

Full Length Research Paper

Euphorbiaceous Plant Formulations *vis-a-vis* Rate of development of the Bruchid *Callosobruchus chinensis* Linn.

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Abstract

Pulses constitute major source of protein in the diet of people of developing countries and play an important role in Indian economy and are traditionally recognized as an indispensable constituent of Indian food. The genus *Callosobruchus* attacks grain legumes during both pre and post harvest stages all over the world. Efficient control of stored grain pests has long been the aim of entomologists and the present work was carried out to screen certain formulations against the pulse beetle *Callosobruchus chinensis* Linn. raised on grains of *Vigna radiata*. The plants selected for the study includes *Euphorbia hirta*, *Phyllanthus amarus* and *Jatropha gossypifolia* all belonging to family Euphorbiaceae. Different formulations using leaf of the plants were employed in the form of crude extract, aqueous suspension, aqueous extract, ethanol extract and diethyl ether extract. The treatments were made using different dose concentrations of 1, 5, 10 and 25% and rate of development was documented. It could be concluded from the study that plants belonging to family Euphorbiaceae do possess chemicals, which are toxic to insects and therefore have a potential to be used against the pest *C. chinensis*. Although, the three select plants showed varied degree of effect on rate of development, *P. amarus* seemed to be the most effective.

Key words: *Callosobruchus chinensis*, Euphorbiaceae, *Euphorbia hirta*, *Phyllanthus amarus*, *Jatropha gossypifolia*, Plant formulations, Rate of development.

Introduction

In India, where the population is predominantly vegetarian, pulses are the most important and rich source of protein and several amino acids. Besides, these also provide energy, minerals and certain vitamins. Pulses contain 20-30% of protein which is almost three times higher than cereals (Doharey et al., 1990). According to reports available, in India, over 200 species of insects have been recorded infesting various pulses (CABI, 2007). Of these, 17 species of bruchids belonging to 11 genera have been recorded infesting different pulses (Arora, 1977). The genus *Callosobruchus* attacks grain legumes during both pre and post harvest stages all over the world; but in India, *C. maculatus*, *C. analis* and *C. chinensis* are the predominant pest species of the genera (Dias, 1988). The bruchid infestation also affects seed quality, market value and can reduce cowpea seed viability to 2% after months of storage (Caswell, 1980).

Efficient control of stored grain pests has long been the aim of entomologists throughout the world and synthetic chemical pesticides have been used for many years to control stored grain pests (Salem et al., 2007; Ani, 2010; Bhalla et al., 2008). However, the persistent use of these insecticides in granaries of small-scale farmers has led to a number of problems such as killing of non-target species, user hazards, toxic residues in food, development of genetic resistance in the treated pest, increased cost of application and the destruction of the balance of the ecosystem (Shaheen & Khaliq, 2005; Boateng & Kusi, 2008). Historical usage of nicotine and pyrethrum has encouraged scientists to focus their attention on alkaloids, flavonoids, terpenoids and other secondary compounds to be used as pest control agents (Rajapakse & Ratnasekera, 2008) and are working for the development and establishment of plant based pesticides. Over 200 plant species have been reported to have insecticidal properties capable of controlling insects (Golob & Webley 1980). Various works screening different plants against stored grain pests include those by Negi et al. (1997), Srivastava & Mann (2002a), Srivastava & Mann (2002b), Kaur & Srivastava (2004), Srivastava & Gupta (2007), Srivastava & Ghei (2007), Gupta & Srivastava (2008), Kiradoo & Srivastava (2010), Kiradoo & Srivastava (2011), Rawat & Srivastava (2011), Rawat & Srivastava (2012), Mann & Srivastava (2013a, b, c, d, e, f), Kosar & Srivastava (2013), Mann & Srivastava (2014).

Callosobruchus chinensis shows clear sexual dimorphism. The female beetle lays eggs on the healthy seed surface and the larva immediately after hatching bores into the seed. By the time it reaches the adult stage it consumes the seed cotyledons. The adult emerges out leaving behind holed grains.

