

**STUDIES ON THE REGULATION OF  
PROGENY PRODUCTION AND SEX-RATIO  
OF *BRACON BREVICORNIS* WESMAEL**

(HYMENOPTERA : BRACONIDAE)

**By**

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**THESIS**

Submitted in partial fulfilment of the  
requirements for the degree of  
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Department of Agricultural Entomology  
COLLEGE OF HORTICULTURE  
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1979

## DECLARATION

I hereby declare that this thesis entitled "Studies on the regulation of progeny production and sex-ratio of Bracon brevicornis Wesmael (Hymenoptera: Braconidae)" is a bonafide record of research work done by me during the course of research work and the thesis had not previously formed the basis for the award to me of any degree, diploma, associateship, fellowship or other similar title of any other University or Society.

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**CERTIFICATE**

Certified that this thesis entitled  
"Studies on the regulation of progeny production  
and sex-ratio of Bracon brevicornis Wesmael  
(Hymenoptera : Braconidae)" is a record of research  
work done independently by Kumari. Susanna Jacob  
under my guidance and supervision and that it has  
not previously formed the basis for the award of  
any degree, fellowship or associateship to her.

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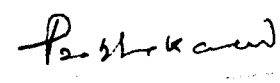
**CERTIFICATE**

We, the undersigned members of the Advisory Committee of Kumari Sossamma Jacob, a candidate for the degree of Master of Science in Agriculture with major in Agricultural Entomology agree that the thesis entitled "Studies on the regulation of progeny production and sex-ratio of Bracon brevicornis Wesmell (Hymenoptera : Braconidae)" may be submitted by Kumari Sossamma Jacob in partial fulfilment of the requirements for the degree.

  
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# INTRODUCTION

## INTRODUCTION

The black headed caterpillar Hephantia gerinopa Heyrick (Lepidoptera: Xyloryctidae) is a major pest of coconut in India occurring more commonly along the West and East-coast regions. Sporadic outbreaks of the pest occur during the dry months from January to May in the West-coast and from April to June in the East-coast. In Kerala, the pest is widely distributed throughout the coastal belt and in pockets around lakes and lagoons.

Along the West-coast, the population of H. gerinopa is found to be minimum during South-West monsoon period from June to August. There is a progressive increase in pest population after abatement of the South-West monsoon and the peak population levels are attained in the summer season. The black headed caterpillar, H. gerinopa supports a wide spectrum of natural enemies and the parasites are relatively more important among the key biotic agents. Periodic inoculative releases of suitable parasites after cessation of the South-West monsoon is considered to be adequate in maintaining the pest populations below the economic threshold.

Serious pest outbreaks are often experienced during the summer months in areas where parasite releases are not carried out on a regular basis. Severe pest outbreaks are to be brought under control by the application of a suitable contact toxicant. Frequent application of insecticides in coconut plantations will lead to pest resurgence due to progressive elimination of the parasites which form the major component of the natural enemy complex associated with the pest in Kerala. It is, therefore, essential that the insecticidal applications are to be followed by parasite releases with a view to compensate for the populations that are eliminated due to insecticidal application.

The important natural enemies of H. scirpota which are mass multiplied and released against the pest in Kerala are the parasites Bracon brevicornis Wesmahl and Trichospilus pupivora Ferriere. The pupal parasite T. pupivora lacks tolerance to high temperature conditions experienced in the summer months and these parasites are, therefore, not quite promising for inoculative or inundative releases during the hot season. B. brevicornis is relatively tolerant to high temperature conditions in the summer months, but a major handicap of this parasite is the production of progeny dominated by the males. The laboratory

stock of this parasite often dwindles substantially due to progressive dominance of males among the offsprings. This is a major obstacle to the adoption of inundative release techniques involving B. brevicornis. In order to build up and maintain large stocks of B. brevicornis in the insectary, a satisfactory level of progeny production consistent with suitable sex-ratio levels in the succeeding generations will have to be ensured.

Ayyar and Ananthanarayanan (1935) reported that B. brevicornis produces greater percentage of males in the hot seasons. Ramachandra Rao et al. (1943) had also recorded the dominance of males in the laboratory cultures of B. brevicornis and suggested that this might be due to the defective fertilization at pairing caused by adverse climate and nutritional factors. That short exposures to high temperature conditions brings about considerable variations in the sex-ratio of the offsprings of Dahlbominus fuliginosus (Nees) (Hymenoptera : Eulophidae) has been reported by Wilkes (1959 and 1963).

Mathew and Hair (1977) and Abraham and Mathew (1978) have indicated that the sex-ratio of the parent parasite populations and the host larval density are

relatively more important in the sex-ratio regulation of B. brevicornis. The importance of host size as a sex-ratio regulating mechanism in parasitic Hymenoptera has been indicated by a few workers (Clausen, 1939; Flanders, 1939; Ulyett, 1943 and 1945).

In order to explore the feasibility of manipulating the sex-ratio and progeny production in laboratory cultures of B. brevicornis, information on the precise influence of different levels of host larval density, weight of host larvae and sex-ratio of the parent parasite population at different temperature-humidity combination is considered to be vital. The present studies were carried out to meet the above desideratum.



# REVIEW OF LITERATURE

## REVIEW OF LITERATURE

Bracon brevicornis Wesmael is an important component of the natural enemy complex associated with the black headed caterpillar Hephantia serinopa Heyrick in Kerala.

Cherian (1929) worked out the biology of this parasite under Coimbatore conditions. The female wasps are yellowish-brown, with dark-brown dorsal bands on the abdomen and with fairly long ovipositor. Males are relatively smaller and duskier. The wasps mate on the day of emergence and egg laying commences the next day. Under confinement, the wasp freely oviposits on several species of noctuid and pyralid caterpillars. The female lays 8 to 12 eggs on the ventral surface of each larva, the mean fecundity being 200 per female. The incubation period is 28 to 36 hours and the larval and pupal periods are 3 to 4 days and 2 to 4 days respectively. The life-cycle is about 7 to 9 days.

B. brevicornis was first recorded in the South Malabar and areas around Cochin as an important parasite on Hephantia serinopa (Venkatasubban, 1932). This parasite was later introduced to North Malabar and

Coimbatore areas, but failed to keep down H. serinopa populations in these regions, perhaps due to the presence of alternative hosts (Ramachandra Rao et al. 1948). The alternate hosts of the parasite in India include Hoorda sorinrae Tams and Adisura atkinsoni Moore (Krishnamurthy, 1944). It also attacks the larvae of Datrachedra anhydrula Meyr., Sphostia castella W., Paria insulana Bois., Phycita infugella Meyr. and Sesania crotica Led., (Cherian and Margabandhu, 1942).

A major handicap of the parasite is the tendency to produce offsprings dominated by males. Quite often, the laboratory stock of the parasite progressively declines due to this handicap.

Work carried out on the factors influencing the regulation of the sex-ratio and progeny production in parasitic Hymenoptera in general and B. brevicornis in particular is briefly reviewed here.

## I GENETIC FACTORS

As in most species of parasitic Hymenoptera, Bracon brevicornis exhibits the phenomenon of arrhenotoky, characterised by the production of females from fertilised diploid eggs and of males from unfertilised haploid eggs.

*and Speicher*  
Speicher (1938) observed that the occasional uniparental females of Bracon hobeter Say originated from patches of tetraploid tissue in a normal diploid ovary.

Whiting (1945) stated that sex determination in Bracon is regulated by a series of multiple alleles of which nine have thus far been identified. The females conform to the  $x_a/x_b$ ,  $x_a/x_c$ ,  $x_c/x_d$  allelic genetic configuration while azygotes of the  $x_a$ ,  $x_b$ ,  $x_c$  types or any diploid homozygote of the types  $x_a/x_a$ ,  $x_b/x_b$  etc., will be males.

Genetic improvement of sex-ratios of some parasitic insects was obtained by Wilkes (1947) and Simmonds (1947). Wilkes (1947) showed that in the case of Dahlbominus fuliginosus (Nees) (Hymenoptera: Mulophidae), a continuous decrease in the number of sterile males from 35 to 2 per cent was brought about by selective breeding. It was shown that continued inbreeding in the laboratory stock was actually responsible for the prevalence of male sterility which resulted in the progressive reduction of female progeny in successive generations.

Simmonds (1947) found that in Mastrus carpedon (Lush), a higher percentage of females was obtained by breeding from males and females selected from high sex-ratio families and he suggested that the sex-ratio in this case is genetically determined.

The absence of fruitful fertilisation of eggs at pairing, possibly as a result of the defective climatic or nutritional factors has been implicated as a factor for male dominance in B. brevicornis (Rameshchandra Rao et al. 1948).

According to White (1954), arrhenotoky is a method of sex determination as well as a form of reproduction, the frequency of occurrence of males in the population being determined on the basis of the frequency of production of unfertilised eggs.

In the arrhenotokous parasitic wasp Dahlbominus fuliginosus, a sex-ratio controlling factor of genetic origin was reported by Wilkes (1964). This factor is sex limited and transmitted by females to their male progeny, its effect on sex-ratio being constant and not influenced by the female parent.

Lee and Wilkes (1965) detected five different types of spermatozoa in Dahlbominus fuscipennis Zett. (Hymenoptera: Eulophidae) on the basis of length and the pattern of spiralling of the helical coil in the head region of the spermatozoa. The spiralling may be of dextral or sinistral types and this dimorphism in the spermatozoa may determine whether fertilization occurs or not.

## II HOST AND HOST NUTRITION

According to Salt (1941), the host is often able to impress its mark upon the parasitoid that systematically destroys it and may bequeath to its parasite striking legacy of morphological, physiological and behaviouristic characters.

Karayanan and Chaudhuri (1955) reported that the sex-ratio of Stenobracon degees (Gan.) (Hymenoptera: Braconidae) varies according to the host and among the adults reared on Chilo zonellus, 28.8% were females, while among those reared on Corcyra larvae only 6.2% were females. It was suggested that the ovipositor enabled the parasite to recognise the host and the stimuli thus received influence the functioning of the spermatheca, which in turn controls fertilisation and sex of the offspring.

Harayanan and Subba Rao (1955) observed that when Bracon gelechiae Ashmead was reared on Platyedra roseovittella, Conocrysa cephalonia and Scelopophaga nivella, the proportion of females produced among the progeny were 55, 74 and 44 per cent respectively. The size of the host had little relation to the sex-ratio of the parasite and the higher incidence of males in some hosts was attributed in part to a lack of external stimulation of the spermatheca in the ovipositing female due to host unsuitability.

House and Jarlow (1961) studied the influence of different diets of a host, Agria affinis (Fall.) (Diptera: Sarcophaginae) on the development of a parasitoid, Aphaereta pallipes (Say.) (Hymenoptera: Braconidae). The diet containing of 3 per cent of amino acids and 0.5 per cent of dextrose caused greater host mortality and unsuccessful emergence of the parasites.

Louis (1961) recorded that when the females of Scambus buolianae (Htg.) (Hymenoptera: Ichneumonidae) were fed on host larvae and pollen in sucrose solution, greatest number of eggs per female was obtained, while females supplied with host larvae alone deposited only fewer eggs.

Katiyar (1962) studied the parasitisation by Trichogramma minutum (Hymenoptera: Trichogrammatidae) and Dragon elechiae of eggs and larvae of Coryvra cephalonica respectively. It was found that C. cephalonica reared on crushed sorghum mixed with 8 per cent yeast was more suitable for parasite development.

Doutt (1960), stated that the host of the parasitoid exerts considerable influence on the rate of development of the parasitoid in a remarkable manner.

Wylie (1966) indicated that females of Masonia vitripennis (Walk.) (Hymenoptera: Pteromalidae) parasitising housefly pupae (Musca domestica L.) gave rise to a smaller percentage of female progeny at high parasite - host ratios than at lower ones due to failure in fertilisation and because more female than male larvae die on superparasitised hosts. The adult females fertilise relatively fewer eggs at high parasite-host ratios since they encounter relatively more previously attacked hosts, on which a smaller percentage of fertilised eggs are laid. The reduction in the percentage of fertilised eggs under high parasite: host ratios is also due to mutual interference by the female parasites which inhibits fertilisation.



Mathai (1971) observed that the fecundity of B. brevicornis was significantly influenced by nutrition of its laboratory host Corycaea cephalonica. The host larvae bred on a media containing wheat flour fortified with skimmed milk powder (25% of the basic diet) or on wheat flower enriched with thiamine at 2 mg per gram was found to increase parasite fecundity. Variations in the nutrition of C. cephalonica did not significantly affect the duration of development of the parasite. A preponderance of males was found among the adult parasites reared on larvae bred on a media containing yeast, bengalgram or skimmed milk powder while the females were dominated in progeny reared out on larvae bred on maize flour diet.

Sangwan (1973) reported that when Stenobracon degees, a parasite of Chilo partellus Swinhoe, was reared in the laboratory on a single fictitious host Corycaea cephalonica, 34.7 per cent of the eggs developed into adults and 65.46 per cent of these were males, as compared with 87.3 per cent and 31.98 per cent respectively obtained from the natural host. However, when additional food in the form of further larvae of C. cephalonica that had been paralysed by B. brevicornis were provided, the survival ratio and the sex-ratio were restored. It was thus concluded that a single larva of C. cephalonica

did not provide sufficient nutrition for females of B. decens to complete their normal development.

Reporting the results of laboratory studies, David et al. (1977) stated that the parasite-host interaction between Bracon hebetor and Sphestia cautella was influenced by migration of host larvae prior to pupation and also by the feeding of the parasite females. Wandering host larvae were attacked more often than confine hosts. Survival and fecundity of adult female parasites were directly related to the frequency of host finding. Host feeding prolonged adult female parasite's survival, but the adult male parasites did not feed on the hosts. The long reproductive life, short life-cycle and reliance on hosts for adult and offspring nutrition should render B. hebetor quite responsive to manipulation of host availability.

Nadarejan and Jayaraj (1977) observed that the sex-ratio of the parasite Tetrastichus israeli Mani and Kurien (Hymenoptera: Eulophidae) was highly influenced by the host species and superparasitism. In Hephaestis gerinora at the density level of 30 parasites, the number of males were slightly more than females. There were more males at the density of 30 parasites per host in Margaronia indica S. The sex-ratio was as high as 2.06

in Coccyx cephalonica even at a level of 10 parasites indicating male preponderance. The male preponderance could be attributed to the differential mortality of sexes during development.

### III

### HOST DENSITY

Flanders (1935) studied the influence of hosts on progeny production in Trichostema evanescens. According to him, with increasing host densities, the number of progeny per parasite increased until a maximum was reached. The number of progeny per parent becomes stable beyond a stage when the number of hosts within the sphere of action of the parasite exceeds the total reproductive potential of the parasite. Under laboratory conditions, progeny per female decreased with the increase in the number of parasites beyond a critical level. This is perhaps due to the phenomenon of superparasitism and the consequential intraspecific competition.

Ulyett (1951) reported that Brachon hebetor tended to deposit a fixed number of eggs during a period of 24 hours regardless of the number of hosts.

Dharmaraju (1952) observed that the maximum fecundity of Perisierola nepentidis Muesbeck (Hymenoptera: Bethyridae) was obtained with one larva of

Cycovra cephalonica, while in the case of Trichoanilus musivora Ferriere (Hymenoptera: Braconidae), the fecundity did not show any increase with increase in the host pupal number.

Pattarudrinh and Usman (1957) found that the decline in the field parasitisation of eggs of Chilo zonellus by Trichorhiza evanescens minutus Riley in the Mysore State was mainly due to low host density levels.

Doutt (1959) concluded from observations on a species of Microbracon that the number of eggs per female per day was dependent on the number of hosts available up to the fourth day. Thereafter, host density was not observed as influencing egg production rate.

According to Flanders (1962), the proportion of host individual which stimulate the spermathecal gland increases as the density of host population decreases.

King (1962) recorded that at low host densities, the females of Nasonia vitripennis produced a smaller percentage of female progeny than at higher levels.

Jayar<sup>a</sup>tnam (1963) established a highly significant positive linear correlation between the weight of host pupa and the progeny production in Trichospilus pupivora.

The progeny production in Bracon brevicornis according to Seshagiri Rao et al. (1967), increased progressively with the number of hosts available but the progeny production decreased in Perisierola nephantidis at high host density levels. Host density had no influence on the sex-ratios of both the parasites.

Viktorov and Azizov (1972) reported that the number of progeny produced in Muscidifurax raptor (Hymenoptera: Pteromalidae) increased with increase in number of host puparia of Musca domestica Vicina (Diptera: Muscidae) which were available.

Fusco and Hower (1974) indicated that reproduction of the arrhenotokous parasite Microctonus aethiops (Nees) (Hymenoptera: Braconidae) produced the maximum number of progeny (average 25.8 per female) when fifty hosts were made available to each mated female. When ~~fifty~~ <sup>one</sup> ~~host~~ <sup>hosts</sup> were exposed to ~~one~~ <sup>parasite,</sup> ~~the~~ <sup>the</sup> progeny production declined and the percentage mortality of the hosts increased substantially. The percentage of females

in the progeny was higher when hosts were supplied discontinuously than when these were offered continuously.

According to Subba Rao et al., (1974), maximum fecundity in Bracon hebetor was obtained when two Coryza larvae of six weeks old were exposed to a single pair of parasites.

Yeargon et al., (1976) tested the effect of density of Hyper postica (Gyllenhal) (Coleoptera : Curculionidae) on parasitism by Bathyplectes curculionis (Hymenoptera: Ichneumonidae). It was observed that host density had no significant effect on the proportion of hosts parasitised. Host mortality was not attributable to normal parasite development but this was positively correlated with parasite density, suggesting some form of injury inflicted by the parasite.

Lathoef et al., (1977) studied the effect of density on host-parasite interactions between Hyper postica and Bathyplectes anura (Thomson). There was no significant relationship between proportion of hosts parasitised and host density. The frequency distribution of the parasite's eggs approximated the poisson distribution in eighty three per cent of the cases. There was

a quadratic relationship between number of super parasitised hosts and parasite density. However, no significant correlation between proportions of hosts parasitised and host density was detected.

Chandurwar (1977) observed that parasitism by Oegilus parvus Turner (Hymenoptera: Braconidae) on the host Phthorimaea operculella (Lepidoptera: Gelechiidae) was higher when the number of hosts per case was increased. However, the increase in parasite density resulted in reduced searching efficiency per individual parasite.

Abraham and Mathew (1978) reported that among the factors regulating the progeny production in Bracon breviscornis, the number of exposed Corycia larvae was relatively more important. The number of host larvae exposed exerts significant influence on the sex-ratio of the offspring, the contribution of this factor being 20.7 per cent.

Cameron and Rodfern (1978) recorded that the densities of the Hymenopteran parasite Forgyus nigritarsis Dalman and Mesopolobus diffinis were determined mainly by the density of available host, Taxomyia taxi (Ichbald) (Diptera: Cecidomyiidae).

Haggai podoler et al., (1978) studied the ovipositional responses of Aphytis holoxanthus (Hymenoptera: Aphelinidae) to increasing density levels of its host, the Florida red scale Chrysomphalus aonidum (Homoptera: Diaspididae). As the host density increased, the number of eggs laid by the female parasite also increased, but at a retarded rate. This type of functional response resulted in reduced parasitism at higher host densities.

Chewyrew (1913) found that the great majority of individuals of Pimpla reared on large hosts were females, whereas those from small hosts were predominantly males.

Holdaway and Smith (1952), studying the sex-ratio of Alysia manducator (Panz.) (Hymenoptera: Braconidae) bred on hosts of different sizes indicated that the quantity of available food determined by the size of the host played an important role in determining the sex of the parasite.

The influence of host size on progeny production has been studied extensively in Trichogramma spp. Salt (1934) stated that the number of eggs laid by



Trichogramma in a single host egg increased with host size, being one in Sitotroga cerealella Olivier, two in Corcyra cephalonica, two to three in Diatraea saccharalis three to four in Chilo sacchariphagus and eight to ten in Panloa demoleus Gsp. (Metcalf, 1959; Breniere, 1965).

According to Clausen (1939) and Flanders (1939), in parasitic Hymenoptera, generally large hosts favour the production of female parasites while the smaller hosts tend to produce males.

Ulyett (1943, 1945) observed that the number of Bracon eggs laid on or near a given host is related to the size of the host and ultimately to its potential as food.

Fusco and Hower (1973) studied the effect of size, sex and age of Hyper postica on the productivity of Microctonus aethiops in the laboratory. According to him, host size had no significant effect on the mean number of progeny or mean percentage of females produced per parasitoid. However, host mortality was significantly higher among the smaller hosts. Host sex had no effect on host selection by M. aethiops nor was any differential mortality observed between male or female hosts.

Garjanova (1974) reported that the sex-ratio of Exenterus abruptorius (Hymenoptera: Ichneumonidae) was determined by the size of the host Neodiprion sertifer and differential infestation rates on its males and females. The parasite lays most of the fertilised eggs on large female larvae and unfertilised eggs on male larvae.

#### IV. SEX OF THE HOST

McGugan (1955) found that the males of Apochthis ontario (Cresson) emerged mainly from male host pupae and females from female pupae, apparently the result of selective oviposition.

Sithanantham and Subramaniam (1977) reported the influence of sex of the larvae of Chilo partellus on the sex-ratio of the associated parasite Stenobracon decessae Cam. The sex-ratio of the parasite was in favour of males in the male hosts while it shifted significantly in favour of females in the female hosts.

#### V AGE OF THE HOST

Branson (1937) stated that females of many hymenopterous species can distinguish between parasitised and unparasitised hosts and fertilise a smaller percentage

of eggs they lay on unsuitable hosts.

Wylie (1962) observed that the age of pupae of the housefly Musca domestica (L.) affect the longevity and fecundity of Nasonia vitripennis that fed on them mainly due to variations in the nutritional status of hosts of different ages. At 24.5°C, both longevity and fecundity of females were greatest when it fed on pupae of less than 48 hours age.

According to Wylie (1963), mortality of N. vitripennis was least on young housefly pupae but increased with increasing host age.

## VI PARASITE DENSITY AND SUPERPARASITISM

According to Narayanan and Subba Rao (1955) superparasitism affected the sex-ratio of the progenies of Dracon gelechiae through selective elimination of females due to intra-specific competition. As the parasite larvae increased from three to thirty, the proportion of females declined drastically from 73.6 to 9.8%.

Narayanan and Mukerji (1956) studied the phenomenon of superparasitism in Trichogramma evanescens minutum and have observed that superparasitism occur at both high and

low host densities.

Harayanan and Chacko (1957) stated that in Trichogramma spp. the occurrence of superparasitism in laboratory breeding led to the production of progeny with impaired vitality, fecundity and longevity.

Iyotomi (1958) concluded that in the case of T. japonicum, superparasitism caused decrease of the percentage of female progeny and fecundity besides the impairment of vitality.

Chacko (1964) demonstrated that in Bracon gelechiae, superparasitism led to a lower proportion of females. He concluded that under crowded conditions, many members of both sexes failed to obtain sufficient food and that the females were the more seriously affected.

Nylie (1965) reported that superparasitism created food shortage and thereby reduced survival and size of adults of Haemaphysalis vitripennis reared on pupae of Bucea domestica. Super parasitism also reduced the percentage of females in the adult progeny.

Reinert and King (1971) found that Bracon hebetor preferred female larvae of Glodia interpunctella (Hubner) (Lepidoptera: Pyralidae) for parasitization. The sex-ratio

of B. hebetor in the F<sub>1</sub> generation appeared to vary with host: parasite density ratio.

Benson (1973) observed that B. hebetor being gregarious in the larval stage, the developing eggs and larvae were subjected to intraspecific competition which leads to direct mortality, size variation and changes in sex-ratio. The size of the surviving progeny and the sex-ratio decreases as parasite density/host-Ephestia cautella increases. It is shown that this change is caused by differential mortality which acts more strongly against the female.

Carpenter (1943) observed that single fertilised females of Bracon mollitor yielded on an average of 7.5 adult offsprings with 20 host larvae maintained at 85°F for 72 hours. But the individual productivity was decreased when batches of 5, 10, 15, 30 and 60 larvae were kept under the same conditions. This decrease was partly due to competition among the parasites and because they had to spend more time in searching for unparasitised hosts.

Dharmaraju (1952) found that the maximum number of fecundity was obtained with one pair of Pezisoteria nephantidis and the same trend was also noticed in the case of Trichospilus pupivora.

Sharma (1956) concluded that the fecundity of B. brevicornis declined with increase in the density of parasite population.

Subba Rao et al. (1974) stated that maximum fecundity was recorded in B. hebetor when one pair of parasites were exposed to two host caterpillars.

#### VII SEX-RATIO OF THE PARENT PARASITE POPULATION

Mathew and Nair (1977) stated that the proportion of females in the progeny of B. brevicornis can be improved to a great extent by adjusting the sex-ratio of the parent parasite population.

Abraham and Mathew (1978) observed that the number of males in the parental population of B. brevicornis significantly influenced the progeny production. The number of females in the parental population exerted significant influence on the sex-ratio of the offspring.

#### VIII MATING AND FERTILISATION

Whiting (1921) observed that the unmated females of Microbracon hebetor produced occasional females.

Flanders (1946) reported that in Macrocentrus anovivorus (Hymenoptera: Braconidae) male offsprings

predominated when multiple mating occurred. According to him, the sex-ratios are determined extrinsically by the effect of the density of oviposition sites on differential mortality during development, pre-ovipositional period, excessive mating, the difference in oviposition response before and after mating, the rate of oviposition etc. Sex-ratios are determined intrinsically by the number of eggs deposited at one insertion of the ovipositor, the number of ovarian eggs ready for deposition and the differential in the polyembryonic tendency.

The fecundity of unmated females was reduced according to Moutia and Courtois (1952) in Trichosyrphus australicus.

Hogelscher and Vinson (1971) observed that offsprings from unmated females of Campoplex nardistinctus (Viereck) (Hymenoptera: Ichneumonidae) were all males and females might not be produced after copulation. The exposure of older females to males for fertilisation yielded offspring with a greater percentage of females.

## IX

### ADULT PARASITE NUTRITION

According to Dharmaraju (1952), ten per cent sugar solution was the best among the food materials tried for improving the fecundity of Perisierola nepentidis and Trichospilus pupivora.

Sharma (1956) indicated that in Bracon brevicornis maximum fecundity was obtained when the parasites were fed with sucrose solution.

Subba Rao et al. (1974) reported that sucrose ten per cent was found to be the best among the different diets tried to increase fecundity of B.hebator.

#### X LIGHT AND PHOTOPERIOD

Martin and Finney (1946) improved the sex-ratio of Macrocentrus anevlivorus by decreasing the light intensity in the mating room.

Ramaoandhra Rao et al. (1948) reported that when sets of pairing adults of B. brevicornis were placed in special cages and kept exposed to diffused light in the open, equal proportions of both sexes of progeny were obtained.

According to Hoelscher and Vinson (1971), a 12:12 (L:D) photoperiod produced offsprings with the greatest percentage of females in Campoletis perdinctus (Viereck) (Hymenoptera: Ichneumonidae).

#### XI TEMPERATURE-HUMIDITY CONDITIONS

Hase (1922) observed that oviposition in Habrobracon brevicornis was inhibited at temperatures below 15°C.



Payne (1931) reported that both the period of development and the duration of adult life of Microbracon hebetor were affected by temperature.

Payne subsequently (1933) studied the differential effect of environmental factors upon M. hebetor. It was shown that at high temperature, the parasite had a decided advantage over the host, but at lower temperatures there was no possibility of the parasite overtaking the host.

Maereks (1933) working on the influence of temperature and air humidity on the embryonic development of M. hebetor, observed that the duration of development depended chiefly on temperature and only slightly on humidity. He noted that under the combined influence of optimum temperature and humidity, all the larvae hatched out simultaneously while under other conditions some larvae hatched first and were then followed by the majority. The effect of low humidities was more than that of high levels and the effect of humidity was more marked at low temperatures than at high ones. High temperatures produced mortality more quickly than lower levels.

Shroad and Garman (1933) reported that temperature affects the sex-ratio of Trichogramma through adverse

effects on the viability of the sperms . They have noticed that an excess of males appeared in the progeny of Trichogramma kept for two weeks at 3° to 8°C.

Payne (1934) found that the proportion of males was higher under low temperature and this was attributed to the suppression of mating activities.

Hanna (1935) indicated that pupae of Euchalcidia garyborzi Hanna. (Hymenoptera: Chalcididae) held at 16°C produced some sterile males, the proportion increasing with the duration of exposure and the females similarly treated produced increasingly fewer eggs.

Ayyer and Ananthanarayanan (1935) reported that B. leucicornis produced greater percentage of males in the hot season.

Ijima (1940) recorded that both males and females of Mucosa achistaceana Snellen, which were held at 32°C during the pupal stage were all found to be sterile.

Ahmad and Ghulamullah (1941) studied the influence of temperature on Marion fabia (Stoll) and its parasite Bracon greeni Lefr. Eudgeon and Gough. At low temperatures, the parasite developed more rapidly than the host, while at high temperatures the host population increased at a faster rate.

Hourai (1946) found that when Nasonia vitripennis were held at temperatures from  $-1^{\circ}\text{C}$  to  $16^{\circ}\text{C}$  for increasing periods of time, there was a gradual decrease in the proportion of females in the progeny.

Georgiann (1949) observed that haploid males in Habrobracon juglandis Ashmead were more vulnerable to environmental adversities.

The sex of the parasite and consequently the sex-ratio in most Hymenoptera is largely a function of the environment (Flanders, 1962). According to him, the remarkable variability in sex-ratios in many species of parasitic Hymenoptera under natural conditions results from the fact that in the progeny of the mated female, the sex of any particular individual is a maternal effect, this being a response to environmental stimuli acting through the spermathecal gland. The control of sex is accomplished by the facultative response of the spermathecal gland to environmental stimuli associated with the process of oviposition. Certain environmental temperatures and humidities and certain species of host may inhibit the stimulation of the spermathecal gland without affecting the number of eggs deposited.

Shalla and Venkataraman (1963) observed an accelerative effect on the rate of egg development with the rise of temperature from 20°C to 40°C in Vinion degees (Gm.) (Hymenoptera: Braconidae), a parasite of Chilo senellus. At 20°C, the larval period was completed on an average in 9.42 days. The developmental period decreased significantly with the rise of temperature till a minimum of 4.03 days was attained at 34°C. At 20°C and 90% RH there was a complete arrest of development in the pre-pupal stage, while at the same temperature and at lower humidities, a smaller percentage pupated and emerged as adults.

Wilkes (1963) reported that in the arrhenotokous insect Rahlbomius fuliginosus, variation in the sex-ratio occurred when unfavourable environmental conditions prevailed before, during and immediately after mating. Temperatures above 27°C during post-embryonic development, sterilised otherwise functional males, but had much less effect on females. Mating was successful only within the normal range for optimum adult activity. Although females were inseminated by more than one male, the sex-ratio of their progeny was not affected. Larval mortality from superparasitism above sixtyfive larvae per host differentially favoured survival of males.

The percentage of females was significantly reduced when oviposition was interrupted or delayed by low temperature.

Pilon (1966) stated that temperature and relative humidity significantly influence the life span of both sexes in the adults of Heodioxion swaini Middleton. The influence of temperature varied with relative humidity in the case of males but not in the case of females.

Rotary and Gerling (1973) studied the influence of some external factors upon the sex-ratio of Bracon habitor and found that numerous females produced male progeny only, perhaps due to internal factors. Cannibalism and mortality due to superparasitism favour male survival. The percentage of females increased significantly when females were allowed to feed only on hosts inside their webbing and also when mating and oviposition were delayed.

Barfield et al. (1977) observed the influence of temperature on the development of immature stages of Bracon mellitor a braconid parasite of the cotton boll weevil Anthonomus grandis Boheman (Coleoptera: Curculionidae) at a series of constant and variable

temperatures. Mean developmental periods were in close agreement with predictions based on a developmental model. They have stated that the adult longevity decreased with increasing temperature.

Nadrajana and Ghanna Dasavanna (1978) studied the effect of different levels of temperature and humidity on the biology and the development of the parasites, Parasitocera nephenitidis and Tetrastichus israeli. They have observed that the incubation period for eggs of P. nephenitidis was not influenced by temperature and humidity variations, while the larval period was highly influenced. There was no effect due to humidities alone but they interacted with temperature in influencing the larval period. There was no effect due to interaction of temperature and humidity and also by humidity alone on the pupal period. The humidity had no influence on the male longevity either alone or with temperature. Male preponderance was noted as the temperatures increased, indicating that the sex-ratio is adversely affected at higher temperature.

