Acaricide resistance among broad mite (*Polyphagotarsonemus latus* (Banks)) populations in Karnataka

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Abstract

The broad mite, *Polyphagotarsonemus latus* (Banks) is a cosmopolitan pest that attacks a wide range of economically important crops like hot and sweet peppers, mulberry, jute, tea and several ornamentals. This study was undertaken to monitor the development of acaricide resistance, if any in five representative field-collected populations of Karnataka. Bioassays were carried out against five acaricide chemistries and resistance ratios were calculated by comparing the LC_{50} values of field populations with the susceptible laboratory population (Pa-Lab). The resistance ratios varied from 26.03 to 81.16 folds for diafenthiuron, 27.35 to 83.47 for dicofol, 9.72 to 45.42 for fenazaquin, 8.77 to 16.84 for propargite and 48.37 to 163.39 for spiromesifen. Resistance to the acaricides is unstable in *P. latus* as a decline in resistance (14.11 to 102.53 folds) was observed over generations in the absence of selection pressure. The results suggested that acaricides should be sprayed at economic threshold levels or on a rotation basis for one or more seasons for better management of *P. latus* by delaying the development of resistance.

Keywords: Acaricide, *Polyphagotarsonemus latus*, stability, susceptibility, bioassay

Introduction

Mite infestations on crops have become an alarming concern in recent years, especially in the tropics and sub-tropics.[1] This can be attributed to several factors *viz*., climate change, rapid host expansion capability, development of acaricide resistance, withdrawal of broad-spectrum pesticides from the markets that were used against the multi-resistant pests, or a combination of these factors. Specifically, the broad mite or yellow mite, *Polyphagotarsonemus latus* (Banks) (Acari: Tarsonemidae) has emerged as a major crop pest attacking several crops of agricultural and horticultural importance.[2,3] Originally it was reported in 1890 from tea plants in Sri Lanka, but later distributed to other countries through trade and commerce. Currently, it is a well-established pest in all six zoogeographical realms of the world including Australia, Asia, Africa, North America, South America and the Pacific Islands.[4] It flourishes well and multiplies faster under tropical, subtropical and greenhouse habitats. The broad mites are known to damage over 250 host plants belonging to 57 families, including hot and sweet peppers, mulberry, jute, tea, sesame, cotton and several ornamentals where the feeding of this pest habitually compromises the yield.[2,3,5,6,7]

The larval and adult stages feed on the sap from the young foliage and budding tips leaving the newest growths severely damaged and plant growth suppressed.[8] The typical symptoms noticed on chilli and capsicum are bronzing, crinkling and downward curling of leaves giving an inverted boat-shaped appearance, elongation of the petiole, crumpled apical shoot, development of abnormal side shoots, thickening of leaves and stunted growth.[9] The damage due to mites in chilli had been estimated to the tune of 60 percent and can cause cent percent yield loss under polyhouse conditions.[10]

The utility of host plant resistance and biocontrol agents like predatory mites, fungi and

bacteria against *P. latus* have been explored with little success. Hence, chemical control with synthetic acaricides remains as the common management strategy followed by farmers in India. More than two dozen acaricides/ insecticides alone or in combination under 12 different modes of action are registered for official use in India for the management of broad mites.[11] Due to its microscopic nature (<0.2mm) and short generation time the damage is being caused even before the mite's presence is detected. This necessitates the repeated application of pesticides that would lead to the rapid development of resistance. Hence, the present study has determined the development of resistance to major acaricides if any, and the resistance stability in *P. latus* populations collected from Karnataka.

Material and Methods

Mite colony maintenance

A colony of *P. latus* (National accession number NBAIR-GR-TAR-01a) is being maintained at ICAR-National Bureau of Agricultural Insect Resources, Bangalore since July 2020. The colony was maintained in insect growth chamber with temperature between 25 ± 5 ℃ and RH of 65 ± 5% on potted mulberry plants (variety V1) that contained a mixture of FYM and red soil at 1:2 ratio. This mite population, designated as Pa-Lab, was retained for more than 70 generations without any exposure to acaricides.

