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1-Aryl-1,2,3,4-tetrahydroisoquinolines as potential antimalarials: synthesis, in vitro antiplasmodial activity and in silico pharmacokinetics evaluation

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In the present study, twenty one 1-aryl-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (THIQ) analogues were synthesized by based-catalyzed Pictet-Spengler reaction, and tested in vitro against P. falciparum using the [3H]hypoxanthine incorporation assay. Two compounds were found to be inactive while seventeen compounds displayed moderate antiplasmodial activity and two compounds were found to be highly active (IC50 < 0.2 µg/ml). The two highly active compounds, 1-(4-chlorophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline and 6-hydroxyspiro[1,2,3,4-tetrahydroisoquinoline-1:1'-cyclohexane], also displayed low cytotoxicity, against rat skeletal myoblast cells, with CC50 values of 257.6 and 174.2 µM respectively. These results justify further investigation of simple 1-aryl-1,2,3,4-tetrahydroisoquinolines as potential anti-malarial agents.

Introduction

Malaria is a life threatening infectious parasitic disease that inflicts both health and economic losses. 1 The disease is transmitted by mosquitoes of the genus Anopheles. The causative agent for malaria is Plasmodium species, and there are four types of human malaria: P. vivax, P. malaria, P. ovale and P. falciparum. P. vivax and P. falciparum are the most common, and P. falciparum is responsible for the most deadly type of malaria infection. 2 Statistics indicate that P. falciparum causes more than 300 to 500 million clinical cases of malaria infection, resulting in 1.7 to 2.5 million deaths annually. 3 Malaria is a particularly important disease in Sub-Saharan Africa where about 90% of cases and death occur, but is also a serious public health burden in South, East and Central Asia, South and Central America, Caribbean, Oceania, and Middle East regions. 4 Many attempts have been made to control the disease by using vector control measures and/or chemoprophylaxis, but they have had limited success. 5 The control of malaria is, among other factors, hampered by the emergence of resistance of the vector (Anopheles sp.) to dichlorodiphenyltrichloroethane (DDT) and other insecticides, and the ongoing spread of strains of P. falciparum that are resistant to currently used anti-malarial drugs. 6 Attempts to address the problem of resistance have resulted in the current shift to artemisinin based combination drug regimens. New anti-malarial drugs, especially those with new structures and novel modes of action against malaria are also needed. This work describes the synthesis of structurally simple tetrahydroisoquinolines (THIQs) and their preliminary evaluation as potential anti-malarial agents. THIQs abound in nature and have attracted significant attention due to their diverse biological activities. The THIQ substructure (Fig. 1) is found in many drugs and alkaloids that exhibit antitumor, cardiovascular, antibacterial, antimicrobial, antitubercular, antifungal, antileishmanial, antitrypanosomal and antiplasmodial activities. 7-20 These compounds include the naphthylisoquinolines, benzylisoquinolines and bisbenzylisoquinolines.