The plant family Euphorbiaceae is a large family of flowering plants with 300 genera and around 7,500 species. A number of plants of this family are of considerable economic importance. Some members of Euphorbiaceae have medicinal properties in

these plants *P. amarus*, *P. nirurim*, *E. hirta*, *J. curcas* and *J. gossypifolia* are included. These plants are also reported to possess insecticidal activity. Among the many plant species that have been used to control stored product pests is the physic nut, *Jatropha curcas* L. plant. The efficacy of *Jatropha* seed oil against insect has been reported by Huis (1991), Adabie- Gomez et al. (2006) and Henning (2007). Aqueous leaf extract of *Ricinus communis* L (Euphorbiaceae), a cultivated plant in tropical countries, showed excellent insecticidal activity against *Callosobruchus chinensis* L (Coleoptera: Bruchidae) as documented by Upasani et al. (2003). Repellency of hydroethanolic extracts of *Ricinus communis* to *Scyphophorus acupunctatus* in the laboratory was studied by Cinthia et al. (2012).

The present work was therefore carried out to screen *Euphorbia hirta*, *Phyllanthus amarus* and *Jatropha gossypifolia* all belonging to family Euphorbiaceae formulations against the pulse beetle *Callosobruchus chinensis* Linn. and document the rate of development (days) of the pest insect.

Materials and Methods

The test insect selected for the investigation pulse beetle *Callosobruchus chinensis* L. was raised on green gram *Vigna radiata* maintained at 28±2°C temperature and 70% relative humidity. The three plants viz., *Euphorbia hirta* Linn., *Phyllanthus amarus* Linn. and *Jatropha gossypifolia* Linn. used for the study were collected from in and around Bikaner city (27°11' & 20° 03' North latitude and 71°54' & 74°12' East longitude). The plant part used was leaves which were picked and washed. For crude extracts fresh leaves were taken while, for rest of the formulations they were shade dried for 10-15 days and were ground separately in electric grinder and kept in airtight plastic container for further use. Formulations of 1, 5, 10 and 25% concentrations were prepared. These included aqueous suspension, aqueous extract, ethanol extract and diethyl ether extract. Normal, control and five experimental sets were laid out. Each set comprised of three replicas. To document the aspect selected, time taken for the development of adult from the day of egg laying, up to the emergence, was recorded in days and expressed as rate of development. The data obtained was subjected to analysis of variance (ANOVA) using MS-Excel software. The critical difference at 1 and 5% level of significance was worked out.

Results

The mean rate of development (days) by the bruchid *C. chinensis* under different treatments of various plants studied has been presented in Table 1a and ANOVA has been presented in Tables 1b to 1n.

During the present study mean rate of development (days) by *C. chinensis* in normal sets it was observed to be 27 (days), while in control sets treated with GDW it was noted to be 29 and in sets treated with ethanol as well as DEE extracts it was observed to be 35 days. Highest rate of development (120 days) was observed in experimental sets treated with crude extract of all concentrations of *P. amarus*. Over all, besides crude extract, aqueous suspension of the same plant was also found to significantly increase the rate of development of the pest insect.

For comparing the effect of different formulations, ANOVA was applied and has been presented in Table 1b. Further, based on this analysis Tables 1c to 1n were constructed. The perusal of the number of days taken to develop by the pest insect was significantly increased in various experimental sets.

Table 1b clearly indicates that the results of rate of development (days) taken by the beetle, pertaining to efficacy of plants (A), extracts (B), concentration, (C) treatments (D) and plants extracts (AXB), plants and concentrations (AXC), plants and treatments (AXD), extracts and concentrations (BXC), extracts and treatments (BXD), concentrations and treatments (CXD), plants, extracts and concentrations (AXBXC), plants, extracts and treatments (AXBXD), plants, concentrations and treatments (AXCXD), extracts, concentrations and treatments (BXCXD) and plants, extracts, concentrations and treatments (AXBXCXD) were highly significant ($p < 0.01$). When comparisons were made on the basis of ANOVA (Table 1c) with respect to the efficacy of the three plants studied on the rate of development of the bruchid, it was found that only treatments of *P. amarus* significantly ($p < 0.05$) increased the development of bruchid and this treatment significantly differed from treatments of *J. gossypifolia* and *E. hirta*.

On the basis of ANOVA (Table 1c) it could further be inferred that various other treatments were also effective in increasing the time taken for development of *C. chinensis*. In crude extract rate of development was observed to be significantly ($p < 0.05$) higher from rest of the formulations, viz., aqueous suspension, aqueous extract, ethanol and DEE extracts.

