Title
CYP51 is an essential drug target for the treatment of primary amoebic meningoencephalitis (PAM).

Permalink
https://escholarship.org/uc/item/74r216hg

Journal
PLoS neglected tropical diseases, 11(12)

ISSN
1935-2727

Authors
Debnath, Anjan
Calvet, Claudia M
Jennings, Gareth
et al.

Publication Date
2017-12-28

DOI
10.1371/journal.pntd.0006104

Peer reviewed
CYP51 is an essential drug target for the treatment of primary amoebic meningoencephalitis (PAM)

Anjan Debnath¹, Claudia M. Calvet¹,², Gareth Jennings¹, Wexu Zhou³, Alexander Aksenov¹, Madeline R. Luth¹, Ruben Abagyan¹, W. David Nes³, James H. McKerrow¹, Larissa M. Podust¹*

¹ Center for Discovery and Innovation in Parasitic Diseases, Skaggs School of Pharmacy and Pharmaceutical Sciences, University of California San Diego, La Jolla, California, United States of America, ² Cellular Ultrastructure Laboratory, Oswaldo Cruz Institute, FIOCRUZ, Rio de Janeiro, RJ, Brazil, ³ Department of Chemistry & Biochemistry, Texas Tech University, Lubbock, Texas, United States of America

* podust@ucsd.edu

Abstract

Primary Amoebic Meningoencephalitis (PAM) is caused by Naegleria fowleri, a free-living amoeba that occasionally infects humans. While considered “rare” (but likely underreported) the high mortality rate and lack of established success in treatment makes PAM a particularly devastating infection. In the absence of economic inducements to invest in development of anti-PAM drugs by the pharmaceutical industry, anti-PAM drug discovery largely relies on drug ‘repurposing’—a cost effective strategy to apply known drugs for treatment of rare or neglected diseases. Similar to fungi, N. fowleri has an essential requirement for ergosterol, a building block of plasma and cell membranes. Disruption of sterol biosynthesis by small-molecule inhibitors is a validated interventional strategy against fungal pathogens of medical and agricultural importance. The N. fowleri genome encodes the sterol 14-demethylase (CYP51) target sharing ~35% sequence identity to fungal orthologues. The similarity of targets raises the possibility of repurposing anti-mycotic drugs and optimization of their usage for the treatment of PAM. In this work, we (i) systematically assessed the impact of anti-fungal azole drugs, known as conazoles, on sterol biosynthesis and viability of cultured N. fowleri trophozoites, (ii) identified the endogenous CYP51 substrate by mass spectrometry analysis of N. fowleri lipids, and (iii) analyzed the interactions between the recombinant CYP51 target and conazoles by UV-vis spectroscopy and x-ray crystallography. Collectively, the target-based and parasite-based data obtained in these studies validated CYP51 as a potentially ‘druggable’ target in N. fowleri, and conazole drugs as the candidates for assessment in the animal model of PAM.
Author summary

The free-living amoeba, *Naegleria fowleri*, is commonly found in water sources including swimming pools having inadequate levels of chlorine, lakes, and rivers. *N. fowleri* can act as an opportunistic pathogen causing severe brain injury called Primary Amebic Meningoencephalitis (PAM), often in healthy children and young adults. With a fatality rate over 97%, *N. fowleri* is considered one of the most deadly human pathogens. Despite the fact that precious lives are lost annually, *N. fowleri* infection is rare and economic incentives to invest in development of anti-PAM drugs by the pharmaceutical industry are lacking. In the absence of economic inducements, drug ‘repurposing’ is a cost effective strategy for re-profiling known drugs for the treatment of neglected diseases. The similarity between pathogenic fungi and free-living amoebae in the sterol biosynthesis pathway encouraged us to assess anti-mycotic drugs, clinically-approved for treatment of a variety of fungal diseases, for anti-*Naegleria* activity. Carrying out biochemical studies and x-ray-crystallography of the target enzyme, we demonstrated that azole anti-fungal drugs, known as conazoles, disrupt sterol biosynthesis in amoebae by competing with the natural substrate for binding in the active site of the sterol 14-demethylase (CYP51). Disruption of sterol biosynthesis leads to a rapid death of pathogenic amoebae.

Introduction

The amphizoic amoeba (existing both in free-living and parasitic forms), *Naegleria fowleri* is commonly found in water resources such as swimming pools having inadequate levels of chlorine, lakes and rivers. It feeds mostly on bacteria, but can also act as an opportunistic pathogen causing infection of the central nervous system (CNS) of humans and animals.[1] *N. fowleri* usually infects people when contaminated water enters the body through the nose. Following infection, *N. fowleri* infiltrates the nasal mucosa and passes along the olfactory neuroepithelial route to invade the brain. *N. fowleri* causes severe primary amebic meningoencephalitis (PAM) resulting in cerebral edema and destruction of brain tissue, mostly in healthy children and young adults.[2] PAM due to *N. fowleri* has a worldwide distribution although it occurs most frequently in tropical areas and during hot summer months.[3] Infection is considered rare in the United States (0–8 infections per year)[4] but PAM cases may go unnoticed among other infections, particularly in the developing countries.[1] Noteworthy, PAM is not on the National Notifiable Diseases Surveillance list; thus, reporting of the national incidence of PAM by the US Centers for Disease Control and Prevention (CDC) depends on individual state health departments to report diseases voluntarily. Despite modern improvements in antimicrobial therapy and supportive medical care, the fatality rate associated with *N. fowleri* PAM is >97%.[4] The disease is particularly problematic due to both its rapid onset and the lack of effective treatments.[5] Currently, there is no single, proven, evidence-based treatment with a high probability of cure. The full recovery of a patient in the summer of 2013, after 35 years without a *Naegleria* survivor in the US, was attributed to early diagnosis and treatment, and the use of combination therapy including the investigational drug miltefosine and induced hypothermia.[6] In the absence of data to estimate the true risk of PAM or to set up and reinforce the measurable standards to protect the human population, early diagnosis and aggressive antimicrobial treatment remain the only option to treat the disease.

The CDC-recommended treatment for patients suspected of PAM currently includes combination therapy consisting of anti-mycotic drugs amphotericin B (AmpB) and fluconazole, antibiotics azithromycin and rifampin, the investigational anti-cancer agent miltefosine and,
finally, an anti-inflammatory drug, dexamethasone, to reduce the cerebral edema.[1] AmpB, a cornerstone of PAM therapy and a standard of care for CNS infections caused by molds, acts via binding ergosterol in cell membranes causing rapid leakage of monovalent ions leading to cell death.[7] Clinical use of AmpB is limited due to its toxicity, including acute infusion-related reactions and dose-related nephrotoxicity.[1] Flucanazole is another anti-mycotic drug that acts via a different mechanism. Flucanazole depletes the ergosterol pool by blocking removal of the methyl group at C-14 position of a biosynthetic precursor catalyzed by sterol 14-demethylase (CYP51).[8–11] Flucanazole belongs to the ‘conazole’ pedigree of antifungal agents targeting CYP51. This drug class also includes miconazole, ketoconazole, voriconazole and itraconazole that were previously reported to exhibit amoebicidal effect against N. fowleri in vitro.[13–17] Flucanazole and miconazole have also been used in combination with other drugs as part of treatment regimens in human PAM patients.[6, 18, 19]

De novo sterol biosynthesis from squalene takes place in most high eukaryotes (although it is lost in certain lineages, e.g. insects and worms) and also in lower eukaryotes with an aerobic life style.[20] The latter include human pathogens such as fungi, kinetoplastids and free-living amoebae. Disruption of sterol biosynthesis by small-molecule inhibitors is a validated interventional strategy against fungal pathogens of medical and agricultural importance. CYP51 is one of the most extensively exploited drug targets for the development of anti-fungal agents. Kinetoplastids are major parasite targets for the development of CYP51 inhibitors outside traditional antifungal drug discovery programs.[21]

Similar to fungi and kinetoplastids, amoebae from the genera Naegleria and Acanthamoeba have an essential requirement for ergosterol.[22–24] In contrast to the lanosterol route in fungi and kinetoplastids,[25] biosynthesis of ergosterol in amoebae occurs via cycloartenol, a sterol biosynthetic precursor typical of photosynthetic organisms, i.e., algae and plants.[22–24] N. fowleri genome encodes CYP51 (NfCYP51; AmoebaDB accession number NF0102700) sharing ~35% sequence identity to human, fungal and kinetoplastid orthologues. Higher sequence identity to plant (<40%), Acanthamoeba (42%), and a non-pathogenic Naegleria gruberi (86%) is consistent with the 'plant-like' substrate specificity of NfCYP51.[23] The similarity of targets raises the possibility of 'repurposing' anti-mycotic drugs, clinically-approved against a variety of fungal diseases, and optimization of azole-based PAM therapy for treatment of PAM.

In this work, we (1) chemically validate sterol biosynthesis pathway as a “druggable” target in N. fowleri and (2) systematically assess the efficacy of the anti-fungal azole drugs, known as conazoles (including the latest additions to the armamentarium of the anti-fungal drugs, posaconazole and isavuconazole) versus both the whole organism and the recombinant molecular target. In the course of the studies, we (i) validated NfCYP51 as an essential biosynthetic enzyme in N. fowleri, (ii) determined the endogenous NfCYP51 substrate, and (iii) characterized drug-target interactions by UV-vis spectroscopy and protein x-ray crystallography.

