

TRIBOLIUM INFORMATION BULLETIN

Number 17

Editor: A. Sokoloff, School of Natural Sciences

California State College, San Bernardino

California

1974

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TRIBOLIUM INFORMATION BULLETIN

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Foreword

It is hoped that *Tribolium* Information 17 will be as useful to others as previous issues of this bulletin have been. In spite of the general scarcity of grant funds available from federal agencies it is apparent from the bibliography and the research notes that Tribolium continues to be researched as intensely as it was some 20 years ago when the editor began his investigations on flour beetles.

While the condition of the *Tribolium* Information Bulletin may be termed good, the same cannot be said for the condition of the *Tribolium* Stock Center. The section on Cellular Biology of the National Science Foundation from which the Stock Center received its support has decided not to provide further support to the *Tribolium* Stock Center as well as the majority of insect stock centers. (Fuller details will be available in the Genetics supplement for 1974).

Except for the timely appearance of a research grant to the Editor which will enable him to maintain some of the more valuable mutant stocks, and an institutional grant which provides for student help to maintain other stocks, the *Tribolium* Stock Center would be in serious difficulty. The responsibility for maintenance of genetic stocks has now been shifted to the section on Systematics, Biology of the National Science Foundation.

It is hoped that this subsection dealing with Biological Resources will recognize that in these times of scarcity a shift

in priorities, particularly toward greater support of centers for maintenance of mutant stocks of useful experimental organisms, is imperative.

For the preparation of TIB 17 I am indebted to Janice Brown, Pat Cavataio, Susan Clark, and Jim Gooch.

The publication of TIB-17 was made possible by Army Research Office Grant RDRD LP-11790-LS.

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ANNOUNCEMENTS

Triboliumists are advised of the following International Conferences or Congresses, to be held October 7-11, 1974:

1. The first International Working Conference on Stored-Product Entomology will be held in Savannah, Georgia, U.S.A. October 7-11, 1974. The Conference will cover all phases of stored-product entomology and will include symposia, panel discussions, and submitted papers. Facilities will be available for informal get-togethers as they are required. A Proceedings will be published following the Conference.

Anyone desiring further information should contact:

Organizers

First International Working Conference on Stored-Product Entomology
c/o Stored-Product Insects Research and Development Laboratory,
ARS-USDA
P.O. Box 5125
Savannah, Georgia 31403
U.S.A.

2. The First World Congress on Genetics Applied to Livestock Production will be held in Madrid, Spain, October 7-11, 1974. The Congress will be held in the Palacio Nacional de Congresos y Exposiciones of Madrid, on the Avenida del Generalisimo, 29, tel. 270 56 00/09, 270 28 01/05 and 270 58 00.

The Palacio, considered as one of the most modern and best equipped halls in all Europe, is essentially made up of three different units according to need, although they are entirely intercommunicable: Congress Area, Exhibition Area and Parking Area.

For further details contact the General Secretariat, located on Departamento de Genetica, Facultad de Veterinaria, Ciudad Universitaria, Madrid-3, Spain, tel. 243 94 59 (Prof. Dr. Carlos Luis de Cuenca). All inquiries regarding information, registration, scientific works and other data should be relayed to the address.

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Stock Lists

BALTIMORE, MARYLAND
THE JOHN HOPKINS UNIVERSITY
SCHOOL OF HYGIENE AND PUBLIC HEALTH

Tribolium confusum: bI and bIV
Tribolium castaneum: cI and cIV-a

Michael Nathanson

BERKELEY, CALIFORNIA
UNIVERSITY OF CALIFORNIA,
DONNER LABORATORY AND LAWRENCE RADIATION LABORATORY

Tribolium confusum

1. "+" - a wild type strain derived from Genetics Department, University of California, Berkeley.
2. Black - an autosomal semi-dominant body color mutant.
From 1.
3. Miniature - an autosomal recessive body size mutant.
From 1.
4. Short elytra - an autosomal dominant elytrum size mutant.
Low viability in adults, indicating a recessive Lethal gene.
5. Blistered elytra - an autosomal recessive mutant. Low viability.

Tribolium brevicornis

Wild type derived from Genetics Department, University of California, Berkeley.

BRIDGEPORT, CONNECTICUT
UNIVERSITY OF BRIDGEPORT,
DEPARTMENT OF BIOLOGY

Tribolium confusum

Wild type strains derived from Dr. Fraenkel's laboratory at the University of Illinois.

(Ed.)

Stock Lists

BURLINGTON, NORTH CAROLINA
CAROLINA BIOLOGICAL SUPPLY COMPANY

Tribolium castaneum

- | | |
|----------|---------|
| 1. black | Chicago |
| 2. jet | |
| 3. pearl | |
| 4. wild | McGill |

Tribolium confusum

- | | |
|---------|----------|
| 1. wild | Carolina |
| | (Ed.) |

CARBONDALE, ILLINOIS
SOUTHERN ILLINOIS UNIVERSITY,
DEPARTMENT OF ZOOLOGY

I. Base populations

1. Purdue + foundation
2. Purdue s foundation (sooty)
3. Purdue b foundation (black)

II. Mutant

1. paddle (pd)
2. spotted (sp)
3. pygmy, red, paddle (py r pd)
4. pygmy (py)
5. red (r)
6. pygmy, red (py r)
7. pygmy, paddle, spotted (py pd sp)
8. pearl (p)
9. white (w)
10. ruby, light ocular diaphragm (rb lod)
11. Short antenna (Sa)
12. chestnut (c)
13. antennapedia (ap)
14. squint (sq)

III. Selected populations

Early: a population subjected to selection for a short larval peroid. Origin in Purdue Wild Foundation.

Stock Lists

Late: a population subjected to selection for a long larval period. Origin in Purdue Wild Foundation.

D. C. Englert

CARLISLE, PENNSYLVANIA
DICKINSON COLLEGE, DEPARTMENT OF BIOLOGY

Tribolium confusum

I. Wild type strains

1. Six strains started from females captured in a feed bin in New York City, 1955.
2. Three strains, one each from T. Park, Chicago; J. Stanley, Montreal; S. Smith, Sault Ste. Marie, Canada.
3. One strain consisting of several above strains mixed together about three years ago.
4. One strain started with individuals taken from (1) above, which has been freed of eye mutations.

NOTE: Some of the wild strains listed in (1) and (2) are known to be carrying pearl-like mutations.

II. Mutant

1. Black - Sault Ste. Marie (1956)
2. Ebony - Chicago (1957)
3. Eyespot - sex-linked - from a wild strain in (I.1) above (1959)
4. Rough - from strain (II.1) above (1957)
5. Split - from a wild strain in (I.1) above (1956)
6. Striped - sex-linked - from (II.1) above (1957)
7. One strain each of Striped/black and split/black

Oryzaephilus surinamensis

One strain started from insects captured in New York City, 1955.

(Ed.)

CHARLOTTESVILLE, VIRGINIA
UNIVERSITY OF VIRGINIA,
DEPARTMENT OF BIOLOGY

Tribolium castaneum

I. Wild type strains

- | | |
|---------------------------------|-----------------------|
| 1. Chicago | University of Chicago |
| 2. Purdue University Foundation | via Stony Brook |
| 3. Synthetic | San Bernardino |

Stock Lists

II. Mutant strains

1. McGill black

University of Chicago
via Stony Brook

CHICAGO, ILLINOIS
UNIVERSITY OF ILLINOIS AT CHICAGO CIRCLE,
DEPARTMENT OF BIOLOGICAL SCIENCES

Tribolium castaneum

"Chicago" - a wild type strain (originally from Thomas Park)

"Brazil" (also known as ci) - a wild type strain (originally from Rio de Janiero)

cII - an inbred strain (derived from "Brazil")

cIII - an inbred strain (derived from "Chicago")

cIV-a - an inbred strain (derived from "Chicago")

Tribolium confusum

"Chicago" - a wild type strain (originally from Thomas Park)

"Ebony" - an autosomal recessive body color mutant

bi - an inbred strain (derived from "Chicago")

bII - an inbred strain (derived from "Chicago")

bIII - an inbred strain (derived from "Chicago")

bIV - an inbred strain (derived from "Chicago")

COLLEGE PARK, MARYLAND
UNIVERSITY OF MARYLAND,
DEPARTMENT OF ZOOLOGY

I. Wild type strains

A. Tribolium castaneum

1. Chicago (via Sokoloff) Berkeley, 1964
2. University del Valle-1 Cali, Colombia, 1964
3. University of Maryland-2

Inbred strains.

4. E 2 (originally from Edinburgh, via Boylan) Manitoba, 1964

B. Tribolium confusum

1. So. Illinois University-1 Carbondale, Ill., 1962

Stock Lists

Inbred strains

2. CFI-11

Berkeley, Calif., 1965

II. Mutant

1. T. confusum

Berkeley, Calif., 1959

2. Ebony (e^{L&H})

(Ed.)

CORAL GABLES, FLORIDA
UNIVERSITY OF MIAMI,
DEPARTMENT OF BIOLOGY

I. Wild type strains

1. Tribolium confusum

Chicago

2. Tribolium castaneum

Chicago

II. Mutant

1. Tribolium castaneum - "jet"

Chicago

2. Tribolium castaneum - pearl type, origin
in local stocks3. Tribolium castaneum - jet x pearl4. Tribolium confusum - "ebony"

(Ed.)

CORVALLIS, OREGON
OREGON STATE UNIVERSITY,
DEPARTMENT OF ZOOLOGY

I. Wild type strains

A. Tribolium castaneum

1. Berkeley

Berkeley, 1966

2. Chicago

Urbana, 1966

3. del Valle

Maryland, 1966

4. Oregon

Urbana, 1967

5. Vivarium

Urbana, 1967

Stock Lists

B. Tribolium confusum

1. Berkeley	Berkeley, 1966
2. Chicago	Urbana, 1966
3. Kansas	Kansas, 1966
4. Oklahoma	Urbana, 1966
5. Oregon	Urbana, 1967

II. Mutant strains

A. Tribolium castaneum

1. sa-2 (+/s)	Berkeley, 1966
2. dve, pd	Berkeley, 1967
3. fas-3a	Berkeley, 1967
4. b, mc, p	Berkeley, 1966
5. bal, c (+/s)	Berkeley, 1966
6. pd	Urbana, 1966
7. Be	Berkeley, 1966
8. mc	Berkeley, 1967
9. aa (+/p)	Berkeley, 1967
10. ctp, ju	Berkeley, 1966
11. Mo	Berkeley, 1966
12. b ^D	Berkeley, 1966
13. ap ^D , s	Berkeley, 1966
14. j	Berkeley, 1966
15. p, s, ap ^D	Berkeley, 1966
16. Fta, c	Berkeley, 1966
17. ser, py, r	Berkeley, 1967
18. Spa (+/c)	Berkeley, 1966
19. r ^s , s	Berkeley, 1967
20. sq	Berkeley, 1967
21. h	Urbana, 1967
22. sh ^s	Berkeley, 1967
23. p, lod	Berkeley, 1967
24. Sa-2, s	Berkeley, 1967
25. r	Urbana, 1968

B. Tribolium confusum

1. sh, b	Berkeley, 1966
2. mag ^{AS}	Berkeley, 1967
3. we	Berkeley, 1966
4. dj, p, lod	Berkeley, 1966
5. thu ^s , a	Berkeley, 1966
6. dj	Berkeley, 1966
7. a	Urbana, 1966
8. thu	Berkeley, 1966
9. p	Berkeley, 1966
10. p ₁ , lod	Berkeley, 1966
11. h ₁	Urbana, 1967
12. thu ₁	Berkeley, 1966
13. hle ₁	Urbana, 1967
14. r ₁	Urbana, 1968
15. dep _c	Urbana, 1969
16. sh _c	Corvallis, 1970

DAVIS, CALIFORNIA
UNIVERSITY OF CALIFORNIA, DEPARTMENT OF ANIMAL HUSBANDRY

I. Wild type strains (T. castaneum)

BC1 T. castaneum Berkeley, 1967
DC1 T. castaneum Davis, 1969

II. Mutant strains

BC2 T. castaneum, sooty Berkeley, 1967
BC114 T. castaneum, sooty, inbred from strain 14a Berkeley, 1967
SCp T. castaneum, pearl eye San Bernardino, 1969

III. Selected strains (all derived from BC1)

6-14 BC1-2, lines 1-8, 10, selected for large 21-day pupa for 23-36 generations.
15 BC1-2, line 9 selected for 48 generations; average 21-day pupa weight 6-10 mg.
16-18 BC1-2L, lines 1-3, selected for small 21-day pupa for 30 generations.

