Honeybee viruses in novel hosts

Studying agrochemical-pathogen stress combination in wild bees.

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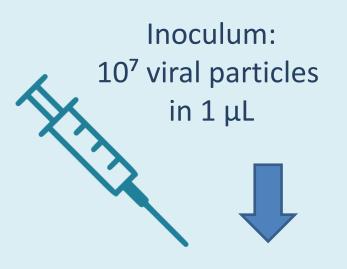
Introduction

Bees face a **combination of stressors** today such as habitat loss, agrochemicals and emerging pathogens¹. Certain agrochemicals, such as neonicotinoids, affect immunity pathways and promote virus replication in honeybees and can potentially synergistically increasing mortality². Honeybee-associated viruses has increased in prevalence across the globe³, and there is evidence of viral spillover from honeybees to wild bees⁴. The effects these viruses can have on wild bee populations, alone and together with agrochemical exposure, is still largely unknown.

Objectives

- To test the biological activity of the three most common honeybee-associated viruses in Europe³ in **new solitary bee hosts** *Osmia bicornis* (Megachilidae) and *Anthophora plumipes* (Apidae), by measuring viral replication over a set timeframe.
- Provide a virus model in wich to test pesticide-pathogen interactive effects in novel wild bee species.

DWV-A Deformed Wing Virus A



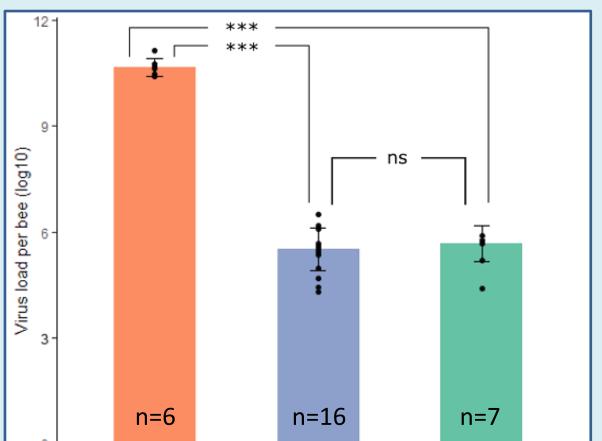
Housing for 3, 6 or 10 days



Apis mellifera

Anthophora plumipes

Results DWV-A





Inoculum:

10⁷ viral particles

in 1 µL

Housing for

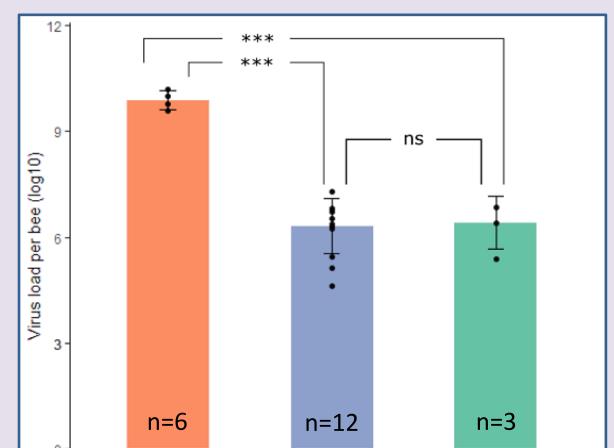
3, 6 or 10 days



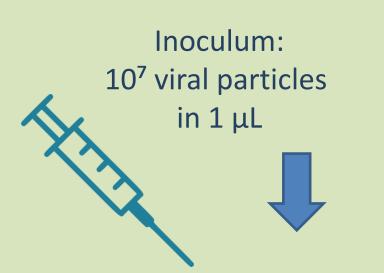




Results DWV-B



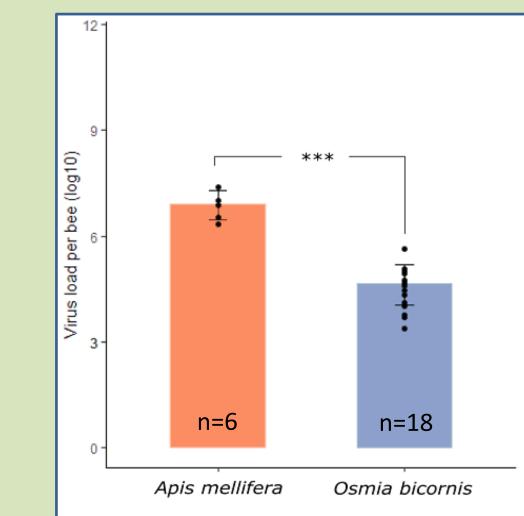
BQCV Black Queen Cell Virus





Housing for 3, 6 or 10 days

Results BQCV





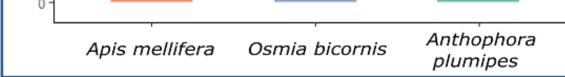


Figure 1. DWV-A virus titre per bee (log10) with median and s.e. Sample day and gender pooled per sp.

Decrease in titre from start inoculum in *O. bicornis* and *A. plumipes*. Virus replication in *Apis mellifera* after 3 days.

No difference in titre between day 6 and 10 for *O. bicornis* and *A. plumipes* and no difference in titre between genders.

Methods

Virus inoculum purified from local Apis colonies, diluted to 10^7 viral particles/µL with potassium sulphate buffer. Control treatment buffer only. 1 µL injected in abdomen using Hamilton syringe. Housing at 22°C (25°C for *A. mellifera*) with ad libitum 50% sugar feed. Entire bees were crushed and viral titren per individual assessed via quantitative real-time PCR (QuantStudio 3). Control bees screened for all three viruses and found to be clean. Virus titres compared using paired Wilcoxon rank-sum test.



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	Apis mellifera	Osmia bicornis	Anthophora plumipes

Figure 1. DWV-B virus titre per bee (log10) with median and s.e. Sample day and gender pooled per sp.

Decrease in titre from start inoculum in *O. bicornis* and *A. plumipes*. Virus replication in *Apis mellifera* after 3 days.

No difference in titre between day 6 and 10 for *O. bicornis* and *A. plumipes* and no difference in titre between genders. **Figure 1**. BQCV virus titre per bee (log10) with median and s.e. Sample day and gender pooled per sp.

Decrease from start inoculum in *O. bicornis*. Weak virus replication in *Apis mellifera* after 3 days – potentially lower virus volume in start inoculum.

No difference in titre between day 6 and 10 for *O. bicornis* and no difference in titre between genders.

Discussion & Way Forward

- No indication of viral replication in O. bicornis or in A. plumipes, in contrast to Bombus terrestris experiments, where injection caused high viral replication levels. Natural exposure route likely oral, where infectivity is likely to be even lower viruses not infective?
- ➢ DWV-B has been confirmed to be replicating in Osmia cornuta through (-)strand detection⁵, but there is no evidence for replication through increase in titre in this trial.
- High mortality in all test groups injection or housing effect?
- > Step 2 measure virus effect on novel host's lifespan, and combine with agrochemical exposure.



Pan-european assessment, monitoring and mitigation of Stressors on the Health of Bees - PoshBee aims to support healthy bee populations, sustainable beekeeping and pollination across Europe.



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