

EFFECT OF ENVIRONMENTAL FACTORS ON THE POPULATION OF *VARROA DESTRUCTOR* IN *APIS MELLIFERA* L. COLONIES

ASHA POONIA*, RACHNA GULATI AND S. K. SHARMA

¹Departments of Zoology and Aquaculture, CCS HAU, Hisar -125 004 (Haryana), INDIA

²Directorate of Research; ³Entomology, CCS HAU, Hisar (Haryana), INDIA

e-mail: asha.poonia@gmail.com

INTRODUCTION

Varroa mite (Acari: Mesostigmata), parasitizing the European honeybee, *Apis mellifera* L. (Hymenoptera: Apidae) is responsible for loss of more than 50% of *Apis mellifera* colonies worldwide (Shaw *et al.*, 2002; Topolska *et al.*, 2010; Martin *et al.*, 2012; Nazzi *et al.*, 2012). 90 per cent apiaries and 50 per cent colonies of state of Haryana are affected by this mite (Gulati *et al.*, 2009). This mite which feeds on haemolymph of brood and adult bees causes colony disorder, weakness, decreasing brood and deforming immature and mature bees (Kotwal and Abrol, 2013)

The temperature relations within honey bee colonies are complex, as honey bees thermoregulate their colonies according to the season and the presence or absence of brood (Seeley and Heinrich, 1981). *Varroa destructor* preferentially reproduces on at 32.5-33.4°C temperature as it is the temperature on which drone brood maintained (Le Conte *et al.*, 1990). Various workers have tried to investigate the relationship of *Varroa destructor* population with abiotic factors in *Apis mellifera* colonies (Martin, 1995; Crane, 1978; Harbo, 2000; Webster *et al.*, 2000; Harris *et al.*, 2003; Underwood and Currie, 2003; Bahreini *et al.*, 2004; De Guzman *et al.*, 2007; Kotpal, 2008). In Hisar, Haryana various abiotic factors (Temperature, rainfall, sunshine hours, relative humidity) shows a wide range in a year. Present investigation was carried with the objective to study the effect these abiotic factors on the population of *Varroa destructor* in *Apis mellifera* colonies in Hisar, Haryana.

MATERIALS AND METHODS

The experiment was conducted in CCS Haryana Agricultural apiary from May 2009 to February 2010. Three colonies statistically comparable in terms of *Varroa* infestation, colony strength, brood, pollen and honey area were taken for the experiment.

There are three standard methods of *Varroa* population estimation (Ritter, 1981; De Jong *et al.*, 1982; Branco *et al.*, 2006; Dietemann *et al.*, 2013) *i.e.*

1. Acaricides to kill all mites in a colony. Mites fall to the bottom of the hive and can be counted.
2. Without the use of acaricides, the natural mortality can be quantified from the bottom of the hive to determine the population size of the live mites.
3. The infestation rates of adults and brood can be estimated from adult and brood samples.

All three methods (using acaricides, monitoring natural mite fall and assessing infestation levels) were found to provide comparable results (Branco *et al.*, 2006). First method could not be chosen as it involved killing of the *Varroa* mites, which could not be used as samples of natural *Varroa* population needed to be taken throughout the year. Third method also could not be chosen as it involved killing

ABSTRACT

Maximum number of mites was observed in second fortnight of May (38 and 51 mites/per hive) which was found significantly positively correlated with maximum ($r = 0.659$) and minimum ($r = 0.648$) temperature. However, *V. destructor* population was found negatively correlated with relative humidity (-0.416) and sunshine hours (-0.023). Rainfall was found none significantly correlated (0.019) with *V. destructor* population. Data suggested that during summer months, when temperature is high and flower availability is less, mite population increases in *Apis mellifera* L. colonies.

KEY WORDS

Varroa destructor
Apis mellifera
temperature, relative humidity, rainfall,
sunshine hours

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*Corresponding author

of brood, thus, interrupted with the natural population of *Apis mellifera*. In this light, monitoring natural mite fall was used to know colony infestation rate of *Varroa destructor* in *Apis mellifera* colonies. In this method mites that die naturally or due to grooming behaviour of bees get dislodged from bees and can be collected from the bottom board of the hive in debris. This natural mite fall count can be used to estimate colony infestation. This method has been used by various workers (Devlin, 1998; Dietemann et al., 2013). Ants were

Table 1: Comparative sampling methods to record *Varroa destructor* population in *Apis mellifera* colonies

Date of observation	Number of mites (Mean \pm S. D.)/ colony*Sticky paper
05.05.08	36.00 \pm 2.00
12.05.08	38.00 \pm 2.00
19.05.08	51.50 \pm 5.00
26.05.08	35.00 \pm 3.00
02.06.08	23.00 \pm 2.00
09.06.08	14.00 \pm 2.00
17.06.08	13.50 \pm 2.00
24.06.08	11.00 \pm 3.00
01.07.08	10.50 \pm 1.00
07.07.08	18.00 \pm 5.00
14.07.08	18.50 \pm 2.00
21.07.08	18.50 \pm 5.00
28.07.08	19.50 \pm 2.00
05.08.08	20.50 \pm 5.00
11.08.08	9.50 \pm 2.00
19.08.08	7.50 \pm 2.00
26.08.08	7.00 \pm 2.00
02.09.08	4.50 \pm 2.00
08.09.08	9.50 \pm 2.00
15.09.08	9.50 \pm 2.00
24.09.08	12.00 \pm 2.00
29.09.08	9.50 \pm 2.00
08.10.08	8.50 \pm 2.00
15.10.08	8.50 \pm 2.00
22.10.08	2.00 \pm 0.00
30.10.08	0.00 \pm 0.00
06.11.08	0.00 \pm 0.00
No mites were noticed from 09.11.08 to 25.12.08	
24.12.08	2.50 \pm 1.00
01.01.09	1.00 \pm 0.00
07.01.09	1.00 \pm 0.00
15.01.09	1.00 \pm 0.00
19.01.09	1.00 \pm 0.00
No mites were noticed from 23.01.09 to 30.04.09	

*Average of three colonies

prevented from access to bottom boards by using a hive stand resting in water containers over which ants cannot walk (Dietemann et al., 2013). Dirt accumulated in water containers was cleared regularly so that ants could not reach the hive.

