

Sunday, November 17

Registration

Convention Center - Lobby I (1st Floor)
Sunday, November 17, 7 a.m. - 10:30 a.m.

Speaker Ready Room

Convention Center - Room 387 (3rd Floor)
Sunday, November 17, 7 a.m. - Noon

Meeting Sign-Up Room

Hilton – Norwich Room and Windsor Room (3rd Floor)
Sunday, November 17, 7 a.m. – 1:15 p.m.

Nursing Mothers Room

Convention Center – Office I120 and Office J121 (1st Floor)
Sunday, November 17, 7 a.m. – 1:15 p.m.

Prayer Room

Convention Center - Room 342 (3rd Floor)
Sunday, November 17, 7 a.m. – 1:15 p.m.



Symposium 165

Reaching and Sustaining Elimination: Global Guidance and Regional Best Practices on Malaria Elimination and the Prevention of Re-establishment (PoR)

Convention Center - Hall I-2 (1st Floor)
Sunday, November 17, 8 a.m. - 9:45 a.m.

The global fight against malaria has led to marked progress towards the long-term goal of malaria eradication, or complete interruption of malaria transmission throughout the world. To date, the WHO has certified a total of 43 countries and one territory as malaria-free and another three countries have achieved the milestone of zero indigenous cases. Until malaria eradication, resurgence remains a threat post-elimination. Despite the risk, countries that have achieved elimination have largely maintained their malaria-free status – a study in 2010 showed that 50 of the 99 historical malaria elimination programs that successfully achieved malaria elimination, only four had experienced resurgence and two of these four were able to eliminate malaria once again. A key component in this success has been strong prevention of re-establishment (POR) programs which are not only a requirement for WHO malaria-free certification but also a critical prerequisite to sustaining subnational and national elimination, and eventually achieving malaria eradication. The Greater Mekong Sub-region (GMS) in Southeast Asia has achieved impressive progress towards malaria elimination over the last decade with cases declining by 70% since 2013. In 2023, Cambodia and Lao PDR reached historical low level of malaria transmission in each

country, reporting 1,384, and 805 cases, respectively. As these countries near elimination, planning for POR has begun to prepare for WHO malaria-free certification and to sustain gains made over the last decade. On the other hand, Thailand, which was targeted by WHO as one of the countries for elimination by 2025 has seen cases increase sharply over the past two years, largely in areas on the border with Myanmar. This increase has stress tested the program's roll out of their POR guidelines and despite the setbacks, has generated several key lessons learned for POR. This symposium will convene representatives from the WHO and three national malaria control programs (NMCPs) in the GMS to share updated global guidelines and recent regional progress on POR. To start, the WHO Global Malaria Program (GMP) will introduce the latest global POR guidelines and their relevance to country elimination programs and malaria-free certification. Cambodia will present their process for developing their newly created POR guidelines. Lao PDR will then share their experience responding to a malaria outbreak in elimination area in 2023. To conclude, Thailand will discuss the challenges they have faced as a leader in the region on designing and implementing POR in the context of the recent rise in cases on the border. #Elimination #Epidemiology #InfectiousDisease

CHAIR

Elkhan Gasimov
World Health Organization, Geneva, Switzerland

Yang Hu
Clinton Health Access Initiative, Phnom Penh, Cambodia

8 a.m. INTRODUCTION

8:10 a.m. NEW WHO GLOBAL PREVENTION OF RE-ESTABLISHMENT (POR) GUIDELINES

Elkhan Gasimov
World Health Organization, Geneva, Switzerland

8:30 a.m. CAMBODIA: LOCALIZING GLOBAL GUIDANCE TO COUNTRY CONTEXT

Siv Sovannaroth
National Center for Parasitology, Entomology, and Malaria Control, Phnom Penh, Cambodia

8:50 a.m. THAILAND: LESSONS LEARNED FROM SUBNATIONAL POR IMPLEMENTATION

Rungrawee Tipmontree
Division of Vector Borne Diseases, DDC, MoPH, Thailand, Bangkok, Thailand

9:10 a.m. LAO PDR: RESPONDING TO OUTBREAKS IN ELIMINATION AREAS

Vonethalom Thongpaseuth
Center of Malariaology, Parasitology and Entomology, Laos, Vientiane, Lao People's Democratic Republic

Scientific Session 166

Bacteriology: Other Enteric Infections

Convention Center - Room 343/344 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

#InfectiousDisease #Diagnostics #Therapeutics
#ClinicalResearch #Epidemiology

CHAIR

Jackie Knee

London School of Hygiene & Tropical Medicine, London, United Kingdom

Stuart Torres Ayala

Universidad San Francisco de Quito, Quito, Ecuador

8 a.m.

8371

EVALUATION OF *IN SILICO* SHIGELLA SEROTYPING TOOLS USING A GLOBAL SHIGELLA COLLECTION

Xiaoliang Ba, Charlotte E. Chong, Fahad Khokhar, Mark A. Holmes, Kate S. Baker
University of Cambridge, Cambridge, United Kingdom

8:15 a.m.

8372

IDENTIFYING OPTIMAL ENDPOINT DEFINITIONS TO MINIMIZE OUTCOME MISCLASSIFICATION IN UPCOMING SHIGELLA VACCINE TRIALS

Maria Garcia Quesada¹, Benjamin A. Lopman¹, Patricia B. Pavlinac², James A. Platts-Mills³, Lance A. Waller¹, Elizabeth T. Rogawski McQuade¹

¹Emory Rollins School of Public Health, Atlanta, GA, United States, ²University of Washington, Seattle, WA, United States, ³University of Virginia, Charlottesville, VA, United States

8:30 a.m.

8373

MULTIPLEX PCR DETECTION OF ENTERIC PATHOGENS IN A COMMUNITY-BASED BIRTH COHORT IN ECUADOR: A COMPARISON OF XTAG-GPP AND TAQMAN ARRAY CARD ASSAYS

Stuart Torres Ayala¹, Lesly Simbaña Vivanco¹, Nikolina Eve Walas², Kelsey Jesser³, Nicolette A. Zhou³, Christine S. Fagnant-Sperati³, Hadley Burroughs⁴, Gwenyth O. Lee⁵, Joseph N.S. Eisenberg⁶, Karen Levy³, Gabriel Trueba¹, Benjamin F. Arnold⁴

¹Microbiology Institute, Universidad San Francisco de Quito, Quito, Ecuador, ²Department of Environmental Health, School of Public Health, University of California Berkeley, Berkeley, CA, United States, ³Department of Environmental & Occupational Health Sciences, University of Washington, Seattle, WA, United States, ⁴Francis I. Proctor Foundation, University of California San Francisco, San Francisco, CA, United States, ⁵Rutgers Global Health Institute, Rutgers, the State University of New Jersey, New Brunswick, NJ, United States, ⁶Department of Epidemiology, School of Public Health, University of Michigan, Ann Arbor, MI, United States

8:45 a.m.

8374

OPTIMAL AZITHROMYCIN TREATMENT RULES FOR CHILDREN WITH WATERY DIARRHEA IN THE ANTIBIOTICS FOR CHILDREN WITH SEVERE DIARRHEA (ABCD) TRIAL

Sara S. Kim¹, Allison Codi², James Platts-Mills³, Patricia B. Pavlinac⁴, Karim Manji⁵, Christopher R. Sudfeld⁶, Queen Dube⁷, Naor Bar-Zeev⁸, Karen Kotloff⁹, Samba O. Sow¹⁰, Sunil Sazawal¹¹, Benson O. Singa¹², Farah Qamar¹³, Judd Walson⁴, Ayesha De Costa¹⁴, David Benkeser², Elizabeth T. Rogawski McQuade¹

¹Department of Epidemiology, Rollins School of Public Health, Emory University, Atlanta, GA, United States, ²Department of Biostatistics, Rollins School of Public Health, Emory University, Atlanta, GA, United States, ³Division of Infectious Diseases and International Health, Department of Medicine, University of Virginia, Charlottesville, VA, United States, ⁴Department of Global Health, Department of Epidemiology, University of Washington,

Seattle, WA, United States, ⁵Department of Pediatrics and Child Health, Muhimbili University of Health and Allied Sciences, Dar es Salaam, United Republic of Tanzania, ⁶Department of Global Health and Population, Harvard T.H. Chan School of Public Health, Boston, MA, United States, ⁷Department of Pediatrics, Queen Elizabeth Central Hospital, Blantyre, Malawi, ⁸International Vaccine Access Center, Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States, ⁹Department of Pediatrics, Department of Medicine, Center for Vaccine Development and Global Health, University of Maryland School of Medicine, Baltimore, MD, United States, ¹⁰Centre pour le Développement des Vaccins, Bamako, Mali, ¹¹Center for Public Health Kinetics, New Delhi, India, ¹²Kenya Medical Research Institute, Nairobi, Kenya, ¹³Department of Pediatrics and Child Health, Aga Khan University, Karachi, Pakistan, ¹⁴Department of Maternal, Child, and Adolescent Health and Aging, World Health Organization, Geneva, Switzerland

9 a.m.

8375

ENTERIC PATHOGEN DETECTION AMONG CHILDREN DISCHARGED FROM OUTPATIENT TREATMENT FOR SEVERE ACUTE MALNUTRITION AND ASSOCIATIONS WITH SUBSEQUENT RELAPSE IN SOUTH SUDAN

Jackie Knee¹, Lauren D'Mello-Guyett¹, Lynn Grignard¹, Alesha Myers¹, Sarah King², John Agong³, Lino Deng³, Mesfin Gose³, Jackson Lwate Hasan³, Nancy Grace Lamaka³, Anastasia Marshak⁴, Dimple Save³, Indi Trehan⁵, Khamisa Ayoub⁶, Heather Stobaugh², Oliver Cumming¹

¹London School of Hygiene & Tropical Medicine, London, United Kingdom, ²Action Against Hunger USA, New York, NY, United States, ³Action Against Hunger South Sudan, Juba, South Sudan, ⁴Tufts University, Boston, MA, United States, ⁵University of Washington, Seattle, WA, United States, ⁶Ministry of Health for the Republic of South Sudan, Juba, South Sudan

9:15 a.m.

8376

HIGH LEVELS OF GUT BIFIDOBACTERIUM ASSOCIATED WITH INTESTINAL INFLAMMATION AND FECAL METABOLITES IN CHILDREN IN RURAL BANGLADESH

Jessica A. Grembi¹, Elizabeth Costello¹, Brian C. DeFelice², Anna Robaczewska¹, Audrie Lin³, Md. Ziaur Rahman³, Shahjahan Ali⁴, Md. Zahir Hussain⁵, Rashidul Haque⁶, Mahbubur Rahman⁶, Leanne Unicomb⁶, Stephen Luby¹, Susan P. Holmes¹, David Relman¹

¹Stanford University, Stanford, CA, United States, ²Chan Zuckerberg BioHub, San Francisco, CA, United States, ³University of California, Santa Cruz, Santa Cruz, CA, United States, ⁴University of Colorado, Denver, CO, United States, ⁵University of Pittsburgh, Pittsburgh, PA, United States, ⁶International Centre for Diarrhoeal Disease Research, Bangladesh, Dhaka, Bangladesh

9:30 a.m.

8377

NEONATAL ACQUISITION OF ESBL-PRODUCING ENTEROBACTERIALES IN MADAGASCAR AND CAMBODIA

Anne-Lise Beaumont¹, Agathe de Lauzanne², Mamitiana Alain Noah Rabenandrasana³, Norohasina Fanja Randriamanga³, Sandrine Bernabeu⁴, Alexis Criscuolo⁵, Perlinot Herindrainy³, Didier Guillemot¹, Jean-Marc Collard³, Long Pring², Laurence Borand², Elsa Kermovant-Duchemin⁶, Muriel Vray¹, Tania Crucitti³, Bich-Tram Huynh¹

¹Institut Pasteur, Université de Versailles-Saint Quentin, INSERM, Paris, France, ²Institut Pasteur du Cambodge, Phnom Penh, Cambodia, ³Phnom Penh, Cambodia, ⁴Institut Pasteur de Madagascar, Antananarivo, Madagascar, ⁵Assistance Publique - Hôpitaux de Paris, Le Kremlin-Bicêtre, France, Paris, France, ⁶Institut Pasteur, Paris, France, ⁷Assistance Publique - Hôpitaux de Paris, Hôpital Necker-Enfants malades, Paris, France, ⁸Institut Pasteur du Sénégal, Dakar, Senegal



Symposium 167

Contemporary Ethical Issues in Tropical Medicine: Content for Research Training from Peru and Mozambique

Convention Center - Room 345 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

Successful biomedical research and international collaboration require strong and well-staffed research support systems and an institutional infrastructure grounded in ethical best practices. While the research environment has matured significantly in low- and middle-income countries (LMICs) like Peru and Mozambique, research ethics training and oversight capacity still has room for improvement. Efforts are underway in many LMICs to close recognized gaps and reinforce sustainable systems that ensure locally led research ethics review as well as forums for LMICs to contribute to global ethical discourse. Investigators and research institutions in LMICs confront many of the same challenging ethical issues faced in high-resource settings, and face similar needs for education in research ethics education. These challenges include assuring research integrity in publications, obtaining meaningful informed consent in contexts of imbalanced power dynamics, and equitable conduct of clinical trials. Recognizing the need to address these challenges, LMIC-based educational programs, such as the NIH Fogarty International Center's International Research Ethics Education and Curriculum Development R25 program, have provided opportunities to reinforce research ethics education and expertise in LMIC institutions. In this symposium, faculty involved in the design and implementation of R25 programs in Mozambique and Peru will share examples of challenges they have faced in research ethics, insights on stakeholder's perceptions of needed research ethics training, and comments about both the benefits and limitations of research ethics education programs in LMIC settings.

#ClinicalResearch #FieldStudies #SocialScience

CHAIR

Richard A. Oberhelman
Tulane School of Public Health and Tropical Medicine, New Orleans, LA, United States

Theresa Ochoa
Instituto de Medicina Tropical Alexander von Humboldt; Universidad Peruana Cayetano Heredia, Lima, Peru

8 a.m. INTRODUCTION

8:10 a.m. LESSONS FROM THE COVID-19 PANDEMIC AND STAKEHOLDER'S PERCEPTIONS OF RESEARCH ETHICS TRAINING NEEDS IN PERU

Cynthia S. Hurtado
Universidad Peruana Cayetano Heredia, Lima, Peru

8:35 a.m.

THE PAPER TRAIL: NAVIGATING THE COMPLEX LANDSCAPE OF AUTHORSHIP AND PAPER MILLS IN GLOBAL HEALTH

Valerie Paz-Soldan
Tulane School of Public Health and Tropical Medicine—Health Office for Latin America, Lima, Peru

8:50 a.m.

ETHICAL LESSONS LEARNED DURING THE COVID-19 PANDEMIC IN MOZAMBIQUE

Esperança Sevene
University Eduardo Mondlane, Maputo, Mozambique

9:05 a.m.

VACCINE INDUCED SEROPOSITIVITY FOLLOWING HIV VACCINE STUDIES IN MOZAMBIQUE. IMPLICATIONS FOR INFORMED CONSENT IN A HIGH HIV BURDEN COUNTRY

D. Troy Moon
Tulane School of Public Health and Tropical Medicine, New Orleans, LA, United States

9:20 a.m.

