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Hydroxylated Di- and Tri-styrylbenzenes a New Class of AntiplasmodialAgents:Discovery and Mechanism of Action†

Naina Sharma,[a,b] Dinesh Mohanakrishnan,[c] Amit Shard,[a] Abhishek Sharma,[a,b] Arun K. Sinha*[a,d] and Dinkar Sahal*[,c]

The first systematic evaluation of antiplasmodial activity of hydroxy stilbene family of natural products and di/tristyrylbenzenes is described. A library of **27**diversely substitutedhydroxy stilbenoids was rapidly synthesized using modified Knoevenagel-Perkin-decarboxylation-Heck sequences from readily available starting materials (i.e. hydroxybenzaldehyde-phenylacetic acid-arylhalide). These compounds were evaluated for *in vitro* antiplasmodial activity against three different strains of *Plasmodium falciparum*. Notably,(2*E*)-1-{4-[(*E*)-2-(4-hydroxy-3,5-dimethoxyph-enyl)ethenyl]phenyl}-3-(2,4,5-trimethoxyphenyl)-prop-2-en-1-one (**27**), an octupolar stilbenoid, showed IC_{50} (μM) of 0.6, 0.5 and 1.36 while a distyrylbenzene (**11**) showed IC_{50} of 0.9, 2.0 and 2.7 against 3D7 (chloroquine sensitive), Dd2 and Indo (chloroquine resistant) strains of *Plasmodium falciparum* respectively Moreover, **27** and **11** which exhibited selectivity indices of 40 and >111 were also found to be nontoxic to HeLa cell line. Microscopic studies revealed that the rings and trophozoites obtained from the **27** and **11** (an octupolar tristyrylbenzene and distyrylbenzene respectively) treated cultures were growth inhibited and morphologically deformed. These cultures also showed DNA fragmentation and loss of mitochondrial membrane potential (ΔΨm) suggestive of apoptotic death of the parasite. Together, these studies introduce di/tristyrylbenzenes as a new class of antimalarial agents

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Introduction

Malaria is one of the most infectious diseases known to mankind and it causes enormous mortality.[1]Although, diverse potent antimalarial agents including quinoline (e.g. *chloroquine) and endoperoxides (e.g. artemisinin)* are available, however resistance against antimalarial drugs have been increasing at an alarming pace. Thus, the search for newer efficacious drugs as well as new molecular scaffolds possessing antimalarial activity remains a vital goal towards achieving control over malaria.[2]

Hydroxylatedstilbenes represent an important class of natural products having special therapeutic significance owing to their various pharmacological activities[3] including anticancer and anti-inflammatory effects. Although there are few reports describing the antimalarial potential of natural prenylated,[4] glycosidic[5] or benzamide[6] containing stilbene derivatives (Figure 1), however, the exploration of higher order stilbenoids (dimeric/trimeric) against *P. falciparum* has not received much attention. For instance, although distyrylbenzenes (DSBs) find important applications in detection[7] and treatment[8] of neurodegenerative disease, but till date DSBs as well as the tristyrylbenzenehave never been investigated as antimalarials.

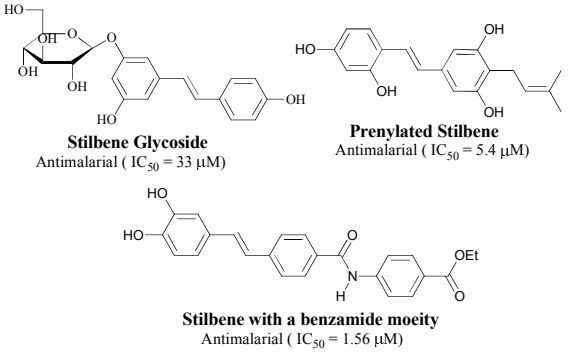

Fig. 1 Example of some previously reported antimalarial stilbenoids.

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† *Electronic Supplementary Information (ESI) available: ^{1}H NMR, ^{13}C NMR data of selected compounds.See DOI: 10.1039/x0xx00000x*

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Pursuant to our interest in synthesis[9-11] and biological activities[12] of hydroxylatedstilbenoids it was envisaged to synthesize a library of monomeric, dimeric and trimerichydroxylatedstilbenoidsand screen them as antimalarials. Herein we show that hydroxylated octupolar[13]stilbenoid (**27**) and a distyrylbenzene (**11**) are highly potent antiplasmodial agents across chloroquine sensitive and chloroquine resistant strains of *P. falciparum*. Mechanistic studies have indicated that these molecules cause morphological deformations, DNA fragmentation and loss of mitochondrial membrane potential ($\Delta\Psi m$) in malaria parasite suggesting apoptotic cell death.

Results and discussion

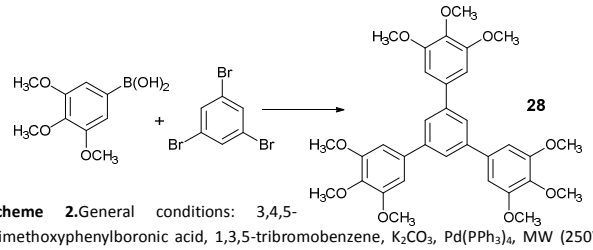
Synthesis of hydroxylatedstilbenoids

Amongst the various possible synthetic routes towards hydroxystilbenoids (A-HC=CH-B), the modified Knoevenagel Heck[9] and Perkin[10] approaches were utilized, as these allowed a rapid access to diversely functionalized stilbenoids using readily available starting materials such as hydroxylatedbenzaldehyde, phenyl acetic acid and halobenzene. A library comprising of 27 diversely substituted hydroxylatedstilbenoids including stilbenes, distyrylbenezenes and tristyrylbenzenes were synthesized. Thus, hydroxylatedstilbenes, **1** to **8** (Scheme 1) were obtained via synthesis of hydroxy substituted styrenes (ring **A**) using modified KnoevenagelDoebner-decarboxylation approach followed by their *in situ* Heck coupling[9] with diversely substituted aryl halides (ring **B**).

