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In vivo active organometallic-containing antimycotic agents†

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Fungal infections represent a global problem, notably for immunocompromised patients in hospital, COVID-19 patient wards and care home settings, and the ever-increasing emergence of multidrug resistant fungal strains is a sword of Damocles hanging over many healthcare systems. Azoles represent the mainstay of antifungal drugs, and their mode of action involves the binding mode of these molecules to the fungal lanosterol 14 α -demethylase target enzyme. In this study, we have prepared and characterized four novel organometallic derivatives of the frontline antifungal drug fluconazole (**1a–4a**). Very importantly, enzyme inhibition and chemogenomic profiling demonstrated that lanosterol 14 α -demethylase, as for fluconazole, was the main target of the most active compound of the series, (N-(ferrocenylmethyl)-2-(2,4-difluorophenyl)-2-hydroxy-N-methyl-3-(1*H*-1,2,4-triazol-1-yl)propan-1-aminium chloride, **2a**). Transmission electron microscopy (TEM) studies suggested that **2a** induced a loss in cell wall integrity as well as intracellular features ascribable to late apoptosis or necrosis. The impressive activity of **2a** was further confirmed on clinical isolates, where antimycotic potency up to 400 times higher than fluconazole was observed. Also, **2a** showed activity towards azole-resistant strains. This finding is very interesting since the primary target of **2a** is the same as that of fluconazole, emphasizing the role played by the organometallic moiety. *In vivo* experiments in a mice model of *Candida* infections revealed that **2a** reduced the fungal growth and dissemination but also ameliorated immunopathology, a finding suggesting that **2a** is active *in vivo* with added activity on the host innate immune response.

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Introduction

Mycoses are one of the most common opportunistic infections worldwide, affecting poor as well as industrialized countries. Skin infections (e.g. athlete's foot), for example, affect 20–25% of the world's population and systemic opportunistic infections are the 4th common cause of bloodstream infections with a lifetime incidence of about 75% (e.g. in the case of *Candidiasis*).^{1–6} Even if such extensive morbidity does not lead to a high mortality rate, local opportunistic infections together with invasive fungal infections (IFI) have a deep impact on the entire health system because they suppress the immune system and thus, increase the risk to develop other/additional pathologies, which can be fatal to immunosuppressed people.^{7–16} Although potent drugs (e.g. clotrimazole) are applied in antifungal therapy, the infection rate is worryingly increasing every year. Moreover, incidences of resistance to conventional antimycotics are steadily increasing, especially for long term therapies that last from several weeks to months.¹⁷ Main mechanisms necessary for antifungal resistance are drug target mutations (e.g. mutations of *CYP51A1*),

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upregulation of efflux channels (e.g. major facilitator super-family (MFS) and ATP-binding cassette (ABC channels)), aneuploidy and loss of heterozygosity (LOH) and modulation of the stress response (e.g. translational signal pathways, RNA dependent response).^{18–21} Surprisingly, despite a pressing need for novel antifungal agent research and drug development, there is an apparent downturn in such activities.²² However, the recent COVID-19 pandemic has witnessed numerous examples of COVID-19 and *C. auris* co-infections, adding further pressure to healthcare systems.²³

Cell wall targeting is at the basis of modern antifungal therapy since it is absent in mammalian cells allowing for selectivity. A selective destabilization or inhibition of cell wall biosynthesis leads to metabolic onset, growth arrest and culminates in fungal death. For decades, Amphotericin B deoxycholate (AmB) has been the only available treatment for invasive fungal infections (IFIs).²⁴ However, during the last ten years, the growing number of IFI incidences has stimulated the introduction of four new classes of antifungal agents, namely polyene-based compounds, azoles, allylamines and echinocandins.^{25–28} Inhibition of cell wall biosynthesis is associated with an accumulation of sterol toxic precursors, cell wall destabilization and cellular stress. During the second half of the 20th century, antifungal research mainly focused on an intensive derivatization of accepted compounds (e.g. from ketoconazole to fluconazole or from AmB to Nystatin and AmB liposomes).^{19–21} Despite these efforts, to date, only seven compounds are included in the World Health Organization Essential Medicine list (WHO-EM, data of October 2013).^{1–6} Moreover, the benchmark drugs were all discovered almost 20 or more years ago and accounts of ineffective therapies have been reported for several of these agents especially towards *Candida albicans* (*C. albicans*) and *Aspergillus terreus*.^{1–6} For this reason, and encouraged by the spectacular results obtained with metal complexes (e.g. salvarsan, cisplatin, auranofin, ferroquine, ferrocifen, etc.) in other medicinal fields,^{29–38} some groups have decided to assess the potential of such compounds to fight fungal infections.^{39,40} Despite such a wealth of promising examples of metal-based drugs, there is a paucity of examples of such compounds applied in the field of antifungal therapy. Here, just sporadic reports confirmed the effectiveness of different transition metal complexes (e.g. Co(II), Ni(II), Cu(II), Pd(II) complexes) bearing a variety of ligands (e.g. azole moieties, thiosemicarbazones,

carboxamides, indoles) against several fungal strains or some organometallic moieties bound to active antifungal agents.^{41–53} However, all these studies generally provided only MIC₅₀ values without any further biological investigation such as target engagement and toxicity. Willing to grasp this opportunity, we embarked on a research program to unveil potent complexes against fungal infections with an emphasis on addressing cellular targets and mode of action. Herein, we present the identification of a metallocene-based lead compound against such infections.

