ABSTRACTS OF ORAL PRESENTATIONS

OC 8173 RAPID DETECTION OF *MYCOBACTERIUM ULCERANS* BY RECOMBINASE POLYMERASE AMPLIFICATION

¹Michael Frimpong*, ¹Hubert Ahor, ¹Francisca Sarpong, ²Ken Laing, ²Mark Wansbrough-Jones, ¹Richard Phillips. ¹Kumasi Centre for Collaborative Research in Tropical Medicine, Knust, Ghana; ²St. George's University of London, London, UK

10.1136/bmjgh-2019-EDC.2

Background There are no primary measures to prevent people from contracting Buruli ulcer, mainly due to poor understanding of its epidemiology. The current control strategy emphasises early diagnosis and prompt treatment, with the goal of avoiding the complications associated with advanced stages of the disease. There is no diagnostic test for the disease appropriate for use at the primary health care level where most cases are detected and treated. Diagnosis based on clinical signs is unreliable in inexperienced hands and complicated by infections that have similar presentations. This study was to develop and evaluate the use of recombinase polymerase amplification (RPA) assay for the detection of *Mycobacterium ulcerans* at the point of patient care.

Methods A specific fragment of IS2404 of *M. ulcerans* was amplified in 15 min at a constant temperature of 42°C, using the RPA assay and analysed on a portable fluorometre. The method was tested for sensitivity and specificity with molecular standard of IS2404 DNA fragment, various *M. ulcerans* strains, other mycobacteria and environmentally associated bacteria. Additionally, the assay performance as a diagnostic tool was tested with archived DNA from symptomatic patients. All results were compared with that of a highly sensitive IS2404 PCR.

Results The detection limit was 50 copies of IS2404 in 15 min using plasmid standard and 125 fg with genomic *Mu* DNA equivalent 25 genomic copies. The assay was highly specific in detecting all strains of *M. ulcerans* with no observed cross reactivity with other mycobacteria and common skin colonising bacteria. The clinical sensitivity and specificity of the BU-RPA assay using clinical samples was 86% and 100% respectively.

Conclusion We have developed a real-time isothermal RPA assay for the detection of *M. ulcerans* as a cheaper alternative to PCR. Combining this assay with a simple extraction protocol will maximise its use as point-of-care test for Buruli ulcer.

OC 8215 DIAGNOSTIC ACCURACY OF XPERT MTB/RIF ULTRA FOR PULMONARY TUBERCULOSIS IN CHILDREN: A MULTICENTRE COMPARATIVE STUDY

¹Issa Sabi, ^{2,3}Andrea Rachow, ¹Daniel Mapamba, ¹Petra Clowes, ¹Nyanda E Ntinginya, ⁴Mohamed Sasamalo, ⁴Lujeko Kamwela, ^{4,5,6}Frederick Haraka, ^{2,3}Michael Hoelscher, ^{5,6}Daniel Paris, ^{2,3}Elmar Saathoff, ^{4,5,6}Klaus Reither*. ¹NIMR-Mbeya Medical Research Center, Tanzania; ²Division of Infectious Diseases and Tropical Medicine, Medical Center of the University of Munich, Germany; ³German Centre for Infection Research (DZIF), Partner Site Munich, Germany; ⁴Ifakara Health Institute, Tanzania; ⁵Swiss Tropical and Public Health Institute, Switzerland; ⁶University of Basel, Switzerland

10.1136/bmjgh-2019-EDC.3

Background Xpert® MTB/RIF (Xpert) has suboptimal sensitivity for the diagnosis of pulmonary tuberculosis in children. The next-generation Xpert MTB/RIF Ultra assay (Xpert Ultra) is substantially more sensitive than Xpert and may allow

improved detection of paediatric tuberculosis. We evaluated the diagnostic accuracy of Xpert Ultra versus Xpert in the detection of pulmonary tuberculosis in children.

Methods From May 2011 to September 2012, children with presumptive pulmonary tuberculosis were enrolled at two Tanzanian research sites in the EDCTP-funded TB CHILD project. Sputum samples were collected and examined for tuberculosis using sputum smear, Xpert and culture. Xpert Ultra analysis was performed between January and June 2017 at both sites using decontaminated sputum pellets which had been stored at -80°C. The diagnostic accuracy of Xpert and Xpert Ultra was determined using well-defined case definitions as reference standard.

Results In total, 215 children were included in the analysis. The median age was 5.4 years (IQR, 1.5 to 9.9), the HIV prevalence was 52%, and 28 children (13%) had culture-confirmed pulmonary tuberculosis. When only the first available sample of each patient was considered, the sensitivity of Xpert Ultra was 64.3% (95% CI: 44.1 to 81.4) while that of Xpert was 53.6% (95% CI: 33.9 to 72.5). The specificity of Xpert Ultra based on the analysis of all available samples was 98.1% (95% CI: 93.4 to 99.7), whereas that of Xpert was 100%.

Conclusion In settings with a high burden of tuberculosis and HIV, Xpert Ultra had a better sensitivity than Xpert in children. However, the specificity was slightly lower than that of Xpert. Thus, Xpert Ultra has the potential to increase the reliability and the speed of tuberculosis diagnosis in children.

OC 8244 QUANTITATIVE LATERAL FLOW ASSAY FOR DETECTION OF *M. LEPRAE* INFECTION USING FINGERSTICK BLOOD

¹Annemiek Geluk*, ¹Anouk Van Hooij, ¹Elisa Tjon El Fat, ²Kosrhd Alam, ⁴Sipho Dlamini, ³Moises Batista Da Silva, ⁵John Spencer, ³Claudio Salgado, ⁶Jan Hendrik Richardus, ¹Paul Corstjens. ¹Leiden University Medical Center, The Netherlands; ²The Leprosy Mission Rural Health programme, Dhaka, Bangladesh; ³Universidade Federal do Pará, Brazil; ⁴Groote Schuur Hospital, Cape Town, South Africa; ⁵Colorado State University, USA; ⁶Erasmus MC, University Medical Center, Rotterdam, The Netherlands

10.1136/bmjgh-2019-EDC.4

Background Leprosy is a debilitating, infectious disease caused by *Mycobacterium leprae* leading to skin and nerve damage and often lifelong handicaps. The unabated rate of new leprosy case detection indicates that transmission of *M. leprae* is persistent and that current measures for prevention and multi-drug therapy (MDT) are insufficient. Contact with *M. leprae*-infected individuals is a risk factor for development of leprosy. Thus, detection of *M. leprae*-infected individuals without clinical symptoms, allowing informed decision making on who needs treatment at a preclinical stage, is vital to interrupt transmission and can help prevent leprosy.

Immunoprophylaxis by vaccination or post-exposure prophylaxis (PEP) with antibiotics provide effective strategies for the prevention of leprosy. To target individuals unknowingly spreading leprosy bacilli, methods allowing objective measurement of *M. leprae* infection are needed. Besides antibody (Ab) levels that correspond with bacterial load and higher risk of progression to leprosy, detection of cytokine profiles can provide significant added value to identify infection.

Methods Quantitative detection of anti-PGL-I IgM antibodies, and cytokines such as IP-10 was performed on lateral flow (LF) test strips utilising the luminescent up-converting particle (UCP) technology. Precise amounts of fingerstick (FS)-blood samples were collected using disposable heparinised capillaries.