When the effect of different concentrations on the rate of development of the pest was compared, it was noted that only 25% extracts resulted in significantly ($p < 0.05$) more time taken by the bruchid for development. While comparing the effect of plants and extract, it was observed that a significantly ($p < 0.05$) higher rate of development was found in experimental sets treated with crude extracts of *P. amarus* as compared to rest of the plant formulations (Table 1d).

While comparing the effect of plants and their different concentrations it was observed that a significantly ($p < 0.05$) higher rate of development of the bruchid was found in experimental sets treated with 25% concentration of *P. amarus* as compared to rest of the plant formulations (Table 1c).

Further, when the effect of plants and treatments on the rate of development of the bruchid was observed it was found that all the treatments resulted in significantly ($p < 0.05$) increased rate of development (Table 1f).

Table 1a. Mean rate of development (days) of *C. chinensis* under different formulations of leaves of select three plants

Treatments	Plants → Conc. ↓	<i>Euphorbia hirta</i>	<i>Phyllanthus amarus</i>	<i>Jatropha gossypifolia</i>
Crude extract	Normal	27.00 ± 0.00	27.00 ± 0.00	27.00 ± 0.00
	Control	29.00 ± 0.00	29.00 ± 0.00	29.00 ± 0.00
	1%	31.20 ± 1.92	120.00 ± 0.00	60.00 ± 0.00
	5%	31.60 ± 0.55	120.00 ± 0.00	58.00 ± 2.74
	10%	31.00 ± 1.00	120.00 ± 0.00	58.00 ± 2.74
	25%	30.40 ± 0.55	119.20 ± 1.10	58.00 ± 2.74
Aqueous suspension	Normal	27.00 ± 0.00	27.00 ± 0.00	27.00 ± 0.00
	Control	29.00 ± 0.00	29.00 ± 0.00	29.00 ± 0.00
	1%	31.40 ± 0.89	106.00 ± 5.48	38.60 ± 2.88
	5%	30.60 ± 0.89	106.00 ± 5.48	38.40 ± 3.13
	10%	30.40 ± 0.89	106.00 ± 5.48	38.80 ± 2.68
	25%	29.40 ± 0.89	106.00 ± 5.48	38.60 ± 2.97
Aqueous extract	Normal	27.00 ± 0.00	27.00 ± 0.00	27.00 ± 0.00
	Control	29.00 ± 0.00	29.00 ± 0.00	29.00 ± 0.00
	1%	30.20 ± 0.45	81.00 ± 2.24	29.60 ± 2.07
	5%	30.20 ± 1.79	83.00 ± 2.74	29.40 ± 1.95
	10%	30.00 ± 1.41	83.00 ± 2.74	29.60 ± 2.07
	25%	29.20 ± 1.79	81.00 ± 2.24	29.20 ± 2.17
Ethanol extract	Normal	27.00 ± 0.00	27.00 ± 0.00	27.00 ± 0.00
	Control	35.00 ± 0.00	35.00 ± 0.00	35.00 ± 0.00
	1%	31.60 ± 0.55	40.00 ± 0.00	32.40 ± 1.52
	5%	31.40 ± 0.55	40.00 ± 0.00	32.20 ± 1.64
	10%	30.60 ± 0.55	39.00 ± 2.24	31.80 ± 1.48
	25%	28.80 ± 0.84	38.00 ± 2.74	31.80 ± 0.45
Di-ethyl ether extract	Normal	27.00 ± 0.00	27.00 ± 0.00	27.00 ± 0.00
	Control	35.00 ± 0.00	35.00 ± 0.00	35.00 ± 0.00
	1%	31.20 ± 0.45	40.60 ± 0.89	34.00 ± 6.56
	5%	31.20 ± 0.84	40.00 ± 2.92	34.00 ± 6.56
	10%	30.00 ± 1.41	40.60 ± 6.19	34.00 ± 6.56
	25%	28.80 ± 0.45	39.80 ± 2.77	34.00 ± 6.60

Values given are mean ± SD

Table 1b. ANOVA for rate of development showing different interactions and level of significance

