



# Emergence and spread of ancestral *Yersinia pestis* in Late-Neolithic and Bronze-Age Eurasia, ca. 5,000 to 2,500 y B.P.

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Evolutionary history of any living organism is as fascinating as it is complex. The causative agent of plague, the bacterium *Yersinia pestis*, is no exception. Having diverged from the enteropathogen *Yersinia pseudotuberculosis*, ancestral strains of *Y. pestis* spread all over Late-Neolithic Eurasia. In their study, Andrades Valtueña et al. (1) present a tour de force by reporting 17 new prehistoric *Y. pestis* genomes from Eurasian human burials (adding to 13 previously published) (1–7). Furthermore, their work, together with previously published data, lays the foundations for a new classification of *Y. pestis* strains and broadens our insight into the dynamics of emergence and spread of *Y. pestis* in prehistoric Eurasia.

Of the ancient genomes, the authors classify the two oldest (previously published) genomes as the “preLNBA–” (pre-Late Neolithic/Early Bronze Age) lineage and 26 others as the “LNBA–” lineage (1). PreLNBA– genomes are from Latvia and Sweden, dated, respectively, to 5,300 to 5,050 and 5,040 to 4,867 y B.P. The low number of reported preLNBA– genomes suggests that these lineages may have died out several centuries after their rise. In contrast to preLNBA– lineages, LNBA– lineages were present over a wide geographic area (from Lake Baikal to central Europe) and existed for at least 2,500 y—from ca. 5,100 y B.P. until 2,736 to 2,457 y B.P. (the date of the latest genome).

The data reported by Andrades Valtueña et al. (1) indicate that the LNBA– lineage had a different evolutionary story compared to later lineages, such as “Branch 1” causing the second and third plague pandemics, characterized by phylogenetic and global focal diversity (8–10). Conversely, the LNBA– lineage is characterized by a genetic monophyly, lacking distinct subbranches, containing many evolutionary dead ends, and showing no correlation between genetic and geographic distance (1). Interestingly, the two earliest LNBA– genomes from central Siberia and North Caucasus are both chronologically concurrent (ca. 4,836 to 4,622 y B.P.) and phylogenetically positioned together. Given the 4,600 km that separate these genomes, it appears that the LNBA– lineage displayed a fast and extensive spread, presumably facilitated first by ox-hauled carts and, later, horse and camel domestication (11–13). Overall, LNBA– strains may have 1) all evolved from a single source deme, 2) spread with a high geographic mobility, and 3) had a limited reservoir.

In addition to new “LNBA–” genomes, the authors report a genome from El Sotillo (Spain), dated 3,361 to 3,181 y B.P. This genome, together with the previously published genome from the Volga region (Russia), dated 3,868 to 3,704 y B.P., represent two separate lineages that emerged at the beginning of the second millennium BCE and designated as “LNBA+” because they acquired *ymt*, which expands the range of mammalian hosts that sustain

flea-borne plague (14). Interestingly, the LNBA+ and LNBA– temporally coexisted among themselves and with other early lineages (0.PE7, 0.PE2, 0.PE4, and 0.PE5), emerging successively in the later third and the second millennia BCE and also harboring *ymt* (Fig. 1) (1).

Both LNBA– and LNBA+ acquired the loci (*pla* and *caf*) important to produce different “modern” forms of plague with high incidence: pneumonic plague transmitted by aerosol between humans and bubonic plague transmitted by fleas (15, 16). Also, LNBA+ would have evolved further to become more virulent in mammals and fleas, notably through the acquisition of the YPMT1.66c and *ymt* genes (14, 17) (Fig. 1).

Some argued that the early clinical manifestation of plague was the pneumonic form (2, 5). This rarest form of plague requires a permanent close contact between humans in densely settled environments to be maintained over space and time (18), which hardly characterizes late Neolithic and Bronze Age Eurasia. While some “proto-urban” sites in Eurasia, such as Cucuteni-Trypillia (northeastern Romania, Moldova, and western Ukraine) or Altyndepe (Turkmenistan) had high population densities (19, 20), most regions, especially the steppe, were sparsely populated. Therefore, one may wonder whether LNBA– and “+” lineages would have been more likely transmitted by fleas (or even lice) rather than inhalation of contaminated aerosols.

