



# Commercial AHAS-inhibiting herbicides are promising drug leads for the treatment of human fungal pathogenic infections

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The increased prevalence of drug-resistant human pathogenic fungal diseases poses a major threat to global human health. Thus, new drugs are urgently required to combat these infections. Here, we demonstrate that acetohydroxyacid synthase (AHAS), the first enzyme in the branched-chain amino acid biosynthesis pathway, is a promising new target for antifungal drug discovery. First, we show that several AHAS inhibitors developed as commercial herbicides are powerful accumulative inhibitors of *Candida albicans* AHAS ( $K_i$  values as low as 800 pM) and have determined high-resolution crystal structures of this enzyme in complex with several of these herbicides. In addition, we have demonstrated that chlorimuron ethyl (CE), a member of the sulfonylurea herbicide family, has potent antifungal activity against five different *Candida* species and *Cryptococcus neoformans* (with minimum inhibitory concentration, 50% values as low as 7 nM). Furthermore, in these assays, we have shown CE and itraconazole (a P450 inhibitor) can act synergistically to further improve potency. Finally, we show in *Candida albicans*-infected mice that CE is highly effective in clearing pathogenic fungal burden in the lungs, liver, and spleen, thus reducing overall mortality rates. Therefore, in view of their low toxicity to human cells, AHAS inhibitors represent a new class of antifungal drug candidates.

acetohydroxyacid synthase | *Candida albicans* | antifungal drug | crystal structure | herbicide

The number of diagnosed pathogenic fungal infections continues to rise significantly each year, with an estimated 1 billion people worldwide now at risk (1) and ~2 million people per annum succumbing to these infections (2), a number that exceeds the annual death toll for tuberculosis or malaria (3). Despite this, major advances in the search for novel antifungal treatments have been a rarity (4). Individuals commonly affected by fungal infections (e.g., candidiasis, cryptococcosis, aspergillosis, or *Pneumocystis pneumonia*) are transplant recipients, the immunocompromised suffering HIV, or patients receiving immunosuppressant therapy. However, existing antifungal therapeutics are limited to just a few classes of drugs (e.g., polyenes, azoles, echinocandins, and fluorinated pyrimidine analogs), and these are costly to produce. Furthermore, treatment using these medications requires high doses to be taken over a prolonged time, which can result in serious side effects and lack of compliance (5, 6). In addition to these factors, several clinically relevant pathogenic fungi, including *Aspergillus fumigatus*, *Candida glabrata*, *Candida krusei*, and *Cryptococcus* spp., are intrinsically resistant to some first-line antifungals (4, 5, 7). Moreover, acquired resistance to all these classes of antifungal drugs has been extensively reported (8–11), and the prevalence of multidrug-resistant strains of pathogenic fungi is continuing to rise (4, 12). Thus, it is clear that new drugs are urgently needed to combat this major threat to human health.

A potential target for the development of new antifungal drugs is acetohydroxyacid synthase (AHAS; EC 2.2.1.6), the first enzyme in

the branched-chain amino acid (BCAA) biosynthesis pathway. BCAAs are synthesized by a common pathway in plants and microorganisms; however, this pathway is absent in animals, making it an especially attractive drug target (13). It has previously been shown that AHAS activity is crucial for survival of *Candida albicans* and *Cryptococcus neoformans* in vitro, and BCAA auxotrophic strains of these two fungal pathogens are avirulent in vivo (14, 15). These findings suggest AHAS inhibitors could be useful as antifungal agents.

It has been 30 y since the first AHAS inhibitors were introduced as commercial herbicides, with five families of compounds [i.e., sulfonylureas (SUs), imidazolinones (IMIs), pyrimidinyl-benzoates (PBs), triazolopyrimidines (TPs), and sulfonylamino-carbonyl-triazolinones (SCTs)] still in use today (16), which is a testament to their robustness as biocidal agents. Further enhancing their credentials as drug candidates, these compounds display low toxicity in mammals ( $LD_{50}$  in rat >5 g/kg), which is expected since animals do not possess the BCAA pathway (13, 17). Studies on AHAS inhibitors as antifungals have been sporadic and inconclusive. Chlorimuron ethyl (CE) (Fig. 14), an SU herbicide, has previously been shown to inhibit the growth of *C. albicans* in culture with a minimum inhibitory concentration, 50% ( $MIC_{50}$ ) of 2  $\mu$ M (18), and several noncommercial TPs have activity against *C. albicans*, *C. neoformans*, and *A. fumigatus* with  $MIC_{50}$  values in the range of 1–4  $\mu$ g/mL (19). However, beyond these

## Significance

Human fungal pathogens resistant to conventional therapeutics pose a major threat to global human health. Thus, there is an urgent need to discover new antifungal drugs that act via novel mechanisms of action. Here, we show that commercial herbicides that inhibit acetohydroxyacid synthase (AHAS) have potent and broad-spectrum antifungal activity in vitro and that chlorimuron ethyl, a member of the sulfonylurea herbicide family, has antifungal activity in a mouse model. Thus, this study shows that AHAS inhibitors have strong potential to be developed into potent antifungal therapeutic agents.

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Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, [www.wwpdb.org](http://www.wwpdb.org) (PDB ID codes 6DEK, 6DEL, 6DEM, 6DEN, 6DEO, 6DEP, 6DEQ, 6DER, and 6DES).

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reports, a comprehensive study into the suitability of the AHAS-inhibiting herbicides as antifungal agents has yet to be undertaken.

## Results

**CaAHAS and CnAHAS Inhibition by Commercial Herbicides.** Representatives of the five families of herbicides that target AHAS (*SI Appendix, Fig. S1*) were tested as inhibitors of *C. albicans* (*Ca*) AHAS and *C. neoformans* (*Cn*) AHAS. Members of the SU and TP families are the most potent inhibitors (Table 1), with CE having a  $K_i$  of 24.26 nM for *Ca*AHAS and 119.7 nM for *Cn*AHAS (Fig. 1*B*), metosulam (MT) a  $K_i$  of 800 pM for *Ca*AHAS, and cloransulam methyl a  $K_i$  of 35.3 nM for *Cn*AHAS. Members of the SCT, PB, and IMI families inhibit *Ca*AHAS, but with  $K_i$  values only in the micromolar range, while the SCTs and PBs inhibit *Cn*AHAS, with  $K_i$  values in the millimolar range (Table 1). *Cn*AHAS is partially resistant to SU herbicides, including bensulfuron methyl (BSM), ethoxysulfuron (ES), and sulfometuron methyl (SM). This lower inhibitory activity appears to be due to the P188A and A191L substitutions in the *Cn*AHAS herbicide binding pocket compared with *Ca*AHAS (discussed below) (*SI Appendix, Fig. S24*). Two noncommercial SUs derived from the chemical structure of CE, iodomuron ethyl (IE) and iodomuron methyl (IM) (18) (*SI Appendix, Fig. S1*), were also tested as inhibitors of the fungal enzymes. Compared with CE, these compounds show a two- to fourfold reduction in the  $K_i$  values against *Ca*AHAS; however, this improvement in binding affinity is not extended to *Cn*AHAS (Table 1).

Recently, we demonstrated AHAS inhibition by commercial herbicides fits a model of time-dependent accumulative inhibition from which first-order rate constants of enzyme inactivation ( $k_{iapp}$ ) and enzyme recovery ( $k_3$ ) can be determined (20). Here, we found that the five families of herbicides show different levels of reversible accumulative inhibition for *Ca*AHAS and *Cn*AHAS (Table 1 and *SI Appendix, Fig. S3*). The  $k_{iapp}$  and  $k_3$  values were determined by curve fitting to Eq. 3 (20) and show that three families of herbicides (i.e., SUs, TPs, and SCTs) are highly effective in promoting the inactivation of *Ca*AHAS ( $k_{iapp}$  values from 0.33 to 21.26  $\text{min}^{-1}$ ). Despite the high level of conservation observed in the herbicide-binding site of *Cn*AHAS and *Ca*AHAS (*SI Appendix, Fig. S24*), the former enzyme is less prone to undergo inactivation by commercial herbicides with  $k_{iapp}$  values in the range of 0.13–3.99  $\text{min}^{-1}$ .  $k_3$  values for *Ca*AHAS in the presence of IE, and for *Cn*AHAS in the presence of pyroxulam (PYS), flumetsulam (FT), or propoxycarbazone (PC) are so low that the inactivation evoked by these herbicides is virtually irreversible (Table 1). This implies that, when  $k_{iapp} \gg k_3$ , the enzyme's ability to recover from herbicide-induced inactivation determines the potency of accumulative inhibition ( $k_{iapp}/k_3$ ).

