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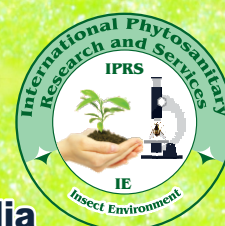
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***Cover Page: *Apoderus tranquebaricus* (Attelabidae) identified by Dr. Chitra N, Professor, Department of Entomology, Tamil Nadu Agricultural University.**

Photo by: Dr. Sevgan Subramanian, Principle Scientist, International Centre of Insect Physiology and Ecology, Kenya

Insect Environment

(Quarterly journal to popularize insect study and
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The first issue of the *Insect Environment* was published in 1996. The sole objective of the *Insect Environment* is to popularize insect study through popular, semi-technical and technical research notes, extension notes for managing insect pests, photographs, short blogs and essays on all aspects of insects. The journal is published quarterly, in March, June, September, and December.

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Editorial

Insect Environment (IE) has been experiencing phenomenal growth in the last two years. The drivers are our authors and the myriad well-wishers in India and around the globe. Our research articles have been touching millions of viewers or rather viewers are making hits into our articles; thanks to CABI (UK), DOI and Research Gate. Many authors message me that interest in their papers from far flung corners of the globe, indeed, made them feel elated, while we felt, our labours were not in vain!

We were the first to whistle-alert *Thrips parvispinous* on chillies in South India through our blogs followed with a couple of articles (IE 24 (4), 2021). Talking of our blogs they have been short ento-thrillers, and every Monday blogs are auto-sent to a few thousands, around the world.

Nobody ever thought of giving amateur insect pictures a chance to see the light of the day. 'Big' journals would accept photographs, relevant to a paper that too with a lot of editorial fuss and sometimes as I have myself experienced, pretence to quality! At IE we make a difference- if an insect picture tells a story by itself or its *oikos* or has aesthetics (all insects, however drab are beautiful to us) we publish it. I am dazed, that android/iphone mobiles nearly capture insects with clarity and presentability, almost matching professional cameras! No wonder, mobile pixels and soft digitization have unfortunately shrunk the genre of camerapersons! But IE is lucky to still have a repertory of professional entomologist-photographers of the likes of Dr. D. N. Nagaraj, Dr. Sevgan Subramanian and many more, who frame excellent pictures to the last setae and bristle!. IE would have been poorer



but for these “lens-persons,” whose watchful eyes, rarely miss a fly! I am also happy to appreciate the efforts of Dr Thirupam Reddy from Jharkhand. His picture on gynandromorphy of tassar silk moth in this issue is mention-worthy.

Once more I reiterate IE is all about capturing the natural history of insects and their environment in words, data and pictures. This phase hopefully- the post-pandemic- is seeing revival of physical meetings. IE would be happy to collaborate in publicity, if poster-pictures are given well on time. It is always ‘fun’ to attend conferences/seminars in India, as the ‘Indianness’ and pluralism injected into such meets, deliver not only scientific discussions, but also unforgettable camaraderie, local cuisines and art-shows.

This was certainly not evident in many international meets I attended round the world. There are excellent discussions (business-style), paid water (!) especially in Europe, and future networking with peers take place only if they have something to gain.

Indians go beyond that (for we are large-hearted!) in generating hospitality and friendship.

In 2012, I attended the ‘grand’ entomological Congress in South Korea. The gathering was large with nine concurrent sessions. Even 10% of the participants could not be met in four days. Big registration fee, but no lunch or tea! We had to buy them. One of the Indian participants quipped, “Many donors, no food!” For us in India, food also decides the output of such meets! And it should be so always! (Ah: I am an indulgent foodie!) One exception has been Thailand. I have attended three international meets here, and all the time my sensuous enjoyment of Thai food has been a hallmark memory.

IE has always been favouring short research notes and observations of scientific value, and to capture important in-betweens of research, which many so called ‘standard’ journals loathe to publish. But articles of longer versions, mini-reviews, and exceptional reports of a group of Insecta have started streaming into IE websites, that our Editorial felt that IE should evolve to give readers a fair share of well written, qualitative, and discursive writing. I am sure commercial

Meleponiculture by Stephen Devanesen will stimulate hiving of stingless bee, *Tetragonella* for pollination services in polyhouses. Those who adopt /recommend this please do give credit to his team from Kerala Agricultural University and ICAR. Likewise Aparna *et al.*’s., Erebid moths of Lonavala and Bakra *et al.*’s., ants of Sunderbans are very useful documentations of longer versions. All articles, finally accepted are notable and useful. I may point out the Butterfly Identification App (BIA) of Ramadevi *et al.* as a useful addendum to native libraries of schools and colleges. Articles/reviews on hormesis, immune priming, insect navigations, etc are useful reading and seminar topics for post graduate students.

We have more than 80 blogs in our websites which hardly lose their topicality as insects exhibit marvellous relevance in wider space and longer temporals.

IE has a unique professional refereeing systems. Our experiences tell us many “peer” reviewers are not up to the mark in gauging the spirit and philosophy of research contributions of authors, so tend to go by physical data in the guise of pretence to some standard! This is often found defeating creativity and stymieing originality in authors. So, our reviewers are mature scientifically empathetic subject-specialists, who goad especially youngsters into higher echelons of research. Many top-notch directors and scientists, vice chancellors and professors have all been once (and still are)

IE authors. So, it is a pleasant worthwhile experience to be an author in IE! So be our contributors, either as articles, blogs or pictures, as these behove well to a splendid career especially for youngsters in future.

So, welcome all, amateurs to professionals, to be part of insect information

dissemination for a better entomological tomorrow!

My special thanks to Dr Romeno Faleiro, FAO Consultant, for hand-holding us through the entire editorial process. Not the least, my special gratitude to IE's anchor-person, Dr M A Rashmi for her multi-tasking-coordinating to get each issue in the web-light!

Dr. Abraham Verghese
Editor-in-Chief

Research articles

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Need for surveillance in case of the invasive thrips, *Thrips parvispinus* (Karny) on chilli (*Capsicum annuum*) (Including IPM recommendations)

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The objective of this article is to sensitize all chilli growers and stakeholders in the light of the recent outbreak of the invasive thrips, *Thrips parvispinus* (Karny) on chilli (*Capsicum annuum*) in Andhra Pradesh, Telangana and Karnataka in 2021 (Sireesha *et al.*, 2021, Anitha *et al.*, 2021 and Nagaraju *et al.*, 2021). There is an urgent need for a coordinated surveillance across the country in all the chilli growing areas.

Ever since Mound and Collins (2020) reported the presence of the South-East Asian thrips, *Thrips parvispinus* (Karny) (Thysanoptera: Thripidae) on *Gardenia* plants in Greece, the fear of it traversing through the Indian sub-continent was not ruled out. Basically, a species of SE Asia with regional distribution starting from Thailand, Malaysia to New Guinea and northern Australia, its introduction to Greece was mainly through flowers of *Gardenia* sp. from Indonesia (Johari *et al.*, 2014). Though a serious pest on chilli, it has also been reported on, papaya (Hawaii) beans and solanaceous crops like chilli, potato,

brinjal (Murai *et al.*, 2009). In India, the alarm bells were rung when Tyagi *et al.* (2015) recorded *T. parvispinus* on papaya in Bengaluru, Karnataka. Subsequently, Rachana *et al.* (2018) and Roselin *et al.* (2021) reported *T. parvispinus* in 2018 on ornamental, *Dahlia rosea*, from Karnataka, India. These two Indian reports amply confirmed that *T. parvispinus* has come to stay and would infest economically important crops, especially chilli (Figs 1 & 2).

The earliest incidence of *T. parvispinus* was noticed in major chilli growing mandals of Andhra Pradesh *ie.* Chilakaluripeta and Prathipadu (16.09N 80.16E and 16.16N 80.22E) during January 2021, when chilli crop was at harvesting stage and subsequently in other major chilli growing mandals (Sireesha *et al.*, 2021). The infestation peaked in the next crop between November and December 2021 (K. Sireesha *et al.*, 2021, Anitha *et al.*, 2021 and Nagaraju *et al.*, 2021). Andhra Pradesh is the largest producer of chilli and contributes 38%, whereas Guntur alone contributes 15 %

of total production in India⁶. In Guntur district, chilli is cultivated in an area of 1,06,656 ha. Here the loss is estimated at 12% of the cropped area (uprooted) and about 50% yield loss on an average (Sireesha *et al.*, 2021).

As chilli is an important domestic and export market crop that is grown extensively as mono-crop in Andhra Pradesh, Telangana and Karnataka together comprising >70% area of the country strict surveillance is 2022 is advocated during the future seasons, using visual and trap catch documentation and additionally in all chilli growing areas.

The risk of *T. parvispinus* potentially able to spread and infest crops in neighbouring chilli growing states in India and in neighboring countries like Sri Lanka, Nepal, Bhutan and Myanmar, cannot be ruled out. An added threat is to capsicums grown in greenhouses, on which too it has been reported (Tan *et al.*, 2015). So, networked surveillance to forewarn farmers and exporters, to lower risk of loss is a high priority.



Integrated control advisory for thrips complex in chilli

1. Seed treatment with Imidacloprid WS@ 12 gm/Kg
2. Drenching liquid Arka Microbial Consortium (AMC) @ 5ml/litre in seed beds and prior to transplanting in main field (This is a product of ICAR-IIHR, Bangalore)
3. Add neem cake @200Kg/acre at transplanting along the rows of the plants
4. Spray Rashvee liquid herbal volatile soap @ 3 ml/litre, 2 weeks after transplanting or neem soap (IIHR) 5-10gms /litre
5. After two weeks of above, spray Rashvee liquid herbal volatile soap @ 3ml/litre or pongamia soap (IIHR) @ 5-10gm/litre
6. Spray vegetable special (IIHR), one month after transplanting @5gm/litre (This can be repeated before flowering).
7. Erect 50 blue sticky traps/acre one month after transplanting
8. If thrips become serious, spray Fipronil 5SC @ 1.5ml/litre
9. Spray delivery should be rotated from below to top, to ensure spray droplets also falling on underside of the leaves
10. After fruit set, if needed spray safer biopesticides as indicated above
11. mbination of Entomopathogenic Nematode+ Beauveria + Metarhizium + Verticillium has shown good results in reducing adults by infecting prepupae and pupae
12. If a second insecticidal spray is required (under expert advice only) Emamectin Benzoate 5% WG @0.4gm/lit or Spinosad @0.3 ml/litre may be given
13. Try to intercrop - for every 20 rows of chillies, one row of tomato.

14. Growing tall maize crops or *Sesbania* along the borders will reduce immigrating adult population

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Protection Advisor for encouragement. We thank Dr. R. R. Rachana, of ICAR-National Bureau of Agricultural Insect Resources (NBAIR) for identification of the thrips.

	
<p>Fig. 1: <i>Thrips parvispinus</i> adults on chilli flower</p>	<p>Fig. 2: Chilli crop highly infested with invasive thrips</p>

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Association of the elusive Fulgorid bug, *Pyrops delesserti* Guérin-Méneville, 1840 (Hemiptera: Fulgoromorpha: Fulgoridae) with pithraj tree *Aphanamixis polystachya* (Meliaceae): an observation from Kerala

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Introduction

Laternflies, the spectacularly coloured large planthoppers belong to the family Fulgoridae Latreille, 1807 (Hemiptera Linnaeus, 1758, Fulgoromorpha Evans, 1946) and has about 774 species in 142 genera according to the FLOW database (Fulgoromorpha Lists On the Web – Bourgoïn, 2022). Mostly distributed in the wet tropics of the globe, these true bugs have caught the attention and imagination of scientists, and bug enthusiasts (Constant & Pham, 2022).

The genus *Pyrops* Spinola, 1839 a member of the family Fulgoridae is set apart from other Fulgoromorphs by the presence of their typical elongated, upward curving cephalic process (Constant, 2015) and has nearly 70 species described so far (Constant & Pham, 2022). It is widely distributed in South East Asia, from Sri Lanka to the Himalayas (north India, south China), eastwards to Taiwan and Vietnam, and southwards to Sulawesi and neighbouring islands through Indonesia and the Philippines (Constant, 2015).

In India, three *Pyrops* species are currently recorded from the Andaman and Nicobar archipelago viz., *Pyrops andamanensis* (Distant, 1880), *P. rogersi* (Distant, 1906) and *P. azureus* (Constant & Mohan, 2017). *Pyrops delesserti* (Guérin Méneville, 1840), the fourth one is recorded from southern India (Constant & Mohan, 2017).

Dearth of information on their biology, natural history and host plant association still remains poorly documented except for a few species (Constant & Pham, 2022; Bourgoïn, 2022). Hence efforts were taken to shed some light into sightings of these beautiful bugs especially *Pyrops delesserti* in Kerala during 2020-2022.

Materials and methods

Purposive surveys were carried out into areas where presences of rare or elusive fulgorid insects were reported by the farmers or officials of the Department of Agriculture and Farmers welfare in the Idukki district of Kerala state, south India. Efforts were taken to

visually identify and confirm the specimens/reports through the photographs given by the farmers or with the information gleaned from the conversations with them to remove misinformation. Live insects were located and their activities were continuously observed and documented without hampering their natural activities. After a reasonable period ranging from few weeks to months of observation, couple of specimens was collected for confirmation of species identity with the help of an expert or available literature. The specimens were pinned and dried for preservation for further studies. Meticulous recording of insect behavior and host plant association were done. Pictures were taken with a Canon EOS 700 D camera with Canon Macro 100 mm lens and Canon EFS 50-250mm zoom lens.

Results and discussion

Heightened attention and media frenzy generated around the invasion of desert locust (*Schistocerca gregaria*) into Indian subcontinent and the emergence of large masses of coffee locust (*Aularches miliaris*) with bright warning colours in the state of Kerala, drove many to report unique insects that they saw. In one such incidence, a farmer confusing the aggregation of brightly colored lanternflies on a tree in his mixed cropping system, as swarms of adult coffee locust took the drastic steps of killing them. On receipt of the information and the picture of the insect through WhatsApp, quick purposive surveys

were conducted to the locality (Nariyampara, Idukki district; 9.7424° N, 77.0939° E).

Adult lanternflies were observed in groups of 3-9 individuals per group. They were observed to remain still at the same spot for more than 3 hours if undisturbed. They were later identified as *Pyrops delesserti* Guérin-Méneville, 1840. Most of the vivid earlier records pertaining to these bugs were made under the name *Fulgora delesserti* especially on their morphology, morphometrics etc (Distant, 1906, Delessert, 1843 and Atkinson, 1885). This species was recorded from Western Ghats of southern India with specific reference on Nilgiris-Malabar, Karwar and Trivandrum (Distant, 1906) and mostly prefers riverine forests (Delessert, 1843).

We could not collect or record any nymphal stages of the insect. In adult insects, cephalic process and head is greenish or brownish with white mottling. Thorax is brown with red bands at the base of the neck. Forewings/tegmina are black with reticulate venation with numerous ochraceous spots arranged in three transverse bands at the basal half with apical one more scattered. The hind wings are bluish green with a broad black border along the margin.

The wild trees on which congregations of *Pyrops delesserti* were observed was locally known as *Chemmmaram* in vernacular (Malayalam) language. Commonly called as pithraj tree (*Aphanamixis polystachya*), these evergreen trees grow to 20 m height having

reddish-brown bark mottled with green. It is widely used for therapeutic uses (Mishra, 2014) and is native to SAARC countries especially India. Even though genus *Pyrops* is well represented by many large specimens with brilliant colours globally, the specimens are hard to collect for their timid nature and long flight on slight disturbance. *Pyrops delessertii* was no exception to this behaviour, which made the documentation of their activities more difficult. The adult lanternflies when disturbed flew away from the host tree to large distances, but returned to the same tree after a gap of 20- 50 minutes. It is curious to note that the bugs dutifully returned to the same tree which stood in the middle of large assorted group of trees. This strongly points to the close affinity of the bugs to *Aphanamixis polystachya*.

Even though excretions of honey dew by the fulgorids and consequent trophobiosis with gecko and cockroaches was already

recorded with *Pyrops whiteheadi* and *P. intricatus* (Constant, 2015), we could observe only opportunistic attendance by ants in the case of *P. delessertii*.

Further dedicated surveys and observations are to be made to record various nymphal stages, host plants, trophobiosis associations and life cycle of this beautiful bug.

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Fig 1: *Pyrops delessertii* Habitus, dorsal view



Fig 2: *Pyrops delessertii* Habitus, ventral view



Fig 3: *Pyrops delessertii* on *Aphanamixis polystachya*



Fig 4: *Pyrops delessertii* aggregation *Aphanamixis polystachya*

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Exotic storage pests intercepted in raw cashew nuts shipments imported into India and cost of salvaging the intercepted material**D. K. Nagaraju*, D. Iyyanar, Maharaj Singh, T. Prema Ranjitham, N. Kasturi, B. Esakkirani, Om Prakash Verma and Ravi Prakash***Directorate of Plant Protection Quarantine and Storage, NH-IV, Faridabad 121 001, India.**Corresponding author: dkn.raju@gov.in*

India is the largest producer, processor and exporter of cashew kernels. India consumes nearly 32% of cashew produced in the world. Cashew ranks high among the Agri-horticultural commodities exported from India, thereby earning foreign exchange to the tune Rs. 4000 crores per annum. Cashew is grown in Kerala, Karnataka, Goa and Maharashtra along the West coast, and Tamil Nadu, Andhra Pradesh, Orissa and West Bengal along the East coast. Cashew is also being cultivated in Chattisgarh, North Eastern states like Assam, Manipur, Tripura, Meghalaya and Nagaland and Andaman and Nicobar Islands to limited extent.

In India, nearly 14 lakh tones of cashew nuts are processed as against annual production of 7 lakh tones. To bridge the gap, India imports raw cashew nuts from different countries. The Indian cashew industry is almost export-oriented; more than 3900 processing units are functioning in India under the organized and unorganized sector. These processing units provide sustainable employment opportunities to 1.5 million people in processing and agrarian sector, especially women thereby contributing substantially to the rural economy. The raw material availability is the

major challenge for cashew processing units. The domestic production of cashew nut meets half of the demand by cashew processing units, while the remaining is met through imports. Over 15 African and Asian countries exported raw cashew to India were cleared from Plant Quarantine Station, Tuticorin, Tamil Nadu alone. The present study records the storage pests intercepted in the raw cashew nuts imported from different countries.

The Plant Quarantine (Regulation of Import into India) Order 2003 issued under Destructive Insects & Pests Act, 1914 (Act 2 of 1914), Government of India regulates the import of all agricultural commodities into India. Imported raw cashew shipments were inspected as per provisions of Plant Quarantine Order. Ship loads of raw cashew nuts are being imported either in bulk or in gunny bags in shipping containers. Consignments in containers were inspected by partially opening the door and examined for flying insects. The outside of gunny bags were examined for any crawling insects. The bags were opened and the interior of the gunny bags were examined. The random samples drawn were spread on a white sheet and examined for insect infestation. The insects thus intercepted were

collected in 70% ethanol for identification. The specimens were sent to the National Bureau of Agricultural Insect Resources (NBAIR), Bangalore for identification.

The raw cashew nuts imported from different countries were intercepted with four species of storage pests namely, *Ahasvesus advena* (Waltl.) (Coleoptera: Silvanidae), *Cryptolestes ferrugineus* (Stephens) (Coleoptera: Cucujidae), *Ephestia cautella* (Walker) (Lepidoptera: Pyralidae) and *Carpophilus* sp. (Coleoptera: Nitidulidae) (Fig. 1).

Shipments from Madagascar and Mozambique were free of *A. advena*; those of Cote d'Ivoire, Madagascar, Tanzania and Togo were free of *Carpophilus* sp. and shipments from Burkina Faso were free of *C. ferrugineus*. *Ephestia cautella* infestation was found in shipments of all the 15 countries. Of the insects intercepted, *A. advena* was not known to occur in India, with *C. ferrugineus* having limited distribution.

During 2018–2020, 8,600 shipments of raw cashew nuts weighing 864,055 MT were imported from 15 countries. More than 80% of the shipments were imported from seven countries namely Benin, Burkina Faso, Cote d'Ivoire, Ghana, Guinea, Guinea Bissau and Senegal. Of the total shipments imported, 370 shipments were intercepted with at least one storage insect and the extent of interception

ranged a minimum of 0.67 to a maximum of 10% (Table 1).

Intercepted shipments were observed to be infested by multiples species of insects. Shipments from Madagascar were intercepted with only two species of insects, those from Togo, Tanzania, Mozambique, Burkina Faso were infested by three species. All the four species of storage insects intercepted were found in shipments from Senegal, Nigeria, Mali, Indonesia, Guinea, Ghana, Gambia, Cote d'Ivoire and Benin (Fig. 2).

The intercepted raw cashew nuts were recommended for fumigation with methyl bromide @ 32 g/m³ for 24 hours at Normal Atmospheric Pressure (NAP) (PQ Order 2003). In India, fumigations are being carried out by the Pest Control Operators (PCOs) accredited by the Directorate of Plant Protection Quarantine and Storage as per the guidelines issued under National Standard for Phytosanitary Measures (NSPM) – 12 (DPPQS, 2022). The shipments are re-inspected after 24 hours of fumigation and released for use, if found free of live infestation.

Such interceptions and subsequent fumigations add to the cost of the imported shipments and become expensive to the consumers. The approximate market rate of fumigating a 20' container is Rs. 1500/-. At this rate, importers altogether might have spent approximately Rs. 28,09,500/- (Twenty-Eight Lakhs Nine Thousand and Five Hundred)

towards treating the infested raw cashew nuts. Such consignments are re-inspected after 24 hours of fumigation to check for pest freedom, which is additional man power and cost. The cost is exclusive of container retention charges. In addition, the infested material has to be on hold for at least 24 hours, which has the cascading effect on the processing industries. Above all, methyl bromide is an Ozone depleting substance, use of which has been restricted except for Plant Quarantine purposes. Interception of insects in the imported shipments leads to use of methyl bromide on the Indian shores, which adds to the cost of imported shipments and environmental pollution (Table 2).

Under India's ambitious "One District One Product (ODOP)" program seven districts of four states namely Andhra Pradesh (Srikakulam), Assam (South Salmora), Chhattisgarh (Kondgaon), Meghalaya (West Garo Hills), Tamil Nadu (Ariyalur, Cuddalore and Pudukottai) are identified for processing of raw cashew nuts. The imported shipments are moved within country to different processing industries located in different states. Intercepted insects are storage pests and exotic, which contaminate the storage products in these processing industries ultimately leading to output of processed products.

Ahasverus advena is found in Angola, Ethiopia, Lesotho, Malawi and Nigeria. Bangladesh, Indonesia, Malaysia, Philippines, Singapore and Sri Lanka in Asia. Widespread

in Bangladesh and Sri Lanka. This species occurs on a wide variety of food stuffs, including grains, cereal products, oil seeds and their products, dried fruit, and spices. *Cryptolestes ferrugineus* found in Chad, Egypt, Ethiopia, Kenya, Namibia, Nigeria, Somalia, Sudan, Zimbabwe. Bangladesh, India (Andhra Pradesh and Haryana), Japan, Saudi Arabia, Singapore, South Korea, Sri Lanka, Taiwan, Turkey, Vietnam. Their original native habitat was probably under the bark of trees and shrubs but they have adapted to commodities in storage including wheat, barley, flour, peanuts, sorghum, oilseeds, cassava root, dried fruits, chillies (CABI, 2022). *Carpophilus*, also known as driedfruit beetles, are a worldwide pest of fruits, both pre- and post-harvest, and grains. Driedfruit beetles attack a wide variety of hosts including stone fruit, persimmons, fallen citrus, apples and figs. Adult *carpophilus* can cause feeding damage on ripening stone fruit and is a vector of the fungal disease brown rot. Most commercial feeding damage is done to ripening stone fruit. Stone fruit can be attacked on the tree, beetles burrow into the fruit, particularly near the stem end suture line. They also enter through splits and mechanical damage. In other fruits such as citrus, apples and figs, only fallen fruit is attacked (DPIRD, 2022).

Interception of large number of exotic storage insects in the imported shipments indicates the poor crop management and storage conditions in the country of origin,

failure of plant quarantine inspections carried out by the NPPOs of exporting countries, and fumigation failure, if carried out. Such instances have to be promptly brought to the notice of the exporting countries as per international norms and plant quarantine inspectors at the point of entry should always be alert while inspecting such consignments. The intercepted insects are biosecurity risk to India. Therefore, all imported shipments intercepted with storage pests were fumigated using Methyl Bromide @ 32 g/ m³ for 24 hrs at Normal Atmospheric Pressure (NAP). The treated shipments were re-inspected prior to release to ensure they were free of live infestation. The non-compliances were

notified to the trading partners on each interception as per the guidelines in the ISPM-13.

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Table 1. Raw Cashew nuts imported and intercepted with storage insects during 2018–2020.

Country	Import		Interception	
	# Shipments	Quantity (MT)	# Shipments*	Quantity (MT)
Benin	594	46,050	36 (6.06)	3,484
Burkina Faso	460	33,287	15 (3.26)	1,288
Cote d Ivoire	1,664	196,227	70 (4.21)	7,357
Gambia	236	16,416	18 (7.63)	1,274
Ghana	2,013	190,080	56 (2.78)	4,123
Guinea	677	60,346	47 (6.94)	5,566
Guinea Bissau	813	127,258	37 (4.55)	3,056
Indonesia	223	18,581	3 (1.35)	710
Madagascar	30	1,483	3 (10)	260
Mali	9	571	5 (55.56)	275
Mozambique	167	21,432	9 (5.39)	1,393
Nigeria	909	60,950	50 (5.50)	3,532
Senegal	306	34,561	17 (5.56)	1,908
Tanzania	297	44,385	2 (0.67)	865
Togo	202	12,428	2 (0.99)	2,364
	8,600	8,64,055	370	37,455

* Figures in parenthesis are percentages

Table 2. Cost of salvaging the infested material

Infested Raw Cashew	# 20' Containers (@ 20 MT/Container)#	MBR used @ 32 g/m³ (Volume of one 20' Container is 33 CuM)*	Amount spent on fumigation (Rs. 1500/20' Container)**
37,455 MT	1873 No.	2019 kg.	Rs. 28, 09, 500/-

Note: Values are indicative for the purpose of calculating approximate cost of salvaging the infested material. # Each 20' container can hold 16-20 MT of raw cashew nuts and raw cashew nuts imported in shiploads is also converted to container volume for ease of calculation. *Actual volume increases based on the sheeting skill of the fumigation operator. ** PCO's approximate market cost of fumigating a 20' container is Rs. 1500.



Raw cashew nuts in gunny bags in containers



Visual examination



Drawal of primary samples



Live beetles crawling on bags



Ahasversus advena (Waltl.)
(Coleoptera: Silvanidae)



Carpophilus sp. (Coleoptera: Nitidulidae)



Cryptolestes ferrugineus (Stephens)
(Coleoptera: Laemophloeidae).



Ephestia sp. (Lepidoptera: Pyralidae).

Fig. 1. Inspection and intercepted insects on raw cashew nuts imported from different countries.

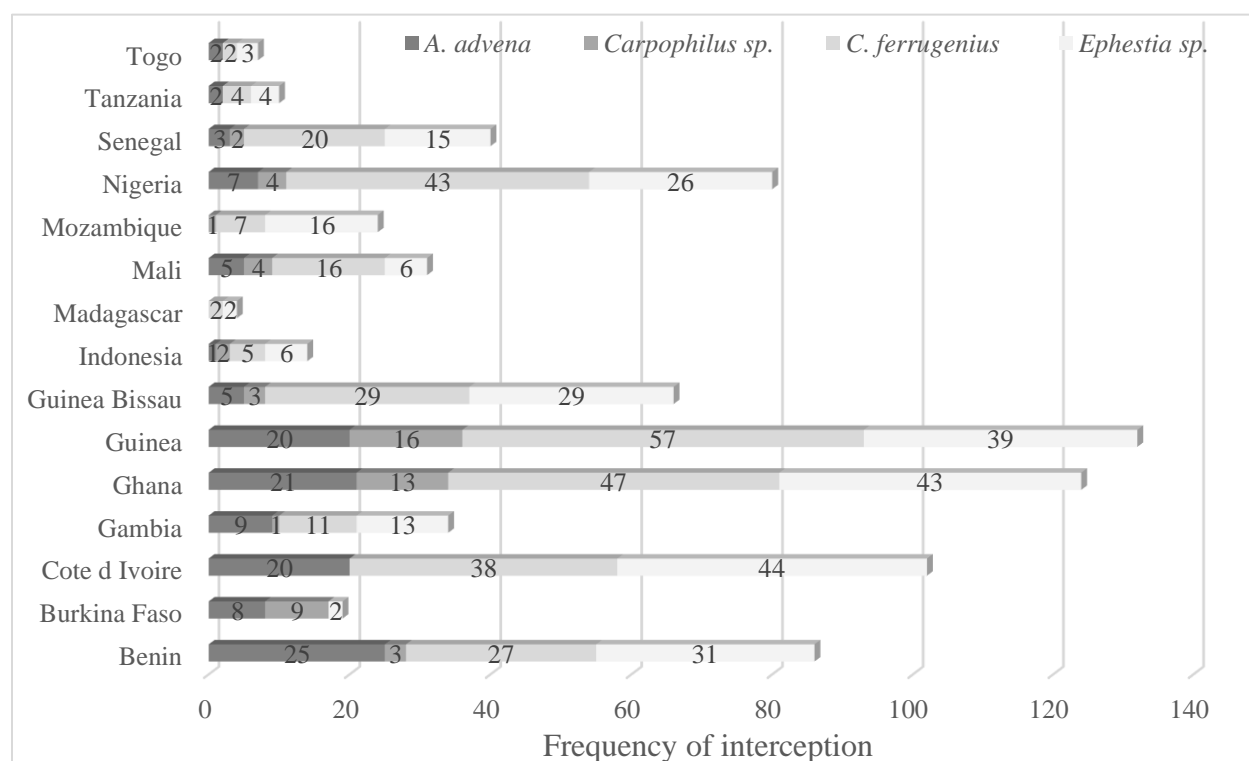


Fig. 2. Frequency of storage pests intercepted in raw cashew nuts.

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	Fruit borer	
TOMATO	Fruit borer	First spray @ 20-25 DAS, follow up spray at 15-20 days after first spray
BRINJAL	Fruit and shoot borer	First spray @ 20-25 DAS, follow up spray at 15-20 days after first spray
CABBAGE	Diamond back moth	First spray @ 15-20 DAS, follow up spray at 15-20 days after first spray
	Tobacco caterpillar, Semilooper	
OKRA	Fruit borer	First spray @ 20-25 DAS, follow up spray at 15-20 days after first spray
	Shoot and fruit borer	
RED GRAM	Spotted Pod borer (Maruca),	First spray @ 45 DAS, follow up spray at 15-20 days after first spray
	Fruit borer	



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Thrips

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Heavy incidence and damage by *Rastrococcus mangiferae* (Green) Ferris in mango**S. Kowsika*, R. Divya, N. Dilipsundar and N. Chitra***Department of Agricultural Entomology, Tamil Nadu Agricultural University, Coimbatore- 641 003, Tamil Nadu, India****Corresponding author: kowsikaentomology@gmail.com***

Mango (*Mangifera indica* L.) the ‘King of Fruits’ is an important fruit crop in tropical and subtropical countries of the world. Mango is vulnerable to a variety of pests including insects, mites, pathogens and vertebrates. Mango originated from Indo-Burma region and is cultivated in India, China, Thailand, Mexico, Pakistan, Philippines, Indonesia, Brazil, Nigeria and Egypt. India contributes about 50% of the world’s mango production with 2.5 million hectares with an annual production of 18.0 million tons (Reddy *et al.*, 2018). Insect pests pose a valid threat to sustainable cultivation of mangoes. An elaborate compilation of insect pests in mango indicates around 400 species of insect pests from different parts of the world (de Laroussilhe 1980; Tandon and Verghese 1985; Veeresh 1989; Pena *et al.* 1998). Among the insects sucking pests (leafhoppers, mealybugs, scales, thrips) and mites form a larger group causing huge yield loss. Sucking insects with shorter life cycles and ability to reproduce asexually result in their huge abundance. Further, frequent outbreaks owing to climatic variation pose serious challenge through both direct and indirect losses (Jayanthi *et al.*, 2014).

Approximately 20 species of mealybugs are reported to infest mango. Among them, *Drosicha mangiferae* (Green), *D. stebbingi* (Green), and *Rastrococcus iceryoides* (Green), are considered to be serious pests and are more frequently reported. They are widely distributed in India, Nepal, Bhutan, China, Pakistan and Bangladesh. *R. iceryoides* is also reported from Malaysia (Tandon and Verghese 1985). Under genus *Rastrococcus* of the 22 species described, three species, *R. invadens* Williams, *R. iceryoides* (Green) and *R. mangiferae* (Green) are documented in mango from India (Narasimham and Chacko, 1988 and 1991).

A survey was conducted in February, 2022 at Tamil Nadu Agricultural University (TNAU) orchard (11.005567 N, 76.930086 E). Fifty to sixty years old trees were observed with heavy infestation of the ornate mealybug species completely covering the foliage. Population densities of the mealybug were noted to be significantly higher in the mango trees on the abaxial leaf surface; approximately 10 to 15 numbers/cm² was observed. Infestation was severe on older leaves later shift towards young tender leaves was noticed. Appearance of sooty mold on leaves and

parasitized mealybug cadavers with exit holes were also observed.

Externally adults were observed to be oval in shape with dorsum pale yellow and with thick white mealy coating. Stout cottony tassels are seen on the margin, increasing in length from anterior to posterior end. Both immatures and adult were present with 30 prominent wax tassels.

The collected specimens were slide mounted based on the protocol given by Bahder *et al.*, (2015) with slight modifications and observed under phase contrast microscope LEICA DM750. Based on the following characters: antennae slender with 9 segments, circulus oval shaped situated in middle of third abdominal segment, posterior pair of ostioles,

anal ring situated a short distance from apex of abdomen, cerarii numbering 15 pairs, each on a more or less round to oval sclerotized base, larger in area than anal ring, trilocular disc pores being present in a ventral marginal zone on thorax, the specimen was identified to be *R. mangiferae* which was in concurrence with the key to species of *Rastrococcus* provided by Williams, 1989.

The incidence of *R. mangiferae* was reported from Coimbatore India by Ramakrishna Ayyar on mango in 1914. Narasimham and Chako (1988) have reported this species to be more common in southern India than *R. invadens*. If left unnoticed this pest may lead to significant loss in size and weight of fresh mangoes and become a growing threat to mango orchards.



***Rastrococcus mangiferae* found on the abaxial leaf surface**



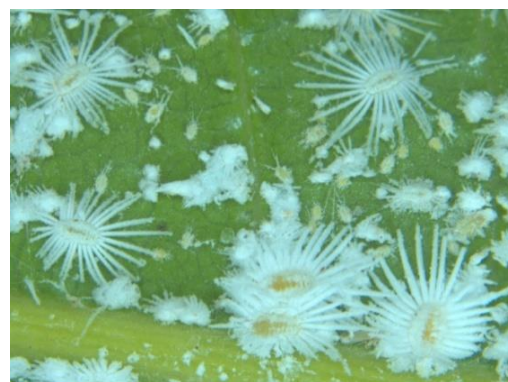
Female



Male



Parasitized



Unparasitized

***Rastrococcus mangiferae* (Green) Ferris 1954**

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Monitoring of major insect pests, correlation and yield loss in pearl millet**R.P. Juneja*, G. M. Parmar, R.J. Chaudhari and K. D. Mungra***Main Pearl Millet Research Station, Junagadh Agricultural University, Jamnagar-361006, Gujarat.**Corresponding author: rajkumarjuneja19@gmail.com***Abstract**

Investigations were carried out at Main Pearl Millet Research Station, Junagadh Agricultural University, Jamnagar (Gujarat) during *kharif* 2021 revealed that shoot fly, *Atherigona soccata*, stem borer, *Chilo partellus*, and ear head worm, *Helicoverpa armigera* were the major insect-pests in pearl millet. Significant correlation was found of shoot fly, stem borer and *Helicoverpa armigera* with different weather parameters. There was a considerable loss in yield due to insect-pest complex in pearl millet.

Key Word: Shoot fly, stem borer, *Helicoverpa armigera*, correlation, yield loss.

Introduction

In India, pearl millet, *Pennisetum typhoides* (Burm.) or *bajra* is grown in Rajasthan, Maharashtra, Gujarat, Uttar Pradesh, Andhra Pradesh, Haryana, Tamil Nadu and Karnataka (Prem Kishore and Solomon, 1989). It occupies an area of 6.93 million ha with an average production of 8.61 million tones and productivity of 1243 kg/ha (Anonymous, 2020). Pearl millet is generally preferred in low rainfall areas and on light soils. This crop has wide spectrum adaptability in respect of rainfall, temperature and soil. It is generally believed that pearl millet either grown as mono crop or mixed crop or in relay cropping system has hardly had any serious problems. However, perusal of literature on insect pest of this crop gives quite a different picture. Twenty six insects and two non-insect

pests were found feeding on pearl millet (Balikai, 2010). Out of these, shoot fly, *Atherigona soccata*, stem borer, *Chilo partellus* Swinhoe and ear head worm, *Helicoverpa armigera* are comparatively more serious pests attacking the crop. The need to study the effect of different weather parameters on insect-pest incidence has also arisen in present situation of climate change. Hence, study was under taken.

Material and methods

The experiment was taken during *kharif* 2021 at Pearl Millet Research Station, Junagadh Agricultural University, Jamnagar (Latitude:22.46816, Longitude:70.02855, Altitude:64 ft). Sowing of released pearl millet variety (GHB 558) was done over an area of 200 m² which was kept free from insecticidal

application during crop season. During, *khariif*-2021, monsoon commenced in the third week of June. Total rainfall *i.e.* 800 mm was received in 28 rainy days. Sowing for entomological trials was done on 20th July, 2021. After sowing there was long dry spell and hence irrigation was given. All the agronomical package of practices was followed from time to time. Incidence (%) and population of various insect pests observed during the crop period was recorded at weekly interval from 20 randomly selected plants seven days after germination (DAG) of the crop till maturity. The presence of bio agents was also recorded simultaneously. Weather data was also recorded on weekly basis (Meteorological Standard Weather Week) for correlation. The correlation of major insect pest was worked out. Simultaneously, one treated plot was maintained and kept insect-pest free by taking recommended package of practices for insect pest management to get the information for losses. The following treatments were adopted for treated plot.

1. Seed treatment imidacloprid 600 FS @ 8.75 ml/kg seed was given for protection against shoot fly, stem borer & white grub at early crop stage. For later stage crop protection (Shoot fly & stem borer) foliar spray of fipronil 5 SC @ 0.01%, at 35 days after germination of the crop was given.
2. Spray of Novaluron 10 EC 0.01%, at ear head stage at pest appearance of the *Helicoverpa armigera* was given.
3. For leaf binder, grass hopper, grey weevil and hairy caterpillar spray of neem seed kernel extract 5% was done.