THE NIH FOGARTY INTERNATIONAL CENTER'S R25 PROGRAM IN INTERNATIONAL RESEARCH ETHICS EDUCATION AND CURRICULAR DEVELOPMENT: GOALS AND LESSONS

Elizabeth Heitman
University of Texas Southwestern Medical Center, Dallas, TX, United States

Scientific Session 168

Kinetoplastida and Other Opportunistic and Anaerobic Protozoa: Genomics, Proteomics and Metabolomics, Molecular Therapeutic Targets, Treatment, Drug Delivery, Drug Repurposing and Drug Discovery

Convention Center - Room 352 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

This session does not carry CME credit.

#InfectiousDisease #Therapeutics #FieldStudies

CHAIR

Byron Arana
Drugs for Neglected Diseases Initiative, Geneva, Switzerland

Louisa Messenger
University of Nevada, Las Vegas, Las Vegas, NV

8 a.m.

8378

EFFICACY AND SAFETY OF THERMOTHERAPY IN COMBINATION WITH MILTEFOSINE IN COMPARISON TO MILTEFOSINE MONOTHERAPY FOR THE TREATMENT OF CUTANEOUS LEISHMANIASIS IN THE AMERICAS: A PHASE III, OPEN LABEL, MULTICENTER, RANDOMIZED TRIAL

Joelle Rode¹, Juan Pascalle², Alejandro Llanos-Cuentas³, Marcia Hueb⁴, Jaime Soto⁵, Glaucia Cota⁶, Paulo Machado⁷, Marina Boni¹, Bethania Blum¹, **Byron Arana**⁸

¹Drugs for Neglected Diseases Initiative, Rio de Janeiro, Brazil, ²Instituto Conmemorativo Gorgas de Estudios de la Salud, Panama, Panama, ³Universidad Peruana Cayetano Heredia, Lima, Peru, ⁴Julio Muller University Hospital, Cuiaba, Brazil, ⁵Fundación Nacional de Dermatología, Santa Cruz de la Sierra, Plurinational State of Bolivia, ⁶Instituto René Rachou, Belo Horizonte, Brazil, ⁷Federal University of Bahia, Salvador Bahia, Brazil, ⁸Drugs for Neglected Diseases Initiative, Geneva, Switzerland

8:15 a.m.

8379

FIRST-IN-HUMAN, RANDOMIZED, DOUBLE BLIND CLINICAL TRIAL OF LXE408 FOR KINETOPLASTID DISEASES

Natasha Hochberg¹, Li Ruobing², Amanda Nguyen¹, Christopher Breen³, Jean Lecot⁴, Andrea de Soyers⁵, Linda Armstrong³, Thierry Diagana⁶, Peter Pertel¹, Megan Barton¹, Florencia Segal¹, Srinivasa Rao⁶

¹Novartis Global Health, BioMedical Research, Cambridge, MA, United States, ²Novartis Global Health, BioMedical Research, Shanghai, China, ³Novartis Global Health, BioMedical Research, East Hannover, NJ, United States, ⁴Novartis Global Health, Analytics-Global Medical Affairs, Basel, Switzerland, ⁵Novartis Global Health, BioMedical Research, Basel, Switzerland, ⁶Novartis Global Health, BioMedical Research, Emeryville, CA, United States

8:30 a.m.

8380

TOPICAL APPLICATION OF AC2-26, AN ANNEXIN A1 PEPTIDOMIMETIC, REDUCES LESIONS AND IMPROVES IMMUNE RESPONSES IN A MURINE MODEL OF LEISHMANIA AMAZONENSIS INFECTION

Liliane M. dos Santos, Tiago Nery, Gisele Goulart, Ana Paula Fernandes
Universidade Federal de Minas Gerais, Belo Horizonte, Brazil

8:45 a.m.

8381

SEVERE ANAEMIA AND HAEMOGLOBIN TRAJECTORY FOLLOWING TREATMENT OF VISCERAL LEISHMANIASIS: AN INDIVIDUAL PATIENT DATA META-ANALYSIS USING THE INFECTIOUS DISEASES DATA OBSERVATORY DATA PLATFORM

Prabin Dahal¹, Abdalla Munir², Rishikesh Kumar³, Sauman Singh-Phulgenda¹, Niyamat Ali Siddiqui³, James Wilson¹, Gemma Buck¹, Caitlin Naylor¹, Matthew Brack¹, Manju Rahi⁴, Paritosh Malaviya⁵, Monique Wasunna⁶, Francois Chappuis⁷, Koert Ritmeijer⁸, Carlos Costa⁹, Gustavo Romero¹⁰, Vassiliki Syriopoulou¹¹, Fabiana Alves¹², Kasia Stepniewska¹, Krishna Pandey³, Ahmed Musa¹³, Philippe J. Guerin¹

¹University of Oxford, Oxford, United Kingdom, ²Infectious Diseases Data Observatory (IDDO), Oxford, United Kingdom, ³ICMR - Rajendra Memorial Research Institute of Medical Sciences (RMRIMS), Patna, India, ⁴Indian Council of Medical Research (ICMR), New Delhi, India, ⁵Infectious Disease Research Laboratory, Department of Medicine, Institute of Medical Sciences, Banaras Hindu University, Varanasi, India, ⁶Drugs for Neglected Diseases initiative, Nairobi, Kenya, ⁷Division of Tropical and Humanitarian Medicine, Geneva University Hospitals, Geneva, Switzerland, ⁸Médecins Sans Frontières, Amsterdam, Netherlands, ⁹Department of Community Medicine, Federal University of Piauí, Piauí, Brazil, ¹⁰Center for Tropical Medicine, University of Brasilia, Brasilia, Brazil, ¹¹Infectious Diseases and Chemotherapy Research Laboratory, Medical School, National and Kapodistrian University of Athens, Athens, Greece, ¹²Drugs for Neglected Diseases initiative, Geneva, Switzerland, ¹³Institute of Endemic Diseases, University of Khartoum, Khartoum, Sudan

9 a.m.

8382

EFFICACY OF SHORT-COURSE TREATMENT FOR PREVENTION OF CONGENITAL TRANSMISSION OF CHAGAS DISEASE: A RETROSPECTIVE COHORT STUDY

Moscatelli Guillermo, Samanta Moroni, Juan Carlos Ramirez, Belen Warszatska, Fernanda Lascano, Nicolas Gonzalez, Andres Rabinovich, Jaime ALTCHER
Hospital de Niños R. Gutierrez, Buenos Aires, Argentina

9:15 a.m.

8383

IMPACT OF ASCARIDOLE ON METABOLIC BIOENERGETICS IN LEISHMANIASIS

Deblina Sarkar, Sritama De Sarkar, Deep Goswami, Chaitali Karmakar, Prof. Mitali Chatterjee
IPGME&R, Kolkata, Kolkata, India

9:30 a.m.

8384

INVOLVING PATIENTS IN DRUG DEVELOPMENT FOR NEGLECTED TROPICAL DISEASES (NTDS): A QUALITATIVE STUDY EXPLORING AND INCORPORATING PREFERENCES OF PATIENTS WITH CUTANEOUS LEISHMANIASIS INTO TARGET PRODUCT PROFILE DEVELOPMENT

Maria del Mar Castro¹, Astrid C. Erber², Byron Arana³, Glauca Cota⁴, Claudia Denkinge⁵, Nicole Harrison⁶, Julia Kutyl⁶, Liliana López-Carvajal⁷, Emma Plugge⁸, Julia Walochnik⁹, Piero Olliaro¹⁰

¹Centro Internacional de Entrenamiento de Investigaciones Médicas (CIDEIM), Cali, Colombia, ²Department of Epidemiology, Center for Public Health, Medical University of Vienna, Vienna, Austria, ³Drugs for Neglected Diseases initiative (DNDi), Geneva, Switzerland, ⁴Instituto René Rachou (IRR), Fundação Oswaldo Cruz (FIOCRUZ), Minas Gerais, Brazil, ⁵Center of Infectious Diseases, Heidelberg University Hospital, Heidelberg, Germany, ⁶Division of Infectious Diseases and Tropical Medicine, Department of Medicine I, Medical University of Vienna, Vienna, Austria, ⁷Programa de Estudio y Control de Enfermedades Tropicales (PECET), Universidad de Antioquia, Medellín, Colombia, ⁸Primary Care, Population Sciences and Medical Education, University of Southampton, Southampton, United Kingdom, ⁹Institute of Specific Prophylaxis and Tropical Medicine, Center for Pathophysiology, Infectiology and Immunology, Medical University of Vienna, Vienna, Austria, ¹⁰International Severe Acute Respiratory and Emerging Infection Consortium, Pandemic Sciences Institute, University of Oxford, Oxford, United Kingdom

Symposium 169

Novel Approaches for the Radical Cure of Vivax Malaria: High Dose Short Course Primaquine and Tafenoquine

Convention Center - Room 353 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

Plasmodium vivax causes between 4 and 11 million cases of malaria annually and represents an increasing proportion of malaria in > 40 countries co-endemic for *P. falciparum* and *P. vivax*. The radical cure of *P. vivax* requires treatment of both the blood stage, which causes acute symptoms, and the dormant liver stage (hypnozoites), which reactivate and cause relapses weeks to months later. Relapsing vivax malaria results in a cumulative risk of anaemia and is associated with direct and indirect morbidity and the potential for further transmission. For the last 60-70 years, *P. vivax* radical cure has relied on chloroquine (for the blood stage) and primaquine (for the liver stage). Primaquine has been used at a low total dose (3.5mg/kg) over 14-days to reduce the risk of drug induced haemolysis in patients who are G6PD deficient; this regimens effectiveness is confounded by poor adherence. Higher total doses of primaquine have been shown to be more efficacious. The therapeutic options for *P. vivax* are rapidly changing with the availability of tafenoquine, shorter course (higher daily dose) primaquine regimens, and the development of point-of-care tests for G6PD activity. Shorter course primaquine regimens are hypothesized to improve adherence, but data to support this has been lacking. Our pragmatic randomized controlled trial (960 patients, 4 countries) provides crucial data on the effectiveness and safety of high dose short course primaquine when unsupervised and compares this to the low dose 14-day treatment regimen. We will also present comparative effectiveness and safety data for single dose tafenoquine. We will also present highly anticipated data on the efficacy of tafenoquine with two different Artemisinin Combination Therapies (ACTs). Previous

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studies of tafenoquine and ACTs have shown poor antirelapse efficacy which has been attributed to a drug-drug interaction, but alternative explanations have also been proposed. Data presented will provide important information on the comparative efficacy of both primaquine and tafenoquine when administered with ACT (DHA-Piperaquine or pyronaridine-artesunate) versus chloroquine. We will provide insights from qualitative research in Cambodia and Ethiopia to inform the design of effective, feasible and acceptable implementation strategies for new radical cure treatment strategies, including G6PD testing and early clinical review. New interventions must be cost-effective, both for local health services and for the global effort towards malaria elimination. In the final talk a cost-effectiveness analysis will be presented. The talks represent as yet unpublished data and will allow time for a panel discussion of the proposed strategies and the challenges of delivering radical cure. #ClinicalResearch, #Elimination #InfectiousDisease #Therapeutics #TranslationalScience

CHAIR

Kamala Thriemer
Menzies School of Health Research, Darwin, Australia

Lek Dysoley
National Center for Parasitology, Entomology and Malaria Control, Phnom Phen, Cambodia

8 a.m. INTRODUCTION

8:10 a.m. EFFECTIVENESS OF UNSUPERVISED HIGH DOSE SHORT COURSE PRIMAQUINE AND TAFENOQUINE

Tamiru T. Degaga
Arba Minch University, Arba Minch, Ethiopia

8:20 a.m. SAFETY OF UNSUPERVISED HIGH DOSE SHORT COURSE PRIMAQUINE AND TAFENOQUINE

Najia Ghanchi
Aga Khan University, Karachi, Pakistan

8:30 a.m. TAFENOQUINE AND ACTS – EVIDENCE FROM INDONESIA AND CAMBODIA

Ayodhia Pitaloka Pasaribu
Universitas Sumatera Utara, Medan, Indonesia

8:40 a.m. INFORMING IMPLEMENTATION STRATEGIES FOR THE SAFE PROVISION OF HIGH DOSE SHORT COURSE PRIMAQUINE AND TAFENOQUINE

Sarah Cassidy-Seyoum
Menzies School of Health Research, Darwin, Australia

8:55 a.m. THE COST-EFFECTIVENESS OF HIGH DOSE SHORT COURSE PRIMAQUINE AND TAFENOQUINE

Angela A. Devine
University of Melbourne, Melbourne, Australia

Scientific Session 170

American Committee of Molecular Cellular and Immunoparasitology (ACMCIP): Parasite Vaccine Development

Convention Center - Room 354/355 (3rd Floor)
Sunday, November 17, 8 a.m. - 9:45 a.m.

Supported with funding from the Burroughs Wellcome Fund

#ClinicalResearch #Vaccinology #TranslationalScience
#Immunology #InfectiousDisease

CHAIR

Nathan Schmidt
Indiana University School of Medicine, Indianapolis, IN, United States

Thalia Pacheco-Fernandez
Food and Drug Administration, Silver Spring, MD, United States

8 a.m. 8385

DEVELOPMENT OF NOVEL HOOKWORM MRNA VACCINE CANDIDATES BY ALTERING THE INTRACELLULAR TRAFFICKING OF *NECATOR AMERICANUS* GST-1 ANTIGEN

Athos Silva de Oliveira¹, Leroy Versteeg¹, Neima Briggs², Rakesh Adhikari¹, Maria Jose Villar¹, Peter Hotez¹, Maria Elena Bottazzi¹, Jeroen Pollet¹

¹Texas Children's Hospital Center for Vaccine Development & Department of Pediatrics, National School of Tropical Medicine, Baylor College of Medicine, Houston, TX, United States, ²Departments of Immunobiology and Internal Medicine, Yale University, New Haven, CT, United States

(ACMCIP Abstract)

8:15 a.m. 8386

CONTROL OF HOOKWORMS USING *BACILLUS THURINGIENSIS* CRY PROTEINS AND VACCINES

Qian Ding¹, Hanchen Li¹, Kelly A. Flanagan¹, Duy Hoang¹, Nicholas R. Cazeault¹, Florentina Rus¹, Ernesto Soto¹, Erich M. Schwartz², Jane Homan³, Wenbin Tuo⁴, Gary R. Ostroff¹, Raffi V. Aroian¹

¹Umass Chan Medical School, Worcester, MA, United States, ²Cornell University, Ithaca, NY, United States, ³ioGenetics LLC, Madison, WI, United States, ⁴United States Department of Agriculture, Beltsville, MD, United States

(ACMCIP Abstract)

8:30 a.m. 8387

OPISTHORCHIS VIVERRINI ANTI-CANCER VACCINE TARGETING THE LIVER FLUKE HOST-PARASITE INTERFACE

Michael J. Smout¹, Sujitra Chaiyadet², Wannaporn Ittiprasert³, Sutas Suttiaprapa², Paul J. Brindley³, Thewarach Laha², Alex Loukas¹

¹James Cook University, Cairns, Australia, ²Khon Kaen University, Khon Kaen, Thailand, ³George Washington University, Washington DC, DC, United States

(ACMCIP Abstract)

8:45 a.m. 8388

A LIVE-ATTENUATED *LEISHMANIA* VACCINE SHAPES THE CELLULAR RESPONSE IN THE BONE MARROW

Thalia Pacheco-Fernandez, Nazli Azodi, Spyros Karaiskos, Laura Klenow, Hannah Markle, Luis DaSilva-Pereira, Luis Santana-Quintero, Alexander Murray, Hira Nakhasi, Sreenivas Gannavaram

Food and Drug Administration, Silver Spring, MD, United States

(ACMCIP Abstract)

9 a.m.