There are two models of flea-borne plague transmission. According to the “early-phase transmission” model, the transmission occurs within a few hours or days after an infectious blood meal and then quickly fades away if uninfected meals are subsequently ingested (21). According to the “blocked-flea model,” transmission requires an extrinsic incubation period of at least 4 to 5 d and occurs even after the ingestion of uninfected meals over 1 mo (22, 23). The mechanism of early-phase transmission is unclear. By contrast, we know that the “blocked-flea transmission” results from *Y. pestis*’ ability to form a solid mass

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Author contributions: P.S. and F.S. wrote the paper.

The authors declare no competing interest.

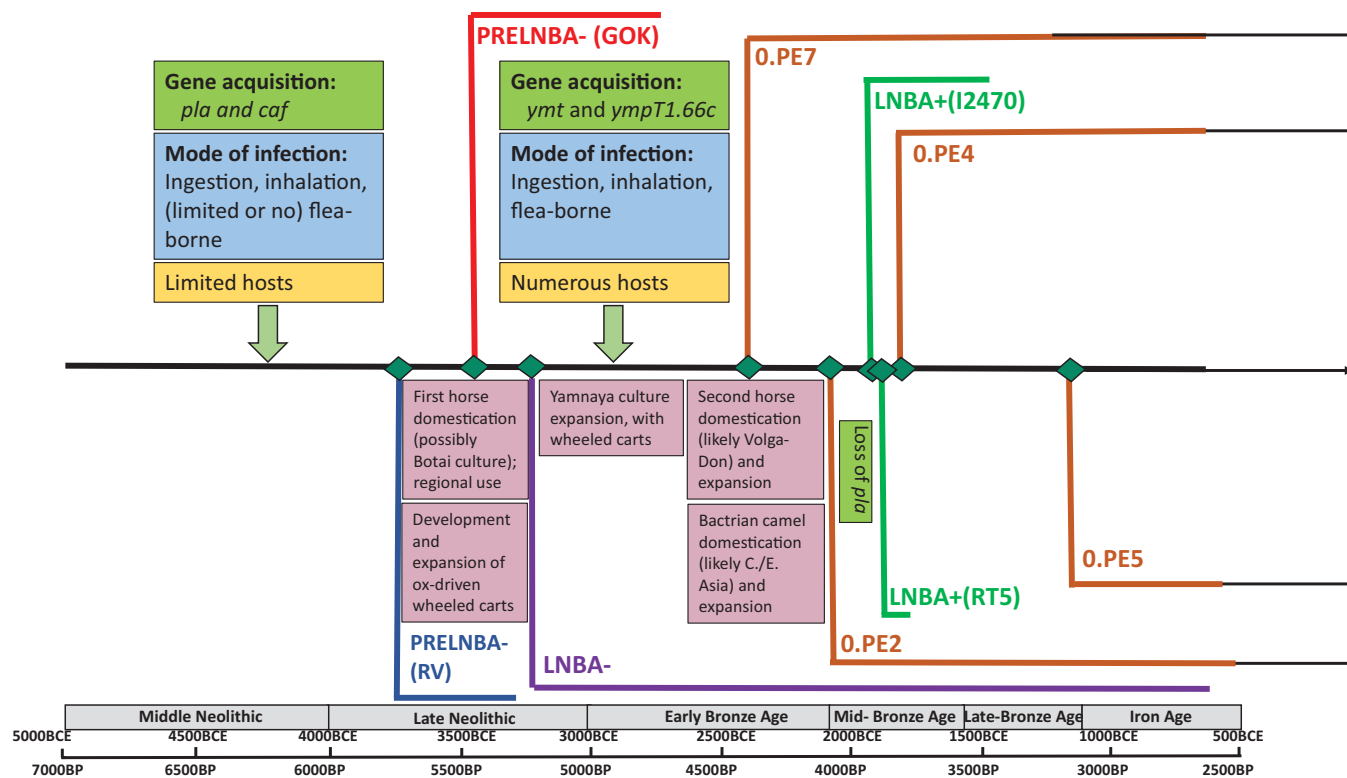
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See companion article, “Stone Age *Yersinia pestis* genomes shed light on the early evolution, diversity, and ecology of plague,” [10.1073/pnas.2116722119](https://doi.org/10.1073/pnas.2116722119).

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Published May 17, 2022.



**Fig. 1.** Early evolutionary history of *Y. pestis* based on the study and refs. 14–17 and 21–24 and its historical contexts, based on refs. 11–13.

in the foregut. This mass obstructs the ingestion of blood into the gut during a feeding attempt and increases the fleabite rate (due to the starving) and, therefore, favors the regurgitation of *Y. pestis* into the mammalian host. The transmission by “blocked” fleas sustains the long-term persistence of the flea-mammal-flea cycle, which may not be possible with early-phase transmission.