Source of Variations	df	SS	MSS	F-cal	S/NS	S.Em.	CD 5%	CD 1%
A	2	51808.03	25904.01	3183.97	**	0.16	0.46	0.60
B	4	12649.05	3162.26	388.69	**	0.21	0.59	0.78
C	3	243.59	81.20	9.98	**	0.19	0.53	0.69
D	3	90512.75	30170.92	3708.43	**	0.16	0.46	0.60
A x B	8	17907.18	2238.40	275.13	**	0.37	1.02	1.35
A x C	6	815.10	135.85	16.70	**	0.33	0.91	1.20
A x D	6	93682.19	15613.70	1919.14	**	0.29	0.79	1.04
B x C	12	593.48	49.46	6.08	**	0.43	1.18	1.55
B x D	12	39611.96	3301.00	405.74	**	0.37	1.02	1.35
C x D	9	487.18	54.13	6.65	**	0.33	0.91	1.20
A x B x C	24	1322.40	55.10	6.77	**	0.74	2.05	2.69
A x B x D	24	38588.90	1607.87	197.63	**	0.64	1.77	2.33
A x C x D	18	1630.20	90.57	11.13	**	0.57	1.58	2.08
B x C x D	36	1186.95	32.97	4.05	**	0.74	2.05	2.69
A x B x C x D	72	2644.80	36.73	4.52	**	1.28	3.54	4.66
Error	660	5369.60	8.14					
Mean	899	359053.36						

* 5% level of significance

** 1% level of significance

S.Em.- standard error of mean

C.D.- Critical difference

MSS- Mean sum of square

SS- Sum of square

A- Plants

B- Extracts

C- Concentrations

D- All Treatments

Table 1c. Comparison of different formulations with respect to overall mean of rate of development (ANOVA) and critical difference

A1	46.59
A2	29.10
A3	32.39
S.Em±	0.16
CD (5%)	0.46
CD (1%)	0.60
B1	42.04
B2	38.23
B3	35.32
B4	32.82
B5	31.73
S.Em±	0.21
CD (5%)	0.59
CD (1%)	0.78
C1	36.92
C2	35.59
C3	35.80
C4	35.80
S.Em±	0.19
CD (5%)	0.53
CD (1%)	0.69
D1	27.00
D2	31.07
D3	50.01
S.Em±	0.33
CD (5%)	0.91
CD (1%)	1.20

A1- *P. amarus*

B1- Crude extract

B4- Ethanol extract

C1- 25%

C4- 1%

D1- Normal

A2- *E. hirta*

B2- Aqueous suspension

B5- DEE

C2- 10%

D2- Control

A3- *J. gossypifolia*

B3- Aqueous extract

C3- 5%

D3- Treatments

Table 1d. Comparison of rate of development of *C. chinensis* with respect to plants and extracts (A x B)

	B1	B2	B3	B4	B5	Mean
A1	58.93	54.33	48.42	37.17	34.08	46.59
A2	29.02	28.82	28.63	29.93	29.10	29.10
A3	38.17	31.53	28.92	31.35	32.00	32.39
Mean	42.04	38.23	35.32	32.82	31.73	
S.Em.±	0.37					
C.D. (5%)	1.02					
C.D. (1%)	1.35					

Table 1e. Comparison of rate of development of *C. chinensis* with respect to plants and concentrations (A x C)

	C1	C2	C3	C4	Mean
A1	49.80	45.37	45.53	45.64	46.59
A2	28.71	29.12	29.27	29.31	29.10
A3	32.24	32.28	32.61	32.44	32.39
Mean	36.92	35.59	35.80	35.80	
S.Em.±	0.33				
C.D. (5%)	0.91				
C.D. (1%)	1.20				

Table 1f. Comparison of rate of development of *C. chinensis* with respect to plants and treatments (A x D)

	A1	A2	A3	Mean
D1	27.00	27.00	27.00	27.00
D2	32.00	29.80	31.40	31.07
D3	80.76	30.50	38.78	50.01
Mean	46.59	29.10	32.39	
S.Em.±	0.29			
C.D. (5%)	0.79			
C.D. (1%)	1.04			

Table 1g. Comparison of rate of development of *C. chinensis* with respect to extracts and concentrations (B x C)

	B1	B2	B3	B4	B5	Mean
C1	41.84	38.11	37.04	36.07	31.51	36.92
C2	42.00	38.24	34.40	31.58	31.73	35.59
C3	42.07	38.22	35.20	31.73	31.80	35.80
C4	42.24	38.33	34.64	31.89	31.87	35.80
Mean	42.04	38.23	35.32	32.82	31.73	
S.Em.±	0.43					
C.D. (5%)	1.18					
C.D. (1%)	1.55					