8389

SEROLOGICAL, CELLULAR, AND BLOOD TRANSCRIPTOMIC RESPONSES TO A RECOMBINANT ONCHOCERCIASIS VACCINE IN CATTLE NATURALLY EXPOSED TO ONCHOCERCA OCHENGI

Ben Makepeace¹, John Graham-Brown¹, Tessa R. Walsh¹, Lisa Luu¹, Germanus Soh Bah², Ndoode Herman Okah-Nnane², David D. Ekale², Rene B. Ayiseh³, David Abraham⁴, Darrick Carter⁵, Sean Gray⁵, Godwin Nchinda⁶, Vincent Tanya², Sara Lustigman⁷
¹University of Liverpool, Liverpool, United Kingdom, ²L'Institut de Recherche Agricole Pour le Développement, Ngaoundéré, Cameroon, ³University of Buea, Buea, Cameroon, ⁴Thomas Jefferson University, Philadelphia, PA, United States, ⁵PAI Life Sciences, Seattle, WA, United States, ⁶Centre International de Référence "Chantal Biya", Yaounde, Cameroon, ⁷New York Blood Center, New York, NY, United States

9:15 a.m.

8390

A PHASE I/II STUDY OF THE SAFETY, IMMUNOGENICITY, AND EFFICACY OF SM-TSP-2/ALYDROGEL WITH OR WITHOUT AP 10-701 FOR INTESTINAL SCHISTOSOMIASIS IN HEALTHY UGANDAN ADULTS

Chrispus M. Bakunda¹, Betty Mwesigwa¹, Grace Mirembe¹, Jacqueline Namugabo¹, Hilda Mutebe¹, Immaculate Nakabuye¹, Douglas Makumbi¹, Amir Wamala¹, Allan Tindikahwa¹, Lara Hoeweler², Proscovia Naluyima¹, Laura Vasquez², Maria Elena Bottazzi³, Peter J. Hotez³, Elissa Malkin², Jeffrey M. Bethony², David J. Diemert², Hannah Kibuuka¹
¹Makerere University Walter Reed Project, Kampala, Uganda, ²George Washington University, Washington, DC, United States, ³Baylor College of Medicine, Houston, TX, United States

(ACMCIP Abstract)

9:30 a.m.

8391

THE ANTI-CIRCUMSPOROZOITE ANTIBODY RESPONSE OF CHILDREN TO SEASONAL VACCINATION WITH THE RTS,S/AS01_E MALARIA VACCINE OVER FIVE YEARS OF FOLLOW-UP (4 BOOSTER DOSES)

M Sanni Ali¹, Lisa Stockdale², Issaka Sagara³, Issaka Zongo⁴, Rakiswendé Serge Yerbanga⁴, Almahamoudou Mahamar³, Frédéric Nikiéma⁴, Amadou Tapily³, Frédéric Sompougou⁴, Modibo Diarra³, Duncan Bellamy², Katie Ewer², Charles Zoungrana⁴, Djibrilla Issiaka³, Alassane Haro⁴, Koualy Sanogo³, Abdoul-Aziz Sienou⁴, Mahamadou Kaya³, Seydou Traore³, Ismaila Thera³, Kalifa Diarra³, Amagana Dolo³, Paul Snell¹, Opokua Ofori-Anyinam⁵, Chris Ockenhouse⁶, Cynthia Lee⁶, Halidou Tinto⁴, Abdoulaye Djimde³, Jean-Bosco Ouedraogo⁴, Alassane Dicko³, Daniel Chandramohan¹, Brian Greenwood¹
¹London School of Hygiene & Tropical Medicine, London, United Kingdom, ²Jenner Institute, University of Oxford, Oxford, United Kingdom, ³The Malaria Research and Training Center, University of Sciences, Techniques and Technologies, Bamako, Mali, ⁴Institut de Recherche en Sciences de la Santé, Bobo-Dioulasso, Burkina Faso, ⁵GSK, Wavre, Belgium, ⁶PATH, Seattle, WA, United States

(ACMCIP Abstract)

Scientific Session 171

Mosquitoes-Biology, Physiology and Immunity

Convention Center - Room 356 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

#Immunology #HostResponse #ClimateChange

CHAIR

Alexandra Francian
University of New Mexico, Albuquerque, NM, United States

Maria Luisa Simoes
Institute of Tropical Medicine Antwerp, Antwerp, Belgium

8 a.m.

8392

EFFECTS OF HUMIDITY AND TEMPERATURE VARIATIONS ON ANOPHELES GENETIC TARGET CANDIDATES FOR MALARIA CONTROL

Thais Lemos-Silva, **Maria Luisa Simoes**
Institute of Tropical Medicine Antwerp, Antwerp, Belgium

8:15 a.m.

8393

TNF SIGNALING ACTIVATES CELLULAR IMMUNITY TO PROMOTE MALARIA PARASITE KILLING IN ANOPHELES GAMBIAE

George-Rafael Samantsidis, Hyeogsun Kwon, Megan Rogers, Catherine Fonder, **Ryan C. Smith**
Iowa State University, Ames, IA, United States

8:30 a.m.

8394

PIXEL INTENSITY OF WING PHOTOS USED TO PREDICT AGE OF ANOPHELES GAMBIAE SENSU LATO CAUGHT DURING THE RIMDAMAL II CLINICAL TRIAL

Greg Pugh¹, Emmanuel Sougué², Ryan Yoe¹, Lyndsey I. Gray¹, Blue Hephaestus³, Anthony Somé², A. Fabrice Somé², Bryce Assay³, Roch K. Dabiré², Sunil Parikh⁴, Brian D. Foy¹
¹Colorado State University, Fort Collins, CO, United States, ²Institute de Recherche en Sciences de la Santé, Bobo-Dioulasso, Burkina Faso, ³Viden Technologies, Laramie, WY, United States, ⁴Yale School of Public Health, New Haven, CT, United States

8:45 a.m.

8395

EXPLOITING MOSQUITO SALIVARY PROTEINS TO DEVELOP VECTOR-TARGETED VACCINES FOR MALARIA AND ARBOVIRUSES

Alexandra Francian¹, Aidan Leyba², Yewel Flores-Garcia³, Robert Taylor⁴, Justin Baca⁴, Pavan Muttli², Fidel Zavala³, Bryce Chackerian¹
¹University of New Mexico School of Medicine, Albuquerque, NM, United States, ²University of New Mexico College of Pharmacy, Albuquerque, NM, United States, ³Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States, ⁴University of New Mexico, Albuquerque, NM, United States

9 a.m.

8396

PLASMODIUM FALCIPARUM INFECTION IN THE HUMAN HOST AND THE VECTOR INFLUENCE NATURAL ANOPHELINE BITING BEHAVIOR AND PARASITE TRANSMISSION

Christine F. Markwalter¹, Zena Lapp¹, Lucy Abel², Emmah Kimachas², Evans Omollo³, Elizabeth Freedman¹, Tabitha Chepkwony², Mark Amunga², Tyler McCormick⁴, Sophie Bérubé⁵, Judith N. Mangeni⁶, Amy Wesolowski⁵, Andrew A. Obala⁶, Steve M. Taylor¹, Wendy P. O'Meara¹
¹Duke University, Durham, NC, United States, ²Academic Model Providing Access to Healthcare, Eldoret, Kenya, ³Duke Global Inc, Nairobi, Kenya, ⁴University of Washington, Seattle, WA, United States, ⁵Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States, ⁶Moi University, Eldoret, Kenya

Sunday
November 17

9:15 a.m.

8397

LANDSCAPES OF INFECTION: REDEFINING THERMAL SUITABILITY OF URBAN MALARIA TRANSMISSION BY THE INVASIVE MOSQUITO SPECIES *ANOPHELES STEPHENSI* IN THE CONTEXT OF RELATIVE HUMIDITY

Courtney Murdock¹, Joel Brown¹, Britny Johnson¹, Paul Huxley², Leah Johnson², Brandon Hollingsworth¹, Michael Wimberly³, Mercedes Pascual⁴, Rajendra Baharia⁵, Ajeet Mohanty⁶, Rajesh Sharma⁷, Sachin Sharma⁸, Keshav Vaishnav⁹, Vijay Kohli⁷, Vikas Desai¹⁰

¹Cornell University, Ithaca, NY, United States, ²Virginia Tech, Blacksburg, VA, United States, ³University of Oklahoma, Norma, OK, United States, ⁴New York University, New York City, NY, United States, ⁵National Institute for Malaria Research, Nadiad, India, ⁶National Institute for Malaria Research, Goa, India, ⁷Ahmedabad Municipal Corporation, Ahmedabad, India, ⁸Cornell In India & National Institutes for Malaria, Delhi, India, ⁹Surat Municipal Corporation, Surat, India, ¹⁰Urban Health And Climate Resilience Centre Of Excellence, Surat, India

9:30 a.m.

8398

ANGIOGENESIS AND CHRONIC EXPOSURE TO *ANOPHELES SALIVARY* PROTEINS

Olayinka M. Olajiga¹, Arley Calle¹, Samuel B. Jameson¹, Brendan H. Carter¹, Dawn M. Wesson¹, Eric Calvo², Jennifer Manuzak³, Berlin Londono-Renteria¹

¹Department of Tropical Medicine and Infectious Disease, School of Public Health and Tropical Medicine, New Orleans, LA, United States, ²Laboratory of Malaria and Vector Research National Institutes of Health (NIH/NIAID), Rockville, MD, United States, ³Department of Microbiology and Immunology, School of Medicine, Tulane University of New Orleans, New Orleans, LA, United States

Symposium 172

What's New? Exploring Emerging Technologies and Innovative Strategies in Advancing Skin NTD Integration

Convention Center - Room 357 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

Many of the 21 diseases on the WHO NTD list primarily result in skin disease – including leprosy, cutaneous leishmaniasis, scabies, yaws, Buruli ulcer, and mycetoma among others. These diseases result in considerable suffering, stigma and financial hardship for patients and affected communities. Worldwide, skin diseases are amongst the five largest contributors to disability adjusted life years lost to ill-health. However, despite this considerable burden of disease and the impact of these conditions, there has been relatively little attention paid to skin NTDs in contrast to those NTD primarily controlled via mass drug administration. In June 2022, the WHO released a pivotal guidance, titled “Ending the neglect to attain the Sustainable Development Goals: A strategic framework for integrated control and management of skin-related neglected tropical diseases”. Moving away from traditional disease-specific approaches, it is anticipated that synergies will be identified and integrated building on this shared feature, where possible, to achieve a greater health impact. The symposium is dedicated to exploring the transformative potential of emerging technologies and innovative strategies in addressing the challenges posed by skin NTDs. As individual diseases, they often face unique obstacles, such as a relatively limited patient population and the necessity for resource-intensive and time-consuming individualized treatments, hindering efficient disease control efforts. This has also led to a lack of interests for technological

advancements, and most yet do not have point-of-care diagnostic tools or have relied on the same medications for years. Following the WHO's strategic framework for skin NTD integration, we are currently at an important juncture. The integration approach not only fosters new possibilities and synergies across multiple skin NTDs, but also serves as a catalyst for the development of novel technologies and strategies. Given the daily advancements in technology, it is imperative to extend their application to the field of skin NTDs. Consequently, we can narrow the gaps in diagnostic capabilities, expand access to quality care, and empower frontline health workers with the tools and resources to effectively combat skin NTDs. This symposium will feature presentations by leading researchers, clinicians, and policymakers in the field, addressing advancements in molecular diagnostics, artificial intelligence (AI), teledermatology, and new models of care delivery. Concluding with a call to action, we urge further commitment to innovation for skin NTD control. At the end of the symposium, the audience will be equipped with knowledge on a state-of-the-art update highlighting the progress made in this emerging field. #Diagnostics #ClinicalResearch #FieldStudies #InfectiousDisease #Prevention

CHAIR

Rie Yotsu

Tulane School of Public Health and Tropical Medicine, New Orleans, LA, United States

Sundeep C. Vedithi

University of Cambridge, Cambridge, United Kingdom

8 a.m.

INTRODUCTION

8:10 a.m.

ADVANCING POINT OF CARE MOLECULAR DIAGNOSTICS FOR SKIN NEGLECTED TROPICAL DISEASES

Sundeep C. Vedithi

Department of Medicine, University of Cambridge, Cambridge, United Kingdom

8:30 a.m.

WHO ARTIFICIAL INTELLIGENCE (AI) PROJECT FOR SKIN NTDS IN KENYA

José Antonio Ruiz-Postigo

World Health Organization (WHO), Geneva, Switzerland

8:50 a.m.

BRIDGING GAPS: TELEDERMATOLOGY'S ROLE IN SKIN NTD DIAGNOSIS AND CAPACITY BUILDING IN CÔTE D'IVOIRE

Aubin Koffi Yao

Hope Commission International, Abidjan, Côte D'Ivoire

9:10 a.m.

NEW MODELS OF CARE DELIVERY FOR SKIN NTDS IN GHANA AND ETHIOPIA

Michael Marks

London School of Hygiene & Tropical Medicine, London, United Kingdom

9:30 a.m.

CALL FOR ACTION FOR ADAPTING EMERGING TECHNOLOGIES FOR SKIN NTD INTEGRATION

Rie Yotsu

Tulane School of Public Health and Tropical Medicine, New Orleans, LA, United States

Scientific Session 173

Viruses – Therapeutic, Antiviral Drugs, and Vaccine Clinical Trials

Convention Center - Room 383/384/385 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

This session does not carry CME credit.

#Therapeutics #Vaccinology

CHAIR

Chasity Elizabeth Trammell
Colorado State University, Fort Collins, CO, United States

Anna P. Durbin
Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States

8 a.m.

8399

USING ANCESTRAL SEQUENCE RECONSTRUCTION FOR GENERATION OF BROAD-SPECTRUM VACCINE PLATFORMS AGAINST TICK-BORNE FLAVIVIRUSES

Chasity E. Trammell, Brian J. Geiss, Gregory D. Ebel
Colorado State University, Fort Collins, CO, United States

8:15 a.m.

8400

PRECLINICAL DEVELOPMENT OF AN ORALLY AVAILABLE NS4B INHIBITOR FOR YELLOW FEVER

Fuxuan Wang¹, Sumangala Darsandhari¹, Biplav Shrestha¹, Abbie Wight², Justin Julander³, Brett Eaton⁴, Michael Murphy⁴, Yu Cong⁴, Manu Anantpadma⁴, Elena Postnikova⁴, Michael Holbrook⁴, Zhengyuan Jiang¹, Yusheng Wu¹, David Renner¹, Jason Clementon¹, Weihong Zhang¹, Yanming Du¹, Ju-Tao Guo¹, **Jinhong Chang¹**
¹Baruch S. Blumberg Institute, Doylestown, PA, United States, ²Utah State University, Logan, UT, United States, ³Utah State University, Logan, UT, United States, ⁴NIAID, Fort Detrick, MD, United States

8:30 a.m.

