

# Plague Phylodynamics and Phylogeography

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## Keywords

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- Plague
- Yersinia pestis
- Phylodynamics
- Phylogeography

# Introduction

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Plague has an impressively long and expansive history as a human pathogen. The earliest evidence of the plague bacterium *Yersinia pestis* comes from ancient DNA studies, dating its emergence to at least the Neolithic [1,2]. Since then, *Y. pestis* has traveled extensively due to ever-expanding global trade networks and the ability to infect a diverse array of mammalian hosts [3,4]. Few regions of the ancient and modern world remain untouched by this disease, as plague has an established presence on every continent except Oceania [5].

Accompanying this prolific global presence is unnervingly high mortality. The infamous medieval Black Death is estimated to have killed more than half of Europe's population [6]. This virulence can still be observed in the post-antibiotic era, where case fatality rates range from 22-71% [7]. As a result, plague maintains its status as a disease that is of vital importance to current public health initiatives.

This high priority disease status is unsurprising given that *Y. pestis* is a member of the Enterobacteriaceae family. This family includes enteric pathogens such as *Escherichia coli* and *Salmonella typhi* that are commonly transmitted by contaminated food and water. In comparison, the plague bacterium is unique among this family due to a striking difference in host habitat and transmission. *Y. pestis* commonly resides in the blood of its mammalian hosts and can be transmitted to new hosts through an infectious fleabite [8]. In addition to these tissues, the plague bacterium is also capable of colonizing parts of the mammalian immune system including the lymphatic and reticuloendothelial systems. The large diversity of media in which *Y. pestis* has adapted to colonize is particularly surprising given that it only recently (within the last 20,000 years) diverged as a clone of its parent species *Yersinia pseudotuberculosis* [9].

Despite a close genetic similarity between *Y. pestis* and *Y. pseudotuberculosis*, in which they share 97% gene identity, they differ widely in their transmission and pathogenicity [10]. Whereas *Y. pseudotuberculosis* causes gastrointestinal disease and is transmitted by the food-borne route, *Y. pestis* is primarily transmitted between mammalian hosts by fleas and causes septicemia, pneumonia, and lymphadenitis. Because of this apparent contradiction of genetic homogeneity and diverse phenotypes, an extensive body of research has formed to address how, when, and where, these epidemiological shifts occurred.

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## TO BE DONE:

- Introduce the topics phylodynamics and phylogeography and what is known so far.
- Introduce the major problem(s) and our objective(s).

## Objectives

1. Curate and critique publicly available *Y. pestis* genomes. Discuss how sampling bias drives our current understanding of plague.
2. To propose a nuanced phylodynamics model.
3. To critique interpretations drawn from phylogeographic approaches?

# Materials and Methods

## Workflow Overview

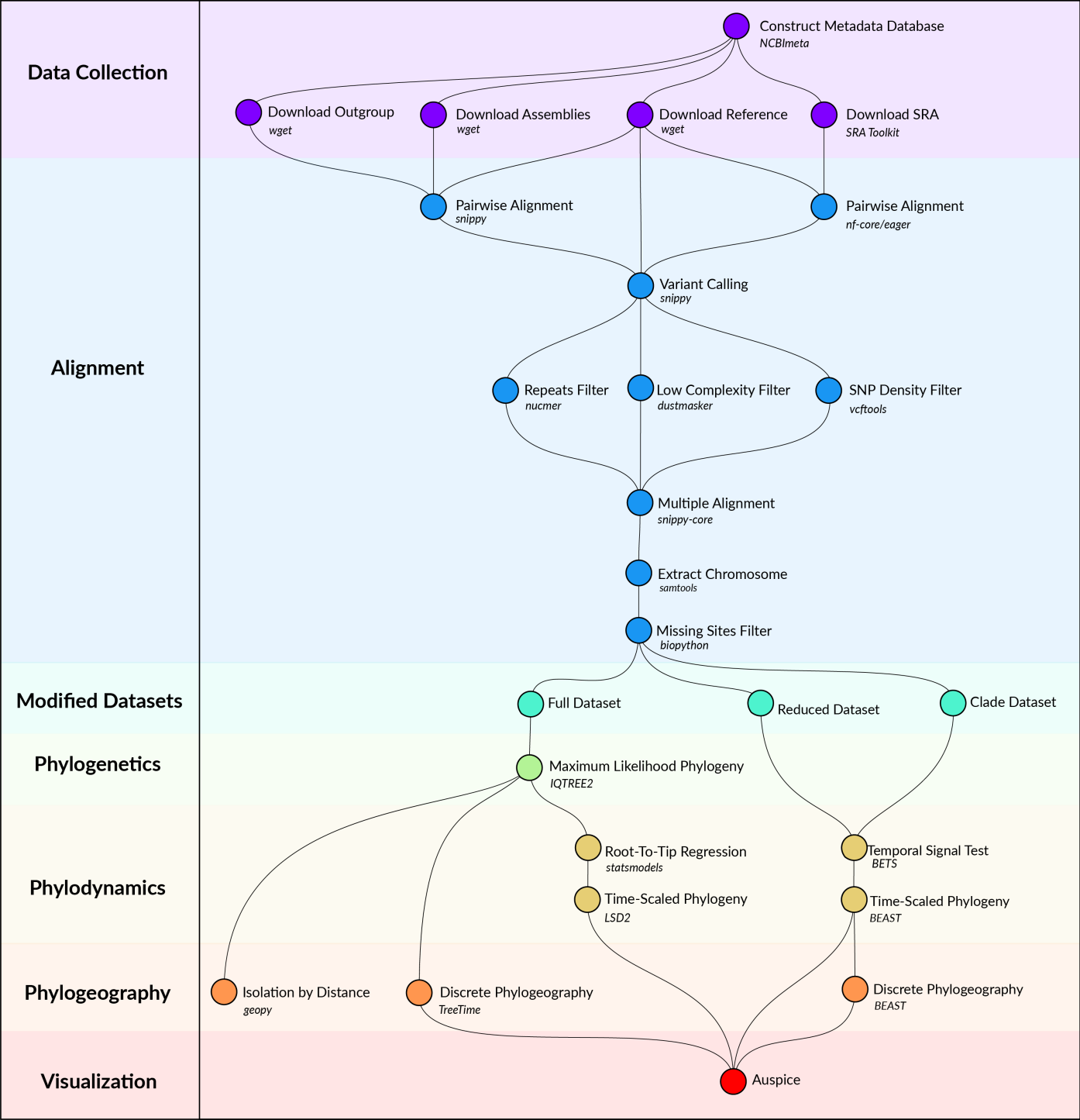


Figure 1: Computational methods workflow.

## Data Collection

*Y. pestis* genome sequencing projects were retrieved from the NCBI databases using NCBImeta [11]. 1657 projects were identified and comprised three genomic types:

- 586 modern assembled
- 184 ancient unassembled
- 887 modern unassembled

The 887 modern unassembled genomes were excluded from this project, as the wide variety of laboratory methods and sequencing strategies precluded a standardized workflow. In contrast, the 184 ancient unassembled genomes were retained given the relatively standardized, albeit specialized, laboratory procedures required to process ancient tissues. Future work will investigate computationally efficient methods for integrating more diverse datasets.

Collection location, collection date, and collection host metadata were curated by cross-referencing the original publications. Collection location was transformed to latitude and longitude coordinates using GeoPy and the Nominatim API for OpenStreetMap [12,13,14]. Coordinates were standardized at a sub-country resolution, taking the centroid of the parent province/state. Collection dates were standardized according to their year, and recording uncertainty arising from missing data and radiocarbon estimates. Collection host was the most diverse field with regards to precision, ranging from colloquial nomenclature (“rat”) to a genus species taxonomy (“*Meriones libycus*”). For the purposes of this study, collection host was recorded as *Human*, *Non-Human*, or *Not Available*, given the inability to differentiate non-human mammalian hosts.

