

Larval age dependent parasitization performance of *Cotesia flavipes* on *Sesamia inferens*

V.K. Sonawane, S.K. Gharde*, K.S. Ghodekar, A.M. Raut, Amine Assouguem¹

Department of Entomology, School of Agriculture, Lovely Professional University, Phagwara, Jalandhar, Punjab,

¹Department of Plant Protection and Environment, National School of Agriculture, Meknes, Morocco

*Corresponding Email: sgharde@rediffmail.com

ABSTRACT

Cotesia flavipes is an important hymenopteran larval parasitoid that belongs to the family Braconidae. Its usage in pest management strategies is promising due to its parasitic impact on the larval stage of lepidopteran pests. The current investigation aims to determine the optimal host age for the parasitoid's mass proliferation and augmentative releases. The experiments showed that the female *C. flavipes* parasitizes all larval age groups of *Sesamia inferens*. Among all the larval ages, *C. flavipes* preferred second to third instars for parasitism during the spring (up to 90%) and kharif (up to 80%) seasons. There was no substantial difference in the development period between stinging, cocoon production, and the adult emergence of parasitoids. The age of the host has a substantial impact on adult longevity, with females taking longer than males. Thus, larval instars (second and third) are also recommended for high-quality mass-rearing larval parasitoids, especially *C. flavipes*, due to their strong parasitism and high net reproductive rate. Therefore, the second and third instars of *S. inferens* will recommend for the mass rearing of *C. flavipes* and the release of these parasitoids in the field as a successful bio-control program.

Key Words	<i>Cotesia flavipes</i> , <i>Sesamia inferens</i> , parasitism, mass rearing, Braconidae
DOI	https://doi.org/10.46488/NEPT.2025.v24i02.B4249 (DOI will be active only after the final publication of the paper)
Citation of the Paper	V.K. Sonawane, S.K. Gharde, K.S. Ghodekar, A.M. Raut, Amine Assouguem, 2025. Larval age dependent parasitization performance of <i>Cotesia flavipes</i> on <i>Sesamia inferens</i> . <i>Nature Environment and Pollution Technology</i> , 24(2), B4249. https://doi.org/10.46488/NEPT.2025.v24i02.B4249

INTRODUCTION

Biological control with various living organisms such as predators, parasitoids, and entomopathogens has significant potential in integrated pest management (Sedaratian-Jahromi 2021). Adult parasitoids are free-living, whereas their egg and larval stages live in or on a single host to complete their life cycle to reach the adult stage (Dodiya & Pathan, 2022; Fei et al. 2023; Holmes et al. 2023). Parasitoids eventually kill their hosts throughout the parasitism

process by inducing paralysis (Quicke 2014; Butcher & Quicke, 2023; Dai et al. 2024). It lays eggs in the hemocoel of the host insect (Jervis et al. 2023), completes larval development inside the host, and emerges just before the pupal formation in the silken cocoons (Van Noort & Broad 2024; Virla et al. 2023; Farahani et al. 2012). However, larval endoparasitoids like *Cotesia* are being used in biological pest control programs worldwide (Chepkemai et al. 2023; Parker & Kingsolver 2024) and parasitize various lepidopteran pests, including *Sesamia inferens*. Pink stem borer, *Sesamia inferens*, is a polyphagous pest reported on wheat, sorghum, oats, rice, pearl millet, barley, sugarcane, finger millet, and grasses (Jeer et al. 2021; Jadhao et al. 2020; Baladhiya et al. 2018). This pest caused damage of up to 78% in maize (Deole et al. 2017; Upadhyay et al. 2023).

Chemical communication is a very important factor in the host selection of parasitoids (Han et al. 2024). When the plant is infested by the insect pest, which emits volatile cues, that are responsible for the identification of the host location for the parasitoid (Penaflor et al. 2011; Uefune et al. 2012; Furstenberg et al. 2013; Bogka et al. 2023; Yi et al. 2023) or emitted by herbivores and their frass after feeding on plants (de Rijk et al. 2016; Liu et al. 2024), host preference could influence the host-parasitoid interaction (Furstenberg-Hagg et al. 2013), the development, success of reproduction, and the percentage of survival of the parasitoid on the hosts in field condition (Kher et al. 2024; Han et al. 2013; Ehteshami et al. 2023; Gomes et al. 2024). Consequently, numerous parasitoids exhibit host preferences based on particular stages of host or species and enhance the percentage of parasitism in invitro conditions (Stacconi et al. 2013; Yazdani et al. 2015; Hegazi et al. 2024; Kher et al. 2024).

The ability of parasitoids to search behaviour is based on the parasitism rate against hosts (Hardy & Wajnberg 2023). After the parasitism of the host by the parasitoid, it goes through various phases such as host habitat location, host location, host acceptance, host suitability, and host regulation (Harris et al. 2012; Kuramitsu et al. 2019; Benrey 2023; Kathirvelu et al. 2024; Fei et al. 2023; Abram et al. 2023). The steps to be followed for effective colonization of a host by a parasitoid according to chemical cues are employed by the male wasps to find conspecific females in the field or an artificial cage (Malek et al. 2021). Certain parasitoids possess the capability to locate potential mates over a considerable distance through the emission of highly volatile pheromones, whereas, in certain instances, less volatile pheromones are employed for short-range (Ruther 2013). It is believed that male wasps may use cues other than female pheromones to locate females, other cues that help to find mates through host-associated volatiles or host-induced plant volatiles (Reddy & Guerrero 2004; Xu et al. 2017). Female parasitoids use plant volatiles to locate host habitats (McCormick et al. 2012; Wäschke et al. 2013; Turlings & Erb 2018). Some insects also take volatile stimuli from the habitat for mating (male dung flies are attracted to the odor of fresh cow manure) (Parker 1978; McAuslane et al. 1990; Webster & Carde 2017).

