

# Real-time analysis and visualization of pathogen sequence data

Richard A. Neher<sup>1,2</sup> and Trevor Bedford<sup>3</sup>

<sup>1</sup>*Biozentrum, University of Basel,  
Switzerland*

<sup>2</sup>*SIB Swiss Institute of Bioinformatics, Basel,  
Switzerland*

<sup>3</sup>*Vaccine and Infectious Disease Division,  
Fred Hutchinson Cancer Research Center, Seattle, WA,  
USA*

(Dated: March 21, 2018)

The rapid development of sequencing technologies has led to an explosion of pathogen sequence data that are increasingly collected as part of routine surveillance or clinical diagnostics. In public health, sequence data is used to reconstruct the evolution of pathogens, anticipate future spread, and target interventions. In clinical settings whole genome sequences identify pathogens at the strain level, can be used to predict phenotypes such as drug resistance and virulence, and inform treatment by linking to closely related cases. However, the vast majority of sequence data are only used for specific narrow applications such as typing. Comprehensive analysis of these data could provide detailed insight into outbreak dynamics, but is not routinely done since fast, robust, and interpretable analysis work-flows are not in place. Here, we review recent developments in real-time analysis of pathogen sequence data with a particular focus on visualization and integration of sequence and phenotypic data.

As pathogens replicate and spread, their genomes accumulate mutations. These changes can now be detected via cheap and rapid whole genome sequencing on unprecedented scale. Such sequence data are increasingly used to track the spread of pathogens and predict their phenotypic properties. Both applications have great potential to inform public health and treatment decisions if sequencing data can be obtained and analyzed rapidly. Historically, however, sequencing and analysis has lagged months-to-years behind sample collection. The results from these studies have taught us much about pathogen molecular evolution, genotype-phenotype maps, and epidemic spread, but have come almost always too late to inform public health interventions or treatment decisions.

The rapid development of sequencing technologies has made routine sequencing of viral and bacterial genomes possible and tens of thousands of whole genome sequences (WGS) are deposited in databases every year (see Fig. 1). Many more, regrettably, are sequenced and not shared. There are currently two major directions in which high-throughput sequencing technologies are used in public health and diagnostics: (i) to track outbreaks and epidemics to inform public health response, and (ii) to characterize individual infections to tailor treatment decisions.

*Sequencing in public health.* The utility of rapid sequencing and phylogenetic analysis of pathogens is perhaps most evident for influenza viruses and food-borne diseases. Due to rapid evolution of its viral surface proteins, the antigenic properties of the circulating influenza viruses change every few years and the seasonal influenza vaccine needs frequent updating<sup>1</sup>. The WHO Global Influenza Surveillance and Response System (GISRS) sequences hundreds of viruses every month and many of these sequences are submitted to the GISAID database ([gisaid.org](http://gisaid.org)) within 4 weeks of sample

collection. Phylogenetic analysis of these data provide an accurate and up-to-date summary of the spread and abundance of different viral variants that is crucial input to the biannual consultations on seasonal influenza vaccine composition.

Such rapid turn-around and data sharing is considerably harder to achieve in an outbreak setting in resource limited conditions. Nonetheless, Quick et al.<sup>2</sup> achieved even shorter turn-around during the tail end of the 2014–2015 West African Ebola outbreak. Similarly, Dyrda et al.<sup>3</sup> analyzed an enterovirus outbreak in Sweden and continuously updated the manuscript until publication with sequences sampled within days of publication included in the analysis.

Molecular epidemiology techniques can reconstruct the temporal and spatial spread of an outbreak. In this case, the accumulation of mutations alongside a molecular clock estimate can be used to date the origin of an outbreak. Similarly, by linking samples that originate from different geographic locations, phylogeographic methods can reconstruct geographic spread and differentiate distinct introductions. The resolution of these inferences critically depends on the rate at which mutations accumulate in the sequenced locus, which increases with the per site evolutionary rate and the length of the locus.

RNA viruses accumulate changes in their genome with a typical rate of 0.0005 to 0.005 changes per site and year. Ebola virus and Zika virus, for example, evolve at a rate of  $\mu \approx 0.001$  per site per year and have a genome length  $L$  of 10kb and 19kb, respectively. The expected time interval without a substitution along a transmission chain is  $1/(\mu L)$ , which is approximately 5 weeks for Zika virus and 3 weeks for Ebola virus. Hence evolution and spread of RNA viruses on the scale of a month can be resolved. While this temporal resolution is typically insufficient to resolve individual transmissions, it is high compared to the duration of outbreaks. Rapid sequencing

and analysis therefore has the potential to inform intervention efforts as outbreaks are unfolding.