8401

B CELL RESPONSES TO A ZIKA PURIFIED INACTIVATED VACCINE ARE SHAPED BY PREVIOUS IMMUNIZATION WITH JAPANESE ENCEPHALITIS AND YELLOW FEVER VACCINES

Samantha Townsley¹, Vincent Dussupt¹, Lauren Smith¹, Rafael De La Barrera¹, Mélanie Merbah¹, Bonnie Slike¹, Yifan Li¹, Annika Schmid¹, Hélène Fradin Kirshner², Lauren Yum¹, Robert Clifford¹, Letzibeth Mendez-Rivera¹, Sam Lieberman¹, Gautam Kundu¹, Aviva Geretz¹, Rasmi Thomas¹, Kevin Wiehe², Morgane Rolland¹, Gregory Gromowski¹, Nelson Michael¹, Natalie Collins¹, **Shelly Krebs¹**
¹Walter Reed Army Institute of Research, Silver Spring, MD, United States, ²Human Vaccine Institute, Duke University, Durham, NC, United States

8:45 a.m.

8402

DEVELOPMENT OF A CONTROLLED ZIKA HUMAN INFECTION MODEL (CHIM)

Anna P. Durbin¹, Ruby Pritchett¹, Rachel Hampton¹, Beulah Sabundayo¹, Megan McKnight¹, Lina Moscarella¹, Xi Fang¹, Jeff Mayfield¹, Helen He¹, Cendy Alacantara¹, Will Stone², Elaine Lamirande², Stephen S. Whitehead²
¹Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States, ²National Institutes of Allergy & Infectious Diseases, National Institutes of Health, Bethesda, MD, United States

9 a.m.

8403

IMOJEV LIVE, ATTENUATED CHIMERIC VACCINE AGAINST JAPANESE ENCEPHALITIS: AN UPDATE AFTER 25 YEARS IN USE

Thomas P. Monath¹, Gray Heppner¹, Tracy Kemp¹, Bertrand Lepine², Marc Vouillamoz², Fabrice Baschiera²
¹Quigley BioPharma LLC, Leominster, MA, United States, ²Substipharm Biologics SA, Geneva, Switzerland

9:15 a.m.

8404

SIMULTANEOUS INHIBITION OF DENGUE VIRUS INFECTION AND NS1-MEDIATED ENDOTHELIAL HYPERPERMEABILITY WITH A NATURAL STEROIDAL SAPOGENIN

E. Vanessa Jimenez Posada, Pedro H. Carneiro, Robert Beatty, Scott B. Biering, Eva Harris
Division of Infectious Diseases and Vaccinology, School of Public Health, University of California, Berkeley, Berkeley, CA, United States

9:30 a.m.

8405

ANTIBODY RESPONSE PROFILE ELICITED BY A LIVE-ATTENUATED TETRAVALENT DENGUE VACCINE IN CHILDREN AND ADOLESCENTS FROM ENDEMIC AREAS

Eduardo J.M. Nascimento¹, Isadora Lape¹, Brooke Norwood¹, Jesuina Fernandes¹, Huahua Wan¹, Apurva Kulkarni¹, Vianney Tricou², Eloi Kpamegan¹, Mayuri Sharma¹
¹Takeda Vaccines Inc., Cambridge, MA, United States, ²Takeda Pharmaceuticals International AG, Zürich, Switzerland

Symposium 174

From Ideation to Implementation: How Malaria and Dengue Early Warning Systems Can Strengthen Health System Adaptation and Response to Climate Change

Convention Center - Room 388/389 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

According to new World Bank data, climate change can lead to “at least 21 million additional deaths by 2050 from just five health risks: extreme heat, stunting, diarrhea, malaria, and dengue.” Climate-informed early warning systems (EWS) have the potential to serve as powerful adaptation tools to the growing threats of climate change to health by increasing the effectiveness of disease control and strengthening the climate resilience of health systems. The combined intelligence of climate, health, mobility, and geospatial data with AI-enabled predictive capacity can be used beyond traditional EWS. In addition to enabling precise and cost-effective health service delivery, these systems can generate evidence to support financing opportunities and facilitate long-term planning. Improved climate-health data platforms and national ownership can enable policymakers to assess health impacts and make informed decisions to build climate resilience. Despite the promising impacts of EWS, challenges and gaps remain. Researchers, scientists, and implementers face key challenges with gaps in local and global climate-health data and fragmented funding. From a survey of 32 EWS experts in the Institute for Health Modeling and Climate Solutions’ (IMACS) community of practice, 40% stated obstacles with operationalization, including delayed data sharing, weak technical

Sunday
November 17

capacity, and scarce funding. To scale and sustain EWS, the operational challenges must be addressed in partnership with those at the frontlines of climate change, particularly the end users of an EWS. The most functional EWS is one with local data and system ownership that considers cost-effectiveness based on country capacity, and is informed by an overall strategic framework. Implementation of all the components of an EWS ecosystem can also provide lessons about data sharing and access, as well as preparation and response activities that are applicable to other diseases and overall public health emergency preparedness. This session brings together diverse stakeholders to review existing tools for malaria and dengue surveillance and health system readiness to adopt data tools for forecasting and response to climate-driven health threats. Attendees will learn and discuss how EWS development for malaria and dengue can better integrate local ownership, data sharing, and stakeholder engagement for the co-design and adoption of an EWS tool for adaptation. #ClimateChange #InfectiousDisease #Modeling

CHAIR

Grace Kim

Malaria No More, Washington, DC, United States

James Colborn

Clinton Health Access Initiative, Maputo, Mozambique

8 a.m.

INTRODUCTION

8:10 a.m.

PERSPECTIVES FROM THE ASIA-PACIFIC REGION: STRENGTHENING TECHNICAL CAPACITY TO USE AND MANAGE SURVEILLANCE AND GEOSPATIAL DATA FOR NATIONAL MALARIA AND DENGUE PROGRAMS

Richard J. Maude

Mahidol Oxford Tropical Medicine Research Unit, Bangkok, Thailand

8:20 a.m.

BRIDGING SCIENTIFIC AND TRADITIONAL KNOWLEDGE: INCLUSION OF COMMUNITY PERSPECTIVES IN UNDERSTANDING CLIMATE-HEALTH LINKAGES

Samson Nzou

Kenya Medical Research Institute (KEMRI), Nairobi, Kenya

8:30 a.m.

PERSPECTIVES FROM THE UGANDA MINISTRY OF HEALTH: UTILITY OF EWS AND HOW TO ADDRESS BARRIERS

Jimmy Opigo

Ministry of Health, Uganda, Kampala, Uganda

8:40 a.m.

ASSESSING THE UTILITY OF EARLY WARNING SYSTEMS THROUGH END-USER ENGAGEMENT

Kate Zinszer

University of Montreal, Montreal, Canada

8:50 a.m.

EXPANDING VECTOR-BORNE EARLY WARNING SYSTEM CAPACITY IN LATIN AMERICA

William Pan

Duke University, Durham, NC, United States

9 a.m.

ENHANCING EARLY WARNING SYSTEMS FOR CLIMATE-SENSITIVE DISEASES: LEVERAGING MULTISECTORAL COLLABORATION, MULTIDIMENSIONAL DATA, AND DIGITAL INNOVATIONS

Kaushik Sarkar

Malaria No More, Washington, DC, United States

Symposium 175

Socio-Cultural Practices, Low Risk Perception and Policy Implementation Gaps Heighten Zoonotic Disease Risk in Indigenous Communities: Observations from Northeast India

Convention Center - Room 391/392 (3rd Floor)

Sunday, November 17, 8 a.m. - 9:45 a.m.

Zoonotic diseases represent an increasing and significant threat to global health, with about one billion cases of illness and millions of deaths occurring annually. An estimated 60% of emerging infectious diseases are caused by zoonotic pathogens. Zoonotic diseases disproportionately affect vulnerable groups in low- and middle-income countries. India has emerged as a hotspot for zoonotic diseases. Northeast India is home to about 40 million people from 200 indigenous tribal communities residing in rural, remote and difficult terrain. These communities have unique cultural practices and a predominantly non-vegetarian diet, including consumption of bushmeat. Over half of the region is covered in forests, with designated 'biodiversity hotspots' linked to higher zoonotic disease risks. Mixed farming, a key economic activity, often involves rearing livestock in close proximity to human dwellings. Compounded by limited veterinary services and irregular animal health assessments, these practices pose inherent risks to animal health and the potential for transmission of zoonotic pathogens to humans. During this symposium, we will present and discuss case-studies that illustrate the heightened risks of zoonoses due to low-risk perception among community members, livestock handlers and traditional healers who treat sick animals. Insights from these case-studies were developed from our ongoing research on Zoonotic Vector Borne Diseases (ZVBD) in Meghalaya, a state in northeast India. We will focus on how socio-cultural beliefs and practices affect intentional and accidental contacts with domestic and wild animals, thereby impacting zoonotic disease risk. Our study has revealed that a significant portion of the population are being exposed to zoonotic pathogens, underscoring the urgent need for collective engagement with these vulnerable, indigenous communities to mitigate their disease risk. The relevance of our findings to other remote, less modern and culturally divergent populations will be explored. Symposium content tags: #EcologicalStudies; #EmergingDiseaseThreats; #FieldStudies; #Prevention; #SocialScience

CHAIR

Rajiv Sarkar

Indian Institute of Public Health Shillong, Shillong, India

Melari S. Nongrum

Indian Institute of Public Health Shillong, Shillong, India

8 a.m.
INTRODUCTION

8:10 a.m.
LOW-RISK PERCEPTION OF ZOOSES AND HEALTH SEEKING BEHAVIOR FOR ANIMALS IN MEGHALAYA, INDIA

Melari S. Nongrum
Indian Institute of Public Health Shillong, Shillong, India

8:30 a.m.
ANIMAL REARING AND HANDLING PRACTICES AMONG RURAL HOUSEHOLDS IN MEGHALAYA, INDIA

Uniqueky G. Mawrie
London School of Hygiene & Tropical Medicine, London, United Kingdom

8:45 a.m.
MEAT HANDLING PRACTICES AND FOOD-SAFETY KNOWLEDGE AMONG BUTCHERS AND ABATTOIR WORKERS IN MEGHALAYA, INDIA

Rajiv Sarkar
Indian Institute of Public Health Shillong, Shillong, India

9 a.m.
THE IMPACT OF SOCIO-BEHAVIORAL PRACTICES ON INCREASED RISK OF ZOO NOTIC DISEASES IN NORTHEAST INDIA

Benedicta S. Kumar
Indian Institute of Public Health Shillong, Shillong, India

9:15 a.m.
PANEL DISCUSSION

Mark Wilson
University of Michigan, School of Public Health, Ann Arbor, MI, United States



Symposium 176

Bartonellosis: A (Re)emerging Ectoparasite-borne Disease of People and Animals

Convention Center - Room 393/394 (3rd Floor)
Sunday, November 17, 8 a.m. - 9:45 a.m.

This session does not carry CME credit.

This session focuses on Bartonellosis, ectoparasite-borne systemic infections caused by various Bartonella species. Bartonellosis has worldwide distribution and diverse disease manifestations. Globally, epidemiologic risks for Bartonellosis depend on the local ecology of vectors and reservoir species associated with each Bartonella species. The session aims to explore the disease's re-emergence due to enhanced human-animal-vector interactions, evolving vector populations, and advancements in diagnostic methods. The symposium will delve into the epidemiology of Bartonella infections, particularly in low- and middle-income countries, and explore the risk factors for human infection and challenges to effective treatment. It will also examine the transmission dynamics of Bartonella species, including the potential role of ticks as vectors and roles for wildlife reservoirs. This session seeks to foster a comprehensive understanding of Bartonellosis, promoting global health and inter-professional collaboration. #EcologicalStudies

#EmergingDiseaseThreats #Epidemiology #InfectiousDisease #TranslationalScience

CHAIR

Erin Lashnits
University of Wisconsin, Madison, WI, United States

Monica Embers
Tulane National Primate Research Center, Tulane University Health Sciences, Covington, LA, United States

8 a.m.
INTRODUCTION

8:10 a.m.
THE NEGLECTED BURDEN OF BARTONELLA QUINTANA INFECTION IN LOW- AND MIDDLE-INCOME COUNTRIES

Carl Boodman
University of Antwerp/ Institute of Tropical Medicine (Belgium); University of Manitoba, Winnipeg, Canada

8:25 a.m.
VETERINARY PROFESSIONALS AND RISK FOR BARTONELLA EXPOSURE

Edward B. Breitschwerdt
North Carolina State University College of Veterinary Medicine, Raleigh, NC, United States

8:40 a.m.
B. BACILLIFORMIS IN PERU

Victor Alberto Jimenez- Vasquez
Instituto Nacional De Salud

8:55 a.m.
BARTONELLA IN WILD ANIMALS FROM BRAZIL: SCRATCHING THE TIP OF THE ICEBERG

Marcos Rogério André
Universidade Estadual Paulista Júlio de Mesquita Filho, UNESP, Jaboticabal, Brazil

9:10 a.m.
BARTONELLA ANIMAL MODELS: ANTIBIOTIC EFFICACY, CO-INFECTION, AND TICKS

Monica Embers
Tulane National Primate Research Center Tulane University Health Sciences, Covington, LA, United States

Scientific Session 177

Schistosomiasis II: Epidemiology and Control

Convention Center - Room 395/396 (3rd Floor)
Sunday, November 17, 8 a.m. - 9:45 a.m.

#ChildHealth #Diagnostics #EcologicalStudies

CHAIR

Charles B. Delahunty
Global Health Labs, Bellevue, WA, United States

Karen Mozo Velazco
Instituto de Medicina Tropical Alexander von Humboldt – Universidad Peruana Cayetano Heredia, Cusco, Peru

8 a.m.

8406

COMBINING WEARABLE GPS LOGGERS WITH ENVIRONMENTAL AND SNAIL DATA TO UNCOVER FINE-SCALE *SCHISTOSOMA MANSONI* TRANSMISSION DYNAMICS IN UGANDAFabian Reitzug¹, Narcis B. Kabatereine², Betty Nabette², Goylette F. Chamii¹¹University of Oxford, Oxford, United Kingdom, ²Vector Control Division, Ministry of Health, Kampala, Uganda

8:15 a.m.

8407

ACCEPTABILITY AND FEASIBILITY OF A ONE-STOP HOME-BASED GENITAL SELF-SAMPLING FOR FEMALE GENITAL SCHISTOSOMIASIS, HUMAN PAPILLOMA VIRUS AND TRICHOMONAS AND HIV SELF-TESTING: BASELINE DATA FROM A LONGITUDINAL COHORT STUDY IN ZAMBIARhoda Ndubani¹, Kwame Shanaube¹, Olimpia Lamberti², Nkatya Kasese¹, Jennifer Fitzpatrick¹, Maina Cheeba¹, Beatrice Nyondo¹, Bronwyn Neufeld², Barry Kosloff², Bonnie Webster⁴, Helen Kelly², Amaya L. Bustinduy²¹Zambart, Lusaka, Zambia, ²London School of Hygiene & Tropical Medicine, London, United Kingdom, ³Longhorn Vaccines and Diagnostics LLC, Bethesda, MD, United States, ⁴Natural History Museum, London, United Kingdom

8:30 a.m.

8408

***SCHISTOSOMA MANSONI* AND *HELICOBACTER PYLORI* CO-INFECTIONS AMONG SCHOOL-AGED POPULATIONS: A STUDY IN NIGERIAN COMMUNITIES**Babatunde Adewale¹, Muinah Fowora¹, Hammed Mogaji¹, Omolola Omolopo¹, Nurudeen Rahman¹, Ayorinde James¹, Herbert De'Broski²¹Nigerian Institute of Medical Research, Lagos, Nigeria, ²University of Pennsylvania, Philadelphia, PA, United States

8:45 a.m.