The biocontrol program is successful when bio-agents are available at ground level and natural enemies are mass culturing in the laboratory. The natural population available in the field will not survive for a longer period due to the application of pesticides to protect the crops. In agriculture,

crops are protected through synthetic pesticides, causing several problems like the development of resistance and resurgence in pests, habitat loss, the decline of ecological service members, environmental pollution, and many more (Onaho et al. 2023; Ahmad & Akhtar 2016; Shimeng et al. 2001; Mruthunjayaswamy et al. 2016; Nwankwo et al. 2016). Therefore, the laboratory mass culture of braconid wasps requires lepidopteran larvae as a feeding source. However, one issue encountered during laboratory production of *Cotesia flavipes* is parasitoid quality (Prezotti & Parra 2002). The parasitoid's inferior quality is the reason for its production failure and lack of demand (Ramalho et al. 2012). Hence, the knowledge of the parasitoid is most important for the suitable host age used for augmentative parasitoid releases (Li et al. 2006). Therefore, the present study mainly focuses on the performance of *Cotesia flavipes* on parasitization and net reproductive rate on different larval instars of the pink stem borer, *Sesamia inferens*, to enhance the mass-rearing process of parasitoids.

Material and Methods:

The experiment entitled Parasitization performance of *Cotesia flavipes* on different larval instars of pink stem borer, *Sesamia inferens* was conducted under laboratory conditions, at the School of Agriculture, Lovely Professional University, Phagwara (Punjab). The experiments were conducted during the spring and kharif seasons of 2023.

Mother culture of *Sesamia inferens* Walker

A fourth and fifth instar of the pink stem borer, *Sesamia inferens* Walker, were collected from a maize field and reared under laboratory conditions. The insects were kept in in-vivo conditions at 26 ± 2 °C, RH $65 \pm 5\%$, and a 12L:12D photoperiod. The emerged adult from this culture was reared on a maize plant in the ovipositional chamber (Kumar et al. 2011), by providing 10 % honey solution to the emerged adults. After five to six days, remove the potted plant from the ovipositional chamber, separate the leaf sheath from the plant where the egg mass was, and rear in the incubator up to the hatching of neonate larvae. These larvae were reared on a maize stem (7×2.5 cm) in tubes, with this method, culture was maintained for future study (Sharma et al. 2017). The old stems were replaced by fresh stems every day. Two generations were reared under laboratory conditions before starting the experiments.

Mother culture of *Cotesia flavipes*

Parasitized cadavers were collected from the maize field and brought into the laboratory for further mass culturing. The cocoons were kept at $26 \pm 2^{\circ}\text{C}$, RH $65 \pm 5\%$, with a photometer for 12 hours of equal day and night length. The cocoons of parasitoids were reared in the rearing chamber by covering them with brown paper (22×15×5 cm). The observations were recorded daily until adults emerged. Emerged adults were then shifted to the rearing cage by providing a 20% honey solution in cotton swabs as food for adult parasitoids. After 24 hours, keep the old larvae separate and the fresh larvae with the parasitoids, and continue this process till the death of the parasitoid. All parasitized larvae of *S. inferens* were reared at $26 \pm 2^{\circ}\text{C}$, RH $65 \pm 5\%$, with a photoperiod of 12 hours equal to the length of day and night. The honey solution was provided daily to the adult parasitoid, and three generations of *Cotesia flavipes* were reared in laboratory conditions with the same procedure.

Preference of *Cotesia flavipes* on different larval instars

To study the preference of female parasitoid *Cotesia flavipes* on different larval instars of pink stem borer, a pair of adult parasitoids were released for 24 hours in a glass tube for mating (Saini et al. 2019) with a 20% honey solution in a cotton swab as food. For mating, a no-choice experiment was carried out, with 10 larvae of each instar in a rearing box (22 × 15 × 5 cm). The separation of the larvae was done (instar-wise) according to their size and colour (Sharma et al. 2017; Nagarjuna et al. 2015). The parasitized host larvae of different instars of pink stem borer were then reared individually and observed until the pupation and emergence of the adult parasitoid.

Impact on developmental biology of *Cotesia flavipes*

The experiment on the developmental biology of the parasitoid *Cotesia flavipes* on the larval instars of pink stem borer was investigated through a series of experiments involving the parasitization of various larval instars of stem borer. The eclosion of adults was sorted by sex, and a pair of parasitoids were exposed separately to batches of ten larvae of host II to VI instars in a no-choice experimental design. After 24 hours, the old larvae were replaced with new ones. The old larvae were reared according to the prescribed procedure to facilitate the emergence of parasitoids (Saini et al. 2019). Since immature stages (egg and larvae) of the parasitoid were present in the body of the host, various parameters like duration of egg and larval stages (stinging to cocoon formation stage), pre-pupa and pupal stage (cocoon stage), longevity of adults, and sex ratio were recorded throughout the experimental period.

Statistical analysis:

Different parameters recorded during the study were subjected to one-way analysis of variance (ANOVA) using SPSS 22.0 software. We investigated the interactions of season and instar on the parasitization and developmental period of *Cotesia flavipes*, considering season and instar as fixed effects. The means that showed significant differences were distinguished using the least significant difference (LSD) method.