Results and discussion

(A) Insect-pest incidence:

Shoot fly: The initiation of shoot fly incidence was found in 31st SWW (5.0%). The highest shoot fly incidence (25.0%) was observed during 40th SWW. The average incidence was 13.18% during the crop period. **Stem borer:** The initiation of stem borer was found from 32nd SSW (5.0%). However, its incidence was found highest (25.0%) during 36th week with an average incidence of 12.73% during the crop period. The overall range of other insect pests *viz.* White grub (0.0-10.0%), leaf roller (0.85-2.50 damage score), grass hopper (0.0-5.0%), grey weevil (0.0-1.85 damage score), hairy caterpillar (0.0-10.0/20 ear heads), chaffer beetles (0.0-7.0/20 ear heads), blister beetle (0.0-6.0/20 ear heads), *Helicoverpa* (0.0-12.0 larvae/20 ear heads). The incidence of fall army worm was not observed during the study period. The natural enemy ladybird beetle population initiated during 32nd SWW (2 adults/20 plants) and was observed till the end of the season ranging from 0.0 to 25.0 adults/20 plants. The average population was 11.73 adults/20 plants. *Chrysopa* population was initiated during 33rd SWW (1 adult/20 plants) and was observed till 36th SWW *i.e.* during mid crop stage. The overall range was 0.0 to 2.0/20 plants with an average of 0.55 adults/20 plants, table-1.0.

(B) Correlation

The correlation of shoot fly with temperature minimum (-0.671^*) was found negatively significant and highly and negatively significant with wind speed (-0.832^{**}). The correlation of stem borer was again negatively significant with temperature minimum (-0.629^*) and highly and negatively significant with wind speed (-0.809^{**}). As far as *Helicoverpa armigera* was concerned its correlation was found negatively significant with wind speed (-0.608^*) and negatively highly significant with evaporation rate (-0.725^{**}), table 1.1. Raghvani *et al.*, 2008 reported that none of the weather parameters showed significant correlation with incidence of shoot fly. However, maximum temperature and difference of minimum and maximum temperature exhibited negative correlations with stem borer. While significant positive association of minimum temperature was observed with larval population of *Helicoverpa* and correlation with rainy days was found significantly negative.

(C) Losses

The treated plot recorded 2343 kg/ha grain and 4566 kg/ha fodder yield (Table-1.2). The losses in grain and fodder yield were 16.41% and 20.01%, respectively. Prem

Kishore, 1996 reported that, shoot fly causes 23.3 to 36.5 percent grain losses and 37.5 per cent fodder loss. Whereas, borers indicated losses varied from 20 to 60 per cent. Juneja and Raghvani (2000) reported that on an average 10 to 15 per cent reduction in yield was observed due to *Helicoverpa armigera*.

Conclusions

The above study on monitoring of major insect pests in pearl millet during *kharif* 2021 revealed that there is huge difference in the intensity of different insect pests. These insect pests are collectively responsible for loss in yield. Moreover, weather played an important role in the pest incidence.

Acknowledgements

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Table-1.0: Incidence/population of insect-pests and natural enemies in pearl millet during *kharif*,2021

No.	SWW	Date of Observation	Days After Germination	Shoot fly % incidence	Stem borer % incidence	White grub % Inci.	leaf roller damage score (0-10)	Grass hopper % damage	FAW % damage	Mean Grey weevil damage score (0-10)	Hairy caterpillar /20 pl.	Blister beetles/ 20 EH	Chaffer beetle 20 / EH	Helicoverpa larvae/ 20 EH	Lady bird beetle/ 20 pl.	Chrysopa/ 20 pl.
1	30	26.07.21	7 DAG	0.00	0.00	0.00	0.85	0.00	0.00	0.00	0	0	0	0	0	0
2	31	02.08.21	14 DAG	5.00	0.00	0.00	1.00	0.00	0.00	0.00	0	0	0	0	0	0
3	32	09.08.21	21 DAG	5.00	5.00	5.00	1.10	5.00	0.00	0.40	0	0	0	0	2	0
4	33	16.08.21	28 DAG	10.00	10.00	10.00	1.45	5.00	0.00	0.60	0	0	0	0	5	1
5	34	23.08.21	35 DAG	15.00	15.00	10.00	1.62	5.00	0.00	0.80	0	0	0	0	10	1
6	35	30.08.21	42 DAG	20.00	20.00	10.00	1.80	5.00	0.00	0.85	0	0	0	0	12	2
7	36	06.09.21	49 DAG	10.00	25.00	10.00	1.95	5.00	0.00	1.00	0	0	0	0	15	1
8	37	13.09.21	56 DAG	15.00	10.00	10.00	2.10	5.00	0.00	1.20	4	3	0	0	18	1
9	38	20.09.21	63 DAG	20.00	15.00	10.00	2.20	0.00	0.00	1.65	7	5	4	5	20	0
10	39	27.09.21	70 DAG	20.00	20.00	10.00	2.25	0.00	0.00	1.75	8	5	6	8	22	0
11	40	04.10.21	77 DAG	25.00	20.00	10.00	2.50	0.00	0.00	1.85	10	6.00	7	12	25	0.00
			Mean	13.18	12.73	7.73	1.71	2.73	0.00	0.92	2.64	1.73	1.55	2.27	11.73	0.55
N.B.: Date of sowing: 20.07.2021, Harvesting: 15.10.2021, Variety: GHB 558																

Table-1.1: Correlation of major insect-pests of pearl millet with different weather parameters

No.	SWW	Date of Observation	Days After Germination	Temp. C Maxi.	Temp. C Mini.	R.H. Morn.	R.H. Even.	Wind speed km/hr	BSS (hrs)	Eo (mm)	Rainfall (mm)	Rainy Days
1	30	26.07.21	7 DAG	32.0	27.1	93	76	15.2	1.4	5.6	30.0	2
2	31	02.08.21	14 DAG	32.1	26.5	85	67	16.2	1.6	5.6	4.5	1
3	32	09.08.21	21 DAG	33.0	25.8	85	66	9.3	5.8	6.6	0.5	0
4	33	16.08.21	28 DAG	33.0	25.4	84	65	10.1	7.0	6.8	0.0	0
5	34	23.08.21	35 DAG	32.7	25.9	86	65	9.8	6.1	6.9	2.0	0
6	35	30.08.21	42 DAG	33.1	25.5	89	72	8.0	5.0	7.2	68.5	2
7	36	06.09.21	49 DAG	31.4	25.6	93	83	8.2	2.7	5.4	91.0	4
8	37	13.09.21	56 DAG	30.3	24.8	95	83	7.2	1.1	4.7	204.0	5
9	38	20.09.21	63 DAG	32.5	26.1	91	76	7.8	6.5	4.8	6.5	1
10	39	27.09.21	70 DAG	31.4	24.8	94	86	4.9	3.3	4.2	244.0	4
11	40	04.10.20	77 DAG	32.6	25.2	90	73	5.1	8.1	4.1	7.0	1
		Correlation	Shoot fly%	0.027 ^{NS}	<u>-0.671*</u>	0.258 ^{NS}	0.276 ^{NS}	<u>-0.832**</u>	0.512 ^{NS}	-0.372 ^{NS}	0.280 ^{NS}	0.153 ^{NS}
			Stem Borer%	-0.051 ^{NS}	<u>-0.629*</u>	0.349 ^{NS}	0.437 ^{NS}	<u>-0.809**</u>	0.360 ^{NS}	-0.205 ^{NS}	0.334 ^{NS}	0.335 ^{NS}
			<i>Helicoverpa</i> /5 EH	0.003 ^{NS}	-0.384 ^{NS}	0.287 ^{NS}	0.305 ^{NS}	<u>-0.608*</u>	0.437 ^{NS}	<u>-0.725**</u>	0.157 ^{NS}	0.047 ^{NS}

Table-1.2: Yield losses in grain & fodder due to insect-pest complex in pearl millet.

Parameters	Yield kg/ha in treated plot	Yield kg/ha in Un-treated plot	% losses
1. Grain	2343	1959	16.41
2. Fodder	4566	3652	20.01

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Butterfly Identification App (BIA): A mobile application for identification and monitoring of butterflies in the state of Karnataka**O.K. Remadevi*, J. Cruz Antony, Jane Maria D Souza and K.H. Vinaya Kumar***Centre for Climate Change, Environmental Management and Policy Research Institute (EMPRI), Bengaluru 560 078, Karnataka, India.***Corresponding author: okremadevi@gmail.com***Abstract**

Butterflies attract the attention of everyone, but it is difficult to identify and know their names instantly. Even for the students and researchers, identifying them in the field itself necessitates capturing and referring field guides. A novel mobile application known as Butterfly Identification App (BIA) has been developed by Environmental Management and Policy Research Institute EMPRI, Bengaluru, India to identify butterflies in the field. It is a colour-based identification application, which can be used by anyone having an android mobile phone. It is not necessary to capture the butterflies but photos taken can be compared with the photos in the colour based groups in the BIA database. The identification of all butterflies reported from Karnataka is possible through the BIA. The usage of this by proficient taxonomists, students and environmentalists can contribute to the monitoring and conservation of butterfly species in their localities. Pilot studies conducted in six green spaces in Bengaluru (12.9716° N, 77.5946° E) with GPS locations (Lalbhag: 12.9487,77.5887; Cubbon Park: 12.9798,77.5968; Doresanipalya: 12.8971,77.5905; IISc: 13.0173,77.5712; GKVK: 13.0808,77.5677; Bannerghatta: 12.785,77.5745) using the BIA is presented in the paper.

Keywords: Android mobile, Butterfly Identification App, EMPRI.**Introduction**

According to Ghazoul (2002), about 19,238 species of butterflies have been documented from all over the world, which includes 1501 species from India (Kunte *et al.*, 1999). Ashish *et al.*, (2007) have reported 332 species from the Western Ghats. In Karnataka, about 323 species of butterflies have been recorded (Remadevi *et al.*, 2020) and around 140 species are found in Bengaluru alone

(Yates, 1933; Remadevi *et al.*, 2018). Climate change and environmental degradation are indicated by butterflies. Butterflies react faster to ecological changes than birds and vascular plants (Thomas *et al.* 2004). They are important components in the food chain of birds, reptiles, spiders and predatory insects. They help in pollinating many economically important crops. The crucial information on the ecology of a particular region is provided by a faunistic survey of butterflies based on

their occurrence and characteristics (Ghazoul, 2002). In general, monitoring butterfly populations are considered an essential means of evaluating change in the ecosystem and also the state of habitation for biodiversity (Paul and Sultana, 2020). Van Swaay *et al.*, (2012) stated that these trends contribute as an indicator of biodiversity status and to interpret the environmental change. Butterflies act as best bio-indicators for effect on climate change (Ghazanfar, et al., 2016). Since butterflies are one of the most eye-catching insects with various colour patterns, the identity of a butterfly could be mistaken.

Utilizing Information and Communication Technologies (ICT) in agriculture and economy can boost production in the shortest and fastest way (Nitin *et al.*, 2020). Smart phone technology helps in field for delivery of information. In contrast to printed publications, the information conveyed through the mobile application can be augmented, restructured, updated, and corrected repeatedly. For extension professionals in turf grass management, the development of application and the delivery of information through this technology present an invaluable, fast, accurate, new resource and delivery of information (McCullough *et al.*, 2011).

Identification of butterflies using mobile applications is being practiced in many countries. Fox *et al.*, (2015) through the scheme “Butterflies for the New Millennium

(BNM)” which runs in partnership with the Biological Records Centre (part of the UK Centre for Ecology & Hydrology) surveys butterfly distributions across the UK. The BNM scheme uses smart mobile technology to aid and strengthen the recording of butterfly distribution. A free mobile app named “iRecord Butterflies” was developed and more than 10,000 users have started using this app since the launch of the app in 2014. Using the app, a butterfly surveyor/recorder can log and submit sightings as a single process and also make use of the built-in GPS facility in their mobile phone to automatically generate high-resolution grid references for their records. The app also acts as a simple identification guide to help new surveyors/recorders and guides them through the process of submitting butterfly records.

Andre Poremski (2019) developed a mobile app named “Leps” which is funded privately by Public Good Projects, United States. Leps was developed using image recognition technology to identify butterflies and moths (Lepidoptera) which is achievable with a photo, date and location of the species observed. The app uses a machine learning framework named “Fieldguide” to help identify the butterflies and moth. The “Butterfly Id” is a paid mobile app developed by Sunbird Images, Germany. This app is used to identify butterfly species very easily with available butterfly records created by professional ecologists. This app also provides information about the anatomy, ecology and

taxonomy of the butterflies. To aid in fast identification of the species, this app provides information about every butterfly's hind and forewings.

Shubhalaxmi *et al.* (2016) have developed a butterfly mobile app under the U.S. State Department's International Exchange Alumni programme. The mobile app characterizes 50 common Indian butterfly species with their images, common and scientific name, host plants and other related facts. Theivaprakasham Hari (2019) has developed an app called "Butterfly Vision" which helps in the identification of nearly 300 butterflies from the Western Ghats region. This app uses a machine learning technique to help identify butterflies in the field. This app was developed to exclusively engage school kids to identify butterflies and thus helps in the documentation of butterfly fauna with the adoption of technology. This app is linked with the website <https://butterflyvision.in/> which also provides facility to upload the captured butterfly images and in return helps to identify the butterfly with similarity index using percentage.

The Nature Web's "Indian Butterflies" app is an offline mobile app which provides butterfly information along with the butterfly's regional names viz., Assamese, Bengali, Marathi, Malayalam and Tamil. "Indian Butterflies" app is easy to use as a field guide by providing information such as habitat, host plant, wingspan, sexual differences, etc. This

app is linked with <http://www.indiabutterflies.com> which acts as the butterfly repository. It also allows creating checklists of the on-field observations and can be further exported into CSV file.

1. Materials and Methods

Thus, BIA has been developed to identify the butterflies based on their major colour in the wings. This mobile app can be used for the continuous monitoring of butterflies in specific areas with the involvement of naturalists, officials from the forest department, eco clubs, school children etc.

1.1 Butterfly Identification Mobile Application (BIA)

For monitoring the diversity and the impact of climate change on butterflies in Karnataka, an android mobile app (BIA - butterfly identification app) was developed. BIA is extremely simple to use and it can be downloaded into any android mobile and the information such as GPS of the location, name of the area, date and time along with the identified butterfly information reaches the BIA dashboard through the internet transfer protocols. The minimum requirement for the operating system to download the BIA is Android 5 Lollipop. BIA allows taking pictures of butterfly resting with open or closed wings in field conditions. BIA app would help in identifying nearly 300 butterfly species of Karnataka. In BIA, the butterflies of Karnataka, a state in south India are classified

based on their major colours like white, blue, yellow, orange, brown and black. Photo taken can be zoomed in and out and matched with details and image of the butterflies that are available in the BIA database, for identification. If the identification is confirmed, by pressing the SUBMIT button the information collected will be submitted to the database. In case the captured butterfly is not present in any of the colour code available, then the identification can be submitted as UNIDENTIFIED. Later this category will be checked and processed by the resource persons available at EMPRI to identify or add the new species into the BIA database.

1.1.1 Steps involved in using BIA app

- 1) Open the app and press the “Start” button
- 2) Notice the butterfly in the field and take photos of open / closed wings of a butterfly. Press the “Skip” button if the second photo cannot be taken
- 3) To acquire only the image of butterfly, crop the background
- 4) Notice the major colour in butterfly wings and press a similar colour button
- 5) Butterflies photos with chosen colour will be displayed below
- 6) Scroll, compare and match with the butterfly photo taken and the butterfly photo available from the database
- 7) Tap and zoom the photos to notice the wing patterns and read about wings description
- 8) After verifying the identification, press the SELECT button
- 9) Identified butterfly details will be displayed for further verification
- 10) Finally, press the SUBMIT button to send the identified butterfly data

1.2 BIA Dashboard

The identified/unidentified information about the butterflies sent from BIA such as the butterfly name and photos along with the geographical information fetched using the mobile phone’s GPS system are transferred to the dashboard available at Karnataka State Climate Change Strategic Knowledge Portal (KSCCSKP) Link - <http://skccempri.karnataka.gov.in>. The portal was developed under the Department of Science and Technology, DST-SPLICE funded project. The main objective of KSCCSKP is to disseminate information on climate change and related activities in the state of Karnataka. The dashboard is developed on the basis of 2-tier architecture using Laravel PHP and MYSQL software application. The dashboard displays various reports on identified butterflies based on the family, region and year-wise using pie charts and displays the butterfly name (scientific and common name) and geographical information. Fig.-2 represents the flow diagram of the complete butterfly identification process.

2. Results and discussion

The BIA app is intended for documenting the diversity in different seasons and has particularly been developed to be simple to use. A training workshop was conducted on how to use BIA android application to eco-club school teachers as a part of NGC activity at EMPRI on 2nd Feb 2019 and a pilot testing using BIA for the identification of butterflies at EMPRI campus was organized. As an outcome, 55 records were received until June 2019. Further, a presentation was provided to Karnataka districts teachers from eco-club schools through SATCOM (Satellite communication) and after that 42 records were received until October 2019. A pilot test study using BIA was made during winter seasons in 2019 and 2020 (October, November, December and January) in the 6 green spaces of Bangalore and a total of 69 species were recorded in different regions.

Fig. 1 depicts the total butterfly species found in different survey areas Doresanipalya has the highest number of species found with 22 per cent (69 species) followed by Bannerghatta with 21 per cent (68 species). IISc has 19 per cent (59 species) and GKVK has 18 per cent (58 species) of butterfly species. Lalbagh has the second least butterfly species with 12 per cent (39 species) and least is Cubbon Park with 8 per cent (26 species) of butterfly species.

Fig. 2 depicts butterfly families found in different survey regions. Nymphalidae is

found to be dominant in all the study areas with 25 species in Doresanipalya and Bannerghatta and 9 species in Cubbon Park which is least among other areas. Lycaenidae is found to be second most dominant with 18 species in Doresanipalya and 17 species in Bannerghatta and least in Cubbon Park with 6 species. Pieridae and Papilionidae are moderate with 15 species in Bannerghatta and 8 species in Doresanipalya, GKVK and IISc respectively. Riodinidae is the least in Doresanipalya and Bannerghatta with one species whereas it is not found in the other four study areas. Hesperidae is second least with 7 species in Doresanipalya and 4 species in Lalbagh and GKVK.

Conclusions

Over the past centuries, many researchers have significantly contributed to the field of butterfly diversity and ecology within the various ecosystems in India. Monitoring diversity in different geographical locations, integrating more than the conventional hotspots, makes it quite feasible to identify changes over an extensive range of habitations. This might provide an enhanced knowledge of the species associations in various vegetation categories. The phenological changes in plants due to climatic variability and changes in different areas is reflected in the diversity of butterflies in the area. A mobile application “Butterfly Identification Application (BIA)” is developed by EMPRI to achieve the dual purpose of helping butterfly field identification

effortlessly and receiving the butterfly diversity information for further studies of butterfly ecology in Karnataka in relation to climatic and environmental changes. BIA helps in easy identification of butterflies by anyone from anywhere in Karnataka. Currently, the app helps in the identification of 323 butterfly species reported from Karnataka. BIA can be downloaded in any Android mobile and the information such as GPS of the location, name of the area, date and time along with the identified butterfly information reaches the BIA dashboard through the internet

transfer protocols. The BIA dashboard is developed using Laravel PHP and MYSQL for database and has linked to both EMPRI website and the Karnataka State Climate Change Strategic Knowledge Portal (KSCCSKP). This BIA dashboard developed for processing the butterfly data is hosted in Karnataka State Data centre (KSDC) and shall be accessible to everyone interested to know the butterfly information of the state. The link to the BIA dashboard is <https://bmpempri.karnataka.gov.in/>.

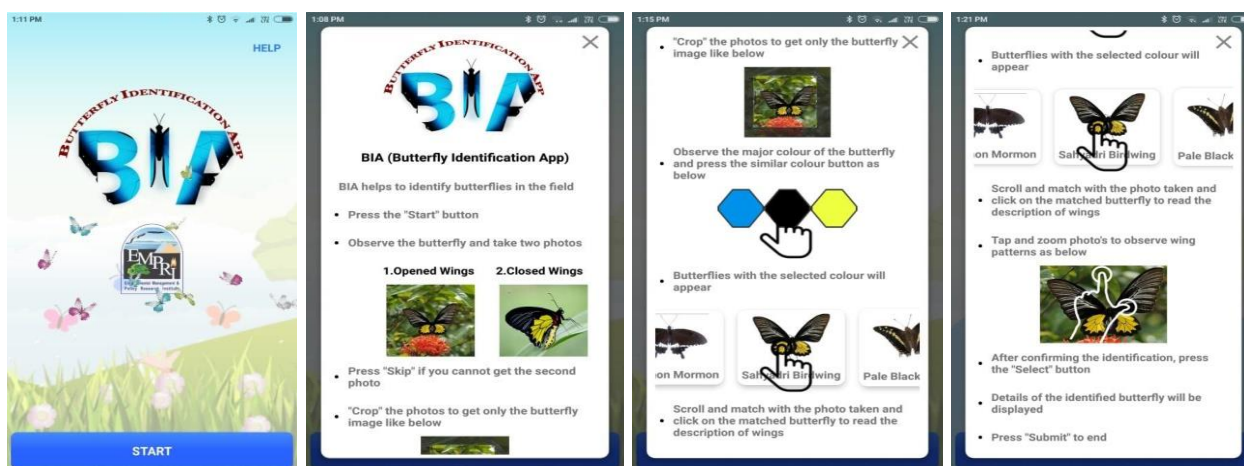


Fig. 1: Butterfly Identification Application screenshots

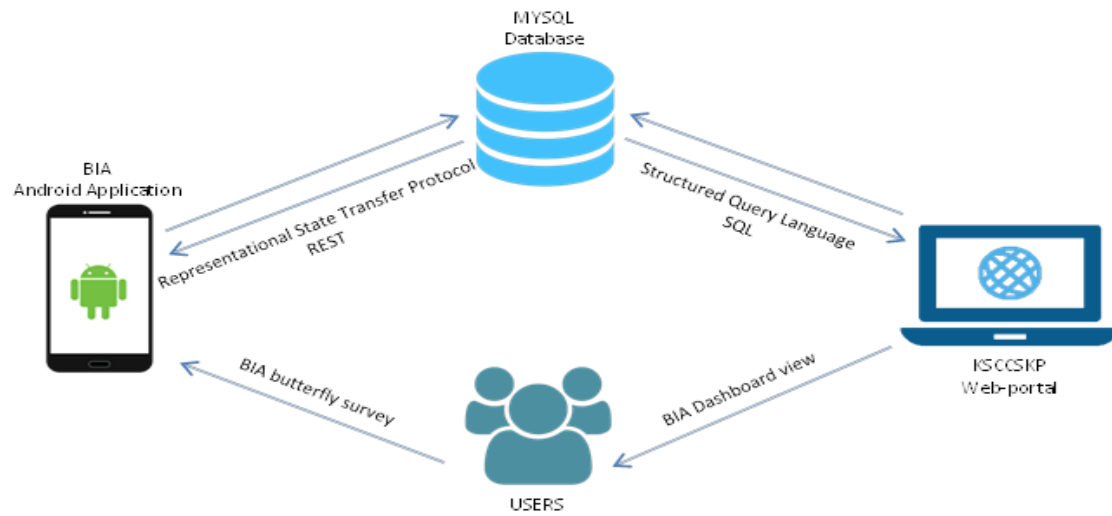


Fig. 2: Flow diagram of Butterfly Identification Application

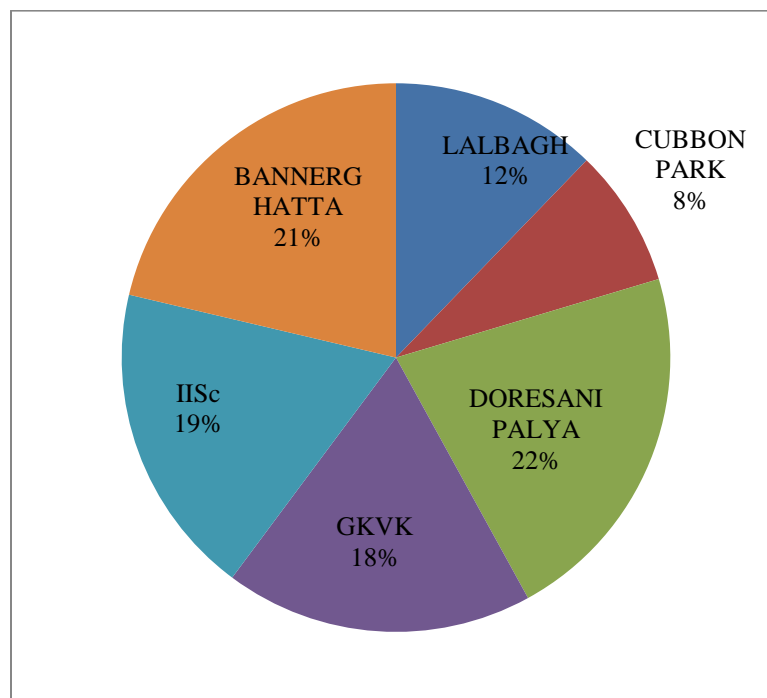


Fig. 3: Butterfly species recorded by BIA in survey areas

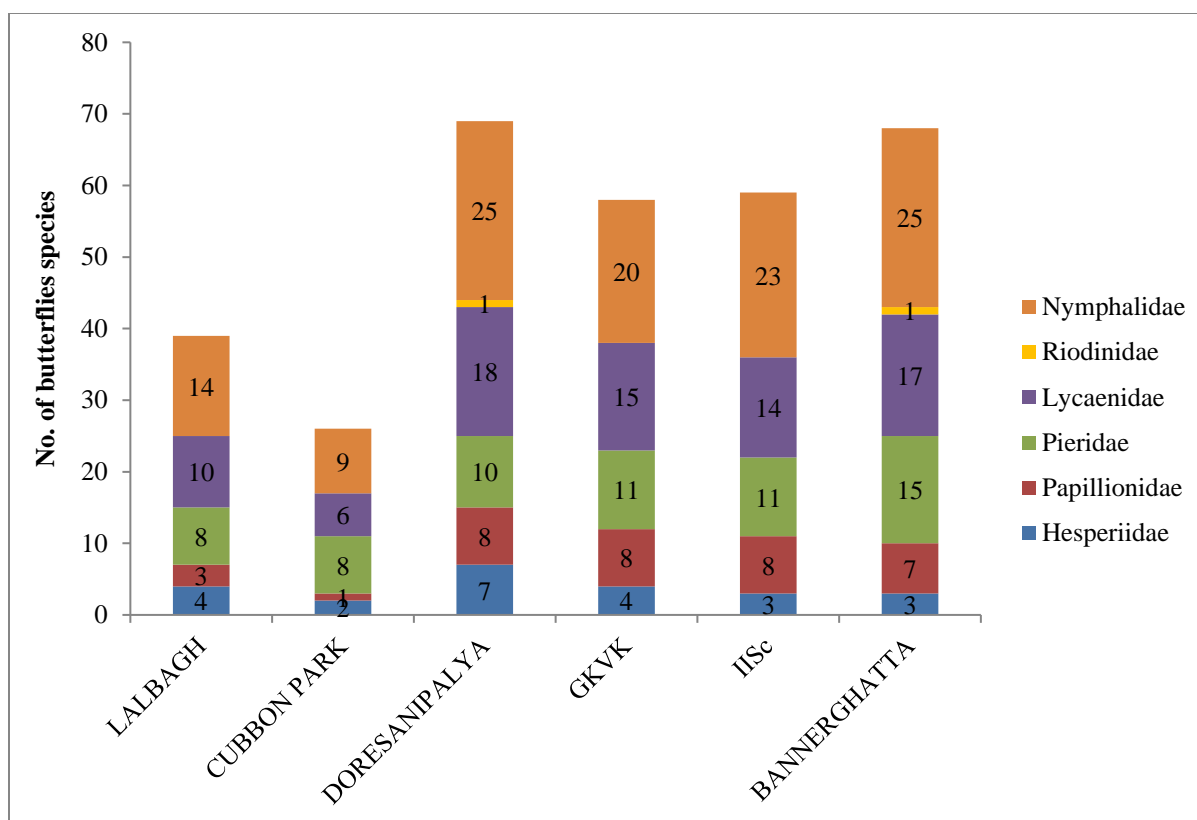


Fig. 4 Butterfly family-wise distribution in different areas

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A note on the biology, host plants and distributional record of Goa's state butterfly- *Idea malabarica***Channabasava Veershetty and R. Maruthadurai***Crop Science Section,**ICAR – Central Coastal Agricultural Research Institute, Ella Old Goa – 403 402***Corresponding author: basava6959@gmail.com***Abstract**

The Malabar tree nymph butterfly, *Idea malabarica* (Moore, 1877) (Nymphalidae: Lepidoptera) was declared as state butterfly of Goa at the fifth Goa bird festival – December 2021 by Government of Goa. This butterfly is familiar for its captivating black and white wing pattern and unique gliding flight hence it is also known “Paperkite”. *Idea malabarica* is endemic to the moist evergreen forests of Western Ghats and is categorized as “near threatened” species by International Union for Conservation of Nature (IUCN). It undergoes complete metamorphosis with six week of life period. Five instar larval phase feeds mainly on *Aganosma cymosa*, a species of Apocynaceae family. Losses of habitat and climate change are the major concerns to be dealt for the conservation of butterflies, which are indicators of healthy ecosystem and healthy environment.

Keywords: Malabar tree nymph, Butterfly, Western Ghats, IUCN, climate change

Introduction

Idea malabarica, the Malabar tree nymph is one of the largest milkweed butterfly confined to wet evergreen forests of Indian peninsula. This attractive butterfly is in current news of India because the state government of Goa (India) has declared Malabar tree nymph butterfly as their state butterfly on December 2021. Goa, a tiny state of India with rich biodiversity of different flora and fauna is sandwiched between the Arabian Sea and Western Ghats. Though being the smallest state of the country, Goa has endowed with 215 species of butterflies (Gaude and

Janarthanam, 2015). The steps like, the state butterfly status will encourage in developing conservation measures of Malabar tree nymph butterfly in this region as it falls under the “Near Threatened” category of International Union for Conservation of Nature (IUCN). This review article emphasizes on the classification, biology, habit, habitat, host plants and importance of Malabar tree nymph butterfly *I. malabarica*. These studies would be helpful in developing broad approaches for conservation of *I. malabarica* and other butterfly diversity.

Scientific classification of Malabar tree nymph butterfly

Kingdom:	Animalia
Phylum:	Arthropoda
Class:	Insecta
Order:	Lepidoptera
Family:	Nymphalidae
Genus:	<i>Idea</i>
Species:	<i>malabarica</i>
Binomial name:	<i>Idea malabarica</i> (Moore, 1877)

Biology

Malabar tree nymph butterfly experiences a holometabolous life cycle consisting of four stages: egg, larva, pupa and adult. The butterflies of genus *Idea* takes around six weeks to complete their life cycle (Cabras et al., 2015).

Egg: Adult butterfly lays oval shaped, white creamy or translucent eggs singly on under surface of the host plant leaves (Figure 1). The cream-colored eggs turn pinkish on the 4th and 5th day when they are about to hatch (Cabras et al., 2015). The maximum egg duration may vary according to the season and climatic condition (Evans, 1932).

Larva: The caterpillar (Figure 2) is stout and smooth which undergoes five instars (Instars I, II, III, IV, V) of developmental stages. After hatching, the neonate larva starts feeding on empty egg shell and epidermis of the host plant leaves. Later instars feed rigorously on the different parts of the host plant and grow

bigger in size. The morphological characters of fully grown matured caterpillar are black body with white stripes and red spots on its lateral side. One can see varying number of long black spikes/horn like structures protruding on the dorsal surface of the body, which are used for protection against predators. Before entering into the pupal stage, the last larval instar stops feeding, becomes lethargic and shrinks in its body size (Pre-pupal stage). Usually, the butterflies of genus *Idea* take around three weeks to complete the larval period (Kumar, 2005; Cabras et al., 2015).

Pupa: The pupal form is known as “Chrysalis” which is initially golden colour ornamented with black shiny spots over its surface (Figure 3). Later from 8th day onwards black pigmentation starts to appear on the surface and on 14th day, a day before adult emergence it turns black. On the under surface of leaves the chrysalis suspends free from the anal hook or cremaster without any silken girdle (Cabras et al., 2015).

Adult: Malabar tree nymph appears as a white butterfly with black markings and the wingspan measures generally about 120-154 mm (Figure 4). The upper sides of both the wings are semitransparent white with powdery black scales. These have long black antennae with rounded clubs at the end, head and thorax spotted and streaked with black and abdomen is white, with broad dusky black streak above (Moore, 1890; Thomas, 1905; Varshney and Smetacek 2015).

Habit

Malabar tree nymph butterfly is slow and week flier with flapping movement of wings. A gliding flight is common habit of the genus *Idea* so these are also called as “Paperkite”. They can be seen often gliding above the forest tree canopy but infrequently move lower down in forest openings (Evans, 1932).

Habitat and distribution

Idea malabarica have specific habitat requirements depending upon their feeding and reproduction needs. It is endemic to the wet evergreen forests with heavy rainfall areas which include Western Ghats stretches of Maharashtra, Goa, Karnataka and Kerala (Bringham, 1905; Manoj and Sharma, 2013). Also found in few parts of Tamil Nadu bordering to Kerala state (Arun, 2003; Dunstan and Raj, 2005; Alagumurugan *et al.*, 2011). In Goa it is found mainly in the areas of swamp vegetation of Ajobachi Tali, Bibtyan, Nirankarachi Rai and Mharinginichi Rai in Sattari and Bhati, and Savari in Sanguem (Gaude and Janarthnam, 2015). Numerical abundance study by Rao *et al.*, 2021 at Rivona near the foothills of Western Ghats - Goa reported Malabar tree nymph under “Very Rare” status with 0.026 relative abundance.

Host Plants

The larva of *Idea malabarica* mainly feed on the species of Apocynaceae *Aganosma cymosa* (Wynter-Blyth 1957; Kunte, 2000) and according to few recent reports the medicinal

plants *Parsonsia spiralis* and *P. alboflavescens* (Apocynaceae) are also considered as larval host plants of *I. malabarica* (Susanth 2005; Aishwarya and Revanna, 2018).

Conclusion

By declaring, Goa’s “State butterfly” status to the Malabar tree nymph butterfly, endemic to Western Ghats is recognized as biodiversity indicator of this region. Butterflies are important component of rich biodiversity, have been on planet around for at least 50 million years and these have special aesthetic, educational, scientific, ecosystem and economic value. Butterflies are widely used as model organism by ecologists to study the impact of habitat loss and fragmentation, and climate change because these are indicators of a healthy environment and healthy ecosystems. In the present global scenario, loss of wildlife habitat in a large scale and unpredictable shifting of climate and weather patterns in response to pollution of the atmosphere are the major issues related to decreasing trend of wildlife including butterflies. In this case, a broad and scientific approach for butterfly conservation is necessary. Butterfly distribution and population status assessment, diagnosing the driving factors causing decline trend of the species, identifying effective species recovery solutions and application of long-term sustainable solutions are the stepping factors for butterfly conservation. By gaining Goa’s “State butterfly” status, the Malabar tree nymph butterfly endemic to Western Ghats is recognized as biodiversity indicator of this region.



Figure 1. Egg

(Source: Vinayraj, Wikimedia)



Figure 2. Larva

(Source: Ashok Senagupta, Wikimedia)



Figure 3. Pupa

(Source: Ashok Senagupta, Wikimedia)



Figure 4. Adult butterfly

(Source: Ajith Unnikrishnan, Wikimedia)

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Efficacy of entomopathogenic fungi against chrysanthemum aphid, *Macrosiphoniella sanborni* (Gillette)

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Abstract

An experiment was carried out to evaluate different entomopathogenic fungi against chrysanthemum aphid, *Macrosiphoniella sanborni* (Gillette) under laboratory conditions at Biocontrol Laboratory, N.M. College of Agriculture, Navsari Agricultural University, Navsari during 2019-20 and 2021-21. The pooled mean corrected percent mortality revealed that among all the entomopathogenic fungi tested at 4g/L, *Verticillium lecanii* was found superior over the rest of the treatments by recording highest percent mortality (53.66%) of aphids, *M. sanborni* and it was followed by *Metarhizium anisopliae* and *Beauveria bassiana* with 36.84 and 32.61 percent mortality, respectively. The lowest per cent mortality of aphid was recorded by *Nomuraea rileyi* (25.55%). The chemical check, Dimethoate 30 EC at 1ml/L recorded highest percent mortality of aphids (77.82 %) among all the treatments.

Keywords: Chrysanthemum aphid, *Macrosiphoniella sanborni*, Entomopathogenic fungi

Introduction

Chrysanthemums (*Dendranthema grandiflora* Tzvelev), sometimes called mums or chrysanthus, are flowering plants of the genus *Chrysanthemum* of the family Asteraceae. Countless horticultural varieties and cultivars of this flower exist. Presently, 2000 varieties are grown around the world and in India, about 1000 varieties are grown (Datta and Bhattacharjee, 2001). As many as seven insect pests are reported damaging this crop right from germination to harvesting of the crop. The important insect pests attacking Chrysanthemum are aphid, *Macrosiphoniella*

sanborni (Gillette); thrips, *Haplothrips ramakrishnae* Krishna; Chrysanthemum caterpillar, *Diacrisia oblique* Walker; grub, *Holotrichia* spp. and leafminer, *Pytomyza syngenesiae* (Hardy). The Chrysanthemum aphid, *M. sanborni* is a widespread pest on cultivated Chrysanthemum throughout the world. It is a holocyclic species of East Asian origin (Heie, 1995). It feeds mainly on young leaves and developing flower buds and could become very abundant on them. In case of high infestation, the aphid causes significant damage which results in deformation and disturbance of flower development and it also acts as a vector to vein mottle and virus B

(Blackman and Eastop, 1984; Chan *et al.*, 1991). All these factors together become hereby significant and responsible for economic damage to the Chrysanthemum by decreasing the beauty and value of cut flowers (Zahedi, 1999). Pal and Sarkar (2009) reported *M. sanborni* as the major sucking pest of Chrysanthemum in hilly regions of West Bengal area by conducting field surveys. Considering the importance of Chrysanthemum aphid, the eco-friendly management and the growing demand for Chrysanthemum in South Gujarat, the present study was taken up to study the efficacy of various entomopathogenic fungi against the Chrysanthemum aphid under laboratory conditions.

Materials and methods

The laboratory experiment on efficacy of various entomopathogenic fungi against Chrysanthemum aphid, *M. sanborni* were carried out in Biocontrol Laboratory, N. M. College of Agriculture, Navsari Agricultural University, Navsari (20.925°N, 72.908°E) following Completely Randomized Design with six treatments and four repetitions during the year 2019-20 and 2020-21. The treatments included four entomopathogenic fungi viz., *Nomuraea rileyi* (1×10^8 cfu/g), *Beauveria bassiana* (1×10^8 cfu/g), *Lecanicillium* (*Verticillium*) *lecanii* (1×10^8 cfu/g) and *Metarhizium anisopliae* (1×10^8 cfu/g) along with chemical check Dimethoate 30 EC (1ml/L) and untreated control. The treatments were imposed by dipping the healthy

Chrysanthemum leaves for five minutes in conidial solution of each treatment. The treated leaves were allowed to dry under ceiling fan. Twentyfive aphids were released on each treated leaf to study the mortality. After 15 minutes of exposure, the aphids on treated chrysanthemum leaves were transferred to fresh leaves. The mortality of chrysanthemum aphid were recorded at 3rd, 5th, 7th, 10th and 14th days after treatment. The turgidity of leaves was maintained by using standard technique (?). At an interval of 24 hrs., each leaf (*i.e.*, untreated) was changed with new fresh leaf in same Petri dish. The data obtained on cumulative dead chrysanthemum aphid counts were summed up and utilized for calculation of percent corrected mortality. The data of cumulative dead aphids were converted into arcsine transformation and analyzed statistically by using completely randomized design. The percent corrected mortality was worked out through utilizing the formula suggested by Henderson and Tilton (1955).

