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A molecular ecology approach using Stable Isotope Probing and metagenomics to study viruses of methanogens' diversity

H. Ngo¹, M. Sotomski¹, M. Krupovic², F. Enault³, O. Chapleur¹, T Bouchez¹, A. Bize^{1*} ¹ Irstea, UR PROSE, 1 rue Pierre-Gilles de Gennes, F-92761 Antony, France ² Institut Pasteur, BMGE, 28, rue du Docteur Roux, 75724 Paris Cedex 15, France ³ UCA, LMGE, 1 impasse Amélie Murat, TSA 60026 CS 60026, 63178 Aubière Cedex, France *ariane.bize@irstea.fr



A. Bize

Context

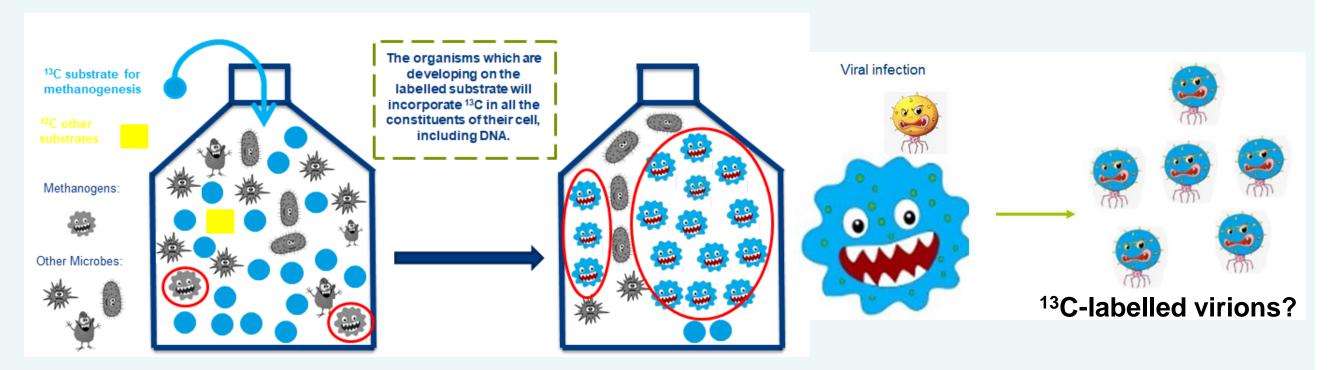
Istea

Identifying hosts from viruses in complex microbial communities is still non-trivial

- Very promising approaches have recently emerged to identify hosts from viruses in complex ecosystems (1, 2)
- However, most of them require either host cultivability or prior knowledge of host and virus partial genome sequence
- Moreover, none of them establish a direct link with the host metabolism

Stable Isotope Probing (SIP) should enable to identify viruses infecting hosts which assimilate a specific substrate, within complex ecosystems