Result

Parasitization rate of *Cotesia flavipes* on different larval instars:

The *C. flavipes* parasitization rate significantly differs between the different larval instars of *S. inferens*. The most preferred stage and the highest level of parasitism (80% and 90%) were observed in third-instar larvae in the spring ($F(4,9) = 3.68$; $p = 0.011$) and kharif ($F(4,9) = 5.07$; $p = 0.002$) seasons, respectively (Fig 1). The parasitization rate gradually decreased after the third instars until the sixth instars during both seasons. However, there were no statistically significant impact of season on parasitization of *C. flavipes* in respect with larvae host age.

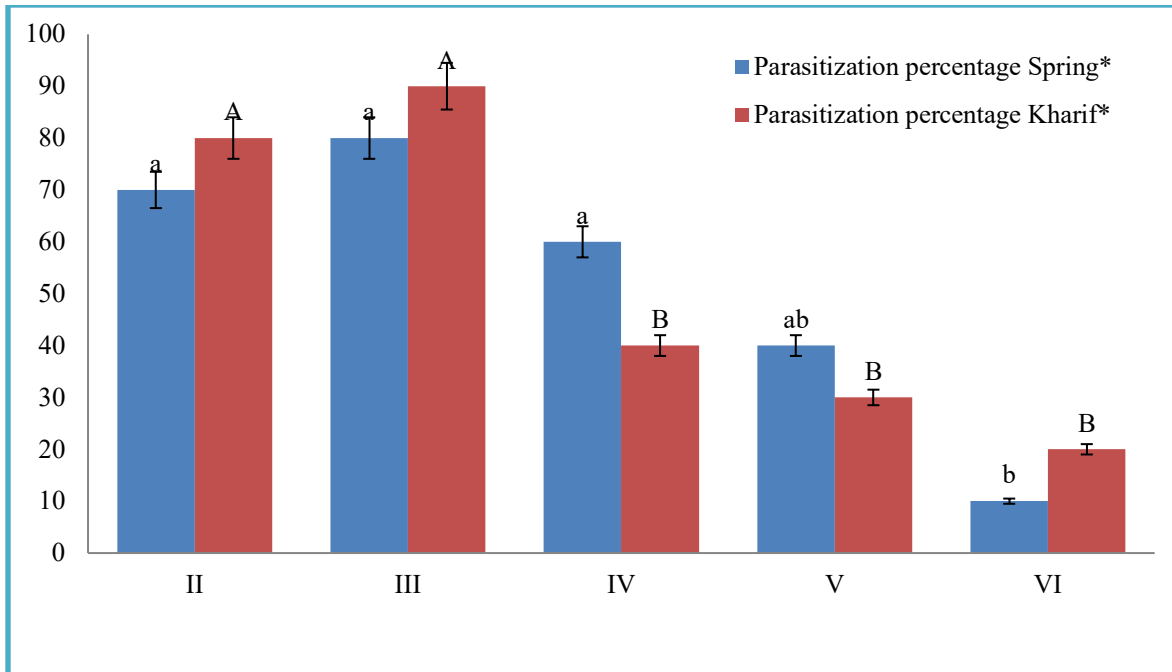


Figure 1 Parasitization capability of *Cotesia flavipes* on different larval instars. ^{a,A}values with same letters in a column are non-significant as per DMRT. ‘*’ significant at 5% level of significance

Impact on developmental biology of *Cotesia flavipes*:

The development period of immature parasitoids on host age was not significantly different ($p > 0.05$) in both seasons (Table 1). The age of the parasitized host showed a substantial effect on adult longevity ($p < 0.05$), with females living longer in both seasons. In *spring* 2023, the third instars had the longest development period, followed by the second, fourth, fifth, and sixth instars and a similar pattern seen in *Kharif* 2023 (Table 1).

Table 1 Development time and adult longevity of parasitoids on different ages of host.

Spring 2023					
Host instars	Development time (days) \pm SD			Adult longevity (days) \pm SD	
	Stinging to cocoon formation	Cocoon stage	Stinging to Adult emergence	Male	Female
II	8.2 ± 4.42^a	3.3 ± 1.83^{ab}	11.5 ± 6.13^a	4.9 ± 3.48^a	5.1 ± 3.57^a
III	8.5 ± 7.40^a	4.0 ± 3.5^a	12.5 ± 10.8^a	5.2 ± 2.86^a	5.5 ± 3.10^a
IV	7.1 ± 5.02^{ab}	3.0 ± 3.89^{ab}	10 ± 7.04^{ab}	4.5 ± 4.00^a	4.7 ± 4.1^a
V	6.2 ± 8.01^{ab}	2.9 ± 2.08^{ab}	9.2 ± 11.89^{ab}	2.9 ± 3.78^b	3.2 ± 4.21^{ab}

VI	1.7 ± 5.38^b	0.8 ± 2.53^b	2.5 ± 7.91^b	0.7 ± 2.21^b	0.9 ± 2.85^b
F _{value}	2.25	2.00	2.17	3.72	3.19
P _{value}	0.116 ^{NS}	0.159 ^{NS}	0.129 ^{NS}	0.022*	0.041*
Kharif 2023					
Host instars	Development time (days)± SD			Adult longevity (days) ± SD	
	Stinging to cocoon formation	Cocoon stage	Stinging to Adult emergence	Male	Female
II	$8.0 \pm 4.37^{a\#}$	3.5 ± 1.90^a	11.5 ± 6.15^a	4.7 ± 2.71^a	5.6 ± 3.17^a
III	8.8 ± 7.60^a	4.0 ± 3.50^a	12.8 ± 11.08^a	5.1 ± 3.03^a	5.3 ± 2.95^a
IV	6.0 ± 3.27^a	3.2 ± 4.16^a	9.2 ± 11.93^a	4.0 ± 3.53^{ab}	4.3 ± 3.74^{ab}
V	6.0 ± 7.79^a	2.9 ± 1.66^a	8.9 ± 4.91^a	2.5 ± 3.37^{ab}	2.8 ± 3.65^{ab}
VI	3.1 ± 6.54^a	1.7 ± 3.59^a	4.8 ± 10.13^a	1.2 ± 2.57^b	1.4 ± 2.95^b
F _{value}	1.27	0.761	1.088	2.803	2.44
P _{value}	0.293 ^{NS}	0.556 ^{NS}	0.374 ^{NS}	0.037*	0.035*

[#]The values with same letters in a column are non-significant as per DMRT.