The broad spectrum of biological activities of tetrahydroisoquinolines, especially reports on the anti-human immunodeficiency virus (HIV) activity of Michellamine B (1) and antiplasmodial activity of Dioncophylline C (3), naphthylisoquinolines (Fig. 2) isolated from Ancistrocladus korupensis, 11 prompted us to investigate 1-aryl-1,2,3,4-tetrahydroisoquinolines for antiplasmodial activity. Naphthylisoquinoline alkaloids are known to exhibit curative potentials against malaria. 12-14 Dioncophylline E (5) has exhibited good antimalarial activity against both chloroquine-sensitive and -resistant strains of P. falciparum in vitro, 15 while dioncophylline C was active against the chloroquine-resistant P. berghei Anka CRS parasites 12 with dioncophylline B (2), dioncophylline D (4) and synthetic naphthylisoquinolines also found to display good in vitro antiplasmodial activities. 14 A noteworthy example is the spirofused compound IVa, which was previously described by Kametani et al., 21 as an abbreviated analogue of erysodine, one of member of the Erythrina class of alkaloids. Plants of the genus Erythrina have been used extensively in ethnomedicine for a number of applications.
including the treatment of liver disorders,\textsuperscript{20} possess antiplasmodial, analgesic, and anti-inflammatory activity,\textsuperscript{21} as well as in the treatment of diseases such as female infertility, stomach pain and gonorrhea.\textsuperscript{22} Moreover, the alkaloids of \textit{Erythrina} type are characterized by their unique skeleton of a tetracyclic spiroamine.\textsuperscript{23} They are classified in two main groups: alkaloids possessing a skeleton of a 6,5,6,6 indoloisoquinoline called erthryrinanes and those with a skeleton of 6,5,7-6 indolobenzazepine called schlehammerans or homoeorthyrrhane alkaloids (Fig. 3). \textit{Erythrina} alkaloids are known to display antiplasmodial activity \textit{in vitro}.\textsuperscript{24} The rationale behind the design of THIQ analogues as potential antimalarials is that they seem to be the common denominator of the naturally occurring naphthoisocoumarinones (1 to 5) which display antimalarial activities (marked in red on Fig. 2). It could therefore be easily inferred that the THIQ scaffold could contain the pharmacophore responsible for the anti-malarial activity. Previous studies have revealed that even simple THIQs such as 1-aryl-6,7-dihydroxy-1,2,3,4-tetrahydroisoquinolines and 1-aryl-6,7-dimethoxy-1,2,3,4-tetrahydroisoquinolines, display anti-HIV\textsuperscript{15} and bronchodilator\textsuperscript{16} properties. The current study, which forms part of a larger investigation, was undertaken to investigate the antiplasmodial activity of 1-aryl-1,2,3,4-tetrahydroisoquinolines. The THIQ skeleton presents eight positions that can accommodate substituents. The choice of positions 1 and 6 for this study was based on the ease of synthesis of 1-substituted-THIQs (which is offered by the Pictet Spengler) and the presence of alkoy groups on the aromatic portion of complex THIQs like the naphthoisocoumarinones, such as 2-S, which display antiplasmodial activity.

Designing compounds with the THIQ moiety would therefore considerably lower the molecular masses and hence derive analogues with properties that fall within the range of those of ‘drug-like’ compounds. The target 6-hydroxy-1,2,3,4-tetrahydroisoquinoline derivatives (IIa to IIIu and IVa) were synthesized via base-catalyzed Pictet-Spengler cyclization as previously reported.\textsuperscript{19} Overall yields over two steps ranged between 4 and 99%. The compounds were characterized by NMR and mass spectrometry. The corresponding hydrochlorides were subsequently tested \textit{in vitro} for antiplasmodial activity against \textit{P. falciparum}. An \textit{in silico} assessment of drug metabolism and pharmacokinetics (DMPK) profiles was also carried out by the use of computed molecular descriptors related to the absorption, distribution, metabolism, excretion and toxicity (ADMET) of drugs.

\textbf{Results and discussion}

\textbf{Antiplasmodial activity versus cytotoxicity}

All compounds were tested \textit{in vitro} against \textit{P. falciparum} as described in the experimental section. The biological data were plotted to obtain IC\textsubscript{50} values which represent the concentration required to cause 50% inhibition of parasite growth. Activity criteria were as follows: inactive, IC\textsubscript{50} > 5 \textmu g/mL; moderately active, 0.5 \textmu g/mL < IC\textsubscript{50} < 5 \textmu g/mL; highly active, IC\textsubscript{50} < 0.5 \textmu g/mL. The antiplasmodial activity, presented as IC\textsubscript{50} values, is provided in Table 1. The \textit{P. falciparum} strain K1, known to be resistant to Chloroquine and Pyrimethamine, was employed for this assay. Chloroquine was used as the reference drug. Of twenty-one compounds tested, three could be described as inactive following the criteria established. The rest were either moderately active (twelve) or active (seven). Among the active compounds at least three (IIb, IIIk and IVa) displayed antiplasmodial activity comparable to that of chloroquine, suggesting the need for further evaluation of these compounds \textit{in vivo}. In general, the more active compounds tended to possess electron withdrawing substituents on the pendant phenyl group (IIIb-c,j,k,q). However, a number of moderately active compounds contained electron donating substituents on the same phenyl group, thereby obscuring the role of electronic substituent parameters as determinants of antiplasmodial activity in this series. The present demonstration of antiplasmodial activity in the compound would therefore suggest that the key pharmacophoric elements for antiplasmodial activity in this class of alkaloids reside in the skeleton as described by IVa.\textsuperscript{19}

When tested for cytotoxicity, several of the compounds were found to be considerably less toxic against rat skeletal mycoblast cells than the reference compound podophyllotoxin. In addition, the tested compounds displayed toxicity at concentrations that were two orders of magnitude higher than the concentrations at which they displayed antiplasmodial activity. Therefore, structurally simplified 1,2,3,4-tetrahydroisoquinolines may be potentially useful as antiplasmodial agents.

