

Methods Urine samples from men, 10 patients with urethritis and >5 PMNL/hpf and 10 healthy controls with <5 PMNL/hpf, were collected. All samples were tested for *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, *Mycoplasma genitalium*, *Ureaplasma urealyticum*, *U. parvum*, *Trichomonas vaginalis*, Herpes Simplex Virus type 1 and 2 and Adenovirus using specific PCR assays. The V3 and V4 regions of the 16S rRNA gene were PCR amplified, tagged and sequenced using the Titanium kit and GS FLX pyrosequencing system (Roche) according to manufacturer's instructions. Sequences were analysed using the RDP Pyrosequencing Pipeline and CLC Genomics Workbench.

Results From each of the 20 samples, 8150 quality filtered sequences were randomly selected. Sequences were assigned to the genus level using the RDP Classifier. A total of 172 gen-era were identified, 133 in patients and 104 in controls. The median number of genera was 35.5 (19–49) in patients with urethritis compared to 25 (16–43) in the controls. No single genus was present in all samples. Members of the genera *Pseudomonas* and *Sphingomonas* were present in all the controls and in most of the patient samples. Also, representatives of the genera *Brevundimonas*, *Micrococcus*, *Bradyrhizobium* and *Chryseobacterium* were present in 15–18 of the samples. Of the 172 genera, 77 were only found in a single sample and 33 were found in two different samples. Rarefaction analysis at the 5 % level (comparable to genus) suggested that all the controls and most urethritis samples harboured each between 31 and 125 different groups. Four of the urethritis samples apparently contained between 150 and 300 different groups.

Conclusions Urine, even from healthy men, contains a very diverse micro-flora. Though not statistically significant, the total and median number of genera was found higher in patients with urethritis than in controls. Several widespread genera are likely to represent commensals and bacteria present in the environment.

P4-S4.02 A 22-ORGANISM MICROARRAY APPROACH FOR DETECTING MICROBIOLOGICAL ASSOCIATIONS WITH SYMPTOMATIC URETHRITIS IN MALES

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Background In addition to its known microbiological aetiology, urethritis in men may be linked with other genital tract organisms, as yet unidentified in its pathogenesis. We used a microarray, with capacity to detect 22 genital tract organisms, in order to determine the association of symptomatic urethritis with infection or carriage of these organisms.

Methods 129 patients were asked to provide an extra first void urine specimen or give permission for their residual urine specimen submitted for *Chlamydia trachomatis* NAAT testing to be utilised. Patients were categorised into three self-reported symptom groups: definite symptoms of urethritis (discharge and/or dysuria), category 1 (C1) n=80; non-specific symptoms of urethritis (eg, minimal urethral discomfort), category 2 (C2) n=26; and asymptomatic category 3 (C3) n=23. Total urine nucleic acid was extracted and subsequently used for PCR coupled microarray analysis. Organisms were defined as present or absent using an online data analysis method. In a pre-planned analysis, the following categories were compared for prevalence of organisms: C1 vs C2 and C3 combined; C1 and C2 combined vs C3 using Fisher's exact test.

Results One or more organisms were detected in 74% (n=95) of patients and two or more organisms in 33% (n=42). The prevalence of organisms known to cause urethritis in this largely symptomatic cohort was: 16% (n=21), 9% (n=12) and 5% (n=6) for *C trachomatis*, *Mycoplasma genitalium* and for *Neisseria gonorrhoeae* respectively. *Escherichia coli* was the most prevalent organism detected with a prevalence of 18% (n=23). The presence of *M genitalium* was stat-

istically associated with C1 and C1 and C2 combined (p=0.03 and 0.01 respectively). In symptomatic patients, *C trachomatis*, *Ureaplasma urealyticum*, and *Gardnerella vaginalis* appeared to be more prevalent than in asymptomatics although not statistically significantly. Lactobacilli were detected in 1.3% and 4% of patients with C1 and C2 symptoms respectively, compared with 17% of asymptomatic patients. The absence of lactobacilli was associated with urethritis symptoms, either C1 alone or C1 and C2 combined (p=0.03 and p=0.01) respectively.

Conclusions Using a polymicrobial microarray approach we have demonstrated that symptomatic urethritis is associated with depletion of lactobacilli. This confirms early work using urethral swabs. The temporal nature of Lactobacilli depletion in relation to the onset of symptomatic urethritis needs to be investigated further.

P4-S4.03 A LOW-COST MICROFLUIDICS-BASED DIAGNOSTIC TEST FOR STDs

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Background Undiagnosed and untreated STIs cause large morbidity and mortality, including birth defects and stillborn babies. Since most STI's have known treatments, the largest barriers for treating patients include high cost of transporting specimens to central labs and lack of access to diagnostic testing. We present data on a portable and low-cost microfluidics device for point-of-care diagnosis of multiple STDs (such as HIV and syphilis) in combination. In order to reduce the cost and size of the assay while maintaining high performance, we incorporated microfluidic designs such as single-use plastic microfluidic cassettes, a passive method for delivering reagents, and an amplification chemistry using gold nanoparticles.