Results and discussion

Synthesis and characterization

The two most common antifungal drugs inscribed in the List of Essential Medicine of the WHO and commercially available, namely clotrimazole and fluconazole (Fig. 1), belong to the azole class. The molecular target for both those azoles was found to be the enzyme lanosterol-14 α -demethylase.⁴⁷ Of utmost interest, mammalian cells use cholesterol instead of ergosterol. This important difference secures high selectivity for the reported antimycotics.⁵⁴ Recently, the broad therapeutic applications of clotrimazole as well as fluconazole against skin, genital and invasive fungal infections have generated comparison of primary or cross resistance, especially for *Candidiasis*.^{55,56} Motivated by the promising results obtained from the ferrocenyl derivatization of organic drugs such as tamoxifen and chloroquine to give ferrocifen and ferroquine, respectively, we envisaged applying the same concept to fight fungal infections.^{32,49,57–59} Fluconazole has been chosen as the parent drug because of its broad spectrum applications (e.g. systemic and topic administration, active against a large number of mycoses). The drug design took into consideration the mode of action of fluconazole and its interaction with the active site of the target enzyme. Computing binding geometry data showed how fluconazole activity is exerted by the interaction of the triazole moiety (*i.e.* the nitrogen atom highlighted in blue in Fig. 1) with the haem group present in the active site of the target enzyme (*i.e.* the iron atom).⁶⁰ This interaction is also favoured by the presence of the difluorophenyl group that is located in the enzyme active site, in (close) proximity to the hydrophobic binding cleft and interacts with it *via* π - π

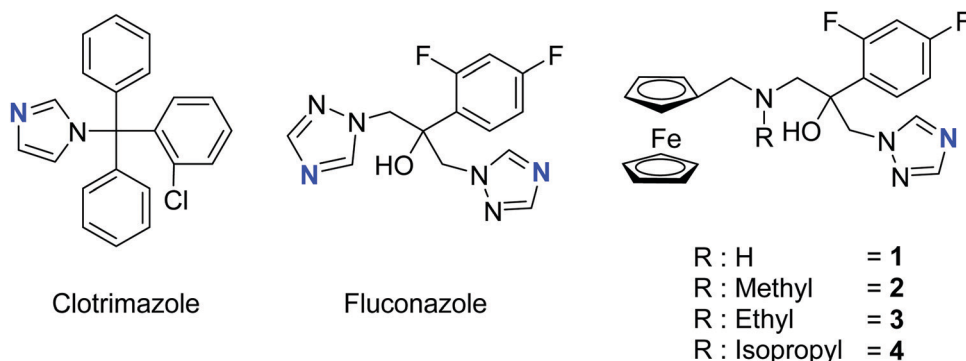
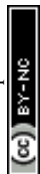


Fig. 1 Chemical structures of the main antifungal therapeutics present on the market (clotrimazole and fluconazole) and our fluconazenes **1–4**, highlighted in blue are the nitrogen responsible for the binding with the haem group.



stacking.⁶⁰ On the contrary, the second triazole moiety is involved in non-bonding interactions with several prosthetic groups present in the enzyme cavity. Interesting work by Sheng and co-workers, involving protein docking studies, demonstrated that organic modification with different functional groups at the C3 atom of fluconazole resulted in an increased activity in relation to the parent drug. This was supported by MIC₈₀ values. Therefore, in the present case, we aimed at derivatizing the fluconazole core on the triazole not involved in the binding pocket interaction, replacing it with a ferrocenyl moiety. Ferrocenyl derivatization is proposed to play an important role for the overall biodistribution and uptake (e.g. increasing the lipophilicity of the parent drug) as well as in potentially allowing for an additional redox-induced mode of action as observed for ferrocifen and ferroquine.^{32,57,58,61} Finally, different alkyl substituents were inserted at the bridging nitrogen in order to garner Structure–Activity Relationship (SAR) information. The target ferrocenyl compounds of this study, namely **1a–4a** can be seen in Fig. 1. Herein, we name them fluconazenes.

The synthesis of the new fluconazenes **1–4** can be visualized in Scheme 1. In the envisioned retrosynthesis (Scheme 1A), the target compounds would be obtained from the addition of a ferrocenylamine (**5**) on the epoxide **6**, easily accessible from 1,2,4-triazolo-ketone **7**, which is commercially available, by a Corey–Chaykovski epoxidation. The ferrocenylmethanamines (**5a–d**) were prepared from ferrocene carboxaldehyde (**8**) in two steps, following adapted procedures by Tice *et al.* and Baramée *et al.*^{62,63} Reaction of the aldehyde (**8**) with hydroxylamine hydrochloride under basic conditions yielded the oxime (**9a**). The imines (**9b–d**) were obtained by stirring **8** in the presence of a solution of the corresponding alkyl amines. The intermediates (**9a–d**) were reduced to amines (**5a–d**) with LiAlH₄ and NaBH₄, respectively. Spectroscopic data of **5a–d** matched those reported in the literature (Scheme 1B).^{62–64} Epoxide **6** was obtained in 76% yield by treating ketone **5** with a solution of trimethylsulfoxonium iodide ylide (Scheme 1C).^{60,65} The subsequent epoxide ring opening of **6** with the primary and secondary ferrocenemethanamines (**5a–d**) yielded compounds **1–4**.^{64,65} The derivative carrying the methylamine linker (**2**) was obtained in noticeably higher yield (46%) than the other derivatives (**1**, 30%; **3**, 31%; **4**, 20%). This can be attributed to the donating inductive effect of the alkyl groups on the N atom, which increases the nucleophilic strength of **9b–d**. In contrast, the growing steric hindrance of the ethyl and isopropyl group can explain the decreasing yield within the series of tertiary amines (**2** > **3** > **4**).

While fluconazole is achiral, with two identical 1,2,4-triazolyl-methyl side chains attached to the C2 carbon, our derivatives **1–4** are chiral. However, according to molecular docking experiments of chiral fluconazole derivatives by Sheng *et al.*, both *R* and *S* isomers, can interact with the active site of the *C. albicans* CYP51 through a similar binding mode,⁶⁶ thus avoiding the need for a chiral separation, or an asymmetric synthesis of a specific enantiomer.⁶⁷ The derivatives **1–4** were converted to the corresponding hydrochloride salts by treatment with HCl in acetone, following an adapted procedure of Bader *et al.* (Scheme 1D).⁶⁸ Overall, the new complexes **1a–4a** were all characterized *via* ¹H,

¹³C, ¹⁹F NMR, MS, IR and their purity was confirmed *via* elemental analysis and UPLC-MS (see Fig. S2–S29 in ESI†).