Results and discussion

Anti-proliferative effect of conazoles in vitro

An assay previously developed and validated [26] was used to assess anti-proliferative activity of the conazoles presented in Table 1. In this assay, conazoles demonstrated anti-N. fowleri activity in a broad range of concentrations—from an EC$_{50}$ of 13.9 μM (fluconazole) to ≤0.01 μM (for itraconazole and posaconazole) (Table 1). Notably, in vitro potency of all conazole drugs exceeded that of miltefosine; itraconazole and posaconazole were an order of magnitude more potent than AmpB, while ketoconazole and isavuconazole were equipotent to AmpB. Across the board, anti-Naegleria potency of conazoles increased with the increase of the molecular
Fig 1. Azole antifungal drugs also known as conazoles.

https://doi.org/10.1371/journal.pntd.0006104.g001
weight (MW), from fluconazole (306.3 g/mol) to itraconazole (705.6 g/mol). Between 340 g/mol and 450 g/mol, anti-proliferative activity of conazoles correlated with lipophilicity per non-hydrogen atom expressed as logP/MW values—more lipophilic drugs demonstrated higher potency (Table 1). This trend is explained by the hydrophobicity of the NfCYP51 binding site favoring binding of lipophilic molecules, as demonstrated by the co-crystal structures determined in this work. The potency observed for fluconazole (EC$_{50}$ of 13.9 μM, equivalent of 4.3 mg/ml) was lower than the MIC$_{50}$ values of 0.5–2.0 mg/ml reported elsewhere for N. fowleri isolates, either from natural water sources[16] or from patients who had died of PAM.[15]

Growth inhibition as a function of time

To answer the question of how fast CYP51 inhibitors kill N. fowleri, we measured growth inhibition dose-response to posaconazole at different time points. N. fowleri trophozoites were exposed to a single dose of posaconazole serially diluted in 96-well format from 25 μM to 0.008 μM in 0.2% sulfobutylether-β-cyclodextrin (SBE-β-CD), also known as Captisol. SBE-β-CD is used as an excipient (a formulating agent) to increase the solubility of poorly soluble drugs, including posaconazole in the Noxafil intravenous formulation (Merck).[31] Growth inhibition curves constructed for different time points (Fig 2) demonstrate an inhibitory effect of posaconazole at the highest concentrations as early as 8 h post-exposure. Inhibition reaches 40% at 16 h post-exposure, ~90% at 24 h (EC$_{50}$ of 2.7 nM) and maximizes at 48 h (EC$_{50}$ of 4.9 nM).

Target engagement in N. fowleri trophozoites

Impact of posaconazole on sterol biosynthesis. To chemically validate NfCYP51 as a therapeutic drug target, we analyzed metabolites accumulated in N. fowleri in response to posaconazole. Lipids extracted from 20 or 50 million trophozoites treated with 0.1% DMSO, 0.2 μM AmpB or 0.2 μM posaconazole for 24 hours were subjected to gas chromatography-mass spectrometry (GC-MS) analysis. The sterol identities were assigned based on relative chromatographic behavior, the characteristic molecular masses and electron ionization (EI) fragmentation patterns of free sterols (Fig 3) or of the trimethylsilyl (TMS)-derivatized sterols (S1 Fig).
Similar to other previously characterized free-living amoebae of Naegleria and Acanthamoeba genera,[22–24] sterol biosynthesis in *N. fowleri* proceeds from cycloartenol (15) via 31-norlanosterol (14) to ergosterol (5) (Fig 4). Ergosterol and its biosynthetic precursor ergosta-5,7-dienol (7) are the two most abundant endogenous sterols dominating lipid extracts of DMSO-treated *N. fowleri* (48% of total sterol mass), whereas 4-monomethyl-, 4,4-dimethyl- and 14α-methylsterols are represented sparsely (Table 2). Cholesterol (1) constituting about 25% of sterol mass in both DMSO- and posaconazole-treated samples (Table 2), is likely ingested from the growth media containing fetal bovine serum (FBS) and liver extract. AmpB, a drug with a different mechanism of action used as a negative control, did not perturb the native sterol pattern (S1 Fig).

Upon treatment with posaconazole, the ratio of metabolites changed dramatically. The total content of the 14α-methylsterols increased from 6.5% to 49.1%, with the major accumulated intermediate being 31-norlanosterol, followed by its C24-C25 hydrogenated product, 4,14-dimethylcholest-8-enol (9) (Table 2). Removal of the 4β-methyl group was unaffected by posaconazole, meaning that it likely occurs prior to 14-demethylation. Based on these data,
31-norlanosterol serves an endogenous substrate of NfCYP51. Accumulation of the 4α-methylsterols in posaconazole-treated *N. fowleri*—increase from 6.8% to 48.5%—indicates that removal of the 4α-methyl group occurs downstream of CYP51. Finally, both C24 methylation catalyzed by sterol C24-methyltransferase, double bond isomerization catalyzed by sterol Δ8-Δ7-isomerase (reaction 4→8), and C22-C23 bond desaturation catalyzed by Δ22-desaturase, also occur downstream of CYP51 (reaction 7→5) (Fig 4).

**Steryl esters.** Along with the sterol intermediates, multiple steryl esters were detected in the lipid extracts (Fig 3). Steryl moieties were identified by the molecular masses calculated from the m/z values of the [M-ROH]+ fragments compared with the fragmentation patterns to

---

**Fig 3.** Gas chromatography separation of the total sterol fractions extracted from *N. fowleri* trophozoites. Chromatogram fragments from the non-derivatized DMSO- (black trace) and posaconazole (POS)-treated (red trace) sterol extracts are shown. Peaks are labeled according to Table 2. The content of 31-norlanosterol (14) remarkably increased in posaconazole-treated samples.

https://doi.org/10.1371/journal.pntd.0006104.g003
the sterol acetate standards in the NIST database (Table 3). The fatty acid identities of steryl esters have not been analyzed due to limited sample size. Upon exposure to posaconazole, the steryl ester pool declined compared to the free sterols, as judged by the steryl ester/free sterol ratio (Table 3). The composition of the steryl ester pool also changed. The notable increase in cholesteryl ester from 4.3 to 13.6\% may be an attempt to compensate for the deficit of endogenous sterols. On the contrary, the content of squalene dropped in the posaconazole-treated
Table 2. Sterol composition in *N. fowleri* treated with 0.1% DMSO or 0.2 μM posaconazole.

| Sterols                        | RTTe | % Composition |
|-------------------------------|------|---------------|
|                               |      | DMSO          | Posaconazole   |
| 4-Desmethylsterols (% of total)|      |               |               |
| 1 Cholesterol                 | 1    | 25.6          | 26.1           |
| 2 7-Dehydrocholesterol        | 1.044| 12.5          | 3.9            |
| 3 Lathosterol                 | 1.061| 0.8           |               |
| 4 Zymosterolb                 | 1.068| 0.4           |               |
| 5 Ergosterol                  | 1.114| 26.7          | 14.2           |
| 6 Ergost-5-enol               | 1.136| 1.4           | 0.7            |
| 7 Ergosta-5,7-dienol          | 1.202| 21.3          | 5.7            |
| 8 Ergost-7-enol               | 1.22 | 1.2           |               |
| 4-Monomethylsterols           |      | 6.8           | 48.5           |
| 9 4a,14a-Dimethylcholesta-8-enol| 1.082| 1.1           | 15.5           |
| 10 4a-Methylergost-8-enol     | 1.095| 3.5           | 0.4            |
| 14 31-Norlanosterol           | 1.143| 2.2           | 29.7           |
| 11 Δ24(25)-31-norlanosterol   | 1.238|               | 2.9            |
| 4,4-Dimethylsterols           |      | 3.2           | 1.0            |
| 12 24-Dihydrolanosterol       | 1.304| 0.2           | 0.2            |
| 13 24-Dihydrocycloartenol     | 1.324| 1.1           |               |
| 14 Parkeol                    | 1.384|               | 0.1            |
| 15 Cycloartenol               | 1.406| 2.0           | 0.7            |
| 14α-Methylsterols             |      | 6.5           | 49.1           |
| Δ24(25)-sterols               |      | 8.1           |               |

*a*relative retention time compared to cholesterol

*b*in bold are highlighted sterols constituting the mainstream of biosynthesis cascade in Fig 4

https://doi.org/10.1371/journal.pntd.0006104.t002

sample from 5.5 to 1.5% suggesting a regulatory loop signaling excess of the downstream intermediates.

Table 3. Composition of the steryl ester pool in *N. fowleri*.

| Steryl ester* | [M-ROH]* | RT (min) | % of Total |
|---------------|-----------|----------|------------|
|               |           | DMSO     | Posaconazole|
| 7-Dehydrocholesterol ester-1 | 366       | 9.3      | 6.1          | 5.7          |
| Squaleneb      |           | 410      | 9.5        | 5.5          | 1.5          |
| Ergosteryl ester|           | 378      | 9.7        | 15.9         | 16.1         |
| 5,7,24-Triene-ergosteryl ester-1 | 378     | 10.1     | 11.6        | 15.6         |
| 5,7-Diene-ergosteryl ester-1   | 380      | 10.2     | 13.0        | 8.2          |
| Cholesteryl ester              | 368      | 10.5     | 4.3         | 13.6         |
| 5,7-Diene-ergosteryl ester-2   | 380      | 10.7     | 12.8        | 8.4          |
| 7-Dehydrocholesterol ester-2   | 366      | 11.1     | 4.6         | 3.6          |
| 5,7,24-Triene-ergosteryl ester-2| 378      | 11.7     | 12.6        | 18.2         |
| 5,7-Diene-ergosteryl ester-3   | 380      | 12.5     | 13.5        | 9.1          |
| Ratio (steryl ester/free sterol)c |           | 0.24     | 0.15        |

*a*The fatty acid moieties were not identified (the MS scan range was from 50 to 450 amu)

*b*Not a steryl ester.

