

# First Record of *Aspergillus flavus* as a Fungal Pathogen of the Predator *Rhynocoris marginatus* (Hemiptera: Reduviidae)

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*EntomoBrasilis* 5 (1): 80-81 (2012)

**Abstract.** An entomopathogenic fungus, *Aspergillus flavus* Raper and Fennell is recorded for the first time as a pathogen of the predator *Rhynocoris marginatus* (Fab.) (Hemiptera: Reduviidae) in natural agro-ecosystems of Tirunelveli District, Tamil Nadu, India.

**Keywords:** Natural enemy; Microbe, Pathogen; Reduviid predator.

## Primeiro Registro de *Aspergillus flavus* como Patógeno do Predador *Rhynocoris marginatus* (Hemiptera: Reduviidae)

**Resumo.** O fungo entomopatogênico *Aspergillus flavus* Raper and Fennell é registrado pela primeira vez como um patógeno do predador *Rhynocoris marginatus* (Fab.) (Hemiptera: Reduviidae) em agroecossistema natural no distrito de Tirunelveli, Tamil Nadu, Índia.

**Palavras-chaves:** Inimigo Natural; Micro-organismo; Predador; Reduviídeo predador.

The reduviid predator, *Rhynocoris marginatus* (Fab.) is an entomophagous insect distributed in many agro-ecosystems and feeding on more than twenty economically important insect pests in India (SAHAYARAJ 2007) (Figure 1). The potential of *R. marginatus* as a biocontrol agent under laboratory (SAHAYARAJ 2000; SAHAYARAJ & BALASUBRAMANIAN 2009) and field conditions (SAHAYARAJ & MARTIN 2003; SAHAYARAJ & RAVI 2007) has been reported earlier. SAHAYARAJ (2007) reported that the egg cluster of this bug was parasitized by a hymenopteran parasitoid, *Trissolcus* sp. So far no information is available about the microbial pathogen of this biological control agent. Here, we recorded a fungal pathogen which infects the life stages of *R. marginatus* both in laboratory and field condition for the first time.



Figure 1. *Rhynocoris marginatus* female feeding on *Phenacoccus solenopsis* Tinsley (Hemiptera: Pseudococcidae).

Six males and four females of *R. marginatus* was collected from agro-ecosystems and maintained under laboratory conditions at 30±1 °C, 70-80 % RH and with a photoperiod of 11: 13 h D: L for one week. Mycelial scarbed out from the dead predator was cultured in 15 mL Sabouraud dextrose agar medium at 30°C for 12 h. Microscopic examination of fungal mass revealed large number of spores born on conidiospores. On the basis of colonial morphology and microscopic characteristics (ABUBACKER & RAMANATHAN 2003 and CAPPUCINO & SHERMAN 2004) the isolated fungus was identified as *Aspergillus flavus* Raper and Fennell. The fungus isolated from such infected predator has satisfied the Koch's postulates. In another study, 10<sup>6</sup> conidia/mL suspensions was prepared from the 5-day old PDA slant culture, sprayed over the body of the healthy adult predators and recorded the mortality if any for a period of seven days. Thirty individuals (15 each of male and female) were maintained for this study.

During our field visit in Sivanthipatti (77° 47' E and 8° 30' N), Tirunelveli District, Tamil Nadu, India on fourth April 2005, live adult *R. marginatus* (six males and four females) were collected and brought to the laboratory. Next day we observed that 20 per cent of the collected predators were dead, and after five day of infection predators were mummified. Initially white mycelial growth was observed in the infected predators. Pale yellow mycelial growth appeared after 48 hours. After four days, the mycelial color turned to yellowish-green as originally in infected predator. Conidia of *A. flavus* were not virulent when applied to the surface of healthy *Galleria mellonella* (Linn.) caterpillars (RAYMOND *et al.* 2000). However, these fungi could, colonize, killed insects and produced spores on reduviid cadavers within 4 days of infection.

The pathogenicity test showed that an average of 96.6% of mortality was recorded within 4 days in the treated adult predators under

laboratory conditions. The mycelial growth appeared fourth day after inoculation. From the dead adult, the fungus was re-isolated and checked with healthy predators. Similar pathogenicity was recorded again. Numbers of previous reports are available on the presence of aflatoxins, secondary metabolites from fungus, *A. flavus* in foods and feeds, cause serious economic loss, apart from their carcinogenic potentials and insecticidal ability (DOYLE *et al.* 1982; HAGLER *et al.* 1983; MISLIVEC *et al.* 1988; KUMAR & PRASAD 1992 and ABUBACKER & RAMANATHAN 2003). SELVARAJ *et al.* (2002) reported that *A. flavus* infect various life stages of red cotton bug, *Dysdercus cingulatus* (Fab.) during laboratory rearing. But there are no records available about the infection of *A. flavus* on reduviid predators.

### ACKNOWLEDGEMENT

K. Sahayaraj is thankful to Council of Scientific and Industrial Research, Govt. of India for the financial assistance (ref. No. 37/1350/08/EMRII). The authors are highly thankful to the authorities of St. Xavier's College (Autonomous), Palayamkottai for providing necessary laboratory facilities and the encouragements.

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Recebido em: 23/07/2011

Aceito em: 28/10/2011

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### Como citar este artigo:

Sahayaraj, K, J.A.F. Borgio & S.M. Kumar, 2012. First Record of *Aspergillus flavus* as a Fungal Pathogen of the Predator *Rhynocoris marginatus* (Hemiptera: Reduviidae). EntomoBrasilis, 5(1): 80-81.

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