

Expert meeting on dengue diagnosis and risk prediction, with a focus on improving clinical management and integration with IMCI/IMAI

Ho Chi Minh City

21st-23rd September 2015

Oxford University Clinical Research Unit - Vietnam (OUCRU-VN) Hospital for Tropical Diseases, 764 Vo Van Kiet, District 5 Ho Chi Minh City, Vietnam

This meeting is being convened on behalf of WHO-TDR, the EC-FP7 funded IDAMS Dengue Consortium, the Oxford University Clinical Research Unit and the Hospital for Tropical Diseases of Ho Chi Minh City. The meeting will be held in the main OUCRU meeting room in the Research Building of the Hospital for Tropical Diseases.

The three main objectives of the meeting are a) to discuss the current progress and expected impact of the clinical observational IDAMS study on early dengue; b) to evaluate the potential need for adaptation of the IMCI algorithm for dengue, and c) to discuss efforts towards developing operational definitions for intermediate and severe dengue for use in therapeutic intervention trials & clinical research.

Background

Agenda

List of participants

Background

Improvements in clinical diagnosis and risk prediction for severe disease are urgently needed in order to improve management of dengue, especially in settings with a high case burden where appropriate allocation of limited resources is crucial to outcome.

In the ongoing clinical observational study currently being carried out by members of the IDAMS consortium (www.idams.eu), outpatients with undifferentiated fever and a history consistent with possible dengue are being recruited within 3 days of fever onset in 5 Asian countries (Bangladesh, Cambodia, Indonesia, Malaysia and Vietnam), and then followed daily until full recovery. The same study is also being carried out across a number of Latin American countries. The main aims of the study are 1) to identify clinical features and simple laboratory parameters that differentiate dengue from other febrile illness in the febrile phase, and 2) among study participants with confirmed dengue, to evaluate risk factors for progression to more severe disease. One of the main objectives of this meeting is to bring together senior clinicians and public health specialists from across the Southeast Asian region to discuss how the data being generated (>5000 participants to date across Asia) can best be utilized to improve dengue guidelines for disease classification and management. As well as contributing to the development of diagnostic and prognostic algorithms this very large clinical cohort has considerable potential towards standardization of procedures for dengue management and reporting.

Guidelines for the integrated management of childhood illness (IMCI) for use in children below 5 years of age, as well as for adolescent and adult illness (IMAI), are published by WHO and then adapted for local use in many resource limited countries, with a particular focus on management at the primary healthcare level. However, evaluations of the utility of IMCI guidelines have indicated that there are significant limitations with respect to the assessment of febrile illness in dengue endemic countries, and it has been suggested that in such countries the guidelines should be adapted to reduce the focus on dengue among children <5 years, while also including/developing a section directed towards identifying dengue among older children and adolescents presenting with febrile illness.

The third objective of the meeting relates to a new initiative aimed at developing a standardized system for classification of moderate and severe dengue specifically for use in research studies. As the burden of dengue increases globally, research to measure the impact of potential interventions (vaccines, therapeutic agents or vector control measures) is becoming ever more important, and it has become clear that consensus is lacking among the scientific community on how to define moderate and severe dengue for research purposes. At an initial meeting in Washington in April 2015, co-sponsored by NIAID/NIH, WHO and PDC, an initiative was launched aiming to develop criteria for moderate and severe dengue that are sufficiently detailed to allow comparability between different studies and across different research groups, but are also integrated with the classification systems used for clinical management and public health related surveillance.

