Effects of Self-Superparasitism and Host Age on Fitness-Correlated Traits in the Solitary Endoparasitoid Wasp Meteorus pulchricornis

Authors: Zhang, Bo, Li, Baoping, and Meng, Ling

Source: Journal of Insect Science, 14(103) : 1-11

Published By: Entomological Society of America

URL: https://doi.org/10.1673/031.014.103

The BioOne Digital Library (https://bioone.org/) provides worldwide distribution for more than 580 journals and eBooks from BioOne's community of over 150 nonprofit societies, research institutions, and university presses in the biological, ecological, and environmental sciences. The BioOne Digital Library encompasses

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

Effects of self-superparasitism and host age on fitness-correlated traits in the solitary endoparasitoid wasp *Meteorus pulchricornis*

Bo Zhang, Baoping Li, and Ling Meng[a]

College of Plant Protection, Nanjing Agricultural University, Agricultural Pests Management Laboratory of Ministry of Education, Nanjing 210095, PR China

Abstract

The domed fitness functions are suggested to describe developmental patterns of progeny parasitoids in relation to host age at oviposition in solitary koinobint parasitoids that are engaged in single parasitism, but few studies have investigated the applicability of the functions as related to superparasitism. The present study was designed to compare fitness functions between single parasitism and superparasitism by examining developmental patterns of *Meteorus pulchricornis* (Wesmael) (Hymneoptera: Braconidae) progeny in relation to the beet armyworm, *Spodoptera exigua* Hübner (Lepidoptera: Noctuidae), either singly parasitized or self-superparasitized as second–fifth instar larvae. Self-superparasitism caused deleterious effects on the fitness-related traits of parasitoid progeny, as demonstrated by a prolonged egg-to-adult emergence time, a smaller body size, and shorter longevity of the emerging adults, and decreased survival to adult emergence. While the domed fitness function was detected for development time, survival, adult body size, and longevity in relation to host larvae that were singly parasitized, the function was observed only for progeny survival in relation to host larvae that were self-superparasitized. This study suggests that developmental fitness functions with self-superparasitism can deviate from those with single parasitism in solitary koinobiont parasitoids.

Keywords: development, fitness, life history, koinobiont, superparasitism, parasitoids
Correspondence: [a] ml@njau.eud.cn
Editor: Michael Strand was editor of this paper.
Received: 10 August 2012 **Accepted:** 10 January 2013 **Published:** 1 August 2014
Copyright: This is an open access paper. We use the Creative Commons Attribution 3.0 license that permits unrestricted use, provided that the paper is properly attributed.
ISSN: 1536-2442 | Vol. 14, Number 103

Cite this paper as:
Zhang B, Li B, Meng L. 2014. Effects of self-superparasitism and host age on fitness-correlated traits in the solitary endoparasitoid wasp *Meteorus pulchricornis*. *Journal of Insect Science* 14(103). Available online: http://www.insectscience.org/14.103

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

Introduction

Among the most important trade-offs faced by organisms is whether to grow larger at the cost of extended development time or to develop more rapidly at the cost of reduced size (Abrams and Rowe 1996). To understand factors influencing offspring development and host selection strategies in parasitoids, the vast majority of studies have focused on the effects of host age, stage, or mass at parasitism on developmental traits in their parasitoids (reviewed by Mackauer and Sequeira 1993; Godfray 1994). The studies observe that the fitness functions differ markedly between (and within) koinobiont and idiobiont parasitoids in response to ecophysiological characteristics of the host species, including host size, growth potential at parasitism, immunity, and feeding ecology (Harvey et al. 2004; Pennacchio and Strand 2006). While a clear-cut host size–parasitoid fitness relationship is recognized for idiobionts, it is not always the case for koinobionts. The relationship between host quality and parasitoid fitness is much more complicated than is the case with idiobionts because it is determined by a range of potentially interacting ecophysiological selection pressures that are less prevalent in the latter group of parasitoids. For koinobionts attacking hosts of limited growth potential relative to maximum parasitoid size, the pattern is similar to that observed in idiobionts (Harvey et al. 1994). In contrast, fitness functions for solitary parasitoids attacking hosts with the potential to grow considerably larger than the adult parasitoid are distinctly dome-shaped (or may even decrease with host size in extreme cases) because progressively larger hosts possess more potent immunological defenses, or else they may be less nutritionally compatible for parasitoid development (and survival) than younger hosts (Strand 2000; Harvey et al. 2004). In the dome-shaped developmental model, fitness for parasitoids is not a linear or increasing function of host size or age at parasitism, but initially increases with host size and then rapidly declines as resources increase beyond the capacity of the parasitoid larva to consume them. The model gains strong support in numerous studies of solitary parasitoids involved in single parasitism (Smilowitz and Iwantsch 1975; Harvey et al. 1994, 1999, 2004; Pandey and Singh 1999; Chau and Mackauer 2001; Elzinga et al. 2003; Harvey et al. 2004; Liu and Li 2006, 2008; Chen et al. 2011), but few studies have yet investigated the applicability of this model to the parasitoids engaged in superparasitism.