The species identity of the mites was confirmed both by morphological[4] and molecular methods.[12] Bioassays were conducted with Pa-Lab population at regular intervals to monitor the progression of susceptibility in the absence of selection pressure. The median lethal doses (LC_{50}) estimated at the 70^{th} generation were used as a reference for comparing the response of field populations to acaricides.

In order to evaluate the status of acaricide resistance in the field populations of *P. latus*, roving surveys were conducted from July 2020 to March 2022 in the major chilli growing places of Karnataka, India where the acaricides and insecticides were used indiscriminately. The data on frequency of acaricide applications and percentage of farmers reporting control failures were collected from each location from at least ten farmers. Mite populations were collected from five locations, *viz*., Bangalore (13.0713° N, 77.5905° E), Chikkaballapur (13.1432° N, 77.6428° E), Haveri (14.6834° N, 75.3849° E), Tumkur (13.4489° N, 77.1678° E) and Ramnagara (12.6254° N, 77.2319° E). The collected mites were released on uninfested mulberry plants under growth chamber conditions for mass multiplication. Adult female mites from the F_1 generation were used in resistance monitoring bioassays.

Acaricides and bioassays

Five acaricides approved by the Central Insecticides Board and Registration Committee (CIBRC),[11] India for the control of broad mites, representing different chemical groups were chosen. Commercial formulations of diafenthiuron 50% WP, dicofol 18.5% EC, fenazaquin 10% EC, propargite 57% EC and spiromesifen 22.9 % SC were procured from local retailers. Different concentrations of each chemical that bracketed five to 95 percent mortality were recognized on the basis of preliminary bioassay studies. A minimum of five required concentrations were prepared by serial dilution from stock solutions of acaricides and used for conducting bioassays.

The leaf dip bioassay (Method No. 4) standardized by the Insecticide Resistance Action Committee (IRAC, 2009)[13] was adopted with appropriate modifications. Fresh mulberry leaf discs were immersed in the respective test chemical solutions for 30 seconds and allowed to air dry at room temperature. An untreated control was maintained by dipping the leaf discs in

distilled water. The leaf discs were placed on wet cotton wads in Petri plates that were kept moistened to maintain leaf disc turgidity and to check the escape of mites. Thirty adult female mites were gently transferred onto each leaf disc and kept in a BOD incubator at a temperature of 25 ± 1 ℃ and relative humidity of 65 ± 10%. Each concentration and the respective control treatments were replicated thrice and observations on mortality were documented after 24 hours of acaricide treatment. Moribund mites that were unable to show any signs of movement when probed with a fine brush were considered dead.

Statistical analysis

The median lethal concentrations (LC_{50}) and their 95 percent confidential limits (CLs) were determined by Probit analysis[14] using Polo Plus 2.0 software.[15] For the susceptible strain, susceptibility indices were calculated on the basis of LC_{50} values obtained for the F_1 and F_{70} generations.[16]

Susceptibility index, SI = LC_{50} of F_1/ LC_{50} of F_{70}

Other parameters namely, the rate of resistance decline (R) and the number of generations required for a ten-fold decrease in LC_{50} (G) were calculated as described by Tabashnik.[17]

Rate of resistance decline, R = [log (final LC_{50}) - log (initial LC_{50})]/n

The rate of resistance decline that quantifies the rate of change in LC_{50} when the selection pressure is withdrawn was estimated by using the number of generations not exposed to insecticide (n), the LC_{50} of the parental generations before 'n' generations (initial LC_{50}) and the LC_{50} after 'n' generations without selection (final LC_{50}).

The number of generations (G) required for a ten-fold decrease in the LC_{50} value was calculated using the formula: G = 1/R.

For the field populations, lethal concentration ratios (LCRs) at LC_{50} (LCR 50) and their 95% confidential limits were calculated and the tests for the hypotheses of equality and parallelism were performed as given by Robertson *et al*.[18] Further, resistance ratios (RR) were calculated by dividing the LC_{50} of the corresponding field population by that of the susceptible population. Based on the RR values, the intensity of resistance was categorized as low (RR <10), moderate (10> RR <40), high (40> RR <160) and extremely high (RR >160).[19] To ascertain cross-resistance, pairwise correlation coefficients between the LC_{50} values of field populations for the acaricides were evaluated by Pearson's correlation analysis and visualized using Graphpad prism.