Methods Our overall device is named mChip (mobile microfluidic chip for immunoassay on protein markers). We demonstrated an ability of mChip to simultaneously detect antibodies against HIV and syphilis in needle-pricked sample volumes. Both commercial specimens and archived specimens from Sub-Saharan Africa were used.

Results The test sensitivities and specificities for detection of HIV-specific antibodies and treponemal-specific antibodies matched the performances from lab-based ELISA. Compared to ELISA, our test can be performed anywhere, uses a very small volume of blood, and is about 10 times faster. Some of these results are in press in *Nature Medicine*, 2010.

Conclusion The mChip provided excellent performance in the diagnosis of HIV using only 1 µl of unprocessed whole-blood and <15 min assay-time, and an ability to simultaneously diagnose HIV and syphilis with sensitivities and specificities equal to lab-based assays. Overall, we demonstrate an integrated strategy for miniaturising complex laboratory assays using microfluidics and nanoparticles to enable POC diagnostics and early detection of infectious diseases in remote settings.

P4-S4.04 TEMPORAL DYNAMICS OF VAGINAL BACTERIAL COMMUNITIES

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Background Dysbiosis of vaginal bacterial communities have been associated with increased risk for sexually transmitted infections and bacterial vaginosis. This is the first observational study to model temporal dynamics of vaginal microbiota using frequently collected samples, behavioural data and culture-independent methods.

LETTER

Unusual increase in reported HIV/AIDS cases among older persons in western Hunan province, China

An unusual increase in HIV/AIDS cases among older people was reported to the Hunan Centers for Disease Control between 2005 and 2007. Cases originated in four rural, western districts of this inland province of China. Given the historical concern for outbreaks of HIV in rural areas due to blood donation,¹ these cases prompted closer examination to understand the reasons for their appearance and to take measures to prevent further spread.

Eighty cases met our investigation criterion of 50 years or older and underwent a structured interview. The median age was 65 years (range 51–82); 42% were female. Most were ethnic minorities, 76% Tujia and 9% Miao, with low education. Nearly all had been married; 43% were widowed. Most (54%) spent time away from their spouse (median >5 years); 10% were currently sexually active with a spouse; few ever used condoms with their spouse.

Investigation of the possible modes of HIV acquisition suggests most infections among men were from female sex workers (83% paid for sex, two-thirds in the last 5 years), and among women through infected husbands. One case had a history of selling blood, most recently in 1981. Eight received a blood transfusion, three before 1976 and five after 1984. All denied drug use; all men denied male–male sex; all women denied extramarital sex. Among men buying sex, 97% never used condoms.

Two-thirds had never heard of HIV prior to their diagnosis. Few (14%) knew HIV could be transmitted sexually, through blood (11%) or from mother to child (4%). Of the men reporting commercial sex contact, 82% had no knowledge that condoms could prevent HIV. Most cases (86%) were detected incidentally during the course of treatment for other diseases or because their spouse was HIV-positive. By interview, 78% indicated their spouse had tested for HIV, of whom 69% were reported to be positive.

Our investigation highlights that basic information on HIV/AIDS is not reaching all parts of China, and may especially lag among rural and older people. Discussion of sex with older people has been taboo in China, presenting special challenges in finding effective ways to reach them. As treatment extends survival, the cohort of persons living with HIV will also age. The movement of people between urban and rural areas, an ageing population and

the shift of the HIV/AIDS epidemic to sexual transmission² are three trends in China that may now have a dangerous intersection.

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Competing interests None.

Patient consent Obtained.

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Contributors XC oversaw and coordinated the fieldwork. JZ, JMH and BYQ conducted the fieldwork. YH completed the survey design, data analysis and drafting of the manuscript. LW and NW provided technical support during and prior to the survey and mobilised some funds to complete the survey.

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REFERENCES

1. Wu Z, Liu Z, Detels R. HIV-1 infection in commercial plasma donors in China. *Lancet* 1995;**346**:61–2.
2. China Ministry of Health, UNAIDS, and World Health Organization (WHO). 2009 Estimates for the HIV/AIDS Epidemic in China. Beijing, China: China Ministry of Health, 2010. <http://www.unaids.org.cn/download/2009%20China%20Estimation%20Report-En.pdf> (accessed 1 May 2011).

CORRECTIONS

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RamaKrishnan A. Symposium 9: Applications of program science in the field of STI: S9.3 The programme science of scale: the Avahan experience. *Sex Transm Infect* 2011;**87**:A10. doi:10.1136/sextrans-2011-050102.38.

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Starnino S, Liao M, Ruben M, Storey A, Dillon JAR, GASP-LAC Network. P1-S1.45 Neisseria Gonorrhoeae Antimicrobial Susceptibility in Latin America and the Caribbean (2000–2009) - A Contribution to the Treatment Guidelines Revision. *Sex Transm Infect* 2011;**87**:A117. doi:10.1136/sextrans-2011-050108.45.

Starnino S, Liao M, Ruben M, Storey A, Dillon JAR, GASP-LAC Network. P1-S4.28 Survey Of Methodology Used For The Identification And Antimicrobial Susceptibility Testing Of Neisseria Gonorrhoeae In Latin America And The Caribbean. *Sex Transm Infect* 2011;**87**:A172. doi:10.1136/sextrans-2011-050108.172.

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