

PRNC100

PRNC 100

PUERTO RICO NUCLEAR CENTER

RADIATION-INDUCED STERILITY FOR POPULATION CONTROL
OF THE SUGARCANE BORER (DIATRAEA

Summary Progress Report

1964 - 1967

Principal Investigator, S. Galka

OPERATED BY UNIVERSITY OF PUERTO RICO UNDER CONTRACT
NO. AT (40-1)-1892 FOR U. S. ATOMIC ENERGY COMMISSION

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PROJECT SUMMARY REPORT: July, 1964 - June, 1967

Potential for Gamma-Induced Sterility

Control of the Sugarcane Borer

(eab.) to

Puerto Rico

uch supported by the USABC Division
of Biology and Medicine under

contract no. AT (40-1) - 1833

Report prepared April, 1967, by David

Walker

Principal Investigator, Puerto Rico Nuclear Center

Mayaguez, Puerto Rico

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24 Original Objectives

1, to determine if the sugarcane borer, *Distraco saccharalis*

(ob.), can be rendered sterile by subjecting it to gamma irradiation,

and

2. to study the bionomics of this species with reference to

phases of the life cycle that are applicable to a mass-release program

in Puerto Rico.

B. The Problem in Relation to the Economy

The sugarcane borer *Digtraea saccharalis* is tropical and sub-

tropics in distribution and is known to infest crops from northern

Louisiana to Peru and Brazil. It is of economic importance to the

Production of sugarcane, corn, milo, Fico, lemongrasses, pasture and

forage grasses. Principal crops affected in Latin America are sugar

cane, lemongrass, citronella, and corn. Serious losses of sugar

production have been reported in the United States, Mexico, Guatemala,

El Salvador, Nicaragua, Costa Rica, Panama, Colombia, Puerto Rico and the rest of the "West Indies. Heavy infestations have been reported in commercial plantings of Lemongrass and citronella grass in Guatemala. World annual loss is difficult to estimate. Mortorell (1965) reports an annual loss of three million dollars to the Puerto Rican sugar industry and Charpentier, et al (1965) estimate six million dollars in United States cane production. The infestation rate is two or three times higher in the continental United States than in Puerto Rico (Charpentier, et al, 1965).

Damage to young plants of less than two months age is more severe than in older cane. Infestation in young plants is called "dead heart" because the terminal bud is killed by the feeding larva. Usually these young plants are killed and when there is widespread damage to young plants the whole field must be replanted. Older plants are more resistant to attack and usually outgrow damage. Yield is greatly

reduced by the borer through stunted growth, reduction in sucrose content, sugar inversion, and increased susceptibility to plant
Atseases ond other Insect posts

In Puerto Rico cane mortality is high when the young succulent
ants are subjected to heavy attack by cane borers and aphids at the
same tine

Under natural conditions, tho sugarcane borer lays ies eget
fon the leaf of the sugsreane plant. After hatching and initial feeding
fon the leaf during the first ond second stages, the third stage Larva
tunnels into the plant stem inside of the leaf axil, where it 1s
Protected from virtually all types of conventional insecticidal

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applications that are of practical use under comercial rowing

Conditions, Pupation takes place in this tunnel within the stem.

During the early summer months when sunlight, temperature, and rainfall are optimal for rapid cane growth the larval period can be completed in twenty-five days, However average Larval growth period is 35 days and during the less favorable times of the year the Larval period is more than 45 days.

When diapause occurs it takes place as fifth or sixth stage larvae and lasts two or three months. In Puerto Rico larvae begin diapause in December. It is believed that the onset of diapause is controlled by hormonal factors that are controlled by exposure to short-day length in the preceding generation. If this is so, the regulatory mechanism must be in very delicate balance because the change in day length in Puerto Rico is less than one hour from summer to winter. Adults and larvae are most abundant during the summer, because population increase is dependent upon the rapid cane growth caused by high rainfall and high temperature. Thus, optimum conditions for larval growth and reproduction occur in Puerto Rico during April, May and June. After August the cane plant is not as suitable for supporting larvae because of its saturation.