Results

Field surveys and pest identity

Field surveys indicated partial to total control failures using acaricides which was highest for spiromesifen (66.67%), followed by diafenthiuron (63.33), fenazaquin (58.33%), dicofol (55.56%) and propargite. On an average, inadequate control of the mite with the selected acaricides was reported by 57.86 % of the farmers interviewed. The taxonomic identity of *P. latus* was confirmed both by morphological and molecular methods. A 631 bp long sequence was deposited in the NCBI-GenBank and BOLD databases (Accession number: ON103156; BIN: AED8321).

Stability of acaricide resistance

Bioassays with Pa-Lab showed a decrease in resistance levels over the generations (Table 1). The initial LC_{50} values at the F_1 generation were 34.48 ppm for diafenthiuron, 37.10 ppm for dicofol, 17.03 ppm for fenazaquin, 5.25 ppm for propargite and 59.16 ppm for spiromesifen. At the 70^{th} generation, the population was found to be highly susceptible to propargite with an LC_{50}

of 0.37 ppm, followed by diafenthiuron (0.40 ppm), fenazaquin (0.44 ppm), spiromesifen (0.58 ppm) and dicofol (0.70 ppm).

The LC_{50} values obtained for F_1 were compared with that of F_{70} to generate the susceptibility index (SI) which was observed to be the highest for spiromesifen (102.53) (Table 2). The susceptibility indices for other chemicals were found to be 86.4 (diafenthiuron), 52.77 (dicofol), 38.88 (fenazaquin) and 14.11 (propargite). Propargite recorded the highest number of generations required for a tenfold decrease in LC_{50} (43.48 generations) which was followed by fenazaquin (31.50), dicofol (29.00), diafenthiuron (25.83) and spiromesifen (24.89).

The values of 'response to selection (R)' varied from -0.023 (propargite) to -0.040 (spiromesifen). The consistent negative value of 'R' indicated a decline in resistance to acaricides over successive generations. The highest values for susceptibility index and response to selection in spiromesifen indicated a rapid reversion of mite population to susceptibility for the chemical when there was no selection pressure.

Acaricide resistance in field mite populations

Significant differences were recorded in LC_{50} of the populations with respect to the susceptible population for all the tested chemicals as the 95 percent confidential limits (CLs) of their LCRs did not contain the value of 1.0 (Table 3).

Observations on resistance of populations to diafenthiuron showed a high level of resistance development in Haveri (LC_{50} 32.38 ppm), Tumkur (LC_{50} 32.00) and Chikkaballapur populations (26.00 ppm). The tests for hypotheses of equality (of slopes and intercepts) and parallelism (of slopes) between the populations disclosed that the regression lines were neither equal ($\chi^2 = 496$; df = 10; $P < 0.05$) nor parallel ($\chi^2 = 15.48$; df = 5; $P = 0.01$).

Three of the populations were highly resistant to dicofol with the highest RR for the populations collected from Haveri (83.47 folds), followed by Chikkaballapur (49.29 folds) and Ramanagara (46.16 folds) (Figure 1). Bangalore and Tumkur populations recorded moderate levels of resistance with RRs of 28.47 and 27.35 folds, respectively. The hypothesis of parallelism was accepted (χ^2 = 4.94; df = 5; P = 0.42) but that of equality was rejected (χ^2 = 467; df = 10; P < 0.05).

The bioassays with fenazaquin indicated a low level of resistance in the Bangalore population (9.72 folds), moderate resistance in Ramanagara (18.62 folds), Tumkur (28.61 folds) and Chikkaballapur (33.42 folds) populations and high level of resistance in Haveri population (45.42 folds). The tests for equality and parallelism hypotheses between the populations revealed that regression lines were not equal (χ^2 = 470; df = 10; P < 0.05) but were parallel (χ^2 = 6.47; df = 5; P = 0.26) for fenazaquin.

Results of dose-responses of *P. latus* populations to propargite revealed a relatively narrow range of LC_{50} values (3.26 ppm to 6.27 ppm) and they differed significantly from the susceptible strain. The resistance ratios varied from 8.77 to 16.84 folds representing low to moderate resistance levels. The tests for hypotheses of parallelism and equality between the populations disclosed that the regression lines were parallel (χ^2 = 4.24; df = 5; P = 0.52) but not equal (χ^2 = 266; df = 10; P < 0.05).

