

Approche hybride de modélisation explicable du métabolisme des écosystèmes microbiens

Hybrid approach for explainable metabolic modelling of microbial ecosystems'

Présenté par Maxime LECOMTE

November 14, 2023

Membres du jury

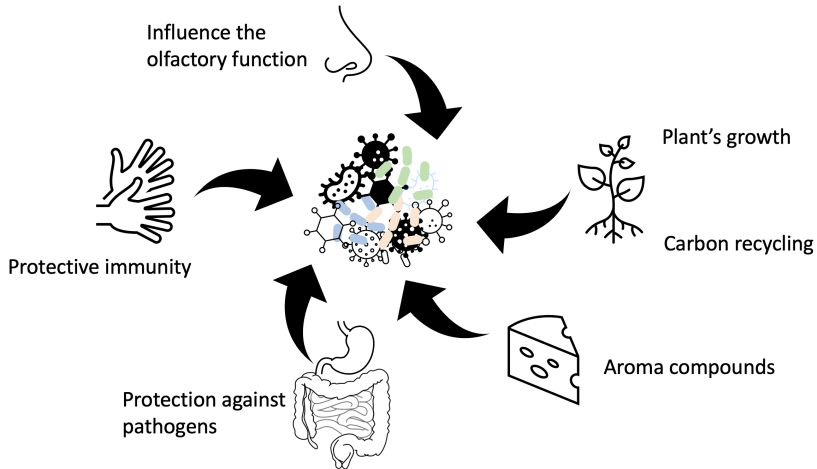
Président: SIMON Laurent

Rapportrices: BAROUKH Caroline
COCAIGN-BOUSQUET Muriel

Examineurs: COTTRET Ludovic
LAROCHE Béatrice
MARKOV Gabriel

Co-direction: David SHERMAN et Hélène FALENTIN
Encadrement: Clémence FRIIOUX

Why the study of microorganisms is relevant ?



- High diversity of microorganisms
- Microorganisms roles specific to the environment (Royet and Plailly, 2004; Belkaid and Hand, 2014; Zhang et al., 2015; Hoorman, 2011; McSweeney and Sousa, 2000)

What underlying mechanisms are responsible of the observed activity ?

Metabolism

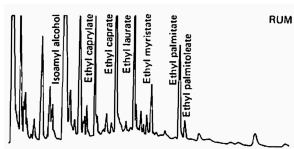


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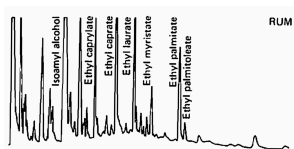


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What is metabolism ?

Set of all biochemical reactions occurring in the cell of an organism that permit the production of energy and metabolic goods. (Sánchez López de Nava A, 2023)

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Metabolism and Bacterial interactions

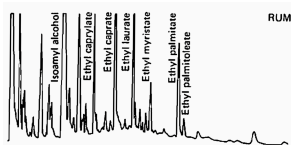


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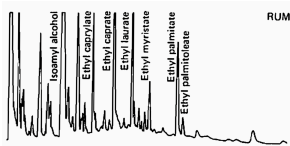


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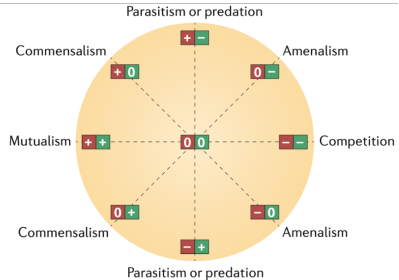


Figure 2: List of different types of bacterial interactions (Faust and Raes, 2012)

- Bacterial interaction can affect positively / negatively other organisms

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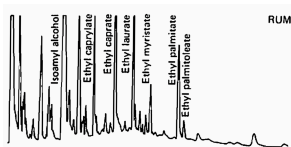


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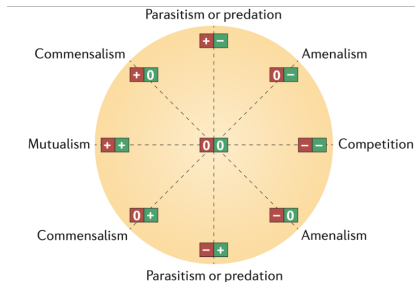


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Bacterial interactions can modulate metabolic goods

How can we study this impact through metabolism?