‘*’significant at 5% level of significance (P<0.05), NS - non-significant

Net reproductive rate:

The maximum oviposition period of *C. flavipes* was up to 96 hours observed in the second and third instars and then gradually decreased with host age in the *spring* season, and the same trend was observed in the *Kharif* season (Fig. 2). However, the female adult who emerged from the cocoon was twice as old as the male adult in all the time periods and the age of the host in both seasons. Overall, there was no significant variation in the cocoon-forming population and adult emergence between seasons (Fig 2) . The sex ratio was 0.40:1 and 0.37:1 (male: female) found in the *spring* and *kharif* seasons, respectively.

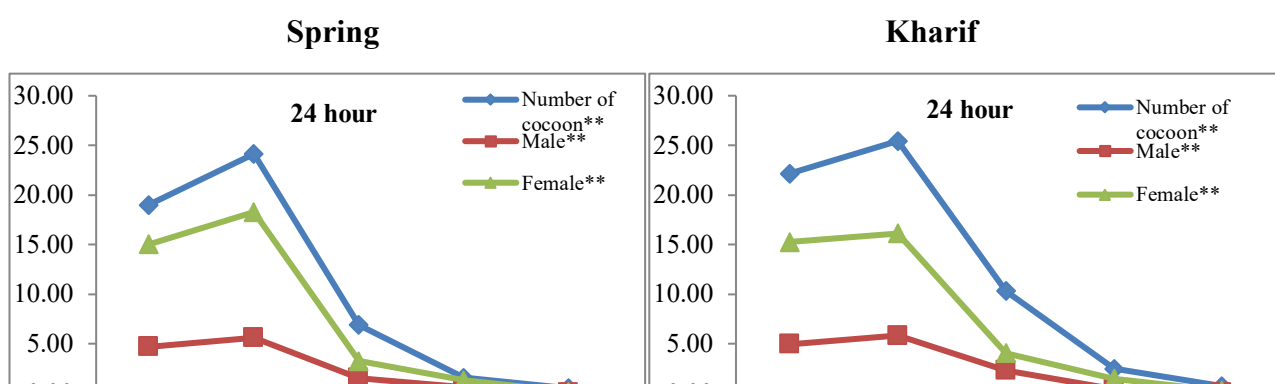


Figure 2 Variation in the larval Parasitoid Reproductive Rate Across Different Host Ages

‘***’ Significant level at $P \leq 0.01$

Discussion

Bio-agents have the potential to check pest populations in the available ecosystem, and therefore they are an important tool in the integrated pest management (IPM) system (Sampaio et al.

2009). The biological control includes predators, parasitoids, and entomopathogens successfully accommodated in the environment by the availability of suitable habitat, the application of ecofriendly pesticides, and etc. Among the natural enemies, parasitoids are usually the best option for pest management due to their high host specificity and free-living behaviours (Colmenarez et al. 2018; Trejo & Contreras 2024). Therefore, mass culturing of parasitoids is an important step, and a temperature of 25– 30°C indicates a faster development rate (Adamo et al. 2012; Huey et al. 2012; Moore et al. 2020). However, the mass production of *Cotesia* species in the laboratory is very cost-effective for pest management (Zang et al. 2021). Our research study showed that the *Cotesia flavipes* preferred the second and third instars of the host for parasitism at a temperature 25°C due to the availability of high protein, sugar, and lipid content in early instars larvae (Shekharappa & Kulkarni 2003; Jervis et al. 2008; Silva-Torres et al. 2009; Sokame et al. 2021), however, the old age instars were not preferred by the female *C. flavipes* for parasitization because the old larval instars developed defense mechanisms themselves and were aggressive in nature, which led to the death of parasitoids (Potting et al. 1997).

Abiotic factors, especially temperature, play a crucial role in parasitoid growth and development (Adamo et al. 2012; Huey et al. 2012). However, parasitoid wasp emergence and survival rate were maximum at 20°C and 25°C but when it exceeded 30 °C exhibited the rate of wasp emergence and survival rate (Moore et al. 2020). The longer development process, cocoon formation, and duration from stung to the emergence of adult parasitoids were observed at 25 °C in the present investigation. A similar result was observed in *Cotesia* species, development periods were longer in the second instar of *Chilo partellus* (Sarkar et al. 2020; Khan et al. 2017), and the formation of the highest number of parasitoid cocoons was recorded in the third instar of *Spodoptera frugiperda* (Obala et al. 2023).

Augmentation is the release of natural enemies or environmental manipulation to increase the effectiveness of naturally occurring natural enemies (Hoy 2008). Environmental manipulation can include supplying alternate hosts or prey, food or nesting locations, or changing agricultural methods to benefit bioagents (Hoy 2008). The periodic release of natural enemies is evaluated based on fitness factors such as offspring sex ratio (Mohamad et al. 2015; Kruitwagen et al. 2018). Our research reported that males emerged early as compared to females, and the population of female adults was almost double than the male adults. This result may be due to mated female parasitoids being able to manage progeny sex ratios by regulating fertilization during oviposition. The present result supported by (Heimpel & de Boer 2008), revealed that females develop from fertilized eggs and are diploid, while males develop from unfertilized eggs and are haploid. However, the previous research showed that the female adult emerged more than the male adult of *Cotesia* sp. when the host was *Spodoptera frugiperda* (Obala et al. 2023) and *S. littoralis* (Agbodzavu et al. 2018).

