

# **Completeness of Active and Passive** Surveillance in San Francisco



## S. Nabity<sup>1</sup>, S. Shin<sup>2</sup>, P. Shiono<sup>1</sup>, D. Vugia<sup>3</sup>, S. Huang<sup>1</sup>

<sup>1</sup> San Francisco Department of Public Health, San Francisco, CA, <sup>2</sup> California Emerging Infections Program, Oakland, CA

<sup>3</sup> California Department of Health Services, Richmond, CA.

#### Introduction

#003

Clinicians, laboratorians, and other mandated reporters are required to notify California local health jurisdictions (LHJ) upon the diagnosis of selected infections. Completeness of these passive notifications has important implications for the ability of LHJs to monitor community health trends and implement effective interventions. Evaluation of core surveillance systems is further considered a fundamental aspect of LHJ preparedness for bioterrorism and infectious disease emergencies.

Valid estimates for completeness of communicable disease notifications to LHJs can be challenging and resourceintensive in the absence of comprehensive integrated lab information systems. The California Emerging Infections Program (CEIP) supplements San Francisco Department of Public Health (SFDPH) passive surveillance through active case finding at 12 clinical labs serving San Francisco clinicians. CEIP active surveillance represents the most complete, readily-available source of data on selected labconfirmed infections for comparison with SFDPH records.

#### Aims

· To quantify the completeness of active and passive notifications to SFDPH for 4 communicable diseases.

· To identify and characterize gaps in notifications that inform potential interventions.

•To evaluate the appropriateness of capture-recapture methods for quantifying the fraction of lab-confirmed cases not captured by either surveillance system.

#### Methods

Cases were SF residents identified separately through passive and active surveillance for specimens collected during the 6 months from January 1 to June 30, 2005. During this period, data from CEIP was not included in the SEDPH database

We evaluated two pathogens reportable by clinicians (Cryptosporidium and Shigella spp.), and two others reportable by both labs and clinicians (Salmonella and Campylobacter spp.).

CEIP active surveillance was conducted by surveillance officers who contacted laboratorians in 12 clinical labs serving SE clinicians at least monthly and/or received computerized printouts to ensure that all relevant cases were ascertained.

SFDPH passively received notifications from clinicians and labs per California Code of Regulations Title 17 for all SF residents from multiple sources including labs, clinicians, and other LHJs,

If an individuals had multiple positive lab results for the same pathogen and the tests were done >30 days apart, then we defined them as separate cases

We merged the SFDPH and CEIP databases using personal identifiers and manually reviewed the matches for accuracy

We evaluated our data sources for adherence to 4 established assumptions of the capture-recapture method: 1) no change to the population during investigation

- 2) individuals can be accurately matched between data sources
- 3) individuals have an equal probability of being in each data source 4) the data sources are independent.

#### Results



Figure 1. Distribution of unique cases (N=299) by pathogen after merging active and passive data sources, SF, Jan - Jun 2006.



Table 2. Completeness of active and passive notifications for all pathogens combined.

85% (95%Cl 80%-88%) of lab-confirmed cases were identified through passive surveillance. Passive notifications for Cryptosporidium, Salmonella, and Shigella were significantly more complete (each ≥ 95%) than Campylobacter (79%, 95%CI 73%-85%).

94% (95%Cl 91%-96%) of lab-confirmed cases were identified through active surveillance. Active notifications did not differ significantly by disease.

· Most cases missed by active and passive surveillance were Campylobacter: 12 (63%) of 19 cases missed by active and 39 (87%) of 45 cases missed by passive surveillance were Campylobacter.



Figure 2. Completeness (with 95% CIs) of active notifications by pathogen, SF, Jan - Jun 2006.



Figure 3. Completeness (with 95% CIs) of passive notifications by pathogen, SF, Jan - Jun 2006.

Capture-recapture methods could not be applied to our data because (1) SFDPH receives notifications from laboratories outside the CEIP network or from clinicians using their diagnostic services so each individual does not have an equal chance of being included in both data bases, and (2) the data sources are not independent because laboratories are obligated to report results to both SFDPH and CEIP. Further, laboratorians are reminded to report to SFDPH as a direct result of collaboration with CĖIP.

### Conclusion

 The fraction of lab-confirmed cases missed by both passive and active surveillance systems in SF labs appears small.

· Completeness of passive case finding was lower for campylobacteriosis than for the other 3 diseases. As SFDPH does not routinely investigate individual cases of campylobacteriosis, these reporting failures did not greatly impact public health response; rather, it affected the valid enumeration of burden of disease trends.

 SEDPH is working with labs to evaluate their information systems and reporting procedures to identify causes for gaps and to prepare for electronic lab reporting (ELR).

 Capture-recapture calculations are not valid when comparing surveillance ascertainment methods that draw from the same source, as demonstrated by clinical laboratories in this analysis. Comparing the cases obtained through both methods is sufficient.

#### More Information

Scott Nabity, Communicable Disease Control Unit San Francisco Department of Public Health Ph: (415) 554-2830 Email: Scott.Nabity@sfdph.org

#### Acknowledgements

We acknowledge the ongoing data collection efforts of SFDPH and CEIP surveillance staff. Reporting clinicians and participating laboratories made these data available for analvsis.