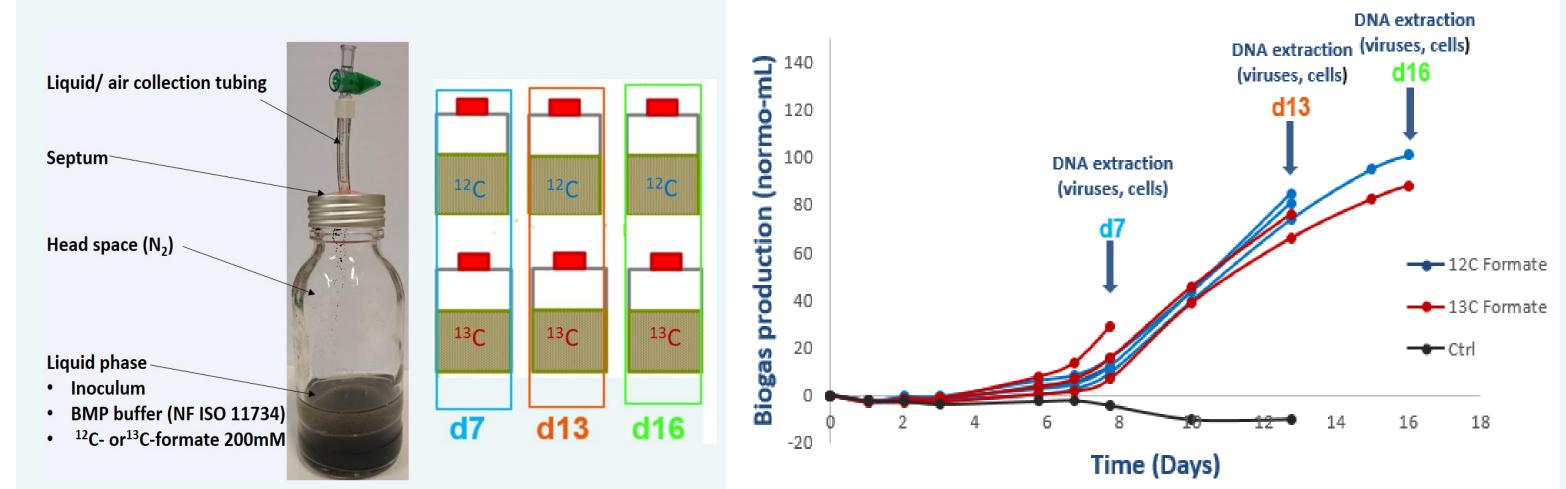
The use of various ¹³C-labeled methanogenesis substrates enables us to activate different populations of methanogenic archaea and enrich them with ¹³C (3). Logically, viruses that infected them should also be labelled. The viral DNA will be separated by ultracentrifugation on a CsCl gradient (4), collected and further studied (sequencing, ...).



Proof of concept with a simple biological model

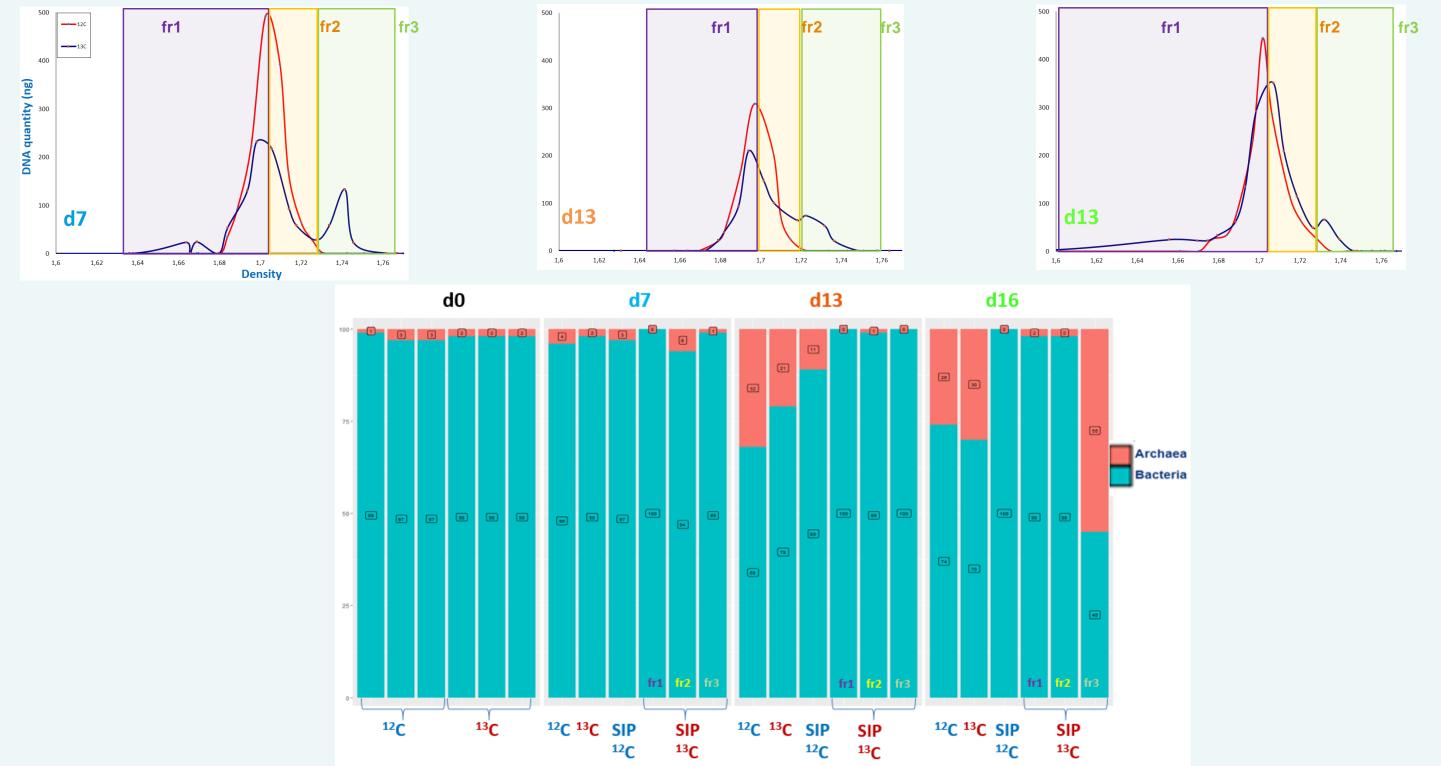
Test of SIP applied to organic waste anaerobic digestion microbial communities