Conclusion

The present investigation concludes that *Cotesia flavipes* preferred the second or third instars of *Sesamia inferens* for parasitization. These larval stages are used for mass rearing of *C. flavipes*

in laboratory conditions and are periodically relived in the field to manage the lepidopteron pest. In laboratory conditions, a temperature of 25–30°C will enhance the fecundity, cocoon formation, and emergence rate of *C. flavipes*. However, for the conservation of parasitoids, we will provide alternate hosts and food by planting nectarine plants and avoiding the application of non-selective pesticides. The conservation of the larval parasitoid *C. flavipes* enhances the efficacy of biological pest management strategies aimed at mitigating the impact of lepidopteran pest infestations in field conditions.

Acknowledgement

All the authors are thankful to Lovely Professional University for providing the facility for conducting the experiments.

Disclosure statement

No potential conflict of interest was reported by the author(s).

Author contribution

VKS- design, conducting the experiment and data collection, SKG- concept formulation, monitoring the experiments and preparation manuscript, KSG- data collection and mother culture maintenance, AMR, SKG and AA - data validation, data analysis, finalizing the manuscript, result interpretation

Reference

- Abram, P. K. Haye, T. Clarke, P. Grove, E. Thiessen, J. & Garipey, T. D. (2023). Partial refuges from biological control due to intraspecific variation in protective host traits. *Ecological Applications*, 33(4), e2796.
- Adamo, S. A. Baker, J. L. Lovett, M. M. & Wilson, G. (2012). Climate change and temperate zone insects: the tyranny of thermodynamics meets the world of limited resources. *Environmental Entomology*, 41(6), 1644-1652.
- Agbodzavu, M. K. Lagat, Z. O. Gikungu, M. Rwomushana, I. Ekesi, S. & Fiaboe, K. K. M. (2018). Performance of the newly identified endoparasitoid *Cotesia icipe* Fernandez-Triana & Fiaboe on *Spodoptera littoralis* (Boisduval). *Journal of applied entomology*, 142(7), 646-653.
- Ahmad, M. & Akhtar, S. (2016). Development of resistance to insecticides in the invasive mealybug *Phenacoccus solenopsis* (Hemiptera: Pseudococcidae) in Pakistan. *Crop Protection*, 88, 96–102.
- Baladhiya, H. C. Sisodiya, D. B. & Pathan, N. P. (2018). A review on pink stem borer, *Sesamia inferens* Walker: a threat to cereals. *Journal of Entomology and Zoology Studies*, 6(3), 1235-1239.
- Benrey, B. (2023). The effects of plant domestication on the foraging and performance of parasitoids. *Current Opinion in Insect Science*, 57, 101031.
- Bogka, G. Anastasaki, E. Milonas, P. G. Psoma, A. Kabourakis, E. M. Zwaan, B. J. Pannebakker, B. A. & Fatouros, N. E. (2023, June 16). Chemical cues involved in the host foraging behavior of *Psytalia concolor* wasps to locate the olive fruit fly *Bactrocera oleae*. *Frontiers in Ecology and Evolution*, 11.

- Butcher, B. A. (2023). *Parasitoid Wasps of South East Asia*. GB: CABI.
- Colmenarez, Y. Corniani, N. Mundstock, S. Sampaio, M. & Vásquez, C. (2018). Use of parasitoids as a biocontrol agent in the neotropical region: challenges and potential. In *Horticultural crops* (pp. 171-185). London: IntechOpen.
- de Rijk, M. Krijn, M. Jenniskens, W. Engel, B. Dicke, M. & Poelman, E. H. (2016). Flexible parasitoid behaviour overcomes constraint resulting from position of host and nonhost herbivores. *Animal Behaviour*, 113, 125-135.
- Deole, S. Dubey, V. K. Rana, D. K. & Gauraha, R. (2017). Evaluation of newer insecticides against maize pink stem borer: major constraint insect pest of maize in Raipur, Chhattisgarh. *Journal of Plant Development Sciences*, 9(4), 335-339.
- Dodiya, R. D. & Pathan, N. P. (2022). Host Searching Behaviour of Parasitoids and Predators. *A Monthly Peer Reviewed Magazine for Agriculture and Allied Sciences*, 14.
- Ehteshami, F. Aleosfoor, M. Allahyari, H. Kavousi, A. & Fekrat, L. (2023). Comparative demography, population projection, functional response and host age preference behavior of the parasitoid *Goniozus legneri* on two lepidopterous insect hosts. *Egyptian Journal of Biological Pest Control*, 33(1), 2.
- Farahani, H. K. Bell, H. & Goldansaz, S. H. (2012). Biology of *Apanteles myeloenta* (Hymenoptera: Braconidae), a larval parasitoid of carob moth *Ectomyelais ceratoniae* (Lepidoptera: Pyralidae). *Journal of Asia-Pacific Entomology*, 15(4), 607-610.
- Fei, M. Gols, R. & Harvey, J. A. (2023). The biology and ecology of parasitoid wasps of predatory arthropods. *Annual review of entomology*, 68, 109-128.
- Fürstenberg-Hägg, J. Zagrobelny, M. & Bak, S. (2013). Plant defense against insect herbivores. *International journal of molecular sciences*, 14(5), 10242-10297.
- Gao, Y. Hu, Y. Fu, Q. Zhang, J. Oppert, B. Lai, F. ... & Zhang, Z. (2010). Screen of *Bacillus thuringiensis* toxins for transgenic rice to control *Sesamia inferens* and *Chilo suppressalis*. *Journal of invertebrate pathology*, 105(1), 11-15.
- George-Onaho, J. a. 1 Moore, J.C.2 *Haastrup, N.O.1 Agboola, I.S.1 Ete, J.A.1 and Ajao, O.I.3 (2023). Insecticidal Efficacy of *Tithonia Diversifolia* Extracts Against Pink Stem Borer (*Sesamia Calamistis* (Hampson)).Ethiopian Journal of Environmental Studies & Management, 16(1), 117-123.” n.d.)
- Gomes, E. Lemaître, J. F. Rodriguez-Rada, V. Débias, F. Desouhant, E. & Amat, I. (2024). Foraging at night under artificial light: impacts on senescence and lifetime reproductive success for a diurnal insect. *Animal Behaviour*, 210, 85-98.
- Gornard L, S. Mougél, F. Germon, I. Borday-Birraux, V. & Kaiser, L. (2024). Cellular Dynamics of Host-Parasitoid Interactions: Insights from the Encapsulation Process in a Partially Resistant Host. *Florence and Germon, Isabelle and Borday-Birraux, Véronique and Kaiser, Laure*.
- HAN, L. B. Huang, L. Q. & Wang, C. Z. (2013). Host preference and suitability in the endoparasitoid *Campoletis chloridae* is associated with its ability to suppress host immune responses. *Ecological Entomology*, 38(2), 173-182.

