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Background

Syphilis testing and treatment of new infections, if detected early, is simple and inexpensive. We previously carried out the Enhanced Syphilis Screening Among HIV-positive Men (ESSAHM) Trial that paired opt-out syphilis tests with routine HIV bloodwork to improve early syphilis case detection (Clin Infect Dis 2022; 74(5):846-853). One of the trial objectives was to determine if routinized syphilis testing reached men at higher risk of infection (Implementation Sci 2015; 11, 8). However, this could not be tested using trial data alone as few patient covariates were collected given the pragmatic trial approach. Thus, we re-analysed ESSAHM Trial intervention effects using data from a concurrent cohort study that ascertained sociodemographic characteristics and sexual histories.

Methods

The ESSAHM Trial used a stepped wedge cluster-randomized trial (SW-CRT) design. Four outpatient HIV clinics introduced a practice change to pair syphilis testing with routine HIV viral loads in a step-wise fashion over 5 six-month periods. All clinics were in the control condition in step 1. Clinics were then randomized to one of four roll-out schedules such that, by step 5, all were in the intervention condition.

Population: HIV-positive men attending HIV outpatient clinics in Toronto and Ottawa, Canada

Intervention: standing orders for syphilis testing with HIV viral loads

Control: usual syphilis testing practice

Outcomes: early syphilis case detection, proportion screened ("coverage"), and screening frequency

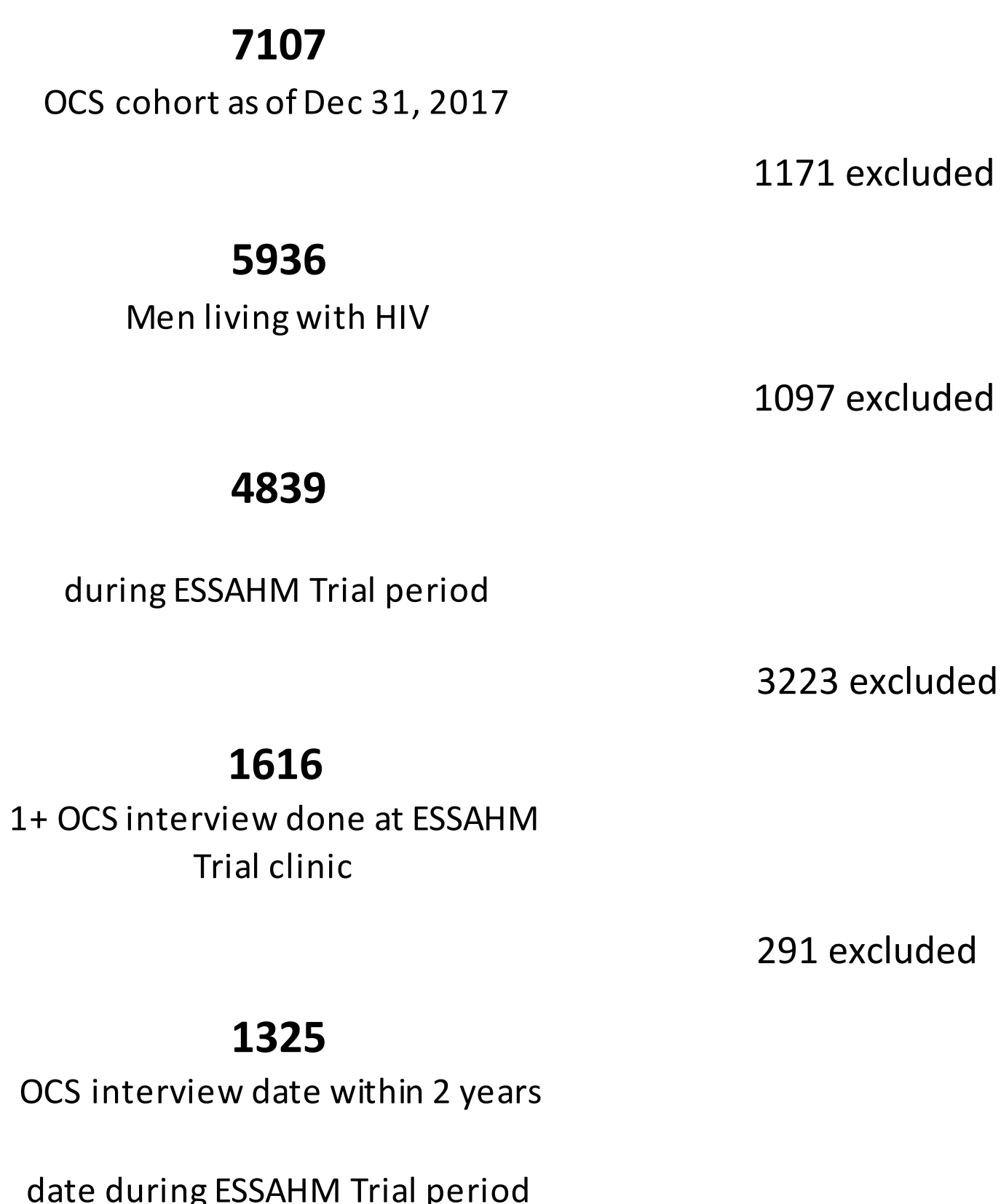
Time: 01/02/2015 to 31/07/2017

The 4 ESSAHM Trial clinics were among the 10 sites of the Ontario HIV Treatment Network Cohort Study (OCS). At annual interviews, OCS participants report income, education, sexual orientation, and number of sexual partners in the past 3 months. We applied analysis inclusion criteria for OCS participants (Figure 1):

- 1) a man (cis- or transgender) living with HIV
2) ≥1 viral load test during trial period
3) ≥1 OCS questionnaire completed within 2 years of the trial period at a participating ESSAHM clinic

We quantified intervention effects among OCS participants using time-adjusted generalized linear mixed-effect models to estimate odds ratios (OR) and rate-ratios (RR) with 95% confidence intervals (CI) and explored evidence for confounding and effect modification by sociodemographic and sexual history covariates.

