

# Detecting Emerging COVID-19 Community Outbreaks

CSTE Spatial Analysis Workgroup  
July 22, 2020

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# Objectives

- General: to prioritize case investigations and contact tracing efforts, to allocate testing and community engagement resources, and to protect the elderly and other vulnerable populations
- Specific: To determine where SARS-CoV-2 percent positivity is increasing faster than jurisdiction-wide trend, or decreasing slower than jurisdiction-wide trend
- Why SaTScan?
  - Scan statistics can detect increased disease activity anywhere at any time, without a priori specification of temporal period or geographic location or size

Do we want  
to detect  
increases in  
case counts  
or in percent  
testing  
positive?

- Why might COVID-19 case counts increase?
  - Ongoing disease transmission requiring public health investigation
  - Proportionally more patients with SARS-CoV-2 are tested
- Why might the percent positive of SARS-CoV-2 tests increase?
  - Ongoing disease transmission requiring public health investigation
  - Proportionally fewer patients without SARS-CoV-2 are tested
- True disease transmission most apparent when both case counts and percent positive tests increase

## Research

## A Large Community Outbreak of Legionnaires' Disease Associated With a Cooling Tower in New York City, 2015

[On This Page](#)

## Practice Full Report

SDC

## Detecting Drop-offs in Electronic Laboratory Reporting for Communicable Diseases in New York City

Sharon K. Greene, PhD, MPH; Erin M. Andrews, DrPH; Pamela Evans Lloyd, MPA;  
Jennifer Baumgartner, MSPH; Eric R. Peterson, MPH

# EMERGING INFECTIOUS DISEASES®

EID Journal > Volume 22 > Number 10—October 2016 > Main Article

Volume 22, Number 10—October 2016

Dispatch

Daily Reportable Disease Spatiotemporal Cluster Detection, New York City, New York, USA, 2014–2015

Sharon K. Greene✉, Eric R. Peterson, Deborah Kapell, Annie D. Fine, and Martin Kulldorff

Morbidity and Mortality Weekly Report (MMWR)

CDC

## Salmonellosis Outbreak Detected by Automated Spatiotemporal Analysis — New York City, May–June 2019

Weekly / July 3, 2020 / 69(26);815–819

Julia Latash, MPH<sup>1</sup>\*; Sharon K. Greene, PhD<sup>1</sup>\*; Faina Stavinsky, MS<sup>2</sup>; Sandy Li<sup>3</sup>; Jennifer A. McConnell, MS<sup>3</sup>; John Novak, PhD<sup>3</sup>; Teresa Rozza<sup>3</sup>; Jing Wu, PhD<sup>3</sup>; Enoma Omoregie, PhD<sup>3</sup>; Lan Li, MPH<sup>1</sup>; Eric R. Peterson, MPH<sup>1</sup>; Bruce Gutelius, MD<sup>1</sup>; Vasudha Reddy, MPH<sup>1</sup> ([View author](#))

# Why different from existing aberration detection processes?

- Different goals:
  - Detect new outbreak in context of minimal or stable disease incidence
- vs.
- Monitor ongoing outbreak
- Case count-only analyses are limited by temporal and spatial shifts in SARS-CoV-2 testing availability and uptake

# Strategy

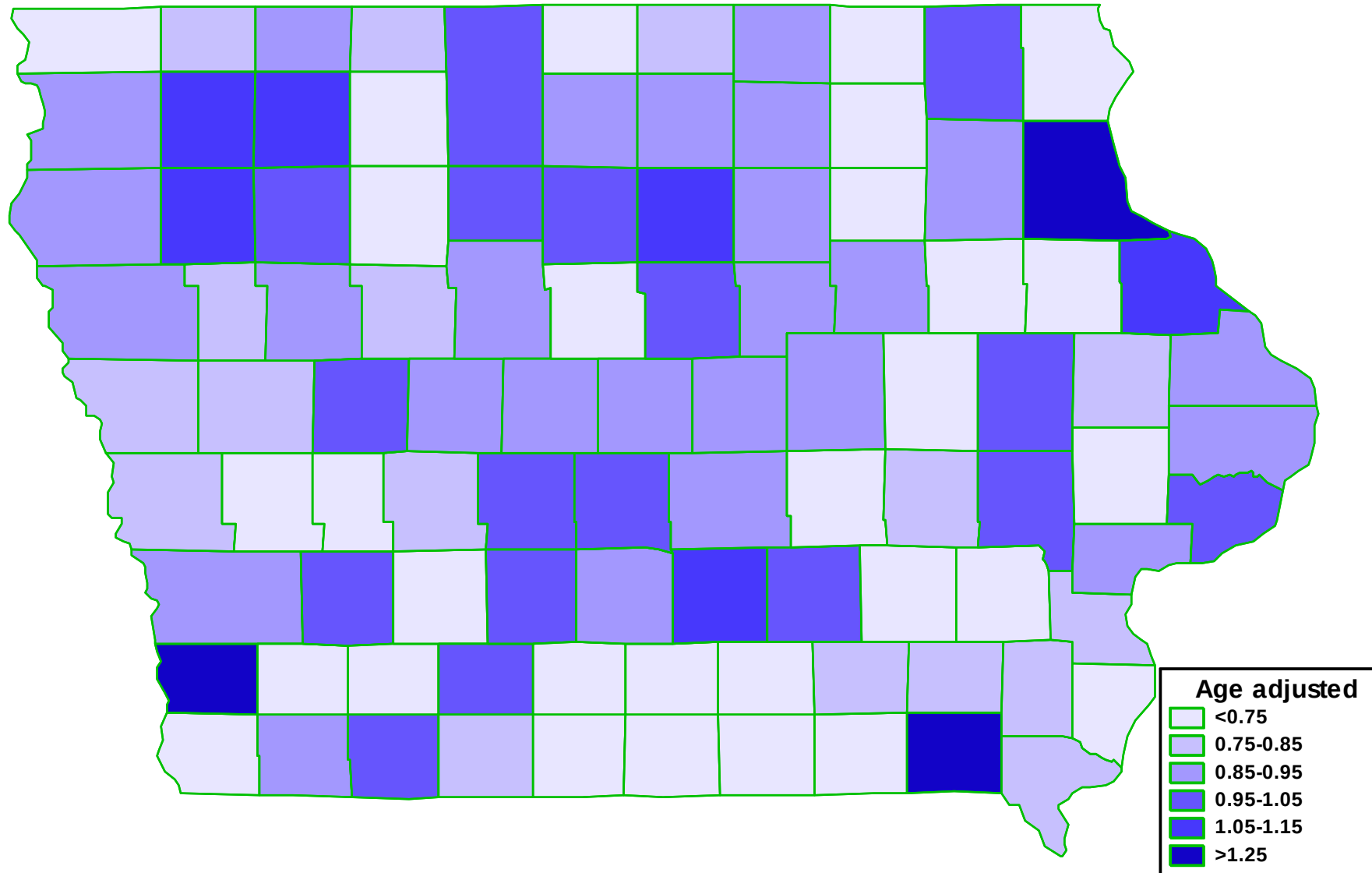
- Automate daily analyses of geocoded lab data to identify community clusters of increased test positivity
- Prospective discrete Poisson-based space-time scan statistic
  - “Population” = unique persons receiving PCR testing
- Adjust for purely spatial variation nonparametrically using spatial stratified randomization
- Adjust for purely temporal trends using log-linear trend adjustment
- Details on [medRxiv](#)