Both LNBA– and LNBA+ could, in theory, have been transmitted by fleas. However, fundamental genetic differences between the two lineages imply that the chances of transmission of the LNBA– by fleas is rather low and most likely limited to early-phase transmission. Indeed, LNBA– lineage has retained 1) a ureolytic activity killing >40% of fleas within the first day of infection and 2) functional ancestral genes that prevent the bacteria from producing a carbohydrate polymer important for flea blockage (23, 24). Furthermore, it lacks *ymt* that expands the range of mammalian hosts that sustains flea-borne plague (14). This genetic pattern may explain the particular LNBA– phylogeny, characterized by a low level of genetic diversification and numerous evolutionary dead ends, and evidently an eventual extinction of this lineage, judging by its absence from modern-day reservoirs.

The putative extinction of nonadapted flea LNBA– lineage occurred at least 2,500 y after its rise. How did this lineage survive and spread for such a long time? Is it possible that it spread via the ingestion of contaminated food, considering it emerged from an enteropathogen? In the Late Neolithic and Early Bronze Age, the Eurasian steppe between the Black Sea and western China was settled primarily by nomads, dwelling in proximity to a wider natural world in which sylvatic rodents abounded. While there is, at present, no paleodiet data on human consumption of

rodents, there is archaeological evidence of their hunting and skinning, as well as using their teeth and bones for tools and crafts in late Neolithic Central Eurasia (5, 25, 26). However, as evidence from 20th-century Central Asia shows, primary outbreak of gastroenteric plague in humans is confined to local communities, rather than spread across regions (27). Moreover, to maintain such a route of transmission would require constant contacts between infected rodents and humans all over Eurasia—hardly a feasible option. Finally, small migratory birds could be incriminated for the dissemination of LNBA– strains, given that a limited number of birds can be infected by *Y. pestis* and some may spread *Y. pestis*-infected fleas (28).

But what about flea-adapted LNBA+ lineages which, despite their theoretical capability to seed multiple reservoirs and become genetically diversified, became extinct, judging by their absence from modern-day reservoirs—in contrast to O.PE branches, thriving today in different Asian foci (11, 29, 30)? All documented O.PE2, O.PE4, O.PE5, and O.PE7 genomes (>100 in total) come from sparsely populated regions in Asia. Could it be that local ecological and sociodemographic landscapes of O.PE branches, marked by nomadic pastoralism, meant that plague was confined primarily to rodents, attacking humans only sporadically, and having less potential to burn itself out than LNBA+, spreading over a much wider geographic range, which included sedentary and more densely populated landscapes of western Eurasia, and circulating more intensively in humans? To appreciate why and when LNBA+ branches, in contrast with O.PE, eventually died out, more genomes from various late Bronze and Iron Age and contexts are needed.

The question of the context in which *Y. pestis* lineages emerged, spread, evolved, and disappear is as important.

Establishing the origins of the early lineages is, at present, not feasible, but a hypothesis may be offered. Given that the two earliest LNBA- genomes (RISE509 from central Siberia and RK1001 from North Caucasus) are both chronologically concurrent and phylogenetically positioned together, despite being situated some 4,600 km apart, may indicate that their common source was somewhere in between: in Central Asia, a home to both several Bronze-Age 0.PE lineages and all Iron-Age and medieval 0.ANT branches. If the preLNBA-, LNBA-, and LNBA+ branches arose in Central Asia, then the local climatic context of their chronologies is to be considered. The successive emergence of the three earliest known lineages between ca. 5,600 and 5,100 y B.P. as well as the 0.PE2, the two LNBA+ lineages and 0.PE4 between ca. 4,000 and 3,800 y B.P. (Fig. 1) occurred in the

context of excessively wet episodes in Central Asia (31–34). Could it be that excessive rainfall, creating abundant grass biomass and facilitating population growth of rodent hosts, provided the optimal conditions for intense bacterial activity leading to divergence events?

Overall, Andrades Valtueña et al.'s (1) work opens the door to some exciting big questions such as when, where, how, and why ancestral strains have emerged, evolved, spread, and sometimes counterselected to extinction, and how they got transmitted from wildlife reservoirs to human populations. The work also invites the question of where we draw a border between attributes defining *Y. pestis* and its ancestor *Y. pseudotuberculosis*. Future collaborative synergistic research will undoubtedly advance our understanding of these fascinating questions.

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