

University of Nebraska Medical Center [DigitalCommons@UNMC](https://digitalcommons.unmc.edu/) 

[Journal Articles: Epidemiology](https://digitalcommons.unmc.edu/coph_epidem_articles) **Executive Contract Articles: Executive Contract Articles: Executive Contract Articles:** 

2020

# Real-Time Public Health Communication of Local SARS-CoV-2 Genomic Epidemiology

Chaney C. Kalinich Yale School of Public Health

Cole G. Jensen Yale School of Public Health

Peter Neugebauer Yale School of Public Health

Mary E. Petrone Yale School of Public Health

Mario Peña-Hernández Yale School of Medicine

See next page for additional authors Tell us how you used this information in this [short survey.](https://unmc.libwizard.com/f/DCFeedback/) Follow this and additional works at: [https://digitalcommons.unmc.edu/coph\\_epidem\\_articles](https://digitalcommons.unmc.edu/coph_epidem_articles?utm_source=digitalcommons.unmc.edu%2Fcoph_epidem_articles%2F72&utm_medium=PDF&utm_campaign=PDFCoverPages)

**C** Part of the Epidemiology Commons

#### Recommended Citation

Kalinich, Chaney C.; Jensen, Cole G.; Neugebauer, Peter; Petrone, Mary E.; Peña-Hernández, Mario; Ott, Isabel M.; Wyllie, Anne L.; Alpert, Tara; Vogels, Chantal B. F.; Fauver, Joseph R.; Grubaugh, Nathan D.; and Brito, Anderson F., "Real-Time Public Health Communication of Local SARS-CoV-2 Genomic Epidemiology" (2020). Journal Articles: Epidemiology. 72.

[https://digitalcommons.unmc.edu/coph\\_epidem\\_articles/72](https://digitalcommons.unmc.edu/coph_epidem_articles/72?utm_source=digitalcommons.unmc.edu%2Fcoph_epidem_articles%2F72&utm_medium=PDF&utm_campaign=PDFCoverPages) 

This Article is brought to you for free and open access by the Epidemiology at DigitalCommons@UNMC. It has been accepted for inclusion in Journal Articles: Epidemiology by an authorized administrator of DigitalCommons@UNMC. For more information, please contact [digitalcommons@unmc.edu.](mailto:digitalcommons@unmc.edu)

#### Authors

Chaney C. Kalinich, Cole G. Jensen, Peter Neugebauer, Mary E. Petrone, Mario Peña-Hernández, Isabel M. Ott, Anne L. Wyllie, Tara Alpert, Chantal B. F. Vogels, Joseph R. Fauver, Nathan D. Grubaugh, and Anderson F. Brito

This article is available at DigitalCommons@UNMC: [https://digitalcommons.unmc.edu/coph\\_epidem\\_articles/72](https://digitalcommons.unmc.edu/coph_epidem_articles/72)

**PERSPECTIVE** 

## Real-time public health communication of local SARS-CoV-2 genomic epidemiology

 $\mathbf{C}$ haney C. Kalinich $\mathbf{D}^{1\mathbf{\odot}\,*}$ , Cole G. Jensen $\mathbf{D}^{1\mathbf{\odot}}$ , Peter Neugebauer<sup>1</sup>, Mary E. Petrone<sup>1</sup>,  $M$ ario Peña-Hernández<sup>2</sup>, Isabel M. Ott<sup>1</sup>, Anne L. Wyllie $\mathbf{O}^{\mathsf{T}}$ , Tara Alpert $\mathbf{O}^{\mathsf{T}}$ , Chantal B.  $\mathsf{F}\text{-}\mathsf{V}$ ogels $\textbf{D}^1$ , Joseph R. Fauver<sup>1</sup>, Nathan D. Grubaugh $\textbf{D}^1*$ , Anderson F. Brito<sup>1</sup>\*

**1** Department of Epidemiology of Microbial Diseases, Yale School of Public Health, New Haven, Connecticut, United States of America, **2** Department of Microbial Pathogenesis, Yale School of Medicine, New Haven, Connecticut, United States of America

 $\odot$  These authors contributed equally to this work.