Table 1h. Comparison of rate of development of *C. chinensis* with respect to extracts and treatments (B x D)

	B1	B2	B3	B4	B5	Mean
D1	27.00	27.00	27.00	27.00	27.00	27.00
D2	29.33	29.33	29.33	34.00	33.33	31.07
D3	69.78	58.35	49.63	37.45	34.85	50.01
Mean	42.04	38.23	35.32	32.82	31.73	
S.Em.±	0.37					
C.D. (5%)	1.02					
C.D. (1%)	1.35					

Table 1i. Comparison of rate of development of *C. chinensis* with respect to concentrations and treatments (C x D)

	C1	C2	C3	C4	Mean
C1	27.00	27.00	27.00	27.00	27.00
C2	31.07	31.07	31.07	31.07	31.07
C3	52.68	48.71	49.35	49.32	50.01
Mean	36.92	35.59	35.80	35.80	
S.Em.±	0.33				
C.D. (5%)	0.91				
C.D. (1%)	1.20				

Table 1j. Comparison of rate of development of *C. chinensis* with respect to plants, extracts and concentrations (A x B x C)

		C1	C2	C3	C4	Mean
A1	B1	58.73	59.00	59.00	59.00	58.93
	B2	54.33	54.33	54.33	54.33	54.33
	B3	54.33	46.00	46.67	46.67	48.42
	B4	47.67	33.33	33.67	34.00	37.17
	B5	33.93	34.20	34.00	34.20	34.08
A2	B1	28.80	29.00	29.20	29.07	29.02
	B2	28.47	28.80	28.87	29.13	28.82
	B3	28.40	28.67	28.73	28.73	28.63
	B4	29.27	30.13	30.13	30.20	29.93
	B5	28.60	29.00	29.40	29.40	29.10
A3	B1	38.00	38.00	38.00	38.67	38.17
	B2	31.53	31.60	31.47	31.53	31.53
	B3	28.40	28.53	30.20	28.53	28.92

	B4	31.27	31.27	31.40	31.47	31.35
	B5	32.00	32.00	32.00	32.00	32.00
	mean	36.92	35.59	35.80	35.80	
S.Em.±	0.74					
C.D. (5%)	2.05					
C.D. (1%)	2.69					

Table 1k. Comparison of rate of development of *C. chinensis* with respect to plants, extracts and treatments (A x B x D)

		D1	D2	D3	Mean	
A1	B1	27.00	30.00	119.80	58.93	
	B2	27.00	30.00	106.00	54.33	
	B3	27.00	30.00	88.25	48.42	
	B4	27.00	35.00	49.50	37.17	
	B5	27.00	35.00	40.25	34.08	
A2	B1	27.00	29.00	31.05	29.02	
	B2	27.00	29.00	30.45	28.82	
	B3	27.00	29.00	29.90	28.63	
	B4	27.00	32.00	30.80	29.93	
	B5	27.00	30.00	30.30	29.10	
A3	B1	27.00	29.00	58.50	38.17	
	B2	27.00	29.00	38.60	31.53	
	B3	27.00	29.00	30.75	28.92	
	B4	27.00	35.00	32.05	31.35	
	B5	27.00	35.00	34.00	32.00	
	Mean	27.00	31.07	50.01		
S.Em.±	0.64					
C.D. (5%)	1.77					
C.D. (1%)	2.33					

Table 1l. Comparison of rate of development of *C. chinensis* with respect to plants, concentrations and treatments (A x C x D)

		D1	D2	D3	Mean	
A1	C1	27.00	32.00	90.40	49.80	
	C2	27.00	32.00	77.12	45.37	
	C3	27.00	32.00	77.60	45.53	
	C4	27.00	32.00	77.92	45.64	
A2	C1	27.00	29.80	29.32	28.71	
	C2	27.00	29.80	30.56	29.12	
	C3	27.00	29.80	31.00	29.27	
	C4	27.00	29.80	31.12	29.31	
A3	C1	27.00	31.40	38.32	32.24	
	C2	27.00	31.40	38.44	32.28	
	C3	27.00	31.40	39.44	32.61	
	C4	27.00	31.40	38.92	32.44	
	Mean	27.00	31.07	50.01		
S.Em.±	0.57					
C.D. (5%)	1.58					
C.D. (1%)	2.08					