8409

MAIN RESULTS FROM A PHASE II RANDOMISED PLACEBO-CONTROLLED TRIAL OF PRAZIQUANTEL IN PRESCHOOL CHILDREN WITH INTESTINAL SCHISTOSOMIASISAmaya Lopez Bustinduy¹, Andrew Edielu², Gloria Kakoba², Jacent Nassuuna², Racheal Nakyesige², Victor Anguajibi², Simon Mpooya², Moses Adriko³, Govert Van Damm⁴, Paul Corstjens⁴, Sophie Pach¹, Hannah Wu⁵, Susannah Colt⁵, Patrice Mawa², Edridah Muheki², Narcis Kabatereine³, Emily Webb¹, Jennifer F. Friedman⁵¹London School of Hygiene & Tropical Medicine, London, United Kingdom, ²MRC-LSHTM Research Unit, Entebbe, Uganda, ³Vector Control Division, Kampala, Uganda, ⁴Leiden University Medical Centre, Leiden, Netherlands, ⁵Rhode Island Hospital, Providence, RI, United States

9 a.m.

8410

THE USE OF WATER EDNA IN SNAIL IDENTIFICATION FOR THE UNDERSTANDING OF *FASCIOLOA* ENVIRONMENTAL BURDEN AND SNAIL DIVERSITY IN THE HIGHLANDS OF PERUMelinda B. Tanabe¹, Martha Vanessa Fernandez-Baca², Rodrigo A. Ore Velarde², Alejandro Castellanos-Gonzalez¹, Miguel M. Cabada¹, Fasciola Peru TMRC Research Group³¹UTMB, Galveston, TX, United States, ²UPCH – UTMB Collaborative Research Center - Cusco, Universidad Peruana Cayetano, Cusco, Peru

9:15 a.m.

8411

INFLAMMATION-ADJUSTED VITAMIN A DEFICIENCY IS ASSOCIATED WITH HEAVY *SCHISTOSOMA MANSONI* INFECTION INTENSITY AMONG PRESCHOOL-AGED CHILDREN IN UGANDASusannah Colt¹, Hannah W. Wu¹, Andrew Edielu², Emily L. Webb³, Racheal Nakyesige⁴, Patrice A. Mawa², Christopher P. Duggan⁵, Jonathan D. Kurtis⁶, Amaya L. Bustinduy³, Jennifer F. Friedman¹¹Rhode Island Hospital, Providence, RI, United States, ²London School of Hygiene & Tropical Medicine Uganda Research Unit, Entebbe, Uganda, ³London School of Hygiene & Tropical Medicine, London, United Kingdom, ⁴Uganda Virus Research Institute, Entebbe, Uganda, ⁵Boston Children's Hospital, Boston, MA, United States, ⁶Warren Alpert Medical School of Brown University, Providence, RI, United States

9:30 a.m.

8412

ASSOCIATION OF *S. HAEMATOBIIUM* INFECTION WITH PREGNANCY IN TANZANIASheridan Bowers¹, Jane K. Maganga², Loyce Mhango², Peter Shigella², Crispin Mukerebe², Humphrey D. Mazigo³, Govert J. van Dam⁴, Danielle de Jong⁵, Paul L. Corstjens⁵, Saidi H. Kapiga², W. Evan Secor⁶, Myung Hee Lee⁷, Jennifer A. Downs⁷, John M. Changalucha²¹Weill Cornell Medicine, New York, NY, United States, ²Mwanza Intervention Trials Unit/ National Institute for Medical Research, Mwanza, United Republic of Tanzania, ³Department of Parasitology, Catholic University of Health and Allied Sciences, Mwanza, United Republic of Tanzania, ⁴Department of Parasitology, Leiden University Medical Center, Leiden, Netherlands, ⁵Department of Cell and Chemical Biology, Leiden University Medical Center, Leiden, Netherlands, ⁶Division of Parasitic Diseases for Malaria, Center for Global Health, Centers for Disease Control and Prevention, Atlanta, GA, United States, ⁷Center for Global Health, Weill Cornell Medicine, New York, NY, United States**Coffee Break**Convention Center - Third Floor Lobby Areas
Sunday, November 17, 9:45 a.m. - 10:15 a.m.**Plenary Session 178****Plenary Session V**Convention Center - Hall I-2 (1st Floor)
Sunday, November 17, 10:15 a.m. - 11:15 a.m.**THIS SESSION DOES NOT CARRY CME CREDIT.****Paulin Basinga, MD, PhD**
Director, Africa
Bill & Melinda Gates Foundation
Nairobi, Kenya

Dr. Paulin Basinga leads the Gates Foundation's work across the African continent.

Assuming this role in March 2024, Paulin brings extensive experience gained through his country and global level engagements. Before this role, he held the position of foundation's Global Director of Program Advocacy and Communications (PAC) based in Seattle. In this role, Paulin orchestrated strategic advocacy initiatives and forged valuable partnerships with advocacy teams in S/E Asia and Africa.



Symposium 179

Malaria without Borders: Understanding the Influence of Cross-Border Migration on Malaria Elimination Strategies

Convention Center - Hall I-2 (1st Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Geopolitical instability, climate change, and other factors make human migration a dynamic and increasingly important factor in disease elimination strategies. Since 2018, 4 countries in the Americas - El Salvador, Belize, Argentina, and Paraguay - have eliminated malaria. Honduras, Panama, and Ecuador (among others) were believed to be on the same elimination trajectory. Between 2014 and 2019, all dipped below 600 annual, locally transmitted cases. Unfortunately, in 2022, these same 3 countries reported 11,984 cases combined, the vast majority of which occurred in the border regions of Honduras-Nicaragua, Panama-Colombia, & Ecuador-Peru. Accelerating and achieving elimination at foci, national, or regional scales will require – among other things - understanding the mechanisms in which cross-border migration is sustaining transmission and finding creative solutions for malaria programs to respond. While data on cross-border movements and international migrant populations can be limited and difficult to analyze, when available, they provide valuable insight into malaria transmission dynamics, the success (or lack there-of) of various interventions, and strategies that could be deployed to reach the most affected populations and reduce transmission. The influence of migration patterns on transmission is highly setting-specific, as it depends both on the political and demographic aspects of people's movement and on the epidemiology of malaria and magnitude of interventions. Leveraging such data with appropriate analytical approaches is key to understanding how case importation and malaria transmission interact specifically in each geography. This symposium will explore four unique approaches to understanding cross-border migration and its impact on malaria elimination. First, we will explore the genetic connectivity of parasites from South to Central America leveraging targeted and whole genome sequencing data from 5+ countries where cross-border migration is jeopardizing elimination. Second, we will dive into a novel *P. vivax* mathematical model that highlights how the success of various intervention strategies and elimination probabilities in Honduras are altered based on reductions in imported cases from Nicaragua. Third, we will present the most recent evidence from research being conducted in communities serving as migration transit points for the over 1 million migrants that have passed through the Darien Gap, Panama the past three years. Lastly, we will showcase how a network analysis along the Ecuador-Peru border is informing bi-national malaria response. Emphasis from each presentation will be given to discussing the implications of the findings for MoH strategies and elimination trajectories across the Americas. #Elimination #Modeling #Genetics #PopulationSurveillance #InfectiousDisease

CHAIR

Sarah Park
Clinton Health Access Initiative, Boston, MA, United States

Before his PAC role, Paulin served as the foundation's Director of Health for Africa, leading a dedicated team in devising and implementing comprehensive country plans to advance health priorities across the continent. As the foundation's country director in Nigeria, Paulin revitalized critical partnerships and advanced health, nutrition, agriculture, gender, and financial inclusion priorities, aligning them with the Nigerian government's Human Capital Development Agenda.

Paulin's journey with the foundation began in 2012 as a Senior Program Officer on the HIV team. Here, he worked on the HIV Efficiency and Effectiveness Initiative, pioneering strategies to maximize the impact of the foundation's investments in combating HIV/AIDS. Subsequently, he transitioned to the Integrated Delivery team, overseeing investments in community and child health before assuming the role of Deputy Director of the Country Primary Health Initiative team - covering Ethiopia, Nigeria, Senegal, Burkina Faso and other countries.

In 2015, Paulin was seconded to Rwanda to assume a pivotal role as Senior Advisor to the Rwanda Biomedical Centre within the Rwandan Ministry of Health. During this tenure, he provided crucial decision support for health sector reforms in Rwanda.

Before joining the foundation, Paulin served as Deputy Director of Research at the National University of Rwanda's School of Public Health. Here, he led groundbreaking research initiatives addressing HIV/AIDS, tuberculosis, and maternal and child health, contributing significantly to efforts to strengthen Rwanda's healthcare systems.

Paulin holds a medical degree from the National University of Rwanda, complemented by a Master's degree and Ph.D. in International Development from Tulane University.

CHAIR

Linnie Golightly
Weill Cornell Medical College, New York, NY, United States

10:15 a.m. INTRODUCTION

Linnie Golightly
Weill Cornell Medical College, New York, NY, United States

10:30 a.m. PLENARY ADDRESS: FROM LOCAL INNOVATION TO GLOBAL COLLABORATION: BREAKING DOWN SILOS TO TACKLE INFECTIOUS DISEASES IN AFRICA

Paulin Basinga
Bill & Melinda Gates Foundation, Nairobi, Kenya

Break

Sunday, November 17, 11:15 a.m. - 11:30 a.m.

Sunday
November 17

11:30 a.m.
INTRODUCTION**11:35 a.m.**
THE ROLE OF CROSS-BORDER CASE IMPORTATION ON P. VIVAX MALARIA ELIMINATION IN HONDURAS: COMPARING INTERVENTION STRATEGIES WITH MATHEMATICAL MODELLING

Emilie Pothin
Swiss Tropical and Public Health Institute, Allschwil, Switzerland

11:55 a.m.
RISK FACTORS OF MALARIA INFECTION IN COMMUNITIES ALONG MIGRATION ROUTES IN PANAMA

Mark Janko
Duke University, Durham, NC, United States

12:15 p.m.
LEVERAGING SOCIAL NETWORK ANALYSIS TO IMPROVING MALARIA RESPONSE ALONG THE ECUADOR-PERU BORDER

Carolina Coombs
Universidad Peruana Cayetano Heredia (UPCH), Lima, Peru

12:35 p.m.
TRENDS IN GENETIC CONNECTIVITY BETWEEN CENTRAL AND SOUTH AMERICAN PLASMODIUM POPULATIONS

Paulo Manrique
Harvard T.H. Chan School of Public Health, Boston, MA, United States

Scientific Session 180**Clinical Tropical Medicine: Disease and Vaccine Safety Surveillance**

Convention Center - Room 343/344 (3rd Floor)
Sunday, November 17, 11:30 a.m. - 1:15 p.m.

#InfectiousDisease #Epidemiology #Diagnostics
#PopulationSurveillance

CHAIR

Eileen Farnon
Brighton Collaboration, Decatur, GA, United States

Polly Yap
Monash University Malaysia, Subang Jaya, Malaysia

11:30 a.m.
8413**PREPARING FOR VACCINE ADVERSE EVENTS OF SPECIAL INTEREST-X (AESI-X): A STANDARDIZED APPROACH APPLIED TO NOVEL VACCINES**

Eileen C. Farnon¹, Barbara J. Law¹, Dale Nordenberg¹, Andy Stergachis¹, Rebecca E. Chandler², Robert T. Chen¹, Brighton Collaboration SPEAC AESI-X Working Group¹
¹Safety Platform for Emergency vACCines (SPEAC) Project, Brighton Collaboration, Decatur, GA, United States, ²Coalition for Epidemic Preparedness Innovations, Oslo, Norway

11:45 a.m.
8414**ENHANCING ACCESS TO HIGHLY MULTIPLEXED DIAGNOSTICS IN LMICS: LEVERAGING OXFORD NANOPORE SEQUENCING FOR DETECTION OF RESPIRATORY VIRUSES AND EMERGING PATHOGENS**

Polly S. X. Yap¹, Ryan C. Chapman², Nathaniel Christy³, Sharifah Syed Hassan¹, Chin Fang Ngim¹, Lian Yih Pong¹, Jessica D. Wiley⁴, Robert D. Hontz³, Anca Selariu³, Amreeta Dhanoa¹, Wan Fadhilah Wan Ismail⁵, Andrew G. Letizia³, Michael R. Wiley²
¹Monash University Malaysia, Subang Jaya, Malaysia, ²University of Nebraska Medical Center, Omaha, NE, United States, ³United States Naval Medical Research Unit- INDO PACIFIC, Singapore, Singapore, ⁴PraesensBio LLC, Omaha, NE, United States, ⁵Mahmoodiah Health Clinic, Johor Bahru, Malaysia

Noon
8415**INTEGRATED SEROLOGICAL SURVEILLANCE FOR INFECTIOUS DISEASES IN ZAMBEZIA PROVINCE, MOZAMBIQUE USING MULTIPLEX BEAD ASSAYS**

Andrea C. Carcelen¹, Celso Monjane², Sophie Bérubé¹, Saki Takahashi¹, Thebora Sultane², Imelda Chelene², Gretchen Cooley³, Brook Goodhew³, Catriona Patterson⁴, Kevin Tetteh⁴, Manuel Mutambe², Melissa Higdon¹, George Mwinnyaa¹, Gilberto Nhapure⁵, Pedro Duce⁵, Diana L. Martin³, Christopher Drakeley⁴, William J. Moss¹, Ivalda Macicame²
¹Johns Hopkins University, Baltimore, MD, United States, ²Instituto Nacional de Saude, Maputo, Mozambique, ³Centers for Disease Control and Prevention, Atlanta, GA, United States, ⁴London School of Hygiene and Tropical Medicine, London, United Kingdom, ⁵Instituto Nacional de Estadística, Maputo, Mozambique

12:15 p.m.
8416**A SEPSIS SYNOPSIS: HETEROGENEITY OF SEPSIS PRESENTATIONS ACROSS THE ACESO NETWORK**

Alyse Wheelock¹, Melissa Bradshaw¹, George Oduro², Hannah Kibuuka³, Stephen Okello³, Andrew Kambugu⁴, Te Vantha⁵, Nehkonti Adams⁶, Subramaniam Krishnan¹, Patrick Blair¹, Joshua Chenoweth¹, Deborah Striegel¹, Qianru Wu¹, Carlyle Gollogly⁷, Kristen Petrone¹, Danielle Clark¹
¹ACESO, the Henry Jackson Foundation, Bethesda, MD, United States, ²Komfo Anokye Teaching Hospital, Kumasi, Ghana, ³Makerere University Walter Reed Project, Kampala, Uganda, ⁴Infectious Diseases Institute, Kampala, Uganda, ⁵Takeo Provincial Hospital, Takeo, Cambodia, ⁶Naval Medical Research Command, Falls Church, VA, United States, ⁷ACESO, Henry Jackson Foundation, Bethesda, MD, United States

12:30 p.m.
8417**SEPSIS-RELATED DEATHS AMONG CHILDREN BELOW 5 YEARS OF AGE ENROLLED IN THE CHILD HEALTH AND MORTALITY PREVENTION SURVEILLANCE (CHAMPS) NETWORK PROGRAM IN SUB-SAHARAN AFRICA AND SOUTH ASIA BETWEEN 2017 - 2022**

Harun Owuor¹, Kyu Hun Lee², Emily S. Gurley³, Shams El Arifeen⁴, Portia C. Mutevedzi², Cynthia G. Whitney², Jeannette Wadula⁵, Mohammad Zahid Hossain⁴, Inacio Mandomando⁶, Nega Assefa⁷, Soter Ameh⁸, Ikechukwu Ogbuanu⁸, Dickens Dickens Onyango⁹, Joyce Akinyi Were¹, Kotloff Karen¹⁰, Shabir madhi⁵, Rosaura Varo⁶, Ziyaad Dangor⁵, Victor Akelo¹¹
¹KEMRI CGHR, Kisumu, Kenya, ²Emory University, Atlanta, GA, United States, ³John Hopkins University, Baltimore, MD, United States, ⁴International Centre for Diarrheal Disease Research, Bangladesh (icddr), Dhaka, Bangladesh, ⁵University of the Witwatersrand, Johannesburg, South Africa, ⁶Centro de Investigacao en Saude de Manhica (CISM), Maputo, Mozambique, ⁷College of Health and Medical Sciences, Haramaya University, Harar, Ethiopia, ⁸Crown Agents, Freetown, Sierra Leone, ⁹Kenya Ministry of Health, Kisumu, Kenya, ¹⁰University of Maryland School of Medicine, Baltimore, MD, United States, ¹¹Liverpool School of Tropical Medicine, London, United Kingdom

12:45 p.m.