- Hardy, I. C. & Wajnberg, E. (2023, December 11). *Jervis's Insects as Natural Enemies: Practical Perspectives*. Springer Nature. <http://books.google.ie/books?>
- Harris, C. M. Ruberson, J. R. Meagher, R. & Tumlinson, J. H. (2012). Host suitability affects odor association in *Cotesia marginiventris*: implications in generalist parasitoid host-finding. *Journal of chemical ecology*, 38, 340-347.
- Hegazi, E. & Khafagi, W. (2024). Host-instar selection, interspecific competition, and reproductive capacity of extant and novel parasitoids (Hymenoptera: Braconidae) on Egyptian cotton leafworm. *Egyptian Journal of Biological Pest Control*, 34(1), 10.
- Heimpel GE, de Boer JG (2008) Sex determination in the Hymenoptera. *Annual Review of Entomology* 53: 209–230.
- Holmes, L. A. Nelson, W. A. & Loughheed, S. C. (2023). Strong effects of food quality on host life history do not scale to impact parasitoid efficacy or life history. *Scientific Reports*, 13(1), 3528.
- Hoy, M.A. (2008). Augmentative Biological Control. In: Capinera, J.L. (eds) *Encyclopedia of Entomology*. Springer, Dordrecht.
- Huey, R. B. Kearney, M. R. Krockenberger, A. Holtum, J. A. Jess, M. & Williams, S. E. (2012). Predicting organismal vulnerability to climate warming: roles of behaviour, physiology and adaptation. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 367(1596), 1665-1679.
- Jadhao, K. R. Bansal, A. & Rout, G. R. (2020). Silicon amendment induces synergistic plant defense mechanism against pink stem borer (*Sesamia inferens* Walker.) in finger millet (*Eleusine coracana* Gaertn.). *Scientific Reports*, 10(1), 4229.
- Jeer, M. Yele, Y. Sharma, K. C. & Prakash, N. B. (2021). Exogenous application of different silicon sources and potassium reduces pink stem borer damage and improves photosynthesis, yield and related parameters in wheat. *Silicon*, 13(3), 901-910.
- Jervis, M. A. Ellers, J. & Harvey, J. A. (2008). Resource acquisition, allocation, and utilization in parasitoid reproductive strategies. *Annu. Rev. Entomol.* 53, 361-385.
- Kathirvelu, C. Manickavasagam, S. & Gopianand, L. (2024). Host Preference of *Xanthopimpla Saussure* Parasitising *Sesamia inferens* Walker. *Indian Journal of Entomology*, 1-7.
- Khan, J. Ali, R. Blouch, A. Javed, H. I. Haq, E. U. Mehmood, T. & Rasool, A. (2017). Biological parameters of *Cotesia flavipes* reared on different larval instars of *Chilo partellus* under laboratory conditions. *Journal of Entomology and Zoology Studies*, 5(3), 829-832.
- Kher, S. V. Kulkarni, S. S. Dosdall, L. M. & Cárcamo, H. A. (2024). Life history and host preferences of *Tetrastichus julis* (Walker)(Hymenoptera: Eulophidae), the principal parasitoid of the cereal leaf beetle, *Oulema melanopus* (L.)(Coleoptera: Chrysomelidae). *Biological Control*, 188, 105432.
- Kruitwagen, A. Beukeboom, L. W. & Wertheim, B. (2018). Optimization of native biocontrol agents, with parasitoids of the invasive pest *Drosophila suzukii* as an example. *Evolutionary Applications*, 11(9), 1473-1497.
- Kumar, P. Suby, S. B. Sekhar, J. C. and Kumar R. S. (2011). A Collapsible insect rearing cage.

