Outcome Report

Assessment of Cost-effectiveness of Mass Testing and Treatment (MTaT) of Malaria in Inaccessible Areas in India



User Department

National Vector Borne Disease Control Programme

HTAIn Regional Resource Hub



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Table of Contents

Contents

Research Team2
Acknowledgements
Table of Contents4
List of Tables5
List of Figures6
List of Abbreviations
Executive Summary
Background of the Project10
Malaria: Global Epidemiology and Economic Burden10
Malaria: Indian Scenario11
Malaria Control Program in India11
Mass Testing and Treatment (MTaT) of Malaria: Global Scenario
Mass Testing and Treatment in Indian Context13
Rationale of the project14
Overall aim16
Specific objectives
Methods17
Population, Intervention, Comparator, and Outcomes (PICO)17
Study settings17
Conceptual framework for decision tree18
Ethical considerations20
Findings
Comparison of Annual Parasite Index Trends in intervention and control sites21
Costing of NVBDCP and MTaT in Intervention blocks22
Costing of NVBDCP in control blocks26
Comparison of cost: NVBDCP+MTaT in intervention and NVBDCP in control blocks
Incremental Cost-Effectiveness Ratio (ICER)28
Conclusions and Policy Implications
References

List of Tables

Tables No.	Page No.
Table 1. Settings and sampling strategy	18
Table 2. Detailed Methods	18
Table 3. Description of testing status in selected blocks	22
Table 4. Demographic and epidemiological parameters of each of the intervention blocks in 2021	22
Table 5: Overall demographic and epidemiological parameters of all intervention blocks in 2021	22
Table 6. Coverage of MTaT through program data (Year 2021)	23
Table 7. Break-up cost of village contact drive for MTaT	23
Table 8. Cost of conducting MTaT in intervention blocks in the year 2021	23
Table 9. Human Resources cost for NVBDCP in intervention blocks (Year 2021)	24
Table 10. Incentive cost and grand total on human resources for NVBDCP in intervention blocks	24
Table 11. Training Cost for NVBDCP in intervention blocks (Year 2021)	24
Table 12. Cost of consumables for NVBDCP in intervention blocks (Year 2021)	25
Table 13. Indirect cost for visiting PHCs by suspected malaria cases in intervention blocks (Year 2021)	25
Table 14. Cost for various behavior change communication activities for NVBDCP in intervention blocks	25
Table 15. Per unit cost of NVBDCP for intervention block (in INR)	25
Table 16. Total unit cost (per person) for NVBDCP + MTaT	25
Table 17. Human Resources cost for NVBDCP in control blocks (Year 2021)	26
Table 18. Incentive cost and grand total on human resources for NVBDCP in control blocks (Year 2021)	26
Table 19. Training Cost for NVBDCP in control blocks (Year 2021)	26
Table 20. Cost of consumables for NVBDCP in control blocks (Year 2021)	27
Table 21. Indirect cost for visiting PHCs by suspected malaria cases in control blocks	27
Table 22. Cost for various behavior change communication activities for NVBDCP in control blocks	27
Table 23. Per unit cost of NVBDCP (in INR) in control blocks	27
Table 24. Comparison of costing for NVBDCP + MTaT in Intervention and NVBDCP in control areas	28
Table 25. Per unit cost of NVBDCP for intervention block (Year 2021)	29
Table 26. Total unit cost (per person) for NVBDCP + MTaT	29
Table 27. Per unit cost of NVBDCP for control block (Year 2021)	29
Table 28. Comparison of costing for NVBDCP + MTaT in Intervention and NVBDCP in control areas	29

List of Figures

Figure No.	Page No.
Figure 1. Major malaria control interventions	12
Figure 2. Gaps in Malaria Elimination Programme	12
Figure 3. Overview of MTaT intervention	14
Figure 4: Map of the intervention and control sites taken in the study	17
Figure 5: Conceptual Framework for Decision tree	19
Figure 6: Trend of Annual Parasite Index (API) 2011-21 in MTaT Interventions and Control districts	21

List of Abbreviations

API: Annual Parasite Index ACD: Active Case Detection **BCC: Behavioural Change Communication** DAMaN: Durgama Anchalare Malaria Nirakarana HTA: Health Technology Assessment **ICERs:** Incremental Cost-Effectiveness Ratios IEC: Information, Education and Communication ICMR: Indian Council of Medical Research MDA: Mass Drug Administration LLINs: long-lasting insecticide-treated nets LMIC: Low-and Middle-Income Countries MTaT: Mass Testing and Treatment NMCP: National Malaria Control Programme NMEP: National Malaria Elimination Programme NVBDCP: National Vector Borne Diseases Control Programme WHO: World Health Organization

Executive Summary

Background

Malaria is a significant health problem in India contributing to >85% of the estimated cases in South East Asia. In 2021, India accounted for 79% of all cases and 83% of all malaria deaths in the WHO South East Asian Region. In India, approximately 539 million people reside in high transmission areas. Malaria is a major challenge in rural/tribal areas of the central eastern and northeastern states of India. These are all inaccessible areas including hilly terrains, forest areas or regions with seasonal cut offs, having large population of ethnic groups. The National Vector Borne Disease Control Programme (NVBDCP) is responsible for malaria control and provides technical and operational guidance whereas the onground services and oversight are provided by the State Vector Borne Disease Control division. The key elements of India's malaria control strategy include early case detection and complete treatment (EDCT) identified through active and passive case surveillance; along with vector control measures, information education and communication (IEC) strategies, and surveillance.

It was observed from the operational research that the persistence of malaria transmission occurs in remote villages/hamlets where there are poor surveillance and the presence of asymptomatic malaria cases (no fever, but they test positive for malaria). These asymptomatic cases act as silent reservoirs for malaria transmission. Hence, the Government of Odisha has initiated a program in 2016-17, called "Durgama Anchalare Malaria Nirakarana (DAMaN); Malaria Control in Inaccessible Areas" to combat the asymptomatic reservoirs through mass testing and treatment (MTaT) of positive cases. The main focus of MTaT was to kill malaria-parasites from the entire population in remote and inaccessible pockets, along-with addressing the equity concerns. Asymptomatic malaria remains a challenge for malaria control programs as it significantly influences the transmission dynamics. Odisha has reduced malaria by 80% between 2017 and 2018 through mass testing and treatment (MTaT) in high endemic regions. The reductions in the incidence of malaria in Odisha are attributed to MTaT intervention. In India, Odisha, Chhattisgarh, Madhya Pradesh, and Jharkhand states account for 74.1% of the total malaria cases reported in the country which requires a cost-effective intervention along with a Budget Impact Analysis, which could be useful for large scale implementation of interventions in other parts of the country where the malaria burden is high. Hence, we reviewed the malaria trend in MTaT interventions and control settings and assessed the cost-effectiveness of this intervention along with routine NVBDCP.

Methods

<u>P: Population</u> – High-risk groups for malaria transmission (all age group)
 <u>I: Intervention</u> – Mass Testing and Treatment (MTaT)
 <u>C: Comparator/Control</u> – Routine malaria control program by NVBDCP
 <u>O: Outcome</u> – Incremental cost-effectiveness ratios (ICERs) for MTaT, averted cases (through API)

<u>T: Time Horizon</u> – Six-years

Findings

An overall significant reduction in the API was observed in intervention districts as compared to control districts post implementation of MTaT intervention. Although, the pace of the slope or reduction plateaued around 2020 which could be due to disruptions caused by COVID-19 pandemic that led to halt in conduction of MTaT camps (social distancing restrictions). The per unit cost of implementing MTaT was 57 INR and that of NVBDCP was 197 INR for the year 2021 in six intervention sites. The total per unit cost of conducting NVBDCP along with MTaT was 254 INR in intervention districts. We found that the per unit cost of conducting NVBDCP in control sites was Rs. 246 INR. Hence, we observed that among intervention sites, there is an additional expenditure of 8 INR on malaria control programme as compared to that of the control site.