Phylogenetic and phylogeographic methods are best established for viral pathogens with high evolutionary rates and small genomes for which large scale sequencing has been possible for years. The evolutionary rates of bacteria are many orders of magnitudes lower than those of RNA viruses. However, bacteria also have about 100 to 1000-fold larger genomes and it is now possible to sequence entire bacterial genomes at low cost. Substitution rate estimates in bacteria come with substantial uncertainty, but they tend to be on the order of one substitution per megabase per year<sup>4</sup>. With a typical genome size of 5 megabases, this translates into 5–10 substitutions per genome and year — similar to many RNA viruses. Hence real-time phylogenetics should be possible for bacterial outbreak tracking in much the same way. Analysis of bacterial genomes, however, is vastly more complicated than that of RNA viruses with short genomes. Bacteria frequently exchange genetic material via horizontal transfer, take up genes from the environments and rearrange their genome. If not properly accounted for, these processes can blur any temporal signal and obscure links between closely related isolates.

While the resolution typically does not allow to make the case for a direct transmission, transmission can be confidently ruled out for divergent sequences, seemingly unrelated cases can be grouped into outbreaks (e.g. an outbreak of drug resistant MtB among migrants arriving in multiple European countries<sup>5</sup>), predominant routes of transmission and likely sources in the environment or animal reservoirs can be identified. GenomeTrakr, for example, is a large federated effort to sequence thousands of genomes from food-borne outbreaks<sup>6</sup>.

These examples illustrate the potential and feasibility of obtaining actionable information from pathogen sequence data<sup>7</sup>, but there are considerable hurdles that need to be overcome to achieve this goal on larger scales for a broad collection of pathogens.

*Sequencing in diagnostics and therapy.* For some pathogens like Zika virus, sequencing the genome has no implications for treatment. In the case of HIV, however, drug resistance profiles derived from sequence data have directly informed treatment for years<sup>8</sup>. As the genetic basis of drug resistance phenotypes are better understood, rapid whole genome sequencing will increasingly be used to diagnose and phenotype pathogens directly from the clinical specimen. Such culture-free methods are particularly important for tuberculosis, in which culture based susceptibility testing takes many weeks. Votintseva et al.<sup>9</sup> have recently shown that high-throughput sequencing directly from respiratory samples can provide drug resistance profiles of *M. tuberculosis* within a day.

The focus of this mini-review is on applications of WGS in molecular epidemiology. Above, we discussed sequencing in clinical settings since data generated for diagnostic purposes can aid surveillance, while surveillance data provides context for the individual case in the clinics. Public health response typically requires recent data with an emphasis on dynamics.

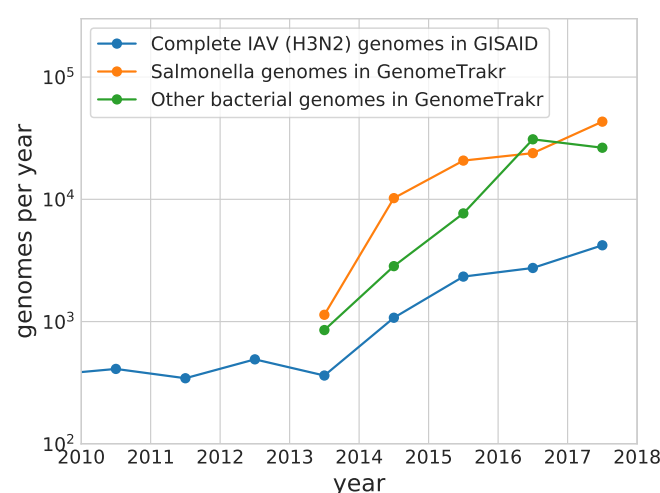


FIG. 1 The number of complete pathogen genomes has increased dramatically over the last few years. More than 4000 complete influenza A (IAV) subtype H3N2 virus genomes have been deposited in GISAID in 2017. The GenomeTrakr network sequenced in excess of 40,000 *Salmonella* genomes and 25,000 other bacterial genomes (mostly *Listeria*, *E. coli*/*Shigella*, and *Campylobacter*) in 2017<sup>6</sup>.

Informing a treatment decision needs a stable database with validated content to make reliable predictions on drug susceptibility, phylogenetic context, and protective measures. Clinical sequencing data, however, should be fed into surveillance databases immediately whenever ethically possible. Only with rapid and open sharing of sequencing data can the full potential of molecular epidemiology be realized<sup>6</sup>.

