



Complete Genome Sequences of the *Arcobacter cryaerophilus* Strains ATCC 43158^T and ATCC 49615

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ABSTRACT *Arcobacter cryaerophilus* was originally recovered from aborted bovine and porcine fetuses, but it has been subsequently isolated from meat, water, and human clinical samples. This study describes the complete whole-genome sequences of two *A. cryaerophilus* strains, ATCC 43158^T (=A 169/B^T =LMG 24291^T) and ATCC 49615 (=CDC D2610 =LMG 10829).

Arcobacter cryaerophilus (formerly *Campylobacter cryaerophila* [1]) was originally recovered from aborted bovine and porcine fetuses (1–3), from the milk of cows with mastitis (4), and from bovine and porcine feces (1). Subsequent studies reported the recovery of *A. cryaerophilus* isolates from meat (5, 6), water (7), and human clinical samples (8, 9). Additionally, based on the results of whole-cell protein and rRNA restriction fragment length polymorphism analyses (10, 11), *A. cryaerophilus* strains have been divided into two subgroups, 1A and 1B. In this study, we report the first closed genome sequences of two *A. cryaerophilus* strains, the type strain ATCC 43158 (subgroup 1A; =A 169/B^T [1] =LMG 24291^T), isolated in the United Kingdom from the brain of an aborted bovine fetus, and strain ATCC 49615 (subgroup 1B; =CDC D2610 [12] =LMG 10829), isolated in Illinois from human blood. A draft genome sequence (91 scaffolds) was deposited previously for *A. cryaerophilus* strain ATCC 43158^T (GenBank accession number [NXGK01000000](https://www.ncbi.nlm.nih.gov/genome/genomes/11530)). Additionally, draft genome sequences for 12 other *A. cryaerophilus* strains have been deposited previously in GenBank (<https://www.ncbi.nlm.nih.gov/genome/genomes/11530>).

Both *A. cryaerophilus* strains were grown aerobically at 30°C for 48 h on anaerobe basal agar (Oxoid) amended with 5% horse blood. Cells were separately removed from each plate using sterile 5- μ l inoculating loops, and genomic DNA was prepared from each loopful of cells using the Promega Wizard genomic DNA purification kit (Madison, WI). Roche 454 shotgun and paired-end libraries were constructed for both strains, using the manufacturer's standard protocols, and sequenced on a GS-FLX+ instrument using the Titanium chemistry. For each strain, the shotgun and paired-end reads were assembled together, using Newbler v. 2.6 (Roche) and default parameters; 454 reads were quality controlled within Newbler. Sequencing metrics for both strains are presented in Table 1. Two scaffolds were obtained for strain ATCC 43158^T, one chromosomal scaffold of 20 contigs and a megaplasmid scaffold of 4 contigs. One chromosomal scaffold of 16 contigs was obtained for strain ATCC 49615. Each scaffold was closed by using the custom Perl script `contig_extender3` (13) to place within the scaffold gaps unique, nonscaffolded contigs and/or contigs representing sequences present at two or more locations. The two chromosomes and the megaplasmid were assembled manually within SeqMan Pro v. 8.0 (DNASTAR, Madison, WI), using the contig order defined above and contig-spanning 454 reads. Assembly of both genomes was verified using an optical restriction map (restriction enzyme XbaI; OpGen, Gaith-

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TABLE 1 Sequencing metrics and genomic data for *Arcobacter cryaerophilus* strains ATCC 43158^T and ATCC 49615

Feature ^b	Value(s) ^a	
	ATCC 43158 ^T	ATCC 49615
Sequencing metrics		
454 (shotgun) platform		
No. of reads	215,129	87,759
No. of bases	90,272,116	36,134,765
Avg length (bases)	419.6	411.7
Coverage (×)	45.0	17.6
454 (paired-end) platform		
No. of reads	118,785	161,571
No. of bases	38,763,675	49,505,772
Avg length (bases)	326.3	306.4
Coverage (×)	19.3	24.1
Illumina HiSeq 2000 platform		
No. of reads	18,366,932	18,769,966
No. of bases	1,836,693,200	1,895,766,566
Avg length (bases)	100	101
Coverage (×)	915.2	922.1
PacBio platform		
No. of reads	117,659	NA
No. of bases	437,690,629	NA
Avg length (bases)	3,720 ^c	NA
Coverage (×)	218.1	NA
Newbler metrics ^d		
Total no. of contigs	45	32
N50ContigSize (454)	195,432	346,474
Q40PlusBases (454) (%)	99.90	99.87
N50ContigSize (HiSeq pool 1)	195,198	301,090
Q40PlusBases (HiSeq pool 1) (%)	99.98	99.96
N50ContigSize (HiSeq pool 2)	195,524	165,874
Q40PlusBases (HiSeq pool 2) (%)	99.96	99.96
N50ContigSize (HiSeq pool 3)	195,196	301,081
Q40PlusBases (HiSeq pool 3) (%)	99.97	99.99
Genomic data		
Size (bp)	2,006,909; 101,431	2,055,914
G+C content (%)	27.46; 24.63	27.51
No. of CDS ^e	1,951; 94	1,994
Assigned function (% CDS)	847 (43.4); 3 (3.2)	857 (43.0)
General function annotation (% CDS)	675 (34.6); 34 (36.2)	694 (34.8)
Domain/family annotation only (% CDS)	131 (6.7); 4 (4.3)	136 (6.8)
Hypothetical (% CDS)	298 (15.3); 53 (56.4)	307 (15.4)
Pseudogenes	23; 6	8
Genomic islands/CRISPR		
No. of genetic islands	8; 0	5
No. of CDS in genetic islands	108 [2]; 0	85 [2]
No. of CRISPR/Cas loci	0	0
Gene content/pathways		
IS elements, mobile elements, or transposases	2; 1	2
Signal transduction		
No. of Che proteins	9; 0	9
No. of methyl-accepting chemotaxis proteins	20; 0	23
No. of response regulators	23; 2	25
No. of histidine kinases	25; 1	27
No. of response regulator/histidine kinase fusions	0	3
No. of diguanylate cyclases	14; 0	15
No. of diguanylate phosphodiesterases (HD-GYP, EAL)	4, 1; 0	5, 1
No. of diguanylate cyclase/phosphodiesterases	9; 0	9
No. of other	10; 1	10
Motility		
Flagellin genes	<i>flaAB</i>	<i>flaAB</i>
Restriction/modification		
No. of type I systems (<i>hsd</i>)	[2]; 0	2
No. of type II systems	1; 0	3
No. of type III systems	0	2