Genomes were removed if no associated date or location information could be identified in the literature, or if there was documented evidence of laboratory manipulation.

Two additional datasets were required for downstream analyses. First, *Y. pestis* strain CO92 (GCA\_000009065.1) was used as the reference genome for sequence alignment and annotation. Second, *Yersinia pseudotuberculosis* strains NCTC10275 (GCA\_900637475.1) and IP32953 (GCA\_000834295.1) served as an outgroup to root the maximum likelihood phylogeny.

## Alignment

Modern assembled genomes were aligned to the reference genome using Snippy, a pipeline for core genome alignments [15]. Modern genomes were removed if the number of sites covered at a minimum depth of 10X was less than 70% of the reference genome.

Ancient unassembled genomes were downloaded from the SRA database in FASTQ format using the SRA Toolkit [16]. Pre-processing and alignment to the reference genome was performed using the nf-core/eager pipeline, a reproducible workflow for ancient genome reconstruction [17]. Ancient genomes were removed if the number of sites covered at a minimum depth of 3X was less than 70% of the reference genome. It is a typical approach to relaxed coverage thresholds for ancient genomes relative to modern genomes (CITE). This threshold is commonly used, and aims to strike a balance between variant confidence and sample representation (CITE).

A multiple sequence alignment was constructed using the Snippy Core module of the Snippy pipeline. The output alignment was filtered to only include chromosomal variants and to exclude sites that had more than 5% missing data.

## Modified Datasets

To investigate the influence of between-clade variation in substitution rates, the multiple alignment was separated into the major clades of *Y. pestis*, which will be referred to as the *Clade* dataset. Clade and subclade labeling was derived from the five-branch population structure accompanied by a biovar abbreviation ([18]). Only one modification was made, in that the subclade associated with the Plague of Justinian (0.ANT4) was considered to be a distinct clade from its parent (0.ANT) due to its geographic, temporal, and ecological uniqueness. In total, 12 clades were considered and are described in Table 2.

To improve the performance and convergence of Bayesian analysis, a subsampled dataset was constructed and will be referred to as the *Reduced* dataset. Clades that contained multiple samples drawn from the same geographic location and the same time period were reduced to one representative sample. The sample with the shortest terminal branch length was prioritized, to diminish the influence of uniquely derived mutations on the estimated substitution rate. An interval of 25 years was identified as striking an optimal balance, resulting in 200 representative samples.

## Phylogenetics

Model selection was performed using Modelfinder which identified the K3Pu+F+I model as the optimal choice based on the Bayesian Information Criterion (BIC) [19]. A maximum-likelihood phylogeny was then estimated across 10 independent runs of IQTREE [20]. Branch support was evaluated using 1000 iterations of the ultrafast bootstrap approximation, with a threshold of 95% required for strong support [21].

## Phylogenetics

### Temporal Signal

To explore the degree of temporal signal present in the data, two categories of tests were performed. The first was a root-to-tip (RTT) regression on collection date. This linear model is a simple approach to explore whether the data follows a strict clock model. Uncertainty in the model parameters, namely the mean substitution rate and tMRCA, were estimated using 1000 iterations of the non-parametric bootstrap on the residuals.

While RTT is a practical approach, it has two main limitations: 1) No rate variation is accounted for, and 2) The data are not independent observations due to shared internal branch lengths. Therefore to complement this approach, a bayesian evaluation of temporal signal (BETS) was performed.

### Rate Variation

A bayesian timetree was estimated using ... as implemented in BEAST.

### Time Tree

A maximum-likelihood timetree was estimated using a least-squares approach as implemented in LSD2 [22]. Rate variation was modeled using a lognormal relaxed clock with default parameters for the mean (1.0) and the standard deviation (0.2). The outgroup *Y. pseudotuberculosis* was used to root the tree and then subsequently removed.

Note: I'm still pondering the best choice of parameters for the LSD2 relaxed clock.

## Phylogeography

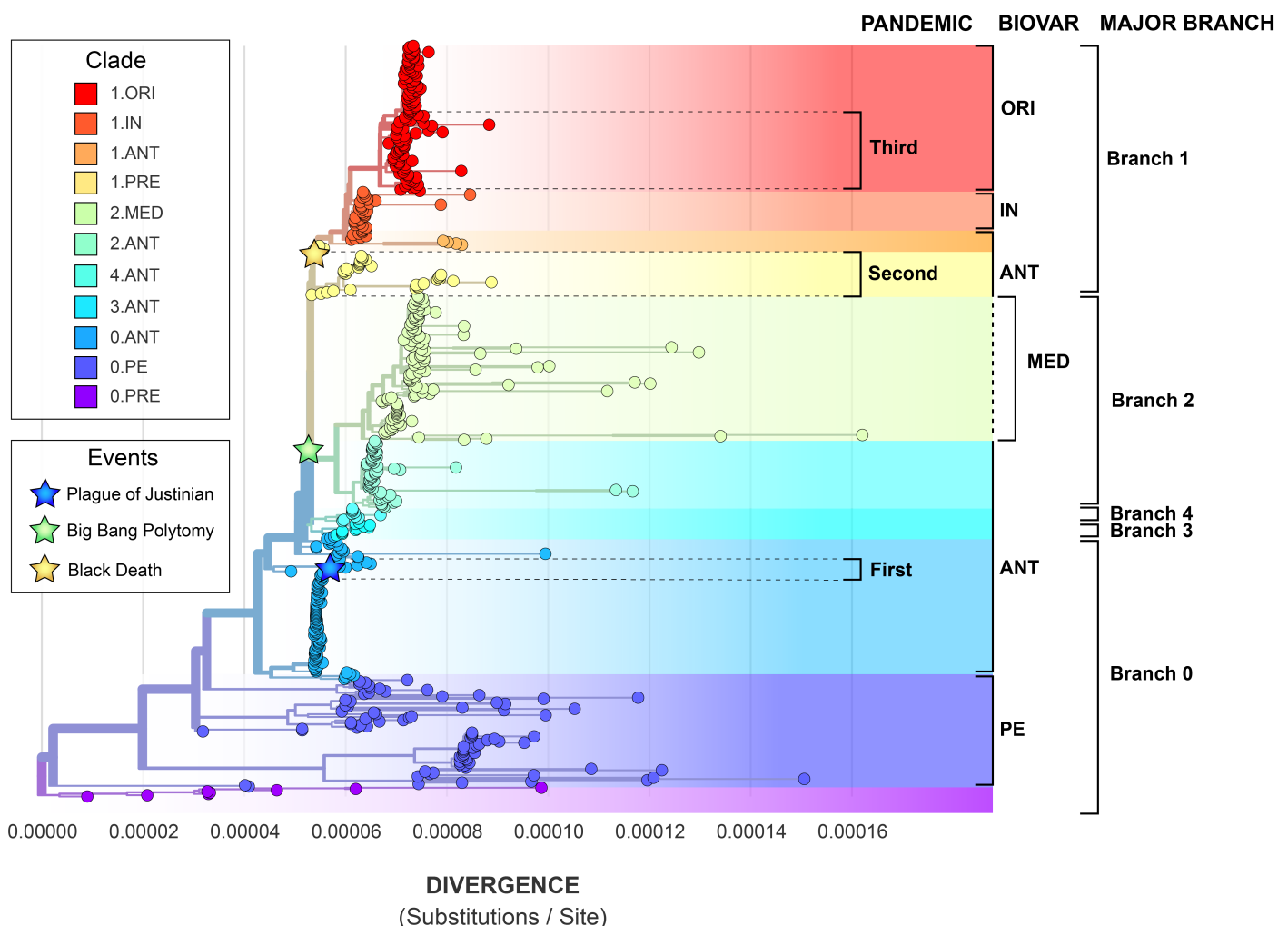
Geographic location was modeled as a discrete state with transitions following a GTR migration model as implemented in TreeTime [\[23\]](#).