Field populations of *P. latus* showed wider variation in susceptibility to spiromesifen with LC_{50} ranging from 27.91 to 94.28 ppm. *P. latus* collected from the Haveri region showed an extremely high level of resistance (163.40 folds) followed by the populations obtained from Bangalore, Chikkaballapur, Ramanagara and Tumkur (55.15, 93.41, 48.37 and 101.29 folds,

respectively). The hypothesis of parallelism of slopes was accepted (χ^2 = 4.89; df = 5; P = 0.43) but that of equality of slopes and intercepts was rejected (χ^2 = 553; df = 10; P < 0.05).

Assessment of cross-resistance by pairwise correlation analysis

To assess the cross-resistance amongst different classes of acaricides, Pearson's correlation analysis was performed (Figure 2). The perusal of data in Figure 2 revealed that resistance to spiromesifen had a positive and significant correlation with that of fenazaquin resistance (correlation coefficient r = 0.929, p < 0.05). On the other hand, a positive but non-significant correlation was observed between resistance to all other acaricides in field populations of *P. latus* tested (p > 0.05).

Discussion

The broad mite, *P. latus* is distributed worldwide and its population build-up is strongly favored by many intrinsic and extrinsic factors such as short life cycle, sex ratio, arrhenotokous reproduction, absence of effective natural enemies, host plants, temperature, humidity and other micro-climatic conditions, especially under protected cultivation.[20,21,22] The indiscriminate application of chemical pesticides in many crop eco-systems and under protected cultivations causes the elimination of natural enemies and development of resistance in many populations of Tetranychid mites like *Tetranychus urticae, Panonychus citri* and *P. ulmi.*[23] A similar situation was perceived in the case of broad mite outbreak in red pepper and capsicum in the sampled places in the current investigation where inadequate control of the mite with any of the selected acaricides was reported by 57.86 % of the farmers interviewed. However, compared to spider mites and other insect pests, there are no field studies on the broad mite resistance to acaricides in India and abroad despite frequent applications of heavy doses of various acaricides.

The reversal of field evolved resistance over the generations (14.11 to 102.53 folds) in the absence of selection indicates the recessive nature of the genes involved in the resistance. The recessive mode of inheritance of resistance could be easily tackled by temporary withdrawal of that particular acaricides from field usage for a few years.[24] Similar results were observed in the red spider mite, *T. urticae* where a significant decline in resistance in the absence of selection pressure under laboratory conditions was reported against acaricides like fenpyroximate,[25] abamectin,[26] spirodiclofen[27] and milbemectin.[28] Specifically, susceptibility was restored to 282, 89, 31 and 221 folds for dicofol, fenazaquin, propargite and spiromesifen, respectively at the end of 91 generations of laboratory rearing.[29]

In the present study, the susceptible population of the mite regained its partial to full susceptibility to the tested acaricides by the 70th generation under laboratory conditions. Similarly, Mohin[30] reported baseline LC_{50} values in *T. urticae* after 128th generation which were 0.18 ppm for fenazaquin, 0.20 ppm for propargite, 0.29 ppm for spiromesifen and 0.30 ppm for diafenthiuron and dicofol. Naveena *et al*.[31] calculated the LC_{50} values of laboratory population of *T. urticae* at the 25th generation and observed the highest susceptibility to fenazaquin (LC_{50} of 0.11 ppm) followed by fenpropathrin (0.12 ppm), chlorfenapyr (0.15 ppm), diafenthiuron (0.22 ppm), propargite (0.91 ppm) and spiromesifen (2.00 ppm), respectively.

The tested field populations showed high to extremely high resistance to spiromesifen (up to 163.39 folds) (Figure 1). Being an insecticide-cum-acaricide, this chemical is frequently used against the sucking pests complex[32,33] especially in chilli and capsicum ecosystems which might have accelerated the resistance development to this chemical. Three populations showed significantly high resistance to diafenthiuron (mean RR of 56.21) which is another broad-

spectrum insecticide-acaricide widely used against sucking pests in Karnataka. Furthermore, the lowest levels of resistance were observed in case of propargite (mean 13.42).

