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Crithidia bombi presence and prevalence in solitary bee species Liz Sykes, Cameron Lamphere, Lynn Adler University of Massachusetts Amherst



INTRODUCTION

- Wild bees increase the economic yield of crops through pollination¹. Solitary bees 0 are not well studied compared to honey and bumble bees but comprise >85% of the wild bee population².
- Wild bees are facing population declines due to habitat degradation and loss, climate 0 change, and parasites and pathogens3. One such pathogen is Crithidia bombi, a trypanosomatid gut parasite that decreases bumble bee survival and foraging4.
- Sunflower (Helianthus annuus) pollen significantly reduces Crithidia infection in 0 the common eastern bumble bee⁵, but effects in solitary bees have not been assessed.
- Halictus ligatus and Melissodes agilis/trinodis are solitary bees that are commonly 0 found on sunflowers6.
- In previous studies, trypanosomatids have been found in Halictus and Melissodes 0 collected from wildflowers7.

What effect does sunflower pollen have on Crithidia infection in Halictus ligatus and Melissodes agilis/trinodis? Does Crithidia replicate in Melissodes agilis/trinodis?

METHODS

Crithidia survey

- Halictus ligatus and Melissodes agilis/trinodis were collected from 10 farms, 9 of which were growing sunflower, in western Massachusetts from mid-July until the end of August (Figure 3).
- Bees were incubated for <24 hours and provided with 30% sucrose.
- Bees were dissected, and the gut solution was assessed for Crithidia.
- o Pollen samples were collected and analyzed for sunflower content.

Infection Experiment

- I screened the feces of 30 Melissodes agilis/trinodis collected from RED (Figure 3) to determine infection levels.
- Bees were then starved for 3 hours before being hand-inoculated with Crithidia bombi and incubated with 30% sucrose.
- Bees were dissected in three intervals: <24 hours, 4 days, and 8 days post-collection.
- The gut solution was assessed for Crithidia.



Figure 3. Melissodes agilis/trinodis and Halictus ligatus were collected from 10 farms across western Massachusetts.



Figure 1. Melissodes agilis/trinodis on sunflower.



RESULTS

Crithidia survey

- 298 total bees were sampled (Figure 4). Of the 170 Halictus, none showed evidence of Crithidia infection. Of the 128 Melissodes, 3 (2.34%) showed evidence of Crithidia infection.
- The infected Melissodes were collected from DVF, NEX, and ROOT, which had average
- 0 Halictus were 95% sunflower pollen, and pollen samples taken from Melissodes were 98% sunflower pollen.

Infection Experiment

- 23 of the 30 bees died before their dissection date.
- o Of the 7 live Melissodes dissected, 1 (14.29%) showed evidence of Crithidia infection. The infected bee was dissected 8 days postcollection, indicating a replicating Crithidia infection.



- fg 30
- inflorescence values ranging from 498 to 641 (Figure 5).
- Overall, pollen samples taken from

Figure 4. The number of *Melissodes* agilis/trinodis and Halictus ligatus collected from each farm.



Figure 5. The average number of sunflower inflorescences at each farm. Not shown: LAR extends beyond the maximum y-value, with 85738 inflorescences.

DISCUSSION

Crithidia survey

- o In a prior field study, Halictus ligatus had a trypanosomatid prevalence of 38%, Melissodes agilis had a prevalence of 60%, and Melissodes trinodis had a prevalence of 47%7.
- In the current study, the much lower prevalence of Crithidia (none in Halictus ligatus and of 2.34% of Melissodes agilis/trinodis) suggests that infection levels are reduced in Halictus and Melissodes collected from sunflowers.
- The pollen samples showed evidence of Halictus and Melissodes feeding primarily on 0 sunflower pollen. However, there was no correlation between the number of sunflower inflorescences and infection levels at a farm.
- Future research could explore the difference in infection intensity between infected Halictus ligatus and Melissodes agilis/trinodis fed sunflower compared to nonsunflower pollen.

Infection Experiment

- o A Melissodes kept for 8 days had Crithidia infection, indicating that Crithidia may replicate in Melissodes as the pathogen did not pass immediately through the gut.
- Future research could test replication of Crithidia in a larger sample size of Melissodes.

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