As hosts represent a limited discrete resource, superparasitism generally results in reduction in survivorship of larvae and/or size, longevity, and fecundity of emerging adults (Visser et al. 1992; Harvey et al. 1993). However, superparasitism commonly is found not only in the laboratory (van Alphen and Marcel 1990; Bai and Mackauer 1992; Visser et al. 1992; Feuster et al. 1993; Yamada and Miyamoto 1998; Chau and Mackauer 1999), but in the field as well (Janssen 1989; Santolamazza-Carbone and Rivera 2003; Fleury et al. 2004; Jarmillo et al. 2006; Zhang et al. 2012). Superparasitism is now generally interpreted as an adaptive strategy in numerous theoretic models and empirical studies (van Alphen and Visser 1990; Visser et al. 1992; Mackauer and Chau 2001; Yamada and Kuzuma 2003; Reynolds and Hardy 2004) or as a result of virus manipulation (Varaldi et al. 2003; Gandon et al. 2006). It is therefore necessary to investigate developmental fitness functions in relation to host age in solitary koinobionts that are engaged in superparasitism so as to fully understand developmental strategies of the parasitoid wasps.

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

This study was designed to investigate the relationship between fitness-correlated traits of *Meteorus pulchricornis* (Wesmael) (Hymneoptera: Braconidae) progeny in either singly parasitized or self-superparasitized *Spodoptera exigua* Hübner (Lepidoptera: Noctuidae) hosts as second–fifth instar larvae, with the goal of testing whether the domed fitness functions apply to the parasitoid engaged in single as well as self-superparasitism. *M. pulchricornis* has a wide host range of exposed-living lepidopteran larvae (Huddleston 1980; Maeto 1989, 1990) and is a potential biological control agent against some lepidopteran pests (Fuester et al. 1993; Liu and Li 2006; Chau and Maeto 2008; Wu et al. 2008). The host suitability for development of the parasitoid progeny varies with host instars at the time of oviposition (Askari et al. 1977; Liu and Li 2006, 2008). Although foraging parasitoids can avoid self-superparasitism on the basis of host movement (Chau and Maeto 2009; Zhang et al. 2012), superparasitism rate still can be as high as 31% in the laboratory (Fuester et al. 1993) and 23% in the field (Zhang et al. 2012).

Materials and Methods

Culture of the parasitoid and host

Meteorus pulchricornis was obtained from *S. exigua* larvae collected in soybean fields and maintained using *S. exigua* as host in the insectary (26 ± 2ºC, 60%–80% RH, 14L:10D). The parasitoid is thelytokous. The host larvae were reared on an artificial diet (Liu and Li 2006).

Experiment

Spodoptera exigua larvae undergo five instars (designated as L1–L5) before pupation. As the first instar larva was not susceptible to self-superparasitism in the exploratory trial (n = 30), it was not used in the experiment. The exploratory trial (n = 50) indicated that females that realized one egg deposition performed a characteristic stinging behavior (flapping wings when withdrawing the ovipositor), which was used to obtain hosts that were either singly parasitized (1 egg) or superparasitized (2 eggs). Host larvae were individually exposed to a parasitoid in a glass vial and continuously observed until they were attacked once (single parasitism) or twice (self-superparasitism). The time interval between the first and second stinging was kept within 30 minutes; within this time, symmetrical competition between offspring larvae can be created in this parasitoid (Chau and Maeto 2008). The host larvae treated above were individually reared on the artificial diet in vials and monitored daily for egression of mature larvae, pupation, and adult emergence of progeny parasitoids. The emerged adults were kept in groups without feeding and hosts until dead, by which longevity was measured. The development time of progeny parasitoids was measured from oviposition to adult emergence (egg–adult emergence time), and the survival from this period was recorded. The right hind tibia length of the adults (a standard measure of body size) was measured under a stereomicroscope. Data were obtained from between 50 and 70 hosts for each parasitism mode/instar combination.

Statistical analysis

Two-way ANOVA was used to compare single parasitism and superparasitism in developmental parameters of offspring parasitoids. Orthogonal polynomial regression was used to analyze the trend in development time, body size, and longevity of offspring according to host stages with either single parasitism or self-superparasitism, with the goal of test on the quadratic term as the manifestation of the domed fitness functions. This regression method can overcome computational difficul-

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

Table 1. Results of orthogonal polynomial regression of life-history traits of *Meteorus pulchricornis* progeny parasitoids in relation to host instar singly parasitized (A) and self-superparasitized (B) as second–fifth instar larvae.