Results and discussion

The results on the efficacy of entomopathogenic fungi against chrysanthemum aphid, *M. sanborni* during the years 2019-20 and 2020-21 were presented in Table 1. The mean corrected percent mortality of 3, 5, 7, 10, 14 DAT (Days after Treatment) revealed that among all the entomopathogenic fungi, the highest mortality was recorded for *V. lecanii* with 52.93 per cent during 2019-20 and 54.40 per cent mortality during 2020-21. The next most effective treatment was *M.*

anisopliae with 35.51 per cent during 2019-20 and 38.17 percent mortality during 2020-21 which was on par with *B. bassiana* with 31.5 per cent mortality during 2019-20 and 33.72 percent during 2020-21. The lowest per cent mortality of aphid was recorded for *N. rileyi* with 25.11 and 26.00 per cent mortality during 2019-20 and 2020-21, respectively. However, chemical check Dimethoate 30 EC recorded 77.65 (2019-20) and 77.99 per cent mortality of aphid, *M. sanborni*.

The overall pooled data on corrected percent mortality of aphid, *M. sanborni* are presented in Table 1. The mean data of 3, 5, 7, 10 and 14 DAT revealed that among all the entomopathogenic fungi, the treatment of *V. lecanii* at proved to be the most effective with highest per cent mortality (53.66%), the next effective treatment was *M. anisopliae* at (36.84%) which was at par with *B. bassiana* at *i.e.* 32.61 per cent mortality. The lowest per cent mortality of aphid was recorded in the treatment with *N. rileyi* (25.55%). Among all the treatments, chemical check Dimethoate 30 EC recorded highest per cent mortality of aphid, *M. sanborni* (77.82%). The results of present study are in agreement with those

obtained by Saranya *et al.* (2010) who reported 100 per cent mortality of cowpea aphid, *Aphis craccivora* by treating with *V. lecanii* and followed by *B. bassiana*, and *M. anisopliae*. Further, Husnain *et al.* (2014) reported *V. lecanii*, *M. anisopliae* and *Paecilomyces lilacinus* as effective against aphids in Pakistan. *Verticillium lecanii* was also reported effective against cotton aphid, while the study of Palthiya and Nakat (2017) indicated that combination of entomopathogenic fungi like *V. lecanii* 1.15% WP + *M. anisopliae* 1.15% WP was very effective against okra aphid. In a study Janu *et al.* (2018) also reported *V. lecanii* and *B. bassiana* effective against mustard aphid, *Lipaphis erysimi*. Gore *et al.* (2021) also concluded that *V. lecanii* was very effective against cotton aphid followed by *M. anisopliae* and *B. bassiana*. All these earlier studies thus, closely support the present findings.

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Table 1: Efficacy of entomopathogenic fungi against Chrysanthemum aphid, *M. sanborni* under laboratory condition

Treatments	Con.	Corrected per cent mortality (Yr. 2019-2020)						Corrected per cent mortality (Yr. 2020-2021)						Corrected per cent mortality (Pooled)					
		3 DAT	5 DAT	7 DAT	10DAT	14DAT	Mean	3 DAT	5 DAT	7 DAT	10DAT	14DAT	Mean	3 DAT	5 DAT	7 DAT	10DAT	14DAT	Mean
<i>B. bassiana</i>	4g/l	20.49 (12.25)	22.9 (15.18)	29.97 (24.97)	42.98 (46.48)	49.99 (58.63)	34.14 (31.5)	21.42 (13.38)	23.5 (15.94)	31.1 (26.68)	43.97 (48.21)	50.37 (59.29)	35.49 (33.72)	20.95 (12.81)	23.2 (15.56)	30.53 (25.83)	43.48 (47.35)	50.18 (58.96)	34.5 (32.61)
<i>M. anisopliae</i>	4g/l	23.84 (16.33)	26.21 (19.54)	32.9 (29.52)	45.65 (51.14)	51.37 (61.01)	36.58 (35.51)	24.58 (17.33)	26.69 (20.2)	33.91 (31.13)	46.62 (52.82)	53.29 (64.23)	38.16 (38.17)	24.20 (16.83)	26.45 (19.87)	33.41 (30.33)	46.14 (51.98)	52.33 (62.62)	37.06 (36.84)
<i>V. lecanii</i>	4g/l	27.55 (21.42)	36.1 (34.72)	45.62 (51.08)	58.89 (73.21)	66.76 (84.23)	46.68 (52.93)	28.96 (23.46)	35.04 (32.97)	47.56 (54.45)	59.09 (73.51)	65.5 (82.74)	47.53 (54.4)	28.25 (22.43)	35.57 (33.85)	46.59 (52.77)	58.99 (73.36)	66.13 (83.48)	46.77 (53.66)
<i>N. rileyi</i>	4g/l	12.88 (5.08)	21 (12.96)	25.93 (19.23)	39.58 (40.64)	43.63 (47.62)	30.06 (25.11)	14.1 (6.08)	20.78 (12.73)	27.32 (21.1)	39.29 (40.14)	42.54 (45.71)	30.65 (26)	13.49 (5.58)	20.89 (12.84)	26.63 (20.17)	39.44 (40.39)	43.09 (46.67)	30.08 (25.55)
Dimethoate 30EC	1ml/L	47.92 (55.08)	53.84 (65.18)	59.26 (73.84)	76.2 (94.16)	90 (100)	61.79 (77.65)	48.76 (56.54)	54.3 (65.94)	58.21 (72.23)	79.35 (95.34)	90 (100)	62.03 (77.99)	48.34 (55.81)	54.07 (65.56)	58.73 (73.03)	77.77 (94.75)	90 (100)	61.88 (77.82)
Control	-	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)	0.00 (0.00)
SE.m ±																			
Treatment (T)		0.56	0.79	0.85	1.16	1.11	0.39	0.78	0.76	0.63	1.93	0.92	0.45	0.47	0.55	0.53	1.13	0.72	0.25
(Y×T)		-	-	-	-	-	-	-	-	-	-	-	-	0.67	0.78	0.75	1.59	1.02	0.36
C.D. 5%																			
Treatment (T)		1.67	2.34	2.51	3.46	3.29	1.15	2.32	2.27	1.87	5.74	2.74	1.32	1.38	1.58	1.51	3.23	2.07	0.73
(Y×T)		-	-	-	-	-	-	-	-	-	-	-	-	NS	NS	NS	NS	NS	NS
C.V. (%)		5.10	5.92	5.24	5.31	4.40	2.23	6.80	5.72	3.81	8.64	3.67	2.50	6.04	5.82	4.57	7.20	4.05	2.04

Note: *Figures in parentheses are original values while those outside are arcsine transformed values, DAT: Days After Treatment

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Record of *Isotima* sp. (Hymenoptera: Ichneumonidae) from pink borer, *Sesamia inferens* (Walker)**Arun Baitha, Ankit K Mishra, Deepmala Kurre, P. K. Bareliya and Anuj Kumar***Division of Crop Protection**ICAR-Indian Institute of Sugarcane Research, R.B. Road, Dilkusha, P.O.**Lucknow-226 002, India****Corresponding author:** *arunbaitha@rediffmail.com*

The pink borer, *Sesamia inferens* (Walker) is a polyphagous pest and attacks many crops viz., rice, wheat, maize, sorghum, sugarcane, finger millet etc (Majumder, 2020). Due to the availability of one or the other host crops round the year, the pink borer is found throughout the year. However, during the peak winter (December to February) its activity is greatly reduced in sugarcane ecosystem but in a situation when paddy is grown near sugarcane field, it gets ready access to the adjoining paddy stubbles (after harvesting) during winter months. The larvae feed gregariously in the tillers of stubbles during winter months (Calora and Reyes, 1971). A large number of parasitoids were recorded on *S. inferens* i.e., *Trichospilus diatraeae*, *Cotesia* (*Apanteles*) *pallipes*, *Tropobracon scoenobii*, *Coccygomimus* (*Piimpla*) *laothoe*; *Devorgilla* sp., *Temelucha* sp., *Stenobracon nicevillae* (Rao *et al.*, 1967 and Shepard *et al.*, 1987).




Surveys were conducted in harvested field of paddy stubbles near sugarcane field at the research farm of ICAR-IISR, Lucknow. Different phases of development of the pink borer (larvae as well as pupae) and its parasitoids were collected in the tillers of

paddy stubbles during November 2020 to February 2021. Further, different instars of larvae were kept in glass jars (15 x 2.5 cm) with tillers of paddy (as food) for development at room temperature. During different phases of development of larvae/pupae, a light brown cocoon was collected from glass jar and kept for emergence of parasitoid at $26 \pm 2^{\circ}\text{C}$ and $75 \pm 5\%$ relative humidity in BOD. On the completion of the development, adult parasitoid found its way out of the cocoon through a circular aperture which it cut on one side of the cocoon and it was identified as *Isotima* sp. This is the first record of *Isotima* sp. reared from stubble-inhabiting larvae of *S. inferens* in rice ecosystem. The males have slender body and absence of the white band in the antennae. It however has an additional white band on the abdominal segment.

Isotima sp. is a solitary ecto prepupal parasitoid of sugarcane top borer in India. It attacks the prepupal stage and deposits an egg either on the prepupa or near its vicinity (Gupta, 1958 and Kalra and David, 1967). The colour and texture of the cocoon of *Isotima* sp. depends upon the season through which it has to pass in the pupal stage (Ahmad and

Mathur, 1945). A long and rigorous winter imposes on the grub an equally long period of hibernation and calls for a compact and tough cocoon which one finds in the field during winter months. The silken materials used by the grub in spinning this cocoon is - light brown in rice ecosystem but dark brown in sugarcane ecosystem. It is interesting to note that the size of the cocoon varies not only with

the environmental conditions, food etc, which affect the size of the grub, but also according to the sex, so much that the sex of the adult that would emerge from a cocoon can easily be predetermined by taking measurements of the cocoon. Under ordinary field conditions, the parasite goes into hibernation in its larval stage along with the host during winter months in sugarcane ecosystem (Baitha *et al.*, 2017).

	
<p>Cocoon of <i>Isotima</i> sp on paddy tiller</p>	<p>Circular hole upper side of cocoon</p>
	
<p>Male of <i>Isotima</i> sp.</p>	

The parasitoid *Isotima* sp. was first recorded and described by Rohwer (1918) from Java (Indonesia) on *Scirpophaga nivella* intact Snellen, the white moth borer of sugarcane and named as *Eripternimorpha javensis*. Indian specimens were erroneously identified as *Melcha ornatipennis* Cameron (1907) and all the work was carried out under this name till the misidentification was pointed by Townes *et al.* (1961) supported by Gupta (1961). It has been established that the ichneumonid parasite of top borer from Java as *Eripternimorpha javensis*, is a species belonging to the genus *Isotima* rather than *Eripternimorpha*. Various workers (Mathur, 1942; Ahmad and Mathur, 1945 and Box, 1953) mentioned that *S. nivella* as its only host but Khanna (1953, 1954 and 1955) and Varma and Maninder (1981) reported as larval parasite of top shoot borer (*S. nivella*), stem borer (*Chilo infuscatellus*) and Gurdaspur borer, *Acigona Steniella* whereas Bennett (1965) opined that this parasite doesn't normally attack stem borers. It is also reported on yellow rice stem borer, *Scirpophaga incertulas* and Gyrinid beetle, *Dineutus unidentatus* (Vazirani, 1952; Gupta, 1964 and Beg and Khan, 1982).

Isotima sp. is a pre pupal parasite, and many times more effective than that of an egg parasitoid. Further studies need to be carried out to develop protocols for its mass rearing in the laboratory on an alternate host, release technique, taxonomical study and survival in the field.

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Outbreak of common banded awl butterfly, *Hasora chromus* (Cramer) (Hesperiidae: Lepidoptera) in Bangalore

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Introduction

Butterflies, the flying jewels, serve human society economically and environmentally because of their undeniable beauty and ability to pollinate, a fundamental ecological process in natural sustainability around the world. Butterflies are important pollinators and are an important part of the life support system (Suryanarayana *et al.*, 2015). Besides acting as pollinators, few butterfly species have economic importance as pests of several crops. Lepidopteran families like Hesperiidae, Papilionidae and Nymphalidae have major role as defoliators of several agricultural and forest crops, causing intensive damage. *Millettia* (= *Pongamia*) *pinnata*, often known as karanja, is a leguminous oil-producing multifunctional tree that can withstand a variety of environments. The tree is famed for its insecticidal properties, but it is also attacked by a variety of insect pests, reducing its overall vigor and oil yielding ability.

The common banded awl, *Hasora chromus*, is one of the principal defoliator pests, causing severe leaf losses and in extreme situations, total defoliation. This pest has

recently been noticed fluttering in large numbers in the University of Agricultural Sciences, Gandhi Krishi Vigyana Kendra (UAS-GKV) Campus (13°04'30.4"N, 77°34'46.4"E) (Figs. 1&2). During the pest occurrence, entire tree looked dried without leaves. All the karanja trees in the campus were almost infested heavily by *H. chromus* with 90 - 100% defoliation. Occurrence of the skipper was characterized by presence of numerous larvae on leaves, pupae in folded leaves and activity of adults were also noticed in and around the karanja planted area. Outbreak of *H. chromus* was also recorded earlier in HSR layout and NBAIR Yelahanka campus (Anonymous, 2016b).

Occurrence, Habit and Habitat

The Indian subcontinent, Southeast Asia (including the Malay Peninsula and the Indonesian archipelago), South China, Okinawa, Japan, Papua New Guinea, and Australia are all home to the common banded awl. It extends up to 7,000 feet in the plains and slopes (2,100 m). It can be found in both the forest and open land, in both light and heavy rainfall locations. The common banded awl, or Coeliadinae subfamily of skippers, is

the most abundant in India. It has a whirring, fast flight that may be heard at close range. It is less sun-sensitive than other awls, and it is frequently seen soaring among shrubs in full sunlight. It can be spotted early in the day visiting flowers and basking on leaves, frequently with its wings slightly separated. It closes its wings and rests.

Life cycle

Eggs: It lays single eggs both above and below on young shoots or fresh leaves. When the egg is laid, it is pinkish white, dome-shaped with a flattened top and minute longitudinal ridges. These have delicate

transverse striations and are bead patterned. As the egg matures, it turns a dirty white color.



Figure 1: Completely defoliated Pongamia/Karanja tree at UAS GKVK campus due to *Hasora chromus* (Image by: Suresh R. Jambagi)



Figure 2: Severity of banded awl skipper, *Hasora chromus* on *Millettia pinnata* (Image by: Suresh R. Jambagi)

Larva: The caterpillar is cylindrical in shape, with a constricted second segment that resembles a neck with a black collar. The head is lobed, spherical, and the colour is yellowish red. The caterpillar is yellowish black in colour with brown sides ranging from light to dark brown. The markings are really varied. The caterpillar's underside is yellow-tinged greenish white. The green, coupled with the

dark patterns, can sometimes cover the entire body. When the caterpillar is first hatched, it eats the eggshell in bits and scurries off to a leaf, where it quickly constructs a cell for itself. It is active when young but becomes inactive as it matures. Only when the light is very low and at night does the caterpillar venture out to feed.



Figure 3: Pupation of *H. chromus* on different vegetation adjacent to karanja trees (Image by: Suresh R. Jambagi)

Pupa: The pupa is robust, pale brown, and has a conspicuous projection on the head between bulbous eyes, as well as a white belly. It will pupate in the folded fresh leaves and show shelter building behavior. During the intense infestation, it is observed that the larva can pupate in the vegetation (weed plants) present in around the main host (Fig. 3).

Adult: The wings of adult butterfly are entirely covered with scales and are brownish black in color. Male butterfly wings are unmarked, whereas female butterfly wings feature two spots on both sides of the fore wings and a horizontal white band on the lower side of the hind wings in both sexes. Adult butterflies are quite active. The antenna is long and progressively grows larger towards the tip, having a hook-like projection on the terminal segment.

Host range: Larva of *H. chromus* recorded on many host plants but its intense defoliation is noticed in castor, Indian beech (karanja/ pongamia), orange climber, *Derris scandens* and *Heynea trijuga*.

Management

Even though the trees were heavily damaged/defoliated by skipper, soon they rejuvenated after receiving early monsoon showers during May and larval population was also reduced drastically. Meanwhile, the activity of many insectivorous birds feeding on caterpillars was noticed. As per the earlier report, *Crinibracon chromusae* Gupta and van Achterberg (Hymenoptera: Braconidae) act as a pupal parasitoid of *H. chromus* (Anonymous, 2016a; Anonymous, 2017; Rani *et al.*, 2020). Hence, there is no need to use any insecticides.

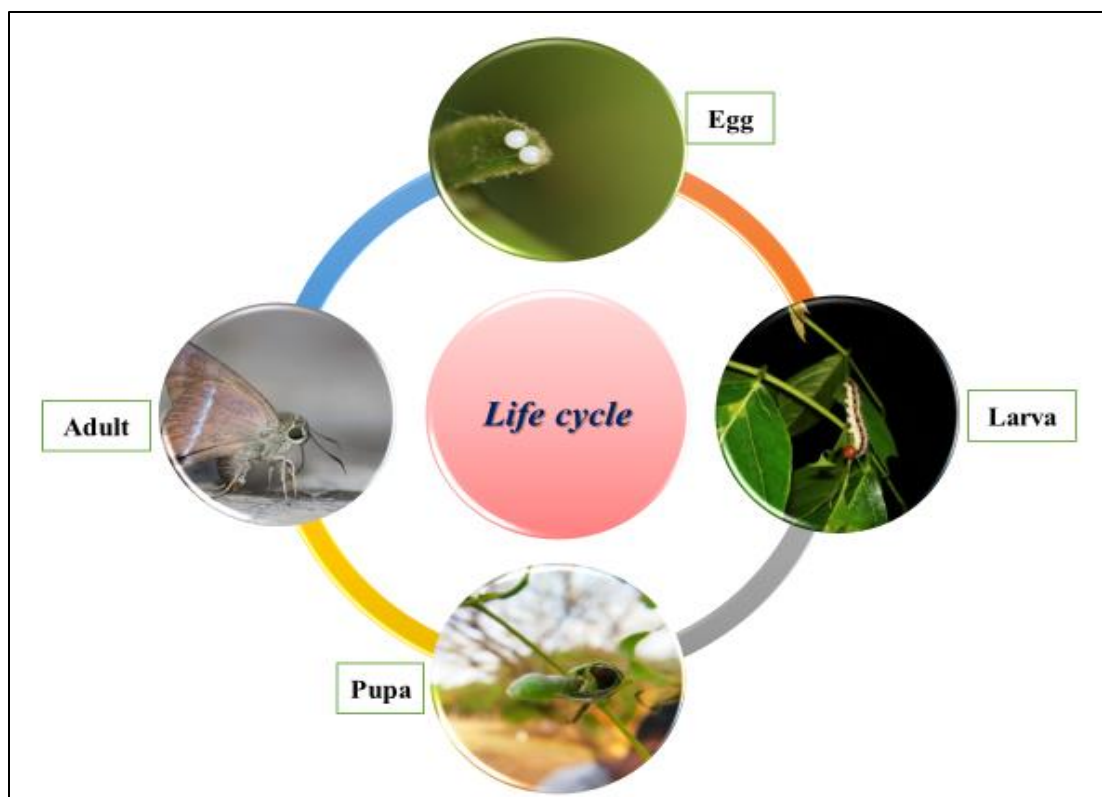


Figure 4: Different life stages of common banded awl skipper, *Hasora chromus* (Image by: Suresh R. Jambagi)

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Efficacy of new molecules of insecticides against aphid, *Aphis gossypii* (Glover) in summer sesame**Mohanlal Pensiya, G. M. Parmar*, Asha C. Detroja and R. M. Vikani***Department Entomology, College of Agriculture, Junagadh Agriculture University, Junagadh, Gujarat-362001***Corresponding author – gmparmar@jau.in***Abstract**

Experiment was undertaken during 2021 at Instructional Farm, Department of Agronomy, College of Agriculture, Junagadh Agricultural University, Junagadh to determine the efficacy of new molecules of insecticides against aphid in summer sesame. The results showed that the seed treatment with (fipronil 40% + imidaclopride 40% WG) @ 5 g/kg + FS of afidopyropen 50 DC @ 2 ml/l and (fipronil 40% + imidaclopride 40% WG) @ 5 g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l were found highly effective against aphid. The highest yield of 692 kg/ha was obtained from the treatment of (ST with fipronil 40% + imidaclopride 40% WG) @ 5g/kg + FS of afidopyropen @ 2 ml/l which was statistically at par with seed treatment of (chlothiodin 50 WDG) @ 7.5 g/kg + FS of afidopyropen 50 DC @ 2ml/l (680 kg/ha) and ST with (fipronil 40% + imidaclopride 40% WG) @ 5 g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l (672 kg/ha). The highest ICBR (1:8.68) was obtained from the treatment of (ST with fipronil 40% + imidaclopride 40% WG) @ 5g /kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l.

Keywords: sesame, aphid, sulfoxaflor, fipronil, afidopyropen, imidacloprid, efficacy

Introduction

Sesame is the ancient oilseed crop of India, is grown from time immemorial. Its seeds contain 52-57 per cent oil and 25 per cent protein (Smith *et al.*, 2000). Its cultivation gained impetus because of high quality edible oil, rich source of carbohydrate, protein, calcium and phosphorus (Prasad *et al.*, 2002). So, known as queen of oil seeds. It is used in confectioneries, cookies, cake, margarine, bread making etc. Sesame is rich in natural antioxidants or lignin's, which are both oil and water soluble provide very long shelf life and

stable characteristics of sesame seed and oil (Ermia *et al.*, 2009).

Sesame is attacked by different species of insect pests but sucking insects have great economic importance to sesame plants. Aphid, *Aphis gossypii* (Glover) is serious pest which suck the cell sap from leaves, flowers and capsules. Due to this downward curling of leaf margins, reddening of leaf margins, stunted growth of the plants, sickly appearance of the crop and subnormal growth of the leaf tissue occur (El-Gindy, 2002). The use of insecticides

has undoubtedly resulted in the maximum production of food grain for the world food supply, but the proliferation of insecticides and their unilateral utilization have posed many problems such as development of resistance in insect pests to insecticides, resurgence of insect pests, outbreak of secondary insect pests, insecticidal residues etc. Frequent use of single pesticide will not provide effective management of these pests. Therefore, in sesame crop, it is a prime need to find out such pesticides which was effectively control the various sucking pest attacking this crop. Presently, various new molecules with different mode of action are available, that necessitate evaluation against the aphid in summer sesame.

Materials and Methods

In order to study the efficacy of different molecules of insecticides against

aphid, the experiment was conducted in summer 2021 at the Instructional Farm, Department of Agronomy, College of Agriculture, Junagadh Agricultural University, Junagadh. Sesame variety G.Til-3 was sown at a spacing of 30 cm x 10 cm. All the recommended agronomical practices were followed. Seed treatment was given at the time of sowing, whereas foliar spray was given at the time of substandard population of aphid. The data on population of aphid was recorded on three leaves (top, middle and bottom canopy of the plants) per plant by randomly selecting five plants from each plot and tagged. The pre-treatment observation was recorded at one day before 1st spray and post treatment observations were recorded at 3, 7 and 10 days after each spray. Statistical analysis was carried out using ANOVA technique given by Panse and Sukhatme (1985).

Treatment details

No.	Treatment*	Dose
T ₁	ST with clothianidin 50% WDG	7.5 g/kg seed
T ₂	ST with fipronil 40% + imidacloprid 40% WG	5 g/kg seed
T ₃	T ₁ +FS of afidopyrofen 50 DC	7.5 g/kg seed+2 ml/l
T ₄	T ₁ +FS of flupyradifurone 200 SL	7.5 g/kg seed+1.5 ml/l
T ₅	T ₁ +FS of sulfoxaflor 24% SC	7.5 g/kg seed+1.5 ml/l
T ₆	T ₂ +FS of afidopyrofen 50 DC	5 g/kg seed+ 2 ml/l
T ₇	T ₂ +FS of flupyradifurone 200 SL	5 g/kg seed+1.5 ml/l
T ₈	T ₂ +FS of sulfoxaflor 24% SC	5 g/kg seed+1.5 ml/l
T ₉	Control	

*ST: Seed Treatment; FS: Foliar Spray; WDG/WG: Water Dispersible Grannules; DC: Dispersible Concentrate; SC: Suspension Concentrate; SL: Soluable Concentrate.

Results and discussion

Results showed that all the treatments having treated seeds were found significantly superior over the control in reducing the incidence of aphid. The result based on mean aphid population are presented in Table 1 indicate the pre foliar spray count of aphid showed that the aphid population in plots having treated seed with fipronil 40% + imidaclopride 40% WG @ 5g /kg varied from (1.58 to 1.68 aphid /3 leaves) and seed treated with clothianidin 50 WDG @ 7.5 g/kg varied from (1.84 to 1.91 aphid /3 leaves). However, significantly higher population of aphid (2.38 aphid /three leaves/plant) was recorded in control plot.

The data on mean number of aphid population after three days of application of insecticides presented in Table 1 indicate that all the treatments were found significantly superior over untreated plot. Seed treatment with fipronil 40% + imidaclopride 40% WG @ 5g /kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l and seed treatment with clothianidin 50 WDG @ 7.5g /kg + FS of sulfoxaflor 24% SC @ 1.5 ml /l were found most effective (0.38 /3 leaves), which was at par with seed treatment of fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l and seed treatment of clothianidin 50 WDG + FS of flupyradifurone 200 SL @ 1.5 ml/l which gave (0.52 aphid /3 leaves). Seed treatments with fipronil 40% + imidaclopride 40% WG @ 5g /kg + FS of afidopyropen 50 DC @ 2

ml/l (1.14/3 leaves) and ST with clothianidin 50 WDG @ 7.5g /kg + FS of afidopyropen 50 DC @ 2 ml /l (1.33/3 leaves) were found medium in their effectiveness.

All the treatments were found significantly superior over untreated plot after seven days of application of insecticides. Seed treatment with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of afidopyropen 50 DC @ 2 ml/l was found most effective which gave 0.26 aphid per three leaves which was at par with seed treatment of clothianidin 50 WDG @ 7.5 g/kg + FS of afidopyropen 50 DC @ 2 ml/l (0.31/3 leaves). While, treatments comprising of ST with fipronil 40% + imidaclopride 40% WG @ 5g /kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l (0.79/3 leaves), ST with clothianidin 50 WDG @ 7.5 g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l (0.96/3 leaves), ST with clothianidin 50 WDG @ 7.5 g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l (1.14/3 leaves) and ST with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l (1.16/3 leaves) were found next best in their effectiveness.

Seed treatment with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of afidopyropen 50 DC @ 2 ml/l was found most effective (0.99 per three leaves) and it was statistically at par with seed treatment of clothianidin 50 WDG @ 7.5 g/kg + FS of afidopyropen 50 DC @ 2 ml/l (1.06/3 leaves) after ten days of application of insecticides.

The seed treatment with clothianidin 50 WDG @ 7.5 g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l (1.27 /3 leaves), ST with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l (1.46/3 leaves), ST with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l and ST with clothianidin 50 WDG @ 5g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l (1.52 /3 leaves) and seed treatment with fipronil 40% + imidaclopride 40% WG @ 5 g/kg (1.66 /3 leaves) were found next effective in order of efficacy. More or less similar trend of aphid population recorded after second spray of insecticides.

Data presented in Table 3 indicate that a difference in grain yield was significant. The highest yield of 692 kg/ha was obtained from the treatment of ST with fipronil 40% + imidaclopride 40% WG @ 5g /kg + FS of afidopyropen 50 DC @ 2 ml/l which was found statistically at par with the seed treatment of clothianidin 50 WDG @ 7.5g/kg + FS of afidopyropen 50 DC @ 2 ml/l (680 kg/ha) and ST with fipronil 40 % + imidaclopride 40% WG @ 5g/kg + FS of sulfoxaflor 24% S.C @ 1.5 ml/l (672 kg/ha). The highest (1:8.66) Incremental Cost Benefit Ratio was obtained from the seed treatment of fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l it is followed by seed treatment of clothianidin 50 WDG @ 7.5 g/kg + FS of sulfoxaflor 24 % SC @ 1.5 ml/l

(1:8.33) and ST with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of flupyradifurone 200 SL @ 1.5 ml/l (1:7.44).

According to Ambarish *et al.* (2017), the lowest number of aphids (1.71) was recorded in the application of sulfoxaflor 30 % @ 108 g. *a.i* /ha. Prasad (2017) recorded that flupyradifurone 200 SL at lower dose of 150g *a.i*/ha was found superior in efficacy against cotton aphid. Garg *et al.* (2018) reported that flupyradifurone 200 SL @ 125, 150, 175 g *a.i*/ha was found effective for managing aphid population in okra. Singh *et al.* (2020) reported the lowest population of aphid 1.33/five leaves/plant after 10 days of spray in flupyradifurone 200 SL @ 2.5 ml/l in okra. Susheel kumar *et al.* (2020) reported that afidopyropen 50 DC @ 2 ml/l showed 82.91, 77.48, and 76.59% aphid reduction in the pest population. So, the results obtained from the present finding are closely fitted with the result reported by earlier worker.

Conclusion

Considering the efficacy, yield and economics of insecticides, seed treatment with fipronil 40% + imidaclopride 40% WG @ 5g/kg + FS of afidopyropen 50 DC @ 2 ml/l were found highly effective against aphid. The treatment comprising of ST with fipronil 40% + imidaclopride 40% WG @ 5 g/kg + FS of sulfoxaflor 24% SC @ 1.5 ml/l and ST with clothianidin 50 WDG @ 7.5 g/kg + FS of afidopyropen 50 DC @ 2ml/l were found moderately effective.

Table 1. Bio-efficacy of different insecticide against aphid after first spray

Sr. No.	Treatment	Dose (g/kg & ml/l)	Mean number of aphid /3 leaves			
			Before spray	3 DAS	7 DAS	10 DAS
T ₁	ST with clothianidin 50WDG	7.5g	1.36 (1.84)	1.31 (1.72)	1.36 (1.86)	1.38 (1.91)
T ₂	ST with Fipronil 40% + Imidaclopride 40% WG	5g	1.26 (1.58)	1.26 (1.59)	1.31 (1.72)	1.29 (1.66)
T ₃	T ₁ +FS of afidopyropen 50DC	7.5g + 2ml	1.38 (1.91)	1.15 (1.33)	0.56 (0.31)	1.03 (1.06)
T ₄	T ₁ +FS of flupyradifurone 200SL	7.5g +1.5ml	1.36 (1.85)	0.72 (0.52)	1.07 (1.14)	1.23 (1.52)
T ₅	T ₁ +FS of sulfoxaflor 24% SC	7.5g +1.5ml	1.36 (1.84)	0.62 (0.38)	0.93 (0.96)	1.13 (1.27)
T ₆	T ₂ +FS of afidopyropen50DC	5g + 2ml	1.30 (1.68)	1.07 (1.14)	0.51 (0.26)	1.00 (0.99)
T ₇	T ₂ +FS of flupyradifurone 200 SL	5g +1.5ml	1.26 (1.58)	0.72 (0.52)	1.06 (1.12)	1.23 (1.52)
T ₈	T ₂ +FS of sulfoxaflor 24% SC	5g +1.5ml	1.28 (1.65)	0.62 (0.38)	0.89 (0.79)	1.21 (1.46)
T ₉	Control		1.54 (2.38)	1.59 (2.53)	1.55 (2.40)	1.57 (2.45)
	S. Em.±	T	0.08	0.06	0.06	0.06
		P	-	-	-	-
		T×P	-	-	-	-
	C. D. at 5 %	T	0.23	0.17	0.18	0.18
		P	-	-	-	-
		T×P	-	-	-	-
	C. V.%		10.21	9.57	9.92	8.62

Figures within parentheses indicate retransform values, while outside are square root transformed value

Table 2. Bio-efficacy of different insecticide against aphid after second spray

Sr. No.	Treatment	Dose (g/kg & ml/l)	Mean number of aphid /3 leaves			
			Before spray	3 DAS	7 DAS	10 DAS
T ₁	ST with clothianidin 50WDG	7.5g	1.46 (2.12)	1.46 (2.12)	1.48 (2.19)	1.52 (2.32)
T ₂	ST with fipronil 40% + Imidaclopride 40% WG	5g	1.38 (1.91)	1.43 (2.05)	1.46 (2.12)	1.46 (2.12)
T ₃	T ₁ +FS of afidopyropen 50DC	7.5g + 2ml	1.43 (2.05)	1.07 (1.14)	0.62 (0.38)	1.00 (0.99)
T ₄	T ₁ +FS of flupyradifurone 200SL	7.5g +1.5ml	1.41 (1.99)	0.80 (0.65)	1.18 (1.39)	1.28 (1.65)
T ₅	T ₁ +FS of sulfoxaflor 24% SC	7.5g +1.5ml	1.38 (1.90)	0.76 (0.58)	1.09 (1.20)	1.26 (1.59)
T ₆	T ₂ +FS of afidopyropen50DC	5g + 2ml	1.34 (1.79)	1.15 (1.33)	0.51 (0.26)	0.84 (0.71)
T ₇	T ₂ +FS of flupyradifurone 200 SL	5g +1.5ml	1.36 (1.84)	0.68 (0.46)	1.13 (1.27)	1.26 (1.59)
T ₈	T ₂ +FS of sulfoxaflor 24% SC	5g +1.5ml	1.34 (1.79)	0.62 (0.38)	1.09 (1.20)	1.23 (1.51)
T ₉	Control		1.48 (2.19)	1.50 (2.25)	1.57 (2.46)	1.63 (2.65)
	S. Em.±	T	0.07	0.07	0.06	0.07
		P	-	-	-	-
		T×P	-	-	-	-
	C. D. at 5 %	T	NS	0.20	0.17	0.22
		P	-	-	-	-
		T×P	-	-	-	-
	C. V.%		8.47	10.86	8.79	9.84

Figures within parentheses indicate retransform values, while outside are square root transformed value

Table 3. Economics of different treatment applied for the control of aphid of summer sesame

Sr	Treatments	Total quantity for 2 sprays (g or ml/ha)	Price of Insecticide (Rs./lit. or kg)	Cost of Insecticide (Rs./ha.)	Total cost of Treatment (Rs./ha)	Yield/ha	Gross Realization (Rs./ha)	Net realization (Rs./ha)	ICBR
T ₁	ST with clothianidin 50WDG	30g	13000	390	890	399	45087	5424	1:6.09
T ₂	ST with fipronil 40% + Imidacloprid 40%WG	20g	15600	312	820	404	45652	5989	1:7.37
T ₃	T ₁ +FS of afidopyropen 50DC	30+1.600	3197	390+5115	6505	680	76840	37177	1:5.71
T ₄	T ₁ +FS of flupyradifurone 200SL	30+1.200	4080	390+4896	6286	652	73676	34013	1:5.41
T ₅	T ₁ +FS of sulfoxaflor 24% SC	30+1.200	2390	390+2868	4258	665	75145	35482	1:8.33
T ₆	T ₂ +FS of afidopyropen 50DC	20+1.600	3197	312+5115	6427	692	78196	38533	1:6.00
T ₇	T ₂ +FS of flupyradifurone 200 SL	20+1.200	4080	312+4896	6208	658	74354	34691	1:5.59
T ₈	T ₂ +FS of sulfoxaflor 24% SC	20+1.200	2390	312+2868	4180	672	75936	36273	1:8.68
T ₉	Control	-	-	-	-	351	39663	-	-

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Commercial meliponiculture- beekeeping with stingless bee, *Tetragonula iridipennis* Smith (Hymenoptera: Apidae: Meliponini)

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Tetragonula iridipennis, belonging to the family Apidae under subfamily Apinae, is known as stingless bees as they do not have sting apparatus (Devanesan & Raakhee 1999). They are also called dammar bees as they construct numerous elliptical cells for storing pollen and honey by using a special material called “cerumen” consisting of wax and resin (Raakhee and Devanesan 2000).

Stingless bees are ancient eusocial bees which populated the earth 65 million years ago, quite earlier to *Apis* species and are limited to tropics and subtropics because of their inability to maintain the hive temperature. Honey bees and the stingless bees are honey producers coming under the order Hymenoptera, family Apidae and subfamily Meliponinae. They lack sting hence the name “stingless bee” and defend by biting with mandibles. Unlike honey bees stingless bees do not use pure wax for nest building and water for nest cooling. Brood production is similar to solitary bees and they follow mass provisioning to rear their larvae. Both groups make honey in perennial nests founded by a swarm of sterile workers and a queen and colonies occasionally produce male bees.

The ICAR All India Co-ordinated Research Project (AICRP) on Honey bees and Pollinators, Vellayani Centre of Kerala Agricultural University, pioneered the research on Meliponiculture in India (Devanesan & Raakhee 1999).

We happened to notice the presence of dammar bees in different districts of Kerala in basements of buildings and compound walls during Thai sac Brood virus (TSBV) disease survey (1992-93) . The photographs were shown to Dr. O.P. Dubey, the then Assistant Director General (PP), who advised us to conduct detailed studies about the bee species. The ICAR sanctioned a project entitled “Bio-ecology, domestication and management of stingless bees in Kerala” (9-7-1999 to 8-1-2003 for INR. 13 lakhs).

We succeeded in standardizing technologies for commercial meliponiculture by a team of scientists (Raghee M, Shailaja K.K, Premila K.S. and Nisha, M. M) under the leadership of Dr. Stephen Devanesan. We developed technologies for hiving, domestication and management of stingless bees with a view to enhance the number of

colonies and this centre has a prime role in popularization of Meliponiculture in the State. As instructed by ICAR, the final report was printed out as “STATUS PAPER ON STINGLESS BEE *TRIGONA IRIDIPENNIS* SMITH” authored by S. Devanesan, K.K. Shailaja and K.S. Premila. The same was released in the Annual Group Meeting held at Rajendra Agricultural University, Pusa, Samasthipur, Bihar in February 2009 by then Assistant Director General (PP), Dr.T.P. Rajendran. It was resolved in the group meeting that the research on stingless bees has to be promoted in all AICRP Centres and Vellayani centre as nodal centre for stingless bee research (Fig. 1).

Out of more than 500 species described worldwide, about 50 species occur in Asia. They have a wide distribution in India with reports of the following nine species. *Lepidotrigona arcifera* (Cockerell, 1929), *Lisotrigona cacciae* (Nurse, 1907), *Lisotrigona mohandasi* Jobiraj & Narendran, 2004, *Tetragonula ruficornis* (Smith, 1870), *Tetragonula bengalensis* (Cameron, 1897), *Tetragonula iridipennis* (Smith, 1854), *Tetragonula praeterita* (Walker, 1860), *Tetragonula aff. laeviceps* (Smith, 1857), *Tetragonula gressitti* (Sakagami, 1978). Asiatic stingless bee *Tetragonula* (*Trigona*) *iridipennis* Smith is the common stingless bees found in Kerala which was identified by Dr. David W Roubik, Smithsonian Tropical Research Institute, USA.

