

Taxonomy and systematics

First reports of *Zoophthora radicans* (Entomophthoraceae) and *Metarhizium rileyi* (Clavicipitaceae) as pathogens of *Plutella xylostella* (Lepidoptera: Plutellidae) in Argentina

Primeros registros de Zoophthora radicans (Entomophthoraceae) y Metarhizium rileyi (Clavicipitaceae) como patógenos de Plutella xylostella (Lepidoptera: Plutellidae) en Argentina

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Abstract

Plutella xylostella (L.), diamondback moth (DBM), is considered among the 20 most resistant arthropods in the world. Entomopathogenic fungi are important mortality factors of Lepidoptera in the field. However, in Argentina entomopathogenic fungi as mortality factors of *P. xylostella* have not been identified. The aim of this study was to identify and isolate fungal pathogens of *Plutella xylostella* in Santa Fe, Argentina. The prevalence and seasonality of fungal infections were also investigated. *Zoophthora radicans* (Bref.) A. Batko (Entomophthoromycota: Entomophthorales) and *Metarhizium rileyi* (Farl.) Kepler, Rehner & Humber (Ascomycota: Hypocreales) as fungal pathogens of *Plutella xylostella* L. are reported for the first time in Argentina. The pathogens were identified from larvae, pupae and adults of *P. xylostella* on cauliflower and on cabbage. *Zoophthora radicans* was the predominant fungal pathogen, while *M. rileyi* was recorded from only one infected larva.

Keywords: Diamondback moth; Fungal pathogens; Prevalence; Seasonality

Resumen

Plutella xylostella (L.), la polilla de las coles, es considerada entre los 20 artrópodos más resistentes en el mundo. Los hongos entomopatógenos son importantes factores de mortalidad de lepidópteros en el campo. Sin embargo, en Argentina no han sido identificados hongos entomopatógenos como factores de mortalidad de *P. xylostella*. El objetivo de este estudio fue identificar y aislar hongos patógenos de *Plutella xylostella* en Santa Fe, Argentina. La prevalencia y la estacionalidad de las infecciones fúngicas fue también investigada. *Zoophthora radicans* (Bref.) A. Batko (Entomophthoromycota: Entomophthorales) y *Metarhizium rileyi* (Farl.) Kepler, Rehner & Humber (Ascomycota: Hypocreales) como hongos patógenos de *Plutella xylostella* L. son registrados por primera vez en Argentina. Los patógenos fueron aislados e identificados desde larvas, pupas y adultos de *P. xylostella* en repollo y coliflor. *Zoophthora radicans* fue el hongo patógeno predominante mientras que *M. rileyi* fue registrado infectando una única larva.

Palabras clave: Polilla de las coles; Hongos patógenos; Prevalencia; Estacionalidad

Introduction

Plutella xylostella (L.), diamondback moth (DBM), is the most destructive pest of Brassicaceae crops (Talekar & Shelton, 1993) and is considered among the 20 most resistant arthropods in the world (Mota-Sánchez et al., 2002). The extraordinary ability of this insect to develop resistance to various active ingredients of chemicals used against it means that other approaches are essential for the long-term management of this destructive pest.

Microbial pest control agents are welcome additions to Integrated Pest Management programs by their specificity against target pests and minimal environmental impacts. Entomopathogenic fungi are important mortality factors of lepidopteran in the field. Several species of fungal pathogens have been isolated from DBM (Cherry et al., 2004; Kirk et al., 2004; Vandenberg et al., 1998), but few have been studied in detail.

Zoophthora radicans (Bref.) A. Batko is an entomopathogenic fungus that has been reported to cause natural epizootics in several different insect species, including *P. xylostella* (Galaini-Wraight et al., 1991; Vandenberg & Soper, 1987). *Metarhizium rileyi* (Farl.) Kepler, S.A Rehner & Humber (Ascomycota: Hypocreales) is able to cause epizootics and dramatically reduce pest populations, as has been recorded in the U.S., Brazil and Australia (Corrêa & Smith, 1975).

In Argentina Bertolaccini et al. (2010) have mentioned entomopathogenic fungi as mortality factors of *P. xylostella* but the pathogenic species have not been identified. Thus, the aim of the present study was to isolate and identify the pathogenic fungi of *P. xylostella*, as well as to determine the prevalence of fungal infections.