# MATERIALS AND METHODS

## MATERIALS AND METHODS

### I. Maintenance of Bracon brevicornis Wesmael stock cultures

The nucleus culture of Bracon brevicornis Wesmael required for the present investigations were obtained from the Parasite Breeding Laboratory of the Kerala Department of Agriculture at Irinjalakuda, Trichur District. The parasites were reared on twenty five days old fourth instar larvae of Coryza cephalonica Stainton, maintained at  $28 \pm 1.5^{\circ}\text{C}$ . Stock cultures of the parasite were maintained in small specimen tubes of size 10 x 2.5 cm. The tubes were thoroughly washed and sterilised in hot air oven at  $50^{\circ}\text{C}$  for one hour and after cooling, were stored in sterilised desiccators. As and when required, the tubes were drawn out from the desiccators. The parasites with adequate number of females were introduced into the specimen tubes and six to ten Coryza larvae of required age were then introduced into these tubes with a camel hair brush. A mixture of equal volumes of honey and 10 per cent glucose solution in water was used for feeding the adult parasites. The mixture was provided on bits of coconut leaflet of size 6.0 cm x 0.5 cm in the form of fine droplets by means of a coconut midrib piece of about 10 cm long with bluntly



pointed ends. The leaflet bits with food droplets were introduced into the specimen tubes which were then closed with cotton plugs. Fresh honey-glucose solution was provided immediately on exhaustion of the solution previously supplied. About five to six days after confinement of the parasites and host larvae, the tubes were cleaned by removing the unparasitised Ceromya larvae and the dead parasites. The stock cultures were maintained at  $28 \pm 1^{\circ}\text{C}$  and 75 per cent relative humidity.

The adult parasites of the succeeding generation emerged in eight to ten days and the culture was continuously maintained to ensure availability of adequate populations for experimental purposes.

## II. Maintenance of stock culture of Ceromya cephalonica Staint.

Stock cultures of Ceromya cephalonica Staint. were established from eggs obtained from the Parasite Breeding Laboratory of M/S SID Parry & Co. LTD., at Nellikuppam, S. Arcot District of the Tamil Nadu State. Adequate cultures were maintained throughout the course of the studies in museum jars of size 13.0 cm x 12.5 cm x 5.0 cm using crushed "jowar" (Sorghum vulgare Pers.) as the food medium. About 0.2 c.c. of freshly laid Ceromya eggs containing about 3500 eggs were initially mixed with 25 gms of jowar meal

and the mixture was then transferred to cylindrical glass bottles of size 28 cm x 21 cm. The bottles were closed with muslin cloth and tightly held in position by rubber bands. Four to five days later the jowar meal containing freshly emerged Goreyva larvae were taken out of the bottles and this was mixed with 350 gms of crushed jowar fortified with 50 gms of split black gram (Phaseolus mungo Linn.). The 'jowar' meal and crushed material as also split blackgram were sterilised at 50°C for one hour in a hot air oven before offering these as food for Goreyva larvae. The high temperature sterilisation was carried out to destroy any latent infestations of Goreyva or of other storage pests. Further rearings were carried out in rectangular museum jars. The moths emerging from these cultures were collected daily and these were confined in oviposition cages. The oviposition cage consisted of a cylindrical glass tube of size 19 cm x 12 cm, both ends of which were open. The two open ends of the glass tube were closed with suitably sized poly-meshes which were very tightly held by rubber bands. The moths were released inside the cage and the eggs collected inside glass dishes of the same diameter held on the bottom of the cylindrical glass tube. The eggs deposited on the poly-meshes were collected by brushing with a painting brush of

size 2.5 cm. The glass dish and the cylindrical glass tube were held together by winding adhesive bandage around. The oviposition cage was kept protected from predatory spiders by enclosing it in an ant-proof wire net cage.

### III. Studies on parasite development under different conditions

The influence of host larval density, weight of the host larvae and sex-ratio of the parent parasite population on the number of eggs laid, duration of development, progeny production and female-male composition of the  $F_1$  progeny at different temperature-humidity combinations were studied inside BOD incubators. The parasites under all treatment combinations were fed with honey-glucose mixture and exposed to semi-dark conditions obtaining inside the incubators.

The fecundity, duration of development, progeny production and female-male composition of the  $F_1$  progeny were studied as follows:-

#### a) Fecundity

The fecundity was recorded by examining the host larvae on successive days after confinement, under Stereomicroscope and directly counting the eggs adhering to the

host outcicle.

**b) Duration of development**

The duration of development was found out by calculating the weighted means of the number of days taken by the parasites of both sexes emerging on successive days. For this, the parasites emerging on successive days were isolated, killed and the female-male composition in relation to the durations of development recorded.

**c) Progeny production**

The total number of males and females which emerged out from each replicate was recorded as the  $F_1$  progeny production under different treatment combinations.

**d) Female-male composition of the  $F_1$  progeny**

The female-male composition of the  $F_1$  progeny produced under each treatment combination was recorded by pooling the data on the parasites of both sexes emerging on successive days.

**IV. Experimental details**

**a) Treatment combinations**

The influence of host larval density (A), weight of host larvae (B) and sex-ratio of the parent parasite

population (G) on the number of eggs laid, duration of development, progeny production and female-male composition of the  $F_1$  progeny, were studied in the present experiments. The levels at which each of these three factor was studied are indicated below.

Factor 1 - A - Host larval density

Levels  $a_0 = 1$  host larva per female parasite  
 $a_1 = 2$  host larvae per female parasite  
 $a_2 = 3$  host larvae per female parasite

Factor 2 - B - Weight of host larvae

Levels  $b_0 =$  Larval weight 30 - 35 mg  
 $b_1 =$  Larval weight 8 - 10 mg  
 $b_2 =$  50 per cent of larvae of the  $b_0$  type +  
 50 per cent of larvae of the  $b_1$  type.

Factor 3 - C - Sex-ratio of parent parasite population

Levels  $c_0 = 1:1$  (Female : Male)  
 $c_1 = 2:1$  ( " )  
 $c_2 = 3:1$  ( " )  
 $c_3 = 1:2$  ( " )  
 $c_4 = 1:3$  ( " )

There were altogether forty five treatment combinations involving the three main factors and these were

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replicated thrice. The treatment combinations are furnished in Table M-1. The statistical design employed was Factorial C.R.D.

b) Maintenance of temperature-humidity levels

The experiments were conducted at the following temperature-humidity combinations which represented near optimum conditions i.e., 28°C and 75% RH and the two progressively adverse conditions viz., 30°C and 60% RH and 32°C and 50% RH.

The different levels of constant temperatures were maintained in electric B.O.D. incubators while constant levels of humidity were obtained in desiccators (30 cm diameter) by keeping calculated quantities of pure saturated potassium hydroxide pellets dissolved in distilled water according to Burton (1931).

c) Selection of Coryvra larvae of the required weight group

Considerable size variations were observed among Coryvra larvae of the same age group. The larvae were classified according to their weight by quickly weighing live individual larvae in physical balances. In order to ensure that the larvae do not wriggle out from the pan, they were taken on coarse emery paper cards of size 4 x 4 cm

Table H-1

Treatment combinations involving host larval density (A), host larval weight categories (B) and sex-ratio of the parent parasite population (C).

Sl. No.	Treatment codes	Host larval density(A)	Host larval weight categories (B)		Sex-ratio of parent parasite population(C)	
			30-35 mg	8-10 mg	Number of females	Number of males
1.	a <sub>0</sub> b <sub>0</sub> c <sub>0</sub>	6	6	-	6	6
2.	a <sub>0</sub> b <sub>0</sub> c <sub>1</sub>	6	6	-	8	4
3.	a <sub>0</sub> b <sub>0</sub> c <sub>2</sub>	6	6	-	9	3
4.	a <sub>0</sub> b <sub>0</sub> c <sub>3</sub>	6	6	-	4	8
5.	a <sub>0</sub> b <sub>0</sub> c <sub>4</sub>	6	6	-	3	9
6.	a <sub>0</sub> b <sub>1</sub> c <sub>0</sub>	6	-	6	6	6
7.	a <sub>0</sub> b <sub>1</sub> c <sub>1</sub>	6	-	6	8	4
8.	a <sub>0</sub> b <sub>1</sub> c <sub>2</sub>	6	-	6	9	3
9.	a <sub>0</sub> b <sub>1</sub> c <sub>3</sub>	6	-	6	4	8
10.	a <sub>0</sub> b <sub>1</sub> c <sub>4</sub>	6	-	6	3	9
11.	a <sub>0</sub> b <sub>2</sub> c <sub>0</sub>	6	3	3	6	6
12.	a <sub>0</sub> b <sub>2</sub> c <sub>1</sub>	6	3	3	8	4
13.	a <sub>0</sub> b <sub>2</sub> c <sub>2</sub>	6	3	3	9	3
14.	a <sub>0</sub> b <sub>2</sub> c <sub>3</sub>	6	3	3	4	8
15.	a <sub>0</sub> b <sub>2</sub> c <sub>4</sub>	6	3	3	3	9
16.	a <sub>1</sub> b <sub>0</sub> c <sub>0</sub>	12	12	-	6	6
17.	a <sub>1</sub> b <sub>0</sub> c <sub>1</sub>	12	12	-	8	4
18.	a <sub>1</sub> b <sub>0</sub> c <sub>2</sub>	12	12	-	9	3
19.	a <sub>1</sub> b <sub>0</sub> c <sub>3</sub>	12	12	-	4	8
20.	a <sub>1</sub> b <sub>0</sub> c <sub>4</sub>	12	12	-	3	9
22.	a <sub>1</sub> b <sub>1</sub> c <sub>1</sub>	12	-	12	6	6
23.	a <sub>1</sub> b <sub>1</sub> c <sub>2</sub>	12	-	12	9	3
24.	a <sub>1</sub> b <sub>1</sub> c <sub>3</sub>	12	-	12	4	8
25.	a <sub>1</sub> b <sub>1</sub> c <sub>4</sub>	12	-	12	3	9
26.	a <sub>1</sub> b <sub>2</sub> c <sub>0</sub>	12	6	6	6	6
27.	a <sub>1</sub> b <sub>2</sub> c <sub>1</sub>	12	6	6	8	4
28.	a <sub>1</sub> b <sub>2</sub> c <sub>2</sub>	12	6	6	9	3
29.	a <sub>1</sub> b <sub>2</sub> c <sub>3</sub>	12	6	6	4	8
30.	a <sub>1</sub> b <sub>2</sub> c <sub>4</sub>	12	6	6	3	9
31.	a <sub>2</sub> b <sub>0</sub> c <sub>0</sub>	18	18	-	6	6
32.	a <sub>2</sub> b <sub>0</sub> c <sub>1</sub>	18	18	-	8	4
33.	a <sub>2</sub> b <sub>0</sub> c <sub>2</sub>	18	18	-	9	3
34.	a <sub>2</sub> b <sub>0</sub> c <sub>3</sub>	18	18	-	4	8
35.	a <sub>2</sub> b <sub>0</sub> c <sub>4</sub>	18	18	-	3	9
36.	a <sub>2</sub> b <sub>1</sub> c <sub>0</sub>	18	-	18	6	6
37.	a <sub>2</sub> b <sub>1</sub> c <sub>1</sub>	18	-	18	8	4
38.	a <sub>2</sub> b <sub>1</sub> c <sub>2</sub>	18	-	18	9	3
39.	a <sub>2</sub> b <sub>1</sub> c <sub>3</sub>	18	-	18	4	8
40.	a <sub>2</sub> b <sub>1</sub> c <sub>4</sub>	18	-	18	3	9
41.	a <sub>2</sub> b <sub>2</sub> c <sub>0</sub>	18	9	9	6	6
42.	a <sub>2</sub> b <sub>2</sub> c <sub>1</sub>	18	9	9	8	4
43.	a <sub>2</sub> b <sub>2</sub> c <sub>2</sub>	18	9	9	9	3
44.	a <sub>2</sub> b <sub>2</sub> c <sub>3</sub>	18	9	9	4	8
45.	a <sub>2</sub> b <sub>2</sub> c <sub>4</sub>	18	9	9	3	9

with uprightly folded edges. The larvae were then classified on the basis of their weights into the two categories viz., those with a weight range of 30 mg to 35 mg and those with a range of 8 mg to 10 mg.

d) Obtaining the required female-male composition of the parent parasite population

In order to obtain the required number of females and males for the different treatment combinations, freshly emerged parasites were drawn from the nucleus culture and counted immediately after anaesthetisation. For anaesthetisation, sterilised absorbent cotton pads were pressed on to the bottom of the specimen tubes of size 10 cm x 2.5 cm and then 2 to 3 drops of chloroform were poured on to the cotton pad. The specimen tube was kept tightly closed with cork and was opened just prior to anaesthetisation. The parasites were anaesthetised by inverting the tube containing chloroform-soaked cotton pad over the open tube containing parasites for a split second. The parasites which were thus immobilised were immediately transferred to glass plates and sexed. The parasites were observed till they recovered from anaesthetisation. Those which did not recover were removed and fresh parasites were introduced to compensate for the loss.



## V. Statistical methods

The analysis of the data on fecundity, progeny production and number of female progeny was carried out by using the chi-square method. The total variability in the data as measured by chi-square assuming equal distribution of counts, has been partitioned into the components reckoning the variation due to levels of the factors and their interactions. Thus the total  $\chi^2$  was computed by summing up  $\chi^2_A$ ,  $\chi^2_B$ ,  $\chi^2_{A \times B}$ ,  $\chi^2_C$ ,  $\chi^2_{A \times C}$ ,  $\chi^2_{B \times C}$  and  $\chi^2_{A \times B \times C}$ . The chi-squares for the factors were used to test the null hypothesis of equal distribution of counts over the levels. The chi-squares for interactions measure the heterogeneity among the levels of one factor. For example,  $\chi^2_{A \times B}$  measures the heterogeneity of distribution of counts according to the levels of A over the levels of B. Here the chi-square for the combination of A x B are partitioned into  $\chi^2_A + \chi^2_B + \chi^2_{A \times B}$ . Here the chi-square for a set of observations  $a_1, a_2, \dots, a_n$  was obtained by the formula  $\frac{\sum a_i^2 - n\bar{a}^2}{n}$  where  $\bar{a}$  is the arithmetic mean of the frequencies.

The analysis of variance technique which is applicable in the case of observations following the normal

law has not been used here, since the observations were in the form of counts. Even after square root transformation normality could not be achieved in such cases and hence chi-square test which is a distribution-free test has been adopted. However, in the case of duration of development the usual method of analysis of variance has been adopted since normality was evident.

The sex-wise counts of the parasites under different treatment combinations have been analysed for the following aspects employing the techniques outlined below.

(a) Testing the significance of the difference between the levels of the main factors in the expression of the female-male composition of the  $F_1$  offspring by transforming the ratio  $\frac{\text{Number of females}}{\text{Total}}$  as angles and by testing the differences between these angles in pairs by applying the formula  $\frac{\theta_1 - \theta_2}{\sqrt{320.7 \left( \frac{1}{n_1} + \frac{1}{n_2} \right)}}$  where  $\frac{320.7}{n_i}$  is the variance of  $\theta_i$ .

(b) Testing the association between the main factors and sex-wise classification of the  $F_1$  progeny, the chi-square values were computed on the basis of the formula  $\frac{(an_2 - a'n_1)^2}{(a+a')(n_1+n_2)}$  where  $a$  and  $a'$  denote the sex-wise frequencies corresponding to the levels and  $n_1$  and  $n_2$  represent the corresponding total counts in each case.

# RESULTS

## RESULTS

The fecundity, progeny production and the female-male composition of the  $F_1$  progeny of Bracon brevicornis Wesm. as influenced by number and weight of host larvae of Coryra cephalonica Stainton and sex-ratio of the parent populations of B. brevicornis were studied at three temperature-humidity combinations which represented near optimum conditions of 28°C and 75% RH and near adverse conditions of 30°C and 60% RH and 32°C and 50% RH. The effects of the three factors on the duration of development were also studied in the present experiments.

The following were the levels at which the effect of the three factors were studied.

### Factor 1:- Host larval density (A)

Levels -  $a_0$  = 1 host larva per female parasite  
 $a_1$  = 2 host larvae per female parasite  
 $a_2$  = 3 host larvae per female parasite

### Factor 2:- Weight of the host larvae (B)

Levels -  $b_0$  = larval weight: 30-35 mg  
 $b_1$  = larval weight: 8-10 mg  
 $b_2$  = half of the hosts of the ' $b_0$ ' category and the other half of the ' $b_1$ ' type.

**Factor 3:- Sex-ratio of parent parasite population (S).**

Levels -	$c_0 = 1:1$ (Female : Male)
	$c_1 = 2:1$ ( " )
	$c_2 = 3:1$ ( " )
	$c_3 = 1:2$ ( " )
	$c_4 = 1:3$ ( " )

There were altogether forty five treatment combinations and these were replicated thrice as outlined in materials and methods.

**FECONDITY****i. Fecundity of B. brevicornis at 18°C and 75% RH**

The fecundity was recorded by examining the host larvae on successive days after confinement under Stereomicroscope and directly counting the eggs adhering to the host cuticle. The data on the mean number of eggs laid by the parasite females in the different treatment combinations at the above temperature-humidity level are furnished in column 3 of Table 1-a and the raw data in Appendix-I. The fecundity was too low (6.57) Under the treatment combination involving one host larva weighing 3-10 mg per female parasite and a composition of one female: one male in the parent population ( $a_0$   $b_1$   $c_0$ ).

Fecundity of *B. bassioicornis* in different temperature and humidity combinations

Sl. No.	Treatments	Mean number of eggs laid		
		20°C and 75% RH	30°C and 60% RH	32°C and 50% RH
1.	a <sub>0</sub> b <sub>0</sub> e <sub>0</sub> *	60.67	29.33	61.33
2.	a <sub>0</sub> b <sub>0</sub> e <sub>1</sub>	45.67	50.67	110.00
3.	a <sub>0</sub> b <sub>0</sub> e <sub>2</sub>	32.00	47.00	122.33
4.	a <sub>0</sub> b <sub>0</sub> e <sub>3</sub>	55.00	81.00	49.67
5.	a <sub>0</sub> b <sub>0</sub> e <sub>4</sub>	12.67	29.00	58.67
6.	a <sub>0</sub> b <sub>1</sub> e <sub>0</sub>	6.67	34.33	27.33
7.	a <sub>0</sub> b <sub>1</sub> e <sub>1</sub>	22.67	58.33	24.67
8.	a <sub>0</sub> b <sub>1</sub> e <sub>2</sub>	78.33	63.33	20.67
9.	a <sub>0</sub> b <sub>1</sub> e <sub>3</sub>	52.33	14.00	25.33
10.	a <sub>0</sub> b <sub>1</sub> e <sub>4</sub>	53.00	39.33	16.33
11.	a <sub>0</sub> b <sub>2</sub> e <sub>0</sub>	57.67	50.00	78.00
12.	a <sub>0</sub> b <sub>2</sub> e <sub>1</sub>	54.00	73.33	80.00
13.	a <sub>0</sub> b <sub>2</sub> e <sub>2</sub>	93.33	71.67	61.00
14.	a <sub>0</sub> b <sub>2</sub> e <sub>3</sub>	64.00	46.33	35.00
15.	a <sub>0</sub> b <sub>2</sub> e <sub>4</sub>	82.00	25.33	43.67
16.	a <sub>1</sub> b <sub>0</sub> e <sub>0</sub>	88.33	38.33	67.33
17.	a <sub>1</sub> b <sub>0</sub> e <sub>1</sub>	66.33	80.33	118.00
18.	a <sub>1</sub> b <sub>0</sub> e <sub>2</sub>	96.00	54.33	119.33
19.	a <sub>1</sub> b <sub>0</sub> e <sub>3</sub>	45.00	71.00	87.67
20.	a <sub>1</sub> b <sub>0</sub> e <sub>4</sub>	51.33	22.67	75.33
21.	a <sub>1</sub> b <sub>1</sub> e <sub>0</sub>	69.33	35.00	27.33
22.	a <sub>1</sub> b <sub>1</sub> e <sub>1</sub>	88.33	61.67	33.33
23.	a <sub>1</sub> b <sub>1</sub> e <sub>2</sub>	39.67	43.33	38.00
24.	a <sub>1</sub> b <sub>1</sub> e <sub>3</sub>	45.00	30.00	46.67
25.	a <sub>1</sub> b <sub>1</sub> e <sub>4</sub>	36.67	39.33	36.67
26.	a <sub>1</sub> b <sub>2</sub> e <sub>0</sub>	68.33	115.00	75.67

27.	a <sub>1</sub> b <sub>2</sub> e <sub>1</sub>	127.33	108.33	108.33
28.	a <sub>1</sub> b <sub>2</sub> e <sub>2</sub>	107.33	98.33	54.33
29.	a <sub>1</sub> b <sub>2</sub> e <sub>3</sub>	28.00	55.00	34.00
30.	a <sub>1</sub> b <sub>2</sub> e <sub>4</sub>	43.00	30.00	105.00
31.	a <sub>2</sub> b <sub>0</sub> e <sub>0</sub>	33.00	91.00	59.67
32.	a <sub>2</sub> b <sub>0</sub> e <sub>1</sub>	54.33	85.67	86.67
33.	a <sub>2</sub> b <sub>0</sub> e <sub>2</sub>	66.00	56.67	112.67
34.	a <sub>2</sub> b <sub>0</sub> e <sub>3</sub>	25.33	21.00	88.67
35.	a <sub>2</sub> b <sub>0</sub> e <sub>4</sub>	33.00	44.00	35.67
36.	a <sub>2</sub> b <sub>1</sub> e <sub>0</sub>	20.67	62.00	30.67
37.	a <sub>2</sub> b <sub>1</sub> e <sub>1</sub>	77.67	36.67	30.33
38.	a <sub>2</sub> b <sub>1</sub> e <sub>2</sub>	45.33	53.67	53.00
39.	a <sub>2</sub> b <sub>1</sub> e <sub>3</sub>	27.67	50.00	17.00
40.	a <sub>2</sub> b <sub>1</sub> e <sub>4</sub>	32.67	15.33	33.00
41.	a <sub>2</sub> b <sub>2</sub> e <sub>0</sub>	64.00	23.00	70.67
42.	a <sub>2</sub> b <sub>2</sub> e <sub>1</sub>	154.00	37.00	69.00
43.	a <sub>2</sub> b <sub>2</sub> e <sub>2</sub>	33.00	36.67	90.33
44.	a <sub>2</sub> b <sub>2</sub> e <sub>3</sub>	45.00	38.33	68.67
45.	a <sub>2</sub> b <sub>2</sub> e <sub>4</sub>	31.00	32.33	31.00

\*a<sub>0</sub> = 1 host larva per female parasite  
a<sub>1</sub> = 2 " " " "  
a<sub>2</sub> = 3 " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
b<sub>1</sub> - " " " 8 - 10 mg  
b<sub>2</sub> - half of the larvae of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

e<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)  
e<sub>1</sub> - " " " " 2:1  
e<sub>2</sub> - " " " " 3:1  
e<sub>3</sub> - " " " " 1:2  
e<sub>4</sub> - " " " " 1:3

Table 1-b

Chi-square values for the effect of the main factors A, B and C on fecundity of *B. brevicornis* at different temperature and humidity combinations

Source	df	Chi-square values		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
* A	2	90.94**	65.41**	81.74**
° B	2	185.94**	77.25**	1009.03**
A x B	4	90.76**	220.97**	40.04**
° C	4	343.35**	337.78**	332.33**
A x C	8	463.72**	79.98**	138.02**
B x C	8	132.29**	84.77**	275.66**
A x B x C	16	554.23**	531.45**	185.67**

\*\* Significant at 1% level

\*A - host density

°B - weight of host larvae

°C - sex-ratio of parent parasite population

Table 1-c

Number of eggs laid by *B. brevicaornis* at different levels of the main factors at different temperature and humidity combinations

	28°C and 75% RH		30°C and 60% RH		32°C and 50% RH	
	Total	Mean	Total	Mean	Total	Mean
a <sub>0</sub> *	2310	51.33	2139	47.53	2442	54.27
a <sub>1</sub>	2844	63.20	2558	56.84	3068	68.18
a <sub>2</sub>	2228	49.50	2050	45.56	2565	57.00
b <sub>0</sub> *	2294	50.98	2406	53.47	3659	81.31
b <sub>1</sub>	2038	46.40	1909	42.42	1402	31.61
b <sub>2</sub>	3000	66.67	2432	54.04	3004	66.76
e <sub>0</sub> *	1406	31.24	1434	31.87	1494	33.20
e <sub>1</sub>	1915	42.56	1686	37.47	1995	44.33
e <sub>2</sub>	1773	39.40	1575	35.00	2015	44.78
e <sub>3</sub>	1162	25.82	1220	27.11	1268	28.18
e <sub>4</sub>	1126	25.02	832	18.49	1303	28.96

a<sub>0</sub>\* - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

b<sub>0</sub>\* - larval weight 30 - 35 mg.  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and other half of the 'b<sub>1</sub>' type.

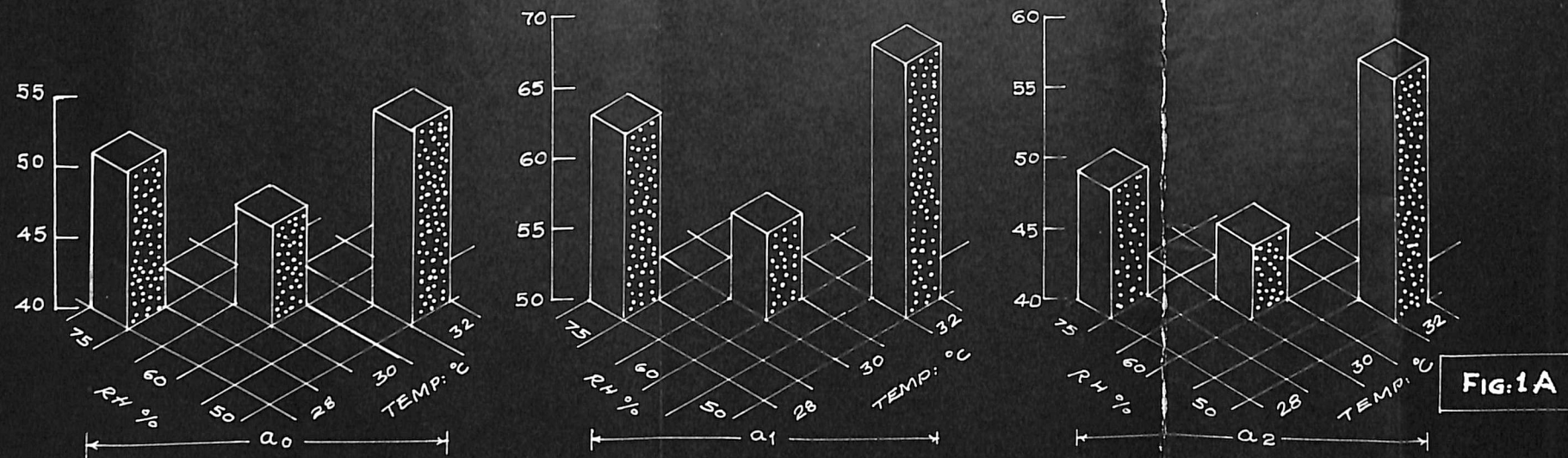
e<sub>0</sub>\* - Sex ratio of parent parasite population - 1:1 (Female: Male)  
 e<sub>1</sub> - " " " " - 2:1 ( " " )  
 e<sub>2</sub> - " " " " - 3:1 ( " " )  
 e<sub>3</sub> - " " " " - 1:2 ( " " )  
 e<sub>4</sub> - " " " " - 1:3 ( " " )



**Fig. 1 A - Fecundity of Bracon brevicornis at different temperature-humidity combinations as influenced by the host larval density**

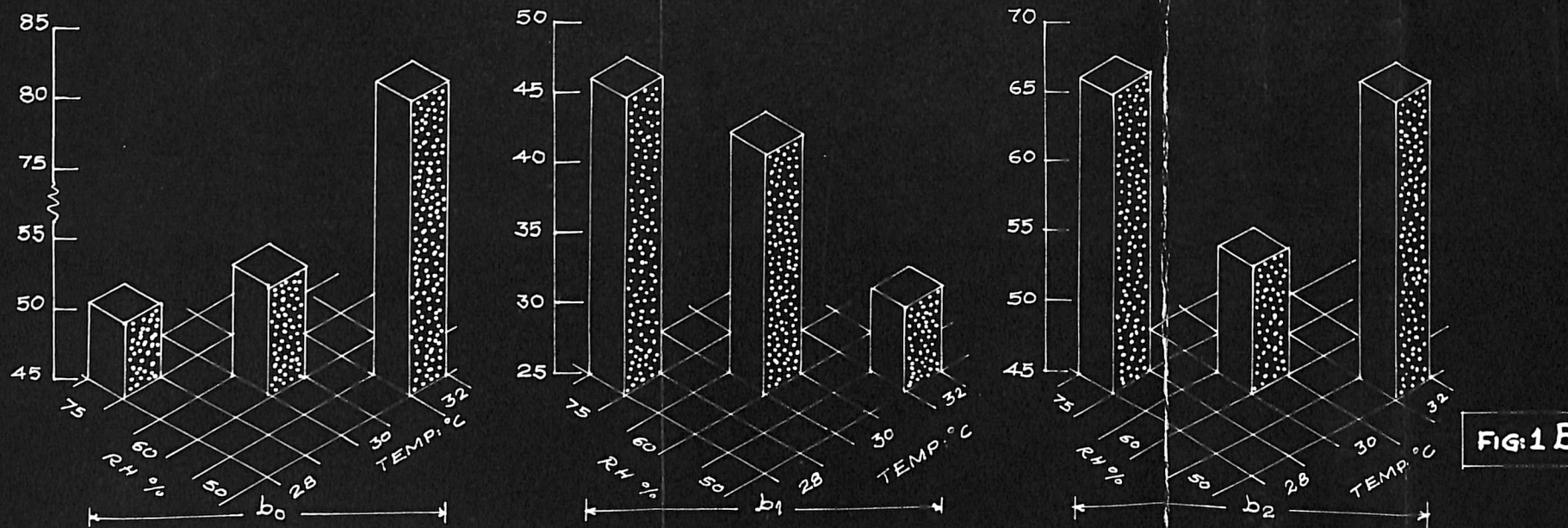
**Fig. 1 B - Fecundity of Bracon brevicornis at different temperature-humidity combinations as influenced by the weight (size) of host larvae**

**Fig. 1 C - Fecundity of Bracon brevicornis at different temperature-humidity combinations as influenced by the sex-ratio of the parent parasite population**



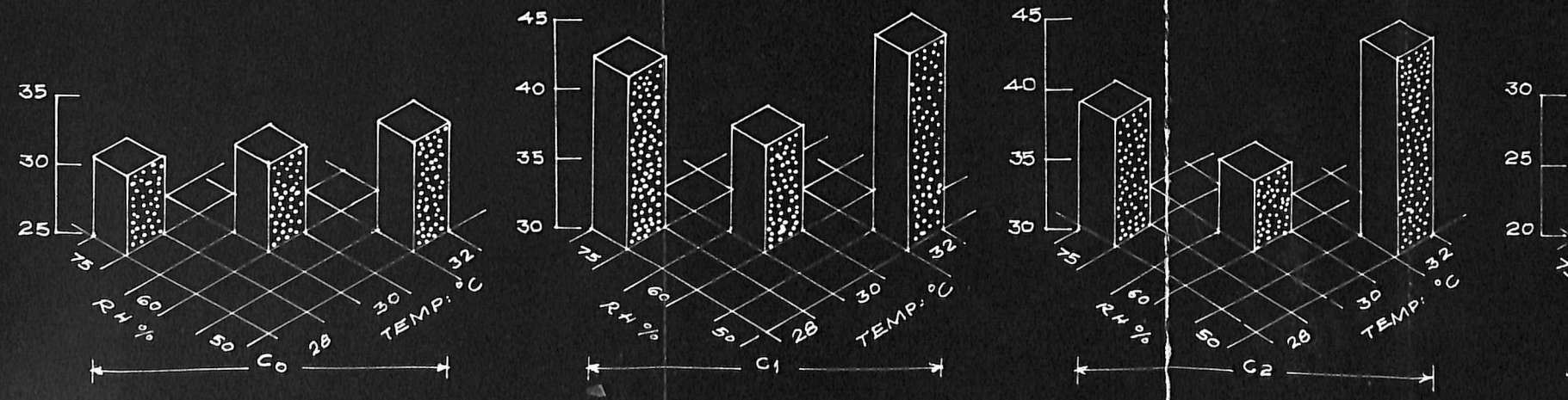
A

a <sub>0</sub>	1 HOST LARVA PER FEMALE PARASITE
a <sub>1</sub>	2 " " " " " "
a <sub>2</sub>	3 " " " " " "



B

b <sub>0</sub>	LARVAL WEIGHT 30-35 mg.
b <sub>1</sub>	" " " " 8-10 "
b <sub>2</sub>	HALF OF THE LARVAE OF THE 'b <sub>0</sub> ' TYPE AND THE OTHER HALF OF THE 'b <sub>1</sub> ' CATEGORY



C

c <sub>0</sub>	1 FEMALE : 1 MALE
c <sub>1</sub>	2 " " 1 " "
c <sub>2</sub>	3 " " 1 " "
c <sub>3</sub>	1 " " 2 " "
c <sub>4</sub>	1 " " 3 " "

Fig. 1 B

Fig. 1

Table 1-d

Chi-square values for comparing the effects of different levels of the main factors A, B and C at different temperature and humidity combinations on fecundity of B. brevicornis

