מי מאביק את הרקפת המצויה?

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תקציר

ההאבקה של הרקפת המצויה תלויה לחלוטין בחרקים מאחר והיא אינה מואבקת על ידי הרוח והאבקה עצמית נמנעת אצלה על ידי מנגנון אי התאם עצמי. מבנה הפרח של הרקפת מתאים להאבקת זמזום הנעשית על ידי דבורים גדולות ונצפתה באופן נדיר במחקר. המאביקים המשניים הם מספר מיני תריפסים (Thysanoptera) וזבובי רחף (Syrphidae) שאינם ספציפיים ואילו המאביק העיקרי הוא העש הפרימיטיבי Lepidoptera, Micropterigidae (מ״מ) חוסה בתוך פרח הרקפת בעת גשם ולילות קרים, ניזון מגרגרי האבקה של הרקפת המצויה בעזרת גפי פה לועסות, מזדווג על הפרחים ומטיל עליהם ביצים.

Pollination by Bees and flies

Although bees and flies were rarely observed as pollinators of *C. persicum* (Table 1) their contribution was significant in comparison with the control since the flower was open and receptive over twenty days. They contributed 24.6% and 19.7% of the fruit set in the hysteranthous and synanthous populations respectively (Table 2).

Pollination by Moths



Introduction

The pendulous flowers of *Cyclamen* suggests buzz-pollination by large bees (Picture 1), as was recently found in *C. repandum*, and *C. hederifolium*. Accordingly, we expected to find the same syndrome in *C. persicum*. Preliminary observations showed that *C. persicum* was visited by various agents such as: small moths, thrips, and very rarely by bees and syrphids. This study evaluated the relative contribution of the flower's visitors to the pollination success of the *C. persicum*.



Picture 1.

Flowers of *C. persicum* — The corolla "mouth" surrounded by a deep pink colored ring that emphasizes the entrance to the reproductive organs.

Picture 4.

The primitive moth *Micropterix berytella* (Micropterigidae) in a feeding position on the bottom of the *C.persicum* flower.

Observations on the dynamics of moths and flowers of

C. persicum through the season showed a similar pattern (Fig.2). Using video taping the moth was observed pollinating the *Cyclamen* flowers by touching the stigma with its mouth-parts. The moth has chewing mandibles (not a proboscis which is common in the Lepidoptera) which enables eating pollen (Picture 4) that was also identified in the moth's stomach contents. The moth's pollination activity contributes 12.6% to the fruit set, which is significant in comparison to the control (Table 2).

Order	Species	Fall*	Winter*
Hymanoptera	Amegilla sp.	4 (4)	0
	Apis mellifera	3 (3)	3 (8)
	"large" bee	0	1 (1)
Diptera	Myiatropa florea	1 (1)	0
	Metasyrphus corollae	1 (1)	2 (2)
	Melinda biseta	1 (1)	0
	Pelecocera latifrons	0	1 (1)
	SUM	10 (10)	7(12)

Table 1. Flies and bees visits in the two populations of *C. persicum* during 1996/7. In parenthesis are the number of flowers that were visited by each pollinator. *Fall refers to hysteranthous population and *winter refers to synanthous population.



Picture 6.

Syrphid fly

Metasyrphus

corrollae, one of

the pollinators of

C. persicum.

Picture 5. Thrips on the corolla "mouth" of *C. persicum*. Picture 7. Honey bee Apis mellifera on flower of *C. persicum*.

Study site

The research was conducted at Ramat-Hanadiv, southern Mt. Carmel in Israel. Synanthous plants (flowering and leafing are simultaneous) of *C. persicum* are very common and flowers in the winter, while hysteranthous types (flowering and leafing are separated), flower in the autumn and are limited to north facing slopes. The flowering seasons of the different populations are almost entirely separated with just a short overlapping (Fig 1).









Dynamics of *C. persicum* flowers and *M. berytella* throughout the winter of 1997.

Pollination by Thrips

The presence of thrips in *C. persicum* continued throughout the flowering season in both populations (Fig.3). Mature thrips were observed walking and mating on the corolla petals, and different stages of young and mature larvae were found inside the stamens. Thrips' pollinating activity contributed to successful fruit-sets at a rate of 8.2% and 9.4% in the hysteranthous and synanthous populations respectively. These results are significant as compared with the control (Table 2).



Relative contribution of the different pollinators

The relative contribution of different pollinators was studied in isolated plants under insect-proof Perspex cages. In each cage we placed three *Cyclamen* plants. At the flowering



Picture 8. Perspex cages.

commencement 20 thrips were introduced once a week into five cages, 10 moths were introduced once a week into five cages and five cages served as a control. Fifteen plants, in the wild, were exposed to natural pollination while 15 other plants were sprayed with an insecticide against thrips and moths to assess the contribution of large pollinators. At the end of each season, we calculated the fruit set which was used as a measure for pollination success.

Treatment no	Treatment	Population	Fruit set	Treat. comparison	Significance
1	Field pollination	Winter	40.1	2 & 4	ns
2	Field pollination	Fall	26.9	7 & 9	s
3	Field pollination + spray	Winter	24.6	2 & 9	s
4	Field pollination + spray	Fall	19.7	5 & 8	s
5	Pollination in cages + moths	Winter	12.6	6 & 8	s
6	Pollination in cages + thrips	Winter	9.4	1 & 3	s
7	Pollination in cages + thrips	Fall	0.82	1 & 8	s
8	Pollination in cages + spray	Winter	0.4	1 & 2	s
9	Pollination in cages + spray	Fall	0.3	6 & 7	ns
				3 & 4	ns

Fig. 1. Flowering course of *C. persicum* in 1996-7. Fall Cyclamen* refers to hysteranthous plants and Winter Cyclamen* refers to synanthous ones.

Fig.3. Thrips presence in *C. persicum* flowers in the two populations of 1996/7. Fall Cyclamen* refers to hysteranthous plants and Winter Cyclamen* refers to of C of final synanthous ones.

Table 2. Relative contribution of the different pollinator groups in two populations of *C. persicum* during 1996/7. The data was subjected to Replications goodness of fit tests that compares the different treatments. Significant differences, P<0.05, is shown with an "s" and nonsignificant results are marked with an "ns".

Conclusion

The paradigm of "pollination syndrome" suggests that a set of flower traits were selected through an evolutionary process to fit typical traits of the pollinator. The typical anther structure with channeled stamens, is suitable for buzz-pollination by bees. However, we seldom observed buzz-pollination by bees and only in the hysteranthous population. It might be that the *C. persicum* coevolved originally with a large bee which performed buzz-pollination and is now extinct. The vacant flower's niche is now occupied by the monolectic moth *Micropterix berytella*. *The C. persicum supplies food, shelter, and* a site for mating and laying eggs in a way that suits the *M. berytella* size, senses, mouth structure organs, and its life cycle.

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