Materials and methods

Surveys were conducted in conventional crop production in Monte Vera ((31°32'58.21" S, 60°41'34.74" W), Santa Fe province, Argentina. *Brassica oleracea* L. var. *botrytis* (cauliflower) and *B. oleracea* L. var. *capitata* (cabbage) were surveyed during the time of crop production. On the cauliflower crop, the survey was made from April 08 to July 20, 2011, whereas on the cabbage crop, the survey was made from March 29 to May 27, 2011. Surveys were conducted from planting to harvest.

Plutella xylostella sampling: *P. xylostella* populations were sampled weekly. Twenty plants of each crop were randomly selected and checked to quantify for larvae, pupae and adults of *P. xylostella*. Dead *P. xylostella* with evidence of external fungal growth (showing sporulation) were collected. Dead *P. xylostella* without external signs

of mycosis were also collected and placed in Petri dishes (60 mm in diameter) with a filter paper moistened with a few drops of distilled water (humid chambers) and maintained at 20 °C for 24-72 h to allow the development of overt mycoses.

To obtain pure cultures, *P. xylostella* infected with *Z. radicans* were placed on a moistened piece of sterile filter paper attached with double coated tape to the lid of a sterile 60 mm Petri dish, and then the lid was inverted over the bottom of a sterile Petri dish containing SEMA (80% Sabouraud dextrose agar and 20% of a mixture of egg yolk and skim milk) (Papierok & Hajek, 2012) plus 40,000 units/mL penicillin G (Merckâ, Germany) and 80,000 units/ml streptomycin (Parafarmâ, Argentina). This assembly was left for 12 h in darkness at 22 ± 1 °C. The lid with the attached *P. xylostella* was replaced with a sterile lid after 12 h. *Metarhizium rileyi* was isolated on Sabouraud maltose agar + 1% yeast extract (SMAY). All isolates were incubated at 22 ± 1 °C with a photoperiod of 16:8 (L: D). Isolates were deposited in the Mycological Collections at the Centro de Estudios Parasitológicos y de Vectores (CEP, La Plata, Argentina). Herbarium materials such as dried infected specimens and microscope slides were also deposited in the Mycological Collections at the CEP (La Plata, Argentina).

Infected *P. xylostella* were examined under a Zeiss DV4 dissecting stereomicroscope. Mycelia of *M. rileyi* were mounted in lactophenol/cotton blue (0.01% w/v), while fungal structures of *Z. radicans* were mounted in aceto-orcein (LPAO) (1:1) or stained with 1% aceto-orcein plus glycerine for semipermanent mounts. Fungal preparations were observed by phase contrast with an Olympus CH3 microscope and were photographed using a Nikon Optiphot microscope equipped with differential interference contrast (DIC) fitted with a Canon PowerShot A80 camera. Measurements of length and width of fungal structures (conidia, conidiogenous cells and mycelia) from fresh infected cadavers were made to enable specific identification. *Metarhizium rileyi* was identified according to taxonomic keys and monographs in Samson (1974), Samson et al. (1988) and Zare and Gams (2001) and *Z. radicans* according to taxonomic keys and monographs of Bałazy (1993), Humber (2012) and Keller (1991).

The numbers of healthy and infected *P. xylostella* in each development stage (larvae, pupae and adults) were compared by parametric Anova and Tukey's (HSD) *post hoc* test with $p = 0.05$ after log transformation of data. The Mann-Whitney *U*-test was used to compare the density of healthy and infected *P. xylostella* between crops and between developmental stages. Analyses were performed using InfoStat statistical software (2004).

Results

Zoophthora radicans and *M. rileyi* were the pathogens identified as mortality factors of *P. xylostella*. *Zoophthora radicans* was the predominant fungal pathogen on both crops and was recorded infecting larvae and pupae. One adult of *P. xylostella* infected with *Z. radicans* was also recorded. Infected adult was found in a lettuce crop located adjacent to the cabbage crop. *Metarhizium rileyi* was recorded from only one infected larva.

Zoophthora radicans (Bref.) A. Batko, Bull. Acad. Polon. Sci., Cl. II. sér. sci. biol. 12: 323 (1964) (Fig. 1).