# Results

## Population Structure

A maximum-likelihood phylogeny was estimated from 603 genomes (600 *Y. pestis* isolates, 1 *Y. pestis* reference, and 2 *Y. pseudotuberculosis* outgroup taxa). A total of 26,282 Single Nucleotide Polymorphisms (SNPs) were identified with 17,333 sites present in at least two genomes and 16,370 sites present in only one. Following removal of the outgroup taxa, 10,249 sites remained with 3,844 sites shared by at least two genomes and 6,405 sites in only one.

The global population structure of *Y. pestis* is organized according to a vast array of historical, ecological, biochemical, and molecular characteristics (Figure 2). Arguably the most important event, in terms of phylogenetic structure, is the Big Bang Polytohy from which arose four monophyletic clades: Branches 1-4. All lineages that diverged prior to this multifurcation are grouped into Branch 0.



**Figure 2:** The maximum-likelihood tree, constructed from 10,249 SNPs, depicts the global population structure of *Y. pestis*.

Each major branch is further subdivided into biovars according to metabolic properties [24,25]. The oldest isolates of plague date to the Late Neolithic Bronze Age following a divergence that pre-dates all known modern lineages [26]. In the absence of metabolic evidence, this clade is designated 0.PRE. The *microtus* biovar, alias *pestoides* (PE), is also a basal clade found in Branch 0 and while it is typically avirulent in humans, sporadic cases can occur (CITE).

The other inhabitant of **Branch 0** is biovar *antiqua* ( **ANT** ) which is the ancestral state prior to the Big Bang Polytomy and continues to be isolated from all major branches. **Branch 1** encapsulates a transition from *antiqua* ( **ANT** ) through the *intermedium* biovar ( **IN** ) and into *orientalis* ( **ORI** ).

**Branch 2** includes the transition from *antiqua* ( **ANT** ) to *medievalis* ( **MED** ), which was once hypothesized to be associated with the Medieval Black Death, but is now known to be a distinct emergence. **Branch 3** and **Branch 4** are exclusively composed of *antiqua* ( **ANT** ) strains.

## Pandemics of Plague

Additionally, several lineages of plague have been associated with historically documented plague pandemics. The First Pandemic (6th - 8th century CE) began with the Plague of Justinian and proceeded to devastate the Byzantine Empire of the Mediterranean world (CITE). A unique emergence of *Y. pestis* within the *antiqua* biovar of **Branch 0** ( **0.ANT4** ) is thought to derive from this pandemic given spatiotemporal overlap of the skeletal remains from which this lineage was retrieved [27,28].

Similarly, variants of the the *antiqua* biovar of **Branch 1** are thought to have given rise to the Second Pandemic. This well-documented pandemic began with the infamous Black Death and swept across most of Eurasia from the 14th to 19th centuries (CITE). The divergence of ancient *Y. pestis* dated to this time period pre-dates all other **Branch 1** lineages, with several samples placed directly at the base of **Branch 1**. To mark this unique phylogenetic positioning, this clade is designated **1.PRE** rather than a subclade of **1.ANT**.

The third documented pandemic of plague, alias the *Modern Pandemic*, spread globally from the end of the 18th Century and until the mid-20th Century. There is little dispute that a new lineage of plague emerging from **Branch 1** as biovar *orientalis* ( **1.ORI** ) was the causative agent of this pandemic. While the World Health Organization (WHO) declared the third pandemic over in 1950 (CITE)), this lineage continues to re-emerge to cause recent epidemics such as the 2010 plague in Peru and the Madagascar Outbreaks of 2017.

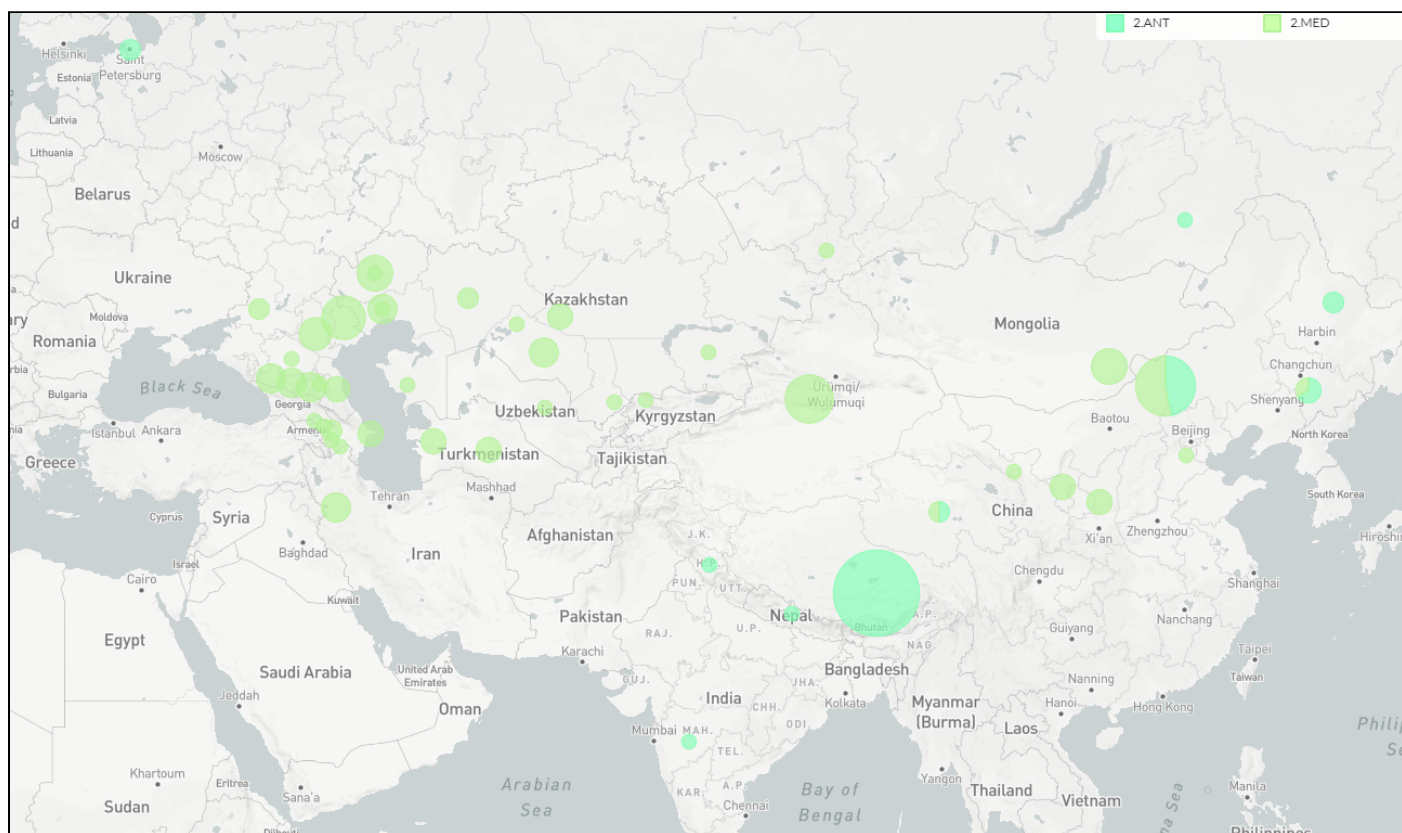
## The Three Pandemic Problem

While the pandemic clade nomenclature provides an excellent foundation for historical discussion, there are several problems with this system. First is the growing awareness of the spatiotemporal overlap of the Second and the Third Pandemic. Previously, the temporal extents of these events were mutually exclusive, dating from the 14th-18th century, and the 19th-20th century respectively. Recent historical scholarship has contested this claim, and demonstrated that these constraints are a product of a Eurocentric view of plague (CITE Nukhet). The Second Pandemic is now known to have extended into the 19th Century in parts of the Ottoman Empire, with the latest epidemics dating to 1819 (CITE). Similarly, the Third Pandemic is now hypothesized to have begun as early as 1772 in southern China [29]. It remains unclear where to draw the distinction, if it even exists, between the Second and Third Pandemic.