*c*The ratio was calculated by dividing a sum of total ion count peak areas of the steryl esters by that of free sterols.

https://doi.org/10.1371/journal.pntd.0006104.t003
Impact of posaconazole on the *N. fowleri* cell ultrastructure

DMSO-treated *N. fowleri* trophozoites displayed normal morphology with several food vacuoles, mitochondria, lipid droplets and a nucleus containing one large nucleolus (Fig 5A). Lipid droplets serve as the energy and carbon reservoirs in all domains of life. From analysis of another free-living amoeba, *Dictyostelium discoideum*, reported elsewhere,[32] we know that amoeba lipid droplets consist of a hydrophobic core of triglycerides, steryl esters and free sterols surrounded by one leaflet derived from the endoplasmic reticulum membrane to which a specific set of proteins is bound. Between 24 and 48 hours of undisrupted growth, *N. fowleri* lipid droplets increased in number and density (Fig 5C).

Treatment with posaconazole led to disorganization of *N. fowleri* membranes, swelling of mitochondria and appearance of multiple autophagic vacuoles engulfing organelle debris and myelin figures (loops of membranes), indicative of disruption of lipid metabolism. (Fig 5B and 5D). Similar ultrastructural alterations were previously reported in *N. fowleri* treated with AmpB, including mitochondrial abnormalities and an increase in autophagic vacuoles with myelin-like membranous whorls.[33] AmpB-treated amoebae also had abnormally shaped nucleus and an increase in rough and smooth endoplasmatic reticulum membranes.[33] alterations not observed with posaconazole. Finally, lipid droplets of the AmpB-treated amoebae were clustered and enclosed in membranous sheet, whereas treatment with posaconazole in these studies resulted in dispersion of lipid droplets throughout the cell with decrease in size and density.

Consistent with the rate of growth inhibition experiment (Fig 2), significant differences in cell ultrastructure are observed after 24 h of posaconazole exposure and no further differences were recorded between 24 h and 48 h of treatment (Fig 5B vs. Fig 5D). This observation distinguishes *N. fowleri* from kinetoplastid parasites where CYP51 inhibitors are notoriously slow-acting.[34] In *T. cruzi*, ultrastructural alterations and increase in autophagic vacuoles with myelin-like membranous whorls are first observed after 72–96 h drug exposure when lipid droplets are largely exhausted.[35, 36] In *N. fowleri*, dispersed lipid droplets albeit of reduced density are present after 48 h in significantly damaged *Naegleria* cells. Similar lipid accumulation after posaconazole exposure is observed in *Leishmania amazonensis*, both by TEM and fluorescent staining by Nile Red.[37]

Interaction of conazoles with the recombinant NfCYP51 target by UV-vis spectroscopy

The recombinant NfCYP51 was authenticated and the drug-target interactions were characterized by UV-vis spectroscopy. The ferric, Fe$^{3+}$, spectrum of NfCYP51 is typical of that of a low-spin P450 (CYP) with a Soret band at 417 nm, while that of the dithionite reduced ferrous, Fe$^{2+}$, species has a Soret band of 411 nm (Fig 6A). The 449 nm Soret band of the dithionite reduced and CO bound NfCYP51 is consistent with that of functional CYP enzymes (Fig 6A, inset). Upon binding to P450, a heterocyclic drug replaces the heme axial water ligand resulting in a red-shift of the iron Soret band, known as type II.[38] Conazoles bound to NfCYP51 produced type II low-spin difference spectra with a trough at 411 nm and a peak at 430 nm, indicative of azole coordination to the heme iron (Fig 6B). 31-Norlanosterol binding produced a type I high-spin difference spectrum with a peak at 388 nm and a trough at 418 nm (Fig 6B), resulting from the expulsion of water molecule ligand from the iron coordination sphere by the incoming substrate.[38] The dissociation constant, $K_D$, of 124±25 nM, was calculated for 31-norlanosterol by fitting plotted spectroscopic data to the standard Michaelis-Menten or Morrison binding equations (Fig 6C), both yielding the same $K_D$ value. The dissociation constants for conazoles could not be calculated by this method due to enzyme saturation reached after the addition of one molar equivalent of a conazole drug to 0.5 μM NfCYP51, as illustrated.
Fig 5. Ultrastructural analysis of posaconazole-treated *N. fowleri* by transmission electron microscopy (TEM). 0.1% DMSO-treated controls (A: A1-A3, 24 h; and C: C1-C3, 48 h, show several food vacuoles (V), mitochondria (M), nucleus (N) and lipid droplets (L). Exposure to 0.2 μM posaconazole (B: B1-B3, 24 h; D: D1-D3, 48 h) led to mitochondrial swelling, accumulation of atypical lipid droplets, alteration of nuclear membrane and appearance of autophagc vacuoles (A) engulfing organelle debris and myelin figures (MF). Bar = 500 nm.

https://doi.org/10.1371/journal.pntd.0006104.g005
for fluconazole and posaconazole (Fig 6D). When sub-stoichiometric concentrations of drugs were titrated into the enzyme solution, a linear increase in signal was observed for all conazoles up until the equivalence point, after which no further increase in signal was detected. Dilution of NfCYP51 below 0.5 μM resulted in a substantial drop in signal-to-noise ratio in the recorded spectra. From the UV-vis data, we conclude that even the smallest azole drug, fluconazole, binds NfCYP51 with affinity roughly an order of magnitude exceeding that of natural substrate, 31-norlanosterol. The binding superiority of the azole drugs over natural substrate is achieved due to the formation of the coordination bond between an aromatic nitrogen of the azole heterocycle and the heme iron (Fig 7A–7D).

Drug-target interactions by x-ray crystallography

**Overall structure.** To further characterize drug-target interactions, we have screened conazoles for co-crystallization propensity with the NfCYP51 target and determined the co-crystal structures of NfCYP51 bound to five conazole drugs: posaconazole (1.71 Å), ketoconazole (1.87 Å), itraconazole (2.6 Å), voriconazole (2.4 Å) and fluconazole (2.7 Å) (Table 4). All conazoles form a coordination bond to the heme iron via an aromatic nitrogen of the heterocycle moiety (Fig 7A–7D). The overall protein scaffold of NfCYP51 is similar to that of previously characterized CYP51 from other eukaryotes with the qualification that certain secondary structure elements sample multiple conformations spontaneously or in response to the incoming ligand.[39–41] A number of the co-crystal structures for human,[42] yeast,[43, 44] fungi
and kinetoplastid [21] orthologues reported to date point at the BC- and FG-loops and the F- and G-helices (nomenclature introduced by Poulos et al. [47]), as the most conformationally variable structural elements whose concerted motion modulates the size and topology of the CYP51 binding site.

**Drug-target interactions.** Both posaconazole and itraconazole (MW ~700 g/mol) occupy the whole length of the CYP51 hydrophobic tunnel spanning from the heme macrocycle to the protein surface (Fig 7A). At Pro213, the tunnel turns forcing the inhibitor’s long moiety to bend at piperazine ring. The bent conformation of posaconazole has been previously reported in *T. brucei* CYP51. [48] As evidenced by the *F*<sub>o</sub>–*F*<sub>c</sub> electron density map (S2 Fig), the 2,4-dichlorophenyl moiety of itraconazole may flip in the vicinity of heme adopting two alternative conformations. Flipping of the 2,4-difluorophenyl moiety was previously observed in fluconazole. [48] The smaller ketoconazole (MW 531 g/mol) also extends into the hydrophobic tunnel but only to reach Pro213 (Fig 7B). Finally, voriconazole and fluconazole (MW 300–350 g/mol), bind in vicinity of heme macrocycle and benefit very little from the tunnel space (Fig 7C and 7D). Well-defined by electron density, the 5-fluoropyrimidinyl moiety of voriconazole makes a series of contacts within 6 Å of Tyr107, Phe109, Met110, Leu358, Met362 and Leu467, plus a H-bond to carbonyl oxygen of M360 (3.6 Å) (Fig 7C). The smaller and more hydrophilic 1,2,4-triazolyl group of fluconazole is less favored by this environment, as evidenced by

![Figure 7. NCYP51-inhibitor binding.](https://doi.org/10.1371/journal.pntd.0006104.g007)
the longer distances and the electron density map progressively less defined for the triazole moiety not involved in the coordination of heme iron (Fig 7D).