=Synonymy:

Empusa radicans Bref. 1870

Classification: Zygomycota, Entomophthoromycotina, Entomophthorales, Entomophthoraceae.

Herbarium access number: CEP 50 (specimen number)

Description. Microscopic characters: conidia uninucleate, primary conidia elongate, papilla generally conical demarcated with a slight protuberance from the body of the conidia: 20.15 ± 2.32 (15.67-24.02) in length $\times 6.19 \pm 0.83$ (4.94-7.44) μm in width. Secondary conidia

as capilliconidia: 20.04 ± 1.83 (18.89-24.4) $\times 7.09 \pm 0.54$ (6.39-7.93) $\times 46.95 \pm 9.70$ (35.81-60.31) μm . Thin rhizoids compounds with specialized adhesive disc. No resistance spores were observed. Macroscopic characters: colony diameters reached 20 mm in 10 d on SMYA at 24 °C in darkness, rather compact, white, with clear yellow on reverse. Isolated but not maintained in pure cultures.

Metarhizium rileyi (Farl.) Kepler, Rehner & Humber, in Kepler, Humber, Bischoff & Rehner, Mycologia 106 (4): 824 (2014) (Fig. 2).

=Synonymy:

Botrytis rileyi Farl. 1883

Classification: Ascomycota, Pezizomycotina, Sordariomycetes, Hypocreomycetidae, Hypocreales, Cordycipitaceae.

Culture collection access number: CEP 367

Description. Microscopic characters: conidiogenous cells usually subglobose to short cylindrical or ampulliform with a globose base, $4.34 \mu\text{m} \pm 0.69$ (3.4-5.3 μm) in length $\times 1.8 \mu\text{m} \pm 0.23$ (1.4-2.2 μm) in width with necks short, smooth-walled. Conidia strongly ellipsoidal, smooth-walled, hyaline $4.2 \mu\text{m} \pm 0.47$ (3.6-5.3 μm) in length \times

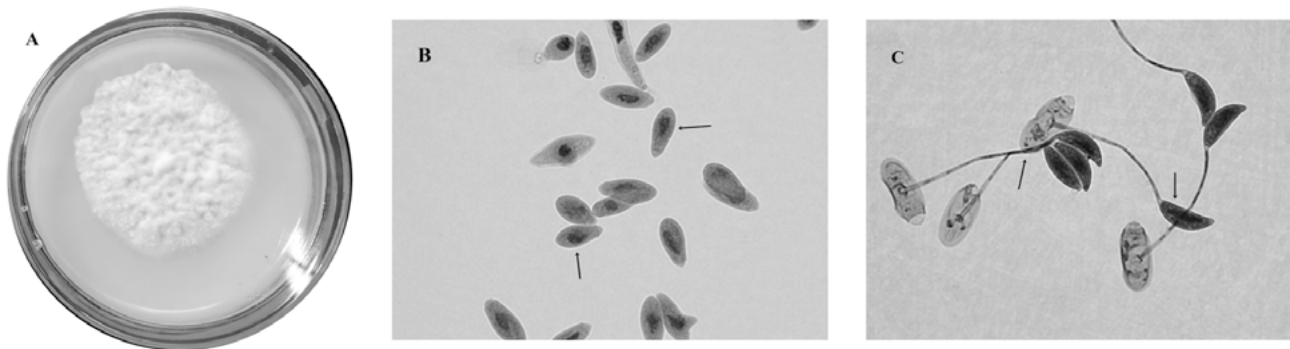


Figure 1. A. Isolate of *Zoophthora radicans* in pure culture; B, primary conidia of *Zoophthora radicans*; C, capilliconidia of *Z. radicans*. Scale bar: 10 μm



Figure 2. A. Isolate of *Metarhizium rileyi* in pure culture; B, conidiophore (cf) and phialides (ph) of *M. rileyi*; c, conidia of *M. rileyi*. Scale bar: 5 μm

2.1 $\mu\text{m} \pm 0.21$ (1.7-2.4 μm) in width (average). Hyphae septate, smooth-walled, hyaline 1.8 $\mu\text{m} \pm 0.33$ (1.0-2.2 μm) wide occasionally branched. Macroscopic characters: colonies growing up to 33.6 mm in diameter after 10 d at 25 °C. Circular colonies, with regular borders, reverse white to slightly yellowish. Spore colonies malachite green in according to the color chart Rayner (1970). Isolated and maintained in pure cultures.