Their small size (2-16 mm) allows them to access varied flowers whose openings are too narrow for nectar and pollen from which the *Apis* bees cannot forage (Devanesan, *et al.*, 2002). They also collect nectar and pollen from low lying herbs and weeds having comparatively smaller flowers, which are not commonly visited by *Apis* spp. including medicinal plants (Premila, *et al.*, 2007). The stingless bees make less honey compared to *Apis* honey bees and are widely distributed in diverse tropical regions of India (Vijayakumar *et al.*, 2013). Although it produces far less honey the honey is unique having great medicinal value fetching high price. It is used in folk medicine, Ayurveda, Sidha, Unani etc for curing many diseases. Stingless bees are excellent pollinators of both rest/agricultural/horticultural flora and crop plants, and exhibit great flower constancy which increases the ecological importance of these bees for conservation of biodiversity and yield enhancement through crop pollination.

Meliponiculture is the art and science of rearing stingless bees on a commercial scale for honey production or pollination. It is reared as backyard beekeeping practice mainly for honey production. It is being popularized in the rural homesteads for additional income generation. In Kerala, the stingless bee colonies are reared in logs, earthen and clay pots, bamboo bits, wooden boxes and coconut shells. Creating awareness about the scientific meliponiculture will be helpful to the farmers for rearing them in homesteads so as to benefit

for the pollination of vegetables and other fruit crops available in and around promoting commercial meliponiculture (Fig. 2). Handling of stingless bees is easier, even children and women can rear stingless bees. Hence it is recommended that “each homestead to have one stingless bee colony”. The production of stingless honey does not match the demand. As it is recommended for the cure of different diseases including cancer, the demand has increased in the market due to its medicinal properties.

Natural domicile and nest architecture

Stingless bees prefer darkness and the natural domicile is the crevices in the basements of old buildings, compound walls, tree trunks, hollow blocks etc. The colonies found in natural conditions are called ‘feral’ colonies. They remain in the same location if there is no disturbance. The nest of stingless bee essentially consists of hive entrance, brood cells, storage pots of honey and pollen, resin dumps, pillars of wax, waste dumps etc.

Openings to the nest are through resin tubes of varying shapes, lengths and sizes, normally built of wax and mud, sometimes sticky in nature. The structure of hive entrance of *T. iridipennis* colonies in wooden boxes, bamboo bits and earthen pots varies in their shape and size (long or short tubular, round or cryptic, half oval in shape etc). Major part of entrance tube material is made up of resin. Some dust materials like soft barks, mud and cobweb like silky material is also found over

the tubes (Fig. 3). The texture is not brittle and the nest consist of five parts; an entrance, brood comb, involucrum, store pots and batumen. In the construction of brood comb, storage pot and involucrum they use cerumen (Raaghee and Devanesan, 2020) (Fig.4).

The arrangement of brood cluster is loose in *T. iridipennis*. They build multi-layered combs one over the other and each expanding concentrically and horizontally. Cell construction starts from the bottom of the pillars and proceeds in an upward direction. At a time, a batch of cells is constructed through successive and intermittent contributions of several workers. Brood cells are dark brown in colour in the early stages and as the pupae mature, the cerumen from walls of the brood cells are removed and became creamy in colour so that the cocoons get exposed.

Most of the removed cerumen is used again to build another brood cell, storage pot or other nest structures. The queen cells of *T. iridipennis* are larger (4.00 mm) than that of worker cells (0.22 mm). Queen cells could be seen in stingless bee colonies from November to March. They are intermixed with worker cells or seen at the periphery of the brood cluster. The food chambers are normally oval in shape and larger than the brood cells. The pollen pots are located near the entrance and seen intermixed whereas the honey pots are seen either near the entrance or far from the entrance. The honey pots are also seen intermixed with pollen pots.

Stingless bees also lead a colonial social life and each colony consists of a single queen few drones more and workers as in the case of *Apis* bees. Usually one queen is present in a colony and the queen lays egg and controls and co-ordinates all the activities in it. The queen is much larger than workers and drones, lacking corbiculae and wax glands. It has longer scape, shorter tongue, smaller mandibles, wings partially covering the swollen abdomen and less distinct glabrous streaks on mesonotum, abdomen is dark brown in colour with white stripes. Queen lays 80-120 eggs per day. The developmental period of a queen (egg to adult) is 65-70 days. The newly emerged queen is called 'gyne' or virgin queen. The new queen makes her nuptial flight, where she is mated with a single male. The virgin queen will mate only once. The sperm is stored in a special sac 'spermatheca' in her body for her entire life. The queen's ovaries initiate egg development, enlarging the abdomen, so that she cannot fly anymore. Egg laying commences within 10 days after mating. She can choose whether or not to fertilize each egg with the sperm. The queen and workers develop from the fertilized eggs, and drones/males develop from the unfertilized eggs. The life span of queen is up to five years, though her stamina for egg laying will be reduced over time. The life span of queen is reported to be up to five years (Raaghee 2000) (Fig.5).

The drones or male bees are developed from unfertilized eggs and are few in a colony and function is mating. Number of drones

increase in the colonies during the brood rearing season. The drones can be identified by straightly arranged ocelli, laterally arranged compound eyes, smallest scape, longest antennae, smallest mandibles, less distinct glabrous streaks on mesonotum, rudimentary corbicula, and wings projecting slightly beyond the blunt abdomen and genitalia (Fig. 6).

Worker bees are females developed from fertilized eggs. Most of the works in a colony are performed by the worker bees. The worker bees are smaller in size than the queen with small abdomen and entire body is black to blackish-brown. Males and workers are very similar and difficult to differentiate. The developmental period of a worker (egg to adult) is found to be 44.5 days. The young workers after emergence perform different jobs, preparation and repairs of the brood chamber, construction of the involucre and provisioning of cells. Some will become nurse bees, producing royal jelly to feed the larvae, young adults and queen. As age advances, perform duty as guards and foragers and with a life span of 80 days - (Fig. 7).

Honey and pollen are the food of stingless bees as *Apis* bees. Stingless bees do not go far distance for foraging as other honey bees. They visit plants within an area of 1 km around the hive. They forage from medicinal plants, agricultural and horticultural crops, spices, vegetables, cash crops, ornamental plants and number of weed plants around the

locality. They start foraging activity in the early morning.

Studies conducted to document the plants visited by the stingless bee revealed that there are 142 plant species, which provide either nectar or pollen or both to stingless bees in the State (AICRP, 2014). Out of the 142 plants, 70 provided nectar alone, 25 provided pollen only and 47 provided both nectar and pollen. These include medicinal plants, plantation crops, condiments and spices, vegetables, field crops, ornamental plants, wild plants and weeds.

The AICRP on Honey bees and Pollinators, Vellayani centre has standardized technologies for hiving, domestication, management such as division of colonies, honey extraction, hygienic honey processing and dearth season management. These techniques helped meliponiculturists to adopt commercial meliponiculture. The centre disseminated to the public by imparting trainers training in different districts of Kerala in which 2074 women and 8322 men were trained. It is estimated that more than 50,000 stingless bee colonies are domesticated in the state for pollination service, ensuring sustainable agriculture and the conservation of biological diversity resulting in food security.

Hiving of feral colonies to artificial hives (earthen pot/wooden hive) will damage the buildings/ walls and the public hesitate to demolish the structures. However, feral colonies can be easily transferred to hives. For

this, the mouth of an earthen pot is to be placed in front of the entrance of a feral colony and its rim is to be firmly attached to the wall using mud. Adequate support to the pot should be provided wherever required. An opening is to be given at the opposite side of the pot as bee entrance and surrounded with wax of stingless bee. The worker bees pass through the hole in the pot using it as new entrance. Gradually the bees realize that there is sufficient space inside the pot to store the brood and food reserves and hence they newly construct pollen and honey pots inside the earthen pot. Eventually they build the brood cells and start brood rearing inside it. The colony and pot should be left undisturbed for at least six months for the feral colony to settle inside the mud pot (Fig. 8). Later the pot with brood, pollen, honey, worker bees and a queen can be detached and shifted to a suitable site to establish as a new colony. Similarly wooden hives can also be used for hiving feral colonies (AICRP, 2011) (Fig. 8).

For transferring feral colonies from electric meter boxes or such structures, tubes/bent tubes are to be used. For this carefully remove the entrance tube of the bee colony. Then fix the detached entrance tube to one end of the plastic tube (1cm diameter). Fix the other end of the tube or bent tube in the original position of the hive entrance. The foragers will pass through the newly fitted plastic entrance hole. After a week time, a new wooden box hive / earthen pot with entrance hole at opposite side is to be attached to the

feral colony. It should be left undisturbed for at least six months.

Kerala Agricultural University (KAU) wooden hive

Among the different types of hives (wooden box, earthen pot, bamboo bit with volume of 1500 cc, 2250 cc, 3000 cc, and 3750 cc) evaluated for domestication, bamboo hive with 1500 cc capacity showed better brood development and maximum storage of pollen and honey followed by earthen pot and wooden hive. Considering the scarcity of bamboo bits, this centre has designed and developed a new 'KAU wooden hive' with two equal halves for the domestication of stingless bees. Volume of the box is 1960 cc and the internal measurements are: length - 35 cm, breadth - 7 cm and height - 4 cm, providing a hole in the centre of two halves on one side of the hive will act as the hive entrance. This hive is suitable for easy hiving, domestication, proper brood development, division of colonies and honey collection (Fig.9).

Seasonal Management of stingless bee colonies

The stingless bees have brood rearing season (October - December) followed by honey flow season (February - May) and a dearth season (June -September). Brood rearing season is best suited for the beginners to start meliponiculture. Acquiring knowledge about scientific meliponiculture through trainings and field practical will help for sustainable meliponiculture. Weekly

observations of hives are not required as in *Apis* bees. Open the colonies for honey extraction during April-May and division of colonies during October-November. Protective measures have to be taken to prevent the attack of ants, predatory spiders, mites etc.

Growth period management

During the natural growth period, worker bees construct pillars of wax, which serve as base to the brood cells, over which oval/elliptical brood cell cups are constructed with the help of several workers. Queen bee lays more eggs during growth period. The laying queen bee, inspect the cup cells and few workers fill the cups with food materials for the growth and development of larvae. Then the queen lays an egg over the food stuff in the cup. This is known as 'mass provisioning.' After oviposition worker will close the cup cell with the resin. Once the first set of brood cells are provisioned and oviposit newer cells are constructed above it (Fig.10). When, the population of worker bees increase, new queen cells (royal cells) are produced naturally resulting in swarming of stingless bees colonies. Keeping empty KAU hive in the apiaries with stingless bees, pasted with stingless bee wax in the hive entrance will help to trap the swarm. Management of colonies properly during the growth period will help to increase the number of colonies through division.

Division of the colonies

Healthy and disease-free, active colonies having sufficient mature brood, young brood, pollen, honey and queen cells are to be selected for division. Division of the colonies is to be done during evening hours. Rainy and cloudy days are unsuitable for opening the hive.

While dividing the colonies many worker bees may be lost. To minimize the loss of worker bees, they are to be safely removed from the colony before opening it, as done during honey extraction. After removing the worker bees, locate the queen/queen cells and transfer the queen along with half the quantity of mature brood, one-fourth quantity of young brood and half the volume of pollen and honey pots to a new hive. A good healthy queen cell is retained in the mother colony. If no queen cell is present in the selected colony a queen cell is to be grafted from another colony. The daughter colony with queen is to be shifted to a new site approximately 1000 ft. (300 m) away from the original site. Mother colony with queen cell is to be maintained in the original site.

The colonies maintained in the newly designed wooden hive with two equal halves is easier to divide. Open the hive and equalize the brood, pollen and honey storage in both the halves and provide an empty half box above the bottom piece of the hive. Similarly, provide the other empty half below the other piece to make a full hive. Care should be taken to

provide either a queen or queen cell in each of the newly divided colonies (Fig.11).

Avoid damage of the honey pots during division which may cause invasion of ants and fermentation of honey. Since the brood cells are very soft, care should be taken to minimize the damage of brood cells, otherwise the colony fails to establish. Presence of excess pollen and honey in the colony make them prone to pests and disease incidence and hence, it is desirable to remove excess pollen and honey periodically to maintain healthy colonies. While handling a hive, the direction of hive opening is to be marked so that the hive entrance can be retained in the original direction itself to avoid the confusion of foraging bees to find the entry.

Dearth season management

Feed can be prepared by using previous year's waste honey (collected during honey processing). A plastic tray used in the fridge for keeping the vegetables, with small ventilation (2-3 mm), is used as feeding tray, through which only the stingless bee workers can pass. Place a rectangular glass piece at the bottom of the tray. Put a cotton layer over the glass plate and drip with honey syrup using a wash bottle. 10-20 ml of syrup can be used for a single colony. Close the device with another piece of glass and keep the tray in a stand with proper ant protection devise (Fig.12).

Honey extraction

The honey from stingless bee colonies is extracted once in a year during April -May. The traditional method of honey extraction from stingless bee colonies is by squeezing the honey pots along with the pollen pots and brood cells resulting in unhygienic honey. The honey extracted by this method contained large quantity of pollen and other extraneous matters, which caused contamination. It also resulted in the destruction of bees, brood and ultimately the colony. Care should be taken to use sterile knife, spoons, vessels and cover head and mouth with mask and use glove.

While opening the colonies for honey extraction many worker bees will bite the beekeeper and may be lost. To minimize the loss of worker bees, before opening the colony for honey extraction the bees are to be removed and protected in empty water bottles. For this, the mouth of an empty water bottle (dried), with adequate small holes is to be placed at the hive entrance tube. Gently tap on the hive so that the worker bees' start coming out from the hive and will enter into the bottle. Two or more bottles can be used according to the strength of the colony. Cap the bottles, keep them aside and open the hive for extraction (Figs. 13, 14).

The pots alone are collected by using a sterile spoon from the nest and are kept in a clean tray. The tray is to be exposed to sunlight in a slanting position. The wax of the honey pots melt due to the heat of the sun and the

honey pouring out is to be filtered and bottled. There is no need for heating stingless bee honey like *Apis* honey. This method is relatively simple and yield clear honey without extraneous matters. After extraction, the bees in the bottles may be released.

Stingless bee honey

The stingless bee honey (SBH) is a natural product produced and marketed worldwide, collected from stingless bees which is only produced in tropical and subtropical regions and features a distinct sweetness, mixed with an acidic taste, fluid texture, high moisture content and minimal crystallization. Stingless bee honey is popular for its unique behavior of collecting nectar from multiple flowers and extra floral nectaries. The nectar then goes through a conversion in the bees' 'honey stomach' to transform into honey and deposited into cerumen pots which are developed from propolis. This gives the honey rich in nutrients and phenolic compounds, mainly flavonoids and phenolic acids, which act as potent antioxidants.

The stingless bee honey (SBH) has high nutritional and therapeutic value due to the diversified plants foraged including medicinal plants and structure of nest, the antimicrobial activity of SBH is little bit stronger than other honeys with unique therapeutic properties with great potential to be developed for modern medical uses. It is used in many medicines in Indian System of

Medicine, the Ayurveda for many diseases, in reducing the heat of the body, inflammation, bleeding in the throat, impurities of the blood, boils and even for cancer. SBH as a therapeutic agent in various health issues such as anti-diabetic wound healing, anticancer, treatment of eye diseases, and also in fertility. Studies have proven that the SBH has excellent potential and portrays beneficial effects as antimicrobial, anticancer agent, improving hypertension, lipid profiles, and with some studies showing better anti-diabetic effects. In order to provide a major comprehensive understanding on the potential uses and benefits of the SBH, more systematic studies need to be carried out. In terms of quality control, methods to authenticate pure SBH need to be developed. A rapid and destructive analysis technique is required to avoid possible adulteration by irresponsible manufacturers. The quality standard can be established by the identification of its bioactive component since SBH is rich in antioxidants and innovative efforts should be taken to fully explore and utilize these benefits. Honey-based products should be diversified, such as making supplement capsules or tablets which contain probiotics isolated from the SBH that can aid in gastrointestinal health. These properties should also be made readily in the form of topical creams or gels for wound healing or other purposes (Fig.15).

Pollination service and playhouses

There is ample potential for utilization of stingless bees in pollination service as they





are better pollinators of some crops than honey bees which thrive much better in tropical areas and are polylectic. The smaller body size, shorter foraging distance, less aggressiveness, floral constancy, higher longevity of the colonies, efficient worker recruitment behavior towards food sources, medium colony size, lesser swarming tendency, tolerant to high temperature and less pest and disease incidence are the advantages of stingless bees as pollinators. Studies conducted at Tamil Nadu Agricultural University (Tej *et al.*, 2017) revealed that keeping stingless bees in greenhouse cucumber can improve its pollination and thus fruit weight and yield. Stingless bees play an important ecological role as pollinators of many wild plant species and seem good candidates for future alternatives in commercial pollination. Eleven stingless bee species across six genera have been found to forage effectively under enclosed conditions, indicating the potential of stingless bees as pollinators of greenhouse crops- over the past six years the number of crops reported to be effectively pollinated by stingless bees has doubled, putting the total figure on 18 crops. (Judith *et al.*, 2006). It is reported in sweet pepper *Capsicum annuum* L. that despite flowers are considered autogamous, this crop benefits from pollination by stingless bee *Melipona subnitida*, producing fruits significantly heavier and wider, containing a greater number of seeds and of better quality (lower percentage of malformed fruits) than self-pollinated sweet pepper. Thus, *M. subnitida* can be considered

an efficient pollinator of greenhouse sweet pepper (Cruz *et al.*, 2006). The potential of stingless bees for seed production, pollination and yield enhancement of various crops under protected cultivation to be explored in India has by conducting scientific studies.

Pests / Enemies of stingless bee

Studies conducted at this centre could identify some enemies of stingless bees, either

attacking while foraging or encountering the colony. They are dipteran fly *Hermetia illucens* L., Nitidulid beetle *Aethina* sp. reduviid bug *Acanthaspis siva*, Assassin Bug *Sycanus* sp., predatory spiders viz., *Thomisus lobosus*, *Thomisus projectus*, *Menemerus bivittatus*, *Neoscona nautica*, *Heteropoda venetoria*, ants *Solenopsis geminata*, a species of megachilid bee, a mite *Amblyseius* sp. etc

	
<p>Fig. 1. Dr.T.P. Rajendran, Assistant Director General (PP), released the book “status paper on stingless bee <i>Trigona iridipennis smith</i>”</p>	<p>Fig. 2. Commercial meliponiculture</p>
	
<p>Fig. 3. Hive entrance helps to detect the presence of feral colony</p>	<p>Fig. 4. Nest architecture</p>


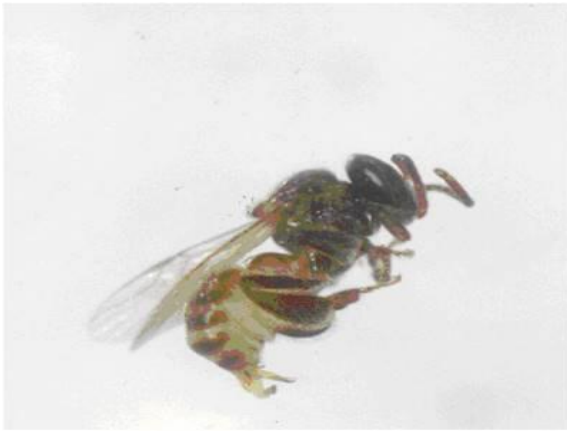




	
<p>Fig. 5. Stingless bee queen</p>	<p>Fig. 6. Stingless drone</p>
	
<p>Fig. 7. Stingless worker</p>	<p>Fig. 8. Hiving feral colonies without damaging compound wall</p>
	
<p>Fig. 9. KAU hive for commercial meliponiculture</p>	<p>Fig. 10. Mass provisioning</p>



Fig. 11. Division of colonies



Fig. 12. Dearth season management-
Artificial feeding of colonies



Fig. 13. Worker bees transferred in the empty
water bottle.



Fig. 14. Honey extraction without harming
the worker bees



Fig. 15. Hygienic pure honey pots

Conclusions

Practicing commercial meliponiculture in the homesteads will provide the service of stingless bees in pollination of various crops particularly vegetables and fruits and their yield enhancement. The services of stingless bees can be utilized for quality seed production too. "If the bee disappears from the surface of the earth, man would have no more than four years to live. No more bees, no more pollination, no more plants, no more animals, no more man." (Albert Einstein)

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Erebid Moths of Lonavala: A tourist spot in the Western Ghats with tremendous anthropogenic pressure, and a new range extension record

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Abstract

The present study was taken up with an aim to document the diversity of Erebid moth from Lonavala, Maharashtra. A total of 44 species of 36 genera in 10 subfamilies belonging to Erebiidae, Lepidoptera have been reported. It was a part of a short project conducted by the Green Skilled Development Programme (GSDP) students of the certificate course on para-taxonomy [including Peoples Bio-diversity Register (PBRs)] at Zoological Survey of India (ZSI), Western Regional Centre (WRC), Pune. This article also presents a new distribution and range extension of *Calesia* sp. One endemic moth, *Olepa clavatus* (Swinhoe, 1885) is also reported from the studied site. Additionally, the data on type species, type locality along with their distributional record from India and outside India, bionomics for each species are also presented.

Key words: Moths, Northern Western Ghats, diversity, anthropogenic, *Calesia*.

Introduction

Insects are the most successful creatures on the earth. Among them, Lepidoptera is one of the most diverse insect orders. Moths and butterflies are the insect groups under Lepidoptera. There are approximately 160,000 species of moths (Nieukerken *et al.*, 2011) and most of them are major and minor pest. Moths are nocturnal feasting on the varied crops, wild trees, ornamental plants, etc. They are of economic importance as the larvae of majority of the moths feed voraciously on the plant parts (leaves, flowers, stem, seeds) and are the

damaging stage responsible for food losses. Erebiidae moths are of immense economic importance as some adults (fruit sucking moth) feed on the commercial fruits like citrus, causing economic loss to the farmers. In case of a widely distributed fruit sucking moth, *Eudocima phalonia* (Linnaeus, 1763), the adults have destructive feeding habits and not the larvae. Larvae of this moth feed on wild trees (Menispermaceae and Fabaceae) (Kumar and Lal, 1983). The damage in citrus orchard in India may vary from 10–55% (Dadmal and Pawar, 2001) and 57% in pomegranate (Mote *et al.*, 1991). Some microorganisms introduced

on the citrus orchards due to the feeding habits of these moths cause rotting and premature fruit fall (Sands *et al.*, 1993). The damage caused by the moth is also severe in economic crops like cotton, brinjal, citrus, tomato, sugarcane, cereals, millets, pulses, vegetables and plantation crops.

Besides, some are indicator taxa that utilize lichens as a food source and are useful in pollution monitoring (Kendrick, 2002). They are preyed in different life stages by bats, birds, lizards, amphibians, dragonflies, spiders, small mammals, fungi, bacteria etc. It is one of the most speciose families with the species count of 24,569 in 1,760 genera (Nieukerken *et al.*, 2011) and are well-studied. From Maharashtra, 86 species have been reported by Mitra *et al.*, (2019); 128 species from northern Western Ghats of Maharashtra by Subhalaxmi *et al.*, (2011); 101 species from the northern Maharashtra (Gurule and Nikam, 2013); 10 species from Pench National Park, Maharashtra (Kalawate & Sharma, 2017); 44 species from northern Western Ghats of Maharashtra (Kalawate, 2018a); two species of erebid moths, *Gurna indica* (Moore 1879) (Kalawate *et al.*, 2019) and *Mecodina metagraptia* Hampson, 1926 (Kalawate, 2018b) were recently rediscovered from India.

The present study was a part of short project undertaken by the GSDP students of Para-Taxonomy [including Peoples Biodiversity Register (PBR)] certificate course of ZSI, WRC, Pune, and hence was a time bound

project (course duration: 03 months). The nearby study site selected was Lonavala, a small town and a hill station in Pune district of Maharashtra and a part of Western Ghats, which is an important Biodiversity Hotspots and a UNESCO world heritage site. It is surrounded by the highly urbanised cities like Mumbai and Pune, thus faces lot of pressure of tourists being a hills station. The biodiversity in Northern Western Ghats faces degradation by human exploitation. The global conservation problem is the loss and fragmentation of tropical rainforest forming a major proportion of the world's biodiversity (Whitmore, 1997; Kapoor, 2006). Fragmentation of natural habitat is a problem in the northern Western Ghats, and moths are sensitive to habitat fragmentation and the species whose larvae are monophagous are more affected by the loss of habitat than the polyphagous (Ockinger *et al.*, 2010).

The distributional records of the species from India and outside India along with the type species and type locality data and bionomics have been provided in this paper. On perusal of literature, it was found that less or no work on the moth fauna of Lonavala in general and Erebidae in particular has been carried out and hence in the present study an attempt has been made to assess the Erebid moth fauna of Lonavala.

Materials and methods

Moths were collected from Lonavala, Maharashtra (Fig. 2) using light trap (Fig. 1).

The light trap is consisting of white cloth measuring 3 m long x 1.5 m and was hung in between the two poles. The light source used was Mercury Vapour Lamp of 160 W powered by portable generator and was hanged middle of the white cloth. The collected specimens were euthanized by ethyl acetate vapors and preserved as dry. The specimens were relaxed, pinned and preserved in the laboratory for further studies. They were studied under Leica EZ 4 HD stereozoom microscope. All identified specimens were labelled, duly registered and deposited at National Zoological Collection, ZSI, WRC, Pune, Maharashtra, India. The latitude and longitude coordinate of the collection site was 18.7546171°N & 73.4062342°E and elevation 626m. The map of the study site (Fig. 2) was prepared using open free software QGIS. Images of the moths are depicted in Fig. 4 to Fig. 5.

The identification of the moths was done with the help of Hampson (1894, 1895). The classification and sequences of the subfamilies followed is as per Kononenko & Pinratana (2013); Zahiri *et al.*, 2011, 2012. The distribution and larval host plants have been consulted from Hampson (1894; 1895); Shubhalaxmi *et al.*, (2011); Gurule & Nikam (2013); Smetacek (2008) and Sivasankaran *et al.*, (2017).

In the foregoing pages, the taxonomic account along with their distributional record and bionomics has been documented.

Abbreviations used: Coll.: Collected by; WRC: Western Regional Centre; ZSI: Zoological Survey of India.

Results and discussion

This study was a part of the short-term project conducted by the students of the Certificate course on Para-taxonomy [including Peoples Bio-diversity Register (PBRs)] of GSDP at ZSI, WRC, Pune. The certificate course was of three months duration and hence was given a short period bound project to the students. Lonavala, a part of the Sahyadri ranges is a famous hills station near metro cities like Mumbai and Pune and hence, faces tremendous anthropogenic pressure. On literature review, it was found that report on moths of Lonavala is lacking.

The present study resulted in enumeration of 44 species in 36 genera belonging to 10 Subfamilies of Erebid moth fauna of Lonavala, Maharashtra. One new range extension records of the species, *Calesia fuscicorpus* Hampson, 1891 is reported in this study. To confirm the new distributional record and its range extension, Hampson (1891) and Sondhi *et al.*, (2018) were consulted. One endemic moth to India has been recorded from the study area. From Fig. 3, it can be seen that the maximum number of species recorded was from the subfamily Erebininae (17 species) followed by Arctiinae (13 species), Aganainae (05 species), Tinoliinae (02 species), Calpinae (01 species), Lymantriinae (01 species), Hypeninae (01 species), Hypocalinae (01 species) and Scoliopteryginae (01 species).

Taxonomic account Superfamily NOCTUOIDEA Latreille, 1809 Family EREBIDAE Leach, [1815] Subfamily EREBINAE Leach [1815]					
Genus	Species	Type locality	Material Examine	Distribution	Bionomics
<i>Hyospila</i> Guenée, 1852 Type Species: <i>Hyospila bolinoides</i> Guenée, 1852.	<i>Hyospila bolinoides</i> Guenée, 1852	Java.	05 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1605).	India: Throughout India including Maharashtra (Mumbai, Satara, Nashik and Dhule). Elsewhere: Australia, China, Japan, Malaysia, Myanmar, New Guinea, Sri Lanka. Elsewhere: Australia, Cambodia, China, Indonesia, Japan, Korea, Malaysia, Nepal, New Guinea, Sri Lanka, Thailand, Vietnam (Mitra <i>et al.</i> , 2019).	The larval host plant is <i>Derris</i> (Leguminosae) (Holloway, 2005). In present study, it is reported in the late July, and Mitra <i>et al.</i> , (2019) recorded it in October from Raigad, Maharashtra. As per Holloway (2005) it is a lowland species and is recorded up to a level of 1930m.
<i>Hamodes</i> Guenée, 1852 Type Species: <i>Ophiura propitia</i> Boisduval, 1832 [= <i>Hamodes propitia</i> (Guerin-Meneville, 1831)].	<i>Hamodes propitia</i> (Guerin-Meneville, 1831)	Nouvelle-Irlande [Bismarck Archipelago].	03 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1585).	Throughout India including Maharashtra (Nashik, Dhule, Jalgaon and Nandurbar). Elsewhere: Australia, Indonesia, Malaysia, Myanmar, New Ireland, Philippines, Solomon Islands, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plant is <i>Dalbergia</i> (Leguminosae) (Holloway, 2005). This species is found in the altitude ranging from sea-level to almost high-altitude level of 2110m and in various forest types and also in cultivated area of agricultural lands.
<i>Erebus</i> Latreille, 1810 Type Species: <i>Phalaena crepuscularis</i> Linnaeus, 1758.	<i>Erebus caprimulgus</i> (Fabricius, 1781)	China.	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1608).	Throughout India including Maharashtra (Amravati, Pune, Sindhudurg, Nashik and Dhule). Elsewhere: Indonesia, Malaysia, Myanmar, Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Smilax macrophylla</i> , <i>Smilax ovalifolia</i> (Smilacaceae) (Leong and Kueh, 2011). Adult feeds on the ripened fruits of <i>Melastoma malabathricum</i> (Melastomataceae) (Leong and Kueh, 2011). The species prefers mostly lowland forest but some are recorded from the high altitudinal areas (Holloway, 2005).
	<i>Erebus macrops</i> (Linnaeus, 1768)	"India Orientali" [India].	02 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1609).	India: Kerala, Maharashtra, Tamil Nadu, Uttarakhand. Elsewhere: China, Indonesia, Malaysia, Myanmar, Nepal, Sri Lanka, Thailand (Kalawate, 2018; Mitra <i>et al.</i> , 2019).	The larval host plant is <i>Acacia</i> (Leguminosae) (Holloway, 2005); adult is a fruit piercer (Bänziger, 1982). It is a common species and found commonly near the human dwellings.
<i>Artena</i> Walker, 1858 Type Species: <i>Artena submira</i> Walker, 1858.	<i>Artena dotata</i> (Fabricius, 1794)	"India Orientali" [India].	04 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1606).	India: Throughout India including Maharashtra (Mumbai, Pune, Sindhudurg, Nashik, Dhule, Jalgaon and Nandurbar). Elsewhere: Indonesia, Japan Malaysia, Sri Lanka, Taiwan (Mitra <i>et al.</i> , 2019).	The larval host plants recorded are <i>Combretum</i> , <i>Getonia</i> , <i>Quisqualis</i> , <i>Terminalia</i> (Combretaceae) (Holloway, 2005). Adult feeds on fruits by piercing it and sucking the juice (Bänziger, 1982). As per Holloway (2005), the species prefers lowland forest areas, with secondary vegetation after logging. The recorded flight period is from April to June and August to December.
	<i>Artena submira</i> Walker, 1858	Hindustan.	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1795).	India: Jharkhand, Kerala, Maharashtra. Elsewhere: Bangladesh, Myanmar, Thailand, Vietnam (Singh & Ranjan, 2016; Mitra <i>et al.</i> , 2019).	The Larval food plants are <i>Getonia floribunda</i> , <i>Quisqualis indica</i> , <i>Terminalia paniculate</i> , <i>T. tomentosa</i> (Combretaceae) (NHM, 2021). It is reported in the month of July in this study.
<i>Thyas</i> Fabricius, 1775 Type Species: <i>Thyas honesta</i> Hübner, 1824.	<i>Thyas honesta</i> Hübner, 1806	[India. Indies]? East	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1607).	India: Throughout India including Maharashtra (Pune, Satara, Nashik, Dhule, Jalgaon, Nandurbar). Elsewhere: Borneo, Indonesia, Malaysia, Myanmar, Philippines, Sumatra, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Careya</i> , <i>Barringtonia</i> , <i>Planchonia</i> (Lecythidaceae); <i>Terminalia</i> Combretaceae (Holloway, 2005); <i>Citrus</i> (Rutaceae) (Ngampongsai <i>et al.</i> , 2005). It is recorded in August in this study and by Sambath (2014) from Jharkhand. It is recorded from 300m to 1620m (Holloway, 2005).

	<i>Thyas coronata</i> (Fabricus, 1775)	China.	03 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1610).	Throughout India including Maharashtra. Elsewhere: Australia, Borneo, Indonesia, Myanmar, Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Quisqualis indica</i> (Combretaceae) (Hampson, 1894); <i>Terminalia</i> (Combretaceae), <i>Litsea</i> (Lauraceae), <i>Anamirta</i> (Menispermaceae), <i>Pinus</i> (Pinaceae), <i>Nephelium</i> (Sapindaceae) (Holloway, 2005). The adult is a piercer of fruit on <i>Citrus</i> (Rutaceae) (Bänziger, 1982) and commonly called as fruit piercing moth. It is mainly found in the forests, disturbed habitats and upto 2600m (Holloway, 2005).
<i>Buzara</i> Walker, [1865] 1864 Type Species: <i>Buzara chrysomela</i> Walker, 1865.	<i>Buzara onelia</i> (Guenee, 1852)	Silhet, Bangladesh.	02 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1807).	Distribution: India: Himachal Pradesh, Maharashtra (Satara, Sindhudurg). Elsewhere: Bangladesh, China, Indonesia, Malaysia, Myanmar, Malay Peninsula, Nepal, Philippines, Singapore, Sri Lanka, South Japan, Taiwan, Thailand, Vietnam, (Kononenko & Pinratana, 2013; Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Breynia</i> , <i>Phyllanthus</i> , <i>Sauropus</i> (Euphorbiaceae) (Holloway & Miller, 2003). This species found from lowland to mid montane forest. In this study the specimen is collected in August.
<i>Polydesma</i> Boisduval, 1833 Type Species: <i>Polydesma umbricola</i> Boisduval, 1833.	<i>Polydesma boarmioides</i> Guenee, 1852	Java.	03 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1661).	India: Sikkim, Uttar Pradesh, Maharashtra (Mumbai, Pune, Dhule, Jalgaon). Elsewhere: Australia, Bangladesh, Fiji, Hawaii, Malaysia, New Caledonia, Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Gmelina</i> (Lamiaceae); <i>Acacia</i> , <i>Albizia</i> , <i>Pithecellobium</i> (Leguminosae); <i>Salix</i> (Salicaceae); <i>Litchi</i> (Sapindaceae) (Holloway, 2005). This species is recorded in August, November and December.
<i>Ericeia</i> Walker, [1858] Type Species: <i>Ericeia sobria</i> Walker, 1858.	<i>Ericeia inangulata</i> (Guenee, 1852)	Silhet.	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1813).	India: Throughout India including Andaman and Nicobar Islands, Himachal Pradesh, Madhya Pradesh, Maharashtra (Mumbai, Pune, Nashik and Jalgaon) and Uttarakhand, West Bengal. Elsewhere: Africa, Australia, China, Myanmar, Sri Lanka (Mitra <i>et al.</i> , 2019).	The Larval host plants are <i>Acacia</i> , <i>Albizia</i> , <i>Cassia</i> , <i>Dalbergia</i> , <i>Mimosa</i> , <i>Paraserianthes</i> , <i>Senna</i> , <i>Xylia</i> (Leguminosae); <i>Adiantum</i> (Adiantaceae); <i>Lagerstroemia</i> (Lythraceae); <i>Citrus</i> (Rutaceae) (Holloway, 2005). The adult moth is a fruit piercer (Bänziger, 1982). Records have been made from the lowlands to 2600m, but the species is commoner above 1500m.
<i>Parallela</i> Hübner, 1818 Type Species: <i>Parallela bistriaris</i> Hübner, 1818.	<i>Parallela stuposa</i> (Fabricius, 1794)	"India Orientali" [India].	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1810).	India: Maharashtra (Mumbai, Pune, Sindhudurg, Satara, Mumbai, Nashik, Dhule, Jalgaon, Nandurbar). Elsewhere: China, Indonesia, Japan, Korea, Philippines, Sri Lanka (Mitra <i>et al.</i> , 2019).	The Larval host plants are Salicaceae; Euphorbiaceae; Rosaceae; Lythraceae (Leley, 2016). In the present study it is reported in August.
<i>Achaea</i> Hübner, [1823] Type Species: <i>Phalaena melicerta</i> Drury, 1773 [= <i>Achaea janata</i> (Linnaeus, 1758)].	<i>Achaea (Acanthodelta) janata</i> (Linnaeus, 1758)	Indiis [India].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1603).	India: Andaman and Nicobar Islands, Himachal Pradesh, Maharashtra (Pune, Sindhudurg, Satara, Mumbai, Nashik, Dhule, Jalgaon, Nandurbar). Elsewhere: Australia, China, Indonesia, Japan, Myanmar, New Guinea, New Zealand, Philippines (Mitra <i>et al.</i> , 2019).	Polyphagous: The larval host plants are <i>Ricinus</i> (Euphorbiaceae); Leguminosae; <i>Agathis</i> , <i>Araucaria</i> (Araucariaceae); <i>Anogeissus</i> , <i>Terminalia</i> (Combretaceae); <i>Ipomoea</i> (Convolvulaceae); <i>Brassica</i> , <i>Raphanus</i> , <i>Cucurbita</i> (Cucurbitaceae); <i>Cupressus</i> (Cupressaceae); <i>Shorea</i> (Dipterocarpaceae); <i>Acalypha</i> , <i>Aleurites</i> , <i>Andrachne</i> , <i>Bischofia</i> , <i>Chamaesyce</i> , <i>Codiaeum</i> , <i>Croton</i> , <i>Euphorbia</i> , <i>Excoecaria</i> , <i>Flueggea</i> , <i>Jatropha</i> , <i>Manihot</i> , <i>Pedilanthus</i> , <i>Phyllanthus</i> , <i>Ricinus</i> , <i>Sapium</i> (Euphorbiaceae); <i>Saccharum</i> (Gramineae); <i>Planchonia</i> (Lecythidaceae); <i>Acacia</i> , <i>Albizia</i> , <i>Arachis</i> , <i>Bauhinia</i> , <i>Dalbergia</i> , <i>Desmanthus</i> , <i>Glycine</i> , <i>Leucaena</i> , <i>Mimosa</i> , <i>Paraserianthes</i> , <i>Phaseolus</i> , <i>Prosopis</i> , <i>Vigna</i> , <i>Zylia</i> (Leguminosae); <i>Strychnos</i> (Loganiaceae); <i>Lagerstroemia</i> , <i>Punica</i> (Lythraceae); <i>Abutilon</i> , <i>Gossypium</i> (Malvaceae); <i>Ficus</i> (Moraceae); <i>Eucalyptus</i> , <i>Psidium</i> (Myrtaceae); <i>Cocos</i> (Palmae); <i>Emex</i> (Polygonaceae);

					<i>Polypodium</i> (Polypodiaceae); <i>Macadamia</i> (Proteaceae); <i>Ziziphus</i> (Rhamnaceae); <i>Rosa</i> (Rosaceae); <i>Litchi</i> , <i>Nephelium</i> , <i>Schleichera</i> (Sapindaceae); <i>Madhuca</i> , <i>Mimusops</i> , <i>Palaquium</i> (Sapotaceae); <i>Capsicum</i> (Solanaceae); <i>Sterculia</i> , <i>Theobroma</i> (Sterculiaceae); <i>Grewia</i> (Tiliaceae); <i>Tribulus</i> (Zygophyllaceae) (Holloway, 2005). The adult makes the fruit unfit for consumption by piercing it and feasting upon it (Bänziger, 1982). The adult moth is attracted to the light traps
<i>Bastilla</i> Swinhoe, 1918 Type Species: <i>Ophiura redunca</i> Swinhoe, 1900 [= <i>Bastilla hamatilis</i> (Guenée, 1852)].	<i>Bastilla joviana</i> (Stoll, 1782)	Cote de Coromandel [India].	02 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (Reg. No. ZSI-WRC-L-1604).	India: Adaman Islands, Maharashtra (Pune, Sindhudurg, Nashik), West Bengal. Elsewhere: Australia, Bangladesh, Bhutan, China, Indonesia, Malaysia, Myanmar, Nepal, New Guinea, Pakistan, Sri Lanka, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Acalypha</i> , <i>Breynia</i> and <i>Phyllanthus</i> (Euphorbiaceae) (Holloway & Miller, 2003). It is like other fruit sucking moths' pierces fruit (Bänziger, 1982). Recorded from lowlands to 1930m and from forested and cultivated areas as per Holloway (2005). In this study it is recorded from 626m.
<i>Grammodes</i> Guenée, 1852 Type Species: <i>Noctua geometrica</i> Fabricius, 1775 [= <i>Grammodes geometrica</i> (Fabricius, 1775)].	<i>Grammodes geometrica</i> (Fabricius, 1775)	India Orientali [India].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1793).	India: Andaman and Nicobar Islands, Assam, Madhya Pradesh, Maharashtra (Mumbai, Pune, Satara, Sindhudurg, Nashik, Dhule, Jalgaon and Nandurbar), Odisha, Punjab, Rajasthan, Sikkim, Tamil Nadu, Uttar Pradesh, West Bengal. Elsewhere: Africa, Australia, Bangladesh, China, Europe, Indonesia, Myanmar, New Guinea, Singapore, Sri Lanka, Taiwan (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Phyllanthus</i> , <i>Sapium</i> (Euphorbiaceae); <i>Cistus</i> (Cistaceae); <i>Diospyros</i> (Ebenaceae); <i>Ricinus</i> (Euphorbiaceae); <i>Oryza</i> (Gramineae); <i>Polygonum</i> (Polygonaceae); <i>Ziziphus</i> (Rhamnaceae); <i>Tamarix</i> (Tamaricaceae) (Holloway, 2005). Adults are fruit piercer (Bänziger, 1982). It is recorded in forests and cultivated area. Adults are attracted to the light trap.
<i>Spirama</i> Guenée, 1852 Type Species: <i>Phalaena retorta</i> Clerck, 1764 [= <i>Spirama retorta</i> (Clerck, 1764)].	<i>Spirama retorta</i> (Clerck, 1764)	Not known.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1611).	India: Andaman and Nicobar Islands, throughout India including Himachal Pradesh, Maharashtra (Mumbai, Pune, Sindhudurg, Matheran, Nashik, Dhule, Jalgaon, Nandurbar). Elsewhere: Japan, China, Sri Lanka, Myanmar, Andamans, Java (Sekhon and Singh, 2015; Mitra <i>et al.</i> , 2019).	The larval host plant are <i>Vitis</i> (Vitaceae); <i>Acacia mangium</i> (Fabaceae) (NHM, 2021). It is attracted to light and recorded at an altitude of 626m.
<i>Mocis</i> Hübner, [1823] Type Species: <i>Phalaena virbia</i> Cramer, 1780 (= <i>Mocis undata</i> (Fabricius, 1775)).	<i>Mocis undata</i> (Fabricius, 1775)	East Indies.	01 ex., Amby valley Road, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1903).	India: Throughout India including Andaman and Nicobar Islands, Himachal Pradesh, Madhya Pradesh, Maharashtra, Uttarakhand, West Bengal. Elsewhere: Africa, throughout Oriental region (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Cytisus</i> , <i>Desmodium</i> , <i>Wisteria</i> (Fabaceae); <i>Arachis</i> , <i>Butea</i> , <i>Cajanus</i> , <i>Calopogonium</i> , <i>Crotalaria</i> , <i>Derris</i> , <i>Desmodium</i> , <i>Glycine</i> , <i>Indigofera</i> , <i>Mucuna</i> , <i>Phaseolus</i> , <i>Pueraria</i> , <i>Rhynchosia</i> , <i>Tephrosia</i> , <i>Vigna</i> (Leguminosae); <i>Shorea</i> (Dipterocarpaceae); <i>Hevea</i> (Euphorbiaceae); <i>Gossypium</i> (Malvaceae); <i>Nephelium</i> (Sapindaceae); <i>Solanum</i> (Solanaceae) (Holloway, 2005). It is recorded mostly from the open habitat, cultivation and disturbed forest. It is usually found from lowlands to 1200m (Holloway, 2005).
EREBINAE incertae sedis					
<i>Ischyja</i> Hubner, [1823] 1816 Type species: <i>Phalaena manlia</i> Cramer, 1776.	<i>Ischyja manlia</i> (Cramer, 1776)	Cote de Coromandel (=Tamil Nadu).	01 ex., Amby valley Road, 23.viii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1584).	India: Throughout India including Maharashtra. Elsewhere: Indonesia, Myanmar, Sri Lanka (Sambath, 2014).	The larval host plants are Lauraceae, Lardizabalaceae, Ebenaceae, Rosaceae, Combretaceae, Rubiaceae (Leley, 2016), Theaceae (Holloway, 2005). As per Kononenko and Pinratana (2013) the flight period is from August – November. But, in the present study, it is reported in late July.