In *T. urticae* populations of Karnataka, the levels of resistance were high to extremely high (143 to 1038 folds) for dicofol, moderate (15.65 to 32.83 folds) for propargite, moderate to high (12.02 to 75.00 folds) for fenazaquin while that for spiromesifen was extremely high (431.26 to 969.10 folds).[34] In Tamil Nadu, the resistance levels in *T. urticae* to major acaricides were monitored and a low level of resistance (2.00 to 8.62 folds) to fenazaquin, low to moderate level of resistance to fenpropathrin (1.86 to 37.28 folds), moderate to a high level of resistance to diafenthiuron (15.81 to 50.53 folds), high level of resistance to propargite (45.16 to 65.10 folds) and extremely high level of resistance to spiromesifen (193.04 to 452.61-fold) were recorded.[31] However, comparatively lower levels of resistance to the aforesaid acaricides were reported previously in *T. urticae* populations of Punjab,[35] Himachal Pradesh[36] and Kerala[37] states.

In the present investigation, the analysis methodology combining LCRs, their 95 percent CLs and the tests of hypotheses of parallelism and equality were used where dose-response regressions are considered as evidence for acaricide resistance.[18] This method is more edifying and has greater statistical power. According to the tests for equality and parallelism hypotheses between the populations, the regression lines were parallel but not equal for all the tested chemicals except diafenthiuron. This is indicative of the heterogeneity of the experimental populations which could imply that the populations are typical of a range of both resistant and susceptible individuals. This might be attributed to polyphagous nature of the pest and its inter-seasonal movement across different crops.

Studies on cross-resistance between pesticides are of utmost importance as alteration, rotation and mixing of pesticides are common strategies to delay or avoid the development of

resistance.[38] The pairwise correlation coefficients assessed between the values of LC_{50} of the tested chemicals revealed that there was a significant positive correlation between resistance to spiromesifen and fenazaquin. The lack of significant cross-resistance in other acaricides could guide their rotation and sequential application in the field.

Conclusion

The present study provides the first concerted effort on the establishment of reference susceptibility data and monitoring of resistance and cross-resistance to major acaricides in *P. latus*. The aforementioned studies are indispensable elements of a sustainable and economically viable management program for this pest and can also aid in reducing the pesticide load in the environment. The results of resistance studies revealed that field populations from five districts of Karnataka have developed resistance to acaricides, especially spiromesifen and diafenthiuron,. Hence, there is an imperative requirement for rotation or alteration of chemicals, especially insecticide-acaricide compounds like spiromesifen and diafenthiuron with others. The plausible biochemical and molecular bases of acaricide resistance in *P. latus* populations necessitate further investigations. Also, studies on the genetic and molecular aspects of *P. latus* needs to be extended to other hosts and different agroecosystems to maintain the efficacy of currently available acaricides.

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Tables

Table 1. Stability of resistance in *Polyphagotarsonemus latus* to different acaricides over generations in the absence of selection pressure

Acaricide	Generation	LC_{50} (95% CL)	Slope ± SEM	χ^2 (df)	Heterogeneity
Diafenthiuron	F_1	34.48 (20.576-56.520)	1.47 ± 0.18	4.47 (3)	1.49
	F_{20}	11.84 (9.491-14.455)	1.50 ± 0.16	1.12 (3)	0.37
	F_{40}	0.69 (0.567-0.828)	1.40 ± 0.12	2.55 (4)	0.64
	F_{60}	0.45 (0.374-0.527)	2.07 ± 0.21	0.43 (3)	0.14
	F_{70}	0.40 (0.339-0.465)	2.02 ± 0.18	1.17 (3)	0.39
Dicofol	F_1	37.10 (25.008-58.153)	1.40 ± 0.16	3.63 (3)	1.21
	F_{20}	9.72 (7.999-11.772)	1.55 ± 0.14	2.64 (4)	0.66
	F_{40}	1.52 (1.259-1.801)	1.76 ± 0.17	1.02 (3)	0.34
	F_{60}	0.77 (0.646-0.916)	1.82 ± 0.18	0.25 (3)	0.19
	F_{70}	0.70 (0.585-0.832)	1.92 ± 0.16	2.42 (3)	0.81
Fenazaquin	F_1	17.03 (14.071-20.514)	1.77 ± 0.19	2.46 (3)	0.82
	F_{20}	3.76 (3.123-4.484)	1.74 ± 0.17	1.21 (3)	0.40
	F_{40}	0.74 (0.615-0.876)	1.75 ± 0.16	0.62 (3)	0.21
	F_{60}	0.42 (0.309-0.548)	2.02 ± 0.19	3.01 (3)	1.00
	F_{70}	0.44 (0.370-0.513)	1.84 ± 0.17	0.34 (3)	0.11
Propargite	F_1	5.25 (3.537-7.351)	1.55 ± 0.16	3.30 (3)	1.10
	F_{20}	2.11 (1.471-2.986)	1.35 ± 0.14	4.32 (4)	1.08
	F_{40}	0.84 (0.632-1.090)	1.12 ± 0.13	3.72 (4)	0.93
	F_{60}	0.39 (0.317-0.472)	1.54 ± 0.14	1.48 (4)	0.37