Another limitation of the pandemic nomenclature is the complete disconnection of **Branch 2** to any pandemic-related events. This is surprising given that several criteria of a pandemic pathogen are fulfilled by **Branch 2** lineages, namely extensive spread and virulence. **Branch 2** genomes of *Y. pestis* have been collected from all throughout Eurasia, stretching from the Caucasus, to India, and to eastern China (Figure 3). Furthermore, clade 2.MED was demonstrated to have the highest spread velocity of any *Y. pestis* clade [30]. And finally, while **Branch 2** isolates are not historically linked to the Third Pandemic Proper, clade 2.MED has been implicated in numerous modern plague outbreaks.



As historical plague scholarship extends beyond the bounds of Western Europe, the traditional narrative of three pandemic plague becomes unstable.



**Figure 3:** The geographic distribution of *Y. pestis* Branch 2 .

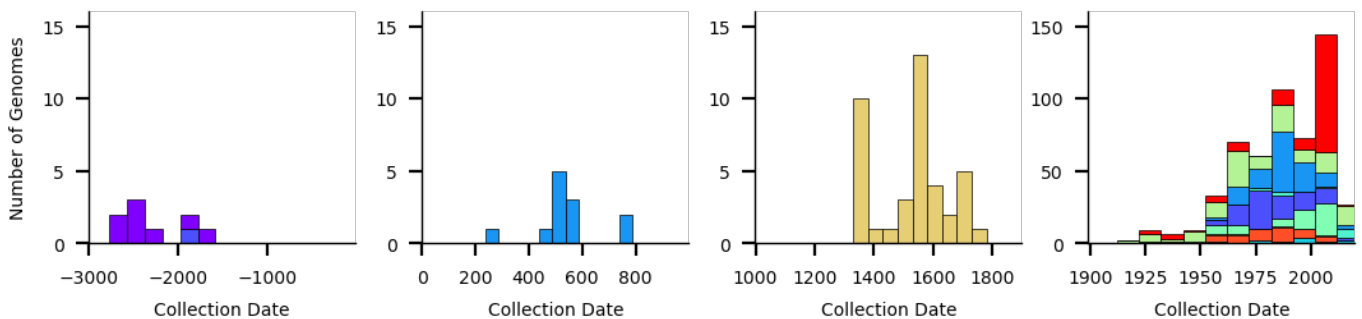
## Time

How has the known diversity of plague changed over time?

In addition to the biovar nomenclature is the categorization of plague according to collection date. Historical strains of plague isolated prior to the 20th century cluster into three time periods: the Late Neolithic Bronze Age (LNBA), Roman Period/Late Antiquity(?), and the Medieval Period (4). For example, all strains of *Y. pestis* isolated in the Roman period group together in Branch 0 and form a distinct subclade within 0.ANT. Similarly, all Medieval strains of *Y. pestis* cluster together in Branch 1 as clade 1.PRE. The LNBA is a notable exception as mixing between clades 0.PE and 0.PRE is observed.

However, collection date and collection location are highly confounded as ancient DNA sampling strategies have predominantly targeted Western Europe. Due to this sampling bias, it is challenging to evaluate whether multiple strains of plague have co-occurred in human populations in the past.

An additional observation is that the temporal sampling strategy of genomic data reflects greater interest in *Y. pestis* as a historical pathogen, rather than a public health threat to modern humans. One example of this is that the Medieval Plague in Western Europe (Clade 1.PRE) has more representative samples than all of the African continent (Clade 1.ANT). Sequencing initiatives are greatly needed that shift the balance away from Eurocentrism and encompass a greater diversity of affected populations.



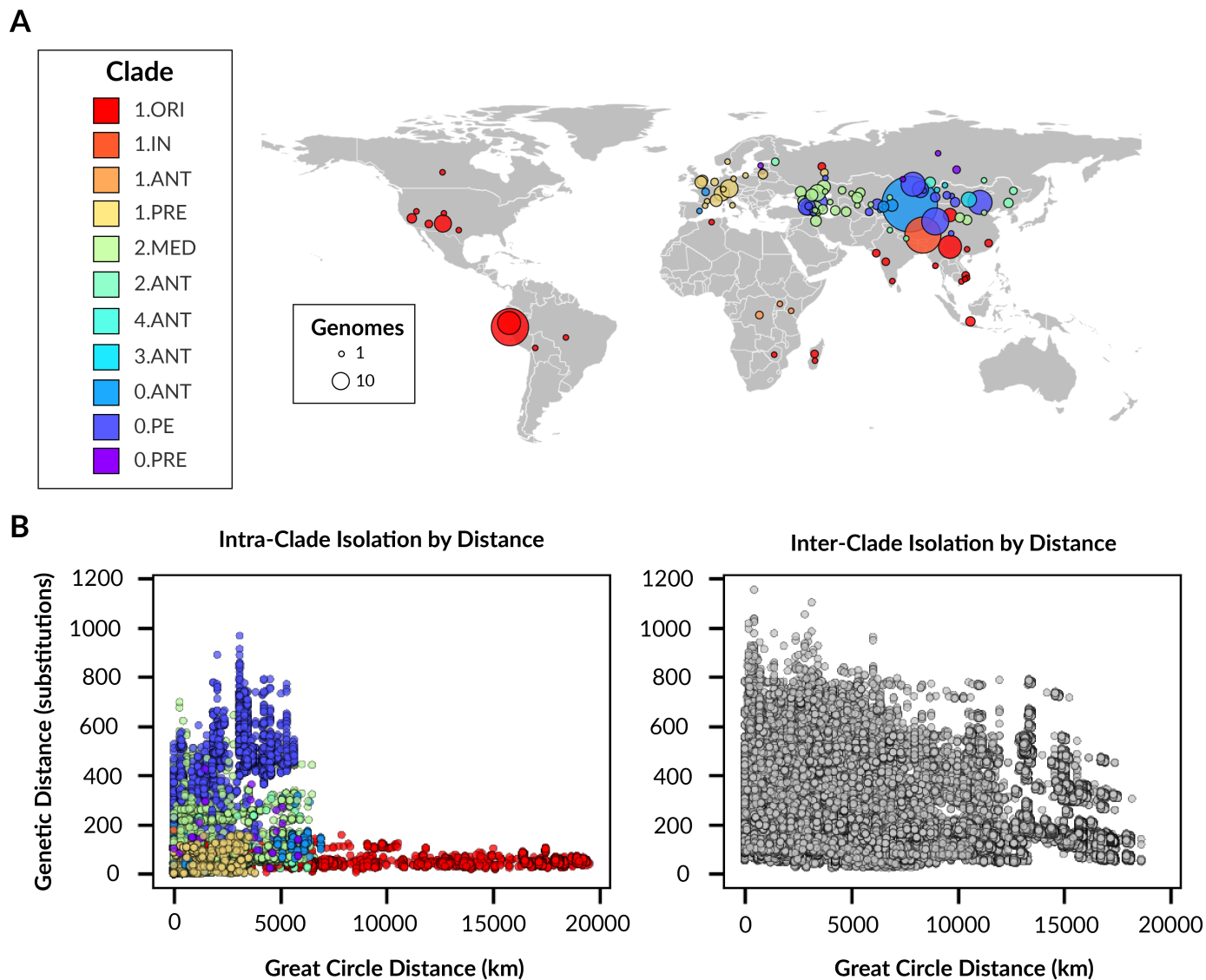
**Figure 4:** Temporal distribution of *Y. pestis* genomes.