Observation variable	Parasitism mode	Polynomial term	Estimate	Std. Error	t value	Pr(>|t|)
Development time	Single parasitism	Intercept	14.14	0.05	259.6	< 0.001
		Linear	- 0.25	0.11	- 2.24	0.03
		Quadratic	0.28	0.1	2.53	**0.01**
		Cubic	0.11	0.1	0.97	0.33
	Superparasitism	Intercept	14.63	0.05	275.6	< 0.001
		Linear	0.23	0.11	2.18	0.03
		Quadratic	0.19	0.11	1.83	**0.07**
		Cubic	0.14	0.11	1.31	0.19
Body size	Single parasitism	Intercept	1.61	0.01	410.2	< 0.001
		Linear	0.07	0.01	8.81	< 0.001
		Quadratic	- 0.02	0.01	- 1.97	**0.05**
		Cubic	- 0.01	0.01	- 1.21	0.23
	Superparasitism	Intercept	1.56	0.01	404.1	< 0.001
		Linear	0.09	0.01	10.98	< 0.001
		Quadratic	- 0.01	0.001	- 1.52	**0.13**
		Cubic	- 0.01	0.01	- 0.32	0.75
Longevity	Single parasitism	Intercept	41.29	0.39	107.2	< 0.001
		Linear	1.28	0.79	1.63	0.1
		Quadratic	- 1.87	0.77	- 2.43	**0.02**
		Cubic	- 0.01	0.75	- 0.01	0.99
	Superparasitism	Intercept	37.53	0.27	139.4	< 0.001
		Linear	1.2	0.54	2.22	0.03
		Quadratic	- 1.02	0.54	- 1.89	**0.06**
		Cubic	0.25	0.54	0.46	0.64

Bold values indicate significant or marginally significant differences.

ties due to collinearity (Armitage et al. 2002). The Cochran-Armitage test of trend was used to approximately test for departures from a non-monotonic, dome-shaped trend in survival proportions across host stages at oviposition for single parasitism or self-superparasitism. Analyses were carried out using R (R Development Core Team 2009).

Results

A two-way ANOVA on development time revealed a significant effect of parasitism mode, where offspring parasitoids extended the egg-to-adult emergence time in self-superparasitized host larvae as opposed to counterparts in singly parasitized hosts (Figure 1). Orthogonal polynomial regression analysis revealed best fits of a quadratic curvilinear function to the relationship between development time and host stage at both single parasitism and self-superparasitism (Figure 1A, B; Table 1).

Offspring adult parasitoids were significantly smaller in body size, as measured by hind tibia length, in self-superparasitized hosts than in singly parasitized hosts (Figure 1C, D). Where the relationship between body size and host stage at oviposition was best fitted by a quadratic curvilinear function for offspring parasitoids engaged in single parasitism, it was best fitted only by a linear function for the counterparts involved in self-superparasitism (Figure 1C, D; Table 1).

Longevity of adults was shortened in self-superparasitized hosts as opposed to singly parasitized hosts (Figure 2). Whereas there was a quadratic effect of host stage on longevity of parasitoids engaged in single parasitism, the quadratic effect was close to statistically significant for parasitoids involved in self-superparasitism ($P = 0.06$) (Figure 2A, B; Table 1).

The survival of offspring parasitoids decreased by 15–21% in self-superparasitized