Incubation with ¹²C- or ¹³C-formate as substrate



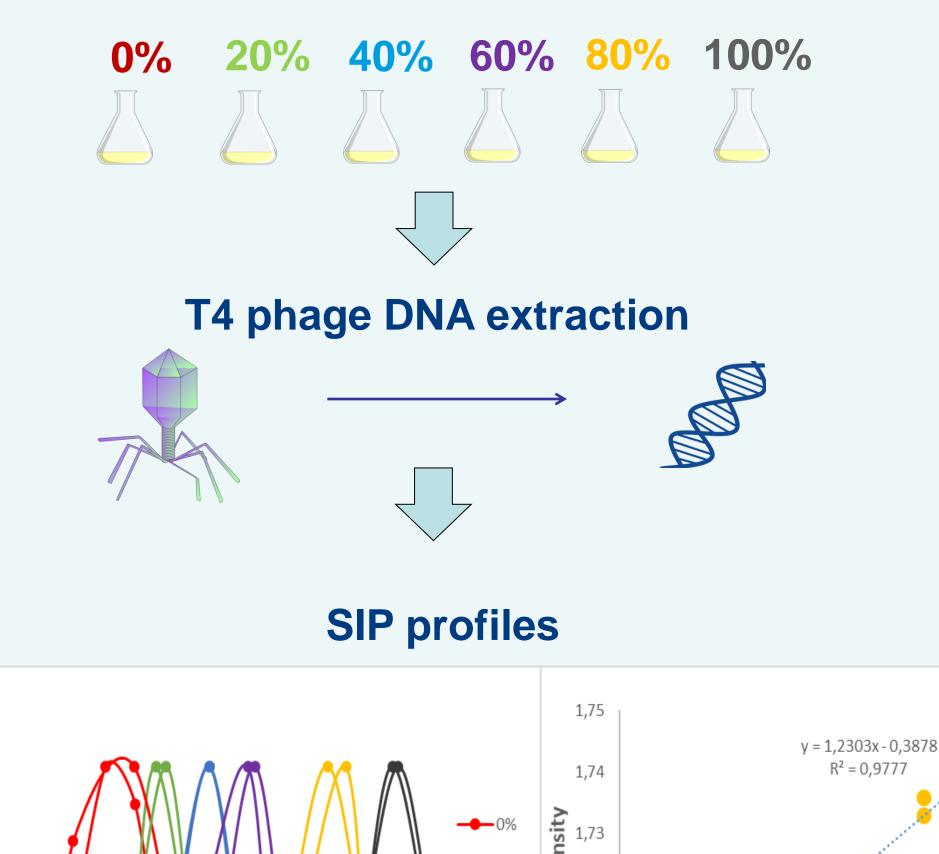
- Good reproducibility of biogas production
- DNA extraction at 3 different time points to select for the samples with the optimal amounts of enriched DNA from archaeal cells or viruses