Meeting Objectives

- 1. To discuss the progress of the clinical observational study in IDAMS and review the implications for diagnosis, classification and clinical management of dengue:
- 2. To discuss the need for adaptation of the IMCI algorithm for dengue
 - a. Extension of the age range for IMCI and/or integration with IMAI
 - b. Harmonization of guidelines
- 3. To discuss efforts towards developing operational definitions of moderate/severe dengue for therapeutic intervention trials & clinical research

AGENDA

Day 1 – September 21 st (Participants arrive the day before)			
Time	Торіс	Presenter	
08.00 - 08.30	Transport from hotels to OUCRU / HTD site		
08.30 - 08.45	Welcome and introduction of participants	Bridget Wills	
08.45 - 09.00	Opening Address by Professor Tran Dac Phu - Director of the Centre for Preventive Medicine, Hanoi, Vietnam		
09.15 - 09.45	Scientific key note: "Adult Dengue in Southeast Asia"	Prof Yee-Sin Leo, Singapore	
09.45 - 10:00	Overview of dengue epidemiology in China	Dr Yu Li, China CDC	
10.00 - 10.15	Overview of dengue epidemiology in Taiwan	Prof CC King, Taiwan	
10.15 - 10.45	Coffee break		
	UPDATE ON THE IDAMS STUDY		
10.45 - 11.15	Update on the current state of the IDAMS study (recruitment, monitoring, data management, timelines)	Thomas Jaenisch	
11.15 – 13.00	Update on the current state IDAMS study (perspectives from the various recruiting sites) - Bangladesh - Cambodia - Indonesia - Malaysia - Vietnam	Malabika Sarkar Chuop Bhopal Ida Safitri Lucy Lum Dong Thi Hoai Tam	
13.00 - 14.00	Lunch break		
14.00 - 15.30	 Group work (4 groups) Dengue versus other febrile illness in the early febrile phase Role of current warning signs for 		
	decision to hospitalize		
	- Hospitalization and IV fluids as a proxy of disease severity		
	- Adult dengue – how different from paediatric?		
15.30 - 16.00	Coffee break		
16.00 - 18.00	Presentation of group work, discussion (plenary)		
19.30	Group Dinner (Hoa Tuc Restaurant)		

Day 2 – September 22 nd			
Time	Торіс	Presenter	
ADAPTATION OF IMCI GUIDELINES			
08.30 - 09.00	Dengue Guidelines in the Context of IMCI	Martin Weber	
09.00 - 09.30	Development and Validation of IMCI Guidelines for Dengue	Jackie Deen	
09.30 - 10.00	Current state – different IMCI and clinical guidelines in Asia	Thomas Jaenisch / Nay Wyine	
10.00 - 10.30	Coffee break		
10.30 - 12.00	Perspectives on IMCI / IMAI from leading clinicians, local MOH / WHO representatives in Asia	Pheaktra Ngoun, Cambodia Doan Thi Ngoc Diep, Vetnam - NN 3	
12.00 - 13.30	Lunch break		
DEVELOPMENT OF MODERATE / SEVERE DENGUE DEFINITIONS			
13.30 - 13.45	Efforts towards developing operational definitions of moderate/severe dengue for dengue intervention trials & clinical research	General Introduction Kay Tomashek	
13.45 - 14.30	Proposed definitions for moderate/severe plasma leakage	Bridget Wills (20 mins) Group Discussion (25 mins)	
14.30 - 15.15	Proposed definitions for moderate/severe bleeding	Lucy Lum (20 mins) Group Discussion (25 mins)	
15.15 - 15.45	Coffee break		
15.45 - 16.15	Proposed definitions for moderate/severe CNS and liver involvement	Kay Tomashek (20 mins) Group Discussion (25 mins)	
16.15 - 16.45	Plans for validating moderate/severe disease definitions for dengue intervention trials	Thomas Jaenisch (20 mins) Group Discussion (25 mins)	
16.45 - 17.30	Plenary discussion: intermediate/severe disease definitions		
19.30	Group Dinner (Cuc Gach Quan)		

Day 3 – September 23rd				
Time	Торіс	Rapporteur		
08.30 - 09.45	Wrap up and closing discussion – IMCI	Jacqueline Deen		
09.45 - 11.00	Wrap up and closing discussion – intermediate/severe disease definitions	Sophie Yacoub		
11.00 - 11.30	Coffee break			
11.30 - 12.30	Future opportunities for regional collaboration?	Bridget Wills		
12.30 - 13.00	Closure of the meeting			
13.00 - 14.00	Lunch			
14.00	Return to hotels and departure of participants			