Figure 1. Flow diagram for study inclusion



Results

Thirty-four percent (1325/3895) of ESSAHM trial participants were also OCS participants who met the analysis inclusion criteria. They had a mean age of 52.0 years and 86% were men who have sex with men (Table 1). At baseline, 25% (n=316) reported 2+, 30% (n=365) reported 1, and 45% (n=556) reported 0 sex partners.

Table 1. Characteristics of OCS participants who were also participants ESSAHM Trial at step 1 when all clinics were in control condition (a)

Table with 2 columns: Characteristic and n (%). Rows include: Number of viral loads per person in 6-month period (Mean (SD) 1.41 (0.74)), Result of (first) HIV viral load (Undetectable (<40 copies/mL) 868 (87.5%), Known past history of syphilis at baseline (b) 148 (14.9%), Age (Mean (SD) 52.0 (11.5)), Personal Income (CAD) (Less than \$20,000 343 (34.6%)), Education (Less than high school 106 (10.7%)), Race (Indigenous 21 (2.1%)), Sexual Orientation (Heterosexual/Straight 210 (21.2%)), Men who have sex with men (MSM) 848 (85.5%), Number of sex partners, past 3 months (None 422 (42.5%)), Number of male sex partners, past 3 months (None 511 (51.5%)), Number of female sex partners, past 3 months (None 836 (84.3%)), Multiple partners living with HIV, past 3 months (None or one partner of any status 680 (68.5%)), Multiple HIV-negative/unknown status partners, past 3 months (None or one partner of any status 680 (68.5%)).

Frequency (percent) unless specified otherwise. CAD = Canadian currency. All proportions may not add to 100% due to missing data (not greater than 2.4%).

(a) Baseline characteristics are shown for male patients under study during the first six-month step of the SW-CRT, when all clinics were in the "control" setting. The 333 men who entered the study in subsequent steps are not shown.

(b) Past history of syphilis as determined by reactive syphilis serology in the year preceding the trial start.

Conclusions

Among all men under follow-up, routine syphilis screening produced an 83% increase in early case detection that was non-statistically significant. But among men with two or more sex partners (who would have been more likely to have had sexual exposure to syphilis), early case detection tripled. Although the magnitude of increases in screening coverage and frequency were less among men with multiple partners compared to usual care. Altogether, these findings support the implementation of routine syphilis screening for men living with HIV.

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There were 75 diagnoses of early syphilis (control: 26, intervention: 49). Comparing intervention to control periods, there were increases in case detection (OR 1.83, CI 0.85, 3.93), the proportion screened (OR 2.81, CI 2.20, 3.59), and number of tests per year (RR 1.73, CI 1.51, 1.97) (Table 2). There was no evidence of confounding by sociodemographic characteristics. However, intervention effects were modified by number of sex partners (Table 3).

Table 2. Intervention effects on early syphilis case detection, screening coverage, and screening frequency

Table with 3 columns: Outcome, Unadjusted Ratio (95%CI), Time-adjusted Ratio (95%CI). Rows include: Early syphilis detection (a) 2.44 (1.45, 4.11), Screening coverage (a) 6.47 (5.50, 7.60), Screening frequency (c) 2.25 (2.08, 2.43).

CI, confidence interval. Referent = control periods. (a) Either (1) seroconversion with prior negative syphilis serology within the previous 12 months; or (2) in men previously diagnosed with syphilis, a 4-fold or greater rise in RPR titre from the last titre within the prior 12 months, indicative of re-infection Odds ratio estimated by general linear mixed model using a logit link. (b) Proportion of men with at least one syphilis test during a six-month step. Rate ratio estimated by general linear mixed model using a log link. (c) Count of syphilis serology test episodes per six-month step. Rate ratio estimated by general linear mixed model using a log link.

Table 3. Multiplicative interactions between sexual behaviours and intervention effects on early syphilis case detection, screening coverage, and screening frequency

Table with 4 columns: Sexual partnership history, Early syphilis detection: Time-adjusted (95%CI), Screening coverage: Time-adjusted (95%CI), Screening frequency: Time-adjusted (95%CI). Rows include: Number of male and female sex partners (≤1 1.46 (0.53, 4.00), 2+ 3.05 (1.14, 8.12)), Multiple partners living with HIV (≤1 of any HIV status 1.42 (0.52, 3.91), 2+ but not living with HIV 1.57 (0.44, 5.57)), Multiple HIV-negative / unknown status partners (≤1 of any HIV status 1.51 (0.54, 4.20), 2+ but not HIV-negative/unknown status 5.13 (1.44, 18.33)).

CI, confidence interval. Reference = control periods. (a) P-value contrast is between 2+ living with HIV (or HIV-negative/unknown status) versus ≤1.