		Chi-square values					
		20°C and 75% RH	Rank- ing	30°C and 60% RH	Rank- ing	32°C and 50% RH	Rank- ing
a <sub>0</sub>	Vs a <sub>1</sub>	55.33**	a <sub>1</sub> a <sub>2</sub>	37.38**	a <sub>1</sub> a <sub>2</sub>	71.12**	a <sub>1</sub> a <sub>2</sub>
a <sub>1</sub>	Vs a <sub>2</sub>	74.81**	a <sub>1</sub> a <sub>2</sub>	55.64**	a <sub>1</sub> a <sub>2</sub>	44.92**	a <sub>1</sub> a <sub>2</sub>
a <sub>0</sub>	Vs a <sub>2</sub>	1.48 NS	a <sub>1</sub> a <sub>2</sub>	1.89 NS	a <sub>1</sub> a <sub>2</sub>	3.02 NS	a <sub>1</sub> a <sub>2</sub>
b <sub>0</sub>	Vs b <sub>1</sub>	9.68**	b <sub>0</sub> b <sub>1</sub>	57.24**	b <sub>0</sub> b <sub>1</sub>	1006.53**	b <sub>0</sub> b <sub>1</sub>
b <sub>1</sub>	Vs b <sub>2</sub>	163.47**	b <sub>0</sub> b <sub>1</sub>	63.01**	b <sub>0</sub> b <sub>1</sub>	582.48**	b <sub>0</sub> b <sub>1</sub>
b <sub>0</sub>	Vs b <sub>2</sub>	94.15**	b <sub>0</sub> b <sub>1</sub>	0.14 NS	b <sub>0</sub> b <sub>1</sub>	64.39**	b <sub>0</sub> b <sub>1</sub>
c <sub>0</sub>	Vs c <sub>1</sub>	78.01**	c <sub>0</sub> c <sub>1</sub>	20.35**	c <sub>0</sub> c <sub>1</sub>	100.84**	c <sub>0</sub> c <sub>1</sub>
c <sub>0</sub>	Vs c <sub>2</sub>	42.37**	c <sub>0</sub> c <sub>1</sub>	6.61*	c <sub>0</sub> c <sub>1</sub>	77.36**	c <sub>0</sub> c <sub>1</sub>
c <sub>0</sub>	Vs c <sub>3</sub>	23.18**	c <sub>0</sub> c <sub>1</sub>	17.26**	c <sub>0</sub> c <sub>1</sub>	18.49**	c <sub>0</sub> c <sub>1</sub>
c <sub>0</sub>	Vs c <sub>4</sub>	30.96**	c <sub>0</sub> c <sub>1</sub>	159.93**	c <sub>0</sub> c <sub>1</sub>	13.04**	c <sub>0</sub> c <sub>1</sub>
c <sub>1</sub>	Vs c <sub>2</sub>	5.47*	c <sub>0</sub> c <sub>1</sub>	3.78 NS	c <sub>0</sub> c <sub>1</sub>	0.10 NS	c <sub>0</sub> c <sub>1</sub>
c <sub>1</sub>	Vs c <sub>3</sub>	184.27**	c <sub>0</sub> c <sub>1</sub>	74.73**	c <sub>0</sub> c <sub>1</sub>	161.98**	c <sub>0</sub> c <sub>1</sub>
c <sub>1</sub>	Vs c <sub>4</sub>	204.71**	c <sub>0</sub> c <sub>1</sub>	289.64**	c <sub>0</sub> c <sub>1</sub>	145.20**	c <sub>0</sub> c <sub>1</sub>
c <sub>2</sub>	Vs c <sub>3</sub>	127.20**	c <sub>0</sub> c <sub>1</sub>	45.09**	c <sub>0</sub> c <sub>1</sub>	169.97**	c <sub>0</sub> c <sub>1</sub>
c <sub>2</sub>	Vs c <sub>4</sub>	144.40**	c <sub>0</sub> c <sub>1</sub>	229.35**	c <sub>0</sub> c <sub>1</sub>	152.79**	c <sub>0</sub> c <sub>1</sub>
c <sub>3</sub>	Vs c <sub>4</sub>	0.57 NS	c <sub>0</sub> c <sub>1</sub>	73.36**	c <sub>0</sub> c <sub>1</sub>	0.48 NS	c <sub>0</sub> c <sub>1</sub>

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and the other half of the 'b<sub>1</sub>' type

c<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>1</sub> - " " " " 2:1  
 c<sub>2</sub> - " " " " 3:1  
 c<sub>3</sub> - " " " " 1:2  
 c<sub>4</sub> - " " " " 1:3

Table 1-c

Chi-square values for comparison of effects of the three main factors A, B and C on fecundity of *B. brevicornis* at different temperature and humidity combinations

			Chi-square for levels of B <sup>o</sup>			Chi-square for levels of C <sup>o</sup>				
			b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>	c <sub>0</sub>	c <sub>1</sub>	c <sub>2</sub>	c <sub>3</sub>	c <sub>4</sub>
TH <sub>1</sub>	Vs	TH <sub>2</sub>	2.67 NS	8.01**	59.39**	0.28 NS	14.56**	11.71**	1.41 NS	44.15**
TH <sub>2</sub>	Vs	TH <sub>3</sub>	257.63**	27.63**	60.19**	1.23 NS	25.94**	53.93**	0.93 NS	103.91**
TH <sub>1</sub>	Vs	TH <sub>3</sub>	312.99**	134.84**	0.002 NS	2.67 NS	1.64 NS	15.46**	4.82*	12.90**
Ranking			TH <sub>3</sub> TH <sub>2</sub> TH <sub>1</sub>	TH <sub>1</sub> TH <sub>2</sub> TH <sub>3</sub>	TH <sub>3</sub> TH <sub>1</sub> TH <sub>2</sub>	TH <sub>3</sub> TH <sub>2</sub> TH <sub>1</sub>	TH <sub>3</sub> TH <sub>1</sub> TH <sub>2</sub>	TH <sub>3</sub> TH <sub>1</sub> TH <sub>2</sub>	TH <sub>3</sub> TH <sub>2</sub> TH <sub>1</sub>	TH <sub>3</sub> TH <sub>1</sub> TH <sub>2</sub>

Chi-square due to heterogeneity

Host larval density (A)	6.51 NS
Weight of host larvae (B)	461.60**
Sex-ratio of parent parasite population (C)	64.66**

<sup>o</sup>B - Weight of host larvae  
 b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>1</sub>' type  
 and the other half of the 'b<sub>0</sub>' type

<sup>o</sup>C - Sex-ratio of parent population  
 c<sub>0</sub> - 1 female : 1 male  
 c<sub>1</sub> - 2 " : 1 "  
 c<sub>2</sub> - 3 " : 1 "  
 c<sub>3</sub> - 1 " : 2 "  
 c<sub>4</sub> - 1 " : 3 "

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

TH<sub>1</sub> - 29°C and 75% RH  
 TH<sub>2</sub> - 30°C and 60% RH  
 TH<sub>3</sub> - 32°C and 50% RH

while under three host larvae per female parasite which is a mixture of heavier and lighter types and a ratio of two females : one male in the parent population ( $a_1, b_2, c_1$ ), the egg production was 154.00.

The data being in the form of counts of eggs, the analysis of variance table was not found applicable even after square root transformation since normality could not be achieved even by transformation. Therefore, chi-square test which is a distribution free test, has been adopted as outlined in the materials and methods and the results of testing of the three main factors A, B and C and their interactions are presented in Table 1-b. It will be seen that all the main effects, viz., host larval density (A), weight of the host larvae (B), sex-ratio of parent parasite population (C) as well as the interactions AxB, AxC, BxC and AxBxC are highly significant at 1 per cent level indicating the importance of these factors operating individually and jointly in the realisation of oviposition potential of the female parasites.

The data pertaining to the total number of eggs laid at different levels of the three main factors host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population (C) are summarised and

presented in Table 1-c and these are graphically depicted in Fig. 1A, 1B and 1C. A maximum number of 63.20 eggs were obtained when two host larvae were offered per female parasite ( $a_1$  level). The chi-square values for the comparison between the different levels of the factors are given in Table 1-d. At the  $a_1$  level (two hosts per female parasite) of host density, which gave maximum number of eggs, the fecundity was significantly higher than those recorded at the  $a_0$  level of one host larva per female parasite and at the  $a_2$  level of three host larvae per female parasite, the mean fecundity values for the two latter cases being 51.33 and 49.50 respectively. Significant differences were not detected between the  $a_0$  and  $a_2$  levels.

Among the levels of the weight of host larvae (B), (Table 1-c) the maximum fecundity of 66.67 was registered when the host larvae comprised of heavier and lighter ones in equal proportions ( $b_2$ ). This was followed by the  $b_0$  level (host larvae with a weight of 30-35 mg) with mean 50.98 and the  $b_1$  level (host larvae with a weight of 8-10 mg) with mean 46.40. All the three levels of the weight of host larvae (B) were significantly different. It will be seen that the weight of the host larvae considerably influences the egg production in B. brevicornis

and that the maximum egg production is registered when relatively heavier and lighter larvae of the same age were offered for parasitisation.

Among the different levels of the sex-ratios of the parent parasite population (C), the ratio of 2 females: 1 male ( $c_1$ ) gave the maximum number of eggs, mean being 42.56 (Table 1-d). Under the levels 3 females: 1 male ( $c_2$ ) and 1 female:1 male ( $c_0$ ), the mean fecundity were 39.40 and 31.74 respectively. When the sex-ratio of the parental population was 1 female: 2 males ( $c_3$ ) and 1 female: 3 males ( $c_4$ ), there was a drastic decline in the number of eggs laid, the mean values under these two levels being 25.82 and 25.02 respectively. Significant differences were not detected between the two levels  $c_3$  and  $c_4$ .

#### Fecundity of B. brevicornis at 30°C and 60% RH

The mean fecundity of the parasites at 30°C and 60% RH are furnished in column 4 of Table 1-a. The range of fecundity was quite wide being 14.00 under  $a_0$   $b_1$   $c_3$  treatment combination (one host larva per female parasite; larval weight 8-10 mg; sex-ratio of the parent parasite population 1:3). The fecundity was the maximum (115) for the treatment combination  $a_1$   $b_2$   $c_0$  (two host larvae per female parasite; half of the host larvae of 30-35 mg type

and the other half of 8-10 mg type; sex-ratio of the parent parasite population 1:1).

Analysis of the data was carried out by employing chi-square test and the results are furnished in Table 1-b. It is indicated that all the main effects (host larval density, weight of host larvae, sex-ratio of parent parasite population) as well as their interactions are highly significant. The data showing the mean fecundity at different levels of the three main factors are summarized in Table 1-c and these are illustrated in Fig. 1A, 1B and 1C. The maximum number of eggs (56.84) were recorded when two host larvae ( $a_1$  level) were offered per female parasite and this was followed by the fecundity registered with one host per female parasite ( $a_0$ ) and three hosts per female parasite ( $a_2$ ). The chi-square values to compare between levels of the three factors are presented in column 3 Table 1-d. It will be noted that the fecundity registered at two hosts larvae per female parasite ( $a_1$ ) is superior (56.84) to those at one host larva per female parasite ( $a_0$ ) and three host larvae per female parasite ( $a_2$ ). Significant differences were, however, not detected between one host larva per female parasite ( $a_0$ ) and three host larvae per female parasite ( $a_2$ ) levels.

As regards the weight of the host larvae of the



same age, it was found that the maximum number of eggs was recorded with a mixture of relatively heavier and lighter larvae in equal proportions ( $b_2$ ) the mean fecundity being 54.04. This was followed by the values corresponding to  $b_1$  (relatively lighter larvae) and  $b_0$  (relatively heavier larvae), the means being 53.47 and 42.42 respectively. No significant difference was evident between  $b_0$  (host larvae with 30-35 mg weight) and  $b_2$  levels (a mixture of heavier and lighter larvae in equal proportions) indicating that heavier host larvae and a mixture of heavier and lighter ones in equal proportions are equally effective for the better realization of oviposition potential of B. brevicornis. The host larval weight level  $b_1$  (8-10 mg) gave significantly lower fecundity.

The maximum fecundity of 37.47 at 30°C and 60% RH was registered when the sex-ratio of the parent population ( $c_1$ ) was maintained at 2:1 (female:male). At the  $c_2$  (3:1) and  $c_3$  (1:2) levels, mean fecundity levels were 35.00 and 27.11 respectively. The number of eggs laid were quite low under the levels  $c_3$  (1:2) and  $c_4$  (1:3) characterised by the dominance of the males, the mean values being respectively 27.11 and 18.49. The fecundity at the  $c_1$  and  $c_2$  levels were on par while  $c_0$ ,  $c_3$  and  $c_4$  gave

significantly different fecundity.

**Fecundity of B. brevicornis at 32°C and 50% RH**

The data on the fecundity of B. brevicornis at 32°C and 50% RH are presented in column 5 of Table 1-a. The number of eggs laid by the parasites was the lowest (16.33) under the treatment  $a_0 b_1 c_4$  (one host larva per female parasite, weight of host larva 8-10 mg; sex-ratio of parent parasite population 1:3) to 122.33 under the treatment  $a_0 b_0 c_2$  (one host larva per female parasite, weight of host larvae 30-35 mg, sex-ratio of parent parasite population 3:1). The results of chi-square test are given in Table 1-b. It is seen that all the main effects A, B and C (host larval density, weight of host larvae and sex-ratio of parent parasite population) as well as their interactions are highly significant.

The data representing the total and mean number of eggs laid at different levels of the main factors are summarized in Table 1-c and these are depicted in Fig. 1A, 1B and 1C. Among the levels of host larval density (A), maximum eggs (68.18) were obtained at two hosts per female parasite ( $a_1$ ). The same trend was also revealed for the other temperature and humidity conditions viz., 23°C and 75% RH and 30°C and 60% RH. The chi-square values

for the treatment comparisons between the different levels of the factors are indicated in Table 1-d.

The fecundity registered under the level  $a_1$  (two hosts per female parasite) was superior to those obtained at the levels  $a_2$  (three hosts per female parasite) and  $a_0$  (one host per female parasite) there being no significant difference between the levels  $a_2$  and  $a_0$ .

Among the levels of the host weight categories, the  $b_0$  level (host larvae weighing 30-35 mg) produced the maximum number of eggs (81.31), followed by the  $b_2$  level (a mixture of larvae with 30-35 mg and 8-10 mg in equal proportions) giving 66.76 eggs and under the level  $b_1$ , when the hosts were all relatively lighter, the number of eggs was quite low, the mean being 31.16. Significant differences were detected among all the three levels of B (weight of host larvae).

Among the levels of the sex-ratio of the parent parasite population (C), the maximum number of eggs were obtained at  $c_2$  (3 female:1 male), the mean being 44.73 (Table 1-e). At the  $c_1$  level (2:1) the fecundity was 44.33 and the two levels  $c_2$  and  $c_1$  were on par. When the sex-ratio was maintained at 1 female: 1 male ( $c_0$ ), the mean fecundity was 33.20. Under the two levels  $c_3$  (1 female: 2 males) and  $c_4$  (1 female: 3 males) a sharp

reduction in the fecundity was recorded, the mean values being respectively 20.18 and 28.96 and these two levels were on par. The low levels of fecundity for the two levels  $c_3$  and  $c_4$  were also recorded for the other two temperature and humidity combinations as well i.e., 23°C and 75% RH and 30°C and 60% RH.

Whether the effects of the different levels of the three main factors viz., host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population (C) were significantly different at different temperature and humidity conditions has been tested employing the chi-square method and the results of testing are furnished in Table 1-e. The chi-square due to heterogeneity of host larval density (A) being non-significant, it is evident that there is no heterogeneity in response to host larval density over the different temperature and humidity levels. The heterogeneity chi-square for the weight of host larvae (B) and sex-ratio of parent parasite population (C) are significant indicating that the responses vary between temperature and humidity combinations.

The  $b_0$  level (host larvae having a weight range 30-35 mg) produced the maximum number of eggs(81.31)

at 32°C and 50% RH (TH<sub>3</sub>). This is significantly higher than those at 30°C and 60% RH (TH<sub>2</sub>) and 28°C and 75% RH (TH<sub>1</sub>). But no significant difference was detected between the fecundity at 28°C and 75% RH (TH<sub>1</sub>) and 30°C and 60% RH (TH<sub>2</sub>).

At the b<sub>1</sub> level of host larval weight (larval weight 8-10 mg) maximum fecundity (46.40) was produced at 28°C and 75% RH (TH<sub>1</sub>). This is significantly higher than those at 30°C and 60% RH (TH<sub>2</sub>) and 32°C and 50% RH (TH<sub>3</sub>).

At the b<sub>2</sub> level (mixture of heavier and lighter larvae in equal proportions) maximum number of eggs were produced at 32°C and 50% RH (66.76). But no significant difference was detected between the fecundities at 32°C and 50% RH (TH<sub>3</sub>) and 28°C and 75% RH (TH<sub>1</sub>), while the fecundity at 28°C and 75% RH (TH<sub>1</sub>) is found to be significantly higher than that at 30°C and 60% RH (TH<sub>2</sub>).

When the sex-ratio of parent parasite population was kept at the c<sub>0</sub> level i.e., 1:1 (Female: male), no significant difference was observed in the egg production between the three temperature and humidity combinations (28°C and 75% RH; 30°C and 60% RH and 32°C and 50% RH) though the maximum fecundity (33.20) was obtained at 32°C and 50% RH (TH<sub>3</sub>).

At the  $c_1$  level of the sex-ratio of parent parasite population (1:2), maximum fecundity was (44.33) obtained at 32°C and 50% RH ( $TH_3$ ). But no significant difference was detected between fecundities at 32°C and 50% RH ( $TH_3$ ) and 28°C and 75% RH ( $TH_1$ ). At the  $c_2$  level (1:3) highest egg production was found to be at ( $TH_3$ ) 32°C and 50% RH (44.78). This is found significantly higher than those at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ). At the  $c_3$  level (1 female: 2 males) also maximum fecundity was found to be at ( $TH_3$ ) 32°C and 50% RH (28.18) which is significantly higher than those at 28°C and 75% RH ( $TH_1$ ). When the sex-ratio of parent parasite population was maintained at the  $c_4$  level also (1 female: 3 males), maximum egg production was obtained at 32°C and 50% RH (28.96). Significant differences were detected between the three temperature and humidity combinations (32°C and 50% RH, 28°C and 75% RH and 30°C and 60% RH).

## II. DURATION OF DEVELOPMENT

The duration of development was found out by calculating the weighted means of the number of days taken by the parasites of both sexes emerging on successive days. For this, the parasites emerging on successive days were isolated, killed and the male-female

composition in relation to the durations of development recorded.

Duration of development of B. brevicornis at 23°C and 75% RH.

Data on the mean duration of development of the parasites under different treatments at the above temperature and humidity combination are furnished in column 3 of Table 2-a and the raw data in Appendix II. The range in the duration of development was from 8.47 days under the treatment combination  $a_1 b_0 c_4$  (two host larvae per female parasite; larval weight 30-35 mg; sex-ratio of parent parasite population 1:3) to 11.73 days under the treatment combination  $a_0 b_0 c_1$  (one host larvae per female parasite; larval weight 30-35 mg; sex-ratio of parent parasite population 2:1).

Analysis of the data using the method of analysis of variance has been carried out and the results of the analysis of variance table is given in Table 2-b. It will be seen that the factors A (host larval density) and C (sex-ratio of parent parasite population) have significant influence on the duration of development of the parasites. The weight of host larvae (B) did not show any influence on the duration of development of the parasite. The interactions  $A \times C$  and  $A \times B \times C$  were also found to be significant.

The mean values of the duration of development at different levels of the three factors host larval density (A); weight of host larvae (B) and sex-ratio of parent parasite population (C) are furnished in Table 2-c and these are depicted in Fig. 2 A, 2B and 2C. The duration of development was shortest at the  $a_2$  level (three host larvae per female parasite) the mean being 9.607 days. With regard to the duration of development under  $a_1$  (two host larvae per female parasite) and  $a_2$  (three host larvae per female parasite) significant difference was not detected and the mean values were 9.764 and 9.607 days respectively.

The duration of development was found to be longest (10.104 days) under the level  $c_3$  i.e., when the parent parasite population was kept at one female : two males. The shortest duration of development of the parasite was recorded at  $c_2$  level (3females: 1 male) the mean duration of development being 9.574 days. The duration of development at the level  $c_2$  (3 females: 1 male) was found significantly superior to that at  $c_3$  level (1:2). The duration of development at the levels  $c_0$ ,  $c_1$  and  $c_4$  were on par.



Duration of development of *B. terrestris* in different temperature and humidity combinations

Sl. No.	Treatments	Mean duration of development in days		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
1.	a <sub>0</sub> b <sub>0</sub> e <sub>0</sub> <sup>a</sup>	9.37	9.88	8.80
2.	a <sub>0</sub> b <sub>0</sub> e <sub>1</sub>	11.73	8.63	8.73
3.	a <sub>0</sub> b <sub>0</sub> e <sub>2</sub>	10.63	9.90	9.67
4.	a <sub>0</sub> b <sub>0</sub> e <sub>3</sub>	9.47	8.63	8.53
5.	a <sub>0</sub> b <sub>0</sub> e <sub>4</sub>	10.77	88.33	9.27
6.	a <sub>0</sub> b <sub>1</sub> e <sub>0</sub>	10.03	11.40	8.07
7.	a <sub>0</sub> b <sub>1</sub> e <sub>1</sub>	9.00	9.60	8.00
8.	a <sub>0</sub> b <sub>1</sub> e <sub>2</sub>	10.13	8.63	8.00
9.	a <sub>0</sub> b <sub>1</sub> e <sub>3</sub>	10.37	8.00	8.43
10.	a <sub>0</sub> b <sub>1</sub> e <sub>4</sub>	10.13	9.73	8.00
11.	a <sub>0</sub> b <sub>2</sub> e <sub>0</sub>	9.73	8.80	8.00
12.	a <sub>0</sub> b <sub>2</sub> e <sub>1</sub>	10.50	9.03	8.30
13.	a <sub>0</sub> b <sub>2</sub> e <sub>2</sub>	9.97	8.67	8.30
14.	a <sub>0</sub> b <sub>2</sub> e <sub>3</sub>	9.70	8.47	8.00
15.	a <sub>0</sub> b <sub>2</sub> e <sub>4</sub>	9.70	9.13	8.37
16.	a <sub>1</sub> b <sub>0</sub> e <sub>0</sub>	9.87	10.43	9.57
17.	a <sub>1</sub> b <sub>0</sub> e <sub>1</sub>	10.03	8.97	8.83
18.	a <sub>1</sub> b <sub>0</sub> e <sub>2</sub>	7.00	8.60	8.70
19.	a <sub>1</sub> b <sub>0</sub> e <sub>3</sub>	9.97	8.70	8.47
20.	a <sub>1</sub> b <sub>0</sub> e <sub>4</sub>	8.67	9.10	9.23
21.	a <sub>1</sub> b <sub>1</sub> e <sub>0</sub>	9.53	8.47	9.13
22.	a <sub>1</sub> b <sub>1</sub> e <sub>1</sub>	9.70	8.70	8.20
23.	a <sub>1</sub> b <sub>1</sub> e <sub>2</sub>	8.47	8.10	8.10
24.	a <sub>1</sub> b <sub>1</sub> e <sub>3</sub>	9.67	8.97	7.93
25.	a <sub>1</sub> b <sub>1</sub> e <sub>4</sub>	9.67	8.77	8.47
26.	a <sub>1</sub> b <sub>2</sub> e <sub>0</sub>	9.97	9.10	8.21
27.	a <sub>1</sub> b <sub>2</sub> e <sub>1</sub>	9.57	8.67	8.40
28.	a <sub>1</sub> b <sub>2</sub> e <sub>2</sub>	9.33	8.00	8.37
29.	a <sub>1</sub> b <sub>2</sub> e <sub>3</sub>	10.47	8.00	9.57
30.	a <sub>1</sub> b <sub>2</sub> e <sub>4</sub>	10.13	9.57	9.70
31.	a <sub>2</sub> b <sub>0</sub> e <sub>0</sub>	9.90	9.07	8.53
32.	a <sub>2</sub> b <sub>0</sub> e <sub>1</sub>	8.80	8.00	8.17
33.	a <sub>2</sub> b <sub>0</sub> e <sub>2</sub>	9.20	8.17	8.13
34.	a <sub>2</sub> b <sub>0</sub> e <sub>3</sub>	10.40	8.97	8.23
35.	a <sub>2</sub> b <sub>0</sub> e <sub>4</sub>	9.80	8.30	8.33
36.	a <sub>2</sub> b <sub>1</sub> e <sub>0</sub>	9.43	8.57	8.30
37.	a <sub>2</sub> b <sub>1</sub> e <sub>1</sub>	9.07	8.30	8.17
38.	a <sub>2</sub> b <sub>1</sub> e <sub>2</sub>	9.00	8.37	8.07
39.	a <sub>2</sub> b <sub>1</sub> e <sub>3</sub>	11.03	8.33	8.83
40.	a <sub>2</sub> b <sub>1</sub> e <sub>4</sub>	9.70	8.60	8.27
41.	a <sub>2</sub> b <sub>2</sub> e <sub>0</sub>	9.60	8.40	8.43
42.	a <sub>2</sub> b <sub>2</sub> e <sub>1</sub>	9.60	8.57	8.60
43.	a <sub>2</sub> b <sub>2</sub> e <sub>2</sub>	9.43	8.93	8.50
44.	a <sub>2</sub> b <sub>2</sub> e <sub>3</sub>	9.80	8.30	9.67
45.	a <sub>2</sub> b <sub>2</sub> e <sub>4</sub>	9.40	8.40	9.97

a<sub>0</sub> - 1 host larva per female parasite  
a<sub>1</sub> - 2 " " " "  
a<sub>2</sub> - 3 " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
b<sub>1</sub> - " " " 8 - 10 mg  
b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

e<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)  
e<sub>1</sub> - " " " " " 2:1 (Female:Male)  
e<sub>2</sub> - " " " " " 3:1 (Female:Male)  
e<sub>3</sub> - " " " " " 1:2 (Female:Male)  
e<sub>4</sub> - " " " " " 1:3 (Female:Male)

Table 2-b

Duration of development of *B. brevicornis* at different temperature and humidity combinations

Analysis of variance table

Source	df	Mean squares		
		29°C and 75% RH	30°C and 60% RH	30°C and 50% RH
● A	2	2.640**	4.594**	1.063**
● B	2	1.342 H.S.	0.661 H.S.	3.196**
A x B	4	0.626 H.S.	1.425**	2.352**
● C	4	1.136*	3.351**	0.922**
A x C	8	1.398**	1.009**	0.589**
B x C	8	0.747 H.S.	1.030**	0.903**
A x B x C	16	0.878*	0.922**	0.337**
Error	90	0.479	0.251	0.152

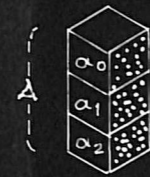
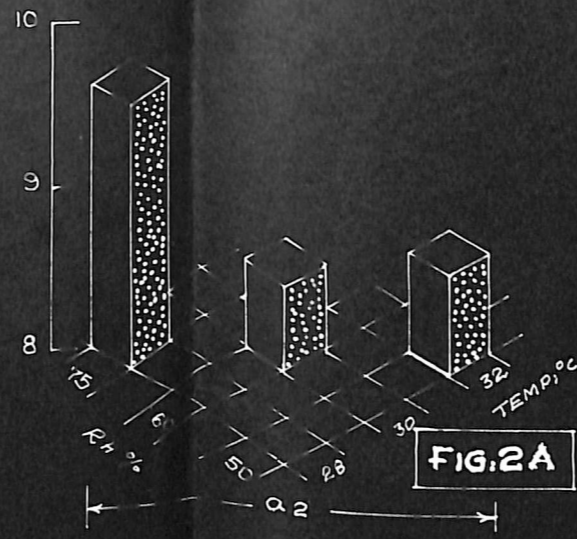
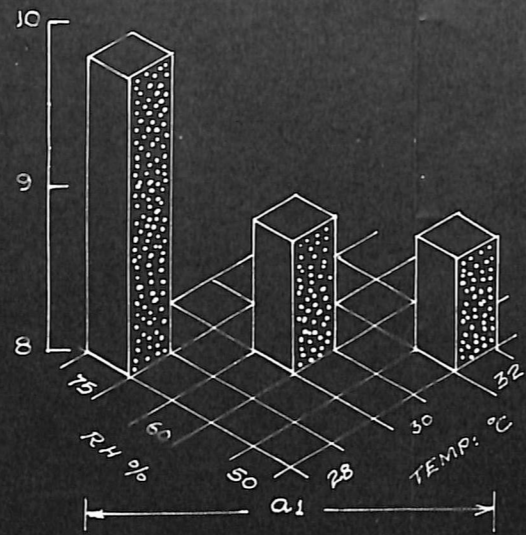
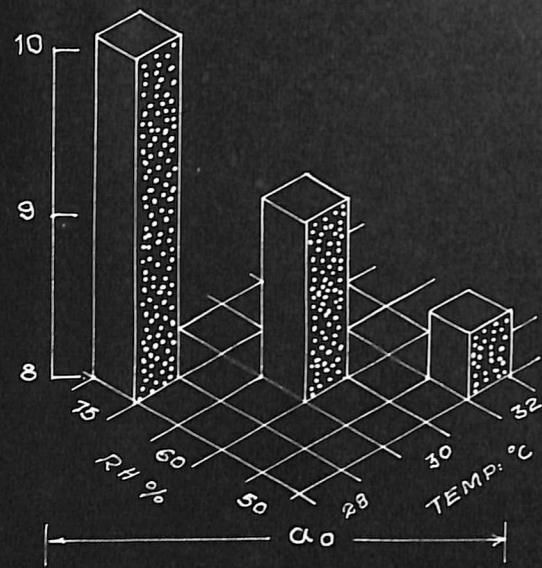
\* Significant at 5% level  
 \*\* Significant at 1% level  
 H.S. Not significant

● A - host larval density  
 ● B - weight of host larvae  
 ● C - sex-ratio of parent parasite population

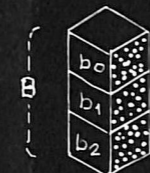
**Fig. 2 A - Duration of development of Bracon brevicornis at different temperature-humidity combinations as influenced by the host larval density**

**Fig. 2 B - Duration of development of Bracon brevicornis at different temperature-humidity combinations as influenced by the weight (size) of host larvae**

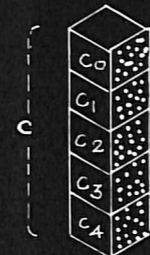
**Fig. 2 C - Duration of development of Bracon brevicornis at different temperature-humidity combinations as influenced by the sex-ratio of the parent parasite population**



1 HOST LARVA PER FEMALE PARASITE  
2 " " " "  
3 " " " "



LARVAL WEIGHT 30-35 mg  
" " 8-10 "  
HALF OF THE LARVAE OF THE 'b<sub>0</sub>' TYPE AND  
THE OTHER HALF OF THE 'b<sub>1</sub>' CATEGORY



1 FEMALE : 1 MALE  
2 " : 1 "  
3 " : 1 "  
1 " : 2 "  
1 " : 3 "

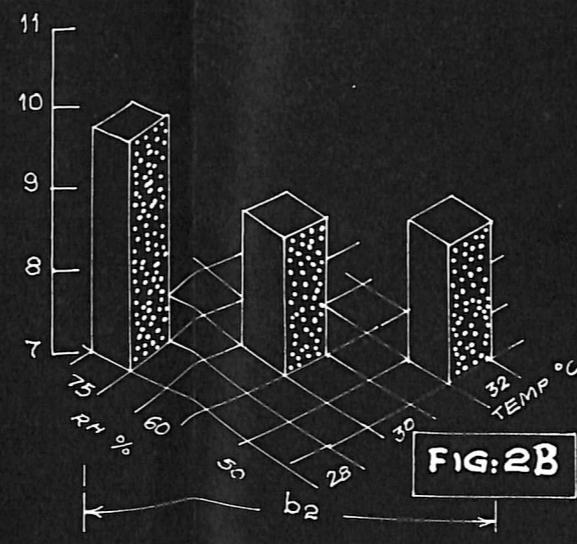
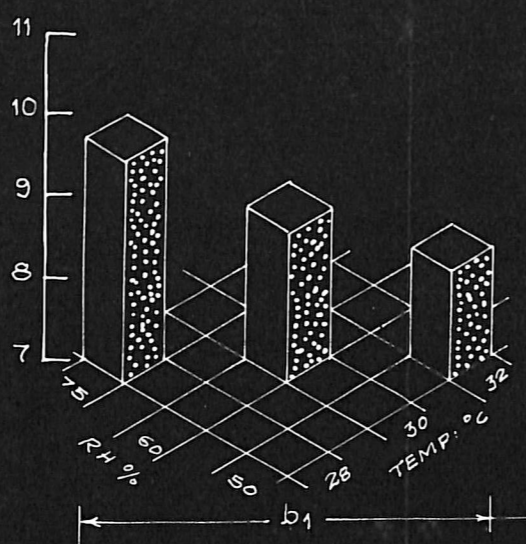
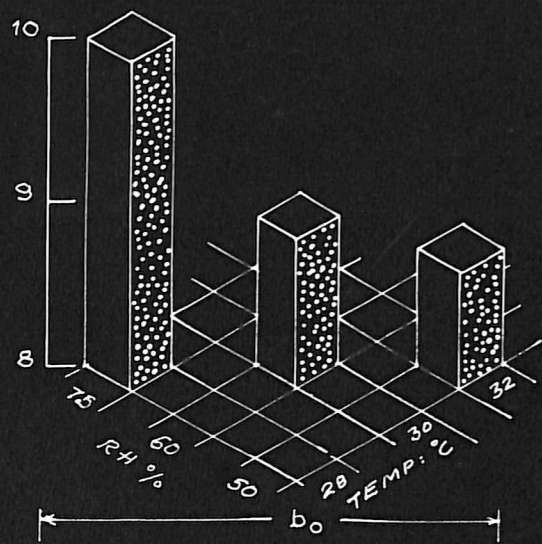