Conclusions and Implication

We observed that the API trend is decreasing in both intervention and control sites, yet there is a steep slope close to zero in the intervention group which highlights that MTaT along with NVBDCP will be more effective in controlling malaria burden. Additionally, we observed that the cost of implementing NVBDCP is slightly higher in intervention sites as compared to control. This suggests that MTaT is a cost-effective intervention which could help in reducing the cost of routine programme (NVBDCP) in long run as lesser the cases, reduced cost of diagnosis and treatment.

States deemed at high risk for malaria transmission are identified for prompt MTaT implementations. According to the experts' opinion, the states are encouraged to classify their districts for MTaT interventions, with a particular emphasis on selecting districts with an API of 2 and above per 1000 population. Furthermore, districts may subdivide their blocks based on API, and within each block identify the sub-centres based on API (more than 2 per 1000) for MTaT intervention. Moreover, the implementation of MTaT, along with NVBDCP, is encouraged in villages that are geographically inaccessible due to factors such as dense forest areas, absence of road connectivity, seasonal cut-off, a shortage of community health workers (ASHA), and a high density of tribal population.

9

Background of the Project

Malaria: Global Epidemiology and Economic Burden

According to World Health Organization (WHO) Malaria Report 2022, globally (in 84 malaria-endemic countries), there were an estimated 247 million cases of malaria in 2021. In there year 2021, there was an overall increase in 2 million malaria cases as compared to that of the year 2020 [1]. The highest burden of the increase was observed in countries based in the WHO African Region. Moreover, this has also increased from the baseline year of the Global Technical Strategy for Malaria 2016-2030 when there were only 230 million cases in the year 2015 [1]. Nonetheless, malaria case incidence decreased from 82 in 2000 to 57 in 2019, but in the year 2020, it again rose to 59 owing to the disruptions caused by the COVID-19 pandemic, with no change in subsequent years i.e. 2021 [1]. An estimated 13.4 million cases were attributed to disruptions owing to COVID-19. However, the world noticed a reduction in the proportion of cases due to *Plasmodium vivax* which reduced from about 8% in 2000 to 2% in 2021 [1]. Twenty-nine countries in the world account for 96% of malaria cases globally which includes India also. Although, WHO African Region has the highest malaria burden with four countries (Nigeria, the Democratic Republic of Congo, Uganda, and Mozambique) accounting for almost half of all cases globally. Nonetheless, Cabo Verde reported zero indigenous cases for consecutive 3 years in 2021, thus ending the malaria epidemic [1]. The WHO South-East Asia Region accounted for 2% of malaria cases globally. Malaria cases reduced significantly in South-East Asia from 23 million in 2000 to about 5 million in 2021 [1]. The incidence of malaria reduced by 82% in South-East Asia i.e. from 18 cases per 1000 population in 2000 to about 3 cases per 1000 population at risk in 2021. Sri Lanka was certified malaria-free in 2016 and remains malaria free. Between 2020 and 2021, there was an increase in 400000 cases in the region with over half of these cases reported in Myanmar [1]. Malaria cases reduced by 37% in the WHO Eastern Mediterranean region by 38% between 2000 and 2015, however the region witnessed an increase of 44% between 2016 and 2021 [1]. The WHO Western Pacific region had a decrease of 49% cases in 2021 as compared to 2000. Papua New Guinea accounted for majority (87%) of cases in the region in 2021. China was certified malaria-free in 2021 and Malaysia had no cases of non-zoonotic malaria for four consecutive years [1]. The WHO Region of the Americas witnessed a reduction of 60% case incidence in 2021 as compared to 2000 with majority of cases being reported from Brazil, Colombia, and The Bolivarian Republic of Venezuela in the region [1]. The WHO European Region has been malaria free since 2015. Globally, malaria deaths reduced over the period of 2000-2019, but then surged in 2020. However, mortality has again declined in 2021 [1]. Globally, an estimated 2 billion cases and 11.7 million deaths were averted in the period 2000-2021 [1]. Haakenstad et al., estimated that global malaria spending—accounting both for government and outof-pocket spending—amounted to \$4.3 billion (95% UI 4.2–4.4) in 2016, which is an 8.6% (95% UI 8.1– 8.9) per year increase over malaria spending in 2000 [2, 3].

Malaria: Indian Scenario

Malaria is a significant health problem in India contributing to >85% of the estimated cases in South East Asia. In 2021, India accounted for 79% of all cases and 83% of all malaria deaths in the WHO South East Asian Region [4, 1]. Although, *Plasmodium falciparum* is the most prevalent malarial parasite globally, India accounts for almost half of the *Plasmodium Vivax* cases globally. In India, approximately 539 million people reside in high transmission areas [5]. Malaria is a major health problem in rural/tribal areas of the central eastern and northeastern states of India [6]. These are all tribal dominated states having large population of ethnic groups [7]. Out of 609 districts, 124 districts with 30% or more tribal population comprising about 8% country's population contributed to 46% of the total malaria cases, 70% of *P. falciparum* and 47% of malarial deaths in the country [8]. Thus, tribal areas are heartlands of malaria and there should be the focus on malaria elimination strategies specific to these areas.

Malaria Control Program in India

India has a long history of success and struggles with malaria control. Prior to the launch of the National Malaria Control Programme (NMCP) in 1953, 75 million cases and about 0.8 million deaths were reported annually for malaria (Figure 1). The widespread use of DDT indoor residual spray (IRS) resulted in a sharp decline in malaria cases, and in 1958, the NMCP was converted to the National Malaria Eradication Programme (NMEP) [9]. The NMEP was initially a great success with the malaria incidence dropping to 0.1 million cases and no deaths were reported due to malaria in 1965. In 1971, the urban malaria scheme (UMS) was also launched to cover 131 cities and towns [10]. These gains were short-lived, and in 1976, 6.4 million cases of malaria re-emerged. The resurgence was attributed to complacency and various operational, administrative, and technical reasons. The emergence of drug resistance in the parasites and insecticide resistance in the vectors also contributed to the resurgence. In 1977, the modified plan of operation (MPO) was initiated with the objectives to reduce malaria morbidity and prevent deaths due to malaria. Under the MPO, in addition to early diagnosis and prompt treatment, IRS was recommended in areas with annual parasite incidence (API) ≥ 2 [11]. The malaria incidence dropped to 1.66 million cases in 1987. The limited resources in many states, however, allowed spray coverage in areas with API > 5 only. In 1977, the eradication goal was officially shelved and the programme was changed to the National Anti-Malaria Programme (NAMP). By 1996, there was another malaria upsurge with reported 3.03 million cases and 2803 deaths [10]. In 2002, the NMCP became a part of the National Vector Borne Disease Control Programme (NVBDCP).