	F_{70}	0.37 (0.233-0.546)	1.20 ± 0.13	4.95 (4)	1.24
Spiromesifen	F_1	59.16 (38.207-97.568)	1.64 ± 0.17	5.62 (3)	1.87
	F_{20}	23.73 (19.568-28.719)	1.61 ± 0.17	1.71 (3)	0.57
	F_{40}	1.31 (1.094-1.551)	1.63 ± 0.13	1.29 (4)	0.32
	F_{60}	0.63 (0.523-0.746)	1.98 ± 0.20	0.50 (3)	0.17
	F_{70}	0.58 (0.485-0.678)	1.91 ± 0.18	0.60 (3)	0.20

df- degrees of freedom

Table 2. Acaricide selection response in *P. latus* over generations

Acaricide	**Susceptibility index (SI)**	**Response to selection (R)**	**Resistance stability (G)**
Diafenthiuron	86.41	-0.039	25.83
Dicofol	52.77	-0.034	29.00
Fenazaquin	38.88	-0.032	31.50
Propargite	14.11	-0.023	43.48
Spiromesifen	102.53	-0.040	24.89

Table 3. Dose-responses of *Polyphagotarsonemus latus* populations to acaricides

Acaricide	Population	LC_{50} (95% CL)	Slope ± SE	χ^2 (df)	h	LCR (95% CL)
Diafenthiuron	Bangalore	12.00 (9.25-15.70)	1.21 ± 0.14	3.29 (4)	0.82	30.12 (22.15-40.94)
	Chikkaballapur	26.00 (13.48-40.04)	1.45 ± 0.17	5.88 (3)	1.96	65.22 (49.43-86.05)
	Haveri	32.38 (25.80-40.52)	1.48 ± 0.17	2.59 (3)	0.86	81.24 (61.70-106.94)
	Ramanagara	10.39 (7.21-14.88)	1.51 ± 0.14	5.74 (4)	1.43	26.05 (20.07-33.83)
	Tumkur	32.00 (26.34-38.57)	1.78 ± 0.18	1.03 (3)	0.34	80.28 (62.64-102.91)
Dicofol	Bangalore	20.01 (16.18-24.65)	1.34 ± 0.13	3.86 (4)	0.97	28.46 (21.62-37.45)
	Chikkaballapur	34.65 (20.45-60.27)	2.52 ± 0.19	4.97 (3)	1.66	49.28 (36.97-65.68)
	Haveri	58.68 (47.17-72.70)	1.61 ± 0.19	1.88 (3)	0.63	83.45 (63.21-110.16)
	Ramanagara	32.45 (17.25-59.39)	1.73 ± 0.20	7.54 (3)	2.51	46.14 (35.32-60.28)
	Tumkur	19.23 (11.49-37.29)	1.45 ± 0.18	5.18 (4)	1.73	27.34 (20.33-36.77)
Fenazaquin	Bangalore	4.26 (2.98-5.80)	1.59 ± 0.14	5.82 (4)	1.46	9.73 (7.57-12.51)
	Chikkaballapur	14.64 (11.87-17.77)	1.68 ± 0.18	2.31 (3)	0.77	33.45 (25.80-43.36)
	Haveri	19.89 (16.18-24.38)	2.84 ± 0.21	0.84 (4)	0.28	45.46 (35.00-59.05)
	Ramanagara	8.16 (6.56-10.03)	1.46 ± 0.16	0.72 (3)	0.24	18.64 (14.26-24.36)
	Tumkur	12.53 (10.31-14.93)	2.02 ± 0.21	1.48 (3)	0.49	28.63 (22.36-36.65)
Propargite	Bangalore	5.27 (4.27-6.48)	1.46 ± 0.13	2.93 (4)	0.73	14.17 (10.16-19.75)
	Chikkaballapur	6.27 (5.02-7.75)	1.53 ± 0.17	2.48 (3)	0.83	16.85 (12.03-23.61)
	Haveri	5.05 (1.97-9.35)	1.40 ± 0.18	6.42 (3)	2.14	13.58 (9.39-19.62)
	Ramanagara	5.11 (3.73- 7.07)	1.44 ± 0.12	5.04 (4)	1.26	13.76 (9.94-19.05)
	Tumkur	3.26 (2.63-4.04)	1.55 ± 0.17	1.64 (3)	0.55	8.78 (6.28-12.27)
Spiromesifen	Bangalore	31.82 (19.01-57.60)	1.63 ± 0.20	5.20 (3)	1.73	55.12 (41.63-72.98)