Stability

As potential drug candidates, the hydrochloride salts **1a–4a** need to be stable in DMSO, which is the administering medium for *in vitro* and *in vivo* studies, as well as in the biological medium, namely water. It was previously demonstrated that this can be problematic for metal-based drugs.^{69–71} The stability of **1a–4a** was assayed by dissolving the salts in deuterated solvents, namely neat DMSO-d₆ and neat D₂O, and by monitoring the samples using ¹H NMR spectroscopy over a period of 48 hours (see ESI† for details). It is worth mentioning that **1a** and **4a** are only sparingly soluble in D₂O. Generally, all four hydrochloride salts were stable for up to 24 hours and only little decomposition was observed for **2a** and **4a** in water after 48 h.

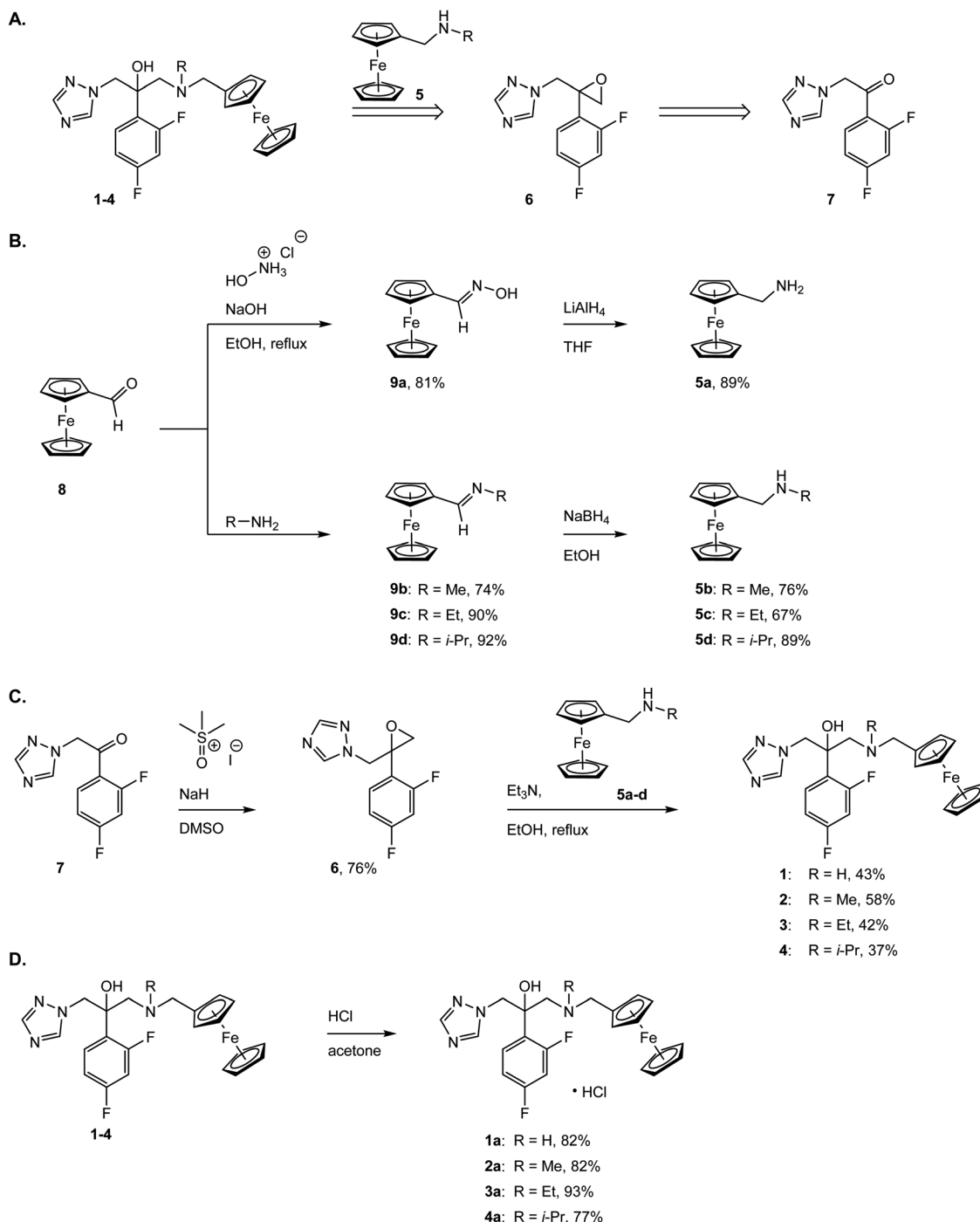
In vitro biological screening

To assess the antifungal activity of the new drug candidates, we employed a well-plate based assay recently developed in our group.³⁹ The first antimycotic investigation was performed on the solid and largely used model system *S. cerevisiae*. Therefore, *S. cerevisiae* was cultured and diluted to reach the beginning of the growing phase just before treatment. YPD agar terrain at increasing concentrations of the target complexes were poured on a 6 or 12-wells plate and the *S. cerevisiae* culture were then spotted on them and incubated for 24 h at 30 °C. Interestingly, all compounds displayed an improved antifungal activity compared to the parent drug (e.g. see Fig. 2). Small alkyl group-substituted tertiary amines (complexes **2a** and **3a**) as linkers, showed an improved profile, between 9–17-fold, compared with secondary amines (see Table 1).

To further evaluate the potential of the compounds, we investigated their host toxicity, *in vitro* mycotoxicity and cellular uptake. An effective inhibitor would need a good therapeutic window, with low host toxicity *versus* effective nM fungal toxicity. The first investigation on the new antimycotic drug candidates involved their possible influence on host cells and their efficiency towards a common fungal model. We investigated the anti-proliferative effect of **1a–4a** *in vitro* on human retinal pigment epithelial cell (RPE-1-hTert) and human fibroblast (MRC-5). As expected, fluconazole displayed only moderate effects on cell viability. All complexes displayed IC₅₀ values in the mid-high micromolar range, following the order of potency: isopropyl (**4a**) > ethyl (**3a**) > methyl (**2a**) > H (**1a**) (see Table 1). This suggests that a decrease in the size of the *N*-substituent (i.e., a decrease in lipophilicity) correlates with a decrease in host toxicity. Overall, while the organometallic compounds are slightly more toxic than fluconazole, the toxicity remains relatively moderate, especially taking into account that the toxicity towards yeast is much higher (see below).