Figure 1: Major malaria control interventions

Responsibility for malaria control was divided between the central and the state governments of India. Technical and operational guidance was provided by the NVBDCP and the on-ground services and oversight were provided by the State Vector Borne Disease Control Division. The key elements of India's malaria control strategy include early case detection and complete treatment (EDCT), based on parasitological diagnosis of all suspected cases and complete treatment of all confirmed cases identified through active and passive case surveillance; along with vector control measures, information education and communication (IEC) strategies, and surveillance (Figure 2). The NVBDCP's commitment to end malaria aligns with rest of the world and responds to the commitment for malaria elimination by 2030 [11]. The National Strategic Plan for Malaria Control in India 2012–2017 was followed by the development of the National Framework for Malaria Elimination in India 2016–2030. The plan details every technical and operational element required for the nation to achieve malaria elimination by 2030.



Figure 2: Gaps in Malaria Elimination programme

Mass Testing and Treatment (MTaT) of Malaria: Global Scenario

The pathogenesis of malaria is complex, and the clinical presentation of disease ranges from severe and complicated, to mild and uncomplicated, to asymptomatic malaria. Asymptomatic malaria remains a challenge for malaria control programs as it significantly influences the transmission dynamics [12]. A study conducted in Ethiopia showed the prevalence of asymptomatic malaria was about six percent; furthermore, it also revealed that larger family size and previous history of malaria had significantly associated with the burden of asymptomatic malaria [13]. Moreover, in sub-Saharan Africa, Zanzibar provided new evidence on sustainable malaria reduction by asymptomatic malaria screening in the high endemic regions [14]. According to Sturrock et al. both hotspots – geographically discrete households and hotpops – demographically discrete populations having high malaria rates are the main target region for asymptomatic malaria screening and treatment. Hence, scaling up the screening and treatment of asymptomatic malaria in endemic regions is one of the priority interventions for the elimination of malaria [14, 15]. MTaT is one of the various recommendations provided by the WHO for eliminating malaria. The recommendations are broadly divided into three categories: i) mass strategies applied to the entire population of a delimited-geographical area, whether it be a village, township or district; these strategies include mass drug administration, MTaT and mass relapse prevention (MRP); ii) targeted strategies applied to people at a greater risk of infection than the general population; and iii) reactive strategies triggered in response to individual cases that includes reactive case detection and treatment to reduce transmission of malaria and reactive IRS.

Mass Testing and Treatment in Indian Context

It was observed from the operational research that the persistence of malaria transmission occurs in remote villages/hamlets where there are poor surveillance and the presence of asymptomatic malaria cases (no fever but they test positive for malaria). These asymptomatic cases act as silent reservoirs for malaria. Hence, the Government of Odisha state has initiated an innovative program in 2016-17, called *"Durgama Anchalare Malaria Nirakarana* (DAMaN); Malaria Control in Inaccessible Areas" to combat the asymptomatic reservoirs through mass screening and treatment of positive cases [16]. DAMaN is a type of MTaT activity with a focus on killing malarial parasites from the entire population in remote and inaccessible pockets, along-with addressing the equity concerns (Figure 3). It is done through mass testing with rapid diagnostic kit (RDK) for all age groups twice a year by organizing camps at the identified sites in villages. After mass testing, complete treatment is provided by the community health workers by following those tested positive. Additionally, information, education and communication activities are also done to increase the uptake of long lasting insecticidal treated nets (LLINs). The elimination interventions will require additional resources and regular monitoring to

sustain the momentum and stop the re-emergence of malaria. Hence, the Government needs support to identify the population at risk for asymptomatic malaria screening and its cost-effectiveness and budget impact analysis for the scaling-up of the program.

- Main focus in <u>DAMaN</u> was to kill malaria-parasites from the entire population in remote and inaccessible areas – addressing equity issues
- Attack on parasite in human
 - ✓ Cleaning the malaria parasites from the population of hard to reach village/hamlets by treating all positive malaria infections with appropriate anti-malaria drugs irrespective of their fever status
- - ✓ Thus malaria infection transmission is blocked from both symptomatic and afebrile (Asymptomatic) malaria cases in the <u>DAMaN</u> or nearby villages situated in an around 3 to 5 km radius.



Figure 3: Overview of MTaT intervention

Rationale of the project

Globally, malaria is considered as one of the severe public health issues, which causes immense morbidity as well as mortality [17]. It is a leading cause of death and disease particularly in low- and middle-income countries (LMIC) including India [17]. The elimination of malaria focusing on prevention and control of infection needs community access to the program as well as large scale coverage of effective interventions [18]. Furthermore, decisions on scale-up of the interventions require strong evidence on effectiveness, cost-effectiveness and equity of the interventions [19-23] along with acceptability of the program by the various stakeholders including community members, service providers, managers and decision-makers [24-27].

Globally, there is evidence that the detection and management of asymptomatic carriers have become an innovative strategy for malaria control programs as the treatment of asymptomatic cases have a potential impact on malaria reduction particularly in endemic regions [28]. A systematic review of the costs and cost-effectiveness of malaria interventions showed that most of the studies were undertaken in sub-Saharan Africa (78%) and limited studies were conducted in Asia (18%) as well as South America (4%) [23]. It was observed that the median cost of diagnosing a case of malaria was \$4.32, however, the median financial cost of treating an episode of severe malaria was \$30.26 [23]. The previous studies showed that MTaT is preferred than mass drug administration (MDA), as the latter involves massive over-use of drugs. MTaT incremental cost-effectiveness ratios (ICERs) were estimated in different sub-Saharan African settings revealed that at low transmission, MSTM probably is not worth considering. Instead, MTaT may be suitable at medium to high levels of transmission regions [22]. However, a cluster randomized control trial in Zambia estimated that population-wide MDA was more cost-effective compared to MTaT for transmission reduction [21]. In low resource settings, the optimal use of limited financial resources perpetrates the cost and cost-effectiveness analyses of MTaT. Consequently, clear evidence on the costs and cost-effectiveness of malaria control interventions is crucial for resource allocation and the selection of ideal interventions by malaria control programs [18, 23].

Community-based intervention on the management of malaria was proven as an effective and promising strategy for the prevention and control of malaria [25, 26]. Moreover, it was worthwhile to break financial and geographical barriers in relation to access healthcare especially in rural, remote, and hilly areas, where a majority of the population are vulnerable to malaria parasite infections. Furthermore, several studies proposed MTaT is an effective strategy to reduce parasite load particularly in high malaria-endemic regions [14, 15]. However, refinement of the large-scale implementation of interventions requires evidence on operational feasibility, cost implications and sustainability before expanding/scale-up the program.

MTaT was evaluated as a potential tool for malaria control and prevention across the globe [25-27]. Although, it was well accepted by the various stakeholders' including community members; still, it is important to understand the operational feasibility of the interventions including contextual needs – acceptance, community mobilization, service delivery and timing [25]. The evidence suggests that community literacy and long-term approach to community-engagement are vital to avoid misconceptions and fear towards MTaT [24-25, 27]; which is crucial for successful implementation of the community-based intervention. Therefore, it is essential to understand all stakeholders' perceptive on asymptomatic screening of malaria along with operational feasibility and cost-effectiveness analysis.

The evidence showed that if timely interventions are not undertaken or the existing surveillance system is not strengthened then it may increase the societal cost in terms of higher out of pocket expenditure for the treatment of positive cases, along with increased mortality and reduced quality-adjusted life years [13, 15, 16, 18, 21, 28]. As per the strategic plan, identification of local hot spots, identification of risk population MTaT and surveillance are essential. Hence, it is crucial to identify the population at risk for asymptomatic malaria screening and its cost-effectiveness and operational feasibility for scaling-up the program along with the budget impact analysis.

