



**HAL**  
open science

## Design of a functional DNA microarray targeting the methanogenic community of complex anaerobic ecosystems

Ourdia Bouzid, Corinne Biderre-Petit, Eric Dugat-Bony, Mohieddine Missaoui, Diego Morgavi, Pascale Mosoni, Evelyne Forano, Gérard Fonty, Pierre Peyret

### ► To cite this version:

Ourdia Bouzid, Corinne Biderre-Petit, Eric Dugat-Bony, Mohieddine Missaoui, Diego Morgavi, et al.. Design of a functional DNA microarray targeting the methanogenic community of complex anaerobic ecosystems. 7. Joint Symposium of Rowett-INRA 2010, Jun 2010, Aberdeen, United Kingdom. 2010. hal-02823797

**HAL Id: hal-02823797**

**<https://hal.inrae.fr/hal-02823797v1>**

Submitted on 6 Jun 2020

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



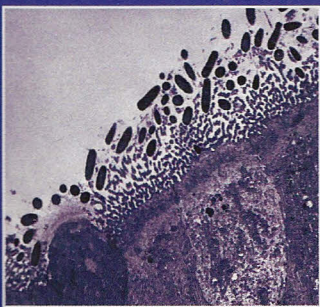
Rowett Institute  
of Nutrition and Health  
University of Aberdeen



**Rowett-INRA 2010**

**23-25 June 2010**

**Aberdeen Exhibition and  
Conference Centre**



**Gut Microbiology:**  
new insights into  
gut microbial ecosystems

7th Joint Symposium organised by the **Rowett Institute of Nutrition and Health**, University of Aberdeen, Scotland (UK) & the **Institut National de la Recherche Agronomique**, Clermont-Ferrand-Theix (France)

### Design of a functional DNA microarray targeting the methanogenic community of complex anaerobic ecosystems

Ouardia Bouzid<sup>1</sup>, Corinne Biderre-Petit<sup>3</sup>, Eric Dugat-Bony<sup>3</sup>, Mohieddine Missaoui<sup>3</sup>, Diego P. Morgavi<sup>1</sup>, Pascale Mosoni<sup>2</sup>, Evelyne Forano<sup>2</sup>, Gérard Fonty<sup>3</sup>, Pierre Peyret<sup>3</sup>  
<sup>1</sup>INRA-UR1213 Herbivores, St Genès Champanelle/Auvergne, France, <sup>2</sup>INRA-UR 454 Microbiology, St Genès Champanelle/Auvergne, France, <sup>3</sup>Laboratoire « Microorganismes : Génome & Environnement », LMGE, UMR CNRS 6023, Aubière/Auvergne, France

Methane is a potent greenhouse gas produced in the environment, e.g. natural anoxic lakes and marshes, but the most important source is due to human activity. Farm animals, particularly ruminants, produce substantial amounts of methane. In gastrointestinal tracts, methane is produced by methanogenic archaea when feeds are fermented. To improve our understanding of the rumen methanogenic community, we developed a DNA microarray based on a functional gene involved in methanogenesis. In this context, 530 oligonucleotides (24 to 59 mers) targeting the methyl-coenzyme M reductase alpha subunit gene (*mcrA*) were designed using Metabolic Design, HISPOD and GoArrays softwares. These dedicated softwares developed in our laboratory allowed the design of specific probes targeting known sequences and also explorative probes capable of hybridizing to sequences not yet available in databases but potentially present in complex communities. This functional microarray covers the diversity of the five known methanogenic archaeal orders: Methanobacteriales, Methanomicrobiales, Methanosarcinales, Methanopyrales and Methanococcales. It was validated using *mcrA* gene fragments amplified from members of the Methanobacteriales, Methanomicrobiales, Methanosarcinales and Methanococcales. Our results showed an excellent specificity and sensitivity of the designed probes towards the species tested, providing a species-specific fingerprint tool. This DNA microarray, developed to study the structure and dynamics of methanogens, can be applied to the rumen and other digestive tracts but also to a wide variety of aquatic and terrestrial methanogenic ecosystems.

### Correlation of rumen microbial community structure with milk fatty acid composition

John Wallace<sup>1</sup>, Graham Horgan<sup>1</sup>, Stefan Muetzel<sup>1</sup>, Kate Crosley<sup>1</sup>, Delphine Paillard<sup>1</sup>, Ian Givens<sup>2</sup>, Kevin Shingfield<sup>3</sup>  
<sup>1</sup>University of Aberdeen Rowett Institute of Nutrition and Health, Aberdeen AB21 9SB, United Kingdom, <sup>2</sup>School of Agriculture, Policy and Development, University of Reading, Reading RG6 6AR, United Kingdom, <sup>3</sup>Animal Production Research, MTT Agrifood Research Finland, 31600, Jokioinen, Finland

The conversion of dietary polyunsaturated fatty acids (PUFA) to saturated fatty acids in the rumen (biohydrogenation) is of concern because of the perceived positive implications that PUFA and conjugated linoleic acid (CLA) in ruminant products have on human health and the negative implications of saturated fatty acids. Bacteria of the *Butyrivibrio* group are thought to be primarily responsible for biohydrogenation. Other bacteria and fungi have sometimes been implicated in biohydrogenation, but the relative importance of these organisms in determining the fatty acid composition of ruminant products has never been quantified. The aim of this work was to compare milk fatty acid composition with the composition of the rumen microbial community, rumen metabolites and feed composition across 7 dairy trials carried out in MTT, Jokioinen, Finland and the University of Reading, UK. The question being asked was - "Can we relate milk fatty acid composition to the composition of the ruminal microflora, even across diets?" Cows received basal diets of maize or grass silage supplemented with concentrates and vegetable or fish oil. Total bacteria, *Butyrivibrio* spp., *Streptococcus bovis*, *Propionibacterium acnes* and *Megasphaera elsdenii* were analyzed by qPCR. All data were compiled into a common dataset comprising >100 individual parameters. A multivariate analysis 'heat-map' was constructed to illustrate correlations between measurements. Within trials, there were associations between bacterial population densities and milk fatty acid composition, but between-trial comparisons revealed a predominant influence of diet rather than microbial numbers. The unequivocal answer to the question posed above is, "No, diet is most important."