The challenges involved in sample collection, processing, sequencing and data sharing have been discussed at length elsewhere<sup>10</sup>. Here, we focus on algorithm and software developments that facilitate the implementation of real-time surveillance platforms.

## RAPID AND INTERPRETABLE ANALYSIS OF GENOMIC DATA

A typical molecular epidemiological analysis aims to date the introduction, detail geographic spread, and identify potential phenotypic change of a pathogen from sequence data. The rapidly increasing numbers of sequenced genomes make comprehensive analysis computationally challenging. While 1000s of viral genomes can be aligned within minutes<sup>11</sup> and the reconstruction of a basic phylogenetic tree typically takes less than one hour<sup>12;13;14</sup>, the most popular tool for phylogenetic inference (BEAST) will often take weeks to finish<sup>15</sup>.

To overcome these hurdles, several tools that use simpler heuristics have been developed to infer time-stamped phylogenies<sup>16;17;18</sup>. Rather than sampling a large number of tree topologies, these tools use the topology of an input tree with little or no modification. Dating of ancestral events tends to be of comparable accuracy to BEAST, at least for simi-

lar sequences that recently diverged. However, these tools do not integrate uncertainty of tree reconstruction and provide limited flexibility to infer demographic models. The computational cost of Bayesian phylodynamics could be mitigated if continuous updating and augmenting of the Markov chain with additional data was developed. For the time being, however, efficient heuristics and sensible approximations deliver sufficiently accurate and reliable results when near real-time analysis is required.

## Viral genomes: nextflu and nextstrain

The number of influenza viruses that are sequenced and phenotyped per month has increased sharply to a point that a comprehensive and timely manual analysis and annotation of the results is no longer feasible. Nextflu is a phylodynamic analysis pipeline that operates on an up-to-date database of sequences and serological information<sup>20</sup>. Primary outputs include a phylogeny, branch-specific mutations, clade frequency trajectories and antigenic phenotypes.

While nextflu is influenza specific, nextstrain aims at analysis and surveillance of multiple diseases and provides a platform for outbreak investigation<sup>19</sup>. Nextstrain uses TreeTime<sup>18</sup> to infer time-scaled phylogenies and conduct ancestral sequence inference. In addition, nextstrain uses the discrete ancestral character inference of TreeTime to infer the likely geographic state of ancestral nodes. Since this approach applies “mutation” models to “migration”, it is often called “migration model”. A phylodynamic/phylogeographic analysis of 1000 sequences of length 10kb takes on the order of an hour.

## Bacterial WGS data

Bacterial WGS data typically comes in the form of millions of short reads that can either be assembled into contigs, mapped against reference sequences, or classified based on kmer distributions.

Tools like WGSa allow the user to upload an assembly and WGSa will detect the species and infer the multi-locus sequence type within a few seconds. In addition, WGSa predicts antibiotic resistance profiles for a number of species. WGSa was developed by the *Center for Genomic Pathogen Surveillance* and is available at [wgsa.net](http://wgsa.net).

To account for the fact that bacteria have a very dynamic gene repertoires and clinically important genes are often horizontally transferred between strains and species, collections of bacterial genomes are often analyzed using pan-genome tools. These tools aim to cluster all genes in the collection of genomes into orthologous groups. Genes that are present in every genome are called *core genes* and genes present in only a fraction of individuals are referred to as *accessory genes*. Early methods for pan-genome analysis scaled poorly with the number of genomes that are analyzed since every gene in every genome needed to be compared to every other gene. The

first tool capable of analyzing 100s of bacterial genomes was Roary<sup>21</sup>. Roary is designed to work with very similar genes (as is common in outbreak scenarios) and accelerates inference of orthologous gene clusters by pre-clustering genomes. A more recent pan-genome analysis pipeline capable of large scale analysis is panX<sup>22</sup> that speeds up clustering by hierarchically building up the complete pan-genome from sub-pan-genomes inferred from smaller batches of genomes.

While the pan-genome tools cluster annotated genes in the collection of genomes, they are of little help to assess the origin and distribution of a particular sequence. Traditional tools for homology search in NCBI only index assembled sequence, but today the majority of sequence data are stored in short read archives rather than in Genbank. Bradley et al.<sup>23</sup> developed a method to search the entire collection of microbial sequence data including meta genomic samples from a wide variety of environment. The ability to search this vast treasure trove of data will likely be transformative in assessing spread and prevalence of novel resistance determinants. The recently discovered mobile colistin resistance gene *mcr-1*, for example, was found in more than 100 datasets where it wasn't previously described.

