

Figure S1: Sequence proportion heat map (white = absent, red = high, gray = no sample) of genus *Vibrio* at six sampling stations (R-Outfall, R-Estuary-1, R-Estuary-2, NS-Marine, OS-Marine and SM-Outfall) in the winter, spring, summer, and autumn sampling during the 2015/2016 and 2018/2019 surveys (Orel et al., 2022).

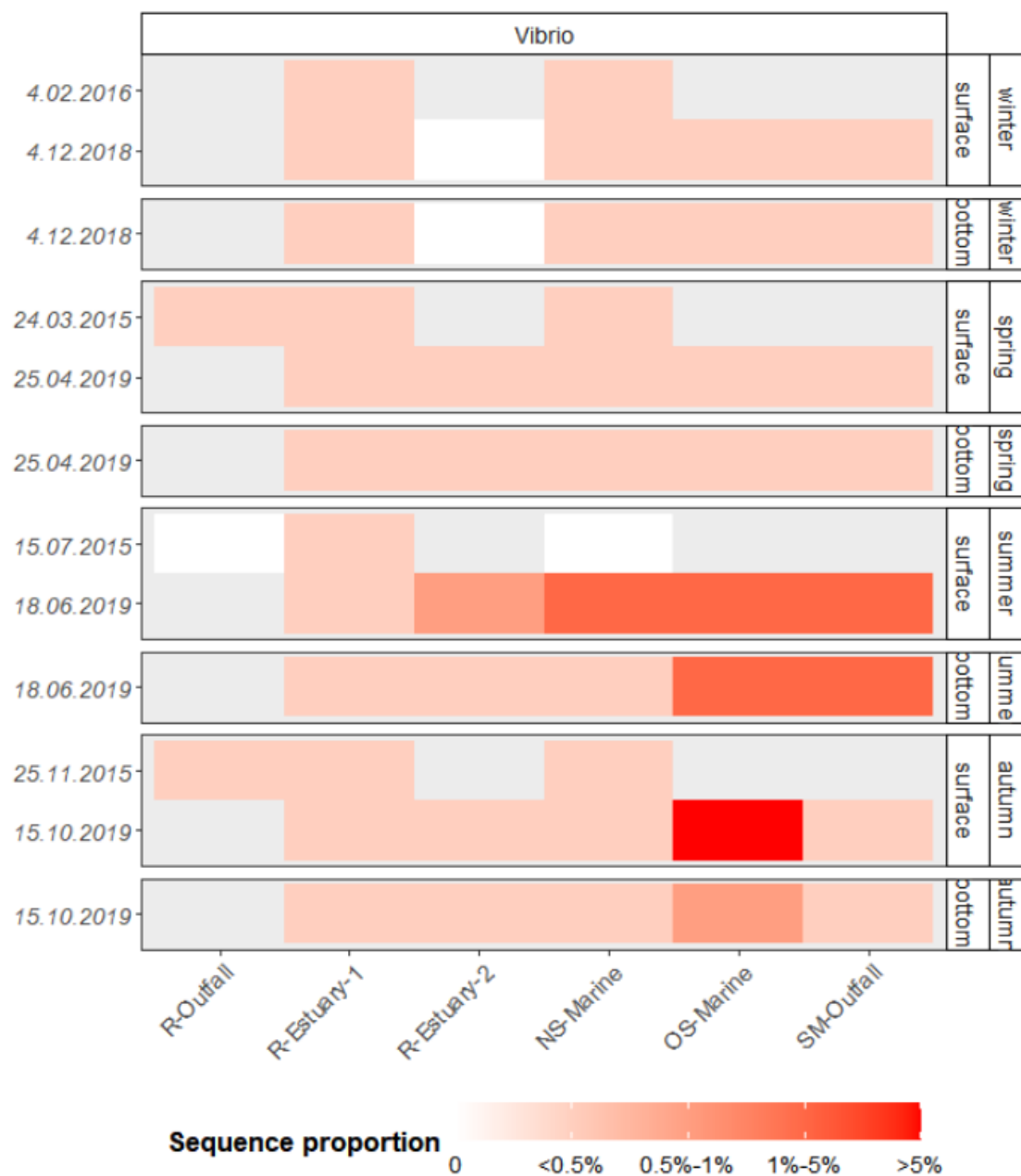
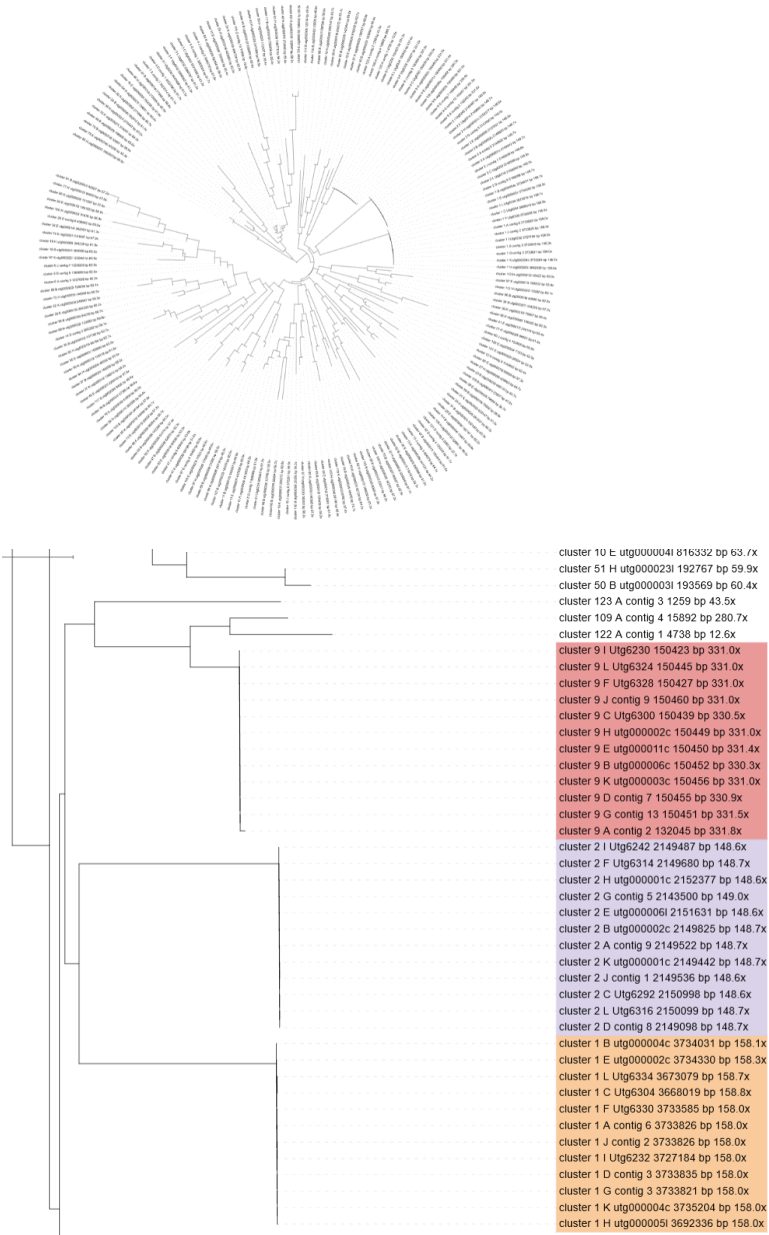


