

# Galactoglucomannan fibres promote a beneficial porcine gut microbiome

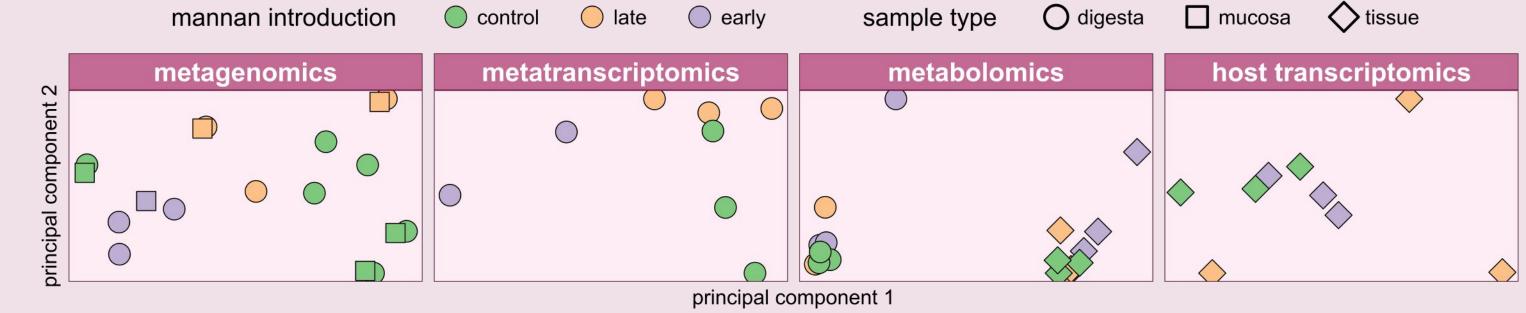
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Most mammals and their microbiomes are codependent, forming a functional unit known as a holobiont. Exchanging metabolites, regulating gene expression, and combating pathogens are vital to the health and performance of the holobiont 1-2. By understanding the interactions occurring within these complex systems, we can more effectively improve animal and feed production, favouring both animal welfare, production efficiency, and the growing needs of the World's increasing human population <sup>3-4</sup>.

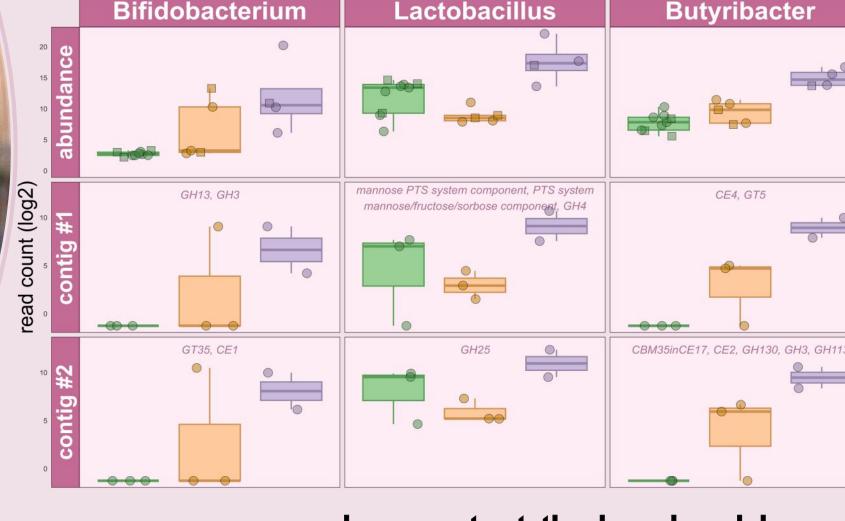
Mannan fibres made from spruce can be host-accessible into metabolised compounds by microbes with carbohydrate-active **enzymes** 5-6. These microbes also ease piglets' transition from milk to solid feed 7. Hence we ask: can we jump-start the young microbiome mannan pre-weaning diet? spoiler! --



principal component analyses show gradients that correspond with duration of mannan exposure. Among the differentially abundant populations in control and mannan-fed piglets are

> Bifidobacterium longum (log<sub>2</sub> fold change 8.3), Lactobacillus johnsonii (LFC 5.9), and Butyribacter sp. (LFC 5.7). Many of their differentially expressed genes yield enzymes for mannan degradation, like glycoside hydrolases and transferases,

and carbohydrate esterases.



Thus yes, we can jump-start their microbiome!

HOLO-OMIC MODEL

## THE TRIAL

We used three groups of 10-day old piglets and gave fibres to two groups: one starting before and one after weaning.