**CaAHAS Overall Structure.** Crystal structures of the catalytic subunit of uninhibited *Ca*AHAS and *Ca*AHAS in complex with eight different herbicidal inhibitors were determined at resolu-

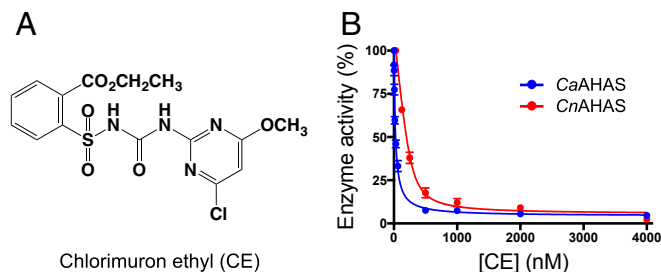
tions ranging from 1.81 to 2.97 Å (*SI Appendix, Table S1*). *Ca*AHAS is homotetrameric (Fig. 2*A*) consistent with *Arabidopsis thaliana* (*At*) AHAS but different from *Saccharomyces cerevisiae* (*Sc*) AHAS, which is homodimeric (*SI Appendix, Fig. S4A*). The overall fold for the polypeptide of uninhibited *Ca*AHAS closely resembles that of uninhibited *Sc*AHAS (*SI Appendix, Fig. S4B*) but differs significantly in the capping region ["mobile loop" (Q576–Y591) and "C-terminal arm" (P646–Y683)] (Fig. 2*B*). In uninhibited *Sc*AHAS the majority of residues in these two regions are not observed in the electron density maps or adopt a different fold (e.g., an antiparallel  $\beta$ -sheet from V583 to H597 is observed in *Sc*AHAS that changes to a helix when herbicide binds) (21) (*SI Appendix, Fig. S4B*). In this respect, *Ca*AHAS more closely resembles that of uninhibited *At*AHAS (rmsd = 0.699 Å when 428 C $\alpha$  of a single polypeptide are superposed) that has a folded capping region and preformed binding pocket for the herbicide (22) (*SI Appendix, Fig. S4B*).

In the uninhibited *Ca*AHAS structure the entire polypeptide is well-resolved, except for the regions between I192–Q198 and N267–A276. Importantly, I192–Q198 is in the vicinity of the active site, forming part of the substrate access channel and herbicide-binding site (*SI Appendix, Fig. S5A*). Upon herbicide binding, these residues are locked in position by direct contacts to the herbicides (*SI Appendix, Fig. S5B*). The N267–A276 region is remote from the active site and herbicide-binding site and not likely to influence the binding of substrates, cofactors, or inhibitors (Fig. 2*B*).

The electron densities for FAD, thiamine diphosphate (ThDP), and magnesium ion ( $\text{Mg}^{2+}$ ) and for the herbicides are all unequivocal, noting that in all of the enzyme-inhibitor complexes the chemical structure of ThDP is chemically modified (Fig. 2*C*). When the SCT, PC, binds ThDP is modified to thiamine thiazolone diphosphate (ThThDP), and when the SUs, BSM, IE, and IM, or the TP, PS, bind the cofactor is altered to thiamine aminoethenethiol diphosphate (ThAthDP). However, the electron densities in the *Ca*AHAS–SM and the *Ca*AHAS–MT complexes suggest both ThThDP and ThAthDP are present. In contrast, the crystallographic data of the *Ca*AHAS–CE complex suggests that ThAthDP along with an intermediate product of degradation of ThDP, thiamine formamidoethenethiol diphosphate (ThFAthDP), are present. The assignment of these chemical modifications was confirmed by MS of the cofactor isolated from a solution of enzyme incubated with inhibitor (*SI Appendix, Fig. S6*). Herbicide (e.g., PB, SCT, and TP families)-induced modifications to ThDP have also been observed for *At*AHAS (23, 24), and these are the result of oxidative reactions that promote the formation of peracetate in the active site (24) and the subsequent alteration of ThDP (Fig. 3). Importantly, ThThDP has been attributed to have antimetabolic activity (ThDP antagonist) (25) and reactive oxygen species, such as peracetate, inhibit *C. albicans* growth via NADH depletion (26). These factors, in addition to ThDP's being replenished at a metabolic cost to cells, could be major contributors to biocidal activity.

**CaAHAS Structures in Complex with Herbicide.** To understand the molecular basis of herbicide inhibition of *Ca*AHAS, we cocrystallized this enzyme in complex with five SUs, CE, BSM, SM, IE, and IM; two TPs, MT and PS; and the SCT, PC (*SI Appendix, Table S1*). The overall fold of these structures is similar to that of the uninhibited enzyme, with rmsd values between 0.145 and 0.271 Å when all C $\alpha$  atoms are superposed, but in these structures the I192–Q198 region, which is involved in herbicide binding (*SI Appendix, Fig. S5*), is now fully resolved.

**Herbicide Binding Mode to CaAHAS.** All eight herbicides are located in a binding pocket (Figs. 2*D* and 4) that is situated above the active site and thus operate by preventing the substrate, pyruvate, access to ThDP (*SI Appendix, Fig. S5A*). This pocket is formed by 12 residues and FAD (*SI Appendix, Fig. S7*) and is the putative binding site of quinones, which inhibit AHAS activity via FAD



**Fig. 1.** Inhibition of *Ca*AHAS and *Cn*AHAS by CE. (A) Chemical structure of CE. (B) Percentage of enzyme activity vs. inhibitor concentration. The solid line represents the best fit to the data using the equation for weak to medium binding (Eq. 1; *Cn*AHAS) or for tight-binding (Eq. 2; *Ca*AHAS). Error bars represent the SD of the mean (SEM) ( $n = 3$ ).

**Table 1. Inhibition constants and kinetic rate constants of accumulative inhibition of commercial herbicides for fungal AHASs**

Inhibitor	<i>C. albicans</i>				<i>C. neoformans</i>			
	$K_i$ , nM	$k_{iapp}$ , min <sup>-1</sup>	$k_3$ , min <sup>-1</sup>	$k_{iapp}/k_3$	$K_i$ , nM	$k_{iapp}$ , min <sup>-1</sup>	$k_3$ , min <sup>-1</sup>	$k_{iapp}/k_3$
SU								
IE	6.3 ± 0.7	0.33 ± 0.05	3.3 × 10 <sup>-11</sup>	10 × 10 <sup>10</sup>	110.5 ± 10.8	0.17 ± 0.02	0.023	7.4
IM	12.7 ± 1.1	1.10 ± 0.15	0.020	55.0	472.7 ± 25.5	0.17 ± 0.03	0.056	3.0
CE	24.3 ± 2.5	1.00 ± 0.32	0.045	22.2	119.7 ± 4.2	0.73 ± 0.03	0.022	33.0
BSM	29.9 ± 2.7	1.82 ± 0.25	0.064	28.4	NI	—	—	—
ES	69.3 ± 6.9	1.69 ± 0.20	0.082	20.6	NI	—	—	—
CS	570 ± 59	NP	NP	—	769 ± 89 × 10 <sup>3</sup>	NP	NP	—
SM	630 ± 51	0.67 ± 0.14	0.081	8.3	NI	—	—	—
TP								
MT	0.8 ± 0.1	3.39 ± 0.69	0.023	147.4	65.4 ± 4.9	1.14 ± 0.06	0.029	39.3
DS	1.7 ± 0.8	2.31 ± 0.20	0.071	32.5	901.0 ± 72.9	0.13 ± 0.02	0.007	18.6
PS	2.4 ± 0.3	15.26 ± 3.42	0.074	206.2	331.1 ± 20.0	2.18 ± 0.11	0.037	58.9
CSM	4.6 ± 1.2	3.86 ± 0.54	0.052	74.2	35.3 ± 3.7	0.94 ± 0.06	0.018	52.2
FS	10.1 ± 1.7	5.97 ± 0.40	0.068	87.8	590 ± 41.0	0.67 ± 0.03	0.025	26.8
PYS	212 ± 18	2.26 ± 0.10	0.067	33.7	320 ± 29 × 10 <sup>3</sup>	3.99 ± 0.60	5.3 × 10 <sup>-10</sup>	7.5 × 10 <sup>9</sup>
FT	773 ± 62	3.78 ± 0.27	0.068	55.6	137 ± 9 × 10 <sup>3</sup>	2.00 ± 0.59	1.5 × 10 <sup>-10</sup>	1.4 × 10 <sup>10</sup>
SCT								
PC	484 ± 37	1.83 ± 0.28	0.080	22.9	9.4 ± 1.6 × 10 <sup>3</sup>	0.19 ± 0.04	3.7 × 10 <sup>-11</sup>	5.1 × 10 <sup>9</sup>
TCM	28 ± 6 × 10 <sup>3</sup>	21.3 ± 4.4	0.109	195.0	1.2 ± 0.2 × 10 <sup>6</sup>	NP	NP	—
PB								
BS	2.6 ± 0.3 × 10 <sup>3</sup>	NP	NP	—	5.8 ± 0.6 × 10 <sup>6</sup>	NP	NP	—
PTB	403 ± 37 × 10 <sup>3</sup>	NP	NP	—	765 ± 65 × 10 <sup>3</sup>	NP	NP	—
IMI								
IQ	716 ± 61 × 10 <sup>3</sup>	NP	NP	—	NI	—	—	—
IT	4.3 ± 0.7 × 10 <sup>6</sup>	NP	NP	—	NI	—	—	—

BS, bispyribac; CS, chloresulfuron; CSM, cloransulam methyl; DS, diclosulam; FS, florasulam; IQ, imazaquin; IT, imazethapyr; NI, no inhibition detected; NP, no reversible accumulative inhibition present; PS, penoxsulam; PTB, pyriithiobac; TCM, thiencazabone methyl.