Figure S2: Trycycler contigs tree. Trycycler tool requires manual intervention; therefore, its output is not deterministic. In the first step of Trycycler workflow, contigs from the input assemblies are clustered based on pairwise Mash distances between contigs and the user must decide which clusters are valid. **(A)** presents FastME tree, built using the pairwise distances of contigs for BF5_0283 sample. Selected clusters (1, 2 and 9) meet the criteria: cluster contain many contigs (ideally 12, one from each assembly), contigs are close to each other and have similar lengths, and similar and realistic read depths. **(B)** presents FastME tree, built using the pairwise distances for Mt009 sample. Clusters were not well defined; selected clusters (1, 2, 3) only partly meet criteria used for BF5_0283.

A



B

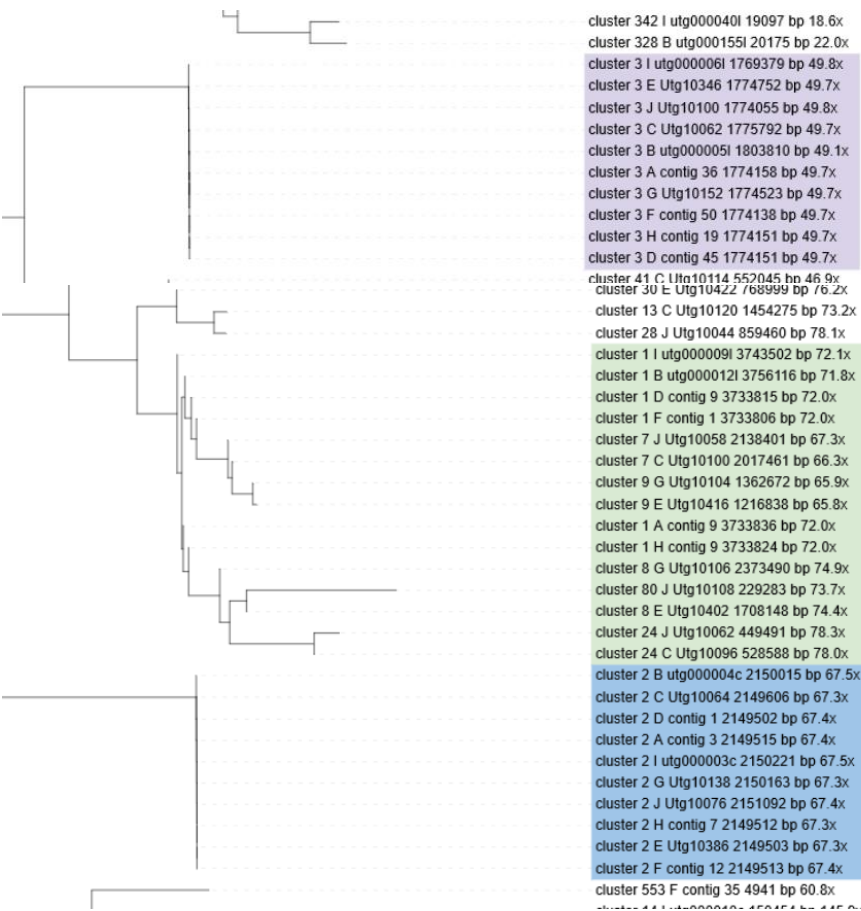
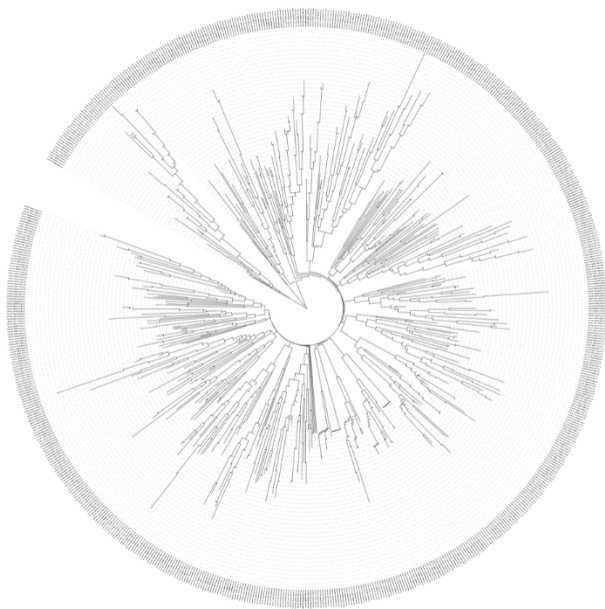
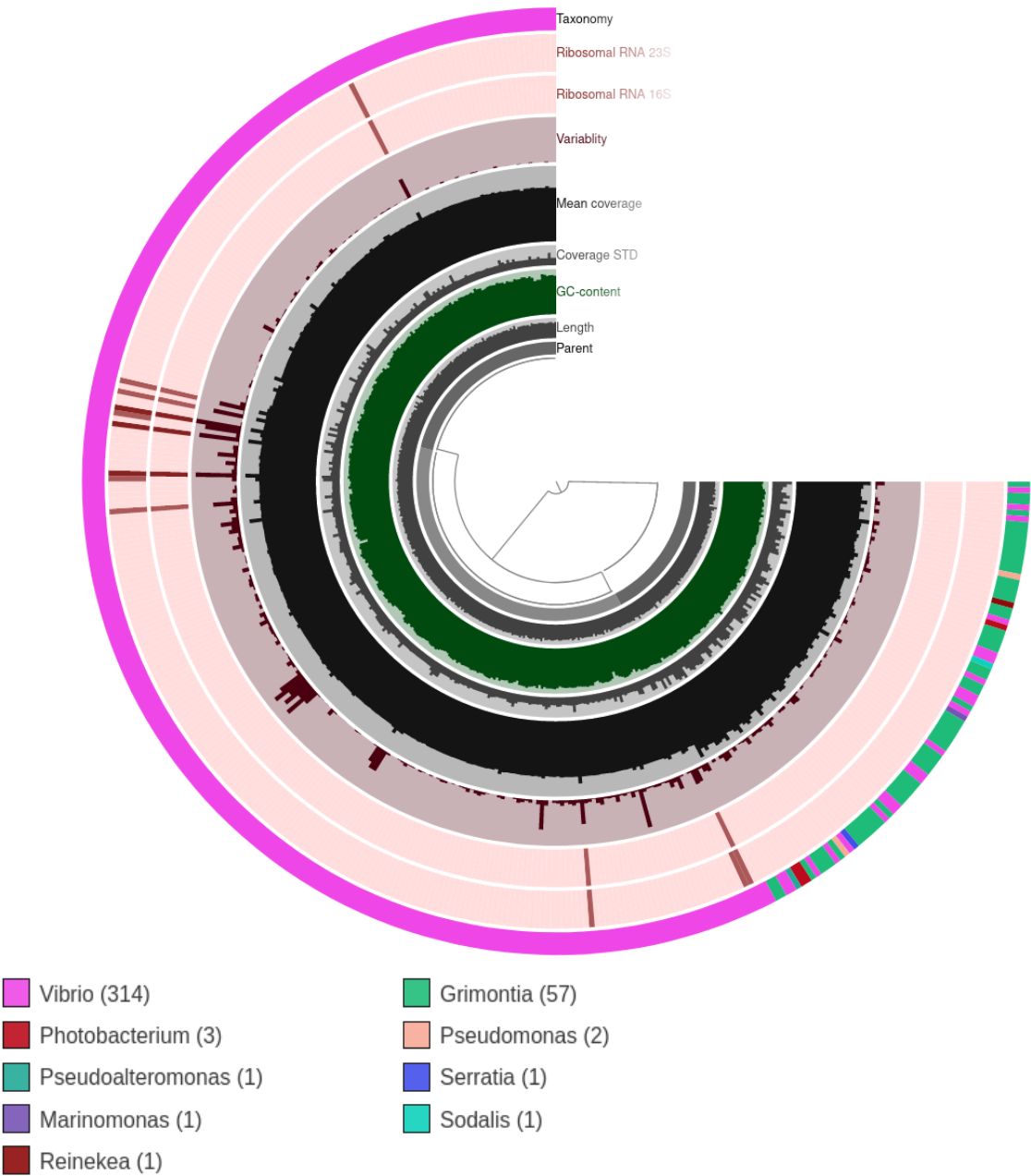
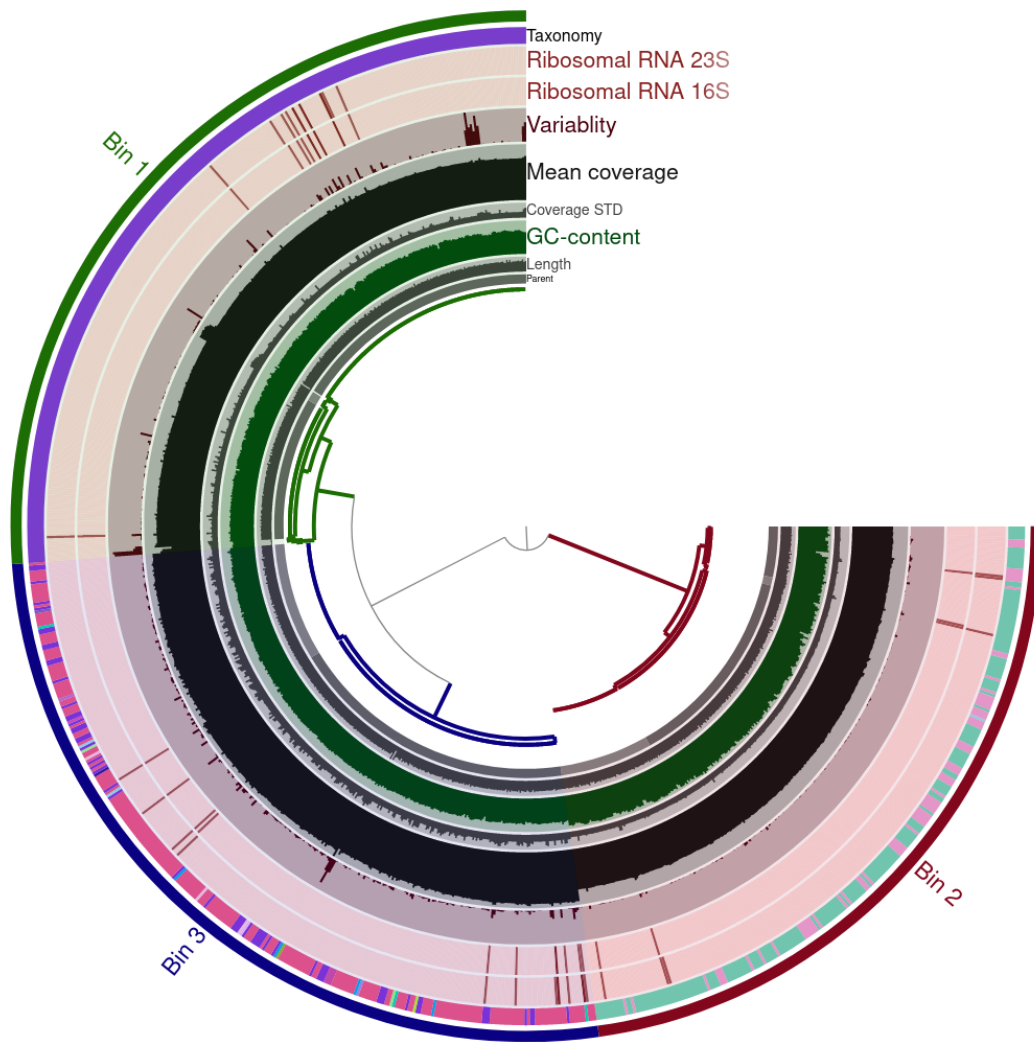