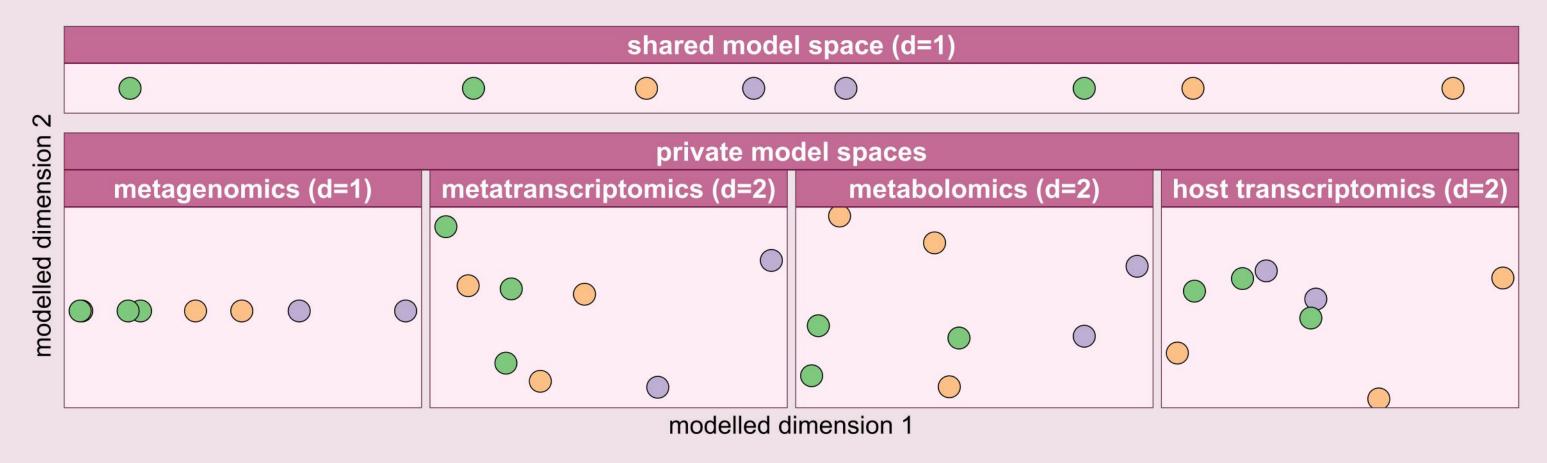


3 x late two weeks with mannan

3 x early two weeks with mannan

After one month, we sampled their caeca and generated four layers: metagenomics, metatranscriptomics, metabolomics, and host transcriptomics. The datasets were analysed both as individual omic layers, and jointly through a holoomic approach. Utilised methods were R-implemented tools for principal component 8 and differential abundance 9 analyses, and a Python-based package for holo-omic modelling through multiset correlation and factor analysis (MCFA) 10.

The fitted MCFA model reconstructs the full dataset as matrices across three feature spaces: one shared by all omic layers; one private to each dataset; and an omic-specific residual. The inferred features fit well with mannan exposure, hence the model will be used to learn more about the interactions across the porcine holo-omic boundary.



**REFERENCES 1** Rosenberg et al. The hologenome concept of evolution after 10 years. Microbiome. 2018;6(1):78. **2** Roughgarden et al. Holobionts as Units of Selection and a Model of Their Population Dynamics and Evolution. Biol Theory. 2018;13(1):44–65. **3** Simon et al. Host-microbiota interactions: from holobiont theory to analysis. Microbiome. 2019;7(1):5. **4** Alberdi et al. Disentangling host-microbiota complexity through hologenomics. Nat Rev Genet. 2022;23(5):281–97. **5** La Rosa et al. The human gut Firmicute Roseburia intestinalis is a primary degrader of dietary β-mannans. Nat Commun. 2019;10(1):905. **6** Michalak et al. Microbiota-directed fibre activates both targeted and secondary metabolic shifts in the distal gut. Nat Commun. 2020;11(1):5773. **7** Vasquez et al. Gut microbiome-produced metabolites in pigs: a review on their biological functions and the influence of probiotics. J Anim Sci Technol. 2022;64(4):671–95. **8** stats [base R, v.4.2.1]. **9** DESeq2 [R package, v. 1.38.3], doi: 10.18129/B9.bioc.DESeq2. **10** MCFA [Python package v. 1.0.1], doi: 10.5281/zenodo.8128339.