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

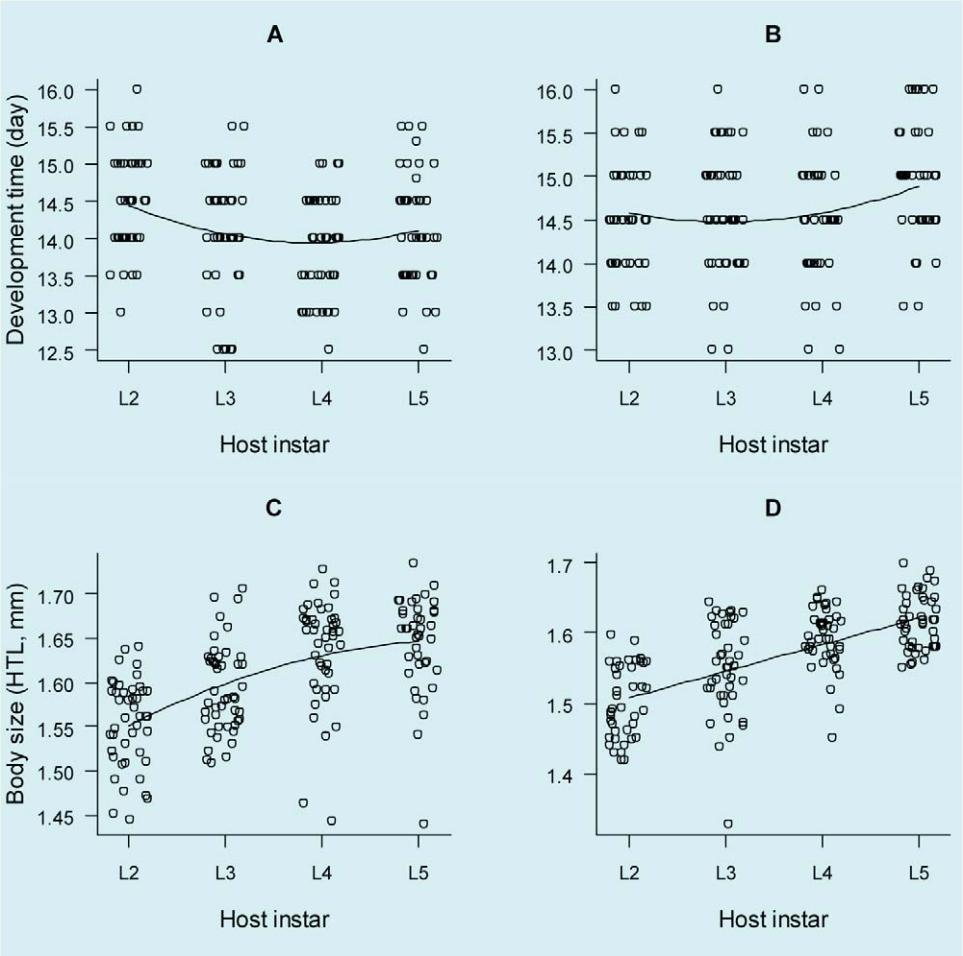

Figure 1. *Meteorus pulchricornis* development time (egg–adult emergence) and body size (hind tibia length, HTL) in relation to host instar singly parasitized (A, C) and self-superparasitized (B, D) as second–fifth instar larvae. High quality figures are available online.

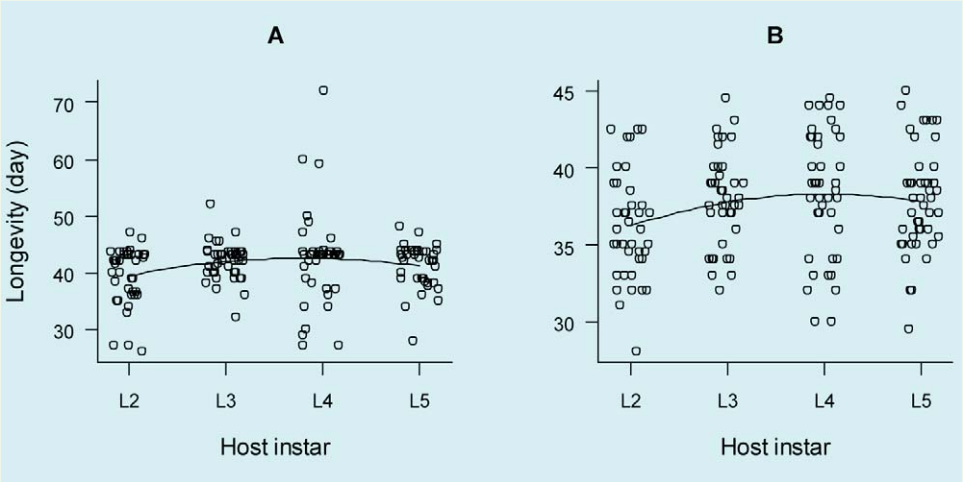

Figure 2. Longevity of *Meteorus pulchricornis* adults in relation to host instar singly parasitized (A) and self-superparasitized (B) as second–fifth instar larvae. High quality figures are available online.

hosts as compared to singly parasitized hosts (Figure 3). The Cochran-Armitage test of trend supported the dome-shaped trend in survival proportions across successive host stages at ovipositon for both single parasitism (χ^2 =

