Eastern and Southern Africa National Malaria Programmes

and Partners Annual Meeting

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Guidance on malaria elimination

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WHO Guidance on Malaria Elimination

Global Malaria Programme

A framework for malaria elimination

World Health Organization

- New guidance for malaria elimination developed in 2017
- The Framework for malaria elimination encourages all countries to accelerate towards malaria elimination, in line with the Global technical strategy for malaria 2016–2030
- Highlights programme actions across the continuum of transmission intensity from high to zero and prevention of re-establishment



Malaria Transmission Continuum





Principles of Malaria Elimination



Guidelines for elimination



4 PREVENTION

- 4.2.6 Mass drug administration (MDA)
 - 4.2.6.3 MDA to reduce transmission of *P. falciparum* in very low to low transmission settings

6 INTERVENTIONS IN THE FINAL PHASE OF ELIMINATION AND PREVENTION OF RE-ESTABLISHMENT

- 6.1 Intervention recommended for mass implementation in delimited geographical areas
- 6.2 Interventions targeting infections in people at higher risk
- 6.3 Interventions in response to detection of confirmed malaria cases



Additional interventions for elimination

Countries or areas that have attained very low to low levels of transmission require additional interventions in order to eliminate malaria. These interventions should:

- accelerate the decline in malaria transmission to a level at which intensive surveillance, i.e. follow-up of every case, is feasible;
- target specific groups at increased risk of infection that may not be reached adequately through routine prevention and treatment services; and
- respond to individual cases and foci to interrupt transmission.



Mass, Targeted and Reactive Strategies for Elimination







Mass strategies

applied to the entire population of a delimited geographical area, whether a hamlet, township or district

Targeted strategies

target specific groups at increased risk of infection that may not be reached adequately through routine prevention and treatment services

Reactive strategies

respond to individual cases and foci to interrupt transmission



Mass Drug Administration

MDA to reduce transmission of *P. falciparum* in very low to low transmission settings is **recommended**

- The effect wanes within 1-3 months
- Other components of a robust malaria elimination programme should be in place to reduce the risk of resurgence
- Considered for areas with limited risk of importation.
- Should not divert the resource of other essential components of an elimination programme
- The frequency of rounds and duration of the MDA programme should consider the local malaria epidemiology, the length of the prophylactic period provided by the antimalarial used, and the feasibility and cost of delivering each additional round.



Mass testing and treatment

MTaT to reduce the transmission of malaria is **<u>not</u> <u>recommended</u>**.

- MTaT has a very limited beneficial impact on malaria prevalence and incidence
- Resources required to implement this strategy are considered large.

However, WHO's conditional recommendation against MTaT notes that there may be exceptional circumstances under which MTaT might be appropriate, such as in very low transmission settings where MDA is not an acceptable or feasible strategy.

Mass strategies are generally not recommended for post-elimination settings unless there is a resumption of local transmission of malaria.

Targeted drug administration

Targeted drug administration to reduce transmission of malaria is **recommended**

- Persons given antimalarials should be those with increased risk of infection.
- Factors identifying individuals or groups at increased risk of infection should be easy to recognize.
- P. vivax: carefully consider how to safely and feasibly administer treatment to prevent relapses.
- Avoid stigmatizing groups at increased risk of infection.
- Additional complementary strategies to eliminate or prevent re-establishment of malaria transmission should be in place.

Targeted testing and treatment

TTaT to reduce transmission of malaria is **not recommended**

- Likely impact of TTaT on malaria transmission in very low to low or post-elimination settings is trivial
- Challenges with detecting very low parasite densities and a lack of diagnostics for hypnozoites.

The GDG noted that there may be limited circumstances under which targeted testing and treatment (TTaT) could be beneficial. For example, TTaT could be used when people at a higher risk of infection can be easily identified and chemoprevention is not acceptable to the population. Additionally, TTaT could be used if safe and effective implementation of radical cure to prevent P. vivax relapses is only feasible for those with confirmed infections.



Testing and treatment at points of entry to reduce importation of malaria

Routine malaria testing and treatment at points of entry is **not recommended**

- No studies of the impact of testing and treatment at points of entry on the rate of malaria importation were found, so no direct evidence
- Impact on importation of malaria was likely to be small
- Acceptability and feasibility of testing and treating for malaria at points of entry would be low given the likely disruption to travel.