IV. Wild type strains (T. confusum)

BF1 T. confusum Berkeley, 1967
DF1 T. confusum Davis, 1967
DF3 T. confusum Davis, 1969

V. Mutant strains

SFp (pearl eyes) San Bernardino, 1969
G. A. E. Gall

DENTON, TEXAS 76204
TEXAS WOMAN'S UNIVERSITY
DEPARTMENT OF BIOLOGY

A. Wild type strains

1. Tribolium confusum Chicago
2. Tribolium castaneum Chicago

B. Mutant strains

1. Tribolium castaneum "sooty" San Bernardino
H. E. Erdman

EAST LANSING, MICHIGAN
MICHIGAN STATE UNIVERSITY, BIOLOGY RESEARCH CENTER

Tribolium castaneum

I. Wild type strain

1. McGill

Stock Lists

II. Mutant strains

1. paddle
2. spotted

Chicago via Berkeley,
Berkeley,

(Ed.)

EAST LANSING MICHIGAN
MICHIGAN STATE UNIVERSITY, DEPARTMENT OF ZOOLOGY

Tribolium confusum

I. Wild type strain

1. Chicago wild, Chi +/-

Berkeley, 1964

II. Mutant strains

1. ruby eyespot (rus)
2. melanotic stink glands (msgHo)
3. light ocular diaphragm, pearl (lod p)
4. black, melanotic stink glands, ruby eyespot (b msg rus)
5. black, ruby eyespot (b rus)
6. McGill black, light ocular diaphragm, pearl (McGill b lod p)

Berkeley, 1964

Berkeley, 1964

Berkeley, 1964

Berkeley, 1964

Berkeley, 1964

Berkeley, 1964

Tribolium castaneum

Wild type strain
black strain

GAINESVILLE, FLORIDA
ARS, USDA
P.O. BOX 14565
INSECT ATTRACTANTS, BEHAVIOR AND BASIS BIOLOGY LAB.

<u>Attagenus megatoma</u>	black carpet beetle
<u>Cadra cautella</u>	almond moth
<u>Cylas formicarius elegantulus</u>	sweet potato weevil
<u>Ladioderma serricornis</u>	cigarette beetle
<u>Oryzaephilus surinamensis</u>	sawtoothed grain beetle
<u>Paramyelois transitella</u>	navel orangeworm
<u>Plodia interpunctella</u>	Indian meal moth
<u>Sitotroga cerealella</u>	Angoumois grain moth
<u>Sitophilus oryzae</u>	rice weevil
<u>Tribolium castaneum</u>	red flour beetle
<u>Trogoderma granarium</u>	khapra beetle
<u>Trogoderma inclusum</u>	

HAMPTON, IOWA
FARMERS HYBRID COMPANY

I. Wild type strain

1. Chicago

via Berkeley, 1965

Stock Lists

II. Mutant strains

1. r py
2. j mc
3. Be/+

HUMACAO, PUERTO RICO
 UNIVERSITY OF PUERTO RICO, COLLEGE OF HUMACAO
 DEPARTMENT OF BIOLOGY

Tribolium castaneum

I. Wild type strain

1. Chicago

II. Mutant strains

1. paddle
2. pearl
3. Microcephalic
4. Bar eye, sooty
5. Short antennae (Sa-2)

HUNTSVILLE, TEXAS
 SAM HOUSTON STATE UNIVERSITY
 BIOLOGY DEPARTMENT

Tribolium castaneum

I. Wild type strains

- A. Purdue University Foundation
- B. Huntsville, Texas wild type - source of squint-like (sl).

II. Mutant stains

- | | |
|------------------------------------------------------|-----------------------------------|
| A. Bar eye (<u>Be</u>) | Berkeley , 1962 |
| B. black (<u>b^D</u>) | Carbondale, Ill., 1961 |
| C. light ocular diaphragm (<u>lod^D</u>) | Carbondale, Ill., 1961 |
| D. maroon (<u>m</u>) | Purdue + Foundation , 1962 |
| E. microcephalic (pearl) (<u>mc</u> , <u>p</u>) | Chazy, New York , 1959 |
| F. paddle (<u>pd</u>) | Chicago , 1955 |
| G. peach (<u>r^{ph}</u>) | Carbondale, Ill., 1961 |
| H. pygmy (<u>py</u>) | Chazy, New York , 1960 |
| I. pink, ivory (<u>p^{Pk}</u> , <u>i</u>) | Chazy, N. Y.: Purdue + Foundation |
| J. ring (<u>rg</u>) | Purdue + Foundation , 1961 |
| K. rose (<u>rs</u>) | Purdue + Foundation , 1964 |
| L. ruby (<u>rb</u>) | Carbondale, Ill., 1961 |
| M. ruby, jet (<u>rb</u> , <u>j</u>) | Carbondale, Ill., 1961 |
| N. ruby, peach (<u>rb</u> , <u>r^{ph}</u>) | Purdue + Foundation , 1956 |
| O. sooty (<u>s</u>) | Chazy, New York , 1960 |
| P. squint (<u>sq</u>) | Huntsville, Texas , 1973 |
| Q. squint-like (<u>sql</u>) | |

Stock Lists

IMMACULATA, PENNSYLVANIA
IMMACULATA COLLEGE, CANCER RESEARCH UNIT

I. Wild type strains

1. <u>Alphitobius diaperinus</u>	PIL
2. <u>Alphitobius laevigatus</u>	PIL
3. <u>Gnathocerus cornutus</u>	PIL
4. <u>Gnathocerus maxillosus</u>	PIL
5. <u>Latheticus oryzae</u>	Berkeley
6. <u>Tenebrio molitor</u>	PIL
7. <u>Tenebrio obscurus</u>	PIL
8. <u>Tribolium anaphae</u>	Berkeley
9. <u>Tribolium brevicornis</u>	Berkeley
10. <u>Tribolium castaneum</u>	Berkeley
11. <u>Tribolium confusum</u>	Berkeley
12. <u>Tribolium destructor</u>	Berkeley
13. <u>Tribolium madens</u>	Berkeley

II. Mutant Strain

1. Tribolium confusum melanotic stink glands (msg)

Note: The insect strains formerly maintained by one of us (S.K.L.) at the John Hopkins University, Chemistry Department, in Baltimore, Md., have been transferred to Immaculata College.

(Ed.)

IRVINE, CALIFORNIA
UNIVERSITY OF CALIFORNIA, DEPARTMENT OF ORGANISMIC BIOLOGY

Tenebrio molitor

(Ed.)

ITHACA, NEW YORK
CORNELL UNIVERSITY, DEPARTMENT OF ANIMAL SCIENCE

Tribolium castaneum

The Purdue Foundation wild type obtained from the Population Genetics Institute in April, 1965.

(Ed.)

Stock Lists

ITHACA, NEW YORK
CORNELL UNIVERSITY, DEPARTMENT OF ENTOMOLOGY AND LIMNOLOGY

I. Wild type strains

1. Tribolium confusum from Dr. H. Ducoff, University of Illinois.
2. Tribolium confusum infected with Nosema whitei.

(Dr. L. V. Knutson, same department, is said to have a wild type strain of T. confusum. Whether this strain is the same as that listed above is not known. Ed.)

JAMAICA, NEW YORK
ST. JOHN'S UNIVERSITY, DEPARTMENT OF BIOLOGY

Tenebrio molitor

(Ed.)

KENT, OHIO
KENT STATE UNIVERSITY, DEPARTMENT OF BIOLOGICAL SCIENCES

I. Wild type strains

A. Tribolium castaneum

Synthetic strain combined from Chicago wild type derived from Dr. Thomas Park and a strain obtained from Dr. Karl Schurr, Department of Biology, Bowling Green State University, Bowling Green, Ohio.

B. Tribolium confusum

Derived from stock maintained by Dr. L. V. Knutson, Department of Entomology, Cornell University, Ithaca, New York.

C. Oryzaephilus surinamensis

From infested flour.

Stock Lists

KINGSTON, R. I. 02881
 UNIVERSITY OF RHODE ISLAND
 DEPARTMENT OF ZOOLOGY

Tribolium castaneum

I. Wild type strains

Purdue Foundation	Via Purdue
Black Foundation	Via Purdue
Pearl Foundation	Via Purdue

II. Mutant strains

Unsaturated fatty acid sensitive (<u>cos</u>)	
Melanotic stink glands (msg)	Berkeley, 1964

Tribolium confusum

<u>T. confusum</u> , Chicago	Park, 1955
	R. F. Costantino

LAFAYETTE, INDIANA 47907
 PURDUE UNIVERSITY
 ANIMAL SCIENCES DEPARTMENT

Tribolium castaneum

I. Wild Type strains

A. Base populations for quantitative genetics studies:

1. Foundation + - wild type population synthesized in 1954 from a broad genetic base and maintained with no artificial selection and minimum of inbreeding.
2. Foundation s - same genetic base as Foundation + but marked with sooty (s).
3. Foundation b - synthesized in 1959 and marked with black (b), unrelated to Foundation +, broad genetic base, no selection, minimum inbreeding.
4. Foundation p - synthesized in 1959 and marked with pearl (p), unrelated to Foundations + and b, broad genetic base, no selection, minimum inbreeding.

Stock Lists

B. Laboratory stocks

5. Arkansas	Fayetteville, 1954
6. Brazil	Vicosa, 1958
7. Capetown	South Africa, 1958
8. Chicago	University of Chicago, 1954
9. Carbondale	Illinois, 1958
10. Columbia	South America, 1958
11. Florida	Gainesville, 1958
12. Georgia	Tipton, 1954
13. Japan	Kyoto, 1958
14. McGill	Montreal, Canada, 1958
15. Minnesota	Minneapolis, 1958
16. Texas	College Station, 1954
17. Virginia	Blacksburg, 1954

II. Mutant strains

18. antennapedia, <u>ap</u>	Purdue <u>Sa</u> Stock, 1962
19. black, <u>b^D</u>	Carbondale, Illinois, 1964
20. chestnut, <u>c</u>	Purdue + Foundation, 1961
21. cordovan, <u>bcd</u>	Purdue + Foundation, 1962
22. corn oil sensitive, <u>cos</u>	Purdue + Foundation, 1966
23. ivory, <u>i</u>	Purdue + Foundation, 1961
24. jet, <u>j^E</u>	Purdue + Foundation, 1961
25. maroon <u>m</u>	Purdue + Foundation, 1962
26. paddle, <u>pd</u>	Chicago, 1955
27. peach, <u>r^{PH}</u>	Carbondale, Illinois, 1964
28. pearl, <u>p^S</u>	Fla. Inbred, (Purdue), 1963
29. pygmy, <u>py</u>	Chazy, New York, 1960
30. red, <u>r^S</u>	Purdue + Foundation, 1964
31. ring, <u>rg</u>	Purdue + Foundation, 1961
32. rose, <u>rs</u>	Purdue + Foundation,
33. ruby, <u>rb</u>	Carbondale, Illinois, 1964
34. Short antenna, <u>Sa</u>	Purdue + Foundation, 1960
35. short antenna, <u>sa₃</u>	Purdue + Foundation, 1966
36. sooty, <u>s</u>	Purdue + Foundation, 1956
37. squint, <u>sq</u>	Chazy, New York, 1960
38. wine <u>r</u>	Purdue + Foundation, 1963

(Ed.)

