

Discovery and Structure–Activity Relationships of Pyrrolone Antimalarials

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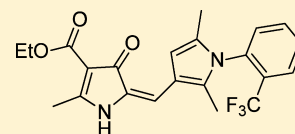
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S Supporting Information

ABSTRACT: In the pursuit of new antimalarial leads, a phenotypic screening of various commercially sourced compound libraries was undertaken by the World Health Organisation Programme for Research and Training in Tropical Diseases (WHO-TDR). We report here the detailed characterization of one of the hits from this process, TDR32750 (**8a**), which showed potent activity against *Plasmodium falciparum* K1 (EC₅₀ ~ 9 nM), good selectivity (>2000-fold) compared to a mammalian cell line (L6), and significant activity against a rodent model of malaria when administered intraperitoneally. Structure–activity relationship studies have indicated ways in which the molecule could be optimized. This compound represents an exciting start point for a drug discovery program for the development of a novel antimalarial.



INTRODUCTION

In recent years, the emergence of drug-resistant pathogens has led to treatment failures for many infectious diseases, such as malaria. Resistance to current antimalarials is a major problem, e.g., chloroquine, sulfadoxine-pyrimethamine, and in some areas, mefloquine.¹ To reduce the spread of resistance, the World Health Organisation (WHO) recommends treatment for malaria to be artemisinin combination therapy; although worryingly there are now reports of increased parasite clearance times with the artemisinins^{2–7} which may herald resistance. Thus, there is an urgent need for novel classes of compounds that are effective against these re-emerging infections. With the elimination of malaria now considered as a goal, it would be useful for scientists and policy makers to have an expanded arsenal of antimalarials that could help make this goal a reality.^{8,9}

As part of a program for the discovery of new starting points for drug discovery programmes, the World Health Organisation Programme for Research and Training in Tropical Diseases (WHO-TDR) phenotypically screened various compound libraries. One of these was a 5000 strong structurally diverse compound collection from ChemDiv, which was screened against *Plasmodium falciparum*, the causative organism for the most pathogenic form of malaria. This led to the identification of pyrrolone (**8a**) shown in Figure 1. This compound had potent activity against *P. falciparum* K1, with an EC₅₀ ~ 9 nM, and was

efficacious in a *P. berghei* mouse model when given intraperitoneally (ip), but was relatively inactive orally.

The screening of several hundred commercially available compounds related to **8a** containing the pyrrolone motif gave some indication of structure–activity relationships (SAR) (Figure 1). Here we report the further evaluation of **8a** as a novel antimalarial lead, with further systematic SAR studies, in parallel with studies to assess drug metabolism and pharmacokinetics (DMPK), with the initial aim of achieving oral efficacy in the *P. berghei* mouse model, to establish the potential for further development of the pyrrolones as antimalarials.

RESULTS AND DISCUSSION

Chemistry. An efficient synthesis of **8a** and congeners was developed (Scheme 1), through acylation of ethyl-3-amino-crotonate (**1**) with chloroacetyl chloride¹⁰ and subsequent cyclization to the pyrrolone **3**,¹⁰ which was not stored, but used immediately, due to its relatively poor stability. The condensation of **3** with 3-formyl pyrroles (**7a–x**) in the presence of potassium hydrogen sulfate^{11,12} generated the (*E*)-isomer predominantly.¹³ This three-step sequence could be carried out in yields of up to ~60%, and only required chromatography at step

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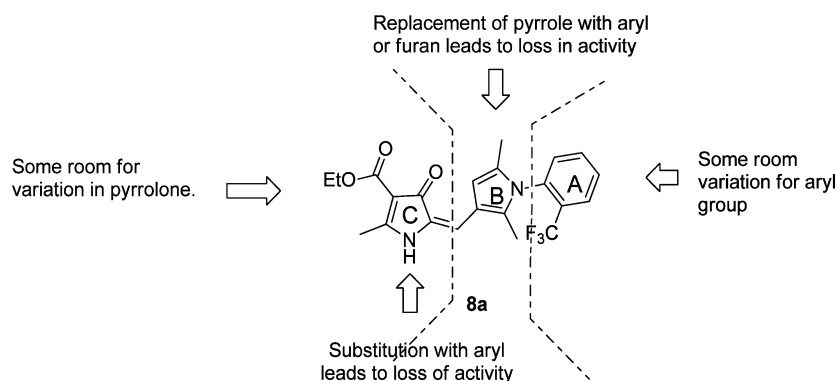


Figure 1. Initial SAR of lead 8a.

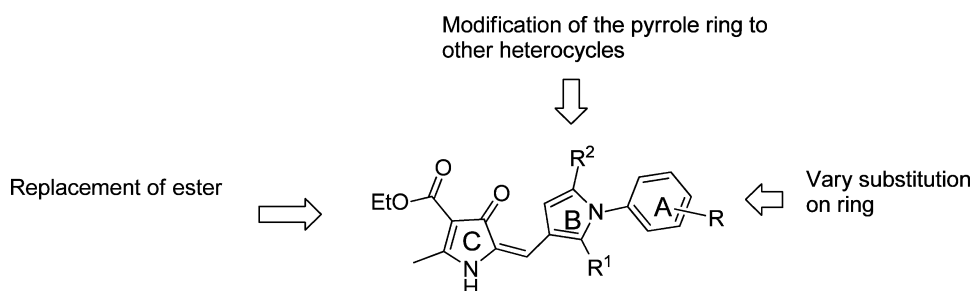
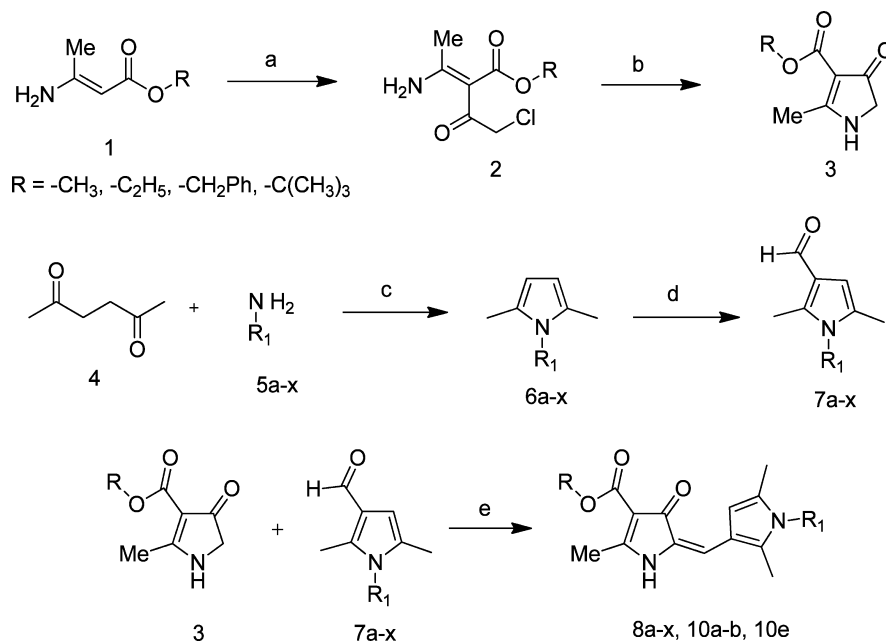


Figure 2. Strategy for elaboration of SAR around 8a.

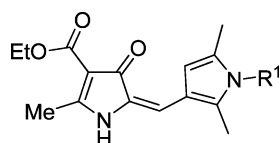
Scheme 1. General Synthetic Approach to Pyrrolones^a

^a(a) Chloroacetyl chloride, pyridine, 0 °C, 30 min, 75%; (b) KOH, EtOH, 95%; (c) *p*-toluenesulfonic acid bound with silica gel, 15–20 min, microwave, 80–90%, or *p*-toluenesulfonic acid, toluene, 90 °C, 3 h Dean–Stark apparatus; (d) POCl₃, DMF, 100 °C, 3 h, 80–95%; (e) KHSO₄, EtOH, 3 h, reflux, 80–95%; R and R¹ as defined in Tables 1 and 3.