List of participants

Expert meeting on dengue clinical management and IMCI

A) Countries in Asia (Clinicians, researchers, MOH representatives)

Bangladesh

1. Dr Firoz Ahmed

Associate Scientist & Head Molecular & Serodiagnostic Lab Clinical Laboratory Services, icddr,b Dhaka, Bangladesh

2. Prof Malabika Sarker

Director Research James P Grant School of Public Health, BRAC University 5th Floor, (Level- 5), icddr,b Building 68 Shahid Tajuddin Ahmed Sharani, Mohakhali Dhaka-1212, Bangladesh

3. Imran Ahmed

Research Associate James P Grant School of Public Health, BRAC University 5th Floor, (Level-6), icddr,b Building 68 Shahid Tajuddin Ahmed Sharani Mohakhali, Dhaka-1212, Bangladesh

4. MD. Mahmubul Alam

Senior Research Officer Molecular& Serodiagnostic Laboratory, icddr,b Clinical Laboratory Services, icddr,b Dhaka, Bangladesh

5. Prof. Ridwanur Rahman

Professor at Shaheed Suhrawardy Medical College and Hospital Sher-E-Bangla Nagar, Dhaka – 1207, Bangladesh

Cambodia

6. Dr. Pheaktra Ngoun

Medical Executive Director Angkor Hospital for Children Tep Vong & Um Chhay Street Mondul 1, Svay Dangkum, PO Box 50, Siem Reap Kingdom of Cambodia

7. Dr Chuop Bophal

Research Clinician Angkor Hospital for Children Tep Vong &Um Chhay Street Mondul 1, Svay Dangkum, PO Box 50, Siem Reap, Kingdom of Cambodia

China

8. Dr Yu Li

Research Assistant, Division for Infectious Diseases Chinese Center for Disease Control and Prevention (China CDC)

Indonesia

9. Dr. Ida Safitri Laksono

Tropical and Infectious Disease Sub Division Paediatric Department Dr. Sardjito Hospital/Faculty of Medicine, Gadjah Mada University Yogyakarta 55281, Indonesia

10. Dr. Yodi Mahendradhata

GMU Centre for Tropical Medicine Gedung PAU-UGM Jl. Teknika Utara, Barek Yogyakarta 55281, Indonesia

Japan

11. Dr Norio Ohmagari,

Director, Disease Control and Prevention Centre National Centre for Global Health and Medicine Tokyo, Japan

Laos

12. Professor Douangdao SOUK ALOUN, MD, PhD.

Deputy Director of Mahosot Hospital Acting President of Council of Medical Sciences P.O. Box: 1720, Vientiane Capital, Lao PDR

13. Dr Bandith Soumphonphakdy

Paediatric Intensive care Unit, Mahosot Hospital, P.O. Box: 1720, Vientiane Capital, Lao PDR

Malaysia

14. Professor Lucy Lum

Department of Paediatrics, Faculty of Medicine University of Malaya Kuala Lumpur, Malaysia

15. Dr Suresh Kumar Chidambaram

Infectious Disease Consultant HOSPITAL SUNGAI BULOH, MoH Malaysia Kuala Lumpur, Malaysia

Myanmar

16. Dr. Nay Yee Wyine

Project Coordinator, Myanmar Neglected Tropical Diseases Research Collaboration University of Public Health No.246, Myo Ma Kyaung Street, Yangon, Myanmar

The Philippines

17. Dr Nimfa M. Putong

Department Head, Public Health Office 237 Rizal Avenue St. Bo. Balite, Montalban, Rizal.