LARAMIE, WYOMING
UNIVERSITY OF WYOMING, DEPARTMENT OF ZOOLOGY AND PHYSIOLOGY

Tribolium castaneum

I. Mutant strains

1. Fta c
2. Be s
3. pd py pte
4. sp
5. Spa s eju
6. p b
7. p lod
8. ap sq

(Ed.)

Stock Lists

LAURINGBURG, NORTH CAROLINA
ST. ANDREWS COLLEGE

Tribolium confusum

A wild stock that is infected with Nosema whitei.

(Ed.)

LEXINGTON, KENTUCKY
AGRICULTURAL EXPERIMENT STATION
UNIVERSITY OF KENTUCKY

Tribolium castaneum

I. Base Populations

- | | |
|---------------------------------------|--------|
| 1. Purdue <u>t</u> foundation | Purdue |
| 2. Purdue <u>s</u> foundation (sooty) | Purdue |
| 3. Purdue <u>b</u> foundation (black) | Purdue |
| 4. Purdue <u>p</u> foundation (pearl) | Purdue |

II. Wild strains

- | | |
|--------------------------------|----------------|
| 1. 4 strains collected locally | Kentucky, 1970 |
|--------------------------------|----------------|

III. Inbred Lines

- | | |
|------------------------------------------------------------------|-------------------------|
| 1. CSI-5 | Berkeley via Minnesota |
| 2. CSI-10 | Berkeley via Minnesota |
| 3. E-1 | Edinburgh via Minnesota |
| 4. E-2 | Edinburgh via Minnesota |
| 5-9. Five Inbred lines derived
from different wild
strains | Purdue |

IV. Selected Strains

Several strains which have been selected for increased 21 day pupa weight.

R. Goodwill

Stock Lists

LIVERMORE, CALIFORNIA
BIOLOGICAL FRONTIERS INSTITUTE

Only wild type strains of T. confusum and T. castaneum are maintained. We have a number of stocks of these species received from Dr. I. Michael Lerner and described by him in Tribolium Information Bulletin #3 (p.28). In addition we have a number of stocks of both species collected locally.

We have also a wild type strain of the saw-toothed grain beetle, Oryzaephilus surinamensis (L.)

(Ed.)

LORETTO, PENNSYLVANIA
ST. FRANCIS COLLEGE, BIOLOGY DEPARTMENT

I. Wild type strain

- | | |
|-------------------------------|----------------------|
| 1. <u>Tribolium confusum</u> | Chicago via Berkeley |
| 2. <u>Tribolium castaneum</u> | Chicago via Berkeley |

(Ed.)

LOS ANGELES, CALIFORNIA
UNIVERSITY OF CALIFORNIA MEDICAL CENTER
DEPARTMENT OF MEDICAL MICROBIOLOGY

I. Wild type strain

- | | |
|------------------------------|----------------------|
| 1. <u>Tribolium confusum</u> | Chicago via Berkeley |
|------------------------------|----------------------|

(Ed.)

MANHATTAN, KANSAS
DEPARTMENT OF ENTOMOLOGY
KANSAS STATE UNIVERSITY

LEPIDOPTERA

Phycitidae

Cadra cautella (Walk.), almond moth, from USDA, Manhattan, Kansas, 1971.

Plodia interpunctella (Hbn.), Indian-meal moth, Kansas.

Gelechiidae

Sitotroga cerealella (Oliv.), Angoumois grain moth, Kansas, about 1970.

- A. Sitotroga cerealella (Oliv.), Red-eyed Angoumois grain moth, from stock cultures, 1967.

Stock Lists

COLEOPTERA

Anobiidae

- Lasioderma serricorne (F.), Cigarette beetle, Kansas, 1966.
Stegobium paniceum (L.), Drugstore beetle, from USDA,
Richmond, Virginia, 1971.

Bostrichidae

- Rhyzopertha dominica (F.), Lesser grain borer, Kansas.

Bruchidae

- Callosobruchus maculatus (F.), Cowpea weevil, Kansas, 1971.

Cucujidae

- Cryptolestes ferrugineus (Steph.), Rusty grain beetle, Kansas.
Cryptolestes pusillus (Schon.), Flat grain beetle, Kansas.
Oryzaephilus surinamensis (L.), Saw-toothed grain beetle,
Kansas.
Oryzaephilus mercator (Fauv.), Merchant grain beetle, from
USDA, Savannah, Georgia, 1964.

Curculionidae

- Sitophilus granarius (L.), Granary weevil, Kansas.
Sitophilus oryzae (L.), Rice weevil, Kansas, 1955.
Sitophilus zeamais Mots., Maize weevil, from Stuttgart,
Arkansas, 1955.
Sitophilus zeamais Mots., Maize weevil, from Veracruz,
Mexico, 1964.

Dermestidae

- Megatoma piceus (Oliv.), Black carpet beetle, Kansas.
Trogoderma inclusum LeC., Larger cabinet beetle, from USDA,
Manhattan, Kansas.
Trogoderma sternale Jayne, Manhattan, Kansas, 1971.
Trogoderma variabile Ballion, Kansas.

Ostomatidae

- Tenebroides mauritanicus (L.), Cadelle, Kansas.

Ptinidae

- Gibbium psylloides (Czemp.), Spider beetle, Chicago, Ill.,
1966.

Stock Lists

Silvanidae

Ahasverus advena (Waltl.), Foreign grain beetle, Manhattan, Kansas, 1969.

Tenebrionidae

Palorus ratzeburgi (Wissm.), Small-eyed flour beetle, Kansas, 1965.

Tenebrio molitor L., Yellow mealworm, Kansas.

Tenebrio obscurus F., Dark mealworm, Manhattan, Kansas, 1971.

Tribolium castaneum (Hbst.), Red flour beetle, Kansas.

Tribolium confusum J. du V., Confused flour beetle, Kansas.

R. B. Mills

MIDLAND, MICHIGAN

THE DOW CHEMICAL COMPANY, BIOPRODUCTS DEPARTMENT

Tribolium confusum

Wild strain maintained in laboratory more than 20 years.

(Ed.)

MILWAUKEE, WISCONSIN 53201
THE UNIVERSITY OF WISCONSIN
ZOOLOGY DEPARTMENT

Wild type strains

1. Purdue Foundation +
2. Purdue Foundation b

Selected strains

1. Late: a population subjected to selection for a long larval period. Origin in Purdue Foundation +.
2. High chaetae: a population subjected to selection for increased pregenital chaetae number. Origin in Purdue Foundation b.
3. Low chaetae: a population subjected to selection for decreased pregenital chaetae number. Origin in Purdue Foundation b.

E. L. Lange

Stock Lists

MOSCOW, IDAHO
UNIVERSITY OF IDAHO, DEPARTMENT OF ENTOMOLOGY

- A. Tribolium castaneum - large and small selections, sooty marked, obtained from Berkeley last October.
- B. Tribolium madens from the Boise Valley area, Idaho, started in November, 1967.
- C. Tribolium confusum - probably of local origin, held under weekly subculturing for about three years.
- D. Tribolium castaneum - of local origin, subcultured largely at weekly intervals for about five years, started from a very few individuals surviving neglect of cultures previously, somewhat sporadically, maintained for several years in the laboratory.

(Ed.)

MUNCIE, INDIANA
BALL STATE UNIVERSITY, DEPARTMENT OF PHYSIOLOGY AND HEALTH SCIENCETribolium castaneum, large stock, from Purdue University.Tribolium castaneum, foundation stock, from Purdue University.NATICK, MASSACHUSETTS
U.S. ARMY NATICK LABORATORIES, PIONEERING RESEARCH LABORATORY

I. Wild type strains

Lepidoptera:

Anagasta kuhniella - USDA Lab., Georgia, 1969
Cadra cautella " " " "
Plodia interpunctella " " " 1964
Sitotroga cerealella " " " 1969
Tineola bisselliella - Univ. New Hampshire, Durham, N.H., 1965

Coleoptera:

Anthrenus flavipes - USDA Lab., Georgia, 1967
Attagenus megatoma " " " 1957
Cryptolestes pusillus - Kansas State Univ., Manhattan, Kansas, 1971
Dermestes maculatus - USDA Lab., Georgia, 1968
Gibbium psyllodes - Kansas State Univ., Manhattan, Kansas, 1971
Lasioderma serricorne - USDA Lab., Georgia, 1968
Oryzaephilus surinamensis - USDA Lab., Georgia, 1968
Palorus ratzeburgi - Kansas State Univ., Manhattan, Kansas, 1971
Rhyzopertha dominica - USDA Lab., Georgia, 1969
Sitophilus granarius - " " " 1968
Sitophilus oryzae " " " 1968

Stock Lists

- Tenebrio molitor - Univ. New Hampshire, Durham, N.H., 1965
- Tenebroides mauritanicus - USDA Lab., Georgia, 1968
- Tribolium audax - Univ. California, Riverside, Calif., 1971
- Tribolium brevicornis - Univ. California, Riverside, Calif., 1971
- Tribolium castaneum - USDA, Georgia, 1956
- Tribolium confusum - USDA, Georgia, 1969
- Tribolium destructor - Univ. California, Riverside, Calif.
- Tribolium madens - " " " "
- Trogoderma variable - NLABS, Natick, Mass., 1968

Mutant:

- Tribolium confusum - Ebony strain, A. Sokoloff, 1968

NORMAN, OKLAHOMA
UNIVERSITY OF OKLAHOMA, DEPARTMENT OF ZOOLOGY

Coleoptera

- Tribolium castaneum (Tenebrionidae) wild type, Chicago; Univ. of Chicago.

F. J. Sonleitner.

NORTHRIDGE, CALIFORNIA
SAN FERNANDO VALLEY STATE COLLEGE, DEPARTMENT OF BIOLOGY

Tenebrio molitor infested with gregarines.

(Ed.)

NOTRE DAME, INDIANA
UNIVERSITY OF NOTRE DAME, DEPARTMENT OF BIOLOGY

I. Wild type strains

- 1. CFI-11 Berkeley, 1965
- 2. CFI-22 Berkeley, 1965
- 3. CFI-11 x CFI-22 Berkeley, 1965
- *4. ND-11 Park, Univ. of Chicago, 1954

*Since 1956, maintained at the Air Force Weapons Laboratory,
Kirtland, A. F. B., New Mexico.

(Ed.)

PITTSBURGH, PENNSYLVANIA
DUQUESNE UNIVERSITY, DEPARTMENT OF BIOLOGICAL SCIENCES

I. Wild type strains

- 1. Tribolium confusum (Chicago) used Via Sokoloff
as interned host for Hymenolepis diminuta.

(Ed.)

Stock Lists

POCATELLO, IDAHO
IDAHO STATE UNIVERSITY, DEPARTMENT OF BIOLOGY

I. Wild type strains

Tribolium castaneum--Synthetic strain marked with sooty
from Berkeley.

Tribolium confusum--Synthetic strain from Berkeley.

(Ed.)

PULLMAN, WASHINGTON 99163
WASHINGTON STATE UNIVERSITY
DEPARTMENT OF ENTOMOLOGY

Tribolium confusum - synthetic wild type
Tribolium castaneum - sooty

Roger Akre

RICHLAND, WASHINGTON
BATTELLE-NORTHWEST, BIOLOGY DEPARTMENT

I. Wild type strains

- | | |
|-------------------------------------------------------|---------------------|
| 1. <u>Tribolium confusum</u> Duval (Chicago Standard) | Univ. of
Chicago |
| 2. <u>Tribolium castaneum</u> Herbst (Brazil cl) | Univ. of
Chicago |

II. Mutant strain

- | | |
|----------------------------------------------|------------------------------|
| 1. <u>Tribolium castaneum</u> Herbst (Sooty) | Univ. of Calif.,
Berkeley |
|----------------------------------------------|------------------------------|

(Ed.)