The pathogenesis of malaria is complex, and the clinical presentation of disease ranges from severe and complicated, to mild and uncomplicated, to asymptomatic malaria. Asymptomatic malaria remains a challenge for malaria control programs as it significantly influences the transmission dynamics [12]. Odisha state has emerged as an innovative step in malaria reduction – reduced above

15

80% between 2017 and 2018 through MTaT in high endemic regions, which is projecting as a future model for malaria eradication in India [16, 29]. The reductions in the incidence of malaria in Odisha are mainly due to DAMaN (MTaT) intervention. The program includes MTaT of malaria along with supportive supervision and community literacy. Hence, it is prudent to address the equity in terms vulnerability mapping – identify the prevalence and map the risk population, which will be considered for MTaT. In India Odisha, Chhattisgarh, Madhya Pradesh, and Jharkhand states account for 74.1% of the total malaria cases reported in the country which requires a cost-effective intervention along with a Budget Impact Analysis, which could be useful for large scale implementation of interventions in other parts of the country where the malaria burden is high.

Overall aim

To review the malaria trend between MTaT intervention and control settings before and after intervention; and to assess the cost-effectiveness of mass testing and treatment (MTaT) of malaria along with other routine NVBDCP programs.

Specific objectives

- To examine the trend of malaria in interventions and control blocks.
- To examine the cost-effectiveness of MTaT intervention along with routine malaria control program.

Methods

Population, Intervention, Comparator, and Outcomes (PICO)

P: Population – High-risk groups for malaria transmission (all age group including newborn children)
 I: Intervention – Mass Testing and Treatment (MTaT) of malaria
 C: Comparator/Control – Routine malaria control program by NVBDCP
 O: Outcome – Incremental cost-effectiveness ratios (ICERs) for MTaT, averted cases (through API)
 T: Time Horizon – Six-year time period

Study settings

We chose six intervention sites from Odisha (MTaT implemented along with regular NVBDCP) and six control sites i.e. three each from Jharkhand and Chhattisgarh (only NVBDCP).



Figure 4: Map of intervention and control sites taken in the study

These intervention and control sites (Table 1) were matched for API in the year 2017 at the time of MTaT implementation (before intervention matched API). The detailed sites are shown in the figure 4. The detailed methods presented in Table 2.

Table 1. Settings and sampling strategy

State	District	API (2017)	Block	Total population	% of Tribal Population	Literacy Rate	Total villages under CHC	Total no. of residential school	Total number of households	Total no of ASHA
	Latehar	12.8	Balumath	206422	27	58	172	5	37848	273
Jharkhand (Control)	West Singhbhum	10.39	Nowamundi	120456	58	63	66	1	19675	114
	Lohardaga	2.29	Kuru	135868	48	67	76	2	23837	102
Chile attice and	Narayanpur	42.05	Narayanpur	122290	24	56	187	3250	22691	365
Chnattisgarn (Control)	Bastar	15.05	Bastanar	55992	92	23	46		11726	317
(Control)	Durg	0.71	Patan	365289	8.6	68	146			497
	Boudh	4.76	Harbhanga	135797	18	74	368	8	32209	184
	Dhenkanal	1	Hindol	178899	9	73	167	3		170
Odisha	Anugul	7.4	Pallahara	145245	48	67	218	13	35320	134
(Intervention)	Keonjhar	16.9	Joda	149185	40	67	215	6	29456	189
	Malkangiri	37.3	Khairput	47903	58	34	214	16	10881	85
	Koraput	26.4	Boipariguda	127632	58	38	650	24	29847	230

Table 2: Detailed Methods

Specific Objectives	Design, Settings, and Participants		Analysis
	 Desk review of programmatic data 	•	Visual
Malaria Trend	 Programmatic data of state National Vector 		Representation
	Borne Diseases Control Programme (NVBDCP)		
	Health system cost	•	Decision Tree
Cost-effectiveness	Prevalence of malaria (API)		
of MTaT	 Data collected from primary and secondary 		
	sources.		

Conceptual framework for decision tree

The existing National Vector Borne Disease Control Program (NVBDCP) has been active against all vector-borne diseases in India for nearly two decades. However, this program has been mixed success in states with high malaria endemicity. To reduce the high malaria burden, the Government of Odisha started a unique initiative malaria control programme in 2017 in hard-to-reach-areas. Mass testing and treatment (MTaT) is the key strategy to control malaria transmission, along with the promotion of long-lasting-insecticidal nets (LLIN), insecticidal residual spray (IRS) and spreading awareness against malaria. A decision tree model with a six-year time horizon (three years each for pre and post MTaT implementation period) was used to evaluate the cost-effectiveness of the MTaT programme. We

defined effectiveness as the reduction in API of the year in interventions and control blocks. In this model, we assumed risk population for malaria infection are screened, diagnosed, treated, and prevented from malaria infection through the NVBDCP program with the MTaT scheme as intervention and routine program of NVBDCP as control group; as NVBDCP is the routine existing universal program implemented in India for malaria prevention and control. These programs identify the malaria-infected persons in high API as well as low API districts. Prevalence of malaria infection (through API) was seen for MTaT and NVBDCP separately in the intervention branch. The intervention and control branch has two branches with end nodes and their expected cost is determined as follows:

- Branch A|A': The cost of preventing and detecting of malaria infection, reflecting with Prevalence of malaria in NVBDCP + MTaT and routine program of NVBDCP implemented blocks of high API district, is included under this branch.
- Branch B|B': The cost of preventing and detecting of malaria infection, reflecting with prevalence of malaria in NVBDCP + MTaT and routine program of NVBDCP implemented blocks of low API district, is included under this branch.

The total costs of MTaT + NVBDCP and the Routine program of NVBDCP branches indicate the total cost of each program for the risk population. Our expected effectiveness for each program is calculated by multiplying the number of risk population entering the model by taking the prevalence of malaria (Figure 5).



Figure 5: Conceptual Framework for Decision tree

The framework for costing was done in five stages: firstly, we collected preliminary data regarding budgetary allocation, programmatic data from NVBDCP and block programme management unit of Department of Health of the respective states. It was followed by segregation of data regarding basic unit costs of human resources, various consumables and non-consumables for the malaria control programs. The costs were then calculated based on services provided for malaria control such as mass screening and treatment. Human resource costing was done according to the time spent for malaria control for the respective programs. NVBDCP also works for diseases other than malaria whereas MTaT specifically works for malaria control so, all the costs were adjusted according to the time spent and prevalence of malaria cases in the region. The budget impact analysis estimates the annual total cost of each of the malaria program from the perspective of the health system for each block regarding screening, and treatment of malaria infection by MTaT. The total cost includes the unit cost per person screened, and treated multiplied by the number of persons at risk of malaria in the blocks.

Ethical considerations

This study was approved by the Technical Appraisal Committee (TAC) of Health Technology Assessment in India, Department of Health Research (DHR), Ministry of Health and Family Welfare, Government of India. The ethical clearance was obtained from the Institutional Ethical Committee of ICMR-RMRC Bhubaneswar and State Research and Ethics Committee, Department of Health and Family Welfare, Govt. of Odisha. Permission was obtained from the concerned local authorities.

