

LONDON
SCHOOL of
HYGIENE
& TROPICAL
MEDICINE



Brady, OJ; Slater, HC; Pemberton-Ross, P; Wenger, E; Maude, RJ; Ghani, AC; Penny, MA; Gerardin, J; White, LJ; Chitnis, N; Aguas, R; Hay, SI; Smith, DL; Stuckey, EM; Okiro, EA; Smith, TA; Okell, LC (2017) Model citizen - Authors' reply. *Lancet Glob Health*, 5 (10). e974. ISSN 2214-109X DOI: [https://doi.org/10.1016/S2214-109X\(17\)30338-8](https://doi.org/10.1016/S2214-109X(17)30338-8)

Downloaded from: <http://researchonline.lshtm.ac.uk/4398440/>

DOI: [10.1016/S2214-109X\(17\)30338-8](https://doi.org/10.1016/S2214-109X(17)30338-8)

Usage Guidelines

Please refer to usage guidelines at <http://researchonline.lshtm.ac.uk/policies.html> or alternatively contact researchonline@lshtm.ac.uk.

Available under license: <http://creativecommons.org/licenses/by/2.5/>



Model citizen

Authors' reply

Tom Peto and colleagues point out, reasonably, that the effect of mass drug administration (MDA) on malaria could be affected substantially by patterns of human movement, and that our Article¹ does not consider the effects of the specific patterns of movement they observed in Cambodia and west Africa. The purpose of our Article¹ was to derive general results about the possible effect of MDA and to test how robust these are to the assumptions in different models, so we avoided assumptions about population movement that are specific to any particular place. Migration is among several factors that the different groups in our collaboration have investigated as potential modifiers of the effects of mass treatment in specific situations. For those analyses, we have used field data or realistic assumptions of migration rates.²⁻⁴ Human migration is a particularly complicated modifier; the endemicity of malaria in the places from which immigrants or temporary visitors originate could be important, not just the season and extent of migration.

A thorough empirical investigation of the implications of different patterns of migration for the effect of MDA would require a prohibitively complex set of field trials. Corresponding *in silico* analyses, parameterised with local field data, are more feasible but still represent an extensive piece of research in their own right, which is well beyond the scope of our Article.¹ Some of our research groups use extensive field data collection for this exact purpose.⁵ The questions addressed in our Article¹ were raised by WHO for their Evidence Review Group on mass treatment of malaria, which contains many experts who work directly on MDA interventions. Furthermore, several ongoing modelling exercises are being carried out by the authors of our

Article¹ in collaboration with control programmes to assess the potential effect of MDA in specific settings.⁶ We agree with Peto and colleagues on the importance of modelling groups and field researchers working closely together to better inform models and improve predictions of intervention outcomes. Involvement of modellers in trial design and operational planning for specific interventions allows questions to be framed as accurately as possible for relevant geographies and broader policy recommendations, and we hope that the number of these exercises will increase in the coming years.

ACG declares grant funding from the UK Medical Research Council (MRC), Bill & Melinda Gates Foundation, the Wellcome Trust, the Medicines for Malaria Venture, and WHO. She has also received consultancy contracts in the past 3 years from the Medicines for Malaria Venture, Oxford Policy Management, and The Global Fund to Fight AIDS, Tuberculosis and Malaria. EMS and EAO are or have been employed by the Bill & Melinda Gates Foundation. LCO declares grant funding from WHO, the Bill & Melinda Gates Foundation, and Medicines for Malaria Venture, and has received consultancy contracts from Medicines for Malaria Venture and WHO. All other authors declare no competing interests.

Copyright © The Author(s). Published by Elsevier Ltd. This is an Open Access article under the CC BY 4.0 license.

Oliver J Brady, Hannah C Slater,
Peter Pemberton-Ross,
Edward Wenger, Richard J Maude,
Azra C Ghani, Melissa A Penny,
Jaline Gerardin, Lisa J White, Nakul
Chitnis, Ricardo Aguas, Simon I Hay,
David L Smith, Erin M Stuckey,
Emelda A Okiro, Thomas A Smith,
*Lucy C Okell

lokell@imperial.ac.uk

Centre for the Mathematical Modelling of Infectious Diseases, Department of Infectious Disease Epidemiology, and Malaria Modelling Consortium, London School of Hygiene & Tropical Medicine, London, UK (OJB); MRC Centre for Outbreak Analysis and Modelling, Department of Infectious Disease Epidemiology, Imperial College, London W2 1PG, UK (HCS, ACG, LCO); Swiss Tropical and Public Health Institute, Basel, Switzerland (PP-R, MAP, NC, TAS); University of Basel, Basel, Switzerland (PP-R, MAP, NC, TAS); Institute for Disease Modeling, Bellevue, WA, USA (EW, JG); Mahidol Oxford Tropical Medicine Research Unit, Faculty of Tropical Medicine, Mahidol University, Bangkok, Thailand (RJM, LJW, RA); Centre for Tropical Medicine and Global Health (RJM, LJW, RA), and Oxford Big Data

Institute, Li Ka Shing Centre for Health Information and Discovery (SIH), University of Oxford, Oxford, UK; Harvard TH Chan School of Public Health, Harvard University, Boston, MA, USA (RJM); Malaria Modelling Consortium (SIH), and Institute for Health Metrics and Evaluation (SIH, DLS), University of Washington, Seattle, WA, USA; Malaria Modelling Consortium, Bill & Melinda Gates Foundation, Seattle, WA, USA (EMS, EAO); and Kemri Wellcome Trust Research Programme, Nairobi, Kenya (EAO)

- 1 Brady OJ, Slater HC, Pemberton-Ross P, et al. Role of mass drug administration in elimination of *Plasmodium falciparum* malaria: a consensus modelling study. *Lancet Glob Health* 2017; **5**: e680-87.
- 2 Crowell V, Briet OJ, Hardy D, et al. Modelling the cost-effectiveness of mass screening and treatment for reducing *Plasmodium falciparum* malaria burden. *Malar J* 2013; **12**: 4.
- 3 Gerardin J, Bever CA, Bridenbecker D, et al. Effectiveness of reactive case detection for malaria elimination in three archetypal transmission settings: a modelling study. *Malar J* 2017; **16**: 248.
- 4 Silal SP, Little F, Barnes KI, White LJ. Hitting a moving target: a model for malaria elimination in the presence of population movement. *PLoS One* 2015; **10**: e0144990.
- 5 Marshall JM, Toure M, Ouedraogo AL, et al. Key traveller groups of relevance to spatial malaria transmission: a survey of movement patterns in four sub-Saharan African countries. *Malar J* 2016; **15**: 200.
- 6 Stuckey EM, Miller JM, Littrell M, Chitnis N, Steketee R. Operational strategies of anti-malarial drug campaigns for malaria elimination in Zambia's southern province: a simulation study. *Malar J* 2016; **15**: 148.