FIG. 2B

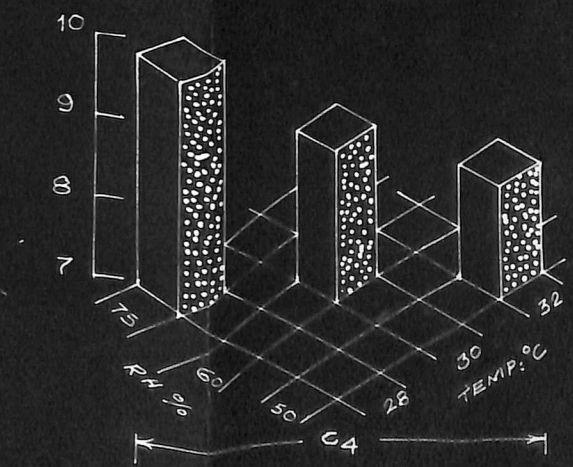
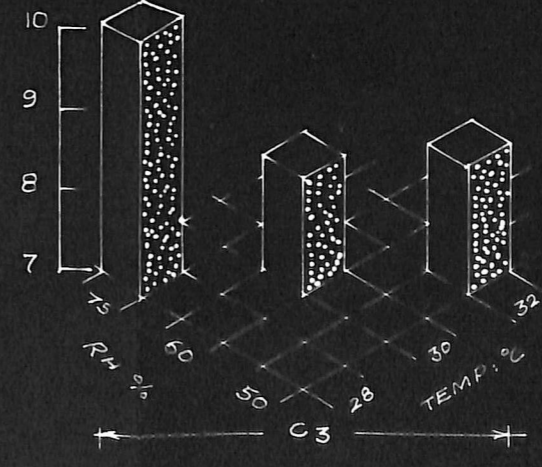
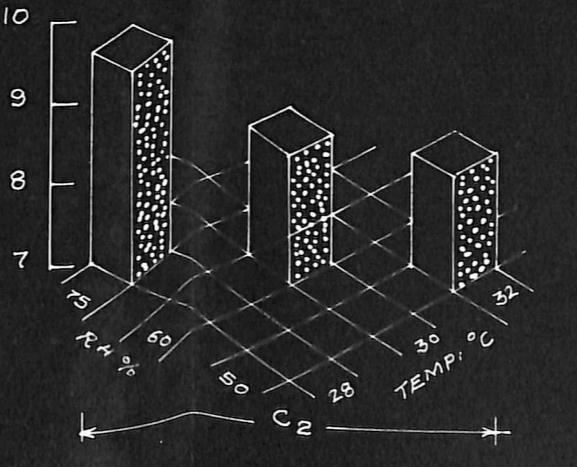
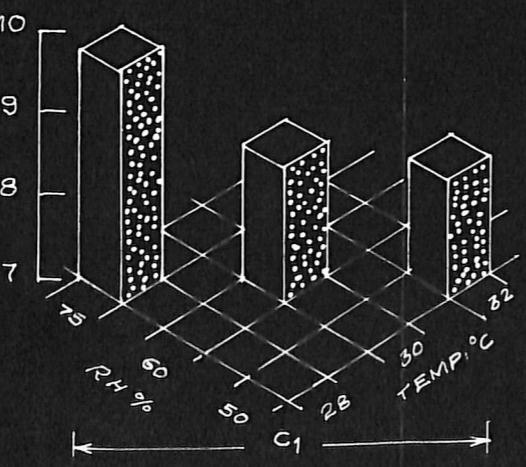
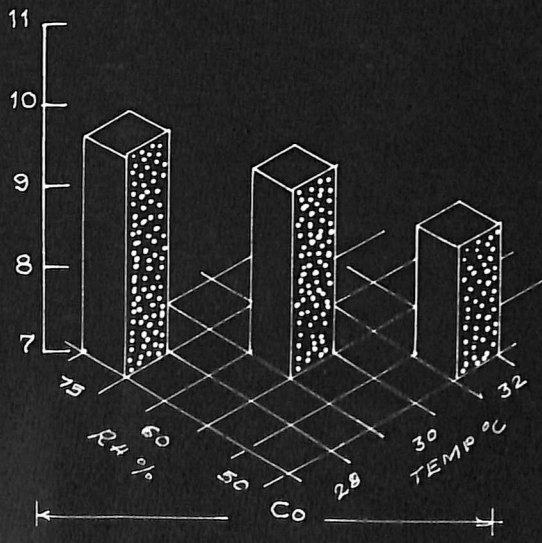


FIG.

**Table 2-c**  
**Duration of development of *B. brevicornis* at 28°C and 75% RH**  
**(Mean table)**

		* A levels			Mean	Ranking
		a <sub>0</sub>	a <sub>1</sub>	a <sub>2</sub>		
* B levels	b <sub>0</sub>	10.393	9.993	9.620	10.002	
	b <sub>1</sub>	9.993	9.407	9.640	9.660	
	b <sub>2</sub>	9.920	9.893	9.560	9.791	
* C levels	c <sub>0</sub>	9.711	9.789	9.611	9.704	
	c <sub>1</sub>	10.411	9.767	9.156	9.778	
	c <sub>2</sub>	10.244	9.267	9.211	9.574	<u>c<sub>2</sub>c<sub>0</sub>c<sub>1</sub>c<sub>4</sub>c<sub>3</sub></u>
	c <sub>3</sub>	9.844	10.053	10.433	10.104	
	c <sub>4</sub>	10.200	9.967	9.622	9.930	
Mean		10.082	9.764	9.607	9.818	<u>a<sub>2</sub> a<sub>1</sub> a<sub>0</sub></u>

		B levels			Mean
		b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>	
C levels	c <sub>0</sub>	9.711	9.667	9.733	9.704
	c <sub>1</sub>	10.189	9.256	9.889	9.778
	c <sub>2</sub>	9.944	9.200	9.578	9.574
	c <sub>3</sub>	9.944	10.356	10.011	10.104
	c <sub>4</sub>	10.224	9.822	9.744	9.930
Mean		10.002	9.660	9.791	9.818

CB for comparing levels of A = 0.282  
 " " B = 0.282  
 " " C = 0.364  
 " combinations of AB = 0.489  
 " " AC = 0.631  
 " " BC = 0.631

\* a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

\* b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and the other half of the 'b<sub>1</sub>' type

\* c - sex-ratio of parent parasite population -1 : 1 (Female:Male)  
 c<sub>0</sub> - " " " " 2 : 1  
 c<sub>1</sub> - " " " " 3 : 1  
 c<sub>2</sub> - " " " " 1 : 2  
 c<sub>3</sub> - " " " " 1 : 3

Table 2-4  
Duration of development of *B. brevicornis* at 30°C and 60% RH  
(Mean table)

		* A levels			Mean	Ranking
		a <sub>0</sub>	a <sub>1</sub>	a <sub>2</sub>		
* B levels	b <sub>0</sub>	9.073	9.160	8.500	8.911	
	b <sub>1</sub>	9.473	8.480	8.433	8.800	
	b <sub>2</sub>	8.820	8.667	8.520	8.669	
* C levels	c <sub>0</sub>	10.022	9.333	8.678	9.344	
	c <sub>1</sub>	9.089	8.778	8.289	8.719	
	c <sub>2</sub>	9.067	8.233	8.489	8.596	c <sub>3</sub> c <sub>2</sub> c <sub>1</sub> c <sub>4</sub> c <sub>0</sub>
	c <sub>3</sub>	8.367	8.356	8.533	8.419	
	c <sub>4</sub>	9.067	9.144	8.433	8.881	
Mean		9.122	8.769	8.484	8.792	a <sub>2</sub> a <sub>1</sub> a <sub>0</sub>
		B levels			Mean	
		b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>		
C levels	c <sub>0</sub>	9.789	9.478	8.767	9.344	
	c <sub>1</sub>	8.533	8.867	8.756	8.719	
	c <sub>2</sub>	8.889	8.367	8.533	8.596	
	c <sub>3</sub>	8.767	8.233	8.256	8.419	
	c <sub>4</sub>	8.578	9.033	9.033	8.881	
Mean		8.911	8.800	8.669	8.792	

OD for comparing levels of	A = 0.211
" " "	B = 0.211
" " "	C = 0.273
" " combinations of	AB = 0.366
" " "	AC = 0.473
" " "	BC = 0.473

\* a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " " " "  
 a<sub>2</sub> - 3 " " " " " "

\* b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and the other half of the 'b<sub>1</sub>' type

\* sex-ratio of parent parasite population - 1:1 (Female:Male)  
 " " " " " " 2:1  
 " " " " " " 3:1  
 " " " " " " 1:2  
 " " " " " " 1:3

**Table 2-c**  
**Duration of development of *B. brevicornis* at 32°C and 50% RH**  
**(Mean table)**

		a <sub>0</sub>	* A levels a <sub>1</sub>	a <sub>2</sub>	Mean	Ranking
* B levels	b <sub>0</sub>	9.000	8.960	8.280	8.747	
	b <sub>1</sub>	8.100	8.367	8.327	8.264	b <sub>1</sub> b <sub>2</sub> b <sub>0</sub>
	b <sub>2</sub>	8.193	8.880	9.033	8.702	
* C levels	c <sub>0</sub>	8.289	9.022	8.422	8.578	
	c <sub>1</sub>	8.344	8.478	8.311	8.378	
	c <sub>2</sub>	8.656	8.389	8.233	8.426	c <sub>1</sub> c <sub>2</sub> c <sub>4</sub> c <sub>0</sub> c <sub>3</sub>
	c <sub>3</sub>	8.322	8.656	8.911	8.630	
	c <sub>4</sub>	8.544	9.133	8.855	8.440	
Mean		8.431	8.736	8.547	8.571	a <sub>0</sub> a <sub>2</sub> a <sub>1</sub>

		B levels			Mean
		b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>	
C levels	c <sub>0</sub>	8.967	8.500	8.267	8.578
	c <sub>1</sub>	8.578	8.122	8.433	8.378
	c <sub>2</sub>	8.833	8.056	8.389	8.426
	c <sub>3</sub>	8.411	8.400	9.078	8.630
	c <sub>4</sub>	8.944	8.244	9.344	8.440
Mean		8.747	8.264	8.702	8.571

CD for comparing levels of	A = 0.165
" " "	B = 0.165
" " "	C = 0.212
" " combinations of	AB = 0.285
" " "	AC = 0.368
" " "	BC = 0.368

\* a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

\* b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and the other half of the 'b<sub>1</sub>' type

\* sex-ratio of parent parasite population - 1:1 (Female:Male)  
 " " " " " " 2:1 " "  
 " " " " " " 3:1 " "  
 " " " " " " 1:2 " "  
 " " " " " " 1:3 " "

Duration of development of B. brevicornis at 30°C and 60% RH.

Data on the mean duration of development under various treatments are given in column 4 of Table 2-a. Minimum duration of development was registered for the combination of one host larva per female parasite which was of lighter type and with one female : three males ratio for the parent parasite population ( $a_0 b_1 c_3$ ) being eight days. The levels  $a_1 b_2 c_0$ ,  $a_1 b_2 c_3$  and  $a_2 b_0 c_1$  also indicated relatively short durations of development. Maximum duration of development was 11.4 days for the treatment combination of one host larva per female parasite, host larvae of lighter type (8-10 mg) and one female: one male ratio of the parent parasite population ( $a_0 b_1 c_1$ ).

The abstract of the analysis of variance table is given in Table 2-b. The effects of host larval density (A) and sex-ratio of parent parasite population (C) were significant while weight of host larvae (B) had no significant effect on the duration of development of the parasite at 30°C and 60% RH. The interactions AxB, AxC, BxC and AxBrC are also found to be significant.

The mean values of the duration of development at different levels of the factors are furnished in



Table 2-d and these are represented in Fig. 2A, 2B and 2C. The longest duration of development (9.122 days) was observed at the  $a_0$  level (one host larva per female parasite). At the  $a_2$  level (two host larvae per female parasite) the duration of development was 8.484 days. The mean duration of development at the  $a_0$  level (one host larva per female parasite) is significantly longer than the durations for  $a_1$  (two host larvae per female parasite) and  $a_2$  (three host larvae per female parasite) levels. The durations of development at the  $a_1$  and  $a_2$  levels were also found to be significantly different.

In relation to the sex-ratio of the parent parasite population, the  $c_0$  level (1 female: 1 male in the parent population) recorded the longest duration of development of 9.344 days. Under  $c_3$  level i.e., when the sex-ratio of parent population was maintained at one female: three males, the developmental period was the lowest being 8.419 days. The mean duration of development for  $c_0$  level (1 female: 1 male) is significantly higher than for the other levels  $c_1$ ,  $c_2$ ,  $c_3$  and  $c_4$ .

Duration of development at 32°C and 50% RH

Data on the mean duration of development under various treatments are furnished in column 5 of Table 2-a.

The duration of development ranged from 7.93 days under the treatment  $a_1 b_1 c_3$  (two host larvae per female parasite; weight of host larvae 8-10 mg sex-ratio of parent population one female; two males) to 9.97 days under the treatment  $a_2 b_2 c_4$  (three host larvae per female parasite; a mixture of host larvae with 30-35 mg weight and 8-10 mg in equal proportions; sex-ratio of parent parasite population - one female; three males).

The results of the analysis of variance is furnished in Table 2-b. It is indicated that the effects of host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population (C) are significant. The mean duration of development at different levels of the factors are given in Table 2-c and are illustrated in Fig. 2A, 2B and 2C.

It will be seen from the Table 2-c that the parasites produced at the  $a_0$  level (1 host larva per female parasite) recorded the shortest duration of development (8.431 days). But no significant difference was detected in the duration of development between the  $a_0$  and  $a_2$  levels.

Among the levels of B, the  $b_1$  level (larval weight 8-10 mg) recorded the shortest duration of

development 8.264 which was significantly superior to that recorded at  $b_2$  and  $b_0$  levels.

The shortest duration of development (8.378 days) was recorded for the  $c_1$  level (2 females : 1 male for the parent population) the duration of development at the  $c_1$  level was significantly superior to those recorded at  $c_4$  (1 female: 3 males) and  $c_3$  (1 female: 2 males) levels.

### III. PROGENY PRODUCTION

Progeny production at 28°C and 75% RH

The total number of males and females which emerged out from each replicate was recorded as the  $F_1$  progeny production under different treatment combinations. The data on the mean number of progenies produced under various treatments at 28°C and 75% RH are furnished in column 3 of Table 3-a and the raw data is given in Appendix III. The mean progeny production varied widely from 2.00 for the treatment  $a_0 b_1 c_1$  (one host larva per female parasite; weight of host larvae 8-10 mg; two females: one male in the parent parasite population) to 25.67 under the treatment  $a_0 b_0 c_3$  (one host larva per female parasite; weight of the host larva 30-35 mg, a ratio of one female: two males in the parent parasite

population). The data being in the form of counts, the analysis of variance technique was not found applicable in this case even after square root transformation. Therefore, the chi-square test, which is a distribution free test has been adopted as indicated in materials and methods and the results of testing are furnished in Table 3-b. All the main effects viz., host larval density (A), weight of the host larvae (B) and sex-ratio of the parent parasite population (C) as well as their interactions were found to be significant.

The data on the total number of progenies produced and their mean values are presented in Table 3-c and these are graphically depicted in Fig. 3A, 3B and 3C. The different levels of the three factors viz., host larval density (A), weight of the host larvae (B) and sex-ratio of the parent parasite population (C) are compared using the chi-square test and the results are given in table 3-d. Maximum number of progeny production was registered with two host larvae per female parasite ( $a_1$ ) the mean being 12.31 and the minimum (9.20) was recorded with three host larvae per female parasite of  $a_2$  level and the difference between the two levels was significant. No significant difference was detected between the  $a_0$  (one host larva per female parasite)

Table 3-a

Progeny production of *B. brevicornis* in different temperature and humidity combinations

Sl. No.	Treatments	Mean number of progenies		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
1.	a <sub>0</sub> b <sub>0</sub> c <sub>0</sub> <sup>a</sup>	18.00	19.33	22.00
2.	a <sub>0</sub> b <sub>0</sub> c <sub>1</sub>	13.33	20.00	31.33
3.	a <sub>0</sub> b <sub>0</sub> c <sub>2</sub>	5.00	3.67	12.33
4.	a <sub>0</sub> b <sub>0</sub> c <sub>3</sub>	26.67	34.67	16.67
5.	a <sub>0</sub> b <sub>0</sub> c <sub>4</sub>	7.67	13.33	11.33
6.	a <sub>0</sub> b <sub>1</sub> c <sub>0</sub>	4.67	9.67	12.67
7.	a <sub>0</sub> b <sub>1</sub> c <sub>1</sub>	2.00	18.33	5.33
8.	a <sub>0</sub> b <sub>1</sub> c <sub>2</sub>	10.00	22.00	8.33
9.	a <sub>0</sub> b <sub>1</sub> c <sub>3</sub>	9.67	4.00	5.67
10.	a <sub>0</sub> b <sub>1</sub> c <sub>4</sub>	7.33	7.00	4.33
11.	a <sub>0</sub> b <sub>2</sub> c <sub>0</sub>	6.67	8.00	35.00
12.	a <sub>0</sub> b <sub>2</sub> c <sub>1</sub>	21.67	6.33	32.33
13.	a <sub>0</sub> b <sub>2</sub> c <sub>2</sub>	10.00	11.33	28.33
14.	a <sub>0</sub> b <sub>2</sub> c <sub>3</sub>	14.33	6.00	21.33
15.	a <sub>0</sub> b <sub>2</sub> c <sub>4</sub>	11.33	8.00	13.33
16.	a <sub>1</sub> b <sub>0</sub> c <sub>0</sub>	18.00	16.33	39.33
17.	a <sub>1</sub> b <sub>0</sub> c <sub>1</sub>	11.00	22.00	64.33
18.	a <sub>1</sub> b <sub>0</sub> c <sub>2</sub>	15.67	11.67	19.33
19.	a <sub>1</sub> b <sub>0</sub> c <sub>3</sub>	7.00	7.67	8.67
20.	a <sub>1</sub> b <sub>0</sub> c <sub>4</sub>	8.33	12.33	9.67
21.	a <sub>1</sub> b <sub>1</sub> c <sub>0</sub>	17.00	13.67	7.33
22.	a <sub>1</sub> b <sub>1</sub> c <sub>1</sub>	17.00	20.33	11.67
23.	a <sub>1</sub> b <sub>1</sub> c <sub>2</sub>	14.67	4.67	6.33
24.	a <sub>1</sub> b <sub>1</sub> c <sub>3</sub>	14.00	11.67	11.00
25.	a <sub>1</sub> b <sub>1</sub> c <sub>4</sub>	7.67	8.00	15.33
26.	a <sub>1</sub> b <sub>2</sub> c <sub>0</sub>	10.67	33.67	12.67
27.	a <sub>1</sub> b <sub>2</sub> c <sub>1</sub>	7.67	20.67	29.33
28.	a <sub>1</sub> b <sub>2</sub> c <sub>2</sub>	15.00	14.33	9.00
29.	a <sub>1</sub> b <sub>2</sub> c <sub>3</sub>	10.33	14.00	18.67
30.	a <sub>1</sub> b <sub>2</sub> c <sub>4</sub>	12.67	10.00	13.33
31.	a <sub>2</sub> b <sub>0</sub> c <sub>0</sub>	11.00	17.33	31.33
32.	a <sub>2</sub> b <sub>0</sub> c <sub>1</sub>	7.00	30.67	20.33
33.	a <sub>2</sub> b <sub>0</sub> c <sub>2</sub>	12.00	12.67	24.67
34.	a <sub>2</sub> b <sub>0</sub> c <sub>3</sub>	13.33	8.67	36.00
35.	a <sub>2</sub> b <sub>0</sub> c <sub>4</sub>	9.67	6.33	9.00
36.	a <sub>2</sub> b <sub>1</sub> c <sub>0</sub>	3.67	29.00	16.00
37.	a <sub>2</sub> b <sub>1</sub> c <sub>1</sub>	12.00	9.33	6.00
38.	a <sub>2</sub> b <sub>1</sub> c <sub>2</sub>	12.00	12.00	7.67
39.	a <sub>2</sub> b <sub>1</sub> c <sub>3</sub>	9.67	9.33	7.67
40.	a <sub>2</sub> b <sub>1</sub> c <sub>4</sub>	10.00	8.00	14.00
41.	a <sub>2</sub> b <sub>2</sub> c <sub>0</sub>	8.33	13.67	40.67
42.	a <sub>2</sub> b <sub>2</sub> c <sub>1</sub>	14.00	19.33	36.33
43.	a <sub>2</sub> b <sub>2</sub> c <sub>2</sub>	5.67	15.00	39.00
44.	a <sub>2</sub> b <sub>2</sub> c <sub>3</sub>	4.67	9.00	26.00
45.	a <sub>2</sub> b <sub>2</sub> c <sub>4</sub>	5.00	19.00	18.00

a<sub>0</sub> - 1 host larva per female parasitea<sub>1</sub> - 2 " " " "a<sub>2</sub> - 3 " " " "b<sub>0</sub> - larval weight 30 - 35 mgb<sub>1</sub> - " " 8 - 10 mgb<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' typec<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)c<sub>1</sub> - " " " " 2:1 { " }c<sub>2</sub> - " " " " 3:1 { " }c<sub>3</sub> - " " " " 1:2 { " }c<sub>4</sub> - " " " " 1:3 { " }

**Fig. 3 A - Progeny production of Bracon brevicornis at different temperature-humidity combinations as influenced by the host larval density**

**Fig. 3 B - Progeny production of Bracon brevicornis at different temperature-humidity combinations as influenced by the weight (size) of host larvae**

**Fig. 3 C - Progeny production of Bracon brevicornis at different temperature-humidity combinations as influenced by the sex-ratio of the parent parasite population**

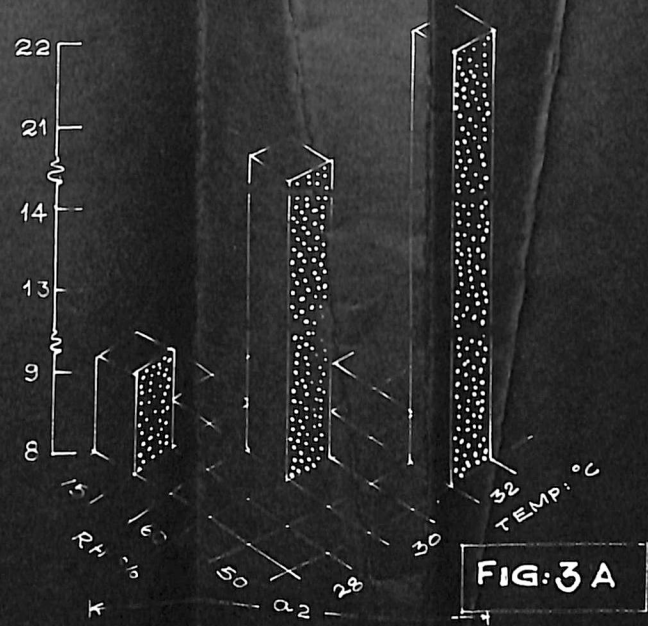
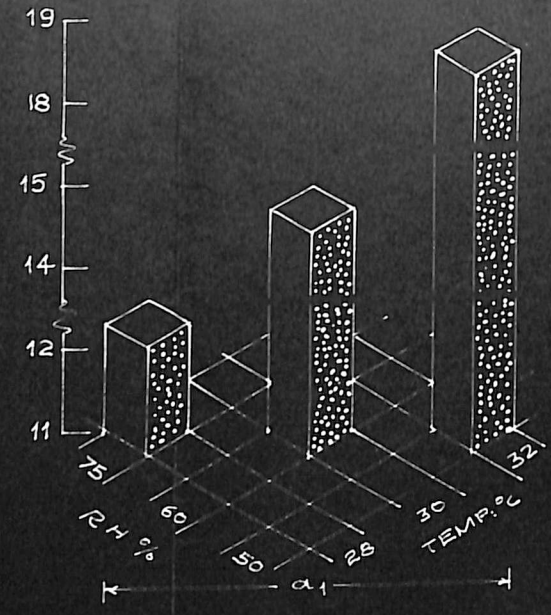
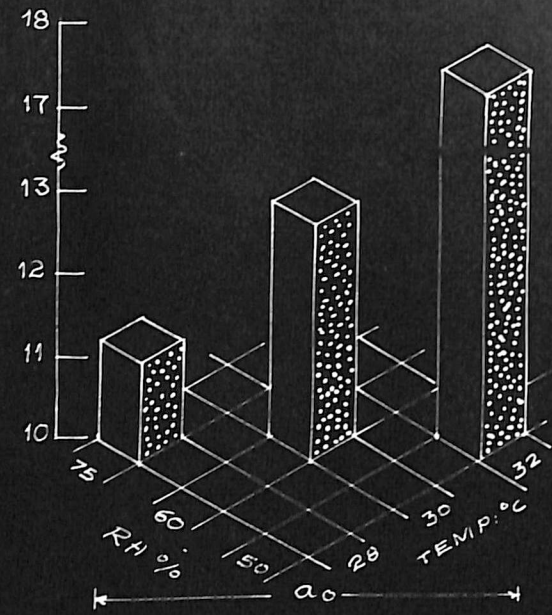
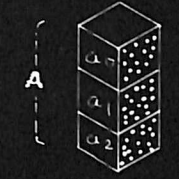


FIG: 3 A



1 HOST LARVA PER FEMALE PARASITE  
2 " " " " " "  
3 " " " " " "

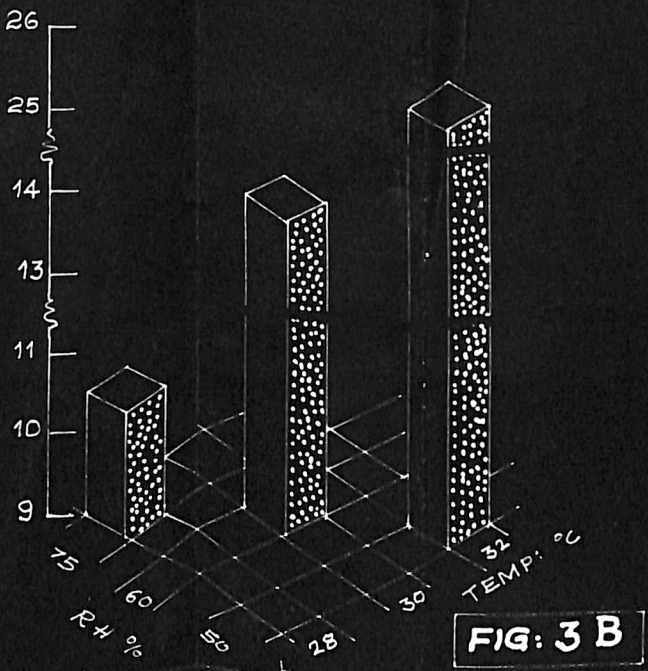
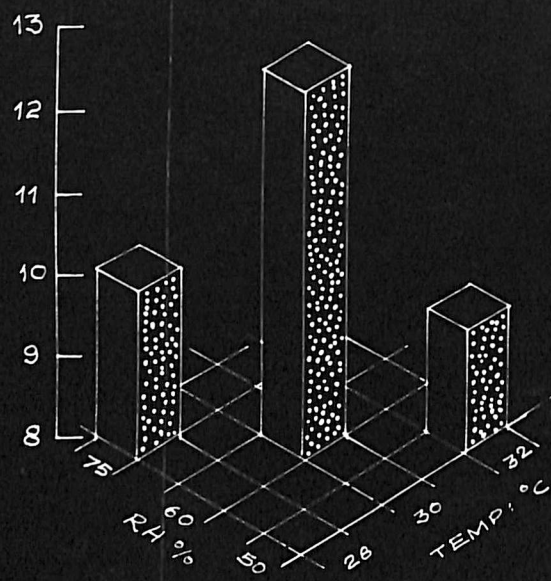
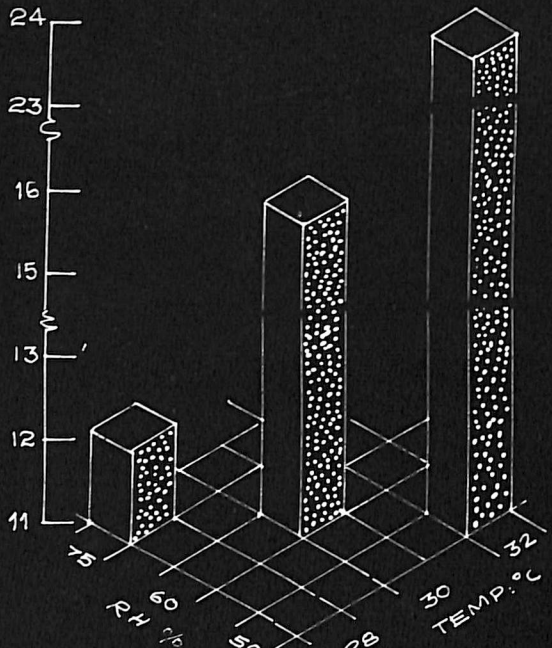
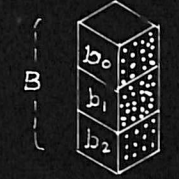
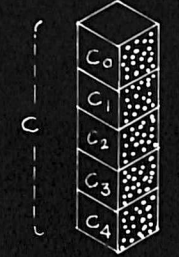


FIG: 3 B



LARVAL WEIGHT 30-35 mg  
" " " " " " 8-10  
HALF OF THE LARVAE OF THE  $b_0$  TYPE AND  
THE OTHER HALF OF THE  $b_1$  CATEGORY



1 FEMALE : 1 MALE  
2 " " " " " "  
3 " " " " " "  
1 " " " " " "  
1 " " " " " "

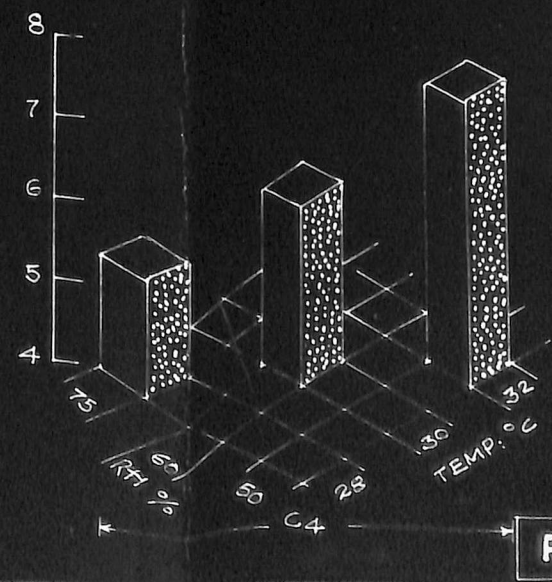
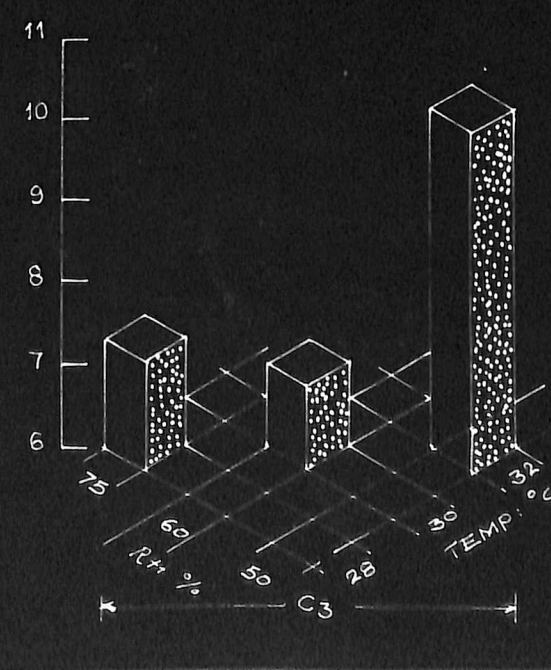
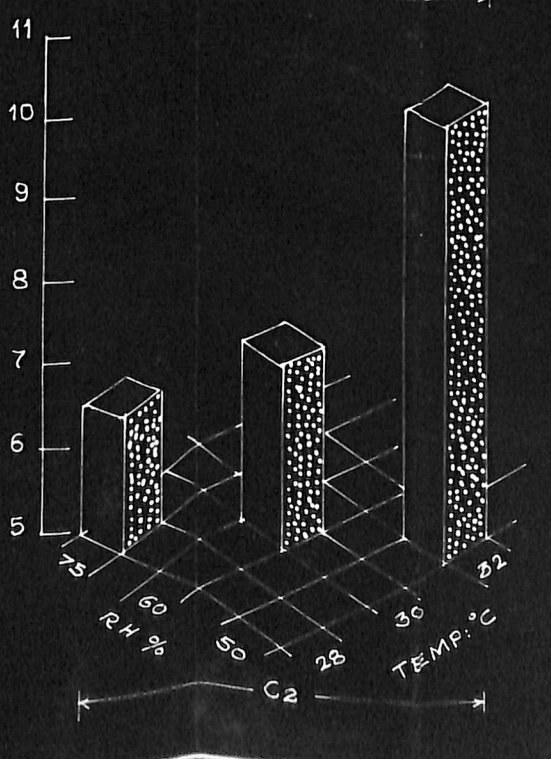
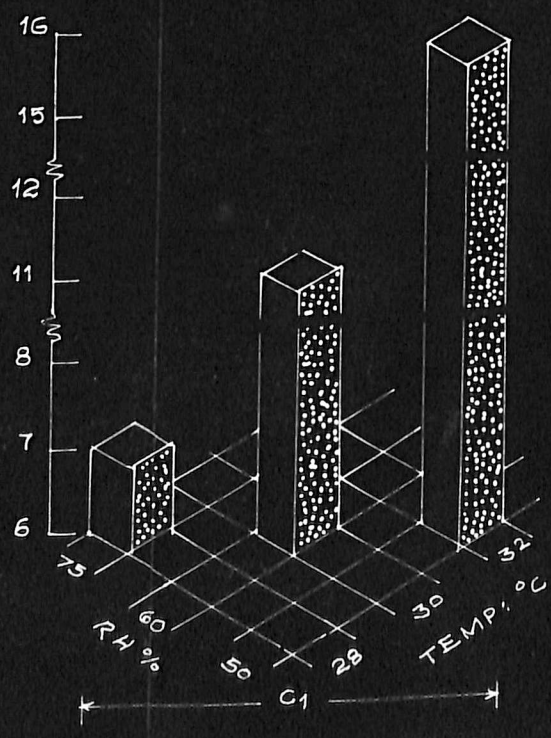
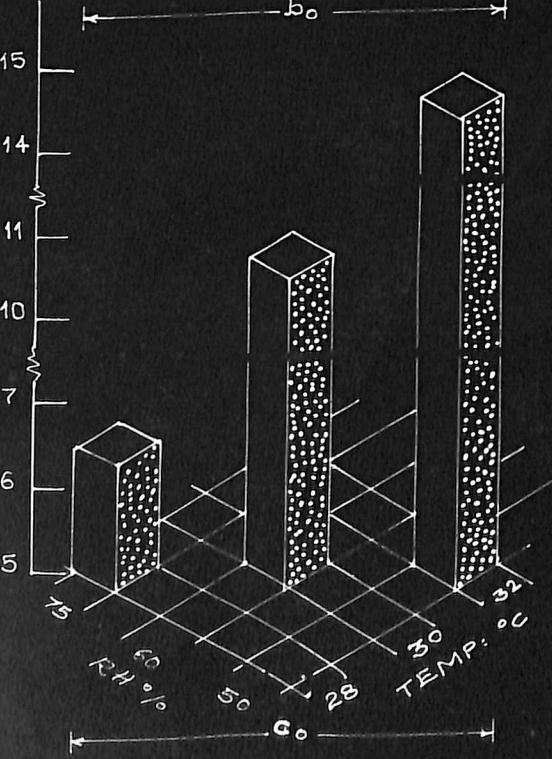


FIG: 3 C

Table 3-b

Chi-square values for the effects of the three factors A, B and C at different temperature and humidity combination on progeny production in B. brachicornis

Source	df	Chi-squares		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
<sup>⊙</sup> A	2	20.56**	7.73*	30.70**
<sup>⊙</sup> B	2	9.32**	17.68**	350.05**
A x B	4	46.49**	67.09**	94.96**
<sup>⊙</sup> C	4	16.22**	114.57**	189.15**
A x C	8	50.64**	50.78**	110.79**
B x C	8	46.75**	55.33**	143.66**
A x B x C	16	98.01**	231.74**	196.77**

\* Significant at 5% level  
\*\* Significant at 1% level

<sup>⊙</sup>A - Host larval density  
<sup>⊙</sup>B - Weight of host larvae  
<sup>⊙</sup>C - Sex-ratio of parent parasite population



Table 3-c

Number of progenies produced by *B. brevicornis* at different levels of the main factors at different temperature and humidity combinations.

