Population Genomics of Legionella longbeachae and Hidden Complexities of Infection Source Attribution

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Legionella longbeachae is the primary cause of legionellosis in Australasia and Southeast Asia and an emerging pathogen in Europe and the United States; however, our understanding of the population diversity of L. longbeachae from patient and environmental sources is limited. We analyzed the genomes of 64 L. longbeachae isolates, of which 29 were from a cluster of legionellosis cases linked to commercial growing media in Scotland in 2013 and 35 were non–outbreak-associated isolates from Scotland and other countries. We identified extensive genetic diversity across the L. longbeachae species, associated with intraspecies and interspecies gene flow, and a wide geographic distribution of closely related genotypes. Of note, we observed a highly diverse pool of L. longbeachae genotypes within compost samples that precluded the genetic establishment of an infection source. These data represent a view of the genomic diversity of L. longbeachae that will inform strategies for investigating future outbreaks.

Legionellosis presents as 2 clinically distinct forms: an influenza-like illness called Pontiac fever and a severe pneumonia known as Legionnaires’ disease (1). In Europe and the United States, most legionellosis cases are caused by Legionella pneumophila serogroup 1 (1,2); <5% of cases are caused by nonpneumophila Legionella spp. (3,4). In Australasia, New Zealand, and some countries in Asia, infections caused by L. longbeachae occur at comparable levels to infections caused by L. pneumophila (5–7). Unlike L. pneumophila infections, which are typically linked to artificial water systems, L. longbeachae infections are associated with exposure to soil, compost, and potting mixes (8).

The number of legionellosis cases caused by L. longbeachae is increasing worldwide (7), with a notable rise reported across Europe (9–11). Within the United Kingdom, most L. longbeachae infections have been identified in Scotland, where 6 cases were diagnosed during 2008–2012 (12) and another 6 were diagnosed in the summer of 2013 and represented a singular increased incidence or cluster with all patients requiring intensive care hospitalization (11). Epidemiologic investigation revealed that most patients from the 2013 cluster were avid gardeners, and L. longbeachae was isolated from respiratory secretions and from samples of the growing media they had used for gardening before becoming ill (11,12). However, an investigation into the provenance of the growing media did not reveal a single commercial or manufacturing source that would suggest a common origin for the L. longbeachae associated with the outbreak (11).

Molecular typing methods used to discriminate between L. longbeachae and other Legionella spp. and between the 2 L. longbeachae serogroups have limited efficacy, and although considerable evidence supports growing media as a source for L. longbeachae infections (13,14), there is still a lack of genetic evidence for an epidemiologic link. Furthermore, a population genomic study involving large numbers of L. pneumophila isolates has been conducted (15,16), but the same has not been done for L. longbeachae, so the diversity of environmental and pathogenic genotypes and the relationship between them remains unknown for L. longbeachae. To examine the etiology of the 2013 cluster of legionellosis cases in Scotland in the context of L. longbeachae species diversity, we analyzed the genomes of 70 Legionella spp. isolates from 4 countries over 18 years.

Materials and Methods

Bacterial Isolates
We sequenced 65 isolates that had previously been identified as L. longbeachae. These isolates were obtained during 1996–2014 from several patients, growing media samples (including compost and soil), and a hot water supply. Of these isolates, 55 were from Scotland (29 from the 2013 cluster of infections and 26 from other clinical and environmental samples) and 10 were from patients and

In our analysis, we also included all publicly available genome sequences for L. longbeachae: L. longbeachae NSW150 (serogroup 1) and L. longbeachae C-4E7 (serogroup 2) isolated from patients in Australia; and L. longbeachae D-4968 (serogroup 1), L. longbeachae ATCC39642 (serogroup 1), and L. longbeachae 98072 (serogroup 2) isolated from patients in the United States (17–19). We sequenced multiple isolates (n = 2 to 5) for each of 3 patients and their linked growing media samples from the 2013 outbreak in Scotland and for 2 additional compost samples. The species of all isolates had been determined by serotyping or macrophage infectivity potentiator (mip) gene sequencing (20,21).

**Bacterial Culture, Genomic DNA Isolation, and Whole-Genome Sequencing**

We cultured Legionella spp. isolates in a microaerophilic and humid environment at 37°C on BCYE (buffered charcoal yeast extract) agar plates for 48 h. We then picked individual colonies from the plates and grew them in ACES-buffered yeast extract broth containing Legionella BCYE Growth Supplement (Oxoid Ltd., Basingstoke, UK) with shaking at 37°C for 24–48 h. We extracted genomic DNA from fresh cultures by using the QIAGEN DNeasy Blood and Tissue Kit (QIAGEN Benelux B.V., Venlo, the Netherlands).

