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Spotlight on the viral diversity of fermented vegetables using viral metagenomics

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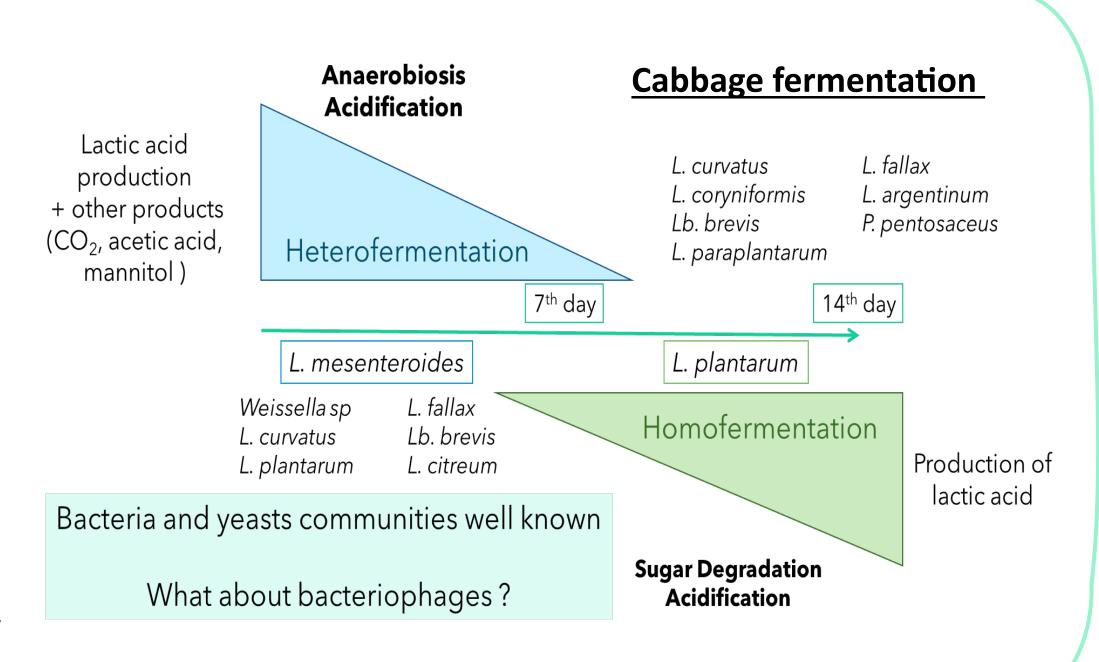
Introduction

Fermented vegetables are usually produced using the natural microorganisms present in the raw material. The process is characterized by a first heterofermentative step, generally carried out by Leuconostoc mesenteroides, which acidifies the medium and creates anaerobic conditions. The second homofermentative step is mainly carried out by Lactiplantibacillus plantarum. The composition of bacterial and fungal communities present in fermented vegetables (cabbage, carrot, turnip) have been widely characterized, whereas only few studies described the viral communities of these ecosystems.

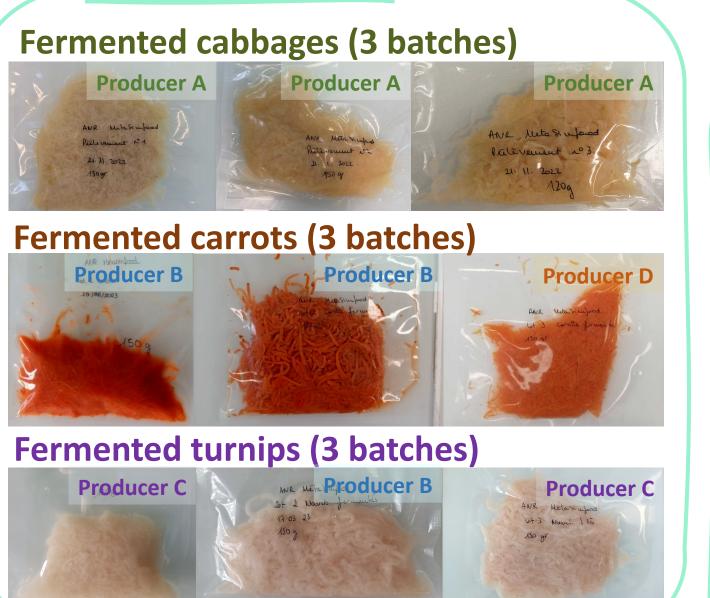
Several phages infecting different bacterial species were previously isolated from sauerkraut during the fermentation process, which revealed for the first time the complex ecology of phages in this ecosystem¹. Later, viral metagenomic analysis performed on different fermented foods including sauerkraut and kimchi highlighted again that complexity and showed that the vast majority of the sequences originated from uncharacterized phages².