Subfamily Arctiinae (Leach, 1815)					
<i>Amata</i> Fabricius, 1807 Type Species: <i>Zygaena passalis</i> Fabricius, 1781.	<i>Amata passalis</i> (Fabricius, 1781)	[India or Sri Lanka?].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1806).	India: Andhra Pradesh, Assam, Karnataka, Maharashtra (Aurangabad, Solapur, Pune, Mumbai), Manipur, Tamil Nadu, West Bengal. Elsewhere: Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Ipomoea</i> (Convolvulaceae); <i>Phaseolus</i> , <i>Cajanus</i> (Leguminosae); <i>Dahlia</i> , <i>Cosmos</i> (Compositae); <i>Santalum album</i> (Santalaceae); <i>Vigna unguiculata</i> (Leguminosae) (Venkatesha and Gopinath, 1992). This species breeds throughout the year and passes through 6-11 generations a year.
	<i>Amata bicincta</i> (Kollar, [1844])	Not known.	03 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1586).	Arunachal Pradesh, Himachal Pradesh, Maharashtra, Meghalaya, North West India, Sikkim, Uttarakhand, West Bengal, Elsewhere: China (Hampson, 1898; Shubhalaxmi <i>et al.</i> , 2011; Singh <i>et al.</i> , 2014).	The larval host plant is not known.
	<i>Amata bicincta</i> (Kollar, [1844])	Not known.	03 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1586).	India: Arunachal Pradesh, Meghalaya, North West India, Sikkim, West Bengal, (Singh <i>et al.</i> , 2014).	The larval host plant is not known.
<i>Cretonotos</i> Hubner, 1819 Type Species: <i>Phalaena interrupta</i> Linnaeus, 1767.	<i>Cretonotos gangis</i> (Linnaeus, 1763)	Not known.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1587).	India: Throughout India including Maharashtra (Nashik, Nandurbar). Elsewhere: Australia, China, Japan, Malaysia, Nepal, New Guinea, Pakistan, Philippines, Sri Lanka, Taiwan, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Arachis hypogaea</i> , <i>Medicago sativa</i> , <i>Vigna mungo</i> (Fabaceae); <i>Eleusine coracana</i> , <i>Oryza sativa</i> , <i>Pennisetum glaucum</i> , <i>Zea mays</i> (Poaceae); <i>Ipomoea batatas</i> (Convolvulaceae); <i>Mimulus gracilis</i> (Phrymaceae) (NHM, 2021). They found in the secondary habitats from the lowlands to the montane region. They are attracted to light. The males of this species have four eversible coremata at the tip of the abdomen which emit pheromones, and is longer than the abdomen. It breeds throughout the year.
	<i>Cretonotos transiens</i> (Walker, 1855)	Assam.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1588).	India: Arunachal Pradesh, Assam, Maharashtra, (Mumbai, Pune, Satara, Sindhudurg, Nashik), Manipur, Meghalaya, Mizoram, Nagaland, Sikkim, Uttarakhand, West Bengal. Elsewhere: China, Indonesia, Japan, Malaysia, Nepal, Philippines (Mitra <i>et al.</i> , 2019).	The larval host plants are Polyphagous: <i>Beta</i> (Chenopodiaceae); <i>Dioscorea</i> (Dioscoreaceae); <i>Paspalum</i> , <i>Zea</i> (Gramineae); <i>Pithecellobium</i> , <i>Vigna</i> , <i>Wisteria</i> (Leguminosae); <i>Toona</i> (Meliaceae); <i>Musa</i> (Musaceae); <i>Salix</i> (Salicaceae); <i>Cayratia</i> , <i>Cissus</i> (Vitidaceae) (Holloway, 1988). The species is common in cultivated agricultural fields, open habitats and secondary vegetation. Adults are commonly attracted to light.
<i>Mangina</i> Kaleka & Kirti, 2001 Type Species: <i>Euprepia argus</i> Kollar, [1844] (= <i>Argina argus</i> Kollar, 1844).	<i>Mangina syringa</i> (Cramer, 1775)	“Coromandel” [India].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1589).	India: Throughout India including Maharashtra (Pune). Elsewhere: Australia, Sri Lanka (Mitra <i>et al.</i> , 2019).	Larval host plants are <i>Crotalaria assamica</i> , <i>Crotalaria juncea</i> , <i>Crotalaria longipes</i> , <i>Crotalaria saltiana</i> (Leguminosae); <i>Musa × paradisiaca</i> (Musaceae) (Kirti and Singh, 2015). In this study it is recorded in August. It is not a frequent visitor at light trap.
<i>Olepa</i> Watson, 1980 Type Species: <i>Alope ocellifera</i> (Walker, 1855) [= <i>Olepa</i>	<i>Olepa clavatus</i> (Swinhoe, 1885)	Bombay (India).	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1590).	India: South India, Maharashtra (Cotes and Swinhoe, 1887; Dubatolov, 2010; Kalawate <i>et al.</i> , 2020).	The larval host plant is not known. It is usually attracted to the light. Remark: Endemic to India.

<i>ocellifera</i> (Walker, 1855)].	<i>Olepa ricini</i> (Fabricius, 1775)	"Indiae orientalis ricino" (India).	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1826).	India: Throughout India including Maharashtra (Mumbai, Pune, Ahmednagar, Satara). Elsewhere: Bangladesh, Nepal, Northern Pakistan, Sri Lanka, Thailand (Cotes and Swinhoe, 1887; Dubatolov, 2010; Kalawate <i>et al.</i> , 2020).	Larval host plants are <i>Calotropis procera</i> (Apocynaceae); <i>Camellia sinensis</i> (Theaceae); <i>Campsis grandiflora</i> (Bignoniaceae); <i>Gossypium</i> (Malvaceae), <i>Ricinus communis</i> (Euphorbiaceae); <i>Helianthus</i> (Asteraceae), <i>Zea mays</i> (Poaceae); <i>Coccinia grandis</i> (Cucurbitaceae); <i>Solanum melongena</i> (Solanaceae); <i>Ipomoea batatas</i> (Convolvulaceae); <i>Musa</i> (Musaceae) (Farooqui <i>et al.</i> , 2020; TNAU, 2021; Shubhalaxmi, 2018). They are attracted to light.
	<i>Olepa zedesi</i> Kalawate, 2020	Pune, India.	01 ex., Lonavala, Pune, 23.viii.2017, coll. A.S. Kalawate (ZSI-WRC-L-2154).	India: Maharashtra (Pune, Lonavala, Satara) (Kalawate, 2021).	Data deficient.
<i>Rajendra</i> Moore, 1879 Type Species: <i>Rajendra lativitta</i> Moore, 1879.	<i>Rajendra biguttata</i> (Walker, 1855)	Canara, Malabar Coast [India].	03 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1591).	India: Arunachal Pradesh, Bihar, Himachal Pradesh, Karnataka, Kerala, Maharashtra, Madhya Pradesh, Orissa, Punjab, Sikkim, Tamil Nadu, West Bengal. Elsewhere: Bangladesh (Dubatolov <i>et al.</i> , 2007).	The larval host plants is <i>Elettaria cardamomum</i> (Zingiberaceae) (Kirti and Singh, 2015). The flight period is from April to December. It is attracted to light.
<i>Argina</i> Hübner, [1819] Type species: <i>Phalaena cribraria</i> Clerck, 1764 (= <i>Argina astrea</i> (Drury, 1773)).	<i>Argina astrea</i> (Drury, 1773)	[Ghana], Africa, Gold Coast.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1595).	India: Throughout India including Maharashtra (Pune, Mumbai, Satara, Amravati) (Dubatolov, 2010; Cotes and Swinhoe, 1887). Elsewhere: China, Myanmar, Mauritius, New Guinea, Sri Lanka (Kaleka and Kirti, 2001).	The larval host plants are <i>Crotalaria</i> spp., <i>Lablab purpureus</i> , <i>Melilotus indica</i> (Leguminosae); <i>Beaumontia</i> (Apocynaceae); <i>Buddleja</i> (Buddlejaceae); <i>Theobroma cacao</i> (Sterculiaceae) (Holloway, 1988; NHM, 2021). The flight period is throughout the year. It is attracted to the light. It is found in the lowlands, cultivated areas and open habitats.
<i>Cyana</i> Walker, 1854 Type Species: <i>Cyana detrita</i> Walker, 1854.	<i>Cyana puella</i> (Drury, 1773)	Madras [India].	02 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1592).	India: Chhattisgarh, Himachal Pradesh, Madhya Pradesh, Maharashtra (Mumbai, Pune), North West Himalayas, Sikkim, South India, Uttarakhand. Elsewhere: Africa, Indonesia, Madagascar, Nepal, Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plant is unknown. In this study, the specimen was collected in August.
<i>Brunia</i> Moore, 1878 Type Species: <i>Lithosia antica</i> Walker, 1854.	<i>Brunia antica</i> (Walker, 1854)	Ceylon (=Sri Lanka).	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1594).	India: Andaman and Nicobar Islands, Maharashtra (Mumbai, Pune, Satara, Nashik, Dhule), West Bengal. Elsewhere: China, Indonesia, Malaysia, Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Hevea</i> (Euphorbiaceae), <i>Terminalia</i> (Combretaceae) and <i>Theobroma</i> (Sterculiaceae) (Holloway, 2005). It's a lowland species and observed frequently in coastal vegetation and mangrove (Holloway, 2001). In this study it is found in August and at an altitude of 626m and is attracted to light.
<i>Nannoarctia</i> Koda, 1988 Type Species: <i>Pericallia takanoi</i> Sonan, 1934. Subgenus <i>Pseudorajendra</i> Dubatolov, 2007 Type Species: <i>Aloa dentata</i> Walker, 1855.	<i>Nannoarctia (Pseudorajendra) dentata</i> (Walker, 1855)	Canara [India].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1593).	India: Eastern India, Karnataka, Maharashtra (Mumbai, Pune), Kerala, Tamil Nadu (Dubatolov, 2010; Mitra <i>et al.</i> , 2019).	The larval host plant is unknown. This species is attracted to light. Remarks: This species is endemic to India.

Subfamily Aganainae Boisduval, 1833					
<i>Mecodina</i> Guenée, 1852 Type Species: <i>Mecodina lanceola</i> Guenée, 1852.	<i>Mecodina metagrapt</i> Hampson, 1926	Bali.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (Reg. No. ZSI-WRC-L-1808).	India: Karnataka, Maharashtra (Lonavala). Elsewhere: Bali, Singapore, Java, Sulawesi (Hampson, 1926; Holloway, 2005; Kalawate, 2018b).	The larval host plant is unknown. The species rarely attracted to the light.
<i>Asota</i> Hubner, [1819] Type Species: <i>Phalaena javana</i> Cramer, [1780] [= <i>Asota javana</i> (Cramer, [1780])].	<i>Asota producta</i> (Butler, 1875)	Not known.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1597).	India: Assam, Sikkim, South India, Maharashtra, Kerala. Elsewhere: Borneo, Burma, Malaysia, Penang, Sri Lanka, Sumatra, (Gurule, 2013; Sondhi <i>et al.</i> , 2018).	The larval host plants is unknown. The species is attracted to light and is recorded from the disturbed cultivated land.
	<i>Asota ficus</i> (Fabricius, 1775)	India.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1597).	India: Throughout India including Maharashtra. Elsewhere: China, Japan, Malaysia, Myanmar, Nepal, Sri Lanka, Taiwan, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Ricinus communis</i> (Euphorbiaceae); <i>Ficus carica</i> , <i>F. hispida</i> , <i>F. racemosa</i> , <i>F. pumila</i> , <i>F. infectoria</i> , <i>F. religiosa</i> , <i>Ficus</i> sp. (Moraceae); <i>Mitragyna diversifolia</i> (Rubiaceae) (ICAR-NBAIR, 2020). It is generally attracted to light trap.
<i>Digama</i> Moore, [1860] Type Species: <i>Digama hearseyana</i> Moore, 1858.	<i>Digama marchali</i> (Guérin-Méneville, 1843)	Not known.	15 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1598).	India: Gujarat, Maharashtra (Mumbai, Pune and Raigad), South India. Elsewhere: Myanmar (Mitra <i>et al.</i> , 2019).	The larval host plant is <i>Carissa carandus</i> (Apocynaceae) (NHM, 2021). The flight period is from January to March and June to November. It is attracted to light.
<i>Psimada</i> Walker, 1858 Type Species: <i>Psimada quadripennis</i> Walker, 1858.	<i>Psimada quadripennis</i> Walker, 1858	Canara.	03 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1599).	India: Andaman and Nicobar Islands, Chhattisgarh, Himachal Pradesh, Madhya Pradesh, Karnataka, Kerala, Maharashtra (Pune, Nashik and Nandurbar). Elsewhere: China, Myanmar, Sri Lanka, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plant is <i>Ficus</i> (Moraceae) (Holloway, 2005). The adult of this species is attracted to the light. They are recorded in forest and cultivated lands.
Subfamily Tinoliinae Moore, [1885]					
<i>Calesia</i> Guenée, 1852 Type Species: <i>Calesia comosa</i> Guenée, 1852 [= <i>Calesia dasypterus</i> (Kollar, 1844)].	<i>Calesia stillifera</i> Felder & Rogenhofer, 1874	Manila.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1812).	India: Throughout India including Maharashtra (Satara, Nashik). Elsewhere: Sri Lanka, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Thunbergia</i> sp., <i>Neuracanthus phaeostachys</i> (Acanthaceae) (Anonymous, 2021a). It is recorded from July to November.
	<i>Calesia fuscicorpus</i> Hampson, 1891	Nilgiri.	08 ex., Lonavala, Pune, 23.vii.2017, A.S. Kalawate & party (ZSI-WRC-L-1801).	India: Nilgiris (Tamil Nadu), Travancore (Kerala). Elsewhere: Ceylon (Sri Lanka) (Hampson, 1894).	The larval host plant is <i>Justicia wynaadensis</i> (Acanthaceae) (NHM, 2021). Not much known about its habitat. Remark: New distributional and Range extension in Northern Western Ghats' Maharashtra.
Subfamily Hypeninae Herrich-Schäffer, [1851]					
<i>Dichromia</i> Guenée, 1854 Type Species: <i>Phalaena orosia</i> Cramer, 1780 [= <i>Dichromia sagitta</i> (Fabricius, 1775)].	<i>Dichromia pullata</i> Moore, 1885	Ceylon Lanka].	07 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1600).	India: Maharashtra (Pune, Sindhudurg, Nashik, Dhule). Elsewhere: Sri Lanka (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Anacardium occidentale</i> (Anacardiaceae); <i>Tylophora</i> , <i>Dregea</i> (Apocynaceae) (Swafvan and Sureshan, 2021).

Subfamily Scoliopteryginae Herrich-schaffer, [1851]					
<i>Rusicada</i> Walker, 1858 Type Species: <i>Rusicada nigratarsis</i> Walker, 1858.	<i>Rusicada fulvida</i> (Anomis) (Guenée, 1852)	Java	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1601).	India: Throughout India including Maharashtra (Nashik, Dhule, Nandurbar). Elsewhere: Africa, Australia, Sri Lanka. (Mitra <i>et al.</i> , 2019)	The larval host plants are <i>Abutilon</i> , <i>Alcea</i> , <i>Gossypium</i> , <i>Hibiscus</i> , <i>Kydia</i> , <i>Pterospermum</i> , <i>Sterculia</i> , <i>Thespesia</i> Urena, <i>Waltheria</i> (Malvaceae); <i>Cissampelos</i> (Menispermaceae) (Anonymous, 2021a). Adult moth appears rarely to light traps.
Subfamily Calpinae Boisduval, 1840					
<i>Eudocima</i> Billberg, 1820 Type Species: <i>Phalaena salaminia</i> Cramer, 1777.	<i>Eudocima phalonia</i> (Linnaeus, 1763)	Africa.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1602).	India: Karnataka, Madhya Pradesh, Maharashtra (Mumbai, Pune, Satara, Sindhudurg, Nashik, Dhule, Jalgaon, Nandurbar), Punjab, Tamil Nadu, Uttar Pradesh, Uttarakhand. Elsewhere: Australia, China, Indonesia, Japan, Korea, New Guinea, New Zealand (Mitra <i>et al.</i> , 2019).	Polyphagous: The larval host plant are <i>Leschenaultia</i> (Goodeniaceae); <i>Anamirta</i> , <i>Arcangelisia</i> , <i>Cissampelos</i> , <i>Cocculus</i> , <i>Coscinium</i> , <i>Cyclea</i> , <i>Diploclesia</i> , <i>Legnephora</i> , <i>Sinomedium</i> , <i>Stephania</i> , <i>Tiliacora</i> , <i>Tinomisium</i> , <i>Tinospora</i> (Menispermaceae); <i>Theobroma</i> (Sterculiaceae); <i>Erythrina</i> (Leguminosae) (Holloway, 2005) preferred for egg laying (Leong and Kueh, 2011). Adult sucks fruit juice from ripe or ripening fruit and is a major pest on <i>Citrus</i> sp. (Rutaceae) (Leong and Kueh, 2011); <i>Punica granatum</i> (Lythraceae) (Jayanthi <i>et al.</i> , 2015); <i>Lycopersicon</i> (Solanaceae); <i>Malus pumila</i> (Rosaceae); <i>Mangifera indica</i> (Anacardiaceae) (Bhumannavar, and Viraktamath, 2000). This species is attracted to light.
Subfamily Lymantriinae Hampson, 1893					
<i>Nygmia</i> Hübner, [1820] Type species: <i>Phalaena icilia</i> Stoll, 1790.	<i>Nygmia icilia</i> (Stoll, [1790])	Coromandel coast.	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1817).	India: Jharkhand, Karnataka, Kerala, Maharashtra (Pune, Mumbai), NW. Himalaya, S. India, Tamil Nadu. Elsewhere: Sri Lanka (Cotes and Swinhoe, 1887; Sambath, 2014).	The larval host plants are <i>Dendrophthoe glabrescens</i> , <i>Loranthus</i> (Loranthaceae); <i>Mallotus paniculatus</i> (Euphorbiaceae) (NHM, 2021). In the present study it was recorded in August.
Subfamily Hypocalinae Guenée, 1852					
<i>Hypocala</i> Guenée, 1852 Type Species: <i>Hyblaea deflorata</i> Fabricius, 1794.	<i>Hypocala deflorata</i> (Fabricius, 1794)	India Orientalis [India].	01 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1794).	India: Himachal Pradesh, Karnataka, Maharashtra (Sindhudurg, Nashik, Dhule, Jalgaon, Nandurbar), North West Himalayas, Tamil Nadu. Elsewhere: Africa, Australia, China, Hong Kong, Indonesia, Japan, Korea, Madagascar, Malaysia, Nepal, New Zealand, Sri Lanka, Taiwan, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plants are <i>Diospyros dichrophylla</i> (Ebenaceae); <i>Pouteria sapota</i> (Sapotaceae) (Holloway, 2005). The adult moth sucks fruit juice making it unfit for consumption.
Erebidae incertae sedis					
<i>Chrysopera</i> Hampson, 1894 Type Species: <i>Achaea combinans</i> Walker, 1858 [= <i>Chrysopera combinans</i> (Walker, 1858)].	<i>Chrysopera combinans</i> (Walker, 1858)	Ceylon Lanka). (Sri	15 ex., Lonavala, Pune, 23.vii.2017, coll. A.S. Kalawate & party (ZSI-WRC-L-1575).	India: Maharashtra (Mumbai, Pune, Satara, Nashik, Dhule), North Western Himalayas, Peninsular India. Elsewhere: Australia, China, Indonesia, Malaysia, Nepal, Sri Lanka, Thailand. Elsewhere: Australia, China, Indonesia, Malaysia, Nepal, Sri Lanka, Thailand (Mitra <i>et al.</i> , 2019).	The larval host plant is not known. In the present study it is recorded from 626m elevation and as per Holloway (2005) the highest report of this species is from 500m.

Erebinae emerged as the dominant subfamily with 39% (Fig. 3) diversity in the present survey and this finding is in consistent with Farooqui *et al.*, (2020). The subfamily Arctiinae resulted as the second dominant with 30% diversity and this is in consistent with the reports of Shivaperuman (2014) and Gurule & Nikam (2013). Recently, Arctiidae family has been incorporated in the Erebidae as the subfamily due to the results of phylogenetic study. Noctuoidea is the highly unstable superfamily hence, studies on this group is warrant to resolve the instability in this economic important group. Noctuoidea is a cosmopolitan superfamily and the highest diversity of these moths is in Oriental tropics. In the present study an attempt has been made to document the diversity of Erebid moth fauna from Lonavala, which comes under an important Biodiversity Hotspots i.e. Western Ghats, a UNESCO world heritage site (Anonymous, 2021b). Diversity study plays a very important role in decision making for planning conservation and management actions. The present study may be helpful to the decision-making authority for making conservation and management plans.

Conclusions

The study resulted in identification of 44 species placed in 36 genera of Erebid moths. One new distributional and range extension record of *Calesia fuscicorpus* Hampson, 1891 has been recorded. To exactly predict the diversity of this important eco-region and other parts in the Sahyadri ranges, more extensive surveys are warranted for all the families of the moth.

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Conflict of Interest: The authors declare that they have no conflict of interest.

Declaration

"We declare that the manuscript has not been published in any journal/book or proceedings or in any other publication, or offered for publication elsewhere in substantially the same or abbreviated form, either in print or electronically.



Fig. 1 Light trap for moth study.

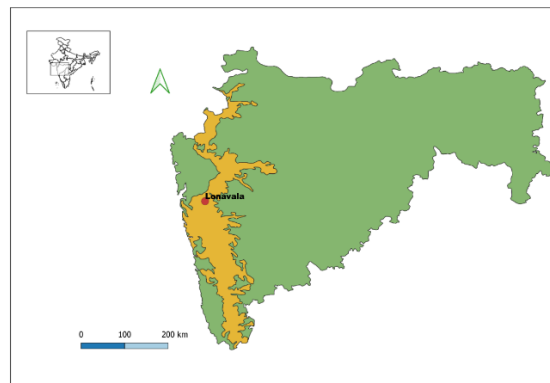


Fig. 2 Map showing survey locality.

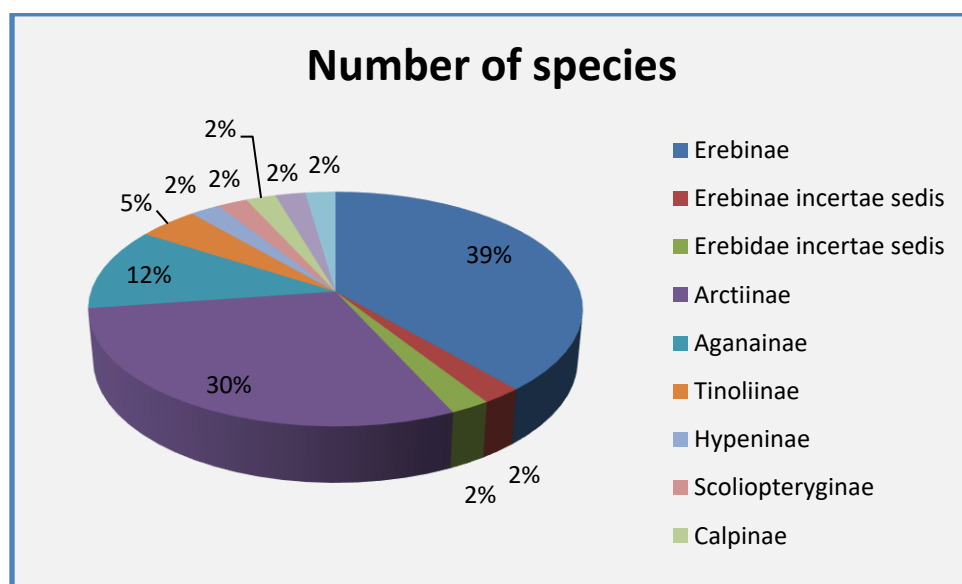


Fig. 3 Number of species recorded from the subfamilies of Erebidae from Lonavala, Maharashtra.

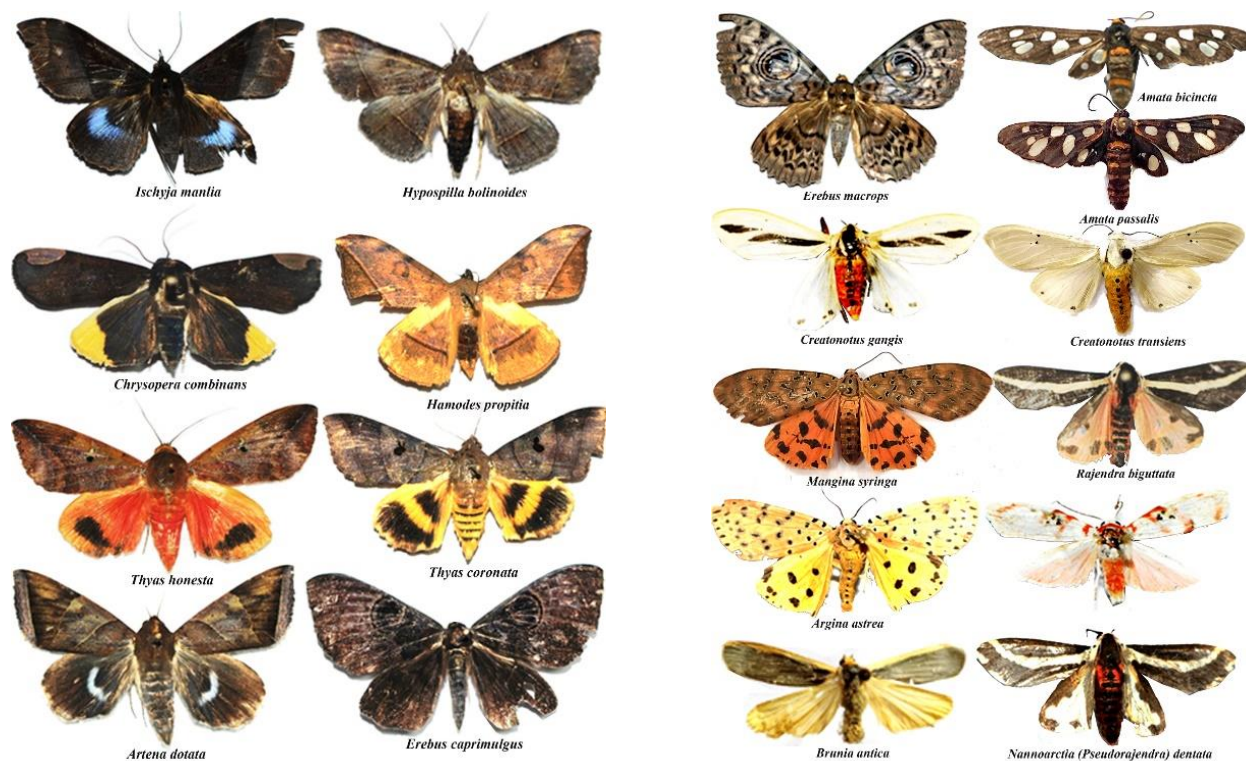


Fig. 4 Some erebid moths from the studied area.



Fig. 5 Some erebid moths from the studied area.

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Diversity of ants (Hymenoptera: Formicidae) in the mangrove patches of reclaimed Sunderbans, West Bengal, India

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Abstract

Ants are highly diverse social insects distributed in every terrestrial habitat except the north and south poles. Even though they are a dominant group in the mangrove ecosystem the study of ant diversity in Sunderbans mangrove, the world's largest mangrove ecosystem is yet to start. The objective of this study was to understand the ant diversity in mangrove patches in the villages of Indian Sunderbans. Ants were collected from the reclaimed areas of Indian Sunderbans. Total of 35 species, 21 genera of ants belonging to 5 subfamilies were found in this study. The highest number of species was found in the subfamily Myrmicinae. *Dilobocondyla gasteroreticulatus* reported first time from West Bengal. Ant species *Paratrechina longicornis*, *Crematogaster spp.*, *Monomorium spp*, *Camponotus spp.* and *Tetraponera rufonigra* were the most abundant species in the mangrove habitat. Most of the ants were found from the mangrove associate plant, *Excoecaria agallocha*. The ant species diversity index in this study is moderate ($H' = 2.37$), while the evenness index classified as moderate ($E = 0.5$), and no species of ants dominate this area ($D = 0.1$).

Key words: ants, mangrove, biodiversity, Sunderbans

Introduction

Ants are most impressive group of social insects. They belong to the order Hymenoptera and class Insecta and are placed in a single family of Formicidae. They are abundant in most of the habitats and trophic levels in several terrestrial ecosystems. Ants play important roles in terrestrial ecosystem as predators, detritivores, mutualists, herbivores and sometime pollinators (Holldobler and Wilson, 1990). Many species of ants contribute

to pest suppression in agricultural systems (Way and Khoo, 1992). Although there are 17 subfamilies, about 344 genera and about 14,150 described species of ants in the world (www.antwiki.org).

Sunderbans is the active deltaic complex of Ganga- Brahmaputra estuarine system which covers an area of approximately 10,000 sq. km. of which 62% lies within Bangladesh and 38% in India (ref). In India, the land area measures about 9,629 sq. km of

which 4,493 sq. km. is inhabited by people and rest is reserve forest. During high tide vast areas are inundated with brackish water, covered by halophytic herbs, shrubs and trees called mangroves. Mangrove swamps occur on the intertidal mudflats of estuaries, creeks and inlets. Mangrove habitats are among the most productive and biologically diverse wetland ecosystem on earth. In comparison to Bangladesh, Indian portion of Sunderbans has a poor formation of mangrove due to higher salinity and human interference (Naskar, Guha & Bakshi, 1987). In spite of these problems, Sunderbans in West Bengal still possess 34 true mangrove plants and 62 mangrove associate plants (Mandal & Nandi, 1989). The most common floras are *Avicennia alba*, *A. officinalis*, *Bruguiera gymnorhiza*, *Ceriops decandra*, *Excoecaria agallocha*, *Rhizophora apiculata*, *Sonneratia apetala*. Beyond this, Sunderbans have rich aquatic and terrestrial faunal diversity. More than 40 species of mammals, 163 species of birds, 56 species of reptiles, 165 species of fish, 23 species of molluscs, 15 species of prawns, 67 species of

crabs have so far been reported in Sunderban Biosphere Reserve (www.UNESCO.org).

Ants are often regarded as the most abundant and influential insect group in mangroves (Hogarth, 2007). It is reported that they play an important role in predation of insect pest on *Sonneratia*, *Rhizophora* and *Bruguiera* (Dakir, 2009). Ants and their pheromone could deter herbivorous insects and can reduce crab herbivory (Offenberg, 2006). Many studies of ants from India have been conducted previously except ant diversity in mangrove ecosystem of Sunderbans. By this study a baseline data is generated on ant biodiversity in Sunderbans mangroves which will bring out more of its associations and activities in further studies.

Material and Methods

Sampling sites

Sampling sites selected from inhabited areas of eastern, central and western parts of Indian Sunderbans are given below (Fig.1):

Location	Block	District	Coordinates
Sahebkhali	Hingalganj	24 Parganas North	22°20'4.000"N/88°58'18.000"E
Dayapur	Gosaba	24 Parganas South	22°7'44.040"N/88°50'24"E
Bhagabatpur	Patharpratima	24 Parganas South	21°43'28.39"N/88°18'35.54"E
Bakkhali	Namkhana	24 Parganas, South	21° 33'40.57"N/88°16'4.49"E
Radhakrisnapur	Sagar	24 Parganas, South	21°42'56 "N/88°03'30"E

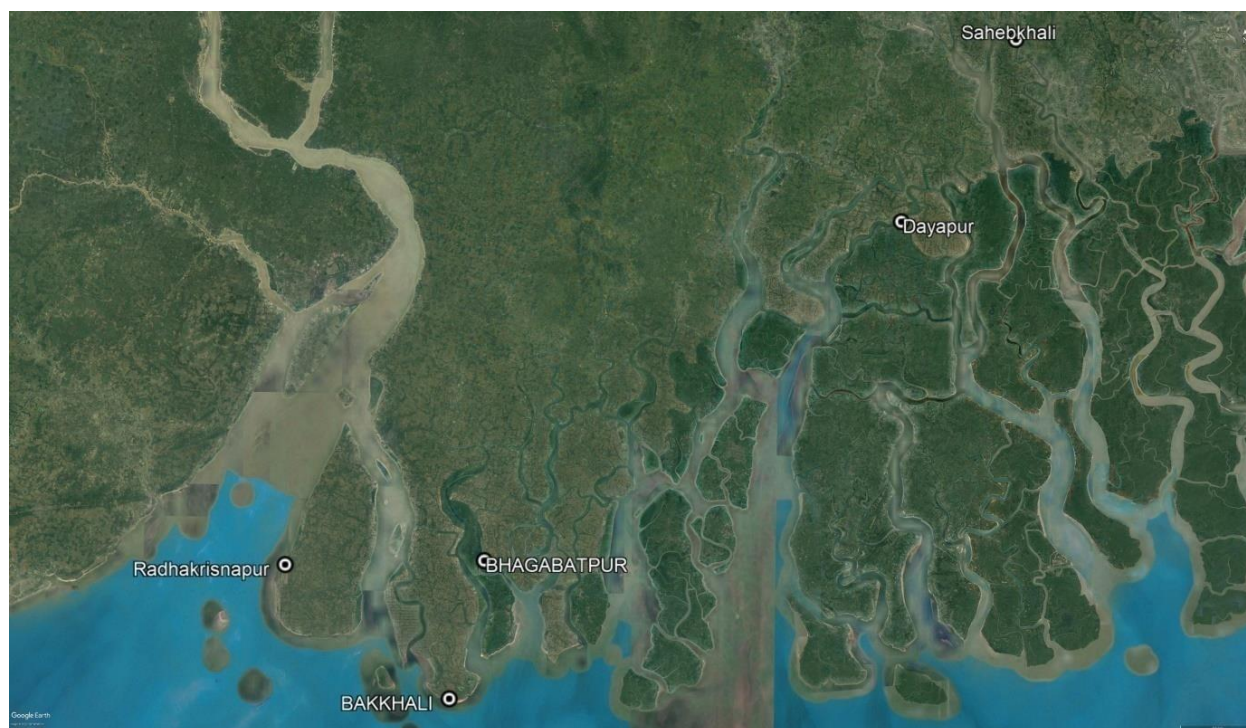


Fig. 1: A satellite image showing five locations of ant sampling in Indian Sunderbans

Methodology

Ants were collected directly by forceps and brush soaked in alcohol from stems, leaves and hollow branches of trees, from the soil, leaf litter and lower vegetation during pre-monsoon (April-June) and post-monsoon (October-November) during 2019-2020. The specimens were identified by referring to Bolton (1995), Bingham (1903), Tiwari (1998) and various keys available.