The borer's long Larval Life-span and short adult life-span

are the main biological obstacles inherent in using the Gass-release
?method for population control of this species.

In order of their effectiveness natural control agents are:

ants (egg predator); *Telchogracna sinutum* (Hymenoptera, Chalcididae)s
(eae. predator); *Lixophags diateses* (Diptera, Larvavoridae); mtcro~
Srganises such as *Cordycepe* spp. (attack larvae and pupae); and other
pathogenic fungi and bacteria. ?The most effective of these natural
Eonerel agents Kill. the Larvae before they have had a chance to

Damage the plant, Natural control by predators and parasites say

Teach 70L during dry periods although this varies considerably through=
ue the year.

?The practice of burning standing cane before harvest and of
burning cane trash after harvest has helped reduce the level of adult
population. Probably re-infestation takes place from alternate hoots
each year under these cultural practices.

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©. Relation of Other Work to PREC Program

?The USDA and other Agencies have several research and operational programs devoted to eradication of insects by the sterile male release technique, These include the following insects under investigation:

Dipters

Scrow-worm fly (operational phase)

Melon fly

Mexican fruit fly

Oriental fruit fly (operational phase)

Mediterranean fruit fly (advanced field testing phase)

Olive fruit fly

Lepidoptera

oypsy soth

Codling moth

Pink bollworm,

Corn earworm

European Cornborer

Oriental fruit moth

This List includes projects in which sterility is radio-

induced or chemically-induced,

The Diptera projects are in a more advanced state than the

Lepidoptera projects. In general the plant-feeding Diptera are much more easily mass-produced under artificial conditions than Lepidoptera.

This is due to their simpler dietary requirements, higher resistance to bacteria and fungi, behavior differences and to other factors. In addition the food requirements of plant-feeding Diptera are better known than those of the Lepidoptera. Yet many of our damaging agricultural pests are Lepidoptera.

The discovery and subsequent evaluation of the importance of sex pheromones has added a new potential in control of Lepidoptera, This method is particularly promising for the suppression of those insects that have a short adult life span and that pass through certain immature stages in the soil, in plant stems and fruits, or in other places that are inaccessible by conventional chemical control methods; of those insects for which conventional control measures are ineffectual or only partly effective. This approach is new and so little knowledge has been gained from studies of it that it is difficult to accurately predict the future potential of its usefulness. Natural chemical attractants produced by the insect are usually very specific in their activity. Most of the preliminary biological testing has been done with natural substances produced by the insects themselves. During

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this phase of development the results have been spectacular. Solution of the problems associated with isolating the chemically-active substances and subsequently evaluating specific activity has been a very painstaking task in the gypsy moth project and in similar programs.

Baye fou of the substances have been isolated and chemically identified. Among those that are known are the cabbage Looper pheromone isolated by Berger (1966). Hensle, is attempting isolation and Chemical identification of the sex pheromone produced by female

D. seccharalis.

?Tests with synthetic pheromone have not been as conclusive as was anticipated. This is due to the fact that the synthetically manufactured substances contain more than the single laevorotatory isomer that serves as sexual releaser in the insect. Trace impurities neutralize or alter the effectiveness of the specific isomer.

Gamma-induced sterility of the codling moth has been studied by Proverbs (1962), spruce budworms by Berryman (1966), the pine shoot moth by Lee (1966), the European Corn Borer by Walker (1966), corn earworm by Garcia (1966), pink bollworm by Ouye, et al (1964) and the havel orange worm by Hussein and Madsen (1964). All of these projects are oriented to applied insect control. Gamma exposure doses required to sterilize the adults have been found for most of these insects, but the radiation effect on longevity, mating capacity, and behavior has not yet been fully studied in all of them.