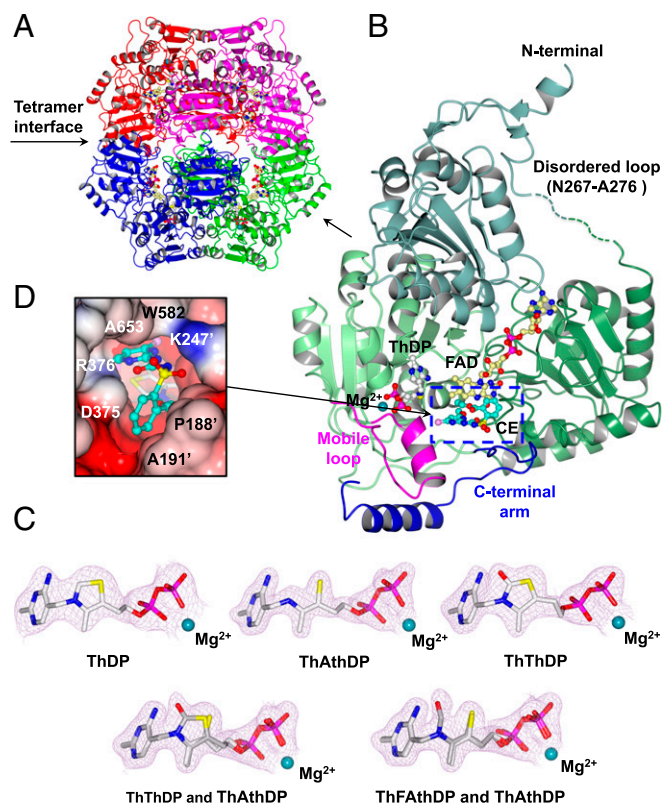
oxidation (27). Key to the binding of the three families of herbicides are the contacts formed with P188, F187, K247, M350, R376, M578, and W582 (Fig. 4 and *SI Appendix, Fig. S8*). Except for P188, all these residues are highly conserved and located in the deepest part of the herbicide-binding site (*SI Appendix, Fig. S2B*). K247 and R376 contribute with up to five hydrogen bonds when the SUs or the SCT, PC, bind (Fig. 4A and C), whereas between four to six hydrogen bonds are formed with the TPs (Fig. 4B). In addition, the guanidinium group of R376 forms a  $\pi$  cation interaction with the aromatic ring of the herbicide. The remaining residues of the herbicide-binding site form a large hydrophobic surface (*SI Appendix, Fig. S7*) that accounts for 30–50 noncovalent interactions. Of these, probably the most important are the  $\pi$  stacking interaction formed between the side chain of W582 and the heterocyclic ring of the herbicide and the carbon- $\pi$  interaction with P188, both of which anchor the herbicide in the binding pocket. A more detailed analysis of the binding mode of each family of herbicides is available as *SI Appendix*.

An overlay of the CE complex with an example of the other herbicide complexes shows that the three different families partially overlap their modes of binding to *Ca*AHAS (*SI Appendix, Fig. S9*). The aromatic rings, and the sulfonyl and carbonyl groups of the SUs and the SCT, PC, are the only components of these two families that adopt a similar position and conformation. In contrast, the TPs have a different orientation, with the aromatic ring overlapping with the ester substituent of the SUs or SCTs and the heterocyclic and linker moieties inserted deepest in the herbicide-binding site.

**Conformational Changes in *Ca*AHAS upon Herbicide Binding.** A comparison of uninhibited *Ca*AHAS and the *Ca*AHAS-inhibitor complexes shows that, in the uninhibited enzyme, R376 and W582 protrude toward the herbicide-binding cavity and occupy

the location taken by the heterocyclic ring of all three families of herbicides (*SI Appendix, Fig. S9*). This implies that these residues would block herbicide binding. However, the side chains of R376 and W582 change their conformation so as to make numerous contacts with the herbicides. The adjustments of W582 are the most important for herbicide binding because these allow the  $\pi$  stacking interaction that locks the herbicides into the binding pocket. Other residues in the herbicide-binding site, including P188, A191, K247, and D375, undergo less pronounced side-chain and backbone adjustments to accommodate the herbicides (*SI Appendix, Fig. S9*). Compared with *At*AHAS (22, 23) the herbicide-binding site in *Ca*AHAS is much more rigid and ready to receive inhibitors with only minor conformational changes, a feature of importance for the rational design of alternative *Ca*AHAS inhibitors.

**Effect of P188A and A191L Substitutions in *Cn*AHAS.** To understand why some SU herbicides are potent inhibitors of *Ca*AHAS but fail to inhibit *Cn*AHAS we performed in silico mutation and energy minimization of P188A and A191L on the structures of uninhibited *Ca*AHAS and for the CE and MT complexes. These models show that in the P188A substitution key nonpolar interactions (including CH- $\pi$  interactions; see *SI Appendix, Fig. S7*) formed between P188 and the aromatic ring of the SUs are no longer present due to the smaller alanine side chain (*SI Appendix, Fig. S2C*). However, the side chain of leucine in the A191L substitution protrudes toward the binding cavity, reducing its volume; therefore, conformational adjustments are required to accept the herbicide (*SI Appendix, Fig. S2C*). Although extra contacts may be formed in the presence of this substitution, the overall effect of the P188A and A191L substitutions is negative as witnessed by the increase in the  $K_i$  value of CE and resistance in the presence of SM, BSM, and ES. In contrast, the  $K_i$  values of the TPs are less affected by these



**Fig. 2.** CaAHAS structure. (A) Overall fold of the uninhibited CaAHAS tetramer in cartoon representation. The cofactors ThDP (white) and FAD (yellow) are drawn as ball and sticks models, and the magnesium ion is depicted as a dark cyan sphere. (B) Cartoon representation of a single subunit of CaAHAS in complex with CE. The three domains:  $\alpha$  (76–272),  $\beta$  (273–454), and  $\gamma$  (468–639) are colored in different shades of green. The two elements of the capping region, the “mobile loop” (Q576–Y591) and “C-terminal arm” (P646–Y683), which are key for trapping the herbicide in its binding pocket, are colored magenta and blue, respectively. The carbon atoms in CE are colored cyan. A green broken line represents the N267–A276 region, which is disordered in all structures. (C)  $2F_o - F_c$  electron density maps (contoured at  $\geq 2.0 \sigma$ ) for ThDP, ThAthDP, ThThDP, ThThDP/ThAthDP, and ThFATHDP/ThAthDP in the uninhibited enzyme (2.9-Å resolution), the IE (1.8-Å resolution), PC (2.4-Å resolution), MT (2.1-Å resolution), and CE (2.1-Å resolution) complexes, respectively. (D) Connolly surface showing CE (cyan) inserted into the substrate access channel thereby blocking the active site, where ThDP (white) is located (enclosed in B with broken lines). Nitrogen, blue; oxygen, red; sulfur, yellow; potassium, magenta; and chlorine, pink.

substitutions (Table 1). In the light of these results, this appears to be because these herbicides rely less on the interactions formed with P188 than on the additional contacts that may occur with the A191L substitution (*SI Appendix*, Fig. S2D).

**Fungal Growth Inhibition by AHAS-Inhibiting Herbicides.** The five classes of commercial herbicide that target AHAS were assessed for their ability to prevent the growth of five *Candida* species (*C. albicans*, *Candida parapsilosis*, *C. glabrata*, *Candida tropicalis*, and *C. krusei*), *C. neoformans*, and *S. cerevisiae* in culture (*SI Appendix*, Table S2). CE was found to display broad-spectrum antifungal activity, being highly effective in preventing the growth of *C. albicans*, *C. parapsilosis*, *C. glabrata*, and *S. cerevisiae* with MIC values as low as 0.03  $\mu\text{g/mL}$  (72.32 nM), 0.003  $\mu\text{g/mL}$  (7.23 nM), 0.005  $\mu\text{g/mL}$  (12.05 nM), and 0.008  $\mu\text{g/mL}$  (19.29 nM), respectively (Table 2), while other herbicides, including MT, are also effective at preventing growth, but not with the same level of broad-spectrum activity (*SI Appendix*, Table S2). Independent of the strain tested, some herbicides reduced growth for 24 h, whereas CE, among