Compounds IIIb, IIIk and IVa were the most active with IC\textsubscript{50} values 0.70, 0.76 and 0.89 \textmu M, respectively, and their activity was in the same range as that of chloroquine. In addition, the cytotoxicity of these compounds against mycoblast cells could be considered as an index of drug effectiveness. Compounds IIIb, IIIk and IVa had selective index values 369.6, 229.1 and > 471.5 respectively. These values are up to tenfold lower than the Selectivity Index value reported for chloroquine but significantly higher than the value of 100 which is required to qualify a compound as a “hit” in the screening of compounds for antiplasmodial activity.

The cytotoxicity of compounds IIa to IIIu and IVa was evaluated on rat myoblast cells grown as described in experimental section. Table 1 shows the results obtained from the assay. Most of the compounds exhibited antiplasmodial activity, with IC\textsubscript{50} values ranging from 0.70 to 20 \textmu M. Two compounds, III and IIIa, were considered inactive (IC\textsubscript{50} > 20 \textmu M). Compound IIIb, IIIk and IVa emerged as the most active analogues with IC\textsubscript{50} values within the range below 2 \textmu M, their activity being close to that of chloroquine. At 0.19 \textmu M, IIIa was moderately active, but the presence of chlorine at C-4 significantly increases, maybe due to the electron withdrawing property by inductive effect and \textit{ortho} and \textit{para} directive effect. This activity is not maintained by IIIc, IIId, IIIe, IIIf, IIIm, IIIn, IIIo, IIIp and IIIq. When a methoxy group in (electron donating by resonance) is introduced in compound IIIa to get IIIb, only leads to a slight drop in activity with the introduction of the methoxy group. When further methoxy groups are introduced, (IIId, IIIf, IIIi) the activity drops even further. We could conclude that when both electron withdrawing and electron donating groups were introduced in the molecule, the activity decreased even further. On the contrary, when a phenyl group, a cyclohexyl group is introduced instead (as in IVa), the anti-malarial activity significantly increases, maybe because of the rigidity introduced in this analogue.
In silico pharmacokinetics profiles

Molecular modelling was carried out on all synthesised molecules as described in the experimental section. The computed molecular descriptors are those related to the prediction of DMPK/ADMET properties like the n-octanol/water partition coefficient, aqueous solubility, brain/blood partition coefficient, Caco-2 cell permeability, serum protein binding, number of likely metabolic reactions, solvent accessible surface area, blockage of human-ether-a-go-go potassium ion (HERG K⁺) channels and the total volume of the molecule enclosed by the solvent accessible surface area. The bioavailability of a compound depends on the processes of absorption and liver first-pass metabolism. 27 Absorption in turn depends on the solubility and permeability of the compound, as well as interactions with transporters and metabolizing enzymes in the gut wall. The computed parameters used to assess oral absorption are the predicted aqueous solubility, log $S_{\text{wat}}$, the conformation-independent predicted aqueous solubility, $\text{CI log } S_{\text{wat}}$, the predicted qualitative human oral absorption, the predicted % human oral absorption and compliance to predict BBB penetration. 28 Absorption in turn depends on the solubility and permeability of the compound, as well as interactions with transporters and metabolizing enzymes in the gut wall. The computed parameters used to assess oral absorption are the predicted aqueous solubility, log $S_{\text{wat}}$, the conformation-independent predicted aqueous solubility, $\text{CI log } S_{\text{wat}}$, the predicted qualitative human oral absorption, the predicted % human oral absorption and compliance to Lipinski’s famous “Rule of Five” (ro5).29 The size of a molecule, as well as its capacity to make hydrogen bonds, its overall lipophilicity and its shape and flexibility are important properties to consider when determining permeability. Molecular flexibility has been seen as a parameter which is dependent on the number of rotatable bonds (NRB), a property which influences bioavailability in rats.29 The blood/brain partition coefficients (logB/B) were computed and used as a predictor for access to the central nervous system (CNS). Madin-Darby canine kidney (MDCK) monolayers, are widely used to make oral absorption estimates, the reason being that these cells also express transporter proteins, but only express very low levels of metabolizing enzymes. They are also used as an additional criterion to predict BBB penetration. Thus, our calculated apparent MDCK cell permeability could be considered to be a good mimic for the BBB (for non-active transport). The efficiency of a drug may be affected by the degree to which it binds to the proteins within blood plasma. Binding of drugs to plasma proteins (like human serum albumin, lipoprotein, glycoprotein, $\alpha$, $\beta$, and $\gamma$ globulins) greatly reduces the quantity of the drug in general blood circulation and hence the less bound a drug is, the more efficiently it can traverse cell membranes or diffuse. The predicted plasma-protein binding has been estimated by the prediction of binding to human serum albumin; the log $K_{\text{HSA}}$ parameter (recommended range is -1.5 to 1.5 for 95% of known drugs). Human ether-a-go-go related gene (HERG) encodes a potassium ion (K⁺) channel that is implicated in the fatal arrhythmia known as torsade de pointes or the long QT syndrome.30 The HERG K⁺ channel, which is best known for its contribution to the electrical activity of the heart that coordinates the heart’s beating, appears to be the molecular target responsible for the cardiac toxicity of a wide range of therapeutic drugs.31 HERG has also been associated with modulating the functions of some cells of the nervous system and with establishing and maintaining cancer-like features in leukemic cells.32 Thus, HERG K⁺ channel blockers are potentially toxic and the predicted IC₅₀ values often provide reasonable predictions for cardiac toxicity of drugs in the early stages of drug discovery.33