Variability of the CYP51 binding site. The first-tier contacts within 5 Å of inhibitors deduced from the NfCYP51 structure have been projected to the CYP51 sequences of other human pathogens (Fig 7E). Predominantly aliphatic and aromatic amino acid residues delineate the binding tunnel in CYP51, with only three of them being invariant across the species. Variable positions with different degree of conservation confer substrate- and inhibitor-binding specificity to the CYP51 orthologues. Consistent with the biochemical data, phylogenetic analysis points at 'plant-like' substrate specificity of NfCYP51 defined by Phe109 (numbering is according to NfCYP51) that is only present in the orthologues converting the 4α-monomethylsterol substrates, obtusifoliol or 31-norlanosterol. In this regard, *Naegleria*, *Acanthamoeba* and *T. brucei* group together with plant CYP51 represented in Fig 7E by *Arabidopsis thaliana*. The *T. cruzi*, fungal and mammalian CYP51 having leucine/isoleucine at 109 metabolize 4α,β-dimethylated sterol substrates, lanosterol, eburicol and 24,25-dihydrolanosterol.

Another critical sequence variability is mapped to proline 213. A non-conserved proline-to-histidine substitution in human CYP51 accounts for the host-pathogen selectivity of conazoles. The histidine moiety protruding into the hydrophobic tunnel of human CYP51 interferes with a long substituent of the high MW conazoles. The substantial genetic divergence between the etiological agents of human diseases require tailored implementation of pathogen-specific drug discovery programs utilizing methodologies specific to the targeted

| Protein | NfCYP51 | NfCYP51 | NfCYP51 | NfCYP51 | NfCYP51 |
|---------|---------|---------|---------|---------|---------|
| Ligand  | Posaconazole X2N | Itraconazole 1YN | Ketoconazole KKK | Voriconazole VOR | Fluconazole TPF |
| PDB ID  | 5TL8 | 6AYC | 6AYB | 6AY6 | 6AY4 |

Data collection

| Space group | C2 | C2 | C2 | C2 | C2 |
|-------------|----|----|----|----|----|
| b, c (Å)    | 120.0, 55.0, 71.6 | 120.3, 55.2, 72.4 | 119.2, 55.3, 71.6 | 120.2, 55.3, 73.2 | 122.1, 55.3, 73.5 |
| α, β, γ (°) | 90.0, 100.1, 90.0 | 90.0, 100.6, 90.0 | 90.0, 100.1, 90.0 | 90.0, 100.8, 90.0 | 90.0, 101.5, 90.0 |

| Wavelength (Å) | 1.11587 | 1.11587 | 1.11587 | 1.11587 | 1.11587 |
|----------------|---------|---------|---------|---------|---------|
| Resolution (Å) | 1.71 | 1.87 | 2.49 | 2.70 |
| R f (%) | 4.1 (144.8)a | 13.3 (286.5) | 20.3 (188.7) | 10.9 (136.0) | 7.0 (160.4) |
| lₙl | 14.4 (0.62) | 7.1 (0.59) | 3.79 (0.53) | 7.0 (0.47) | 7.94 (0.57) |
| Completeness (%) | 98.3 (85.6) | 99.7 (99.8) | 95.1 (68.4) | 97.3 (66.5) | 97.7 (99.8) |
| Redundancy | 3.6 (2.4) | 6.5 (6.9) | 6.0 (4.7) | 5.5 (2.2) | 2.6 (2.6) |

Refinement

| No. reflections | 47085 | 13616 | 34450 | 17255 | 12443 |
|----------------|------|------|------|------|------|
| R work / R free (%) | 19.5/24.9 | 21.8/29.0 | 24.3/28.4 | 21.9/29.0 | 25.7/31.3 |
| No. atoms | 3799 | 3731 | 3712 | 3660 | 3670 |
| Wilson | 36.1 | 86.0 | 42.9 | 82.1 | 85.6 |
| Mean B value | 39.9 | 95.1 | 46.0 | 85.3 | 97.8 |
| R.m.s deviations | 0.018 | 0.011 | 0.019 | 0.009 | 0.008 |

aValues for highest-resolution shells are in parentheses
pathogens. For *N. fowleri*, the crystal structures determined in this work may be indispensable for designing molecules combining potency against NfCYP51 with brain permeability.

While UV-vis spectroscopy could not prioritize conazoles by binding affinity due to insufficient sensitivity of the experimental setup, the co-crystal structures unambiguously link the decline in conazole’s potency to the decrease in a number of drug-target interactions as a function of MW. The highest MW conazoles, posaconazole and itraconazole, making a large number of interactions in the CYP51 hydrophobic tunnel, are broad spectrum. Fluconazole, with the least number of drug-target contacts, is a narrow spectrum antifungal agent with the activity limited to *Candida albicans* (but not *C. krusei* or *C. glabrata*) and *Cryptococcus neoformans*. Despite of being part of the drug formulation used for the treatment of PAM patients, fluconazole has the lowest activity against *N. fowleri* (*EC₅₀* of 13.9 μM) among azole drugs. Differences in the first-tier residues directly contacting fluconazole in *N. fowleri* and *C. albicans* (Fig 7E) may account for the differential activity. Also, high glycine content (positions equivalent to Ala54, Ala289 and Ala293 in *N. fowleri* are taken by glycine in *C. albicans*) may render *C. albicans* CYP51 more adaptive to fluconazole.

**Perspectives of conazoles in the treatment of PAM**

Although the potency of some conazoles against cultured *N. fowleri* was equal or exceeded that of AmpB (Table 1), to attain parasitological cure in PAM patients, a drug must cross the blood-brain barrier (BBB). Conazoles exhibit variable physicochemical characteristics and differ with regard to cerebrospinal fluid (CSF) and brain parenchymal penetration.[30, 49] Across the board, brain permeability of conazoles is inversely related to the MW and to their in vitro anti-*Naegleria* potency. Thus, fluconazole, having the lowest activity against *N. fowleri*, is known to rapidly distribute through body tissues, including different CNS compartments, where it achieves concentrations greater than MIC₉₀ of common fungal pathogens.[49, 50] The intermediate MW conazoles penetrate CSF and brain tissue to a different extent. Voriconazole brain permeability has been reported in human studies of meningitis patients.[51–53] Miconazole is effective in treatment of human fungal meningitis not susceptible to AmpB; high brain concentrations of miconazole are achieved by intrathecal (IT) administration.[54, 55] Rabbit studies have shown that ravuconazole[56] and ketoconazole[57] penetrate brain tissue; intermediate concentrations of ketoconazole were found in CSF and are modestly increased in the presence of meningeal inflammation.[57]

Finally, the highest MW conazoles—posaconazole and itraconazole—have CNS pharmacokinetics similar to that of a cornerstone of PAM therapy, AmpB. In rabbit studies, none of the AmpB formulations produced measurable concentrations in the CSF regardless of CNS inflammation.[58, 59] At the same time, detectable brain parenchymal AmpB concentrations were observed even in the absence of CNS infection (3–27% of serum concentrations) and increased two- to four-fold in presence of infection.[59] Animal model and human studies reported nearly undetectable CSF concentrations of itraconazole,[60–62] while posaconazole showed striking differences in the CSF-to-plasma ratios ranging from the below limit of detection to 2.4.[63] Disturbance of the BBB tight junctions by inflammation may facilitate passage of posaconazole into the CSF.[64] Despite the lack of appreciable CSF concentrations, itraconazole accumulates in brain parenchyma as an active hydroxylated metabolite at concentrations higher than the MIC of the infecting fungi.[65] The ability of itraconazole and posaconazole to accumulate in brain tissue may account for successful use of these drugs for CNS invasive fungal infections in humans.[57, 66–68] If brain parenchymal kinetics are valid predictor of antifungal efficacy in the treatment of CNS mycoses compared to CSF concentrations,[49] the same may be true for anti-*Naegleria* efficacy. Systematic assessment of conazoles
in an animal model of PAM would single out the most efficacious drug of this class for the treatment of PAM.

**Summary.** Based on the collective evidence of the target-based and whole-parasite studies, we conclude that amoebicidal effect of conazoles in *N. fowleri* is due to depletion of the ergosterol pool concomitant with accumulation in large amounts of sterol intermediates with molecular structure/physicochemical properties incompatible with normal cell physiology. Disruption of CYP51 function induces massive autophagocytosis leading to rapid *N. fowleri* cell death. These data validate CYP51 as an essential enzyme and potentially a druggable target in *N. fowleri*. However, the amoebicidal activity of conazoles is inversely related to their brain permeability: it increases with the increase of drug molecular weight (MW), reaching low nanomolar potency for posaconazole and itraconazole. The low anti-*N. fowleri* activity of the brain-penetrant fluconazole questions its role in a drug combination currently recommended by CDC for the PAM treatment.

**Materials and methods**

**Materials**

The *Naegleria fowleri* strain KUL originally isolated from human cerebrospinal fluid in Belgium in 1973 [69] was obtained from ATCC. KUL is type 3 strain based on the length of the internal transcribed spacers 1 (ITS1), with the T at position 31 in the 5.8S rDNA sequence.[3]

Azole inhibitors were purchased from commercial sources: fluconazole from Cayman Chemical, clotrimazole, miconazole (racemic mix) and voriconazole from Sigma-Aldrich, ketoconazole and itraconazole from Alfa Aesar, voriconazole and isavuconazole from Stru-Chem (China). Posaconazole was purified from a Noxafil (Merck) suspension purchased from a pharmacy, as previously described.[35] Miltefosine and amphotericin B were purchased from Sigma-Aldrich. Sulfobutylether-β-cyclodextrin (SBE-β-CD), also known as Captisol, was from MedChem Express and methyl-β-cyclodextrin (M-β-CD) was from Sigma-Aldrich. 31-Norlanosterol was purified from *Candida albicans* treated with both CYP51 and SMT inhibitors; the structure of the sterol was authenticated by both the GC-MS and NMR methods.