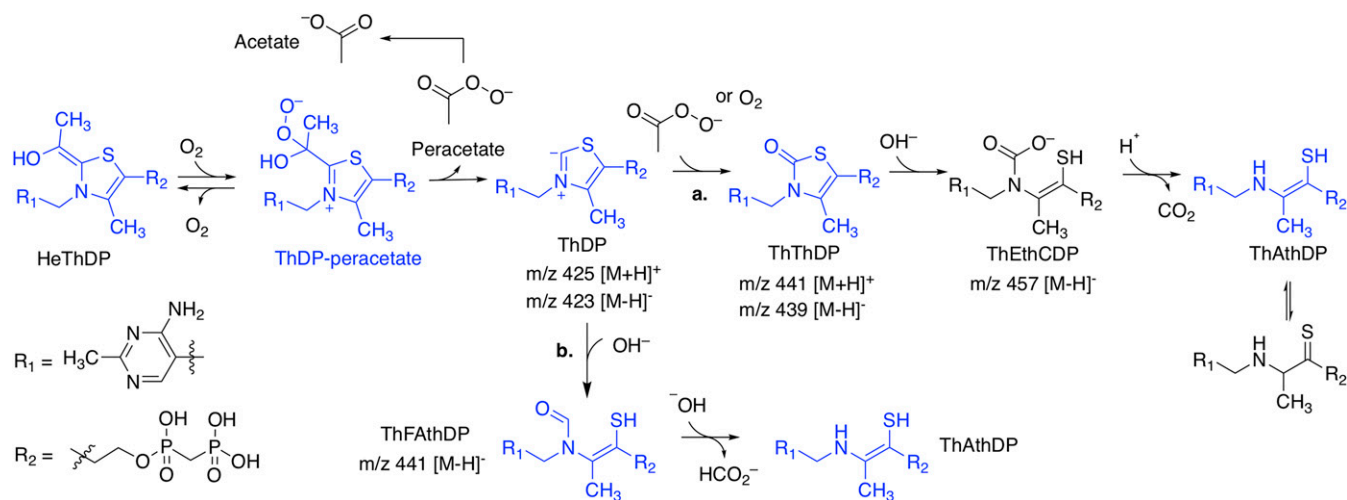
others, maintained potency for up to 72 h (Fig. 5A and *SI Appendix*, Table S2). CE has a minimum fungicidal concentration (MFC) for *C. albicans* that is four times MIC, and thus is considered fungicidal rather than fungistatic (Table 2).

**Antifungal Activity of CE Is Enhanced by Itraconazole.** To test if the current first-line antifungal compounds, fluconazole, itraconazole, or amphotericin B, can influence the activity of CE in reducing the growth of *C. albicans*, a grid experiment was performed. This showed that CE in combination with itraconazole had a synergistic effect, resulting in a fractional inhibitory concentration (FIC) value of 0.63 (Fig. 5B, Top). In addition, an additive interaction was observed when CE and amphotericin B were combined (FIC = 1). In contrast, the combination of CE with fluconazole shows an antagonistic interaction (FIC = 1.5). It is worth noting that fluconazole (a fungistatic agent) is an antagonist of other antifungal drugs, for example amphotericin B (a fungicidal) (28), an interaction usually observed when the drugs have different mechanisms of action.

A method that plants adopt to detoxify AHAS-inhibiting herbicides is to use P450 enzymes. This is achieved by adding or removing functional groups, thereby preventing binding to the enzyme (29, 30). Azoles are potent inhibitors of the fungal cytochrome P450 lanosterol 14- $\alpha$ -demethylase (CYP51A1). They also have activity toward human P450 enzymes (CYP3A4 and CYP2C9), increasing the serum concentration of several drugs (31). Therefore, we hypothesize itraconazole may inhibit CE degradation via P450s, thus maintaining the concentration of the compound in the cell for a longer time period. This implies existing fungal P450s can play a role in reducing the antifungal activity of those herbicides that are susceptible to modification. To answer this question, a synergy grid experiment was performed using PS, a TP that has a low MIC value for *C. albicans* growth of 6.41  $\mu\text{g/mL}$  after 48 h, but which loses activity after 72 h (MIC = 185.5  $\mu\text{g/mL}$ ) (Fig. 5A), in combination with fluconazole, itraconazole, or amphotericin B. The results show that only itraconazole enhances the longevity of antifungal activity of PS (synergistic effect, FIC = 0.31) (Fig. 5B, Bottom), resulting in a >15-fold reduction in MIC value for PS after 72 h. However, adding fresh PS alone after 24, 48, and 72 h to concentrations equal to those used in the microdilution tests or additive self-drug combination of PS did not produce an improvement in the MIC values. These data suggest that itraconazole could be coadministered to minimize the effect of herbicide metabolism.

#### Effect of BCAA Supplementation on the Antifungal Activity of CE.

Next, we investigated whether BCAA supplementation affects the efficacy of CE in reducing the growth of five *Candida* species and *C. neoformans* in culture media. Compared with the control [yeast nitrogen base (YNB) broth supplemented with 10 mM proline as nitrogen source], the addition of BCAAs (10 mM; >15-fold the concentration found in human blood) as sole nitrogen source produces a shift in the MICs of CE for all five *Candida* strains. *C. neoformans* is unable to grow under these conditions (15) (Fig. 5C). The effect of BCAAs on *C. albicans* growth in the presence of CE is significantly lower (2.3-fold increase of MIC) compared with that of the other *Candida* strains (>20-fold increase of MIC). The inclusion of proline along with BCAAs partially restores the antifungal activity of CE against *C. albicans*, *C. parapsilosis*, and *C. tropicalis* (Fig. 5C). In this situation, *C. neoformans* growth was viable (15) with a twofold increase in the MIC of CE. Given that serum albumin could be a potential source of BCAAs during fungal infections, we next investigated the effect of BSA (4 g/L) on the performance of CE. These results showed that only *C. glabrata* and *C. parapsilosis* can partially bypass AHAS inhibition through scavenging BCAAs from BSA (8.3- and 2.9-fold increase of MIC, respectively). Remarkably,



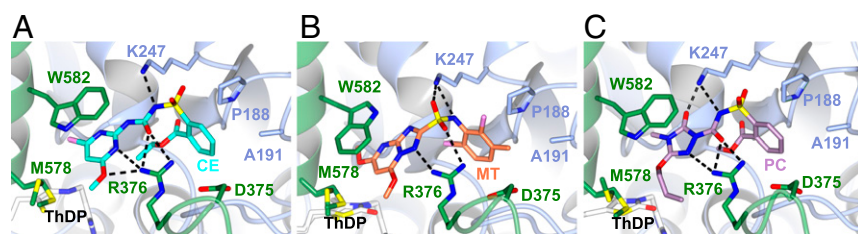
**Fig. 3.** Proposed mechanism for AHAS modification of ThDP. Our kinetic, structural, and MS data suggest that, upon herbicide binding, ThDP is primarily altered into two analogs: ThThDP or ThAthDP. Turnover conditions and molecular oxygen [which binds AHAS nearby ThDP (59)] are required for this to occur (23, 24), implying that the presence of the hydroxyethyl (He) ThDP intermediate in the active site is required for either herbicide binding and/or the generation of peracetate, a product of the oxygenase activity of AHAS (60). From these results, the following mechanism has been developed. In pathway **a**, free peracetate or molecular oxygen (61, 62) (both trapped in the active site by the herbicide) (24) react with the C2 carbene of ThDP, producing ThThDP. The thiazolone ring is then hydrolyzed by hydroxide and ring-opened, leading to intermediate thiamine ethenethiolcarbamate diphosphate (ThEthCDP), which undergoes decarboxylation of the carbamic acid group to give ThAthDP (63). Alternatively, in pathway **b**, ThAthDP is produced by a parallel mechanism that involves direct nucleophilic ring opening of the thiazolium ring of ThDP, producing formamidoethenethiol diphosphate (ThFATHDP) (62, 64). Hydroxide then induces cleavage of the N-CHO bond to give ThAthDP (65). The ThDP related molecules depicted in blue have been observed in the crystal structures of AHAS in complex with the herbicide, except for intact ThDP, which is observed in the crystal structure of uninhibited CaAHAS.

the MIC of CE against *C. neoformans* was reduced a further 5.5-fold compared with the control under this assay condition (Fig. 5C).

To identify the requirements of these fungal pathogens to circumvent AHAS inhibition in the presence of BCAAs, we next assessed the growth of *C. albicans* in YNB broth with leucine, valine, isoleucine, proline, and/or ammonium sulfate as the only nitrogen source or in combinations, and in the presence of different concentrations of CE (Fig. 5D). These experiments showed that the addition of 10 mM valine, leucine, or isoleucine as the sole nitrogen source does not affect the efficacy of CE in preventing *C. albicans* growth. Indeed, the MICs of CE determined under these conditions are lower compared with the MIC determined using YNB media supplemented with 10 mM ammonium sulfate or proline. However, the combination of valine and isoleucine suppresses significantly the antifungal activity of CE (MIC = 5.6  $\mu\text{g}/\text{mL}$ ), while this effect is nullified by addition of leucine, proline, or ammonium sulfate (Fig. 5D). Amino acid transport is highly regulated by the nitrogen catabolite repression or the Ssy1p-Ptr3p-Ssy5p systems, the intracellular and extracellular amino acid ratio, and the presence of nitrogenated molecules in the environment (32, 33). Ammonium diminishes BCAA transport via Gap1 (Gap2 in *C. albicans*) and Bap2 repression (34, 35), a mechanism which explains why ammonium sulfate restores the antifungal activity of CE in the presence of BCAAs. In agreement with this result, 100  $\mu\text{g}/\text{mL}$  eugenol, an inhibitor of BCAA permeases (36), synergizes with CE (FIC 0.38; 48-h MIC<sub>CE</sub> = 0.125  $\mu\text{g}/\text{mL}$ ) against *C. albicans* when BCAAs are present in the growth media (SI Appendix, Fig. S10). However, while it has been shown that proline derepresses BCAA transport (34), here we observed that it also diminishes the ability of *C. albicans* to bypass AHAS inhibition by taking up BCAAs from the media. In addition, we observed when leucine, valine, and isoleucine are added in combination in the media, growth is reduced (Fig. 5E). This could be due to reciprocal competition as the three BCAAs try to use the same method of transport into the cell (37) (Fig. 5E). It has been shown that absence of valine is highly detrimental for *C. albicans* AHAS (*ilv2Δ*) mutant growth (14). Here, we show valine