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

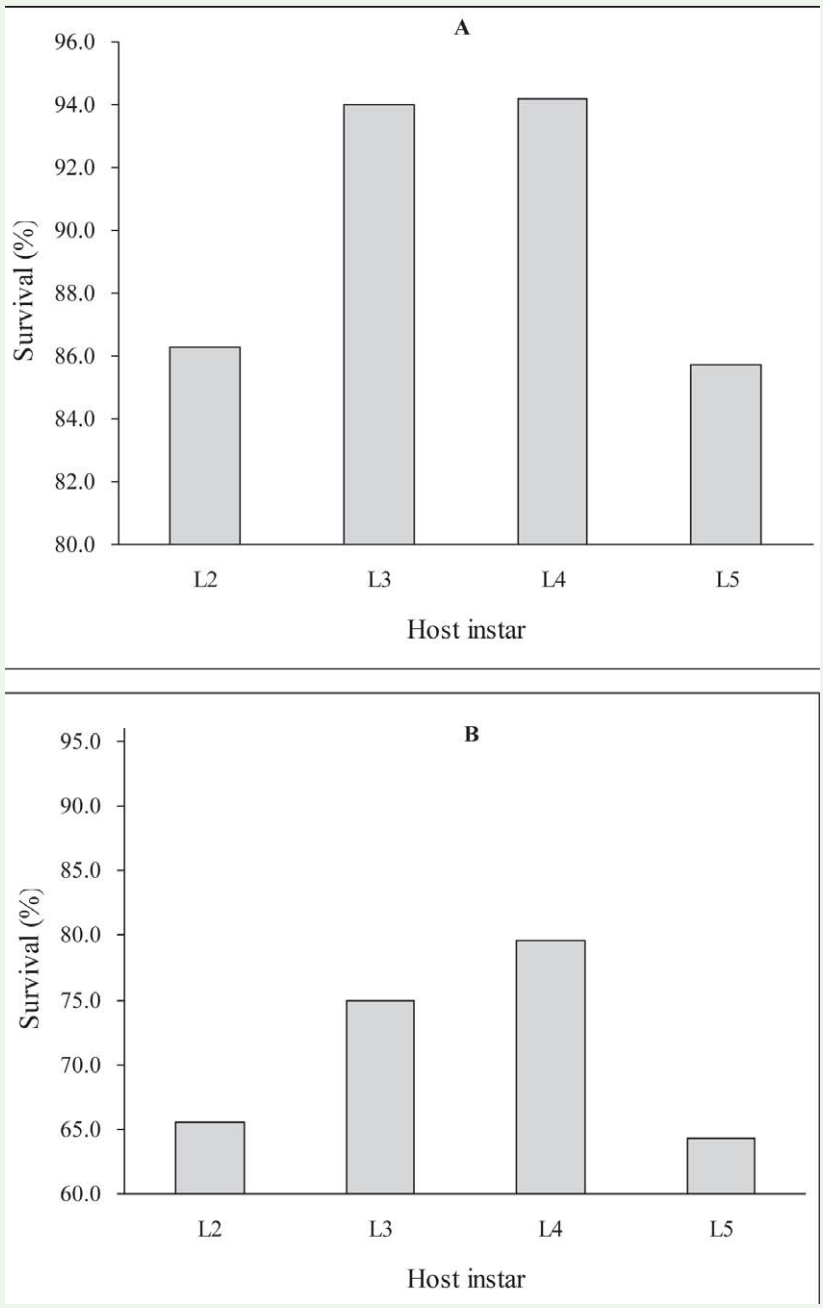

Figure 3. Percentage survival to adult emergence of *Meteorus pulchricornis* from hosts singly parasitized (A) and self-superparasitized (B) as second–fifth instar larvae. High quality figures are available online.

18.81, P <0.001) and self-superparasitism (χ^2 = 4.41, P = 0.04) (Figure 3).

Discussion

Our experiments show that self-superparasitism in *M. pulchricornis* had deleterious effects on the development of parasitoid progeny, leading to prolonged egg-to-adult emergence time, decreased survival to adult emergence, smaller body size, and shorter longevity of the emerging adults. The results provide data in support of some other studies, which observed that superparasitism in solitary parasitoids delays the development of the progeny (Eller et al. 1990; Harvey et al. 1993; Hegazi and Khafagi 2005; Chau and Maeto 2008; Tunca and Kilinger 2009), increases larval mortality (Vinson and Spoka 1978; Tunca and Kilinger 2009), and results in offspring small in body size (Harvey et al. 1993; Tunca and Kilinger 2009). The fitness costs in development of parasitoid progeny in a superparasitized host are generally assumed to be due to increased competition for limited resources among progeny inside the host (van Lenteren 1981; Waage and Godfray 1986). Moreover, some other competing behaviors may also bear fitness costs for the winning parasitoid. Chau and Maeto (2008) found, in the study of larval competition in *M. pulchricornis*, that fierce physical fighting between first-instar larvae occurred inside the superparasitized host. However, some fitness costs for progeny parasitoids from superparasitism may be offset. For example, studies of the solitary aphid parasitoid *Monoctonus paulensis* show that progeny parasitoids are larger in body size in superparasitized aphid hosts than counterparts in singly parasitized aphids without a corre-

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

sponding increase in development time (Bai and Mackauer 1992; Mackauer and Chou 2001).

The results of this study show that fitness correlates in development of progeny parasitoids engaged in single parasitism do not linearly increase with host stage at oviposition. The analysis of orthogonal polynomial regression revealed a quadratic effect of host stage on development time, body size, and longevity of progeny parasitoids, respectively; the Cochran-Armitage test on survival proportions showed a dome-shaped trend from L2 to L3 hosts at oviposition. These results provide data in support of the findings not only with this parasitoid parasitizing other host species (Liu and Li 2008), but also with other koinobiont parasitoids that exhibit developmental patterns illustrating clear trade-offs between body size, development time, and pre-adult mortality (Smilowitz and Iwantsch 1975; Harvey et al. 1994, 1999; Pandey and Singh 1999; Chau and Mackauer 2001; Elzinga et al. 2003; Harvey et al. 2004). Such domed fitness functions may be an adaptive response that optimizes several traits affecting parasitoid fitness (Strand 2000).