1. The 3-formylpyrroles (7a–x) were obtained through condensation of the appropriate aniline with 2,5-hexanedione (4) (Paal–Knorr pyrrole synthesis) and subsequent Vilsmeier–Haack formylation.^{13,14} The use of silica-supported *p*-toluenesulfonic acid in the condensation of anilines with 4, in a solvent-free procedure using microwave heating, provided a considerable rate advantage over classical procedures (i.e., reaction complete within 15–20 min compared to over 12 h with external heating);

even relatively non-nucleophilic amines were condensed smoothly in the absence of Lewis acid catalysts.

Ring A. Using the approach outlined in Scheme 1, a number of analogues at the A ring (Table 1) were prepared by varying the aniline or amine 5a–x, with a particular aim of increasing the solubility. This provided a means of incorporating solubilizing groups either within the aromatic ring (e.g., pyridine analogues 8q and 8r), or as pendant amines (8y and 8aa), morpholines

Table 1. LogD/LogP, Kinetic Solubility, *in Vitro* Intrinsic Clearance and Activity against *P. falciparum* for Ring A Variants^d

Compound	R ₁	<i>P. fal</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	LogD ^a pH 7.4	cLogP	Solubility ^b (μg/ml)		CLint ^c (μL/min /mg)
						pH 2.0	pH6.5	
8a		0.009	34	3.3	4.9	6.3- 12.5	3.1-6.3	50
8b		0.019	220	3.4	4.9	<6.3	<3.1	53
8c		0.021	100	3.4	4.9	1.0- 2.1	2.1-4.2	97
8d		0.065	140	2.9	3.9	6.3- 12.5	3.1-6.3	94
8e		0.095	14	3.0	3.9	3.1- 6.3	6.3- 12.5	77
8f		0.038	100	3.4	4.5	1.6- 3.1	1.6-3.1	72
8g		0.032	100	3.4	4.6	2.1- 4.2	2.1-4.2	62
8h		0.065	140	3.0	3.8	3.1- 6.3	3.1-6.3	54

Table 1. continued

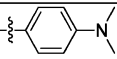
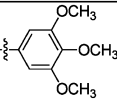
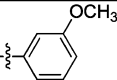
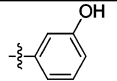
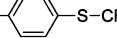
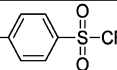
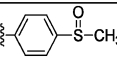
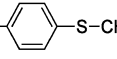
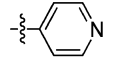
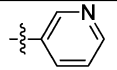
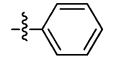
Compound	R ₁	<i>P. fal</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	LogD ^a pH 7.4	cLogP	Solubility ^b (μg/ml)		CLint ^c (μL/min /mg)
						pH 2.0	pH6.5	
8i		0.027	180	3.4	3.9	12.5- 25	<1.6	56
8j		0.063	46	2.6	3.5	6.3- 12.5	3.1-6.3	63
8k		0.026	56	2.9	3.8	6.3- 12.5	<1.6	150
8l		4.5	69	n.d.	3.6	n.d.	n.d.	n.d.
8m		0.008	140	n.d.	5.3	1.0- 2.1	1.0-2.1	76
8n		0.029	120	n.d.	4.3	n.d.	n.d.	58
8o		6.1	220	n.d.	3.5	n.d.	n.d.	n.d.
8p		0.045	54	n.d.	4.2	3.1- 6.3	1.6-3.1	66
8q		1.6	100	3.2	2.7	>100	50-100	61
8r		2.3	160	2.1	2.7	50- 100	25-50	82
8s		0.071	95	n.d.	3.8	n.d.	n.d.	n.d.

Table 1. continued

Compound	R ₁	<i>P. fal</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	LogD ^a pH 7.4	cLogP	Solubility ^b (μg/ml)		CLint ^c (μL/min /mg)
						pH 2.0	pH6.5	
8t		6.0	310	n.d.	2.6	n.d.	n.d.	n.d.
8u		0.19	210	3.3	3.8	<0.8	<0.8	31
8v		1.7	230	n.d.	2.4	n.d.	n.d.	n.d.
8w		0.12	72	n.d.	5.3	1.6- 3.1	1.6-3.1	100
8x		0.30	50	n.d.	4.2	n.d.	n.d.	n.d.
8y		0.089	81	8.5	4.5	50- 100	12.5- 25.0	60
8z		0.077	57	6.7	3.5	50- 100	12.5- 25.0	56
8aa		0.48	120	8.3	3.8	50- 100	12.5- 25.0	31

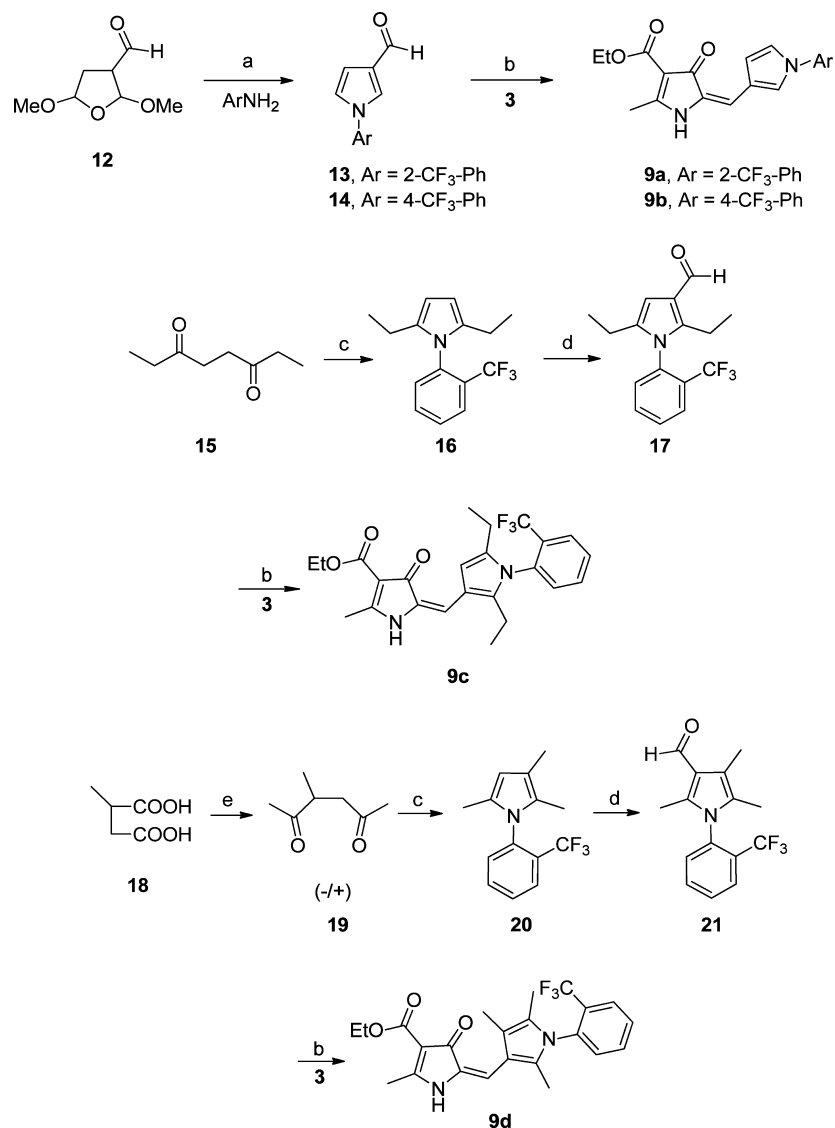
^aValue measured using a chromatographic gLogD technique. ^bEstimated using nephelometry. ^cIn vitro intrinsic clearance determined in mouse hepatic microsomes. ^dChloroquine, EC₅₀ *P. falciparum* K1 0.095–0.172 μM; podophyllotoxin EC₅₀ L6 cells 0.009–0.022 μM. The EC₅₀ values are the means of two independent assays and varies less than ±50%. n.d = not determined.

(**8u**), or sulfones (**8n**). The aryl ring was also replaced with a simple alkyl group (**8t**).