## Outlook

Most current analysis pipelines require rerunning the entire analysis even when only a single sequence is added. While this strategy is still feasible today, this will likely become unsustainable in the future. Applications that support cheap updating of datasets and on-line addition of user data will likely replace current versions.

## VISUALIZATION AND INTERPRETATION

With increasing dataset sizes, interpretation and exploration of data become increasingly challenging. Phylogenetic trees can be visualized as familiar planar graphs, but the tree alone only shows genetic similarity between isolates and becomes quickly unintelligible as the number of sequences increases. To make pathogen sequence data truly useful, it needs to be integrated with other types of information, ideally in an interactive way. A suitable platform to do so is the web-browser and several powerful web applications have emerged over the last few years. In addition, browser-based visualization are naturally disseminated online.

## Microreact

Microreact is a web application based on React (a JavaScript framework for interactive applications), D3.js (a JavaScript library for producing dynamic, interactive data visualizations), Phylocanvas (a JavaScript flexible tree viewer), and Leaflet (a JavaScript mapping toolkit). Microreact allows exploration of a phylogenetic tree, the geographic locations,

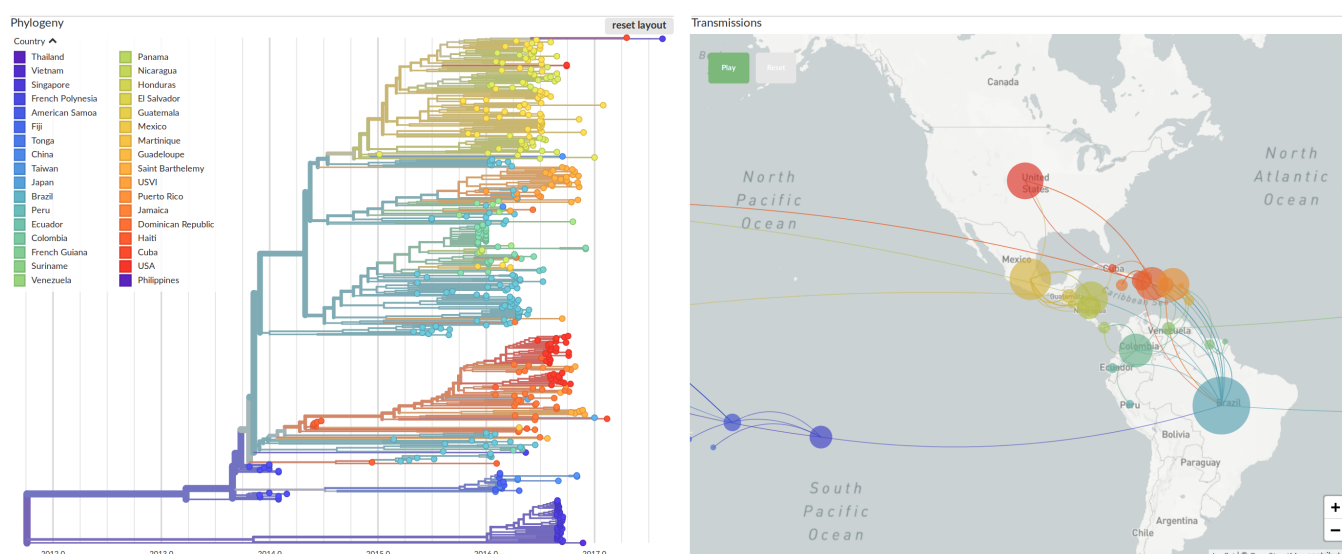


FIG. 2 Phylogeographic analysis of Zika virus sequences on nextstrain.org<sup>19</sup>. Whole genomes sequences sampled between 2013 and 2017 were processed using the nextstrain pipeline. Nextstrain reconstructs likely time and place of each internal node of the tree and from this assignments infers possible transmission patterns that are displayed on a map. Molecular analysis of this sort reveals for example multiple introductions of Zika virus into Florida originating most likely from viruses circulating in the Caribbean in 2015-2016.

and a time line of the samples<sup>24</sup>. Custom data sets can be loaded into the application in the form of a Newick tree and a tabular file containing information such as geographic location or sampling data.

## nextstrain

Nextstrain was developed as a more generic and flexible version of nextflu<sup>19;20</sup>. Similar to Microreact, nextstrain uses React, D3.js, and Leaflet, but uses a custom made tree viewer that has flexible zooming and annotation options. The tree can be decorated with any discrete or continuous attribute, both on tips of the tree and inferred values for internal nodes (for example geographic location). Nextstrain maps individual mutations to branches in the tree and thereby allows to associate mutations with phenotypes or geographic distributions. The map in nextstrain shows putative transmission events and a panel indicates genetic diversity across the genome (Fig. 2).