On the other hand, **9&10** possessing bromo functional group on ring **B** (Scheme 1) were synthesized via modified Perkin condensation-decarboxylation[10] reaction. Hydroxy substituted symmetrical DSBs (**11-19**) with different substitution patterns on rings **A** and B (**11-17**) or with variations on ring **C** (**18&19**) were synthesized via sequential Knoevenagel/decarboxylation-double Heck reaction in one-pot.[9]

Unsymmetrical DSBs **20-26**(Scheme 1) having a 4-hydroxy substitution at ring B and different electron releasing groups at ring **A**, were synthesized by Perkin-Knoevenagel-condensation-decarboxylation-Heck strategy under microwave irradiation.[11]

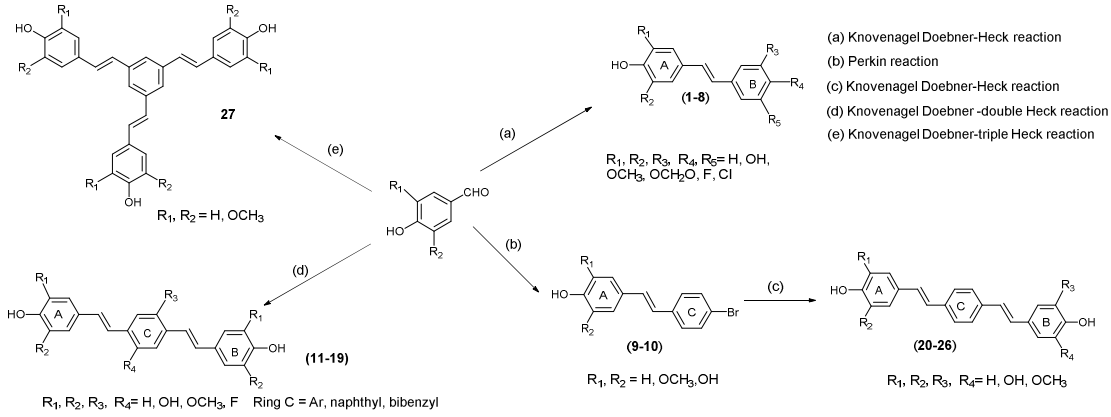
In the course of our efforts to further enhance the diversity of our panel of higher order stilbenoids, the oligophenylenevinylene (OPV, **27**)[9] with 4-hydroxy-3,5-dimethoxysubstitution (Scheme 1) was prepared via Heck reaction of *in situ* formed 4-hydroxy-3,5-dimethoxystyrene with 1,3,5-tribromobenzene in DMF under reflux conditions. In order to gauge the specific effect of octupolar moiety, an analogue[14] of **27** possessing biaryl linkage (**28**, Scheme 2) was synthesized via palladium catalyzed triple Suzuki coupling of 1,3,5-tribromobenzene with 3,4,5-trimethoxy- phenylboronic acid under microwave in the presence of $Pd(PPh_3)_4$, K_2CO_3 in dioxane: water (5:1).



Scheme 2. General conditions: 3,4,5-trimethoxyphenylboronic acid, 1,3,5-tribromobenzene, K_2CO_3, $Pd(PPh_3)_4$, MW (250W, 115°C) in dioxane: water (5:1) for 35 min.

Biological activity of monomeric/dimeric/trimericstilbenes

The hydroxylated (**1-8**) and brominated (**9-10**) stilbenes were subjected to micro titre plate high-throughput format SYBR Green I fluorescence based antiplasmodial screen.[15] As shown in Table 1, 4,4′-dihydroxystilbene (**1**) displayed an IC_{50} of 92 μM against the 3D7 strain of *P. falciparum*. Replacement of one of the hydroxy groups with methoxy i.e. 4-hydroxy-4′-methoxystilbene (**2**) caused a decrease in potency (IC_{50}>100 μM).



Scheme 1. Reagents and conditions: (a) Substituted iodobenzene, $CH_2(COOH)_2$, $Pd(PPh_3)_4$, piperidine, LiCl, DMF, reflux 10 h (b) p-bromophenylacetic acid, N-methylimidazole, piperidine, PEG-200, Microwave (160°C) for 25 min (c) Malonic acid, substituted *p*-hydroxybenzaldehyde, $Pd(PPh_3)_4$, piperidine, LiCl, DMF, MW (150°C) for 45 min (d,e) $CH_2(COOH)_2$, piperidine, $Pd(PPh_3)_4$, LiCl, substituted 1,4-diiodobenzene in condition d and 1,3,5-tribromobenzene in condition e were refluxed in DMF for 14

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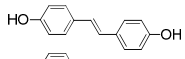
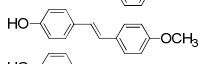
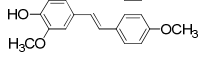
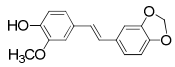
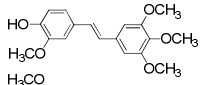
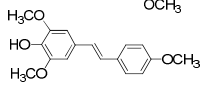
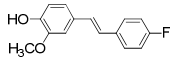
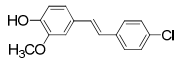
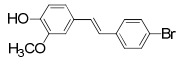
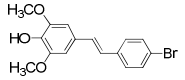
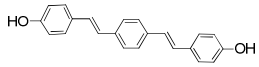
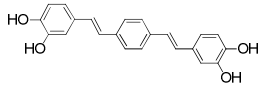
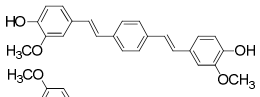
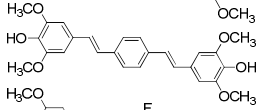
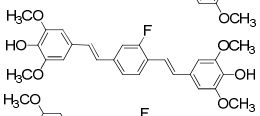
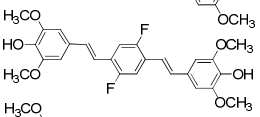
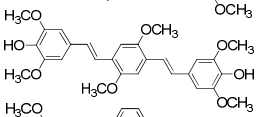
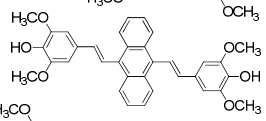
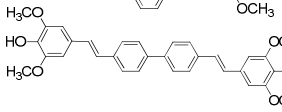
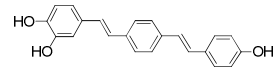
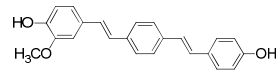
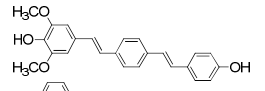
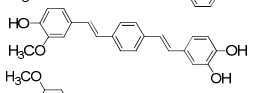
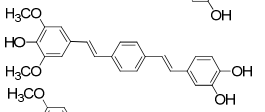
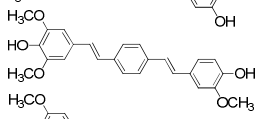
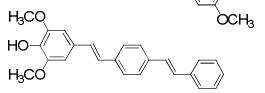
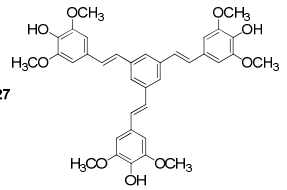
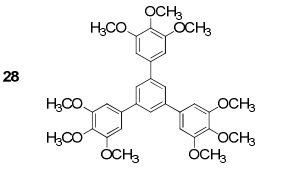
Table 1. Antiplasmodial activities of unsymmetrical and symmetrical stilbenoids against *P. falciparum* 3D7 strain.

