## Reversal of Chloroquine and Mefloquine Resistance in *Plasmodium* falciparum by the Two Monoindole Alkaloids, Icajine and Isoretuline

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The chloroquine-potentiating activities of the Strychnos myrtoides alkaloids strychnobrasiline and malagashanine has been demonstrated in vitro and in vivo by Rasoanaivo et al. in 1994 [1]. In the continuation of our search for new antiplasmodial indole alkaloids [2] and with the aim of finding new resistance-modifiers agents, eight naturally occurring monoindole alkaloids (icajine, strychnobrasiline, isoretuline, retuline, vomicine, novacine, holstiine and dolichantoside) were evaluated in vitro for their ability to inhibit Plasmodium falciparum growth and, in drug combination, to reverse the resistance of a chloroquine-resistant strain of Plasmodium falciparum. None of these indole alkaloids has significant intrinsic antiplasmodial activity (IC50 > 10  $\mu$ M or 5  $\mu$ g/ml). Nevertheless, three alkaloids (icajine, isoretuline and strychnobrasiline) reverse chloroquine resistance at concentrations between 2.5 and 25  $\mu$ g/ml (IF of 12.82 for isoretuline on W2 strain). The Interaction Factor (IF) equals 2, <2, or>2 for additive, antagonistic or synergistic effects of alkaloids on chloroquine inhibition, respectively. Icajine and isoretuline were also assessed in vitro for their mefloquine potentiating activity on a mefloquine-resistant strain of Plasmodium falciparum. Only icajine proved to be synergistic with mefloquine (IF = 15.38).

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