The analyses by nextstrain and nextflu critically depend on timely and open sharing of sequence information that many laboratories around the globe contribute. To incentivize early pre-publication sharing of data, platforms like nextstrain need to explicitly acknowledge the individual contributions. Ideally, such platforms should provide added value to authors, such as for example deep links that show data by a particular group in the context of the outbreak.

## Phandango

Phandango is an interactive viewer for bacterial whole genome sequencing data<sup>25</sup> and combines a phylogenetic

tree with metadata columns and gene presence-absence maps or recombination events. Phandango is available at [phandango.net](http://phandango.net) and can ingest the output of a number of analysis tools commonly used for the analysis of bacterial WGS data such as Gubbins, Roary and BRAT.

## panX

PanX is a pan-genome analysis pipeline that is coupled to a web-browser based visualization<sup>22</sup>. Similar to Phandango, it displays a core genome SNP phylogeny but is otherwise more centered on genetic variation in individual genes. The tree and alignment of each gene in the pan-genome can be accessed rapidly by a searching a table of gene names and annotations. PanX then displays gene and species tree side by side and maps gene gain and loss events to branches in the core genome tree and mutations to branches in the gene tree. Trees can be colored by arbitrary attributes such as resistance phenotypes and associations between genetic variation and these phenotypes can be explored.

## Other tools

Spread3 allows of visualization of phylogeographic reconstructions from models implemented in the software package BEAST<sup>26</sup>. PhyloGeoTool is a web-application to interactively navigate large phylogenies and to explore associated clinical and epidemiological data<sup>27</sup>. TreeLink displays phylogenetic trees alongside metadata in an interactive web application<sup>28</sup>.



## CHALLENGES IN DATA INTEGRATION AND VISUALIZATION

Epidemiological investigation of a novel outbreak typically seek to identify the sources, track the spread, and detect transmission chains. In this case, a generic combination of map, tree, and time line will often be an appropriate and sufficient visualization. Nextstrain and Microreact both follow this paradigm.

However, when analyzing established pathogens that continuously adapt to treatments, vaccines, or pre-existing immunity, more specific applications will be necessary since case data, phenotype data, and clinical parameters differ wildly by pathogen. Such data will generally have a common core such as sample date and location, but other parameters such as drug resistance phenotypes, disease severity, host age, risk group, serology, etc., are pathogen specific. While these data are often stored in non-standardized formats and ethical and technical reason can impede data sharing, these data are often at least as important for interpretation of the epidemiological dynamics as phylodynamic inference from sequence data. The value of either data is greatly increased by seamless integration, but the idiosyncrasies require flexible analysis and visualization frameworks that can be tailored to specific pathogens.

One such example is the serological characterization of influenza viruses via hemagglutination inhibition (HI) titers using antisera raised in ferrets. Such titers are routinely measured as part of GISRS to monitor the antigenic evolution of influenza viruses are a good example how phenotype information can be interactively integrated with phylogeny and molecular evolution. HI titers are reported in large tables and have been traditionally visualized using multidimensional scaling without any reference to the phylogeny. In Bedford et al.<sup>29</sup> and Neher et al.<sup>30</sup>, we developed methods to integrate the molecular and antigenic evolution of influenza virus. This integration allows association of genotypic changes with antigenic evolution and suggests an intuitive and interactive visualization of HI titer data on the phylogeny. A screenshot of this integration is shown in Fig. 3. Due to data sharing restrictions, most HI titer data are not openly available, but historical data by McCauley and colleagues are visualized along with the molecular evolution at [hi.nextflu.org](http://hi.nextflu.org).

In addition to phenotype integration, it is crucial to choose the right level of detail for a specific application. This is particularly true for bacteria where the relevant information might be the core genome phylogeny, the presence/absence of particular genes or plasmids, or individual mutations in specific genes. If the analysis tool and the visualization does not provide a fine grained analysis at the relevant level, the most important patterns might stay hidden. On the other hand, sifting through every gene or mutation is prohibitive. The primary aim should be to highlight the most important and robust patterns upfront and provide flexible methods to filter and rank variants (e.g. by recent rise in frequency, association with host, resistance or risk group, etc). The user should then have