Drug stock solutions were freshly made either in DMSO or 40% SBE-β-CD. The DMSO stock was used for preparation of the electron microscopy and GC-MS samples, while the SBE-β-CD stock was used for serial dilutions in the growth inhibition experiments. The SBE-β-CD stock solutions were prepared based on the compositions for the posaconazole intravenous administration provided in the US Patent 2013/0096053 A1.[31] Briefly, a 40% solution of SBE-β-CD was prepared by dissolving 4 grams of SBE-β-CD in 10 ml of ddH2O. Five μmoles of each drug was added to 1 ml of 40% SBE-β-CD and sonicated until dissolved. In order to solubilize posaconazole and itraconazole, the drug-SBE-β-CD suspension was acidified to pH 2 by the addition of 15% HCl and then sonicated.

**Validation of azole inhibitors for amoebicidal activity**

To determine EC50 values, conazoles were tested for dose-response against *N. fowleri* trophozoites axenically cultured in Nelson’s medium supplemented with 10% fetal bovine serum at 37°C.[70] all the experiments were performed in triplicate using trophozoites harvested during the logarithmic phase of growth.[71] Drug concentration ranges of 0.4–50 μM and 0.008–25 μM in 0.2% SBE-β-CD were generated by transferring 0.5 μl of serially diluted compounds to a corresponding well of the 96-well plate followed by addition of 99.5 μl of *N. fowleri* trophozoites (10,000 amoebae). Assay plates were incubated for 48 h and cell viability was determined by the CellTiter-Glo Luminescent Cell Viability Assay.[26, 71] The experiments using
trophozoites were conducted in a biosafety cabinet following the BSL2 procedures as specified in the UCSD Biosafety Practices Guidelines.

Ultrastructural analysis by transmission electron microscopy (TEM)

*N. fowleri* trophozoites (2x10^6) were treated with posaconazole at 0.2 μM, for 24 h and 48 h, washed with PBS and then fixed overnight at 4˚C in modified Karnovsky’s fixative (2.5% glutaraldehyde and 2% paraformaldehyde in 0.1 M sodium phosphate, pH 7.2)[72] 0.1% DMSO-treated controls were simultaneously processed. The samples were then post-fixed for 1 hour with 1% osmium tetroxide in 0.15 M sodium cacodylate, pH 7.4, dehydrated with an ascending series of ethyl alcohol and propylene oxide, and finally embedded in an Epon resin (Scipoxy 812, Energy Beam Sciences). Thin sections (50–60 nm) were cut using Leica UCT ultramicrotome, mounted on the Formvar and carbon-coated copper grids, and counterstained for 5 min with 2% uranyl acetate followed by Sato’s lead stain for 1 min. *Naegleria* thin sections were examined using a Tecnai G2 Spirit BioTWIN transmission electron microscope (TEM) equipped with an Eagle 4k HS digital camera (FEI, Hillsboro, OR).

GC-MS analysis of the *Naegleria* sterols

*N. fowleri* sterols were analyzed by the use of GC-MS, wherein the lipids extracted from *N. fowleri* trophozoites grown in the presence of a vehicle or inhibitor at concentrations that produce ultrastructural changes without destroying a parasite cell, were separated by gas chromatography and subsequently analyzed by electron-ionization mass-spectrometry (EI). 2x10^7 or 5x10^7 trophozoites per sample were treated with 0.1% DMSO alone or 0.2 μM posaconazole dissolved in DMSO. AmpB, a drug with a different mechanism of action, was used as a negative control at 0.2 μM. To avoid parasite death, drug exposure was terminated after 24 h when the amoebae were pelleted by centrifugation, washed three times with PBS (3x10 ml) and, finally, 2 ml of chloroform/methanol 2:1 solution was added to the cell pellet. The organic solvents were evaporated under N₂ flow, and the pellet was incubated for 24 h with 3 ml chloroform. Polar molecules were removed by several extractions with water (3x10 ml). The organic solvent was then subsequently changed to chloroform/methanol 9:1 and then acetonitrile (3 ml each) through evaporation under N₂ flow; each step followed by triple washes in water as described above.

Extracted sterols were either directly analyzed as free sterols, or first derivitized with TMS group. For free sterol analysis, extracted dry sterols were re-dissolved in chloroform (100 μl) and 2 μl of each sample were injected into the analytical column of the Agilent 6890 gas chromatograph (the inject port temperature was controlled at 250˚C), coupled to a 5973 mass selective detector (MSD). The sterols were separated using a ZB5 capillary column (30m X 250um X0.25um) with helium carrier gas flow rate set at 1.2 ml/min and temperature profile beginning at 170˚C for 1 min, then increased by 20˚C/min to 280˚C, and then hold at 280˚C for 20 min. The mass spectrometer scanned m/z 50–500 during the course of analysis. The sterols were identified by comparing the GC retention time to that of the internal cholesterol [73] and the fragmentation patterns to that of the authentic standards and the NIST (2008) mass spectral library. A forward and reverse match score of 800 and above was considered a correct match. The sterols were quantified based on the total ion current peak areas of each sterol.

For chemical derivatization, extracted dry sterols were dissolved in 30 μl of hexane and 70 μl of N,N-bis(trimethylsilyl)-2,2,2-trifluoroacetamide (BSTFA), and incubated for 2 h at 37˚C. Three microliters of the TMS-derivatized lipid mixture was injected directly into an Agilent 7820A gas chromatography system coupled to a mass selective detector. The inject port
temperature was controlled at 250˚C, the helium carrier gas flow rate was set at 13 ml/min. The lipids were separated on the analytical column using a temperature profile that begins at 200˚C for 3 min, increases by 15˚C/min to 270˚C, and then holds at 270˚C for 30 min, finishing with post run 280˚C 4 min. The mass spectrometer scanned m/z 50–750 during the course of analysis.

**NfCYP51 expression and purification**

NfCYP51, codon-optimized for bacterial expression, had a coding sequence with 34 N-terminal membrane anchoring residues replaced with the MAKKTSSKGKL to increase recombinant protein recovery during purification. ([S1 Data](#)). This construct was generated synthetically (GenScript) and cloned into the pCW-LIC expression vector obtained from the non-profit plasmid repository (Addgene, Cambridge, MA). NfCYP51 was expressed in DH5α E. coli strain co-transformed with the pGro7 plasmid (Takara) carrying GroEL/ES chaperones with induction by 0.5 mM isopropyl-β-D-thiogalactopyranoside (IPTG) for 40–48 hours at 25˚C. All purification steps were carried out at 4˚C. Cells were pelleted, re-suspended in the lysis buffer (50 mM K-PO₄, pH 8.0; 100 mM NaCl, 10% glycerol, 1 mM EDTA, 1 mM DTT and 0.5 mM PMSF) and then disrupted using the fluid processor Microfluidics M-110P (Microfluidics Inc.). Non-ionic detergent CHAPS was added to 0.5% and cell lysate was incubated for 30 min prior to centrifugation. The crude extract was separated from cell debris. Cleared lysate was loaded to a Ni-NTA column and after a series of washes NfCYP51 was eluted by increasing the imidazole concentration from 0 to 500 mM. Fractions containing NfCYP51 were pooled and passed through Q-Sepharose and then S-Sepharose (GE Healthcare Life Sciences). NfCYP51 flowed-through both ion exchange columns was then bound to hydroxyapatite (HAP) column (BioRad) and eluted from it in the gradient of K-PO₄ concentrations from 0.02 M to 0.8 M supplemented with 10% glycerol, 0.5 mM EDTA and 1 mM DTT. Fractions containing pure NfCYP51 were pooled, concentrated, aliquoted and frozen at -80˚C.

**UV-vis spectroscopy of NfCYP51**

Recombinant NfCYP51 was characterized spectrally for integrity of the heme prosthetic group and for substrate and inhibitor binding. All spectra were recorded using a Cary 1E (Varian) dual beam UV-visible spectrophotometer. Purified NfCYP51 was diluted to 3.5 μM in 50 mM K-PO₄ (pH 7.4) and 10% glycerol buffer and allowed to equilibrate to room temperature for 10 min prior to readings. Spectra were recorded from 250–700 nm for the ferric and dithionite reduced ferrous NfCYP51 with buffer in the reference cuvette. The CO difference spectrum was recorded by splitting dithionite reduced ferrous NfCYP51 into the sample and reference cuvettes. A baseline was recorded and CO was bubbled into the sample cuvette, after which the difference spectrum was recorded. The concentration of NfCYP51 was calculated using the extinction coefficient ε₄₅₀ = 91 mM⁻¹cm⁻¹.