8418

RICKETTSIOSIS AND SCRUB TYPHUS AMONG HOSPITALIZED PATIENTS WITH ACUTE FEBRILE ILLNESS IN RURAL NORTHEASTERN AND NORTH BORDER PROVINCES OF THAILAND, 2017-2020

Adelaide F. Green¹, Saithip Bhengsr², Betsy Cadwell¹, Emily Bloss², Nuttagarn Chuenchom³, Supphachoke Khemla⁴, Sumonmal Uttayamakul⁵, Pongpun Sawatwong², Ornuma Sangwichian², Duangkamon Siludjai², Phanthaneeya Teepruksa², Beth Skaggs², Cecilia Y. Kato¹, Christopher J. Gregory², John R. Macarthur², James D. Heffelfinger², Carol Y. Rao¹

¹Centers for Disease Control and Prevention, Atlanta, GA, United States, ²Centers for Disease Control and Prevention - Thailand, Nonthaburi, Thailand, ³Mae Sot Hospital, Nakhon Phanom, Thailand, ⁴Nakhon Phanom hospital, Nakhon Phanom, Thailand, ⁵Bamrasnaradura Infectious Diseases Institute, Department of Disease Control, Ministry of Public Health and National Institute of Health, Department of Medical Sciences, Ministry of Public Health, Nonthaburi, Thailand

1 p.m.

8419

MICROSTRATIFICATION OF VISCERAL LEISHMANIASIS ENDEMIC AREAS TO IDENTIFY HOTSPOTS AND DISEASE SHIFTING PATTERNS IN NEPAL

Krishna Pant¹, Megha R. Banjara¹, Dharmanand Joshi¹, Govinda R. Paneru¹, Abraham Aseffa², Rudra Marasini³, Gokarna Dahal³, Anand B. Joshi¹

¹Public Health and Infectious Disease Research Center, Kathmandu, Nepal, ²Special Program for Research and Training in Tropical Diseases, Geneva, Switzerland, ³Epidemiology and Disease Control Division, Ministry of Health, Kathmandu, Nepal

Symposium 181

The Role of Molecular Xenomonitoring in Transmission Risk Assessment and Post-Validation Surveillance After Mass Drug Administration for Lymphatic Filariasis

Convention Center - Room 345 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

To eliminate lymphatic filariasis (LF) as a public health problem, the mass drug administration (MDA) strategy aims to reduce infection levels within the population to a point where transmission is no longer sustainable. However, confirming that treatment areas meet the established thresholds to suggest LF elimination in countries with less-developed health infrastructure can take time and effort. Community members may be unwilling or unable to participate in post-treatment surveillance for personal or logistical reasons. Molecular xenomonitoring (MX) may help overcome some of these limitations. This symposium will explore whether MX can monitor transmission interruption following MDA. Key areas of focus include 1) the correlation between data derived by MX and human monitoring studies after MDA for LF; 2) the required threshold of mosquito infection prevalence and the necessary number of mosquitoes to be sampled to determine cessation of transmission reliably; 3) the dependability of using surrogate mosquito species (e.g., *Culex*) in areas with *Anopheles* as the major LF vectors; and 4) the role and cost-effectiveness of MX in post-validation surveillance. These and other questions will be examined in three lectures, followed by a panel discussion and Q&A session. The symposium's organizers, speakers, and panelists

have experience in LF elimination in India, Africa, and Papua New Guinea. #Elimination, #FieldStudies, #PopulationSurveillance, #Infectious Diseases.

CHAIR

Christopher L. King
Case Western Reserve University, Cleveland, OH, United States

Stephan Karl
James Cook University, Cairns, Australia

11:30 a.m.

INTRODUCTION

11:40 a.m.

WHAT IS THE SENSITIVITY OF MX, AND IS IT BETTER THAN HUMAN SURVEYS AT DETECTING POST-MDA RESERVOIRS OF INFECTION IN LF?

Lisa Reimer
Centers for Disease Control, Atlanta, GA, United States

Noon

WHAT IS THE STATUS OF MX IN LF ENDEMIC AREAS WHERE CULEX IS THE PRINCIPAL VECTOR?

K. Krishnamoorthy
Vector Control Research Centre (retired). Formerly Senior Scientist, Pondicherry, India

12:20 p.m.

POST-VALIDATION LF MONITORING WITH MX WORKFLOWS FOR INTEGRATION WITH OTHER MOSQUITO-BORNE PATHOGENS AND THE DESIGN OF "TRAPPING NETWORKS" TO MAXIMIZE DETECTION OF LF SIGNALS

Brian Johnson
Queensland Institute for Medical Research (QIMR) Burghofer, Brisbane, Australia

12:40 p.m.

POST-VALIDATION LF MONITORING WITH MX WORKFLOWS FOR INTEGRATION WITH OTHER MOSQUITO-BORNE PATHOGENS AND THE DESIGN OF "TRAPPING NETWORKS" TO MAXIMIZE DETECTION OF LF SIGNALS

Gregor Devine
QIMR Burghofer, Brisbane, Australia

Scientific Session 182

Kinetoplastida and Other Opportunistic and Anaerobic Protozoa: Immunology, Invasion, Cellular and Molecular Biology

Convention Center - Room 352 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

#InfectiousDisease #Immunology #HostResponse
#CellBiology

CHAIR

Hira Nakhasi
FDA, Silver Spring, MD, United States

Abhay Satoskar
Ohio State University, Columbus, OH, United States

11:30 a.m.

8420

BANGLADESHI CHILDREN HAVE IMMUNITY TO CRYPTOSPORIDIA-ASSOCIATED DIARRHEA BUT NOT TO INFECTION

Carol A. Gilchrist¹, William A. O. Petri¹, Biplob Hossain², Md. Mamun Kabir², Hannah H. So¹, G. Brett Moreau¹, Uma Nayak¹, Jennie Z. Ma¹, Zannatun Noor², Abu S. G. Faruque², Md. Masud Alam², Rashidul Haque², William A. Petri¹

¹University of Virginia, Charlottesville, VA, United States, ²icddr, Dhaka, Bangladesh

(ACMCIP Abstract)

11:45 a.m.

8421

MECHANISM OF INTESTINAL BARRIER REPAIR IN GIARDIASIS

Rita Teye Kosile, Vanessa Angelova, Evan Pannkuk, Steven Singer
Georgetown University, Washington, DC, United States

(ACMCIP Abstract)

Noon

8422

UNWINDING THE IRONY OF SEVERE ANEMIA IN ANTIMONY-RESISTANT *LEISHMANIA DONOVANI* INFECTION AT THE NEXUS OF OXIDATIVE OUTBURST AND IRON PURSUIT

Souradeepa Ghosh, Krishna Vamshi Chigicherla, Budhaditya Mukherjee
Indian Institute of Technology, Kharagpur, Kharagpur, India

(ACMCIP Abstract)

12:15 p.m.

8423

UNIQUE IMMUNE AND TISSUE REPAIR MARKERS IN CONGENITAL CHAGAS

Sneider Alexander Gutierrez Guarnizo¹, Carolina Duque¹, Jill Hakim¹, Martín Obregón², Jessi Condori², Paloma Samame³, Emily Arteaga³, Clariza Roxana³, Jean Karla Velarde³, Edith Malaga², Andrea Diestra², Alejandra Pando², Manuela Verastegui², Monica Pajuelo², Maritza Calderon², Freddy Tinajeros³, Natalie Bowman⁴, Robert Gilman¹, Monica Mugnier¹

¹Johns Hopkins University, Baltimore, MD, United States, ²Universidad Peruana Cayetano Heredia, Lima, Peru, ³Hospital de La Mujer Dr. Percy Boland Rodriguez, Santa Cruz, Plurinational State of Bolivia, ⁴University of North Carolina at Chapel Hill School of Medicine, Chapel Hill, NC, United States

(ACMCIP Abstract)

12:30 p.m.

8424

CO-DELIVERY OF IL-12 AND *LEISHMANIA* PEPCK AS A VACCINATION STRATEGY TO INCREASE EXPRESSION OF SKIN HOMING MOLECULES AND RESIDENT MEMORY T CELL DEVELOPMENT

Anabel Zabala-Peñafiel¹, Claudia Gonzalez-Lombana¹, Anthony Phan¹, Jude Uzonna², Chris Hunter¹, Phil Scott¹

¹University of Pennsylvania, Philadelphia, PA, United States, ²University of Manitoba, Winnipeg, MB, Canada

12:45 p.m.

8425

IMMUNE SIGNATURES PREDICT TREATMENT RESPONSE IN CUTANEOUS LEISHMANIASIS PATIENTS

Rúbia Costa¹, Lucas P. Carvalho¹, Paulo Machado², Edgar M. Carvalho¹, Augusto M. Carvalho¹

¹Gonçalo Moniz Institute, SALVADOR, Brazil, ²Federal University of Bahia, SALVADOR, Brazil

(ACMCIP Abstract)

1 p.m.

8426

MALNUTRITION CONTRIBUTES TO VISCERAL LEISHMANIASIS SEVERITY BY EXACERBATING LIVER PATHOLOGY

Lais Amorim Sacramento¹, Claudia Lombana¹, Lisa Mattei¹, Daniel Beiting¹, Phillip Scott²

¹School of Veterinary Medicine University of Pennsylvania, PHILADELPHIA, PA, United States, ²School of Veterinary Medicine University of Pennsylvania, Philadelphia, PA, United States

(ACMCIP Abstract)

Symposium 183

Closing the Typhoid Surveillance Gap: Operationalizing Novel Surveillance Methods to Describe Enteric Fever Burden and Support Vaccine Decision-Making

Convention Center - Room 353 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Typhoid fever continues to impose a significant health burden worldwide, particularly in low- and middle-income countries where access to clean water and sanitation can be limited. Typhoid affects up to 20 million people annually, claiming 8 million years of healthy life each year. Despite the existence of effective typhoid conjugate vaccines (TCVs), many high-risk communities remain unvaccinated due to the lack of precise, locally relevant epidemiological data. Passive, clinic-based case detection underestimates typhoid incidence, as a consequence of a sub-optimal laboratory reference standard (blood culture) and barriers to health care utilization. Consequently, the cases presenting to a health facility represent only a fraction of the true burden of illness, which makes it difficult for policymakers to accurately estimate the cost effectiveness of TCV introduction. Many Gavi-eligible countries interested in adding TCV to their routine immunization schedules have very limited disease burden data to support their application for Gavi funding, and countries that have introduced TCV are challenged to evaluate the true impact of the vaccine. Our symposium seeks to bridge these data gaps by describing the latest advancements in deploying low-cost, scalable tools to estimate enteric fever disease burden where blood-culture based estimates are not implementable. Seroepidemiology leverages antibody response data to characterize infection burden from cross-sectional population surveys that can be implemented quickly and efficiently. Detection of *Salmonella* Typhi bacteriophages in community water sources is a low-cost method of environmental surveillance to identify geographic areas of potential typhoid transmission. The combination of seroepidemiology and environmental surveillance offers a powerful suite of tools that public health systems can use to monitor disease burden and transmission hotspots. Typhoid intestinal perforations (TIP) are a severe and devastating complication of typhoid disease, with significant medical, economic and social consequences for patients and their families. In settings with limited typhoid surveillance, the presence of TIP can function as an indicator of underlying disease burden in a community, even

in the absence of laboratory confirmation of milder infections. We will present some of the latest findings from studies in Afghanistan, Cote d'Ivoire, India and Niger that are implementing these novel surveillance methods. #Epidemiology #FieldStudies #InfectiousDisease

CHAIR

Jessica Seidman
Sabin Vaccine Institute, Washington, DC, United States

Kristen Aiemjoy
University of California, Davis School of Medicine, Davis, CA, United States

11:30 a.m. INTRODUCTION

11:40 a.m. ENTERIC FEVER SEROSURVEILLANCE IN EASTERN AFGHANISTAN

Ghulam Rahim Awab
Nangarhar University, Nangarhar, Afghanistan

11:55 a.m. PAIRED PHAGE-BASED TYPHI ENVIRONMENTAL SURVEILLANCE AND COMMUNITY-BASED SEROSURVEILLANCE IN COTE D'IVOIRE

Jean Coulibaly
University Félix Houphouët-Boigny, Abidjan, Côte D'Ivoire

12:10 p.m. COMPREHENSIVE S. TYPHI SURVEILLANCE IN VELLORE, INDIA: INTEGRATING WASTEWATER, CLINICAL AND SEROLOGICAL DATA

Dilip Abraham
Christian Medical College Vellore, Vellore, India

12:25 p.m. ADVANCING ENVIRONMENTAL SURVEILLANCE METHODS: LYOPHILIZED REAGENTS FOR PHAGE-BASED DETECTION

Kesia da Silva
Stanford, Palo Alto, CA, United States

12:40 p.m. TYPHOID INTESTINAL PERFORATIONS AS AN INDICATOR OF ENTERIC FEVER DISEASE BURDEN IN GALMI, NIGER

Yakoubou Sanoussi
SIM Galmi Hospital, Madaoua-Kyara, Niger

Scientific Session 184

American Committee of Molecular Cellular and Immunoparasitology (ACMCIP): Parasite Immunology - Insights from Natural and Experimental Infections

Convention Center - Room 354/355 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Supported with funding from the Burroughs Wellcome Fund

#InfectiousDisease #CellBiology #Immunology #HostResponse #Pathogenesis

CHAIR

Usheer Kanjee
Harvard T.H. Chan School of Public Health, Boston, MA, United States

Michelle Boyle
Burnet Institute, Melbourne, Australia

11:30 a.m.

8440

TICK INNATE IMMUNE RESPONSE TO PATHOGEN INFECTION AT SINGLE-CELL RESOLUTION

Abdulsalam Adegoke¹, Jose M. c. Ribeiro², Ryan C. Smith³, Shahid Karim⁴
¹Department of Microbiology and Immunology, University of Michigan Medical School, Ann Arbor, MI, United States, ²Vector Biology Section, Laboratory of Malaria and Vector Research, National Institute of Allergy and Infectious Diseases, National Institutes of Health, Rockville, MD, United States, ³Department of Plant Pathology, Entomology, and Microbiology, Iowa State University, Ames, IA, United States, ⁴School of Biological, Environmental, and Earth Sciences, The University of Southern Mississippi, Hattiesburg, MS, United States

11:45 a.m.

