

IDENTIFICATION OF A CORE METAGENOME THROUGH FUNCTIONAL ANNOTATION IN SWINE AND WILD BOARS



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- Hypothesis 1

A functional core metagenome can be defined for both species

- Hypothesis 2

Swine and wild boars have a different functional core metagenome

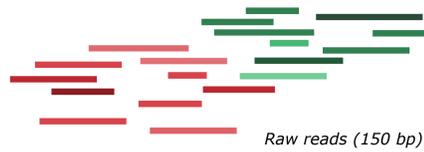
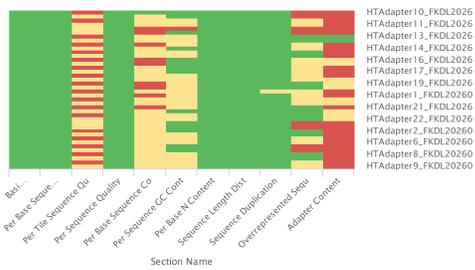


Wild boar samples from Poland
Number of samples: 6

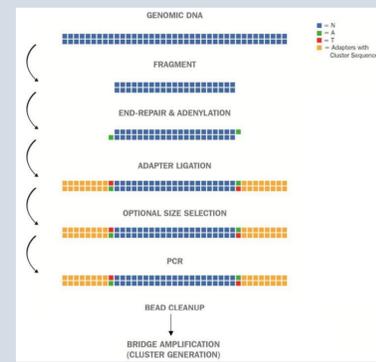


Swine samples from France
Number of samples: 12

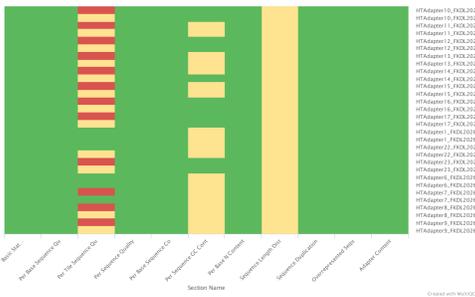
Multi FastQC before preprocessing [1,2]



LAB WORK AND SEQUENCING



Multi FastQC after preprocessing [1,2]



Host removal:
- Swine (*Sus scrofa domestica*) → [GCA_000003025.6](#)
- Wild boar (*Sus scrofa scrofa*) → [GCA_006511355.1](#)



- Filter those reads with predicted gene ontology
- Use elim algorithm with Kolmogorov-Smirnov method to find the most specific (down in the hierarchy) gene ontologies that are significant.
- Count number of samples where a significant ontology appears

