

Single-low dose Primaquine (SLD PQ) in Sinda, Zambia

Zambia Medical Association Scientific Conference and AGM

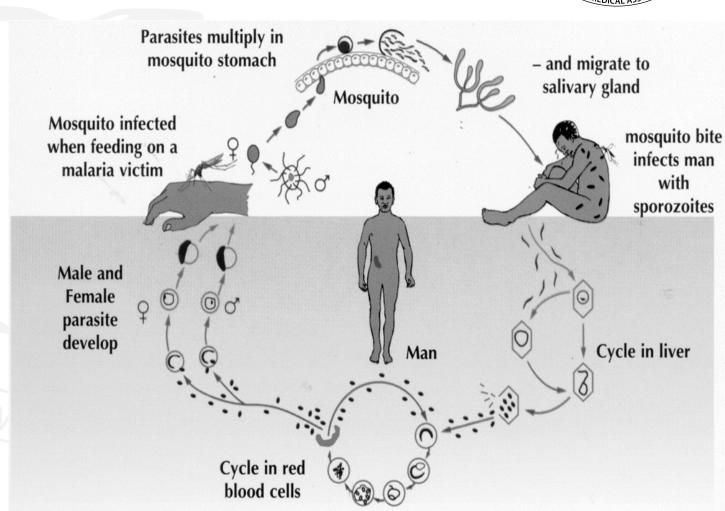
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Plasmodium falciparum life cycle



- Asexual forms of the Plasmodium sp. are responsible for causing Malaria symptoms
- Sexual forms (male and female gametocytes) are responsible for infecting the mosquito
- Gametocytes do not cause clinical symptoms of malaria, but are essential for human-tohuman transmission of malaria



Single-low dose primaquine (SLD PQ)



- Antimalarial medication, effective against Plasmodium falciparum gametocytes
- Primaquine poses a dose dependent risk of causing acute-hemolytic anemia (AHA) in people with Glucose-6-phosphate dehydrogenase deficiency (G6PD)
- G6PD deficiency is an X-linked recessive disease due to defective G6PD enzyme activity, which makes red blood cells susceptible to oxidative damage leading to AHA
- WHO (2015) recommended SLD PQ in low malaria transmission areas, as a single dose of 0.25 mg/kg primaquine with ACT, without need for G6PD testing



Single-low dose primaquine (SLD PQ)



- Renewed global interest in SLDPQ as an intervention that could curb the transmission of artemisinin resistant parasites – a threat of global concern.
- SLD PQ recommended in Zambian health facility catchment areas with malaria case incidence of less than 125 cases per 1,000 population
- SLD PQ included in the upcoming updated versions of the national malaria treatment guidelines (6th edition) as an addition to standard ACTs

SLDPQ in the 2022 to 2026 NMESP, Page 31

The intervention packages/activities planned for each level of transmission are shown in table 5.

Table 5: Intervention Package/Activities

Epidemiologic	Malaria	Intervention Package/Activities
Level (Stratum)	Indicator	
0	0 cases, No local transmission	No malaria - Maintenance of malaria-free zone Ensure uninterrupted availability and rational use of malaria commodities in health facilities and communities Maintain quality case management at facility and community levels – Malaria case investigation Reactive case detection Primaquine administration
		ITN continuous distribution ITN mass campaigns, except in Lusaka city Responsive IRS in eligible HFCAs LSM in select urban sites Entomologic surveillance Enhanced epidemiologic surveillance Social and Behaviour Change
1	1–49 cases/1,000 population, <1% parasite prevalence	Very low malaria transmission Ensure uninterrupted availability and rational use of malaria commodities in health facilities and communities Maintain quality case management at health facility and community levels Maintain and scale up community case management – Malaria case investigation Reactive case detection Targeted Mass Drug Administration Primaquine administration ITN continuous distribution ITN mass campaigns, except in Lusaka city Responsive IRS in eligible HFCAs

Exclusion criteria for SLDPQ



- The following have been excluded from the Sinda SLDPQ Pilot
 - Pregnant women
 - Infants under 6 months
 - Mothers of lactating infants under 6 months
 - Patients with severe anaemia Hb < 7g/dL
 - Patients from outside the catchment area implementing the SLDPQ pilot

SLDPQ in the 2022 to 2026 NMESP, Page 31

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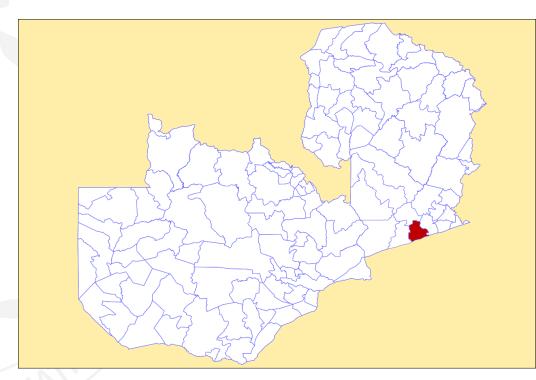
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The SLD PQ pilot in Sinda, Zambia



- Implemented by the Zambian Ministry of Health, National Malaria Elimination Program, supported by the U.S. President's Malaria Initiative Project (PMI) PAMO Plus Project
- Objectives
 - Pilot SLD PQ with active pharmacovigilance
 - Advice for nationwide rollout of SLD PQ in Zambia
- Geographical area
 - Zambia population: 20,966,971, 100% at risk of Malaria
 - Sinda District Population: 226,780.
 - 13 of 33 health facilities selected for the SLD PQ pilot (Pop: 122,658)



The SLD PQ pilot in Sinda, Zambia



- Pharmacovigilance for hemolytic anaemia when patients receive standard anti-malarial therapy + SLDPQ
 - Day 0 Urine scale on Hillmen Chart + Hb on Hemocue
 - Day 7 Urine scale on Hillmen Chart + Hb on Hemocue
- Preliminary findings (patient enrolment since 8 April 2024)
 - 250+ patients documented
 - 1 adverse event recorded, not hemolytic anaemia
- Timelines
 - Mid year pilot evaluation November 2024
 - End of year evaluation May 2025
- Challenges
 - Patient follow up on Day 7 (esp. patients outside catchment area)



Anticipated steps after the SLDPQ Pilot



- Evaluation of pharmacovigilance data
- Scale up implementation of SLDPQ to all eligible Health Facility Catchment Areas (HFCAs) in Zambia with an incidence of less that 125 malaria cases per 1,000 population.
- Transition from active to passive pharmacovigilance
- SLDPQ to be administered by Community Health Workers
- Evaluation of the impact of SLDPQ on malaria transmission and elimination in low burden areas





Thank you for your attention