In order to investigate the pharmacodynamic profile of the target antimycotic drug candidates, a series of uptake studies in *S. cerevisiae* were performed. The uptake measurements were performed *via* ICP-MS by detecting the free iron content in the culture medium and normalizing it against the colony density





Scheme 1 (A) Retrosynthetic analysis; (B) preparation of the primary (**7a**) and secondary (**7b–d**) ferrocenylmethanamines *via* the oxime (**9a**) and imine (**9b–d**) intermediates. Overall yields, comprising both steps: **7a**, 71%; **7b**, 56%; **7c**, 60%; **7d**, 81%; (C) synthesis of the organometallic fluconazole derivatives (**1–4**), which feature an amine-linked ferrocene moiety; (D) synthesis of the hydrochloride salts of **1–4** (**1a–4a**).

at different incubation times (1 h, 6 h and 18 h). The results were then compared with the survival rate measured *via* OD (see Fig. 3). The uptake of the target complexes followed the order **4a** < **1a** < **3a** < **2a** and fits very well with their time-dependent mycotoxicity. This suggests that complexes with sterically-hindered groups like an isopropyl or the less lipophilic complex **1a** showed a lower efficacy than ones with short chain alkyl groups (**2a**, **3a**).

In order to assess the possible increase in reactive oxygen species (ROS) level induced by **1a–4a**, possibly due to the presence of the ferrocenyl moiety, we performed a series of *in vitro* experiment to measure the ROS level upon treatment on *S. cerevisiae*. Briefly, *S. cerevisiae* was cultured in YPD medium overnight before treatment. After that, the fungal concentration was measured and normalized and the cultures were treated



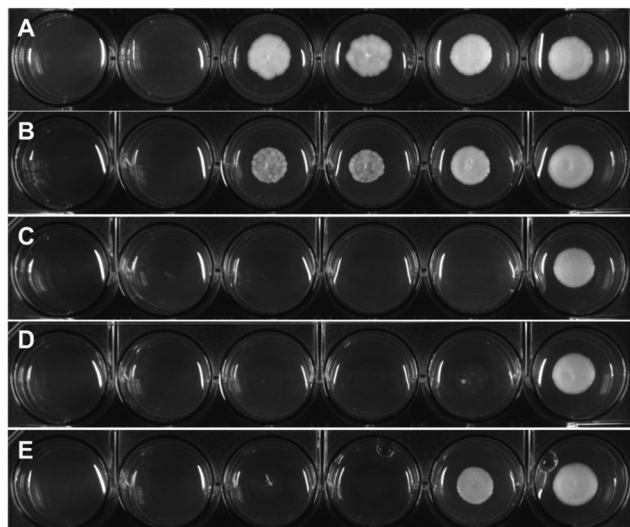


Fig. 2 Depicted example of a colony formation inhibition experiment; (A) *S. cerevisiae* treated with fluconazole; (B) *S. cerevisiae* treated with **1a**; (C) *S. cerevisiae* treated with **2a**; (D) *S. cerevisiae* treated with **3a**; (E) *S. cerevisiae* treated with **4a**. *S. cerevisiae* has been treated with increasing concentrations of the target drug candidates ranging from 2.5 up to 200 μM (higher concentration not depicted) and incubated at 30 $^{\circ}\text{C}$ for 24 h. The experiments with the drug candidates and with negative control of untreated colony have been performed in triplicate.

Table 1 Host toxicity of the new fluconacenes against RPE and MRC-5 human cell lines (expressed as IC_{50}) and inhibition of colony formation in *S. cerevisiae* treated with increasing concentrations of the **1a–4a** for 24 h at 30 $^{\circ}\text{C}$; fluconazole was used as parent drug for comparative purposes; values are expressed in μM . The activity of compounds **1a–4a** on *S. cerevisiae* was evaluated via a newly established colony formation assay³⁹

Compound	MRC5	RPE	<i>S. cerevisiae</i>
Fluconazole	> 100	> 100	51.0 \pm 3.7
1a	94.6 \pm 2.4	> 100	13.2 \pm 3.0
2a	32.2 \pm 0.1	74.7 \pm 5.2	3.02 \pm 2.18
3a	70.4 \pm 17.9	51.2 \pm 0.1	5.86 \pm 0.41
4a	59.4 \pm 2.8	48.4 \pm 2.9	9.62 \pm 0.56

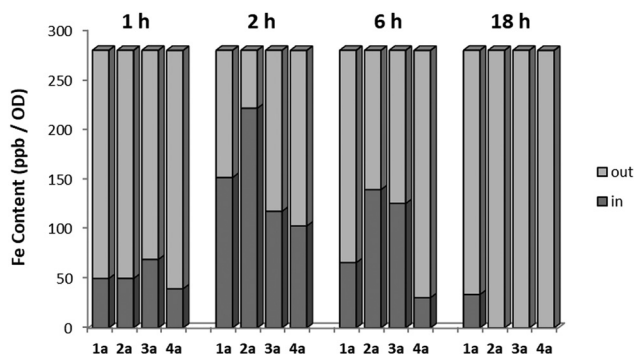


Fig. 3 Uptake studies compared to the colony survival rate measured in OD. The experiment has been performed on the medium of incubation and corrected with OD in order to take into account of the possible biological material loss due to toxicity of the compounds and which would lead to decrease in uptake. Negative controls of the untreated and of the medium alone have been also performed.

with YPD containing 5 μM of the different drug candidates at different time frames (1 h, 6 h and 18 h). After incubation, the colonies were washed, fresh medium containing H2DCF was added and the fluorescence of the hydrolyzed DCF was measured (see Fig. S1, ESI[†]). While the parent drug fluconazole displayed weak action at early stages, which increased with incubation time, the new drug candidates showed a higher ROS levels at an early stage. However, although the complexes proved to induce an increase of ROS levels, the relative ROS induction still remains moderate and it is probable that ROS production is not responsible for fungi death.