## Geography

The geographic distribution of *Y. pestis* reflects a complex dispersal history (Figure 5 .B). Many regions have been colonized by diverse strains of *Y. pestis*. This diversity can be contemporaneous, such as endemic foci in the Caucasus and Western China (Clade 0.PE). Alternatively, this diversity may occur over multiple centuries through distinct re-introductions and extinctions, as seen in the historical epidemics of Europe (Clades 0.ANT and 1.PRE). In these examples, a relatively large amount of genetic diversity appears in a small geographic range. In contrast, regions such as the Americas have been colonized by a single strain of *Y. pestis* (Clade 1.ORI) which shows a relatively small amount of genetic diversity over a tremendously large geographic range.

An important consideration is that the geographic sampling strategy of *Y. pestis* genomes (Figure 5 .A) does not reflect the known distribution of modern plague [30], let alone historical pandemics. Nor does it adequately characterize the most heavily affected regions of the world, namely Madagascar and the Democratic Republic of the Congo [5]. The over-sampling of East Asia has been previously described by 31 and considerably drives the hypothesis that *Y. pestis* originated in China [18,32]. This once established hypothesis is now in contention, as the most basal strains of *Y. pestis* (Clades 0.PRE and 0.PE) have been isolated from all across Eurasia.

The geographic distribution of *Y. pestis* genomes reflects the *sampling history* more so than the *natural history* of the disease.



**Figure 5:** Spatiotemporal distribution of *Y. pestis* genomes. **A:** Geographic distribution, **B:** Isolation by distance as a function of geographic distance and genetic distance.

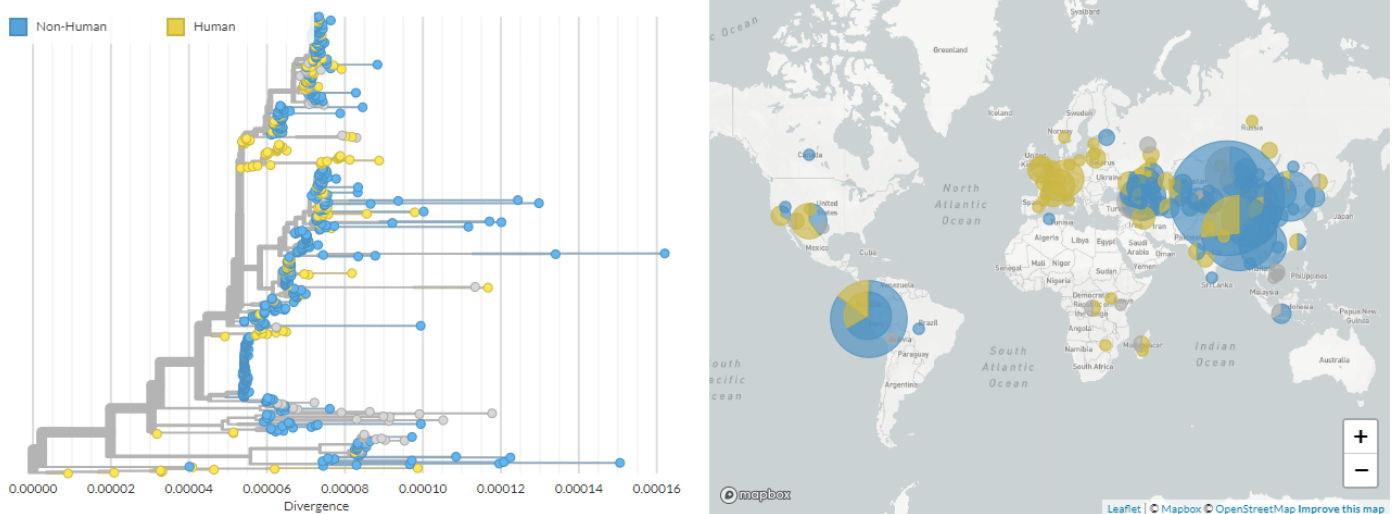
## Hosts

Clades of *Y. pestis* are additionally defined by virulence in particular hosts. For example, the Pestoides clade is frequently avirulent in humans, although sporadic cases of human plague do still occur (CITE). In general, all branches of *Y. pestis* are capable of causing plague in humans and the species barrier between wild rodents and human populations is crossed frequently (FIGURE).

While clades of ancient *Y. pestis* are exclusively associated with humans, this is more likely due to the sampling strategies of ancient DNA studies which have prioritized human skeletal remains over zooarchaeological remains. Given that no other clades across the *Y. pestis* phylogeny show a specificity for human hosts, ... isolate aDNA *Y. pestis* from rats.

Plague can cause disease in humans at any time, and from anywhere. There are virtually no lineages that are “safe” for humans. Thus plague as a disease is treated with exceptional caution.

No lineage of modern plague has been observed to exclusively infect humans (?) and thus ... attention to multi-host ecology.



**Figure 6:** Distribution of human vs. non-human samples.

## Pandemics

Finally, *Y. pestis* can be structured according to historically documented human pandemics. Ancient isolates of *Y. pestis* that date to the Late Antiquity are hypothesized to be associated with First Pandemic which includes the Plague of Justinian. Medieval isolates have been linked to the Second Pandemic and the Black Death based on spatiotemporal overlap.

While the three-pandemic nomenclature is undeniably useful in discourse, it does pose several problems...

Are the Second and Third Pandemics genetically separate? If so, what does this mean for clade 1.ANT? Is it Pandemic 2.5? Why are there no pandemics attribute to branch 2? The two clades within branch 2 stretch across Asia, from the Caucasus to Eastern China to South Asia.

# Phylodynamics

- How does the evolutionary rate of *Y. pestis* compare to other human pathogens?

*Y. pestis* has one of the slowest substitution rates observed in a bacterial pathogen (Table tbl:table\_bacterial\_rate\_comparison). Given the tremendous variation observed in modern plague ecology, it is surprising that the evolutionary rate does not reflect this need to rapidly adapt to changing environments. However, this slow rate makes perfect sense when viewed in the context of rate *time-dependency*, wherein the observed substitution rate decreases as the sampling time frame increases.

**Table 1:** Substitution rates of bacterial pathogens.

Organism	Disease	Substitution Rate (subs/site year <sup>-1</sup> )	Sampling Time (years)	Study
<i>Yersinia pestis</i>	Plague	1.42 x 10 <sup>-8</sup>	4687	This Study
<i>Mycobacterium leprae</i>	Leprosy	1.56 x 10 <sup>-8</sup>	1993	[33]
<i>Mycobacterium tuberculosis</i>	Tuberculosis	5.39 x 10 <sup>-8</sup>	895	[33]
<i>Neisseria meningitis</i>	Meningitis	6.05 x 10 <sup>-8</sup>	59	[33]
<i>Salmonella enterica</i>	Typhoid	7.60 x 10 <sup>-8</sup>	84	[33]
<i>Pseudomonas aeruginosa</i>	Pneumonia	3.36 x 10 <sup>-7</sup>	35	[33]

- Lack of temporal signal in Wagner et al. (2014)
- *Y. pestis* has substantial rate variation, and estimates in the literature have ranged from 1E-8 to 2.85E-8 [18,34]

The mean substitution rate observed here falls within this described range. Using a strict clock model for all of *Y. pestis* likely underestimates the mean substitution rate (1E-8) .