	28°C and 75% RH		30°C and 60% RH		32°C and 50% RH	
	Total	Mean	Total	Mean	Total	Mean
a <sub>0</sub> *	505	11.22	575	12.78	781	17.36
a <sub>1</sub>	554	12.31	665	14.73	838	18.62
a <sub>2</sub>	414	9.20	658	14.62	1004	22.31
b <sub>0</sub> *	545	12.11	710	15.78	1069	23.76
b <sub>1</sub>	454	10.09	561	12.47	424	9.42
b <sub>2</sub>	474	10.53	625	13.89	1130	25.11
c <sub>0</sub> *	294	6.53	482	10.71	661	14.69
c <sub>1</sub>	317	7.04	501	11.13	711	15.80
c <sub>2</sub>	294	6.53	322	7.16	465	10.33
c <sub>3</sub>	329	7.31	315	7.00	455	10.11
c <sub>4</sub>	239	5.31	276	6.13	331	7.26

a<sub>0</sub>\* - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

b<sub>0</sub>\* - larval weight 30 - 35 mg.  
 b<sub>1</sub> - " " " 8 - 10 mg.  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and other half of the 'b<sub>1</sub>' type

c<sub>0</sub>\* - Sex-ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>1</sub> - " " " " - 2:1 " "  
 c<sub>2</sub> - " " " " - 3:1 " "  
 c<sub>3</sub> - " " " " - 1:2 " "  
 c<sub>4</sub> - " " " " - 1:3 " "

Table 3-d

Chi-square values for comparing the progeny production under different levels of the main factors A, B and C at different temperature and humidity combinations in B. brevicornis

Chi-square values						
	28°C and 75% RH	Rank- ing	30°C and 60% RH	Rank- ing	32°C and 50% RH	Rank- ing
a <sub>0</sub> Vs a <sub>1</sub> <sup>⊙</sup>	2.07 NS	a <sub>1</sub> a <sub>0</sub>	6.26*	a <sub>1</sub> a <sub>0</sub>	2.01 NS	a <sub>1</sub> a <sub>0</sub>
a <sub>1</sub> Vs a <sub>2</sub>	20.25**	a <sub>1</sub> a <sub>2</sub>	0.02 NS	a <sub>1</sub> a <sub>2</sub>	14.96**	a <sub>1</sub> a <sub>2</sub>
a <sub>0</sub> Vs a <sub>2</sub>	9.01**	a <sub>1</sub> a <sub>2</sub>	5.59*	a <sub>1</sub> a <sub>2</sub>	27.86**	a <sub>1</sub> a <sub>2</sub>
b <sub>0</sub> Vs b <sub>1</sub> <sup>⊙</sup>	0.23**	b <sub>2</sub> b <sub>0</sub>	17.47**	b <sub>2</sub> b <sub>0</sub>	278.65**	b <sub>2</sub> b <sub>0</sub>
b <sub>1</sub> Vs b <sub>2</sub>	0.43 NS	b <sub>2</sub> b <sub>0</sub>	3.45 NS	b <sub>2</sub> b <sub>0</sub>	320.74**	b <sub>2</sub> b <sub>0</sub>
b <sub>0</sub> Vs b <sub>2</sub>	4.95*	b <sub>2</sub> b <sub>0</sub>	5.41*	b <sub>2</sub> b <sub>0</sub>	1.69 NS	b <sub>2</sub> b <sub>0</sub>
c <sub>0</sub> Vs c <sub>1</sub> <sup>⊙</sup>	0.07 NS		0.37 NS		1.82 NS	
c <sub>0</sub> Vs c <sub>2</sub>	0 NS		31.84**		34.12**	
c <sub>0</sub> Vs c <sub>3</sub>	1.97 NS		34.99**		33.03**	
c <sub>0</sub> Vs c <sub>4</sub>	5.68*		55.98**		103.70**	
c <sub>1</sub> Vs c <sub>2</sub>	0.87 NS		38.93**		51.46**	
c <sub>1</sub> Vs c <sub>3</sub>	0.22 NS		43.31**		56.21**	
c <sub>1</sub> Vs c <sub>4</sub>	5.44*		65.15**		130.53**	
c <sub>2</sub> Vs c <sub>3</sub>	1.97 NS		0.08 NS		0.11 NS	
c <sub>2</sub> Vs c <sub>4</sub>	5.68*		3.54 NS		22.56**	
c <sub>3</sub> Vs c <sub>4</sub>	14.26**		2.57 NS		10.56**	

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

a<sub>0</sub><sup>⊙</sup> - 1 host larva per female parasite  
 a<sub>1</sub><sup>⊙</sup> - 2 " " " "  
 a<sub>2</sub><sup>⊙</sup> - 3 " " " "

b<sub>0</sub><sup>⊙</sup> - larval weight 30 - 35 mg  
 b<sub>1</sub><sup>⊙</sup> - " " " 8 - 10 mg  
 b<sub>2</sub><sup>⊙</sup> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

c<sub>0</sub><sup>⊙</sup> - sex-ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>1</sub><sup>⊙</sup> - " " " " - 2:1 " "  
 c<sub>2</sub><sup>⊙</sup> - " " " " - 3:1 " "  
 c<sub>3</sub><sup>⊙</sup> - " " " " - 1:2 " "  
 c<sub>4</sub><sup>⊙</sup> - " " " " - 1:3 " "

Table 3-e  
Chi-square values for comparison of the three main factors A, B and C on progeny production of B. brevicornis at different temperature and humidity combinations

Ranking	Chi-square for levels of A <sup>a</sup>			Chi-square for levels of B <sup>b</sup>			Chi-square for levels of C <sup>c</sup>				
	a <sub>0</sub>	a <sub>1</sub>	a <sub>2</sub>	b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>	c <sub>0</sub>	c <sub>1</sub>	c <sub>2</sub>	c <sub>3</sub>	c <sub>4</sub>
	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>2</sub>
	4.54*	9.76**	55.54**	21.69**	11.28**	20.75**	45.55**	41.39**	1.27 NS	0.30 NS	2.66 NS
	31.29**	20.40**	72.03**	72.45**	19.05**	145.31**	28.03**	36.39**	25.98**	25.45**	4.98*
	59.23**	57.94**	245.49**	170.12**	1.03 NS	268.29**	141.03**	151.01**	38.53**	20.25**	14.85**

Chi-square due to heterogeneity

Host larval density (A)	44.72**
Weight of host larvae (B)	40.68**
Sex-ratio of parent parasite population (C)	57.86**

0A - Host larval density  
 a<sub>0</sub> - 1 larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

0B - Weight of host larvae  
 b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

0C - Sex-ratio of parent population  
 c<sub>0</sub> - 1 female : 1 male  
 c<sub>1</sub> - 2 female : 1 male  
 c<sub>2</sub> - 3 female : 1 male  
 c<sub>3</sub> - 1 female : 2 males  
 c<sub>4</sub> - 1 female : 3 males

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

TH<sub>1</sub> - 28°C and 75% RH  
 TH<sub>2</sub> - 30°C and 60% RH  
 TH<sub>3</sub> - 32°C and 50% RH

and  $a_1$  (two host larvae per female parasite) levels.

With regard to the weight of host larvae, the maximum progeny production was obtained with the  $b_0$  level having host larvae of weight 30-35 mg, with a mean value of 12.11. The larvae with 8-10 mg weight range ( $b_1$ ) produced minimum number of progenies having a mean value 10.09. There was no significant difference in progeny production between the  $b_1$  and  $b_2$  levels. The progeny production from heavier larvae ( $b_0$ ) was significantly better than from larvae of the level  $b_2$  i.e. heavier and lighter larvae in equal proportions.

Among the levels of the sex-ratio of parent parasite population (C), the maximum progenies (7.31) were produced by the  $c_3$  level (1 female: 2 males for the parent parasite population). The level  $c_4$  (1 female: 3 males) gave a much lower (5.31) number of progenies. The level  $c_3$  (1 female: 2 males) was found to be on par with the levels  $c_1$  (2 females: 1 male),  $c_2$  (3 females: 1 male), and  $c_0$  (1 female: 1 male).

Progeny production at 30°C and 60% RH

The data on the mean number of progenies produced under various treatments at the above temperature and humidity combination are furnished in column 4 Table 3-a.

The production of progeny ranged from 3.67 under the treatment  $a_0 b_0 c_2$  (one host per female parasite; larval weight range 30-35 mg; sex-ratio of parent parasite population 3 females: 1 male) to 34.67 under the treatment  $a_0 b_0 c_3$  (one host larva per female parasite; larval weight 30-35 mg; sex-ratio of parent population one female: two males).

Chi-square test involving the factors (Table 3-b) have indicated that all the main effects viz., host larval density, weight of host larvae and sex-ratio of parent population as well as their interactions  $A \times B$ ,  $A \times C$ ,  $B \times C$  and  $A \times B \times C$  are significant. The total and mean values of progeny production at different levels of the factors are summarised in Table 3-c and these are illustrated in Fig. 3A, 3B and 3C. The different levels of the factors have been compared using chi-square test and the results are given in Table 3-d.

Among the levels of host density (A) maximum progeny production (14.75) was produced with two hosts per female parasite ( $a_1$ ). No significant difference was detected between  $a_1$  (two host larvae per female parasite) and  $a_2$  (three host larvae per female parasite). The  $a_0$  level (one host larva per female parasite) gave significantly lower number of progenies of mean 12.75.

With regard to the levels of weight of host larvae (B), larvae with a weight range of 30-35 mg ( $b_0$ ) recorded the maximum number of progenies, the mean being 15.78 and this was significantly higher than those registered at  $b_1$  (larvae with a weight range 8-10 mg) and  $b_2$  (larvae consisted of lighter and heavier ones in equal proportions) levels. No significant difference was observed between the levels  $b_1$  and  $b_2$ .

Among the levels of G (sex-ratio of parent parasite population) progeny production was found to be maximum at the  $c_1$  level (2 females: 1 male) with a mean 11.13. This was found to be not significantly different from the progeny production at  $c_0$  level (1 female: 1 male). The level  $c_4$  (1 female: 3 males) recorded very low number of progenies (6.13). The  $c_2$  and  $c_3$  levels were on par.

Progeny production at 32°C and 50% RH

The data pertaining to the mean progeny production under the various treatment combinations at the above temperature and humidity combination are furnished in column 5 of Table 3-a. The maximum number of progenies (64.33) was registered under the treatment  $a_1 b_0 c_1$  (two host larvae per female parasite; host larvae with a weight range of 30-35 mg, sex-ratio of parent population

two females: one male), while the treatment combination  $a_0 b_1 c_4$  (one host larva per female parasite; host larvae with a weight range 8-10 mg; sex-ratio of parent parasite population one female: three males) recorded the minimum number of progenies (4.33). The total as well as the mean number of progenies produced at different levels of the factors at 32°C and 50% RH are given in Table 3-c and these are diagrammatically represented in Fig. 3A, 3B and 3C. The results of the analysis employing the chi-square test are summarised in Table 3-b.

It will be seen that all the main effects viz., host density (A), weight of host larvae (B), and sex-ratio of parent parasite population (C) as well as their interactions  $A \times B$ ,  $A \times C$ ,  $B \times C$  and  $A \times B \times C$  are significant. The different levels of the above three factors are compared using the chi-square test and the results are presented in Table 3-d. It is evident that the maximum number of progenies (21.31) were produced with three host larvae per female ( $a_2$  level) and this was significantly higher than those produced at  $a_0$  (one host larva per female parasite) and  $a_1$  (two host larvae per female parasite) levels. No significant variation in progeny production was detected between the  $a_0$  and  $a_1$  levels.

Among the levels of B (weight of host larvae), the  $b_2$  level (larvae consisted of heavier and lighter

ones in equal proportions) registered maximum number of progenies (25.11). The  $b_2$  level recorded significantly higher number of progenies than the  $b_1$  level (larvae with a weight range 9-10 mg). The  $b_1$  level recorded a very low number of progenies (9.42). The  $b_2$  and  $b_0$  levels were *par*.

In relation to the levels of sex-ratio of parent parasite population (C), maximum progeny production of 15.80 was registered as the sex-ratio of the parent population was kept at 2 females: 1 male ( $c_1$ ). But as the parent population was maintained at 1 female : 3 males ( $c_4$ ) very low number of progenies were produced the mean being 7.56. The levels  $c_2$  (3 females : 1 male) and  $c_3$  (1 female : 2 males) have registered more number of progenies than the  $c_4$  level (1 female : 3 males).

Comparisons of progeny production between different levels of temperature and humidity conditions at different levels of the factors A, B and C are presented in Table 3-c. Heterogeneity chi-square is significant for the three factors viz., host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population indicating that the responses vary with the temperature and humidity conditions.



At the  $a_0$  level (one host larva per female parasite), the progeny production was maximum (17.36) at 32°C and 50% RH ( $TH_3$ ). This was significantly higher than those at 30°C and 60% RH ( $TH_2$ ) and 28°C and 75% RH ( $TH_1$ ). The same trend was observed at the  $a_1$  (2 hosts per female parasite) and  $a_2$  (three hosts per female parasite) levels also. At the three levels of host larval density (A), an increasing trend in the progeny production was observed with increasing temperature and decreasing humidity within the levels tested.

At the  $b_0$  level (host larval weight in the range 30 to 35 mg), maximum progeny production (13.76) was recorded at 32°C and 50% RH ( $TH_3$ ) which is significantly higher than those corresponding to 30°C and 60% RH ( $TH_2$ ) and 28°C and 75% RH ( $TH_1$ ). The progeny production was increased with increasing temperature and decreasing humidity. But at the  $b_1$  level (host larval weight in the range 8 to 10 mg), maximum progeny production was obtained at 30°C and 60% RH ( $TH_2$ ) which was significantly higher than those at 28°C and 75% RH ( $TH_1$ ) and 32°C and 50% RH ( $TH_3$ ). But no significant difference was detected between the counts of progeny at 28°C and 75% RH ( $TH_1$ ) and 32°C and 50% RH ( $TH_3$ ). The  $b_2$  level (a mixture of heavier and lighter larvae in equal proportions) revealed the same trend as that of  $b_0$ .

With respect to the progeny production at one female: one male ratio ( $e_0$ ) of the parent parasite population, the maximum (14.69) was recorded at 32°C and 50% RH ( $TH_3$ ). This is significantly higher than those corresponding to 30°C and 60% RH ( $TH_2$ ) and 28°C and 75% RH ( $TH_1$ ). There is a progressive increase in the progeny production with increasing temperature and decreasing humidity. The same trend was observed with two females: one male ( $e_1$ ) and three females: one male ( $e_2$ ) ratios of parent parasite population. But no significant difference was detected between the counts of progeny at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ) at  $e_2$  level. At  $e_3$  level (1 female: 2 males) maximum progenies were produced at 32°C and 50% RH ( $TH_3$ ) which is significantly higher than those at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ). But no significant difference was detected between the number of progenies recorded at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ). The same trend was observed at  $e_4$  level also (1 female: 3 males).

#### IV. NUMBER OF FEMALE PROGENY

Number of female progeny at 28°C and 75% RH

The total number of females which emerged out from each replicate was recorded under different treatment

combinations. Data on the mean number of females produced under various treatments at 28°C and 75% RH are furnished in column 3 of Table 4-a and the raw data given in Appendix-IV. The mean number of females ranged from 1.00 under the treatment  $a_0, b_1, c_1$  (one host larva per female parasite; weight of host larvae 8-10 mg; sex-ratio of parent population - one female: two males) to 14.00 under the treatment  $a_0, b_0, c_1$  (one host larva per female parasite; larvae with a weight of 30-35 mg and 8-10 mg in equal proportions; sex-ratio of parent population one female: two males).

The data being in the form of counts, chi-square test has been adopted to analyse the data as outlined in materials and methods and the results are presented in Table 4-b. It will be seen that all the main effects viz., host density (A); weight of host larvae (B) and sex-ratio of parent parasite population (C) as well as their interactions are significant.

The total as well as the mean values of the number of females produced by the different levels of the factors, host density (A); weight of host larvae (B) and sex-ratio of parent parasite population (C) are shown in Table 4-c and these are graphically depicted in Fig. 4A, 4B and 4C. The levels of the above factors are compared in Table 4-d. One host larva per female parasite ( $a_0$ ) produced the

maximum number of female progenies, the mean being 5.58. This level is not significantly different from the number of progenies produced corresponding to  $a_1$  level (two host larvae per female parasite). But when there were three host larvae per female parasite ( $a_2$ ), the number of females was significantly lower than those corresponding to one host larva per female ( $a_0$ ) and two host larvae per female parasite ( $a_1$ ).

Among the three levels of the weight of the host larvae (B), larvae consisting of 50% of heavier ones with 30-35 mg weight and the remaining 50% with 8-10 mg weight ( $b_2$ ) produced maximum number of females, with a mean value of 5.51. This was followed by larvae with a weight range of 30-35 mg ( $b_0$ ) and the two levels  $b_0$  and  $b_2$  being statistically on par. Larvae with a mean weight of 8-10 mg ( $b_1$ ) produced significantly lower counts of females than those with the  $b_0$  and  $b_2$  levels.

Among the levels of sex-ratio of parent parasite population (C), parent population with a ratio of two females: one male ( $c_1$ ) have produced the maximum number of female progeny, the mean being 3.64. The number of females at this level ( $c_1$ ) is significantly higher than those obtained with one female:one male ( $c_0$ ) and one female: three males ( $c_2$ ) ratio of the parent population.

Table 4-a

Number of female progeny of *B. brevicornis* in different temperature and humidity combinations

Sl. No.	Treatments	Mean number of females		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
1.	a <sub>0</sub> b <sub>0</sub> c <sub>0</sub>	6.33	7.33	15.00
2.	a <sub>0</sub> b <sub>0</sub> c <sub>1</sub>	7.00	8.00	17.00
3.	a <sub>0</sub> b <sub>0</sub> c <sub>2</sub>	2.33	1.67	3.00
4.	a <sub>0</sub> b <sub>0</sub> c <sub>3</sub>	12.00	14.67	10.00
5.	a <sub>0</sub> b <sub>0</sub> c <sub>4</sub>	2.67	5.33	6.00
6.	a <sub>0</sub> b <sub>1</sub> c <sub>0</sub>	2.00	2.67	7.33
7.	a <sub>0</sub> b <sub>1</sub> c <sub>1</sub>	1.00	9.00	4.00
8.	a <sub>0</sub> b <sub>1</sub> c <sub>2</sub>	4.00	9.33	3.00
9.	a <sub>0</sub> b <sub>1</sub> c <sub>3</sub>	4.33	3.00	3.33
10.	a <sub>0</sub> b <sub>1</sub> c <sub>4</sub>	3.33	2.33	2.00
11.	a <sub>0</sub> b <sub>2</sub> c <sub>0</sub>	3.00	4.00	18.00
12.	a <sub>0</sub> b <sub>2</sub> c <sub>1</sub>	14.00	2.33	7.00
13.	a <sub>0</sub> b <sub>2</sub> c <sub>2</sub>	7.33	4.00	17.67
14.	a <sub>0</sub> b <sub>2</sub> c <sub>3</sub>	6.33	3.33	8.67
15.	a <sub>0</sub> b <sub>2</sub> c <sub>4</sub>	8.00	2.67	9.00
16.	a <sub>1</sub> b <sub>0</sub> c <sub>0</sub>	7.67	9.33	26.67
17.	a <sub>1</sub> b <sub>0</sub> c <sub>1</sub>	5.67	9.00	40.00
18.	a <sub>1</sub> b <sub>0</sub> c <sub>2</sub>	8.00	3.00	9.33
19.	a <sub>1</sub> b <sub>0</sub> c <sub>3</sub>	3.67	2.33	5.67
20.	a <sub>1</sub> b <sub>0</sub> c <sub>4</sub>	3.00	8.67	5.00
21.	a <sub>1</sub> b <sub>1</sub> c <sub>0</sub>	7.33	6.67	2.00
22.	a <sub>1</sub> b <sub>1</sub> c <sub>1</sub>	8.33	12.00	5.67
23.	a <sub>1</sub> b <sub>1</sub> c <sub>2</sub>	5.00	2.33	2.33
24.	a <sub>1</sub> b <sub>1</sub> c <sub>3</sub>	3.00	4.00	5.33
25.	a <sub>1</sub> b <sub>1</sub> c <sub>4</sub>	6.00	2.22	2.22
26.	a <sub>1</sub> b <sub>2</sub> c <sub>0</sub>	5.00	10.33	7.00
27.	a <sub>1</sub> b <sub>2</sub> c <sub>1</sub>	3.00	8.67	14.33
28.	a <sub>1</sub> b <sub>2</sub> c <sub>2</sub>	6.00	8.00	3.33
29.	a <sub>1</sub> b <sub>2</sub> c <sub>3</sub>	4.33	8.67	7.00
30.	a <sub>1</sub> b <sub>2</sub> c <sub>4</sub>	6.67	5.00	8.33
31.	a <sub>2</sub> b <sub>0</sub> c <sub>0</sub>	4.00	8.00	13.00
32.	a <sub>2</sub> b <sub>0</sub> c <sub>1</sub>	2.67	10.00	6.67
33.	a <sub>2</sub> b <sub>0</sub> c <sub>2</sub>	4.33	6.67	5.00
34.	a <sub>2</sub> b <sub>0</sub> c <sub>3</sub>	7.33	6.00	16.00
35.	a <sub>2</sub> b <sub>0</sub> c <sub>4</sub>	4.33	3.00	6.00
36.	a <sub>2</sub> b <sub>1</sub> c <sub>0</sub>	2.00	9.67	5.33
37.	a <sub>2</sub> b <sub>1</sub> c <sub>1</sub>	6.00	5.33	3.67
38.	a <sub>2</sub> b <sub>1</sub> c <sub>2</sub>	6.33	4.33	4.33
39.	a <sub>2</sub> b <sub>1</sub> c <sub>3</sub>	3.67	3.33	2.67
40.	a <sub>2</sub> b <sub>1</sub> c <sub>4</sub>	2.67	5.33	9.33
41.	a <sub>2</sub> b <sub>2</sub> c <sub>0</sub>	5.33	4.67	13.67
42.	a <sub>2</sub> b <sub>2</sub> c <sub>1</sub>	7.00	5.00	13.00
43.	a <sub>2</sub> b <sub>2</sub> c <sub>2</sub>	2.67	3.33	18.33
44.	a <sub>2</sub> b <sub>2</sub> c <sub>3</sub>	2.33	5.00	9.67
45.	a <sub>2</sub> b <sub>2</sub> c <sub>4</sub>	1.67	9.33	11.00

a<sub>0</sub> - 1 host larva per female parasitea<sub>1</sub> - 2 " " " "a<sub>2</sub> - 3 " " " "b<sub>0</sub> - larval weight 30 - 35 mgb<sub>1</sub> - " " " 8 - 10 mgb<sub>2</sub> - half of the larvae of the 'b' type

Table 4-b

Chi-square values for the effects of the three factors A, B and C at different temperature and humidity combinations on the number of female progeny in D. brevicornis.

Source	df	Chi-square		
		28°C and 75% RH	30°C and 60% RH	32°C and 50% RH
⊙ A	2	11.41**	8.51*	3.49 NS
⊙ B	2	7.49*	16.37**	180.13**
A x B	4	16.63**	24.78**	83.51**
⊙ C	4	9.58*	44.76**	86.92**
A x C	8	27.05**	24.46**	104.66**
B x C	8	27.51**	35.34**	131.74**
A x B x C	16	82.84**	110.13**	149.45**

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

⊙ A - host larval density  
 ⊙ B - weight of host larvae  
 ⊙ C - sex-ratio of parent parasite population

Table 4-c

Total and mean number of female progeny of *B. brevicornis* produced at different levels of the main factors at different temperature and humidity combinations

	28°C and 75% RH		30°C and 60% RH		32°C and 50% RH	
	Total	Mean	Total	Mean	Total	Mean
$a_0^*$	251	5.58	239	5.31	393	8.73
$a_1$	248	5.51	304	6.76	445	9.89
$a_2$	187	4.16	291	6.47	407	9.04
$b_0^*$	243	5.40	333	7.40	553	12.29
$b_1$	195	4.33	248	5.51	194	4.31
$b_2$	248	5.51	53	5.62	498	11.07
$c_0^*$	128	2.84	188	4.18	324	7.20
$c_1$	164	3.64	232	5.16	334	7.42
$c_2$	136	3.02	128	2.84	199	4.42
$c_3$	141	3.13	151	3.36	205	4.56
$c_4$	115	2.56	135	3.00	183	4.07

$a_0^*$  - 1 host larva per female parasite  
 $a_1$  - 2 " " " "  
 $a_2$  - 3 " " " "

$b_0^*$  - larval weight 30 - 35 mg  
 $b_1$  - " " " 8 - 10 mg  
 $b_2$  - half of the hosts of the 'b' type and other half of the 'b<sub>1</sub>' type

$c_0^*$  - Sex-ratio of parent parasite population - 1:1 (Female:Male)  
 $c_1$  - " " " " - 2:1 " "  
 $c_2$  - " " " " - 3:1 " "  
 $c_3$  - " " " " - 1:2 " "  
 $c_4$  - " " " " - 1:3 " "

**Fig. 4 A -** Number of female progeny of Bracon bravicornis produced at different temperature-humidity combinations as influenced by the host larval density

**Fig. 4 B -** Number of female progeny of Bracon bravicornis produced at different temperature-humidity combinations as influenced by the weight (size) of host larvae

**Fig. 4 C -** Number of female progeny of Bracon bravicornis produced at different temperature-humidity combinations as influenced by the sex-ratio of the parent parasite population



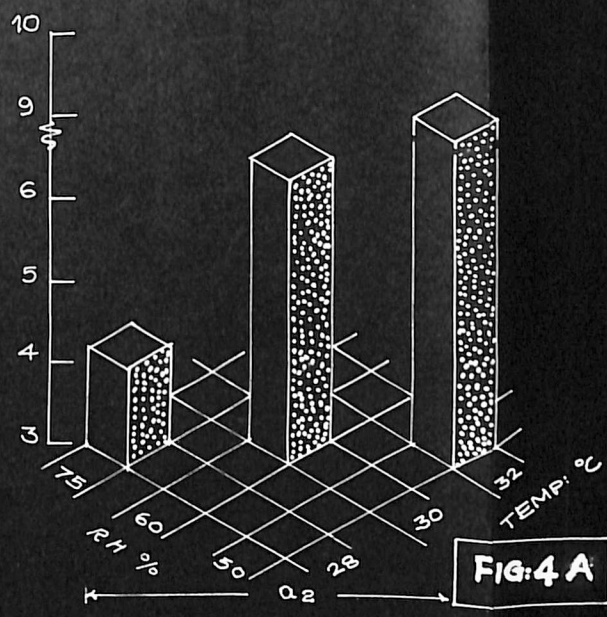
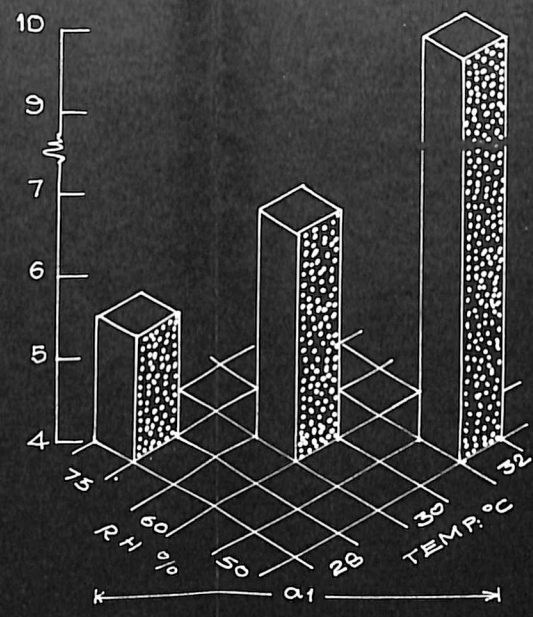
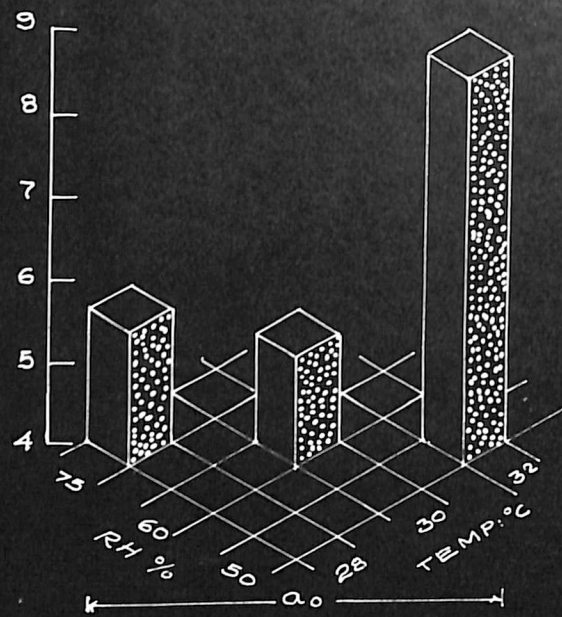