We prepared sequencing libraries by using the Nextera XT kit for MiSeq or HiSeq (all from Illumina, San Diego, CA, USA) sequencing at Edinburgh Genomics, University of Edinburgh (Edinburgh, Scotland, UK). For each isolate, one 2 × 250–bp or two 2 × 200–bp paired-end sequencing runs were carried out using the MiSeq and HiSeq technologies, respectively. Raw reads were quality checked using FastQC v0.10.1 (22), and primers were trimmed by using Cutadapt (23). We used wgsim software (24) to simulate sequence reads for publicly available, complete whole-genome sequences.

**Bioinformatic Analysis and Data Deposition**

A detailed description of the bioinformatic analyses is available in the online Technical Appendix. The sequence data for the 65 genomes of Legionella spp. sequenced in this study were deposited in the SRA database (accession no. PRJEB14754).

**Results**

**Limitations of Current Typing Approaches for Legionella spp. Identification**

We sequenced 65 isolates obtained from several patients and environmental samples over 18 years in different countries and previously identified as L. longbeachae. To confirm the species identity of the Legionella isolates, we constructed a phylogenetic tree that included all Legionella type strains for which cultures are available, based on the 16S rRNA gene sequence (25). We also built phylogenetic trees based on the whole-genome content and core-genome diversity. For each approach, 64 of the 70 isolates examined co-segregated within the L. longbeachae–specific clade, 4 isolates clustered with Legionella anisa, and 2 belonged to a separate clade that was distinct from all known Legionella spp. (Figure 1; online Technical Appendix Figures 1, 2). The species identities were further supported by determination of the average nucleotide identity values (online Technical Appendix Figure 3), a widely used method for bacterial species delineation based on genomic relatedness (26). Of note, L. anisa is the most common nonpneumophila Legionella spp. in Europe (27–29). In addition, L. longbeachae isolates 13.8642 (from a compost sample from Scotland) and 13.8295 (from a patient in New Zealand) belong to a putative novel Legionella spp. Overall, the data indicate that current serotyping methods and mip gene sequencing are limited in their capacity to identify L. longbeachae to the species level.

To investigate the genetic relatedness of L. longbeachae strains associated with the 2013 outbreak to temporally and geographically distinct isolates, we constructed a core genome–based neighbor-joining tree of the 64 confirmed L. longbeachae isolates obtained from 4 countries over 18 years (online Technical Appendix Figure 4). This phylogenetic tree presents a comet-like pattern, with 2 distinct clades separated by 9,911 single-nucleotide polymorphisms, representing the major serogroups (serogroups 1 and 2) previously identified for L. longbeachae (20), each containing isolates from patient and environmental samples from different years. In contrast with findings from a previous analysis of 2 isolates of L. longbeachae serogroup 1 (20), we observed a higher diversity among the 56 isolates within serogroup 1 (online Technical Appendix Figures 1, 4); this finding is not unexpected, given the difference in the number of genomes examined. Nevertheless, compared with isolates from the same serogroup in other Legionella spp., such as L. pneumophila serogroup 1 (2% polymorphism) (20), L. longbeachae serogroup 1 exhibits a lower diversity (<0.1% polymorphism). Although serogroup 1 and 2 clades contained isolates from Scotland, Australasia, and the United States, 96% of the isolates from Scotland (including all of the 2013 outbreak isolates) belonged to serogroup 1, suggesting that serogroup 1 may be more clinically relevant in Scotland than in some other countries where L. longbeachae is a more established cause of legionellosis. However, analysis of more isolates from different countries would be required to investigate this observation further.
Effect of Recombination on *L. longbeachae* Serogroup 1 Population Structure