Compound No.	Structure	IC_{50}(μM)
1		92
2		>100
3		>50
4		>100
5		47.5
6		>50
7		>100
8		88
9		99
10		63
11		0.9
12		66
13		18
14		11
15		50
16		26
17		15
18		16
19		>100
20		>100
21		1.9
22		2.4
23		6.0
24		1.4
25		15.0
26		>100
27		0.62
28		>100
Chloroquine		40 nM

Later on, 4-hydroxy-3-methoxy substitution on ring **A** was kept constant and the effects of different electron releasing (Table 1, **3-5**) and electron withdrawing groups (**7-9**) on the ring **B** were evaluated. Interestingly, **5** (IC_{50} 47.5 μM) possessing greater electron density on both rings showed slightly better antiplasmodial activity than **1-4** and **7-9** (IC_{50}> 50 μM). However none of these monomeric stilbenoids showed significant antimalarial potential.

Hence, it was decided to extend the conjugation of the core stilbene monomers (Table 1, **1-10**) and evaluate the antiplasmodial potency of resulting distyrylbenzenes (**11**). It is evident from the screening results that a dimer of the inactive 4,4′-dihydroxy monomeric stilbene (**1**, IC_{50} 92μM) i.e. 4,4′-dihydroxy substituted symmetrical distyrylbenzene (**11**) displayed promising activity (IC_{50} 0.9 μM) against the 3D7 strain of *P. falciparum*. An increase in hydroxy substitution on rings **A** and **B** of dimericstilbene led to decreased antiplasmodial potential (**12**, IC_{50} 66.0 μM) while introduction of methoxy substitution (**13**) resulted in comparatively better activity (IC_{50}18 μM) than **12**. Interestingly, a further increase in electron density (**14**) slightly improved the activity (IC_{50} 11 μM). On the other hand dimericstilbenes**15** (IC_{50} 50 μM) and **16** (IC_{50} 26 μM) having the same substituents as **14**, but having the electron withdrawing fluoro groups on ring **C** (**15** &**16**) showed decrease in antiplasmodial potency. Further, the replacement of central ring **C** with 2,5-dimethoxybenzene (**17**) &anthracenyl (**18**) led to moderate antiplasmodialactivity (**17**, IC_{50}15 μM&**18**, IC_{50} 16 μM), whereas the presence of a biaryl ring (**19**) diminished the activity (IC_{50}>100 μM). It is evident from above discussion that the presence of either 4-hydroxy or4-hydroxy-3,5-dimethoxy substitution on terminal rings **A** &**C** augments antiplasmodial activity.

After the antiplasmodial analysis of the symmetrical DSBs (Table 1, **11-19**), it was decided to evaluate the effect of

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unsymmetrical substitution pattern of distyrylbenzenes (Table1, **20-26**). The antiplasmodial evaluation results for the unsymmetrical DSBs (**20-26**) indicated that among the three compounds (**20-22**), the DSB possessing 3, 4-dihydroxy functionality at ring **A** (**20**) (IC_{50} >100 µM) was inactive (Table 1). Surprisingly, slight variation of substituents i.e. having 4-hydroxy-3-methoxy (**21**, IC_{50} 1.9 µM) and 4-hydroxy-3,5-dimethoxy (**22**, IC_{50} 2.4 µM) substitutions at ring **A**, significantly enhanced the antiplasmodial activity (Table 1).

Interestingly, a further increase in electron density on ring **B** (**23 & 24)** of dimericstilbenesled to decreased activity in **23** (IC_{50} 6 µM) while increase in potency was observed in case of **24** (IC_{50} 1.4 µM). Further, replacement of a hydroxy group with methoxy in ring **B** (**25**) decreased the activity (IC_{50} 15 µM). Similarly, **26** (IC_{50} >100 µM) having an unsubstituted ring **B** was also found to be inactive. In view of above results, potent activity seems to be associated with the presence of 4-hydroxy-3-methoxy (ring **A**) and 4-hydroxy (ring **B**) (**21**) or 4-hydroxy-3,5-dimethoxy i.e. syringol (ring **A**) and 3,4-dihydroxy (ring **B**) (**24**). Thereafter, the tristyrylbenzene**27** (Table 1) was also evaluated for antimalarial activity. Interestingly it was found to be the most potent *Pf*3D7 inhibitor with IC_{50} 0.62 µM. In contrast to octupolarstilbene (**27**, IC_{50} 0.62 µM), the triarylbenzene (**28)** with IC_{50}>100 µM (Table 1, Scheme 2) was found to be inactive suggesting a specific role of tristyrylbenzenemoiety in *P. falciparum* inhibition.

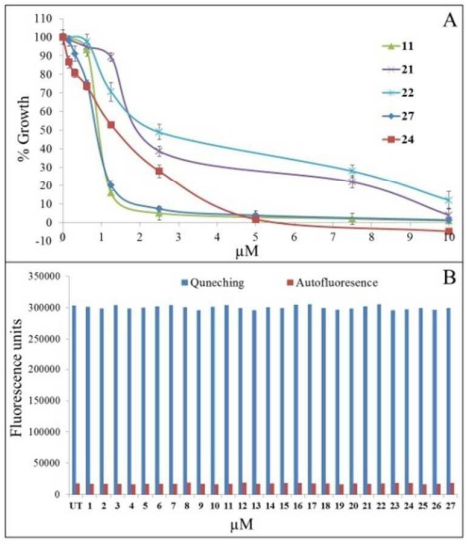

Fig. 2 (A) Dose dependent antiplasmodial activities of potent stilbenoids estimated by SYBR green assay against *P. falciparum* 3D7. Compound numbers and the corresponding color codes are indicated in the strip on the right. Standard deviation bars at each data point have been calculated from triplicate observations. (B) Data validation. Fluorescence intensity test of autofluorescent or quenching nature of test molecules: Untreated (UT) or drug (1-27) treated cultures were subjected to fluorescence intensity measurements using excitation/emission of 480/535 nm (red bars). The nearly identical intensities across control and test samples indicate that the test compounds caused no interference due to autofluorescence. Control and test molecule treated cultures were read for fluorescence intensities following lysis by SYBR Green I containing lysis buffer (blue bars). The nearly identical intensities across control and test samples indicate absence of quenching effects.

*It is worthwhile to mention that unlike DSBs (**11-26**) which have been used for the detection/treatment of Alzheimer's disease, such octupolarstilbenoids (**27**) have not yet been explored as biologically active agents* even though their application in material science[16] and non-linear optics[17] is well documented. The dose dependent growth inhibition of malarial parasites by potent stilbenoids is shown in Fig 2A. Since test molecules can interfere with SYBR green fluorescence[15], all the test molecules were checked for (a) auto fluorescence and (b) quenching effects in cultures of malaria parasites. Measurement of auto florescence helps avoid false negatives among the active molecules which may be fluorescent. Evaluation of quenching effects helps to avoid false negatives from test molecules that may be quenchers of fluorescence but may have no antiplasmodial action. As shown in Fig 2B, none of the **27**molecules studied by us (a) had intrinsic fluorescence or (b) had the ability to quench the fluorescence due to SYBR Green. This gives validity to the results obtained by the SYBR Green assay. The results of SYBR Green assay were further corroborated by examining the time dependent effects of **11**&**27** on the malaria parasite using microscopy on Giemsa stained smears of parasitecultures (Figure 3).