FIG:4 A



1 HOST LARVA PER FEMALE PARASITE  
 2 " " " " " "  
 3 " " " " " "

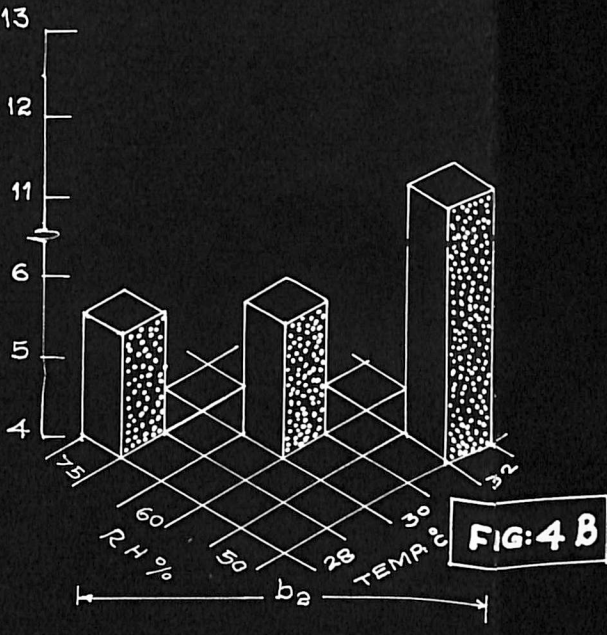
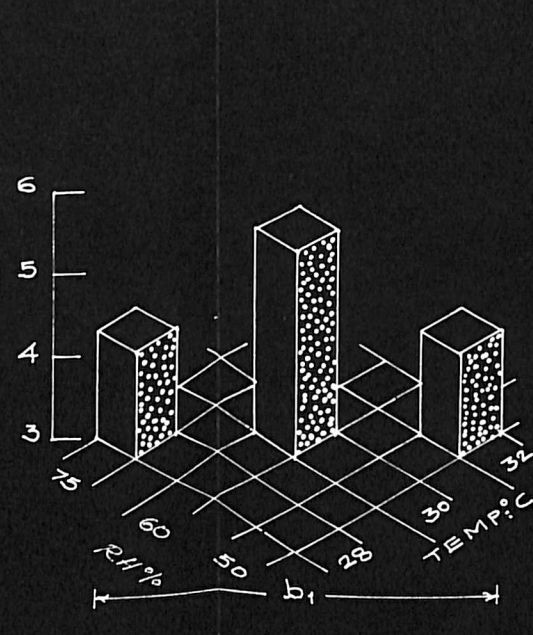
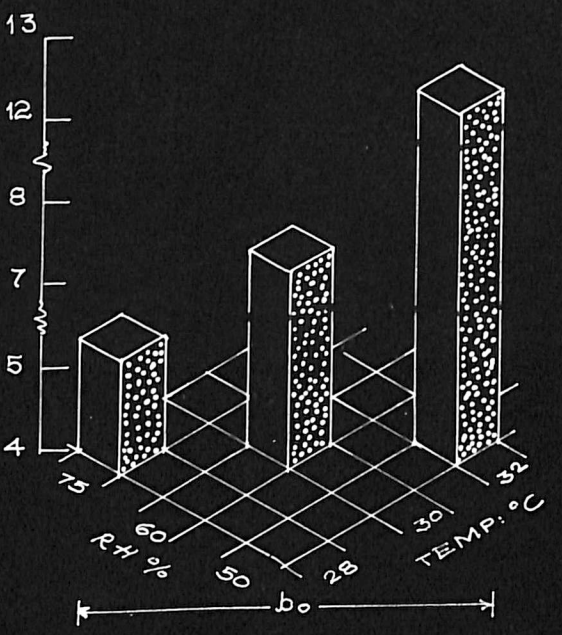
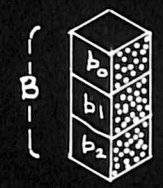
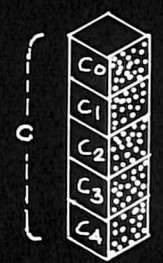


FIG:4 B



LARVAL WEIGHT 30-35 mg.  
 " " " " " " 8-10 "  
 HALF OF THE LARVAE OF THE 'b0' TYPE  
 THE OTHER HALF OF THE 'b1' CATEGOR



1 FEMALE : 1 MALE  
 2 " : 1 "  
 3 " : 1 "  
 1 " : 2 "  
 1 " : 3 "

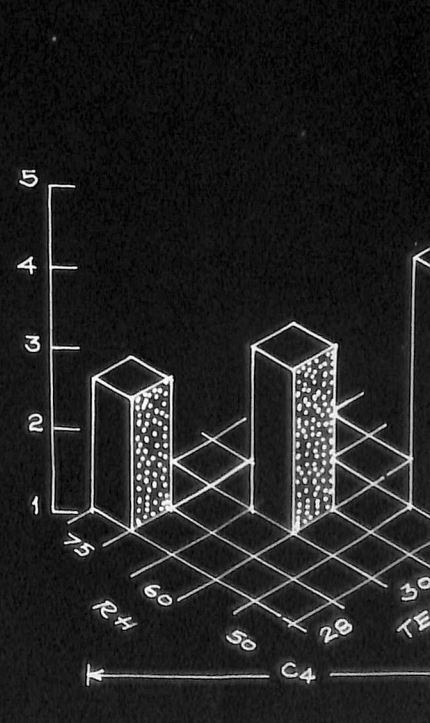
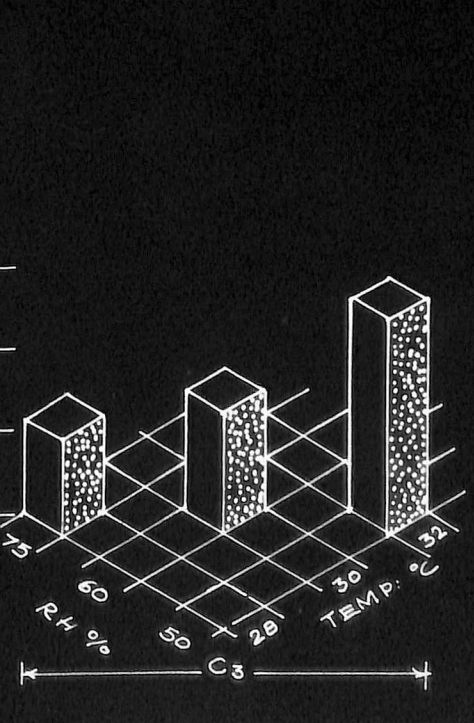
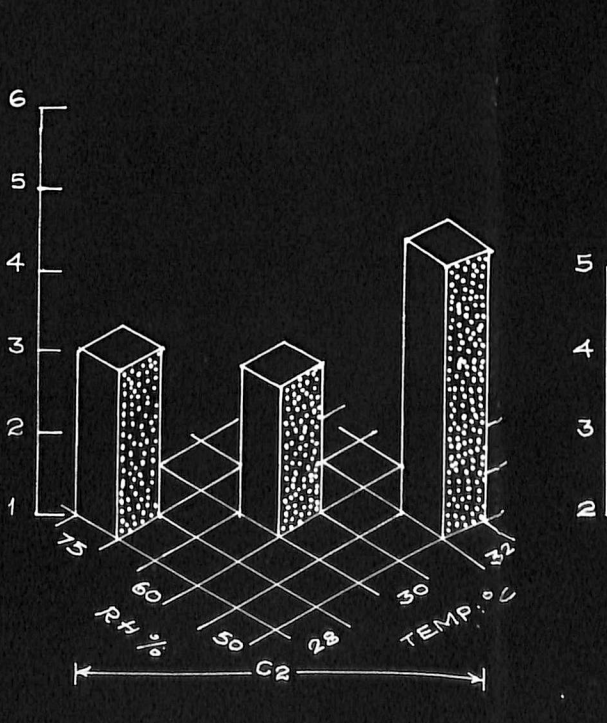
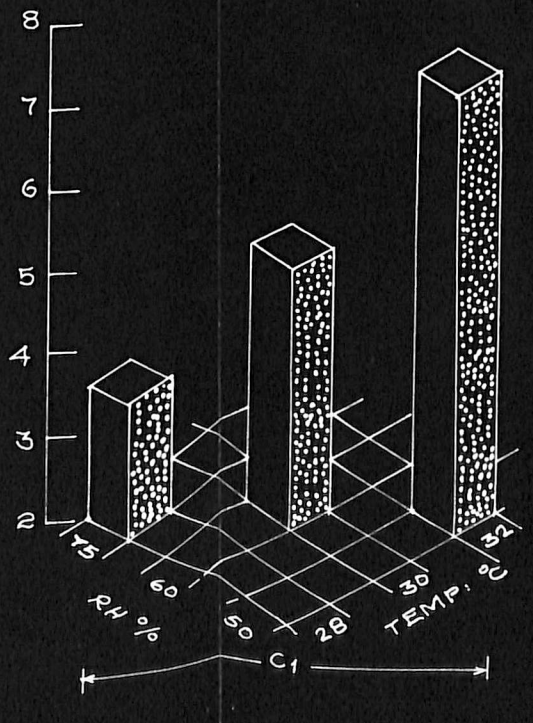
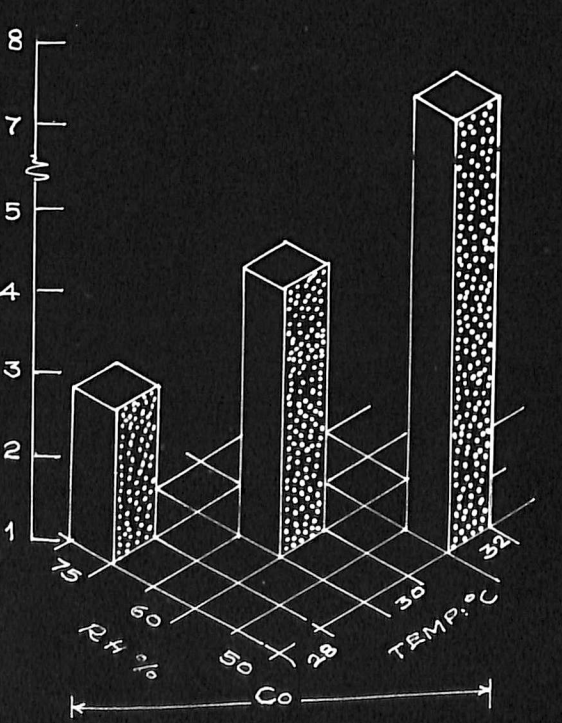


Table 4-d

Chi-square for comparing the effects of different levels of the three factors A, B and C at different temperature and humidity combinations on the number of female progeny of *B. brachycephala*

		Chi-square values					
		20°C and 75% RH	Rank- ing	30°C and 60% RH	Rank- ing	30°C and 50% RH	Rank- ing
a <sub>0</sub>	Vs a <sub>1</sub>	0.02 NS	10000	7.78**	10000	3.23 NS	
a <sub>1</sub>	Vs a <sub>2</sub>	8.55**		0.28 NS		1.69 NS	
a <sub>0</sub>	Vs a <sub>2</sub>	9.35**		5.10*		0.25 NS	
b <sub>0</sub>	Vs b <sub>1</sub>	5.26**	10000	12.44**	10000	172.53**	10000
b <sub>1</sub>	Vs b <sub>2</sub>	6.34*		0.05 NS		133.55**	
b <sub>0</sub>	Vs b <sub>2</sub>	0.05 NS		10.92**		0.88 NS	
c <sub>0</sub>	Vs c <sub>1</sub>	4.44*	10000	4.61*	10000	0.15 NS	10000
c <sub>0</sub>	Vs c <sub>2</sub>	0.24 NS		11.39**		29.88**	
c <sub>0</sub>	Vs c <sub>3</sub>	0.63 NS		4.04*		26.77**	
c <sub>0</sub>	Vs c <sub>4</sub>	0.70 NS		8.70**		39.21**	
c <sub>1</sub>	Vs c <sub>2</sub>	2.61 NS		30.04**		34.19**	
c <sub>1</sub>	Vs c <sub>3</sub>	1.73 NS		17.13**		49.09**	
c <sub>1</sub>	Vs c <sub>4</sub>	8.61**		25.64**		44.10**	
c <sub>2</sub>	Vs c <sub>3</sub>	0.09 NS		1.90 NS		0.09 NS	
c <sub>2</sub>	Vs c <sub>4</sub>	1.76 NS		0.19 NS		0.67 NS	
c <sub>3</sub>	Vs c <sub>4</sub>	2.64 NS		0.90 NS		1.25 NS	

\*Significant at 5% level  
 \*\*Significant at 1% level  
 NS Not significant

a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type  
 and the other half of the 'b<sub>1</sub>' type

c<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female: Male)  
 c<sub>1</sub> - " " " " - 2:1 " "  
 c<sub>2</sub> - " " " " - 3:1 " "  
 c<sub>3</sub> - " " " " - 1:2 " "  
 c<sub>4</sub> - " " " " - 1:3 " "

Table 4-9

Chi-square values for the comparison of the effects of the three main factors A, B and C on the number of female progeny in *B. brevicornis* at different temperature and humidity combinations.

	Chi-square for levels of A <sup>ⓐ</sup>			Chi-square for levels of B <sup>ⓑ</sup>			Chi-square for levels of C <sup>ⓒ</sup>				
	a <sub>0</sub>	a <sub>1</sub>	a <sub>2</sub>	b <sub>0</sub>	b <sub>1</sub>	b <sub>2</sub>	c <sub>0</sub>	c <sub>1</sub>	c <sub>2</sub>	c <sub>3</sub>	c <sub>4</sub>
	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>2</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>	TH <sub>1</sub> Vs TH <sub>3</sub>	TH <sub>2</sub> Vs TH <sub>3</sub>
TH <sub>1</sub> Vs TH <sub>2</sub>	0.29 NS	5.68*	22.63**	14.06**	6.54*	0.05 NS	11.39**	11.39**	0.24 NS	0.34 NS	1.6 NS
TH <sub>2</sub> Vs TH <sub>3</sub>	37.53**	26.54**	19.28**	54.63**	6.60*	79.93**	36.13**	18.38**	15.42**	8.19**	7.25**
TH <sub>1</sub> Vs TH <sub>3</sub>	31.31**	56.00**	81.58**	120.73**	0.002 NS	83.78**	84.99**	58.03**	11.85**	11.84**	15.52**
Ranking	TH <sub>3</sub>   TH <sub>1</sub>   TH <sub>2</sub>	TH <sub>3</sub>   TH <sub>2</sub>   TH <sub>1</sub>	TH <sub>3</sub>   TH <sub>2</sub>   TH <sub>1</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>	TH <sub>1</sub>   TH <sub>2</sub>   TH <sub>3</sub>

Chi-square due to heterogeneity

Host larval density (A) 14.17\*\*  
 Weight of host larvae (B) 64.03\*\*  
 Sex-ratio of parent parasite population (C) 24.48\*\*

ⓐ - Host larval density  
 a<sub>0</sub> - 1 larva per female parasite  
 a<sub>1</sub> - 2 " " "  
 a<sub>2</sub> - 3 " " "

ⓑ - Weight of host larvae  
 b<sub>0</sub> - larval weight 30 - 35 mg.  
 b<sub>1</sub> - " " " 8 - 10 mg.  
 b<sub>2</sub> - half of the hosts of the b<sub>0</sub> type and the other half of the b<sub>1</sub> type

ⓒ - Sex-ratio of parent population  
 c<sub>0</sub> - 1 female : 1 male  
 c<sub>1</sub> - 2 " : 1 "  
 c<sub>2</sub> - 3 " : 1 "  
 c<sub>3</sub> - 1 " : 2 "  
 c<sub>4</sub> - 1 " : 3 "

\* Significant at 5% level  
 \*\* Significant at 1% level  
 NS Not significant

TH<sub>1</sub> - 28°C and 75% RH  
 TH<sub>2</sub> - 30°C and 60% RH  
 TH<sub>3</sub> - 32°C and 50% RH

But no significant difference was detected between 2:1 ( $e_1$ ) and 3:1 ( $e_2$ ) ratios of the parent population; and also between the 2:1 level ( $e_1$ ) and the 1:2 level ( $e_3$ ). The number of females produced was minimum when the sex-ratio of parent population was 1:3, the mean being 2.56.

#### Number of female progeny at 30°C and 60% RH

Treatment-wise data on the mean number of females produced at the above temperature and humidity condition is furnished in column 4 of Table 4-a and the raw data in Appendix IV. The number of female progeny varied from 1.67 under the treatment  $a_0 b_0 c_2$  (one host larva per female parasite; weight of host larva in the range 30 to 35 mg; sex-ratio of parent parasite population 3 female: 1 male) to 18.00 under the treatment  $a_2 b_0 c_1$  (two host larvae per female parasite; weight of host larvae in the range 30 to 35 mg; sex-ratio of parent parasite population 2 females:1male). The results of analysis of the data using chi-square test is presented in Table 4-b. It is indicated from the Table that host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population (C) have significant effects on the number of female progeny. The interactions of the factors are also significant. The total

and mean values of the number of female progeny produced are presented in Table 4-c and these are represented in Fig. 4A, 4B and 4C. Comparisons of the different levels of the factors A, B and C by chi-square method is given in Table 4-d.

Among the levels of the host larval density (A), the maximum female progeny was produced when two host larvae were provided per female per site ( $a_1$  level) with mean number 6.76. But at the  $a_2$  level of three host larvae per female parasite, there was no significant difference from the  $a_1$  level. Minimum number of female progeny was produced when one host larva was provided per female parasite ( $a_0$  level), the mean being 5.31.

Among the levels of the weight of host larvae (B),  $b_0$  level i.e., larvae with a weight range of 30 to 35 mg produced maximum number of female progeny having a mean of 7.40. This is significantly higher than those corresponding to  $b_2$  (a mixture of heavier ( $b_0$ ) and lighter ( $b_1$ ) larvae in equal proportions) and  $b_1$  levels (larvae with a weight range 8 to 10 mg). Larvae having a weight of 8 to 10 mg ( $b_1$ ) produced the minimum number of 5.51 female progeny.

Among the levels of the sex-ratio of parent parasite population (C), the ratio 2 females : 1 male ( $c_1$ )

gave the maximum number of females (5.16) this being significantly higher than those obtained at the other levels. With 3 females: 1 male ( $c_2$ ) in the parent population, minimum number of females were obtained. No significant difference was detected between 1 female : 2 males ( $c_3$ ) and 1 female: 3 males ( $c_4$ ) ratios of the parent population.

Number of female progeny at 32°C and 50% RH

Data on the mean number of females produced under various treatments are furnished in column 5 of Table 4-a. Minimum of 2.00 females were recorded by the treatments  $a_0$ ,  $b_1$ ,  $c_4$  (one host larva per female parasite; weight of host larvae in the range 8 to 10 mg; sex-ratio of parent population 1 female: 3 males) and  $a_1$ ,  $b_1$ ,  $c_0$  (two host larvae per female parasite; weight of host larvae in the range 8 to 10 mg; sex-ratio of parent population 1 female: 1 male) while maximum number of 40.00 females were recorded by the treatment  $a_1$ ,  $b_0$ ,  $c_1$  (two host larvae per female parasite; weight of host larvae in the range 30 to 35 mg; sex-ratio of parent population 2 females : 1 male).

Analysis of the data by the chi-square test is given in Table 4-b. The effects of weight of host larvae (B), sex-ratio of parent parasite population (C) and the

interactions  $A \times B$ ,  $A \times C$ ,  $B \times C$  and  $A \times B \times C$  are found to be significant whereas host larval density (A) had no significant effect on the number of female progeny. The total and mean number of females produced at different levels of the factors A, B and C are expressed in Table 4-c and these are illustrated in Fig. 4A, 4B and 4C. Comparisons of the different levels of the factors are carried out by the chi-square test which are presented in Table 4-d.

Among the levels of the weight of host larvae (B), maximum number of females (12.29) were produced with host larvae of weight range 30 to 35 mg ( $b_0$  level). Host larvae having a weight range of 8 to 10 mg ( $b_1$ ) produced very low number (4.31) of females. No significant difference was detected between  $b_0$  (larval weight range 30 to 35 mg) and  $b_2$  levels (50% of 30 to 35 mg type + 50% of 8 to 10 mg type).

Among the levels of the sex-ratio of parent parasite population (C), 2 females: 1 male ratio ( $c_1$ ) produced maximum number of 7.42 females. Parent population in the ratio 1 female : 3 males ( $c_4$ ) produced the minimum number of 4.07 female progeny. No significant variation was observed between parent parasite populations with

1 female: 1 male ( $a_0$ ) and 2 females : 1 male ( $a_1$ ) sex-ratios. Parent parasite population with a sex-ratio of 3 females: 1 male ( $a_2$ ) is not found to be significantly different from those with 1 female : 2 males ( $a_3$ ) and 1 female : 3 males ( $a_4$ ) sex-ratios with reference to the number of females produced. But parent population with 1 female : 2 males ( $a_3$ ) and 1 female : 3 males ( $a_4$ ) are statistically on par.

Comparisons between different levels of temperature and humidity combinations at different levels of the factors A, B and C are given in Table 4-e. The heterogeneity chi-square for A, B and C are significant indicating that the responses vary with temperature and humidity conditions. The number of females produced at  $a_0$  level (one host per female parasite) was maximum at 32°C and 50% RH ( $TH_3$ ). This is significantly higher than those at 29°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ). But no significant difference was detected between the counts of females at 29°C and 75% RH ( $TH_1$ ) and at 30°C and 60% RH ( $TH_2$ ).

When two hosts per female parasite were given ( $a_1$ ) the maximum number of female progeny was again recorded at 32°C and 50% RH ( $TH_3$ ). This is significantly



higher than those at 30°C and 50% RH (TH<sub>2</sub>) and 28°C and 75% RH (TH<sub>1</sub>). Thus when the host larval density is two per female parasite (a<sub>1</sub>) there is a progressive increase in the number of females with increasing temperature and decreasing humidity. The same trend was observed when there were three hosts per female parasite (a<sub>2</sub>).

There is an increasing trend of female progeny production with increasing temperature and decreasing humidity when the host larvae had a mean weight of 30 to 35 mg (b<sub>0</sub>), all the differences between the three temperature and humidity combinations being significant. But when lighter larvae with a weight range of 8 to 10 mg (b<sub>1</sub>) were given, the maximum number of female progeny was recorded at 30°C and 60% RH (TH<sub>2</sub>), this being significantly higher than those at 28°C and 75% RH (TH<sub>1</sub>) and 32°C and 50% RH (TH<sub>3</sub>). No significant difference in the female progeny number was detected between 28°C and 75% RH (TH<sub>1</sub>) and 32°C and 50% RH (TH<sub>3</sub>). Corresponding to b<sub>2</sub> level (a mixture of heavier and lighter larvae in equal proportions), there is an increase in the number of females with increasing temperature and decreasing humidity levels, the difference between the counts of females at 30°C and 60% RH (TH<sub>2</sub>) and 28°C and 75% RH (TH<sub>1</sub>) being not significant.

An increasing trend was observed in the number of

females produced with increasing temperature and decreasing humidity level when there was 1 female : 1 male ratio ( $c_0$ ) for the parent parasite population. The same trend was observed when the sex-ratio of parent parasite population was 2 females: 1 male ( $c_1$ ). But when the sex-ratio of parent population is 3 females: 1 male ( $c_2$ ) minimum number of females were produced at 30°C and 60% RH ( $TH_2$ ). At  $c_3$  (1 female : 2 males) and  $c_4$  (1 female : 3 males) levels, the maximum number of females were produced at 32°C and 50% RH ( $TH_3$ ). This was significantly higher than those produced both at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ). But no significant difference was detected between the counts of females at 28°C and 75% RH ( $TH_1$ ) and 30°C and 60% RH ( $TH_2$ ) both at  $c_3$  and  $c_4$  levels.

## V. FEMALE-MALE COMPOSITION

The sex-wise counts of the parasites under different treatment combinations have been analysed for the following aspects employing the techniques outlined below.

A. Testing the significance of the difference between the levels of the main factors in the expression of the female-male composition of the  $F_1$  progeny by

transforming the ratio  $\frac{\text{Number of females}}{\text{Total}}$  as angles and by testing the differences between these angles in pairs by applying the formula

$$\frac{\theta_1 - \theta_2}{\sqrt{820.7 \left( \frac{1}{n_1} + \frac{1}{n_2} \right)}} \quad \text{where } \frac{820.7}{n_i}$$

is the variance of  $\theta_i$

B. Testing the association between the main factors and sex-wise classification of the  $F_1$  progeny, the chi-square values were computed on the basis of the formula  $\frac{(an_2 - a'n_1)^2}{(a+a')n_1n_2}$  where  $a$  and  $a'$  denote the sex-wise frequencies corresponding to the levels, and  $n_1$  and  $n_2$  represent the corresponding total counts in each case.

Data on the mean number of females and males recorded at the three temperature-humidity combinations under various treatments are furnished in Table 5-a and the replication-wise data is given in Appendix V. The mean number of females and males produced at the different levels of the three main factors are graphically depicted in Fig. 5A, 5B and 5C. The female-male composition of the  $F_1$  progeny of B. brevicornis under different levels of



the main factors A, B and C under the three temperature-humidity combinations are furnished in Tables 5-b, 5-c and 5-d.

**Female-Male composition at 28°C and 75% RH**

It is indicated that out of the three factors tested, the weight of host larvae (B) alone has shown significant influence on the female-male composition of  $F_1$  progeny at 28°C and 75% RH (Table 5-b). The  $b_2$  level has recorded the highest percentage of females (52.32). Significant difference in the sex-ratio was detected between the  $b_1$  (host larval weight 8-10 mg) and the  $b_2$  (heavier and lighter ones in equal proportions) levels.

**Female-Male composition at 30°C and 60% RH**

At this temperature and humidity combination, the sex-ratio of the parent parasite population (C) alone had significant effect on the female-male composition of the  $F_1$  progeny, while host larval density (A) and host larval weight (B) had no significant effect (Table 5-c). The  $c_4$  level (1 female : 3 males) recorded the highest percentage of females (48.91) while the  $c_0$  level (1 female : 1 male) recorded a low percentage of females (39.00). The sex-ratio recorded at the  $c_4$  level was significantly better than those at  $c_2$  and  $c_0$  but the sex-ratios at the  $c_2$  and  $c_0$  levels were on par.



Table 5-b

Female-Male composition of *B. brevicornis* under different levels of the main factors A, B and C at 28°C and 75% RH

Levels	Females	Males	Chi-square
⊙ a <sub>0</sub>	251 (49.70)	254 (50.30)	
a <sub>1</sub>	248 (44.80)	306 (55.20)	3.04 NS
a <sub>2</sub>	187 (45.20)	227 (54.80)	
⊙ b <sub>0</sub>	243 (44.60)	302 (55.40)	
b <sub>1</sub>	195 (42.95)	259 (57.05)	9.55**
b <sub>2</sub>	248 (52.32)	226 (47.68)	
⊙ c <sub>0</sub>	128 (43.54)	166 (56.46)	
c <sub>1</sub>	164 (34.10)	317 (65.90)	6.54 NS
c <sub>2</sub>	136 (46.26)	158 (53.74)	
c <sub>3</sub>	141 (42.86)	188 (57.14)	
c <sub>4</sub>	115 (48.12)	124 (51.88)	

Figures in parenthesis indicate the related percentage values.

\*\* Significant at 1% level

NS Not significant

⊙ a - 1 host larva per female parasite  
 a<sub>0</sub> - 2 " " " "  
 a<sub>1</sub> - 3 " " " "

⊙ b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the host larvae of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

⊙ c - sex-ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>0</sub> - " " " " 2:1 { " }  
 c<sub>1</sub> - " " " " 3:1 { " }  
 c<sub>2</sub> - " " " " 1:2 { " }  
 c<sub>3</sub> - " " " " 1:3 { " }  
 c<sub>4</sub> - " " " " " { " }

Table 5-c

Female-Male composition of *B. brevicornis* under different levels of the main factors A, B and C at 30°C and 60% RH

Levels	Females	Males	Chi-square
⊙ a <sub>0</sub>	239 (41.57)	336 (58.43)	
a <sub>1</sub>	304 (45.85)	359 (54.15)	2.32 NS
a <sub>2</sub>	291 (44.22)	367 (55.78)	
⊙ b <sub>0</sub>	333 (46.90)	377 (53.10)	
b <sub>1</sub>	248 (44.21)	313 (55.79)	5.58 NS
b <sub>2</sub>	253 (40.48)	372 (59.52)	
⊙ c <sub>0</sub>	188 (39.00)	294 (61.00)	
c <sub>1</sub>	232 (46.31)	269 (53.69)	17.44**
c <sub>2</sub>	128 (39.75)	194 (60.25)	
c <sub>3</sub>	151 (47.94)	164 (52.06)	
c <sub>4</sub>	135 (48.91)	141 (51.09)	

Figures in parenthesis indicate the related percentage values.

\*\* Significant at 1% level

NS Not significant

⊙ a - 1 host larva per female parasite  
 a<sub>0</sub> - 2 " " " "  
 a<sub>1</sub> - 3 " " " "

⊙ b - larval weight 30 - 35 mg  
 b<sub>0</sub> - " " 8 - 10 mg  
 b<sub>1</sub> - half of the host larvae of the 'b' type  
 b<sub>2</sub> - and the other half of the 'b<sub>1</sub>' type.

⊙ c - sex-ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>0</sub> - " " " " 2:1  
 c<sub>1</sub> - " " " " 3:1  
 c<sub>2</sub> - " " " " 1:2  
 c<sub>3</sub> - " " " " 1:3  
 c<sub>4</sub> - " " " " }

Table 5-d

Female-Male composition of *B. brevicornis* under different levels of the main factors A, B and C at 32°C and 50% RH

Levels	Females	Males	Chi-square
⊙ a <sub>0</sub>	393 (50.32)	398 (49.68)	
a <sub>1</sub>	445 (53.10)	393 (46.90)	32.56**
a <sub>2</sub>	407 (40.54)	597 (59.46)	
⊙ b <sub>0</sub>	553 (51.73)	516 (48.27)	
b <sub>1</sub>	194 (45.75)	230 (54.25)	13.52**
b <sub>2</sub>	498 (44.07)	632 (55.93)	
⊙ c <sub>0</sub>	324 (49.02)	337 (50.98)	
c <sub>1</sub>	334 (46.98)	377 (53.02)	13.95**
c <sub>2</sub>	199 (42.80)	266 (57.20)	
c <sub>3</sub>	205 (45.05)	250 (54.95)	
c <sub>4</sub>	183 (56.13)	143 (43.87)	

Figures in parenthesis indicate the related percentage values.

\*\* Significant at 1% level

NS Not significant

⊙ a<sub>0</sub> - 1 host larva per female parasite

a<sub>1</sub> - 2 " " " "

a<sub>2</sub> - 3 " " " "

⊙ b<sub>0</sub> - larval weight 30 - 35 mg

b<sub>1</sub> - " " 8 - 10 mg

b<sub>2</sub> - half of the host larvae of the 'b<sub>0</sub>' type

and the other half of the 'b<sub>1</sub>' type.