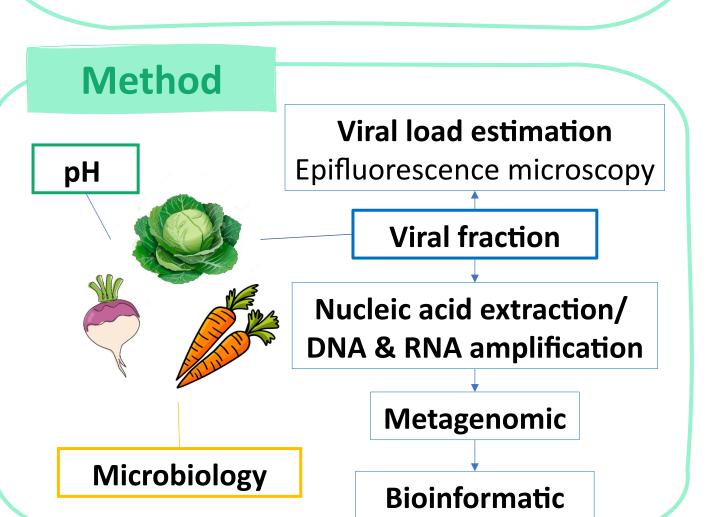
The objective of the present study was to describe the viral composition of commercial fermented vegetables (sauerkraut, fermented carrots and fermented turnips) as a first stone to build future research on the phage ecology in this peculiar ecosystem.

Viral particles extraction method



Material





Epifluorescence microscopy

Viral composition

Epifluorescence microscopy - viral load estimation	Viral load estimation (phages/mL)
Cabbage batch 1 (pH 3,39)	2.14E+08
Cabbage batch 2 (pH 3,36)	1.02E+08
Cabbage batch (pH 3,53)	4.47E+07
Carrot batch 1 (pH 3,34)	6.16E+07
Carrot batch 2 (pH 3,84)	1.43E+08
Carrot batch 3 (pH 3,47)	2.70E+07
Turnip batch 1 (pH 3,67)	1.27E+08
Turnip batch 2 (pH 3,55)	6.93E+07
Turnip batch 3 (pH 3,30)	NA
Control buffer	4.01E+04

Extraction and concentration of viral fraction from 10 g of fermented vegetables Homogenization in buffer **Matrix dilution** 1 L Tris 10 mM, NaCl 200 mM pH 7,5 2. Incubation Overnight at 4°C 3. Centrifugation: 7000 x g 15 min at 4°C **Eliminating microbial** cells 4. Filtration: 0.45 μm (PES) 5. Concentration: PEG precipitation 10% Overnight at 4°C **Concentration of the** 6. Centrifugation 5000 x g 30 min at 4°C viral fraction 7. Pellet dilution in 1 mL of SM buffer (Tris-HCl 50 mM, MgSO₄ 10 mM, NaCl 200 mM) Enumeration of viral particles by Metagenomic