SIP profiles of cellular DNA and 16S metabarcoding analysis

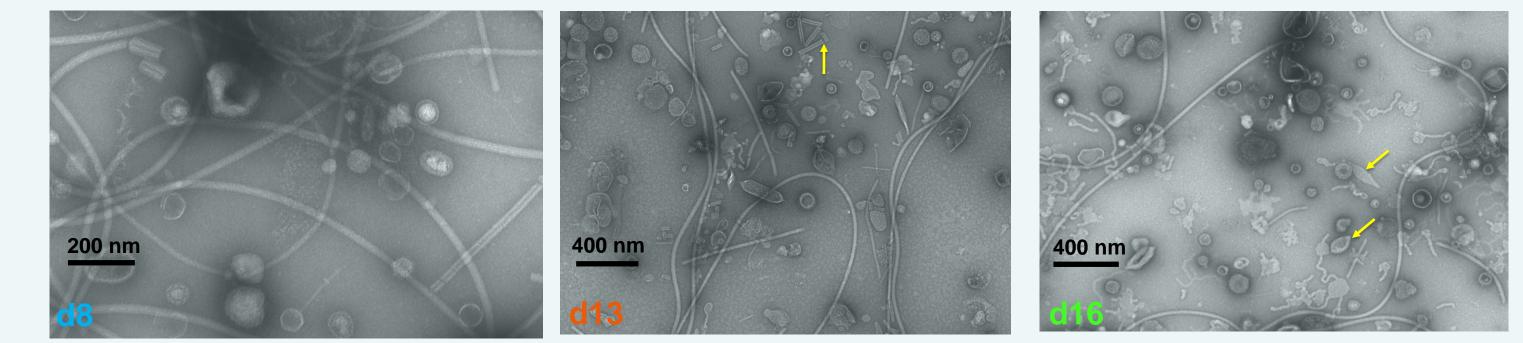


T4 bacteriophage (DSM 613) production on Escherichia coli B cells (DSM 4505) grown in M9 minimal medium

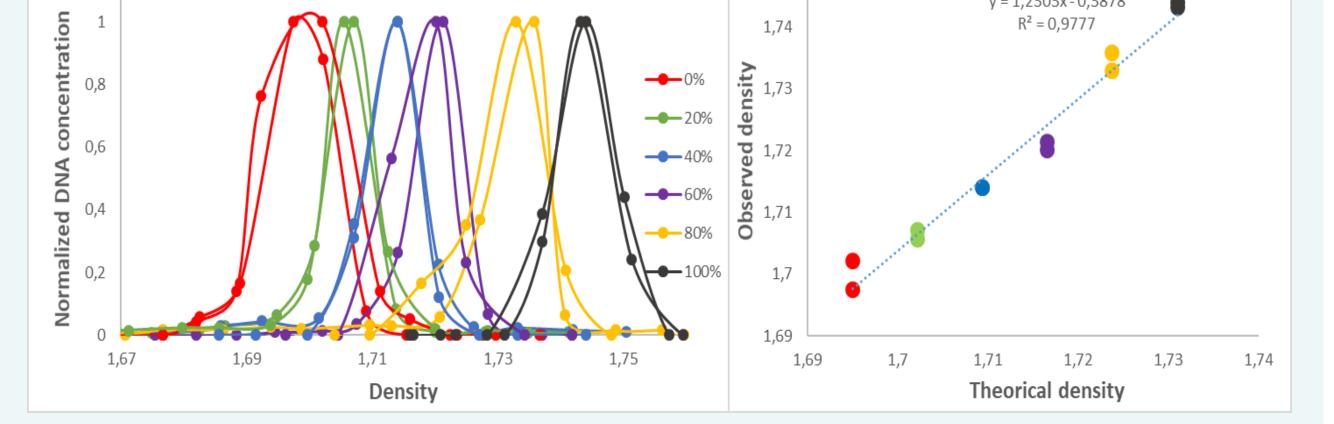
% of ¹³C-glucose in the growth medium



- Archaea were mostly methanogens and their proportion increased over time
- The highest proportion of archaea was reached in the fr3 ¹³C fraction at d16 Virions observation (TEM)



At d13-d16, virus-like particles with morphotypes similar to archaea-specific



- Significant differences in positions of density peaks \bullet
- Good agreement between the expected and observed peak \bullet positions, according to the ¹³C-glucose percentages

viral families were observed

No significant differences in viral DNA quantities collected at d7, d13 or d16

Perspectives

Perform the SIP analysis of viral DNA and sequence the fractions of interest Identify contigs of archaeal viruses thanks to bioinformatics analysis of the metavirome sequences (pipeline development).

References

1,2

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Project