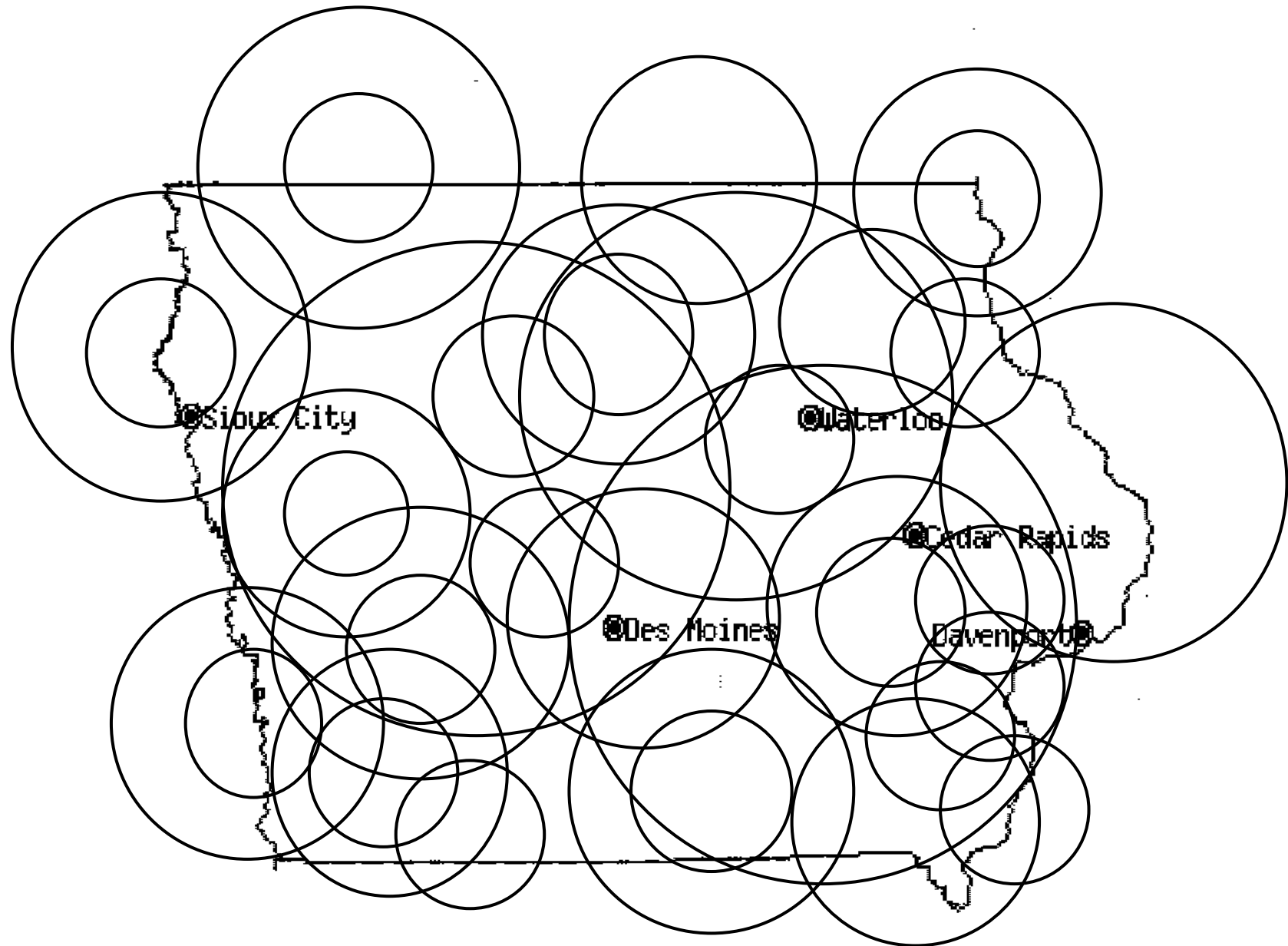
# Statistical Analysis

- Spatial Scan Statistic
- Space-Time Scan Statistic
- Different Probability Models
- Adjusting for Covariates, Geographical Variation and Temporal Trends
- Free SaTScan software