Subsequently, trimeric**27** (Table 1) and some potent DSBs (IC_{50} < 2.5 µM) (Table 1, **11, 21**, **22** & **24**) were explored against two chloroquine resistant strains of *P. falciparum* (*Pf*Dd2 &*Pf*INDO) (Table 2). Importantly, **27** displayed a broad spectrum of antiplasmodial potential with IC_{50} 1.36 µM (*Pf* INDO) and 0.5 µM (*Pf* Dd2). Further, **27** also showed a selectivity index of 40 against HeLa cell lines there by indicating it to be non-toxic.

Table 2. Antiplasmodial potential, Resistance and Selectivity indices for potent stilbenoids

	SYBR green Assay IC_{50}(µM)			Resistance Index		Selectivity Index	
Comp. No.	*Pf*3D7	*Pf*3D2	*Pf*Indo	IC_{50}Dd2/ IC_{50}3D7	IC_{50}Indo/ IC_{50}3D7	IC_{50}HeLa/ IC_{50}3D7	IC_{50}L929/ IC_{50}3D7
11	0.9	2.0	2.7	2.2	3	>111.11	>111.11
21	1.9	3.2	2.2	1.5	1.2	15.8	>52.6
22	2.4	3.2	2.8	1.3	1.2	4.7	>41.6
24	1.4	6.25	6.1	4.4	4.5	18.6	39.3
27	0.62	0.5	1.36	0.8	2.2	40.3	29.0

Probing the antiplasmodial action of DSB 11 and octupolarstilbenoid 27

Microscopic examination of **11** and **27** treated parasite cultures revealed stressed and shrunken "crisis forms" (Figure 3). This led us to explore if these two lead compounds were triggering apoptotic programmed cell death in the parasite. Synchronized parasite cultures of *Pf*3D7 were incubated with **27** and **11** at their respective IC_{90}{1.3 µM (**27**) & 1.8 µM (**11**)} for 12, 24, 48 hrs (rings) and 6, 12, 24 hrs (trophozoites). In contrast to the

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healthy appearances of the untreated control cultures and their sequential transition to the subsequent stages, the pycnotic

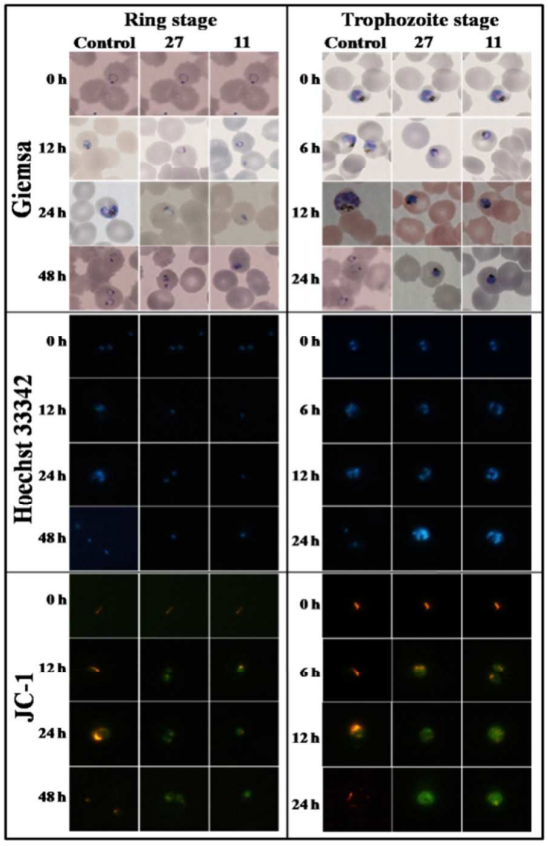

Fig. 3: Microscopic studies indicating apoptotic cell death in **11** and **27** treated malaria parasites. Ring and trophozoite stages of *Pf*3D7 were treated with compounds **27** and **11** at their IC_{90}values for different time points indicated against each panel. Giemsa staining showed stressed and shrunken "crisis forms in **27** &**11** treated cultures. Drug treated ring and trophozoite stages lagged behind their respective controls in terms of ring to trophozoite and trophozoite to schizont transitions. Increased fluorescence of Hoechst 33342 in treated trophozoites (24 h) suggests DNA condensation and fragmentation in malaria parasites. JC-1 staining indicates loss of mitochondrial membrane potential in **27** and **11** treated ring and trophozoites stages. For zoom of these images please see Supplementary Figures S1a-S1c

appearances of the rings and the trophozoites in the treated cultures suggested that stress and probable death had prevented the parasite to transit to the respective next stages of its life cycle. It may be noted that cell shrinkage (crisis form) which can be readily observed under the microscope, is one of the major characteristic features of apoptotic cell death.[18] The shrunken, condensed and darkly stained nuclei of **27** and **11** treated rings and trophozoites stages as observed by Giemsastaining (Figure 3 and Supplementary Figure S1a for zoom) led us to look for other indicators of apoptotic cell death.

DNA fragmentation and condensation are among the major features of apoptosis.[18] Minor groove binding DNA stain, Hoechst 33342 was used to monitor DNA condensation in drug treated parasite cultures. As shown (Figure 3), the control rings showed progressively greater staining as the rings matured to become early trophozoites (12 h) and late trophozoites (24 h). In contrast, **11** and **27** treated rings failed to become trophozoites and appeared as highly condensed dot like bodies till 48 hrs. Staining with Hoechst 33342 in control parasite cultures owes its low intensity to the fact that highly compact and condensed native chromatin exposes very few sites for the binding of dyes like the Hoechst 33342. DNA fragmentation results in higher degree of exposure and greater number of binding sites resulting in brighter fluorescence associated with cells undergoing apoptosis. The Hoechst 33342 staining of **11** and **27** treated trophozoites showed strong fluorescenceat 12 h. At 24 h while the control culture showed dull staining in the form of small dots corresponding to rings, the treated cultures showed strong fluorescence originating from arrested trophozoitesharboring fragmented chromatin spread over a large area (Fig. 3 and Supplementary Fig.S1b for zoom). It is interesting to note that the chromatin fragmentation observed here bears resemblance to the pattern of chromatin fragmentation we have earlier observed in trophozoite stage cultures treated with Stilbene-Chalcone hybrids.[12]While Hoechst 33342 staining based results described above were suggestive of DNA fragmentation, a more direct proof of DNA fragmentation was obtained from agarose gel electrophoretic analysis of genomic DNA obtained from 11/27 treated vs control parasite cultures (Fig 4). While the untreated culture showed the only intense band of genomic band, the DNA obtained from 11/27 treated cultures showed a diminished intensity of genomic DNA together with a streak of low molecular weight DNA fragments. Interestingly, the intensity of these fragmented pieces of genomic DNA was higher in cultures from 12 hr exposure than from analogous cultures resulting from 24 hr exposure to 11/27. This data suggests that 11 and 27 may induce DNA fragmentation at times earlier than 12 hrs and 24 hrs may be the time corresponding to extensive DNA breakdown to sizes too small to be detected by this gel based staining method. Indeed this time dependent phenomenon observed with 11/27 found a good match with chloroquine (used as positive control in this experiment) which also showed a higher intensity of fragmented DNA bands at 12 hrs than at 24 hrs.

