



**QUEEN'S
UNIVERSITY
BELFAST**

Editorial: Methane: A Bioresource for Fuel and Biomolecules

Kalyuzhnaya, M. G., Kumaresan, D., Heimann, K., Caetano, N. S., Visvanathan, C., & Parthiba Karthikeyan, O. (2020). Editorial: Methane: A Bioresource for Fuel and Biomolecules. *Frontiers in Environmental Science*, 8, [9]. <https://doi.org/10.3389/fenvs.2020.00009>

Published in:
Frontiers in Environmental Science

Document Version:
Publisher's PDF, also known as Version of record

Queen's University Belfast - Research Portal:
[Link to publication record in Queen's University Belfast Research Portal](#)

Publisher rights

© 2020 The Authors.

This is an open access article published under a Creative Commons Attribution License (<https://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution and reproduction in any medium, provided the author and source are cited.

General rights

Copyright for the publications made accessible via the Queen's University Belfast Research Portal is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The Research Portal is Queen's institutional repository that provides access to Queen's research output. Every effort has been made to ensure that content in the Research Portal does not infringe any person's rights, or applicable UK laws. If you discover content in the Research Portal that you believe breaches copyright or violates any law, please contact openaccess@qub.ac.uk.



Editorial: Methane: A Bioresource for Fuel and Biomolecules

Marina G. Kalyuzhnaya¹, Deepak Kumaresan², Kirsten Heimann³, Nidia S. Caetano⁴, Chettiyappan Visvanathan⁵ and Obulisamy Parthiba Karthikeyan^{6,7*}

¹ Department of Biology, San Diego State University, San Diego, CA, United States, ² School of Biological Sciences, Queen's University of Belfast, Belfast, United Kingdom, ³ Centre for Marine Bioproducts Development, College of Medicine and Public Health, Flinders University, Adelaide, SA, Australia, ⁴ Department of Chemical Engineering, Instituto Superior de Engenharia Do Porto, Porto, Portugal, ⁵ Department of Energy, Environment, and Climate, School of Environment, Resources and Development, Asian Institute of Technology, Pathumthani, Thailand, ⁶ Department of Civil and Environmental Engineering, University of Michigan, Ann Arbor, MI, United States, ⁷ Department of Engineering Technology, College of Technology, University of Houston, Houston, TX, United States

Keywords: methane, methanotrophs, electron transfer, bioreactor, value addition and sustainability

OPEN ACCESS

Edited by:

Hyung-Sool Lee,
University of Waterloo, Canada

Reviewed by:

Seung Gu Shin,
Pohang University of Science and
Technology, South Korea
Wei-Min Wu,
Stanford University, United States

*Correspondence:

Obulisamy Parthiba Karthikeyan
opkens@gmail.com;
pobulisa@umich.edu

Specialty section:

This article was submitted to
Microbiotechnology, Ecotoxicology
and Bioremediation,
a section of the journal
Frontiers in Environmental Science

Received: 18 October 2019

Accepted: 13 January 2020

Published: 05 February 2020

Citation:

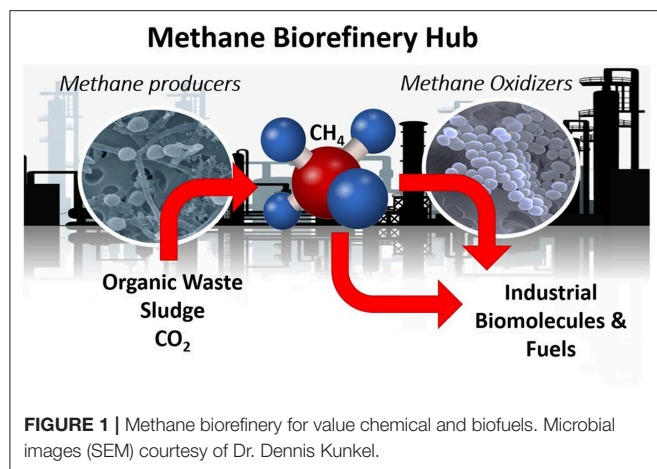
Kalyuzhnaya MG, Kumaresan D,
Heimann K, Caetano NS,
Visvanathan C and
Parthiba Karthikeyan O (2020)
Editorial: Methane: A Bioresource for
Fuel and Biomolecules.
Front. Environ. Sci. 8:9.
doi: 10.3389/fenvs.2020.00009