# Iowa Breast Cancer Incidence

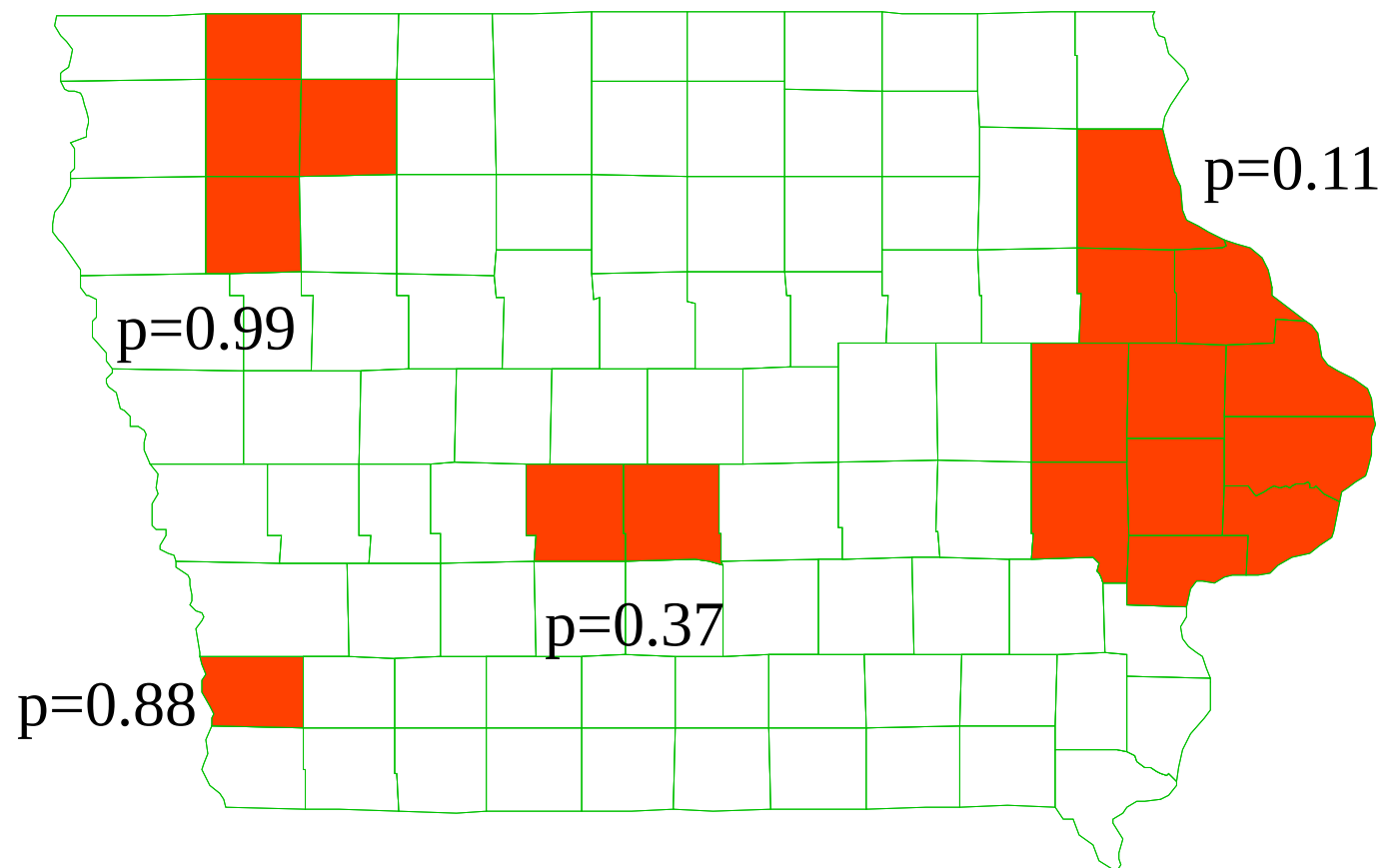
## Age-Adjusted Relative Risks

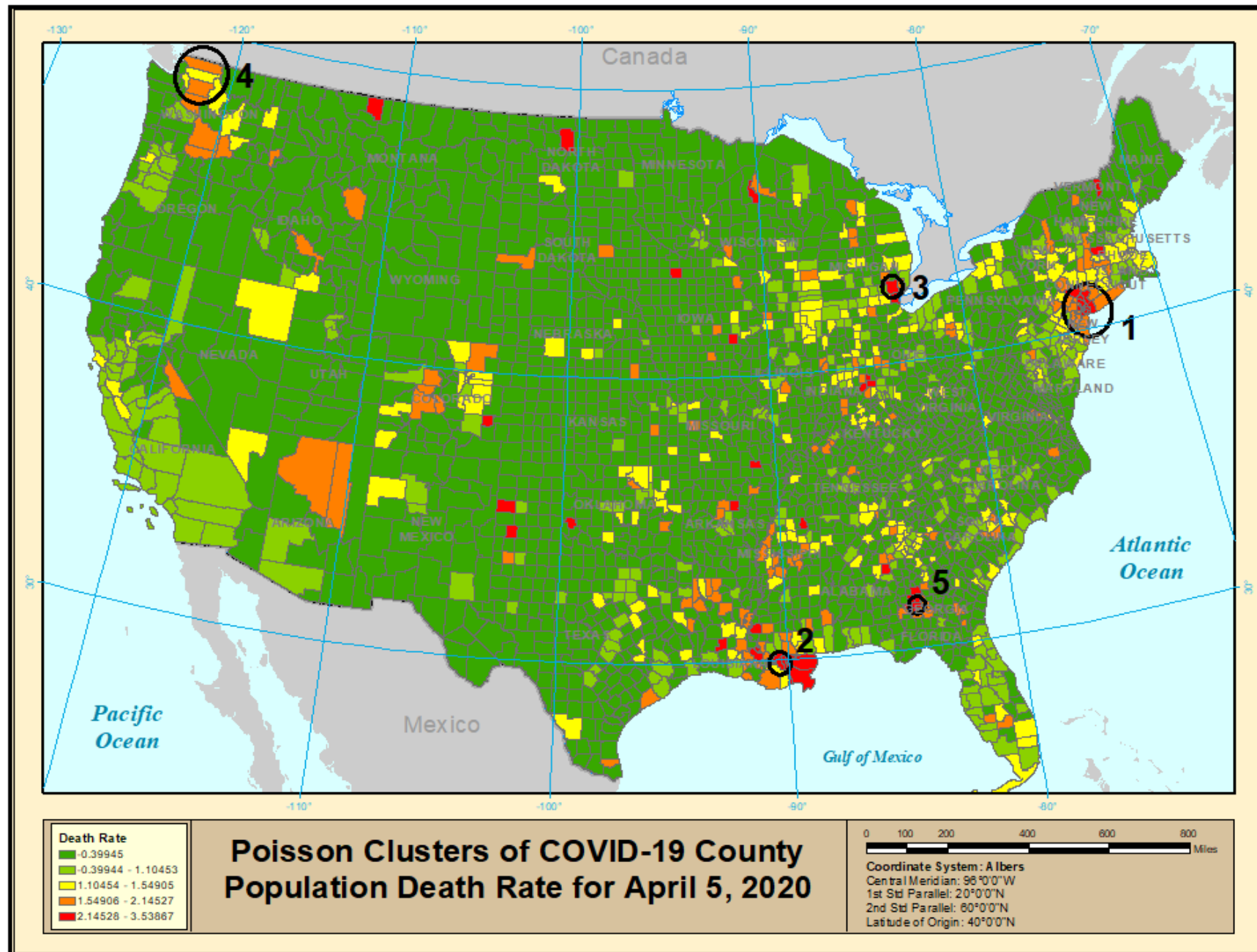




Spatial Scan Statistic: A small sample of the circles used

# Four Most Likely Clusters





Raid Amin et al, Univ West Florida

# Space-Time Scan Statistic

- Use a cylindrical window, with the circular base representing space and the height representing time.
- Retrospective surveillance: One analysis on a static data set, to detect historical and/or current clusters.
- Prospective surveillance: Daily or weekly analyses as new data accrues, to quickly detect emerging clusters. Only consider cylinders that reach the present time.

