Data Article

Microbiome dataset analysis from a shrimp pond in Ninh Thuan, Vietnam using shotgun metagenomics

Lan-Anh Le\textsuperscript{a}, Thien-Phuc Nguyen-Hoang\textsuperscript{b}, Van-Phuc Huynh\textsuperscript{a}, Tan-Huy Nguyen\textsuperscript{a}, Thuy-Vy Nguyen\textsuperscript{b}, Thuy-Duong Ho-Huynh\textsuperscript{a,b,c,}\textsuperscript{*}

\textsuperscript{a} Khoa Thuong Biotechnology Company Limited, No 17, Street 4, Phong Phu, Binh Chanh, Ho Chi Minh, Vietnam
\textsuperscript{b} Department of Genetics, Faculty of Biology and Biotechnology, University of Science, VNU-HCM, 227 Nguyen Van Cu Street, Ward 4, District 5, Ho Chi Minh, Vietnam
\textsuperscript{c} Research Center for Genetics and Reproductive Health, School of Medicine, VNU-HCM, 6 Residential area, Linh Trung, Thu Duc, Ho Chi Minh, Vietnam

\textbf{A R T I C L E   I N F O}

\textbf{Article history:}
Received 16 April 2020
Accepted 13 May 2020
Available online 21 May 2020

\textbf{Keywords:}
Metagenomics
Shotgun sequencing
Oxford Nanopore MinION sequencing
Aquaculture
Shrimp pond

\textbf{A B S T R A C T}

Vietnam is one of the top shrimp producing and exporting countries in the world [1]. However, viral and bacterial epidemic diseases cause severe damages to shrimp farming, resulting in millions of US dollars losses annually [2]. Furthermore, inappropriate use of antibiotics in shrimp rearing lead to increased emergence of drug resistant pathogens [3]. Current practices for water quality control, mostly based on chemical and physical parameters; neglected biological criteria necessary for maintaining pond health.

Ninh Thuan is a region situated in the South Central Coast of Vietnam. Due to its geographic location, a large part of this region is dedicated to shrimp (\textit{Litopenaeus vannamei}) post-larvae production and rearing. This article presents a microbiome dataset from two water samples collected in a shrimp rearing pond in Ninh Thuan. We used Oxford Nanopore Technologies (ONT) for metagenomic sequencing of the samples to characterize microbial communities and antibiotic resistance profiles. The metagenome dataset generated will provide an understanding and comparison framework of the microbial diversity and functionality among shrimp ponds with

\textsuperscript{*} Corresponding author.
\textit{E-mail address:} hhtduong@hcmus.edu.vn (T.-D. Ho-Huynh).

\url{https://doi.org/10.1016/j.dib.2020.105731}

\textcopyright 2020 The Author(s). Published by Elsevier Inc. This is an open access article under the CC BY license. (http://creativecommons.org/licenses/by/4.0/)
potential application in health management and shrimp rearing industry.

© 2020 The Author(s). Published by Elsevier Inc.
This is an open access article under the CC BY license. (http://creativecommons.org/licenses/by/4.0/)

| Specifications Table                                      |
|-----------------------------------------------------------|
| **Subject** Applied Microbiology and Biotechnology        |
| **Specific subject area** Metagenomics                    |
| **Type of data** Figures and Fastq files                   |
| **How data were acquired** MinION (ONT)                    |
| **Data format** Raw and analyzed                           |
| **Parameters for data collection** Water samples collected from a shrimp pond at 0.3-m depth |
| **Description of data collection** Total DNA was extracted from water samples, and shotgun metagenomic sequencing was performed using ONT MinION platform |
| **Data source location** Village/Town/City: Phuoc Dinh, Thuan Nam, Ninh Thuan Country: Vietnam |
| Latitude and longitude coordinates for collected samples: 11°24′46.6″N, 108°58′19.0″E and 11°24′46.4″N, 108°58′17.0″E |
| **Data accessibility** Data is available at the NCBI with Bioproject PRJNA552940 and SRA accession numbers SRR9648445 (https://trace.ncbi.nlm.nih.gov/Traces/sra/?run=SRR9648445) and SRR9648446 (https://trace.ncbi.nlm.nih.gov/Traces/sra/?run=SRR9648446). |

**Value of the Data**

- The analysis revealed an insight of microbiome present in water of a shrimp rearing pond in Vietnam.
- Data obtained from this article provided the scientific community with microbial taxonomic profiles and functional characteristics of a shrimp pond using metagenomics.
- Data can be used for the comparison of taxonomic and functional profiles among shrimp ponds with different geographic distribution.
- Data can serve future analyses of the relationship between microorganism profiles of shrimp pond water and risks of disease outbreak as well as health threats due to antibiotic resistance genes spread.