\* chaney.kalinich@yale.edu(CCK); nathan.grubaugh@yale.edu(NDG); anderson.brito@yale.edu (AFB)

### Abstract

Genomic epidemiology can provide a unique, real-time understanding of SARS-CoV-2 transmission patterns. Yet the potential for genomic analyses to guide local policy and community-based behavioral decisions is limited because they are often oriented towards specially trained scientists and conducted on a national or global scale. Here, we propose a new paradigm: Phylogenetic analyses performed on a local level (municipal, county, or state), with results communicated in a clear, timely, and actionable manner to strengthen public health responses. We believe that presenting results rapidly, and tailored to a nonexpert audience, can serve as a template for effective public health response to COVID-19 and other emerging viral diseases.

#### **Genomic epidemiology**

Pathogen genomics has played a key role in understanding the emergence and rapid global spread of SARS-CoV-2. Since the beginning of the COVID-19 pandemic, viral genomes have been generated at an unprecedented rate, with over 60,000 sequences on GISAID by early July 2020 (gisaid.org). Genomic epidemiology, using pathogen genomes to explore patterns of transmission, is a growing field. Genomic epidemiological analyses infer evolutionary relationships among pathogens using genome sequences and phylogenetic approaches [1]. These analyses can now occur in near real-time  $[1,2]$  and have yielded important insights into the early COVID-19 pandemic in the United States, Brazil, and South Africa  $[3-7]$ . In the context of a devolved public health system, genomic epidemiology of SARS-CoV-2 performed at a local level can provide (at least) two key insights. First, it can reveal how transmission transcends specific administrative boundaries. Second, it can capture idiosyncratic factors driving transmission, such as patterns of human movement and local and state policies, to quickly provide the type of actionable targets for mitigation strategies that generalized models or country-level analyses cannot achieve.

Hyper-local, situational sources of spread have been shown to play a crucial role in driving outbreaks of other respiratory illnesses [8]. Genomic epidemiology has demonstrated potential to elucidate these sources. Effective deployment of rapid sequencing and analysis for public



### **G** OPEN ACCESS

**Citation:** Kalinich CC, Jensen CG, Neugebauer P, Petrone ME, Peña-Hernández M, Ott IM, et al. (2020) Real-time public health communication of local SARS-CoV-2 genomic epidemiology. PLoS Biol 18(8): e3000869. https://doi.org/10.1371/ journal.pbio.3000869

**Published:** August 21, 2020

**Copyright:** © 2020 Kalinich et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

**Funding:** Funding for covidtrackerct.com was provided by the Huffman Family Donor Advised Fund (NDG), Fast Grant funding support from the Emergent Ventures at the Mercatus Center, George Mason University (https://fastgrants.org/; NDG), Yale Institute for Global Health (https://medicine. yale.edu/yigh/; NDG), and NWO Rubicon 019.181EN.004 (https://www.nwo.nl/en/funding/ our-funding-instruments/nwo/rubicon/index.html; CBFV). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

**Competing interests:** The authors have declared that no competing interests exist.

health interventions depends on researchers' ability to clearly and accurately convey results, their implications, and knowledge gaps within an actionable timeframe [9]. Yet the majority of these detailed genomic epidemiological analyses are finished weeks after the outbreaks, and are not updated as new data arrives. They are intended to be case studies aimed at understanding how viruses spread through an area, providing information with potential application to future outbreaks [1]. Such delay, however, hinders the use of genomic epidemiology to inform the response to viral outbreaks as they unfold.

Some fast-evolving RNA viruses can spread rapidly, but their genetic changes (mutations) do not always accumulate at the same speed as viral transmission. This means that when viral genomes are sequenced during short timeframes as outbreaks unfold, their sequences can be remarkably similar. This low genetic resolution often leads to uncertainties in the phylogenetic reconstructions, generating nuanced results that require careful analysis to prevent misinterpretations [10]. Some of these concerns can be mitigated by pairing phylogenetic findings with the best available epidemiologic data and other supporting data [10]. Despite these trade-offs between speed and accuracy, real-time phylogenetic analyses that are clearly communicated to the general public can support policies to prevent further spread, and foster actions and behavior changes based on scientific evidence.

#### **Science communication**

A number of websites have played key roles in communicating the dynamics of SARS-CoV-2 spread in real-time by aggregating, analyzing, and displaying data. Dashboards with aggregated data such as case counts, tests, and hospitalizations became popular  $[11]$ . These dashboards often communicate data at global, national, and in the US, state and county levels, but do not (yet) incorporate any genomic data or analysis. Weekly situation reports of SARS-CoV-2 genomic epidemiology on global and continental levels have been published by Nextstrain (nextstrain.org) since January 23, 2020 [12], presenting the status of the pandemic across large geographical regions. Given the breadth of data covered by Nexstrain, however, there is a need for more fine-scale analysis and communication to impact policies at a local scale, as conducted by initiatives in the UK (COG-UK, cogconsortium.uk/news); Brazil (CADDE project, caddecentre.org/covid19) [6]; and South Africa (KRISP, krisp.org.za/covidnews.php) [7]. With these initiatives in mind, we propose three guiding principles for genomic epidemiologists looking to communicate their locally focused results in the context of the current COVID-19 pandemic (Box 1).