Ti ACCOMPLISHMENTS

he reader is referred to Technical Reports 1 (1965), 2 (1966), and 3 (1967) for complete reports of this Project. Conclusions may be summarized as follows:

AL Sterility, Adult Longevity, Mating Behavior and Vigor:

sterilized by exposure to 30 Kilorads, and

?No apparent somatic damage occurred when? adults.

of either sex are exposed to 50 Kilorads or less (Walker, 1965).

Thirty-five Kilorads were chosen for sterilizing adults to be liberated

In cage tests. There was no apparent effect on longevity, oviposition

rate, fecundity nor mating ability by their treatment. There is

[No] evidence that mating capacity was actually increased by radi-

ation (Walker, 1967), we do not anticipate that there will be problems

associated with adverse effects due to irradiation in an eradication

program with the sugarcane borer,

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B. Delayed Leihal Effect:

Survival and egg hatchability have been measured in the offspring of adults subjected to exposure to sub-sterilizing level of radiation by Walker (1957). Hussein and Madeen (1964) report a similar effect in the navel orange worm moth, *Paranyelois transitella*, and North (1966) cites similar response with the eaBbage looper moth, *Tetchnoplusts af*. At exposures as low as 4 kr. there is a striking reduction in hatchability of the offspring from irradiated borer parents. This effect occurs regardless of whether the male or female parent (in the P generation) was treated. We believe that the delayed lethal effect is due to the character of the diffuse kinetochore (centromere) in Lepidoptera chromosomes. In the preliminary series of tests in which forty-nine matings were made the net hatch was 31 percent of the eggs produced by the P generation and 26 percent of the eggs produced by the F₁ generation,

At the present time the results of a second series of tests
are being evaluated. The following matings were made in the parental
(P) generation in this series

Mating combinations of P generation Adults

8

Series Male

a R (irradiated)

2 n

c R

D[®]

Exposures were made at 1, 2 and 4 Kr. in this second series.

The survival rate of Fy larvae as well as rate of growth are

being scored. After ecdysis these F₁ adults have been used to test the effect on egg hatch in the F₂ generation.

The F₁ mating tests were made in April and May, 1967 and the F₂ rate of egg hatch 12 nov being observed.

In these tests the lethal effect was greater on egg hatch= ability of the F₂ generation than F₁ generation. In F₂ matings we expect the greatest lethal effect in C x C sibling matings, and the least effect in the A or B with normal outbreeding matings, If we have enough time to do so, we will continue the delayed lethal effect into the fourth generation. Our preliminary results indicate that the Lethal effect can be observed in larval survival of the F₂ generation.

©. Cage Testing:

Eight cages were built on the College Farm in April, 196

These were built to observe population change under field conditions

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+6.

Each cage is 40 ft. wide, 40 ft. long and 10 ft. high. The supports are made of 1 1/2" galvanized steel pipe and are covered with parafilm screen. The screen has a 37 percent shade factor. Mesh size is 60 openings per square inch. We release sterile and normal adults into these cages and measure the reproductive rate. Reproductive rate is

measured by harvesting the corn plants and extracting the Fy Larvae from them,

Results from 14 tests are shown in Table 1. Mean increase

in tests where normal males and females were released was approximately six fold. In comparison competition tests yielded less variability. By comparison the rate of infestation in corn plants

outside the cage were nearly double the highest infestation within the cage (18.9% of stalks infested outside as compared with 11.0% inside the cage).

Hose Rang:

Two hundred species of grass have been examined for
Biatraca larvae utilizing specimens from the living grass auseua at
the University in Mayaguez subject to heavy borer attack. Larvae were
found in twelve species of these grasece a5 shown in Table 2,

E. Diet Toot

Our ultimate goal is to develop an inexpensive food that
fulfills the following criteria:

high rate of survival to aduie

2. rapid development through Larval stages

3. production of vigorous adults as measured by length of
Life span, oviposition rato and mating frequency.