The domed fitness functions, however, are demonstrated only in parasitoids engaged in single parasitism. There remains a need to know if these functions apply to the parasitoids engaged in superparasitism, in consideration of superparasitism as an adaptive strategy in parasitoids (van Alphen and Visser 1990). The results of this study do not reveal a significantly quadratic function in development time, body size, and longevity of progeny parasitoids engaged in self-superparasitism with regard to host stages at oviposition, but support a quadratic trend in survival proportions across successive host stages. The discrepancy in developmental fitness functions between single parasitism and self-superparasitism, as exhibited in *M. pulchricornis*, may be explained by several factors. The eggs of parasitoids in several families (especially in the Braconidae) produce large cells, teratocytes, which are released to the host haemocoel when the parasitoid first instar ecloses (Dahlman and Vinson 1993). Among other functions, teratocytes have the function of nutrition for developing parasitoid larvae. Bai and Mackauer (1992) reported that larvae of the solitary koinobiont endoparasitoid *Aphidius ervi* readily feed on them along with host tissues, and that the larger size of progeny emerging from superparasitized hosts compared with singly parasitized hosts may be a result of increased numbers of teratocytes circulating in the haemolymph of superparasitized hosts. Our study also provides some corroborative evidence for this factor. The body size of progeny initially increased with host instar, but rapidly declined when the host at oviposition was the final instar with single parasitism, whereas it linearly increased with host instar at oviposition with self-superparasitism.

Alternatively, the discrepancy may be explained by the same mechanism for the domed fitness functions in single parasitism. Strand (2000) suggested that with single parasitism, offspring fitness initially increases with host size, but as resources increase beyond the capacity of the parasitoid larva to consume them, fitness rapidly declines. We assumed that the resource threshold for the consumption of parasitoid larvae may be higher with self-superparasitism, so as to shift the increasing trend in some life-history traits, such as body size, with host size (or stage) at oviposition. In this respect, self-superparasitism may be an adaptive strategy in solitary koinobionts to exploit suboptimal host stages. This as-

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

sumption has gained some support in the study of *Microplitis rufiventris*, where more parasitoid eggs survive to mature larvae when laid in the sixth-instar larva, the suboptimal stage for single parasitism, than the fifth-instar host, which is the optimal stage for single parasitism (Khafagi and Hegazi 2005; Hegazi and Khafagi 2008). Furthermore, self-superparasitism may be enhanced by host sharing in solitary endoparasitoids toward the evolution of gregariousness over time (Riddich 2002; Khafagi and Hegazi 2005). Pexton and Mayhew (2005) reported that competition for hosts may have contributed to the evolution of gregarious development in the braconid genus *Aphaereta*.

Acknowledgements

We thank Lulu Li, Sheng Sheng, Zhu Yu, and Lu Feng for their help in the study, and the Editor for the valuable comments on the manuscript. This study was supported by the Natural Science Fund of China (NSFC-30871670) and Special Research Fund for Agricultural Technique for the Public Interests (201103002).

References

Abrams PA, Rowe L. 1996. The effects of predation on the age and size of maturity of prey. *Evolution* 50: 1052–1061

Armitage P, Berry G, Matthews JNS. 2002. *Statistical Methods in Medical Research.* Fourth edition. Blackwell Science Ltd.

Askari A, Mertins JW, Coppel HC. 1977. Development biology and immature stages of Meteorus pulchricornis in the laboratory. *Annals of the Entomological Society of America* 70: 655–659.

Bai B, Mackauer M. 1992. Influence of superparasitism on development rate and adult size in a solitary parasitoid wasp, *Aphidius ervi*. *Functional Ecology* 6: 302–304.

Chau A, Mackauer M. 1999. Self-superparasitism in the solitary parasitoid *Monoctonus paulensis* (Hymenoptera: Braconidae, Aphidiinae): Proximate mechanisms. *Canadian Entomologist* 131: 769–777.

Chau A, Mackauer M. 2001. Preference of the aphid parasitoid *Monoctonus paulensis* (Hymenoptera: Braconidae, Aphidiinae) for different aphid species: Female choice and offspring survival. *Biological Control* 20: 30–38.

Chau NNB, Maeto K, 2008. Intraspecific larval competition in *Meteorus pulchricornis* (Hymenoptera: Braconidae), a solitary endoparasitoid of lepidopteran larvae. *Applied Entomology and Zoology* 43: 159–165.

Chau N, Maeto K. 2009. Temporary host paralysis and avoidance of self-superparasitism in the solitary endoparasitoid *Meteorus pulchricornis*. *Entomologia Experimentalis et Applicata* 132: 250–255.

Chen W, Li B, Meng L. 2011. Host selection and offspring developmental performances of *Meteorus pulchricornis* on *Spodoptera litura* larval instars. *Chinese Journal of Ecology* 30(7): 1317–1321 (In Chinese with English abstract).