Ring B. The methyl groups on the pyrrole were predicted computationally to be likely sites of metabolic vulnerability, and it was therefore of interest to investigate the effect of modifications of the pyrrole, including removing the methyl groups (**9a–b**) and replacing them with ethyl groups (**9c**). The 3-formyl pyrrole intermediates were prepared according to Scheme 2. Variants in which the B-ring pyrrole was replaced with other 5-ring heterocycles (imidazole, pyrazole, triazole, or thiazole) were synthesized as in Scheme 3. For the furan (**9n–o**) and chloro pyrazole (**9p**) derivatives, the requisite aldehydes were commercially available.

A number of B-ring variants in which an aryl ring replaces the pyrrole (**9j–m**) were prepared using a Suzuki coupling to make the requisite aldehyde intermediates (Scheme 4, Table 2).

Ring C. Concern that hydrolysis of the ethyl ester in **8a** might adversely affect oral bioavailability and the elimination half-life led to an investigation of potentially more stable ester (Scheme 1) and amide derivatives, accessible from the carboxylic acid **10c** (Scheme 5, Table 3). Base catalyzed hydrolysis of the ethyl ester **8a** to generate **10c** proved problematic due to decarboxylation to **10d** under the conditions required. Decarboxylation could be reduced to a minimum (~10%) by utilizing acid-catalyzed deprotection of the *tert*-butyl ester derivative **10a**, with **10d** readily removed by chromatography. The effect of methylating the pyrrolone NH was also investigated.

Scheme 2. Synthesis of Pyrrole B-ring Variants^a

^a(a) Acetic acid, 90 °C, 3 h; (b) KHSO₄, EtOH, reflux, 3 h; (c) 2-trifluoromethylaniline, *p*-toluenesulfonic acid bound with silica gel, 90 °C, reflux, 3 h; (d) POCl₃, DMF, 100 °C, 3 h; (e) MeLi (3 M) in DME, THF, 0 °C – rt, 23 h, 75 °C for 17 h.

In Vitro Biological Activity. Compounds were evaluated against *P. falciparum* (K1) strain and counter-screened for cytotoxicity against mammalian L6 cells (Tables 1–3). Compound **8a** retained activity across a variety of drug-sensitive and -resistant strains (Table 4) and showed a high degree of selectivity for *P. falciparum* relative to L6.

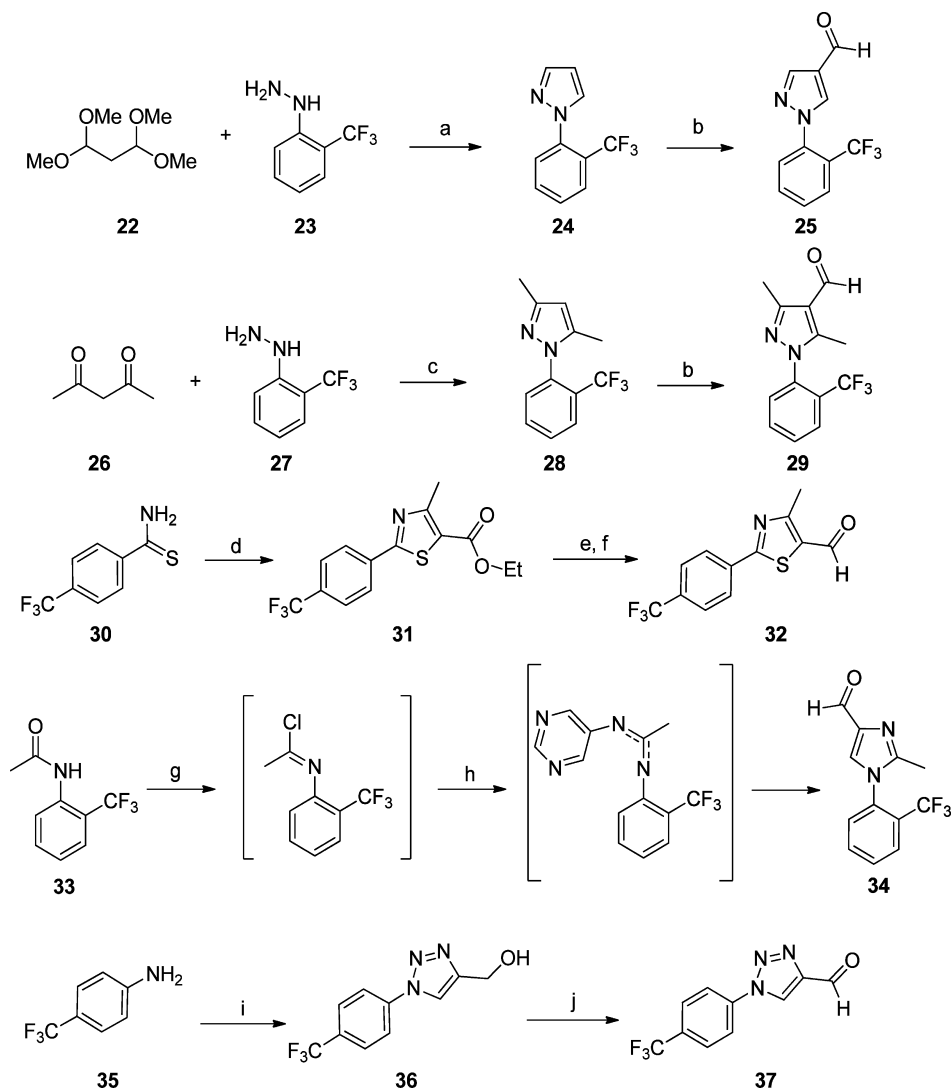
The activities of the compounds varied from 0.008 μM to 12 μM; the most active **8a–c**, **8m** and **9c** had *P. falciparum* (K1) EC₅₀ values of 0.008–0.021 μM, superior to chloroquine and the majority of the compounds showed good selectivity (>1000-fold) for *P. falciparum* compared to L6 mammalian cells. From the variations across rings A–C the following SAR was noted:

Ring A (Table 1). (1) Replacing the phenyl ring with a methyl group (**8t**) led to a significant loss of activity (600-fold), suggesting that a hydrophobic group was required. Replacing the phenyl (**8s**) with a benzyl (**8x**) resulted in a small drop in activity (4-fold), indicating that the phenyl ring did not need to be attached directly to the pyrrole.

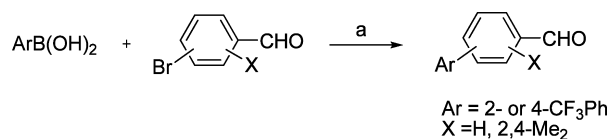
(2) In varying the position of the CF₃ on the aryl ring there seemed to be relatively little difference in activity comparing the

ortho (**8a**), *meta* (**8c**), or *para* (**8b**) derivatives. An *ortho* CF₃ would be expected to reduce the conformational flexibility of ring A by hindering coplanarity of the pyrrole and phenyl rings, but this did not seem to have an effect on either activity or solubility (see below). Compounds **8a–c** were only ~5–10-fold more active than the unsubstituted derivative **8s**, consistent with the idea that the lipophilicity contributed by ring A is the most significant variable in this part of the pyrrolones affecting *in vitro* activity.

(3) Further variation of substitution on the aromatic ring focused on the *para* position, as this is likely to be the most susceptible aryl position to cytochrome P450 (CYP)-mediated oxidation. In general, a wide variety of substituents were tolerated with relatively slight variation in potency, with the notable exception of the sulfoxide (**8o**), which was ~100-fold less active than the corresponding sulphide (**8p**) and sulfone (**8n**). Similarly, the *meta*-substituted hydroxyl derivative (**8l**) was not tolerated, while the *meta*-substituted methoxy (**8k**) was. Solubilizing substituents were tolerated to a degree: although the pyridine derivatives (**8q–r**) and dimethylamine (**8aa**)

Scheme 3. Synthesis of Formyl Pyrazole, Thiazole, Imidazole, and Triazole Intermediates^a

^a(a) aq EtOH, HCl 90 °C, 3 h; (b) POCl₃, DMF, 100 °C, 3 h; (c) acetylacetone, 90 °C, 3 h; (d) ethyl-2-chloroacetate, EtOH, reflux; (e) LiAlH₄, THF, 0 °C; (f) PCC, CH₂Cl₂, rt; (g) PCl₅ in CHCl₃, reflux, 3 h; (h) tetrachloroethylene, 5-aminopyrimidine, few drops of POCl₃; (i) *t*-BuONO/TMSN₃, propargyl alcohol, MeCN; (j) Dess–Martin Reagent, CH₂Cl₂.

