

Methods for combining data from multi-jurisdiction sentinel surveillance projects

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Objective

To identify best practices for combining public health data for multi-jurisdiction surveillance projects.

Introduction

Sentinel surveillance, where selected jurisdictions follow standardized protocols to collect and report enhanced public health data not available through other routine surveillance efforts, is a key part of national surveillance of sexually transmitted diseases (STDs). Although four STDs are nationally notifiable conditions (chlamydia, gonorrhea, syphilis and chancroid), the burden of these conditions (over 2.3 million cases were reported in 2017) limits the amount of detailed clinical and demographic data available for all cases. Sentinel surveillance in clinical settings serving at-risk populations, such as STD clinics, provides an opportunity to collect enhanced data elements on persons seeking STD-related services, such as sex of sex partners and anatomic site of infection.

However, there are challenges in combining data across jurisdictions as estimated effect measures may vary by jurisdiction (e.g., some may have higher observed burden of disease among certain populations) and the amount of data contributed by jurisdiction may vary; combined this could lead to biased estimates if heterogeneity is not taken into account.

Methods

Using data from the STD Surveillance Network (SSuN), a sentinel surveillance project implemented in 10 jurisdictions, we investigated the effect of using different statistical methods to combine data across jurisdictions. We evaluated 5 methodologies:

- "Fully stratified" where estimates were provided separately for each jurisdiction;
- "Aggregated" where numerators and denominators were summed across jurisdictions without weighting;
- "Mean estimate" where the mean of the jurisdiction-specific estimates was estimated;
- "Random effects" where jurisdiction-specific estimates were combined using an inverse variance weighted random effects model to adjust for heterogeneity between jurisdictions; and
- "Stratified random effects" where a possible effect modifier was identified and used to group jurisdictions prior to calculating the estimate from the random effects model.

Through SSuN, jurisdictions collect visit-level data on patients attending selected STD clinics and report clinical and demographic data. As an illustrative example, we estimated rectal gonorrhea positivity among gay, bisexual, and other men who have sex with men (MSM) attending participating clinics. Jurisdiction-specific positivity was estimated as the # of unique MSM testing positive at least once for rectal gonorrhea divided by all MSM tested 1 or more times for rectal gonorrhea in all of the clinics in the jurisdiction. The stratifying variable for the stratified random effects method was the percent of MSM screened in the jurisdiction's clinics, as low screening coverage may reflect targeted testing of MSM likely to be infected which may inflate observed positivity. For each of the five methods, we estimated rectal gonorrhea positivity and the corresponding 95% confidence interval (CI).

Results

In 2017, 123,210 patients attended 30 STD clinics participating in the 10 SSuN jurisdictions, of which 31,052 (25.2%) were identified as MSM (jurisdiction-specific range: 8.8% to 70.0%). (Table 1) One jurisdiction (I) accounted for 39% of all MSM



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included in the analysis while one jurisdiction (J) accounted for only 1.6% of MSM included. The proportion of MSM tested for rectal gonorrhea at least once varied by jurisdiction, ranging from 44.3% to 76.9%. The fully stratified method identified differences in rectal gonorrhea positivity across jurisdictions, with jurisdiction-specific positivity ranging from 9.9% to 24.1%. Aggregating across jurisdictions masked this heterogeneity and provided a single summary estimate of 15.2% (95% CI: 14.7, 15.7). Taking the mean across the jurisdiction-specific estimates also provided a summary estimate; however, the uncertainty of the estimate increased (15.8%, 95% CI: 13.3, 18.7). Accounting for the heterogeneity by using a random effects model resulted in an estimate of 15.5% (95% CI: 13.9, 17.2). After stratifying by a likely confounder (% of MSM screened); the random effects estimate among 3 jurisdictions with lower screening coverage (<60%) was 19.7% (95% CI: 14.6, 24.8) and among 7 jurisdictions with higher screening coverage (\geq 60%) was 14.3% (95% CI: 12.9, 15.7).

Conclusions

In a sentinel surveillance project implemented in 10 jurisdictions, there was substantial heterogeneity in the observed proportion of MSM testing positive for rectal gonorrhea in selected STD clinics. Although a stratified analysis captured the heterogeneity across jurisdictions, it may not be feasible to present fully stratified estimates for all analyses (e.g., surveillance reports likely provide metrics for multiple diseases). Additionally, it limits the ability to succinctly communicate key findings. Aggregating numerators and denominators across jurisdictions to calculate a single summary estimate masks this heterogeneity and biases estimates toward high volume jurisdictions. Taking the mean across jurisdictions ensures that high-volume jurisdictions. Using a random effects model accounted for both varying sample sizes and differences in observed heterogeneity; although the summary estimate was similar to the aggregate in this example, the wider 95% CI more accurately reflects the uncertainty in the estimate. Finally, stratifying by a likely effect measure modifier (% of MSM screened) prior to estimating the measure from the random effects model captured key differences in jurisdictions and it is not feasible to provide fully stratified estimates, analysts could consider using methods to account for heterogeneity and minimize bias due to differing sample sizes, such as stratified random effects models.

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| Jurisdictio | # of | % of | % of MSM | % MSM testing positive for rectal gonorrhea | | | | | |
|-------------|-----------|----------------------------------|--|---|--|--|----------------------------|---------------------------------------|--|
| n | s patient | patients identified as MSM | patients screened for rectal gonorrhea | Fully stratifie d | Aggregate d across jurisdiction s | Mean of jurisdictio n estimates | Random effects model | Stratified random effects model | |
| А | 6,262 | 14.3% | 44.3% | 14.4% | n/a | n/a | n/a | 19.7% | |
| В | 8,160 | 8.8% | 46.4% | 24.1% | n/a | n/a | n/a | (95% CI: | |
| С | 13,519 | 15.7% | 58.5% | 20.9% | n/a | n/a | n/a | 14.6, 24.8) | |
| D | 6,017 | 44.6% | 65.6% | 18.2% | n/a | n/a | n/a | 14.3% | |
| E | 15,083 | 16.8% | 65.8% | 14.7% | n/a | n/a | n/a | (95% CI: | |
| F | 9,188 | 48.4% | 67.8% | 15.8% | n/a | n/a | n/a | | |

Table 1. Trends in rectal gonorrhea among MSM tested in STD clinics participating in the STD Surveillance Network (SSuN),2017



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| G | 10,657 | 30.1% | 68.1% | 13.5% | n/a | n/a | n/a | 12.9, |
|-------|--------|-------|-------|-------|----------|----------|-------|-------|
| Н | 3,665 | 46.2% | 70.5% | 12.3% | n/a | n/a | n/a | 15.7) |
| Ι | 49,792 | 24.4% | 71.7% | 14.3% | n/a | n/a | n/a | |
| J | 867 | 70.0% | 76.9% | 9.9% | n/a | n/a | n/a | |
| TOTAL | 123,21 | 25.2% | 67.5% | n/a | 15.2% | 15.8% | 15.5% | n/a |
| | 0 | | | | (95% CI: | (95% CI: | (95%) | |
| | | | | | 14.7, | 13.3, | CI: | |
| | | | | | 15.7) | 18.7) | 13.9, | |
| | | | | | | | 17.2) | |
| | | | | | | | 17.2) | |

MSM: Gay, bisexual and other men who have sex with men; CI: Confidence interval



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