

Dipicolinic acid as a tracer for thermophilic endospores and hydrocarbon seeps in deep water marine sediments

Jayne. E. Rattray¹, Gretta Elizondo¹, Anirban Chakraborty¹, Bernie Bernard², James Brooks², Adam MacDonald⁴, Calvin Campbell⁴, Alexandre Normandeau⁵, Martin Fowler⁶, Jamie Webb⁶ and Casey R.J. Hubert¹.

¹University of Calgary, Canada, ²TDI Brooks International, USA, ⁴Nova Scotia Department of Energy and Mines, Canada, ⁵Bedford Institute of Oceanography, Canada, ⁶Applied Petroleum Technology Ltd., Canada.

Introduction

Understanding the sediment biogeography of dormant marine thermophilic bacterial endospores (thermospores) has the potential to assist locating and characterising working petroleum systems. The presence of thermospores in cold ocean environments suggests that distribution is governed by spore dispersal via advective hydrocarbon seepage sourced from deep hot oil reservoirs. Low abundance and endospore coat physiology mean nucleic acid based microbiological surveillance techniques have limited success for in situ detection of thermospores. The biomarker 2,6-pyridine dicarboxylic acid (dipicolinic acid or DPA) is specific to endospore-forming bacteria from the phylum Firmicutes, and constitutes a significant percentage of endospore dry weight. DPA is therefore a potential biomarker for sediment dwelling endospores, and in particular for detecting anomalies due to oil reservoir-derived thermospores. If so, DPA could have utility in locating seabed hydrocarbon seeps, however its suitability for such seabed screening has so far not been tested.

Method

To address the efficacy of DPA biomarker analysis as a tool for hydrocarbon seep location we established a modified Tb³⁺ chelation method (Lomstein and Jørgensen, 2012). Sediment samples were extracted using complete digestion with acid hydrolysis, chelated with Tb³⁺ and analysed using HPLC coupled to a fluorescence detector measuring at 270 nm emission and 545 nm excitation.

Results

DPA distribution was assessed in deep seabed sediment samples from 97 locations in the Eastern Gulf of Mexico (Figure 1). 16S rRNA gene amplicon libraries of thermophilic spore formers from high temperature sediment incubations (Chakraborty et al., 2018) were used to assess whether DPA detection in the sediments could be associated with the presence of different thermophilic spore forming bacteria (i.e. assessing the likelihood of DPA originating from mesophilic vs thermophilic endospores). DPA concentrations were compared with

GeoConvention 2020

geochemistry and available seep data. Additional sediment samples from along the Scotian shelf and Laurentian channel in Atlantic Canada provided both hydrocarbon positive and negative sediment cores, and enabled higher resolution down-core DPA depth profiles. Sediment cores from deeper water showed higher and more variable concentrations of DPA down core at hydrocarbon-positive stations compared to hydrocarbon-negative stations.

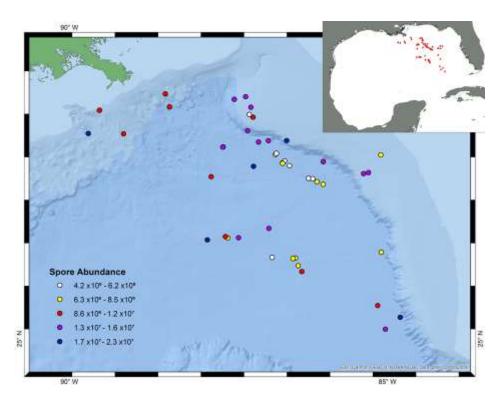


Figure 1. Distribution of endospore abundance in the Eastern Gulf of Mexico surface sediment calculated using spore specific DPA as a biomarker.

Novel Information

The efficacy of DPA for tracing thermospores associated with hydrocarbon seeps in marine sediments has undergone preliminary assessment and we propose that DPA has potential as a biomarker for assisting in locating hydrocarbon systems in deep water marine environments, based on hydrocarbon seeps being point sources for thermophilic endospores.

GeoConvention 2020 2



Acknowledgements

This work was supported by Genome Canada (GAPP), Canada Foundation for Innovation 568 (CFI-JELF 33752) and Campus Alberta Innovates Program Chair funding to CRJH.

References

Chakraborty, A., Ellefson, E., Li, C., Gittins, D., Brooks, J. M., Bernard, B. B., and Hubert, C. R. J.: Thermophilic endospores associated with migrated thermogenic hydrocarbons in deep Gulf of Mexico marine sediments, The ISME journal, 12, 1895-1906, 2018.

Lomstein, B. A. and Jørgensen, B. B.: Pre-column liquid chromatographic determination of dipicolinic acid from bacterial endospores, Limnol. Oceanogr. Meth. 10, 227-233, 2012.

GeoConvention 2020 3