Scheme 4. Synthesis of Aldehyde Intermediates for B-Ring Aryl Derivatives: (a) Tetrabutyl Ammonium Bromide (TBAB), K₂CO₃, Pd(OAc)₂, Dioxane/Water (1:1)

derivatives lost significant activity, the sulphone (8n) retained activity, while the piperidine (8y) and the morpholine (8z) were only ~5–10-fold less active than 8a.

Ring B (Table 2). The methyl substituents on the pyrrole were identified as potential points of CYP-mediated metabolism.

1. Removal of the methyl substituents (9a, 9b) resulted in a significant (~20–25-fold) loss in activity.

2. Replacing both methyls with ethyl (9c) did not significantly affect activity.

3. Replacement of the pyrrole with imidazole (9e), pyrazole (9f, 9g, 9p, 9q, and 9s), triazole (9h), thiazole (9i), aryl (9j–m),

furan (9n–o) or isoxazole (9q) gave a significant loss of activity (~20–1000-fold).

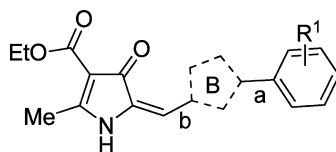
Ring C (Table 3).

1. Methylation of the NH (11a) resulted in a significant reduction of potency (>100-fold) relative to 8a, consequently no further N-substituted derivatives were investigated.

2. The potential for degradation of 8a through cleavage of the ester by esterases was of concern, although 8a was fairly resistant to chemical hydrolysis. Both the free acid 10c and the product from decarboxylation (10d) were significantly less active (~50–150-fold) than 8a.

3. All modifications of the ester in 8a, including amides 10f–m, resulted in considerably reduced activity, although with the benzyl ester 10b the reduction in activity was relatively moderate (~5-fold). The amides were inactive, which may be due to them adopting different conformations.

4. Reduction of the double bond linking the pyrrolone and pyrrole rings would be expected to improve solubility by reducing planarity. However the reduced derivative 11b proved

Table 2. cLogP, Kinetic Solubility, *in Vitro* Intrinsic Clearance and *in Vitro* Activity against *P. falciparum* for Ring B Variants

Cmpd	B	R ¹	<i>P. fal.</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	cLog P	Solubility ^a pH 6.5 (μg/ml)		Clint ^b (μL/ min/mg)
						pH 2.0	pH 6.5	
8a		2- CF ₃	0.009	34	4.9	6.3- 12.5	3.1-6.3	50
8b		4- CF ₃	0.019	215	4.9	<6.3	<3.1	53
9a		2- CF ₃	0.190	47	4.2	3.1-6.3	3.1-6.3	160
9b		4- CF ₃	0.47	18	4.2	1.6-3.1	1.6-3.1	140
9c		2- CF ₃	0.020	17	5.6	3.1-6.3	<1.6	140
9d		2- CF ₃	1.5	1.1	5.3	n.d.	n.d.	n.d.
9e		2- CF ₃	0.82	13	3.9	n.d.	n.d.	n.d.
9f		2- CF ₃	2.5	150	3.5	n.d.	n.d.	n.d.

Table 2. continued

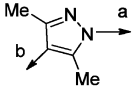
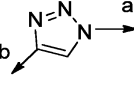
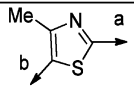
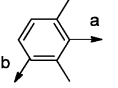
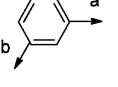
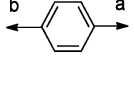
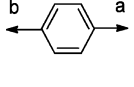
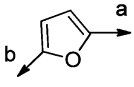
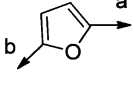
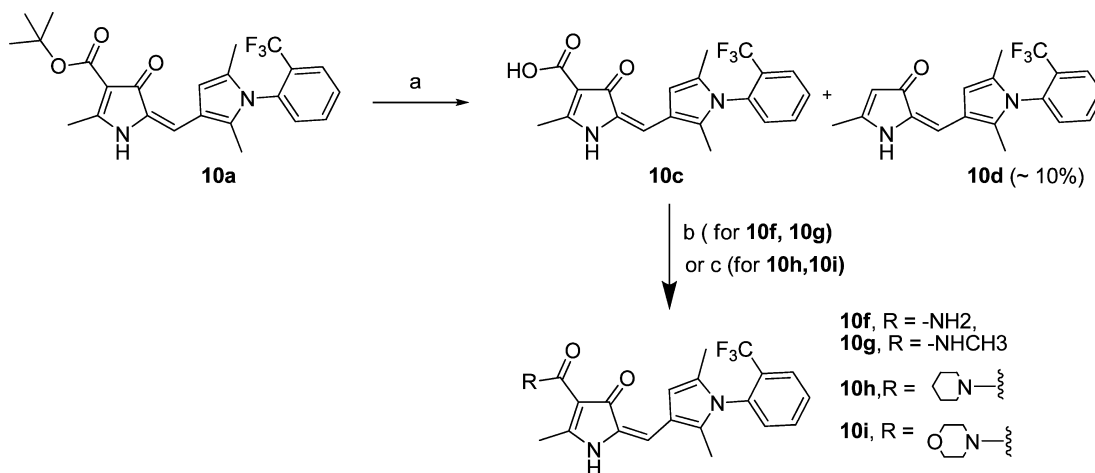
Cmpd	B	R ¹	<i>P. fal.</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	cLog P	Solubility ^a pH 6.5 (μg/ml)		Clint ^b (μL/ min/mg)
						pH 2.0	pH 6.5	
9g		2- CF ₃	5.5	120	4.3	n.d.	n.d.	n.d.
9h		4- CF ₃	2.1	14	3.0	n.d.	n.d.	n.d.
9i		4- CF ₃	4.4	62	4.6	n.d.	n.d.	n.d.
9j		2- CF ₃	0.29	10	5.8	n.d.	n.d.	n.d.
9k		2- CF ₃	1.6	13	5.1	n.d.	n.d.	n.d.
9l		2- CF ₃	0.55	7	5.1	n.d.	n.d.	n.d.
9m		4- CF ₃	8.2	130	5.1	n.d.	n.d.	n.d.
9n		2- CF ₃	0.27	99	4.4	n.d.	n.d.	n.d.
9o		3- CF ₃	2.1	13	4.4	n.d.	n.d.	n.d.

Table 2. continued

Cmpd	B	R ¹	<i>P. fal.</i> (K1) EC ₅₀ (μM)	L6 cells EC ₅₀ (μM)	cLog P	Solubility ^a pH 6.5 (μg/ml)		Clint ^b (μL/ min/mg)
						pH 2.0	pH 6.5	
9p		-	5.7	13	3.4	n.d.	n.d.	n.d.
9q		4- CH ₃	4.9	42	3.0	n.d.	n.d.	n.d.
9r		4-F	17.7	110	3.4	n.d.	n.d.	n.d.
9s		4-F	33.7	130	2.3	n.d.	n.d.	n.d.

^aEstimated using nephelometry. ^b*In vitro* intrinsic clearance determined in mouse hepatic microsomes, n.d. = not determined.

