





Draft Genome Sequence of *Vibrio splendidus* DSM 19640

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ABSTRACT Here, we present the draft genome sequence of *Vibrio splendidus* type strain DSM 19640. *V. splendidus* is an abundant species among coastal vibrioplankton. The assembly resulted in a 5,729,362-bp draft genome with 5,032 protein-coding sequences, 6 rRNAs, and 117 tRNAs.

Vibrio splendidus, an opportunistic marine pathogen, is a dominant member of the coastal vibrioplankton and one of the causative agents of vibriosis (1). Vibriosis affects marine wildlife, as well as aquaculture, and thus leads to major economic losses (2). *V. splendidus* can cause high mortality rates in larval and juvenile marine animals, including turbot, oysters, clams, and scallops (3, 4) and can even spread to humans through the consumption of infected seafood.

The pathogenicity of *V. splendidus* is multifactorial and regulated by intrinsic and extrinsic factors. While increasing water temperatures favor its transmission and proliferation, an extracellular metalloprotease (Vsm) has been shown to be the major determinant of toxicity in Pacific oysters (5). Despite a significant body of empirical research, our understanding of the factors that contribute to the virulence of *V. splendidus* is far from being complete.

Genomic DNA of *V. splendidus* DSM 19640 was extracted with the MasterPure complete DNA purification kit (Epicentre, Madison, WI, USA) and used to generate Illumina shotgun paired-end sequencing libraries. Sequencing was performed employing the MiSeq system and the MiSeq version 3 reagent kit (600 cycles), as recommended by the manufacturer (Illumina, San Diego, CA, USA). Quality filtering using Trimmomatic version 0.32 (6) resulted in 2,483,315 paired-end reads. *De novo* genome assembly was performed with the SPAdes version 3.11.0 genome assembler (7). The assembly resulted in 66 contigs (>500 bp) and an average coverage of 18,876-fold. The assembly was validated, and the read coverage was determined with QualiMap version 2.1 (8).

The draft genome of *V. splendidus* DSM 19640 consisted of 5,729,362 bp with an overall GC content of 43.91%. Genome annotation was performed using Prokka (9). In total, 6 rRNAs, 117 tRNAs, 3,416 protein-coding sequences with predicted functions and 1,616 genes with unknown functions were determined.

A search for toxins in the genome revealed the following three putative virulence factors: (i) a gene encoding a protein identical to the *V. cholerae* zona occludens toxin (VSPL_25760), (ii) a protein identical to the transmembrane regulatory protein ToxS involved in the regulation of *V. cholerae* toxins, and (iii) a protein with 93.65% identity to the metalloprotease Vsm known from *V. splendidus* strain JZ6 (VSPL_47630).

This is the first report of the genome sequence of *V. splendidus*. It will provide further insights into the virulence mechanisms of this pathogen and will also serve as a platform to facilitate comparative genomics of other *Vibrio* species.

Received 3 November 2017 Accepted 6 November 2017 Published 30 November 2017

Citation Chibani CM, Poehlein A, Roth O, Liesegang H, Wendling CC. 2017. Draft genome sequence of *Vibrio splendidus* DSM 19640. Genome Announc 5:e01368-17. <https://doi.org/10.1128/genomeA.01368-17>.

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Accession number(s). This whole-genome shotgun project has been deposited at DDBJ/ENA/GenBank under the accession number [PDMZ00000000](https://doi.org/10.1007/s10482-009-9398-4). The version reported here is the first version, PDMZ01000000.

ACKNOWLEDGMENTS

This project was funded by a grant from the Deutsche Forschungsgemeinschaft (DFG) within the priority program SPP1819 given to C.C.W., O.R., and H.L., and by a grant from the Cluster of Excellence 80 “The Future Ocean” given to C.C.W. and O.R. “The Future Ocean” is funded within the framework of the Excellence Initiative by the DFG on behalf of the German federal and state governments.

REFERENCES

1. Balcázar JL, Gallo-Bueno A, Planas M, Pintado J. 2010. Isolation of *Vibrio alginolyticus* and *Vibrio splendidus* from captive-bred seahorses with disease symptoms. *Antonie Van Leeuwenhoek* 97:207–210. <https://doi.org/10.1007/s10482-009-9398-4>.
2. Lacoste A, Jalabert F, Malham S, Cueff A, Gélébart F, Cordevant C, Lange M, Poulet SA. 2001. A *Vibrio splendidus* strain is associated with summer mortality of juvenile oysters *Crassostrea gigas* in the Bay of Morlaix (North Brittany, France). *Dis Aquat Organ* 46:139–145. <https://doi.org/10.3354/dao046139>.
3. Gómez-León J, Villamil L, Lemos ML, Novoa B, Figueras A. 2005. Isolation of *Vibrio alginolyticus* and *Vibrio splendidus* from aquacultured carpet shell clam (*Ruditapes decussatus*) larvae associated with mass mortalities. *Appl Environ Microbiol* 71:98–104. <https://doi.org/10.1128/AEM.71.1.98-104.2005>.
4. Gay M, Renault T, Pons AM, Le Roux F. 2004. Two *Vibrio splendidus* related strains collaborate to kill *Crassostrea gigas*: taxonomy and host alterations. *Dis Aquat Organ* 62:65–74. <https://doi.org/10.3354/dao062065>.
5. Binesse J, Delsert C, Saulnier D, Champomier-Vergès MC, Zagorec M, Munier-Lehmann H, Mazel D, Le Roux F. 2008. Metalloprotease Vsm is the major determinant of toxicity for extracellular products of *Vibrio splendidus*. *Appl Environ Microbiol* 74:7108–7117. <https://doi.org/10.1128/AEM.01261-08>.
6. Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. *Bioinformatics* 30:2114–2120. <https://doi.org/10.1093/bioinformatics/btu170>.
7. Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 19:455–477. <https://doi.org/10.1089/cmb.2012.0021>.
8. Okonechnikov K, Conesa A, García-Alcalde F. 2016. Qualimap 2: advanced multi-sample quality control for high-throughput sequencing data. *Bioinformatics* 32:292–294. <https://doi.org/10.1093/bioinformatics/btv566>.
9. Seemann T. 2014. Prokka: rapid prokaryotic genome annotation. *Bioinformatics* 30:2068–2069. <https://doi.org/10.1093/bioinformatics/btu153>.