_{\text{obs}} - 1F_{\text{calc}}$ electron density map contoured at 1σ for compound **19** in complex with TgENR/NAD⁺ after refinement in REFMAC5. (b) Superposition of TgENR in complex with triclosan (blue) and compound **19** (green) shown in cartoon format, to show the conservation of the overall fold between the two complexes. Both the NAD⁺ cofactor and compound **19** are displayed in stick format and coloured yellow, red, blue, green and purple for carbon, oxygen, nitrogen, chlorine and phosphorus, respectively. (c) Stereo diagram of the TgENR triclosan (green) and compound **19** (blue) complex structures superposed. The NAD⁺ cofactor and inhibitors are shown in stick format in addition to Phe243 which display the greatest change between the different complexes.

Table 1

Data collection/processing	
Space group	P3 ₂ 21
Resolution (Å)	2.7
Unique reflections	16525
Completeness (%)	90.85
I/σ(I) >3 (%)	26
R _{merge} (%)	0.378
Refinement statistics	
R _{cryst} (%) ^a /R _{free} (%) ^b	0.26/0.32
R.m.s.d. values	
Bond length (Å)	0.0054
Bond angle (deg.)	1.00
Ramachandran plot ^c	
Most favoured (%)	93.0
Additionally allowed (%)	7.0
Generously allowed (%)	0.0
Disallowed (%)	0.0
Protein/substrate atoms/waters	4486/128/51
Mean B-values (Å ²)	
Protein	38
Co-factors	38 (compound 19) 36 (NAD)
Water molecules	43

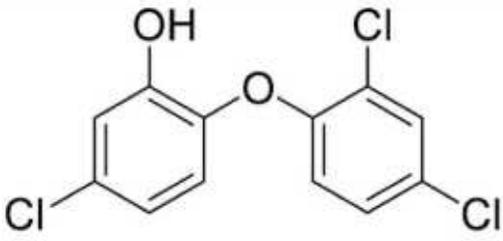
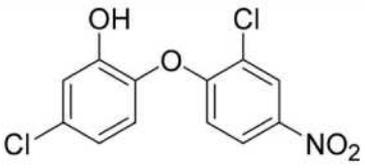
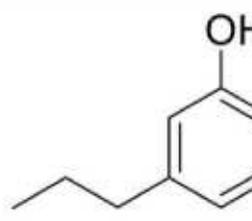
Refinement statistics for TgENR/NAD⁺/compound **19** complex.

$$^a R_{\text{cryst}} = \frac{\sum hkl (|F_{\text{obs}}| - |F_{\text{calc}}|)}{\sum hkl |F_{\text{obs}}|}$$

^b R_{free} was calculated on 5% of the data omitted randomly.

^c Percentage of residues in regions of the Ramachandran plot, according to PROCHECK.

Table 2

	 <p style="text-align: center;">Triclosan</p>	 <p style="text-align: center;">2</p>	
ClogP ^a	5.5	4.5	
ALOGpS ^b	-4.68	-4.44	
tPSA ^c	29.46	75.28	

^a calculated with ChemDraw Ultra 7.0, CambridgeSoft;

^b Predicted water solubility calculated by using ALOGPS 2.1 (www.vcclab.org/lab/alogs/);

^c total Polar Surface Area calculated by using <http://www.molinspiration.com/cgi-bin/properties>.

Reference for vcclab

Tetko, I. V.; Gasteiger, J.; Todeschini, R.; Mauri, A.; Livingstone, D.; Ertl, P.; Palyulin, V. A.; Radchenko, E. V.; Zefirov, N. S.; Makarenko, A. S.; Tanchuk, V. Y.; Prokopenko, V. V. *J. Comput. Aid. Mol. Des.*, 2005, 19, 453-63.