Population variations: the averages of healthy and infected *P. xylostella* on each crop were calculated per sample date and are shown in Table 1.

On cabbage, the peak in population density of *P. xylostella* was observed on April 8, 2011 (\bar{X} = 1.70 insects/plant), and the first infected insects were observed on May 13, 2011. Significant differences were not recorded in the number of infected-*P. xylostella* among sampling dates ($p > 0.05$) (Table 1). The highest prevalence of fungal infection (40%, $n = 5$) was observed on May 19, 2011 and occurred with 18.1 °C temperature and 87% relative humidity (Fig. 3). After this date, there was a sharp decline in the number of healthy and infected insects.

On cauliflower, the peak in population density of *P. xylostella* was observed on May 13, 2011 (\bar{X} = 1.25 insects/plant) (Table 1). There were significant differences between study weeks on the number of infected-*P. xylostella* ($p < 0.05$) (Table 1). The first infected insects were observed on May 5, 2011, and the highest prevalence of fungal infection (90%, $n = 10$) was observed on May 27, 2011 (Table 1). Also, important prevalences were recorded until the end of the crop cycle. The highest percentage of infection occurred with 12.8 °C temperature and 82% relative humidity (Fig. 4).

The number of infected insects was significantly different among both crops ($p < 0.05$; Mann Whitney

$U = 4,612.0$). On cabbage, there were significant differences in the abundance of infected-*P. xylostella* between developmental stages (larvae and pupae) ($p < 0.05$; Mann Whitney $U = 756.00$). On cauliflower, the abundance of infected insects among larvae and pupae were not significantly different ($p > 0.05$; Mann Whitney $U = 1,515.00$).

Discussion

In Argentina there are reports of *Z. radicans* and *M. rileyi* as fungal pathogens of lepidopteran moths. López-Lastra and Scorsetti (2006) have recorded *Z. radicans* as a pathogen of *Epinotia aporema* (Walsingham) (Lepidoptera: Tortricidae) and *Tuta absoluta* Meyrick (Lepidoptera: Gelechiidae), while *M. rileyi* has been identified from *Anticarsia gemmatalis* (Hübner) on soybean crops (Toledo et al., 2004). *Zoophthora radicans* and *M. rileyi* as pathogens of *P. xylostella* are first records for Argentina.

In this study, we recorded infection levels of *Z. radicans* of 40 to 90% on *P. xylostella*. Aragón et al. (1997) cited prevalences of 40 to 80% of *Pandora gammae* (Entomophthoraceae) on *Rachiplusia nu* (Guen) on soybean crops.

Entomopathogenic fungi possess distinct advantages over other microbial control agents because they are able to attack all the developmental stages of insects, including pupal stages (Ferron, 1978). *Zoophthora radicans* infects insects of several orders and is a widespread pathogen of diamondback moths, usually attacking larvae but also sometimes pupae (Kanervo, 1946). In the present study, we found infections of *Z. radicans* on larvae and pupae of *P. xylostella*. The prevalence was higher in larvae than in

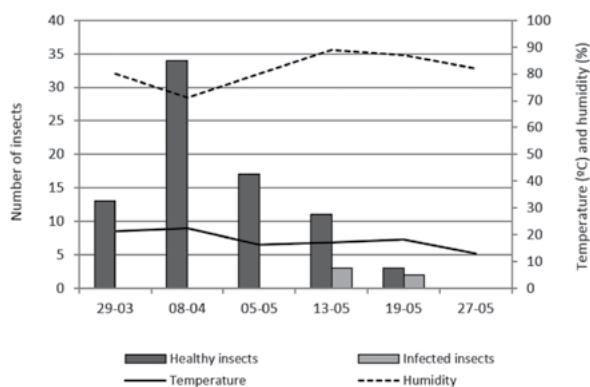


Figure 3. Seasonality and prevalence of pathogenic fungi on *Plutella xylostella* in *Brassica oleracea* L. var. *capitata* in 2011.