Scheme 5. Synthetic Approach to Amide Derivatives^a



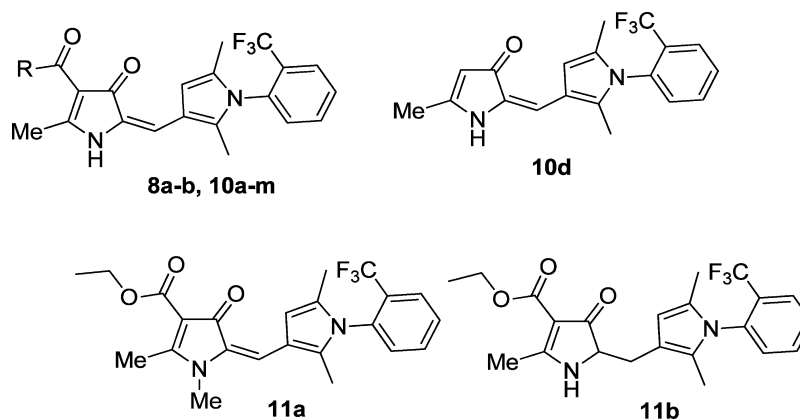
^a(a) HCl (6 M), 3 h, reflux, 80–95% (ratio **10c**/**10d** ~ 90:10); (b) *N,N*-carbonyldiimidazole, THF, reflux, excess amine in MeOH, 0° C, 18–24 h; (c) TBTU, DIPEA, secondary amine, DMF, 0 °C; 18–24 h.

significantly less active (>10-fold) than **8a**; furthermore **11b** was less stable chemically.

Thus there appears to be scope for modification of the aryl A-ring, with less room for maneuver around the 2,5-dimethyl pyrrole in ring B. As only fairly conservative modifications to ring C have been investigated there may be further potential in modifying the pyrrolone.

In Vivo Efficacy Studies in *P. berghei* Mouse Model. Compounds **8a** and **8b** showed good activity in the mouse model

of *P. berghei* when dosed ip at 100 mg/kg once daily for 4 days, with >99% reduction in parasitaemia, comparable to chloroquine at 10 mg/kg; no dose-ranging experiments were undertaken. Chloroquine was dosed ip to give comparison to a known drug by the same route. Both **8a** and **8b** were relatively inactive when given orally (Table 5). Although full PK studies were not undertaken in mice, the oral exposure of **8a** was evaluated in mice at a single dose of 50 mg/kg (formulated as a suspension in PEG400). This confirmed that relatively low plasma levels of **8a**

Table 3. LogD/LogP, Kinetic Solubility, *in Vitro* Intrinsic Clearance and *in Vitro* Activity against *P. falciparum* for Ring C Variants

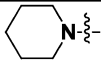
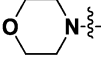
Compound	R	<i>P. fal.</i> (K1) EC ₅₀ (μM)	L6 EC ₅₀ (μM)	LogD ^a pH 7.4	cLog P	Solubility ^b (μg/ml)		CLint ^c (μL/min /mg)
						pH 2.0	pH 6.5	
8a	C ₂ H ₅ O-	0.009	34	n.d.	4.9	6.3- 12.5	3.1- 6.3	50
10a	t-butylO-	0.30	110	n.d.	5.4	3.1- 6.3	1.6- 3.1	86
10b	BenzylO-	0.05	57	n.d.	5.9	1.6- 3.1	<1.6	560
10c	HO	1.5	31	5.8	4.7	3.1- 6.3	1.6- 3.1	22
10d	-	0.43	54	n.d.	4.8	n.d.	n.d.	23
10e	CH ₃ O-	0.007	27	n.d.	4.5	n.d.	n.d.	n.d.
10f	H ₂ N-	0.62	30	n.d.	3.7	6.3- 12.5	6.3- 2.5	n.d.
10g	CH ₃ NH-	1.4	40	n.d.	4.2	n.d.	n.d.	n.d.
10h		0.58	26	n.d.	4.7	n.d.	n.d.	n.d.

Table 3. continued

Compound	R	<i>P. fal.</i> (K1) EC ₅₀ (μM)	L6 EC ₅₀ (μM)	LogD ^a pH 7.4	cLog P	Solubility ^b (μg/ml)		CLint ^c (μL/min /mg)
						pH 2.0	pH 6.5	
10i		1.0	36	n.d.	3.7	n.d.	n.d.	n.d.
10j	C ₂ H ₅ O- CH ₂ CH ₂ NH-	1.7	54	n.d.	4.5	n.d.	n.d.	n.d.
10k	HO-CH ₂ CH ₂ NH-	3.9	49	n.d.	3.5	n.d.	n.d.	n.d.
10l	(C ₂ H ₅) ₂ N- CH ₂ CH ₂ NH-	1.0	23	n.d.	4.7	n.d.	n.d.	n.d.
10m	(CH ₃) ₂ N- CH ₂ CH ₂ NH-	0.65	47	n.d.	4.0	n.d.	n.d.	n.d.
11a		1.3	102	n.d.	4.8	n.d.	n.d.	n.d.
11b		0.13	79	n.d.	5.2	n.d.	n.d.	n.d.

^aValue measured using a chromatographic gLogD technique. ^bEstimated using nephelometry. ^c*In vitro* intrinsic clearance determined in mouse hepatic microsomes. n.d. = not measured.

Table 4. *In Vitro* Antiplasmodial and Cytotoxic Activities of 8a

	<i>P. falciparum</i> EC ₅₀ (nM)				cytotoxicity EC ₅₀ (nM)	
	K1 ^a	D6 ^b	W2 ^c	TM90C2B ^d	TM91C235 ^e	L6
8a	2–28 (<i>n</i> = 7)	2.2–3.8	8.4	11–14	8.6–9.3	15000
chloroquine	63–430	14	260	250	140	
mefloquine		11	4.9	9.8	27	
artemesinin	6.0					

^aK1 = chloroquine and pyrimethamine resistant (STI). ^bD6 = chloroquine sensitive. ^cW2 = chloroquine resistant (WRAIR). ^dTM90C2B = atovaquone resistant (WRAIR). ^eTM91C235 = multidrug-resistant + atovaquone sensitive (WRAIR).

were achieved (<0.01 μM after ~5 h), which is consistent with the lack of oral efficacy in the mouse *P. berghei* model.

Physicochemical Properties and *in Vitro* DMPK. The molecules generally showed reasonable physicochemical properties (MW in the region of 350–450, 1 HBD, 5–7 HBA, PSA in the range 60–100 Å²), compatible with good membrane permeability, but the aqueous solubility of most compounds was generally very low (Tables 1–3). The low solubility is likely a consequence of the conjugated planar nature of the molecules, and in some cases the lipophilicity. For some compounds, particularly (**8i**, **8k**, **8r**) and (**9c**), there was a reduction in the solubility between pH 2.0 and pH 6.5. In the case of (**8i**) and

(**8r**), this is probably due to protonation at pH 2.0, but not at pH 6.5; although the reason for the high pH dependence on solubility is not clear for (**8k**) and (**9c**). Similarly compounds with a pendant amine also showed greater solubility (**8y**, **8z**, **8aa**) as did those with a pyridine replacement for the phenyl ring (**8q**, **8r**).

Attempts to improve the solubility were made as follows:

- Addition of solubilizing groups such as morpholine (**8u**, **8z**) and piperidine (**8y**) on ring A.
- Conversion of ester to amide linkages with various solubilizing groups such as methyl amine, piperidine,

Table 5. *In Vivo* Antimalarial Activity of 8a in Mice Infected with *P. berghei*

			% reduction parasitaemia			MSD ^a
	dose (mg/kg/ day × 4)	route	day 4	day 5	day 6	
Experiment 1						
8a	100	<i>ip</i>	97.72	98.35	99.6	11
8a	100	<i>po</i>	25.51	5.3	0	7.7
chloroquine	10	<i>ip</i>	99.9			11
control						5.7
Experiment 2						
8a	100	<i>ip</i>	99.93			10.7
8b	100	<i>ip</i>	99.91			11.7
8b	100	<i>po</i>	37.5			7
chloroquine	10	<i>ip</i>	99.97			20
chloroquine	100	<i>po</i>	99.91			>30
control						7

^aMSD = mean survival time (days).

morpholine, ethanolamine, dimethylethane-1,2-diamine, diethylethane-1,2-diamine, 2-ethoxyethanamine.

Compound **8a** is predicted to be uncharged at physiological pH as are those compounds where amines are appended directly to the phenyl ring C (**8i**, **8u**) or the pyridine analogues (**8q**, **8r**). The more basic derivatives, where the amine is appended via a methylene linker (**8y**, **8z**, **8aa**), are predicted to have varying degrees of ionization, which correlates with their significantly higher aqueous solubility.