Patent Application No. 0923/DEL/2011

- Kuramitsu, K. Vicencio, E. J. M. & Kainoh, Y. (2019). Differences in food plant species of the polyphagous herbivore *Mythimna separata* (Lepidoptera: Noctuidae) influence host searching behavior of its larval parasitoid, *Cotesia kariyai* (Hymenoptera: Braconidae). *Arthropod-plant interactions*, 13(1), 49-55.
- Li, J. Coudron, T. A. Pan, W. Liu, X. Lu, Z. & Zhang, Q. (2006). Host age preference of *Microplitis mediator* (Hymenoptera: Braconidae), an endoparasitoid of *Mythimna separata* (Lepidoptera: Noctuidae). *Biological Control*, 39(3), 257-261.
- Liu, G. Wang, Q. Chen, H. Wang, Y. Zhou, X. Bao, D. ... & Fu, J. (2024). Plant-derived monoterpene S-linalool and β -ocimene generated by CsLIS and CsOCS-SCZ are key chemical cues for attracting parasitoid wasps for suppressing *Ectropis obliqua* infestation in *Camellia sinensis* L. *Plant, Cell & Environment*, 47(3), 913-927.
- Malek, R. Kaser, J. M. Anfora, G. Ciolli, M. Khimian, A. Weber, D. C. & Hoelmer, K. A. (2021). *Trissolcus japonicus* foraging behavior: Implications for host preference and classical biological control. *Biological Control*, 161, 104700.
- McAuslane, H. J. Vinson, S. B. & Williams, H. J. (1990). Influence of host plant on mate location by the parasitoid *Campoletis sonorensis* (Hymenoptera: Ichneumonidae). *Environmental Entomology*, 19(1), 26-31.
- McCormick, A. C. Unsicker, S. B. & Gershenson, J. (2012). The specificity of herbivore-induced plant volatiles in attracting herbivore enemies. *Trends in plant science*, 17(5), 303-310.
- Mohamad, F., Mansour, M., & Ramadan, A. (2015). Effects of biological and environmental factors on sex ratio in *Ascogaster quadridentata* Wesmael (Hymenoptera: Braconidae), a parasitoid of *Cydia pomonella* L.(Tortricidae). *Journal of Plant Protection Research*. 55(2):151-155.
- Moore, M. E. Kester, K. M. & Kingsolver, J. G. (2020). Rearing temperature and parasitoid load determine host and parasitoid performance in *Manduca sexta* and *Cotesia congregata*. *Ecological Entomology*, 45(1), 79-89.
- Mruthunjayaswamy, P. Thiruvengadam, V. & Kumar, J. S. (2016). Resistance in *Maconellicoccus hirsutus* (Green) in India to selected insecticides and quantification of detoxifying enzymes imparting resistance. *Crop Protection*, 89, 116-122.
- Nagarjuna, B. Manjunatha, M. & Latha, M. (2015). Biology of maize stem borer, *Sesamia inferens* (Walker) Noctuidae: Lepidoptera. *Journal of eco-friendly Agriculture*, 10(1), 90-91.
- Nwankwo, E. N. Onuseleogu, D. C. Ogbonna, C. U. & Okorochoa, A. O. E. (2016). Effect of neem leaf extracts (*Azadirachta indica*) and synthetic pesticide (Carbofuran) on the root-knot nematode (*Meloidogynes* pp.) of cowpea (*Vigna unguiculata* L. Walp). *Int. J. Ento. Res*, 3(1), 01-06.
- Obala, F. Mohamed, S. A. Magomere, T. O. & Subramanian, S. (2023). Old and new association of *Cotesia icipe* (Hymenoptera: Braconidae) with alien invasive and native *Spodoptera* species and key stemborer species: implication for their management. *Pest Management Science*, 79(12), 5312-5320.

- Parker, A. L. & Kingsolver, J. G. (2024). The interactive effects of heat stress, parasitism and host plant quality in a host–parasitoid system. *Functional Ecology*.
- Parker, G. A. (1978). Evolution of competitive mate searching. *Annual review of entomology*, 23(1), 173-196.
- Penaflor, M. F. G. V. Erb, M. Miranda, L. A. Werneburg, A. G. & Bento, J. M. S. (2011). Herbivore-induced plant volatiles can serve as host location cues for a generalist and a specialist egg parasitoid. *Journal of Chemical Ecology*, 37, 1304-1313.
- Potting, R. P. J. Overholt, W. A. Danso, F. O. & Takasu, K. (1997). Foraging behavior and life history of the stemborer parasitoid *Cotesia flavipes* (Hymenoptera: Braconidae). *Journal of Insect Behavior*, 10, 13-29.
- Prezotti L and Parra JP. 2002. Controle de qualidade em criações massais de parasitoides e predadores In: Parra, JRP, Botelho PSM, Corrêa-Ferreira BS & Bento JMS (Eds.). Controle biológico no Brasil: parasitoides e predadores. Manole, São Paulo, 295-296.
- Quicke, D. L. (2014). The braconid and ichneumonid parasitoid wasps: biology, systematics, evolution and ecology. *John Wiley & Sons*.
- Ramvalho, D. G., Viel, S. R., Vacari, A. M., De Bortoli, S. A., Lopes, M. M., Laurentis, V. L., & Veiga, A. C. P. (2012). Criteria for optimization of mass rearing of the parasitoid *Cotesia flavipes* in the laboratory. *Journal of Research in Biology*, 2(5), 529-536.
- Reddy, G. V. & Guerrero, A. (2004). Interactions of insect pheromones and plant semiochemicals. *Trends in plant science*, 9(5), 253-261.
- Ruther, J. (2013). Novel insights into pheromone mediated communication in parasitic hymenopterans. *Chemical ecology of insect parasitoids*, 112-144.
- Saini, A. Sharma, P. L. & Chandel, R. S. (2019). Host age influence on the parasitism of the species *Cotesia vestalis* (Haliday) (Hymenoptera: Braconidae). *Egyptian Journal of Biological Pest Control*, 29, 1-6.
- Sampaio, M. V. Bueno, V. H. P. Silveira, L. C. P. & Auad, A. M. (2009). Biological control of insect pests in the tropics. *Tropical biology and conservation management*, 3, 28-70.
- Sarkar, S. Babu, A. Chakraborty, K. & Deka, B. (2020). Biology and life history of *Cotesia ruficrus* (Hymenoptera: Braconidae) a potential parasitoid of *Hyposidra talaca* (Lepidoptera: Geometridae) larvae, a major tea pest. *Journal of Biopesticides*, 13(1), 79-84.
- Sedaratian-Jahromi, A. (2021). Effects of entomopathogens on insect predators and parasitoids. *Microbes for Sustainable Insect Pest Management: Hydrolytic Enzyme & Secondary Metabolite*—Volume 2, 183-231.
- Sharma, H. Jaglan, M. S. & Yadav, S. S. (2017). Biology of pink stem borer, *Sesamia inferens* (Walker) on maize, *Zea mays*. *Journal of Applied and Natural Science*, 9(4), 1994-2003.
- Shekharappa, S. & Kulkarni, K. A. (2003). Cultural practices for the management of stem borer, *Chilo partellus* (Swinhoe) in sorghum.
- Shimeng, H. Shijie, X. Shangang, X. & Guomeng, M. (2001). Resistance of rice stem borer (*Chilo*

