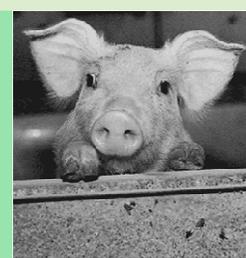




Discrepancies in microbiota composition along the pig gastro-intestinal tract between *in vivo* observations and an *in vitro* batch fermentation model



C. Boudry*, C. Poelaert*, D. Portetelle†, A. Thewis*, J. Bindelle*

*Animal Science Unit, Gembloux Agro-Bio Tech, University of Liege, Belgium

†Animal and Microbial Biology Unit, Gembloux Agro-Bio Tech, University of Liege, Belgium

Introduction

In vitro fermentation models are increasingly used to assess prebiotic potential of novel indigestible carbohydrates (CHO)

However, the evolution of the microbiota during *in vitro* fermentation and along the gut intestinal tract *in vivo* have never been properly compared

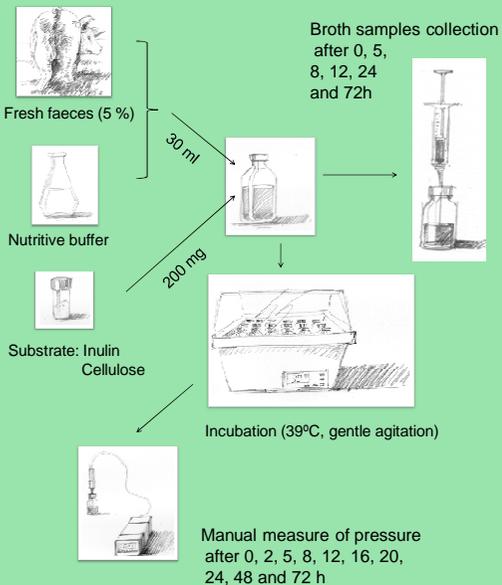
Material and Methods

***In vivo* trial:** 3 X 4 pigs received 3 diets for 3 weeks :

- (i) Semi-purified diet + 5 % Inulin
- (ii) Semi-purified diet + 5 % Cellulose
- (iii) Semi-purified diet + 2.5 % Inulin + 2.5 % Cellulose

At slaughter: Digesta collected along the digestive tract (Jejunum, Ileum, Caecum and 3 parts of the Colon)

***In vitro* trial:** gas fermentation with fresh faeces of the *in vivo* trial



Analyzes

Fermentation Kinetics (modelisation)

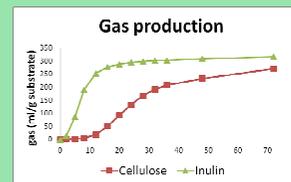
DNA Extraction on digesta and fermentation broth samples

Microflora quantification by qPCR :

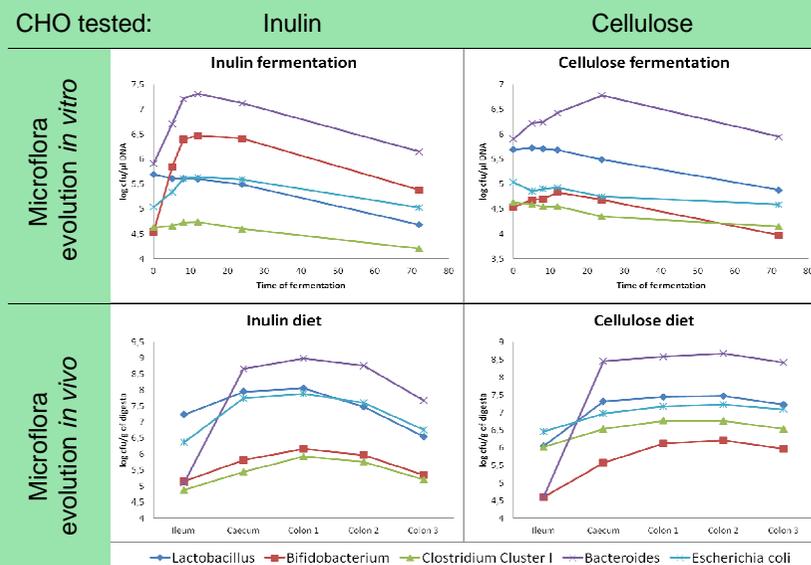
- ✓ *Lactobacillus* sp.
- ✓ *Bifidobacterium* sp.
- ✓ *Clostridium* Cluster I
- ✓ *Bacteroides*
- ✓ *Escherichia coli*

Results

Contrasted fermentation kinetics were observed for the 2 substrates *in vitro*



Microflora evolutions in both systems were consistent for *Bifidobacteria*, *Bacteroides* and *E. coli* in the first part of the digestive tract but not for *Lactobacillus* and *Clostridium* Cluster I.



Conclusions

Lactobacillus and *Clostridium* CL I evolutions are different between both systems probably due to specific bacterial properties.

➔ Improves to the *in vitro* model are needed to assess properly prebiotic potential of new CHO.