Figure S3: Graphical presentation of contigs from Mt009 assemblies along with associated data with “anvi-interactive” function. **(A)** presents Trycycler assembly, **(B)** presents Unicycler assembly and **(C)** presents metaFlye assembly. Tree in the middle shows contigs presented in assemblies. Bins in **(B)** were manually refined based on differences in mean coverage (mapping of Illumina short reads), differences in GC content and genes taxonomy.

A.

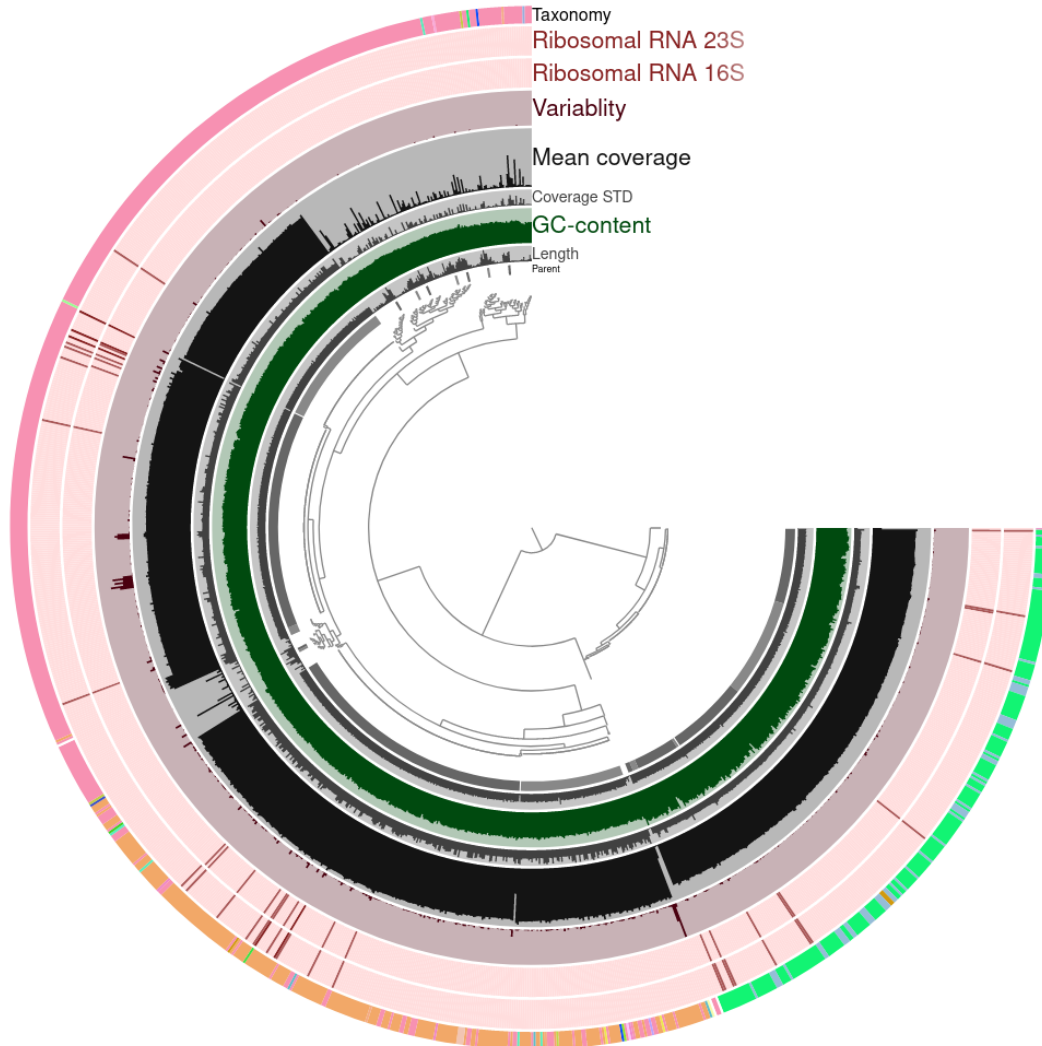


B.