Morphology studies

The activity of **2a** at the biochemical level is also mirrored by morphologic features at the ultra-structural level. The morphologic effect on *S. cerevisiae* upon treatment with the target complex could be conveniently monitored by transmission electron microscopy (TEM). As can be seen in Fig. 4, the resulting images (selection of a large pool) suggested that **2a** induces a loss in wall integrity as well as intracellular features ascribable to late apoptosis or necrosis.

In vitro antifungal screening on clinical isolates

Motivated by the promising results obtained for the *in vitro* antifungal activities against the non-pathogenic model system *S. cerevisiae*, we performed further screening on clinical isolates. We again chose complex **2a** since it displayed the best mycotoxicity and, on the other side, showed low cytotoxicity against human cells. We tested this antifungal drug candidate towards *C. albicans* and non *albicans* strains, including fluconazole-resistant ($\text{MIC}_{50} > 100 \mu\text{M}$) and other mycotoxin-producing and pathogenic fungi, such as *Penicillium paneum*, *Aspergillus glaucus* and *Trichosporon asahi* (see Table 2). Of utmost interest, the new antifungal drug candidate displayed a very strong activity towards almost all investigated strains (only in the case of *A. glaucus*, *C. parapsilosis*, and *C. albicans* MFB005 and SC5314 was the MIC_{50} not in the nanomolar range). Moreover, the antimycotic potency of the complex did overcome the values of the parent drug up to a factor of 400-fold (e.g. *P. paneum*) and also showed activity towards azole resistant strains.

Fluconazole is a fungistatic drug and growth curve analysis performed on the *C. albicans* reference strain SC5314, a frequently used wild-type control, indicate a fungistatic effect of compound **2a** at MIC_{50} concentration for this specific strain (1.98 μM , Fig. S2, ESI[†]).

AOX enzyme inhibition

In order to further define the inhibitory capabilities of the compounds, they were also tested in the mitochondrial complex II and III pathway (succinate quinol reductase pathway (SQR)), two common targets for fungicidal treatment.^{72,73} As can be seen in Fig. 5, compound **1a** demonstrated no inhibitory activity towards either complex, confirming that the antifungal properties of these compounds are not due to off-target effects within mitochondrial respiration. However, while the other analogues demonstrate slightly improved inhibitory properties against the pathway when compared to **1a**, it is unlikely to be the source of fungicidal activity.



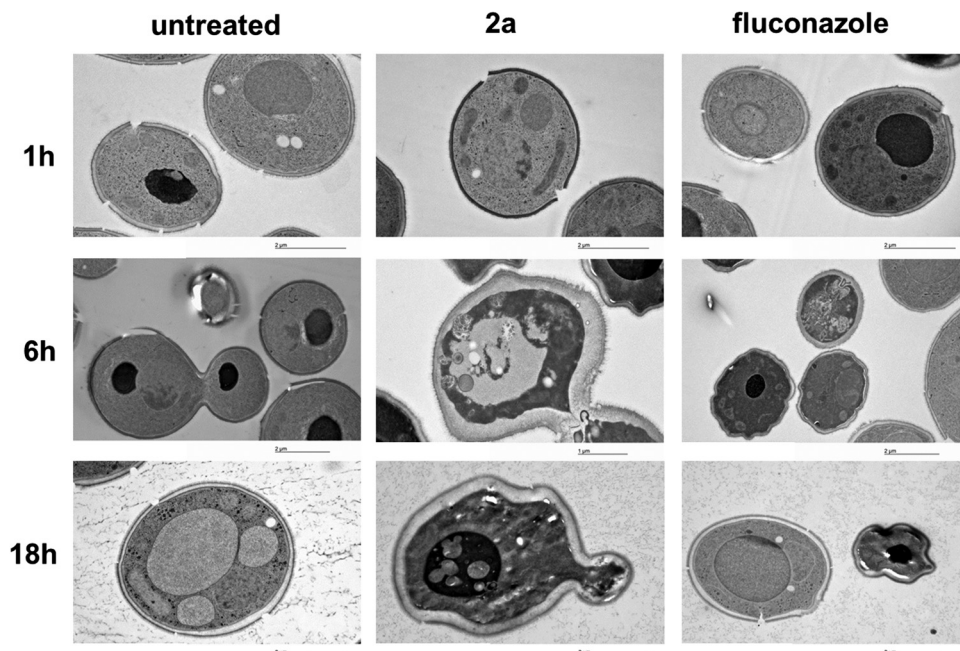


Fig. 4 Ultra-structural studies of untreated *S. cerevisiae*, treated with **2a** or with fluconazole tested at 5 μ M and 30 $^{\circ}$ C incubation temperature. The pictures showed starting with a 6 h incubation formation of vesicles, ascribable as apoptotic bodies and loss of membrane integrity. After longer incubation times a shrinking effect as well as nuclear blebbing could be also noted.

Table 2 MIC₅₀ values on a pool of fungal pathogens treated with **2a**

Strains	MIC ₅₀ 2a (μ M)	MIC ₅₀ fluconazole (μ M)
<i>C. albicans</i> SC5314	> 1	0.82
<i>C. albicans</i> MFB005 FS3	> 1	0.41
<i>C. albicans</i> YMS 102-2	1.00	> 100
<i>C. albicans</i> YMS 102-6	0.06	0.82
<i>C. glabrata</i> MFB005 FS4	0.06	0.41
<i>C. glabrata</i> RTT 199_3	1.00	> 100
<i>C. parapsilosis</i> MFB005 FS5	0.50	0.41
<i>C. parapsilosis</i> MFB070 N1	> 1	> 100
<i>C. tropicalis</i> RTT35-1	0.69	> 100
<i>C. tropicalis</i> RTT35-3	0.50	> 100
<i>P. paneum</i> MFB042 N1	0.25	> 100
<i>A. glaucus</i> MFB027 N1	> 1	0.82
<i>T. asahii</i> MFB034 N1	0.13	1.63

The compounds were also tested against alternative oxidases (AOX) from both *C. albicans* and *C. auris*. The proteins were expressed in a recombinant *E. coli* system to allow for more accurate data acquisition due to their low expression levels proving problematic from a detection standpoint. Again, compound **1a** demonstrated minimal inhibitory activity towards the AOX's, however compounds **3a** and **4a** showed moderate activity against both AOX's, comparable to values typically seen with non-competitive AOX inhibitors such as salicylhydroxamic acid (SHAM).⁷⁴ The parent compound, fluconazole, displayed no inhibition towards either pathway up to a concentration of 100 μ M.