Results and discussion

A total of 35 species of ants were collected from mangrove patches of reclaimed areas of Indian Sunderbans belonging to 5 sub families, 21 genera and 752 individuals (Table1). The highest number of species were found in Myrmicinae (16 species, 46%),

followed by Formicinae (8 species, 23%), Ponerinae (11%) and Pseudomyrmicinae (11%) both had 4 species while Dolichoderinae had 3 species (9%) (Table 2 & Fig.2). *Monomorium* was most common genus with highest number of species collected. The genus *Tetraponera* is usually an arboreal species which represents two common species, *T. rufonigra* and *T. allaborans*. Here we found two more species *T. nitida* and *T. nigra* in which *T. nitida* is not a common species and reported in India, only from Andamans. The most abundant genus were *Paratrechina* (RA=18%), *Crematogaster* (RA=17%), *Monomorium* (RA=14%), *Camponotus* (11%) and *Tetraponera* (9%) (Table3). We identified *Dilobocondyla gasteroreticulatus* (the species is so named due to the presence of fine

reticulations on the gaster) from Patharpratima CD block of South 24 Parganas district on the stem of mangrove trees. *Dilobocondyla gasteroreticulatus* was first reported from Northwest Shivalik range of the Northwest Himalayas (Bharti & Kumar, 2013). We identified four tramp species named *Tetramorium simillimum*, *Paratrechina longicornis*, *Trichomyrmex destructor* and *Solenopsis geminata* from mangrove forest. Primarily these tramp species are closely associated with human and have a detrimental effect on the original inhabitants of ant species of mangroves. We also found that *Excoecaria agallocha*, a mangrove plant is most preferred microhabitat of ants (Table 4 and 5). It was observed that a total 18 genera of ants were using this plant for various purposes. Furthermore, *Camponotus compressus* lives exclusively in twigs of Acacia in a mutualistic relationship with tree hoppers (Fig. 3). *Crematogaster anthracina* found in a small nest cavities and hollow branches of Gum Arabic tree (Fig. 4). Dakir (2009) conducted a study on diversity of ants in mangrove forest of Kolaka, South-East Sulawesi and Muare Angke, Jakarta but he found 18 species. Arryanto *et al.* (2018) studied mangrove forest of North Kayong and recorded only 8 species. Further, Roy *et al.* (2018) reported 12 species from mangroves of Purba Medinipur district, West Bengal.

We calculated species diversity by using Shannon Diversity Index (H'), dominance by using Simpson's Dominance

Index (D) and Evenness by Shannon Evenness Index (E) (Pielou 1966; Shannon & Weaver 1963; Simpson 1949). We found that the ant species diversity index is moderate ($H' = 2.37$), while the evenness index classified as moderate ($E = 0.5$) and no species of ants dominate this area ($D = 0.1$). Therefore, the diversity index indicates that the mangrove areas are rich in biodiversity of ants and this will impact the floral and faunal diversity of mangrove ecosystem.

Conclusions

Mangroves are diverse and highly productive ecological communities at the land-sea interface. The paper deals with the Formicidae fauna collected from five different localities like Sagar Island, Bakkhali, Patharpratima, Gosaba and Hingalganj in Sunderban delta of West Bengal during 2019-2020. The study revealed that ants are very much diverse in mangrove habitat with highly diverse vegetation. They can adapt to extreme physiological condition during high tide and cyclonic storms. Consequently, it is evident that the biological resources rely one another to sustain a functioning environment.

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Table 1. List of ant species recorded from mangrove patches of Indian Sunderbans

Sub family	Genus	Species
Dolichoderinae	<i>Tapinoma</i>	<i>Tapinoma indicum</i> Forel
		<i>Tapinoma melanocephalum</i> Fabricius
	<i>Iridomyrmex</i>	<i>Iridomyrmex anceps</i> Roger
Formicinae	<i>Camponotus</i>	<i>Camponotus compressus</i> Fabricius
		<i>Camponotus sericeus</i> Fabricius
	<i>Lepisiota</i>	<i>Lepisiota sericea</i> Forel
		<i>Lepisiota opaca</i> Forel
	<i>Nylanderia</i>	<i>Nylanderia indica</i> Forel
	<i>Oecophylla</i>	<i>Oecophylla smaragdina</i> Fabricius
	<i>Paratrechina</i>	<i>Paratrechina longicornis</i> Latreille
Myrmicinae	<i>Polyrachis</i>	<i>Polyrachis rastellata</i> Latreille
	<i>Crematogaster</i>	<i>Crematogaster anthracina</i> Smith
		<i>Crematogaster rogenhoferi</i> Mayr
		<i>Crematogaster aberrans</i> Forel
	<i>Carebara</i>	<i>Carebara affinis</i> Jerdon
	<i>Monomorium</i>	<i>Monomorium atomum</i> Forel
		<i>Monomorium indicum</i> Forel
		<i>Monomorium latinode</i> Mayr
		<i>Monomorium floricola</i> Jerdon
	<i>Trichomyrmex</i>	<i>Trichomyrmex destructor</i> Jerdon
		<i>Trichomyrmex scabriceps</i> Mayr
	<i>Pheidole</i>	<i>Pheidole watsoni</i> Forel
		<i>Pheidole sagei</i> Forel
		<i>Pheidole parva</i> Mayr
	<i>Solenopsis</i>	<i>Solenopsis geminata</i> Fabricius
	<i>Meranoplus</i>	<i>Meranoplus bicolor</i> Guerin-Meneville
	<i>Dilobocondyla</i>	<i>Dilobocondyla gasteroreticulatus</i> Bharti & Kumar
Ponerinae	<i>Anochetus</i>	<i>Anochetus madaraszi</i> Mayr
	<i>Diacamma</i>	<i>Diacamma rugosum</i> Le Guilou
	<i>Leptogenys</i>	<i>Leptogenys hystérica</i> Forel
	<i>Pseudoneoponera</i>	<i>Pseudoneoponera rufipes</i> Jerdon
Pseudomyrmicinae	<i>Tetraponera</i>	<i>Tetraponera allaborans</i> Walker
		<i>Tetraponera nigra</i> Jerdon
		<i>Tetraponera nitida</i> Smith
		<i>Tetraponera rufonigra</i> Jerdon

Table 2. Percentage contribution of various subfamilies of mangrove ants in Sunderbans

Subfamily	Genus (%)	Species (%)
Dolichoderinae	2(10%)	3(9%)
Formicinae	6(29%)	8(23%)
Myrmicinae	8(38%)	16(46%)
Ponerinae	4(19%)	4(11%)
Pseudomyrmicinae	1(5%)	4(11%)
Total	21	35

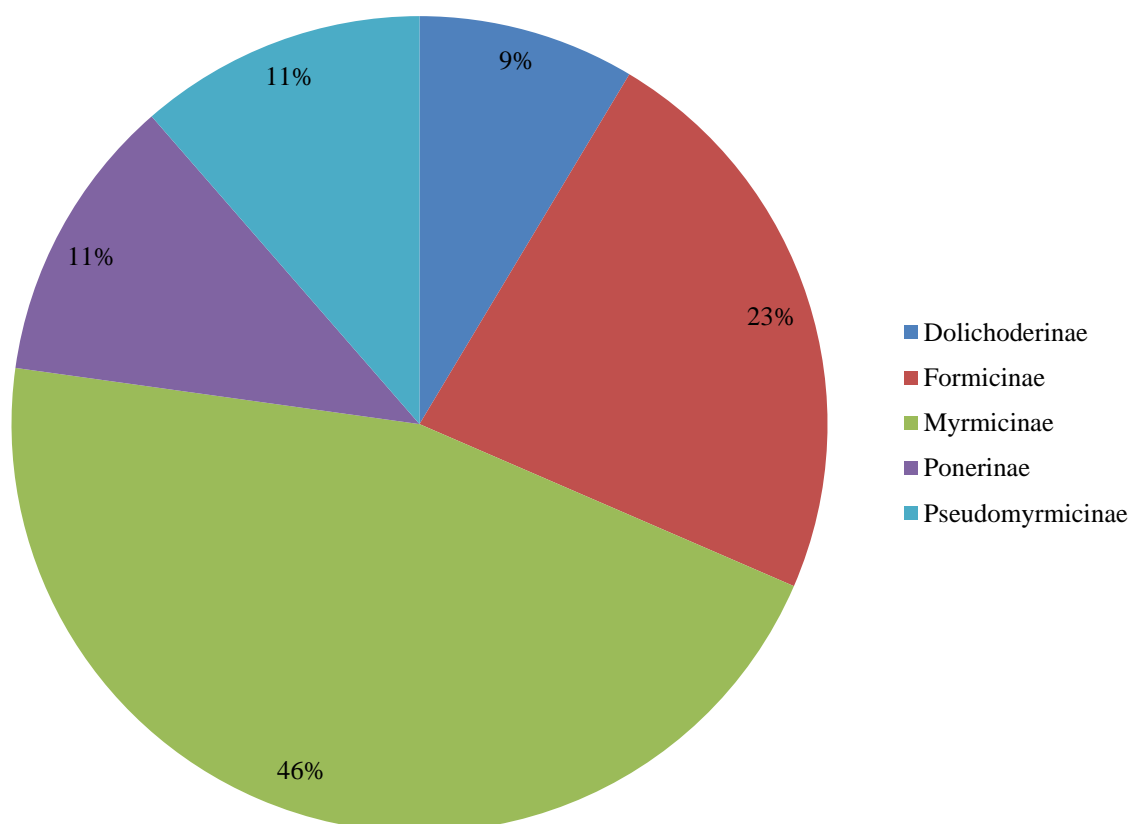
Fig. 2: Subfamilies of mangrove Ants found in reclaimed Sunderbans, India

Table 3: Calculation of Relative abundance (RA), Simpson Dominance Index (H'), Shannon Diversity Index (D) and Evenness Index(E) of ants in Sunderbans mangrove

Genus	Relative Abundance	Simpson Dominance Index(H')	Shannon Diversity Index(D)	Shannon Evenness Index (E)
<i>Paratrechina</i>	18.21809	0.116395	2.37236	0.5106
<i>Crematogaster</i>	17.42021			
<i>Monomorium</i>	13.56383			
<i>Camponotus</i>	11.03723			
<i>Tetraponera</i>	8.510638			
<i>Pheidole</i>	7.845745			
<i>Lepisiota</i>	6.117021			
<i>Nylanderia</i>	4.654255			
<i>Tapinoma</i>	4.255319			
<i>Diacamma</i>	2.792553			
<i>Pseudoneoponera</i>	1.06383			
<i>Solenopsis</i>	0.930851			
<i>Trichomyrmex</i>	0.930851			
<i>Leptogenys</i>	0.531915			
<i>Carebara</i>	0.398936			
<i>Meranoplus</i>	0.398936			
<i>Anochetus</i>	0.265957			
<i>Calalaucus</i>	0.265957			
<i>Dilobocondyla</i>	0.265957			
<i>Iridomyrmex</i>	0.265957			
<i>Tetramorium</i>	0.265957			

Table 4: Preferred host plants of ants in Sunderbans mangrove

ANT GENUS \ PLANT	Gewa (<i>Excoecaria</i> sp.)	Baen (<i>Avicennia</i> sp.)	Kankra (<i>Bruguiera</i> sp.)	Gum Arabic tree (<i>Vachellia</i> sp.)	<i>Casuarina</i>	Ear leaf acacia (<i>Acacia</i> sp.)
<i>Anochetus</i>	P	-	-	-	-	-
<i>Catalacus</i>	P	-	-	-	-	-
<i>Camponotus</i>	P	P	P	P	P	P
<i>Carebara</i>	P	-	-	-	-	-
<i>Cardiocondyla</i>	P	-	-	P	-	-
<i>Crematogaster</i>	P	P	-	P	P	-
<i>Diacamma</i>	P	-	P	-	-	-
<i>Dilobocondyla</i>	P	-	-	-	-	-
<i>Iridomyrmex</i>	P	-	-	-	P	-
<i>Lepisiota</i>	P	P	-	P	-	P
<i>Leptogenys</i>	P	-	-	-	-	-
<i>Monomorium</i>	P	P	-	P	P	P
<i>Meranoplus</i>	-	-	-	P	-	-
<i>Nylanderia</i>	P	-	-	P	P	-
<i>Paratrechina</i>	P	P	-	-	P	P
<i>Pheidole</i>	P	P	-	P	P	P
<i>Pseudoneoponera</i>	P	-	-	P	P	-
<i>Solenopsis</i>	-	P	-	-	-	-
<i>Tapinoma</i>	P	P	-	P	P	P
<i>Tetramorium</i>	-	-	-	P	-	-
<i>Tetraoponera</i>	P	P	-	P	P	-
<i>Trichomyrmex</i>	P	P	-	-	-	-

Table 5: Species richness of ants in different pants in Sunderbans mangrove

Microhabitat	Species richness
Gewa tree (<i>Excoecaria agallocha</i>)	18
Gum Arabic tree	12
Casuarina tree	10
Baen tree (<i>Avicennia</i> sp.)	10
Ear Leaf Acacia tree	6



Fig. 3: *Camponotus compressus* feeding on the excretion of tree hoppers i at Bakkhali, Sunderbans



Fig. 4: *Crematogaster antracina* nest in cavity of gum Arabic tree at Patharpratima, Sunderbans

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Incidence of *Oxycetonia versicolor* Fabricius on sunflower from West Bengal, India – A new report**Sabyasachi Ray¹ and A. Banerjee²**¹ *Department of Agricultural Entomology, Bidhan Chandra Krishi Viswavidyalaya, Mohanpur, Nadia-741252, W.B., India*² *AICRP on MULLaRP (Mungbean, Urdbean, Lentil, Lathyrus, Rajmash & Pea) Directorate of Research, B.C.K.V., Mohanpur, Nadia-741252, W.B., India****Corresponding author: amitavakvk@gmail.com***

Sunflower, botanically known as *Helianthus annuus* and representing the family Asteraceae (formerly known as Compositae), is a crucial oil yielding crop that originated in north America. Seeds are the main economic portion of the crop that is processed for several products and commercialization of these products are executed in cooking purpose as well as feed supply for the livestock animals (Yegorov *et al.*, 2019). Significant health benefits are provided by the crop through the supply of premium oil and dietary fibre (Khan *et al.*, 2015). Seed of the crop contains unsaturated fatty acids like palmitic acid (5%), stearic acid (6%), monounsaturated fatty acid like oleic acid (30%) and polyunsaturated fatty acid like linoleic acid (59%) (Anon., 2005). As per the scenario, worldwide sunflower oil production reached up to 18 million tonnes. Ukraine and Russia are the leading sunflower oil producing country among the world accounting for 53% of the total global production. In the Indian context it was observed that total production of sunflower seed during the fiscal year 2021 was 185 thousand metric tonnes (Anon, 2022). In West Bengal during 2016-2017 the production of

sunflower seed was 0.02 million tonnes from an area of 0.01 million ha where as during 2017-2018 the sunflower growing area in West Bengal remains almost same but production level reduces considerably and became 0.01 million tonnes (Anon., 2018). During the past years sunflower production decreased due to several biotic and abiotic constraints in India as well as in West Bengal. Previous investigation revealed that sunflower has been attacked by 43 insect species in India (Sandhu *et al.*, 1973; Makhdoomi *et al.*, 1984). Hassan *et al.* (1984) reported that about 19 insect pests infested sunflower crop and among them *Helicoverpa armigera* Hubn., *Aphis gossypii* Glover., *Bemisia tabaci* Genn., *Amrasca devastans* (Ishi.) and *Atractomorpha crenulata* (Fab.) causes economic damage to the crop (Satter *et al.*, 1984). Climate change over the years may bring changes in the pest and natural enemy complex of crop ecosystem including sunflower in different growing regions (Geetha and Hegde, 2018). Besides this mono-cropping with intensive cultivation using high yielding varieties, using of higher external inputs including fertilizers and pesticides may lead to the change in pest incidence on a particular

crop. Under the changed circumstances some pest which were previously not known for causing economic damage to the concerned crop are becoming a matter of concern as well as some new insects also are appearing on a particular crop of a concerned region which have no previous history of association with that crop of that particular region (Taggar *et al.*, 2012). One such insect is flower chafer beetle, *Oxycetonia versicolor* Fabricius (Scarabaeidae: Coleoptera).

The flower chafer beetle (*O. versicolor*) feeds on floral parts of several crops causing considerable damage as the reproductive parts gets damaged. It is belonging to the family Scarabaeidae and subfamily Cetoniinae under the order Coleoptera. The insect is also known as *Cetonia versicolor* and *Gametis versicolor* Fabricius, 1775. The existence of this insect in the Seychelles was first noticed by Fairmaire (1893) who reported it as *Glycyphana versicolor*. Dupont in 1917 noticed the infestation of this insect in the rose bushes during day (Matot, 2000). Afterwards several provinces and regions of the world like Samoa, China, India, Sri Lanka, Chagos, Madagascar, Mascarenes and South Africa reported this insect (Matot, 2000). In the Indian subcontinent the earliest incidence of this insect was recorded from Indore district of Madhya Pradesh (Arrow, 1910).

The occurrence of these beetles on sunflower cv. KBSH 1 was first noticed in the

farmer's field located at Bhayna (23.3110° N, 88.6386° E) under the district Nadia, West Bengal (Fig. 1). Incidence of the adult beetles was observed during last week of March, 2021 when the crop was at fifty per cent flowering stage coinciding with the peak flowering from March to April. The infestation was low to moderate (2-3 beetles/ flower) and their occurrence on the flowers was recorded after 11 am in the day time. Feeding of the adult beetles was noticed on the floral head (Fig. 2) from where they fed on the petals, pollens (Fig. 3), nectars and ovaries. Feeding of those floral parts causes substantial feeding injury and chaff in the grain. Association of this insect with sunflower is not previously reported from West Bengal. This is the first observation on the occurrence of chafer beetle (*O. versicolor*) on sunflower crop from West Bengal and also from Eastern India as per the available literature.

The chafer beetles are mainly known for infesting the floral parts particularly they feed on the pollen grains of several crop flowers and most of their activity takes place during daytime. Immature stages of these beetles generally harbored the decomposed soil organic materials and few of them infest plant roots (Taggar *et al.*, 2012). The mouthparts of the adult chafer beetles lack enough potential to penetrate resistant and sclerified plant tissues as they are less sclerotized. For that reason the main damage is caused by the dentate tibiae rather than the mouthparts as they use the tibiae for making

slits in the floral parts which results in the destruction of the flower. This situation happens frequently when the flower inhabits more population causing competition in between the individuals (Viggiani, 1926; Tremblay, 2000).

There are several reports about the insect infesting different crops having economic importance. The insect was found to be distributed among different parts of South India and some northern parts also. In these regions several crops like sorghum, maize, ragi, ground nut were reported to be infested by this insect (Fletcher, 1914). Previous investigation suggested that the adult chafer beetle not only damaged the flowers but also the leaves and shoots of groundnut and okra whereas grubs were the root feeders but no considerable damage was caused by them (Fletcher, 1917). Grubs generally get nourished from grass compost, sawdust and organic matter (Hinckley, 1967). Bhatnagar (1970) discovered the existence of the insect as a pest in different crops like citrus, cotton, okra, sorghum, maize, and groundnut cultivated in some parts of Rajasthan. Different red gram growing parts of India were infested with this beetle where they mainly fed on the pollen grains (Reed *et al.*, 1989). Voracious feeding by these insects was observed from pearl millet panicles in Rajasthan (Nwanze and Youm, 1995). From Tamil Nadu these chafer beetles were noticed to inflict damage the brinjal crop by devouring the tender shoots, floral buds and flowers (Ambethgar, 2000).

The insects were also known for causing injury to *Jatropha curcas* in mild level (Shanker and Dhyani, 2006). Incidence of chafer beetle (2.08%) was also recorded from Doddaballapur, Lalbagh (Bangalore) where they damaged the rose flowers and leaves also; preferably they fed on the flowers (Kumar *et al.*, 2009). The pest was also recorded from green gram at flowering and pod developing stage of the crop (Duraimurugan and Srinivasan, 2009). Incidence of this beetle was also reported from red gram and green gram from Punjab (Taggar *et al.*, 2012). Recently cotton growing regions of central India became infested with this insect which caused low (0.40%) to moderate (12.92%) damage in Rajkot (Gujarat) and Wardha (Maharashtra), respectively (Naik *et al.*, 2017). Daravath *et al.* (2020) recorded the incidence of flower chafer beetle from Telengana and Southern India, where they found it mainly damaging the cotton. Besides they also noticed some other host plants of this beetle viz. brinjal, sesamum, maize, sorghum and *Parthenium hysterophorus* from that region.

It may be concluded that the incidence of flower chafer beetle, *Oxycetonia versicolor* on sunflower as documented above is the first-time report from West Bengal in particular and from Eastern India as a whole as per the existing scientific literatures. In future, further investigations will be needed for assessing the probable population build up of this insect with a pest risk analysis. A comprehensive knowledge on the nature of damages produced

by the insect along with its bionomics and way of sustainable management if required should be studied.

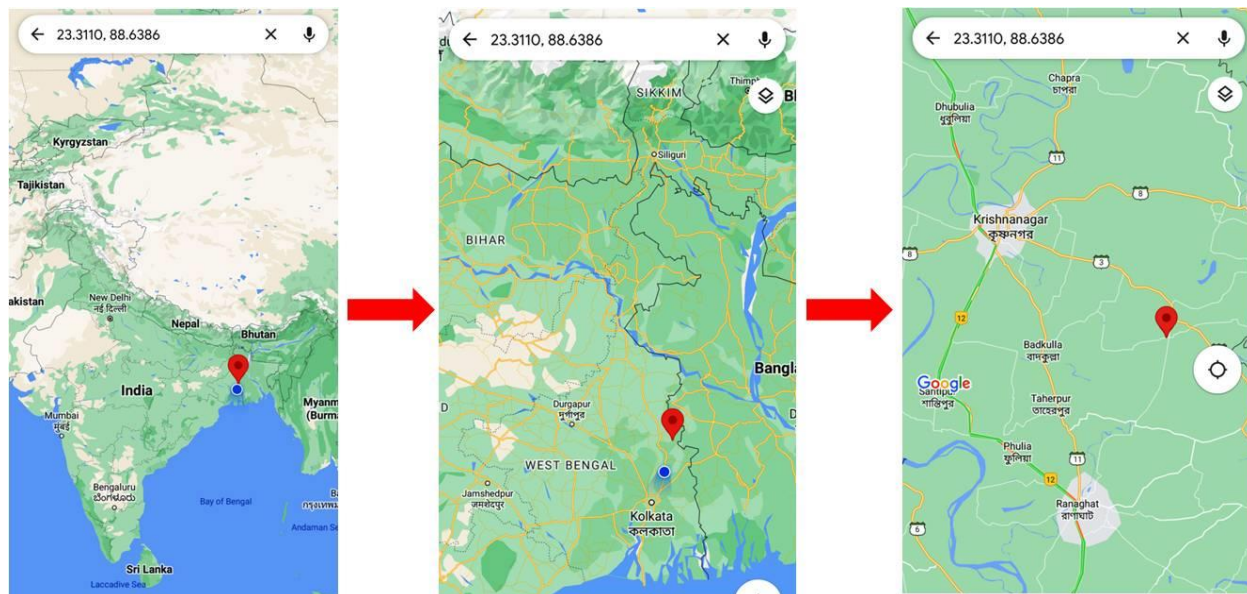


Fig. 1: Place of record as per GPS



Fig. 2: Incidence of flower chafer beetle (*Oxycetonia versicolor* Fab.) on the flower head of sunflower

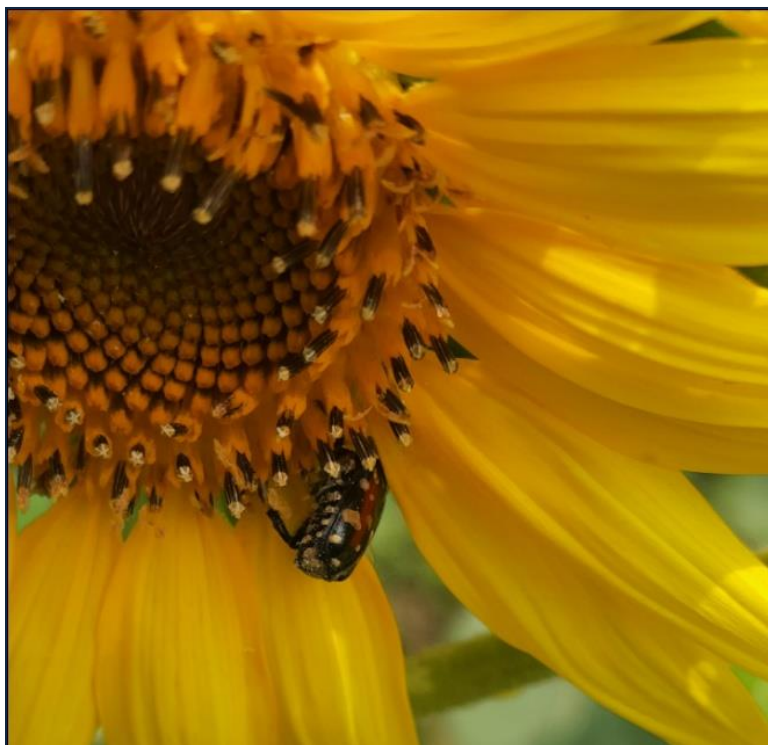


Fig. 3: Flower chafer beetle (*Oxycetonia versicolor* Fab.) feeding on sunflower pollens

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Review articles & Short notes

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Hormesis- a weapon for improving insect natural enemies**Aishwarya Ray^{1,2} and Basana Gowda G^{2*}***1 Department of Entomology, College of Agriculture, Indira Gandhi Krishi Vishwavidyalaya, Raipur, Chattisgarh, 492001, India**2 Crop Protection Division, ICAR-National Rice Research Institute, Cuttack, Odisha, 753006, India****Corresponding author e-mail: basanagowda.g@icar.gov.in**

Constant pressure to feed the world's growing population has led to development of high-yielding varieties which caused crops to be stressed by various pests (Singh *et al.*, 2014). Although the call for using integrated pest management (IPM) is very sound and clear, still chemical management has been the first preference. This has led to excessive use of non-selective pesticides, thereby resulting in ecological backlash like natural enemies' destruction, resistance and resurgence of pests along with secondary pest outbreaks (Wang *et al.*, 2012; Gowda *et al.*, 2021a). Although pesticides are applied at sufficient concentrations to manage the target insect pests, certain spatio-temporal changes in concentrations due to abiotic and biotic factors lead to altered targeted doses thereby causing lethal as well as sublethal effects in the arthropods exposed (Desneux *et al.*, 2007, Ullah *et al.*, 2019). This biological phenomenon is called hormesis or dose response phenomenon. It is a biphasic dose-response to low doses of stress that can stimulate biological activities (Fig.1) (Cutler and Rix, 2015). This hormesis has been seen in

a wide range of organisms including insects (Calabrese, 2005; Cutler and Rix, 2015). In insects, it includes stimulation such as enhancement in longevity, fecundity at any life stage and with any pesticide active ingredient, thereby enhancing the growth of organisms (Cohen, 2006; Cutler, 2013; Ayyanath *et al.*, 2013; Guedes and Cutler, 2014). Because of various ecological backlash and a huge impact on IPM-deciding strategy, the study of insect hormesis has been the center of research for insecticide toxicology with relatively less importance on toxicological impacts in natural enemies. The insecticide-induced insect hormesis is very detrimental whereas for the natural enemies it is a puissant weapon in optimizing mass rearing and enhancing the quality of bio-agents. Due to the high value of bio-agents in IPM, hormetic study or sublethal effect has garnered more attention.

It is very evident that commercial mass rearing and management of beneficial insects is a multi-billion dollar exclusive industry. Insects reared for biological and medical research, sterile insect release program, bio-control and many others have paved way for

the money-making industry. The collaboration of rearing with the hormetic principles can elevate the mass culture programs of insects such as by improving insect longevity, fecundity and parasitization rate (Cutler, 2013). The urge to study the insecticide induced hormesis in natural enemies, especially on the predators, started with the lacewing, *Chrysopa californica*, coccinellid beetle and *Habrobracon hebetor* (Fleschner and Scriven, 1957; Atallah and Newson, 1966; Grosch and Vacovic, 1967; Guedes and Cutler, 2013; Gowda *et al.*, 2021b). The hormetic effects are more profound in insect predators than in parasitoids. For example, an increase in the reproductive outputs and reduction of the generation time of the predatory bug *Podisus distinctus* was seen with a single exposure to the sublethal dose of permethrin (Guedes *et al.*, 2009). A similar trend was observed with the predator *Supputius cincticeps* (Zanuncio *et al.*, 2005). In the case of parasitoids, *Trichogramma* which is an effective bioagent against various lepidopteran pests (Orr *et al.*, 2000; Cabello *et al.*, 2012; Chailleux *et al.*, 2013; Gontijo *et al.*, 2019; Nozad-Bonab *et al.*, 2021; Gowda *et al.*, 2021c), the knowhow about the potential of hormesis on the mass rearing and quality has now become a priority. For example, exposure to low lethal concentrations (LC₃₀) of chlorflurazuron and tebufenozide on *Trichogramma chilonis* Ishii increased the adult longevity and female fecundity (Wang *et al.*, 2012). Such advancement in rearing programs with

hormetic principles can elevate the development of natural enemies.

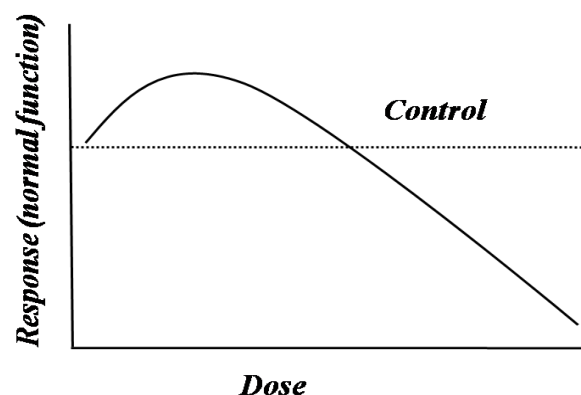


Fig. 1. Hormesis: A biphasic phenomenon (Guedes and Cutler, 2013)

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Immune priming: An overview in insects**Chaitali Banerjee***Assistant Professor in Zoology, Vidyasagar College for Women, 39, Sankar Ghosh Lane
Kolkata 700006, West Bengal, India**Corresponding author: chaitali.banerjeeibl@gmail.com***Abstract**

A characteristics feature of living systems is the ability to provide protection from different kinds of infection and various toxins. Importantly in insects, innate immune response is the most fascinating phenomenon. Though vertebrates' specific adaptive immune system is lacking in invertebrates including insects, defence priming is integral to many insect species. Immune priming is defined as the improved protection by the host to the same infectious agent upon a second encounter. This protection could be species/strain-specific, may be rendering lifetime protection or can be transgenerational. These attributes hence are effective protective system against a range of pathogens. Transgenerational immune priming can be attained primarily by two ways - direct parental transfer or elevated endogenous offspring immunity. The exact mechanisms for direct parental transfer is poorly understood though ways proposed involves direct transfer of antimicrobial peptides and mediators like lysozymes in the cytoplasm of the egg, epigenetic phenomenon like genomic imprinting or transfer of microbial fragments directly by the parents. Hence immune priming in insects are important line of defence and herein in this article an attempt on the elucidation of this phenomenon in the context of insects is done.

Keywords: Immune, Priming, Transgenerational**Introduction**

Class Insecta (Latin *insectum*, "notched or divided body", that implies "cut into") is the largest representative of Phylum Arthropoda, which in turn are the most predominant phyla (Snodgrass, 1960). They have 3 major divisions of the body – head comprising the mouthparts, eyes, and a pair of antennae; thorax (segmented) and abdomen that involves important systems like excretory and reproductive. They harbour very

prominent and primitive innate immune system which exhibit both cellular and humoral immune responses. The insect immune system is known to possess fat bodies that secrete effector molecules into the circulating fluid (hemolymph) and diverse range of hemocytes. The fat body (a substitute of vertebrate liver and adipose tissue) is reported to secrete several soluble factors like antimicrobial peptides into the hemolymph (Lemaitre and Hoffmann, 2007). Hemocytes are versatile in forms and function.

Phagocytosis, encapsulation, nodule formation are some of the important roles worth mentioning. The most prevalent cell types are adipohemocytes, oenocytoids, prohemocytes, spherule cells, granulocytes, secretory, plasmatocytes, crystal cells and lamellocytes (Rosales, 2017).

Immune priming is another important immune mechanism shown by several insect species. It is a defence strategy in which a prior encounter to the pathogen or any of its derived material leads to profound immune response upon further encounter. Hence this strategy protects the insect by rendering it resistant to subsequent infections. This process involves an increase in the density of circulating hemocytes and subsequent enhanced production of antimicrobial peptides (Sheehan *et al.*, 2020). Insects have always been an indispensable model to study microbial pathogen infection in human and hence detailed mechanisms of immune priming is integral to assess several parameters.

Immune priming – the detailed aspect

Immune priming is a fascinating aspect of invertebrate immunology which refers to swift, prompt and improved response of immune system to microbial population if they are exposed to the same pathogen previously at a relatively sub-lethal dose. Immune priming can result because of several factors like exposure to anti-microbial agents, different kinds of thermal and physical stress amongst others (Sheehan *et al.*, 2020). Immune priming

hence bestows insects to counteract potential infections with a history of previous exposure. The armamentarium involved in these process are different components of cellular and humoral immune system. Precisely immune priming is a complicated mechanisms and its effects may vary amongst insects of different species (Cooper and Eleftherianos, 2017).

Various experiments have been performed to demonstrate and understand immune priming. The classical biological model, *Drosophila melanogaster* have served as an integral model to study immune priming. The fruit flies were infected with sub-lethal dose of *Streptococcus pneumoniae* or its heat-killed form and a week later were re-exposed to the lethal dose of same pathogen (Pham *et al.*, 2007). It was observed that the flies successfully withstood the infection. Similar observation were also noted with *Beauveria bassiana*, a natural fly pathogen. In another set of experiments with the bumblebee, *Bombus terrestris* were exposed to gram-negative *Pseudomonas fluorescence* or closely related gram-positive bacteria (*Paenibacillus alvei* and *P. larvae*) followed by repeat exposure with either the same bacteria (homologous) or one of the two bacteria (heterologous). The results showed that primed bees significantly withstand the homologous repeat infection than a heterologous repeat infection (Sadd & Schmid-Hempel, 2006).

These and several other infection studies in insects model have suggested and

indicated the involvement of immune priming in insects. Further studies exploring the mechanism of priming have established the involvement of important signalling pathways like Toll and Imd in the phenomenon (Tanji *et al.*, 2007). Both of these pathways are highly conserved signalling paths and follows similar patterns. The immune deficiency pathway or Imd primarily regulates antimicrobial peptides' expression. The Imd pathway is similar to TNFR and TLR signalling pathways. Precisely TNF- α requires *NF- κ B* for the execution of functioning and Imd pathway relies on *Relish* signalling molecule. Relish is reported to

contain *Rel* homology domain (that explains the nomenclature *Relish*) and an IkappaB-like domain that has ankyrin repeats (6 in numbers). Thus, Relish is a dual domain protein. This explains similarity of Relish to mammalian NF-kappaB precursors - p100 and p105 (Silverman & Maniatis, 2007). The Figure 1 explains the mechanism executed by Imd and TNFR pathways respectively. The conserved signalling molecules have been highlighted by similar colour and shapes (Myllmaki *et al.*, 2014). It is worth mentioning here that *Relish* is typically involved in the humoral immune responses.

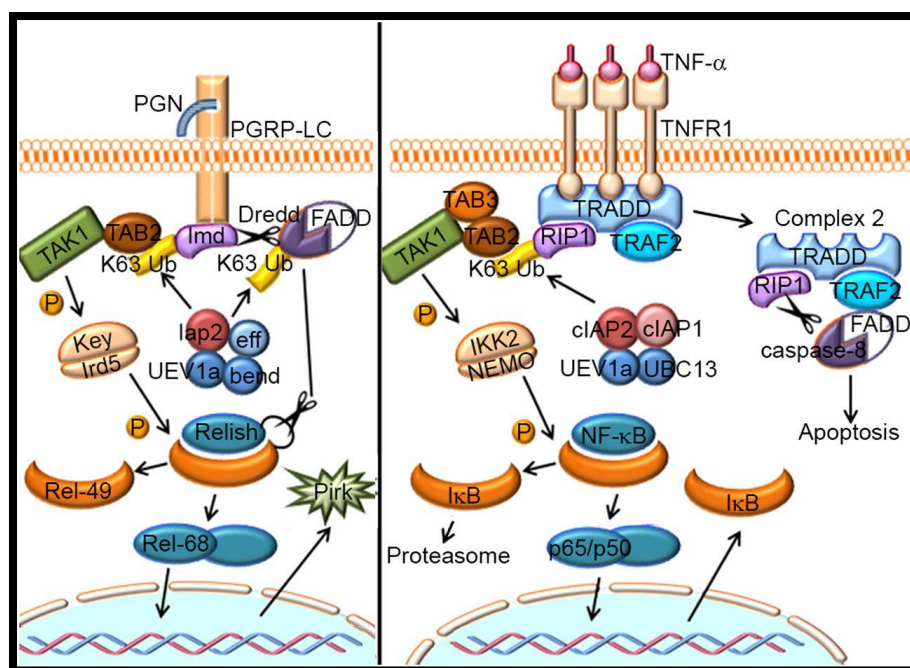


Figure 1: Schematic representation of the *Drosophila* Imd pathway and human TNFR signaling. Conserved components are indicated by similar shapes and colours (Myllmaki *et al.*, 2014).

Drosophila Toll pathway is primarily involved in cellular immune responses and act as important signalling

molecules during phagocytosis, parasite encircling and nodule formation, encapsulation etc. The pathogen sensing is done by molecules extracellularly placed that triggers proteolytic cascade involving hydrolysis of the Toll receptor like prospatzle to spatzle (Lemaitre *et al.*, 1996). The active spatzle binds to Toll receptor for the Toll pathway activation which in turn recruits a cascade of proteins consisting of MyD88, Tube and Pelle. As a consequence, there is a degradation of Cactus, eventually freeing Dorsal, and Dorsal-related immunity factor (DIF) facilitating their entry into nucleus to regulate the expression of AMPs such as drosomycin (Valanne *et al.*, 2011).

Besides immune priming, both these pathways in downstream are severely implicated in disease pathogenesis and are the major determinants of immune evasion or protection against microbial infection. They are important initiators of several cascades like apoptosis, autophagy, necrosis to name a few. However, the Imd pathway is characterized for sensing diaminopimelic acid-containing peptidoglycan (DAP-type PGN) prevalent in Gram-negative bacteria. As mentioned earlier, they mainly regulate the expression of anti-microbial proteins like Diptericin involving Relish, an important transcription factor (Kleino and Silverman, 2014). Toll pathway operates in a different way by sensing lysine-containing peptidoglycan (Lys-type PGN) found in Gram-positive bacteria and by β -glucans characteristic of fungal cell walls. As

a result they activate a different set of AMPs like Drosomycin that triggers downstream cascade *via* the involvement of dorsal-related immune factor (DIF), another important transcription factor.