Spectral binding titrations were performed at 25˚C using 0.5 μM NfCYP51 for conazole and 0.2 μM for 31-norlanosterol. All conazoles with the exception of itraconazole were dissolved in 40% SBE-β-CD and then diluted to 100 μM stocks in 0.8% SBE-β-CD. Itraconazole was dissolved in DMSO. 31-Norlanosterol was dissolved in 20% M-β-CD and diluted to 100 μM. For each titration 2 ml of NfCYP51 was split equally for the reference and sample cuvettes with ligand or inhibitor being added to the sample cuvette while vehicle alone was added to the reference cuvette, with the total added volume being less than 1% of the total volume. Spectra were recorded from 350 to 500 nm. A binding isotherm for 31-norlanosterol was generated by plotting the absorbance minimum subtracted from the absorbance maximum as
function of drug concentration. The spectral dissociation constant $K_D$ was estimated using the Curve Fitting Tool in MATLAB (MathWorks, Natick, MA) by fitting the binding isotherm using the hyperbolic Michaelis-Menten equation $\Delta A = \Delta A_{\text{max}}/[L]/(K_D + [L])$ or the quadratic Morrison equation $\Delta A = (\Delta A_{\text{max}} / 2[E])((K_D + [L] + [E])—(K_D + [E] + [L])^2-4[E][L])^{0.5}$ where $\Delta A$ is the difference between absorbance maximum and minimum, $\Delta A_{\text{max}}$ is the extrapolated maximum absorption difference, $[L]$ is the ligand concentration and $[E]$ is the enzyme concentration.

**Crystallization and x-ray structure determination**

Prior to crystallization, NfCYP51 was diluted to 0.5 mM by mixing with 0.6 mM inhibitor added to the desired volume of water from the 10 mM or 100 mM stock solutions in DMSO, depending on compound solubility. Screening of crystallization conditions for each inhibitor complex was performed using commercial high-throughput screening kits available in deep-well format (Hampton Research or Qiagen), a nanoliter drop-setting Mosquito robot (TTP LabTech) operating with 96-well plates, and a hanging drop crystallization protocol. For diffraction quality, crystals were further optimized in 96-well plates configured using the Dragonfly robot (TTP LabTech) and the Designer software (TTP LabTech). All crystals were harvested from the narrow grid of crystallization conditions: 30–33% PEG MME 550, 30 mM CaCl$_2$, 0–4% Jeffamine M-600, 0.1 M bis-Tris propane, pH 7.1–7.5.

Diffraction data were collected at 100–110 K at beamline 8.3.1, Advanced Light Source, Lawrence Berkeley National Laboratory, USA. Data indexing, integration, and scaling were conducted using XDS.[74] The high-resolution crystal structure of the NfCYP51-posaconazole complex was determined by molecular replacement using as a search model *T. cruzi* CYP51, PDB ID 2X2N. The initial model was built using the BUCCANEER[75, 76] and COOT[77] programs. Refinement was performed by using REFMAC5 software.[76, 78] The newly determined crystal structure (PDB ID 5TL8) was subsequently used for other NfCYP51-conazole complexes reported in this work. Data collection and refinement statistics are shown in Table 4.

**Accession codes.** The atomic coordinates and structure factors (5TL8, 6AY4, 6AY6, 6AYB and 6AYC) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics, Rutgers University, New Brunswick, NJ (http://www.rcsb.org/)

**Supporting information**

**S1 Data. Recombinant NfCYP51.** NfCYP51 codon-optimized DNA sequence synthetically generated (GenScript, Piscataway, NJ) for bacterial expression—with 34 N-terminal residues replaced with the MAKKTSSKGKL leading sequence (to improve protein expression and purification)—and cloned into the pcW-LIC expression vector obtained from the non-profit plasmid repository (Addgene, Cambridge, MA).

(DOCX)

**S1 Fig. Gas chromatography separation of the total sterol fractions extracted from *N. fowleri* trophozoites.** Chromatogram fragments from the TMS-derivatized DMSO-, posaconazole- and Amphotericin B-treated *N. fowleri* lipid extracts are shown. Peaks are labeled according to Table 2. The sterol identities were assigned based on relative chromatographic behavior, the characteristic molecular masses and electron ionization (EI) fragmentation patterns by comparing them to the authentic standards and the NIST (2008) mass spectral library. In contrast to posaconazole, Amphotericin B, a drug with a different mechanism of action used as a negative control, did not perturb the native sterol pattern.

(TIF)
S2 Fig. Flipping of the 2,4-dichlorophenyl moiety of itraconazole evidenced by electron density.

Itraconazole in a single confirmation (yellow sticks) is shown in the fragment of the 2.6 Å 2Fo-Fc electron density map countered at 1.0 σ (cyan mesh) overlapped with a fragment of the Fo-Fc electron density countered at -3.0 σ (red mesh). "Negative" peak at 2-chloro-substituent suggests a possibility of partial occupancy of this site due to flipping of the 2, 4-dichlorophenyl moiety of itraconazole. Heme is shown in van der Waals spheres. Heteroatoms are colored according chemical elements: oxygen–red, nitrogen–blue, chlorine–green, iron–ochre.

(PNG)

Acknowledgments
We thank the staff members of beamline 8.3.1, James Holton, George Meigs and Jane Tanamachi, at the Advanced Light Source at Lawrence Berkeley National Laboratory, for assistance with data collection. We also thank Dr. Dionicio Siegel, UCSD, for posaconazole purification.

Author Contributions

Conceptualization: Larissa M. Podust.
Data curation: W. David Nes, Larissa M. Podust.
Formal analysis: Larissa M. Podust.
Funding acquisition: James H. McKerrow, Larissa M. Podust.
Investigation: Anjan Debnath, Claudia M. Calvet, Gareth Jennings, Wenxu Zhou, Alexander Aksenov, Madeline R. Luth, Larissa M. Podust.
Methodology: Anjan Debnath, Claudia M. Calvet, Gareth Jennings, Wenxu Zhou, Alexander Aksenov, Ruben Abagyan, W. David Nes, James H. McKerrow, Larissa M. Podust.
Project administration: Larissa M. Podust.
Resources: Anjan Debnath, W. David Nes, James H. McKerrow, Larissa M. Podust.
Software: Ruben Abagyan.
Supervision: W. David Nes, Larissa M. Podust.
Visualization: Claudia M. Calvet.
Writing – original draft: Larissa M. Podust.
Writing – review & editing: Anjan Debnath, Claudia M. Calvet, Gareth Jennings, Wenxu Zhou, Alexander Aksenov, James H. McKerrow, Larissa M. Podust.

References

1. Siddiqui R, Ali IK, Cope JR, Khan NA. Biology and pathogenesis of Naegleria fowleri. Acta Trop. 2016; 164:375–94. https://doi.org/10.1016/j.actatropica.2016.09.009 PMID: 27616699
2. Visvesvara GS. Infections with free-living amebae. Handb Clin Neurol. 2013; 114:153–68. https://doi.org/10.1016/B978-0-444-53490-3.00010-8 PMID: 23829906
3. De Jonckheere JF. Origin and evolution of the worldwide distributed pathogenic amoeboflagellate Naegleria fowleri. Infect Genet Evol. 2011; 11(7):1520–8. https://doi.org/10.1016/j.meegid.2011.07.023 PMID: 21843657
4. Center for Disease Control and Prevention. Naegleria fowleri—Primary Amebic Meningoencephalitis (PAM)—Amebic Encephalitis. Accessed May 8, 2017 [Available from: https://www.cdc.gov/parasites/naegleria/.
5. Zysset-Burn DC, Muller N, Beuret C, Heller M, Schurch N, Gottstein B, et al. Genome-wide identification of pathogenicity factors of the free-living amoeba *Naegleria fowleri*. BMC genomics. 2014; 15:496. https://doi.org/10.1186/1471-2164-15-496 PMID: 24950717

6. Linam WM, Ahmed M, Cope JR, Chu C, Visvesvara GS, da Silva AJ, et al. Successful treatment of an adolescent with *Naegleria fowleri* primary amebic meningoencephalitis. Pediatrics. 2015; 135(3):e744–8. https://doi.org/10.1542/peds.2014-2292 PMID: 25667249

7. Lemke A, Kiderlen AF, Kayser O. Amphotericin B. Appl Microbiol Biotechnol. 2005; 68(2):151–62. https://doi.org/10.1007/s00253-005-1955-9 PMID: 15821914

8. Fischer R, Trzaskos J, Magolda R, Ko S, Brosz C, Larsen B. Lanosterol 14 alpha-methyl demethylase. Isolation and characterization of the third metabolically generated oxidative demethylase intermediate. J Biol Chem. 1991; 266(10):6124–32. PMID: 2007571

9. Fischer RT, Stam SH, Johnson PR, Ko SS, Magolda RL, Gaylor JL, et al. Mechanistic studies of lanosterol 14 alpha-methyl demethylase: substrate requirements for the component reactions catalyzed by a single cytochrome P-450 isozyme. J Lipid Res. 1989; 30(10):1621–32. PMID: 2614264

10. Trzaskos J, Fischer R, Favata M. Mechanistic studies of lanosterol C-32 demethylase. Conditions which promote oxysterol intermediate accumulation during the demethylase process. J Biol Chem. 1986; 261(36):16937–42. PMID: 3782148

11. Aoyama Y, Yoshida Y, Sonoda Y, Sato Y. Deformylation of 32-oxo-24,25-dihydrolanosterol by the purified cytochrome P-45014DM (lanosterol 14 alpha-demethylase) from yeast evidence confirming the intermediate step of lanosterol 14 alpha-demethylation. J Biol Chem. 1989; 264(31):18502–5. PMID: 2509459