Loss of mitochondrial membrane potential ($\Delta\Psi m$), a hallmark of apoptosis[19] was observed using JC-1 dye fluorescence microscopy in **27** and **11** treated malarial parasites. This dye is known to acquire a red color upon aggregation in the ambience of a high $\Delta\Psi m$. When $\Delta\Psi m$ is abolished by drugs that trigger apoptosis, the disaggregation of the dye causes its color to change from red to green. Ring stage parasites treated with **27** and **11** showed complete loss of $\Delta\Psi m$ (fully green) at 12 hrs and 24 hrs respectively (Figure 3 and Supplementary Figure S1c for zoom). However, the vulnerability of trophozoites to **11**and **27** appeared to be identical since 12 hrs treatments with each one of them resulted in complete loss of $\Delta\Psi m$.

Apoptosis, a process of programmed cell death involving features of cell shrinkage, DNA fragmentation, chromatin condensation, formation of apoptotic bodies, translocation ofphosphatidyl serine from inner to the outer leaflet of the plasma membrane and the loss of mitochondrial membrane potential has found vivid description in multicellular

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organisms.[20,21]However it is now becoming increasingly apparent that the evolutionary origins of Apoptosismaygo back to

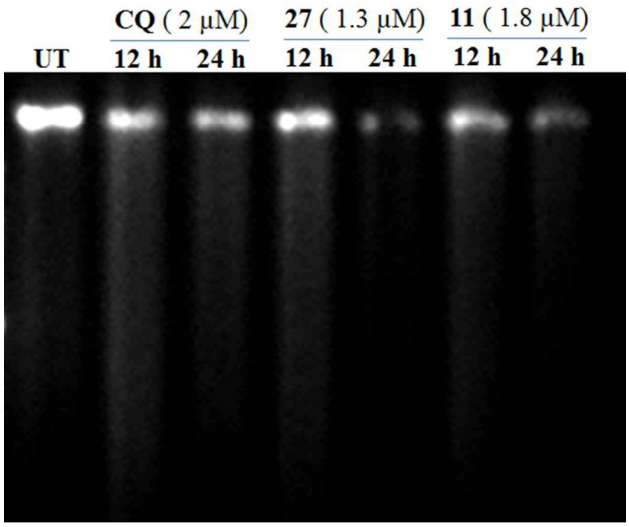

Figure 4. Detection of DNA fragmentation in 11 and 27 treated *Pf*Indo by Agarose Gel Electrophoresis: Early trophozoite stage cultures were treated with indicated concentrations of CQ (+ control), 11 and 27 for 12 and 24 hrs. The DNA isolated from parasites was subjected to electrophoretic separation. Note the strong intensity of genomic DNA band in untreated (UT) culture. Drug treated cultures band intensities of genomic DNA is diminished with simultaneous appearance of low MW DNA fragments.

unicellular organisms like Leishmania, Plasmodium, yeast, bacteria, blastocystis, Trypanosoma, and Trichomonas.[22,23] Our observation of apoptotic death in stilbenoid treated *Plasmodium falciparum* represents a new avenue of targeting a sensitive niche of the malaria parasite with a new pharmacophore. The high selectivity with which **11** and **27** have been found to target the blood stage malaria parasite but not the mammalian cells like the HeLa and L929 suggests that the apoptotic machinery of Plasmodium, a primitive protozoan, may be characteristically different from the corresponding machinery in the highly evolved mammalian cells. It is worth noting that CQ has also been found to inflict apoptotic death in malaria parasite.[24,25] Moreover the report of a putative *P. falciparum*metacaspase (*Pf* MCA-1)[25,26] and apoptotic features in *P. bergheiookinetes*[27,28] endorse the presence of apoptotic machinery in the malaria parasite. Thus apoptosis machinery of the malaria parasite appears to be a valid target worthy of attack by novel pharmacophores like the stilbenoids in the present study.

Conclusions

In conclusion, the first systematic antiplasmodialevaluation of a library of hydroxy substituted monomeric and oligomericstilbenoids (stilbenes, distyrylbenzenes and tristyrylbenzene) was conducted. Importantly, the above study led to the introduction of distyrylbenzenes and tristyrylbenzene as a novel class of potent antiplasmodial scaffolds. Compound **11**, a dimeric form of hydroxyl stilbene {IC_{50}:*Pf* 3D7 0.9 μM, *Pf*Dd2 2.0 μM, *Pf*Indo 2.7 μM, selectivity Index>111 (HeLa and L929)} remarkably shows /high antiplasmodial potency and high selectivity index. Likewise, compound **27**: (2*E*)-1-{4-[(*E*)-2-(4-hydroxy-3,5-dimethoxy-phenyl) ethenyl] phenyl-}-3-(2,4,5-trimethoxy-phenyl)prop-2-en-1-one, displayed highly promising antiplasmodial activity (IC_{50} : *Pf* 3D7 0.62 μM, *Pf* Dd2 0.5 μM, *Pf* Indo 1.36 μM) as well as good selectivity indices of 40.3 (HeLa) and 29 (L929). Further mechanistic investigations have revealed that distyrylbenzene (**11**) and octupolarstilbenoid (**27**) trigger selective apoptotic cell death in malaria parasite.

Experimental Section

Materials & Instruments: All the starting materials were reagent grade. The palladium catalyst was purchased from Aldrich and used as such. The substituted benzaldehydes, haloarenes, 4-bromophenylacetic acid and all other reagents were obtained from commercial sources (Merck and Aldrich). The solvents used for isolation/purification of compounds were obtained from commercial sources (Merck) and used without further purification. Column chromatography was performed using silica gel (Merck, 60-120 mesh size). The chromatographic solvents are mentioned as volume: volume ratios. ^{1}H (300 MHz) and ^{13}C (75.4 MHz) NMR spectra were recorded on a Bruker Avance-300 spectrometer. The following abbreviations have been used to designate chemical shift multiplicities: s = singlet, d = doublet, t = triplet, m = multiplet. The ^{13}C NMR spectra are proton decoupled. The melting points were determined on a digital BarnstedElectrothermal 9100 apparatus and are uncorrected. HRMS-ESI spectra were determined using micromass Q-TOF ultima spectrometer and reported as m/z (relative intensity). A CEM Discover® focused microwave oven (2450 MHz, 300 W) was used for reactions.