Viral metagenomic data

epifluorescence microscopy

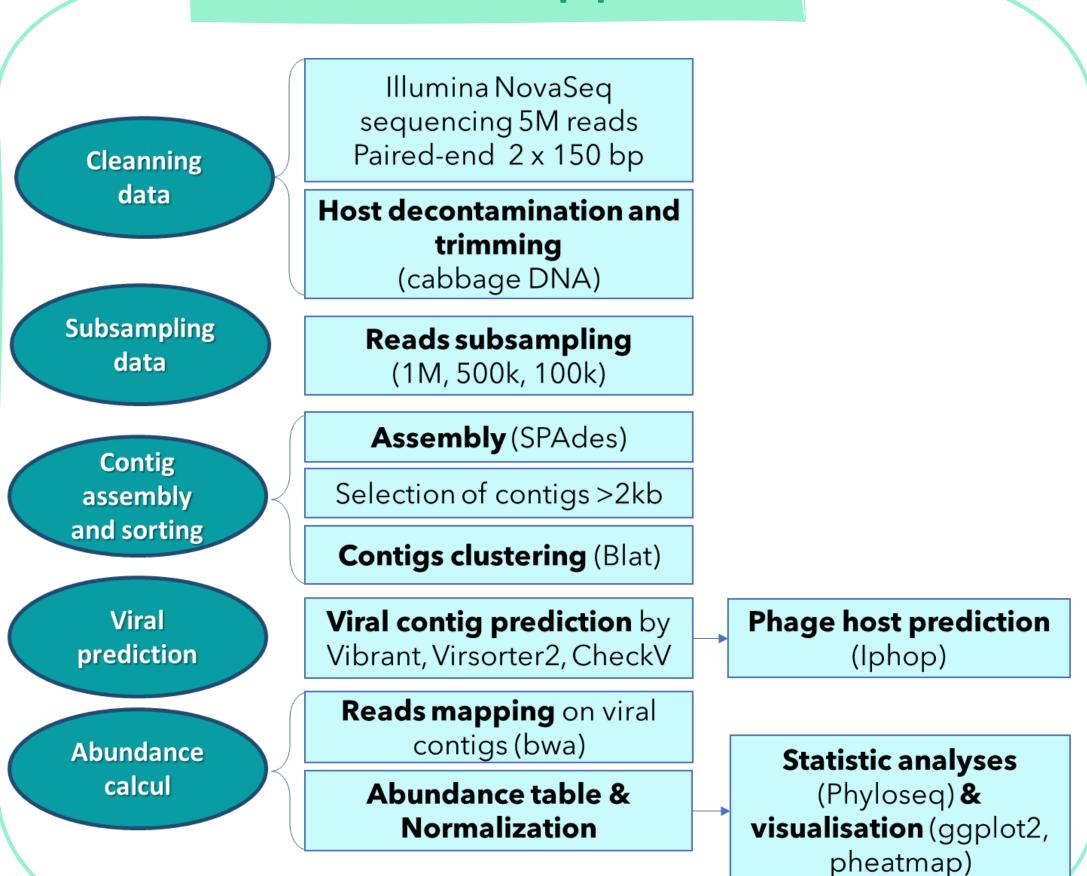
Samples	No. Contigs assembled >2kb	Longest contig
C1_1D	308	31 kb
C2_1D	35	27 kb
C3_1D	295	46 kb
Eau_CD	307	5 kb
Tampon_CD	219	8 kb
N1D	611	33 kb
N2D	194	18 kb
N3D	0	_
Ct1D	21	3 kb
Ct2D	169	49 kb
Ct3D	40	10 kb
WD (water control)	11	2 kb
BD (buffer control)	179	6 kb