Dahlman DL, Vinson SB. 1993. Teratocytes: developmental and biochemical characteristics. In: Beckage NE, Thompson SN, Federici BA, eds. 1993. *Parasites and Pathogens of Insects*. pp. 145–65. New York: Academic.

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

Eller FJ, Tumlinson JH, Lewis WJ. 1990. Intraspecific competition in *Micropitilis croiceps* (Hymenoptera: Braconidae), a parasitoid of *Heliothis* species (Lepidoptera: Noctuidae). *Annals of the Entomological Society of America* 83: 504–508.

Elzinga JA, Harvey JA, Biere A. 2003. The effects of host weight at parasitism on fitness correlates of the gregarious koinobiont parasitoid *Microplitis tristis* and consequences for food consumption by its host, *Hadena bicruris*. *Entomologia Experimentalis et Applicata* 108: 95–106.

Fleury F, Ris N, Allemand R, Fouillet P, Carton Y, Boulétreau M. 2004. Ecological and genetic interactions in *Drosophila* parasitoids communities: a case study with *D. melanogaster*, *D. simulans* and their common *Leptopilina* parasitoids in south-eastern France. *Genetica* 120: 181–194.

Fuester R, Taylor P, Peng H, Swan K. 1993. Laboratory biology of a uniparental strain of *Meteorus pulchricornis* (Hymenoptera: Braconidae), an exotic larval parasite of the gypsy moth (Lepidoptera: Lymantriidae). *Annals of the Entomological Society of America* 86: 298–304.

Godfray H. 1994. *Parasitoids: Behavior, evolution, and evolutionary ecology.* Princeton University Press (Princeton, N.J.), Princeton.

Gandon S, Rivero A, Varaldi J, 2006. Superparasitism evolution: adaptation or manipulation? *The American Naturalist* 176: 1–22.

Harvey JA. 2005. Factors affecting the evolution of development strategies in parasitoid wasps: the importance of functional constraints and incorporating complexity. *Entomologia Experimentalis et Applicata* 117: 1–13.

Harvey JA, Strand MR. 2002. The developmental strategies of endoparasitoid wasps vary with host feeding ecology. *Ecology* 83(9): 2439–2451.

Harvey JA, Harvey IF, Thomson DJ. 1993. The effect of superparasitism on development of the solitary parasitoid wasp, *Venturia canescens* (Hymenoptera: Ichneumonidae). *Ecological Entomology* 18: 203–208.

Harvey JA, Harvey IF, Thompson DJ. 1994. Flexible larval growth allows use of a range of host sizes by a parasitoid wasp. *Ecology* 75: 1420–1428.

Harvey JA, Bezemer TM, Elzinga JA, Strand MR. 2004. Development of the solitary endoparasitoid *Microplitis demolitor*: host quality does not increase with host age and size. *Ecological Entomology* 29: 35–43.

Harvey JA, Jervis MA, Gols R, Jiang NQ, Vet LEM. 1999. Development of the parasitoid, *Cotesia rubecula* (Hymenoptera: Braconidae) in *Pieris rapae* and *Pieris brassicae* (Lepidoptera:Pieridae): evidence for host regulation. *Journal of Insect Physiology* 45: 173–182.

Hegazi E, Khafagi W. 2005. Developmental interaction between suboptimal instars of *Spodoptera littoralis* (Lepidoptera:Noctuidae) and its parasitoid *Microplitis rufiventris* (Hymenoptera: Braconidae). *Archives of Insect Biochemistry* 60: 172–184.

Huddleston T. 1980. A revision of the western Palaearctic species of the genus *Meteorus* (Hymenoptera, Braconidae). *Bulletin of the*

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

British Museum (Natural History), Entomology 41: 1–58.

Janssen A. 1989. Optimal host selection by *Drosophila* parasitoids in the field. *Functional Ecology* 3: 469–479.

Jaramillo J, Borgemeister C, Setamou M. 2006. Field superparasitism by *Phymastichus coffea*, a parasitoid of adult coffee berry borer, *Hypothenemus hampei*. *Entomologia Experimentalis et Applicata* 119: 231–237.

Khafagi WE, Hegazi EM. 2008. Does superparasitism improve host suitability for parasitoid development? A case study in the *Microplitis rufiventris–Spodoptera littoralis* system. *BioControl* 53: 427–438.

Liu Y, Li B. 2006. Developmental interactions between *Spodoptera exigua* (Noctuidae: Lepidoptera) and its uniparental endoparasitoid, *Meteorus pulchricornis* (Braconidae: Hymenoptera). *Biological Control* 38: 264–269.

Liu Y, Li B. 2008. Effects of *Helicoverpa armigera* (Noctuidae, Lepidoptera) host stages on some developmental parameters of the uniparental endoparasitoid *Meteorus pulchricornis* (Braconidae, Hymenoptera). *Bulletin of Entomological Research* 98: 109–114.

Mackauer M, Sequeira R. 1993. Patterns of development in insect parasites. In: Beckage NE, Thompson, SN, Federici BA. Editors. *Parasites and Pathogens of Insects.* pp. 1–20. AcademicPress.