The results for fifteen (15) relevant descriptors have been shown in Table 2, while 2D scatter plots of selected molecular descriptors are shown in Fig. 4. It was observed that only the points of subfigure C fell outside the recommended range for the dipole moment parameter. Subfigures A and B showed that all the compounds complied with Lipinski’s ro5, since the molecular weights (MW) < 500 Da, logarithm of n-octanol/water partition coefficient (log P) < 5, number of hydrogen bond acceptors (accHB) < 10 and number of hydrogen bond donors (donHB) < 5. It was observed that the other computed molecular descriptors of all the THIQ analogues fell within the recommended range for 95% of known drugs (recommended parameters are shown in the ESI), except for two compounds (IIIo and IIIq), as can be seen in subfigures 4D-L, corresponding respectively to the scatter plots for MW against the number of rotatable bonds (#rotor), cohesion index in solids, blood/brain barrier partitioning, HERG K⁺ ion channel blockage, binding affinities to human serum albumin, skin permeability, aqueous solubilities, as well Caco-2 and MDCK permeabilities. As for compound IIIo, the dipole moment was computed to be 0.524 Debye, below the recommended range (1.0 to 12.5 Debye), while for compound IIIq, the computed value was 0.890 Debye, only slightly below the recommended minimum for 95% of known drugs. The weak dipole moments could be explained by the absence of polar groups in the pendant phenyl ring of the two aforementioned compounds. Dipole moments as high as > 6 Debye were observed for compounds IIIe, IIIr and IIIt containing the polarising nitro group in the pendant phenyl ring. It is however important to mention that the most active compounds (IIIb, IIIk and IVa) had interesting predicted pharmacokinetic profiles (with all computed parameters falling within the recommended range for 95% of known drugs, i.e., #stars = 0). We can therefore conclude that these three compounds could be subjected to further investigation as potential anti-malarials.

Experimental

General experimental procedures

The synthesis of organic compounds was carried out with commercially available chemicals that were used as received (scheme 1). All target compounds were tested in the form of the hydrochlorides which were obtained by dissolution of the corresponding free base in ethanolic HCl. Nuclear Magnetic Resonance spectroscopy data were recorded using UNITYplus – 500 spectrometer ($^1$H NMR at 500 MHz). Chemical shifts are denoted in unit parts per million (ppm) relative to the solvent ($^1$H NMR peak: 3.30 for CD₃OD). The following abbreviations are used to indicate peak multiplicities and characteristics: s (singlet), d (doublet), dd (doublet of doublets), t (triplet), q (quartet), m (multiplet). Mass spectra were recorded on a BIOTO FIEST mass spectrometer. TLC analyses were carried out on aluminum plates (Merck) precoated with silica gel 60 F254 (0.2mm thickness). Visualization of spots was performed with UV light or by treatment with iodine.

