



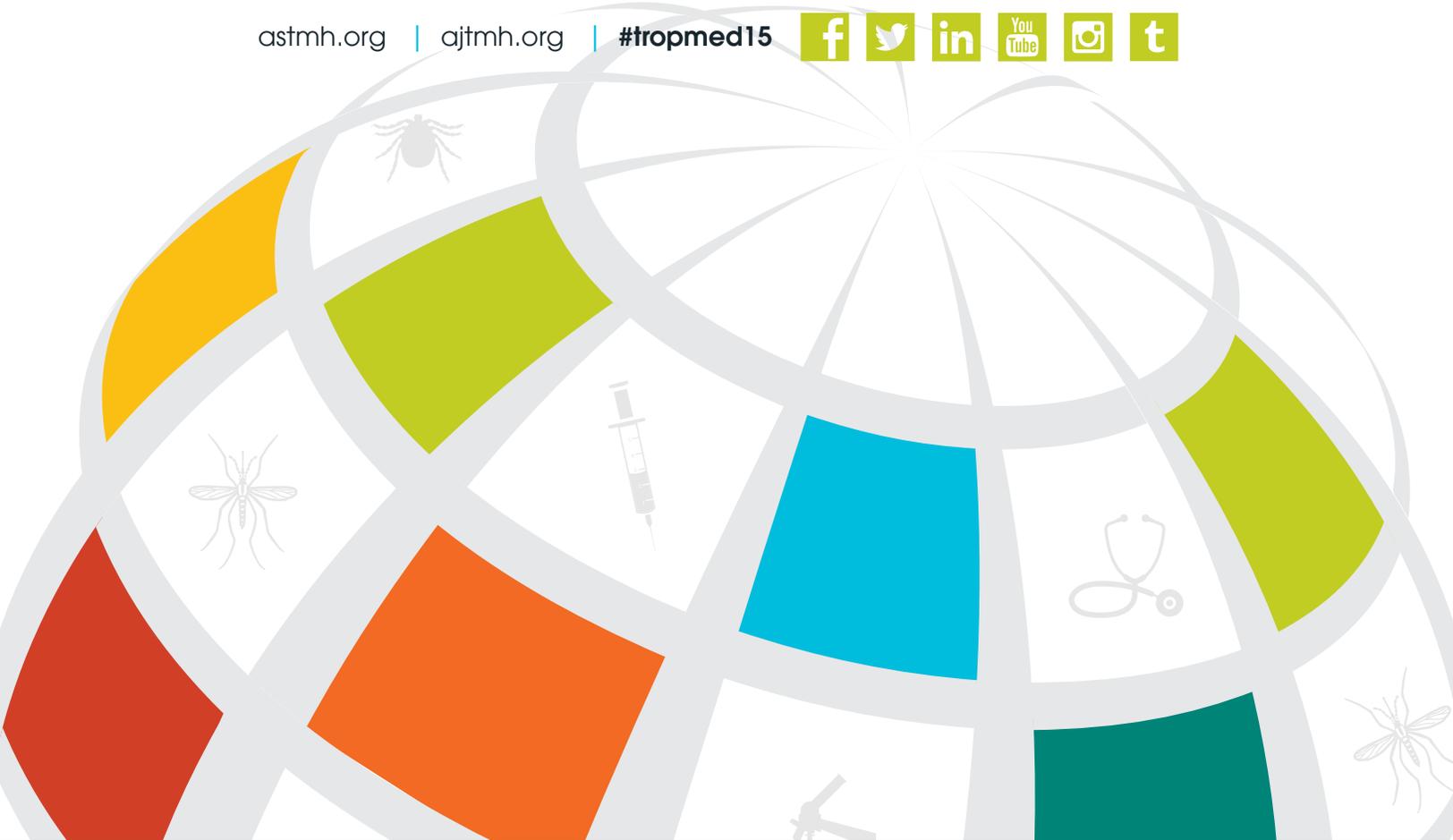
AMERICAN SOCIETY OF TROPICAL MEDICINE & HYGIENE
ADVANCING GLOBAL HEALTH SINCE 1903

VOLUME 93 OCTOBER 2015 NUMBER 4 SUPPLEMENT

64th Annual Meeting **October 25–29, 2015**

PHILADELPHIA MARRIOTT DOWNTOWN, PHILADELPHIA, PENNSYLVANIA USA

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Supplement to

The American Journal of Tropical Medicine and Hygiene

Sponsored Symposia

Dihydroartemisinin/Piperaquine and the Latest Achievements in Uncomplicated Malaria Treatment

Sponsored by Sigma Tau/Pierre Fabre

Monday, October 26, 7:15 p.m. – 9 p.m.

Marriott – Grand Ballroom Salon AB

Considerable increases in the availability and use of artemisinin-based combination therapies (ACTs) together with increased deployment of insecticide treated bed-nets (ITN) have resulted in a substantial fall in global malaria morbidity and mortality. These gains and the prospects for malaria elimination are now threatened by the emergence of artemisinin and/or partner drug resistance in *Plasmodium falciparum*. In the context of these resistance scenarios, the role of high quality artemisinin-based combinations will be discussed. As the only stringent-regulatory-approved Dihydroartemisinin/Piperaquine offering in the market today, it has a significant role to play in malaria endemic countries. Both prior to and subsequent to Dihydroartemisinin/Piperaquine's approval by the EMA, this medicine has been the focus of robust clinical data, not only in terms of efficacy, but also of safety. At this symposium, new safety data will be presented: daily clinical practice with falciparum malaria treatment in several African countries on over 10,000 patients, as well as on over 4,000 pregnant women with Dihydroartemisinin/Piperaquine. Recent studies in Southeast Asia and in Africa have focused on the pharmacokinetics and the pharmacodynamics of piperaquine, the partner drug that is crucial for cure. These studies indicate that the dose of piperaquine recommended in young children is suboptimal, because the kinetics of this drug are different in young children compared to adults. To cover this issue, the recent clinical outcomes of novel Dihydroartemisinin/Piperaquine pediatric dispersible formulations will be reported.

CHAIR

George Jagoe, Executive Vice-President

Medicines for Malaria Venture, Geneva, Switzerland

Modelling the Impact of DHA-PQP on Malaria Transmission and Comparison with other ACTs

Lucy C. Okell

MRC Centre for Outbreak Analysis and Modelling, Department of Infectious Disease Epidemiology, Imperial College, London, United Kingdom

The Safety and Efficacy of Four Artemisinin-Based Combination Treatments in Pregnant African Women with Malaria

Umberto D'Alessandro

Medical Research Council, Banjul, The Gambia

Clinical Development of the Antimalarial Dihydroartemisinin/Piperaquine in Manhiça, Mozambique; Past, Present and Future

Quique Bassat

Barcelona Institute of Global Health (ISGlobal) and Centro de Investigación e Saude da Manhiça, Barcelona, Spain

The Burden of Drug Resistant *Plasmodium falciparum* Malaria: Where Does Dihydroartemisinin/Piperaquine Stand?

Rick Fairhurst

National Institute of Allergy and Infectious Diseases, Malaria Genetics Section (HNM2A3), Rockville, MD, United States