Representative procedure for the synthesis of (*E*)-4,4'-(ethene-1,2-diyl)diphenol(1)via Knoevenagel condensation-double decarboxylation-Heck coupling reaction (Table 1)

Malonic acid (0.64 g, 6.15 mmol) was taken in a round bottom flask and piperidine (0.45 ml, 4.6 mmol) added gradually. The above mixture was stirred in DMF (15 ml) for 2 min. at room temperature. Thereafter, 4-hydroxybenzaldehyde (1a, 1.51 mmol), 4-iodophenol (0.2 g, 0.90 mmol), $Pd(PPh_3)_4$ (0.025 mmol), piperidine (3.1 mmol) and LiCl (0.07 mmol) were added, and the reaction mixture allowed to reflux for 10 h. The above mixture was cooled to room temperature and filtered through celite and washed with ethyl acetate. The filtrate was poured into water (100 ml, acidified with dilHCl, pH= 5-6) and extracted with ethyl acetate (2x40 ml). The combined organic layer was washed with water (1x30 ml), brine (1x10 ml), dried over Na_2SO_4 and vacuum evaporated. The obtained residue was subsequently purified by column chromatography on silica gel (60-120 mesh size) using hexane: ethyl acetate (9.5: 0.5) to give a solid which was further recrystallized in methanol to provide pure 4,4'-dihydroxystilbene (**1**, Table 1).

(*E*)-4,4'-(ethene-1,2-diyl)diphenol (1) [29]

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White solid (33% yield), m.p.200-202°C, ^{1}H NMR δ (CD_3COCD_3, 300 MHz), 8.54 (2H, s), 7.39 (4H, d, *J* = 8.7 Hz), 6.96 (2H, s), 6.82 (4H, d, *J* = 8.7 Hz); ^{13}C NMR δ (75.4 MHz, CD_3COCD_3), 157.3, 129.9, 127.8, 125.9 and 115.9.

The same procedure was also followed for synthesis of other hydroxylatedstilbenes including **2-8** (Table 1).

Representative procedure for the synthesis of (*E*)-4-(4-bromostyryl)-2-methoxyphenol(9, Table 1) *via* modified Perkin condensation-decarboxylation reaction

A stirred mixture of 4-bromophenylacetic acid (3.6 mmol), methylimidazole (4.9 mmol), piperidine (4.9 mmol)and 4-hydroxy-3-methoxybenzaldehyde (0.5 g, 3.28 mmol) in polyethylene glycol-200 (3-4 mL) was irradiated under focused monomode microwave (150 W, 160°C) fitted with reflux condenser for 25 min. After the completion of reaction, the reaction mixture was cooled and acidified with dilHCl (pH=5). Then the aqueous layer was extracted with ethyl acetate (2x20 mL) and the organic layer was dried over sodium sulfate, vacuum distilled to obtain crude product which was further purified by column chromatography using sillica-gel (60-120 mesh size) with a 0.5: 9.5 mixture of ethyl acetate : hexane to give the pure stilbene (**9**).

(*E*)-4-(4-bromostyryl)-2-methoxyphenol(9)[30]

Cream solid (42% yield),m.p.120-122°C, ^{1}H NMR δ ($CDCl_3$, 300 MHz), 7.50 (2H, d, *J*= 8.5 Hz), 7.38 (2H, d, *J*= 8.5 Hz), 7.10-7.02 (3H, m), 6.95 (2H, d, *J*= 8.3 Hz), 5.75 (1H, s), 3.97 (3H, s); ^{13}C NMR δ (75.4 MHz, $CDCl_3$), 147.1, 146.2, 136.9, 132.1, 130.0, 129.7, 128.1, 125.6, 121.2, 121.0, 115.0, 108.6 and 56.3.

The above procedure was also followed for synthesis of 4-Bromo-4'-hydroxy-3',5'-dimethoxystilbene (**10**, Table 1)

Representative procedure for the synthesis of 4,4'-((1*E*,1'*E*)-1,4-phenylenebis(ethene-2,1-diyl))diphenol11, *via*Knoevenagel condensation-double decarboxylation-double Heck coupling reaction (Table 1)

Malonic acid (21.6 mmol) was taken in a round bottom flask and piperidine (16.45 mmol) added gradually. The above mixture was stirred in DMF (25 ml), for 2 min. at room temperature. Thereafter, 4-hydroxybenzaldehyde (2.7 mmol), 1,4-diiodobenzene (0.25 g, 0.755 mmol), $Pd(PPh_3)_4$ (0.045 mmol), piperidine (10.55 mmol) and LiCl (0.12 mmol) were added and the reaction mixture allowed to reflux for 14 h. The above mixture was cooled to room temperature and filtered through celite. The filtrate was poured into water (250 ml, acidified with dilHCl, pH= 5) and extracted with ethyl acetate (3x50 ml). The combined organic layer was washed with water (1x50 ml), brine (1x20 ml), dried over Na_2SO_4 and vacuum evaporated. The resultant residue was subsequently purified by column chromatography on silica gel (60-120 mesh size) using hexane: ethyl acetate (6:4) and obtained solid was recrystallized with methanol to provide pure product (**11**).

4,4'-((1*E*,1'*E*)-1,4-phenylenebis(ethene-2,1-diyl))diphenol (11)[31]

Grey solid (61% yield), m.p. 362-366°C (lit. m.p. 310-312°C), ^{1}H NMR (DMSO, 300 MHz), δ (ppm) 9.68 (2H, s), 7.51-7.45 (8H, m), 7.13-7.04 (4H, m), 6.78 (4H, s); ^{13}C NMR (75.4 MHz, DMSO), δ (ppm) 157.7, 136.7, 128.6, 128.5, 128.3, 126.8, 125.3, 116.0.