12. Heeres J, Meerpold L, Lewi P. Conazoles. Molecules. 2010; 15(6):4129–88. https://doi.org/10.3390/molecules15064129 PMID: 20657432

13. Duma RJ, Finley R. In vitro susceptibility of pathogenic *Naegleria* and *Acanthamoeba* species to a variety of therapeutic agents. Antimicrob Agents Chemother. 1976; 10(2):370–6. PMID: 984777

14. Schuster FL, Guglielmo BJ, Visvesvara GS. *In vitro* activity of miltefosine and voriconazole on clinical isolates of free-living amebas: *Balamuthia mandrillaris*, *Acanthamoeba* spp., and *Naegleria fowleri*. J Eukaryot Microbiol. 2006; 53(2):121–6. https://doi.org/10.1111/j.1550-7408.2005.00082.x PMID: 16579814

15. Tiewcharoen S, Junnu V, Chinabut P. *In vitro* effect of antifungal drugs on pathogenic *Naegleria* spp. Southeast Asian J Trop Med Public Health. 2002; 33(1):38–41. PMID: 12118457

16. Tiewcharoen S, Junnu V, Suvoutho S. Effect of antifungal drugs on pathogenic *Naegleria* spp isolated from natural water sources. J Med Assoc Thai. 2003; 86(9):876–82. PMID: 14649973

17. Thong YH, Rowan-Kelly B, Shepherd C, Ferrante A. Growth inhibition of *Naegleria fowleri* by tetracycline, rifamycin, and miconazole. Lancet. 1977; 2(8043):876.

18. Seidel JS, Harmatz P, Visvesvara GS, Cohen A, Edwards J, Turner J. Successful treatment of primary amebic meningoencephalitis. N Engl J Med. 1982; 306(6):346–8. https://doi.org/10.1056/NEJM198202113060607 PMID: 7054710

19. Vargas-Zepeda J, Gomez-Alcala AV, Vasquez-Morales JA, Licea-Araya A, De Jonckheere JF, Lares-Villa F. Successful treatment of *Naegleria fowleri* meningoencephalitis by using intravenous amphotericin B, fluconazole and rifampicin. Arch Med Res. 2005; 36(1):83–6. PMID: 15900627

20. Desmond E, Gribaldo S. Phylogenomics of sterol synthesis: insights into the origin, evolution, and diversity of a key eukaryotic function. Genome biology and evolution. 2009; 1:364–81. https://doi.org/10.1093/gbe/evp036 PMID: 20333205

21. Choi JY, Podust LM, Roush WR. Drug strategies targeting CYP51 in neglected tropical diseases. Chem Rev. 2014; 114(22):11242–71. https://doi.org/10.1021/cr5003134 PMID: 25337991

22. Raederstorff D, Rohmer M. Sterol biosynthesis de nova via cycloartenol by the soil amoeba *Acanthamoeba polyphaga*. Biochem J. 1985; 231(3):609–15. PMID: 4074328

23. Raederstorff D, Rohmer M. Sterol biosynthesis via cycloartenol and other biochemical features related to photosynthetic phyla in the amoebae *Naegleria lovaniensis* and *Naegleria gruberi*. Eur J Biochem. 1987; 164:427–34. PMID: 3569274

24. Raederstorff D, Rohmer M. The action of the systemic fungicides tridemorph and fenpropimorph on sterol biosynthesis by the soil amoeba *Acanthamoeba polyphaga*. Eur J Biochem. 1987; 164(2):421–6. PMID: 3569273

25. Nes CR, Singha UK, Liu J, Ganapathy K, Villalta F, Waterman MR, et al. Novel sterol metabolic network of *Trypanosoma brucei* procyclic and bloodstream forms. Biochem J. 2012; 443(1):267–77. https://doi.org/10.1042/Bj20111849 PMID: 22176028

26. Debnath A, Tunac JB, Galindo-Gomez S, Silva-Olivares A, Shibayama M, McKerrow JH. Corifungin, a new drug lead against *Naegleria*, identified from a high-throughput screen. Antimicrob Agents Chemother. 2012; 56(11):5450–7. https://doi.org/10.1128/AAC.00843-12 PMID: 22869574
27. Dohta Y, Yamashita T, Horiike S, Nakamura T, Fukami T. A system for LogD screening of 96-well plates using a water-plug aspiration/injection method combined with high-performance liquid chromatography-mass spectrometry. Anal Chem. 2007; 79(21):8312–5. https://doi.org/10.1021/ac070978z PMID: 17910417

28. Herbig ME, Evers DH. Correlation of hydrotrropic solubilization by urea with logD of drug molecules and utilization of this effect for topical formulations. Eur J Pharm Biopharm. 2013; 85(1):158–60. https://doi.org/10.1016/j.ejpb.2013.06.022 PMID: 23958327

29. Lombardo F, Shalaeva MY, Tupper KA, Gao F. ElogD(oct): a tool for lipophilicity determination in drug discovery. 2. Basic and neutral compounds. J Med Chem. 2001; 44(15):2490–7. PMID: 11448232

30. Felton T, Troke PF, Hope WW. Tissue penetration of antifungal agents. Clin Microbiol Rev. 2014; 27 (1):68–88. https://doi.org/10.1128/CMR.00046-13 PMID: 24396137

31. Heimbecher SK, Plains M, Monteith D, inventors; Merck Sharp & Dohme Corp., assignee. Posaconazole intravenous solution formulations stabilized by substituted beta-cyclodextrin. US 2013/0096053 A12013.

32. Du X, Barisch C, Paschke P, Herrfurth C, Bertinetti O, Pawolleck N, et al. Dictyostelium lipid droplets host novel proteins. Eukaryot Cell. 2013; 12(11):1517–29. https://doi.org/10.1128/EC.00182-13 PMID: 24036346

33. Schuster FL, Rechthand E. In vitro effects of amphoterin B on growth and ultrastructure of the amoeboflagellates Naegleria gruberi and Naegleria fowleri. Antimicrob Agents Chemother. 1975; 8(5):591–605. PMID: 12119114

34. Moraes CB, Giardini MA, Kim H, Franco CH, Araujo-Junior AM, Schenkman S, et al. Nitroheterocyclic compounds are more efficacious than CYP51 inhibitors against Trypanosoma cruzi: implications for Chagas disease drug discovery and development. Sci Rep. 2014; 4:4703. https://doi.org/10.1038/srep04703 PMID: 24736467

35. Doyle PS, Chen C-K, Johnston JB, Hopkins SD, Leung SSF, Jacobson MP, et al. A nonazole CYP51 inhibitor cures Chagas’ Disease in a mouse model of acute infection. Antimicrob Agents Chemother. 2010; 54(6):2480–8. https://doi.org/10.1128/AAC.00281-10 PMID: 20385875

36. Veiga-Santos P, Barrias ES, Santos JF, de Barros Moreira TL, de Carvalho TM, Urbina JA, et al. Effects of amiodarone and posaconazole on the growth and ultrastructure of Trypanosoma cruzi. Int J Antimicrob Agents. 2012; 40(1):61–71. https://doi.org/10.1016/j.ijantimicag.2012.03.009 PMID: 22591838

37. de Macedo-Silva ST, Visbal G, Urbina JA, de Souza W, Rodrigues JC. Potent in vitro antiproliferative synergism of combinations of ergosterol biosynthesis inhibitors against Leishmania amazonensis. Antimicrob Agents Chemother. 2015; 59(10):6402–18. https://doi.org/10.1128/AAC.01150-15 PMID: 26239973

38. Luthra A, Denisov IG, Sligar SG. Spectroscopic features of cytochrome P450 reaction intermediates. Arch Biochem Biophys. 2011; 507(1):26–35. https://doi.org/10.1016/j.abb.2010.12.008 PMID: 21167809

39. Calvet CM, Vieira DF, Choi JY, Kellar D, Cameron MD, Siqueira-Neto JL, et al. 4-Aminopyridyl-based CYP51 inhibitors as anti-Trypanosoma cruzi drug leads with improved pharmacokinetic profile and in vivo potency. J Med Chem. 2014; 57(16):9899–7005. https://doi.org/10.1021/jm500448u PMID: 25101801

40. Vieira DF, Choi JY, Calvet CM, Siqueira-Neto JL, Johnston JB, Kellar D, et al. Binding mode and potency of N-indolyloxopyridinyl-4-aminopropyl-based inhibitors targeting Trypanosoma cruzi CYP51. J Med Chem. 2014; 57(23):10162–75. https://doi.org/10.1021/jm501566b PMID: 25393646

41. Vieira DF, Choi JY, Roush WR, Podust LM. Expanding the binding envelope of CYP51 inhibitors targeting Trypanosoma cruzi with 4-aminopyridyl-based sulfonamide derivatives. ChemBioChem. 2014; 15 (8):1111–20. https://doi.org/10.1002/cbic.201402027 PMID: 24771705

42. Strushevich N, Usanov SA, Park HW. Structural basis of human CYP51 inhibition by antifungal azoles. J Mol Biol. 2010; 397(4):1067–78. https://doi.org/10.1016/j.jmb.2010.01.075 PMID: 2049798