Findings

Comparison of Annual Parasite Index Trends in intervention and control sites

The implementation of MTaT encompasses universal screening for malaria has led to a reduction in the API of Odisha. However, there is lack of evidence that whether it is only in MTaT implemented other parts of the country have also experienced similar reductions in API. Hence, we compared the API of intervention (Odisha) and control (Jharkhand and Chhattisgarh) groups post MTaT implementation to garner evidence on the same. We collected data on the Annual Parasite Index (API) of the intervention district and its control from the year 2011 to 2021. Although, we intended to collect block-level data, but API at block were not available at many places especially of the past years due to which we had to resort to district level API. The MTaT intervention was started in the year 2017. The API matched (for 2017) intervention and control sites were compared for the trends in API post implementation of MTaT in Odisha with the regular NVBDCP implemented control sites. We used graphical representation to show the trends of each if the intervention-control groups (Figure 6).



Figure 6: Trend of Annual Parasite Index (API) 2011-21 in MTaT Interventions and Control districts

We observed that there is a sharp decline in the API between 2011-2017 in the intervention district which kept declining more as compared with the control. However, it plateaued around 2020 (COVID-19/pandemic year). However, its decline continued till 2021. We saw a sharp and continuous decline in API between the second intervention and control pair districts during 2017-2021. We observed a very steep reduction in the API of intervention district post MTaT implementation which continued till 2021, although it plateaued around 2019-2021. We saw a significant reduction in the API of the intervention district during 2017-2018 which was levelled during 2019 but again an upsurge was

observed during 2019-2021. In the sixth district, a significant reduction in API was observed from 2017-2021. An overall significant reduction in the API was observed in intervention districts as compared to control districts post implementation of MTaT programme. Although, the pace of the slope or reduction plateaued around 2020 which could be due to disruptions caused by COVID-19 pandemic that led to halt in conduction of MTaT camps (social distancing restrictions).

Costing of NVBDCP and MTaT in Intervention blocks

Table 3: Demographic and epidemiological parameters of each of the intervention blocks in 2021

MTaT (Round 1 + Round 2)	11	12	13	14	15	16
Demographic and epidemiological parameters						
No. of target villages	90	24	138	30	149	231
Villages covered	90	24	138	30	149	149
Total target population	10588	3512	41568	11329	27908	26964
Population covered during camp	4824	3236	21498	10097	26302	25010
No. of camps (Target)	52	14	132	30	140	55
No. of camps (Conducted)	52	14	132	30	140	55
Total Screened	4824	3236	21498	10097	26302	25010
No. of asymptomatic malaria cases diagnosed	98	0	29	3	225	157
No. of asymptomatic malaria treated	98	0	29	3	225	157

Site	State	Block	Year 2021			
Site	State	DIOCK	RDK	Microscopy	Positive	
Control		C1	1133	5639	21	
	Jharkhand	C3	8918	11764	517	
		C4	20049	22422	277	
		C2	15126	26771	13	
	Chhattisgarh	C5	20444	20176	1181	
		C6	12813	752	1058	
		11	29846	3683	331	
Intervention		12	21422	3262	7	
	Odiaha	13	27844	4090	34	
	Ouisila	14	29639	0	18	
		15	24515	4772	632	
		16	24108	3259	176	

Table 4: Description of testing status in selected blocks

*RDK: Rapid Diagnostic Kit

Table 5: Overall demographic and epidemiological parameters of all intervention blocks in 2021

Demographic and epidemiological parameters	Ν
No. of target villages	662
Villages covered	662
Target population (Total population of the target villages)	121869
Population covered	90967
No. of camps (Target)	423
No. of camps (Conducted)	423

Total screened	90967
No. of asymptomatic malaria cases diagnosed	512
No. of patients asymptomatic received treatment	512

Table 6: Coverage of MTaT through program data (Year 2021)

Target population	Population Covered	Percentage of coverage
121869	90967	74.64%

Table 7: Break-up cost of village contact drive for MTaT

Break-up cost of village contact drive for MTaT	Cost per camp (INR)
Advance intimation to community	100
Preparatory meeting	300
Printing and Use of IEC material	700
One wall painting	1000
School sensitization meeting	200
Drum beating	200
Documentation of MTaT activities	100
Total	2600

Table 8: Cost of conducting MTaT in intervention blocks in the year 2021

	COSTING	Unit Cost	Staff (N)	Days (N)	Camps/ Cases	Total
		(INR)			(N)	
Human	Incentives for volunteers	250	3	3	423	951750
Resources	TA & DA for health staffs	100	2	3	423	253800
	(MPHS/MPW)					
Total Human F	Resources cost (1)					1205550
Transport to	Mobility support to camp site	950		3	423	1205550
camp site (2)						
Training	Training of field staff	1200			423	507600
IEC/BCC	Village contact drive	2600			423	1099800
	Contingency (IEC & BCC materials, refreshment)	1000			423	423000
Meetings	Orientation (District @4000/- & Block @1000/-)	5000				5000
	Review meeting (District level @3000/-)	3000				3000
Total cost for t	training, IEC/BCC and meeting (3)					2038400
Screening cost	: (total target population*RDK cost) (4)	20			121869	2437380
Treatment cost [Asymptomatic cases*(10 tabs cost@ 10 INR + ASHA incentive for treatment follow-		85			512	43520
up @Rs75 INR						
Total Cost (1+2					6930400	
MTaT cost per individual (Total MTaT cost/Total target population)						57
IEC/BCC: Information, Education and Communication/Behavioral Change Communication						

Costing of NVBDCP in intervention blocks

Human Resource	Monthly salary	No. of units	Total working	% of hours	Hours dedicated	Annual cost for
	as per		hours/	dedicated	for	malaria
	7th CPC		month	to	malaria	(INR)
	(INR)		(48*30days)	malaria	(month)	
Medical Officer	65637	36	1440	20%	288	5671037
MPHS (M/F)	9600	28	1440	30%	432	967680
MPWs	9600	218	1440	30%	432	7534080
VBDC (Excluding PI i.e.	50468	6	1440	70%	1008	2543587
25% of Base Salary)						
VBDTS (Excluding PI i.e.	20130	6	1440	70%	1008	1014552
25% of base salary)						
Laboratory Technicians	14568	16	1440	40%	576	1118822
Sentinel site malaria	20130	6	1440	100%	1440	1449360
technician &						
Coordinator (Excluding PI						
i.e. 25% of Base Salary)						
Finance & Logistic	23192	6	1440	20%	288	333965
Assistant (NVBDCP)						
(Excluding PI i.e. 25% of						
Base Salary)						
Total (IA)	216825					20633083

Table 9: Human Resources Cost for NVBDCP in intervention blocks (Year 2021)

Table 10: Incentive cost for NVBDCP in intervention blocks (Year 2021)

Incentives	Incentive per case (INR)	Total Number Test	Total Incentives (INR)
ASHA Incentive for testing	15	No. of RDK Test Done: 135952	2039280
ASHA Incentive for treatment per case	75	No. of cases: 1198	89850
Total (IB)			2129130

Table 11: Training cost for NVBDCP in intervention blocks (Year 2021)

Training (District Level)	Unit Training	No. of	Frequency of	Training Cost
	Cost (INR)	Units	training/ Year	(INR)
Medical Officer	5400	36	1	194400
ASHA/ CHWs	1000	992	1	992000
MPHS (M/F) Induction Training	1000	28	1	28000
MPWs Refresher Training	1000	218	1	218000
VBDC	4500	6	1	27000
VBDTS (Refresher training, IHIP), 2 days	6000	6	1	36000
LTs induction training, 10 days	19000	16	1	304000
LTs refresher training, 5 days	8400	16	1	134400
Total (IC)	46300	1318		1933800

