# Understanding the role of mycoviruses in vine fungal communities

**A. Jaccard<sup>1,2</sup>, J. Brodard<sup>2</sup>, I. Kellenberger<sup>2</sup>, S. Schnee<sup>2</sup>, D. Sanglard<sup>1</sup>, K. Gindro<sup>2</sup> and O.Schumpp<sup>2</sup>** <sup>1</sup> Institute of Microbiology, UNIL, Lausanne, Switzerland | <sup>2</sup> Department of Plant protection, Agroscope, Nyon, Switzerland

## 1- Contexte

The SARS-CoV-2 pandemic shows that viruses are infectious and are expected to be transmitted among a population. However, in the fungal endophytic population, viruses called mycoviruses are not easily transmitted to one another, even among fungi of the same species.

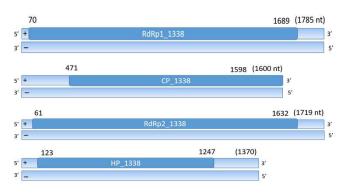
In this project, we seek to **understand why a virus can be present in one isolate but not in other isolates from the same species and same environment**. We will evaluate if a longer adaptation period has an influence on the prevalence of mycovirus among isolates of a fungal species.

### 3- Results

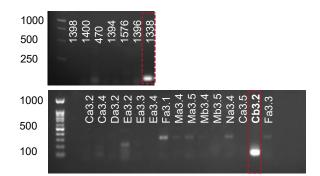
Swiss National Stud Farm SNSF

Agroscope

- Two mycovirus identified in one fungal isolates (Fig. 2).
- 1/21 isolate contained a viral segment identical to the previously identified segment (Fig. 3).
- Among the 583 fungi isolated, 268 were different isolates, of 47 different species (Fig. 4).



**Figure 2:** Four mycoviral genomic segments of 2 partitivirus identified in one isolate of *C. Cladospiroides* 



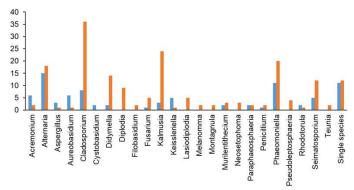
**Figure 3:** Screen for RdRp1\_1338 among endophytic *Cladosporium sp.* isolated from vines of the same parcel same cultivar (1398,1394,1396,1400), and different parcels same cultivar (bottom), different cultivar (1576) and other plant species (470, 1576)

# 2- Material and method

- Fungal endophytes were obtained from vine wood samples deposited on growth medium. The species of fungal isolates were identified by ITS amplification and sanger sequencing.
- **Illumina sequencing** was performed for the initial identification of mycoviruses from a cultured fungal isolate of *Cladosporium Cladospiroides*. The full sequence was reconstructed with **RACE PCR**.
- The identified mycoviruses were screened in isolates of the same fungal species with **RT-PCR**.
- The fungal community was obtained from fungi living in nongrafted vine for a long time compared to fungi living more recently in grafted vines. Plants were growing in the **same pedoclimatic area** and were from the **same cultivar**.



Figure 1: isolation of fungal endophytes from vine wood samples



**Figure 4:** Number of fungal isolates from old (blue) and young (orange) vines per OTU.

### 4- Conclusion and ongoing work

- The presence of mycovirus in the grapevine was confirmed from the identification of four mycoviral segments in one isolates. It shows that mycoviruses can be numerous in a single fungal isolate. However, the screen of the 21 fungal isolates stresses that although the presence of a similar mycoviral segment in isolates from different origines has been found, it is a rare situation.
- The constructed fungal community from old and young plants presents similar species, permitting to identify the prevalence of mycovirus in a same host species among the two variables.
- **dsRNA** will be extracted from all isolates and sequenced with **Illumina sequencing** technology, for a mycovirome identification.





Schweizerische Eidgenossenschaft Confédération suisse Confederazione Svizzera Confederaziun svizra Federal Department of Economic Affairs, Education and Research EAER Agroscope

Swiss Confederation