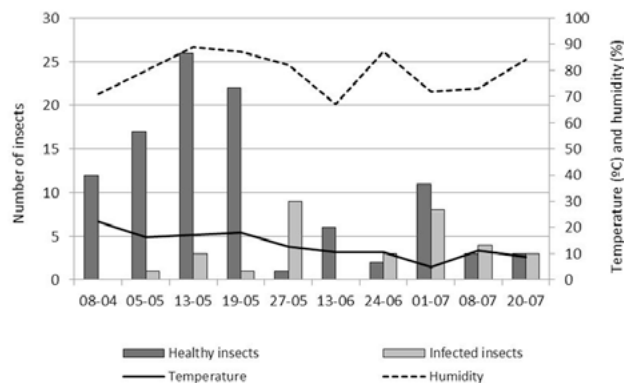


Figure 4. Seasonality and prevalence of pathogenic fungi on *Plutella xylostella* in *Brassica oleracea* L. var. *botrytis* in 2011.

Table 1

Mean number of healthy and infected *P. xylostella* and percentage of infection by sampling date on each crop^a.

Crops	N	Sample date	Healthy <i>P. xylostella</i>	Infected <i>P. xylostella</i>	Percentage of infection
<i>Brassica oleracea</i> var. <i>capitata</i> (cabbage)	13	29.03.2011	0.65 (0.74) a	0.00 (0.00) a	0
	34	08.04.2011	1.70 (1.86) b	0.00 (0.00) a	0
	17	05.05.2011	0.80 (1.64) a	0.00 (0.00) a	0
	14	13.05.2011	0.55 (1.05) a	0.10 (0.30) a	21.4
	5	19.05.2011	0.15 (0.36) a	0.10 (0.30) a	40
<i>Brassica oleracea</i> var. <i>botrytis</i> (cauliflower)	12	08.04.2011	0.65 (0.93) abd	0.00 (0.0) a	0
	18	05.05.2011	0.85 (1.18) ade	0.05 (0.22) a	5.6
	29	13.05.2011	1.25 (1.20) e	0.15 (0.67) ab	10.3
	23	19.05.2011	1.10 (1.41) de	0.05 (0.22) a	4.3
	10	27.05.2011	0.05 (0.22) c	0.45 (0.60) b	90
	6	13.06.2011	0.30 (0.57) abc	0.00 (0.0) a	0
	5	24.06.2011	0.10 (0.30) bc	0.15 (0.36) ab	60
	19	01.07.2011	0.60 (0.68) abcd	0.40 (0.68) b	42.1
	7	08.07.2011	0.10 (0.30) bc	0.20 (0.41) ab	57.1
	6	20.07.2011	0.15 (0.48) bc	0.15 (0.36) ab	50

^a Entries show the mean number (\pm SE) of healthy and infected *P. xylostella*. Means sharing the same lower-case letter are not statistically different ($p > 0.05$).

pupae probably because the thick and sclerotized cuticle of the pupa is a major barrier to fungal infection (De La Rosa et al., 2002; Hajek & St. Leger, 1994).

In the present study, the highest prevalences of fungal infection on both crops were recorded during May, with high humidity percentages (over 80%). In accordance with Steinkraus (2006), relative humidity is the single most critical factor in transmission of fungal pathogens. Fungal pathogens generally require high relative humidity for survival, germination and sporulation. In Argentina, previous studies on the phenology of entomophthoroid fungi in populations of insects have reported that fungal infections are more common in the autumn-winter season (in the southern hemisphere, from March to August) (López-Lastra et al., 2003; Toledo et al., 2008). In Uruguay, Alzugaray et al. (2010) recorded *P. neoaphidis* as one of the main mortality agents of aphids but emphasized that its action is restricted to autumn and winter. The low temperatures and humid conditions prevalent during the winter growing season tend to be favorable for the occurrence of entomophthoroid mycoses (Poprawski & Wraight, 1998; Wraight et al., 1993).

Our results showed that *Z. radicans* is an important factor in the mortality of *P. xylostella* under natural conditions. This study is the first record of *Z. radicans* and *M. rileyi* from *P. xylostella* in Argentina.

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