(Continued on next page)

TABLE 1 (Continued)

Feature ^b	Value(s) ^a	
	ATCC 43158 ^T	ATCC 49615
Transcription/translation		
No. of transcriptional regulatory proteins	32; 4	32
Non-ECF σ factors	σ^{70}	σ^{70}
No. of ECF σ factors	0	0
No. of tRNAs	49; 0	51
No. of ribosomal loci	5; 0	5
CO dehydrogenase (<i>coxSLF</i>)	No	No
Ethanolamine utilization (<i>eutBCH</i>)	No	No
Nitrogen fixation (<i>nif</i>)	No	No
Osmoprotection	<i>ectABC</i>	No
Pyruvate \rightarrow acetyl-CoA		
Pyruvate dehydrogenase (E1/E2/E3)	Yes	Yes
Pyruvate:ferredoxin oxidoreductase	No	No
Urease	No	No
Vitamin B ₁₂ biosynthesis	No	No

^a Numbers in square brackets indicate pseudogenes or fragments. Strain ATCC 43158^T values before a semicolon are for the chromosome, while values after the semicolon are for the pACRY43158 plasmid. NA, not applicable.

^b CDS, coding sequences; ECF, extracytoplasmic function; acetyl-CoA, acetyl coenzyme A.

^c Maximum length, 25,867 bases.

^d Features and values taken from largeContigMetrics within 454NewblerMetrics.txt for each assembly. Large contigs were defined as having ≥ 500 bases. Due to the large number of HiSeq reads, the total reads were split into three pools and assembled independently. N50ContigSize value is in number of bases.

^e Numbers do not include pseudogenes.

ersburg, MD). Illumina HiSeq reads were obtained from SeqWright (Houston, TX) and assembled and quality controlled within Newbler v. 2.6, as above. The Illumina contigs were assembled automatically (unique contigs) or positioned manually (repeated contigs) within the 454 SeqMan assembly. The Illumina contigs and reads were used to verify the 454 base calls, as described previously (14). A repeat region within the type strain chromosome required long reads for validation. Therefore, a 20-kb PacBio library was constructed, using standard methods (15), and sequenced on an RS II instrument, using standard protocols, as described previously (14). PacBio reads were assembled with RS_HGAP_Assembly v. 3 (Pacific Biosciences) and default settings, which yielded single chromosomal and megaplasmid contigs. The PacBio chromosomal contig was manually placed within the SeqMan assembly to confirm the 454 contig order across the repeat region.

Genomic data for the two *A. cryaerophilus* strains are presented in Table 1. The average genome size for the two strains is approximately 2 Mbp, with an average G+C content of 27.5%. Protein-, rRNA-, and tRNA-encoding genes were identified using GeneMark, RNAmmer, and ARAGORN (16–18), respectively, and annotated as described previously (14). The ATCC 43158^T genome also contains the 101,431-bp megaplasmid pACRY43158.

Data availability. The complete genome sequence of *A. cryaerophilus* strain ATCC 43158^T has been deposited in GenBank under the accession numbers [CP032823](#) (chromosome) and [CP032824](#) (pACRY43158), and the complete genome sequence of *A. cryaerophilus* strain ATCC 49615 has been deposited in GenBank under the accession number [CP032825](#). All 454, HiSeq, and PacBio sequencing reads for strain ATCC 43158^T and 454 and HiSeq reads for strain ATCC 49615 have been deposited in the NCBI Sequence Read Archive (SRA) under the accession numbers [SRP164716](#) and [SRP164722](#), respectively.

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