The above procedure was also followed for synthesis of other symmetrical distyrylbenzenes**12-19**(Table 1)

Representative procedure for thesynthesisof4-((*E*)-4-((*E*)-4-hydroxystyryl)styryl)benzene-1,2-diol (20, Table 1) from 3,4-dihydroxy benzaldehyde

To a stirredmixtureof 4-Bromophenylacetic acid (1.5 g, 7.0 mmol), methylimidazole (0.46 ml, 5.8 mmol) andpiperidine (0.62 ml, 6.30 mmol) in PEG-200 (5 ml), 3,4-dihydroxybenzaldehyde (3.94 mmol) wasaddedandthereactionmixtureirradiatedundermicrowave (180 W, 150°C) for 20 min. Thereafter, a mixtureofmalonicacid (4.08 g, 39.2 mmol), piperidine (3.34 ml, 39.2 mmol), 4-hydroxybenzaldehyde (1.2 g, 9.82 mmol), $Pd(PPh_3)_4$ (0.136 g, 0.12 mmol), K_2CO_3 (0.54 g, 3.9 mmol, LiCl (0.014 g, 0.32 mmol) in DMF (10 ml) wasaddedtotheabovepotandirradiatedundermicrowave (180 W, 150°C) for 45 min. The reaction mixture was cooled to room temperature and filtered through celite and washed with little amount of ethylacetate. The filtrate was poured into water (150 ml), acidified with dilHCl, (pH = 5-6) and extracted with ethyl acetate (2x40 ml). The combined organic layer was washed with water (2x15 ml), brine (1x10 ml), dried over Na_2SO_4 and vacuum evaporated. The residue was subsequently purified by column chromatography on silica gel (60-120 mesh size) using hexane: ethylacetate (9.4: 0.6) to give product (**20**).

4-((*E*)-4-((*E*)-4-hydroxystyryl)styryl)benzene-1,2-diol (20)[32]

Green solid (40% yield), m.p. 293-295°C, ^{1}H NMR (CD_3COCD_3: DMSO-d_6 (7:3)300 MHz), δ (ppm) 7.98 (4H, s), 7.91 (2H, d, *J* = 8.3 Hz), 7.60 (2H, d, *J* = 18.1 Hz), 7.54-7.53 (1H, m), 7.51 (2H, d, *J* = 18.1 Hz), 7.35-7.34 (1H, m), 7.30 (3H, d, *J* = 8.7 Hz); ^{13}C NMR (75.4 MHz, CD_3COCD_3: DMSO-d_6 (7:3)), δ (ppm) 158.3, 146.5, 146.3, 137.2, 135.5, 129.7, 129.2, 129.0, 128.8, 128.4, 127.0, 126.7, 125.5, 125.4, 119.3, 116.2, 115.7 and 113.9.

The above procedure was also followed for synthesis of other unsymmetrical DSB's **21-26** (Table 1).

Representative procedure for the one pot synthesis of 4,4',4''-((1*E*,1'*E*,1''*E*)-benzene-1,3,5-triyltris(ethene-2,1-diyl))tris(2,6-dimethoxyphenol)(27, Table 1)

Malonic acid (16.05 g, 154.2 mmol) was taken in a round bottom flask and piperidine (12.75 ml, 128.7 mmol) added gradually. The above mixture was stirred in DMF (30 ml) for 2 min at room temperature. Thereafter, 4-hydroxy-3,5-dimethoxybenzaldehyde (2.36 g, 12.96 mmol), 1,3,5-tribomobenzene (0.75 g, 2.38 mmol), $Pd(PPh_3)_4$ (0.247 g, 0.21 mmol), piperidine (6.37 ml, 63.7 mmol) and LiCl (0.025 g, 0.058 mmol) were added, then reaction mixture allowed to reflux for 16h. The above mixture was cooled to room temperature and filtered through celite. The filtrate was poured into water (250 ml, acidified with dil. HCl, pH= 5) and extracted with ethyl acetate (3x50 ml). The combined organic layer was washed with water (1x50 ml), brine (1x20 ml), dried

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over Na_2SO_4 and vacuum evaporated. The residue was subsequently purified by column chromatography on silica gel (60-120 mesh size) using hexane: ethyl acetate (6:4) and obtained solid was washed with methanol to provide pure ((E,E,E)-1,3,5-Tris(4-hydroxy-3,5-dimethoxy)styrylbenz-ene (**27**).

4,4',4''-((1*E*,1'*E*,1''*E*)-benzene-1,3,5-triyltris(ethene-2,1-diyl))tris(2,6-dimethoxyphenol) (27) [9]

Yellow solid (30% yield), m.p. 223-225°C, ^{1}H NMR ($CDCl_3$, 300 MHz), δ (ppm) 7.53 (3H, s), 7.17 (3H, d, J = 15.7 Hz), 7.04 (3H, d, J = 15.7 Hz), 6.81 (6H, s), 5.62 (3H, s), 3.98 (18H, s); ^{13}C NMR (75.4 MHz, $CDCl_3$), δ (ppm) 147.4, 138.3, 135.1, 129.4, 129.0, 126.7, 123.4, 103.6 and 56.5. HRMS-ESI: m/z $[M+H]^+$ for $C_{36}H_{36}O_9$, calculated 613.2432; observed 613.2432.

Representative procedure for the synthesis of 3,3'',4,4'',5,5''-hexamethoxy-5'-(3,4,5-trimethoxyphenyl)-1,1':3',1''-terphenyl(28, Table 1)

To a stirred mixture of 1,3,5-tribromobenzene (0.95 mmol) in dioxane: water (5:1, 10ml), 3,4,5-trimethoxyphenylboronic acid (3.42 mmol), $Pd(PPh_3)_4$ (0.085 mmol), K_2CO_3 (1.4 mmol) were added and the reaction mixture was irradiated under MW (250W, 115°C) for 35 min. The above mixture was cooled to room temperature and was poured into water (250 ml, acidified with dil. HCl, pH= 5) and extracted with ethyl acetate (3x50 ml). The combined organic layer was washed with water (1x50 ml), brine (1x20 ml), dried over Na_2SO_4 and vacuum evaporated. The resulting residue was subsequently purified by column chromatography on silica gel (60-120 mesh size) using hexane: ethyl acetate (9:1) to provide pure 1,3,5-tris(3,4,5-trimethoxyphenyl)benzene (**28**).

3,3'',4,4'',5,5''-hexamethoxy-5'-(3,4,5-trimethoxyphenyl)-1,1':3',1''-terphenyl(28)

White solid (45% yield), m.p. 278-280°C, ^{1}H NMR ($CDCl_3$, 300 MHz), δ (ppm) 7.68 (3H, s), 6.87 (6H, s), 3.96 (27H, s); ^{13}C NMR (75.4 MHz, $CDCl_3$), δ (ppm) 154.0, 143.1, 138.4, 137.5, 125.7, 105.1, 61.4 and 56.7. HRMS-ESI: m/z [M+H]+ for $C_{33}H_{36}O_9$, calculated 577.2432; observed 577.2456.

