Malaria RDTs in the private retail sectors of malaria endemic countries A review of the evidence







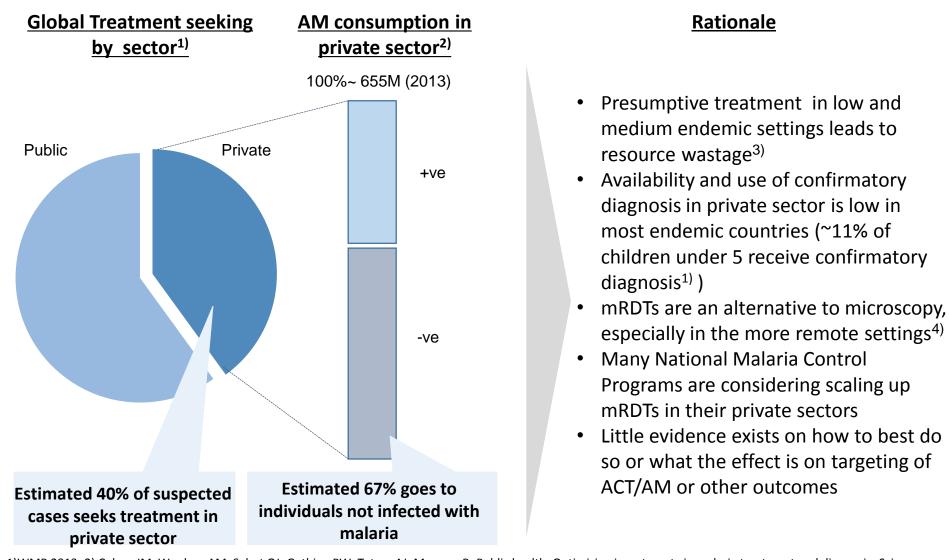




Malaria rapid diagnostic tests (RDTs) and fever case management in the private health care sector in Africa: a consultative working meeting

Entebbe, Uganda Oct 20 &21, 2015

The scale up of malaria Rapid Diagnostic Tests (mRDT) has potential to restrict antimalarial drugs to confirmed cases of malaria



1)WMR 2013, 2) Cohen JM, Woolsey AM, Sabot OJ, Gething PW, Tatem AJ, Moonen B: Public health. Optimizing investments in malaria treatment and diagnosis. *Science* 2012, 338:612-614., 3) D'Acremont V, Lengeler C, Mshinda H, Mtasiwa D, Tanner M, Genton B: Time to move from presumptive malaria treatment to laboratory-confirmed diagnosis and treatment in African children with fever. *PLoS Med* 2009, 6:e252., 4) Maltha J, Gillet P, Jacobs J: Malaria rapid diagnostic tests in endemic settings. *Clin Microbiol Infect* 2013, 19:399-407.

In April 2013, ACT Consortium and the RBM CMWG brought together evidence for preliminary guidance to RBM Board

Objectives

- To share the results of research activities and early pilot interventions to introduce diagnostic testing for malaria into the retail private sector
- To extract lessons learned and identify key bottlenecks and success factors from operational research and limited experiences to inform the future design of pilot projects for deploying diagnostic testing in the private sector in malaria endemic countries

Approach

 Review results of more than a dozen small-scale studies and pilots from eight countries in Africa and Southeast Asia

Emerging Themes¹⁾

- Ensure availability for affordable, quality assured mRDTs to be scaled-up sustainably in the market
- Ensure safe practices in mRDT use
- Support uptake of mRDTs and adherence to results by providers
- Encourage population demand for mRDTs
- Recognize contextual factors not easily amenable to change

Next steps

- Organize follow up meetings to discuss additional findings from ongoing and new studies and pilots to develop more definitive recommendations for countries on best practices for scaling-up comprehensive fever case management in the private retail sector
- Systematically review the evidence (published and unpublished) of mRDT introduction in private retail sectors (Revised version to be re submitted by 12/31)

Preliminary findings

Study characteristics

- 12 studies included, all in Sub-Saharan Africa, except a study in Cambodia
- Almost all studies included registered but small-scale retailers with limited biomedical training (e.g. Drugshops, PPMV, ADDO, Chemical shops)
- Special permission was given to sell or perform mRDTs in most studies
- Substantial human and commodity resource support (i.e. training, supervision, referral program, demand generation, waste disposal)
- Studies showed variety in evaluation design but similar outcome focus:
 - RCTs (6), quasi experimental (4), observational (3)
 - Outcomes were targeting ACT treatment (RDT uptake, adherence, cost) rather than management of febrile illnesses

Outcomes

- Safe performance of RDTs: Range 16-99% (median ~93%)
- Uptake: Range 9-98% patients getting an RDT (median ~56%)
- Adherence to positive RDTs: Range 30-99% RDT positive patients getting an ACT (median ~86%)
- Adherence to negative RDTs: Range 2-51% RDT negative patients getting an antimalarial (median ~41%)
- Antibiotic use for RDT negative febrile cases reported: Range 4-58% (median ~13% v small numbers)

Conclusions

Performance

- Generally, providers were able to accurately and safely perform the mRDT
- Providers also seemed able to interpret mRDT results correctly

Uptake & Adherence

- All of the studies with both an intervention and a control arm showed a reduction in ACT usage in the intervention arm compared to the control arm, suggesting that increased testing rates can help reduce resource wastage
- Although a larger proportion of patients that tested positive received ACTs
 compared to those testing negative, the variety in adherence to positive and
 negative results show that it remains unclear whether mRDTs can improve
 targeting ACTs to only those that have malaria or not

Informing scale up

- Simply introducing mRDTs with minimal oversight appears to be ineffective in promoting appropriate (and therefore cost-effective) use
 - Studies with longer and interactive trainings, close and frequent supervision, low or no cost for the mRDT to the patient generally achieved higher uptake and adherence levels
- It remains unclear what makes up an optimal and cost effective training and supervision program that can meet 'real world' challenges (i.e. high staff turnover, referral system, waste management or data monitoring needs)

The studies to date have mainly been small-scale, closely controlled pilots and therefore do not provide sufficient insights into these requirements for large scale implementation.

ACKNOWLEDGEMENTS

Thank you to all contributors to this paper

Clare Chandler (ACTc), Jane Cunningham (WHO), Lawrence Barat (PMI) & Members of the Informal Private Sector Task Force of the RBM Case Management Working Group Members who contributed to the conceptualization/data, analysis/interpretation and writing of the paper: Richard Allan (Mentor Initiative), Evelyn Ansah (Ghana Health Service), Jennifer Anyanti (SFH) Ian Boulton (TropMed Pharma Consulting), Sian Clarke (LSHTM), Jessica Cohen (Harvard Public School of Health), Caitlin Dolkart (CHAI), Katie Eves (Mentor Initiative), Günther Fink (Harvard Public School of Health), Catherine Goodman (LSHTM)Sham Lal (LSHTM), Kathleen Maloney (CHAI), Anthony Mbonye (MoH UgandaErnest Nwokolo (SFH), Obi Onwujekwe (University of Enugu), Nora Petty (CHAI), Julie Pontarollo (Mentor Initiative), Edmund Rutta (MSH), David Schellenberg (LSHTM), Elizabeth Streat (MC), Abigail Ward (CHAI), Virginia Wiseman (LSHTM),), Shunmay Yeung (LSHTM)