8427

HUMAN FILARIAL INFECTION RESHAPES THE TRANSCRIPTIONAL AND FUNCTIONAL PROGRAMMING OF CD8 T CELLS AT HOMEOSTASIS AND IN RESPONSE TO CYTOMEGALOVIRUS (CMV) IN FILARIAL/CMV COINFECTED INDIVIDUALS

Camila Queiroz Glauss, Thomas B. Nutman
National Institutes of Health, Bethesda, MD, United States

Noon

8428

LATENT CYTOMEGALOVIRUS INFECTION DISRUPTS INNATE AND ADAPTIVE IMMUNITY TO *PLASMODIUM FALCIPARUM* DURING PRIMARY MALARIA INFECTION

Reena Kiran Mukhiya¹, Jessica R. Loughland¹, Jo-Anne Chan¹, Megan Soon², Nick Dooley¹, Luke Quigely¹, Fabian de Labastida Rivera², Dean Andrew², James Beeson¹, James McCarthy³, Bridget E. Barber², Alejandro Lopez⁴, Christian Engwerda², Michelle J. Boyle¹

¹Burnet Institute, Melbourne, Australia, ²QIMR Berghofer Medical Research Institute, Brisbane, Australia, ³Walter and Eliza Hall Institute of Medical Research, Melbourne, Australia, ⁴School of Environmental Sciences, Griffith University, Brisbane, Australia

(ACMCIP Abstract)

12:15 p.m.

8429

HOST DIRECTED THERAPY TO IMPROVE ANTI-PARASITIC IMMUNITY IN VOLUNTEERS EXPERIMENTALLY INFECTED WITH BLOOD STAGE MALARIA

Bridget Barber¹, Rebecca Webster¹, Luzia Bukali¹, Damian Oyong², Fabian de Labastida Rivera¹, Jessica Engel¹, Dean Andrew¹, Megan Soon¹, Nick Dooley², Jessica Loughland¹, James McCarthy³, Gerlinde Obermoser⁴, Michael Leipold⁴, Julianne Jamelink², Reena Mukhiya², Holder Maecker⁴, Christian Engwerda¹, **Michelle Boyle**²

¹QIMR-Berghofer, Brisbane, Australia, ²Burnet Institute, Melbourne, Australia, ³University of Melbourne, Melbourne, Australia, ⁴Stanford University, Stanford, CA, United States

(ACMCIP Abstract)

12:30 p.m.

8430

DIFFERENT MICRORNA PROFILES IN THE CIRCULATING CD4+T CELLS ARE ASSOCIATED WITH DIFFERENT CLINICAL PRESENTATIONS OF LEISHMANIA DONOVANI INFECTION

RITIRUPA ROY¹, Cinthia Hudachek², Shashi Bhushan Chauhan¹, Sundaram Pandey¹, Rajiv Kumar³, Madhukar Rai¹, Mary E. Wilson², Shyam Sundar¹

¹INSTITUTE OF MEDICAL SCIENCES, BANARAS HINDU UNIVERSITY, VARANASI, India, ²University of Iowa and the Iowa City VA Medical Center, Iowa, IA, United States, ³Centre of Experimental Medicine and Surgery, Banaras Hindu University, Varanasi, India, VARANASI, India

(ACMCIP Abstract)

12:45 p.m.

8431

ISOLATION AND CHARACTERIZATION OF A-GAL-CONTAINING EXTRACELLULAR VESICLES FROM TRYPANOSOMA CRUZI: UNVEILING NEW BIOMARKERS FOR CHAGAS DISEASE

Priscila Silva Grijo Farani, Nasim Karimi Hosseini, Susana Portillo, Maria Tays Mendes, Brian Grajeda, Colin Knight, Cameron Ellis, Igor Almeida
The University of Texas at El Paso, El Paso, TX, United States

(ACMCIP Abstract)

1 p.m.

8432

EXTRACELLULAR VESICLES FROM TAENIA SOLIUM DAMPENS PI3K-AKT-MTORC1 SIGNALING AND AMELIORATES DSS-COLITIS IN MICE

Suraj Singh Rawat, Amit Prasad
Indian Institute of Technology Mandi, Mandi, Himachal Pradesh, India

(ACMCIP Abstract)

Symposium 185

The Challenge of Schistosomiasis Diagnosis and Treatment in Young Children and Pregnant Women – The Example of freeBILy

Convention Center - Room 356 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

The WHO NTD 2030 Roadmap targets schistosomiasis for elimination as a public health problem by 2030. There is a need to align national guidelines and strategies in endemic countries and to include all at-risk groups in targeted interventions such as preventive chemotherapy. Particularly, two risk groups are neglected in schistosomiasis prevention and treatment campaigns: preschool-aged children and pregnant women. Both groups are severely affected by schistosomiasis but are neglected in research and policy. To overcome reluctance to include these two vulnerable groups into preventive chemotherapy programs,

a test-and-treat strategy using easy-to-use and affordable diagnostics would be an appropriate approach. The EDCTP-funded freeBILy project included two clinical trials conducted in Gabon and Madagascar from 2018 to 2023. The aim of the project was to assess the safety, efficacy, feasibility and cost-effectiveness of test-based praziquantel treatment of schistosomiasis in pregnant women and young children. In Madagascar, a POC-CCA test-based treatment strategy was implemented in routine maternal health care settings and compared to the highly accurate UCP-LF CAA test. The latter was also implemented in Gabon, along with other diagnostic tests to evaluate efficacy of praziquantel treatment in pregnant women. In this session, we will present the first findings of the project in order to discuss the next steps needed to further push endemic countries to adapt strategies to include young children and pregnant women for preventive praziquantel treatment. After a brief introduction of the freeBILy project, the main findings will be presented within four talks. Specifically, we will discuss the diagnostic outcomes and challenges, the safety and tolerability and efficacy of praziquantel in young children and pregnant women, and the cost-effectiveness of the test-and-treat strategy for schistosomiasis. A final debate between the speakers and the audience will help to identify challenges and opportunities resulting from the freeBILy project and to align them with the goals and targets of the WHO NTD 2030 Roadmap for schistosomiasis (www.freebily.eu). #Diagnostics #Therapeutics #InfectiousDisease #MNCH #ClinicalResearch

CHAIR

Govert van Dam
Leiden University Medical Center, Leiden, Netherlands

Yabo Josiane Honkpehedji
Centre de Recherches Médicale de Lambaréné, Lambaréné, Gabon

11:30 a.m.

INTRODUCTION AND PURPOSE OF THE SESSION

Andrea Kreidenweiss
Universitätsklinikum Tübingen, Tübingen, Germany

Daniela Fusco
Bernhard Nocht Institute for Tropical Medicine, Hamburg, Germany

11:50 a.m.

THE DIAGNOSTIC CHALLENGES OF SCHISTOSOMIASIS: WHY DO WE NEED A POC?

Pytsje T. Hoekstra
Leiden University Medical Center, Leiden, Netherlands

Noon

LESSONS LEARNED FROM THE FIELD - EVALUATION OF THE PERFORMANCE OF CIRCULATING ANODIC ANTIGEN(CAA) MEASUREMENT AS A DIAGNOSTIC TOOL FOR THE DETECTION OF SCHISTOSOMA HAEMATOBIIUM INFECTIONS IN PREGNANT WOMEN AND THEIR NEWBORN AND CHILD IN LAMBARÉNÉ, GABON

Yabo Josiane Honkpehedji
Centre de Recherches Médicales de Lambaréné, Lambaréné, Gabon

12:20 p.m.

LESSONS LEARNED FROM THE FIELD - SCHISTOSOMIASIS AMONG PREGNANT WOMEN IN RURAL MADAGASCAR: A CALL FOR ACTION FOR ADAPTING PUBLIC HEALTH STRATEGIES

Raphael Rakotozandrindrainy
Madagascar Institute for Vaccine Research, University of Antananarivo, Antananarivo, Madagascar

12:30 p.m.

LESSONS LEARNED FROM THE FIELD - EVALUATION OF THE PERFORMANCE OF THE POINT OF CARE CIRCULATING CATHODIC ANTIGEN (POC-CCA) MEASUREMENT AS A DIAGNOSTIC TOOL FOR THE DETECTION OF SCHISTOSOMA MANSONI INFECTIONS IN PREGNANT WOMEN IN RURAL MADAGASCAR

Rivonirina Andry Rakotoarivelo
Université de Fianarantsoa, Fianarantsoa, Madagascar

12:40 p.m.

MODELLING THE COST-EFFECTIVENESS OF TEST AND TREAT STRATEGIES FOR SCHISTOSOMIASIS

Elisa Sicuri
Fundación Privada Instituto de Salud Global Barcelona (ISGlobal), Barcelona, Spain

Symposium 186

Will HAT be the Next NTD to Achieve Global Elimination?

Convention Center - Room 357 (3rd Floor)
Sunday, November 17, 11:30 a.m. - 1:15 p.m.

In the late 1990s, the international community faced a surge in cases of gambiense Human African Trypanosomiasis (gHAT), marking the peak of an epidemic that had persisted since the 1970s. Responding to this public health challenge, global efforts gained momentum in the new millennium, culminating in the publication of the World Health Organisation's (WHO) 2020 and 2030 roadmaps. This strategic framework set ambitious goals for gHAT control, first targeting the disease for elimination as a public health problem (EPHP) then interrupting transmission to humans (EoT). Over the past two decades, national programmes and their international partners have achieved commendable success. Côte d'Ivoire, Uganda, Togo, Benin and Equatorial Guinea have now officially validated EPHP of gHAT. However, as we move past 2020, attention is shifting towards the critical phase of verifying EoT to humans. Despite substantial progress, the rate of decline of cases has slowed since 2018. This underscores the challenges in progressing from very low to zero cases, especially in regions with inadequate infrastructure, fragile healthcare systems or ongoing conflicts. The symposium aims to delve into key aspects of the ongoing battle against gHAT with a particular focus on the challenges faced in achieving EoT. Bringing together experts, researchers and policymakers we will discuss themes including what makes gHAT unique and a target for EoT, reasons to be optimistic, the challenges that lie ahead, the diverse toolkit for gHAT control and transmission interruption, current challenges for control and surveillance and ways to strengthen them, the potential impact of a new single-dose oral drug, acoziborole, current diagnostics and the necessity for new tools to reach EoT,

the extent to which tsetse control is needed to reach EoT, the challenges with current deployment methodologies and alternative approaches, the importance of sustained vigilance after achieving EoT and transmission and health economic modelling as a tool to support the operationalisation and application of interventions. We will also highlight uncertainties and old vs. new strategies as we approach elimination including (i) recommendations for scale-up, where and why (ii) where current strategies should be maintained (iii) where scale-back would be recommended and (iv) where completely new strategies should be introduced and what these constitute. Addressing these key focus areas, the symposium intends to highlight for others in the NTD field the challenges, successes and lessons learned by the HAT community whilst in parallel looking forward to building a comprehensive strategy for the successful elimination of the disease in humans. #Elimination #Epidemiology #InfectiousDisease #Modeling

CHAIR

Emily H. Crowley
University of Warwick, Coventry, United Kingdom

Rachel Bronzan
Bill & Melinda Gates Foundation, Seattle, WA, United States

11:30 a.m.

INTRODUCTION

11:40 a.m.

MOVING TOWARDS THE ELIMINATION OF SLEEPING SICKNESS

Gerardo Priotto
World Health Organization, Geneva, Switzerland

11:55 a.m.

PREPARING FOR THE LAST MILE: INNOVATIONS, OPPORTUNITIES AND CHALLENGES IN THE GHAT ELIMINATION EFFORTS IN THE DRC

Erick Mwamba Miaka
Programme National de Lutte contre la Trypanosomiase Humaine Africaine (PNLTHA), Kinshasa, Democratic Republic of the Congo

12:10 p.m.

LESSONS LEARNED AND CHALLENGES OF VECTOR CONTROL AS PART OF AN INTEGRATED STRATEGY TO ELIMINATE THE TRANSMISSION OF GHAT

Inaki Tirados
Liverpool School of Tropical Medicine, Liverpool, United Kingdom

12:25 p.m.

ACOZIBOROLE: THE DEVELOPMENT OF A SINGLE-DOSE ORAL TREATMENT FOR SLEEPING SICKNESS

Stéphane Hugonnet
Drugs for Neglected Diseases initiative (DNDi), Geneva, Switzerland

12:40 p.m.

WHAT INSIGHTS AND SUPPORT CAN MODELERS PROVIDE ON THE SHORT- AND LONG-TERM BATTLE TOWARDS ELIMINATION?

Kat S. Rock
Warwick University, Coventry, United Kingdom

Sunday
November 17

Symposium 187

Bridging the Rift: Interdisciplinary Approaches to Rift Valley Fever Virus Research

Convention Center - Room 383/384/385 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

This symposium will feature a panel of experts detailing new findings in Rift Valley fever virus (RVFV) research. RVFV periodically re-emerges in epizootics in sub-Saharan Africa that can cause subclinical to severe infection in both ruminants and humans. Indicators of outbreaks include abortion storms in pregnant ewes and cattle and hemorrhagic fever, encephalitis or retinitis in humans. Viral transmission occurs primarily by *Aedes* and *Culex* mosquitoes. However, it can also occur via contact or exposure to infected animal tissue. Vertical transmission of RVFV in mosquitoes has been demonstrated, however, there is an overall lack of understanding of the impact on disease cycles. Climate change and other factors have caused concern that this virus may emerge into new regions, such as Europe or the US. This has stimulated multiple efforts: 1) human vaccine development, 2) study of disease dynamics in US mosquitoes, animals and wildlife and 3) redoubling of surveillance efforts in endemic countries, e.g. Tanzania. Highlights from the field will include gaps in the current understanding of RVF disease ecology and the status of clinical development of a human vaccine. Discussion of surveillance will feature the ways in which inter-epidemic transmission of RVFV across ecologies in Tanzania is addressed by community- and hospital-based surveillance, cross-sectional and sentinel animal epidemiological studies, and how ongoing transmission in absence of epidemic/epizootics fits into the bigger picture of RVF disease ecology. Molecular highlights will cover advances in tools used to understand RNA reassortment between co-infecting strains in mosquitoes and the ways in which viral reassortment affects viral kinetics. Our data demonstrates reassortment between virulent and vaccine strains occurs in mosquito midgut and salivary tissue. Reassortant virus recovered from these tissues, however did not lead to enhanced replication phenotypes compared to non-reassortant parental strains. Another presentation will report on animal and *in vitro* studies that reveal alterations in interferon lambda and key cytokines/chemokines during the inflammatory response to RVFV. scRNA-seq of *Culex tarsalis* ovaries was performed to generate tools for improved characterization of RVFV vertical transmission. Thirteen cell clusters were identified, representing nurse cells, germ cells, stretch cells and hemocytes. *In vivo* characterization by smFISH reveals that, following a round of egg-laying, ovarian stretch cells engulf nurse cells, presumably for nutrient recycling, as indicated by the presence of vitellogenin receptor mRNA in specific cell clusters. Lastly, exploring genomic regulation of RVFV infection in mosquitoes promises to reveal the underpinnings of vector competence. #EmergingDiseaseThreats, #Genomics, #Vaccinology, #MolecularBiology