- A root-to-tip regression is a poor measure of temporal signal in *Y. pestis*.

This is due to several known limitations, namely the underlying assumption of strict clock-behavior and the non-independence of data points [35].

- BETS can detect temporal signal when the RTT R<sup>2</sup>.
- Using a strict clock model for all of *Y. pestis* underestimates the mean substitution rate.
- Several clades of *Y. pestis* demonstrate strong evidence of temporal signal.
- All pandemic clades have temporal signal and the least rate variation. The Big Bang Polytoimy is surrounded by clades with varying degrees of temporal signal. The branches surrounded the Big Bang Polytoimy [18] show some of the most dramatic rate acceleration to accomodate.

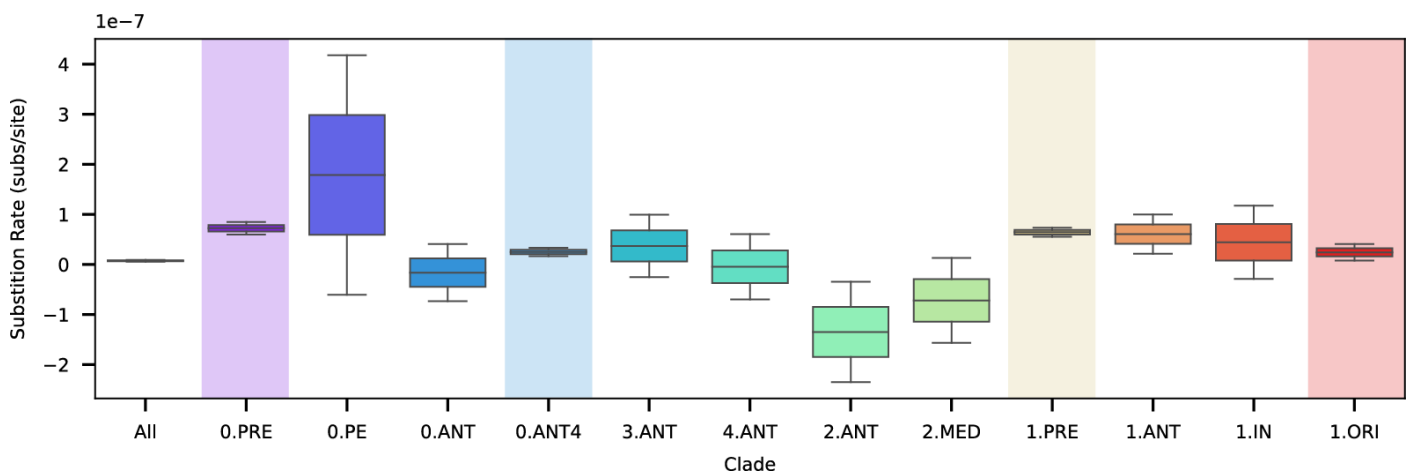
## Rate Variation

- A Root to Tip Regression on collection date confirms this, as the Coefficient of Determination (R<sup>2</sup> ) is 0.09, revealing a poor fit to a simple linear model (Table 2).
- To some extent, this variation can be explained by examining the clades in isolation (Figure 7).

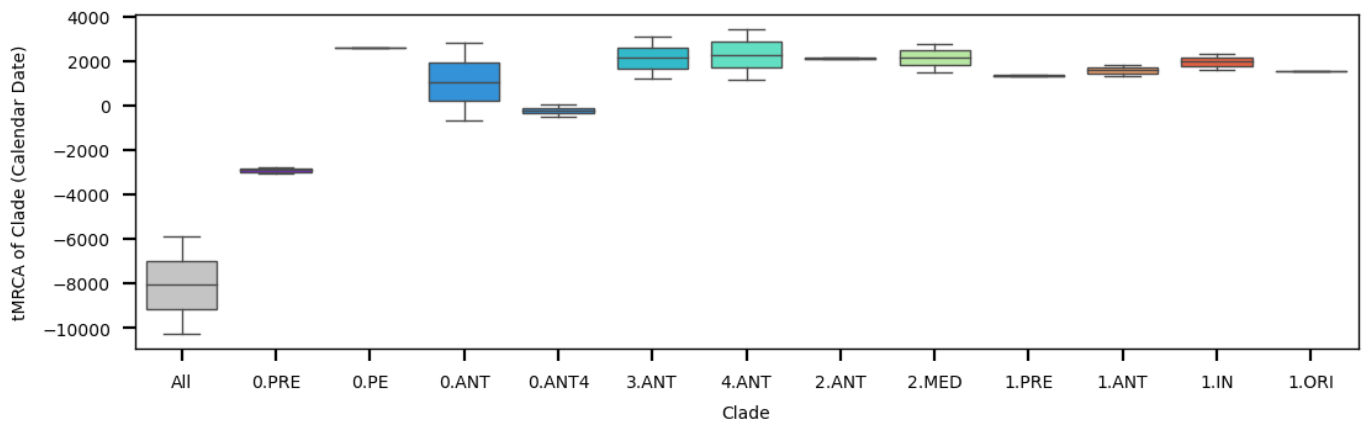
- Finding an appropriate evolutionary model is key to estimating historic events, like clade emergence (Figure 8).

**Table 2:** Temporal signal statistics by clade based on a root-to-tip linear regression. \* indicates a significant p-value or bayes factor.

Branch	Clade	Origin	RTT R <sup>2</sup>	RTT p-value	Strict Clock BF	Relaxed Clock BF	Clock Model
All	All	Ancient, Modern	0.09	3.81E-14*	–	–	–
0	0.PRE	Ancient	0.91	1.53E-04*	83.0*	-2.9	Strict?
0	0.PE	Modern	0.01	2.25E-01	-82.1	12.4*	Relaxed
0	0.ANT4	Ancient	0.66	7.84E-04*	5.3*	5.9*	Relaxed
0	0.ANT	Modern	-0.01	7.35E-01	-2.3	-6.5	None
1	1.ANT	Modern	0.45	2.03E-01	8.9*	12.6*	Relaxed
1	1.IN	Modern	0.0	3.24E-01	-3.9	-10.2	None
1	1.ORI	Modern	0.04	1.32E-02*	29.6*	35.7*	Relaxed
1	1.PRE	Ancient	0.76	1.68E-13*	10.1*	44.1*	Relaxed
2	2.ANT	Modern	0.05	5.96E-02	-20.8	-13.7	None
2	2.MED	Modern	0.01	1.86E-01	–	–	–
3	3.ANT	Modern	-0.04	4.39E-01	-9.6	-11.4	None
4	4.ANT	Modern	-0.11	8.80E-01	-2.9	3.7*	Relaxed