Consumables (Testing kits and medicines)	Unit cost (INR)	No of tablets/ case	No of cases	% age share of cases	Total No. of cases	Total Cost
Chloroquine phosphate tablets (per unit/tablet; 1 strip=10 tabs)	1	10	1198	8%	96	960
Primaquine tablets (per unit/tablet)	1	6	1198	92%	1102	6612
Testing kit (RDK)	20	135952				2719040
Glass slides for microscopy	2	19066				38132
Chemical and reagents						50000
Total (ID)						2814744

Table 12: Cost of consumables for NVBDCP in intervention blocks (Year 2021)

Table 13: Indirect cost/Societal cost for visiting Primary Health Centers by suspected malaria patients in intervention blocks (Year 2021)

Indirect cost category	Unit Cost	Units (cases with microscopy test)	Total
Travel to Primary Health Centres	150	19066	2859900
Total (IE)			2859900

Table 14: Cost for various behavior change communication activities for NVBDCP in intervention blocks (Year 2021)

IEC/BCC Activities	Unit cost	Units	Total
Advocacy at the district Level (World Malaria Day	15000	6	90000
GKS sensitization at the sub-centre level	600	142	85200
Total (IF)			175200

Table 15: Per unit cost of NVBDCP for intervention block (Year 2021)

Total NVBDCP cost (IA+IB+IC+ID+IE+IF) [for 155018 individuals]			
NVBDCP cost (INR) per individual (Total NVBDCP cost/Total number of RDK and	197		
Microscopy test)			

Table 16: Total unit cost (per person) for NVBDCP+ MTaT

MTaT (per unit cost) (INR)	57
NVBDCP (per unit cost) (INR)	197
Total cost per person (NVBDCP+MTaT) in INR	254

Costing of NVBDCP in control blocks

Human Resource	Monthly Salary as per 7th CPC (INR)	No. of units	Total working hours/ month (48*30days)	% of hours dedicated to malaria	Hours dedicated to malaria (month)	Annual towards Malaria (INR)
Medical Officer	65637	58	1440	20%	288	9136670
MPHS (M/F)	9600	22	1440	30%	432	760320
MPWs	9600	154	1440	30%	432	5322240
VBDC (Excluding PI i.e. 25% of Base Salary)	50468	6	1440	70%	1008	2543587
VBDTS (Excluding PI i.e. 25% of Base Salary)	20130	10	1440	70%	1008	1690920
Laboratory Technicians	14568	26	1440	40%	576	1818086
Sentinel site malaria technician and coordinator (Excluding PI i.e. 25% of base salary)	20130	6	1440	100%	1440	1449360
Finance & Logistic Assistant (NVBDCP) (Excluding PI i.e. 25% of base salary)	23192	6	1440	20%	288	333965
Total (CA)						21271824

Table 18: Incentive cost for NVBDCP in control blocks (Year 2021)

Incentives	Incentive per case (INR)	Total Number	Total Incentives (INR)
ASHA Incentive for testing	15	No. of RDK Test Done: 78483	1177245
ASHA Incentive for treatment per case	75	No. of cases: 3067	230025
Total (CB)			1407270

Table 19: Training cost for NVBDCP in control blocks (Year 2021)

Health System Cost: Training (District Level)	Unit	No. of	Frequency	Training Cost
	Cost (INR)	Units	Year	(INK)
Medical Officer	5400	58	1	313200
ASHA/CHWs	1000	1667	1	1667000
MPHS (M/F) Induction Training	1000	22	1	22000
MPWs Refresher Training	1000	154	1	154000
VBDC	4500	6	1	27000
VBDTS (Refresher training, IHIP), 2 days	6000	10	1	60000
LTs induction training, 10 days	19000	26	1	494000
LTs refresher training, 5 days	8400	26	1	218400
Total (CC)	46300	1969		2955600

Table 20: Cost of consumables for NVBDCP in control blocks (Year 2021)

Consumables						
Drugs	Cost (INR)	No of Tablets/case	No of cases	%age share of cases	Total No. of cases	Total Cost
Chloroquine phosphate tablets (Per	1	10	3067	8%	245	2450
unit/Tablet) (1 strip=10 tabs)						
Primaquine tablets (per	1	6	3067	92%	2822	16932
unit/tablet)						
Testing kit (RDK)	20	78483				1569660
Glass slides for microscopy	2	87524				175048
Chemical and Reagents						50000
Total (CD)						1814090

Table 21: Indirect cost for visiting Primary Health Centres by suspected malaria patients in control blocks (Year 2021)

Indirect cost category	Unit Cost	Units (Cases with microscopy test)	Total cost
Travel to Primary Health Centres	150	87524	13128600
Total (CE)			13128600

Table 22: Cost for various behaviours change communication activities for NVBDCP in control blocks

IEC/BCC Activities	Unit cost	Units	Total
Advocacy at the district Level (World Malaria Day	15000	6	90000
GKS Sensitization at sub-centre level	600	171	102600
Total (CF)			192600

Table 23: Per unit cost of NVBDCP (in INR) in control blocks

Total NVBDCP cost (IA+IB+IC+ID+IE+IF) [for 166007 individuals]	40769984
NVBDCP cost (INR) per individual (Total NVBDCP cost/Total number of RDK and	246
Microscopy test)	

Comparison of cost: NVBDCP+MTaT in intervention and NVBDCP in control blocks

Costing Heading	Total NVBDCP Cost	Total NVBDCP Cost	
	Interventions blocks	Control blocks	
	(Year 2021), INR	(Year 2021), INR	
Human Resource (A)	2,06,33,083	2,12,71,824	
Incentives (B)	21,29,130	14,07,270	
Training cost (C)	19,33,800	29,55,600	
Cost of consumables – testing and treatment (D)	28,14,744	18,14,090	
Indirect cost (Travel to PHCs)	28,59,900	13128600	
IEC/BCC activities (E)	1,75,200	1,92,600	
Total NVBDCP cost	3,05,45,857	4,07,69,984	
Total individual tested (RDK + Microscopy)	1,55,018	1,66,007	
NVBDCP cost per individual	197	246	
Total cost for MTaT	69,30,400	NA	
MTaT (per unit cost)	57	NA	
Total cost (NVBDCP+MTaT)	3,74,76,257	NA	
Total unit cost (per person) (NVBDCP+MTaT)	254	NA	

Table 24: Comparison of cost for NVBDCP + MTaT in Intervention and NVBDCP in control blocks

The per unit cost of implementing MTaT along with the routine NVBDCP programme is 254 INR as compared to 246 INR among control sites where only the NVBDCP programme is implemented. We observed that among intervention sites, there is an additional expenditure of Rs. 8.4 per individual on the malaria control programme as compared to that of the control site.

Incremental Cost-Effectiveness Ratio (ICER)

$$ICER = \frac{Cost(A) - Cost(B)}{Benefit(A) - Benefit(B)}$$

ICER=37476257-40769984/2.5-3.3 ICER=-3293727/-0.8 ICER=4117159

Sensitivity analysis for overestimating the total number of cases (RDK + Microscopy)

For calculating the total number of tests done, we used the sum of total RDK tests and total microscopy tests done in the population. However, there is always an overlap between the number of tests done by RDK and microscopy as few individuals would have got both tests done. Hence, we decided to do a sensitivity analysis by considering a reduction of 20% individuals from the total of RDK and microscopy tests.