Data on weather parameters viz., maximum and minimum temperature ($^{\circ}$ C), relative humidity (%), rainfall (mm) and bright sunshine hours (h) was collected from Department of Meteorology, CCS Haryana Agricultural University, Hisar (Haryana) in order to correlate these factors with seasonal abundance of *V. destructor* in *A. mellifera* colonies. Correlation matrix was calculated between *V. destructor* incidence and abiotic factors to see their effect on population build up of mite.

RESULTS AND DISCUSSION

Table 1 shows the population of *V. destructor* from May 2008 to February 2009. To find the relationship between biotic (number of *V. destructor* in *A. mellifera* colonies) and abiotic factors (maximum and minimum temperature, relative humidity, rainfall and sunshine hours), correlation matrix was calculated and presented in Table 2. The study revealed that among abiotic factors, maximum and minimum temperature played a significant role in the population build up of *V. destructor*. The mite population was significantly positively correlated with the maximum ($r = 0.659$) and minimum ($r = 0.648$) temperature. The positive values of correlation coefficient revealed that as the maximum temperature with in a range from 18.6 to 39.6 $^{\circ}$ C and minimum temperature within a range from 3.3 to 26.3 $^{\circ}$ C increased, there was a corresponding increase in *V. destructor* population. The maximum population of *V. destructor* was found in the month of May when maximum temperature fluctuated between 36.4 to 39.6 $^{\circ}$ C which indicated that a higher temperature around 38 $^{\circ}$ C favoured the buildup of mite population. It is also evident from Table 2 that there is a negative correlation between *V. destructor* population and relative humidity, sunshine hours, but it was non significant. Similarly, non-significant relation between *V. destructor* population and rainfall was recorded in present investigation.

Differences in mite population due to climatic variation are reported by several workers. Earlier studies reported that level of *Varroa* mite infestation was lower in tropical regions as compared to temperate regions. Underwood and Currie (2003) observed maximum mite fall at 35 $^{\circ}$ C when various doses of

Table 2: Correlation matrix between *Varroa destructor* population and weather parameters

Treatments	No of mites on Sticky paper	Temperature ($^{\circ}$ C)		Relative Humidity (32 to 89%)	Sunshine) hours (0.04 to10.2h)	Rainfall (0.0 to 11 mm)
		Maximum (18.6 to39.6 $^{\circ}$ C)	Minimum (3.3 to25.8 $^{\circ}$ C)			
No of mites on Sticky paper	1.00					
Maximum Temperature ($^{\circ}$ C)	0.659*	1.00				
Minimum Temperature ($^{\circ}$ C)	0.648*	0.887*	1.00			
Relative Humidity (%)	-0.416 ^{NS}	-0.224 ^{NS}	0.077 ^{NS}	1.00		
Sunshine hours (h)	-0.023 ^{NS}	-0.063 ^{NS}	-0.162 ^{NS}	-0.492*	1.00	
Rainfall (mm)	0.019 ^{NS}	0.132 ^{NS}	0.243 ^{NS}	0.269 ^{NS}	-0.449*	1.00

*Significant at 0.05 % level; NS = Non significant

formic acid at different temperatures were applied. Populations of *Varroa* increase very rapidly in some regions such as Europe but remain at a low level in some tropical countries (De Jong *et al.*, 1984; Ruttner *et al.*, 1984). De Jong *et al.* (1984) attributed this to a lower rate of reproduction in temperate conditions. Woyke (1987) reported a high infestation rate of *A. mellifera* colonies by *Tropilaelaps clareae* in Southern Vietnam and a low rate in northern Vietnam. Harris *et al.* (2003) also reported correlation between ambient temperature and relative humidity with growth of mite populations during 10 year study period (1993-2002). Webster *et al.* (2000) also reported that total mite fall follow temperature regime. A modest positive correlation was found between proportion of mite fall and mean daily high temperature ($r = 0.515$), overall mean weekly temperature ($r = 0.508$) and weekly mean low temperature ($r = 0.452$). The reason attributed was that weather has sub-lethal effect on mites as in case of heat treatment which is used to kill mites in brood cells (Martin, 1995; Harbo, 2000) and in entire hives (Crane, 1978). Bahreini *et al.* (2004) calculated correlation between *Varroa* mortality with temperature and relative humidity. Kotwal *et al.* (2013) has also observed similar results of *Varroa* population in various months in 2006-07 and 2007-08 with peak of population in March.

Kotwal (2008) in Jammu and Kashmir (India) recorded significant negative correlation with rainfall and evening relative humidity during 2006-07 whereas significant negative correlation was obtained with minimum temperature in 2007-08. However, in the present study, relative humidity and rainfall had no significant relation with the population build up of *V. destructor* in *A. mellifera* colonies.

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