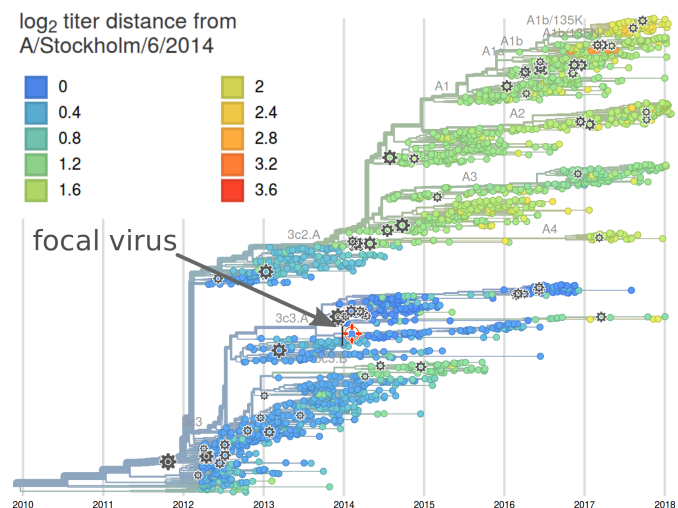


FIG. 3 Integration of HI titer data with molecular evolution influenza virus. Each year, influenza laboratories determine thousands of HI titers of test viruses relative to sera raised against several reference viruses (indicated by gray cogs). These data can be integrated with the molecular evolution of the virus and visualized on the phylogeny (here showing inferred titers using a model). The reference virus with respect to which titers are displayed can be chosen by clicking on the corresponding symbol in the tree<sup>30</sup>. The visualization exposes both raw data (via tool tips for each virus) as well as a model inference that integrates many individual measurements ([hi.nextflu.org](http://hi.nextflu.org)).

the possibility to expose detail on demand when a deeper exploration is required.

Similarly, parameter inferences and model abstractions are very useful to get a concise summary of the data, but should be complemented by the ability of interrogate the raw data (e.g. an estimate of the evolutionary rate should be accompanied by a scatter plot of root-to-tip divergence and sampling time). This is particularly important in outbreak scenarios when methods are applied to an unusual pathogen in a developing situation.

For clinical applications, the presentation of the results of an analysis should be focused on the sample in question and only provide reliable and actionable information, while suggestive and correlative results tend to be a distraction<sup>31</sup>.

## CONCLUSIONS

High-throughput and rapid sequencing is revolutionizing infectious disease diagnostics and epidemiology. Sequence data can be used to unambiguously identify pathogens, to link related cases, to reconstruct the spread of an outbreak, and will soon allow detailed prediction of a pathogen's phenotype.

The Global Influenza Surveillance and Response System (GISRS) is a good example of a near real-time surveillance system. Hundreds of viruses are sequenced and phenotyped every month and the sequence data is shared in a timely manner. A global comprehensive analysis of these data, updated about once a week, is available at [nextflu.org](http://nextflu.org). These

analysis directly inform the influenza vaccine strain selection process<sup>1</sup>.

The FDA has embraced whole genome sequencing and collectively the GenomeTrakr network now sequences and openly releases about 5000 bacterial genomes per month<sup>6</sup>. To our knowledge, these data are currently not comprehensively analyzed on a timely bases. Such a study, would sure provide unprecedented insights into the spread of food-borne pathogens.

These two examples clearly show that near real-time genomic surveillance is possible and valuable and all the individual components to implement such surveillance are in place. However, to realize this potential for many more pathogens, sample collection and sequencing has to be streamlined, data need to be shared along with the relevant meta data, and specific analysis methods and visualizations need to be implemented and maintained.

## ACKNOWLEDGMENTS

We are indebted to scientists around the globe for timely and open sharing of sequence data, without which real-time visualization would not be worthwhile. We are grateful to James Hadfield for critical reading of the manuscript.