Editorial on the Research Topic

Methane: A Bioresource for Fuel and Biomolecules

Methane (CH₄), a highly reduced C1 compound, is one of the long-lived atmospheric gases with high global warming potential i.e., 28–36 times that of CO₂ over 100 years. The atmospheric levels of CH₄ reached ~1863 part per billions (ppb) in 2014, and annual increase of atmospheric CH₄ level thereafter measured as ~10 ppb. The CH₄ is projected to drive the rise in global temperature of ~4–6°C by 2050, and thus it is currently considered as the main target for global climate stabilization and mitigation (COP-21, 2015). Capturing anthropogenic CH₄ to produce value products is highly feasible, but the great challenge is that to tap, concentrate, purify, store, transport, and utilize the CH₄ from different point emission sources is presently not economically viable.

In this special issue, a conceptual model of “Methane-Biorefinery Hub” is proposed as a sustainable development to mitigate the CH₄ emissions from the most significant anthropogenic sources. The idea of “Methane-Biorefinery Hub” is deeply rooted in natural CH₄ production and consumption processes (Figure 1). While in general, with a few exceptions, the natural CH₄ cycle is balanced, the anthropogenic disturbances have typically led to increase the CH₄ emissions. Thus, better understanding of mechanisms that control CH₄ cycle in nature can be used to engineer better systems in human-built environments. For example, Holmes et al. established that the bacteria (donors) and archaea (acceptors) communicate through nano-wires or electron transfer molecules i.e., the electrons transfer through e-pili (i.e., direct interspecies electron transfer), while protons diffusion (direct interspecies hydrogen transfer) is regulated by the partial pressure of the bioreactor system. Mimicking natural processes in laboratory/pilot-scale bioreactors that are designed to optimize specific operational conditions to regulate such communication may lead to successful implementation of technology to effectively utilize the waste materials e.g., municipal solid waste, food waste, industrial organics and wastewaters, or low-grade coals with/without the aid of external carbon/electron sources to produce CH₄ as bio-energy, as shown by Yang et al. As pointed out by Wojcieszak et al., well-balanced microbial consortium are crucial for efficient biogas production, and inoculum sampled from typical methanogenic environments can be gradually adapted to industrial installations to allow effective biogas production. Addition of microbial supplements, metals/nutrients, organic sources (Zaldívar Carrillo et al.), electron



conductive materials or electrodes, and altering the head-space gas composition (and partial pressure) are proposed as options for facilitating electron transfer and microbial interactions, while it may also change the equilibrium between dissolved vs. gas phase CH₄ concentrations within the system.

Another unique approach for recovering dissolved CH₄ using degassing membrane contactors was proposed by Velasco et al.. While still at the stage of infancy, this unique technology highlights limitless opportunities for innovative approaches in CH₄ capturing. If not recovered effectively, the dissolved CH₄ and manipulating conditions may be expected to fuel the anaerobic methane oxidizers (ANME), and also sulfate reducing bacteria (SRB), to thrive and compete with methanogens for electrons/protons that may lead to low CH₄ yield. The ANME possess a reverse methanogenesis process, i.e., utilizing CH₄ (and CO₂) as carbon sources to produce acetate or other products. On the other hand SRB compete with methanogens for similar substrates or electron donors. So, there may not be a competition between SRB and ANME for substrate, but it required investigation. On the other hand, the industrial applications of ANME are challenging, and still limited by the number of unresolved biochemical questions. However, an example of how a solid understanding of enzyme kinetics and energy transfer between the microbial communities can be used to manipulate the operating conditions to either facilitate or eliminate methane production was presented by Grisewood et al.. Alternatively, establishment of co-cultures of methanogens and ANME to produce value chemicals from anaerobic digestion processes was proposed and validated using a newly developed mathematical model by Nazem-Bokaei and Maranas, which, however, still relies on the electron coupling theory. Once fully understood, the ANME-based approach could be a viable option for reducing CH₄ emissions from natural settings, while aerobic oxidation is recommended for industrial-scale organic digestion facilities.