RIVERSIDE, CALIFORNIA
UNIVERSITY OF CALIFORNIA, DEPARTMENT OF ENTOMOLOGY

- | | |
|---------------------------------|------------------|
| A. <u>Cryptolestes turcicus</u> | PIL via Berkeley |
| B. <u>Gnathocerus cornutus</u> | PIL via Berkeley |
| C. <u>Tribolium anaphe</u> | PIL via Berkeley |
| D. <u>Tribolium destructor</u> | PIL via Berkeley |
| E. <u>Tribolium madens</u> | PIL via Berkeley |
| F. <u>Tribolium brevicornis</u> | PIL via Berkeley |

(Ed.)

SALT LAKE CITY, UTAH
UNIVERSITY OF UTAH, DEPARTMENT OF ZOOLOGY AND ENTOMOLOGY

I. Wild type strains

- | | |
|-------------------------------|-----------------------------------------|
| 1. <u>Tribolium confusum</u> | Park, Chicago, 1962 |
| 2. <u>Tribolium castaneum</u> | J. Laurie, Utah, 1962 |
| 3. <u>Tenebrio molitor</u> | W. P. Larsen, via S. Muliak, Utah, 1961 |
| 4. <u>Oryzaephilus</u> sp. | wild, Utah, 1962 |

II. Mutant strain

1. melanotic stink glands

(Ed.)

SAN BERNARDINO, CALIFORNIA
CALIFORNIA STATE COLLEGE, NATURAL SCIENCES DIVISION

I. Wild type strains

A. Tribolium castaneum

- | | |
|----------------------|---------------------------------|
| 1. Arkansas | Bell, 1970 |
| 2. Brazil | ex Park via Howard Erdman, 1963 |
| 3. Capetown | Bell, 1970 |
| 4. Chicago | Park, 1955 |
| 5. Columbia | Bell, 1970 |
| 6. Consejo | Spain, 1968 |
| 7. Davis | Davis, Calif., 1961 |
| 8. Georgia | Bell, 1970 |
| 9. Florida | Bell, 1970 |
| 10. Japan | Bell, 1970 |
| 11. McGill | Stanely, 1958 |
| 12. Sacramento | 1961 |
| 13. Texas | 1958 |
| 14. Veracruz, Mexico | 1963 |
| 15. Virginia | 1958 |

B. Tribolium confusum

- | | |
|-------------------|----------------|
| 1. Chicago | Park, 1955 |
| 2. Davis | 1961 |
| 3. McGill | Stanley, 1958 |
| 4. New York | 1961 |
| 5. Pennsylvania | McDonald, 1963 |
| 6. Sacramento | 1961 |
| 7. San Bernardino | 1968 |

C. Tribolium audax

- | | |
|--------|--------------|
| 1. PIL | Slough, 1971 |
|--------|--------------|

D. Tribolium anaphe

- | | |
|--------|--------------|
| 1. PIL | Slough, 1963 |
|--------|--------------|

E. Tribolium brevicornis

- | | |
|--------------|--------------|
| 1. Riverside | Calif., 1965 |
|--------------|--------------|

Stock Lists

- F. Tribolium destructor
1. PIL Slough, 1963
- G. Tribolium madens
1. PIL Slough, 1963
2. PIL Slough, 1971
- H. Latheticus oryzae
1. Kansas 1970
2. Savannah Georgia, 1970
3. Tifton Georgia, 1970
- I. Oryzaephilus surinamensis
1. Synthetic from Cold Spring, Harbor, N.Y. and Oakland, Calif. populations. 1968
2. San Bernardino 1968
- J. Cryptolestes turcicus
1. PIL Slough, 1963
- K. Stegobium paniceum San Bernardino, 1969
- L. Trogoderma inclusum USDA Lab., Fresno, 1968
- II. Synthetic strains
- A. Tribolium castaneum
1. Berkeley. Synthetic strain from six different laboratory strains marked with sooty. Prepared in 1958.
2. Berkeley. Synthetic strain from seven laboratory strains not marked with body color genes. Prepared in 1964.
- B. Tribolium confusum
1. Berkeley. Synthetic strain from six wild type laboratory strains not marked with body color genes. Prepared in 1958.
- III. Inbred lines
- A. Tribolium castaneum
1. Started 1971 from synthetic strain now in the 9th generation of brother-sister mating and not marked with sooty.
- B. Tribolium confusum
1. Started October, 1958, from the Berkeley synthetic strain (now in 90-99 generation of brother-sister mating, not marked with body color genes).
a. CFI-1
b. CFI-2
c. CFI-5
d. CFI-8
e. CFI-11
f. CFI-12

2. Started in 1964 from the Berkeley synthetic strain, (now in 56-58 generation of brother-sister mating, not marked with body color genes.)
- CFI-13
 - CFI-14
 - CFI-15
 - CFI-23
 - CFI-24

IV. Mutants

A. Tribolium castaneum

Chromosome I

- | | |
|---------------------------------------------|-----------------------|
| 1. paddle (<u>pd</u>) | Park, 1955 |
| 2. paddle-1 (<u>pd-1</u>) | Berkeley, 1965 |
| 3. red (<u>r</u>) | Chazy, New York, 1959 |
| 4. red (<u>r^{Ho}</u>) | Berkeley, 1962 |
| 5. red (<u>r^D</u>) | Berkeley, 1963 |
| 6. pygmy (<u>py</u>) | Chazy, New York, 1959 |
| 7. spotted (<u>sp</u>) | Chazy, New York, 1959 |
| 8. divergent elytra (<u>dve</u>) | Chazy, New York, 1959 |
| 9. truncated elytra (<u>te</u>) | Chazy, New York, 1959 |
| 10. platinum eye (<u>pte</u>) | Berkeley, 1965 |
| 11. pokey (<u>pok</u>) (as heterozygotes) | Berkeley, 1962 |
| 12. red modifier (<u>r^{Mr}</u>) | Berkeley, 1961 |
| 13. serrate (<u>ser</u>) | Berkeley, 1963 |
| 14. <u>pte pd</u> | |
| 15. <u>py pd</u> | |
| 16. <u>sp pd</u> | |
| 17. <u>py r pd</u> | |
| 18. <u>py r</u> | |
| 19. <u>te r</u> | |
| 20. <u>sp r</u> | |
| 21. <u>r pd</u> | |
| 22. <u>py r^{Mr}</u> | |
| 23. <u>pte py pd</u> | |
| 24. <u>r te Mr</u> | |
| 25. <u>sp dve py pd</u> | |
| 26. <u>ser py r</u> | |
| 27. <u>te-1</u> | |

Chromosome II

- | | |
|------------------------------------|-----------------------|
| 28. pearl (<u>p</u>) | Park, 1955 |
| 29. pink (<u>p^{Pk}</u>) | Chazy, New York, 1959 |
| 30. pegleg (<u>pg</u>) | Chazy, New York, 1959 |
| 31. <u>p pg</u> | |

Chromosome III

- | | |
|----------------------------------------------------------|-----------------------|
| 32. aureate (<u>au</u>) | Berkeley, 1965 |
| 33. McGill black (<u>mcGb</u>) | Stanley, 1964 |
| 34. Chicago black (<u>Cb</u>) | Park, 1955 |
| 35. Synthetic (<u>McGb/Cb</u>) | Chazy, New York, 1958 |
| 36. black (<u>b^{S-1}</u>) (Brazil background) | Berkeley, 1963 |
| 37. black (<u>b^S</u>) (Chicago background) | Chazy, New York, 1960 |

- | | | |
|-----|-------------------------------------------------------------|----------------|
| 38. | light ocular diaphragm (<u>lod</u>)
(pearl background) | |
| 39. | light ocular diaphragm (<u>lod^d</u>) | Deweese, 1971 |
| 40. | melanotic stink glands (<u>msg</u>) | Berkeley, 1964 |
| 41. | scar (<u>sc</u>) | Purdue, 1964 |
| 42. | tawny (<u>bt</u>) | PIL, 1965 |

Chromosome IV

- | | | |
|-----|---------------------------------------------|-----------------------|
| 39. | cut prothorax (<u>ctp</u>) | Berkeley, 1962 |
| 40. | elongated juvenile urogomphi (<u>eju</u>) | Berkeley, 1965 |
| 41. | fused antennal segments-2 (<u>fas-2</u>) | Berkeley, |
| 42. | incomplete mesosternum (<u>ims</u>) | Berkeley, 1962 |
| 43. | juvenile urogomphi (<u>ju</u>) | Berkeley, 1962 |
| 44. | reduced juvenile urogomphi (<u>rju</u>) | Berkeley, 1963 |
| 45. | Spatulate (<u>Spa</u>) | Berkeley, 1964 |
| 46. | deformed legs (<u>dfl</u>) | Chazy, New York, 1959 |
| 47. | sternites incomplete (<u>sti</u>) | Berkeley, 1963 |
| 48. | <u>fas-2s</u> | |
| 49. | mahogany (<u>my</u>) | |

Chromosome V

- | | | |
|-----|---------------------------------------------------|-----------------------|
| 50. | jet (<u>j</u>) | Park, 1955 |
| 51. | microcephalic (<u>mc</u>) | Chazy, New York, 1959 |
| 52. | fused antennal segments-3 (<u>fas-3</u>) (=agg) | Berkeley, 1961 |
| 53. | fused antennal segments-3a (<u>fas-3a</u>) | Berkeley, 1963 |
| 54. | <u>j spl mc</u> | |
| 55. | maroon (<u>m</u>) | Eddleman, 1970 |

Chromosome VI

- | | | |
|-----|------------------------------|-----------------------|
| 56. | Microphthalmic (<u>Mo</u>) | Chazy, New York, 1959 |
|-----|------------------------------|-----------------------|

Chromosome VII

- | | | |
|-----|-----------------------------------------|-----------------------|
| 57. | Short antenna (<u>Sa</u>) | |
| 58. | Short antenna (<u>Sa-1</u>) (=Gn) | Berkeley, 1959 |
| 59. | Short antenna (<u>Sa-2</u>) (=Ds) | Berkeley, 1962 |
| 60. | Short antenna (<u>Sa-3</u>) (=Cua) | Chazy, New York, 1959 |
| 61. | short antenna (<u>sa</u>) (=ca) | 1961 |
| 62. | chestnut (c) (ex Eddleman) | Berkeley, 1962 |
| 63. | blistered elytra (<u>ble</u>) | Berkeley, 1962 |
| 64. | short antenna (<u>sa-2</u>) (=vg) | Berkeley, 1962 |
| 65. | Fused tarsi and antennae (<u>Fta</u>) | Berkeley, 1962 |
| 66. | <u>Fta ble</u> | |
| 67. | <u>sa c</u> | |
| 68. | <u>Fta c</u> | |
| 69. | <u>Sa c</u> | |
| 70. | <u>Fta ca c</u> | |
| 71. | <u>ble c</u> | |

Stock Lists

Chromosome VIII

72. antennapedia (ap^D) Berkeley, 1962
 73. antennapedia (ap^S) (=fas-6) Berkeley, 1963
 74. squint (sq) Chazy, 1959

Chromosome IX

75. missing abdominal sternites (mas) Berkeley, 1964
 76. prothoraxless (ptl) Chazy, New York, 1959
 77. prothoraxless-1 (ptl-1) Berkeley, 1965
 78. partially pointed abdominal sternites (ppas) Berkeley, 1963

Chromosome X

79. abbreviated appendages (aa) Cold Spring Harbor, N. Y., 1961
 80. abbreviated appendages-1 (aa-1) Chazy, New York, 1960
 (=csp1)

Multichromosomal

Note: The Roman numerals indicated the linkage groups involved.
 The symbol ? means the linkage group for that gene has not been established.