increases the growth of *C. albicans* and that both leucine and isoleucine compete with valine, as seen by a 15% reduction in fungal growth when these are in combination (Leu-Val or Ile-Val). Isoleucine also inhibits growth in presence of leucine and the combination of the three BCAAs yields 35% less growth compared with that observed when valine is the nitrogen source (Fig. 5E). These results suggest that, even though *C. albicans* is able to scavenge BCAAs from the environment and, to some extent, bypass AHAS inhibition by herbicide to support growth, the intricate interactions between substrates and amino acids limit the absorption of BCAAs and therefore add further support for the concept of using AHAS-inhibiting herbicides to treat human fungal infections.

**Antifungal Activity of CE in Vivo.** We next assessed if CE could be an effective antifungal agent in mice. First, we performed a toxicity study where CE was injected i.p. twice per day and showed that by this procedure over a 7-d period doses as high as 500  $\text{mg}\cdot\text{kg}^{-1}\cdot\text{d}^{-1}$  were tolerated by the mice, with no adverse effects. To assess the efficacy of CE we infected groups of 10 mice via tail vein with *C. albicans* (strain SC5314; 24-h MIC of CE = 0.05  $\mu\text{g}/\text{mL}$ ) and tested their ability to survive given different doses of CE (100 and 500  $\text{mg}\cdot\text{kg}^{-1}\cdot\text{d}^{-1}$ ) delivered in two IP injections, each including half of the daily dose of CE. PBS and fluconazole (200  $\text{mg}/\text{kg}/\text{d}$ ) were included as negative and positive controls and delivered in the same way as CE. Weight loss is a gold-standard symptom indicative of disseminated fungal infection (38) and we used this reference to assess the severity of the infection. With this premise, only one mouse in the PBS group did not show signs consistent with the disease (Fig. 6A) and therefore was not considered in the final statistical analysis, while all mice receiving CE or fluconazole showed signs consistent with the disease (i.e., weight loss) (Fig. 6A). Compared with the control group receiving PBS, the results show that administration of CE prolongs the life of the infected mice in a concentration-dependent manner (Fig. 6A). Organ fungal burden analysis of kidneys, spleen, liver, lungs, and brain showed that all mice that survived until the end of the experiment, except one individual receiving fluconazole, had cleared the infection



**Fig. 4.** SU, TP, and SCT binding sites in CaAHAS. (A) CE, (B) MT, and (C) PC. The herbicides and key residues for herbicide binding are depicted in stick models, whereas the polypeptide is shown in ribbon display with each subunit colored differently. Hydrogen bonds formed between the herbicides and the polypeptide are presented as dashed lines.

(Fig. 6B). In addition, CE significantly decreased fungal load in the spleen, liver, lungs, and brain compared with those mice without receiving antifungal treatment, suggesting that the herbicide is able to reach its target via an adequate distribution of the herbicide in these organs. Importantly, CE is effective in clearing infection from the lungs and liver, even at low concentrations (Fig. 6B). Previous studies have shown that the SUs are readily absorbed following oral administration (17, 39); therefore, here we additionally show that i.p. injection is also a viable route for drug administration. The SUs, including CE, and most AHAS-inhibiting herbicides show a pH-dependent solubility in aqueous solutions, with these compounds becoming insoluble at low pH. We therefore hypothesize the weaker effect of CE in reducing the fungal burden in the kidneys could be due to precipitation of the compound in urine (40).

## Discussion

Since the discovery that AHAS is essential for virulence and survival of *C. albicans* and *C. neoformans* cells in vitro and in vivo (14, 15), this enzyme has been proposed as a potential target for developing antifungal compounds. Preliminary studies have shown that inhibitors targeting AHAS have antifungal activity against *C. albicans*, *C. neoformans*, and *A. fumigatus* in culture (18, 19). Here, we have performed a comprehensive study to identify drug lead candidates with potent antifungal activity within the five families of AHAS-inhibiting herbicides. Our kinetic assays concluded that CE and several other AHAS inhibitors of the SU and TP families show potent inhibition of CaAHAS and CnAHAS that also correlates with strong antifungal activity. The ability of the herbicides to induce accumulative inhibition via ThDP alteration and FAD oxidation (23, 24) is an important determinant in how effective these compounds are in biological assays, so should be a key factor in consideration of the design of AHAS inhibitors to act as antifungal drugs. Importantly, CE stands out over all commercial herbicides tested in this study because it possesses a broad spectrum of antifungal activity against five different *Candida* species and *C. neoformans* and is fungicidal against *C. albicans* in vitro. Our cell-based studies suggest several fungal pathogens have the ability to detoxify some AHAS-inhibiting herbicides via P450s. It is worth noting that CE and the other commercial AHAS inhibitors assayed here were developed to possess crop vs. weed selectivity via herbicide detoxification. *O*-dealkylation of the methoxy substituents at the heterocyclic ring via P450s is an example of how plants detoxify AHAS inhibitors by affecting their performance (16), suggesting a similar mechanism could occur in fungi. CE has a chlorine atom attached to the heterocyclic ring, a unique feature among the commercial herbicides tested (SI Appendix, Fig. S1). Accordingly, CE is effective in reducing the growth of a number of fungal pathogens over a period of 72 h, indicative that this herbicide is less affected by detoxification. In this context, we have shown itraconazole enhances the antifungal activity of AHAS inhibitors, suggesting there is potential to exploit/develop specific fungal P450 inhibitors to further maximize the performance of CE and other herbicides that target AHAS.

The broad-spectrum antifungal activity of CE is a result of the high conservation of the AHAS herbicide-binding site across fungi (SI Appendix, Fig. S2). Of those residues forming direct

contacts with the herbicides, only four have been changed during evolution and are located at the entrance of the binding cavity (SI Appendix, Fig. S2B). The A653G substitution observed in ScAHAS and *C. parapsilosis* AHAS has little effect, since ScAHAS (41) and *S. cerevisiae* and *C. parapsilosis* cells are highly susceptible to AHAS inhibitors. However, *C. neoformans* is resistant to the SU herbicide, SM (15). Modeling suggests this is in part due to the P188A and A191L substitutions in CnAHAS (SI Appendix, Fig. S2 C and D). Based on these findings it was advocated AHAS inhibitors might not be suitable for treating infections caused by this pathogen. While we have confirmed CnAHAS exhibits a natural insensitivity to several AHAS-inhibiting herbicides of the SU (including SM, BSM, and ES), SCT, and IMI families, we also observed that CE (a SU) and MT and CSM (TPs) inhibit this enzyme at the low-nanomolar level. Models of the P188A and A191L substitutions in CaAHAS and our kinetic data show that CE is effective in inhibiting CnAHAS because it possesses a longer ethyl ester substituent at the aromatic ring and this contributes to anchoring the herbicide in its binding cavity even in presence of these two unfavorable substitutions (SI Appendix, Fig. S2C). The TPs, including MT and CSM, are potent inhibitors of CnAHAS because this family of herbicides form fewer contacts with P188, their linker has more flexibility (SI Appendix, Fig. S2D), and these locate the aromatic ring deeper in the herbicide-binding site compared with the SUs (SI Appendix, Fig. S9). For future drug design efforts, a focus should be placed on developing inhibitors that bind only to the highly conserved residues in the AHAS binding site. This should reduce the propensity for resistance to develop.

Previous studies point out the possibility that fungal pathogens might scavenge BCAA from the hosts' serum and bypass AHAS inhibition by herbicides (15, 19). Here, we show the antifungal activity of CE against *C. albicans*, *C. parapsilosis*, *C. tropicalis*, and *C. neoformans* is not affected by BCAA supplementation, which is a product of the complex regulation of BCAA absorption by BCAAs themselves and by other molecules that could serve as nitrogen source (e.g., ammonium ions or proline).