## **For each cylinder:**

- Obtain actual and expected number of cases inside and outside the cylinder.
- Calculate likelihood function.

## **Compare Cylinders:**

- Pick cylinder with highest likelihood function as Most Likely Cluster.

## **Inference:**

- Generate random replicas of the data set under the null-hypothesis of no clusters. May e.g. use 9,999 random replicas.
- Compare most likely clusters in real and random data sets (Likelihood ratio test). If  $R$ =rank of the real data set, then  $p=R/(1+9999)$

# Space-Time Scan Statistic: Properties

- Simultaneously tests for outbreaks of any size at any location, by using a cylindrical windows with variable radius and height.
- Accounts for multiple testing.
- Aggregated or non-aggregated data, such as counties, zip-codes, census tracts, hospitals or residences.

# Probability Models

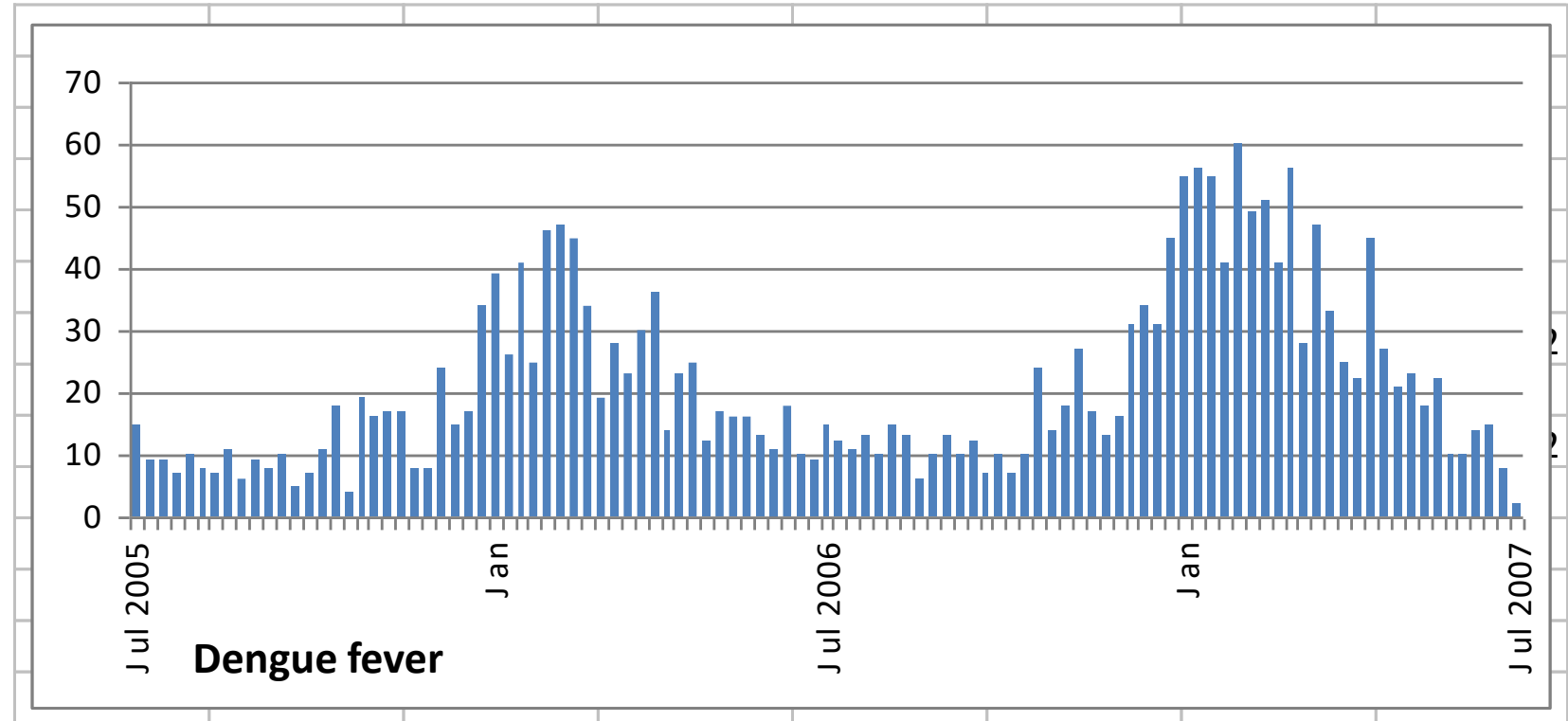
- **Bernoulli Model** (binomial): For 0/1 count data, such as tested positive vs negative.
- **Poisson Model**: For count data with expected counts under the null, such as incidence and age-adjusted population based expected counts.
- **Space-Time Permutation Model**: For count data without a denominator, automatically adjusting for purely spatial variation and purely temporal trends
- *Normal Model: For continuous data, such as birth weights*
- *Exponential Model: For time-to-event data, such as cancer survival*
- *Ordinal Model: For ordered categorical data with 3+ categories, such as cancer stage*
- *Multinomial Model: For un-ordered categorical data with 3+ categories, such as salmonella serotypes*

# Adjusting for Covariates

- Done in different ways for different probability models
- May or may not be appropriate or important
- When looking for unknown risk factors, adjust for age, smoking, etc, to find clusters not explained by those variables
- When allocating public health resources, do not adjust for age, as interest is in finding areas with more cases
- When implementing public health education, do not adjust for e.g. smoking, in order to find areas with a high-risk population

# Adjusting for Purely Temporal Trends

- Adjust if there is no interest in detecting region-wide temporal changes
- Non-parametric adjustments
- Log-linear adjustments, for a specific percent increase or decrease



- Log-linear adjustments may be calculated from the data, or provided by user from external information

# Adjusting for Purely Geographical Variation

- To adjust for background population, when census data is lacking
- To adjust for inherent differences in data collection, such as geographical variation in screening or testing
- When there is only an interest in emerging problem, rather than long-term geographical differences
- Adjustment is normally done non-parametrically
- Also possible to adjust for known geographical differences in the relative risks.

