reviewed. Site-specific gonorrhoea positivity was also calculated.

**Results** 191 heterosexual contacts (102 males and 89 females) were reviewed. The median age was 28 [IQR=24–33] years. The gonorrhoea positivity in male was significantly higher at the oropharynx compared to urethra (17.6% [95% CI: 10.8–26.4%] versus 2.0% [0.2–6.9%]; p<0.001); and higher at the oropharynx in female compared to cervicovaginal site (46.1% [35.4–57.0%] versus 36.0% [26.1–46.8%]; p=0.056). Of the 100 males who did not have genital gonorrhoea, 17 (17.0% [10.2–25.8%]) tested positive at the oropharynx. Of the 55 females who did not have genital gonorrhoea, 21 (23.6% [15.2–33.8%]) tested positive at the oropharynx. Infection at both the oropharynx and genital sites was not associated with sex worker status in females. Overall, 89.5% and 39.6% of gonorrhoea in males and females were detected only in the oropharynx, respectively.

**Conclusion** Multiple sites of gonococcal infection are more common in female contacts than in male contacts. Approximately 90% and 40% oropharyngeal infections would have been missed in males and females, respectively, by genital-only screening among heterosexuals reporting contact with sexual partners with gonorrhoea. Oropharyngeal gonorrhoea screening among heterosexual contacts of gonorrhoea is important to prevent ongoing transmission.

**Disclosure** No significant relationships.

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**002.3 ANTIMICROBIAL RESISTANCE IN PHARYNGEAL NEISSERIA GONORRHOEAE INFECTION: A CROSS-SECTIONAL STUDY IN MEN WHO HAVE SEX WITH MEN**

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**Background** Pharyngeal Neisseria gonorrhoeae (NG) infections are common among men who report sex with men (MSM). The pharynx is an important anatomic reservoir for the development of antimicrobial resistance due to the potential for exchange of genetic material between NG and other commensals of this niche, including other Neisseria species. We investigated whether there was an association between pharyngeal infection and reduced susceptibility (RS) or resistance to antimicrobials used to treat NG compared to genital and rectal infection.

**Methods** Logistic regression of odds ratios (OR) was used to model the association between the anatomical site of infection and RS or resistance to azithromycin, ceftriaxone, cefixime and ciprofloxacin using isolates from MSM in England and Wales collected within the Gonococcal Resistance to Antimicrobials Surveillance Programme (GRASP), 2012–2017.

**Results** Among 5,448 isolates from MSM, 729 (13.4%) were pharyngeal, 2,365 (43.4%) were rectal and 2,354 (43.2%) were genital samples. Pharyngeal infections were more likely to be associated with azithromycin resistance (minimum inhibitory concentration (MIC) >0.5 mg/L (adjusted OR (aOR) 1.62, 95%CI:1.13–2.31, P<0.001) and RS to ceftriaxone (MIC ≥0.015 mg/L) (aOR 1.25, 95%CI:1.03–1.52, P=0.023) compared to genital infections. Pharyngeal infections were also more likely to be associated with azithromycin resistance (aOR 1.49, 95%CI:1.06–2.11, P<0.001) and RS to ceftriaxone (aOR 1.21, 95%CI:1.00–1.47, P=0.045) compared to rectal infections. No significant association was found between site of infection and cefixime or ciprofloxacin resistance.

**Conclusion** Pharyngeal NG infection among MSM are more likely to be RS to ceftriaxone and resistant to azithromycin compared to rectal and genital infections. Poor pharyngeal tissue drug penetration may lead to persistent infections, which would provide more time for exchange of genetic material that confer AMR. This highlights the importance of extra-genital testing and antimicrobial susceptibility testing in this population, to reduce the risk of treatment failure and onward transmission of resistant strains.

**Disclosure** No significant relationships.

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**002.4 INCIDENCE AND DURATION OF PHARYNGEAL AND RECTAL GONORRHEA AND CHLAMYDIA AMONG HIGH- RISK MEN WHO HAVE SEX WITH MEN (MSM)**

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**Background** The duration of untreated extragenital gonococcal (GC) and chlamydial infection (CT) infection is not well defined.

**Methods** From March 2016 to December 2018, we enrolled 140 MSM in a 12-month cohort study. Men ≥18 years were eligible if they reported receptive anal intercourse and had ≥1 following risks in ≤12 months: 1) diagnosis of GC, CT or syphilis; 2) methamphetamine or poppers use; or 3) ≥2 sex partners in ≤2 months or >5 in ≤12 months. Enrolled men either tested negative for GC/CT at enrollment, or, tested positive, were treated and waited 2–3 weeks prior to data collection. Each week, men self-collected pharyngeal and rectal specimens and completed an electronic diary. Specimens were tested after study completion (Aptima, Hologic Inc). We defined incident infection as two consecutively positive tests and clearance as ≥2 consecutively negative tests. We used Kaplan Meier curves to estimate duration of infection censoring subjects for receipt of pathogen-specific antibiotic, positive swab in final week of study, or loss-to-follow-up.

**Results** Forty-eight men were observed for 1,687 weeks and contributed 3,579 tested specimens. Twenty-four (50%) MSM had ≥1 incident GC/CT infection; 13 (27%) had >1 infection. Overall extragenital GC/CT incidence was 129 (95%CI: 94–172) infections per 100 person-years. Pharyngeal GC, and rectal GC and CT incidence were 35 (95%CI: 20–61), 37 (95%CI: 22–64) and 59 (95%CI: 38–91) per 100 person-years, respectively. 46% (6/13) pharyngeal GC, 43% (6/14) rectal GC, 81% (17/21) rectal CT were censored. The estimated median duration of pharyngeal GC, rectal GC and rectal CT were 15 (95%CI 3 – undefined), 12 (95%CI 2 – undefined) and >20 (95% CI 12 – undefined) weeks.

**Conclusion** Among high-risk MSM, incident extragenital GC/CT occur frequently: >1 infection per person per year. Untreated, these infections persist for a median of 3 to 5 months.

**Disclosure** No significant relationships.