■ Vibrio (360)	■ Grimontia (224)
■ Unknown_Enterobacteriaceae (194)	■ Klebsiella (70)
■ Shewanella (4)	■ Unknown_Gammaproteobacteria (3)
■ Pseudoalteromonas (3)	■ Photobacterium (3)
■ Salinivibrio (3)	■ Pseudomonas (1)
■ Psychromonas (1)	■ Aeromonas (1)
■ Moritella (1)	■ Marinobacterium (1)
■ Bacterioplanes (1)	■ Rhodoferax (1)
■ Lacimicrobium (1)	■ Marinomonas (1)
■ Plesiomonas (1)	

C.



Vibrio (518)	Grimontia (224)
Unknown_Enterobacteriaceae (210)	Klebsiella (53)
Photobacterium (5)	Pseudomonas (3)
Unknown_Gammaproteobacteria (3)	Aliivibrio (3)
Salinivibrio (3)	Shewanella (3)
Lactobacillus (2)	Moritella (2)
Unknown_Vibrionaceae (1)	Colwellia (2)
Delftia (1)	Plesiomonas (1)
Escherichia (1)	Catenovulum (1)
Psychromonas (1)	Marinomonas (1)
Tenacibaculum (1)	Caulobacter (1)
Bacterioplanes (1)	Arsenophonus (1)
None (3)	Legionella (1)

Figure S4: Results of PCR reaction with taxa-specific primers. Agarose gel electrophoresis of amplicons visualized by ethidium bromide staining and UV light. Samples on the gel are 1kb ladder, C- (negative control of PCR reaction), BF5_0283, BF5_0283, Mt009, Mt009, Mt009, C+ (positive control of PCR reaction), and C2- (negative control of DNA extraction). All samples were tested with 16S universal primers, *V. campbellii* and *K. pneumoniae* taxa-specific primers.