Transgenerational immune priming

Transgenerational immune priming is an important aspect of insect immunology. Here the parental encounter to the pathogen provides protection to the progenies; hence suggesting immune protection persisting across generations. This dynamic phenomenon has been demonstrated by several representatives of different orders of class Insecta (Tetreau *et al.*, 2019). A Transgenerational mechanism involves transfer of specific signalling molecules through developing eggs. These molecules could be some bacterial peptides translocated from mother's gut to the egg. Often this signal translocation involves enhanced expression of immune-related genes in the egg. Specifically in case of social insects *Apis mellifera*, vitellogenin is greatly implicated in the process. Vitellogenin is a lipoglycoprotein (egg-yolk precursor). They are synthesized and stored in the fat body but are secreted into the hemolymph. Though they are female specific proteins, in honeybees they are also expressed in the workers. Vitellogenin categorically recognises Pathogen associated molecular pattern like LPS and PGN. This can provide immunization to the progenies through many pathways including social immunization. In this later instance, bacterial fragments from the

gut gets transferred to the worker bees' glands involved in producing the royal jelly that the queen bee feeds. This eventually reaches the eggs produced by the queen hence benefitting the colony (Salmella *et al.*, 2015). In insect models like *Galleria mellonella* and *Tribolium castaneum*, transfers of specific microbial proteins occur by crossing the midgut epithelium. It is then entrapped into nodules in the haemocoel which is followed by deposition in the eggs (Freitak *et al.*, 2014).

In insects there are different types of ovary reported - panoistic ovary, telotrophic ovary and polytrophic meroistic ovary. The later one is characterized by the arrangement of nurse cells and oocytes alternatively along the length of the ovariole whereas telotrophic ovary is characterized by placement of nurse cells at the apex of the ovariole. This facilitates feeding the oocyte through a nutritive cord as it descends down. In both these ovarioles, the trophocytes provides the oocyte with RNA, proteins, and ribosomes through much of their development that are otherwise provided only by the oocyte itself in panoistic ovarioles. Insects exhibiting telotrophic ovary (Orders - Hemiptera and Coleoptera) or polytrophic meroistic ovary (Orders - Hymenoptera, Lepidoptera, and Diptera) involve transferring maternal mRNAs in developing eggs during oogenesis. Many of these mRNAs apart from playing important role in development helps in the early immunological protection to the progenies (Johnstone and Lasko, 2001).

Another important transfer method involved is through epigenetic modifications of the parents like acetylation/deacetylation of histone proteins and/or by the methylation/demethylation of immune-related genes. Besides, females are capable of direct transfer of immune effector proteins to their eggs passively through the diffusion or sequestration into the proteins present in the mother's hemolymph. Also the transfer *via* active process takes place through specialized cells, such as nurse cells that mediates transfer to oocytes (Harwood *et al.*, 2019).

Thus these are the different ways of how the developing embryo within the egg gets primed to different microbial patterns from exposure that the parents encountered during their life.

Conclusion and future prospects

Immune response against microbial pathogen is integral to survival of the host. In this respect immune priming is important as it primes the defence machinery and promotes increased protection to further challenges. In insects immune priming has been found as a widespread phenomenon owing to its prevalence in several insect species. This is further elaborated in insects as an event within generation and that persists across generations as well. Transgenerational immunity is a dynamic phenomenon in invertebrates and signifies the memory like response of vertebrates. However, monitoring several successive generations is crucial to establish

the sustainability of transgenerational priming. Additionally, detailed mechanisms of immune priming in insects with special reference to transgenerational impact have to be studied. Further the studies has to be substantiated by molecular approaches. An elaborate transcriptomic and proteomic studies could decipher the intricate mechanisms. Several insect species shows metamorphosis of different extent. Hence it is crucial to investigate stage-wise analysis of transgenerational priming.

Equally important is the differentiation of hemocytes. The different sub-population of hemocytes keep circulating in the hemolymph that confers innate immune responses and as required differentiates to counteract the infection. Overall it can be concluded that immune priming is a conserved protective mechanisms though some more critical and elaborate studies may be required for more detailed understanding.

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Entomophagy: a step towards food security**Laya A. C. and Haseena Bhaskar**Department of Agricultural Entomology, College of Agriculture, Kerala Agricultural University,
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The world population surges ahead and is expected to reach nine billion by 2050 (FAO, 2012). Food security becomes a challenge due to growing population and limited resources, necessitating rethinking about food patterns and habits, particularly those relating to meat consumption. The word entomophagy is derived from the Greek word, *Entomon* means 'insect' and *Phagein* means 'to eat.' Thus, the practice of eating insects is known as entomophagy. Insects form a part of the human diet in many tropical countries and the earliest citing of entomophagy can be found in religious literature of Christian, Jewish and Islamic faiths (Huis *et al.*, 2013). Loaded with proteins, fats and minerals, insects offer enormous scope as an alternative source of food. About 31 per cent of all edible insects around the world belong to the order Coleoptera followed by Lepidoptera, Hymenoptera, Orthoptera and Hemiptera.

The opportunities of insects as alternate food sources are due to its environmental, nutritional and socio-economic benefits. The major environmental benefits include high feed conversion efficiency, lower emission of greenhouse gases and lower water as well as land requirement (Huis, 2013). Higher feed conversion efficiency as well as lower

emission of methane, nitrous oxide and ammonia has been reported in the edible insects *Tenebrio molitor*, *Acheta domesticus*, *Locusta migratoria*, *Pachnoda marginata* and *Blaptica dubia*, when compared to conventional livestock (Oonincx *et al.*, 2010).



Fig. 1. Fried silkworm pupae for sale in China

(Source: <https://images.app.goo.gl/BLiPwMsYUuA5Lmkc9>)

Edible insects are highly nutritious as they are rich in carbohydrates, proteins, amino acids, fatty acids and micronutrients. A comparative study of amino acid content in beef and mealworm larva revealed that mealworms had higher isoleucine, leucine, valine, tyrosine, alanine, glycine and proline content (Huis *et al.*, 2013). Harvesting and raising of insects involve low technology and capital investment and provides opportunities

for subsistence for both urban and rural population, which makes it socio-economically viable.

Edible insects can be obtained by wild harvesting, semi-domestication and rearing/farming. Ninety two per cent of known species of edible insects are obtained through wild harvesting, six per cent by semi-domestication and two per cent by rearing (Yen, 2009). Rearing and semi domestication are considered to be the most productive method as they have huge potential to provide a more stable supply. Bamboo caterpillar, *Omphisa fuscidentalis* and weaver ant, *Oecophylla smaragdina* are collected from the wild, while the palm weevil, *Rhynchophorus palmarum* serve as a classical example of semi-domestication (Govorushko, 2019). Rearing of insects in captivity, isolated from their natural populations and provided with controlled living conditions and diet is referred as insect farming. House crickets, palm weevils and mealworms are being successfully farmed in Thailand (Hanboonsong *et al.*, 2013).

After being harvested, the insects are processed and consumed as whole insects, in ground or paste form or as extracted protein. Blanching and drying are the common processing techniques followed, which minimizes the microbial risk and increases the shelf life of edible insect products. The effect of processing technology and storage conditions on the microbial characteristics of

mealworm larva, field cricket nymph and migratory locust adult revealed that storage after drying at 103°C for 12 hours reduced the total microbial count, as well as counts of enterobacteria, yeast and mould significantly (Adamek *et al.*, 2018).



Fig. 2. Insect food product

(Source: <https://images.app.goo.gl/px3ErwK6KjgKx1Co6>)

Even though use of insects as food confers many advantages, not all insects are safe to eat. Microbial, chemical, physical and allergic risk should be considered while selecting insects for human consumption. In future, meat centric diets will become increasingly expensive and grain-livestock systems environmentally unsustainable. Though entomophagy is still in its early stages, given due support, it can play a significant role in assuring global food security.

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Occurrence of the red spider mite, *Tetranychus* sp on red amaranthus in terrace garden in Bangalore

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Red amaranthus is a delicious leafy vegetable. It is becoming popular in kitchen gardens and in pots on terraces in urban Bangalore city (12.9784° N, 77.6408° E). The mite *Tetranychus* sp which had attacked Moringa (Verghese *et al.*, 2022) in the same area migrated to the amaranthus. The plants

showed white to brown discoloration on the leaves (Fig 2). The leaves showed downward curling (Fig 3) with presence of mites and web on the ventral sides (Fig 2,) as compared to an uninfested plant (Fig.1). As these were organic, no sprays were given and affected plants were uprooted and destroyed.



Fig. 1 Uninfested plant

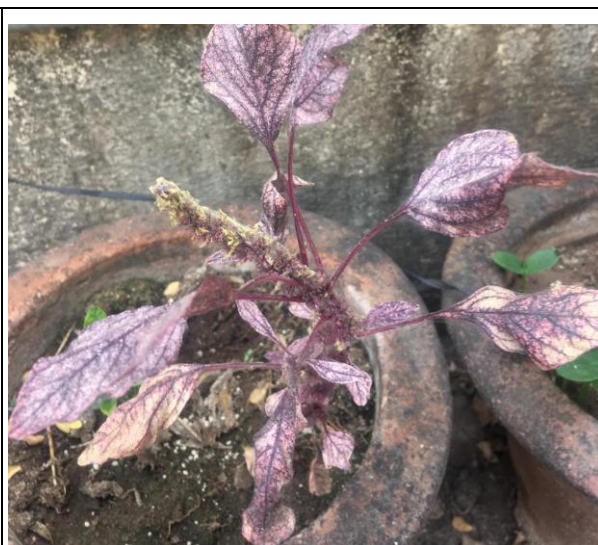


Fig. 2 White to brown discoloration on the leaves, downward curling



Fig. 3 Downward curling with presence of mites

Verghese, A., Rashmi. M. A and Deepak. S,
Liquid pongamia soap unsettles
Tetranychus sp. on Moringa DOI:
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**A report on positive phototaxis exhibited by newly emerged bagworm caterpillars
(Lepidoptera: Psychidae)**

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Abstract

The bagworm, *Thyridopteryx ephemeraeformis* (Haworth) (Lepidoptera: Psychidae) is a polyphagous moth that defoliates ornamental shrubs and trees. It has been reported as an emerging pest in the coastal agroecosystem of Odisha, India. A small experiment demonstrated that bagworm's newly emerged larva (NEL) was positively phototactic under white and UV light. After emerging in the rearing box, the larva was exposed to 3 volts of LED white and UV lights. After which, its phototactic behaviour was observed. Considering this report, light traps can be used to catch this nuisance species, which may now be more manageable. Hanging light sources on infested plant branches may be one technique, and when larvae congregate on that area of the branch, it may be easy to remove and kill them.

Keywords: *Thyridopteryx ephemeraeformis*, bagworm, phototaxis, Psychidae, Lepidoptera

Introduction

The bagworm, *Thyridopteryx ephemeraeformis* (Haworth), is a polyphagous moth that defoliates ornamental shrubs and trees. (Jones and Parks 1928). The adult female of this species has vestigial appendages and mouthparts, small eyes, apterous with no antenna. She spends her entire life upside-down inside the bag constructed by the larva during developmental stages (Kaufmann 1968; Neal 1982). On the other hand, males have wings and are diurnal and get attracted to females with a sex-pheromone scent they emit. Females deposit 1-methyl butyl decanoate in

the cocoon at the bottom of their larval sack until they mate (Leonhardt et al. 1983, Klun et al. 1986). After reaching the female bag, the male inseminates the female by pneumatically stretching his abdomen inside her pupal shell, past the length of her body, to reach her caudal genitalia. The female oviposits inside her pupal shell and then falls from the bag. The eggs overwinter, and the larvae emerge in the spring. Young larvae typically make a silk thread and drift away on the breeze. (Jones and Parks 1928). This action is most likely one of the most important ways of distribution in the environment. On the other hand, late instar

larvae have been seen to crawl away from their host plant shortly before pupation. Extensive research was done by Kaufmann (1968) on the *T. ephemeraeformis* and observed its biology and behaviour.

Over 50 families of deciduous and evergreen trees and shrubs can be targeted by *T. ephemeraeformis* (also known as the evergreen bagworm, common bagworm, eastern bagworm, and common basket worm) (Rhainds *et al.*, 2009) and have been reported as an emerging pest in coastal agroecosystem of Odisha, India (Srivastva and Attri 2004). Still, research on its behaviour and other Psychidae is poorly known (Kaufmann 1968).

Insects use a variety of methods to find their way around. Bees' utilisation of the sun and polarised light for detecting food sources and reorienting to the hive is widely documented. Apart from chemical trails, ants have been observed using the sun for orienting (Romoser 1973). Dung beetles (Smolka *et al.*, 2016), aphids (Hajong and Varman 2002) are known to be phototactic as well. However, artificial light at night (ALAN) strongly impacts the insect population (Grubisic *et al.*, 2018). Luminosity, exposure time, and wavelengths of light-emitting diodes (LEDs) affect agricultural insects and stored-product insects. Agricultural insects are particularly attracted to green or blue LEDs because of their phototactic tendencies, and they are the most effective entrapment methods for these pesky pests (Park and Lee 2017).

Reports on adult lepidopteran attraction towards light are many (Park and Lee 2017). However, there are only a few observations of positive phototactic movements of lepidopteran caterpillars recorded, which is again used to trap the pest (Rao *et al.*, 2016). ALAN was tested in a deciduous forest by Welbers *et al.* (2017), who used street lamps that emitted various colours of light to modify the lighting conditions. Caterpillar abundance peaked substantially higher in trees illuminated with green or white light than it did in trees illuminated with red light or in the absence of light, according to this study. In addition, male caterpillars exposed to green and white light had lower body mass and pupated quicker than those exposed to red or dark treatments, according to Van Geffen *et al.* (2014). Another research by Gotthard (2000) indicated that the length of the illumination could also influence the abundance of caterpillars because of its effect on the level of predation by predatory insects.

A small experiment confirmed the positive phototactic behaviour of the bag worm *Thyridopteryx ephemeraeformis* (Haworth), which was earlier doubtful by Kaufmann (1968) in his study. He observed that the newly emerged larva of the bagworm emerge out of the bag but did not confirm whether it is a positive phototactic response or not.

The bags were collected from the host *Ziziphus jujuba* shrubs from different localities in the agronomy field of Orissa University of

Agriculture and Technology, Bhubaneswar, Odisha (20°15'54.9"N 85°48'25.3"E), in the year 2017. The bags were chosen according to the indications provided by Kaufmann (1968). The matured bag (having the last instar larva) were collected by seeing the moulted head on the outer surface of the bag. This caterpillar moults seven times and becomes an adult (Kaufmann 1968). So the bag whose outer case had around 5-6 moulted head attached were chosen. Total 5-10 bags were collected and kept in a box of size 15cm x 15cm x 15 cm so that after the emergence of the adult winged male, it will find the female bagworm case and try to fertilise it. After copulation, the female bag cases were separated out and each one was kept in a small cylindrical box of 6cm (ht.) x 7cm (dia). Each small box was marked with A (left), B (middle), C (right), D (back) (as shown in figure 1). 3V white and UV LED lights were used to study the behaviour of the newly emerged larva (NEL).

Observations and Discussions

The female, after fertilisation, oviposits eggs inside the pupal case, which is held inside the bag case. After NEL comes out from the bags, it gets scattered all over the box by spinning a thin layer of silk on the inner surface of the box. The NEL move from one place to another by lifting its soft abdomen. Soon after emergence, within minutes, they start to build their bags with the help of the silk formed inside the inner layer of the box. After that, the LEDs were used to study the phototactic

response. When the LEDs were switched off, all the larva were found to be scattered evenly all over the box in every marked position, i.e., A, B, C, and D (Fig 1.1). Soon after 'switching on' of the LED at different box positions (Fig 1.2, 1.3, 1.4, 1.5), the 100s of larvae tend to aggregate on the focus point of the light intensity. The shift in the LED's position tends to shift the aggregation of the bagworm larva. Then after the LED were 'switched off' the larvae get evenly scattered in the whole box again (Fig. 1.1; 2. a-f). This result was accurate for both the LED lights white and UV. Welbers et al. (2017) observed the same results by using green or white lights, increasing the caterpillar abundance. Also, UV light does the same Rao et al. (2016).

An opening cut in the bag's midsection prompted a larva's head to travel toward this opening, where it penned its skin and emerged as a pupa. The same results were produced when the posterior end was left open and the new centre aperture was created larger than the posterior one. Because the larvae oriented their heads toward the opening that provided the bag with the most light, these findings suggest they were positively phototropic (Kaufmann 1968).

According to this report, using light traps to catch this nuisance species may now be more manageable. Hanging light sources on infested plant branches may be one technique, and when larvae congregate on that area of the branch, it is easy to remove and kill them all.

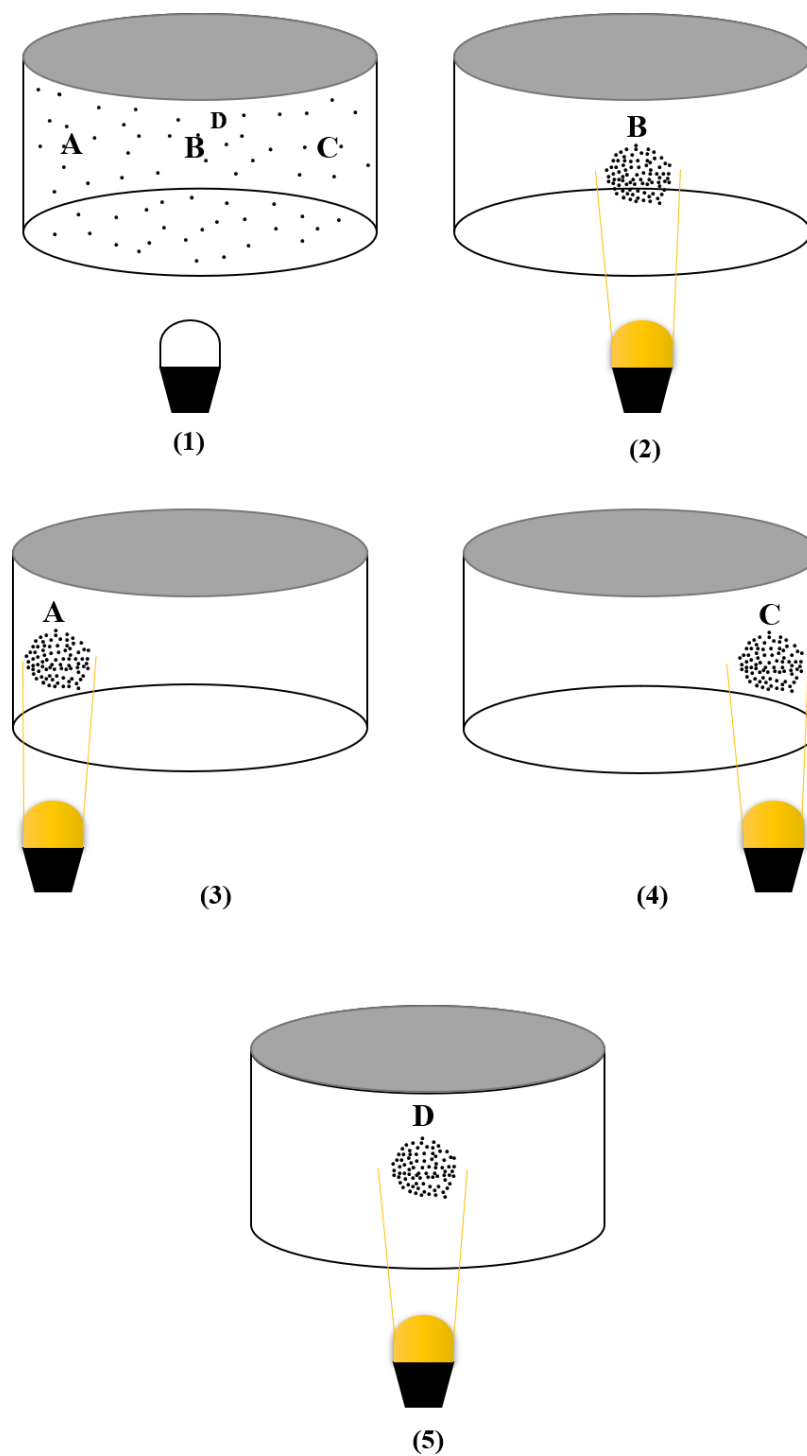


Fig. 1. (1) Evenly scattered bagworm larva when no light intensity was present (2) Aggregation of larva in 'B' (3) Relocation to 'A' (4) Again relocation to 'C' (5) Again aggregation of the larva after placing the light on the back of the box in 'D' position

Conflicts of Interests

The authors have no conflict of interest

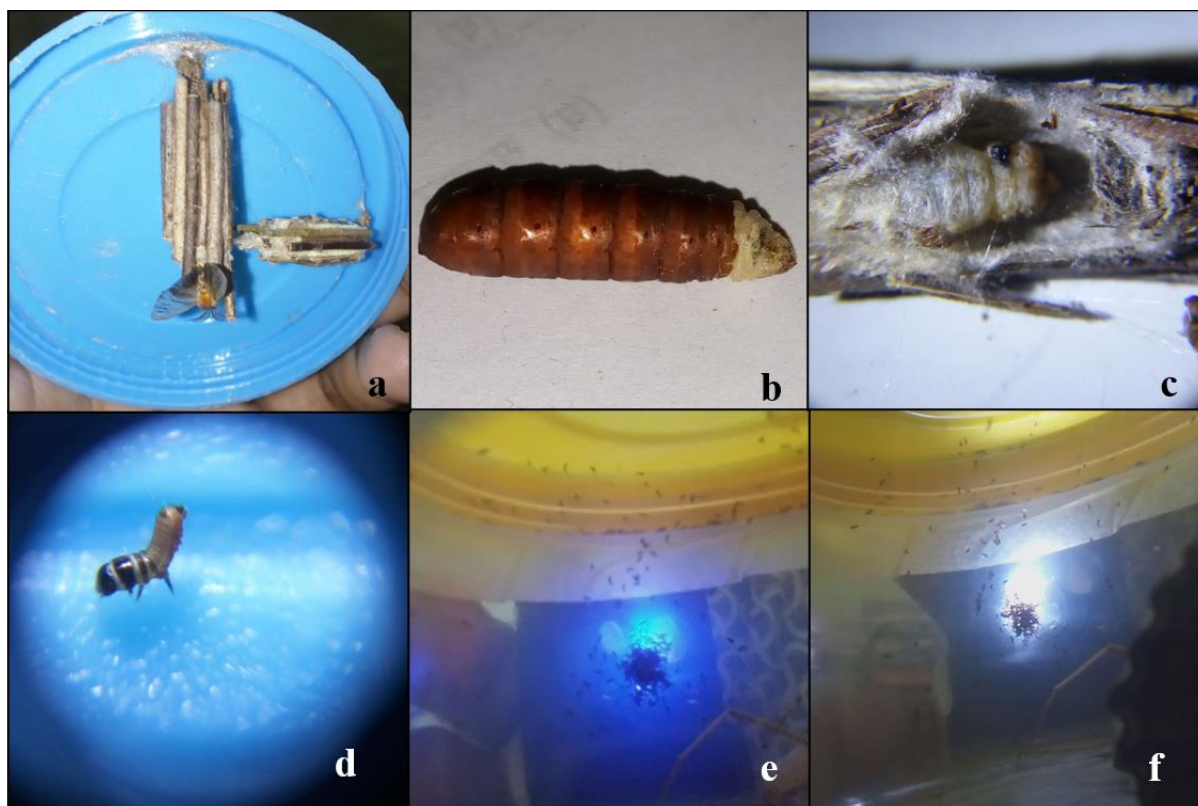


Fig. 2. (a) Mating of *Thyridopteryx ephemeraeformis*; (b) Adult female with the pupal case attached; (c) Adult female inside the bag without the pupal case; (d) Newly emerged larva (NEL) moving with its raised abdomen; (e) Aggregation of NEL under UV light; (f) Aggregation of NEL under white light

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Parasite on Asian rice gall midge - *Orseolia oryzae* (Wood-Mason) in light trap collection**A Sunny Rao^{1,2}, A.P. Padmakumari ^{1*} and Gajendra Chandrarkar²**¹ICAR-Indian Institute of Rice Research, Rajendranagar, Hyderabad, Telangana 500030 India.² Indira Gandhi Krishi Vishwa Vidyalaya, Raipur, Chhattisgarh. 492012, India.***Corresponding author: E mail: padmakumario@gmail.com**

Adult rice gall midge, *Orseolia oryzae* (Wood mason) (Diptera: Cecidomyiidae) insects were collected from light traps during September 2021 and brought to the laboratory. On observation it was found that many insects were dead. While observing the dead adults under a microscope, a peculiar structure was observed to be attached to the female adult on the ventral side. Further microscopic examination revealed that an organism was attached to the intersegmental area between the thorax and abdomen and was feeding on the adult gall fly (Fig1). We were not sure where and how the parasite had attacked the adult insect. It had devoured only the gall midge insect tissues but not the eggs which were in

the abdomen. Two days after feeding it had detached itself from the host tissues. Since it was feeding from outside we presume it be an external parasite.

The parasite was observed to possess an oval shaped body having projections on the dorsal surface (Fig 2). On the ventral surface it had a longitudinal groove running from one end to another end (Fig 3) and possessed a mouth.

This is the first report. Perusal of available literature could not throw light on the external parasites of gall midge.

**Fig. 1: Parasites feeding on the gall midge adult****Fig. 2: Dorsal view of the parasite.**



Fig. 3: Ventral view of the parasite

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Foraging and influence of season on *Apis dorsata* in an urban garden**Anitha A Abraham***Department of Life Science, Bangalore University, Bangalore -560004, India.****Corresponding author: abrahamanitha@yahoo.com***

The Asian honeybee *Apis dorsata* Fabricus (Apidae, Apinae) is known for its ecosystem service. Foraging is an important aspect of insect life. In the case of *A. dorsata*, searching for suitable floral resources is essential for the sustenance of the honeybee. This honeybee is an important crop pollinator known to visit several plants for its food source mainly nectar and pollen (Robinson, 2012). The honeybee is also found foraging on flowers in urban gardens.

An observational study was made in the year 2017-2018 at the nursery in Lalbagh Botanical garden (12.95°N 77.59°E) in Bangalore. The bee was observed from January to June in the garden. It visited many native and non-native plants in the garden. The foraging choices made by this bee are dependent on the seasonal availability of plant resources. It was observed that they visited a wide variety of plants from January to June. The flowers of the plants they foraged on were *Pentas lanceolate*, *Hibiscus rosa sinensis* (Anitha *et al.*, 2017). It was seen foraging on *Nymphae* sp. They were seen foraging in large numbers during February and March. The seasonal availability of nectar and pollen was important in their foraging choices. Its body size and large hives demand high energy.

Their foraging decisions are based on these factors. Many ornamental trees were foraged by *A. dorsata* during early summer like *Tabebuia argentea*, *Samanea saman*, and *Pongamia*. The abundance of floral resources seems to be a reason for higher foraging in summer and the seasonal availability of floral resources. Observations over a temporal scale showed floral fidelity in *A. dorsata*. This shows it can learn complex floral colors and odors of flowers of various plants ((Mogily *et al.*, 2020). This ability enables them to forage on the same floral patch seasonally over a temporal scale.

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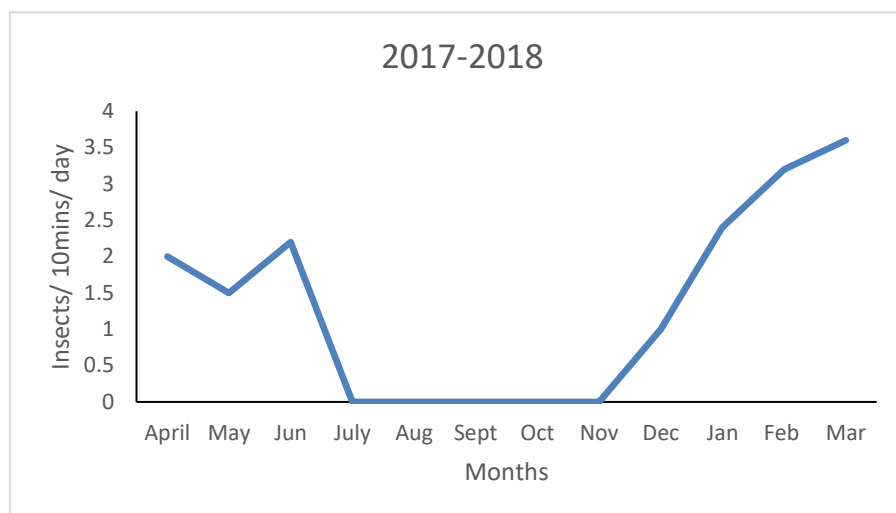


Fig 1: Population of *Apis dorsata* in the year 2017-2018



Fig 2: *Apis dorsata* on *Nymphaeae* sp, with hind leg showing pollen collected

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Diversity of silk production in insects**Athira G. Menon and Haseena Bhaskar***Department of Agricultural Entomology,**College of Agriculture, Kerala Agricultural University, Thrissur-680656, Kerala, India***Corresponding author: athiragmenon94@gmail.com**

Silk is one of the earliest natural fibres discovered by man. Silks are fibrous proteins containing highly repetitive sequences of amino acids stored in the animal as a liquid, and configure into fibres when sheared or spun at secretion (Craig, 1997). The cocoon silk of domesticated silkworm, *Bombyx mori*

(Lepidoptera: Bombycidae) is prized over millennia for textiles, and most of our understanding about silk production is from this species. However, silk is known to occur in many arthropod classes, and a few molluscs and fishes also.

**Fig 1: Spider silk**

(Source: natural-japan.net)

**Fig 2: Embioidae silk**(Source: <https://smbasblog.com/2020/07/05/the-curious-webspinner-insect-knits-a-cozy-home-deep-look-video/>)**Fig 3: Insect silk**(Source: <https://uwm.edu/field-station/the-wonders-of-webs-ii-insect-silk/>)

Among the class Insecta, 16 out of 30 orders produce silk for a variety of purposes, which include reproduction, shelter, protection from predators, etc. (Walker *et al.*, 2012). Study conducted by Gurr and Fletcher (2011) revealed that the silk produced by Australian endemic leafhopper, *Kahaono montana* Evans (Cicadellidae: Typhlocybinae) provided protection from the lacewing predator,

Mallada signata (Schneider) (Neuroptera: Chrysopidae).

In insects, silk is produced and stored in dedicated glands. On the basis of location, Sehna and Akai (1990) classified silk glands into labial glands, Malpighian silk glands and dermal glands. Labial gland accounts for most silks and is the prominent feature in caterpillar and pupal biology of Lepidoptera. Antlions,

mayflies and thrips produce silk from, their Malpighian glands whereas, some water beetles and lacewings synthesize silk from dermal glands.

Insect silks have high levels of amino acids like glycine, alanine and serine. The abundance of these amino acids is most likely due to their non-essential character and intermediate hydrophobicity. Insect silks are semi-crystalline materials whose structure differs from species to species in molecular organization. Hydrogen bonds within and between the structures contribute to the mechanical strength and stability of silk fibres. Accordingly, five different structures have been identified *viz.*, extended β -sheets, cross beta, α -helices, collagen helices and polyglycine-II (Rudall and Kenchington, 1971).



Fig. 7: Silk refuges produced by *K. montana*

(Source: Gurr and Fletcher, 2011)

Sutherland *et al.* (2007) identified a silk gene, *DFibroin* from highly expressed mRNA extracted from the prothoracic basitarsus of male hilarine flies (Diptera: Empididae). The silk gene from the basitarsi cDNA library matched an approximately 220 kDa protein

from the silk-producing basitarsus. The hilarine silk protein is high in glycine and asparagine, and adopts an extended β -sheet conformation.



Fig 8: Silk produced by Hilarine fly

(Source: <https://bugguide.net/node/view/1049683/bgimage>)

Insect silk possesses extraordinary mechanical properties in terms of strength, extensibility and stiffness. The obvious example of the use for silk is cloth, which also takes up the highest proportion of silk consumption. Scientists from Khan Koen University, Thailand, invented a silk bicycle using silk with resin at one third the cost of carbon-fiber or aluminium frame. The silk frame can take five times the pressure and eight times the tension of an aluminium frame. As the silk frame has 30 times more elasticity, it also serves as a natural shock absorber. The versatility and sustainability of silk-based materials attract its use in food packaging, medicine, automobile industry, dietary and cosmetic supplements, optics, art, craft, *etc.* (Huang *et al.*, 2018).

Though the term ‘silk’ encompasses a wide range of distinct materials, it is

remarkable that certain features are common among silk production systems in insects. Today, insect silk has taken on new importance to society beyond fabric. Mechanically enhanced silk is expected to open up possibilities for numerous novel applications.

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Armoring maize from invasive fall armyworm

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Introduction

In India the invasive pest, maize fall armyworm (FAW) (*Spodoptera frugiperda* J. E. Smith; Noctuidae, Lepidoptera) was first recorded in May 2018 (Sharanabasappa *et al.*, 2018). It is the second most destructive agricultural pest, native to tropical and subtropical America (CABI, 2018). Due to its migratory behavior, high dispersal capacity and fast multiplication rate the pest spread quickly to many countries worldwide. The insect is capable of migrating 500 km to 1,000 km during its lifetime. Fall armyworm is a polyphagous pest, It is reported on 353 host plant that belongs to 76 plant families (Montezano *et al.*, 2018). Among them, maize is highly favorite host plant. Fall armyworm attacks the maize crop from the first week of germination till maturity and causes severe damage throughout the year. In maize it causes 70-100% yield loss (Acharya *et al.*, 2020). It also causes economic damage in other cereals and millets.

Life cycle

The life cycle (30 days) of the fall armyworm is mostly influenced by climatic

factors. In hot climatic condition fall army worm undergoes 12 generation per year. Its life cycle comprises of eggs, larvae, pupae, and adult stages. Images of several life stages of FAW were collected from mass culture in our laboratory, AC&RI, Killikulam. As an egg laying substrate, nerium plants were used. The institute is located at an altitude of 40 metres above mean sea level, at 8° 46' North Latitude and 77° 42' East Latitude, in the Semi-Arid Tropics, with mean temperatures ranging from 21 to 37 degrees Celsius and an annual rainfall of 786 mm.

Egg

The female moth often deposits her egg masses on the under surface of the two to third leaves of seedlings or inner side of the tender central whorl. It lays egg masses in a single layer or a few layers (2-3 layers) and then covers them with scales. A single egg mass includes 100- 200 eggs. A female lays 1500 to 2000 eggs in her lifetime. The egg period lasts between 2 to 3 days.



Egg mass covered with scales



Egg mass



Eggs ready to hatch



Neonate emergence



Larva (Caterpillar)

The larval stage is the destructive stage on crop plants. After the egg hatches, masses of neonate start to disperse on leaf surface. The neonate used to be white in colour with black head. There are six larval instars. The larval group spread to the surrounding plants by hanging off from the silken thread secreted by them, due to ballooning effect caused by wind force larvae carried over to numerous plants from a single egg mass. Cannibalistic behavior has been documented from third instar, therefore from the third instar onwards each

plant occupied by single larva rarely we can observe two larvae at different region of the same plant. The FAW larva can be easily identified by its morphological features such as 'Y' shape inverted pale marking on its head. On every segment of the larval body, there is a symmetrical distribution of dark black colour raised dots (2 dorsal pairs and lateral pairs). Each spot consist of single hair. The terminal segment has four dark dots that are distributed evenly and in a square form. The larval body has longitudinal pale and dark bands on the dorsal and lateral sides. Larval period is for 14 to 22 days.



First instar



Second instar



Third instar



Fourth instar



Fifth instar



Sixth instar

Larval stages

Pupa

The mature larvae forms earthen cocoon inside the soil at a depth of 2 to 8 cm. Sometimes it pupates within the plant itself. Pupa is reddish brown in color. The pupa measures 2-3 cm in length and 4.5 mm in width. During the summer, the pupal stage lasts for 7 to 13 days.















Adult

Similar to other noctuid moths it possess scaly wings with variegated colours such as black, light to dark brown, grey and straw colour. The forewing of male moth has different markings. The terminal end of the wing contains white inverted triangular marking. At one third portion of the forewing from thorax has golden yellow colour oval/kidney shape marking. The female moths forewing is uniformly grey in colour. Adult moth is nocturnal, hiding under vegetation or inner side of the leaf whorl during the morning hours. The hind wing of both male and female moths used to be small and silvery in appearance. Female moths begin to lay eggs 3 to 4 days after mating (pre oviposition period). Moths can live for about 7-15 days.

**Male moth****Female moth****Damage symptoms**

- ❖ The first and second instar larvae feed on leaf surface epidermal tissue by scraping chlorophyll and make the leaf surface papery in nature.
- ❖ The third to sixth instar larvae are voracious defoliators; they mainly feed and damage the tender developing central whorl and make small to big size holes. It also feeds on tassel and immature cob.
- ❖ Because of the damage to the unopened whorl, it makes parallel circular window holes and random irregular holes on leaves. The holes expand in size as the plant grows.
- ❖ Heavy whorl damage causes ragged, torn or shredded appearance to newly opened top-most leaves.
- ❖ FAW infestation can be easily identifiable by the presence of mass larval dropping on leaf surface and central whorls.
- ❖ During vegetative stage, larvae hide inside the leaf whorl and cause whorl damage. At the time of tassel and cob emergence, larvae hide between stem and leaf base.
- ❖ Fall armyworm damage to tassel has an impact on pollination.
- ❖ Larvae affect the tip and inner side of the cob and feed on developing kernels during milky stage, affecting seed development and reducing crop productivity and nutritional status.
- ❖ FAW infestation on maize cob invites secondary fungus infection, resulting in aflatoxin production, which reduces grain quality and quantity.

		
Leaf chlorophyll scraped by the neonates		Leaf holes
		
Whorl damage	Whorl covered by insect excreta	
		
Shredding of leaves	Tassel damage	Silk damage
		
Cob damage		
Crop damages		

Scouting: 'W' pattern of scouting from early seedling stage helps to find FAW

Economical Threshold Limit for FAW damage: 10% whorl and cob damage

Integrated fall armyworm management





The Tamil Nadu Agricultural University has conducted a wide range of research and has recommended the following strategies to control the maize fall armyworm.

- ❖ Deep summer ploughing to expose FAW pupa to predators and to kill by scorching sun.
- ❖ In the last ploughing, apply neem cake at the rate of 100 Kg / acre and plough well for providing pest resistance to plants.
- ❖ Avoid staggered sowing; timely and uniform sowing over large area reduces FAW incidence.
- ❖ Seed treatment: Seed treatment should be done with the mixture of Cyantraniliprol 19.8% and Thiamethoxam 19.8% FS at the rate of 4 ml per kg of maize seed. This will reduce the damage caused by fall armyworm as soon as the crop germinates.
- ❖ In irrigated condition, cowpea, sesame, sunflower and redgram should be cultivated as border crop, grow the fodder sorghum as in case of rainfed maize cultivation. Thus increasing the number of natural enemies.
- ❖ Installing sex pheromone traps at the rate of 5 traps per acre can be used to monitor the movement of fall army worm.
- ❖ Regular hand picking and destroying of FAW egg masses and larvae prevents severe damage.
- ❖ Augmentative release of egg parasitoid *Telenomus remus* @ 50,000 per acre based on pheromone trap catches prevents FAW multiplication.
- ❖ Crop spacing: Irrigated condition- 60x25 cm, Rainfed condition- 45x20 cm.
- ❖ Biopesticide such as *Metarhizium anisopliae*, *Nomuraea rileyi*, *Beauveria bassiana* and *Bacillus thuringiensis* also controls fall armyworm growth and development.
- ❖ Recommended chemical pesticides only recommended pesticides should be used at the respective time. Doing so may reduce the insect resistance to the pesticides.
- ❖ Recommended insecticides should be sprayed at the recommended rate. The entire whorl region should be sprayed properly.