Table 1m. Comparison of rate of development of *C. chinensis* with respect to extracts, concentrations and treatments (B x C x D)

		D1	D2	D3	Mean
B1	C1	27.00	29.33	69.20	41.84
	C2	27.00	29.33	69.67	42.00
	C3	27.00	29.33	69.87	42.07
	C4	27.00	29.33	70.40	42.24
B2	C1	27.00	29.33	58.00	38.11
	C2	27.00	29.33	58.40	38.24

	C3	27.00	29.33	58.33	38.22
	C4	27.00	29.33	58.67	38.33
B3	C1	27.00	29.33	54.80	37.04
	C2	27.00	29.33	46.87	34.40
	C3	27.00	29.33	49.27	35.20
	C4	27.00	29.33	47.60	34.64
	C1	27.00	34.00	47.20	36.07
B4	C2	27.00	34.00	33.73	31.58
	C3	27.00	34.00	34.20	31.73
	C4	27.00	34.00	34.67	31.89
	C1	27.00	33.33	34.20	31.51
B5	C2	27.00	33.33	34.87	31.73
	C3	27.00	33.33	35.07	31.80
	C4	27.00	33.33	35.27	31.87
	Mean	27.00	31.07	50.01	
S.Em.±	0.74				
C.D. (5%)	2.05				
C.D. (1%)	2.69				

Table 1n. Comparison of rate of development of *C. chinensis* with respect to plants, extracts, concentrations and treatments (A x B x C x D)

			D1	D2	D3	Mean
A1	B1	C1	27.00	30.00	119.20	58.73
		C2	27.00	30.00	120.00	59.00
		C3	27.00	30.00	120.00	59.00
		C4	27.00	30.00	120.00	59.00
	B2	C1	27.00	30.00	106.00	54.33
		C2	27.00	30.00	106.00	54.33
		C3	27.00	30.00	106.00	54.33
		C4	27.00	30.00	106.00	54.33
	B3	C1	27.00	30.00	106.00	54.33
		C2	27.00	30.00	81.00	46.00
		C3	27.00	30.00	83.00	46.67
		C4	27.00	30.00	83.00	46.67
	B4	C1	27.00	35.00	81.00	47.67
		C2	27.00	35.00	38.00	33.33
		C3	27.00	35.00	39.00	33.67
		C4	27.00	35.00	40.00	34.00
	B5	C1	27.00	35.00	39.80	33.93
		C2	27.00	35.00	40.60	34.20
		C3	27.00	35.00	40.00	34.00
		C4	27.00	35.00	40.60	34.20
A2	B1	C1	27.00	29.00	30.40	28.80
		C2	27.00	29.00	31.00	29.00
		C3	27.00	29.00	31.60	29.20
		C4	27.00	29.00	31.20	29.07
	B2	C1	27.00	29.00	29.40	28.47
		C2	27.00	29.00	30.40	28.80
		C3	27.00	29.00	30.60	28.87
		C4	27.00	29.00	31.40	29.13
	B3	C1	27.00	29.00	29.20	28.40
		C2	27.00	29.00	30.00	28.67
		C3	27.00	29.00	30.20	28.73
		C4	27.00	29.00	30.20	28.73
	B4	C1	27.00	32.00	28.80	29.27
		C2	27.00	32.00	31.40	30.13

		C3	27.00	32.00	31.40	30.13
		C4	27.00	32.00	31.60	30.20
	B5	C1	27.00	30.00	28.80	28.60
		C2	27.00	30.00	30.00	29.00
		C3	27.00	30.00	31.20	29.40
		C4	27.00	30.00	31.20	29.40
A3	B1	C1	27.00	29.00	58.00	38.00
		C2	27.00	29.00	58.00	38.00
		C3	27.00	29.00	58.00	38.00
		C4	27.00	29.00	60.00	38.67
	B2	C1	27.00	29.00	38.60	31.53
		C2	27.00	29.00	38.80	31.60
		C3	27.00	29.00	38.40	31.47
		C4	27.00	29.00	38.60	31.53
	B3	C1	27.00	29.00	29.20	28.40
		C2	27.00	29.00	29.60	28.53
		C3	27.00	29.00	34.60	30.20
		C4	27.00	29.00	29.60	28.53
	B4	C1	27.00	35.00	31.80	31.27
		C2	27.00	35.00	31.80	31.27
		C3	27.00	35.00	32.20	31.40
		C4	27.00	35.00	32.40	31.47
	B5	C1	27.00	35.00	34.00	32.00
		C2	27.00	35.00	34.00	32.00
		C3	27.00	35.00	34.00	32.00
		C4	27.00	35.00	34.00	32.00
			Mean	27.00	31.07	50.01
	S.Em.±	1.28				
	C.D. (5%)	3.54				
	C.D. (1%)	4.66				