# Adjusting for both Geography and Time

- Possible to adjust the Poisson model for geography non-parametrically and time with a log-linear trend.
- Adjusting for both space and time non-parametrically, requires use of the space-time permutation model
- Missing data can be adjusted for by setting  $RR=0$  for the relevant locations and time periods. Should never be used when the true counts are zero. Only when the count is unknown.

# SaTScan Software

- SaTScan is a software that analyzes spatial, temporal and space-time data using the spatial, temporal, or space-time scan statistics.
- Free
- Download: [www.satscan.org](http://www.satscan.org)
- Thousands of users, including many at CDC and at least 44 of the 50 state health departments
- Funded by NCI, NICHD, NIGMS, CDC, Sloan Foundation
- Upcoming COVID-19 improvements funded by Alfred P. Sloan Foundation, CDC Foundation, ELC CARES, and Open Society Foundations
  - With support of Fund for Public Health in New York City

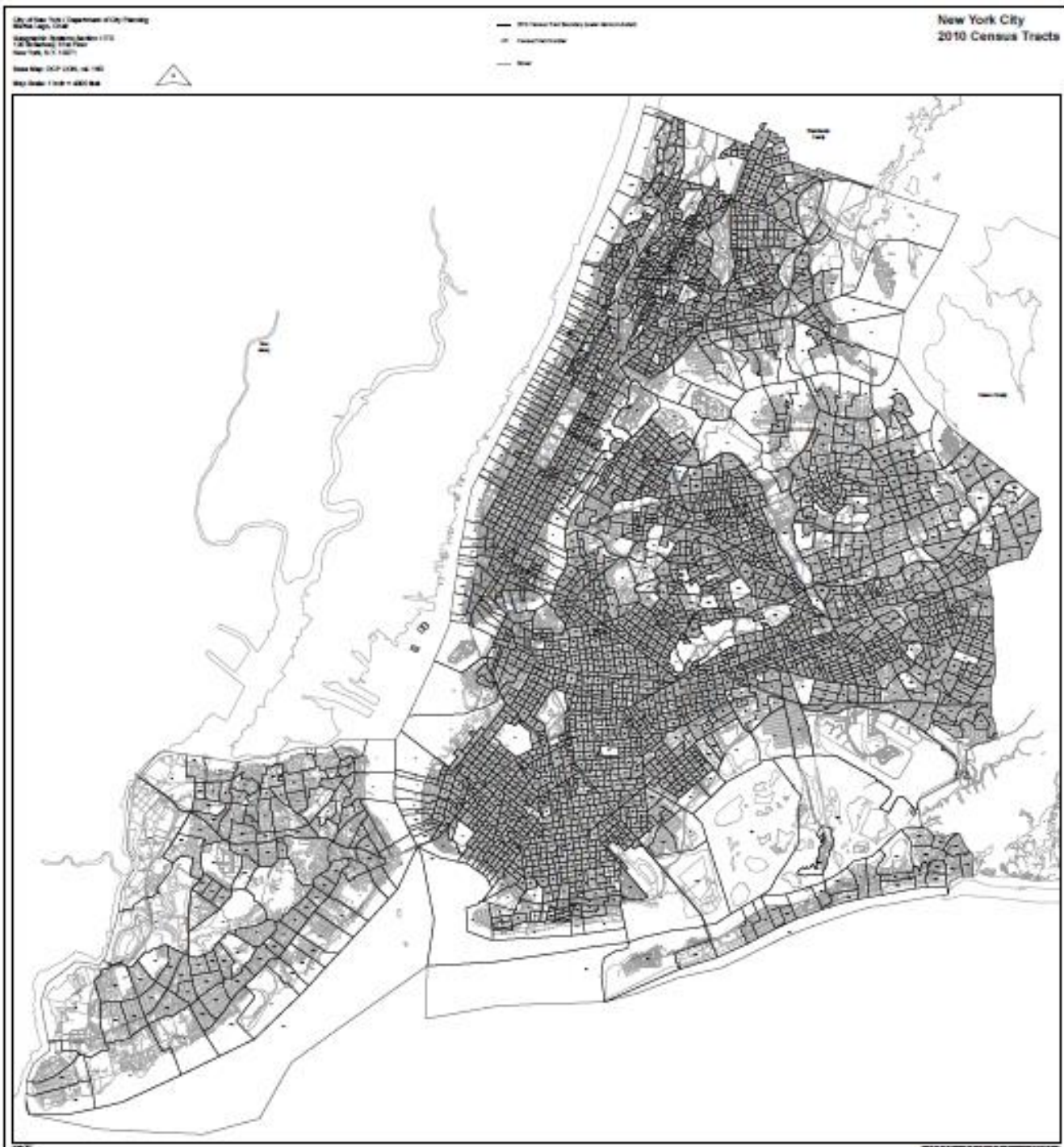


# Input files

- Numerator: Confirmed COVID-19 cases
- Denominator: Unique persons in electronic laboratory reports with PCR test for SARS-CoV-2
- Study period:
  - To prioritize case investigations daily: 21 days
  - To allocate testing and community engagement resources: since June 1, or shorter if discontinuity in citywide percent positivity trend
- Temporal element: specimen collection date
- Lag for data accrual: 3 days

# Input files

- Exclusions, to focus on recent community transmission:
  - Residents of long-term care facilities, correctional facilities, facilities housing people with developmental disabilities, homeless shelters
  - Home address matches hospital or laboratory
  - Diagnosed in 14 days prior to a more recent case residing in same building
  - Illness onset (where available from patient interview) prior to first date of study period
- Spatial resolution: Census tract 2010 (N=2165)
  - Automated geocoding process



# Analyses

- Detect disease clusters in space–time cylinders centered on every census tract centroid
- Circular base: represents space
  - Maximum geographic cluster size: 50% of persons tested
- Height: represents time
  - To prioritize case investigations daily:
    - Min–max temporal window: 3–7 days
    - Study period 21 days
  - To allocate testing and community engagement resources:
    - Min–max temporal window: 7–21 days
    - Study period: since June 1, or shorter if discontinuity in citywide percent positivity trend
- Minimum number of cases: 5

Adjust for purely spatial variation

- To detect areas with relative increases from baseline, even if still lower than the citywide average
- Will not detect areas with consistently high percent positivity
- Spatial adjustment: nonparametric, with spatial stratified randomization