Compounds showed moderate to high rates of degradation when incubated with mouse liver microsomes (see Supporting Information for methodology). The main exceptions to this were (**8u**, **8aa**) and (**10d**), which appeared to be slightly more stable. Putative metabolites having molecular weight consistent with the products of mono-oxygenation (P+16 for **8a**, **8d**, **8e**, **8h**, **8m**, **8p**, **8q**, **9b**, **9c**, **10a**, **10f**), bis-oxygenation (P+32 for **8q**, **10a**), O-demethylation (P-14 for **8i**), and/or morpholine ring cleavage (P-12 for **8u**), were detected for a number of analogues (see Supporting Information). Compound **10c** showed an increased rate of degradation in microsomes containing the dual cofactors NADPH and uridine-5'-diphospho-glucuronic acid (UDPGA) (the cofactors for CYP450-mediated metabolism and glucuronidation, respectively) relative to NADPH alone, suggesting that this compound is susceptible to primary glucuronidation in the microsomal test system. A putative glucuronide metabolite (P+176) was also detected for **10c** (see Supporting Information).

It was predicted computationally that the methyl groups on the pyrrole (ring B) were likely to be metabolically unstable. Hence, these were removed (**9a**, **9b**) or replaced with an ethyl group (**9c**), but these compounds were found to have even lower metabolic stability. All attempts to replace the pyrrole with other heterocycles resulted in much reduced antimalarial activity.

In addition to investigating their stability with mouse liver microsomes, key compounds (**8a** and **8b**) were investigated in rat and human liver microsomes, in order to obtain an idea of species variability in metabolic degradation (Table 6). These results suggested only marginal differences in microsomal stability between species.

Protein binding was investigated for selected compounds (Supporting Information). With the exception of the pyridine containing compounds (**8q**) (80% bound) and (**8r**) (79%

Table 6. Metabolic Stability Parameters for 8a and 8b Based on NADPH-Dependent Degradation Profiles in Human/Rat/Mouse Liver Microsomes

compound	species	$T_{1/2}$ ^a (min)	CL_{int} (μ L/min/mg)	E_H	metabolites detected
8a	human	58	30	0.62	none
	rat	44	40	0.56	P+16
	mouse	35	50	0.68	none
8b	human	35	49	0.73	none
	rat	56	31	0.50	none
	mouse	32.9	53	0.70	none

^aHalf-life. ^b*In vitro* intrinsic clearance.

bound), compounds were found to have moderate to high plasma protein binding (93.7–98.8% bound).

Compound **8a** showed insignificant reactivity on incubation with reduced glutathione ethyl ester, indicating that it does not act as a Michael acceptor (which could adversely affect stability *in vivo* and might give rise to toxicity), and no adducts resulting from conjugation with glutathione were detected in the urine of a rat dosed intravenously with **8a**.

The ester in both **8a** and **8b** is a potential point of metabolism through cleavage by esterases (Figure 3). In rat blood and plasma, ~25% of **8a** was hydrolyzed to the acid **10c** over 4 h at 37 °C. Compound **10c** was accompanied by traces of the decarboxylated product **10d**, consistent with the observed chemical instability of **10c** (although the formation of **10d** during the extraction and/or analytical procedure cannot be ruled out).

In Vivo DMPK Studies. In order to better understand the efficacy data, PK studies for compounds **8a** and **8b** were conducted in rats, with iv, ip, and po administration (Table 7; Figure 4; see Supporting Information for Methodology). No sign of toxicity was observed when **8a** or **8b** were administered orally or ip at doses up to ~20 mg/kg. There was a slight degree of hemolysis when the compounds were dosed iv, most likely arising from the vehicle required for formulation. Both compounds had terminal half-lives of at least 8 h, likely due to the high volumes of distribution. Compound **8a** exhibited high plasma clearance (73 mL/min/kg), which *in vitro* studies suggest is due to a combination of moderate hepatic metabolic clearance and blood-mediated degradation, but not renal elimination given that there was minimal recovery of **8a** excreted unchanged in urine. While plasma/blood stability studies were not conducted with **8b**, similar degradation would be expected given the structural similarity to **8a**.

Both **8a** and **8b** had low oral bioavailability (7 and 5%, respectively) which is likely due to the combined effect of high first pass clearance and poor absorption resulting from the low aqueous solubility. There is also the potential for hydrolysis of the ester linkage in the gut, and during passage through the enterocytes and the liver. Despite the low oral bioavailability, the C_{max} of both compounds when dosed at 20 mg/kg po was about 10-fold higher than the EC_{50} against parasites *in vitro*.

CONCLUSIONS

From phenotypic screening a series of pyrrolone derivatives have been identified with good *in vitro* activity against *P. falciparum* combined with good selectivity relative to the L6 mammalian cell line. A lack of cross resistance with standard antimalarials suggests they may have a novel mode of action, although this has not been investigated. While some of the pyrrolones have shown

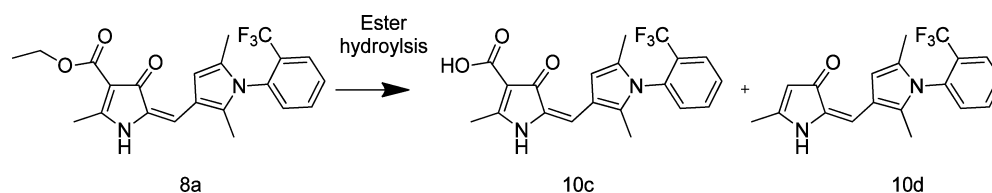


Figure 3. Degradation of 8a *in vivo* through ester hydrolysis.

Table 7. Pharmacokinetic Properties of 8a and 8b in Sprague Dawley Rats^a

	8a			8b	
	oral ^b	iv ^c	ip ^d	oral ^b	iv ^c
dose (mg/kg)	20	4.5	19	19	1.0
apparent $t_{1/2}$ (h)	7.9	12	9.1	c.n.c.	87
% dose in urine	0.08	0.05	0.01	n.d.	n.d.
C_{max} (μ M)	0.11		3.3	0.19	
T_{max} (min)	200		15	100	
oral bioavailability F%	6–7		45.7	4–5	
Vdss (L/kg)		13			19
plasma clearance (mL/min/kg)		73			22

^an.d. = not determined. c.n.c. = could not calculate. ^bOral suspension formulated with aqueous HPMC. ^civ solution formulated in aqueous vehicle with 40% (v/v) propylene glycol. ^dip suspension formulated in aqueous vehicle with 10% DMSO.

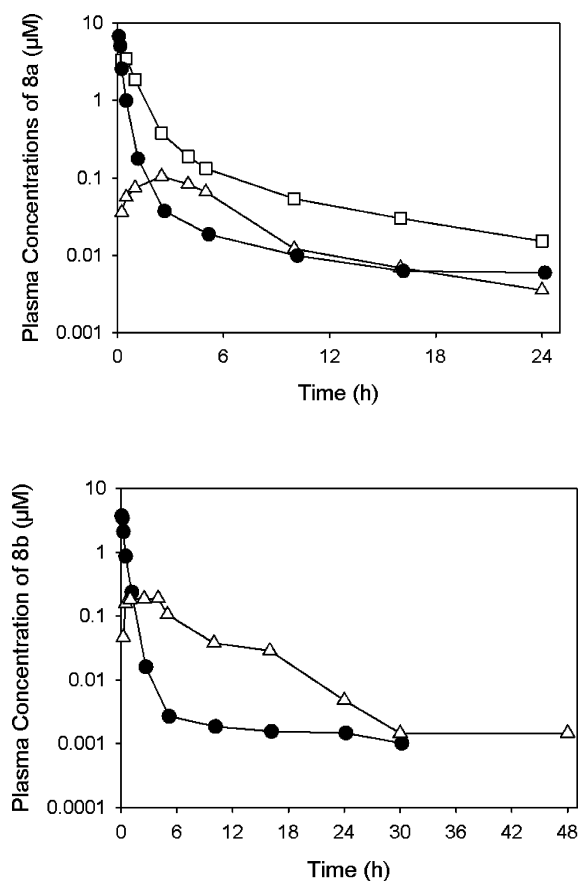


Figure 4. Plasma concentrations of 8a (top) and 8b (bottom) following iv (filled circles), oral (open triangles), and ip (open squares, 8a only) administration to male Sprague–Dawley rats at nominal iv doses of 5 mg/kg (8a) and 1 mg/kg (8b), and oral and ip doses of 20 mg/kg. Each profile represents the average of $n = 2$ rats.

good *in vivo* activity in a *P. berghei* mouse model when administered by the intraperitoneal route, oral activity has so far proved elusive, likely resulting from a combination of poor absorption due to the low aqueous solubility and rapid first-pass metabolism through cytochrome P450-mediated oxidation and/or esterase cleavage. The lack of oral activity is a key hurdle that needs to be overcome in establishing the potential for further development of this series. SAR studies in conjunction with *in vitro* microsomal stability studies have indicated that there is scope for modification at several points on the lead compounds.