General procedure for the synthesis of 1-aryl-6-hydroxy-1,2,3,4-tetrahydroisoquinolines.

6-hydroxy-1-phenyl-1,2,3,4-tetrahydro-isoquinoline (IIIa).

The synthesis followed the procedure earlier described by Kametani et al.21 A mixture of 3-methoxyphenethylamine (2.0 g, 13 mmol), hydrobromic acid (10 mL) and acetic acid (10 mL) was stirred and heated under reflux for six (6) hours. The cooled mixture was concentrated under reduced pressure to obtain a residue. To this residue, benzaldehyde (1.96 g, 14.5
mmol), triethylamine (1 mL), and ethanol (10 mL) were added. The resulting mixture was stirred, heated under reflux for 10 hours and concentrated to remove the solvent. The residue obtained after evaporation was diluted with methylene chloride (100 mL) and water (100 mL) with resulting precipitation of solid. The solid was collected by filtration, washed with acetone (3x10 mL) and air dried to afford IIIa as a white solid (2.80mg, 10%); mp 260°C. 'H NMR (CDOD) δ 2.99-3.69 (m, 4, -(CH2)2), 4.89 (s, 1, OH), 6.89 (s, 1, -CH-N), 6.58-7.09 (m, 6, aryl). MS [M+H]+ 286.146 corresponding to (C17H18NO3).

1-(2,5-Dimethoxyphenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIb).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 2,5-dimethoxybenzaldehyde (2.5 g, 14.5 mmol)] (3.6 g, 98%); mp, 260°C. 'H NMR (CDOD) δ 3.06-3.46 (m, 4, -(CH2)2), 3.69-3.83 (s, 6, methoxyl), 4.89 (s, 1, OH), 5.82 (s, 1, -CH-N), 6.58-7.09 (m, 6, aryl). MS [M+H]+ 286.146 corresponding to (C17H18NO3).

1-(2,6-Dimethoxyphenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIc).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 2,6-dimethoxybenzaldehyde (2.81 g, 14.5 mmol), (3.7 g, 99%); mp, 260 – 266°C. 'H NMR (CDOD) δ 2.99-3.69 (m, 4, -(CH2)2), 3.79 (s, 6, methoxyl), 4.89 (s, 1, OH), 6.05 (s, 1, -CH-N), 6.42-7.46 (m, 6, aryl). MS [M+H]+ 286.146 corresponding to (C17H18NO3).

6-Hydroxyxipr1,2,3,4-tetrahydroisoquinoline-1:1′-cyclohexane (IVa).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and cyclohexanone (4 g, 14.5 mmol)], 250 mg (10%); mp, 240°C for the hydrochloride analogue. 'H NMR (CDOD) δ 4.9-6.0 (m, 10, cyclohexyl), 3.05-3.47 (t, 4, -(CH2)2), 4.89 (s, 1, OH), 6.60-7.26 (m, 3, aromatic protons). MS [M+H]+ 304.154 corresponding to (C18H18NO)2.

1-(4-Bromophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (III).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 4-bromobenzaldehyde (0.875g, 4.73mmol)], (0.68g, 52%). 'H NMR (CDOD) δ 3.09-3.52 (m, 4, -(CH2)2), 4.89 (s, 1, OH), 5.71 (s, 1, -CH-N), 6.20-7.59 (m, 7, aryl). MS [M+H]+ 286.146 corresponding to (C17H18NOBr).

1-(α,α,α-Trifluoromethylphenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIi).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and o,a,a-triflorotolualdehyde (0.63g, 3.63mmol)], 0.12 g (16%). 'H NMR (CDOD) δ 3.11-3.52 (m, 4, -(CH2)2), 4.90 (s, 1, OH), 5.80 (s, 1, -CH-N), 6.62-7.81 (m, 7, aryl). MS [M+H]+ 295.1108 corresponding to (C16H15NOF3).