## Adjust for purely temporal trends



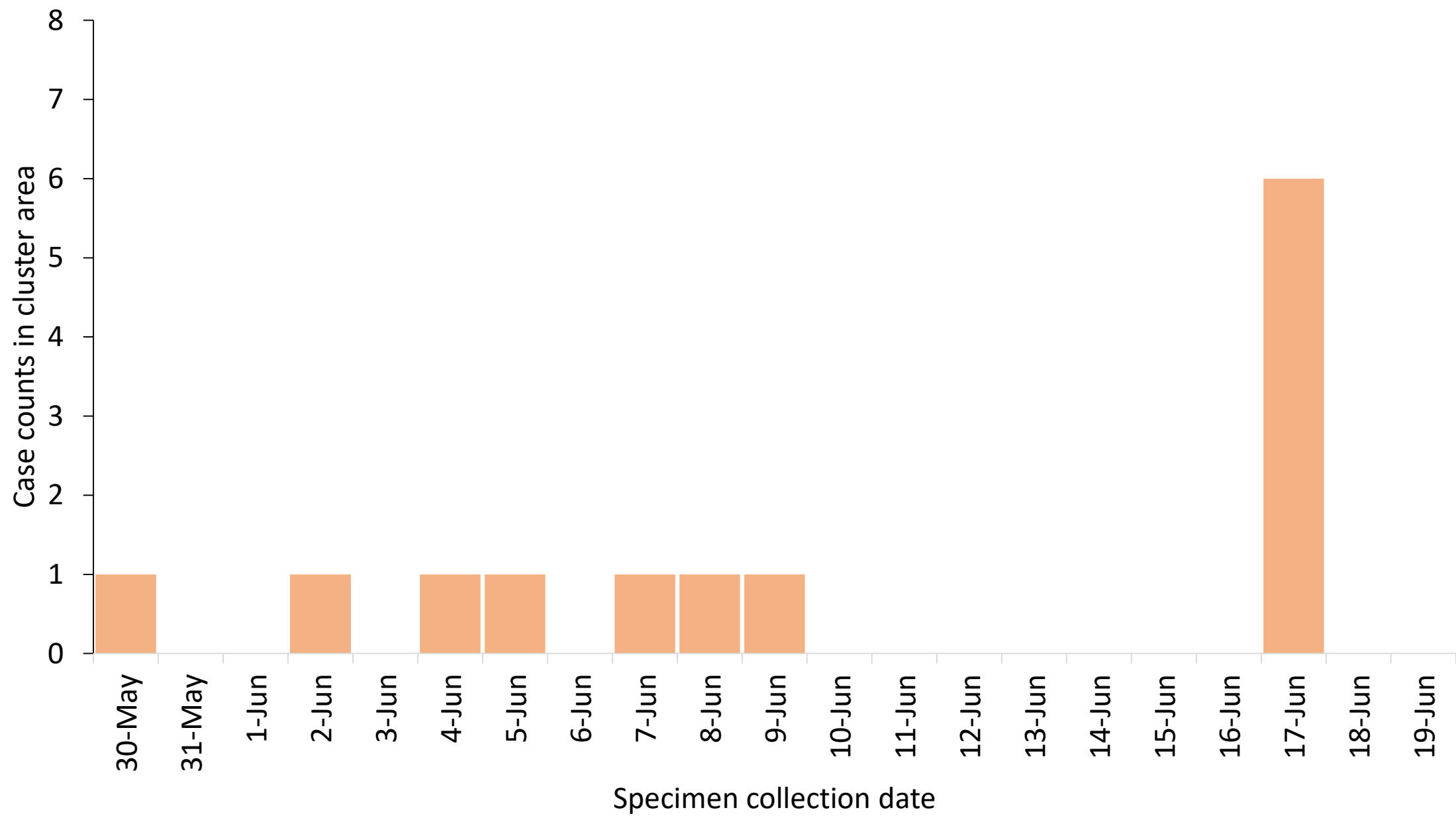
- If test % positivity is decreasing in NYC overall, detect areas where the decrease is less than citywide average
- If test % positivity is increasing in NYC overall, detect areas where the increase is more than citywide average
- Temporal adjustment: log-linear with automatically\* calculated trend
  - \*Specify input files in years, not days
  - SaTScan v9.6 estimates temporal trends at annual time scale
  - Pending SaTScan enhancement to accommodate rapidly changing trend

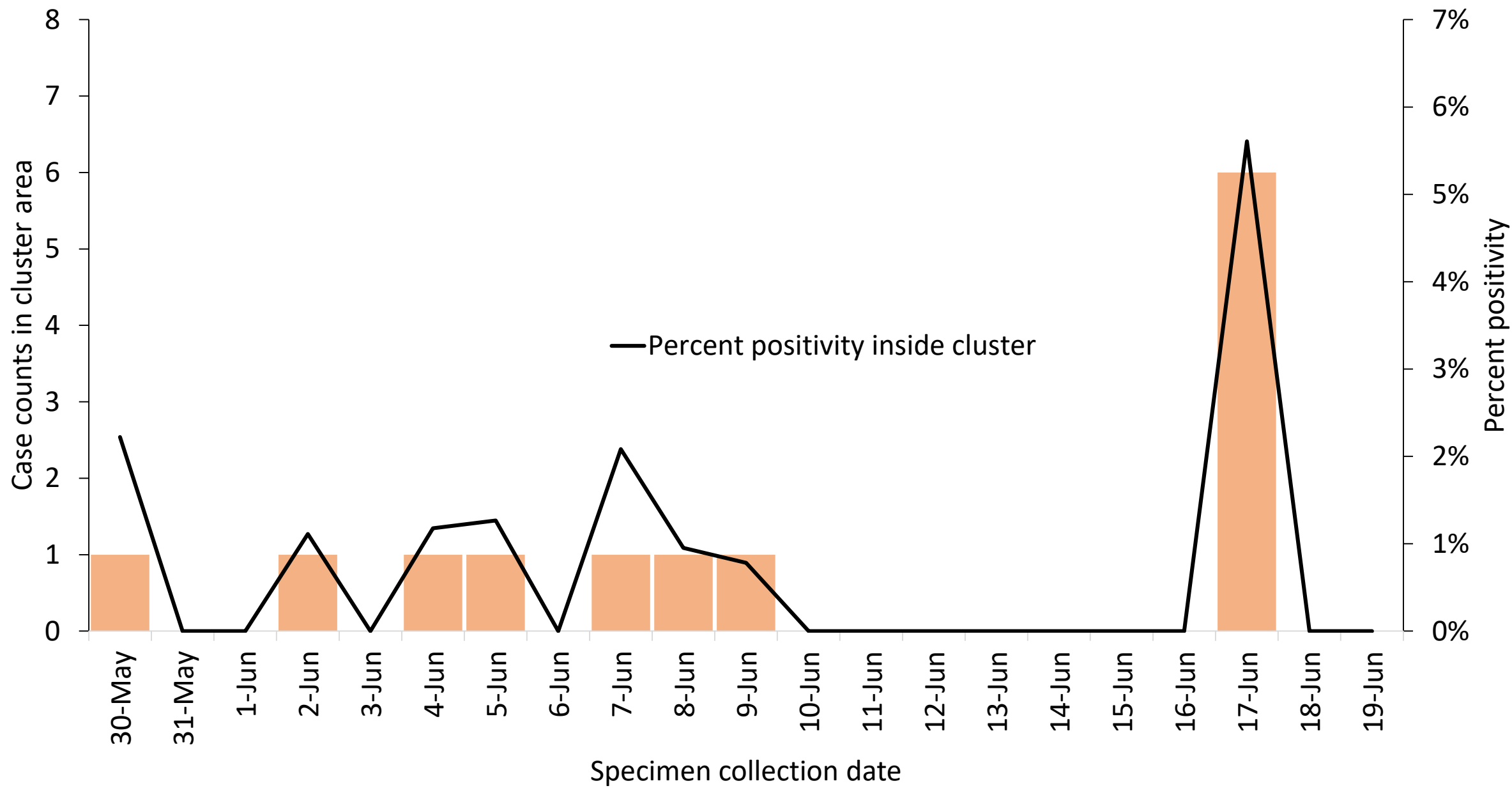
# Percent positivity spatiotemporal cluster detected June 22, 2020