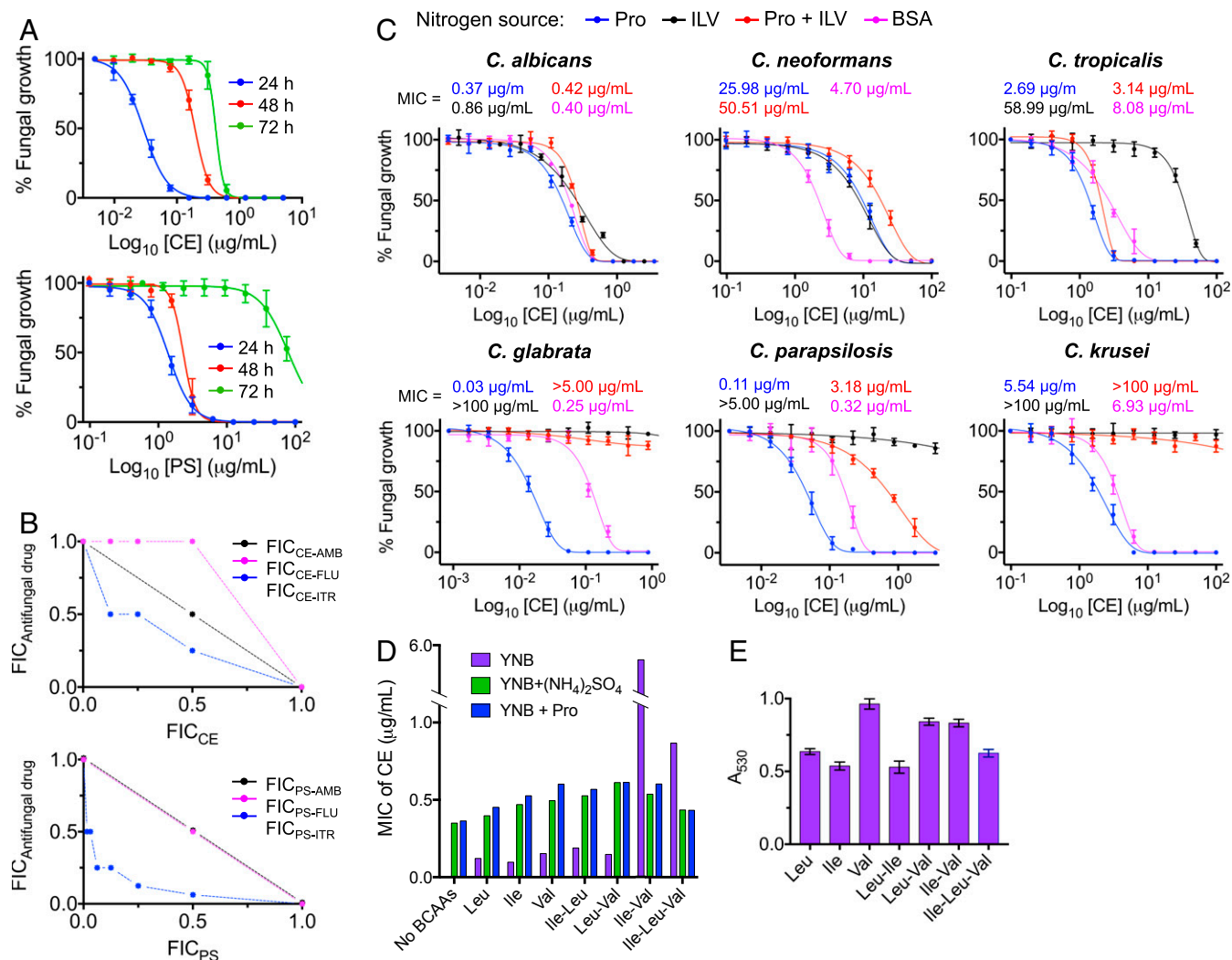
**Table 2.** MIC and MFC of CE for fungal pathogen growth in cell culture

Fungal species	MIC <sub>50</sub>		MIC*		MFC/MIC
	μg/mL	nM	μg/mL	nM	
Ca	0.030	72.32	0.070	168.7	4
Cg	0.005	12.05	0.010	24.1	>8
Cp	0.003	7.23	0.004	9.6	8
Ct	0.550	1.33 × 10 <sup>3</sup>	1.450	3.5 × 10 <sup>3</sup>	8
Ck	0.430	1.04 × 10 <sup>3</sup>	1.960	4.7 × 10 <sup>3</sup>	>8
Cn	2.750	6.63 × 10 <sup>3</sup>	7.980	19.2 × 10 <sup>3</sup>	>8
Sc†	0.008	19.29	0.02	48.2	2–4

MIC<sub>50</sub> and MIC values determined after 24-h culture.

\*The MIC values for fluconazole were 0.31 μg/mL (*C. albicans*, Ca), >100 μg/mL (*C. glabrata*, Cg), 4.66 μg/mL (*C. parapsilosis*, Cp), 6.22 μg/mL (*C. tropicalis*, Ct), 25.81 μg/mL (*C. krusei*, Ck), 15.34 μg/mL (*C. neoformans*, Cn), and 17.03 μg/mL (*S. cerevisiae*, Sc).

†MIC<sub>50</sub> and MIC values determined after 48-h culture.



**Fig. 5.** Fungal growth inhibition by herbicide. (A) Percentage of *C. albicans* cell growth vs.  $\log_{10}$  [CE] or [PS] after 24, 48, and 72 h. The solid lines represent the best fit to the data using the modified Gompertz function (57). (B) Isobolograms showing the interactions between CE (*Top*) or PS (*Bottom*) and three front-line antifungal drugs against *C. albicans* growth. The FIC<sub>CE</sub> are defined as the MIC of CE in combination with an antifungal drug (itraconazole, fluconazole, or amphotericin B) divided by the MIC of CE alone. A straight line represents an additive interaction (CE–amphotericin B combination), a concave curve indicates synergy (CE–itraconazole combination), and a convex curve shows antagonism (CE–fluconazole interaction). The FIC values for PS and each antifungal drug were determined in the same way. Amb, amphotericin B; Flu, fluconazole; Itr, itraconazole. (C) Fungal growth inhibition by CE in presence of different nitrogen sources: (i) 10 mM proline (blue lines); (ii) 2.5 mM proline, 2.5 mM leucine, 2.5 mM valine, and 2.5 mM isoleucine (black lines); (iii) 3.3 mM leucine, 3.3 mM valine, and 3.3 mM isoleucine (magenta lines); and (iv) 4 g/L BSA (magenta lines). The MIC values were determined at 48-h incubation using the modified Gompertz function. Note that the substitution of ammonium sulfate for proline, as the nitrogen source, does not produce a significant change in the MIC values reported in *SI Appendix, Table S2*. (D) Inhibition of *C. albicans* growth by CE in presence different combinations of BCAAs. YNB media was supplemented with 5 mM valine, 5 mM leucine, and/or 5 mM isoleucine in the presence or absence of 10 mM ammonium sulfate or 10 mM proline. (E) *C. albicans* growth in the presence of different combinations of BCAAs (5 mM each) and in the absence of herbicide. Error bars represent the SD of the mean (SEM) ( $n = 3$ ).

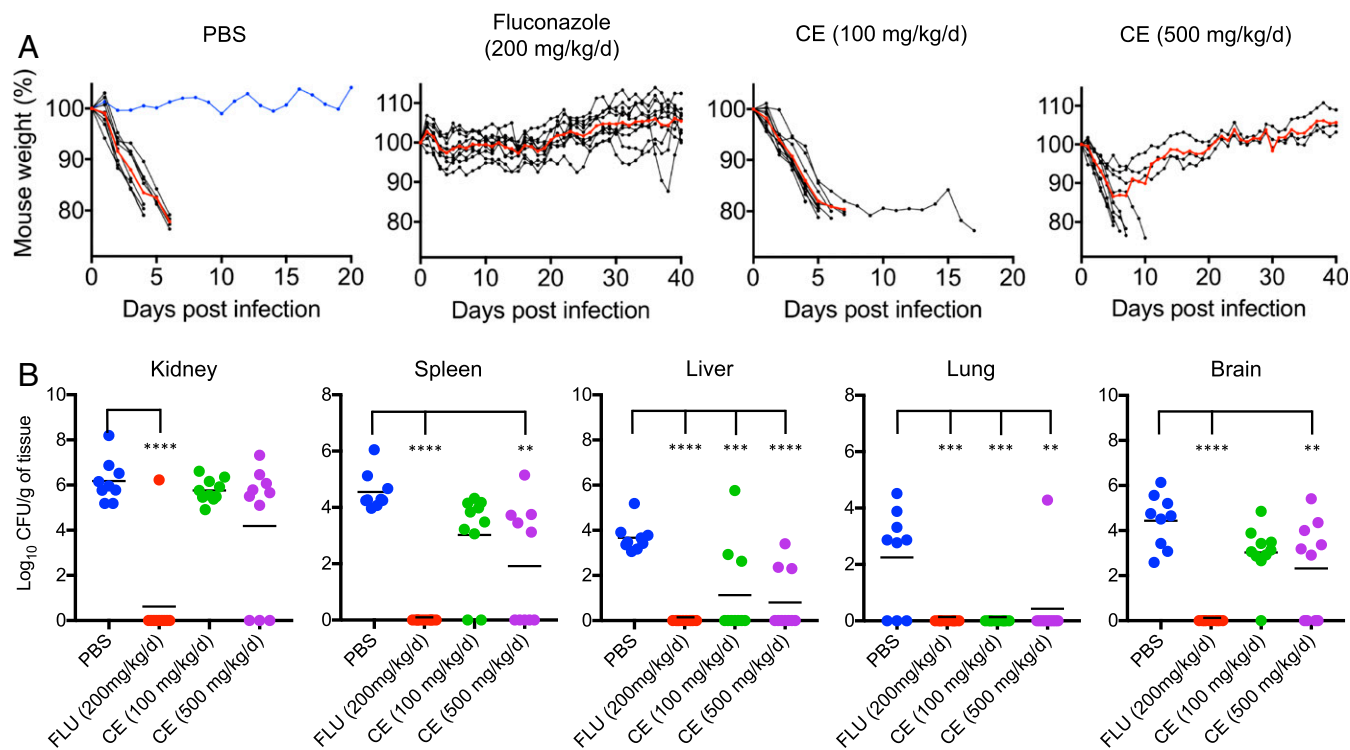
Further supporting our observation, there is evidence during protein synthesis inhibition, which is the latter effect of blocking BCAA biosynthesis by herbicide, endogenous amino acids accumulate inhibiting amino acid uptake below detection limits (42). Conclusively, CE is effective in reducing fungal burden in the spleen, liver, lungs, and brain of mice systemically infected with *C. albicans*, confirming that the hosts' BCAAs absorbed by this pathogen during infection cannot completely bypass AHAS inhibition by CE. Therefore, we identify the SU herbicides as having great potential to be developed as a novel class of antifungal drugs to treat human fungal pathogenic infections.