Cytochrome P450 enzyme inhibition

Based on the very promising data obtained with complex **2a**, further biological experiments were performed on this complex

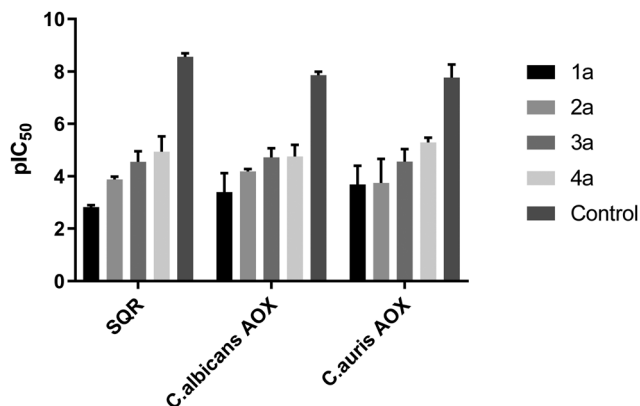


Fig. 5 pIC₅₀ values ($-\log$ IC₅₀) for all compounds against the mitochondrial complex II and III pathway (SQR) from rat liver mitochondria, and rAOX's from *C. albicans* and *C. auris*. Control denotes control compounds for each specific pathway, with antimycin A for SQR and ascofuranone for the AOX pathways. Data was collected in a 96-well plate format, following cytochrome c reduction (ΔA^{550}) for the SQR pathway, and NADH oxidation (ΔA^{340}) for the AOX pathway. All data are a mean average of 3 biological replicates \pm SEM.

to understand further its mechanism of action. Enzyme inhibition studies were performed on different cytochrome P450 enzymes by the service provider Cyprotex GmbH. The new fluconacene **2a** displayed an extremely strong inhibition profile throughout all the series of enzymes, with a larger efficacy in comparison with the parent drug fluconazole, between 4 and 150 fold (see Table 3) suggesting that, if this series of molecule are optimised towards clinical use, like fluconazole,⁷⁵ potential drug-drug interactions will need to be taken into account.



Table 3 Enzyme Inhibition studies of different cytochrome P450 enzyme by the compound **2a** and its parent drug (IC₅₀ in μ M). Standard deviation shown in upper script

Compound	CYP2C9	CYP2C8	CYP2C19	CYP3A4	CYP1A	CYP2D6
Fluconazole	13.9 ± 1.63	97.2 ± 27.9	3.02 ± 0.34	25.8 ± 2.45	> 100	> 100
2a	3.33 ± 0.71	0.64 ± 0.09	<0.4	0.4 ± 0.06	10.4 ± 0.41	2.71 ± 0.12

Chemogenomic studies

In order to investigate the mechanism of action of the new organometallic complex **2a**, chemogenomic assays were performed. The strength of the genome-wide chemogenomic screen is reflected in the clear identification of *ERG11*, encoding lanosterol 14- α -demethylase, as the main target of the drug (see Fig. 6 and Table S1, ESI[†]). *ERG11* is a key enzyme of the ergosterol biosynthesis pathway and the primary target of azole antifungals to which the tested compound **2a** belongs.

Azoles are known to induce *ERG11* gene expression,⁷⁶ which also holds true for compound **2a** yet to a lower extent than fluconazole as revealed by a screen in both susceptible *C. albicans* strains (Fig. S3A, ESI[†]) and in the highly fluconazole resistant strain YMS102-2 (Fig. S3B, ESI[†]). The latter strain showed a high expression of *ERG11* when grown in the absence of drug and *ERG11* gene expression was significantly ($FC = 1.69$; $P = 0.000086$) increased upon fluconazole exposure, while decreased ($FC = -1.81$; $P = 0.000193$) when grown in the presence of compound **2a** (Fig. S3B, ESI[†]). Hence, the improved antifungal activity of compound **2a** for the strain YMS102-2 might be partially explained by the significant reduction of *ERG11* gene expression, because overexpression of *ERG11* is known as an azole resistance mechanism in *Candida*.^{16,77}

Hence, future work should be directed to study if compound **2a** is able to reduce *ERG11 gene* expression also in other *ERG11* overexpressing *Candida* isolates in a similar way.

Our chemogenomic assay identified five other heterozygous deletion strains that in previous chemogenomic screens have been shown to be extremely sensitive to azole antifungals; these are *SET6*, *PDR5*, *PDR16*, *MED4* and *CDC39*. The latter two are essential nuclear genes involved in the regulation of transcription. *CDC39* acts as a component of the CCR4-NOT core complex⁷⁸ and *MED4* encodes a subunit of the RNA polymerase II mediator complex. In *Saccharomyces cerevisiae* and *C. glabrata* the mediator complex is a coactivator of the multiple drug resistance regulator *PDR1* that controls the activation of *PDR5* and *PDR16*.^{79–81} Similarly, also in *C. albicans* the mediator activator complex is recruited to the *CDR1* promoter,⁸⁰ which is the *C. albicans* ortholog of *PDR5*. *PDR5* is a multidrug ABC efflux pump that confers resistance to several chemicals including azoles and it mediates transmembrane transport of steroids.⁸² *PDR16* (ortholog of the *C. albicans* *PDR17*) is a phosphatidylinositol transfer protein involved in sterol biosynthetic process and in resistance to azole drugs.⁸³ Notably, the phosphatidylinositol signal transduction pathway that controls amongst others cell membrane and cell wall remodelling as well as downstream targets such as DNA repair, was recently suggested to