1-(4-Biphenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (III).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 4-biphenylcarboxaldehyde (1.33g, 7.27 mmol)], (1.40g, 70%); mp, 201°C. 'H NMR (CDOD) δ 2.92-3.53 (m, 4, -(CH2)2), 3.92 (s, 1, OH), 5.07 (s, 1, -CH-N), 5.27-5.80 (m, 6, aryl). MS [M+H]+ 286.146 corresponding to (C17H18NO3).
1, -CH-N), 6.57-7.81 (m, 7, aryl). MS [M+H]^+ 304.0593 corresponding to C₁₅H₁₃NOBr.

\[\text{I-(2-Flourophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIa).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 2-flurobenzaldehyde (1g, 7.27mmol)], 1g (63%). \(^1\)H NMR (CD₃OD) δ 3.09-3.35 (m, 4, -CH₂), 4.89 (s, 1, OH), 5.72 (s, 1, -CH-N), 6.64-7.54 (m, 7, aryl). MS [M+H]^+ 304.0626 corresponding to C₁₅H₁₃NOBr.

\[\text{I-(3-Bromophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIo).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 3-bromobenzaldehyde (1.35g, 7.27mmol)], 1.1g (69%). \(^1\)H NMR (CD₃OD) δ 3.09-3.54 (m, 4, -CH₂), 4.89 (s, 1, OH), 5.75 (s, 1, -CH-N), 6.65-7.44 (m, 7, aryl). MS [M+H]^+ 304.1575 corresponding to C₁₅H₁₃NOBr.

\[\text{I-(3-Fluorophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIp).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 3-flurobenzaldehyde (1g, 7.27mmol)], 1.1g (69%). \(^1\)H NMR (CD₃OD) δ 3.07-3.52 (m, 4, -CH₂), 4.91 (s, 1, OH), 5.88 (s, 1, -CH-N), 6.69-8.03 (m, 6, aryl).

\[\text{I-(4-Fluorophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIq).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 4-flurobenzaldehyde (1g, 7.27mmol)], 1.2g (75%). \(^1\)H NMR (CD₃OD) δ 3.07-3.53 (m, 4, -CH₂), 4.92 (s, 1, OH), 5.73 (s, 1, -CH-N), 6.65-7.44 (m, 7, aryl). MS [M+H]^+ 304.1385 corresponding to C₁₅H₁₃NOBr.

\[\text{I-(4-Chloro-3-nitrophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIr).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 4-chloro-3-nitrobenzaldehyde (1.35g, 7.27mmol)], 0.7g (35%). \(^1\)H NMR (CD₃OD) δ 3.07-3.53 (m, 4, -CH₂), 4.91 (s, 1, OH), 5.88 (s, 1, -CH-N), 6.69-8.03 (m, 6, aryl).

\[\text{I-(3-Bromo-2-methylphenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIs).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 5-bromo-o-anisaldehyde (1.6g, 7.27 mmol)], 1.0g (46%). \(^1\)H NMR (CD₃OD) δ 3.10-3.45 (m, 4, -CH₂), 3.90 (s, 3, methoxyl), 4.90 (s, 1, OH), 5.88 (s, 1, -CH-N), 6.69-7.62 (m, 6, aryl).

\[\text{I-(2-Hydroxy-5-nitrophenyl)-6-hydroxy-1,2,3,4-tetrahydroisoquinoline (IIIi).}\]

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and 2-hydroxy-5-nitrobenzaldehyde (1.22g, 7.27 mmol)], 0.32g (17.7%). \(^1\)H NMR (CD₃OD) δ 3.07-3.52 (m, 4, -CH₂), 4.90 (s, 2, OH), 5.96 (s, 1, -CH-N), 6.68-8.27 (m, 6, aryl).

Yield [from 3-methoxyphenethylamine (2.0 g, 13 mmol) and p-tolualdehyde (0.9g, 7.27mmol)], 1.2g (76%). \(^1\)H NMR (CD₃OD) δ 2.37 (s, 3, methyl), 3.06-3.53 (m, 4, -CH₂), 4.92 (s, 1, OH), 5.63 (s, 1, -CH-N), 6.63-7.31 (m, 7, aryl).

\[\text{In vitro sensitivity assays of P. falciparum ([^1]H)-hypoxanthine incorporation).}\]