## Conclusions

Here, AHAS-inhibiting commercial herbicides, developed to efficiently kill weeds over crops, were assessed as antifungal

agents. This is possible by taking advantage of the fact that the sequences of AHAS are highly conserved across both kingdoms and the activity of this enzyme is essential for the survival of plants and microorganisms. The absence of this enzyme in mammals and the low toxicity of AHAS inhibitors in humans (17) make them an appealing class of compound to be optimized into therapeutic agents to treat infections due to microbial pathogens.

The kinetics of inhibition of CaAHAS and CrAHAS by representatives of the five families of commercial herbicides were determined and these served to identify several potent accumulative inhibitors of these enzymes. CaAHAS inhibition by these herbicides is different from that observed when they bind to ScAHAS. In ScAHAS much of the polypeptide has to fold around these compounds to optimize binding. However, in CaAHAS no major conformational changes occur. This is a key



**Fig. 6.** Survival of BALB/c mice infected i.v. with *C. albicans*. (A) Percentage of mouse weight postinfection. Mice ( $n = 10$  per group) received PBS, fluconazole ( $200 \text{ mg}\cdot\text{kg}^{-1}\cdot\text{d}^{-1}$ ), CE ( $100 \text{ mg}\cdot\text{kg}^{-1}\cdot\text{d}^{-1}$ ), or CE ( $500 \text{ mg}\cdot\text{kg}^{-1}\cdot\text{d}^{-1}$ ) via i.p. injection over a 7-d period after i.v. infection with *C. albicans* (Methods). The weight of the mice was recorded every day for the duration of the experiment (40 d), and mice were killed by  $\text{CO}_2$  asphyxiation when they showed  $\geq 20\%$  weight loss or signs consistent with disseminated candidiasis. Each mouse showed weight loss for at least four consecutive days postinfection. Black lines represent each individual within groups, red lines the mean of each group, and a blue line represents an outlier in the PBS group that did not show signs of infection (i.e., weight loss). (B) Organ burden analysis calculated in colony forming units (CFU) per gram of tissue. One-way ANOVA test was performed to determine statistical significance (kidney:  $F = 17.82$ ,  $P < 0.0001$ ; spleen:  $F = 18.26$ ,  $P < 0.0001$ ; liver:  $F = 14.87$ ,  $P < 0.0001$ ; lung:  $F = 8.93$ ,  $P = 0.0002$ ; and brain:  $F = 17.7$ ,  $P < 0.0001$ ; degrees of freedom = 3 for treatments and 35 for residuals), followed by Tukey test ( $\alpha = 0.05$ ,  $**P < 0.01$ ;  $***P < 0.001$ ;  $****P < 0.0001$ ).

finding because it allows the rational structure-based design of inhibitors based on the native structure of CaAHAS. Previously, we have demonstrated AHAS inhibitors are able to take full advantage of the high reactivity of ThDP and to induce reactions that lead to the ultimate inactivation of this cofactor (23, 24). Here, the crystallographic data and MS allowed us to describe the mechanism by which the herbicides convert ThDP into ThThDP and ThAthDP.

Our cell-based assays provide unique insights into the pathogens' response to the potent antifungal activity of several commercial herbicides that target AHAS and the potential use of P450 inhibitors to further enhance antifungal activity and to diminish the prospects of the development of resistance. More importantly, CE possesses antifungal activity in mice infected with *C. albicans*, which illustrates CE is a drug lead suitable for further development and that AHAS inhibitors have great promise to treat human pathogenic fungal infections.

## Methods

**Cloning of CnAHAS and CaAHAS.** The ORF for the catalytic subunit of the CnAHAS gene was amplified from *C. neoformans* cDNA. The first 101 aa corresponding to the mitochondrial transit peptide were deleted using the forward primer 5'-AAAGGATCCATGCCACTAATATGCC-3', which contains a BamHI site at the N terminus to facilitate its cloning into pET30A(+) vector. The reverse primer was 5'-TCTGCGGCGCTTACTCAGAGCCGTTCCG-3' and contains a NotI restriction site after the stop codon. This construct therefore encodes the first 50 residues of the vector, including a hexahistidine tag, followed by the CnAHAS polypeptide from M102 to E718. Cloning of the CaAHAS catalytic subunit into pET30A(+) vector was reported previously (18). The CaAHAS and CnAHAS constructs were then transformed into

*Escherichia coli* strain Rosetta BL21(DE3) and BL21(DE3)pLysS, respectively, for protein expression.

**Protein Expression and Purification.** Expression and purification of the catalytic subunits of CaAHAS and CnAHAS was carried out as reported previously for AtAHAS (23). The enzymes were stored in aliquots of 30  $\mu\text{L}$  at  $-70^\circ\text{C}$ . Protein concentration was measured using the Direct Detect system (Merck Millipore).

**AHAS Assay.** For all assays, unless specifically stated, the standard reaction buffer contained 200 mM potassium phosphate (pH 7.2), 100 mM sodium pyruvate, 10 mM  $\text{MgCl}_2$ , 1 mM ThDP, and 10  $\mu\text{M}$  FAD and assays were conducted at  $30^\circ\text{C}$ . Measurement of  $K_i$  values was carried out in triplicate using a colorimetric method (43) and partial anaerobic conditions as described previously (23). These values were then calculated by nonlinear regression using Eq. 1 [weak to medium binding (44)] or Eq. 2 [tight binding (18)]:

$$v_i = v_o / (1 + [I]/K_i) \quad [1]$$

$$v_i^2[E]_o + v_o v_i ([I]_o - [E]_o + K_i) - v_o^2 K_i = 0. \quad [2]$$

For accumulative inhibition assays, 1  $\mu\text{M}$  enzyme was assayed in triplicates using a continuous method (45). The reaction buffer for these assays contained 50 mM pyruvate. The inhibitors were added 15 min after the enzyme so that inhibition could be measured when maximum activity of the enzyme was attained. The reaction was then monitored for a further 35 min. The data were fitted using Eq. 3 (20) for the calculation of  $k_{iapp}$  and  $k_3$  involved in accumulative inhibition:

$$[P] = V_{\max} \left( F \left( \frac{k_3}{k_{iapp} + Fk_3} \right) t + \left( \frac{k_{iapp}}{k_{iapp} + Fk_3} \right) \frac{1}{k_{iapp}/F + k_3} \right) \times \left( 1 - \exp \left[ - \left( \frac{k_{iapp}}{F} + k_3 \right) t \right] \right). \quad [3]$$



For BS, CS, DS, FS, FT, IM, PC, PS, PTB, PYS, SM, and TCM, their relatively high  $K_i$  values were taken into account for the estimation of the value  $F$  (free enzyme/enzyme-inhibitor complex ratio). The effective concentration of enzyme-inhibitor was calculated using Eq. 4 (inhibition constant formula), the known initial concentrations of enzyme and inhibitor, and the  $K_i$  value of Table 1:

$$K_i = ([E] \cdot [I]) / [EI]. \quad [4]$$

**Protein Crystallization and Structure Determination.** Crystallization trials were performed using the hanging-drop vapor-diffusion method as described previously (23). The protein solution included freshly thawed protein (30 mg/mL), 5 mM  $MgCl_2$ , 1 mM ThDP, 1 mM FAD, 5 mM DTT, 3 mM sodium pyruvate, and, when appropriate, 1 mM herbicide. The hanging drops consisted of protein and well solution in a 2:1 ratio. For uninhibited CaAHAS and the CaAHAS–MT complex, the well solution contained 1 M Na/K tartrate, 0.1 M 2-(*N*-cyclohexylamino)ethanesulfonic acid, and 0.2 M  $Na_2SO_4$ . The pH of the solution ranged from 9.4 to 9.8. Crystals for the remaining CaAHAS-herbicide complexes were obtained in similar way, with the only difference being the well solution contained  $(NH_4)_2SO_4$  instead of  $Na_2SO_4$ . The crystallization experiments were kept at 290 K and protected from light until the crystals reached their maximum size. For cryocooling the crystals were transferred to a drop containing well solution, 35% vol/vol ethylene glycol, cofactors, and inhibitor (when appropriate), mounted in cryoloops and frozen in liquid nitrogen.

