

BLOCKS

Evidence on primaquine

- Pharmacology
 - » What's the active metabolite?
 - » Suitable drug partner?
- Safe and efficacious dose
- Safety by G6PD phenotype



Phase 1

Ex-vivo studies for efficacy and pharmacodynamics and pharmacokinetics

Phase 2

Dose escalation for safety and efficacy in high risk groups

Phase 3

Community trials with surrogate marker endpoints

Phase 4

Determine safety in pregnancy through review of primaquine in pregnancy in countries using primaquine



Common endpoints



Unsafe in G6PD—need to develop PoC test for phenotype



Safe and efficacious dose for mass drug administration alone or in combination

Drug supply and regulation



Identify stakeholders



Inclusion of stakeholders in road map



Implemented

Alternatives



Development of gametocytocidal drugs, e.g., tafenoquine and methylene blue



Product pipeline moving

ROAD MAP

GOAL