⊙ c<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)

c<sub>1</sub> - " " " " 2:1 ( " )

c<sub>2</sub> - " " " " 3:1 ( " )

c<sub>3</sub> - " " " " 1:2 ( " )

c<sub>4</sub> - " " " " 1:3 ( " )



**Fig. 5 A - Female-male composition of the F<sub>1</sub> progeny of Bracon brevicornis at different temperature-humidity combinations as influenced by the host larval density**

**Fig. 5 B - Female-male composition of the F<sub>1</sub> progeny of Bracon brevicornis at different temperature-humidity combinations as influenced by the weight (size) of host larvae**

**Fig. 5 C - Female-male composition of the F<sub>1</sub> progeny of Bracon brevicornis at different temperature-humidity combinations as influenced by the sex-ratio of the parent parasite population**

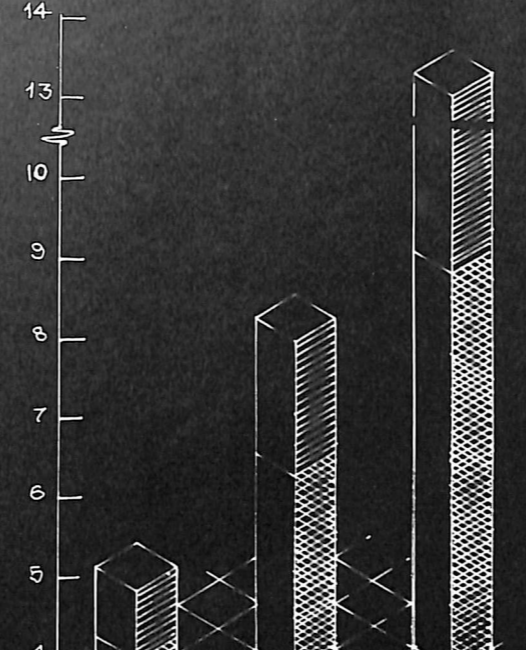
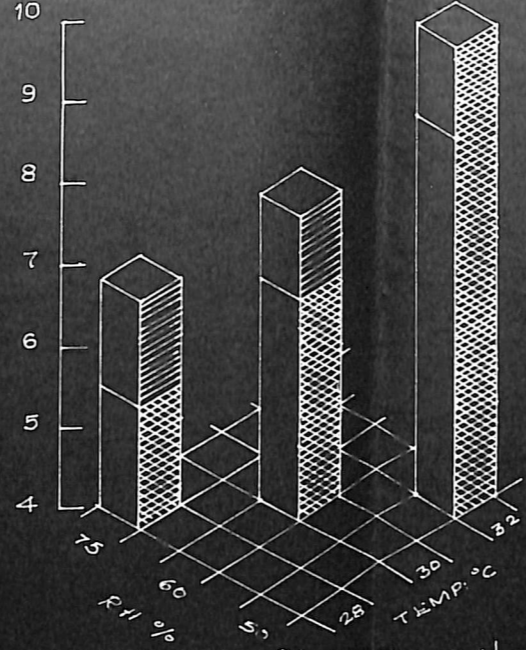
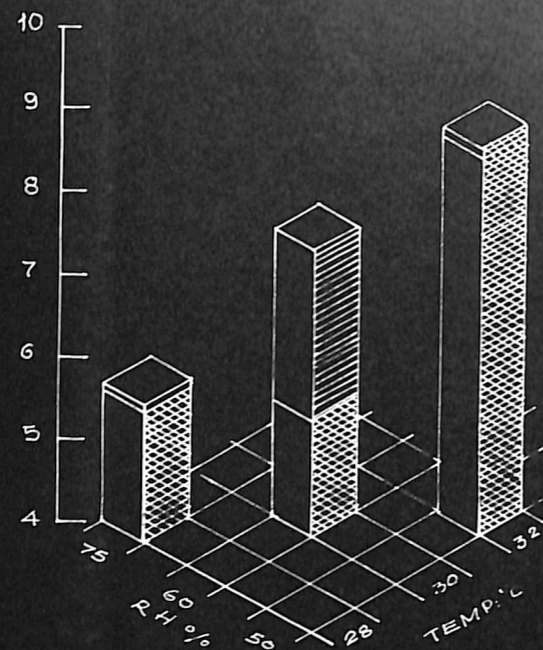


FIG. 5A

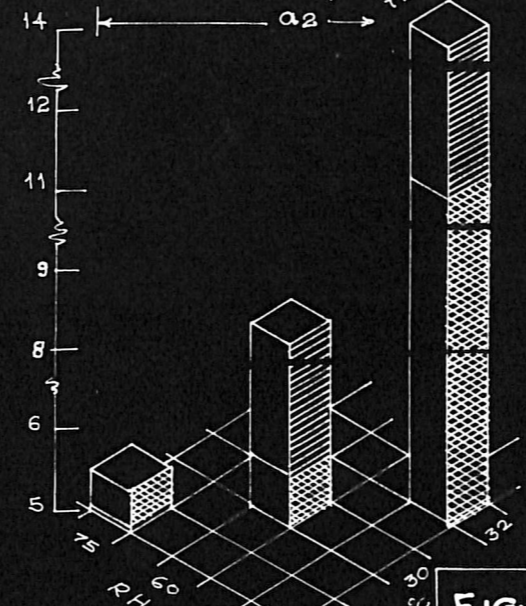
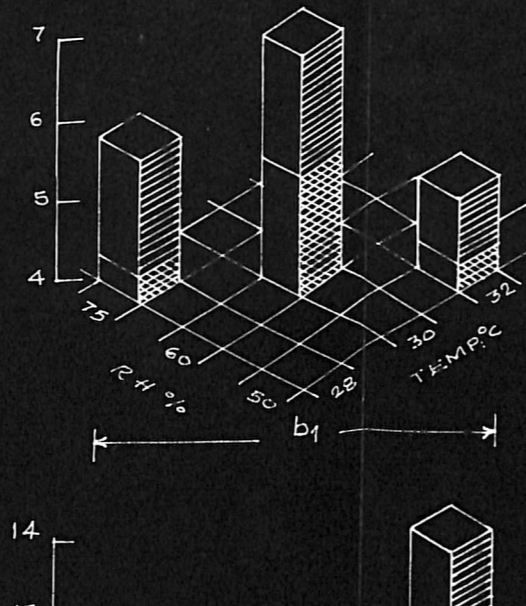
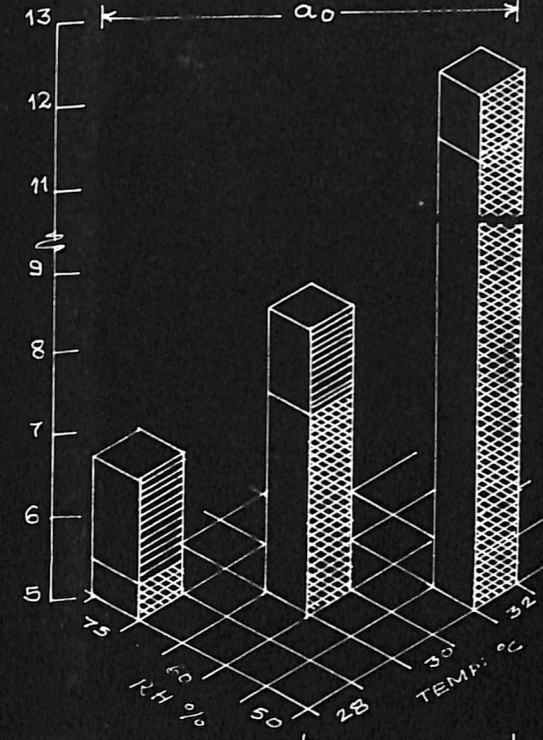
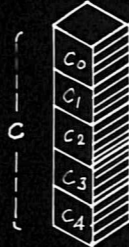
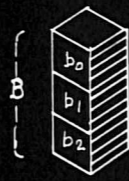


FIG. 5B

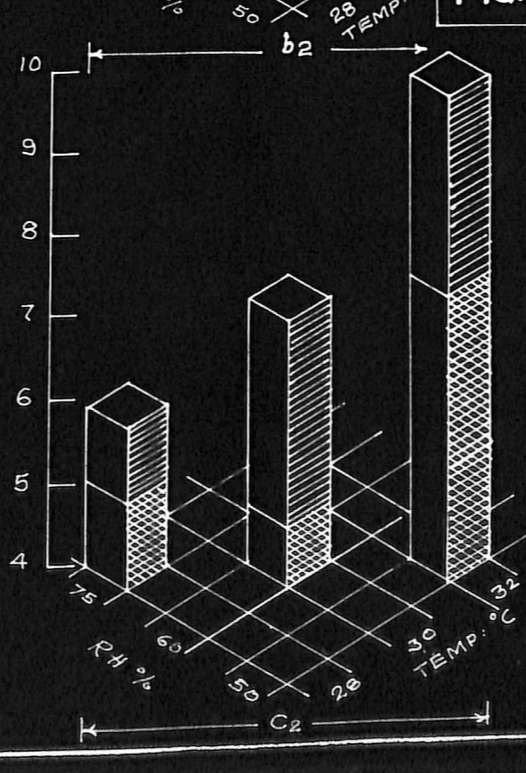
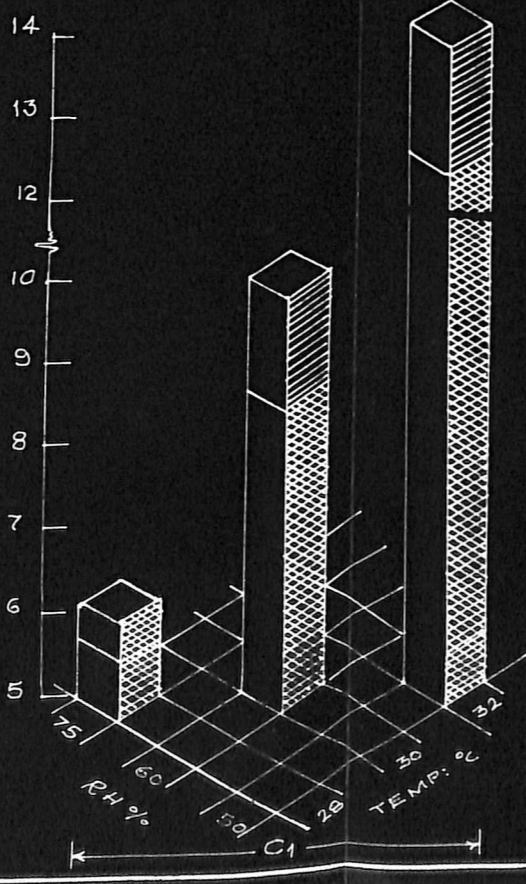
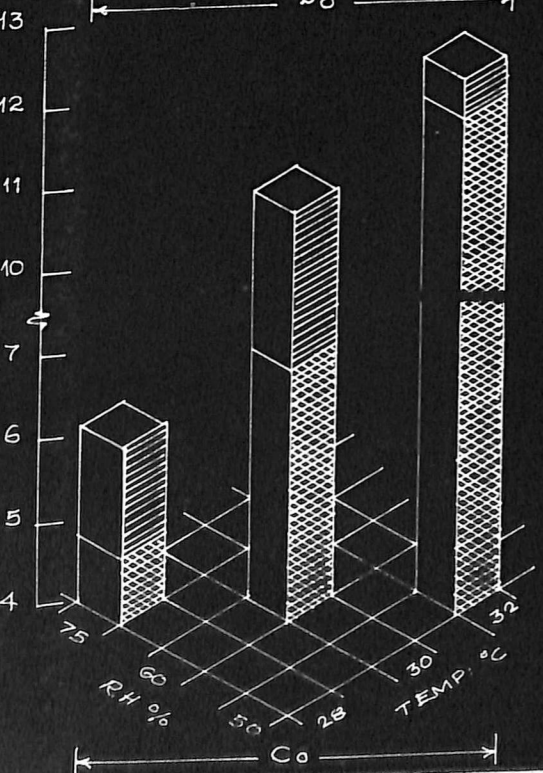


FIG. 5C

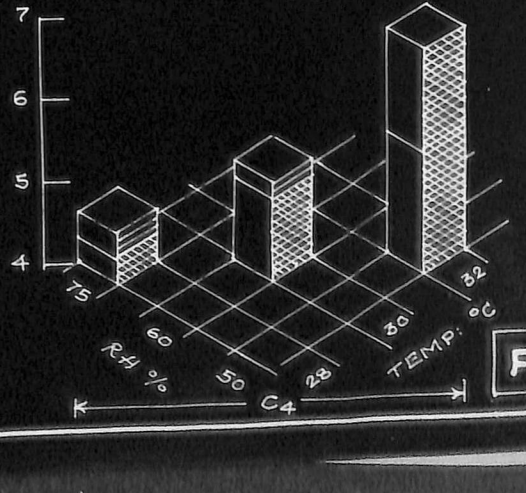
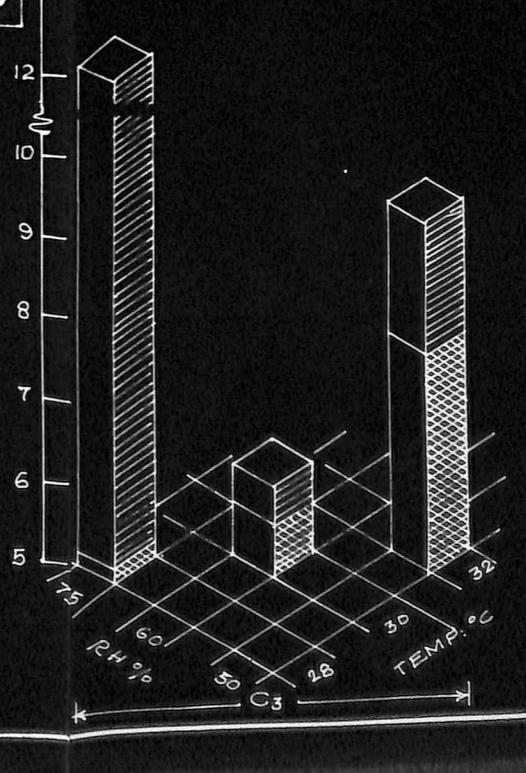


FIG. 5C

**Female-Male composition at 32°C and 50% RH**

At the above temperature and humidity combination, all the three main factors A (host larval density), B (weight of host larvae) and C (sex-ratio of parent parasite population had significant effects on the sex-ratio of the  $F_1$  progeny (Table 5-d).

Among the levels of A (host larval density), the  $a_1$  level (2 host larvae per female parasite) recorded the highest percentage of females (53.10). The sex-ratio recorded at  $a_1$  level was significantly better than those recorded at the  $a_2$  level (3 host larvae per female parasite). No significant difference in the sex-ratio was detected between the levels  $a_0$  (1 host larva per female parasite) and  $a_1$ . But the sex-ratio at the  $a_0$  level was significantly better than that at  $a_2$  level.

Among the levels of B (weight of host larvae), the  $b_0$  level (host larval weight 30-35 mg) recorded the highest percentage of females (51.73) and the sex-ratio at the  $b_0$  level was significantly better than those at  $b_1$  and  $b_2$  levels. But no significant difference was detected between the sex-ratios recorded at  $b_1$  and  $b_2$  levels.

Among the levels of  $\bar{C}$  (sex-ratio of parent parasite population), the  $c_4$  level (1 female : 3 males) recorded the highest percentage of females (56.15) and the sex-ratio was significantly better than those recorded at  $c_3$ ,  $c_2$  and  $c_1$  levels.

# DISCUSSION

## DISCUSSION

The fecundity, progeny production and female-male composition of Bracon brevicornis Wesmael as influenced by the number and weight of host larvae of Coxoza cephalonica Stainton and the sex-ratio of the parent parasite population were studied at three temperature-humidity combinations which represented near optimum condition of 28°C and 75% RH and near-adverse conditions of 30°C and 60% RH and 32°C and 50% RH and these combinations are henceforth referred to as TH<sub>1</sub>, TH<sub>2</sub> and TH<sub>3</sub> respectively. The effects of the above three factors on the duration of development were also studied. The following were the levels at which the effects of the three factors were studied.

Factor 1: Host larval density (A)

Levels a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " "  
 a<sub>2</sub> - 3 " "

Factor 2: Weight of host larvae (B)

Levels b<sub>0</sub> - host larval weight 30 to 35 mg  
 b<sub>1</sub> - " 8 to 10 mg  
 b<sub>2</sub> - half of the hosts of the b<sub>0</sub> type  
 and the other half of the b<sub>1</sub> type

**Factor 3: Sex-ratio of parent parasite population**

Levels	$\alpha_0$	-	1:1	(Female: Male)
	$\alpha_1$	-	2:1	( " )
	$\alpha_2$	-	3:1	( " )
	$\alpha_3$	-	1:2	( " )
	$\alpha_4$	-	1:3	( " )

**FECONDITY OF B. BREVICORNIS**

The fecundity was recorded by examining the host larvae on successive days after confinement with the parasites and directly counting the eggs adhering to the host body. Data on the mean number of eggs laid by the female parasite under the different treatments at the three temperature-humidity levels are furnished in table 1-a and the results of chi-square test are presented in table 1-b. The number of eggs laid at different levels of the three factors are summarised and presented in table 1-c and these are graphically depicted in Fig.1 A, 1 B and 1 C.

All the main effects viz., host larval density (A), weight of host larvae (B) and sex-ratio of parent population were found to be significant.

**Influence of host larval density on fecundity of B. brevicornis**

A significantly higher number of eggs were obtained at all the three TH levels when the host larval density was maintained at two per female parasite ( $a_1$  level) than at the  $a_0$  (1 host larva per female parasite) and  $a_2$  (3 host larvae per female parasite) density levels.

That the fecundity of Bracon brevicornis and Microbracon sp. increased progressively with the number of hosts available is already reported by Douth (1959) and Sheshagiri Rao et al. (1967). The present finding closely agrees with the results obtained by Subba Rao et al. (1974) in the case of B. hebetor. In this case, maximum fecundity was registered with two Corovra larvae per parasite pair. The increased fecundity at high host density levels might be partly due to increase in the density of available ovipositional sites (Flanders, 1946) and partly due to increase in the availability of food for the female parasite (Ulliyett, 1943; 1945). Perhaps in B. brevicornis the optimum density of ovipositional sites and adequate adult nutrition are ensured with two hosts per parasite female. This is, however, not in consonance with Ulliyett's (1951) observation that in B. hebetor, a fixed number of eggs are laid regardless of the number of hosts.



The chi-square due to heterogeneity of host larval density was found to be non-significant indicating lack of heterogeneity in respect of this factor over the different TH levels (Table 1-e).

**Influence of weight of host larvae (B) on the fecundity of B. brevicornis**

Heavier larvae (larval weight 30 to 35 mg) and lighter larvae (larval weight 8 to 10 mg) of the same age and an admixture of both types in equal proportions were used in the present studies to ascertain whether parasitic development is influenced by the weight of larvae.

It was revealed that the weight of host larvae exerted significant influence on the fecundity of parasite at all the TH levels. At TH<sub>1</sub> and TH<sub>2</sub> levels, maximum fecundity was registered when host larvae comprising of heavier and lighter ones in equal proportions (b<sub>2</sub>) were exposed for parasitisation. The trend at the TH<sub>3</sub> level was, however, different and there was maximum egg production under the b<sub>0</sub> level where relatively heavier larvae were alone used.

At all the TH levels, the exclusive use of light weight larvae as hosts for the parasite (b<sub>1</sub>) was found to produce significantly less number of eggs. This

clearly indicates that when lighter larvae are used for rearing B. brevicornis, the oviposition potential is not fully realised. The low egg production by the parasite in lighter larvae may be explained on the basis of the lack of proper ovipositional stimulus in the lighter larvae or based on the reduced availability of body fluid exuding from the oviposition wounds of lighter larvae, for adult feeding as reported by Wylie (1962) in Nasonia vitripennis which is parasitic on Musca domestica.

The chi-square due to heterogeneity of host larval weight (B) over the different TH levels was found to be significant. At the  $b_0$  level, the  $TH_2$  combination was superior to the other two TH levels in obtaining higher fecundity. At the  $b_2$  level also maximum number of eggs were obtained at the  $TH_2$  level but there was no significant difference between the  $TH_2$  and  $TH_1$  levels. However, for the lighter larvae ( $b_1$ ), the superiority of the  $TH_2$  level was not manifested. For this level, the maximum fecundity was obtained for the  $TH_1$  level and the least for  $TH_3$ .

The results thus indicate that when heavier host larvae ( $b_0$ ) were used, maximum fecundity was obtained under  $TH_2$  combination (32°C and 50% RH) while in the case

of lighter larvae ( $b_1$ ) the  $TH_1$  combination ( $23^{\circ}C$  and 75% RH) was found to be relatively more suitable for better egg production. When heavier and lighter larvae were used in equal proportions ( $b_2$ ) both  $TH_3$  and  $TH_1$  levels were found to be equally efficient in obtaining maximum egg production. The studies reveal that the preferred levels of temperature-humidity combination for oviposition by B. brevicornis varies with the weight of host larvae. At high temperature - low humidity levels, the host body fluid exuding from oviposition wounds is likely to get dried up faster and hence heavier hosts might be essential for ensuring sufficient body fluid exudates in the required consistency for adult feeding. High fecundity obtained at the  $TH_3$  combination for heavier larvae is perhaps due to this reason. In the present studies all the adult parasites were offered honey-glucose mixture as food, but it is quite probable that the host body fluid will be preferred by the adult females to the honey-glucose solution.

**Influence of sex-ratio of parent parasite population (C) on fecundity of B. brevicornis**

At all the  $TH$  levels tested, the parental sex-ratio levels  $c_1$  (2:1) and  $c_2$  (3:1) were found to produce more number of eggs than under the sex-ratio levels  $c_0$  (1:1).

$c_3$  (1:2) and  $c_4$  (1:3). This clearly shows that a dominance of females in the parent parasite population ( $c_1$  and  $c_2$ ) is essential for ensuring higher fecundity and that when the proportion of females remained the same ( $c_0, c_3, c_4$ ), the fecundity do not increase with increase in the number of males. In arrhenotokous parasites, since eggs are laid even without mating and fertilisation, the sex-ratio of the parental parasite population is apparently not expected to influence fecundity. The present results indicate that the parental sex-ratio influence fecundity and this may be due to effective mating and successful fertilisation and resultant improvement of the stimulus for oviposition leading to better realisation of egg laying potential. It is quite possible that the optimum sex-ratio for successful fertilisation in B. brevicornis is  $c_1$  (2:1) and  $c_2$  (3:1).

The heterogeneity chi-square for the factor C (sex-ratio of parent parasite population) over the different TH levels being significant, it is revealed that the response to this varied over the different TH levels.

#### PROGENY PRODUCTION

The total number of males and females which emerged out from each replicate was recorded as the  $F_1$  progeny production under different treatments. Data on the

mean number of progenies produced under various treatments at the three TH levels are furnished in Table 3-a and the results of chi-square test are presented in Table 3-b. The number of progenies produced at different levels of the three main factors are summarised and presented in Table 3-c and these are illustrated in Fig. 3 A, 3 B and 3 C.

All the main effects viz., host larval density (A), weight of host larvae (B) and sex-ratio of parent parasite population (C) as well as their interactions were found to be significant.

#### Influence of host larval density on progeny production of B. brevicornis

The maximum number of progenies were produced when two host larvae were exposed per female parasite ( $a_1$ ) at TH<sub>1</sub> and TH<sub>2</sub> levels and this was significantly higher than the  $a_2$  level of three host larvae per female parasite at TH<sub>1</sub> level. But at the TH<sub>2</sub> level, significantly higher number of progenies were produced when three host larvae were offered per female parasite ( $a_2$ ).

It was found that the fecundity of the parasite was significantly higher at all the three TH levels for the level  $a_1$  (2 hosts per female parasite). For the TH<sub>1</sub>

and  $TH_2$  levels, progeny production was also found to be better under the  $a_1$  level and this is only expected based on the fecundity trend. It is evident that temperature-humidity combination for realisation of fecundity potential also operates in the same manner for embryonic and post-embryonic development leading to similar trend in the progeny production. That at  $TH_3$  level maximum fecundity was obtained at two host larvae per female ( $a_1$ ) while maximum progeny production was obtained at  $a_2$  level of three host larvae per female parasite is perhaps due to parasite mortality occurring in the immature stages.

The chi-square due to heterogeneity for the host larval density (A) was significant indicating that the responses to host larval density vary with the TH levels. At all the three host density levels  $a_0$ ,  $a_1$  and  $a_2$ , the progeny production was found to be significantly higher at the  $TH_3$  level as compared to the  $TH_1$  and  $TH_2$  levels obviously due to the improvement in embryonic and post-embryonic development leading to better progeny production. When  $TH_3$  condition is considered separately, it is seen that higher host density ( $a_2$ ) was essential for getting maximum progeny production while an overall consideration shows the superiority of  $TH_3$  level for all the three host density levels.

According to Flanders (1935), in Trichogramma evanescens, with increasing host densities, the number of progeny per parasite increased until a maximum was reached. The number of progeny per parent becomes stable beyond a stage when the number of hosts within the sphere of action of the parasites exceeds the total reproductive potential of the parasite. Seshagiri Rao et al., (1967) reported that progeny production in B. brevicornis increased progressively with the number of hosts available, but it showed a decreasing trend in the case of Perisporola nephandidis at high host density levels.

Abraham and Mathew (1978) observed that the number of exposed Coryza larvae is relatively more important in the progeny production of B. brevicornis. Viktorov and Azisov (1972) recorded that progeny production in Muscidifurax raptor increased with increase in the number of host puparia of Musca domestica. The present results are in general agreement with the early observations, subject to the detection of an optimum level of two larvae per female parasite at  $TH_1$  and  $TH_2$  and three larvae per host at  $TH_3$ . Under all host density levels,  $TH_3$  combination was found the best in getting higher progeny production.

**Influence of weight of host larvae (B) on progeny production of B. brevicornis**

It is indicated that significantly higher number of progenies were produced when heavier larvae ( $b_0$ ) were exposed for parasitisation both at  $TH_1$  and  $TH_2$  conditions. Under the  $TH_3$  level, an admixture of heavier and lighter larvae in equal proportions ( $b_2$ ) and the heavier larvae ( $b_0$ ) were on par with reference to progeny production. The better progeny production from heavier larvae can be explained in the light of better larval and pupal development due to better nutritional status of such larvae.

The heterogeneity chi-square for the weight of host larvae (B) was found to be significant indicating differential responses of the factor B over the three TH levels. The  $b_0$  (heavier larvae) and  $b_2$  (mixture of heavier and lighter larvae in equal proportions) levels registered maximum progeny production at the  $TH_3$  level (32°C and 50% RH). However, for the  $b_1$  level (lighter larvae), maximum progeny production was registered at the  $TH_2$  combination (30°C and 60% RH). The trend in higher progeny production at the  $b_0$  and  $b_2$  levels is similar to the trend in higher egg production at the  $TH_3$  level. The variations in the trend of progeny



production at different temperature-humidity levels with host weight changes is due to the differential impact of these abiotic environmental components in relation to host nutritional status indicated by host weight.

As in Exenterus abruptorius (Hymenoptera: Ichneumonidae) (Gurjanova, 1974), it is quite possible that E. brevicornis lays fertilised eggs on large larvae and unfertilised eggs on small larvae and thus the male producing eggs will be found only on lighter larvae. The differential response of the weight of host larvae over the different TH levels will then be due to the variations in the temperature-humidity tolerances of the two sexes.

**Influence of sex-ratio of parent parasite population (O) on progeny production of E. brevicornis**

A sex-ratio of 1:2 (females:male) in the parent parasite population ( $c_3$ ) has recorded a significantly higher progeny production as compared to the  $c_4$  level (1:3) at  $TH_1$  combination. At the  $TH_2$  and  $TH_3$  levels, the trend in progeny production was in favour of better progeny production under  $c_1$  (2:1) than under  $c_4$  (1:3). At all the tested TH levels, the parental sex-ratio of

1:3 ( $c_4$ ) was found inferior to the rest of the sex-ratio levels with reference to progeny production. The  $c_1$  (2:1) and  $c_2$  (3:1) levels of parental sex-ratios characterized by the dominance of females were already found to be consistently better in the production of eggs at all the TH levels. However, similar consistent trends in progeny production at all TH levels were not detected. Thus at TH<sub>2</sub> and TH<sub>3</sub> levels,  $c_1$  (2:1) sex-ratio was found to be superior to  $c_4$  (1:3) and this indicates that for better progeny production also a dominance of females ( $c_1$ ) rather than of males ( $c_4$ ) is desirable. This finding is not in full agreement with the earlier report by Abraham and Mathew (1978) that the number of males in the parental population of B. brevicornis is relatively of more important.

The results obtained at the TH<sub>1</sub> level (28°C and 75% RH) are not in consonance with the trend under TH<sub>2</sub> and TH<sub>3</sub> conditions and the superiority of the  $c_3$  ratio (1 female: 2 males) indicates that at relatively low temperature-high humidity conditions the preferred nature of sex-dominance is tilted towards male dominance at the above level. The finding of Abraham and Mathew (1978) that the number of males in the parental population

is relatively more important under room temperature conditions around  $28 \pm 1^\circ\text{C}$  is in line with the present observation.

The heterogeneity chi-square was found to be significant and this indicates that the responses of the factor  $\theta$  vary with temperature-humidity combinations. At all the levels of  $\theta$  ( $\theta_0, \theta_1, \theta_2, \theta_3$  and  $\theta_4$ ), significantly higher progeny production was consistently obtained at the  $\text{TH}_3$  level of  $32^\circ\text{C}$  and 50% RH. The improvement of progeny production at the  $\text{TH}_3$  condition for all the parental sex-ratios indicates that for embryonic and post-embryonic development of the parasite this particular temperature-humidity combination<sup>is</sup> relatively more favourable.

#### NUMBER OF FEMALE PROGENY

The total number of female emergents was recorded under different treatment combinations. Data on the mean number of females produced by the parasite under various treatments in different TH combinations are furnished in Table 4-a and the results of chi-square test are presented in Table 4-b. The mean number of females produced by the different levels of the three

main factors are indicated in Table 4-c and these are represented in Fig. 4A, 4B and 4C.

**Influence of host larval density (A) on female progeny production of B. brevicornis**

It is indicated that the host larval density exerted significant influence on the number of females produced by the parasite at  $TH_1$  (28°C and 75% RH) and  $TH_2$  (30°C and 60% RH) levels only. When one host larva per female parasite ( $a_0$ ) was exposed for parasitisation at the  $TH_1$  level, a significantly higher number of female progeny was obtained than those obtained from three host larvae ( $a_2$ ). It is thus observed that as the number of hosts per female parasite increased from one to three, the number of female progeny decreased. Similar trend of production of progeny with a small proportion of females at low host densities was also reported by King (1962) in the case of Nasonia vitripennis.

The trend in female progeny production at  $TH_2$  combination of 30°C and 60% RH was quite different and under this condition significantly more number of female progeny was produced at the  $a_1$  level (two host larvae per female parasite) than at the  $a_0$  level (one host larva per female parasite).

The female progenies are expected to arise from diploid eggs and the variations in the number of female progenies are thus caused by the fertilisation and proportion of diploid eggs that are laid on the hosts. Even after fertilisation, the deposition of diploid eggs is regulated by the presence of adequate stimulus picked up through the ovipositor and/or the antennae and the divergent trends of female progeny production in different TH conditions from varying number of hosts is perhaps due to the modification of ovipositional stimulus under temperature-humidity variations. This is substantiated by the significant heterogeneity chi-square for the host larval density (A) over the different TH combinations.

**Influence of weight of host larvae (B) on female progeny production of B. brevicornis**

The weight of host larvae had significant influence on the number of female progeny produced at all the three TH combinations. At the TH<sub>1</sub> level, an admixture of heavier and lighter larvae in equal proportions (b<sub>2</sub>) produced the maximum number of female progeny. This was followed by the heavier larvae with a weight range of 30 to 35 mg (b<sub>0</sub>). Lighter larvae with a weight range of 8 to 10 mg (b<sub>1</sub>) produced significantly lower number of female progeny.

Both at the TH<sub>2</sub> and TH<sub>3</sub> combinations, the heavier larvae (b<sub>0</sub>) produced significantly higher number of female progenies as compared to the b<sub>1</sub> level. It is thus observed that at all the three TH levels, minimum number of females were produced by lighter larvae (b<sub>1</sub>). The production of significantly higher number of female progeny from heavier hosts is explicable on the basis of better nutritional status of such larvae and the variations in the nutritional requirements of the parasite larvae belonging to the two sexes. It is thus possible that more number of females develop on large hosts than in lighter hosts which is in consonance with the report by Chewyrov (1913) that the great majority of individuals of Pimpla from large hosts were females whereas those from small hosts were predominantly males. This finding also agrees with Clausen (1939) and Flanders (1939) who have reported that in the parasitic Hymenoptera, generally large hosts favour the production of female parasites, while the smaller hosts tend to favour the production of males. But Fusco and Hower (1973) stated that size of the host Hymer postica had no significant effect on the mean percentage of females produced by the parasite Microctonus aethiops in the laboratory. Brunson (1937) found that the female parasite Tiphia popillivora had the ability to control sex of progeny at the time of

oviposition and that the stimulus to which the female responds in controlling sex of the progeny is definitely associated with the size of the host on which eggs are placed.

In addition to the nutritional factor, the modification of ovipositional stimulus also might be responsible for variations in female progeny production under the influence of host weight. The modification of the ovipositional stimulus may be brought about through the stimulus perceived either by the antennae or by the ovipositor from the host, leading to a change in the ratio of fertilised and unfertilised eggs incited by changes in the activity of the spermathecal gland (Narayanan and Choudhuri, 1955).

The chi-square due to heterogeneity for the weight of host larvae was found to be significant indicating that the responses due to this factor (B) vary with TH combinations. At the  $b_0$  level, (heavier host larvae with a weight range of 30 to 35 mg) and  $b_2$  level (heavier and lighter larvae in equal proportions), significantly higher number of females were produced at the  $TH_2$  level (32°C and 50% RH) while at the  $b_1$  level (lighter larvae) the  $TH_2$  condition produced maximum number of female progeny. There is thus an increasing

trend of female progeny production with increasing temperature and decreasing humidity levels both at  $b_0$  and  $b_2$  levels.