The datasets of the uninhibited CaAHAS and the CaAHAS–herbicide complexes were collected at the Australian Synchrotron. The datasets were indexed, integrated, and scaled using XDS (46) and Aimless (47). The crystal structure of uninhibited CaAHAS was solved by molecular replacement with PHASER (48) using ScAHAS in complex with chlorsulfuron (Protein Data Bank ID code 1T9B) as the starting model. Model building was carried out using Coot 0.8.1 (49) and refinement was with PHENIX 1.9-1692 (50). The crystal structures of CaAHAS in complex with herbicides were then solved by the same method using the uninhibited CaAHAS as the starting model. Figures were generated with CCP4mg (51).

**Protein Sequence Alignment Protein–Ligand Interaction Analysis.** The amino acid sequences for the AHAS catalytic subunits of *C. albicans* (XP\_721571.1), *C. glabrata* (XP\_448375.1), *C. parapsilosis* (CCE44821.1), *C. tropicalis* (XP\_002548114.1), *C. krusei* (KGGK39551.1), *C. neoformans* (AAR29084.1), *A. fumigatus* (EAL92550.1), and *S. cerevisiae* (NP\_013826.1) were obtained from GenBank and sequence alignment was performed with Clustal Omega (52). Interactions with each herbicide in CaAHAS were analyzed using Arpeggio (53).

**CnAHAS Herbicide-Binding Site Model.** The P188A and A191L substitutions present in CnAHAS were modeled in the 3D coordinates of uninhibited CaAHAS and the CaAHAS-CE/MT complexes using Coot 0.8.1 (49) and structure optimization was carried out with Yasara (54).

**MS.** Samples for MS analysis of ThDP were prepared as described previously (23). One microliter of each sample was then injected onto an Agilent InfinityLab Poroshell 120 EC-C18 column (2.1 × 50 mm). Liquid chromatography gradient was 0–98% B over 7 min, run at 100  $\mu$ L/min, with the column oven set to 45 °C. Eluted samples were sprayed using a heated electrospray ionization source and analyzed on an Orbitrap Elite mass spectrometer. Source parameters included capillary temperature = 320 °C, source heater temperature = 200 °C, sheath gas = 25 L/min, auxiliary gas = 8 L/min, source voltage = 4.5 kV, and s-lens = 60%. Samples were analyzed with the Orbitrap analyzer set to 30,000 resolution.

**Cell-Based Studies.** Antifungal susceptibility testing was performed using the broth microdilution method according to a standard protocol (55). All experiments were performed in 96-well plates using YNB broth without amino acids and ammonium sulfate (BD Difco) supplemented with 2% glucose, 100 mM  $(NH_4)_2SO_4$ , and 50 mM Hepes. The pH of the media was adjusted to 7.2 to ensure inhibitor solubility and stability (56). The herbicides were dissolved in DMSO and then serially diluted in 96-well plates using YNB. The herbicide solutions were used fresh and kept at –20 °C for no more than 1 wk. The latter was particularly important to guarantee accurate MIC measurements, since some of the herbicide tested had degraded when stored in DMSO for prolonged periods (>1 wk). The well-established antifungal drugs fluconazole and itraconazole were used as controls. All studies were conducted using *C. albicans* ATCC 90028, *C. glabrata* ATCC 90030, *C. parapsilosis* ATCC 22019, *C. tropicalis* ATCC 750, *C. krusei* ATCC 6258, *C. neoformans* ATCC 90113, and *S. cerevisiae* Sigma 1278b. Growth was

determined by measuring the OD<sub>530</sub> every 24 h using a Molecular Devices Spectramax 250 microplate reader (Marshall Scientific). The MIC<sub>50</sub> and MIC values were determined using the modified Gompertz function (57). All experiments were conducted in triplicate, with each performed on a different day. The MFC was determined as described previously (58).

The FIC index of CE and PS in combination with fluconazole, itraconazole, or amphotericin B against *C. albicans* growth was determined using a synergy grid experiment in a microdilution format as described above. The FIC values for CE (FIC<sub>CE</sub>) were defined as the MIC of CE in combination with an antifungal drug (itraconazole, fluconazole, or amphotericin B) divided by the MIC of CE alone. The FIC values for PS and each antifungal drug were determined in the same way. The FIC index for each drug combination was calculated as FIC<sub>herbicide</sub> + FIC<sub>antifungal drug</sub>. Synergism is defined as FIC index values <1, antagonism by FIC index values >1, and additivity by FIC index values = 1. All experiments were conducted in triplicate.

The effect of BCCAs and BSA on the antifungal activity of CE was measured in triplicate using broth microdilution in a 96-well plate format as described above, with the only difference being  $(NH_4)_2SO_4$  was replaced by (i) 10 mM proline; (ii) 2.5 mM proline in addition to different combinations of BCAAs: 2.5 mM leucine, 2.5 mM valine, and/or 2.5 mM isoleucine; (iii) 3.3 mM of each BCAA as sole nitrogen source; or (iv) 4 g/L BSA as sole nitrogen source. Inhibition of *C. albicans* growth by CE in the presence different combinations of BCAAs was measured using YNB broth supplemented with 5 mM valine, 5 mM leucine, and/or 5 mM isoleucine in the presence or absence of 10 mM ammonium sulfate or 10 mM proline.

**Mouse Studies.** A mouse model study of disseminated *C. albicans* infection was conducted following the requirements of animal welfare in accordance with the Australian Code for the Care and Use of Animals for Scientific Purposes by the National Health and Medical Research Council, and after approval by the Molecular Biosciences Animal Ethics Committee (AEC) of The University of Queensland (AEC approval no. SCMB/175/17).

For toxicity assays of CE in mice, 25 6-wk-old BALB/c mice received 0.1-mL PBS injections via the tail vein using a 1-mL tuberculin syringe and a 27-gauge, 0.5-inch needle. Subsequently, five mice received one of two different doses of CE (100 or 500 mg·kg<sup>-1</sup>·d<sup>-1</sup>) delivered by two i.p. injections (morning and evening), each containing half of the daily dose dissolved in 100  $\mu$ L PBS (adjusted to pH 8). Fluconazole (200 mg/kg/d) and PBS alone were used as controls and delivered in the same way as CE. All injections were applied using a 1-mL tuberculin syringe and a 27-gauge, 0.5-inch needle. Weight and aspect of the mice were examined on day 1 and every day after for 2 wk, at which point all mice were killed via CO<sub>2</sub> inhalation. Kidneys, spleen, liver, lungs, and brain were collected to record organ weight and aspect. Mice were provided with food and water for ad libitum intake and with cardboard as environmental enrichment.

For murine infection assays, 40 6-wk-old female BALB/c mice were infected by tail-vein injection of 0.1 mL PBS containing 10<sup>6</sup> viable cells (38) using a 1-mL tuberculin syringe and a 27-gauge, 0.5-inch needle. Mice were separated in groups of 10 and treated during a 7-d period, beginning the same day of infection with two different doses of CE (100 and 500 mg·kg<sup>-1</sup>·d<sup>-1</sup>), fluconazole (200 mg·kg<sup>-1</sup>·d<sup>-1</sup>), or PBS delivered in 100- $\mu$ L i.p. injections twice per day (morning and evening) as described above. A maximum of five mice were housed per individually ventilated cage and provided with food and water for ad libitum intake and cardboard. The weight and aspect of the mice were monitored daily during the duration of the experiment (40 d). Mice were killed via CO<sub>2</sub> asphyxiation once they declined to  $\geq$ 80% of preinfection body weight or when symptoms consistent with infection were obvious. The date of death was recorded as the following day. Kidneys, spleen, liver, lungs, and brain were collected and homogenized in 1 mL PBS using a TissueLyser II (Qiagen) and plated on YPD agar containing 25  $\mu$ g/mL chloramphenicol, 50  $\mu$ g/mL kanamycin, and 100  $\mu$ g/mL ampicillin in 1:10 serial dilutions. The colony-forming units per gram of tissue were determined after 24 h incubation at 30 °C. One-way ANOVA was performed to determine statistical significance within organ fungal burden datasets, followed by the Tukey test.

The inoculum for mice infection was prepared from fresh cultures of *C. albicans* strain SC5314 plated on YPD agar from a glycerol stock and incubated at 30 °C for 48 h. A single colony was transferred to 10 mL YPD broth and further incubated at 30 °C with shaking (150 rpm) for 20 h. Cells were harvested by centrifugation (800 × g) for 1 min at room temperature and washed twice with sterile PBS and the cell density was adjusted to 1 × 10<sup>7</sup> cells per mL using a hemocytometer. The cell suspension was kept on ice and used immediately. Cell concentration was confirmed by plating 1:10 serial dilutions on YPD agar and incubated for 24–48 h at 30 °C.

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