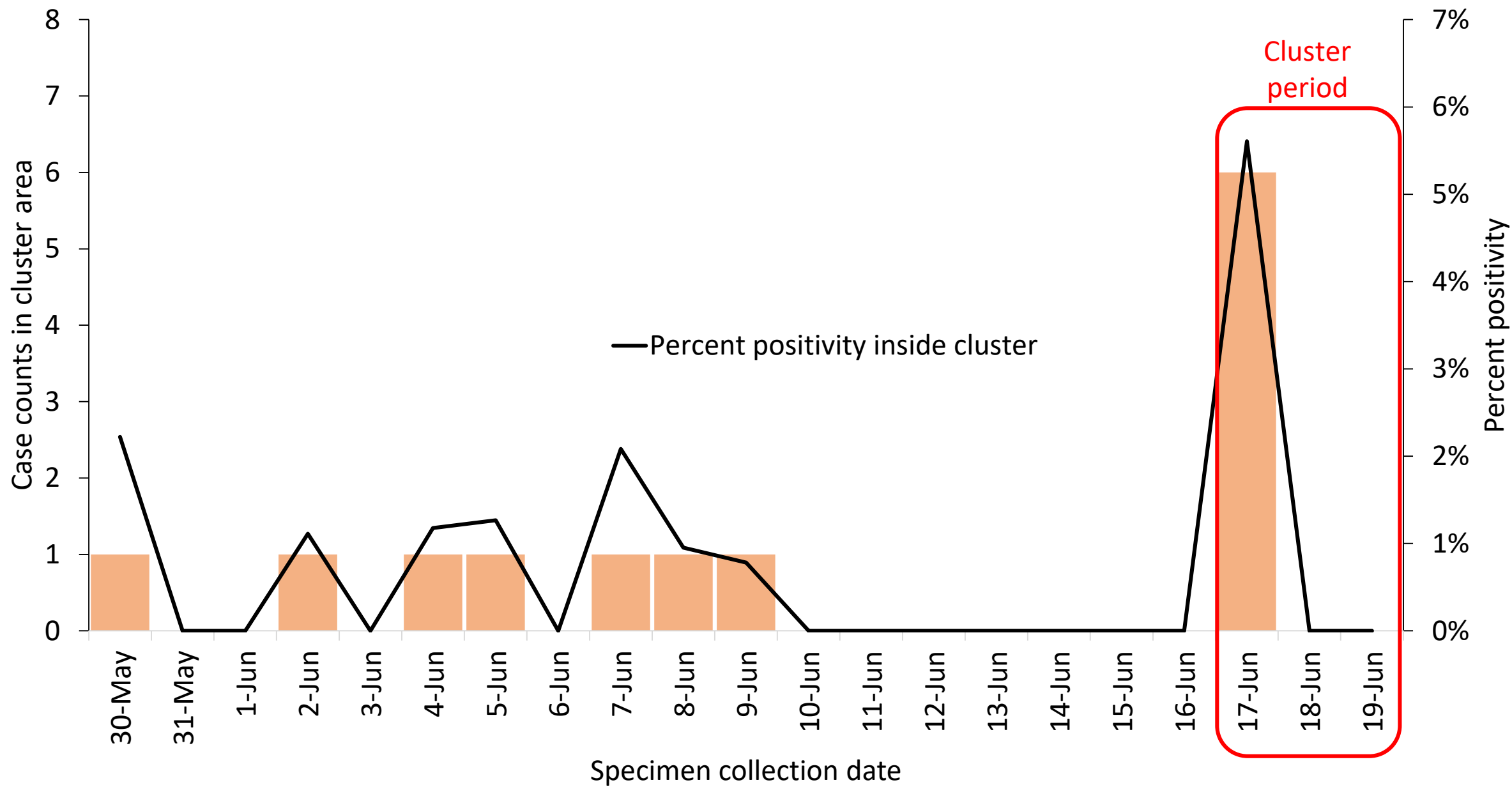
- During June 17–19:

| Radius<br>(km) | Observed<br>cases | Relative<br>risk | Recurrence<br>interval<br>(days) | % positivity<br>inside<br>cluster | % positivity<br>outside<br>cluster |
|----------------|-------------------|------------------|----------------------------------|-----------------------------------|------------------------------------|
| 0.6            | 6                 | 4.0              | 365                              | 2.2                               | 0.8                                |

- Prompted investigation identifying a gathering with inadequate social distancing where viral transmission likely occurred
- Response included testing, contact tracing, community engagement, and health education







SaTScan v9.6

Program run on: Mon Jun 22 05:17:52 2020

Prospective Space-Time analysis  
scanning for clusters with high rates  
using the Discrete Poisson model.  
Adjusted for time with a decrease of 6.42984% per year.  
Adjusted for purely spatial clusters by stratified randomization.