RESULTS

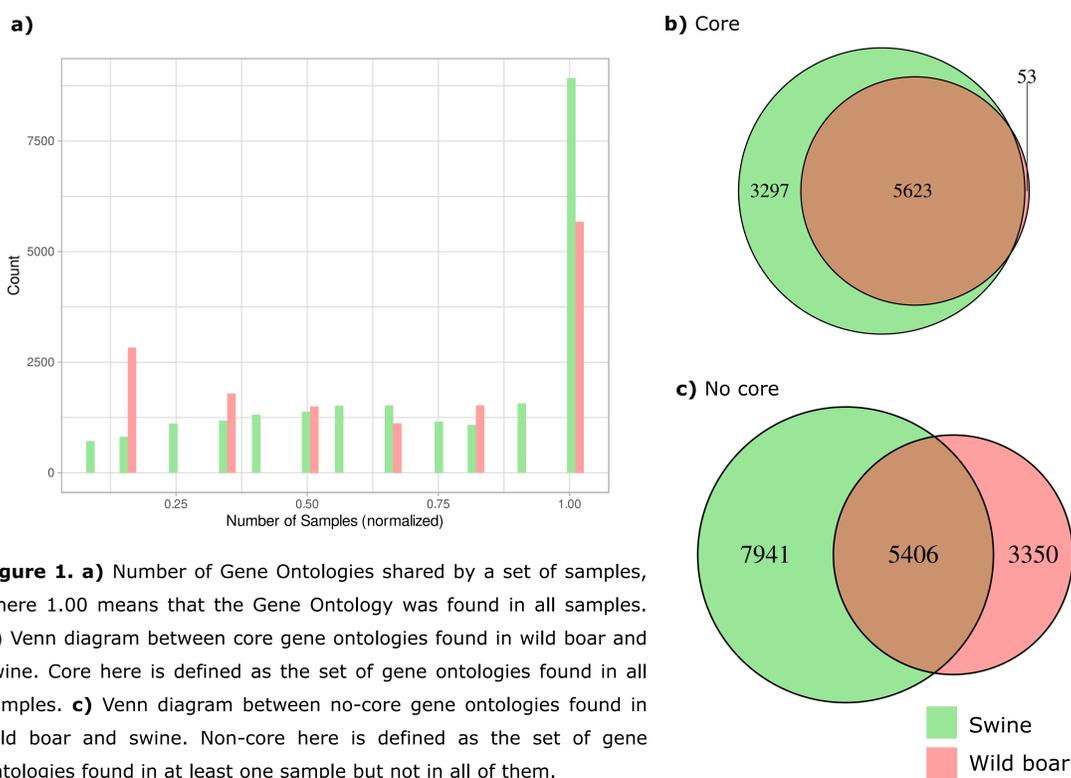


Figure 1. a) Number of Gene Ontologies shared by a set of samples, where 1.00 means that the Gene Ontology was found in all samples. b) Venn diagram between core gene ontologies found in wild boar and swine. Core here is defined as the set of gene ontologies found in all samples. c) Venn diagram between non-core gene ontologies found in wild boar and swine. Non-core here is defined as the set of gene ontologies found in at least one sample but not in all of them.

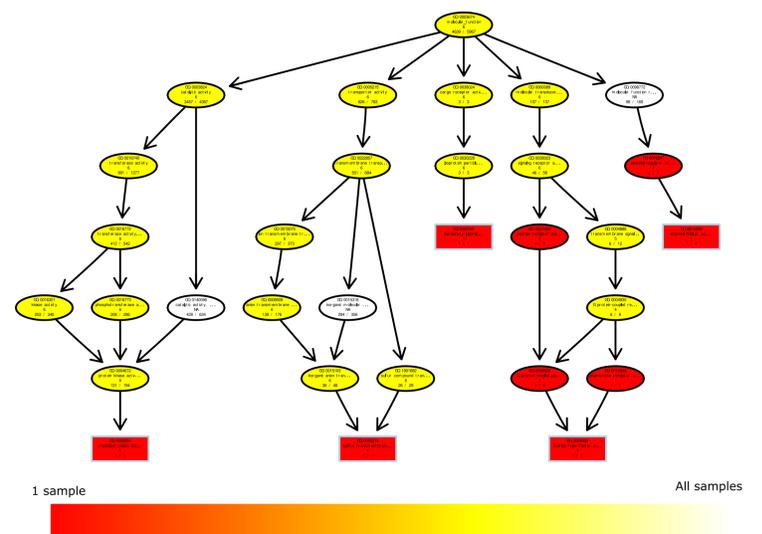


Figure 2. Wild boar gene ontology tree of significant Gene Ontologies scored by number of samples where they appear. Here the number of tips is 5, but for it to be meaningful this number should be increased. However, it was impossible for us to keep it readable with more tips using topGO.

- Hypothesis 1

A functional core metagenome can be defined for both species. Most of the Gene Ontologies are shared by all samples for both species.

- Hypothesis 2

The core Venn diagram suggests that, if more samples were provided, swine and wild boars would not have a different functional core metagenome. A deeper analysis should be done to differentiate alpha and beta diversity within and between the two organisms gut metagenomes.

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