#### Box 1. Guiding principles for public communication of genomic epidemiology finding

- 1. **Communicate methods, analyses, and results in an accessible way.** For effective policy implementation, individual behavior adjustments, and to build trust in public health guidance, it is vital that genomic epidemiology findings are widely accessible, communicated in a jargon-free language, with appealing data visualizations and multilingual content (as appropriate).
- 2. **Speed must be balanced with accuracy, and uncertainty must also be communicated clearly.** Patterns revealed by available data may be inconclusive, or require more time or data to be informative. Even so, we believe that the benefit of sharing data when it can be acted upon in real-time outweighs the potential harm, as long

as the uncertainties are explained. It also engages the public in critically thinking about the public health response to a rapidly developing crisis.

3. **Results should be communicated at a local level, with actionable conclusions.** Rapid insight is useful only insofar as it is also communicated rapidly and clearly to those with the power to act on it. Local public health implications explicitly tied to policy and behavior should be a consistent focus of any report.

#### **Case Study: CovidTrackerCT**

We launched the Yale SARS-CoV-2 Surveillance Initiative to uncover the patterns of the spread and ongoing transmission of SARS-CoV-2 in Connecticut, USA. The public face of our initiative is CovidTrackerCT (covidtrackerct.com) (Fig 1A), a platform launched in late April 2020 to rapidly communicate our findings and translate our research into tangible public health benefits. CovidTrackerCT is based on the aforementioned framework (Box 1; Fig 1B). We present easily digestible descriptions of our analysis and major conclusions in both English and Spanish, alongside interactive figures facilitated by the Nextstrain visualization toolkit (Fig 1C and 1D). From April-July, 2020, our platform has attracted nearly 6,000 visitors.

Speed is emphasized, and results are made public within days of generating complete genomes (Fig 1C and 1D). To achieve a balance with accuracy, any caveats and limitations are disclosed. Also, supplemental information explaining genomic epidemiology concepts and protocols used to generate the results are made available in specific subpages. Results presented are framed in an actionable way; for example, discussing how the patterns of viral spread from neighboring states and between local towns highlight the importance of coordinated testing, contact tracing, and other non-pharmaceutical interventions.

The primary audience of the platform are those for whom the results are most consequential: state residents and those shaping public policy at the state and local level. The format and content of the weekly reports can be adapted in response to new information, needs, and tools. It is important to note that such science communication platforms are not intended to replace more traditional pathways of sharing this work, for example, peer-reviewed research papers, but serve a separate purpose of sharing it in real time, in a jargon-free way, to take advantage of the speed at which we can now generate and analyze data.

#### **Conclusions**

Many US states, including Connecticut, are looking to establish policies to mitigate virus spread for the coming months and years. As the outward-facing arm of the Yale SARS-CoV-2 Genomic Surveillance Initiatives, CovidTrackerCT played a role in informing local, state, and (Northeastern US) regional policy. Specifically, those behind the initiative have been contacted by mayors, state senators, and governors about our conclusions. After demonstrating how SARS-CoV-2 spread in Connecticut is highly influenced by outbreaks in surrounding areas, especially New York City, our results likely influenced policymakers to (**1**) enact greater coordination of control efforts between the adjacent states of Connecticut, New York, and New Jersey and (**2**) implement quarantine procedures for travelers arriving from affected states. Approaches to establishing these policies and communicating the reasoning behind them must be driven by data, including genomic data. Clear communication directly from experts provides an opportunity for closer interaction and trust-building between researchers and the public and helps protect against the spread of misinformation.