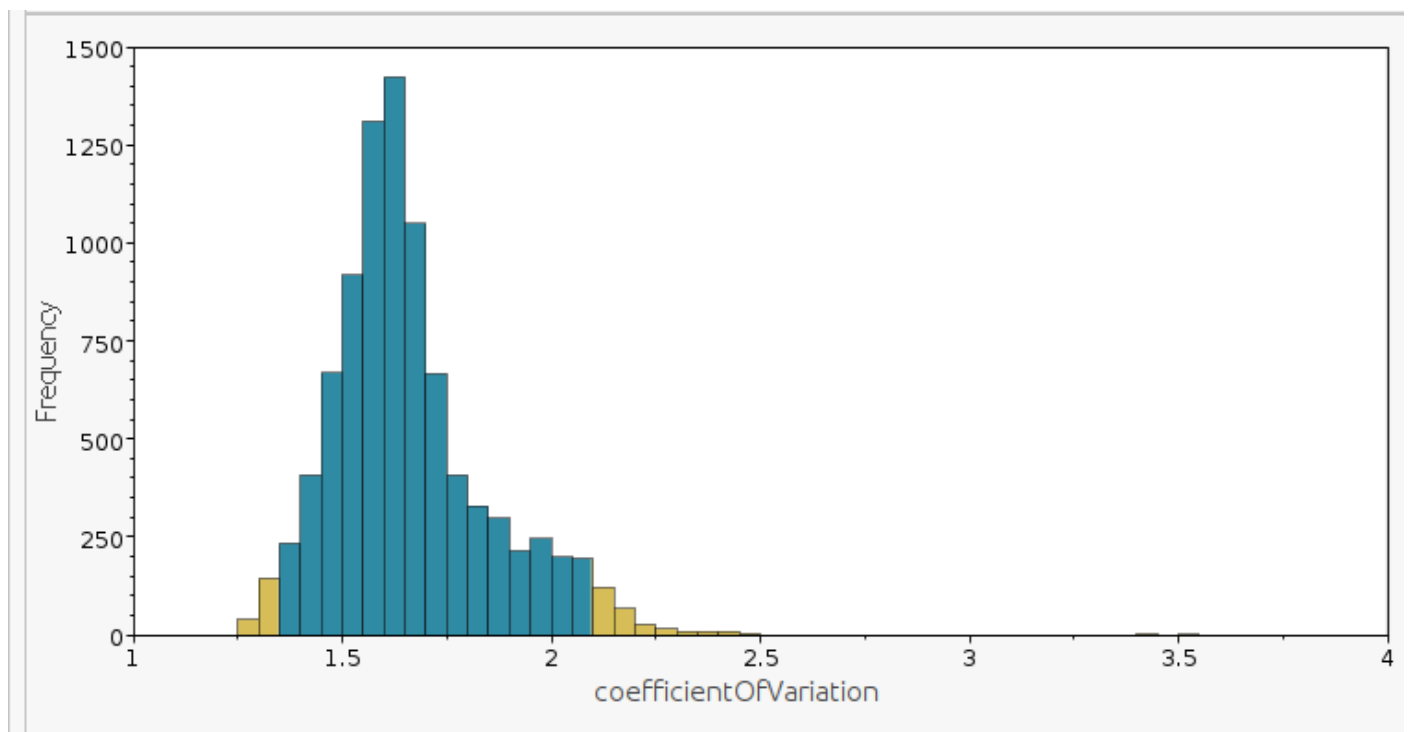
**Figure 7:** Substitution rate uncertainty by clade based on a root-to-tip linear regression. Highlighted clades are associated with historically documented pandemics.



**Figure 8:** Substitution rate uncertainty by clade based on a root-to-tip linear regression. Highlighted clades are known to be associated with human pandemics.

## Clock Model

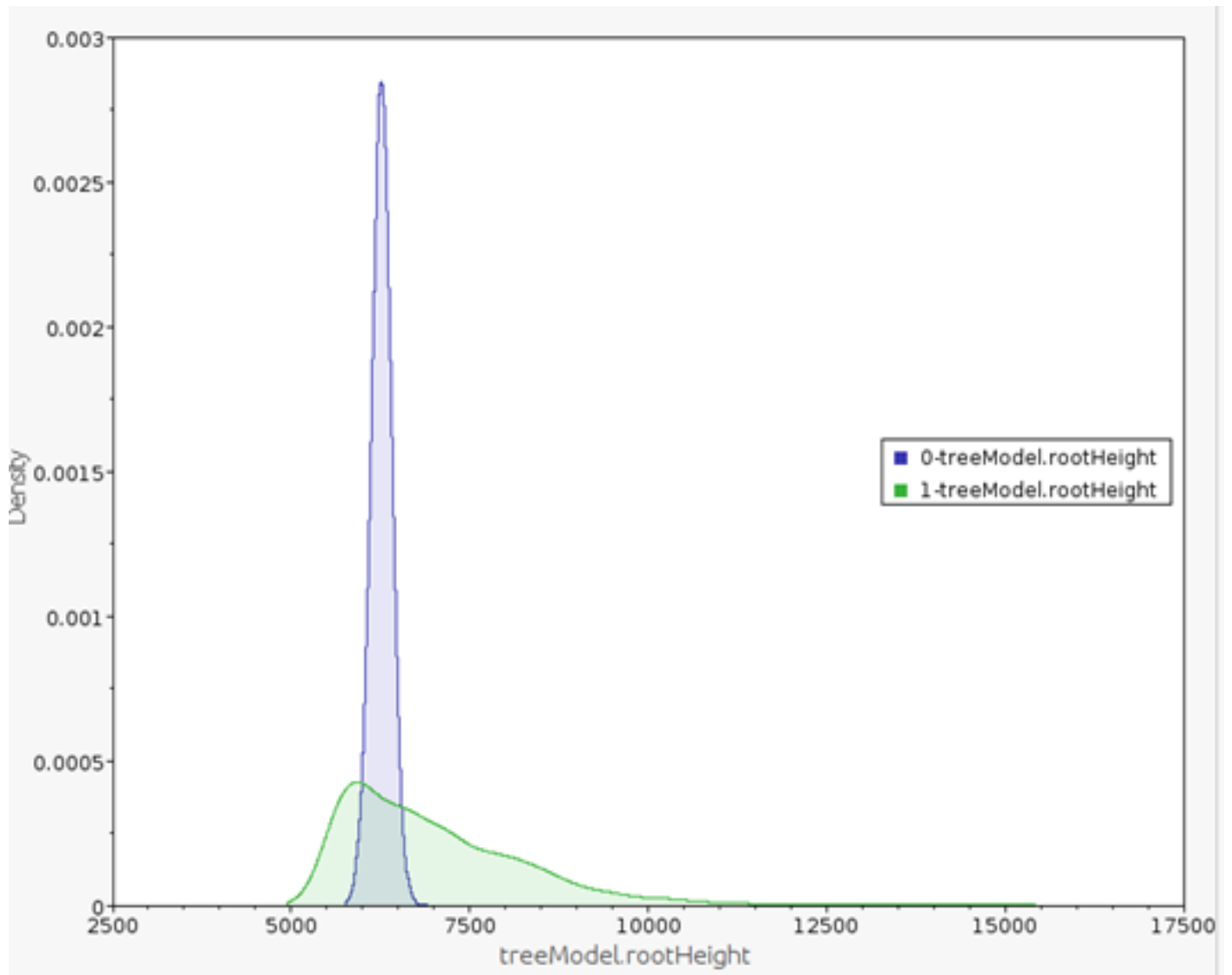
- Relaxed clock MCMC runs produce a high Coefficient of Variation indicating a relaxed model is favored over a strict model (Figure 9). However, these runs do not converge, suggesting there is too much rate variation to confidently estimate key parameters such as the mean Substitution Rate or tMRCA.