	Chikkaballapur	53.90 (30.19-103.44)	1.85 ± 0.19	9.07 (3)	3.02	93.37 (72.81-119.72)
	Haveri	94.28 (64.93-136.55)	2.07 ± 0.19	4.68 (4)	1.56	163.31 (128.38-207.75)
	Ramanagara	27.91 (15.93-49.46)	1.63 ± 0.18	6.74 (3)	2.25	48.35 (37.22-62.81)
	Tumkur	58.48 (28.09-149.49)	1.51 ± 0.18	9.18 (3)	3.06	101.25 (76.56-133.90)

h- heterogeneity; df- degrees of freedom

LCR- Lethal concentration ratios at LC_{50}

Figures

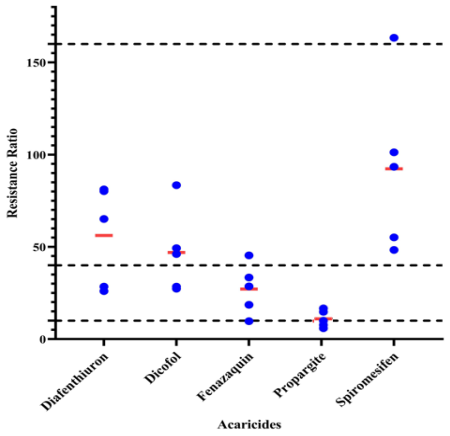

Figure 1. Resistance levels of field populations of *P. latus* to major acaricides. The red horizontal lines denote the mean of the resistance ratios of the different field populations.

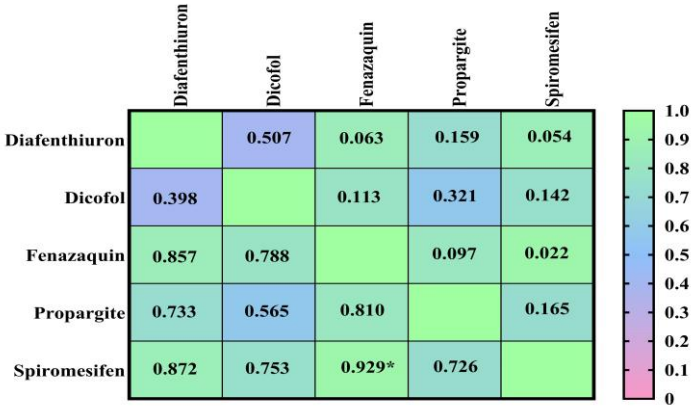

Figure 2. Pairwise correlation analysis of LC_{50} values for five acaricides in the field populations of *P. latus*. The values below the unfilled diagonal squares indicate the correlation coefficient r and those above the unfilled diagonal squares indicate the corresponding p values. The scale colors of the filled boxes indicate the magnitude of the correlation. * represents a significant correlation at $p < 0.05$.