1. **Data Description**

The data in this dataset described the taxonomic and functional profiles of two metagenomic samples from a shrimp pond in Ninh Thuan, Vietnam. We performed shotgun sequencing on ONT MinION platform and analyzed data using MG-RAST server and KmerResistance. A total of 28,837 (sample 1) and 24,434 (sample 2) reads were classified out of 60,944 (sample 1) and 51,784 (sample 2) reads sequenced. Data was presented as taxonomic and functional profiles in Figs. 1 and 2, respectively. Additionally, antimicrobial resistance determinants against beta-lactamase, aminoglycosides, and sulphonamide were observed in both samples from the same template acquired from *Pseudomonas aeruginosa* (aadA2b, *sul1_2_U12338*), *Klebsiella pneumoniae* (*sul1_EU780013*), *Vibrio cholerae strain MO10* (*sul2_2_AY034138*) (Supplementary Table 1).
Fig. 1. Taxonomic metagenomic profile of a water sample from shrimp pond - Ninh Thuan, Vietnam. The most abundant domain was bacteria (94.28%), followed by viruses (3.68%), eukaryota (1.77%), and archaea (0.08%). Among bacteria, phylum observed in a descending order of magnitude included Proteobacteria, Actinobacteria, Bacteroidetes, Firmicutes, Cyanobacteria. Of the 146 bacterial orders detected, the most abundant were Burkholderiaceae (21.07%), Rhodobacteraceae (17.85%) and Gallionellaceae (6.64%). Moreover, a total of 252 families and 515 genus was identified.

Fig. 2. Functional metagenomic profile of a water sample from shrimp pond - Ninh Thuan, Vietnam. The most abundant function corresponded to phages, transposable elements and plasmids (29.97%); followed by clustering-based subsystems (14.02%), carbohydrates (8.96%), amino acids and derivatives (8.51%), protein metabolism (6.23%); and other categories (32.78%).
2. Experimental Design, Materials, and Methods

2.1. Sample collection and DNA extraction

Two water samples were collected at two different sites, approximately at 0.3-m depth, from a shrimp rearing pond located at Ninh Thuan Province, Vietnam in March 2019. Samples were delivered to the laboratory on ice in 9 hours. Filtered samples (50 ml) were submitted to DNA extraction using the QIAamp Blood DNA mini kit (Qiagen, Germany). DNA quality and concentration were assessed by spectrophotometer and Qubit dsDNA HS assay kit (Life Technologies).

2.2. Metagenome library preparation and sequencing

Sequencing library was prepared using 1D Rapid PCR Barcoding Kit (SQK-RPK004) according to the manufacturer's instructions. PrimeSTAR GXL DNA polymerase (Takara, Japan) was utilized for amplification. The two libraries were sequenced for 14 hours on MinION device with FLO-MIN106 (R9.4) flowcell.

2.3. Taxonomic and functional analysis

Reads were preprocessed by base-calling and demultiplexing using Guppy v.2.3.5 (ONT). Those reads were quality-checked, filtered to remove sequence reads shorter than 500 bp and reads with an average quality score less than 7 using NanoFit v.2.5.0 [4], and MinKnow v.19.10.1 (ONT). For taxonomic identification and functional prediction, the filtered sequences were submitted to MG-RAST online server [5] using the M5nr database and subsystems based annotation with default setting, respectively. Cut-offs included a maximum E-value of $1 \times 10^{-5}$, and a minimum alignment length of 15 were used. KmerResistance v.2.2 [6] was used for antibiotic resistance genes analysis with 90% identity and 10% depth corr.

Declaration of Competing Interest

There is no conflict of interest to declare.

Acknowledgements

The project was supported by a research grant from the School of Medicine, Vietnam National University Ho Chi Minh City. The funder had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi: 10.1016/j.dib.2020.105731.

References

[1] Food and Agriculture Organization of the United Nations. Fisheries and Aquaculture Department. The state of world fisheries and aquaculture 2018: meeting the sustainable development goals. http://www.fao.org/documents/card/en/c/95404EN (accessed 14 April 2020).
[2] R. V. Pakingking Jr, E.G.T de Jesus-Ayson, B.O. Acosta, Addressing acute hepatopancreatic necrosis disease (AHPND) and other transboundary diseases for improved aquatic animal health in Southeast Asia: Proceedings of the ASEAN Regional Technical Consultation on EMS/AHPND and Other Transboundary Diseases for Improved Aquatic Animal Health in Southeast Asia, 22-24 February 2016, Makati City, Philippines. Tigbauan, Iloilo, Philippines: Aquaculture Dept., Southeast Asian Fisheries Development Center. (2016) 109. http://hdl.handle.net/10862/3096.

[3] K. Thornber, D. Verner-Jeffreys, S. Hinchliffe, M. M. Rahman, D. Bass, C. R. Tyler, Evaluating antimicrobial resistance in the global shrimp industry. Rev Aquac (2019) 1-21. doi:10.1111/raq.12367.

[4] W. De Coster, S. D’Hert, D. T. Schultz, M. Cruts, C. Van Broeckhoven, NanoPack: visualizing and processing long-read sequencing data. Bioinformatics 34(15) (2018) 2666–2669. doi:10.1093/bioinformatics/bty149.

[5] F. Meyer, D. Paarmann, M. D’Souza, R.A. Edwards, The metagenomics RAST server—a public resource for the automatic phylo- genetic and functional analysis of metagenomes, BMC Bioinform. 9 (2008) 386. https://doi.org/10.1186/1471-2105-9-386.

[6] P.T. Clausen, E. Zankari, F.M Aarestrup, O. Lund, Benchmarking of methods for identification of antimicrobial resistance genes in bacterial whole genome data. J Antimicrob Chemother 71(9) (2016) 2484-2488. doi:10.1093/jac/dkw184.