EXPERIMENTAL SECTION

Parasitology and DMPK methods are described in the Supporting Information.

Profiling Software. StarDrop (www.optibrium.com) was used to predict the sites of metabolism of the compounds.

Chemistry. All commercially available reagents, solvents, and starting materials were purchased from Aldrich Chemical Co. (UK). Where necessary a Biotage FLASH 25+ column chromatography system was used to purify mixtures; reagent-grade solvents used for chromatography were purchased from Fisher Scientific (UK) and flash column chromatography silica cartridges were obtained from Biotage (UK). Analytical thin-layer chromatography (TLC) was performed on precoated TLC plates (layer 0.20 mm silica gel 60 with fluorescent indicator UV254, from Merck). Developed plates were air-dried and analyzed under a UV lamp (UV 254/365 nm). Microwave irradiation was conducted using a BIOTAGE INITIATOR unit. The machine consists of a continuous focused microwave power delivery system with operator-selectable power output (0–400 W at 2.45 GHz). All ¹H and ¹³C NMR spectra were recorded on a Bruker ARX-500 spectrometer (500 and 125 MHz for ¹H and ¹³C NMR, respectively). Chemical shifts (δ) are reported in ppm relative to the residual solvent peak or internal standard (tetramethylsilane), and coupling constants (J) are reported in hertz (Hz). Data are reported as follows: chemical shift, multiplicity (br = broad, s = singlet, d = doublet, t = triplet, m = multiplet), integration. LC–MS analyses were performed with either an Agilent HPLC 1100 series connected to a Bruker Daltonics MicroTOF or an Agilent Technologies 1200 series HPLC connected to an Agilent Technologies 6130 quadrupole spectrometer, where both instruments were connected to an Agilent diode array detector. LC–MS chromatographic separations were conducted with a Waters X bridge C18 column, 50 mm \times 2.1 mm, 3.5 μ m particle size; mobile phase, water/acetonitrile +0.1% HCOOH, or water/acetonitrile +0.1% NH₃; linear gradient from 80:20 to 5:95 over 3.5 min and then held for 1.5 min; flow rate of 0.5 mL min⁻¹. All assay compounds had a measured purity of $\geq 95\%$ (by TLC and UV) as determined using this analytical LC–MS system. High resolution electrospray measurements were performed on a Bruker Daltonics MicroTOF mass spectrometer.

Ethyl (E)-3-Amino-2-(2-chloroacetyl)but-2-enoate (2). A solution of ethyl-3-aminocrotonate (2.0 g, 0.015 mol, 1.0 equiv) and pyridine (1.2 g, 0.015 mol, 1.0 equiv) in diethylether (10 mL) was cooled to 0 °C, and a solution of chloroacetylchloride (4.1 g, 0.037 mol, 2.4 equiv) in diethylether (5 mL) was added dropwise over 30 min, maintaining the temperature at 0 °C. After the mixture was stirred for a further 3 h at 0 °C the solvent was removed *in vacuo*. The resultant solid was washed with cold water to yield 2 as a cream yellow powder (2.7 g, 87% yield), mp 131–132 °C. ¹H NMR (500 MHz; DMSO-*d*₆): δ 5.92 (br s, 2H, NH₂), 4.57 (s, 2H, –CH₂Cl), 4.27 (q, 2H, –OCH₂, $J = 7.2$ Hz),

2.36 (s, 3H, CH₃), 1.36 (t, 3H, OCH₂CH₃, *J* = 7.2 Hz); ¹³C NMR (125 MHz, DMSO-*d*₆): δ 190.6, 169.9, 168.3, 100.8, 60.5, 49.6, 24.6, 14.3; MS (ESI) *m/z* 206.1 [M + H]⁺ 100%.

Ethyl 5-Methyl-3-oxo-1,2-dihydropyrrole-4-carboxylate (3). Ethyl (*E*)-3-amino-2-(2-chloroacetyl)but-2-enoate (**2**) (2.0 g, 0.0097 mol) was dissolved in absolute ethanol (5 mL) and cooled to 0 °C. Potassium hydroxide (1.09 g, 0.019 mol) was added, and the mixture was stirred for 3 h at 0 °C and then acidified to pH 2.0 using 2 M HCl to afford a yellow precipitate, which was washed with cold water to yield **3** as a yellow solid (1.6 g, 99% yield), mp 215 °C. ¹H NMR (500 MHz; DMSO-*d*₆): δ 10.7 (br s, 1H, -OH), 9.4 (br s, 1H, -NH), 7.64 (s, 1H, -NH), 6.05 (d, 1H, *J* = 2.4 Hz), 4.27 (q, 2H, -OCH₂, *J* = 7.1 Hz), 4.08 (m, 2H, -OCH₂), 3.80 (d, 2H), 2.4 (t, 3H, *J* = 1.6 Hz), 2.3 (s, 3H), 1.8 (t, 3H, *J* = 7.1 Hz), 1.2 (t, 3H, *J* = 7.0 Hz, OCH₂CH₃); ¹³C NMR (125 MHz, DMSO-*d*₆): δ 189.2, 171.5, 159.9, 99.3, 86.1, 60.4, 28.02, 14.2; MS (ESI) *m/z* 170.14 [M + H]⁺ 100%.

General Procedure for the Microwave-Accelerated Synthesis of 2,5-Dimethyl-1-aryl-1H-pyrroles (6a-6aa). A mixture of 2,5-hexandione (**4**) (1 mmol), the appropriate aniline (**5a-5aa**) (1.2 mmol) and *p*-toluenesulfonic acid bound with silica gel (0.4 equiv) was stirred in an oven-dried pressure vial fitted with a magnetic stir bar. The vial was then placed in a microwave oven and heated twice (180 °C, 5 min) under microwave irradiation (0–400 W at 2.45 GHz). After being stirred for 15 min at room temperature, the mixture was filtered and the residual silica was washed with DCM (10 mL). The solvent was removed under reduced pressure, affording the pyrrole **6a-6aa** (purity >95%, 80–90% yield), which was used without further purification.

General Procedure for the Synthesis of 2,5-Dimethyl-1-aryl-3-formylpyrroles (7a-7aa). Phosphorus oxychloride (6 mmol) was added dropwise to ice-cooled DMF (12 mL) stirred under a N₂ atmosphere. The mixture was kept at room temperature for 15 min, and then a solution of the requisite pyrrole **6a-6aa** (1 mmol) in DMF (5 mL) was added and the mixture was heated at 100 °C for 3 h. After cooling, 30% NaOH was added dropwise to adjust to pH 10. The resultant precipitate was filtered and washed with water, affording the 2,5-dimethyl-1-aryl-3-formylpyrroles **7a-7aa** (80–95% yield), which were used without further purification.

General Procedure for the Synthesis of Ethyl (5E)-5-[[2,5-dimethyl-1-[Substituted phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylates (8a-8aa, 9a-9s). To a solution of ethyl 5-methyl-3-oxo-1,2-dihydropyrrole-4-carboxylate **3** (1.0 equiv/mol) in absolute ethanol (3 mL) was added the requisite 2,5-dimethyl-1-aryl-3-formylpyrrole **7a-7aa** (1.0 equiv/mol) and potassium hydrogen sulfate (0.2 equiv/mol). The mixture was heated at 70–80 °C for 3 h resulting in the formation of a yellow precipitate. The mixture was poured onto ice and filtered to afford the ethyl (5E)-5-[[2,5-dimethyl-1-[Substituted phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate **8a-8aa** as a yellow powder (80–95% yield). In general no further purification by column chromatography was required.