Crop age	Recommended pesticides	Hand sprayer	Power sprayer	Pesticide Per acre
15-20 Days after emergence	Chlorantraniliprole 18.5 SC	0.4 ml/lit	1.2 ml/lit	80 ml/acre
	Flubendiamide 480 SC	0.5 ml/lit	1.5 ml/lit	100 ml/acre
30-35 Days after emergence	Azadirachtin 1500 ppm (If necessary, when FAW damage crosses the ETL 10% level)	5 ml/lit	15 ml/lit	1 lit/acre
40-45 Days after emergence	Emamectin benzoate 5 SG	0.4 g/lit	1.2 g/lit	80 g/acre
	Spinetoram 11.7 SC	0.5 ml/lit	1.5 ml/lit	100 ml/acre
	Novaluraon 10 EC	1 ml/lit	3 ml/lit	200 ml/acre

- ❖ 60 DAE: Any one of the insecticides which is not sprayed previously, Emamectin benzoate 5 SG @ 0.4 g/lit or Spinetoram

11.7 SC @ 0.5 ml/lit or Novaluraon 10 EC @ 1ml /lit

	
<i>Telenomus remus</i> - egg parasite	
	
FAW egg mass parasitized by <i>Telenomus remus</i>	
	
Pulses as border crop	Pheromone trap

Fall armyworm naturally infected by some beneficial microbes



Bt infected FAW larva

Metarhizium infected FAW
larva

Fungus infected FAW egg
mass

Note: Field photoes were taken from AC&RI, Killikulam research plot.

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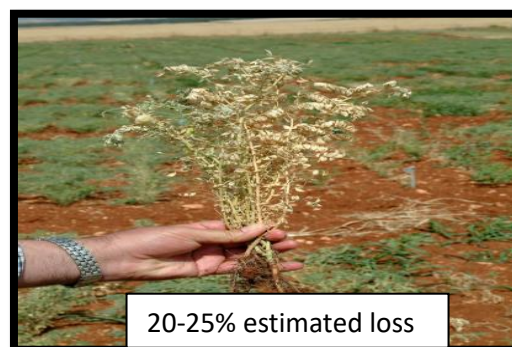
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Molecular approach towards resistance to chickpea pod borer : *Helicoverpa armigera***Smitha S.***Department of Plant Biotechnology**University of Agricultural Sciences, GKVK, Bengaluru-560065 Karnataka, India****Corresponding author: smithaswathickm@gmail.com**

Chickpea (*Cicer arietinum* L.) is a rabi season legume crop and third most extensively cultivated pulse crop globally. *Helicoverpa armigera* is considered as one of the most devastating key pest severely affecting chickpea, due to its high rate of reproduction with short life span (Kumar and Singh, 2014). It is a vigorously feeding, polyphagous pest and is also called as corn ear worm, American cotton boll worm, tobacco bud worm, fruit borer of tomato, carnation worm *etc.* The

prime reason for the lower yield of chickpea is infestation by pod borer (*Helicoverpa armigera*) from the vegetative phase to pod formation period (Dhingra *et al.*, 2003). Single larva of the *Helicoverpa* damages 30-40 pods prior to its maturity. Yearly pod damage caused by *Helicoverpa* alone is about 150 to 200 million tonnes. With the predictable yield loss only due to gram pod borer is 10-90 per cent.



Even though efforts have been made towards developing resistant genotypes through conventional method, especially conferring pod borer resistance it has not been successful up to requisite level due to the lack of genetic resource conferring resistance to chickpea pod borer. This is the similar case in cotton boll worm also where successfully genetic engineering was used to transfer genes from bacteria, *Bacillus thurengiensis* known to

produce proteins toxic to *Helicoverpa armigera*. Now, in the world more than 95 per cent of the commercial cotton growing area is covered with genetically modified cotton, called as *Bt* cotton, resistant to boll worms. Similar efforts have been made in chickpea at Assam Agricultural University, Jorhat.

Although the potential productivity of chickpea has been scientifically or experimentally proved to be 20 – 22 quintals

per hectare, farmers are harvesting only 5 – 10 quintals per hectare. This low and variable productivity level of chickpea in farmers' field is due to loss of crop because of incidence of pod borer and *Fusarium* wilt disease. If these two biotic stresses are not managed, then the extent of loss will be 80 – 90 per cent. To decrease the cost of cultivation and reducing environmental pollution by limiting the application of pesticides and fungicides to manage these two biotic stress problems, developing genotypes tolerant or resistant to

pod borer (*Helioverpa armigera*) and *Fusarium* wilt (*Fusarium oxysporum* f. sp. *ciceri*) is the best alternate option.

Thus it was planned to introgress pod borer and *Fusarium* wilt resistance in Chickpea by involving Bt event of Chickpea (Acharjee *et al.*, 2010) and wilt resistant Super Annigeri-1 genotypes by using Simple Sequence Repeats (SSR markers for screening specific to SA-1).



An overview of experimental site

Hybridization experiment was conducted at transgenic green house during rabi season, when donor parents (Bt events having Cry genes resistance to pod borer pest) and recipient parent (Super Annigeri-1) were sown. Crossing for artificial hybridization was carried out using recipient parent as female parent. F₁ seeds obtained from these crosses were harvested. The harvested F₁ seeds were sown in pots during late kharif season then

checked for the presence of Cry genes in the hybrids.

The genomic DNA was extracted from young leaves of 20 days old seedlings of both parents and hybrids using CTAB method and quality of DNA for each sample was assessed on agarose gel. DNA amplification was carried out using pair of 177 SSR primers developed by Nayak *et al.* (2010) in polymerase chain reaction (PCR). Whereas, Cry genes specific

markers namely *Cry1Ac* and *Cry2Aa* were also involved in the study in order to confirm the presence of cry genes in hybrids. The hybridity of F₁ plants was confirmed when they showed

presence of both male and female parent alleles with the help of parental polymorphic markers along with cry gene specific markers.



Donor parents (1 and 2) having cry genes resistance to pod borer pest, recipient parent Super annigeri-1



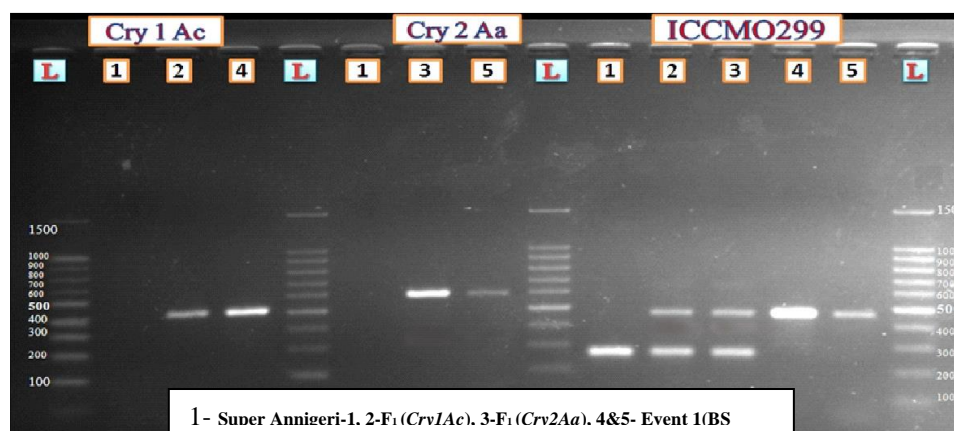
F₁ hybrid plant crossed between Super annigeri-1 × Bt event BS 100B (having cry gene resistance to pod borer)

The presence of both the parental alleles in hybrid plants confirmed their origin from the two parents used in the present study, as well as genuineness of hybrid plants. Plants were confirmed as true hybrids on the basis of amplification pattern of SSR marker. Results of the present study are in agreement with the

conclusions of Hipi *et al.*, (2013) in maize. Sharma *et al.* (2018) confirmed the hybridity of Indian mustard (*Brassica juncea* L.) using SSR markers. Out of 20 random SSR primers used for the screening of parental polymorphism, 5 primers were found polymorphic. BR_A04_9627743 and

BR_A01_13393871 were identified as the specific markers for parents RSPR-01 and

Donskaja-IV which enable to distinguish and identify hybrid form their parental lines.



Confirmation of true F₁ using gene specific primers (cry genes)

F₁ hybrid was confirmed with both donor gene and recipient gene through one of the parental polymorphic marker ICCM0299

This study showed that SSR markers are more reliable and robust for assessing genetic purity as compared to morphological marker. The results of study are expected to be useful in the verification of genetic purity of hybrid seeds in chickpea accurately. Identified polymorphic markers between parents are good source for recovering recurrent parent genome in early backcross generations.

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Insect Navigation - en route to the goal**Kattumulla Tejaswee and Ranjith M. T.***Department of Agricultural Entomology**Kerala Agricultural University, College of Agriculture, Vellanikkara 680656, Thrissur, kerala****Corresponding author: tejasweekattumulla@gmail.com***

Insect navigation refers to the ability of insects to find their way accurately to a specific location. Insects, just like many other moving animals, have evolved navigational abilities to move around their environment accurately and to exploit the resources available (Schultheiss et al., 2015). The three major navigational strategies recognized primarily in social insects are route following/visual piloting, path integration and map like spatial distribution.

Route following is the most straight forward mode of navigation found in the majority of ants, bees and termites, which follow a pheromone trail deposited on the surface by a conspecific and do not require previous knowledge on the path (Wolf, 2011). Similarly, they also employ terrestrial objects as navigational landmarks (Collett, 1996).

Path integration sums the vectors of distance and direction travelled from a starting point to calculate the current position and, thus, the path back to the start. Social insects such as bees, wasps and ants **employ different inputs like odometry, compass and optical flow to find their target in this strategy. In odometry, the insect calculates the distance travelled based on the number of steps taken to reach the**

target site. The compass input, on the other hand, includes celestial cues such as the sun, moon, magnetic field, and wind direction. The honey bee uses sunlight to find food and communicate this information to other members of the hive through various dances and Karl von Frisch was awarded the Nobel Prize in Physiology and Medicine in 1973 for deciphering the bee dance. Fleishmann et al. (2018) demonstrated that in ants, a geomagnetic compass cue is both necessary and sufficient for accomplishing a well defined navigational task. Many insects use optic flow to measure the distance of objects from themselves based on the velocity of movements relative to nearby objects.

The assembly of landmarks, local and global, vectors into a two-dimensional spatial arrangement that yields a true map (mental map) in bees. Though location and mode of memory storage of maps remain elusive, the central complex of the brain in insects is considered to play a vital role in forming the mental map (Menzel *et al.*, 2005).

Insect navigation plays a significant role in insect survival such as to chart food bound routes, locate mate and ovipositional site, survive under adverse conditions,

communicate among nestmates and escape from the predatory pressure (Cohen, 2019)). The basic information about navigational strategies in insect pests also helps to employ different pest management tactics such as trap crops, light traps, pheromone traps and repellents. Integrated Vector Management (IVM) programme proposed by WHO in 2020 uses semiochemicals to redirect the movement of mosquitoes from human habitation (Woodling et al., 2020). The navigational strategies in insects are also employed in modelling of robots used in surveillance programmes (Lambrinos *et al.*, 2000).

The basic strategies of insect navigation is well explicitly studied in social insects, but their importance in the host plant selection process needs to be explored in detail. Though the navigational strategies in insects are employed in various fields, it is imperative to have a multi-dimensional understanding of navigation in insect pests to formulate various strategies to minimize their incidence.

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Observations on the Indian Wax Scale on *Michelia champaka* avenue trees in Bengaluru**Chetan S, Allan Britto and Jayashankar M****Department of Zoology, School of Life Sciences,**St. Joseph's College (Autonomous), Bengaluru-560027, Karnataka, India****Corresponding author: jayashankar.m@sjc.ac.in***

The Indian wax scale, *Ceroplastes ceriferus* (Fabricius) (Homoptera: Coccidae) an invasive polyphagous pest of economic importance to many ornamentals is a soft scale insect with a characteristic thick wax test covering the body of the adult female (<https://www.cabi.org/isc/datasheet/12342>).

Infestations weaken the infested plants, with reduced growth; also deposits of honeydew from these sucking pests enable sooty mold growth hampering growth of plants. Observations on the incidence *C. ceriferus* (Fabricius) on *Michelia champaka* (Linneus) avenue trees on the Museum Road, Bengaluru was undertaken during April 2022. The method used in the current survey involved manual counting. The scales present at

different heights on each tree were counted. The data is restricted to 14 feet height of the trees considering the visual feasibility. Of the 32 trees observed, 21 had the presence of the scale insect and a *Terminalia catappa* tree adjacent to the *M. champaka* had the scale (Figs. 1 & 2). The wax scales were observed on the branches compared to the main stem. The trees infested by a large number of scale insects were observed to be malnourished, with thinning of branches. It was also noticed that the plant part where the wax scales were most prevalent was in the height range of 6-8 feet (Fig. 3). Proper care in terms of monitoring the spread and impact of the pest needs to be initiated.



Fig. 1: Indian wax scale (Star stage) infestation on *M. champaka* in the location



Fig. 2: Spread of wax scale from *M. champaka* to adjacent *T. catappa*

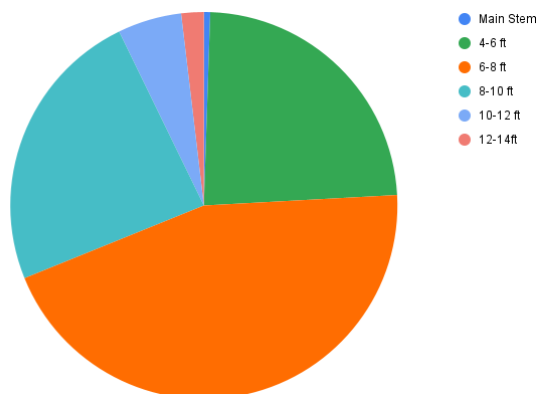


Fig. 3: Graph showing wax scales at different heights

Reference

<https://www.cabi.org/isc/datasheet/12342>

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(c) Subramanian Sevgan

Leaf rolling weevil, Parapoderus submarginatus (Coleoptera: Attelabidae)

Parapoderus submarginatus are small beetles of size 2-8 mm and easily recognisable by their square elytra, that does not cover the last abdominal segment. They are slow moving, but good flyers. Adults feed on leaves and buds and they develop either in leaf rolls or leaf mines, stems or flower heads.

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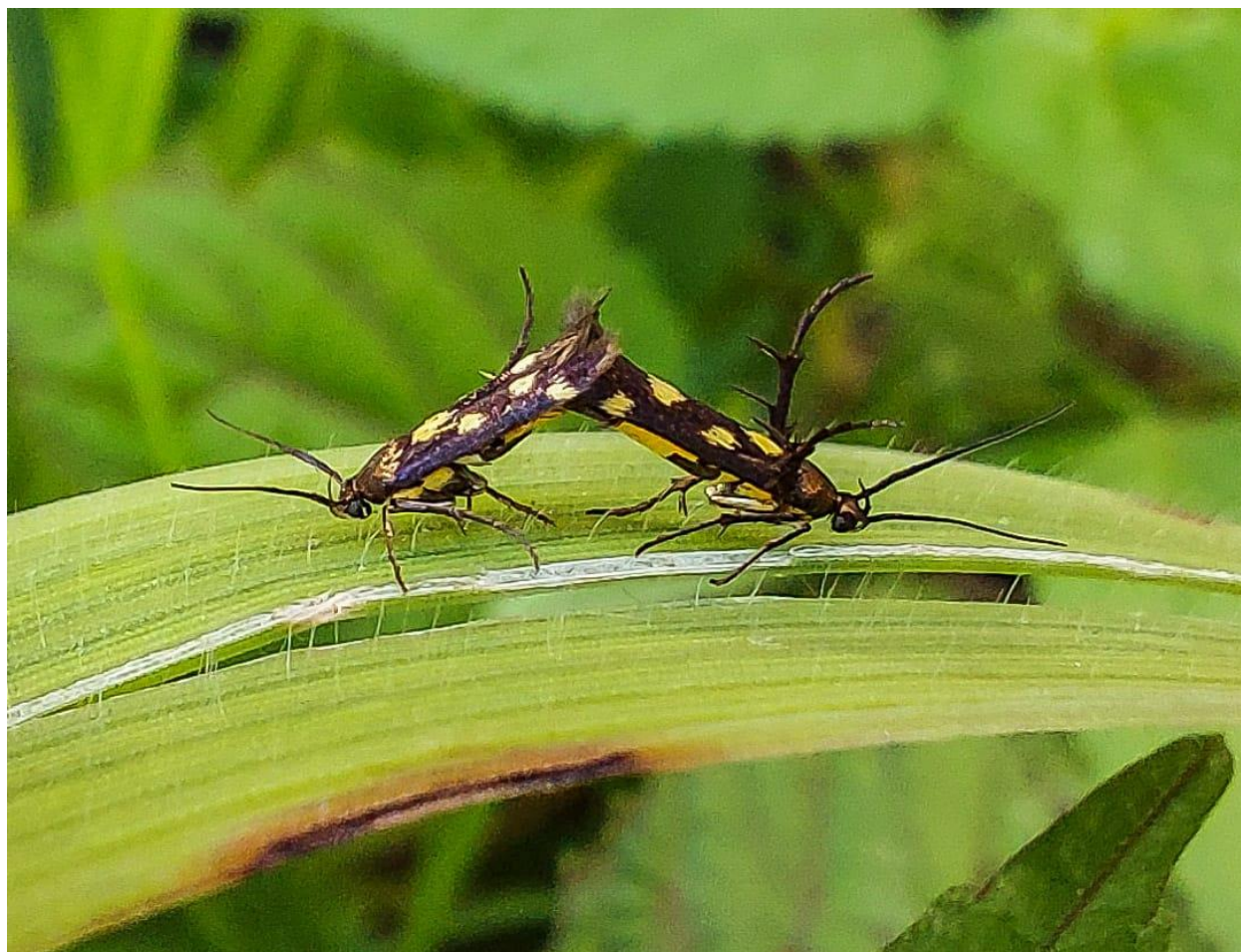
Mud wasp (Hymenoptera)

Here wasp is bringing a caterpillar for its progeny as mass provisional feeding behaviour.

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Leaf webber, Eretmocera impactella (Scythrididae: Lepidoptera)

Leaf webber is sporadic pest of various Amaranthaceae and other food plants distributed in the Indian subcontinent. Caterpillars web leaves with white silken threads and remain hidden in folds feeding from inside.

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Fulgorid Bug (Fulgoridae: Hemiptera)

The family Fulgoridae is a large group of hemipteran insects, sometimes referred to as lanternflies or lanthorn flies, though they do not emit light.

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Cuckoo wasp, Chrysis angolensis (Chrysididae: Hymenoptera)

Like the cuckoo bird, cuckoo wasps will infiltrate the nests of other wasps and bees and lay their eggs alongside their hosts' offspring. Once these eggs hatch, the young cuckoo wasp's larvae will feed on food stored in the nest for the host's offspring, such as a paralysed spider or caterpillar.

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Giant

scales



Monophlebids (Monophlebidae: Hemiptera)

Monophlebids is a family of scale insects commonly known as the giant scales or monophlebids. They occur in most parts of the world but more genera are found in the tropics than elsewhere.

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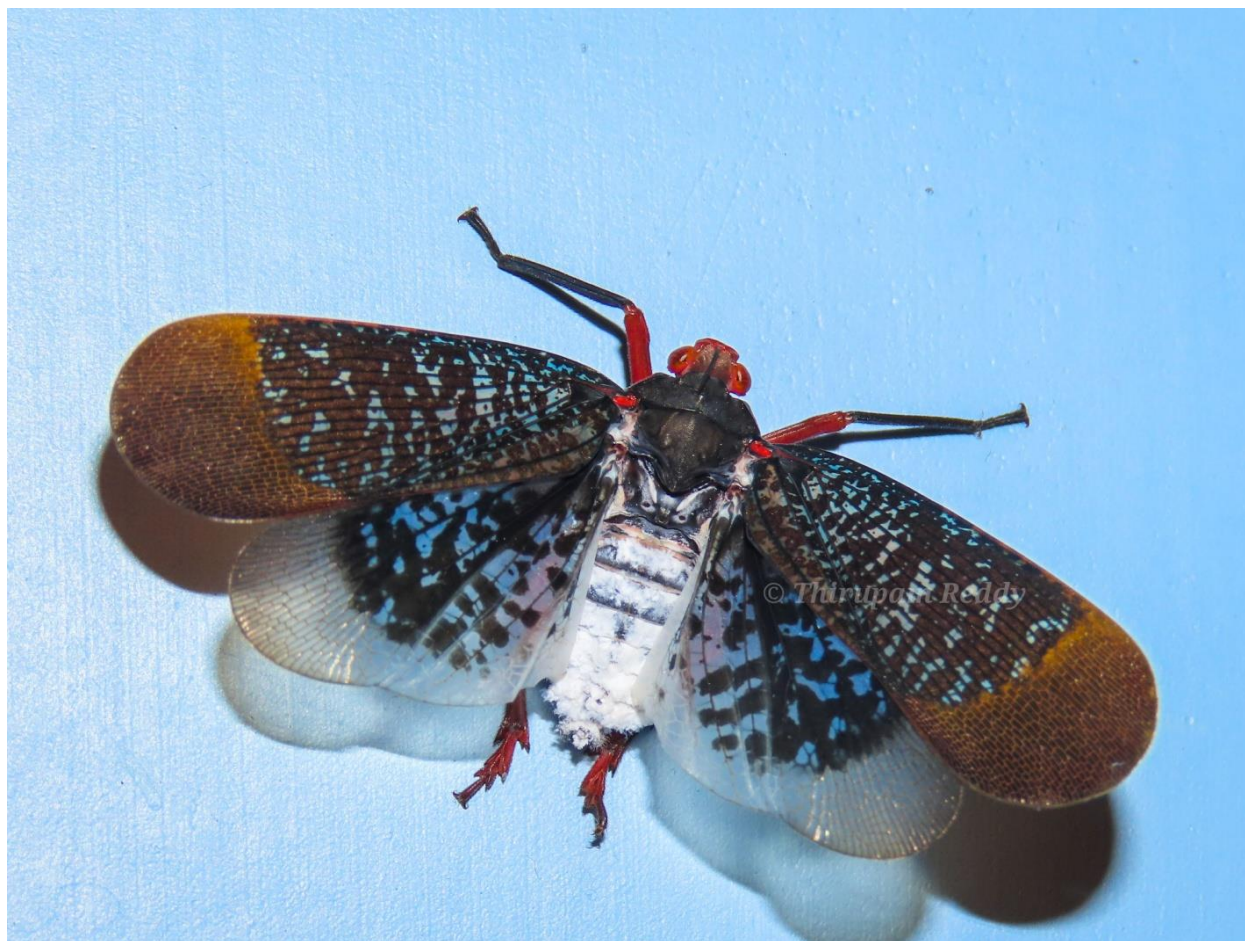
Oleander hawk moth, Daphnis nerii (Sphingidae: Lepidoptera)

Oleander hawk moth was basking in the sun on turmeric leaf stalks. Its larvae are notably green with blue eyespots, often seen munching on nerium leaves.

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Red lanternfly, *Kalidasa lantana* (Fulgoridae: Hemiptera)

The red lanternfly is identified by a slender and flexible stalk-like outgrowth arising from above the tip of its snout.

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Red spider mite: *Tetranychus* sp. (Tetranychidae: Acari)

*The red spider mites (*Tetranychus* sp.) spin the three-dimensional webs to protect themselves from biotic and abiotic factors and it is called as 'ballooning'. This mechanism also aids in their dispersal from plant to plant. The host plant is garden pea, *Pisum sativum*.*

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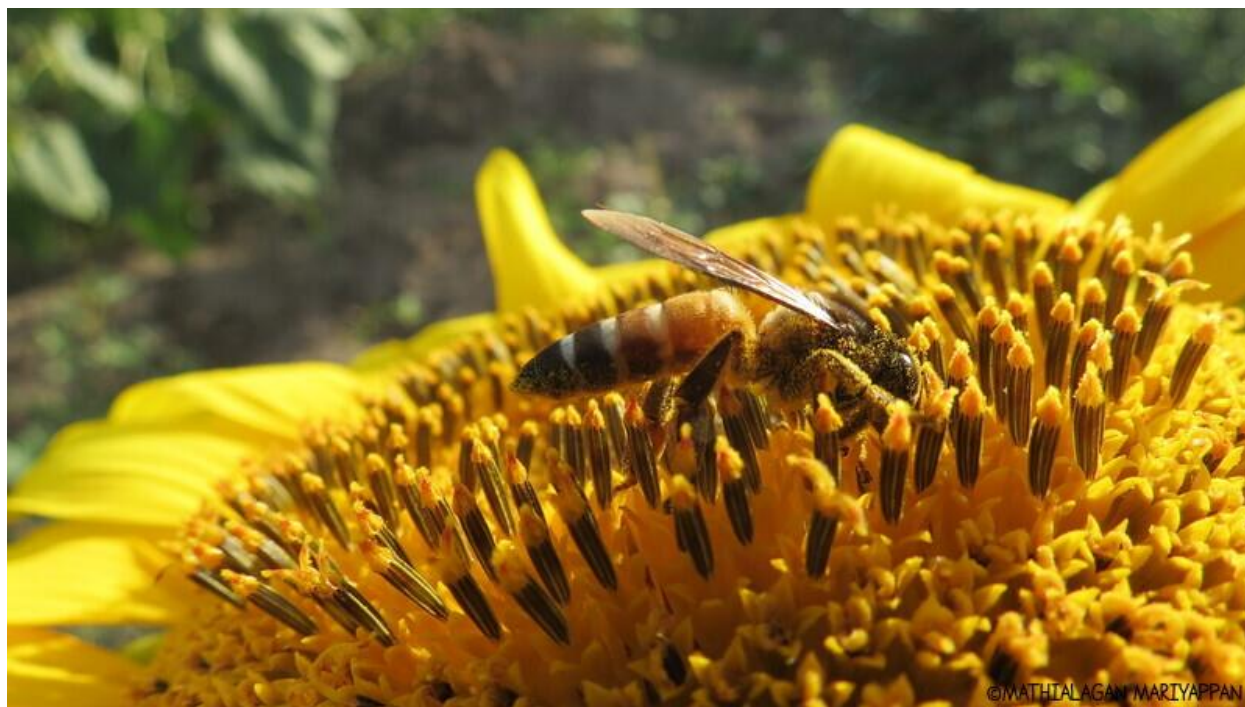
Gynandromorph of tasar silk moth, *Antheraea mylitta* (Saturniidae: Lepidoptera)

In the environment, gynandromorph phenomenon is very rare. Here in the Tasar silk moth, left side represents female (gyn) characters whereas, male (andro) characters in the right side. Gynandromorphy is thought to occur when female egg cells develop with two nuclei- so that one nucleus contains a single Z chromosome and the other contains a single W.

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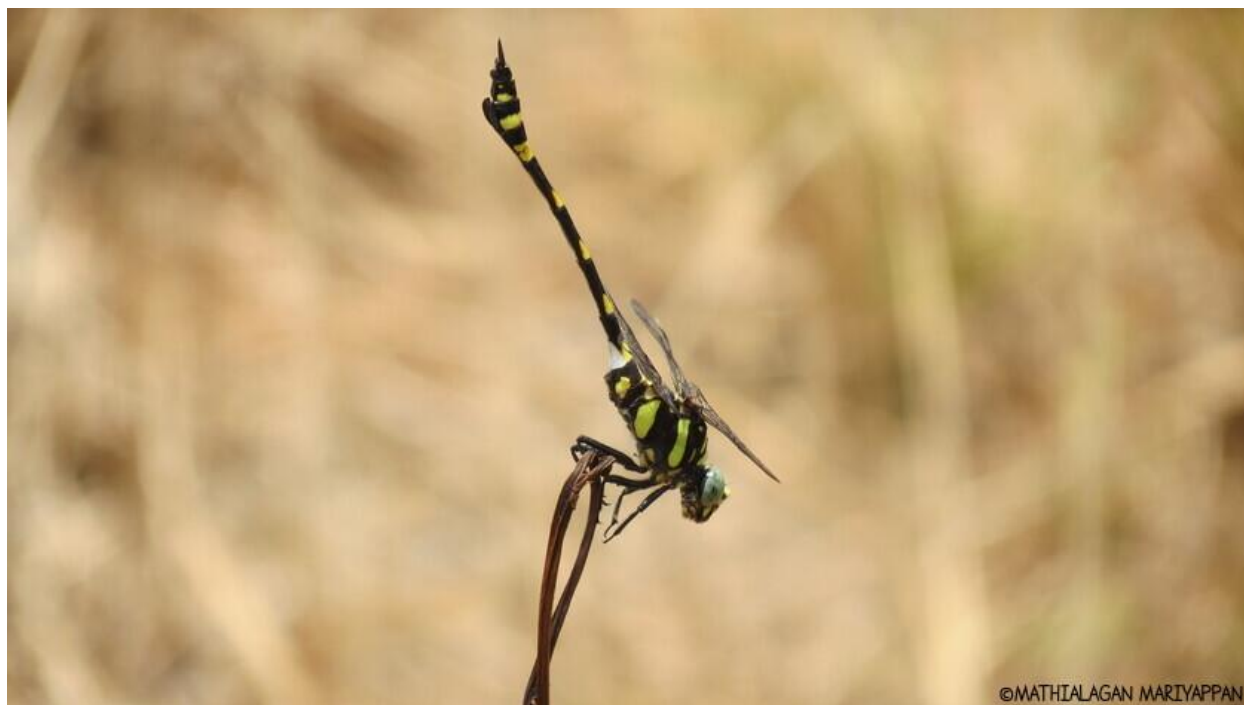
Rock bee, Apis dorsata (Apidae: Hymenoptera)

Apis dorsata is one of the most dangerous animals of the south east Asian jungles due to their threatening defensive behaviours. However, the bees are not managed for pollination, many crops throughout southern Asia depend on rock bees for pollination viz., cotton, mango, coconut, coffee, pepper, star fruit, and macadamia.

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Indian common club-tail dragonfly, Ictinogomphus rapax (Gomphidae: Odonata)

Indian common club-tail dragonfly usually perches on a bare twig facing the water, commonly found in ponds, tanks and rivers. It breeds in running and still water.

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(c) Subramanian Sevgan

The African Fig fly (AFF), *Zaprionus indianus* (Drosophilidae: Diptera)

Z. indianus is considered as cosmopolitan and measures approximately 2.5 to 3.0 mm with red eyes. *Z. indianus* is a generalist that breeds on fallen fruit and fruit on the tree, but does not attack unripe and undamaged fruits.

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Blister beetle, Mylabris pustulata (Meloidae: Coleoptera)

The beetle gets its name from its defensive secretion of Cantharidin, a blistering agent used in the treatment of warts, unwanted tattoos and the papules of Molluscum Contagiosum.

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Stalked eggs of Green Lace wing, Chrysoperla carnea

Single female of lace wing can produce 100–200 eggs. Eggs are laid on plants, usually near where aphids are present in more numbers. Each egg is hung on a slender stalk about 1 cm long, usually on the underside of a leaf. Eggs are distributed as they are highly aggressive and cannibalistic. Gardeners can attract these lacewings by growing companion plants viz., calliopsis (Coreopsis), cosmos (Cosmos), sunflowers (Helianthus) and dandelion (Taraxacum) dill (Anethum) and angelica (Angelica).

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Brinjal Epilachna beetle, Henosepilachna vigintioctopunctata (Coleoptera: Coccinellidae)

Henosepilachna vigintioctopunctata is commonly known as the 28-spotted potato ladybird https://en.wikipedia.org/wiki/Henosepilachna_vigintioctopunctata - cite note-1 or the Hadda beetle. It feeds on the foliage of potatoes and other solanaceous crops. The larvae and adults feed on the leaves by scraping the leaf cuticle, reducing the leaf surface by skeletonising the surface area, resulting in russet browning of the leaves.

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Leaf mining buprestid beetle, Trachys sp. (Buprestidae: Coleoptera)

'Trachys sp. is a beetle of the small size of 3 to 3.5 milli meters. The tiny jewel beetle mines host leaves as a larva, creating large blotches. The female lays eggs on the leaves of deciduous trees, especially elm. The larvae eat the green tissue between the upper and lower layer of the leaves making cavities called mines.'

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Red-tailed spider wasp, Tachypompilus analis (Pompilidae: Hymenoptera)

Red-tailed spider wasp found in most of tropical and subtropical partsof Asia. They prey on spiders from the families Sparassidae, Agelenidae, and Amaurobiidae.

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Crimson Marsh Glider, Trithemis aurora (Libellulidae: Odonata)

A very common dragonfly, yet beautiful in every way is Trithemis aurora. They are widely distributed species and found throughout the year across the Indian subcontinent and Southeast Asia. Common habitat of this marsh glider are weedy tanks and ponds, marshes, channels, and slow flowing streams and rivers in the lowlands and mid-hills.

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Varroa Mite, Varroa jacobsoni (Varroidae: Acari)

Varroa mite is a natural ecto-parasite of honeybee, Apis cerana. However, after having recently jumped from its natural host to the European honeybee, Apis mellifera and has emerged as a potential serious pest causing bee colony losses.

Author: Dr. M. Mathialagan, Assistant Professor (Agrl. Entomology), Sethu Bhaskara Agricultural College and Research Foundation, Karaikudi, Tamil Nadu, India.

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Rock Bee, *Apis dorsata* (Apidae: Hymenoptera)

Apis dorsata build a single, large and exposed comb under tree branches or under cliffs, instead of in cavities. They form dense aggregations at one nesting site, sometimes with up to 200 colonies in one tree. Each colony can have up to 100,000 bees and is separated by only a few centimetres from the other colonies in an aggregation.

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Mango Stem Borer, Batocera rufomaculata (Cerambycidae: Coleoptera)

Mango Stem Borer is a serious pest of fig, mango, guava, jackfruit, pomegranate and walnut in different parts of the world. Infestation may lead to yield losses and even to the death of trees. Most damage is caused by the larvae that initially bore in the tree's sub-cortex and later move deeper into the tree.

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Larva of Fruit Sucking Moth Eudocimaphalonia (Erebidae: Lepidoptera)

The adult is an agricultural pest feed on many fruit crops by piercing its proboscis into fruits and sucks the juice. The larvae tend to feed on foliage of wild host plants of the family Menispermaceae and Fabaceae.

Author: Dr. M. Mathialagan, Assistant Professor (Agrl. Entomology), Sethu Bhaskara Agricultural College and Research Foundation, Karaikudi, Tamil Nadu, India.

Location: Tamil Nadu Agricultural University, Coimbatore, Tamil Nadu, India.

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Blister beetle, Mylabris pustulata Thunberg (Meloidae: Coleoptera)

Blister beetle, as the name are known for their defensive secretion of blistering agent, cantharidin, used in folk medicine as vesicant for treating warts.

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Obituary

Dr Balwinder Singh (01/06/1956- 29/05/2022)

**The entomology fraternity mourns the demise of
India's pesticide residue and biosafety expert**

Dr Balwinder Singh, an eminent entomologist (insect toxicologist) who made notable contributions to Punjab agriculture and especially Punjab Agricultural University (PAU), Ludhiana passed away on May 29, 2022. Dr Balwinder Singh was a bright student, an outstanding teacher, a distinguished researcher in the field of pesticide residue analysis and biosafety, an inspiring mentor, an able administrator, and an academician with impeccable credentials.



Dr Balwinder Singh was born in the village Behman Jassa Singh, Tehsil Talwandi Sabo, District Bathinda, Punjab on 1st June 1956 in a traditional farming family. After graduation from Punjabi University, Patiala in 1975, he moved to the Punjab Agricultural University, Ludhiana. He distinguished himself during his Master's (1978) and Doctorate (1984) programmes in Entomology under the watchful eyes of his major advisor and renowned insect toxicologist, Dr R.L. Kalra. Starting as a research associate in 1979, he moved up to become the Senior Residue Analyst in 1998. He was largely instrumental in upgrading the Pesticide Residue Analysis Laboratory with the latest modern analytical instruments to set up a 'Centre of Excellence in Pesticide Residues' at PAU, Ludhiana. The laboratory was the first such laboratory at any State Agricultural University (SAU) to be

accredited by the NABL. The laboratory is acting as a 'core laboratory' under the Central Sector Scheme, 'Monitoring of Pesticide Residue Analysis at National Level' and as a 'referral laboratory' under the 'All India Network Project on Pesticide Residues'. Besides, the laboratory is engaged in providing regular training to the participating laboratories. He continued to head the laboratory till 2014.

Dr Singh was elevated to the position of the Head of the department of entomology in August 2011 and Director of Research, PAU, Ludhiana in October 2014. He distinguished himself in both these positions. After his superannuation at the end of May 2016, he joined as Consultant, Monitoring of Pesticide Residues at the National Level, Ministry of Agriculture and Farmers Welfare,

Government of India and continued in this position till his sad and untimely demise.

Dr Singh contributed 40 recommendations in the university Package of Practices, 91 research papers in international journals and 70 in national journals besides 6 books. He was conferred Fellowship by the Indian Society for the Advancement of Insect Science and was Vice president, of the Society of Pesticide Science India and President of the Indian Society for the Advancement of Insect Science.

Dr Singh was a Member-FAD-14, Pesticide Residue Committee, Bureau of Indian Standards, New Delhi; Member-Food

Safety Panel, BIS, New Delhi; Member Secretary-Committee constituted by Govt. of India to shortlist NABL accredited private laboratories for pesticide residue analysis in food and Member-Task force constituted by Ministry of Agriculture under the project “Monitoring of Pesticide Residues at National Level. He was also a member of the Board of Management of PAU, Ludhiana.

Dr Balwinder Singh is survived by his wife, Mrs Rupinder Kaur, daughter Sonia and son Gagandeep. With his sudden demise, the country has lost an esteemed teacher/scientist/mentor, administrator, and a wonderful human being.

Dr Manmeet B. Bhullar

Department of Entomology
Punjab Agricultural University
Ludhiana-141004

Dr Ramesh Arora

Faculty of Agriculture
Sri Guru Granth Sahib World University
Fatehgarh Sahib-140407

Dr Balwinder Singh was an outstanding scientist and always had very pleasing manners. As a close friend I will miss him. Condolences from INSECT ENVIRONMENT to his family members.

—Abraham Verghese.



From L to R: Dr Balwinder Singh, Dr BS Dhillon, Dr SP Bhardwaj and Dr HC Sharma, at the 4th Congress on Insect Science, April 16, 2015, PAU, Ludhiana



From L to R: Dr PK Chakrabarty, Dr KK Sharma, Sh. UK Singh, Dr Balwinder Singh and Dr BS Dhillon at the Annual workshop on monitoring of pesticide residues at national level, May 25, 2016, PAU, Ludhiana