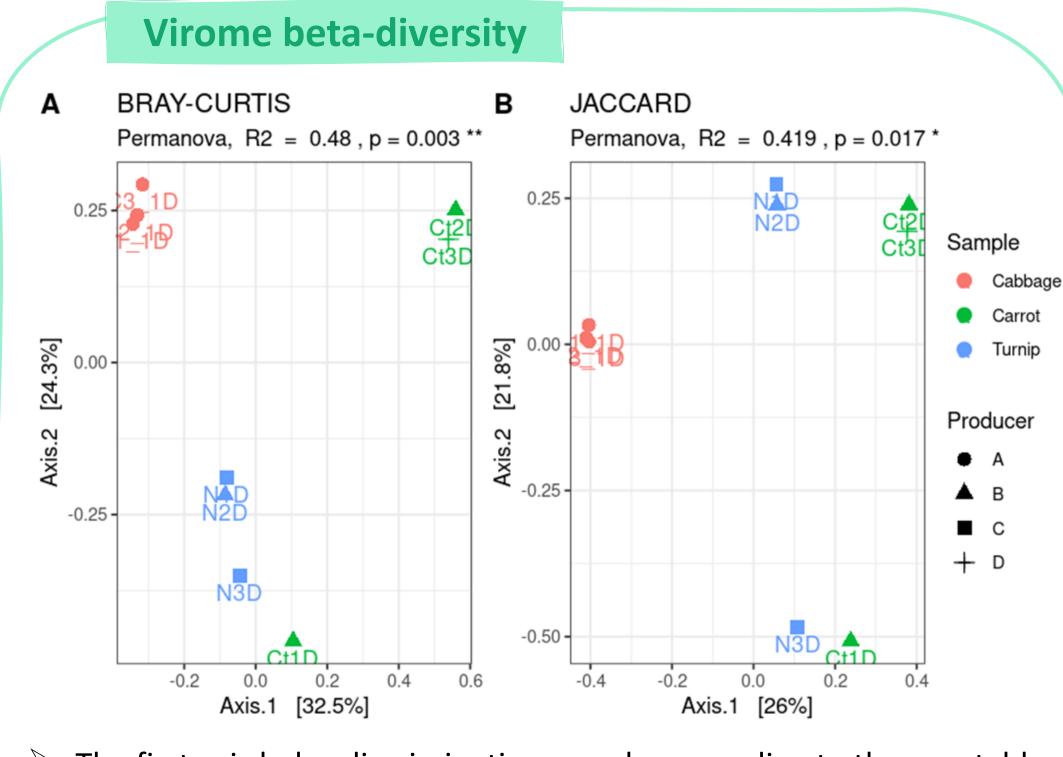
No. Contigs > 2kb	No. Clustered contigs	No. Predicted viral contigs
2832	1293	541

Final dataset

and Ct1D samples: low sequencing quality and assembly statistics poor other compared to samples.