81. py pd; p i, II
 82. sp; p i, II
 83. px; b i, III
 84. py r; lod i, III
 85. sp; j i, V
 86. pd; Mo i, VI
 87. sp; p; j i, II, V
 88. r; lod ii, III
 89. r; b ii, III
 90. r; b; Mo ii, III, VI
 91. r; b; mc ii, III, V
 92. r; mc ii, V
 93. b; Mo iii, VI
 94. j; Mo v, VI
 95. ju ctr c IV, VII
 96. Mo; sa vi, VII
 97. b (p) apt iii, (II), ?
 98. mc apt v, ?
 99. apt i v, ?
 100. Mo (c) mas vi, (VII), IX
 101. (p) b mas (ii), III ?
 102. p Bamp/+ ii, III ?
 103. Bamp/+ap^D iii ?, IX
 104. Bamp/+ ptlHoy iii ?, IX
 105. b max iii, ?

Stock Lists

106. i max V, ?
 107. au Npp IV, ?
 108. ap Npp VIII, ?
 109. Be au IV
 110. Fta ppa VII, ?
 111. nc ppa V, IX
 112. fas-3a pti^{Hoy} III, IX
 113. b ap^S III, VIII
 114. au ctp IV
 115. i ppa V, IX
 116. ppk V, IX
 117. rb m ?; V

Unassigned (but possibly in II)

- | | |
|----------------------------------------------------------------------------------------|-----------------------|
| 118. creased abdominal sternites (<u>cas</u>) | Berkeley, 1963 |
| 119. abnormal abdominal sternites (<u>aas</u>) | Berkeley, 1965 |
| 120. akimbo (<u>akb</u>) | Berkeley, 1964 |
| 121. alate prothorax (<u>apt</u>) | Berkeley, 1964 |
| 122. antennae and tarsi fused (<u>atf</u>) | Berkeley, 1961 |
| 123. ballooned (<u>bal</u>) | Berkeley, 1963 |
| 124. banjo (<u>bj</u>) | Chazy, New York, 1960 |
| 125. bead (<u>bd</u>) | Bell, 1967 |
| 126. bent tibia (<u>bt</u>) | Berkeley, 1961 |
| 127. Blunt abdominal and metathoracic
projections (<u>Bamp</u>) (possibly in III) | Berkeley, 1965 |
| 128. bowed femur (<u>bf</u>) | Berkeley, 1963 |
| 129. bowleg | Bell, 1967 |
| 130. bumpy (<u>by</u>) | Bell, 1966 |
| 131. Charcoal (<u>Chr</u>) | Berkeley, 1966 |
| 132. deflected epimera (<u>dep</u>) | Berkeley, 1964 |
| 133. deformed femur (<u>dff</u>) | Berkeley, 1964 |
| 134. deformed tibia (<u>dft</u>) | Berkeley, 1964 |
| 135. dented | Bell, 1967 |
| 136. diminutive appendages (<u>dim</u>) | Berkeley, 1966 |
| 137. elbowed antennae-1 (<u>elb-1</u>) | Berkeley, 1964 |
| 138. elongated elytra (<u>ele</u>) | Berkeley, 1964 |
| 139. elytra and tarsi affected (<u>eta</u>) | Berkeley, 1963 |
| 140. extra urogomphi (<u>eu</u>) (black) | Chazy, New York, 1960 |
| 141. fused antennal segments-1 (<u>fas-1</u>) | Chazy, New York, 1959 |
| 142. Fused antennal segments-4 (<u>Fas-4</u>) | Berkeley, 1963 |
| 143. Fused antennal segments-5 (<u>Fas-5</u>) | Berkeley, 1963 |
| 144. jagged antecoxal piece (<u>jac</u>) | Berkeley, 1964 |
| 145. knobby prothorax (<u>knp</u>) | Berkeley, 1966 |
| 146. lopped median groove (<u>lmg</u>) | Berkeley, 1964 |
| 147. maxillopedia (<u>max</u>) | Berkeley, 1965 |
| 148. miniature appendages (<u>ma^D r</u>) | Bell, 1967 |
| 149. Multi-urogomphi (<u>Mu</u>) | Bell, 1966 |
| 150. Nonpunctate prothorax (<u>Npp</u>) | Berkeley, 1965 |
| 151. padded prothorax (<u>pdp</u>) | Berkeley, 1965 |

Stock Lists

152.	pectinate antennae (<u>pec</u>)	Berkeley, 1964
153.	reduced gin traps (<u>rgt</u>)	Berkeley, 1965
154.	reduced pleurosternal suture (<u>rps</u>)	Berkeley, 1965
155.	reduced tarsi and antennae (<u>rta</u>)	Berkeley, 1966
156.	rough (<u>ro</u>)	Berkeley, 1964
157.	ruby (<u>rby</u>)	Berkeley, 1962
158.	rugose elytra (<u>rue</u>)	Berkeley, 1966
159.	scalloped prothorax (<u>scp</u>)	Berkeley, 1965
160.	short median abdominal projection (<u>smp</u>)	Berkeley, 1966
161.	short split spinasternum (<u>ssa</u>)	Berkeley, 1965
162.	split (<u>sp</u>)	1963
163.	split-back (<u>sb</u>)	Bell, 1966
164.	stumpy (<u>stu</u>)	Berkeley, 1965
165.	Tetra urogomphi (<u>Tu</u>)	Berkeley, 1965
166.	tiny (<u>ti</u>) (=ty)	1962
167.	umbilicus (<u>umb</u>)	Berkeley, 1964

B. Tribolium confusum

Chromosome I

1.	Striped (<u>St</u>)	McDonald, 1961
2.	eyespot (<u>es</u>)	McDonald, 1961
3.	light eyespot (<u>eslt</u>)	Berkeley, 1963
4.	red (<u>r</u>)	Berkeley, 1962
5.	antennae and elytra reduced (<u>aer</u>)	Berkeley, 1962
6.	labiopedia (<u>lp</u>)	Berkeley, 1962
7.	pointed abdominal segments (<u>pas</u>)	Berkeley, 1963
8.	thickened elytral tips (<u>tet</u>)	Berkeley, 1963
9.	lethal-1 (<u>l1</u>) (in heterozygotes)	Berkeley, 1962
10.	crumpled (<u>cru</u>)	Berkeley, 1964
11.	prothoraxless-like (<u>ptll</u>)	Berkeley, 1964
12.	<u>St es</u>	
13.	<u>es lp</u>	
14.	<u>es lp</u> (synthetic background)	
15.	<u>eslt lp</u>	
16.	<u>St es lp</u>	
17.	alate prothorax (<u>apt</u>)	Berkeley, 1965

Chromosome II

18.	pearl (<u>p</u>)	PIL, via Stanley, 1960
19.	pearl (<u>pS</u>)	Berkeley, 1962
20.	ebony-2 (<u>e2</u>)	PIL, via Stanley, 1960
21.	creased abdominal sternites (<u>cas</u>)	Berkeley, 1963
22.	dirty pearl eye (<u>dpe</u>) (=fro)	Berkeley, 1963
23.	<u>e2p</u>	PIL, via Stanley, 1960
24.	<u>p cas</u>	

Chromosome III

25.	Yugoslavian black (=b2)	Yugoslavia, 1969
-----	-------------------------	------------------

Stock Lists

- | | |
|---------------------------------------------|-----------------------|
| 26. McGill black (McGb) (=b ^{Ho}) | Stanley, 1960 |
| 27. black-3 (b-3) | Berkeley, 1964 |
| 28. ruby spot (<u>rus</u>) | Chazy, New York, 1960 |
| 29. melanotic stink glands (<u>msg</u>) | Berkeley, 1962 |
| 30. <u>rus msg</u> | |
| 31. <u>b rus</u> | |
| 32. <u>b msg</u> | |

Chromosome IV

- | | |
|-----------------------------------------------------------------------|----------------|
| 33. thumbled (<u>thu</u>) | Berkeley, 1963 |
| 34. thumbled ^S (<u>thus</u>) (=rsp ^P . S. D.) | Berkeley, 1963 |

Chromosome V

- | | |
|----------------------------------------------------------|-------------------------|
| 35. ebony (<u>e</u>) | Park, via Stanley, 1960 |
| 36. ebony (<u>e</u> ^{L&H}) | Berkeley, 1959 |
| 37. synthetic (<u>e</u> / <u>e</u> ^{L&H}) | Berkeley, 1961 |
| 38. blistered elytra (<u>ble</u>) | Chazy, New York, 1960 |
| 39. <u>e ble</u> | |

Chromosome VI

- | | |
|-----------------------------|----------------|
| 40. disjoined (<u>dj</u>) | Berkeley, 1963 |
|-----------------------------|----------------|

Unassigned (but possibly in III)

- | | |
|------------------------------------------------------------|----------------|
| 41. light ocular diaphragm (<u>lod</u>) (<u>pearl</u>) | Berkeley, 1961 |
|------------------------------------------------------------|----------------|

Multichromosomal

- | | |
|--------------------------------------------------|----------------|
| 42. <u>p; lod</u> | |
| 43. <u>p; rus</u> | |
| 44. <u>b; sp</u> | |
| 45. <u>rus; sp</u> | |
| 46. <u>rus; ble</u> | |
| 47. <u>b</u> (;) <u>lod; p</u> | |
| 48. <u>b twa</u> | |
| 49. <u>ems dt msg</u> | |
| 50. <u>jac dt b</u> | |
| 51. McGill <u>b p</u> | |
| 52. bent femur (<u>btf</u>) | Berkeley, 1964 |
| 53. bent tibia (<u>btt</u>) | Berkeley, 1962 |
| 54. black-3 (b-3) | Berkeley, 1964 |
| 55. crumpled elytra (<u>cru</u>) | Berkeley, 1964 |
| 56. creased abdominal sternites (<u>cas-1</u>) | Berkeley, 1963 |
| 57. deflected epimera (<u>dep</u>) | Berkeley, 1964 |
| 58. deformed legs (<u>dfl</u>) | Berkeley, 1965 |
| 59. elongated elytra (<u>ele</u>) | Berkeley, 1963 |
| 60. fused antennal segments-1 (<u>fas-1</u>) | Berkeley, 1962 |
| 61. fused antennal segments-2 (<u>fas-2</u>) | Berkeley, 1963 |

- | | |
|--------------------------------------------------------|----------------|
| 62. incomplete meso-metathoracic suture (<u>ims</u>) | Berkeley, 1965 |
| 63. incomplete metathoracic projections (<u>imp</u>) | Berkeley, 1964 |
| 64. knobby prothorax (<u>knp</u>) | Berkeley, 1964 |
| 65. legless (<u>lgl</u>) | Berkeley, 1966 |
| 66. medial abdominal groove (<u>mag</u>) | Berkeley, 1964 |
| 67. nude (<u>nd</u>) | Berkeley, 1964 |
| 68. pockets (<u>poc</u>) | Berkeley, 1965 |
| 69. prosternumless (<u>psl</u>) | Berkeley, 1966 |
| 70. Reduced eyes (<u>Re</u>) | Berkeley, 1965 |
| 71. rough (<u>ro</u>) (black) | McDonald, 1960 |
| 72. ruby (<u>rby</u>) | Berkeley, 1962 |
| 73. scar (<u>sc</u>) (=engraved metasternum) | Berkeley, 1962 |
| 74. separated epimera (<u>sep</u>) | Berkeley, 1964 |
| 75. short elytra (<u>sh</u>) | Berkeley, 1961 |
| 76. split (<u>sp</u>) | Berkeley, 1961 |
| 77. sternites incomplete (<u>sti</u>) | Berkeley, 1963 |
| 78. stilted legs (<u>stl</u>) | Berkeley, 1962 |
| 79. stunted (<u>stt</u>) | Berkeley, 1966 |
| 80. tiny (<u>ty</u>) | Berkeley, 1961 |
| 81. twisted abdomen (<u>twa</u>) | Berkeley, 1965 |
| 82. umbilicus (<u>umb</u>) (=dent) | Berkeley, 1962 |
| 83. warped elytra (<u>we</u>) | Berkeley, 1962 |
| 84. wingless (<u>wgl</u>) | Berkeley, 1965 |
|
 | |
| C. <u>Tribolium anaphe</u> | |
| None | |
|
 | |
| D. <u>Tribolium audax</u> | |
| None | |
|
 | |
| E. <u>Tribolium brevicornis</u> | |
| 1. creased abdominal sternites (<u>cas</u>) | |
| 2. split (<u>spl</u>) | |
| 3. fused antennal segments (<u>fas</u>) | |
|
 | |
| F. <u>Tribolium destructor</u> | |
| None | |
|
 | |
| G. <u>Tribolium madens</u> | |
| 1. fused antennal segments-1 (<u>fas-1</u>) | Berkeley, 1964 |
| 2. split (<u>spl</u>) | Berkeley, 1964 |
| 3. bent tibia (<u>btt</u>) | Berkeley, 1964 |
|
 | |
| H. <u>Latheticus oryzae</u> | |
| None | |
|
 | |
| I. <u>Oryzaephilus surinamensis</u> | |
| None | |

Stock Lists

J. Cryptolestes turcicus

Chromosome I

1. red (r)

PIL, 1963

K. Stegobium paniceum

None

A. Sokoloff

SANTA FE, NEW MEXICO
SANTA FE PREPARATORY SCHOOL

I. Wild type strain

- A. Tribolium castaneum
B. Tribolium confusum

Chicago via Berkeley
McGill via Berkeley

(Ed.)