**Fig 1. Open Science and Communication.** (**A**) The COVIDTracker home page (coviditrackerct.com). Using an easy to maintain online platform, background information and regular reports are shared with the general public. (**B**) Our workflow from data generation, to rapid testing and communication, following principles of Open Science. (**C**) A maximum-likelihood phylogenetic tree showing 1048 SARS-CoV-2 genomes, including the first 200 SARS-CoVi2 genomes sequenced by our group. (**D**) The phylogeographic reconstruction of the genomes shown in (C) reveals the potential sources of introductions and patterns of viral spread within Connecticut. The size of the nodes represents the number of genomes sampled in the corresponding area, and the edges depict viral spread events between these areas. The results, which are embedded into our reports (available on covidtrackerct.com/portfolio/current/), can also be viewed standalone by loading output files hosted on GitHub (https://github.com/ grubaughlab/CT-SARS-CoV-2).

https://doi.org/10.1371/journal.pbio.3000869.g001

#### **Ethics statement**

Collection of clinical specimens at the Yale-New Haven Hospital for sequencing was approved by the Institutional Review Board of the Yale Human Research Protection Program (FWA00002571, Protocol ID. 2000027690).

#### **Acknowledgments**

We thank the Yale IMPACT Research Team, Yale-New Haven Hospital, and Connecticut Department of Public Health for their partnerships in our collective COVID-19 response.

#### **References**

- **1.** Grubaugh ND, Ladner JT, Lemey P, Pybus OG, Rambaut A, Holmes EC, et al. Tracking virus outbreaks in the twenty-first century. Nature Microbiology. 2019; 4: 10–19. https://doi.org/10.1038/s41564-018- 0296-2 PMID: 30546099
- **2.** Gardy JL, Loman NJ. Towards a genomics-informed, real-time, global pathogen surveillance system. Nat Rev Genet. 2018; 19: 9–20. https://doi.org/10.1038/nrg.2017.88 PMID: 29129921
- **3.** Deng X, Gu W, Federman S, du Plessis L, Pybus OG, Faria N, et al. Genomic surveillance reveals multiple introductions of SARS-CoV-2 into Northern California. Science. 2020. https://doi.org/10.1126/ science.abb9263 PMID: 32513865
- **4.** Gonzalez-Reiche AS, Hernandez MM, Sullivan MJ, Ciferri B, Alshammary H, Obla A, et al. Introductions and early spread of SARS-CoV-2 in the New York City area. Science. 2020. https://doi.org/10. 1126/science.abc1917 PMID: 32471856
- **5.** Fauver JR, Petrone ME, Hodcroft EB, Shioda K, Ehrlich HY, Watts AG, et al. Coast-to-Coast Spread of SARS-CoV-2 during the Early Epidemic in the United States. Cell. 2020; 181: 990-996.e5. https://doi. org/10.1016/j.cell.2020.04.021 PMID: 32386545
- **6.** Candido D da S, Claro IM, de Jesus JG, Souza WM de, Moreira FRR, Dellicour S, et al. Evolution and epidemic spread of SARS-CoV-2 in Brazil. medRxiv; 2020. https://doi.org/10.1101/2020.06.11. 20128249
- **7.** Giandhari J, Pillay S, Wilkinson E, Tegally H, Sinayskiy I, Schuld M, et al. Early transmission of SARS-CoV-2 in South Africa: An epidemiological and phylogenetic report. medRxiv. 2020. https://doi.org/10. 1101/2020.05.29.20116376 PMID: 32511505
- **8.** Dowdy DW, Azman AS, Kendall EA, Mathema B. Transforming the fight against tuberculosis: targeting catalysts of transmission. Clin Infect Dis. 2014; 59: 1123–1129. https://doi.org/10.1093/cid/ciu506 PMID: 24982034
- **9.** Black A, MacCannell DR, Sibley TR, Bedford T. Ten recommendations for supporting open pathogen genomic analysis in public health. Nat Med. 2020. https://doi.org/10.1038/s41591-020-0935-z PMID: 32528156
- **10.** Villabona-Arenas CJ, Hanage WP, Tully DC. Phylogenetic interpretation during outbreaks requires caution. Nat Microbiol. 2020. https://doi.org/10.1038/s41564-020-0738-5 PMID: 32427978
- **11.** Dong E, Du H, Gardner L. An interactive web-based dashboard to track COVID-19 in real time. Lancet Infect Dis. 2020; 20: 533–534. https://doi.org/10.1016/S1473-3099(20)30120-1 PMID: 32087114
- **12.** Hadfield J, Megill C, Bell SM, Huddleston J, Potter B, Callender C, et al. Nextstrain: real-time tracking of pathogen evolution. Bioinformatics. 2018; 34: 4121–4123. https://doi.org/10.1093/bioinformatics/ bty407 PMID: 29790939