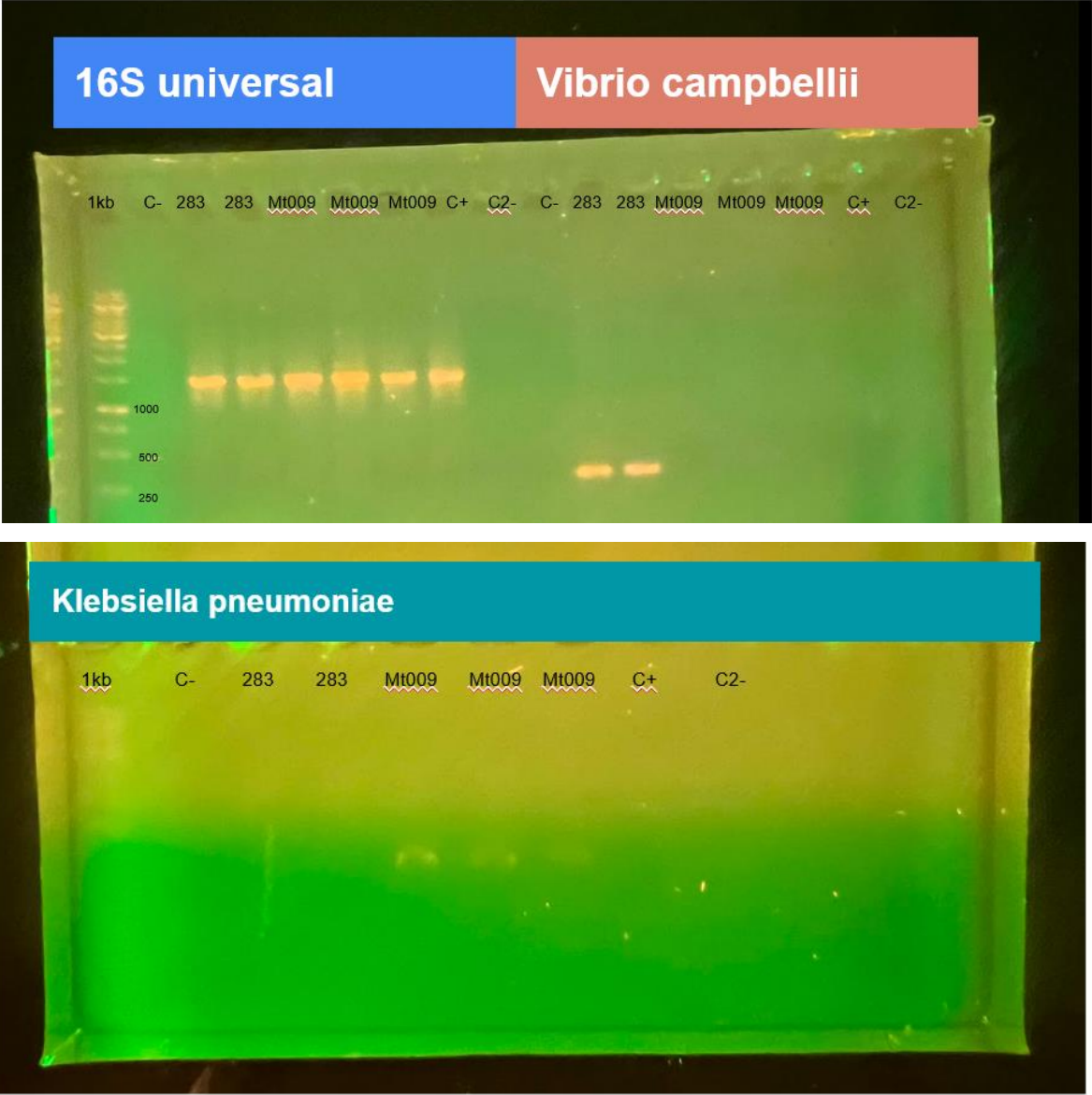


Figure S5: Additional pangenome analyses of BF5_0283 and Mt009_b1 assemblies. Clustering based on the presence/absence patterns of 5293 gene clusters (GCs) resulted in Core (present in both assemblies) and Singletons collection (present in only one