Costing of NVBDCP in intervention blocks

Table 25: Per unit cost of NVBDCP for intervention block (Year 2021)

Total NVBDCP cost (IA+IB+IC+ID+IE+IF) [for 155018- (20% of 155018)= 124014	30545857
individuals]	
NVBDCP cost (INR) per individual (Total NVBDCP cost/Total number of RDK and	246
Microscopy test)	

Table 26: Total unit cost (per person) for NVBDCP+ MTaT			
MTaT (per unit cost) (INR)	57		
NVBDCP (per unit cost) (INR)	246		
Total cost per person (NVBDCP+MTaT) in INR	303		

Costing of NVBDCP in control blocks

Table 27: Per unit cost of NVBDCP (in INR) in control blocks

Total NVBDCP cost (IA+IB+IC+ID+IE+IF) [for 166007- (20% of 166007)= 132806	40769984
individuals]	
NVBDCP cost (INR) per individual (Total NVBDCP cost/Total number of RDK and	307
Microscopy test)	

Comparison of cost: NVBDCP+MTaT in intervention and NVBDCP in control blocks

Table 28: Comparison of cost for NVBDCP + MTaT in Intervention and NVBDCP in control blocks

Costing Heading	Total NVBDCP Cost	Total NVBDCP Cost
	Interventions blocks	Control blocks
	(Year 2021), INR	(Year 2021), INR
Human Resource (A)	2,06,33,083	2,12,71,824
Incentives (B)	21,29,130	14,07,270
Training cost (C)	19,33,800	29,55,600
Cost of consumables – testing and treatment (D)	28,14,744	18,14,090
Indirect cost (Travel to PHCs)	28,59,900	13128600
IEC/BCC activities (E)	1,75,200	1,92,600
Total NVBDCP cost	3,05,45,857	4,07,69,984
Total individual tested (RDK + Microscopy)	1,24,014	1,32,805
NVBDCP cost per individual	246	307
Total cost for MTaT	69,30,400	NA
MTaT (per unit cost)	57	NA
Total cost (NVBDCP+MTaT)	3,74,76,257	NA
Total unit cost (per person) (NVBDCP+MTaT)	303	NA

Conclusions and Policy Implications

Malaria is a major public health concern in India, with inaccessible terrain posing additional challenges that require special attention. Furthermore, asymptomatic malaria continues to pose a challenge to malaria control programmes because it has a significant impact on transmission dynamics. In India, Odisha, Chhattisgarh, Madhya Pradesh, and Jharkhand account for 74.1% of all malaria cases reported, indicating the need for a cost-effective malaria control intervention in accordance with the National Framework for Malaria Elimination in India 2016-2030.

We used API-matched similar settings in both the intervention and control groups to investigate malaria API trends before and after the intervention. We observed that the API trend is declining in both intervention and control sites; however, there is a significant decline in API in the intervention group, which is approaching zero. This demonstrates that MTaT combined with routine NVBDCP is more effective in controlling malaria burden.

In terms of cost, despite the similarity of the intervention and control settings, the intervention group spent less overall than the control sites. In 2021, the per-unit cost of implementing MTaT was 57 INR, while that of NVBDCP was 197 INR at six intervention sites. At intervention districts, the total cost per unit for NVBDCP and MTaT was 254 INR. We discovered that the per-unit cost of performing NVBDCP at six control sites was 246 INR. Thus, the intervention sites spent 8 INR per person more than the control sites. It should be noted that the MTaT intervention began in 2017 and has been followed by mass screening and treatment for the last five years, resulting in a significant reduction in asymptomatic parasite reservoirs and thus lower transmission. Furthermore, MTaT screens and treats everyone before and after the monsoon season, reducing transmission during the season and thus lowering the financial burden on the routine programme, i.e. NVBDCP, as fewer cases require diagnosis and treatment. This demonstrates that MTaT not only reduced the API but also the NVBDCP cost in the long run. Furthermore, similar settings (inaccessible areas) may use MTaT in conjunction with NVBDCP to reduce the API and financial burden on states.

The cost-effectiveness of implementing mass testing and treatment of malaria versus relying solely on routine fever testing and treatment to reduce the API is a crucial consideration in the fight against malaria. The effectiveness of these strategies varies depending on the specific context, prevalence of malaria, available resources, and the healthcare infrastructure of a given region. Mass testing and treatment initiatives can be highly effective in rapidly reducing the API, especially in areas with high malaria transmission rates. They provide a proactive approach to identifying and treating malaria

cases, thereby reducing the overall disease burden. However, they can be resource-intensive and may not be sustainable in the long term, particularly in resource-constrained settings. On the other hand, routine fever testing and treatment, while more cost-effective and sustainable, may not have the same immediate impact on reducing the API. This approach relies on passive case detection and may miss asymptomatic carriers, contributing to ongoing transmission.

The most effective strategy likely lies in a balanced approach that considers the specific needs and circumstances of each region. Mass testing and treatment can be employed as an initial intervention to rapidly reduce the API, followed by a shift towards routine fever testing and treatment for long-term sustainability. This combination allows for the identification and treatment of both symptomatic and asymptomatic cases while optimizing the allocation of limited resources. Moreover, ongoing research and investment in innovative malaria diagnostics, treatment options, and preventive measures are essential to enhance the cost-effectiveness of malaria control efforts. In the end, the goal remains the same: to reduce the API, save lives, and work towards the eventual elimination of malaria in a manner that is both economically viable and sustainable in the long run.

The global malaria strategy by the World Health Organization (WHO) emphasizes the need for all malaria-endemic countries to expedite their efforts toward achieving the elimination goal. The latest guidelines for the conclusive phase of elimination, as outlined in the WHO Guidelines for Malaria, present recommendations categorized into three types of interventions: "mass" strategies, "targeted" strategies, and "reactive" strategies. In instances where malaria transmission is widespread throughout the population in a specified geographical area, such as a district or village, comprehensive strategies may be essential to curtail transmission through Mass Testing and Treatment (MTaT). MTaT involves conducting malaria tests for the entire population within a defined geographic region and administering effective antimalarial treatment to all individuals who tested positive, typically within a synchronized timeframe.

MTaT is regarded as an integral component of the immediate response within the context of endemic control, aligning with various other malaria prevention and control initiatives outlined in the National Vector Borne Disease Control Program (NVBDCP). Given that malaria transmission tends to be localized and focal, the district emerges as the pivotal implementing unit. Consequently, program managers should function as the primary entities for both planning and executing these initiatives. States deemed at high risk for malaria transmission, including Odisha, Chhattisgarh, Jharkhand,

Madhya Pradesh, Maharashtra, Tripura, Meghalaya, and Mizoram, are identified for prompt MTaT implementations. According to the experts' opinion, the states are encouraged to classify their districts for MTaT interventions, with a particular emphasis on selecting districts with an Annual Parasite Incidence (API) of 2 and above per 1000 population. Furthermore, districts may subdivide their blocks based on API, and within each block identify the sub-centres based on API (more than 2 per 1000) for MTaT intervention. Moreover, the implementation of MTaT, in conjunction with the NVBDCP, is encouraged in villages that are geographically inaccessible due to factors such as dense forest areas, absence of road connectivity, seasonal cut-off, a shortage of community health workers (ASHA), and a high density of tribal population.