CHAIR

Corey Campbell
Colorado State University, Fort Collins, CO, United States

Emma Harris
Colorado State University, Fort Collins, CO, United States

11:30 a.m.
INTRODUCTION

11:40 a.m.
RIFT VALLEY FEVER VIRUS: RISK OF EMERGENCE INTO EUROPE AND NORTH AMERICA

Chad Mire
USDA-ARS-National Bio-Agro-Defense Facility, Manhattan, KS, United States

11:55 a.m.
RIFT VALLEY FEVER IN TANZANIA: INTER-EPIDEMIC TRANSMISSION ACROSS ECOLOGIES AND PROSPECTS FOR ITS CONTROL

Robert David Sumaye
Ifakara Health Institute, Ifakara, United Republic of Tanzania

12:10 p.m.
EVALUATION OF IMMUNOMODULATORY TREATMENT STRATEGIES FOR RIFT VALLEY FEVER VIRUS

Darci R. Smith
Naval Medical Research Command, Biological Defense Research Directorate, Frederick, MD, United States

12:25 p.m.
DEFINING REGULATION OF RVFV INFECTION IN VECTOR MOSQUITOES

Corey L. Campbell
Colorado State University, Fort Collins, CO, United States

12:40 p.m.
CO-INFECTION OF CULEX TARSALIS WITH RIFT VALLEY FEVER PHLEBOVIRUS PRODUCES EFFICIENT SEGMENTAL REASSORTMENT ACROSS MIDGUT AND SALIVARY GLAND TISSUES

Emma Harris
Colorado State University, Fort Collins, CO, United States

12:55 p.m.
THE HALI PROJECT: RIFT VALLEY FEVER VACCINE DEVELOPMENT AND SURVEILLANCE IN TANZANIA - GLOBAL PARTNERSHIPS FOR GLOBAL GOOD

Brian Bird
Viral Special Pathogens Branch, University of California, Davis, Davis, CA, United States

Symposium 188

Integrating Anthropology/Social Science Approaches Into Epidemic Outbreak Response

Convention Center - Room 388/389 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Anthropological and social science approaches have gained recognition in outbreak response since the 2014-16 West African Ebola outbreak, and the importance of understanding social dynamics in order to design effective outbreak response interventions has been reconfirmed during the Covid-19 pandemic. As a result, demand for social scientists, and anthropologists in particular, in outbreak response operations has increased significantly and we have successfully integrated these new data streams into epidemic responses to cholera in Malawi, Zimbabwe and Zambia over the past year alone. While largely

sitting under the RCCE pillar in the IMS structure, the findings and recommendations of social science research have much broader operational potential and have been taken up across pillars in these responses. We would like to take this opportunity to reflect on the successes and challenges of embedding social science approaches across the response pillars; the standardization of tools and frameworks; and the broader operational implications for developing sustainable approaches to epidemic preparedness and response. The panelists will discuss their experience of having participated in recent outbreak responses in different capacities – as government officials leading outbreak pillars, as academics joining outbreak investigation assessments for rapid action, and as technical experts representing major agencies' surge teams. Topics to be discussed revolve around methodological, political and operational concerns, including: • How to integrate community-based feedback data and anthropological arguments into the evidence base for decision-making in a rapidly shifting environment? This will require a discussion about the difficulties of navigating rapid research in emergency mode when traditional social science methods promote slow, long-term research and the challenges of working within an environment where large sample sizes often carry more weight than small-scale qualitative investigations, and where social-science and community-based data streams are still fairly new. • What are the political and programmatic implications of giving voice to the most marginalized in society, who often bear the heaviest burden of cases and case fatalities? Panelists will debate the challenges produced when these voices may highlight uncomfortable truths that make government and international partners' shortcomings visible, and the dilemmas created when actions recommended as critical by the community fall outside of planned response activities. The panelists will reflect on the value of social science insights and develop a roadmap for integrating social science data streams into the structure of epidemic preparedness and response. #SocialScience #InfectiousDisease

CHAIR

Nadine Beckmann
LSHTM, London, United Kingdom

11:30 a.m. **INTRODUCTION**

11:35 a.m. **RAPID QUALITATIVE ASSESSMENT AND COMMUNITY FEEDBACK AS DATA STREAMS FOR OUTBREAK RESPONSE**

Christina Craig
US CDC, Atlanta, United States

11:55 a.m. **INTEGRATING COMMUNITY FEEDBACK MECHANISMS INTO OUTBREAK RESPONSE**

Ngonizadshe Nyambarawo
IFRC/Collective Service, Pretoria, South Africa

12:15 p.m.

SOCIAL SCIENCE RESEARCH FOR COMMUNITY ENGAGEMENT IN THE 2023 CHOLERA OUTBREAK IN ZIMBABWE

Sophie Everest
UKHSA, London, United Kingdom

12:35 p.m. **OPERATIONALIZING SOCIAL SCIENCE RESEARCH IN THE 2024 CHOLERA OUTBREAK IN ZAMBIA**

Hanna Woldemeskale
Unicef, Lusaka, Zambia

12:55 p.m. **OPERATIONALIZING AND BUILDING CAPACITY FOR RAPID QUALITATIVE ASSESSMENTS (RQAS) FOR CRISIS RESPONSE IN ZIMBABWE**

Loveness Chimombe
Unicef, Harare, Zimbabwe

Symposium 189

Scaling the Optimal Use of Multiple ACT's to Prevent Antimalarials Drug Resistance: Progress and Challenges

Convention Center - Room 391/392 (3rd Floor)
Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Prompt treatment with effective antimalarials is a crucial element of malaria control. WHO currently recommends the use of Artemisinin-Based Combination Therapies (ACTs) as the first-line treatment. However, partial artemisinin resistance has been confirmed in a number of African countries, notably Rwanda, Uganda, Tanzania and Ethiopia, threatening the longevity of these treatments. Fortunately, despite reduced efficacy of artemether-lumefantrine in some countries, to date ACTs remain largely effective in Africa and there have been no confirmed reports of partner drug failure. Nevertheless, innovative solutions are required to diversify the use of and protect the efficacy of currently available ACTs until new compounds become available. Multiple First-line Therapies (MFTs) in which several treatments are used concurrently, have been shown to slow down resistance evolution compared with using a single therapy for the whole population. Pilot studies on country readiness to implement MFTs have been conducted in Burkina Faso and Kenya. Several other countries including Rwanda have adopted MFTs as strategy in their national strategy and have begun implementation. This symposium is composed of five presentations that outline the evidence on the effectiveness of MFTs as well as supply, demand, implementation, and operational research issues relevant to the adoption and scale up of MFTs. The first presentation will provide an overview of drug resistance and the need for innovative strategies. The second presentation will share the evidence of the effectiveness of various MFT strategies in Central and East Africa derived from mathematical models. The third presentation will highlight how Rwanda has adopted, generated demand and begun implementation of this strategy. The fourth presentation will outline the operational research being undertaken by Nigeria

to document best practices that can be applied to the adoption and scale up of new therapies and strategies including MFTs in other countries in Africa wishing to implement this strategy. The last presentation will discuss the supply issues for new therapies that form part of the strategy to diversify the use of ACTs. The symposium will be moderated by Dr. Estée Török from the Bill & Melinda Gates Foundation. #InfectiousDisease #Resistance #Modeling

CHAIR

Rima Shretta
Jhpiego, Baltimore, MD, United States

Estée Török
Bill & Melinda Gates Foundation, London, United Kingdom

11:30 a.m. INTRODUCTION

11:40 a.m. OVERVIEW OF DRUG RESISTANCE AND THE NEED FOR INNOVATIVE TREATMENT STRATEGIES

Rima Shretta
Jhpiego, Baltimore, MD, United States

11:45 a.m. APPROACHES TO SLOWING DOWN THE CURRENTLY CIRCULATING KELCH13 VARIANTS IN CENTRAL AND EASTERN AFRICA: EVIDENCE ON THE EFFECTIVENESS OF MULTIPLE FIRST-LINE THERAPIES USING MODELING

Maciej F. Boni
Temple University, Philadelphia, PA, United States

Noon FROM ADOPTION TO IMPLEMENTATION OF A POLICY OF MULTIPLE FIRST-LINE THERAPIES IN RWANDA

Aimable Mbituyumuremyi
Rwanda Biomedical Centre, National Malaria Program, Kigali, Rwanda

12:15 p.m. OPERATIONAL RESEARCH TO INFORM THE MULTIPLE FIRST-LINE THERAPY STRATEGY IN NIGERIA

Olugbenga A. Mokuolu
University of Ilorin Teaching Hospital, Ilorin, Nigeria

12:30 p.m. GLOBAL EFFORTS TO DIVERSIFY THE USE OF ARTEMISININ-BASED COMBINATION THERAPIES: SUPPLY CHALLENGES

Andre M. Tchouatieu
Medicines for Malaria Venture, Meyrin, Switzerland

Symposium 190

Clinical Development of Monoclonal Antibodies that Target Malaria Sporozoites

Convention Center - Room 393/394 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

The aim of this symposium is to bring together leaders in the field to present results from Phase 2 trials of L9LS in Mali and Kenya, and a Phase 1 Challenge Study in the US for MAM01. The symposium will also explore delivery and cost of good considerations. The symposium will provide a forum for the community to discuss next steps in the clinical development and registration of mAbs including potential target populations. Each presenter will provide an overview on prior experience exploring some of these concepts in prior and current projects, but also will discuss important concepts to explore in future mAb clinical trials. Ample time will be provided at the end of the symposium to promote Q&A and discussion from the audience. #Immunology #Prevention #InfectiousDisease #Pediatrics #TranslationalScience

CHAIR

Kayla Andrews
Bill & Melinda Gates Medical Research Institute, Cambridge, MA, United States

Aissata Ongoiba
MRTC/ICERMALI, Bamako, Mali

11:30 a.m. INTRODUCTION

11:40 a.m. COST OF GOODS OF PROPHYLACTIC MONOCLONAL ANTIBODIES FOR *PLASMODIUM FALCIPARUM* MALARIA

Hong Liu
Bill & Melinda Gates Medical Research Institute, Cambridge, MA, United States

11:55 a.m. SAFETY AND EFFICACY OF SUBCUTANEOUS ADMINISTRATION OF L9LS IN MALIAN WOMEN OF CHILD-BEARING POTENTIAL

Aissata Ongoiba
MRTC/ICERMALI, Bamako, United States

12:10 p.m. SAFETY, PHARMACOKINETICS AND EFFICACY OF REPEATED DOSING OF L9LS IN MALIAN CHILDREN

Safiatou Doumbo
MRTC/ICERMALI, Bamako, Mali

12:25 p.m. SAFETY, PHARMACOKINETICS, AND EFFICACY OF REPEATED DOSING OF L9LS IN KENYAN INFANTS AND CHILDREN

Titus Kwambai
US Centers for Disease Control, Kisumu, Kenya

12:40 p.m. MAM01: SAFETY, PK AND PRELIMINARY EFFICACY OF A NEW 1ST GEN CSP MONOCLONAL ANTIBODY

Kirsten Lyke
University of Maryland School of Medicine, Baltimore, United States

Symposium 191

The Changing Face of Mapping: Exploring Disease Monitoring Challenges and Innovations After Long-Term Implementation of Helminth Control Programs

Convention Center - Room 395/396 (3rd Floor)

Sunday, November 17, 11:30 a.m. - 1:15 p.m.

Since the advent of Preventative Chemotherapy (PC) as a key control measure for soil-transmitted helminths (STH) and schistosomiasis (SCH), the prevalence of both diseases has reduced globally. Many previously endemic countries now face a very low or highly heterogeneous prevalence landscape. Current guidelines suggest that the frequency of treatment be slowly scaled down until suspension in line with disease prevalence. Guidelines however, lack sufficient nuance around cost-effectively scaling back programs as decision making at a national or equivalent scale may miss endemic pockets of infection or areas of risk where continued or resurging infection may occur. A key determining factor for the future of PC programs is how monitoring is performed. Mapping in such environments is hindered primarily by three key issues 1) given the investment needed to test and treat children for STH and SCH at scale, and that most infections do not tend to trigger individuals to seek testing at health establishments, large-scale representative surveys remain necessary for programmatic decision making. 2) Traditional diagnostics may face issues around precision as prevalence and intensity of infections reduce. Current available alternatives are hindered by prohibitive costs. 3) Given the heterogeneity of prevalence, insufficient guidance exists on how to map sub-nationally and at what level optimal decision making should occur. In addition, geographies may now have multiple goals across a single country. The symposium will explore key issues and novel solutions for monitoring disease patterns as prevalence reduces, focusing on programmatically relevant decision-making and cost-effectiveness. The overarching goal of the symposium is to foster discussion on the issues currently facing decision making for future STH/SCH mapping. Differing from most other discussions a key focus will be what is achievable with cost-effectiveness in mind. Further goals include: Identifying novel survey design which can cost-effectively deal with multiple competing goals presenting helminth control programs after many years of treatment; high light how impactful novel diagnostics which can be cost-effective at scale and their potential for use in maturing helminth control programs; explore case-studies which researchers and program managers can use to approach their own cost-effective programming as programs mature. #Epidemiology #InfectiousDisease #PopulationSurveillance

CHAIR

Mark Minnery

Evidence Action, Washington, DC, United States

Judd Walson

Johns Hopkins University, Baltimore, United States

11:30 a.m.

INTRODUCTION

11:40 a.m.

THE CHANGING FACE OF HELMINTH MAPPING: KEY ISSUES WITH SURVEY DESIGN AND ANALYSIS FOR COST-EFFECTIVE SUB-NATIONAL MAPPING

Kate McCracken

Evidence Action, Washington DC, United States

11:55 a.m.

TRANSITIONING FROM DISTRICT TO SUB DISTRICT MONITORING FOR SCHISTOSOMIASIS: CHALLENGES AND WAY FORWARD

Ndellejong Cosmas Ejong

Sight Savers, Calgary, Canada

12:10 p.m.

EXPLORING SOIL-TRANSMITTED HELMINTH INFECTION PREVALENCE HETEROGENEITY: INSIGHTS FROM RECENT IMPACT ASSESSMENTS IN TANZANIA

Mohamed Ramadhan Nyati

Neglected Diseases Control Program (NTDCP), Ministry of Health, United Republic of Tanzania

12:20 p.m.

EXPLORING SOIL-TRANSMITTED HELMINTH INFECTION PREVALENCE HETEROGENEITY: INSIGHTS FROM RECENT IMPACT ASSESSMENTS IN TANZANIA

Stephen Mbwambo

Tanzania Ministry of Public Health, Dodoma, United Republic of Tanzania

12:35 p.m.

UTILITY AND FEASIBILITY OF HIGH THROUGHPUT MOLECULAR DIAGNOSTICS TO ENHANCE STH SURVEILLANCE

Sitara Swarna Rao Ajjampur

CMC-Vellore, Tamil Nadu, India

12:50 p.m.

GEOSTATISTICAL METHODS FOR SURVEY DESIGN IN LOW PREVALENCE SETTINGS

Freya Clark

Lancaster University, Lancaster, United Kingdom

Sunday, November 17, 1:15 p.m.

ASTMH 2024 Annual Meeting Adjourns

See you next year at the 2025 Annual Meeting
November 9-13, 2025 (Sunday through Thursday)

Metro Toronto Convention Centre

Toronto, Ontario, Canada

Sunday
November 17







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