Artificial diet studies have gone through three develop:

?mental phases since the project began. ?In the first phase a number of diets made from sugarcane fiber, cane stem, and cane leaf extracts were tested. Some of these contained supplementary ingredients for increased vitamin and protein content. Also tested were numerous modifications of diets that had been successfully used for rearing other species of Lepidoptera. Resulting survival was very low and the few survivors that reached adulthood developed very slowly. The work began when Dr. Kenneth Dagen of the University of Tennessee recommended carrot powder supplement. Fifty formulations of this basic diet were tested. From the best three formulations further modifications were tested to determine if any specific ingredient could be substituted or eliminated. The resulting PRN diet was the culmination of this second phase. This diet contains brewer's yeast, carrot powder, corn stalk fiber, ascorbic acid, agar, and mold inhibitor.

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Table 2

Known Hosterange of *Diatraea sacchsralis* in Puerto Rico

Spanish English

Scientific Name Common name Use

1. Zea mays corn

2. Saccharum officinarum sugarcane Fe

3. Sorghum vulgare milo

4. Cynodon dactylon crabgrass Fe

5. Cymbopogon nardus lemongrass re

6. Eleusine indica barnyard grass P

7. Cynchiza latifolia ?

8. Paspalum conjugatum wild rice c

9. Pennisetum glaucum pearl millet c

10. Setaria viridis foxtail 7 ©

11. Pennisetum polystachyon guinea grass | P

12. Oryza sativa rice re

© of no commercial value. ?Grasses used as pasture grass (P), field crop (FC), and wild grass

(©) of no commercial value.

Plants are Listed in the order of susceptibility to attack by
Diatrsea Larvae. Some of these plants are not common comercial
Varieties here. No doubt there are other hosts among other native
Brasses not in the collection.

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In the third phase we are sousing to improve survival rate

and promote faster development. Formulations to which we have added

ingredients known to be required by other Lepidoptera are being

tested, Tests have been made with alpha-tocopherol, wheat germ oil,

corn oil, and mineral supplements. None of these have improved our
efficiency.

Further diet costs 1411 be based upon the known nutritional
requirements of phytophagous lepidoptera. Vanderzant and other USDA
Investigators are testing essential requirements by working with
holistic diets (holly defined). then it becomes possible to predict

the general nutritional requirements of phytophagous Lepidoptera, we
will be able to select natural food materials that supply large
quantities of the specific requirements.

In general, Lepidoptera larvae are considerably more fastidious in their food requirements than diptera larvae. Probably the specific nutritional requirements change during growth, since substances required during early larval stages may not be essential later, and vice versa. The narrow host-range of many species of lepidoptera confirms this.

The long larval growth period is a particularly serious problem in artificially rearing this species. The food decomposes during the twenty to thirty day period required for larval growth. Because the larvae tunnel into the food, extracting them from the food is difficult and time-consuming, it is necessary to extract them when the food decomposes and they must be transferred to fresh food. There are two possible solutions to this problem:

(Q)_to develop a diet that {# nutritionally superior so that
Larvae complete development before the food decomposes,

(2) to select a fi
Laboratory culture.

growing genetic strain of Diatraea for the

We have been studying both of these approaches. Relative to
the first, larval life span during winter has been shortened from 45

days in nature to 30 days or less in the laboratory as a result of
Improved diet. Even under best conditions the fastest developing Larvae
required 17 days. In relation to the second possibility, tests

have been made with 197 lines with short larval life span. Apparently
larval life span is a function of many genetic factors. Even within

the best pure strain there was a great deal of variation in development
rate among offspring. Lines were also selected for longer adult life
span. Variation among these lines was less than in the larval experiment.

It will be difficult to develop a laboratory strain that has a short and uniform growth-rate and is long-lived in the adult stage.