assembly). GC are ordered based on geometric homogeneity index. GCs with geometric homogeneity index lower than 0.999 are marked with red line in outermost circle. Note: This figure is not discussed in the manuscript.

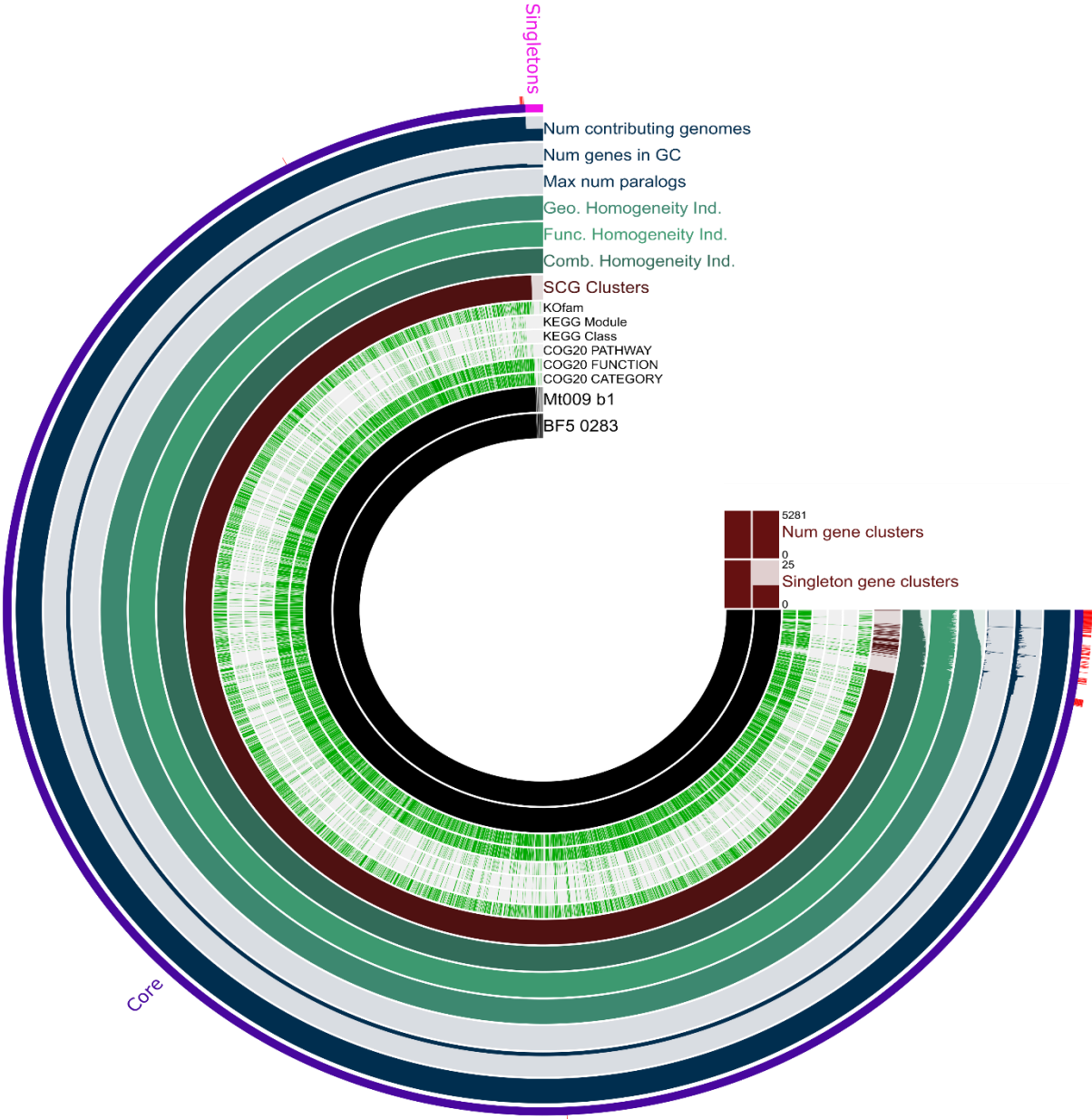
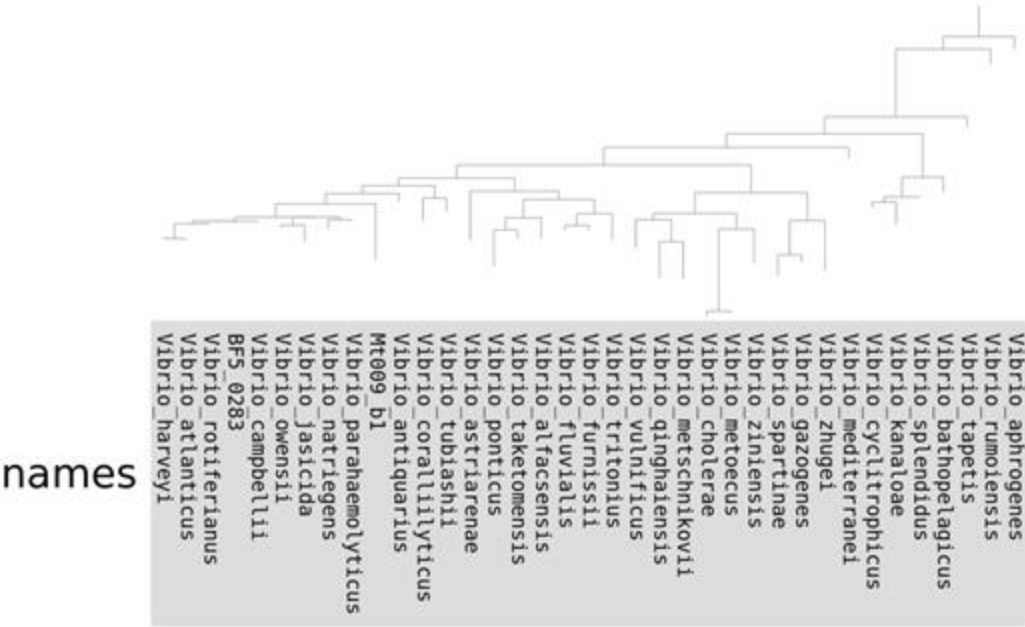