- suppressalis* Walker) to insecticides in Cixi, Zhejiang Province. *Acta Agriculturae Zhejiangensis*, 13(1), 38-41.
- Silva-Torres, C. S. Barros, R. & Torres, J. B. (2009). Efeito da idade, fotoperíodo e disponibilidade de hospedeiro no comportamento de parasitismo de *Oomyzus sokolowskii* Kurdjumov (Hymenoptera: Eulophidae). *Neotropical Entomology*, 38, 512-519.
- Sokame, B. M. Obonyo, J. Sammy, E. M. Mohamed, S. A. Subramanian, S. Kilalo, D. C. ... & Calatayud, P. A. (2021). Impact of the exotic fall armyworm on larval parasitoids associated with the lepidopteran maize stemborers in Kenya. *BioControl*, 66, 193-204.
- Stacconi, R. Grassi, A. L. B. E. R. T. O. Dalton, D. T. Miller, B. Ouantar, M. Loni, A. ... & Anfora, G. (2013). First field records of *Pachycrepoideus vindemiae* as a parasitoid of *Drosophila suzukii* in European and Oregon small fruit production areas. *Entomologia*, 1(e3), 11-16.
- Tang, L. D. Smagghe, G. Wang, S. Lü, Z. X. & Zang, L. S. (2024). Dead-end trap plants as an environment-friendly IPM tool: A case study of the successful use of vetiver grass in China. *Entomologia Generalis*.
- Turlings, T. C. & Erb, M. (2018). Tritrophic interactions mediated by herbivore-induced plant volatiles: mechanisms, ecological relevance, and application potential. *Annual review of entomology*, 63, 433-452.
- Uefune, M. Choh, Y. Abe, J. Shiojiri, K. Sano, K. & Takabayashi, J. (2012). Application of synthetic herbivore-induced plant volatiles causes increased parasitism of herbivores in the field. *Journal of Applied Entomology*, 136(8), 561-567.
- Upadhyay, A. Ranjitha, M. R. & Mishra, P. K. (2023). Major Pests of Maize and their Control.
- Van Dam, N. M. Qiu, B. L. Hordijk, C. A. Vet, L. E. & Jansen, J. J. (2010). Identification of biologically relevant compounds in aboveground and belowground induced volatile blends. *Journal of Chemical Ecology*, 36, 1006-1016.
- Van Noort, S. & Broad, G. (2024). *Wasps of the World: A Guide to Every Family* (Vol. 8). Princeton University Press.
- Virla, E. G. Moya-Raygoza, G. & Guglielmino, A. (2023). A review of the biology of the pincer wasps (Hymenoptera: Dryinidae). *Austral Entomology*, 62(3), 274-299.
- Wäschke, N. Meiners, T. & Rostás, M. (2013). Foraging strategies of parasitoids in complex chemical environments. *Chemical ecology of insect parasitoids*, 37-63.
- Webster, B. & Cardé, R. T. (2017). Use of habitat odour by host-seeking insects. *Biological Reviews*, 92(2), 1241-1249.
- Xu, H. Desurmont, G. Degen, T. Zhou, G. Laplanche, D. Henryk, L. & Turlings, T. C. (2017). Combined use of herbivore-induced plant volatiles and sex pheromones for mate location in braconid parasitoids. *Plant, cell & environment*, 40(3), 330-339.
- Yazdani, M. Feng, Y. Glatz, R. & Keller, M. A. (2015). Host stage preference of *Dolichogenidea tasmanica* (Cameron, 1912)(Hymenoptera: Braconidae), a parasitoid of *Epiphyas postvittana* (Walker, 1863)(Lepidoptera: Tortricidae). *Austral Entomology*, 54(3), 325-331.
- Yi, S. C. Wu, Y. H. Yang, R. N. Li, D. Z. Abdelnabby, H. & Wang, M. Q. (2023). A Highly

Expressed Antennae Odorant-Binding Protein Involved in Recognition of Herbivore-Induced Plant Volatiles in *Dastarcus helophoroides*. *International Journal of Molecular Sciences*, 24(4), 3464.

Zang, L. S. Wang, S. Zhang, F. & Desneux, N. (2021). Biological control with *Trichogramma* in China: History, present status, and perspectives. *Annual review of entomology*, 66, 463-484.