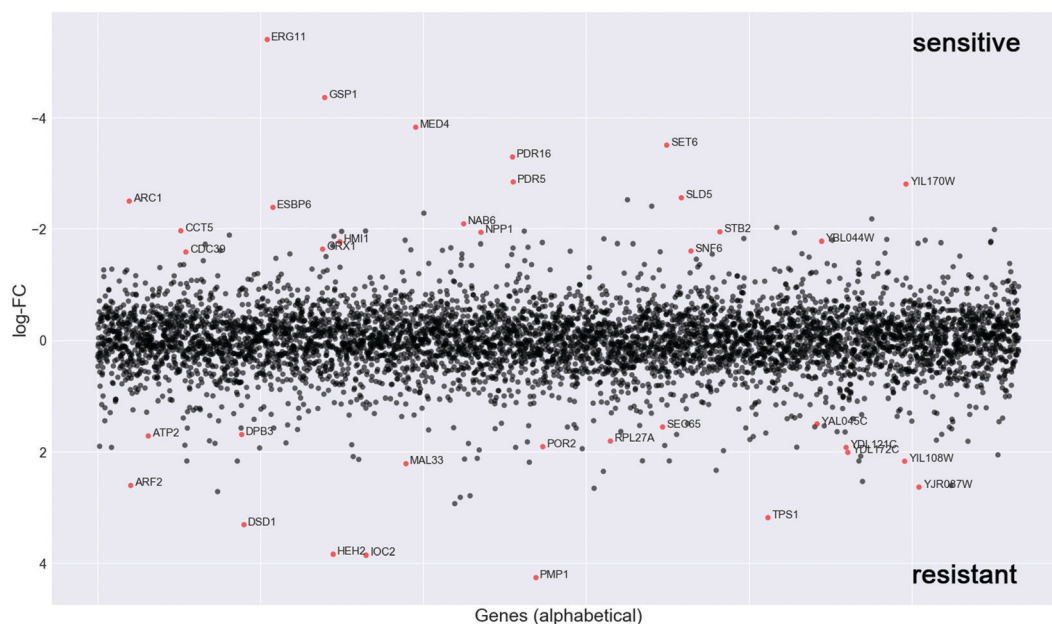


Fig. 6 Chemogenomic screen. To identify potential drug targets the pool of tagged 5936 *S. cerevisiae* heterozygous deletion mutants was grown competitively for 20 generations in the presence and absence of compound **2a**. Strain fitness was determined *via* high-throughput barcode sequencing and normalization to the untreated control. log₂ ratio (control intensity/treatment intensity) was calculated and plotted as a function of gene. The genome-wide readout of heterozygous mutants highly sensitive to compound **2a** included the known target of azoles, *ERG11* and novel targets such as the RAN GTPase *GSP1* (red dots: log₂FC > 1.5 and P value > 0.01).

play an important role in phenotypic antifungal drug resistance related to protein synthesis.¹⁸ Indeed many of the identified deletion strains that are highly sensitive to compound **2a** were heterozygous for genes involved in protein synthesis (e.g. *ARC1*, *RPS5*, *MED4*, *CDC39*, *SNF6*, *CCT5*, *GSP1*) (Fig. 4 and Table S1, ESI†).

Furthermore, compound **2a** targets nuclear genes that are not only essential for RNA transcription, processing and

transport (*GSP1*, *MED4*, *CDC39*) but also essential for DNA replication (*SLD5*), and these together with gene targets functioning in the correct folding of actin and tubulin (*CCT5*) (Fig. 6) point to a possible interaction of compound **2a** with cell division.

Taken together, the genome-wide profiling of the *in vivo* cellular response to compound **2a** identified candidate protein targets (Fig. 6 and Table S1, ESI†), which hint to a mechanism