The test samples were assayed on a P. falciparum clone designated K1 which is chloroquine/pyrimethamine resistant. Test samples of 10 mg were prepared in dimethyl sulfoxide (DMSO). The testing procedure was as follows: A serial dilution factor of 1:2 was prepared with 100 µL of medium (RPMI 1640 without hypoxanthine 10.44 g/L; HEPES 5.94 g/L; Albumax® 5 g/L; Neomycin 10 mL/L (100U/ml); NaHCO₃ 50 g/L stock 42 mL/L (2.1 g/L). The final concentration was distributed into 96 wells of the Costar TM 96 – wells microtiter plates. To these 96 wells, 100 µL of medium, washed human red blood cells A+ (RBC) and P. falciparum mix, were added. The plates were incubated for 48 hours at 37°C in a chamber and gassed with a 4% CO₂, 3% O₂, 93% N₂. After 50 µL of medium and [^1]H)-hypoxanthine (0.5 µCi) were added to each well. The plates were then put back into the chamber and gassed with a 4% CO₂, 3% O₂, and 93% N₂ mix. The chamber was then placed in the incubator for 24 hours at 37°C. The assay was terminated by harvesting the content of each microtiter plate. Data were transferred into a graphic program (Excel) and analyzed to determine the IC₅₀. The test scores were grouped as follows; inactive (no repeat) IC₅₀ > 5 µg/mL; moderate activity (repeat) 0.5 µg/mL < IC₅₀ < 5 µg/mL; high activity (repeat) IC₅₀ < 0.5 µg/mL, 6253.

\[\text{In vitro sensitivity assays (cytotoxicity).}\]

The test samples were submitted to L-6 (rat skeletal myoblast cells). Compounds were prepared in DMSO the testing procedure was as follows. A sample of 100 µL medium (RPMI + 10% FCS + 1.7 µML - Glutamine (850 µL 200 mM for 100 mL), was added to wells of a microtiter plate. Then 100 µL of a cell suspension of 4 x 10⁶ cells/ml was added. After a serial dilution factor of 1:3 was prepared into the wells. The plates were then incubated for 70 hours at 37°C/5%CO₂.

\[\text{In silico modelling.}\]

Each compound was sketched using the GaussView software and geometry optimisation was carried out using the Gaussian 09W package with the density functional theory (DFT) using the B3LYP/6-31+G(d,p) approach until convergence was reached. This was necessary in order to generate good starting geometries for the 3D models. The low-energy 3D chemical structures were saved in mol2 format and initially treated with LigPrep, distributed by Schrodinger, Inc. This implementation was carried out with the graphical user interface (GUI) of the Maestro software package, using the OPLS force field. The treatment with LigPrep was necessary in order to correct protonation states and set the right parameters necessary for running QikProp. Protonation states at biologically relevant pH were correctly assigned (group I metals in simple salts were disconnected, strong acids were deprotonated and strong bases protonated, while explicit hydrogens were added). A set of the ADMET-related properties (a total of 46 molecular descriptors).
were calculated using the QikProp program running in normal mode. QikProp generates physically relevant descriptors and uses them to perform ADMET predictions. An overall ADMET-compliance score, drug-likeness parameter (indicated by #stars), was used to assess the pharmacokinetic profiles of the compounds. The #stars parameter indicates the number of property descriptors computed by QikProp, which fall outside the optimum range for 95% of known drugs. The methods implemented were developed by Jorgensen et al. All molecular modelling was carried out on a Linux workstation with a 3.5 GHz Intel Core2 Duo processor.

Conclusions
A series of 1-aryl-6-hydroxy-1,2,3,4-tetrahydroisoquinoline analogues was synthesized using simple methodology. To the best of our knowledge this is the first time the antiplasmodial activity of these analogues are reported. Following the criteria established by the WHO/TDR screening program, a compound is considered active against P. falciparum when in vitro IC₅₀ is < 1 µM (or 0.2 µg/ml) and SI > 100. In the present study, three compounds (Ⅲb, Ⅲk and Ⅳb) fell within this category and could be described as antimalarial hits from which more potent THIQ analogues could be obtained by further investigation. Additionally, in vivo and mechanistic studies of these compounds are underway. The study clearly demonstrates that 1-aryl-6-hydroxy-1,2,3,4-tetrahydroisoquinolines display sufficient antiplasmodial activity and favourable predicted pharmacokinetic properties to warrant further investigation of their antiplasmodial activity.