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F. Rearing Method and Handling Techniques

The rearing and handling techniques presently utilized to maintain over a colony of approximately 10,000 larvae are the result of testing several factors, as these bear on feeding response. Larval feeding behavior suggests that the stimulant to taste food (phagostimulant) is an olfactory response that develops after hatching, whereas taste stimulates actual feeding. Immediately after hatching young larvae venter about over the food surface. Later

they feed vigorously after first sampling the food. Conditions of food content, Light intensity, temperature, humidity and carbon dioxide level have been studied, to provide relatively satisfactory larval feeding response

Table 3 summarizes the life span of *Diatraça sacch*:

in the Laboratory on PRK diet. In artificial culture more larvae die in the first stage before feeding than at any other time during their life. During the period of early larval growth (first to third larval stadia) a large surface area with rough texture is needed to promote vigorous feeding. At the third stage when the larvae bore tunnels in the food they need large volume of food. Overcrowding, causes accidental cannibalism if there is insufficient food mass for [larval] making.

We have tested the PANG diet in containers of different size

?and shape to obtain the optimn relation of surface area and volune.

Glass shell-vials (32 ml.) with cotton stoppers have produced the best

yield. ?Rearing in trays would be easier since they can be filled and

cleaned easily. Mortality has been high in tray teats because of

dessication. Dr. Frank Novell (1965) suggested pouring the food in

trays and spraying the surface with paraffin wax to prevent drying.

?This method is successful for laboratory rearing the codling woth

Survial wi aca Larvae in pareffin-coated

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Table 3

Synopsis of Ltfe-span of *D. saccharalie*

tn the laboratory?

Lifespan of Length Activity at this Main Cause of

Stage Stadium? = Stage Death

(ays)

Fee. 4 to 9 0.6 to 0.8 embryonic development would

o

First 4 to 18 2 to 4 surface feeding diet and

Larval) starvation

Second 3 to 18 3 to 7 tunnel forming bacteria

Larval (6),

Third 3 to 18 6 to 18 tunnel forming would and bacteria

Larval (7)

Fourth 5 to 21 15 to 29 tunnel forming, stops would and bacteria

Larval = (11) feeding before pupating _ starvation

Pupal 5 to 10-18 to 27 quiescent in tunnel accidental cannot be

© by tunneling Larval,

desiccation, bacteria

Adult 1 to 11 15 to 25 rests during day, nuptial old age, desiccation

Male (3.5) Flight and mating at night, old, starvation

Adult 1 to 1318 to 29 same as males oviposited old age, desiccation,

Female (5.2) during day and night old, starvation

Total 25 to 107-2 to 29 -

as)

Yerom Walker, 1966

FRange, the average (8 shown in parenthesis).

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TIT- PUBLICATIONS

The following have been published:

1964. Bionomics of the Sugarcane Borer *Diatraea*

saccharalis (Fab.). TIT. Oviposition Rate,

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PP. (in press)

1966, Improved Kenic diets for che Sugarcane Borer

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1966, Potential for control of the Sugarcane Borer

Through Radio-induced etertlity, TABA/FAO

Symposium on Radiation, Radiotsotopes and Rearing

Methods in the Control of Ingect Poste, Oct. 17-21,

1966, Tel Aviv, Israel, in press.

B. ?The following are in preparation:

1, Gaumarinduced sterility in *D. saccharalis*.

2. Mortality-staging in eggs of 2

Lectures have been given in Central America in cooperation with the Atoms for Peace Program. Lectures were given at the National Universities and Graduate Instructional Centers in Guatemala, El Salvador, Nicaragua and Costa Rica. In addition, S. Garcia-Rivera has been trained to study the potential for controlling the corn earworm, *Heliothis virescens* in El Salvador and Juan Rodriguez in Nicaragua has been given similar training to evaluate the potential with *Laphygma frugiperda*, the "cogollero del maiz". Both of these pests cause serious damage to crops in Central America. Training was also given for study of the coffee leaf miner, and the pests affecting lemon grass in Guatemala and El Salvador.