The aerobic CH₄ oxidation process is easily coupled with digesters, while the process can yield a number of value products such as polymers, organic acids, single cell protein

(SCP), compatible solutes, short/long chain fatty acids, omega fatty acids, vitamins, methanol, formate, etc. Methanotrophs are classified into Group I and Group II based on their physiology. Group I methanotroph e.g., *Methylococcus capsulatus* (Bath) is reported to produce SCP, while Group-II methanotrophs (e.g., *Methylosinus trichosporium* OB3b) are capable of accumulating biopolymers/SFA/LFA from CH₄. An example of coupled production is highlighted by the work of Henard et al.. Metabolic modeling can further empower application of natural CH₄ consuming bacteria, as exemplified by Lieven et al.. While it is widely acknowledged that the natural capacities of microbial systems (as axenic or mixed cultures) in CH₄-consumption and accumulation of value products are governed by a number of factors e.g., carbon, nutrients, and metals, we are only now applying the knowledge for improving their industrial potential. Further, an example of how nitrogen starvation activates polyhydroxyalkanoate accumulation and alters the fatty acid compositions in biomass is provided by Tays et al.. The metal-switch impacts on the key enzyme activities, kinetics, the internal electron pool, and the carbon flux e.g., are described in the work by Akberdin et al..

Nevertheless, the industrial applications and innovations remain to be challenged by long-standing fundamental questions regarding CH₄ biocatalysts; including: (a) source of electron donors for CH₄ activation as well as electron acceptors for process intensification; (b) regulation of contaminations or development of efficient strategies for controlling natural communities or synthetic co-cultures; (c) improved genetic traceability of methanotrophs in a mixed consortium and (d) coupling of CH₄-conversion potential with efficient nitrogen fixation and denitrification. We would like to acknowledge significant progress in developing systems biology toolbox for manipulating methanotrophic bacteria, as well as new advances in overcoming technological bottlenecks related to CH₄ mass-transfer limitation. Yet, some challenges still remain. New developments highlight additional need for further research, specifically in areas of O₂-capturing, cell immobilization, CH₄ and CO₂ conversion by coculture of methanotrophs and algae/methylotrophs, coupling CH₄ conversion and electrocatalysis. Algae and methanotrophs exchange nutrients during co-culturing are beneficial, while methylotrophs helps to alleviate any methanol toxicity.

The papers published in this special issue confirm that CH₄ emission/production is related to the microbiomes of the system, which are easy to be manipulated through biological/chemical augmentation methods. By coupling CH₄ production with either ANME or methanotrophs, emissions could be significantly reduced and high-value products could be recovered through an integrated “Methane-Biorefinery” approach. Interestingly, in the work by Burton et al. it was concluded that the aerobic methanotrophs, methylotrophs and methanogenic archaea found to have common lineage i.e., use tetrahydromethanopterin (H₄MPT) and/or tetrahydrofolate (H₄F) as coenzymes in one-carbon (C1) transfer pathways that have been overlooked in the past and the relationships need to be well-studied for harnessing the benefits under “Microbiome” theory.

AUTHOR CONTRIBUTIONS

OP and MK written this editorial. NC, KH, CV, and DK edited the final text. All authors approved the final version.

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2020 Kalyuzhnaya, Kumaresan, Heimann, Caetano, Visvanathan and Parthiba Karthikeyan. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.