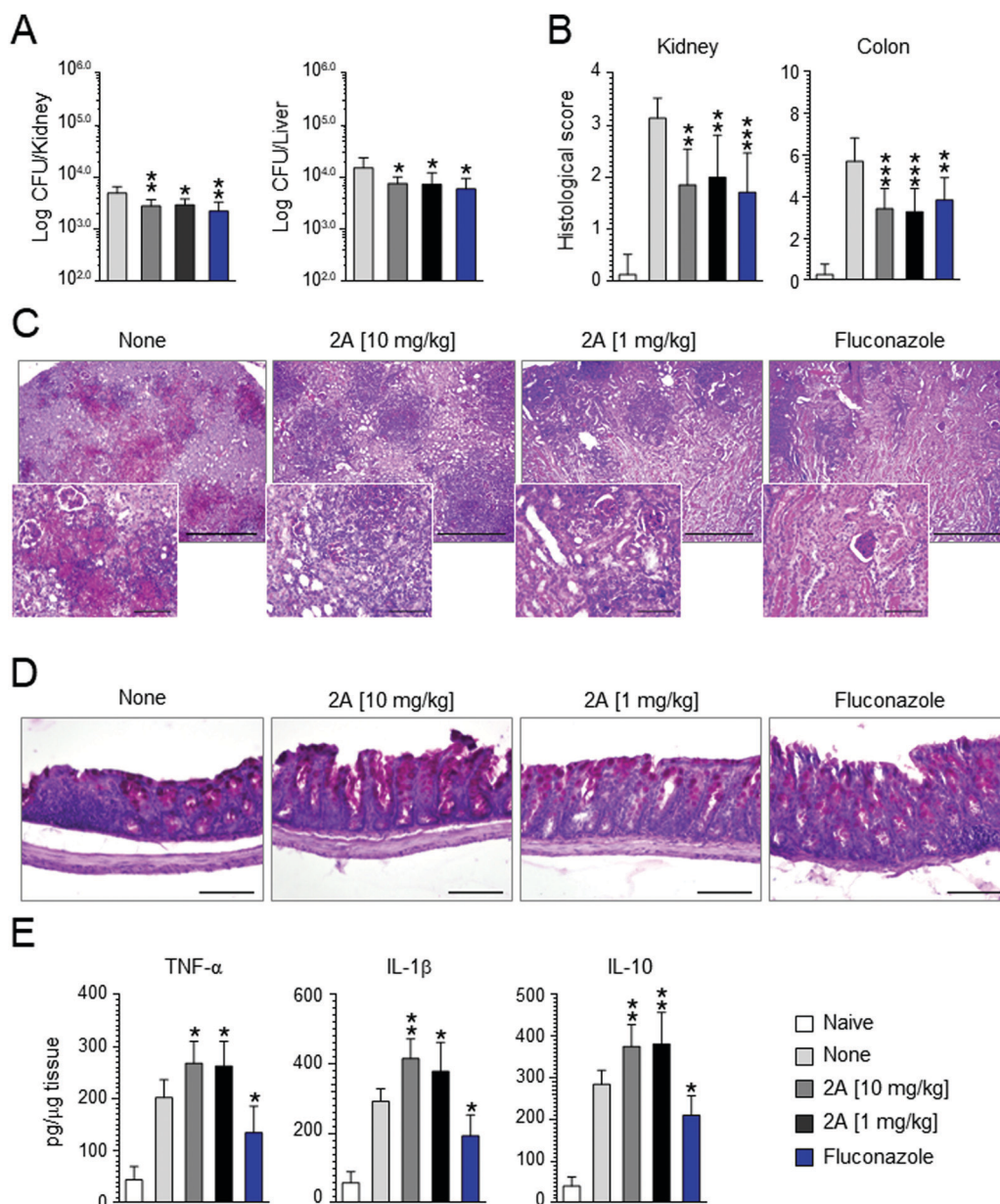


Fig. 7 Effect of the compound **2a** on *C. albicans* (SC-5314) systemic infection. C57BL/6 mice were infected *via* systemic route with 1×10^6 *C. albicans* yeasts. Fluconazole and compound **2a** were administered intragastrically daily, beginning the day of the infection until mice sacrifice (4 days after the infection). Control mice received the diluent alone. (A) Fungal growth. (B) Histological scores and (C and D) analysis (10× magnification and 40× in the inset) of the kidneys and colon of infected and treated mice. (E) Levels of cytokines in kidney homogenates. For histological scores of kidneys, score 0 = no inflammation, score 1 ≤ 3 foci of inflammation, score 2 = 4 to 6 foci of inflammation, score 3 ≥ 6 foci of inflammation, but less than 25% of kidney affected, score 4 ≥ 25% of kidney affected. For histological scores of colon, the total histological score was obtained by summing the four histological components scores: 'inflammation extent', 'damage in crypt architecture', 'hyperemia/edema', 'grade of accumulation with inflammatory cells'. Data are expressed as mean ± SD. **P* < 0.05, ***P* < 0.01, ****P* < 0.005, treated vs. untreated C57BL/6 mice (*n* = 7 mice per group, representative of three experiments). Naive, uninfected mice. None, untreated mice.



of drug action that goes beyond the main targets of azole antifungals, which might explain its elevated activity towards azole resistant strains (Table 1).

In vivo studies

Motivated by these promising results, we decided to perform *in vivo* experiments in a mice model of *Candida* infection. As positive control, fluconazole was used. It significantly improved pathology and cured mice from the infection, as judged by the decreased fungal growth in the kidney and dissemination to the liver (see Fig. 7A), reduction of the inflammatory pathology in the kidney and colon (see Fig. 7B–D) and downregulation of pro-inflammatory cytokine (IL-1 β and TNF- α) levels (Fig. 7D). Similar results were obtained with compound **2a** that proved to be effective in reducing fungal growth and dissemination (see Fig. 7A) as well as tissue inflammatory pathology (see Fig. 6B for histological scores) at a dose as low as 1 mg kg⁻¹ (Fig. 7C and D). Of great interest, at variance with fluconazole, compound **2a** also promoted a vigorous pro-inflammatory cytokine response promptly controlled by the associated high levels IL-10 (Fig. 7E). These findings indicate that compound **2a** is active *in vivo* and that its activity may encompass an action on innate immune response, consistent with the well known synergistic activity of azoles, including fluconazole, with antifungal effector activities, such as phagocytosis, radicals production and Toll-like receptors activation.⁸⁴

Conclusion

Four novel ferrocene-based derivatives (**1a–4a**) of the frontline antifungal drug fluconazole have been synthesized and characterized. All four organometallic derivatives were tested for antifungal activity against the model system *S. cerevisiae* and showed an impressively improved activity compared to fluconazole. Importantly, this activity was confirmed on a panel of clinical isolates with the best candidate of this study (complex **2a**). An improvement of the minimum inhibitory concentration of up to 400 times compared to fluconazole was observed and activity against azole-resistant strains was clearly demonstrated for **2a**. During *in vivo* experiments in mice model of *Candida* infections, compound **2a** was found to diminish the growth of the fungus and to greatly improve the inflammatory pathology in the kidney and colon. Overall, this study further demonstrates the potential of organometallic compounds in medicine. By merely inserting a ferrocenyl moiety in a well-known drug, namely fluconazole, resistance can be overcome, although the main target of the novel antimycotic agent is the same as that of the organic drug (lanosterol 14 α -demethylase), as demonstrated by chemogenomic profiling. The ferrocenyl insertion allows for an additional mode of action.

Ethical statement

Mouse experiments were performed according to Italian Approved Animal Welfare Authorization 360/2015-PR and Legislative Decree 26/2014 regarding the animal license obtained by the Italian

Ministry of Health lasting for 5 years (2015–2020). Animals were treated as the guidelines of EAMC.

Author contributions

All authors were involved with the design and interpretation of experiments and with the writing of the manuscript. The synthesis and characterization of the compounds were performed by R. R., L. M., S. J., J. N., Y. L., and J. H. TEM studies were carried out by A. K. Biological experiments were performed by R. R., T. W., N. T., A. L. M., and L. Y. Chemogenomic studies were carried out by T. W. and M. M. Animal testing was carried out by R. G., M. P. and L. R. All authors have given approval to the final version of the manuscript.

Conflicts of interest

The authors declare no competing financial interests.

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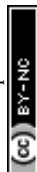
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