## REFERENCES

- [1] Morris DH, Gostic KM, Pompei S, Bedford T, Luksza M, Neher RA, Grenfell BT, Lässig M, McCauley JW. Predictive Modeling of Influenza Shows the Promise of Applied Evolutionary Biology. *Trends microbiology* 2017; .
- [2] Quick J, Loman NJ, Duraffour S, Simpson JT, Severi E, Cowley L, Bore JA, Koundouno R, Dudas G, Mikhail A, et al. Real-time, portable genome sequencing for Ebola surveillance. *Nature* 2016; 530(7589):228–232.
- [3] Dyrda R, Grabbe M, Hammas B, Ekwall J, Hansson KE, Luthander J, Naucier P, Reinius H, Rotzén-Östlund M, Albert J. Outbreak of enterovirus D68 of the new B3 lineage in Stockholm, Sweden, August to September 2016. *Eurosurveillance* 2016; 21(46).
- [4] Duchêne S, Holt KE, Weill FX, Le Hello S, Hawkey J, Edwards DJ, Fourment M, Holmes EC. Genome-scale rates of evolutionary change in bacteria. *Microb. Genomics* 2016 Nov; 2(11). <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5320706/>.
- [5] Walker TM, Merker M, Knoblauch AM, Helbling P, Schoch OD, Werf MJvd, Kranzer K, Fiebig L, Kröger S, Haas W, Hoffmann H, Indra A, Egli A, Cirillo DM, Robert J, Rogers TR, Groenheit R, Mengshoel AT, Mathys V, Haanperä M, Soolingen Dv, Niemann S, Böttger EC, Keller PM, Avsar K, Bauer C, Bernasconi E, Borroni E, Brusin S, Dévis MC, Crook DW, Dedicoat M, Fitzgibbon M, Gagneux S, Geiger F, Guthmann JP, Hendrickx D, Hoffmann-Thiel S, Ingen Jv, Jackson S, Jaton K, Lange C, Stalder JM, O'Donnell J, Opota O, Peto TEA, Preiswerk B, Roycroft E, Sato M, Schacher R, Schulthess B, Smith EG, Soini H, Sougakoff W, Tagliani E, Utpatel C, Veziris N, Wagner-Wiening C, Witschi M. A cluster of multidrug-resistant *Mycobacterium tuberculosis* among patients arriving in Europe from the Horn of Africa: a molecular epidemiological study. *The Lancet Infect. Dis.* 2018 Jan; 0(0). [http://www.thelancet.com/journals/laninf/article/PIIS1473-3099\(18\)30004-5/abstract](http://www.thelancet.com/journals/laninf/article/PIIS1473-3099(18)30004-5/abstract).
- [6] Stevens EL, Timme R, Brown EW, Allard MW, Strain E, Bunning K, Musser S. The Public Health Impact of a Publically Available, Environmental Database of Microbial Genomes. *Front. Microbiol.* 2017; 8. <https://www.frontiersin.org/articles/10.3389/fmicb.2017.00808/full>.
- [7] Gardy J, Loman NJ, Rambaut A. Real-time digital pathogen surveillance — the time is now. *Genome Biol.* 2015; 16:155.
- [8] Beerenwinkel N, Däumer M, Oette M, Korn K, Hoffmann D, Kaiser R, Lengauer T, Selbig J, Walter H. Geno2pheno: estimating phenotypic drug resistance from HIV-1 genotypes. *Nucleic Acids Res.* 2003 Jul; 31(13):3850–3855. <https://academic.oup.com/nar/article/31/13/3850/2904197>.
- [9] Votintseva AA, Bradley P, Pankhurst L, del Ojo Elias C, Loose M, Nilgiriwala K, Chatterjee A, Smith EG, Sanderson N, Walker TM, et al. Same-day diagnostic and surveillance data for tuberculosis via whole-genome sequencing of direct respiratory samples. *J. clinical microbiology* 2017; 55(5):1285–1298.
- [10] Gardy JL, Loman NJ. Towards a genomics-informed, real-time, global pathogen surveillance system. *Nat. Rev. Genet.* 2018 Jan; 19(1):9. <https://www.nature.com/articles/nrg.2017.88>.
- [11] Katoh K, Misawa K, Kuma Ki, Miyata T. MAFFT: a novel method for rapid multiple sequence alignment based on fast Fourier transform. *Nucleic Acids Res.* 2002 Jul; 30(14):3059–3066. <http://nar.oxfordjournals.org/content/30/14/3059>.
- [12] Nguyen LT, Schmidt HA, von Haeseler A, Minh BQ. IQ-TREE: A Fast and Effective Stochastic Algorithm for Estimating Maximum-Likelihood Phylogenies. *Mol. Biol. Evol.* 2015 Jan; 32(1):268–274. <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4271533/>.
- [13] Price MN, Dehal PS, Arkin AP. FastTree 2—approximately maximum-likelihood trees for large alignments. *PLoS ONE* 2010; 5(3):e9490.
- [14] Stamatakis A. RAxML version 8: a tool for phylogenetic analysis and post-analysis of large phylogenies. *Bioinformatics* 2014 May; 30(9):1312–1313.
- [15] Drummond AJ, Suchard MA, Xie D, Rambaut A. Bayesian Phylogenetics with BEAUti and the BEAST 1.7. *Mol Biol Evol* 2012 Aug; 29(8):1969–1973.
- [16] To TH, Jung M, Lycett S, Gascuel O. Fast Dating Using Least-Squares Criteria and Algorithms. *Syst. Biol.* 2016 Jan; 65(1):82–97.
- [17] Volz EM, Frost SDW. Scalable relaxed clock phylogenetic dating. *Virus Evol.* 2017 Jul; 3(2). <https://academic.oup.com/ve/article/3/2/vex025/4100592>.
- [18] Sagulenko P, Puller V, Neher RA. TreeTime: Maximum-likelihood phylodynamic analysis. *Virus Evol.* 2018 Jan; 4(1). <https://academic.oup.com/ve/article/4/1/vex042/4794731>.
- [19] Hadfield J, Megill C, Bell SM, Huddleston J, Potter B, Callender C, Sagulenko P, Bedford T, Neher RA. Nextstrain: real-time tracking of pathogen evolution. *bioRxiv* 2017 Nov; p. 224048. <https://www.biorxiv.org/content/early/2017/11/22/224048>.
- [20] Neher RA, Bedford T. nextflu: real-time tracking of seasonal influenza virus evolution in humans. *Bioinformatics* 2015 Nov; 31(21):3546–3548.