43. Monk BC, Tomasiak TM, Keniya MV, Huschmann FU, Tyndall JDA, O’Connell III JD, et al. Architecture of a single membrane spanning cytochrome P450 suggests constraints that orient the catalytic domain relative to a bilayer. Proc Natl Acad Sci U S A. 2014; 111(10):3865–70. https://doi.org/10.1073/pnas.1324245111 PMID: 24613951

44. Sagatova AA, Keniya MV, Wilson RK, Monk BC, Tyndall JD. Structural insights into binding of the antifungal drug fluconazole to Saccharomyces cerevisiae lanosterol 14α-demethylase. Antimicrob Agents Chemother. 2015; 59(8):4982–9. https://doi.org/10.1128/AAC.00926-15 PMID: 26058392

45. Hargrove TY, Wawrzak Z, Lamb DC, Guengerich FP, Lepeshevab GI. Structure-functional characterization of cytochrome P450 sterol 1αα-demethylase (CYPS1B) from Aspergillus fumigatus and molecular basis for the development of antifungal drugs. J Biol Chem. 2015; 290(39):23916–34. https://doi.org/10.1074/jbc.M115.677310 PMID: 26269599
46. Hargrove TY, Friggeri A, Wawrzak Z, Qi A, Hoekstra WJ, Schotzinger RJ, et al. Structural analyses of Candida albicans sterol 14alpha-demethylase complexed with azole drugs address the molecular basis of azole-mediated inhibition of fungal sterol biosynthesis. J Biol Chem. 2017; 292(16):6728–43. https://doi.org/10.1074/jbc.M117.778308 PMID: 28258218

47. Li H, Poulos TL. Crystallization of cytochromes P450 and substrate-enzyme interactions. Curr Top Med Chem. 2004; 4(16):1789–802. PMID: 15579108

48. Chen C-K, Leung SSF, Guilbert C, Jacobson MP, McKerrow JH, Podust LM. Structural characterization of CYP51 from Trypanosoma cruzi and Trypanosoma brucei bound to the antifungal drugs posaconazole and fluconazole. PLoS Negl Trop Dis. 2010; 4:e651. https://doi.org/10.1371/journal.pntd.0000651 PMID: 20386598

49. Kethireddy S, Andes D. CNS pharmacokinetics of antifungal agents. Expert Opin Drug Metab Toxicol.

50. Lutsar I, Roffey S, Troke P. Voriconazole concentrations in the cerebrospinal fluid and brain tissue of guinea pigs and immunocompromised patients. Clin Infect Dis. 2003; 37(5):728–32. https://doi.org/10.1086/377131 PMID: 12953033

51. Sung JP, Campbell GD, Grendahl JG. Miconazole therapy for fungal meningitis. Arch Neurol. 1978; 35(7):443–7. PMID: 580889

52. Lee JW, Amantea MA, Francis PA, Navarro EE, Bacher J, Pizzo PA, et al. Pharmacokinetics and safety of a unilamellar liposomal formulation of amphotericin B (AmBisome) in rabbits. Antimicrob Agents Chemother. 2004; 48(1):713–8. PMID: 12780341

53. Groll AH, Mickiene D, Pettitais V, Pettitais R, Kelaber A, Sarafand A, et al. Comparative pharmacokinetics and tissue distribution of the antifungal triazole ravuconazole following intravenous administration of its di-lysine phosphoester prodrug (BMS-379224) in rabbits. J Antimicrob Chemother. 2005; 56(5):899–907. https://doi.org/10.1093/jac/dki287 PMID: 16172108

54. Perfect JR, Durack DT. Penetration of imidazoles and triazoles into cerebrospinal fluid of rabbits. J Antimicrob Chemother. 1985; 16(1):81–6. PMID: 2995303

55. Lee JW, Amantea MA, Francis PA, Navarro EE, Bacher J, Pizzo PA, et al. Pharmacokinetics and safety of a unilamellar liposomal formulation of amphotericin B (AmBisome) in rabbits. Antimicrob Agents Chemother. 1994; 38(4):713–8. PMID: 8031034

56. Groll AH, GirI N, Pettitais V, Pettitais R, Candelario M, Bacher JS, et al. Comparative efficacy and distribution of lipid formulations of amphotericin B in experimental Candida albicans infection of the central nervous system. J Infect Dis. 2000; 182(1):274–82. https://doi.org/10.1086/315643 PMID: 10882607

57. Sorensen KN, Sobel RA, Clemons KV, Pappagianis D, Stevens DA, Williams PL. Comparison of fluconazole and itraconazole in a rabbit model of coccidioidal meningitis. Antimicrob Agents Chemother. 2000; 44(6):1512–7. PMID: 10882607

58. Perfect JR, Savani DV, Durack DT. Comparison of itraconazole and fluconazole in treatment of cryptococcal meningitis and candida pyelonephritis in rabbits. Antimicrob Agents Chemother. 1986; 29(4):579–83. PMID: 3010846

59. Kamberi P, Sobel RA, Clemons KV, Waldvogel A, Striebel JM, Williams PL, et al. Comparison of itraconazole and fluconazole treatments in a murine model of coccidioidal meningitis. Antimicrob Agents Chemother. 2007; 51(3):998–1003. https://doi.org/10.1128/AAC.00332-06 PMID: 17178793

60. Calcagno A, Baietto L, De Rosa FG, Tettoni MC, Libanore V, Bertucci R, et al. Posaconazole cerbrospinal concentrations in an HIV-infected patient with brain mucormycosis. J Antimicrob Chemother. 2011; 66(1):224–5. https://doi.org/10.1093/jac/dkj385 PMID: 20961910

61. Ruping MJ, Albermann N, Ebinger F, Burckhardt I, Beisel C, Muller C, et al. Posaconazole concentrations in the central nervous system. J Antimicrob Chemother. 2008; 62(6):1468–70. https://doi.org/10.1093/jac/dkn409 PMID: 18824458

62. Haynes RR, Connolly PA, Durkin MM, LeMonte AM, Smedema ML, Brizendine E, et al. Antifungal therapy for central nervous system histoplasmosis, using a newly developed intracranial model of infection. J Infect Dis. 2002; 185(12):1830–2. https://doi.org/10.1086/340825 PMID: 12085335
66. Mellinghoff IK, Winston DJ, Mukwaya G, Schiller GJ. Treatment of Scedosporium apiospermum brain abscesses with posaconazole. Clin Infect Dis. 2002; 34(12):1648–50. https://doi.org/10.1086/340522 PMID: 12032903

67. Pitisuttithum P, Negroni R, Graybill JR, Bustamante B, Pappas P, Chapman S, et al. Activity of posaconazole in the treatment of central nervous system fungal infections. J Antimicrob Chemother. 2005; 56 (4):745–55. https://doi.org/10.1093/jac/dki288 PMID: 16135526

68. Al-Abdely HM, Alkhunaizi AM, Al-Tawfiq JA, Hassounah M, Rinaldi MG, Sutton DA. Successful therapy of cerebral phaeohyphomycosis due to Ramichloridium mackenziei with the new triazole posaconazole. Med Mycol. 2005; 43(1):91–5. PMID: 15712614

69. Van den Driessche E, Vandepitte J, Van Dijck PJ, De Jonckheere J, Van de Voorde H. Letter: Primary amoebic meningoencephalitis after swimming in stream water. Lancet. 1973; 2(7835):971.

70. Lee J, Kim JH, Sohn HJ, Yang HJ, Na BK, Chwae YJ, et al. Novel cathepsin B and cathepsin B-like cysteine protease of Naegleria fowleri/excretory-secretory proteins and their biochemical properties. Parasitol Res. 2014; 113(8):2765–76. https://doi.org/10.1007/s00436-014-3936-3 PMID: 24832815

71. Debnath A, Parsonage D, Andrade RM, He C, Cobo ER, Hirata K, et al. A high-throughput drug screen for Entamoeba histolytica identifies a new lead and target. Nat Med. 2012; 18(6):956–60. https://doi.org/10.1038/nm.2758 PMID: 22610278

72. Hayat M. Chemical fixation. Basic techniques for transmission electron microscopy. San Diego, California: Academic Press, Inc.; 1986. p. 1–55.

73. Zhou W, Nguyen TT, Collins MS, Cushion MT, Nes WD. Evidence for multiple sterol methyl transferase pathways in Pneumocystis carinii. Lipids. 2002; 37(12):1177–86. PMID: 12617472

74. Kabsch W. XDS. Acta Crystallogr D Biol Crystallogr. 2010; 66(Pt 2):125–32. https://doi.org/10.1107/S0907444909047337 PMID: 20124692

75. Cowtan K. The Buccaneer software for automated model building. 1. Tracing protein chains. Acta Crystallogr D Biol Crystallogr. 2006; 62(Pt 9):1002–11. https://doi.org/10.1107/S0907444906022116 PMID: 16929101

76. Collaborative Computational Project, Number 4. Acta Crystallogr D. 1994; 50:760–3.

77. Emsley P, Cowtan K. Coot: model-building tools for molecular graphics. Acta Crystallogr D Biol Crystallogr. 2004; 60(Pt 12 Pt 1):2126–32.

78. Murshudov GN, Vagin AA, Dodson EJ. Refinement of macromolecular structures by the maximum-likelihood method. Acta Crystallogr D Biol Crystallogr. 1997; 53(Pt 3):240–55. https://doi.org/10.1107/S0907444996012555 PMID: 15299926