Ethyl (5E)-5-[[2,5-dimethyl-1-[2-(trifluoromethyl)phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate (8a). Yellow powder (0.22 g, 89%); mp 230–235 °C. ¹H NMR (500 MHz, DMSO-*d*₆): δ 10.3 (s, 1H, -NH), 8.01 (t, 1H, *J* = 7.7 Hz), 7.91 (t, 1H), 7.8 (t, 1H, *J* = 7.7 Hz), 7.51 (t, 1H, *J* = 7.7 Hz), 6.71 (s, 1H), 6.69 (s, 1H), 4.13–4.09 (q, 2H, *J* = 6.9 Hz), 2.58 (s, 3H), 1.99 (s, 3H), 1.90 (s, 3H), 1.23 (t, 3H, *J* = 7.1 Hz); ¹³C NMR (125 MHz, DMSO-*d*₆): δ 180.5, 170.2, 163.3, 135.9, 134.36 (2C), 131.6, 131.6, 130.43 (2C), 129.17 (2C), 127.5, 113.5, 169.7, 105.9, 102.4, 58.2, 15.7, 14.4, 11.9, 10.3. IR (KBr) ν 3500–2000 (max at 3176.47, 2926.66, and 2340.11 N–H, and C–H st), 1661.80 and 1583.65 (C=O, ar–C–C and ar–C–N st) cm^{−1}. LCMS *m/z* 419.15 (M + 1); HRMS *m/z* calculated (C₂₂H₂₂N₂O₃F₃): 419.1577 (M + H)⁺; found: 419.1566. (Delta PPM: 2.40 ppm; acquired on Microtof; mass resolution: 10000 (fwhm)).

Ethyl (5E)-5-[[2,5-Dimethyl-1-[4-(trifluoromethyl)phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate (8b). Yellow powder, mp 252–255 °C; ¹H NMR (500 MHz, DMSO-*d*₆): δ 10.27 (s, 1H, -NH), 7.96 (d, 2H), 7.65 (d, 2H), 6.75 (s, 1H), 6.70 (s, 1H), 4.16 (q, 2H, *J* = 5.4 Hz), 2.59 (s, 3H), 2.13 (s, 3H), 2.04 (s, 3H), 1.23 (t, 3H, *J* = 1.0, 1.1 Hz); ¹³C NMR (125 MHz; DMSO-*d*₆): δ 180.4, 170.2, 163.2, 140.6, 134.5, 130.4, 129.2, 128.9, 128.6, 126.67, 126.64, 124.9, 122.8, 113.9, 109.4, 106.6, 102.4, 58.1, 15.8, 14.4,

12.6, 10.7; LCMS *m/z* 419.1527 (M + 1); HRMS *m/z* 419.1579 ([M + H]⁺; calcd for C₂₂H₂₁F₃N₂O₃⁺ 419.1577.

Ethyl (5E)-5-[[2,5-Diethyl-1-[2-(trifluoromethyl) phenyl] pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate (9c). Yellow powder, mp 245–250 °C; ¹H NMR (500 MHz, DMSO-*d*₆): δ 10.31 (s, 1H, -NH), 8.01 (d, 1H), 7.92 (t, 1H), 7.83 (t, 1H), 7.60 (d, 1H), 6.71 (s, 1H), 6.69 (s, 1H), 4.13 (m, 2H, *J* = 5.6, 1.1 Hz), 2.61 (br s, 2H), 2.50 (s, 2H), 2.16 (m, 3H, *J* = 6.2, 7.6 Hz), 1.23 (t, 3H); ¹³C NMR (125 MHz; DMSO-*d*₆): δ 187.7, 180.5, 170.2, 163.3, 141.6, 137.7, 134.4, 134.0, 132.0, 132.0, 129.1, 127.6, 126.9, 112.8, 109.3, 104.5, 102.4, 58.2, 19.3, 17.6, 15.8, 15.3, 14.4, 12.4; IR (KBr) ν 3500–2000 (max at 3159.66, 2926.40, and 2358.21 N–H, and C–H st), 1661.05 and 1582.35 (C=O, ar–C–C and ar–C–N st) cm^{−1}; LCMS *m/z* 447.18 [M + H]⁺; HRMS *m/z* 447.1882 ([M + H]⁺; calcd for C₂₄H₂₆F₃N₂O₃⁺ 447.1890).

tert-Butyl-(5E)-5-[[2,5-dimethyl-1-[2-(trifluoromethyl)-phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate (10a). Yellow powder (99% yield), mp 201–205 °C. ¹H NMR (500 MHz; DMSO-*d*₆): δ 10.13 (br s, 1H, -NH), 8.02 (d, 2H, *J* = 7.0 Hz), 7.93 (t, 1H, *J* = 7.6), 7.83 (t, 1H, *J* = 7.7 Hz), 6.7 (s, 1H), 6.63 (s, 1H), 4.3 (m, 2H, *J* = 5.0 Hz), 1.98 (s, 3H), 1.9 (s, 3H), 1.4 (s, 9H), 1.07 (t, 3H, *J* = 7.0 Hz); ¹³C NMR (125 MHz, DMSO-*d*₆): δ 181.6, 180.8, 168.6, 169.6, 162.4, 135.5, 131.7, 129.2, 127.6, 123.8, 113.6, 112.1, 108.9, 106.0, 78.1, 55.9, 28.2(3C), 18.5, 16.6, 12.0(2C) 10.3; LCMS *m/z* 447.18 (M + 1); HRMS *m/z* 447.1888 ([M + H]⁺; calcd for C₂₄H₂₆F₃N₂O₃⁺ 447.1890).

Benzyl (5E)-5-[[2,5-Dimethyl-1-[2-(trifluoromethyl)phenyl]pyrrol-3-yl]methylene]-2-methyl-4-oxo-1H-pyrrole-3-carboxylate (10b). Yellow powder (95% yield); mp 233–236 °C. ¹H NMR (500 MHz; DMSO-*d*₆): δ 10.36 (br s, 1H, -NH), 8.02 (d, 2H, *J* = 7.7 Hz), 7.9 (t, 1H, *J* = 7.0 Hz), 7.8 (t, 1H, *J* = 7.5 Hz), 7.54 (d, 2H, *J* = 7.5 Hz), 7.47 (d, 1H, *J* = 7.2 Hz), 7.39 (t, 1H, *J* = 7.2 Hz), 7.32 (t, 1H, *J* = 7.3 Hz), 6.72 (s, 1H), 6.70 (s, 1H), 5.19 (br s, 2H, CH₂Ph), 2.5 (s, 3H), 2.0 (s, 3H), 1.9 (s, 9H); ¹³C NMR (125 MHz, DMSO-*d*₆): δ 180.4, 170.5, 163.1, 137.2, 136.0, 134.4, 131.7(2C), 131.6, 130.5, 129.2, 128.3(3C), 127.5(2C), 127.3(3C), 113.6, 109.8, 106.0, 102.2, 63.7, 15.8, 12.0, 10.3; LCMS *m/z* 481.17 (M + 1); HRMS *m/z* 481.1736 ([M + H]⁺; calcd for C₂₇H₂₄F₃N₂O₃⁺ 481.1734).

■ ASSOCIATED CONTENT

Supporting Information

Synthetic routes for the synthesis of intermediates, methods for physicochemical evaluation of the compounds and the data generated, methods for the drug metabolism, pharmacokinetics, in vitro and in vivo parasite testing; chemistry experimental for compounds not included in the main text. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

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■ ABBREVIATIONS

CLint, intrinsic clearance; CYP, cytochrome P450; DMPK, drug metabolism and pharmacokinetics; SAR, structure–activity

relationship; UDPGA, uridine-5'-diphospho-glucuronic acid; WHO, World Health Organisation; WHO-TDR, World Health Organisation Programme for Research and Training in Tropical Diseases

■ ADDITIONAL NOTE

^aSome of the compounds had a trace, inseparable amount of Z isomer. This could not be detected in the ¹³C NMR spectra and was only detected using UPLC.

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