Occurrence of the Entomopathogenic Fungi Nomuraea rileyi (Farlow) Samson on Acontia graellsii F. (Noctuidae: Lepidoptera) and Beauveria bassiana (Balsamo) Vuill. on Myllocerus subfaciatus G. (Curculionidae: Coleoptera)

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The bhendi semilooper, Acontia graellsii F. though a minor pest, at times causes severe damage to the bhendi leaves, while Myllocerus subfaciatus G. the brinjal ash weevil, causes damage to the leaves. The damage caused by the larvae in soil to the roots often leads to wilting of the affected plants. During the course of our survey for entomopathogens on insect pests of horticultural crops at Indian Institute of Horticultural Research Farm, Hessaraghatta, Bangalore, we isolated two fungal pathogens from the infected insects of A. graellsii on bhendi and M. subfaciatus on brinjal.

The fungus infected caterpillars of *A*. graellsii became hard and mummified and found adhering to the leaves with their prolegs fixed and the anterior head and abdominal region slightly raised (Fig. 1). The fungus showed white mycelial growth ramifying the entire body of the caterpillar. The infected caterpillars when kept on moist filter paper in a pair of petri dish at 25-27°C the fungus sporulated profusely. The spores were pale green in colour.

The fungus from A. graellsii was isolated into pure culture on Sabouraud maltose agar enriched with 1% yeast, where it grew well and sporulated profusely. The spores were pale green in colour and ellipsoidal in shape measuring 3.0-4.5 μ m. The fungus from M. subfaciatus was isolated into pure culture on Sabouraud dextrose agar where it grew well and sporulated profusely and the spores were muscardine in colour and globose in shape

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measuring 1.7-2.5 μ m. Based on the detailed morphological characters, the fungal pathogens were identified as *Nomuraea rileyi* (Farlow) Samson on *A. graellsii* and *Beauveria bassiana* (Balsamo) Vuill. on *M. subfaciatus*. Our preliminary identification was later confirmed by Dr. Humber Boyce Thompson Institute, New York (Personal communication).



Fig. 1. Nomuraea rileyi-infected semilooper A. graellsil

The pathogenicity tests with the fungal pathogens have proved that they were highly virulent inflicting cent per cent mortality of their host insects in 5-8 days at 25-27°C and relative humidity 80-90%. Both the fungal pathogens were reisolated from such infected insects satisfying the Koch's postulates.

The occurrence of N. rileyi has been reported on Diacrisia obliqua (Singh and Gangrade, 1975) and on tobacco leaf eating caterpillar, Spodoptera litura (Rao and Padke, 1977) and B. bassiana on cabbage semilooper Trichoplusid ni (Urs et al., 1965), rice rests (Rao, 1975; Srivastava and Nair, 1978; Nayak and Srivastava, 1979; Israel and Padmanabhan, 1980), banana leaf beetle (Roy and Pujari, 1979) and sugarcane shoot borer Chilo infuscatellus Snell. (Easwaramoorthy and Santhalakshmi, 1988) from India. This communication is the first report of N. rileyi on A. graellsii and B. bassiana on M. subfaciatus from India.

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KEY WORDS : Acontia graellsii, Nomuraea rileyi, Myllocerus subfaciatus, Beauveria bassiana.

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Effect of Leaching on the movement of Nuclear polyhedrosis virus of *Heliothis armigera* in soil.

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Studies on some insect polyhedral inclusion viruses have shown that inclusion viruses may persist for long periods in the soil (Jaques, 1964; David and Gardiner, 1976). Polyhedral occlusion bodies (POB) appear to be adsorbed quite firmly onto soil particles (Hukuhara and Namura,1971; Hukuhara and Wada,1972; Narayanan *et al.*, 1987). Leaching is one of the several factors which influence the persistence of an insect virus in soil. The present study reports the effect of leaching on

*Scientist S₃, National Centre for IPM, Bellary Road, Bangalore 560 024 the persistence of nuclear polyhedrosis virus (NPV) of *Heliothis armigera* in a column of black soil in the laboratory, using polyhedra prelabelled with an isotope ^{32}p .

The isotope was obtained from Bhaba Atomic Research Centre (BARC), Trombay, Bombay. A suitable technique of incorporating the labelled ³²p into the larval semisynthetic diet (Narayanan, 1979) was developed as follows. "Carrier-free" ³²p as orthophosphoric acid having a specific activity of 15-20 μ ci, was applied @ 0.02 ml per diet disc of