SAVANNAH, GEORGIA
STORED-PRODUCT INSECTS RESEARCH AND DEVELOPMENT LABORATORY

I. Wild type strains

A. Lepidoptera

- | | |
|------------------------------------------|--------------------------------------------------|
| 1. <u>Anagasta kuehniella</u> (Zeller) | N.C. State at Raleigh, N.C. |
| 2. <u>Cadra cautella</u> (Walker) | Tifton, Ga. |
| 3. <u>Cadra figulilella</u> (Gregson) | Unknown |
| 4. <u>Ephestia elutella</u> (Hübner) | Richmond, Va. |
| 5. <u>Plodia interpunctella</u> (Hübner) | Modesto, Calif. |
| 6. <u>Sitotroga cerealella</u> (Olivier) | Manhattan, Kansas |
| 7. <u>Tineola bisselliella</u> (Hummel) | Savannah, Ga.; Ottawa, Can., and
Durham, N.H. |

B. Coleoptera

- | | |
|-----------------------------------------------------|------------------------------------------------------------------|
| 1. <u>Anthrenus flavipes</u> LeConte | Savannah, Ga.; and Durham, N.H. |
| 2. <u>Attagenus megatoma</u> (Fab.) | CSMA strains |
| 3. <u>Callosobruchus maculatus</u> (Fab.) | Fresno, California |
| 4. <u>Cathartus quadricollis</u> (Guérin-Méneville) | Unknown |
| 5. <u>Cryptolestes pusillus</u> (Schönherr) | Tifton, Ga. |
| 6. <u>Dermestes maculatus</u> De Geer | Madison, Wisconsin |
| 7. <u>Gibbium psylloides</u> (Czenpinski) | Unknown |
| 8. <u>Lasioderma serricorne</u> (Fab.) | Unknown |
| 9. <u>Oryzaephilus mercator</u> (Fauvel) | Unknown |
| 10. <u>Oryzaephilus surinamensis</u> (L.) | Manhattan, Kansas |
| 11. <u>Rhyzopertha dominica</u> (Fab.) | Unknown |
| 12. <u>Stegobium paniceum</u> (Linnaeus) | Madison, Wisconsin |
| 13. <u>Sitophilus granarius</u> (L.) | Manhattan, Kansas |
| 14. <u>Sitophilus oryzae</u> (L.) | Arkansas; California; Kansas;
Louisiana; Minnesota; and Texas |

Stock Lists

- | | | | |
|-----|---------------------------------|-------------|-------------------------------------------------|
| 15. | <u>Sitophilus zeamaze</u> | Motschulsky | Estill, S.C. |
| 16. | <u>Stegobium paniceum</u> | (L.) | Madison, Wisconsin |
| 17. | <u>Tenebrio molitor</u> | (L.) | Manhattan, Kansas; and Durham,
New Hampshire |
| 18. | <u>Tenebroides mauritanicus</u> | (L.) | Savannah, Ga. |
| 19. | <u>Tribolium castaneum</u> | (Herbst) | Unknown |
| 20. | <u>Tribolium confusum</u> | Jacquelin | duVal Manhattan, Kansas |
| 21. | <u>Tribolium madens</u> | Charpentier | Tifton, Ga. |
| 22. | <u>Trogoderma glabrum</u> | (Herbst) | Madison, Wisconsin; Riverside,
Calif. |
| 23. | <u>Trogoderma inclusum</u> | LeConte | Madison, Wisconsin; Riverside,
Calif. |
| 24. | <u>Trogoderma variabile</u> | Ballion | Fresno, Calif; Riverside, Calif. |

II. Mutant strains

A. Plodia interpunctella

- | | | |
|----|-----------------|---------------|
| 1. | Scaleless (scl) | Savannah, Ga. |
| 2. | Melanic (m) | Savannah, Ga. |

B. Tribolium castaneum

- | | | |
|----|--------------|---------------|
| 1. | Black mutant | Ocilla, Ga. |
| 2. | Black mutant | Savannah, Ga. |

C. Tribolium confusum

- | | | |
|----|-------------------------|---------------|
| 1. | Fused antennal segments | Savannah, Ga. |
| 2. | Short elytra | Savannah, Ga. |
| 3. | Crumpled elytra | Savannah, Ga. |
| 4. | Blade elytra | Savannah, Ga. |
| 5. | Umbilicus | Savannah, Ga. |
| 6. | Red eye pupae | Savannah, Ga. |

D. New mutants

- | | | |
|----|---------------------------------------------------------------------------------------------------------|---------------|
| 1. | <u>T. confusum</u> , peg-leg (pl) - an autosomal recessive with appendages extremely reduced in length. | Savannah, Ga. |
| 2. | <u>T. confusum</u> , separated elytra (sep) - elytra divergent from proximal end. | Savannah, Ga. |
| 3. | <u>T. confusum</u> , creased elytra (cr) - elytra creased and distal portion divergent. | Savannah, Ga. |

R. Davis

Stock Lists

SOUTH LANCASTER, MASSACHUSETTS
ATLANTIC UNION COLLEGE, BIOLOGY DEPARTMENT

Tribolium castaneum

I. Wild type strains

1. Brazil (C-1)
2. Chicago
3. McGill
4. Sacramento
5. Texas
6. Veracruz, Mexico
7. Virginia

II. Mutant strains

1. red (r^D)
2. red (r)
3. red (r^{Ho})
4. red modifier (MF)
5. McGill black (McGb)
6. Chicago black (Ch)
7. black (BS-1), Brazil black
8. sooty (s)
9. jet (j)
10. chestnut (cS)

(Ed.)

SOUTH ORANGE, NEW JERSEY
SETON HALL UNIVERSITY, DEPARTMENT OF BIOLOGY

I. Wild type strains

A. Laboratory strains

1. Tribolium castaneum - McGill Montreal, Canada via
University of California
2. Tribolium castaneum - Seton Hall South Orange,
New Jersey
3. Tribolium castaneum - inbred - 20 generations
4. Tribolium confusum Fordham University

B. Base Populations for quantitative studies (Tribolium castaneum)

1. Foundation b - marked with black (b) body color -
obtained via Purdue University, Lafayette,
Indiana.
2. Foundation p - marked with pearl (p) eye color - obtained
via Purdue University, Lafayette, Indiana.

Stock Lists

II. Mutant strains

A. Tribolium castaneum

- | | |
|---------------------------------|------------------------------|
| 1. ca | via University of California |
| 2. fas-3 | via California State College |
| 3. paddle | via California State College |
| 4. red ^{no} | via California State College |
| 5. Short antennae (<u>Sa</u>) | Purdue + Foundation, 1960 |
| 6. white | via California State College |

Eliot Krause

STATE COLLEGE, MISSISSIPPI
 USDA, ARS, BOLL WEEVIL RESEARCH LABORATORY

Anthonomus grandis

A. Wild type strains

- | | |
|--------------|------------------------|
| 1. A & M | College Station, Texas |
| 2. Oktibbeha | State College, Miss. |
| 3. Thurberia | Tucson, Ariz. |
| 4. Iguala | Iguala, Mexico |

B. Mutant strains

- | | |
|------------------------|---------------|
| 1. yellow (<u>y</u>) | A & M strain |
| 2. slate (<u>e</u>) | Acala, Mexico |
| 3. ebony (<u>e</u>) | A & M strain |
| 4. pearl (<u>p</u>) | A & M strain |

C. Insecticide resistant

- | | |
|---------------------|-------------------|
| 1. Endrin Resistant | Auburn University |
| ca. 20 g/weevil | |

W. Ivey

(Ed.)

STONY BROOK, NEW YORK
 STATE UNIVERSITY OF NEW YORK, DIVISION OF BIOLOGICAL SCIENCES

Tribolium castaneum

I. Wild type

- | | |
|-------------------|--------------------------|
| 1. UPF Foundation | Purdue University |
| 2. CS-4 | University of California |

II. Mutants

- | | |
|-------------------------|-----------------------|
| 1. Sooty (<u>s</u>) | |
| 2. paddle (<u>pd</u>) | University of Chicago |

Stock Lists

- | | |
|----------------------------------------------------------------------------------------------------------|--------------------------|
| 3. pearl (p) | University of Chicago |
| 4. McGill black (McGb) | University of Chicago |
| Stock | University of California |
| 5. McGill black (McGb) with UPF genetic background obtained by backcrossing to UPF for nine generations. | University of Kansas |

Tribolium confusum

- | | |
|-------------------------|--------------------------|
| I. Wild type | |
| 1. Chicago (Sonleitner) | University of Chicago |
| 2. New York | University of Chicago |
| II. Mutants | |
| 1. McGill black (McGb) | University of California |
| 2. ebony (e) | University of Chicago |
| | Robert R. Sokal |

ST. BERNARD, ALABAMA
ST. BERNARD ABBEY

- | | |
|-------------------------------|--------------------|
| I. Wild Type strains | |
| A. <u>Tribolium castaneum</u> | |
| 1. Chicago | via San Bernardino |
| B. <u>Tribolium confusum</u> | |
| 1. New York | via San Bernardino |
| II. Mutant strains | |
| A. <u>Tribolium castaneum</u> | |
| 1. McGill black | via San Bernardino |
| 2. jet | via San Bernardino |
| 3. Sooty | via San Bernardino |
| 4. Chicago black | via San Bernardino |
| B. <u>Tribolium confusum</u> | |
| 1. pearl | via San Bernardino |
| 2. McGill black | via San Bernardino |
| 3. Ebony (Smith) | via San Bernardino |

Stock Lists

ST. PAUL, MINNESOTA
UNIVERSITY OF MINNESOTA, DEPARTMENT OF ENTOMOLOGY, FISHERIES AND WILDLIFE

Stock list of stored product insects

Tribolium confusum
Sitophilus oryzae (large strain)
Sitophilus granarius
Oryzaephilus surinamensis
Troscoderna parabili
Rhyssopertha dominica
Plodia interpunctella
Ephestia spp.
Attageus maratona

Ernesto De las Casas
Phillip K. Harein

ST. PAUL, MINNESOTA 55101
UNIVERSITY OF MINNESOTA
DEPARTMENT OF GENETICS AND CELL BIOLOGY

Tribolium castaneum

A. Synthetic populations

All selected populations originated from a cross of inbred lines CSI-5 and CSI-10 obtained in 1963 from the Tribolium stock center (marked with sooty).

1. S lines - populations selected for 85 generations for increased pupa weight.
2. C lines - stabilized selected populations for 21 day pupa weight (85 generations).
3. R lines - random bred and random selected for 60 generations.
4. T lines - populations selected for 21 day pupa weight for 52 generations followed by 30 generations of relaxed selection.