**Figure 9:** Coefficient of variation.

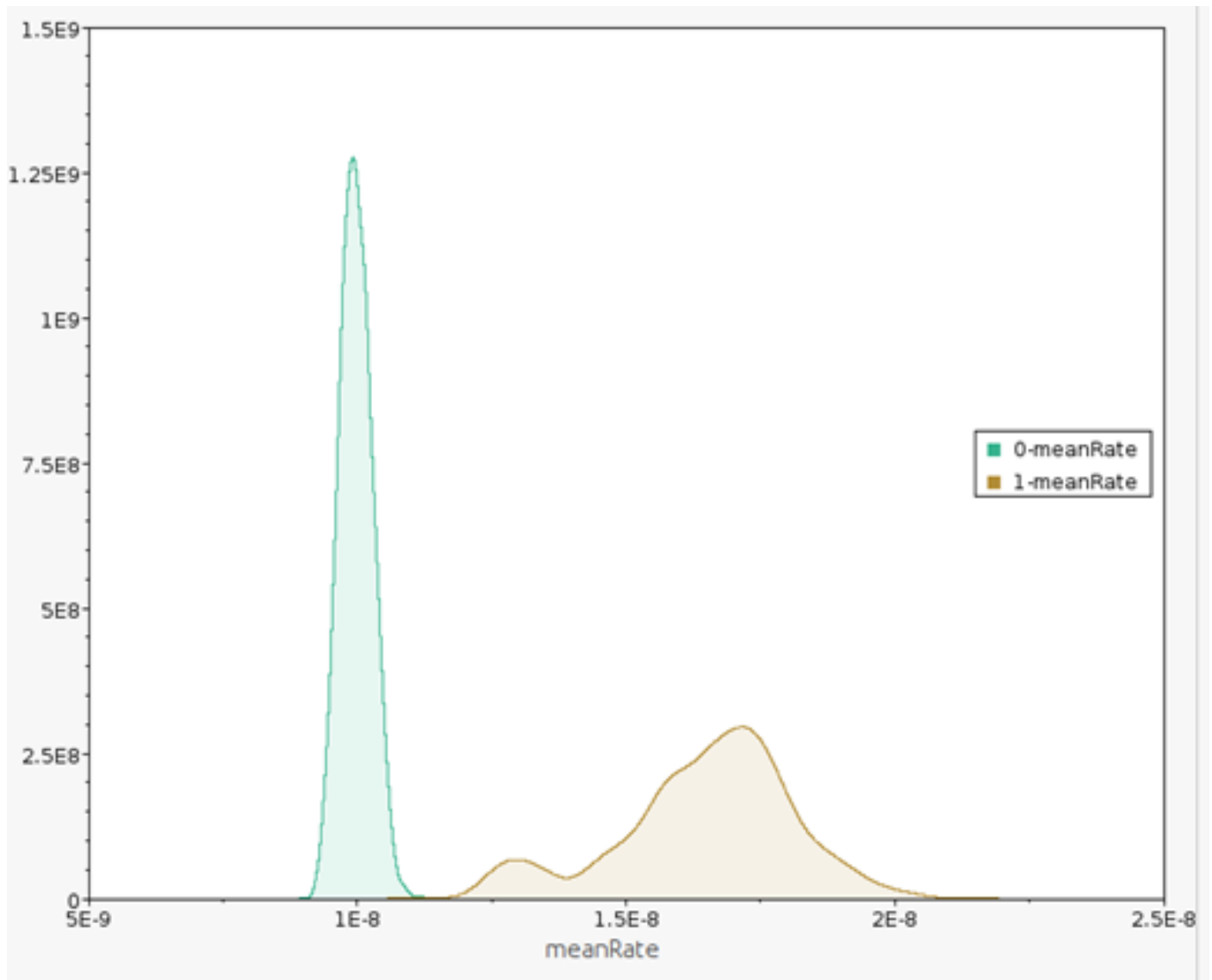
- A strict clock and relaxed clock have overlapping distributions with similar peaks for the Tree Height (blue: strict, green: relaxed) (Figure 10).





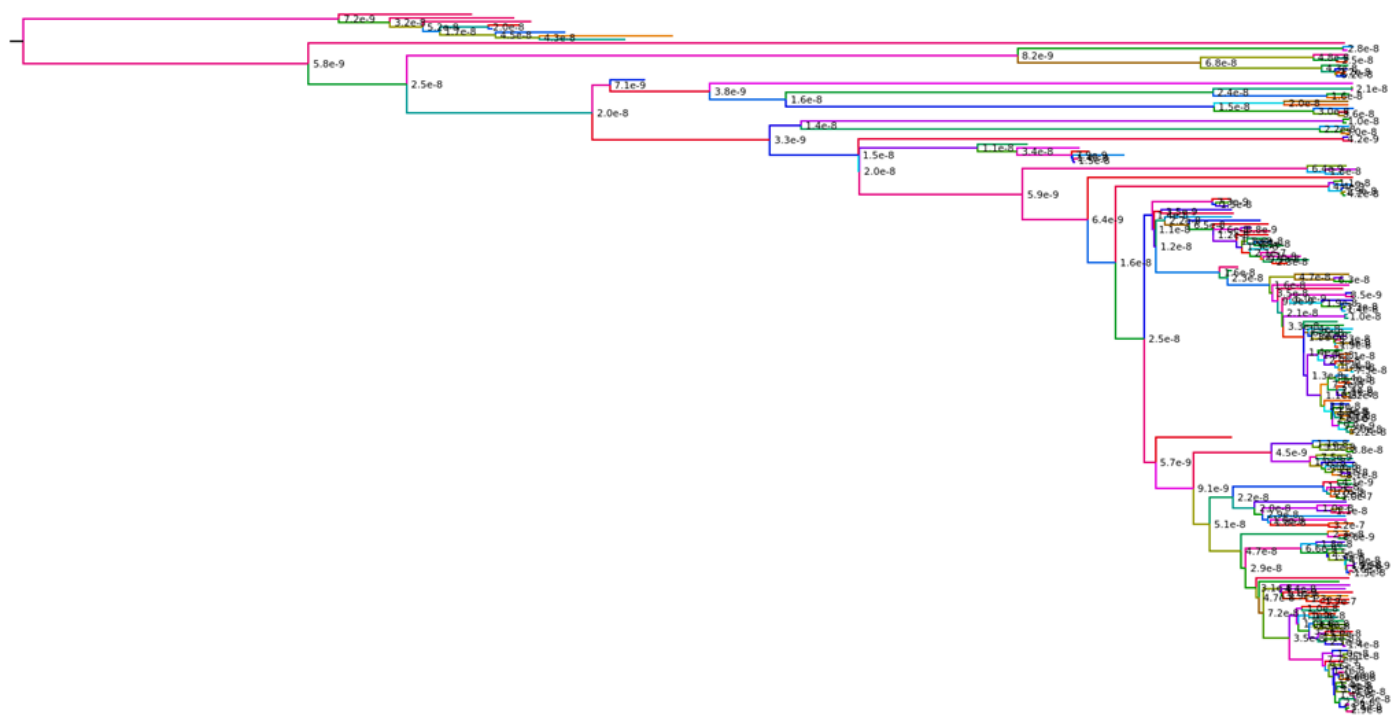
**Figure 10:** Tree height comparison.

- When estimating a Substitution Rate for all of *Y. pestis*, a [[Clock Model | strict clock]] and relaxed clock produce different estimates (green: strict, orange: relaxed) (Figure [11](#)).



**Figure 11:** Substitution rate comparison.

- There doesn't appear to be clustering of rates. Branches with high rates are next to those with low rates (Figure [12](#)).



**Figure 12:** Time tree colored by rate.

## Phylogeography

# Conclusion

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# Appendix

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