Bioinformatic pipeline

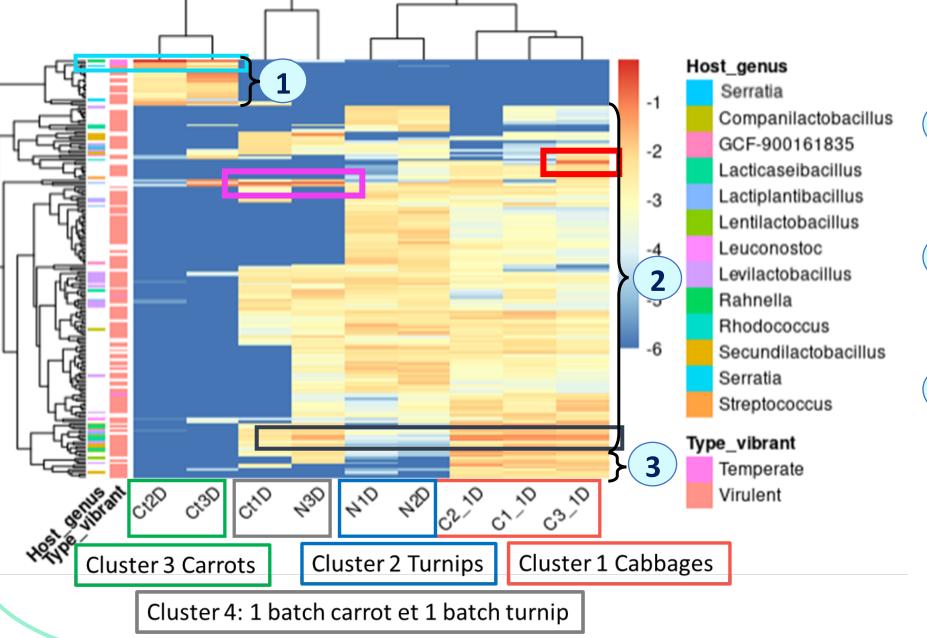




- The first axis helps discriminating samples according to the vegetable
- Samples N3D and Ct1D cluster together (low quality samples)

Viral composition depending on the plant matrix

Host_genus



3 contig groups

- 1 Contigs unique to two carrots samples (cluster 3)
- 2 Contigs common to most samples
- 3 Contigs mostly abundant in cabbage samples (cluster 1)

Most abundant viral contigs

Contigs identity	Infection cycle	Contig quality	Host order	Host family	Host genius	score_iphop
Ct1D_125k_NODE_1_length_2064	-	-	Lactobacillales	Lactobacillaceae	Lactiplantibacillus	94.0
Ct2R 125k NODE 1 length 41237	Temperate	High quality	Enterobacterales	Enterobacteriaceae	Rahnella	91.4
C1_1D_25k_NODE_2_length_23337	Virulent	low quality	Lactobacillales	Lactobacillaceae	Lentilactobacillus	91.4
Ct2D_250k_NODE_59_length_2554	-	-	-	-	-	-
C2_1D_250k_NODE_1_length_27078	Virulent	high quality	Lactobacillales	Lactobacillaceae	Lacticaseibacillus	99.4
C3_1D_25k_NODE_2_length_16738	Virulent	medium quality	Lactobacillales	Lactobacillaceae	Lacticaseibacillus	99.4
N2D_125k_NODE_42_length_2495	-	-	-	-	-	-
C3_1D_25k_NODE_3_length_5026	-	-	-	-	-	-
C2_1D_250k_NODE_2_length_13954	Virulent	low quality	Lactobacillales	Lactobacillaceae	Lentilactobacillus	91.1
C1_1D_25k_NODE_1_length_31726	Virulent	high quality	Lactobacillales	Lactobacillaceae	Lacticaseibacillus	96.6
C3_1D_125k_NODE_58_length_3209	- -	-	Lactobacilles	Lactobacillaceae	Lentilactobacillus	91.6

The most abundant contigs are different between samples

The hosts for these contigs, when predicted, correspond to bacteria usually found during the late stage of the fermentation process. Phages infecting Leuconostoc were also detected but at a lower abundance (not present in the top 10)

Conclusion and perspectives

This study provides an overview of the composition of viral communities in different commercial fermented vegetables, with a particular focus on extracellular particles (virion forms). We identify phages in high abundance that infect bacteria belonging to genera found in the late stage of the fermentation process (e.g. Lactiplantibacillus, Lentilactobacillus,...), and at a lower abundance phages infecting bacteria of the early stage (e.g. Leuconostoc) or spoilage bacteria (e.g. Enterobacter). Our results suggest that several phages infect most of the dominant lactic acid bacteria involved in the fermentation process and, therefore, may impact the bacterial successions occurring in this system. Since phages are likely to modify fermentation kinetics and consequently the quality of end-products, it is now necessary to better understand their individual ecological roles and ultimately set-up technological strategies to better control their activity and secure fermentation.

1 Lu et al., Bacteriophage Ecology in Commercial Sauerkraut Fermentations, 2003, DOI: 10.1128/AEM.69.6.3192-3202.2003 2 Park et al., Metagenomic Analysis of the Viral Communities in Fermented Foods, 2011, DOI:10.1128/AEM.01859-10

















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