It is established that recombination has played a key role in shaping the evolutionary history of *L. pneumophila*, but its effect on *L. longbeachae* population structure is unknown (22,30). This knowledge is critical because for highly recombinant bacteria, recombination networks may represent evolutionary relationships more explicitly than traditional phylogenetic trees. Therefore, we constructed a recombination network of all serogroup 1 isolates by using the neighbor-net algorithm of SplitsTree4 (31). The resultant network displayed a reticulate topology with an extensive reticulated background from which clusters of isolates emerge, supporting an evolutionary history involving recombination (p< 0.01 by \(\phi\) test) (32), followed by clonal expansion and subsequent additional recombination events among some lineages (online Technical Appendix Figure 5). Using BratNextGen (33), we identified a total of 94 predicted recombination events affecting more than half of the core genome (1.74 Mb of 3.36 Mb) and representing recent and ancient recombination events of different sizes (range 1,350 bp–350 Kbp) distributed across the phylogeny (online Technical Appendix Figure 6). Given the reported limitation in sensitivity of BratNextGen for the identification of all recombination events (34), we also used ClonalFrameML (35), an algorithm that uses maximum likelihood inference to simultaneously detect recombination in bacterial genomes and account for it in phylogenetic reconstruction. The estimated average length of the recombined fragments was 8,047 bp, and the ratio of recombination to mutation was 1.42, indicating a greater role for recombination over mutation in the diversification
of *L. longbeachae*. This estimate is in accordance with early estimates for *L. pneumophila* based on multiple gene sequence data (36), but it is low compared with recent estimates based on whole-genome sequence data [recombination to mutation ratios of 16.8 (30) or 47.93 (37)]. Differences in the clonal diversity of *Legionella* spp. sequence datasets used to determine recombination rates could affect the estimates. Reconstruction of the phylogeny after removal of all predicted recombinant sequences resulted in a tree with largely similar clusters of isolates but with reduced branch lengths and variation in the position of nodes deep in the phylogeny (Figure 2).

**Accessory Genome Analysis Indicates Extensive Interspecies and Intraspecies Gene Flow**

The extent to which horizontal gene transfer occurs among *L. longbeachae* isolates and between *L. longbeachae* and other *Legionella* spp. is unknown. In our study, the pangenome of *L. longbeachae* represented by the 56 serogroup 1 isolates was 6,890 genes, including a core genome of 2,574 genes; the average gene content was 3,558 genes per strain. The accessory genome, which included only strain-dependent genes varied from 809 to 1,155 genes, depending on the strain. A parsimony clustering analysis based on the presence or absence of all genes classified the isolates in a manner distinct from that in a core genome–based maximum-likelihood tree, suggesting extensive horizontal gene transfer among *L. longbeachae* isolates (online Technical Appendix Figures 1, 2). BLAST (https://blast.ncbi.nlm.nih.gov/Blast.cgi) analysis of all assembled contigs was used to filter for plasmid-related homologous sequences, revealed 2 major plasmids: pLLO, described previously in *L. longbeachae* NSW150 (20), and pLELO, originally identified...
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In *L. pneumophila* subsp. *pneumophila* (22). Of the 55 serogroup 1 isolates, 36 contained sequences for the pLLO and pLELO plasmids. Of note, the distribution of these plasmids among the *L. longbeachae* isolates correlated with the gene content–based clustering, whereas the distribution of plasmids in the core genome–based tree was independent of the phylogeny (Figure 2). In addition, 11 isolates appeared to contain plasmids with sequences homologous to those for pLLO and pLELO, which is indicative of recombinant forms of the plasmid. Further examination of plasmid diversity using a modified version of PLACNET (38), a program enabling reconstruction of plasmids from whole-genome sequence datasets, confirmed that some plasmids consisted of a mosaic of recombinant fragments homologous to pLELO, pLLO, or other unknown plasmids (Figure 3). Overall, these data indicate the high prevalence of specific plasmids among *L. longbeachae* isolates and reveal extensive recombination and horizontal gene transfer among different *Legionella* spp (39). The high prevalence of plasmids in *L. longbeachae* is notable, considering these elements may be less common in *L. pneumophila* (30).

To examine the possibility that clinical and environmental isolates of *L. longbeachae* contained genomic differences reflecting their distinct origins, we compared their accessory genome content. For isolates obtained from a single patient sample, the accessory genome was highly conserved compared with those for environmental isolates from a single compost sample or closely related environmental isolates from distinct compost samples (Figure 4, panel A). In addition, considering the average gene content of all sequenced isolates (28 clinical and 27 environmental), the gene content for *L. longbeachae* from growing media samples (3,586 genes) was significantly higher than that for isolates from patients (3,533 genes; 2-sample *t*-test, *t* = 2.5213; d.f. = 53; *p* = 0.01474) (Figure 4, panel B). The data imply that gene loss occurs during human infection or that *L. longbeachae* strains with reduced gene content have enhanced human infectivity. However, we did not identify a specific enriched gene or functional category in clinical or environmental samples (data not shown).