Discussion

The average number of days taken by *C. chinensis* to develop from egg and emerge out as adult has been considered as rate of development during the present study. Highest rate of development was observed in experimental sets treated with extracts of *P. amarus*. Bashir & El Shafie (2013), suggested *Jatropha* oil at 10% concentration resulted in delay of the development time of nymphal instar of *S. gregaria* from 5th to 6th nymphal instar by 5 days. Earlier Kiradoo (2009) found that treatments of *M. spicata* enhanced the time taken for development by *C. chinensis*. Kathuria & Kaushik (2006) also reported that the leaves of *O. sanctum L. adversely* affected the larval development, number of days to pupation and percentage normal pupation. Pandey et al. (1985) observed development period of *C. cephalonica* to range from 45.33-48.08 days in control as compared to 46-65.08 days when treated with neem oil, powder of kernel, cakes, leaves, flower and babul gum. Negi (2007) observed the time taken by *C. chinensis* to develop and emerge out as adults to go up to 40 days as compared to 26–28 days in normal and control sets when treated with *P. juliflora* bark + *P. cineraria* fruit. Ghei (2001) also observed the treatments of plants *Trigonella* to significantly increase the time taken for the development by the bruchid. Certain Solanaceous plants were employed by Gupta (2004), who also reached to a similar conclusion when extracts of *S. surattense* were used. Modgil & Essential oils of four species of *Mentha* were found to be effective against stored product insects by Tripathi et al. (2001). All these reports support the present findings. Morallo-Rejaesus et al. (1990) observed retardation in development of *C. chinensis* when treated with fruit of *C. frutescens*; Juneja & Patel (1994) observed no population build up of *C. analis* up to four months of storage of green gram when treated with seed powder of custard apple, black pepper, peel of orange and leaves of mint; Gupta (2004) found fruit formulation of certain Solanaceous plants to increase the time taken for development; Ghei (2001) observed extracts of *Trigonella* to significantly increase rate of development of *C. chinensis*, all these reports corroborate the present study. Similar effects have also been evinced by Schumutterer & Rembold (1980) and Mathur et al. (1989), who noted seeds of *A. indica* to act as growth disruptors.

Different formulations employed during the present study were found to affect the rate of development. Over all, besides crude extract, aqueous suspension also effectively enhanced the time taken by the beetle to develop and emerge as adult. These findings are in close proximity with those of Mann (1997), who observed powder suspension of stem, root and fruit and ether extract of stem of *P. harmala* to significantly lengthen the time taken for development by the bruchid; Ghei (2001), who found ether extracts of roots and leaves, aqueous extract of pods of certain leguminous plants to effect the rate of development of *C. chinensis*; Gupta (2004), who also found ether extract to be most effective in retarding the development of *C. chinensis*, while, Chiranjeevi & Sudhakar (1996) who suggested *A. indica* and *L. camara* to effect the development of pulse beetle only moderately.

During the present study, the rate of development was also found to be significantly affected by the extract concentration and, the time taken for development by the bruchid was particularly enhanced in sets treated with 25% extract. The present findings are in agreement with earlier reports of Mann (1997), Ghei (2001), Gupta (2004) and Negi (2007), who all observed an increase in the time taken by the pest insect to emerge as adult with an increase in dose concentration of different plant extracts. The efficacy of different concentration of commercial neem based insecticides 'Nimbecidine' was evaluated against *C. castaneum* by Das et al. (2006), who observed significant reduction in growth with lengthened developmental period. Khan & Borle (1985) documented clay and powdered rhizome of *A. calamus* at 0.1 and 0.5% concentrations to be most effective in arresting the development of *C. chinensis*.

Conclusion

It could therefore be concluded that, plants belonging to family Euphorbiaceae do possess chemicals, which are toxic to insects and therefore have a potential to be used against the pest *C. chinensis*. Although, the three select plants showed varied degree of effect on rate of development, *P. amarus* seemed to be the most effective.

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