F. D. Enfield

ST. PAUL, MINNESOTA
UNIVERSITY OF MINNESOTA, DEPARTMENT OF ANIMAL SCIENCE

A. Inbreds

1. CSI-10
2. E 1
3. E 2

University of California, Berkeley
Institute of Animal Genetics, Edinburgh
Institute of Animal Genetics, Edinburgh

B. Purdue Foundation, p

- C. Segregating population selected for pupa weight, synthesized by crossing CSI-10 and E 2 lines.

Stock Lists

TEMPE, ARIZONA
ARIZONA STATE UNIVERSITY, DEPARTMENT OF ZOOLOGY

I. Synthetic strains

- A. Tribolium castaneum
1. Berkeley, 1964 via San Bernardino
- B. Tribolium confusum
1. Berkeley, 1958 via San Bernardino

II. Mutant strains

- A. Tribolium castaneum
1. melanotic stink glands (msg),
Berkeley, 1964 via San Bernardino
- B. Tribolium confusum
1. melanotic stink glands (msg),
Berkeley, 1962 via San Bernardino
- Harry E. Wistrand

TIFTON, GEORGIA
ABRAHAM BALDWIN AGRICULTURAL COLLEGE

Tribolium castaneum

- A. Wild type strain
1. Chicago
- B. Mutant strains
1. black
2. squint

(All derived from stocks maintained at Berkeley. Ed.)

URBANA, ILLINOIS
UNIVERSITY OF ILLINOIS, DEPARTMENT OF ZOOLOGY

I. Wild type strains

- A. Tribolium castaneum
1. Berkeley Berkeley, 1966
2. Chicago Urbana, 1966
3. Carbondale Maryland, 1966
4. del Valle Maryland, 1966
5. Kansas Kansas, 1966
- B. Tribolium confusum
1. Berkeley Berkeley, 1966

Stock Lists

2. Chicago
3. Kansas
4. Maryland
5. Minnesota
6. Oklahoma

- Urbana, 1966
 Kansas, 1966
 Maryland, 1966
 Minnesota, 1966
 Urbana, 1966

II. Inbred lines

A. Tribolium castaneum

1. CSI-2
2. CSI-3
3. CSI-5
4. CSI-10
5. CSI-12
6. CSI-14
7. CSI-15
8. CSI-16
9. CSI-22

- Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966

B. Tribolium confusum

1. CFI-1
2. CFI-2
3. CFI-3
4. CFI-5
5. CFI-7
6. CFI-8
7. CFI-11
8. CFI-12
9. CFI-13
10. CFI-14
11. CFI-15
12. CFI-16
13. CFI-18
14. CFI-19
15. CFI-20
16. CFI-21
17. CFI-22
18. CFI-23
19. CFI-24

- Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966
 Berkeley, 1966

III. Mutant strains

A. Tribolium castaneum

1. sa-2 (+/s)
2. i
3. w
4. b, mc, p
5. bal, s
6. pd
7. Be
8. mc
9. aa (+/p)

- Berkeley, 1966
 Purdue, 1967
 Purdue, 1967
 Berkeley, 1966
 Berkeley, 1966
 Urbana, 1966
 Berkeley, 1966
 Berkeley, 1967
 Berkeley, 1967

Stock Lists

10.	<u>r^{Ho}</u>	Berkeley, 1966
11.	<u>Mo</u>	Berkeley, 1966
12.	<u>b</u>	Berkeley, 1966
13.	<u>ap^D, s</u>	Berkeley, 1966
14.	<u>i</u>	Berkeley, 1966
15.	<u>r (+/py)</u>	Berkeley, 1966
16.	<u>Fta/+, c</u>	Berkeley, 1966
17.	<u>c</u>	Berkeley, 1966
18.	<u>Spa/+, +/c</u>	Berkeley, 1966
19.	<u>p</u>	Berkeley, 1967
20.	<u>sq</u>	Berkeley, 1967
21.	<u>msg</u>	Berkeley, 1967
22.	<u>sh^s</u>	Berkeley, 1967
23.	<u>p, lod</u>	Berkeley, 1967
24.	<u>Sa-2, s</u>	Berkeley, 1967
25.	<u>rg</u>	Berkeley, 1967
26.	<u>fas-3a</u>	Berkeley, 1967
27.	<u>r^D, s</u>	Berkeley, 1967
28.	<u>dve, pd</u>	Berkeley, 1967
29.	<u>h</u>	Berkeley, 1967
30.	<u>rs</u>	Urbana, 1967
31.	<u>rb</u>	Purdue, 1967
32.	<u>i, m</u>	Purdue, 1967
33.	<u>ctp, ju</u>	Purdue, 1967
		Berkeley, 1967

(Ed.)

URBANA, ILLINOIS 61801
 UNIVERSITY OF ILLINOIS AT URBANA CHAMPAIGN
 DEPARTMENT OF PHYSIOLOGY AND BIOPHYSICS

Tribolium castaneum

1. Wild type

(maintained since 1960)

Tribolium confusum

1. Wild type from Oklahoma
2. Mutant, ebony
3. Mutant, McGill black

Dawson, 1967
 Dawson, 1967
 Sokoloff, 1966

Tribolium brevicornis

1. Wild type

Yang, 1970

Stock Lists

AUSTRALIA

BRISBANE, QUEENSLAND
DEPARTMENT OF PRIMARY INDUSTRIES, ENTOMOLOGY LABORATORY

COLEOPTERA

- A. Tribolium castaneum
 - 1. Wild type strains
 - 2. Black mutant (reoccurrence of b)
 - 3. Lindane resistant

- B. Sitophilus oryzae
 - 1. Wild type strains
 - 2. DDT resistant (single semi-dominant sex-linked factor)
 - 3. Lindane and dieldrin resistant (single and multi-factor strains.)
 - 4. Black strain

- C. Sitophilus zeamais--wild type

- D. Sitophilus granarius--wild type

- E. Oryzaephilus surinamensis
 - 1. Wild type strains
 - 2. Lindane resistant strains (impure)

- F. Lasioderma serricorne--wild type

- G. Rhizopertha dominica--wild type

- H. Mezium americanum--wild type

LEPIDOPTERA

- A. Cadra cautella--wild type

- B. Phthorimaea operculella
 - 1. DDT-endrin resistant
 - 2. Red-eyed mutant (single autosomal recessive)

HYMENOPTERA

- A. Microchelonus sp.--wild type

BELGIUM

GE'PLOUX
 INSTITUT AGRONOMIQUE DE L'ETAT, ZOOLOGIE GENERALE

Tenebrio molitor

F strain - selected for small weight since 1950
 G strain - selected for large weight since 1950

(Ed.)

LOUVAIN
 F.A. GANSSENS MEMORIAL LABORATORY FOR GENETICS
 AGRICULTURAL INSTITUTE OF THE UNIVERSITY

Tenebrio molitor

Wild type

Belgium

Tribolium confusum

Two inbred and a wild type

Berkeley, 1965
(Ed.)

BRAZIL

CAMPINAS, SAO PAULO
 INSTITUTE AGRONOMICO, SECAO DE ENTOMOLGIA

Anobiidae

Lasioderna serricorne (F) - Campinas, SP - wild type

Bostrochidae

Rhizopertha dominica (F) - Campinas, SP - wild type

Bruchidae

Acanthoscelides obsoletus (Say) - Campinas, SP - wild type

March, 1974

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Stock Lists

Curculionidae

Sitophilus oryzae (L.) - Campinas, SP - wild type

Silvanidae (Cucujidae)

Oryzaephilus surinamensis (L.) - Campinas, SP - wild type

Tenebrionidae

Tribolium castaneum (Herbst.) - Campinas, SP - wild type
(Ed.)

CANADA

BURNLEY, VICTORIA
VICTORIAN PLANT RESEARCH INSTITUTE, DEPARTMENT OF AGRICULTURE

COLEOPTERA

- A. Tribolium castaneum
1. Wild type strains
 2. Malathion specific resistant strain
 3. Malathion non-specific resistant strain

- B. Tribolium confusum
1. Wild type strains
 2. Malathion specific strain

Stock Lists

- C. Oryzaephilus surinamensis
 - 1. Wild type strain
 - 2. Malathion resistant strain
- D. Gnathocerus cornutus wild type strain
- E. Cryptolestes ferrugineus wild type strain
- F. Sitophilus oryzae wild type strain
- G. Sitophilus zeamais wild type strain
- H. Sitophilus granarius wild type strain
- I. Rhyzopertha dominica wild type strain

LEPIDOPTERA

- A. Plodia interunctella wild type strain
- B. Ephestia figulilella wild type strain

EDMONTON, ALBERTA
UNIVERSITY OF ALBERTA, DEPARTMENT OF ANIMAL SCIENCE

- | | |
|------------------------|------------------------|
| A. Brazil | Purdue, 1965 |
| B. Capetown | Purdue, 1965 |
| C. Chicago | Chicago, 1965 |
| D. Consejo | Madrid, 1965 |
| E. Japan | Kyoto and Purdue, 1965 |
| F. Kano | Scotland, 1965 |
| G. Kenya | Scotland, 1965 |
| H. Kingston | Scotland, 1965 |
| I. Lisbon | Portugal, 1965 |
| J. Purdue Foundation + | Manitoba, 1963 |
| K. Scotland | Edinburgh, 1965 |
| L. Seychelles | Scotland, 1965 |
| M. Surrey | England, 1965 |
| N. Veracruz | Berkeley, 1965 |

(Ed.)

GUELPH, ONTARIO
UNIVERSITY OF GUELPH, DEPARTMENT OF ANIMAL AND POULTRY SCIENCE

Tribolium castaneum

Wild type - mass mated stock derived from Purdue
University Foundation Stock.

R. Fairfull

Stock Lists

GUELPH, ONTARIO
UNIVERSITY OF GUELPH, DEPARTMENT OF ZOOLOGY

- A. Sitophilus granarius (L)
 - 1. GG strain, dark, heavy and symbiotic
 - 2. MW strain, paler, lighter and aposymbiotic
 - 3. Two new strains as yet unclassified
- B. Sitophilus oryzae (L.)--small rice weevil
- C. Sitophilus zea-mais (Mots.)--large rice weevil.

(Ed.)

MONTREAL, P. Q.
MCGILL UNIVERSITY, DEPARTMENT OF BIOLOGY

Tribolium castaneum

- | | |
|-----------------------------------------|------------------|
| 1. Berkeley CSI-3F | Sokoloff, 1970 |
| 2. Berkeley CS--synthetic, <u>sooty</u> | Sokoloff, 1970 |
| 3. Berkeley CS--synthetic, <u>PYAMY</u> | Sokoloff, 1970 |
| 4. McGill <u>black</u> | Sokoloff, 1970 |
| 5. Chicago <u>black</u> | Sokoloff, 1970 |
| 6. Purdue Foundation | Scheinberg, 1967 |

OTTAWA, ONTARIO
ANIMAL GENETICS SECTION, ANIMAL RESEARCH INSTITUTE
CENTRAL EXPERIMENTAL FARM

Tribolium castaneum

Purdue Foundation

- RSILW - A population selected for high larval weight for 10 generations restricting developmental time and pupal weight (derived from Purdue Foundation).
- RSIDT - A population selected for short developmental time for 10 generations restricting larval weight and pupal weight (derived from Purdue Foundation).
- RSIPW - A population selected for short developmental time for 10 generations restricting larval weight and developmental time (derived from Purdue Foundation).

Stock Lists

TSLW - A populations selected for high larval weight for 10 generations (derived from Purdue Foundation).

TSDT - A population selected for short developmental time for 10 generations (derived from Purdue Foundation).

TSPW - A population selected for high pupal weight for 10 generations (derived from Purdue Foundation).

(Ed.)