References

- 1. World Health Organization. World malaria report 2022. World Health Organization; 2022 Dec 8.
- Haakenstad A, Harle AC, Tsakalos G, Micah AE, Tao T, Anjomshoa M, Cohen J, Fullman N, Hay SI, Mestrovic T, Mohammed S. Tracking spending on malaria by source in 106 countries, 2000–16: an economic modelling study. The Lancet infectious diseases. 2019 Jul 1;19(7):703-16.
- Andrade MV, Noronha K, Diniz BP, Guedes G, Carvalho LR, Silva VA, Calazans JA, Santos AS, Silva DN, Castro MC. The economic burden of malaria: a systematic review. Malaria journal. 2022 Oct 5;21(1):283.
- 4. Rahi M, Sharma A. For malaria elimination India needs a platform for data integration. BMJ Global Health. 2020 Dec 1;5(12):e004198.
- 5. Lal AA, Rajvanshi H, Jayswar H, Das A, Bharti PK. Malaria elimination: Using past and present experience to make malaria-free India by 2030. J Vector Borne Dis. 2019 Jan 1;56(1):60.
- Chourasia MK, Raghavendra K, Bhatt RM, Swain DK, Valecha N, Kleinschmidt I. Burden of asymptomatic malaria among a tribal population in a forested village of central India: a hidden challenge for malaria control in India. Public health. 2017 Jun 1;147:92-7.
- 7. Sharma RK, Thakor HG, Saha KB, Sonal GS, Dhariwal AC, Singh N. Malaria situation in India with special reference to tribal areas. The Indian journal of medical research. 2015 May;141(5):537.
- Kumar A, Valecha N, Jain T, Dash AP. Burden of malaria in India: retrospective and prospective view. Defining and Defeating the Intolerable Burden of Malaria III: Progress and Perspectives: Supplement to Volume 77 (6) of American Journal of Tropical Medicine and Hygiene. 2007 Dec.
- Strategic plan for malaria control in India 2012–2017: A five year strategic plan. Delhi: Directorate of National Vector Borne Disease Control Programme, Government of India 2019; p.126. Available from: http://www.nvbdcp.gov.in/Doc/Strategic-ActionPlan-Malaria-2012-17-Co.pdf (Accesed on July 27, 2023).
- 10. Sharma V. Re-emergence of malaria in India. Indian J Med Res 1996; 103: 26–45.
- National Framework for Elimination of Malaria in India 2016– 30. Delhi: Directorate of National Vector Borne Disease Control Programme, Government of India; p. 43
- 12. Laishram DD, Sutton PL, Nanda N, Sharma VL, Sobti RC, Carlton JM, Joshi H. The complexities of malaria disease manifestations with a focus on asymptomatic malaria. Malaria journal. 2012 Dec;11(1):29.
- Ameha A, Taffere GR, Zenebe D. Asymptomatic Malaria A Missed Opportunity in the Planned
 2030 Malaria Elimination Programme in Ethiopia. Research & Reviews: A Journal of Immunology.
 2019 May 24;9(1):4-9.
- 14. Björkman A, Shakely D, Ali AS, Morris U, Mkali H, Abbas AK, Al-Mafazy AW, Haji KA, Mcha J, Omar

R, Cook J. From high to low malaria transmission in Zanzibar—challenges and opportunities to achieve elimination. BMC medicine. 2019 Dec;17(1):14.

- 15. Sturrock HJ, Hsiang MS, Cohen JM, Smith DL, Greenhouse B, Bousema T, Gosling RD. Targeting asymptomatic malaria infections: active surveillance in control and elimination. PLoS medicine. 2013 Jun 18;10(6):e1001467.
- Pradhan MM, Meherda PK. Malaria elimination drive in Odisha: Hope for halting the transmission. Journal of vector borne diseases. 2019 Jan 1;56(1):53.
- 17. World Health Organization. *Compendium of WHO malaria guidance: prevention, diagnosis, treatment, surveillance and elimination*. World Health Organization 2019.
- 18. White MT, Yeung S, Patouillard E, Cibulskis R. Costs and cost-effectiveness of Plasmodium vivax control. The American journal of tropical medicine and hygiene. 2016 Dec 28;95(6_Suppl):52-61.
- 19. Lai S, Sun J, Ruktanonchai NW, Zhou S, Yu J, Routledge I, Wang L, Zheng Y, Tatem AJ, Li Z. Changing epidemiology and challenges of malaria in China towards elimination. Malaria journal. 2019 Dec;18(1):107.
- Samby K, Ramachandruni H, Banerji J, Burrows JN, Daumerie PG, van Huijsduijnen RA, Duparc S, Wells TN. Partnering to fight malaria in India: Past, present and future. Journal of vector borne diseases. 2019 Jan 1;56(1):15.
- Silumbe K, Yukich JO, Hamainza B, Bennett A, Earle D, Kamuliwo M, Steketee RW, Eisele TP, Miller JM. Costs and cost-effectiveness of a large-scale mass testing and treatment intervention for malaria in Southern Province, Zambia. Malaria journal. 2015 Dec;14(1):211.
- 22. Crowell V, Briët OJ, Hardy D, Chitnis N, Maire N, Di Pasquale A, Smith TA. Modelling the costeffectiveness of mass screening and treatment for reducing Plasmodium falciparum malaria burden. Malaria journal. 2013 Dec;12(1):4.
- 23. White MT, Conteh L, Cibulskis R, Ghani AC. Costs and cost-effectiveness of malaria control interventions-a systematic review. Malaria journal. 2011 Dec 1;10(1):337.
- 24. Ndong IC, Okyere D, Enos JY, Amambua-Ngwa A, Merle CS, Nyarko A, Koram KA, Ahorlu CS. Challenges and perceptions of implementing mass testing, treatment and tracking in malaria control: a qualitative study in Pakro sub-district of Ghana. BMC public health. 2019 Dec;19(1):695.
- 25. Peto TJ, Tripura R, Sanann N, Adhikari B, Callery J, Droogleever M, Heng C, Cheah PY, Davoeung C, Nguon C, von Seidlein L. The feasibility and acceptability of mass drug administration for malaria in Cambodia: a mixed-methods study. Transactions of The Royal Society of Tropical Medicine and Hygiene. 2018 Jun 1;112(6):264-71.
- 26. Jaiteh F, Masunaga Y, Okebe J, D'Alessandro U, Balen J, Bradley J, Gryseels C, Ribera JM, Grietens KP. Community perspectives on treating asymptomatic infections for malaria elimination in The

Gambia. Malaria journal. 2019 Dec;18(1):39.

- 27. Shuford K, Were F, Awino N, Samuels A, Ouma P, Kariuki S, Desai M, Allen DR. Community perceptions of mass screening and treatment for malaria in Siaya County, western Kenya. Malaria journal. 2016 Dec;15(1):71.
- 28. Das NG, Dhiman S, Talukdar PK, Goswami D, Rabha B, Baruah I, Veer V. Role of asymptomatic carriers and weather variables in persistent transmission of malaria in an endemic district of Assam, India. Infection ecology & epidemiology. 2015 Jan 1;5(1):25442.
- Pradhan S, Pradhan MM, Dutta A, Shah NK, Joshi PL, Pradhan K, Sharma SK, Daumerie PG, Banerji J, Duparc S, Mendis K. Improved access to early diagnosis and complete treatment of malaria in Odisha, India. PloS one. 2019 Jan 2;14(1):e0208943.