**Influence of sex-ratio of parent parasite population(C) on female progeny production of B. brevicornis**

At  $28^{\circ}\text{C}$  and 75% RH, with 2 females : 1 male ratio ( $c_1$ ) among the parent parasite population, significantly higher number of female progenies were produced than those obtained with 1 female: 1 male ( $c_0$ ) and 1 female : 3 males ( $c_4$ ).

At  $\text{TH}_2$  ( $30^{\circ}\text{C}$  and 60% RH) and  $\text{TH}_3$  ( $32^{\circ}\text{C}$  and 50% RH) combinations, maximum number of females were produced when the sex-ratio of parental population was kept at 2 females: 1 male ( $c_1$ ). The superiority of the parental sex-ratio of  $c_1$  (2:1), in improving the fecundity at all the TH levels and in enhancing the progeny production at the  $\text{TH}_2$  and  $\text{TH}_3$  levels has been already established in the present studies. It is now revealed that the same parental sex-ratio is consistently superior at all the TH levels in the production of female progeny also. The female progeny production in successive generations being of vital importance in the sustenance of laboratory cultures at high population



levels, the present finding is considered to be of practical importance. The results reported by Mathew and Kair (1977) that the female-male ratios of 1:3, 2:1, 1:6, 3:1 and 1:4 in the parental populations improved the percentage of female progeny in the offspring, indicated that particular proportions of both sexes are more important than dominance of males or females. The present studies show that dominance of females is definitely essential for ensuring better female progeny production. The influence of female dominance on female progeny production is perhaps due to the necessity of female dominance in the parental population for effective fertilisation. More number of females might also be required to compensate for the sterility among the females.

The heterogeneity chi-square for the sex-ratio of parent parasite population (G) was significant indicating differential responses of this factor over three TH levels. At all levels of parental sex-ratios, the female progeny production showed significant increase at the TH<sub>3</sub> combination of 32°C and 50% RH as compared to the other combinations. This may be due to the selective adverse effect of high temperature

low humidity condition on male offsprings during the embryonic or post-embryonic development. Similar selective susceptibility of the haploid males of Habrobracon juglandis to environmental adversities has been reported by Georgiana (1949). Pilon, (1966) also recorded that the influence of temperature varied with relative humidity in the case of males of Neodiprion swainsei but not in the case of females. Kochetrova (1978) had also indicated the possibility of unequal mortality of sexes in the ontogeny of entomophagous insects due to endogenous and exogenous factors.

#### FEMALE-MALE COMPOSITION OF THE $F_1$ PROGENY

Data on the mean number of females and males recorded at the three TH levels under various treatments are furnished in Table 5-a. The female-male composition of the  $F_1$  progeny under different levels of the main factors at the three TH combinations are presented in Tables 5-b, 5-c and 5-d and these are graphically illustrated in Fig. 5A, 5B and 5C.

#### Influence of host larval density (A) on female-male composition of the $F_1$ progeny

The host larval density (A) had no significant influence on the female-male composition of the  $F_1$

offspring of B. brevicornis at  $TH_1$  and  $TH_2$  levels. But at the  $TH_3$  level, sex-ratio of the  $F_1$  progeny was significantly influenced by the host larval density. Highest percentage of females were obtained when two host larvae per female parasite ( $a_1$ ) were offered for parasitisation. The sex-ratio of the  $F_1$  progeny recorded at the  $a_1$  level was significantly better than those recorded at the  $a_2$  level of three host larvae per female parasite. The female-male composition of the progeny is largely influenced by the proportion of diploid eggs by the inseminated females and also by the selective mortality of the sexes during embryonic and post-embryonic development. The host larval density factor might be of importance as ovipositional sites for the parasite and also in modifying the ovipositional stimulus. It is seen that at  $TH_1$  and  $TH_2$  levels the increase or decrease in larval density does not cause changes in the proportion of male and female producing eggs. In this studies, it was already found that at the  $TH_1$  and  $TH_2$  levels, larval density affected the female progeny production. The apparent anomaly relating to the importance of host density in deciding the number of female progenies on the one hand and the non-significance of the host density factor as a mechanism regulating female-male composition

of the progeny on the other hand can be resolved on the basis of possible concomitant variations in male progeny production leading to the restoration of female-male proportion in such situations. The influence of host larval density on the sex-wise composition of the  $F_1$  progeny at the  $TH_2$  level is perhaps due to the effect of the level of temperature and humidity in modifying the stimulus of depositing haploid and diploid eggs. This is substantiated by the finding of Abraham and Mathew (1978) that host larval density is relatively important among other factors, in the regulation of sex-ratio of the offsprings.

#### Influence of weight of host larvae (B) on female-male composition of $F_1$ progeny

At the  $TH_1$  and  $TH_2$  levels, the sex-ratio of the  $F_1$  progeny was significantly influenced by the weight of host larvae. The highest percentage of females was recorded under the  $b_2$  level (mixture of heavier and lighter larvae in equal proportions) at the  $TH_1$  combination. The sex-ratio registered at the  $b_2$  level was significantly better than those recorded at the  $b_1$  level (lighter larvae). At the  $TH_2$  combination, heavier host larvae ( $b_0$ ) has recorded a significantly better sex-ratio than those recorded at the  $b_1$  and  $b_2$  levels. However, at the  $TH_2$

level the host weight was found not to have any significant effect on the sex-ratio.

In the present studies, the relative superiority of the heavier host larvae ( $b_0$ ) in the realisation of egg production potential and female progeny production under  $M_2$  level (32°C and 50% RH) has been already established. It is also revealed that the heavier host larvae are relatively better for a favourable female-male composition of the  $F_1$  offsprings characterised by the dominance of females. These results indicate that for successful laboratory rearing of the parasite, relatively heavier larvae (weight range 30 to 35 mg) are to be preferred under ambient conditions characterised by relatively high temperature (32°C) and low humidity (50% RH). The influence of host weight on female progeny production in B. brevicornis might be either due to differences in the nutritional status of larvae of different weights or the variations in the nutritional requirements of males and females. This influence may also be due to the sex of the host larvae as reported by Sithanathan and Subramanian (1977) in the case of Chilo partellus and Stenobracor deesae. The sex-ratios of S. deesae was in favour of males in male hosts while it shifted significantly in favour of females in the

female hosts. In B. brevicornis it is possible that the relatively heavier larvae ( $b_0$ ) are the females and the lighter ones are males. The better production of female progeny from heavier hosts might then be the influence of host sex possibly manifested in the form of selective mortality of the opposite sex due to some antagonistic hormonal influence. It is further revealed that when the changes in ambient temperature and humidity conditions are varied the preferred level of the host weight also changes. Thus it is observed that at  $TH_1$  combination, an admixture of heavier and lighter larvae in equal proportions ( $b_2$ ) is to be preferred for the production of a higher proportion of females among the offsprings. The influence of the heavier host larvae on fecundity, female progeny production and the female-male composition of the progeny is found to be inconsistent perhaps due to the changes in the ovipositional stimulus provided by the hosts at different temperature and humidity conditions.

**Influence of sex-ratio of the parent parasite population (J) on female-male composition of  $F_1$  progeny.**

The female-male composition of the  $F_1$  progeny in the form of proportion of females of B. brevicornis was found to be significantly influenced by the sex-ratio

of the parent parasite population at the  $TH_2$  and  $TH_3$  combinations. The  $c_4$  level (1 female: 3 males) has produced a significantly better proportion of females than those recorded at the levels  $c_2$  (3:1) and  $c_0$  (1:1) under the  $TH_2$  combination. At the  $TH_3$  level also the  $c_4$  level has produced a significantly better proportion of females than under the levels  $c_3$ ,  $c_2$  and  $c_1$ . It has been already established that a dominance of females in the parental population ( $c_1/c_2$ ) is superior for better egg production, improved progeny production and for the production of more number of female offsprings. It is now found that for ensuring a high proportion of females among the progeny at the  $TH_2$  and  $TH_3$  conditions, the dominance of males ( $c_4 = 1:3$ ) is essential. This is in general agreement with the finding reported by Mathew and Hair (1977) that particular sex-ratios among the parents ensured satisfactory sex-ratios among the progenies including the one characterised by male dominance. In the present studies, dominance of females was found essential for better fecundity, progeny production and number of female progeny production but the dominance of females is not found critical in deciding the proportion of females among the progeny.

An overall consideration from the point of view of improved fecundity, progeny production and number of

female offsprings is essential for deciding the desirability of a particular sex-ratio among the parental population. The  $c_1$  and  $c_2$  levels characterised by the dominance of females were found superior in the above context, and since the influence of  $c_4$  level characterised by dominance of males cannot be rated as desirable considering the inconsistencies in its regulating influence on other biological attributes such as fecundity, progeny production and female progeny production. Further, the higher number of female progeny production is to be preferred to the higher proportion of females. It is possible that with a few females among a limited progeny, the proportion may appear as quite high, but, this will not be much practical utility.

#### DURATION OF DEVELOPMENT

The duration of development was found out by calculating the weighted means of the number of days taken by both sexes of B. brevicornis emerging on successive days. For this the emerging parasites were isolated, killed and the female-male composition in relation to the duration of development recorded on each day.



Data on the mean duration of development of the parasites under different treatments at the three TH combinations are furnished in Table 2-a. The results of the analysis of variance are given in Table 2-b and the mean values of the duration of development under the treatment combinations are presented in Table 2-c, 2-d and 2-e. These are illustrated in Fig. 2A, 2B and 2C.

#### Influence of host larval density (A)

The host larval density had significant influence on the duration of development of the parasites at all the three TH combinations. It was observed that the shortest duration of development was revealed by the  $a_2$  level (3 host larvae per female parasite) at the TH<sub>1</sub> condition. When only one host larva was offered per female parasite ( $a_0$ ), the duration of development was relatively longer. The above trend in shortening of the duration of development with increase in host density level was also detected for the TH<sub>2</sub> level. Metcalfe and Breniere (1969) in Trichogramma spp. had reported that a high level of parasitism was rapidly attained under high host density levels due to shortening of the life-cycle.

However, at the  $TH_3$  combination, the shortest duration of development was registered at low host density level of one host larva and this was significantly better than the duration for the  $a_1$  level (2 host larvae). The variations in the duration of development at different host larval density levels might be due to the amount of food materials that would be provided by the larvae. Thus shortening of the duration of development at high densities will be possible due to normal development under adequate nourishment. The shortening of duration of development at  $TH_3$  under low density level is perhaps due to the modification of the nutritional requirements under influence of high temperature-low relative humidity levels.

**Influence of weight of host larvae (B) on duration of development of A. brevicornis**

At the  $TH_1$  and  $TH_2$  combinations, the duration of development was not influenced by the host larval weight (B). But weight of host larvae exerted significant influence at the  $TH_3$  combination. It was observed that the parasites emerging from the lighter larvae ( $b_1$ ) had a significantly shorter duration of development. The nutritional status

of lighter larvae will be relatively lower and under such condition, the parasitic larvae might be constrained to complete the life cycle faster as an adaptive mechanism. The duration of development of the parasite at the levels  $b_0$ ,  $b_2$  were not significantly different.

**Influence of sex-ratio of the parent parasite population (C) on the duration of development of B. brevicornis**

At the  $TH_1$  combination the shortest duration of development was noticed when the sex-ratio of the parent parasite population was kept at  $c_2$  (3 females: 1 male) while at  $c_3$  level (1 female: 2 males), duration of development was the longest. The  $c_3$  level (1:2) has indicated the shortest duration of development at the  $TH_2$  combination. At the  $TH_3$  combination the shortest duration of development was recorded by the  $c_1$  (2:1) and  $c_2$  (3:1) levels and the  $c_3$  level (1:2) has registered the longest duration of development at  $TH_1$  and  $TH_3$  combinations.

The divergent trends in the duration of development at different parent sex-ratio levels at various  $TH$  conditions is probably due to the proportion of diploid and haploid eggs laid by parasite females and due to

inherent variations in the speed of development of both sexes. This may ultimately be reflected by causing variations in the duration of development of the parasite.

#### Overall considerations

The nature of influence of the density and weight of host larvae of Cerovra cephalonica and the sex-ratio of the parent parasite population of Bracon brevicornis on the fecundity, progeny production and the female-male composition of the  $F_1$  parasite offsprings at 28°C-75% RH, 30°C-60% RH and 32°C-50% RH is found to be varying in most of the cases, obviously due to the differential effect of these abiotic components on host parasite interaction.

The implications of the trend in fecundity, progeny production, female-male composition of the progeny and the number of female progeny as regulated by the density and weight of host larvae and the sex-ratio of the parent parasite population under diverse temperature-humidity conditions have been discussed.

Among the different host larval density levels tested, the density of two to three larvae per female

parasite is found to be relatively more effective in attaining higher egg production and improved progeny production.

The studies conclusively proved that among host larvae of the same age, lighter larvae were consistently inferior to the heavier ones with regard to the fecundity, progeny production and number of female progeny.

The desirability of maintaining the parental sex-ratio levels at 2:1 and 3:1 (female:male) for ensuring improved fecundity, progeny production and higher number of female offsprings at the different RH levels is brought out in the present studies.

The identification of temperature-humidity levels suitable for the realisation of the innate capacity for numerical increase as revealed by the biological attributes such as parasite fecundity, progeny production and female-male composition of the  $F_1$  progeny leads to the possibility of manipulating the ambient temperature-humidity conditions in insectaries for the sustenance of laboratory cultures of B. brevicornis at the maximum possible levels.

# SUMMARY

### SUMMARY

The fecundity, progeny production, female-male composition of the progeny and duration of development of Bracon brevicornis Wesmael as influenced by the density and size (weight) of host larvae of Coxoyle cephalonica Stainton and the sex-ratio of the parent parasite population were studied at three temperature-humidity (TH) combinations viz., 28°C and 75% RH (TH<sub>1</sub>), 30°C and 60% RH (TH<sub>2</sub>) and 32°C and 50% RH (TH<sub>3</sub>).

### FECONDITY OF B. BREVICORNIS

The host larval density, weight of host larvae and the sex-ratio of the parent parasite population were found to influence the fecundity of the parasite at all the TH levels. The maximum fecundity was registered consistently at all the three TH levels and a host density level of two larvae per female parasite.

The exclusive use of light weight larvae (weight range 8 to 10 mg) as hosts for the parasite produced significantly less number of eggs at all the three temperature-humidity conditions. At the TH<sub>1</sub> and TH<sub>2</sub> levels an admixture of heavier and lighter larvae in equal proportions gave more number of eggs while at the TH<sub>3</sub> level, egg

production was maximum when the heavier host larvae (weight range 30 to 35 mg) were used.

The parental sex-ratio levels of 2:1 and 3:1 (female:male) characterised by the dominance of females led to the production of higher number of eggs than under the sex-ratio levels of 1:1, 1:2 and 1:3.

#### PROGENY PRODUCTION OF B. BREVICORNIS

The host larval density, weight of host larvae and sex-ratio of parent parasite population were found to exert significant influence on the number of progeny of B. brevicornis. The maximum progeny production at the TH<sub>1</sub> (28°C and 75% RH) and TH<sub>2</sub> (30°C and 60% RH) levels was attained under a host density level of two larvae per female parasite. However, at the TH<sub>3</sub> level (32°C and 50% RH) the progeny production was maximum at the larval density level of three hosts per female parasite.

With regard to the weight of host larvae, it was found that the progeny production was higher at TH<sub>1</sub> and TH<sub>2</sub> levels when heavier larvae were exposed for parasitisation. Under the TH<sub>3</sub> level, the heavier larvae or an admixture of heavier and lighter larvae, were found to be equally effective in the production of higher number of progenies.



The superiority of the parental sex-ratio of 2:1 (female:male) in the production of higher number of progeny was established at the TH<sub>2</sub> and TH<sub>3</sub> levels, while the parental sex-ratio of 1:3 with a preponderance of males was found to be distinctly inferior to the other levels.

#### NUMBER OF FEMALE PROGENY OF B. BREVICORNIS

Significant influence of the host larval density was detected only at the TH<sub>1</sub> and TH<sub>2</sub> levels. At 30°C and 60% RH (TH<sub>2</sub>), higher number of female progeny was produced at a host larval density level of two per female parasite, while at 28°C and 75% RH (TH<sub>1</sub>) the female progeny production was higher at a host density level of one larva per female parasite.

Relatively heavier host larvae produced significantly higher number of female offsprings at the TH<sub>2</sub> and TH<sub>3</sub> combinations.

The parental sex-ratio level of 2:1 (female:male) produced maximum number of female progeny at all the tested temperature-humidity combinations.

#### FEMALE-MALE COMPOSITION OF THE F<sub>1</sub> PROGENY OF B. BREVICORNIS

The influence of host larval density was pronounced only at the TH<sub>3</sub> level (32°C and 50% RH) and under this

condition, a host density of two larvae per female parasite was found to be better with reference to the production of a higher proportion of females among the offsprings.

The highest percentage of females among the progeny was registered at  $TH_1$ , in treatments comprising heavier and lighter larvae in equal proportions, while at the  $TH_2$  level, the heavier host larvae recorded a higher proportion of females.

The sex-ratio of the parent parasite population influenced the proportion of females in the  $F_1$  progeny and a ratio of 1:3 (female:male) produced significantly higher proportion of females at the  $TH_2$  and  $TH_3$  combinations.

#### DURATION OF DEVELOPMENT OF B. BRAVICORNIS

The host larval density, weight of host larvae and the sex-ratio of the parent parasite population exerted significant influence on the duration of development of the parasite at various temperature-humidity conditions.

#### OVERALL CONSIDERATIONS

The nature of influence of the density and weight of host larvae of Cercyxa cephalonica and the sex-ratio of the parent parasite population of Bracon brevicornis

of the fecundity, progeny production and the female-male composition of the  $F_1$  parasite offsprings at  $28^\circ\text{C} - 75\% \text{RH}$ ,  $30^\circ\text{C}-60\% \text{RH}$  and  $32^\circ\text{C}-50\% \text{RH}$  is found to be varying in most of the cases, obviously due to the differential effect of these abiotic components on host-parasite interaction.

Among the different host larval density levels, the density of two to three larvae per female parasite is found to be relatively more effective in attaining higher egg production and improved progeny production.

The studies conclusively proved that among host larvae of the same age, lighter larvae were consistently inferior to the heavier ones in the realisation of fecundity and progeny production potentials and also with reference to the production of female progeny.

The desirability of maintaining the parental sex-ratio levels at 2:1 and 3:1 (female:male) for ensuring consistency in improved fecundity, better progeny production and higher number of female offsprings at the different TH levels is brought out in the present studies.

The feasibility of manipulating the ambient temperature-humidity conditions for maintaining laboratory cultures of B. brevianis at the maximum possible levels has been discussed.

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APPENDIX - I

Fecundity of *B. brevicornis* at different temperature and humidity combinations

Sl. No.	Treatments	28°C and 75% RH			30°C and 60% RH			32°C and 50% RH		
		RI	RII	RIII	RI	RII	RIII	RI	RII	RIII
1.	a	80	44	58	30	36	22	60	32	92
2.	a	49	48	40	78	36	38	75	95	160
3.	b	41	15	40	98	35	8	125	120	122
4.	b	72	52	41	56	88	99	22	44	83
5.	c	15	2	21	35	20	32	53	58	65
6.	c	4	14	2	13	48	42	35	42	5
7.	d	16	27	25	60	45	70	25	22	27
8.	d	85	50	100	45	70	75	15	40	9
9.	e	82	40	35	20	5	17	50	8	18
10.	e	50	52	57	20	40	58	15	24	10
11.	e	80	53	40	80	22	48	65	79	90
12.	e	50	60	52	75	60	85	92	53	95
13.	e	95	100	85	35	40	140	79	11	93
14.	e	73	55	64	48	46	45	40	15	50
15.	e	100	108	38	20	38	18	40	42	49
16.	e	85	100	80	35	55	25	77	80	45
17.	e	45	4	150	97	82	62	140	120	94
18.	e	108	115	65	35	28	100	101	107	150
19.	e	45	30	60	65	15	133	78	95	90
20.	e	40	56	58	50	8	10	103	50	73
21.	e	70	25	113	35	20	50	34	20	28
22.	e	85	110	70	50	65	70	35	25	40
23.	e	49	20	50	60	20	50	40	32	42
24.	e	35	60	40	15	20	55	42	48	50
25.	e	30	48	32	35	28	55	35	38	34
26.	e	73	54	78	95	90	160	67	85	75
27.	e	55	68	103	90	75	70	75	115	125
28.	e	95	45	182	70	140	85	40	28	95
29.	e	38	21	25	45	40	80	24	35	43
30.	e	15	55	59	45	15	30	75	115	125
31.	e	38	3	58	68	75	130	44	50	85
32.	e	35	48	80	97	60	100	63	55	142
33.	e	90	50	58	95	30	45	80	125	133
34.	e	12	44	20	15	18	30	60	30	86
35.	e	44	45	10	55	72	5	64	24	19
36.	e	18	27	17	75	55	56	30	35	27
37.	e	48	105	80	15	45	50	26	30	59
38.	e	24	59	53	36	70	55	54	20	85
39.	e	24	16	43	90	42	18	8	18	25
40.	e	13	40	45	20	14	12	30	33	36
41.	e	42	90	60	20	24	25	50	70	92
42.	e	30	250	182	42	35	34	80	52	75
43.	e	3	68	28	8	28	74	50	160	61
44.	e	40	38	57	25	68	22	47	44	115
45.	e	43	35	15	30	49	18	20	45	18

a<sub>0</sub> - 1 host larva per female parasite  
 a<sub>1</sub> - 2 " " " "  
 a<sub>2</sub> - 3 " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " 8 - 10 mg  
 b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

c<sub>0</sub> - sex ratio of parent parasite population - 1:1 (Female:Male)  
 c<sub>1</sub> - " " " " - 2:1 ( " )  
 c<sub>2</sub> - " " " " - 3:1 ( " )  
 c<sub>3</sub> - " " " " - 1:2 ( " )  
 c<sub>4</sub> - " " " " - 1:3 ( " )



APPENDIX - III

Progeny production of *B. brevicornis* at different temperature and humidity combinations

Sl. No.	Treatments	28°C and 75% RH			30°C and 60% RH			32°C and 50% RH		
		RI	RII	RIII	RI	RII	RIII	RI	RII	RIII
1.	a <sub>0</sub>	19	22	13	22	20	16	29	15	22
2.	a <sub>1</sub>	11	26	3	29	11	20	24	36	34
3.	a <sub>2</sub>	3	9	3	7	2	2	13	3	21
4.	b <sub>0</sub>	36	31	13	15	55	34	9	19	22
5.	b <sub>1</sub>	9	2	12	26	9	5	4	7	23
6.	b <sub>2</sub>	4	8	2	13	5	11	21	15	2
7.	c <sub>0</sub>	2	2	2	27	8	20	6	7	3
8.	c <sub>1</sub>	17	6	7	16	37	13	9	6	10
9.	c <sub>2</sub>	15	3	11	3	4	5	11	3	3
10.	c <sub>3</sub>	8	10	4	4	5	12	7	2	34
11.	c <sub>4</sub>	4	14	2	8	2	14	41	17	47
12.	c <sub>0</sub>	19	9	37	3	4	12	19	29	49
13.	c <sub>1</sub>	7	21	2	6	5	23	45	3	37
14.	c <sub>2</sub>	11	13	19	2	4	12	24	13	27
15.	c <sub>3</sub>	11	12	11	3	14	7	5	31	4
16.	c <sub>4</sub>	18	19	17	12	13	24	69	24	25
17.	c <sub>0</sub>	12	3	18	17	23	26	32	79	82
18.	c <sub>1</sub>	4	18	19	7	10	18	24	26	8
19.	c <sub>2</sub>	5	6	10	7	4	12	8	3	15
20.	c <sub>3</sub>	3	4	18	30	3	34	16	7	6
21.	c <sub>4</sub>	24	7	20	9	9	23	10	3	9
22.	c <sub>0</sub>	21	20	10	15	25	21	7	5	23
23.	c <sub>1</sub>	15	9	20	7	3	4	4	10	5
24.	c <sub>2</sub>	5	19	18	12	13	10	10	5	18
25.	c <sub>3</sub>	4	10	9	9	9	6	31	8	7
26.	c <sub>4</sub>	11	16	5	39	32	30	18	19	11
27.	c <sub>0</sub>	7	5	11	17	26	19	30	34	24
28.	c <sub>1</sub>	6	9	30	5	15	23	7	5	15
29.	c <sub>2</sub>	17	12	2	15	7	20	9	25	22
30.	c <sub>3</sub>	5	6	27	11	5	14	2	25	13
31.	c <sub>4</sub>	8	2	23	10	18	24	22	24	48
32.	c <sub>0</sub>	9	7	5	26	56	10	10	18	33
33.	c <sub>1</sub>	32	2	2	19	17	2	12	20	42
34.	c <sub>2</sub>	7	22	11	7	6	13	36	24	48
35.	c <sub>3</sub>	22	5	2	15	2	2	14	10	3
36.	c <sub>4</sub>	5	3	3	24	22	41	16	12	20
37.	c <sub>0</sub>	15	7	14	4	11	13	8	4	6
38.	c <sub>1</sub>	6	19	11	12	10	14	3	7	13
39.	c <sub>2</sub>	10	3	16	13	9	6	6	5	12
40.	c <sub>3</sub>	8	10	12	9	8	7	8	13	21
41.	c <sub>4</sub>	2	15	8	12	15	14	38	38	46
42.	c <sub>0</sub>	6	18	18	20	17	21	47	24	38
43.	c <sub>1</sub>	2	6	9	3	9	33	20	68	29
44.	c <sub>2</sub>	7	2	5	10	7	10	34	9	35
45.	c <sub>3</sub>	6	3	6	24	25	8	13	33	8

a<sub>0</sub> - 1 host larva per female parasite  
a<sub>1</sub> - 2 host larvae per female parasite  
a<sub>2</sub> - 3 host larvae per female parasite

b<sub>0</sub> - larval weight 30-35 mg  
b<sub>1</sub> - " " 8-10 mg  
b<sub>2</sub> - half of the hosts of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

c<sub>0</sub> - Sex ratio of parent population - 1:1 (Female : Male)  
c<sub>1</sub> - " " " - 2:1 ( " )  
c<sub>2</sub> - " " " - 3:1 ( " )  
c<sub>3</sub> - " " " - 1:2 ( " )  
c<sub>4</sub> - " " " - 1:3 ( " )

APPENDIX - IV

Number of female progeny of *B. brevicornis* produced at different temperature and humidity combinations

Sl. No.	Treatments	28°C and 75% RH			30°C and 60% RH			32°C and 50% RH		
		RI	RII	RIII	RI	RII	RIII	RI	RII	RIII
1.	a	9	8	2	11	10	1	16	12	17
2.	a	3	17	1	6	3	15	14	10	27
3.	a	2	4	1	3	1	1	6	1	2
4.	a	21	9	6	5	36	3	4	13	13
5.	a	3	1	4	12	3	1	1	5	12
6.	a	2	3	1	3	4	1	10	11	1
7.	a	3	1	1	17	5	5	5	5	2
8.	a	4	1	4	10	7	11	2	3	4
9.	a	4	1	8	2	3	4	7	1	2
10.	a	3	5	2	1	4	2	3	1	2
11.	a	4	5	1	4	1	7	18	11	25
12.	a	14	2	26	1	1	5	7	13	1
13.	a	6	15	1	5	2	5	24	1	28
14.	a	5	4	10	1	2	7	17	2	7
15.	a	8	7	9	2	2	4	4	22	1
16.	a	5	10	10	7	10	11	52	13	15
17.	a	5	1	11	1	14	12	26	51	43
18.	a	2	14	8	2	1	6	13	12	3
19.	a	2	3	5	1	3	3	3	1	13
20.	a	2	2	5	22	2	2	8	6	1
21.	a	15	2	3	2	4	14	1	1	4
22.	a	11	7	7	9	13	14	4	2	11
23.	a	6	3	6	1	2	4	2	4	1
24.	a	2	4	3	5	3	2	66	3	7
25.	a	3	8	7	6	5	1	10	6	3
26.	a	3	6	1	7	10	14	4	15	2
27.	a	1	6	5	7	6	10	13	17	13
28.	a	2	3	5	10	3	17	2	1	7
29.	a	10	7	9	4	3	15	3	11	7
30.	a	3	2	1	7	4	10	1	16	8
31.	a	7	5	12	1	4	12	14	4	21
32.	a	1	1	4	4	8	1	4	1	15
33.	a	11	3	4	17	36	1	2	7	6
34.	a	4	1	6	8	11	1	16	10	22
35.	a	9	12	1	4	2	12	12	5	1
36.	a	2	3	1	7	1	1	4	5	7
37.	a	4	2	2	7	11	16	4	5	4
38.	a	4	5	9	3	9	4	5	2	8
39.	a	5	9	5	8	1	4	1	4	8
40.	a	3	1	6	6	1	3	2	2	4
41.	a	3	5	1	8	2	6	4	9	9
42.	a	1	9	6	8	3	9	8	25	8
43.	a	2	10	9	7	6	2	6	6	16
44.	a	1	3	4	1	4	5	10	37	17
45.	a	1	1	3	13	12	5	7	24	2

a<sub>1</sub> - 1 host larva per female parasite  
 a<sub>2</sub> - " " " "  
 a<sub>3</sub> - " " " "

b<sub>0</sub> - larval weight 30 - 35 mg  
 b<sub>1</sub> - " " " 8 - 10 mg  
 b<sub>2</sub> - half of the larvae of the 'b<sub>0</sub>' type and the other half of the 'b<sub>1</sub>' type

o<sub>0</sub> - sex-ratio of parent parasite population - 1:1 (Female:Male)  
 o<sub>1</sub> - " " " " 2:1  
 o<sub>2</sub> - " " " " 3:1  
 o<sub>3</sub> - " " " " 1:2  
 o<sub>4</sub> - " " " " 1:3



**STUDIES ON THE REGULATION OF  
PROGENY PRODUCTION AND SEX-RATIO  
OF *BRACON BREVICORNIS* WESMAEL**

(HYMENOPTERA : BRACONIDAE)

**By**

**SOSAMMA JACOB**

**ABSTRACT OF A THESIS**

Submitted in partial fulfilment of the  
requirements for the degree of  
MASTER OF SCIENCE IN AGRICULTURE  
Faculty of Agriculture  
Kerala Agricultural University

Department of Agricultural Entomology  
COLLEGE OF HORTICULTURE  
Vellanikkara :: TRICHUR

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## ABSTRACT

The fecundity, progeny production, female-male composition of the progeny and duration of development of *Bracon brevicornis* Wesmael as influenced by the density and size (weight) of host larvae of *Corycaea canaliculata* Stainton and the sex-ratio of the parent parasite population were studied at three temperature-humidity (TH) combinations, viz., 28°C and 75% RH (TH<sub>1</sub>), 30°C and 60% RH (TH<sub>2</sub>) and 32°C and 50% RH (TH<sub>3</sub>).

The maximum fecundity of the parasite was registered consistently at all the three TH levels at a host density level of two larvae per female parasite. The exclusive use of light weight larvae (weight range 8 to 10 mg) as hosts produced significantly less number of eggs at the TH<sub>1</sub> and TH<sub>2</sub> levels, while at the TH<sub>3</sub> level the fecundity was maximum when the heavier host larvae (weight range 30 to 35 mg) were used. The parental sex-ratio levels of 2:1 and 3:1 (female:male) led to the production of higher number of eggs than under the ratios of 1:1, 1:2 and 1:3.

The maximum progeny production in *B. brevicornis* at the TH<sub>1</sub> and TH<sub>2</sub> levels was attained under a host density level of two larvae per female parasite. The progeny

production was relatively higher at the TH<sub>1</sub> and TH<sub>2</sub> levels when heavier larvae were exposed for parasitisation. The superiority of the parental sex-ratio of 2:1 (female:male) in the production of higher number of progeny was established at the TH<sub>2</sub> and TH<sub>3</sub> conditions.

Significant influence of the host larval density on the female progeny production was detected at the TH<sub>1</sub> and TH<sub>2</sub> levels. The female progeny production was maximum at a host larval density level of two per female parasite at the TH<sub>2</sub> combination, while this was found to be the highest at a density level of one larva per female parasite at the TH<sub>1</sub> level. Relatively heavier host larvae produced significantly higher number of female offsprings at the TH<sub>2</sub> and TH<sub>3</sub> combinations. The parental sex-ratio level of 2:1 (female:male) consistently produced maximum number of female progeny.

The influence of host larval density on the proportion of females was pronounced only at the TH<sub>3</sub> level and a density level of two larvae per female parasite was found to be better with reference to the production of a higher proportion of females. The sex-ratio of the parent parasite population influenced the proportion of females in the F<sub>1</sub> progeny and a ratio of 1:3 (female:male)



produced higher proportion of females at the TH<sub>2</sub> and TH<sub>3</sub> combinations.

The host larval density, weight of host larvae and the sex-ratio of the parent parasite population exerted significant influence on the duration of development of *B. brevicornis*.

The feasibility of manipulating the ambient temperature-humidity conditions for maintaining laboratory cultures of *B. brevicornis* at the maximum possible levels has been discussed.