Mackauer M, Chau A. 2001. Adaptive self-superparasitism in a solitary parasitoid wasp: the influence of clutch size on offspring size. *Functional Ecology* 15: 335–343.

Maeto K. 1989. Systematic studies on the tribe Meteorini from Japan (Hymenoptera, Braconidae) VI. The pulchricornis group of the genus *Meteorus*. *Japanese Journal of Entomology* 57: 768–777.

Maeto K. 1990. Phylogenetic relationships and host associations of the subfamily *Meteorinae cresson* (Hymenoptera, Braconidae). *Japanese Journal of Entomology* 58: 383–396.

Pandey S, Singh R. 1999. Host size induced variation in progeny sex ratio of an aphid parasitoid *Lysiphlebia mirzai*. *Entomologia Experimentalis et Applicata* 90: 61–67.

Pennacchio F, Strand MR. 2006. Evolution of developmental strategies in parasitic Hymenoptera. *Annual Review of Entomology* 51: 233–258.

Pexton JJ, Mayhew PJ. 2005. Clutch size adjustment, information use and the evolution of gregarious development in parasitoid wasps. *Behavioral Ecology and Sociobiology* 58: 99–110.

R Development Core Team, 2009. *R: A Language and Environment for Statistical Computing*. R Foundation for Statistical Computing, Vienna, Austria. Available online: www.r-project.org.

Riddick EW, 2002. Superparasitism occasionally predisposes *Cotesia marginiventris* (Cresson) (Hymenoptera: Braconidae) to develop gregariously in *Spodoptera exigua* (Hubner) (Lepidoptera: Noctuidae). *Journal of Entomological Science* 37: 1–9.

Reynolds KT, Hardy ICW. 2004. Superparasitism: a non-adaptive strategy?

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use

Trends in Ecology and Evolution 19: 347–348.

Santolamazza-Carbone S, Rivera CA. 2003. Superparasitism and sex ratio adjustment in a parasitoid wasp: results at variance with local mate competition? *Oecologia* 136: 365–373.

Smilowitz Z, Iwantsch GF. 1975. Relationships between the parasitoid, *Hyposoter exiguae*, and the cabbage looper, *Trichoplusia ni*: effects of host age on ovipositional rate of the parasitoid and successful parasitism. *Canadian Entomologist* 107: 689–694.

Strand M. 2000. Life history variation and developmental constraints in parasitoids. In: Hochberg M, Ives AR, Editors. *Population Biology of Parasitoids.* pp. 139-162. Princeton University Press.

Tunca H, Kilinger NE. 2009. Effect of superparasitism on the development of the solitary parasitoid *Chelonus oculator* Panzer (Hymenoptera: Braconidae). *Turkey Journal of Agriculture and Forestry* 33: 463–468.

van Alphen J, Visser M. 1990. Superparasitism as an adaptive strategy for insect parasitoids. *Annual Review of Entomology* 35: 59–79.

Varaldi J, Fouillet P, Ravallec M, Lopez-Ferber M, Bouletreau M, Fleury F. 2003. Infectious behaviour in a parasitoid. *Science* 302: 1930.

Vinson SB, Spoka P. 1978. Effects of superparasitism by a solitary endoparasitoid on the host, parasitoid and field samplings. *Southwestern Entomologist* 3: 299–303.

Visser ME, van Alphen JJM, Hemeril L. 1992. Adaptive superparasitism and patch time allocation in solitary parasitoids: An ESS model. *Journal of Animal Ecology* 61: 93–101.

Wu H, Meng L, Li B. 2008. Effects of feeding frequency and sugar concentrations on lifetime reproductive success of *Meteorus pulchricornis* (Hymenoptera: Braconidae). *Biological Control* 45: 353–359.

Yamada YY, Miyamoto K. 1998. Payoff from self and conspecific superparasitism in a dryinid parasitoid, *Haplogonatopus atratus*. *Oikos* 81: 209–216.

Yamada YY, Kazuma S. 2003. Evidence for adaptive self-superparasitism in the dryinid parasitoid *Haplogonatopus atratus* when conspecifics are present. *Oikos* 103: 175–181.

Zhang B, Huang L, Meng L, Li B. 2012. Host discrimination ability and the influencing factors in *Meteorus pulchricornis* (Hymenoptera: Braconidae). *Acta Entomologica Sinica* 54: 1391–1398 (In Chinese with English abstract)

Zhang B, Meng L, Huang L, Li B. 2012. Oviposition selection behavior of *Meteorus pulchricornis* (Hymenoptera: Braconidae) in the field. *Journal of Nanjing Agricultural University* 35: 78–82 (In Chinese with English abstract).

Downloaded From: https://staging.bioone.org/journals/Journal-of-Insect-Science on 25 Mar 2025
Terms of Use: https://staging.bioone.org/terms-of-use