Measurement of inhibition of *P. falciparum* growth in culture: In this study, chloroquine sensitive 3D7 and chloroquine resistant Dd2 and INDO strains of *P. falciparum* were cultivated in vitro by the method of Trager and Jensen[33]with minor modifications. Cultures were maintained in fresh O+ human erythrocytes at 4% hematocrit in complete medium (RPMI 1640 with 0.2% sodium bicarbonate, 0.5% Albumax, 45 mgL-1 hypoxanthine and 50 mgL-1 gentamicin) at 37°C under reduced O_2 (gas mixture 5% O_2, 5% CO2 , and 90% N_2). Stock solutions of chloroquine were prepared in water (milli Q grade) and test compounds were dissolved in DMSO. All stocks were then diluted with culture medium to achieve the required concentrations (in all cases the final concentration contained 0.4% DMSO, which was found to be non-toxic to the parasite). Drugs and test compounds were then placed in 96-well flat-bottom tissue culture grade plates to yield triplicate wells with drug concentrations ranging from 0 to 100 μM in a final well volume of 100 μL. Chloroquine was used as a positive control in experiments (100 nM with 3D7 and 1000 nM with the chloroquine resistant strains). Parasite culture was synchronized at ring stage with 5% sorbitol. Synchronized culture was aliquoted to a drug containing 96-well plates at 2% hematocritand 1% parasitemia. After 48 hrs of incubation under standard culture conditions, plates were harvested and read by the SYBR Green I fluorescence-based method[15] using a 96-well fluorescence plate reader (Victor, Perkin Elmer), with excitation and emission wavelengths at 485 nm and 530 nm, respectively. The fluorescence readings were plotted against drug concentration, and IC_{50} values obtained by visual matching of the drug concentration giving 50% inhibition of growth. In view of the fluorescence basis of the SYBR Green assay, it was important to assess artefacts due to autofluorescence or quenching effects of each test molecule. To measure the auto fluorescence of test molecules, the parasites were treated with 100 μM of all test molecules and incubated for 1 hr at 37 °C following which the cultures were lysed by lysis buffer {20 mMTris; 5 mM EDTA; 0.008 %(w/v) Saponin; 0.08% (v/v) Triton x, pH 7.5} and read at 485/530 nm (excitation/emission). To determine possible quenching effects, untreated parasite cultures or parasite cultures treated with the test molecules (100 μM) were lysed by 1X SYBR Green I containing lysis buffer and read for their fluorescence values at 485/530 nm (excitation/emission). Comparison of fluorescence counts (+/- test molecule) was used as a measure of quenching or lack of quenching.

Measurement of cytotoxic activity against mammalian cell lines in culture: Animal cell lines (HeLa and fibroblast L929) were used to determine drug toxicity by using MTT assay for mammalian cell viability assay as described by Mosmann[34] using HeLa and fibroblast L929 cells cultured in complete RPMI containing 10% fetal bovine serum, 0.2% sodium bicarbonate and 50 μg ml-1 gentamicin. Briefly, cells (104cells/200 μL/well) were seeded into 96-well flat-bottom tissue-culture plates in complete culture medium. Drug solutions (in all cases the final concentration contained 0.4% DMSO) were added after overnight seeding and incubated for 24 h in a humidified atmosphere at 37◦C and 5% CO_2. DMSO (final concentration 10%) was added as +ve control. An aliquot of a stock solution of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) (5 mg mL-1 in 1X phosphate buffered saline) was added at 20 μL per well, and incubated for another 4 h. After spinning the plate at 1500 RPM for 5 min, supernatant was removed and 100 μL of the stop agent DMSO was added to each well. Formation of formazon, an index of growth, was read at 570 nm and IC_{50} values were determined by analysis of dose–response curves. Selectivity index was calculated as IC_{50} mammalian cell/IC50 *Pf* 3D7.

Microscopic evaluation of morphological changes in *P. falciparum*: Morphological changes of *P. falciparum* were monitored by microscopy based Giemsa staining method. Briefly, synchronized ring and trophozoites stage cultures (1% parasitemia, 2% hematocrit) were incubated with $IC_{90}$of 27 (1.3 μM) and 11 (1.8 μM) for 12, 24, 48 (rings) and 6, 12, 24 hrs (trophozoites) in wells of 96 well plate. Thin blood smears were prepared from treated and untreated cultures, methanol fixed,

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stained by Giemsa (Sigma, India) and examined by bright field optical microscopy at 100X.

Detection of DNA fragmentation and Chromatin condensation by Hoechst 33342: DNA fragmentation and condensation were detected by Hoechst 33342 (2′-[4-ethoxyphenyl]-5-[4-methyl-1-piperazinyl]-2,5′-bis-1H benzimidazoletrihydrochloridetrihydrate, Molecular Probes, USA). Hoechst 33342 stains the condensed chromatin of apoptotic cells far more brightly than is the case with native chromatin of live healthy cells. The assay was performed according to the manufacturer's instruction. Briefly, synchronized ring and trophozoite stage cultures (1% parasitemia and 2% hematocrit) were incubated with IC_{90} of **27** (1.3 μM) and **11** (1.8 μM) for 12, 24, 48 (rings) and 6, 12, 24 hrs (trophozoites) in wells of 96 well micro titer plate. Cultures were incubated with Hoechst 33342 stain (λmax emission 460 nm) for 20 min at ice temperature. Following transfer of cells to microfuge tubes, the cells were centrifugally washed (200 μL, twice) with PBS and wet mount slides were prepared. The slides were observed by using a fluorescence microscope (Nikon 50i).

Detection of *P. falciparum* mitochondrial trans-membrane potential (ΔΨm): Changes of *P. falciparum* mitochondrial membranepotential (ΔΨm) were detected by staining with the fluorescent cell-permeable cationic carbocyanine dye JC-1 (5,5′,6,6′-tetrachloro-1,1′,3,3′-tetraethylbenzimidazolylcarbocyanine iodine) (Molecular Probes, USA). JC-1 shows membrane potential dependent transition from a green (λmax emission 525 nm) monomeric form (at low transmembrane potential) to a red (λ max emission 590 nm) aggregated oligomeric form (at higher trans - membrane potential). The assay was performed according to the manufacturer's instructions. Briefly, synchronized ring and trophozoites stage cultures (1% parasitemia and 2% hematocrit) were incubated with $IC_{90}$of 27 (1.3 μM) and 11 (1.8 μM) for 12, 24, 48 (rings) and 6, 12, 24 hrs (trophozoites) in wells of 96 well plate. The cultures were incubated with JC-1 dye for 20 min at 37 °C and washed with PBS. Wet mount slides were prepared and observed by using afluorescence microscope (Nikon 50i) using FITC-TRITC dual band excitation filters.

Detection of drug induced DNA fragmentation by agarose gel electrophoresis: Drug induced DNA fragmentation was examined by agarose gel electrophoresis. Early trophozoite stage cultures (4% parasitemia) of *Plasmodium falciparum* (Indo) were treated with $IC_{90}$of **27** (1.3 μM) and **11** (1.8 μM) for 12 and 24 hrs. **CQ** (2 μM) was used as positive control. After treatment, the parasites were isolated by the 0.05 % saponin treatment procedure, genomic DNA was isolated and samples were analysed by 1 % agarose gel electrophoresis method[35]. The gels were photographed using Alpha Imager EC.

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