Case counts in cluster area

— Percent positivity inside cluster

..... Percent positivity outside cluster

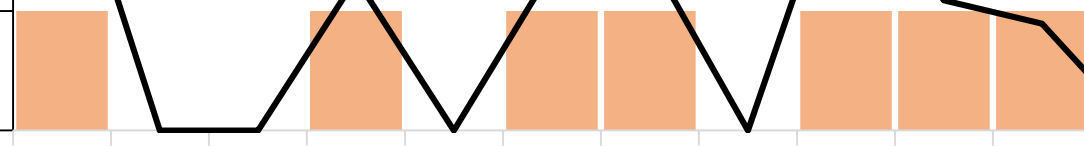
Percent positivity

8  
7  
6  
5  
4  
3  
2  
1  
0

7%  
6%  
5%  
4%  
3%  
2%  
1%  
0%

30-May  
31-May  
1-Jun  
2-Jun  
3-Jun  
4-Jun  
5-Jun  
6-Jun  
7-Jun  
8-Jun  
9-Jun  
10-Jun  
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15-Jun  
16-Jun  
17-Jun  
18-Jun  
19-Jun

Specimen collection date

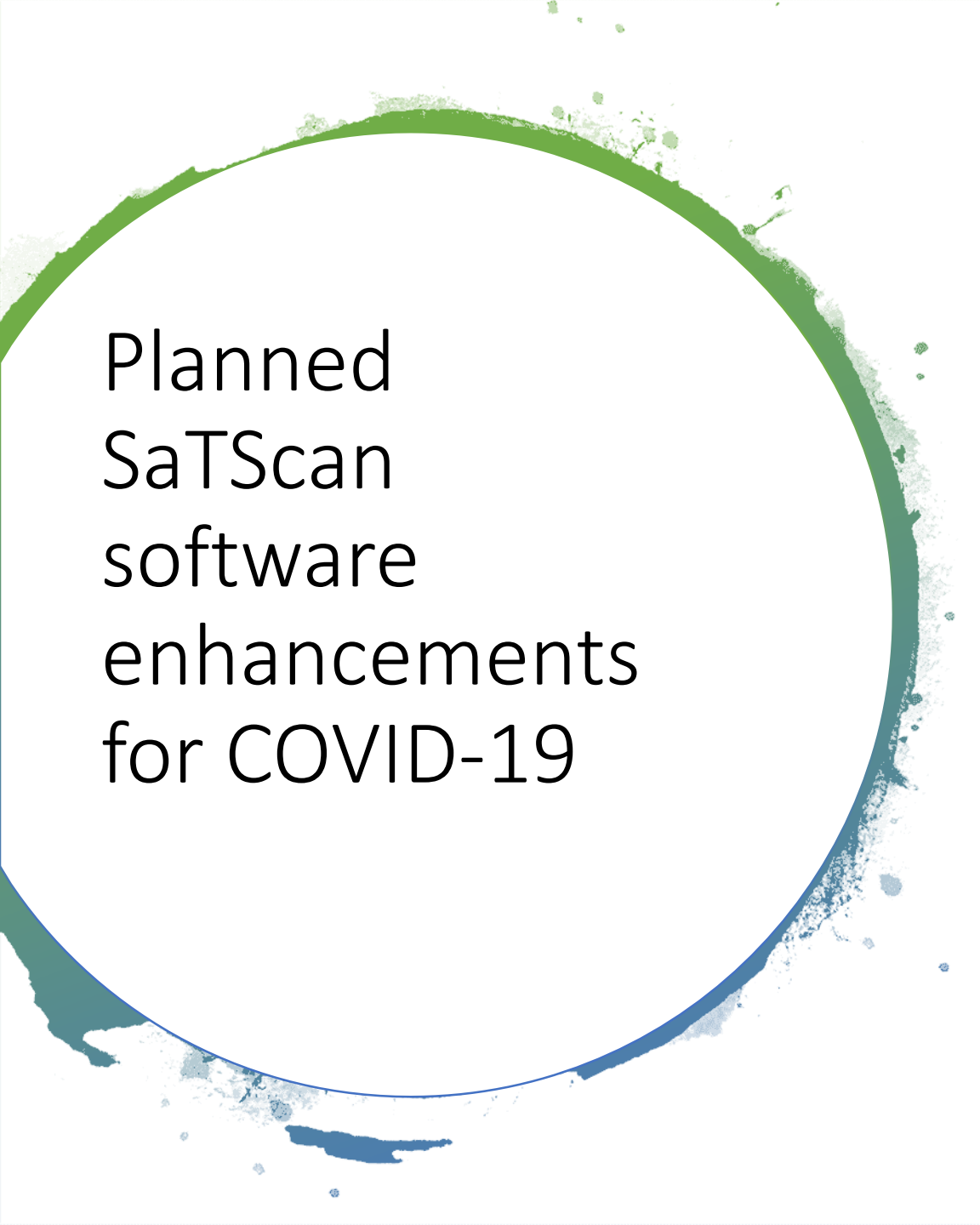


# Cluster Interpretation

- Prioritize clusters with:
  - High percent positivity
  - High relative risk
  - High recurrence interval
    - Areas with many persons tested can have highly significant clusters of only minor increases in percent positivity
- If searching for explicit epi-links:
  - Consider secondary clusters with their overlapping primary cluster
  - Track evolving clusters

# Automation

- Generate input files
- Invoke SaTScan in batch mode
- Output files to secured folders
- Define signals (e.g., clusters with recurrence interval  $\geq 100$  days), and generate patient linelist, visualizations, and investigator notification email
- NYC DOHMH, using SAS:  
<https://github.com/CityOfNewYork/communicable-disease-surveillance-nycdohmh>
- Cook County Dept of Public Health, using R:  
<https://github.com/kb230557/rsatscan-cluster-detection-covid>



# Planned SaTScan software enhancements for COVID-19

- Prioritize short emerging clusters
- Accommodate irregular geographies
- Detect clusters along a distance matrix/network
- Support weekly prospective analysis using data with daily resolution
- Temporal and spatial adjustments for Bernoulli model
- Descriptive and analytic outbreak investigation tools
- Software interface for prospective surveillance
- Automated outbreak alert notices