Genome-scale metabolic network (GEMs) reconstruction

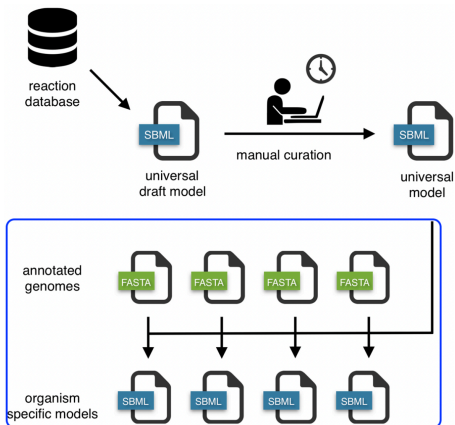


Figure 3: Top down genome-scale metabolic network reconstruction approach (modified from Machado et al., 2018)

- For bacteria: average of 1500 reactions, 1000 genes, 800 metabolites
- Informatic can help to resolve combinatorial problem

How can we study this impact through metabolism?

Systems biology

System biology

Associate an organism to a system and study the all system (Kitano, 2002)

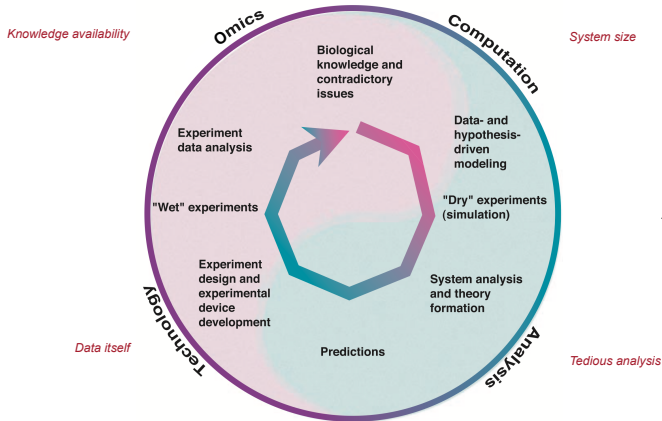
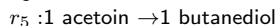
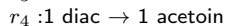
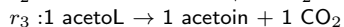
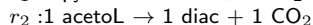
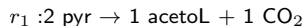


Figure 4: System biology modified from Kitano, 2002

- System biology combines biology and informatic analysis for studying bacterial behavior

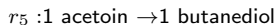
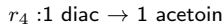
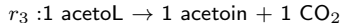
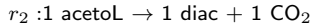
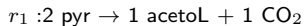
How the computational phase is concretely achieve ?

Metabolic network representation



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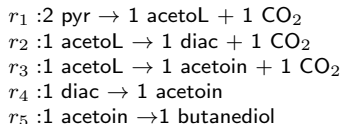


Stoichiometry matrix

	r_1	r_2	r_3	r_4	r_5
pyr	-2	0	0	0	0
acetoL	1	-1	-1	0	0
diac	0	1	0	-1	0
CO ₂	1	1	1	0	0
acetoin	0	0	1	1	-1
butanediol	0	0	0	0	1

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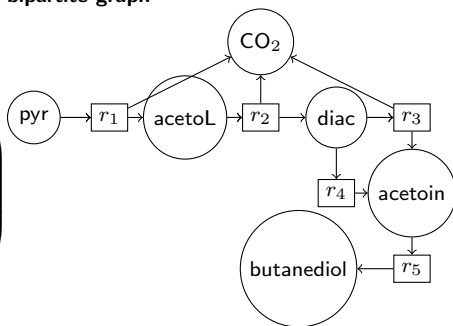
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bipartite graph



Stoichiometry matrix is commonly used for quantitative analysis instead of **graph**, more focused on topology analysis

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Build a metabolic model

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Constraint-based approaches

$$\frac{dx}{dt} = S.v = 0$$

Objectifs de la thèse

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