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Iv- sumaRY

?The ultimate objective of the project has been to determine if the sugarcane borer could be eradicated by the release of sterile adults, Implicit in this objective are a number of related aspects

1. to determine the stage cut point within the stage to be sterilized,

2. to determine if the life-span, fecundity, and general vigor is affected by sterilizing,

ing capability

3. to develop a laboratory method for mass-producing this agent capable of producing large numbers of individuals at a low cost,

4. to study the Life cycle in the Laboratory and under field conditions with particular reference to growth rate, reproductive potential, mating behavior, and other aspects of the bionomes ecology relevant to a successful overflooding program.

n

?The first objective has been completed. Only the adult stage can be sterilized without producing excessive somatic damage and subsequent high mortality. Larval and pupal stages can be sterilized at much lower exposures. The dose/sterility effect relationship appears to be roughly exponential, but there is considerable variation in individual tests at sub-sterilizing exposures.

Life span and oviposition rate are not affected by sterilizing the adults of either sex. Mating capability of sterile adults is equal or slightly superior to normal adults. This is true whether the individuals have been reared on the host-plant or from artificial diet. We have succeeded in rearing this species in the Laboratory at small numbers at high labor cost.

The artificial diet developed is well-adapted to rearing the insect. If the insect is consistently provided with fresh food. Since 17 to 30 days are passed in various stages it is not possible to maintain the food free of bacterial and mold growth. If larvae are transferred to fresh food before contamination becomes too heavy survival is improved. The high labor investment of manually removing larvae from contaminated so fresh food is a problem given considerable attention but is not yet solved.

The second major problem in mass-rearing is to develop a low cost diet. The ingredients for our present diet cost approximately two dollars per thousand larvae. This is about one hundred fold greater than the cost of producing phytophagous diptera and about three hundred times the cost of mass-producing the screw worm. High food-cost and

contamination problems are inherent in rearing lepidoptera under artificial conditions:

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We have obtained considerable data on growth rate in Laboratory rearing but there have not been extensive field studies on this insect in Puerto Rico. We sorely need data on the population of adults in

the field, and the variation of adult populations throughout the year.

Unfortunately we have not been able to study this because of lack of personnel. Some indirect inferences on the net natural reproductive rate can be made from our field tests of cages, but it would be risky to

extrapolate this data to field conditions of cane production. It is expected that the study of Hensley (1965) on the sex pheromone will provide us with greatly improved methods for trapping adult males and that this method will enable us to make accurate estimates of field population. Hensley's group expects to be able to make a chemical analysis of the pheromone in the near future.

The potential for eradicating this species through a combined program of trapping using synthetic pheromone in combination with the release of sterile adults appears to be promising.

Limitations of this program are: 1) lack of a good rearing method, and 2) incomplete knowledge of the biology of this species

under field conditions. Since we were not responsible for these under the 1964 Memorandum of Understanding we did not direct our attention to these problems. Unfortunately the cooperating agencies were unable to solve these problems before terminating their programs.

The PRNC commitment proposed in project objectives under the Memorandum of Understanding between the USDA Entomology Research Division, UPR Agricultural Experiment Station, and PRNC will be completed at the end of Fiscal Year 1967. Further field work and a mass rearing method will have to be developed before the program will be ready for large scale field testing. A logical sequence of necessary stages is shown in Table 4 of the Appendix.

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/~ RECOMMENDATIONS

Let the work be continued with the objective of population suppression or eradication by release of sterile adults. I recommend concentrating on these objectives:

1. Field studies, to determine

a. fluctuation of the adult population in nature,

BI hose range 12 Puerto Rico,

1 rate of population suppression by sterile releases in nature,

4, effect of multiple releases.

for subsequent

2. Laboratory studies, with the aim

a. to develop an improved diet for mass-rearing,

BI to develop rearing and handling techniques for mass

rearing,

to study the lethal effect over several successive

generation

4, to select an approved genetic strain for laboratory rearing.

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VIT- APPENDIX

?TABLE 4

Developnant Stages in a Species

Eradication Prograa®

seep = 30 so,00ne0 [1 7 7

sen s0 w sotoonexe [# _]

seep = 100 e sx0.0mee [>

Series

for species that have not been raised under artificial
?conditions and species for which the biology is poorly known.