**Source Attribution Confounded by Complex Serogroup 1 Populations within Environmental Samples**

Having accounted for the influence of recombination on the phylogeny of *L. longbeachae*, we investigated the diversity of isolates associated with 5 patients and their linked compost samples obtained during 2008–2014, including

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**Figure 3.** *Legionella longbeachae* plasmid analysis: contigs networks reconstructions for 6 representative *L. longbeachae* types of plasmid content. The networks of the contigs representing the main chromosome and plasmids comprising the genome obtained by using PLACNET (38), a program enabling reconstruction of plasmids from whole-genome sequence datasets. The sizes of the contig nodes (in gray) are proportional to their lengths; continuous lines correspond to scaffold links. Dashed lines represent BLAST (https://blast.ncbi.nlm.nih.gov/Blast.cgi) hits to the *L. longbeachae* (blue) or *L. pneumophila* (red) strains; intensity of the line is proportional to the hit (white indicates low, black indicates high). Green lines correspond to plasmid contigs. Background colors indicate species relatedness for the main chromosome and plasmids (blue for *L. longbeachae*, red for *L. pneumophila*, pink for a combination of both, and yellow for previously unidentified genomic content).
3 patients from the 2013 outbreak in Scotland. Of note, isolates from the 2013 outbreak were distributed across several subclades of the tree, indicating that the infections were caused by different strains (Figure 2). However, all isolates from a single patient clustered together, consistent with a monoclonal etiology of each infection. Of note, for all 5 patients, clinical isolates were not closely allied to the environmental isolates obtained from linked compost samples, and therefore a genetic link between patient and compost samples could not be established. Most subclades included isolates of diverse geographic origin, consistent with a wide distribution for \( L. \) longbeachae strains; however, 3 \( L. \) longbeachae isolates originating from Australasia (strains 13.8294, 13.8293, and NSW150) belonged to their own region-specific cluster (Figure 2).

We hypothesized that the lack of genetic relatedness between \( L. \) longbeachae isolates from patients and linked compost samples could be explained by a highly diverse population of \( L. \) longbeachae in growing media samples compounded by a sampling strategy consisting of a single sequenced isolate. All 5 compost samples for which we had >1 isolate contained isolates distributed across multiple clades in the phylogenetic tree. In particular, 5 isolates from the same growing media sample linked to a patient infected in Edinburgh in 2014 were distributed across 4 distinct clades, demonstrating that within a single environmental sample, considerable species diversity may be represented (Figure 2). Taken together, these data suggest that for future outbreak investigations, extensive sampling of environmental samples may be required to identify genotypes responsible for episodes of legionellosis infection, if indeed they are present.

**Discussion**

Our findings reveal the population genomic structure for \( L. \) longbeachae, an emerging pathogen in Europe and the United States, and includes a genome-scale investigation into an outbreak of \( L. \) longbeachae legionellosis. We provide evidence for extensive recombination and lateral gene transfer among \( L. \) longbeachae, including the presence of widely distributed mosaic plasmids that have likely recombined with plasmids from other \( Legionella \) spp., suggesting an ecologic overlap or shared habitat. Our analysis highlights the need to account for recombination events when determining the genetic relatedness of \( L. \) longbeachae isolates.

Our application of whole-genome sequencing for diagnostic purposes revealed the misidentification, using current serotyping methods, of several \( L. \) anisa isolates as \( L. \) longbeachae and led to the identification of a putative novel \( Legionella \) sp. linked to legionellosis. These findings highlight the limitations of current typing methods for differentiation of \( Legionella \) spp. and accurate identification of legionellosis etiology.

We used whole-genome sequencing to attempt to establish a genetic link between legionellosis infections and associated compost samples. Our inability to establish a link probably reflects the traditional strategy of single isolate sampling, which when applied to a highly diverse pool of \( L. \) longbeachae genotypes fails to detect the infecting genotype. We suggest that the approach to investigating the source of future legionellosis cases linked to growing media will require a radical revision of sampling protocols to maximize the chances of isolating the infecting strain, if present. Taken together, our findings provide a view of the population structure of \( L. \) longbeachae and highlight the complexities of tracing the origin of legionellosis associated with growing media. Overall, our findings demonstrate the resolution afforded by whole-genome sequencing for understanding the biology underpinning legionellosis and provide information that should be considered for future epidemiologic investigations.
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