- [21] Page AJ, Cummins CA, Hunt M, Wong VK, Reuter S, Holden MTG, Fookes M, Falush D, Keane JA, Parkhill J. Roary: rapid large-scale prokaryote pan genome analysis. *Bioinformatics* 2015 Nov; 31(22):3691–3693. <http://bioinformatics.oxfordjournals.org/content/31/22/3691>.
- [22] Ding W, Baumdicker F, Neher RA. panX: pan-genome analysis and exploration. *Nucleic Acids Res.* 2017; <https://academic.oup.com/nar/article/doi/10.1093/nar/gkx977/4564799/panX-pan-genome-analysis-and-exploration>.
- [23] Bradley P, Bakker Hd, Rocha E, McVean G, Iqbal Z. Real-time search of all bacterial and viral genomic data. *bioRxiv* 2017 Dec; p. 234955. <https://www.biorxiv.org/content/early/2017/12/15/234955>.
- [24] Argimón S, Abudahab K, Goater RJE, Fedosejev A, Bhai J, Glasner C, Feil EJ, Holden MTG, Yeats CA, Grundmann H, Spratt BG, Aanensen DM. Microreact: visualizing and sharing data for genomic epidemiology and phylogeography. *Microb. Genomics* 2016; 2(11). <http://mgen.microbiologyresearch.org/content/journal/mgen/10.1099/mgen.0.000093>.
- [25] Hadfield J, Croucher NJ, Goater RJ, Abudahab K, Aanensen DM, Harris SR. Phandango: an interactive viewer for bacterial population genomics. *Bioinformatics* 2018 Jan; 34(2):292–293. <https://academic.oup.com/bioinformatics/article/34/2/292/4212949>.
- [26] Bielejec F, Baele G, Vrancken B, Suchard MA, Rambaut A, Lemey P. Spread3: interactive visualization of spatiotemporal history and trait evolutionary processes. *Mol Biol Evol* 2016; 33(8):2167–2169.
- [27] Libin P, Vanden Eynden E, Incardona F, Nowé A, Bezenchek A, Group ES, Sönnernborg A, Vandamme AM, Theys K, Baele G. PhyloGeoTool: interactively exploring large phylogenies in an epidemiological context. *Bioinformatics* 2017; 33(24):3993–3995.
- [28] Allende C, Sohn E, Little C. Treelink: data integration, clustering and visualization of phylogenetic trees. *BMC Bioinforma.* 2015; 16(1):414.
- [29] Bedford T, Suchard MA, Lemey P, Dudas G, Gregory V, Hay AJ, McCauley JW, Russell CA, Smith DJ, Rambaut A. Integrating influenza antigenic dynamics with molecular evolution. *eLife* 2014 Feb; 3. <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3909918/>.
- [30] Neher RA, Bedford T, Daniels RS, Russell CA, Shraiman BI. Prediction, dynamics, and visualization of antigenic phenotypes of seasonal influenza viruses. *Proc. Natl. Acad. Sci. United States Am.* 2016 Mar; 113(12):E1701–1709.
- [31] Crisan A, McKee G, Munzner T, Gardy JL. Evidence-based design and evaluation of a whole genome sequencing clinical report for the reference microbiology laboratory. *PeerJ* 2018 Jan; 6:e4218. <https://peerj.com/articles/4218>.