Acknowledgements
The academic license is acknowledged from Schrodinger Inc., for running the QikProp software.

Notes and references
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Figure Captions

Figure 1: 1,2,3,4 – tetrahydroisoquinoline

Figure 2: michellamine B (1) and dioncophyllines B (2), C(3), D(4) and E(5) with the
tetrahydroisoquinoline scaffold shown in red

Figure 3: homoerythrinane alkaloids

Figure 4: 2D pairwise scatter plots of selected ADMET properties: (A) Molecular weight against lipophilicity, (B) Molecular weight against number of hydrogen bond acceptors, (C) Molecular weight against dipole moments, (D) Molecular weight against number of rotatable bonds, (E) Molecular weight against cohesion indices, (F) Molecular weight against blood-brain barrier penetration coefficient, (G) Molecular weight against predicted logIC_{50} of HERG channel blockage, (H) Molecular weight against predicted binding coefficient to human serum albumin, (I) Molecular weight against skin penetration parameter, (J) Molecular weight against aqueous solubility, (K) Molecular weight against Caco-2 penetration, (L) Molecular weight against Madin-Darby canine kidney parameter,

Scheme 1: Synthesis of 6 - hydroxyl - 1,2,3,4 - tetrahydroisoquinoline analogues

Fig. 1

Fig. 2
Fig. 3

Reagents:
a: acetic acid (HOAc) and HBr (1:1), reflux;
b: benzaldehyde or substituted benzaldehyde, triethylamine and ethanol, reflux;
c: cyclohexanone, triethylamine and ethanol, reflux.

Scheme 1
Table 1: *In vitro* cytotoxicity and antiplasmodial activities of the synthesized 1-aryl-1,2,3,4-tetrahydroisoquinolines, along with selectivity indices.

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<th>Substituent (R)</th>
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<th>IC_{50} P. falciparum K1 (µM)</th>
<th>Cytotoxicity L6 (µg/ml)</th>
<th>Cytotoxicity L6 (µM)</th>
<th>Selectivity index</th>
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Table 2: Predicted ADME-related properties of the THIQs (Fig. 3) computed by QikProp program.\textsuperscript{31}

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<th>NRB</th>
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<th>V&lt;sub&gt;mol&lt;/sub&gt; (Å&lt;sup&gt;3&lt;/sup&gt;)</th>
<th>LogS&lt;sub&gt;cut&lt;/sub&gt;</th>
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*Property which falls outside the recommended range for 95% of known drugs; \#Number of computed properties which fall outside the required range for 95% of known drugs; \#Number of violations of Lipinski’s “Rule of Five”; Molar weight (range for 95% of drugs: 130–725 Da); Logarithm of partitioning coefficient between n-octanol and water phases (range for 95% of drugs: -2 to 6.5); Number of hydrogen bonds accepted by the molecule (range for 95% of drugs: 2–20); Number of hydrogen bonds donated by the molecule (range for 95% of drugs: 0–6); Number of rotatable bonds (range for 95% of drugs: 0–15); Logarithm of predicted blood/brain barrier partition coefficient (range for 95% of drugs: -3.0 to 1.0); Predicted apparent Caco-2 cell membrane permeability in Boehringer–Ingelheim scale, in nm/s (range for 95% of drugs: < 5 low, > 500 high); Total solvent-accessible molecular surface, in Å<sup>2</sup> (probe radius 1.4 Å) (range for 95% of drugs: 300–1000 Å<sup>2</sup>); Hydrophobic portion of the solvent-accessible molecular surface, in Å<sup>2</sup> (probe radius 1.4 Å) (range for 95% of drugs: 0–750 Å<sup>2</sup>); Total volume of molecule enclosed by solvent-accessible molecular surface, in Å<sup>3</sup> (probe radius 1.4 Å) (range for 95% of drugs: 500–2000 Å<sup>3</sup>); Logarithm of aqueous solubility (recommended rang is -6.0 to 0.5); Logarithm of predicted binding constant to human serum albumin (range for 95% of drugs: -1.5 to 1.5); Predicted IC<sub>50</sub> value for blockage of HERG K<sup>+</sup> channels (concern < -5); Number of likely metabolic reactions (range for 95% of drugs: 1–8).