Figure S6: Phylogeny of *Vibrio* genomes based on 16S rRNA genes **(A)** and single-copy core orthologues **(B)**. An approximately-maximum-likelihood phylogenetic tree represented 32 *Vibrio* genomes from NCBI and genomes assembled in this study (BF5_0283 and Mt009_b1). The tree was rooted to the outgroup *V. aphrogenes*.

A



B

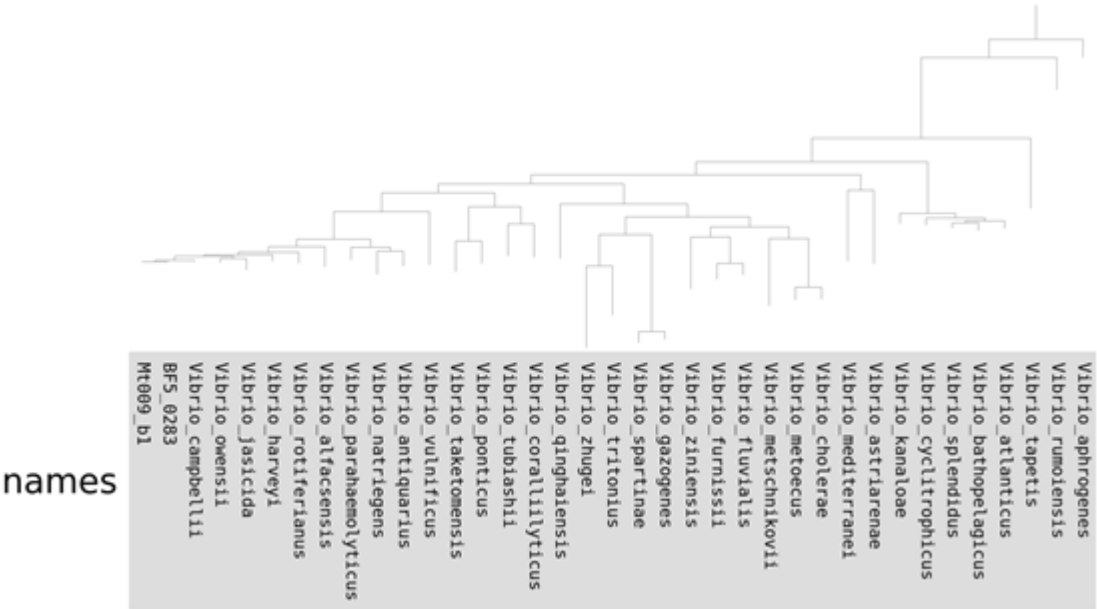
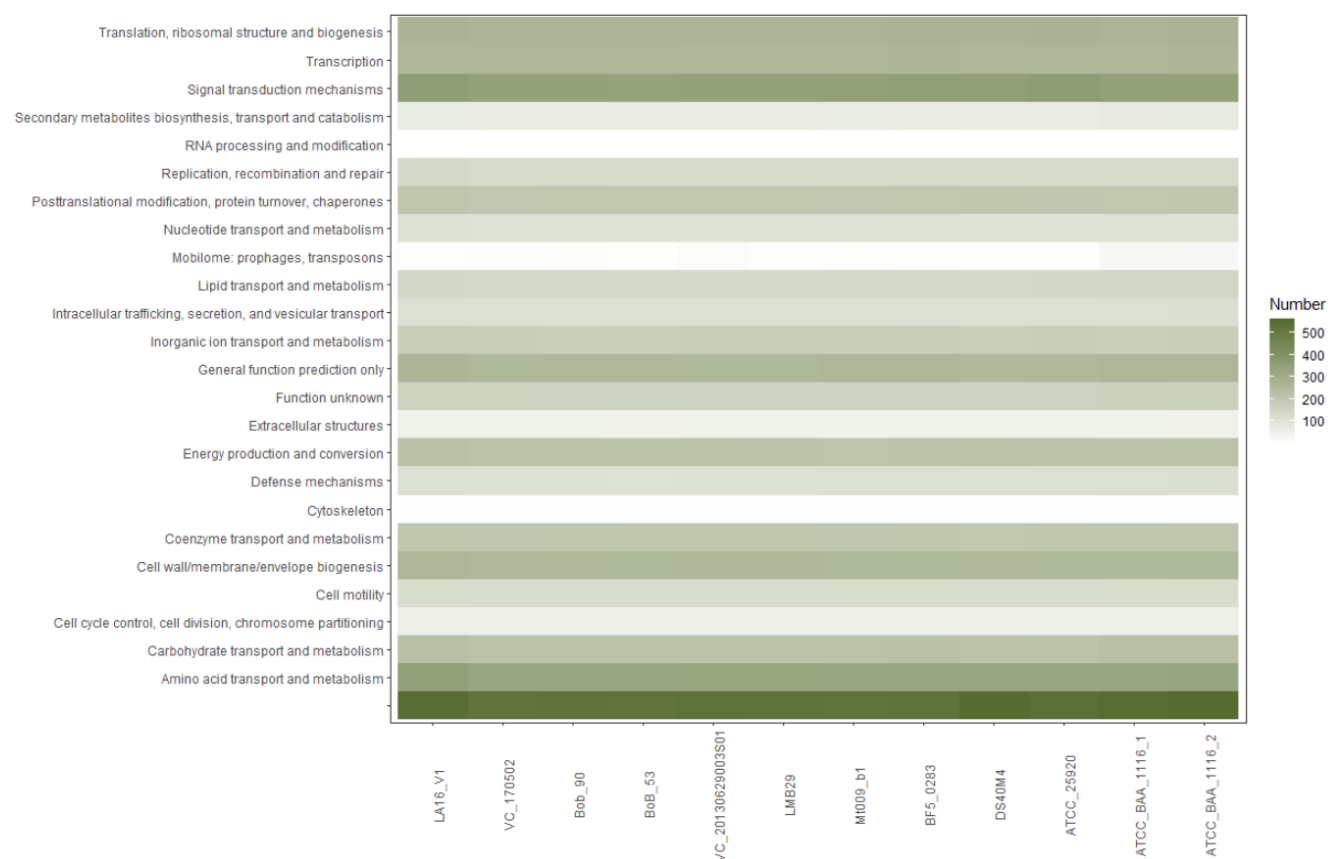
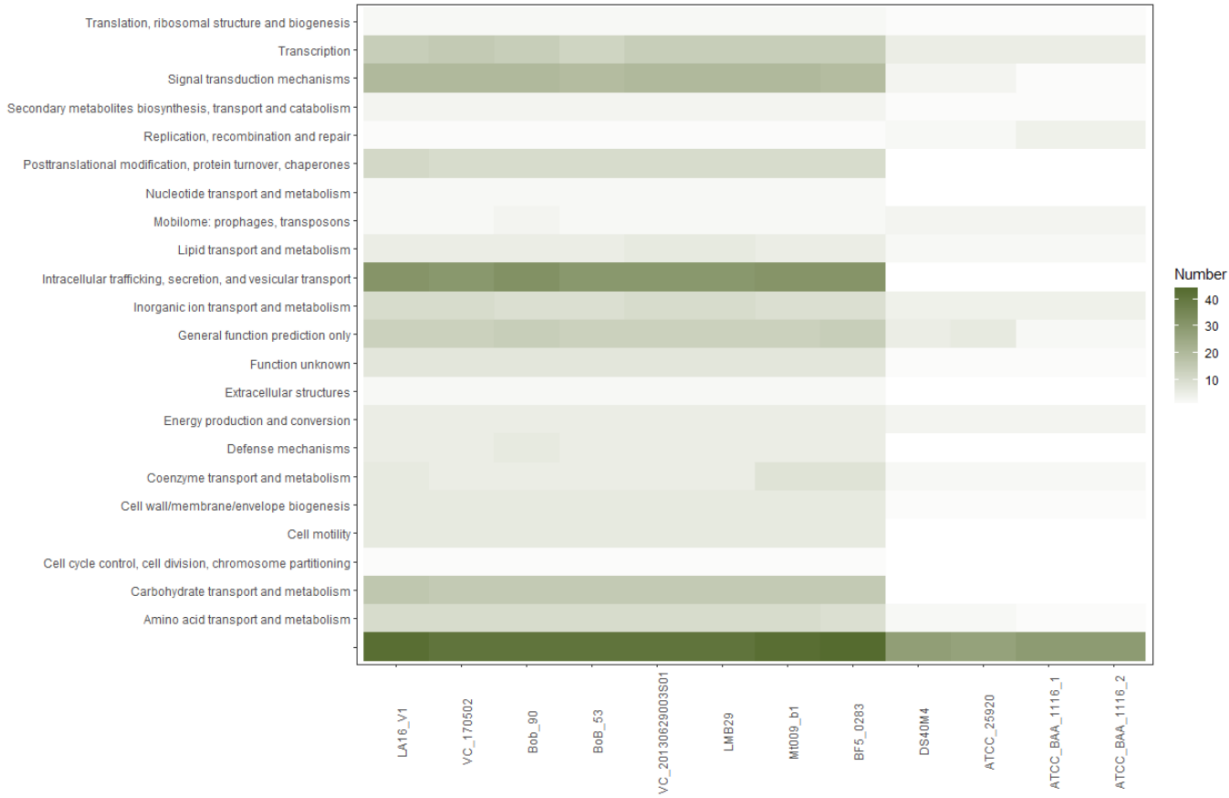


Figure S7: Gene abundance heat map (white = absent, green = high), representing abundance of genes in *V. campbelli* genomes, belonging to different COG categories. **(A)** presents annotations of genes in core collection (present in all genomes according to pangenome analyses), **(B)** presents annotations of genes in accessory Group 1 and Group 2 collections and **(C)** presents annotations of genes unique for BF_0283 and Mt009_b1.



B



C