Series B: for species that can be grown in the laboratory and for
which the biology is fairly well-known,

Series C: for species that can be mass-reared in the laboratory
and for which the biology is well-known.

?This was prepared by John Munro and David Walker at the request
of the participants of the Symposium on Insect Sterility and
Mass-rearing Methods in Tel Aviv, 1967. It will be included as
part of the published proceedings of this meeting.

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Step 1. Using field-collected material should test the following

against all stages of Life cycle, and should include differences in response within stage (if possible):

2. Dose response of:

1) lethality and sterility

2) Lifespan

3) oviposition rate

4) mating behavior

Bb. normal performance for the above as a basis for comparison.

Step 2, Laboratory Rearing

The principal objective of this phase of the program is to develop a reliable method of producing the insect in the Laboratory under artificial conditions. One should strive to develop a method that produces predictable results, not necessarily mass-rearing method. Cost of production should not be an important criterion, and it is recommended that natural (host plant) food substances be tested.

a. Diets

Starting with the natural food material, prepare diets using added food substances to augment nutritional level. Diet tests should be performed on the neonatal larvae, and all instars should be tested because specific nutritional requirements are apt to change during the course of the development.

Criteria to be measured should include: Length of time required for each stage (i.e. time between molts), rate of weight increase (length and diameter), amount of food consumed, pupal weights, longevity of adult life span and of the pupal stage, observation of morphological and behavior aberrations in adults (crumpled wings, active movement, mating behavior, etc.), and survival rate over several generations of laboratory rearing.

b. Develop handling techniques, and test containers in preparation for larger production. With the techniques and diet developed at this stage, one should strive for results that will give the assurance of a predictable survival. The survival rate obtained at this phase may not be as high as needed for mass-production.

. If possible a laboratory strain of the species should be selected for the qualities desired. ?Those qualities might include

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short larval life-span, long adult Life-span, high reproductive rate,

high rate of survival and vigorous adults having high mating capability.

4. In addition, if a distinct phenotype, f.e, one that can be easily recognized in the field by casual observation, can be selected, it would greatly help in later field work.

Step 3. Bionomics and Ecology in nature

Diurnal behavior, including daily soverment, host-range and host-preference, time of day and frequency of mating in nature, natural abundance, seasonal population fluctuation, and other pertinent aspects of the ecology should be emphasized at this stage of the progr

It is of no use to be able to produce sterile adults if these adults are not suitable for their intended use. The sterile adults released should be equal to or superior to adults in nature in respect to longevity and mating capacity. High reproductive rate in the laboratory strain is desirable as it facilitates laboratory rearing.

Step 4. Small Scale Field test

?Test the mating ability, behavior and longevity of sterile adults under natural conditions in contained field environment. The test site should be isolated from the greater environment, preferably Ae should be a small island or an isolated land mass surrounded by suitable geographic barriers such as mountains.

Size and seasonal change in the natural population should be measured before, during and after this test. Trapping methods, marking techniques? and distribution studies can be the natural tools? {oF ?measuring population changes in this phase of the prope

Step 5. Mass-rearing Methods

?The diet selected for mass-culture must be cheap, and should be of materials of uniform quality that are readily available. The food selected for mass-rearing should be easy to prepare in large quantities.

Disposal of food after use may be a problem in some Programs. Attention should be given to obtaining large numbers of vigorous adults.

Step 6. Eradication Progr:

Measurement of effectiveness, stringent quarantine measures for the future, suitable release and recapture methods, low-cost, and

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related factors are likely to be the most important considerations in this phase of the program,

General Comments

Cost will escalate drastically as the program proceeds from phase 4 to 5 and again from phase 5 to 6, Rough estimates of increase are about @ five-fold increase from phase 4 to 5, and 100 to 200 fold from phase 5 to 6.

The policy of determining the limiting factors is most likely to prevent success should receive the most attention in early stages of the program rather than leaving them till later,

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