Malaria testing and treatment of **organized or identifiable groups** arriving or returning from malaria-endemic areas is **recommended**

- Testing and treatment of positive of organized or identifiable groups of people (e.g. military, migrant laborers or religious pilgrims) can help countries nearing elimination or preventing re-establishment by reducing importation.
- Acceptability and feasibility of this strategy is
 considered higher than routine TaT at points of entry

In post-elimination settings, preventing infections in non-immune residents travelling to malaria-endemic areas through chemoprophylaxis would likely be a more effective approach than treating them upon return. International travel and health https://www.who.int/health-topics/travel-and-health#tab=tab_1

Reactive drug administration

Reactive drug administration for reducing malaria transmission is **recommended**

- capacity to conduct case investigations at the residence to determine the likely location of infection and to identify those individuals coexposed with the index case.
- capacity to enumerate and provide antimalarials to the people residing with or near a confirmed malaria case and others that share the same risk of infection.
- people given antimalarial medicine should share the same risk of having acquired infection as the index case
- if the infection was imported and the residence is not located in a receptive area, there may be no benefit from RDA.

Reactive case detection and treatment

Reactive case detection and treatment to reduce transmission of malaria is **recommended**

- Until an area is nearing elimination or is postelimination, it is unlikely that RACDT will have any effect on malaria transmission.
- RACDT becomes an essential component of surveillance when countries are nearing interruption of transmission to monitor progress towards elimination.
- When countries are post-elimination and working towards certification, RACDT can strengthen a country's claim that it has reached and maintained zero indigenous cases.
- RACDT is an essential part of surveillance and response to prevent re-establishment of malaria.



Reactive indoor residual spraying

Reactive indoor residual spraying is **recommended**

- Proactive IRS can be substituted by reactive IRS only
- Adding reactive IRS on top of proactive IRS should balance the potential added benefit with increasing cost and the risk of insecticide resistance.
- ➢ If no IRS is occurring, initiating reactive IRS may be beneficial.
- If the index infection was imported and the residence is not located in a receptive area, there may be no benefit from reactive IRS.



New recommendations overview

Recommendation	For / Against
MDA to reduce transmission of <i>P. falciparum</i>	For
MTaT to reduce transmission of malaria	Against
TDA to reduce transmission of malaria	For
TTaT to reduce transmission of malaria	Against
Routing malaria testing and treatment at points of entry to reduce importation	Against
MTaT of organized or identifiable groups at points of entry to reduce importation of malaria	For
RDA to reduce malaria transmission	For
RACDT to reduce transmission of malaria	For
Reactive indoor residual spraying to prevent or reduce transmission of malaria	For
Organization	

How to access WHO malaria guidance







WHO Global Malaria Programme website









Sections

ABBREVIATIONS		Conditional recommendation for, Very low certainty evidence	!
EXECUTIVE SUMMARY		Targeted drug administration to reduce transmission of malaria (2022)	
INTRODUCTION		In areas with very low to low transmission or post-elimination settings preventing re-establishment of transmission, antimalarial medicine can be given as chemoprevention to people with increased risk of infection relative to the general population to reduce transmission.	
PREVENTION	~		
CASE MANAGEMENT	~	 Persons given antimalarials should be those with increased risk of infection compared to the general population and their infections should constitute a large proportion of the parasite reservoir in the area. The factors identifying individuals or groups at increased risk of infection should be easy to recognise, thereby improving the acceptability and feasibility of the intervention. Programmes considering implementing targeted drug administration for <i>P. vivax</i> should carefully consider how to safely and feasibly administer treatment to prevent relapses. Care should be taken to avoid stigmatizing groups at increased risk of infection. Additional complementary strategies to eliminate or prevent re-establishment of malaria transmission should be in place. Research evidence (1) Evidence to Decision Justification Practical info Decision Aids More Info Feedback	
INTERVENTIONS IN THE FINAL PHASE OF ELIMINATION AND PREVENTION OF RE- ESTABLISHMENT	~		
SURVEILLANCE			
METHODS			
GLOSSARY			



Malaria elimination course





Spanish



https://openwho.org/courses/malaria-elimination

Thank you