Singapore

18. Professor Yee Sin Leo

Head of the Institute of Infectious Diseases and Epidemiology Tan Tock Seng Hospital 11 Jalan Tan Tock Seng Singapore 308433

19. Prof. Annelies Wilder-Smith

Professor of Infectious Diseases Director, Global Health and Vaccinology Programme Lee Kong Chian School of Medicine 11 Mandalay Road Singapore 308232

Sri Lanka

20. Dr Hasitha Tisera

Epidemiologist, Ministry of Health Sri Lanka 231, De Saram Place, Colombo 10 Sri Lanka

Taiwan

21. Prof. King, Chwan-Chuen

Institute of Epidemiology & Preventive Med College of Public Health National Taiwan University

Thailand

22. Dr. Piyarat Suntarattiwong

Department of Pediatrics Queen Sirikit National Institute of Child Health 420/8 Rajavithi Rd. Bangkok, Thailand

23. Ms. Rasana Waleerattanapa

Chief Nurse, Dengue Unit, WHO Collaborating Centre for Case Management of Dengue/DHF/DSS, Queen Sirikit National Institute of Child Health (QSNICH) Bangkok, Thailand

Vietnam

24. Assoc. Prof. Dr. Tran Dac Phu

Director General, Department of Preventive Medicine Ministry of Health, Vietnam 135 Nui Truc, Giang Vo, Ba Dinh, Ha Noi, Viet Nam

25. Dr Ha Manh Tuan,

Director, Children's Hospital Number 2 4 Lý Tự Trọng, Quận 1, Ho Chi Minh City, Viet Nam

26. Dr Nguyen Thanh Hung

Director, Children's Hospital Number 1 341 Sư Vạn Hạnh, Quan 10 Hồ Chí Minh City, Viet Nam

27. Prof. Dr Phan Trong Lan

Director, Pasteur Institute/Ho Chi Minh City Ministry Of Health 167 Pasteur Street, Quan 3, Ho Chi Minh City, Viet Nam

B) WHO representatives

28. Dr. Martin Weber

Programme Manager, Child and Adolescent Health WHO Regional Office for Europe Marmorvej 51, 2100 Copenhagen, Denmark

C) <u>Representatives from countries in Europe</u>

Germany

29. Dr. Thomas Jäenisch

Coordinator IDAMS Section Clinical Tropical Medicine Department of Infectious Diseases Heidelberg University Hospital INF 324, 69120 Heidelberg Germany

30. Dr. Kerstin Rosenberger

Section Clinical Tropical Medicine Department of Infectious Diseases Heidelberg University Hospital INF 324, 69120 Heidelberg Germany

31. Marius Wirths MA

Section Clinical Tropical Medicine Department of Infectious Diseases & Department of Biometrics Heidelberg University Hospital INF 324, 69120 Heidelberg Germany

32. Dr. Silvia Runge-Ranzinger

Consultant to WHO/TDR Affiliated to the Institute of Public Health University of Heidelberg INF 365, 69120 Heidelberg, Germany

D) External

33. Dr. Jacqueline Deen

Independent Consultant Bangkok 10400, Thailand

34. Dr. Kay Tomashek

Division of Microbiology and Infectious Diseases National Institute of Allergy and Infectious Diseases National Institutes of Health 5601 Fisher Lane, Bethesda, MD, USA

E) Local Organizers / Representatives

- 35. Prof. Bridget Wills
- 36. Prof. Cameron Simmons
- 37. Prof. Dong Thi Hoai Tam
- 38. Dr Nguyen Tran Nam
- 39. Dr Doan Thi Ngoc Diep
- 40. Dr Marcel Wolbers
- 41. Dr Phung Khanh Lam
- 42. Dr Nguyen Minh Tuan
- 43. Dr Hoang Quoc Cuong
- 44. Dr Sophie Yacoub
- 45. Dr Dinh The Trung
- 46. Ms Nguyen Tan Thanh Kieu
- 47. Dr. Nguyen Thanh Ha Quyen
- 48. Dr. Elisabeth Giger
- 49. Ms Trang Vo Hong Yeng