QUEBEC, P.Q.
UNIVERSITE LAVAL, DEPARTMENT OF BIOCHEMISTRY

Tribolium confusum Duval

Strain: Laval
Origin: Quebec City

A. Lemonde

QUEBEC, P.Q.
UNIVERSITE LAVAL, DEPARTMENT OF BIOLOGY

Tribolium confusum Duval

Strain: Laval
Origin: Quebec City

L. Huot

VANCOUVER, B. C.
UNIVERSITY OF BRITISH COLUMBIA, DEPARTMENT OF POULTRY SCIENCE

I. Wild type strains

A. Tribolium confusum inbred lines

- | | |
|-----------|----------------|
| 1. CFI-2a | Berkeley, 1965 |
| 2. CFI-3 | Berkeley, 1965 |
| 3. CFI-5 | Berkeley, 1965 |
| 4. CFI-7 | Berkeley, 1965 |
| 5. CFI-8b | Berkeley, 1965 |

II. Mutant strains

A. Tribolium confusum

- | | |
|----------------------------------------------------|----------------|
| 1. eyespot (<u>es</u>); chromosome I | Berkeley, 1965 |
| 2. red (<u>r</u>); chromosome I | Berkeley, 1965 |
| 3. dirty pearl eye (<u>dpe</u>); chromosome IV | Berkeley, 1965 |
| 4. ebony-2 (<u>e₂</u>); chromosome II | Berkeley, 1965 |

Stock Lists

- | | | |
|-------------------------------|---------------------------------------------------------------------|----------------|
| 5. | pearl riboflavinless (<u>pr</u>);
chromosome II | Berkeley, 1965 |
| 6. | pearl slough (<u>p</u>); chromosome II | Berkeley, 1965 |
| 7. | ruby spot (<u>rus</u>); chromosome III | Berkeley, 1965 |
| 8. | light ocular diaphragm (<u>lod</u>);
chromosome III | Berkeley, 1965 |
| 9. | <u>p</u> ; <u>dre</u> ; <u>cas</u> ; multichromosomal | Berkeley, 1965 |
| 10. | <u>r</u> <u>s</u> ; <u>b</u> ; multichromosomal | Berkeley, 1965 |
| 11. | <u>St</u> ; <u>b</u> ; multichromosomal | Berkeley, 1965 |
| B. <u>Tribolium castaneum</u> | | |
| 1. | red (<u>r</u>); chromosome I | Berkeley, 1965 |
| 2. | pearl (<u>p</u>); chromosome I | Purdue, 1967 |
| 3. | pearl riboflavinless (<u>p^r</u>)
(formerly "ivory") | Berkeley, 1965 |
| 4. | pink (<u>p^{pk}</u>); chromosome II | |
| 5. | light ocular diaphragm (<u>p</u>
background); chromosome III | Berkeley, 1965 |
| 6. | jet H.L.E.; chromosome V | Berkeley, 1867 |
| 7. | Chestnut (<u>c</u>); chromosome VII | Berkeley, 1965 |
| 8. | <u>s</u> ; <u>r^D</u> ; multichromosomal | Berkeley, 1965 |

VANCOUVER, B.C.

UNIVERSITY OF BRITISH COLUMBIA, POULTRY SCIENCE GENETICS LABORATORY

Tribolium confusum

Wild type

- U.B.C. wild type - Vancouver, B.C.

Mutants

- Riboflavinless, pearl-eye (p^r)

C.W. Roberts

WINNIPEG, MANITOBA

CANADA DEPARTMENT OF AGRICULTURE, RESEARCH STATION

I. Wild type strains

A. Coleoptera

- | | | |
|----|-------------------------------------------------------|-----------------------|
| 1. | <u>Acanthoscelides obtectus</u> (Say)
Bruchidae | Winnipeg |
| 2. | <u>Ahasverus advena</u> (Waltl)
Silvanidae | Manitoba |
| 3. | <u>Alphitobius diaperinus</u>
Panzer Tenebrionidae | Saskatchewan |
| 4. | <u>Anthicus floralis</u> L.
Anthicidae | Manitoba |
| 5. | <u>Cryptolestes ferrugineus</u> (Steph.)
Cucujidae | Manitoba |
| 6. | <u>Cryptolestes ferrugineus</u> (Steph.)
Cucujidae | PIL United
Kingdom |
| 7. | <u>Cryptolestes ferrugineus</u> (Steph.)
Cucujidae | Australia |

Stock Lists

- | | | |
|-----|---------------------------------------------------------|--------------------|
| 8. | <u>Cryptolestes turcicus</u> (Grouv.)
Cucujidae | Ontario |
| 9. | <u>Cryptolestes turcicus</u> (Grouv.)
Cucujidae | PIL United Kingdom |
| 10. | <u>Cybaeus angustus</u>
Leconte Tenebrionidae | Manitoba |
| 11. | <u>Oryzaephilus mercator</u> (Fauvel)
Silvanidae | Ontario |
| 12. | <u>Oryzaephilus surinamensis</u> (L.)
Silvanidae | Manitoba |
| 13. | <u>Rhyzopertha dominica</u> (Fab.)
Bostrichidae | Australia |
| 14. | <u>Sitophilus granarius</u> (L.)
Curculionidae | Manitoba |
| 15. | <u>Sitophilus oryzae</u> (L.)
Curculionidae | Montreal |
| 16. | <u>Sitophilus zea-mais</u>
Motschulsky Curculionidae | Japan |
| 17. | <u>Stegobium paniceum</u> (L.)
Anobiidae | Winnipeg |
| 18. | <u>Tenebroides mauritanicus</u> (L.)
Ostomidae | Manitoba |
| 19. | <u>Tenebrio molitor</u> (L.)
Tenebrionidae | Manitoba |
| 20. | <u>Tribolium castaneum</u> (Herbst)
Tenebrionidae | Manitoba |
| 21. | <u>Tribolium confusum</u> (DuVal) | Manitoba |
| 22. | <u>Tribolium madens</u>
Charp. Tenebrionidae | Manitoba |
| 23. | <u>Troxoderma variabile</u>
Ballion Dermestidae | Alberta |

B. Lepidoptera

- | | | |
|----|---------------------------------------------------|----------|
| 1. | <u>Nemapogon granella</u> (L.)
Tineidae | Ontario |
| 2. | <u>Plodia interpunctella</u> (Hbn.)
Phycitidae | Winnipeg |
| 3. | <u>Pyralis farinalis</u> L.
Pyralidae | Manitoba |

II. Mutant strain

A. Coleoptera

- | | |
|-----------------------------------------------------------------------------------------|--------------------------|
| <u>Tribolium castaneum</u> Herbst
Tenebrionidae
(Black body color from lab stock) | Winnipeg |
| <u>Tribolium confusum</u> DuVal | Winnipeg, Manitoba, 1963 |

L.B. Smith

Stock Lists

DENMARK

LYNGBY
STATENS SKADEDYRLABORATORIUM
(DANISH PEST INFESTATION LABORATORY)

Alphitobius diaperinus
Anobium punctatum
Anthrenus museorum
Anthrenus vorax
Attagenus alfieri
Attagenus piceus
Dermestes frichi
Hylotrupes bajulus
Lasioderma serricorne
Oryzaephilus mercator
Oryzaephilus surinamensis
Rhizopertha dominica
Sitochilus granarius
Sitophilus oryzae
Stegobium (Sitodrepa) paniceum
Tenebrio molitor
Tenebrioides mauritanicus
Taylodrias contractus
Tribolium confusum
Tribolium destructor
Trogoderma granarium

EASTERN NIGERIA

PORT HARCOURT
THE NIGERIAN STORED PRODUCTS RESEARCH INSTITUTE

I. Wild type strains

- | | |
|-------------------------------------------|-------------------------------------|
| 1. <u>Dermestes maculatus</u> De Geer | Port Harcourt Strain, 1966 |
| 2. <u>Oryzaephilus mercator</u> Fauv. | Port Harcourt Strain, 1966 |
| 3. <u>Sitophilus zea-mais</u> Motschulsky | Kano Strain, 1965 |
| | (Ex Kano Lab. Stock) November, 1965 |
| 4. <u>Tribolium castaneum</u> Hbst. | Kano Strain, 1965 |
| | (Ex Kano Lab. Stock) October, 1965 |
| 5. <u>Tribolium confusum</u> DuVal. | Kano Strain, 1965 |
| | (Ex Kano Lab. Stock) December, 1965 |
| 6. <u>Trogoderma granarium</u> Everts | Kano Strain, 1965 |
| | (Ex Kano Lab. Stock) November, 1965 |

(Ed.)

Stock Lists

GIZA

PLANT PROTECTION DEPARTMENT, MINISTRY OF AGRICULTURE

I. Wild type strains

- | | |
|---------------------------------|---------------|
| 1. <u>Bruchus rufimanus</u> | Egypt, U.A.R. |
| 2. <u>Corcyra cephalonica</u> | Egypt, U.A.R. |
| 3. <u>Ephestia kuhniella</u> | Egypt, U.A.R. |
| 4. <u>Latheticus oryzae</u> | Egypt, U.A.R. |
| 5. <u>Rhizopertha dominica</u> | Egypt, U.A.R. |
| 6. <u>Silvanus surinamensis</u> | Egypt, U.A.R. |
| 7. <u>Sitophilus granarius</u> | Egypt, U.A.R. |
| 8. <u>Sitophilus oryzae</u> | Egypt, U.A.R. |
| 9. <u>Tribolium castaneum</u> | Egypt, U.A.R. |
| 10. <u>Tribolium confusum</u> | Egypt, U.A.R. |

Note: Dr. M. A. Hafeez is at present in London. Fate of above stocks is unknown.

(Ed.)

FRANCE

LYON, RHÔNE

LABORATOIRE DE ZOOLOGIE GÉNÉRALE, FACULTÉ DES SCIENCES

Tribolium castaneum

Wild type strain from Alès, France.

(Ed.)

VILLEURBANNE (LYON) RHÔNE

INSTITUT NATIONAL DES SCIENCES APPLIQUÉES, LABORATOIRE DE BIOLOGIE

- | | |
|-----------------------------------------------------------------------|---------------------------------------------|
| A. <u>Acanthoscelides obsoletus</u> --wild type | France |
| B. <u>Blabera fusca</u> | |
| C. <u>Clitumnus extradentatus</u> | |
| D. <u>Galleria mellonella</u> | Saint Cyr au Mont d'Or |
| E. <u>Oryzaephilus surinamensis</u> --from imported
dried apricots | |
| F. <u>Periplaneta americana</u> | |
| G. <u>Pseudococcus citri</u> | Antibes |
| H. <u>Sitophilus granarius</u> | Infestation Control Laboratory,
Surbiton |

Stock Lists

I. <u>Sitophilus oryzae</u>	P.I.L., Slough
J. <u>Sitophilus sasakii</u> --wild type	Lyon
K. <u>Stegobium paniceum</u>	P.I.L., Slough
L. <u>Tenebrio molitor</u>	
M. <u>Tenebrio obscurus</u>	P.I.L., Slough
N. <u>Tribolium castaneum</u> --wild type	Alès

(Ed.)

GERMANY

MUNICH
 BAYER, LANDESANSTALT FÜR BODENKULTUR
 PFLANZENBAU U. PFLANZENSCHUTZ

Coleoptera

Cucjidae

Cryptolestes turcicus (Grouv.) Munich, 1966

Curculionidae

Sitophilus granarius (L.) Munich, 1966
Sitophilus zea-mais (Motsch.) 1966

Ptinidae

Gibbium psylloides (Czemp.) Regensburg, 1960

Silvanidae

Oryzaephilus mercator (Fauv.) Munich, 1966
Oryzaephilus surinamensis (L.) Munich, 1959

Tenebrionidae

Gnathocerus cornutus (F.) Munich, 1966
Tribolium confusum (Duv.) Munich, 1960
Tribolium destructor (Uyttenb.) Munich, 1957

Lepidoptera

Phyticidae

Anagasta kuehniella (Zell.) Munich, 1966

(Ed.)

Stock Lists

BIRMINGHAM, ENGLAND
 THE UNIVERSITY OF BIRMINGHAM
 DEPARTMENT OF ZOOLOGY AND COMPARATIVE PHYSIOLOGY

Tenebrio molitor
Tenebrio obscurus
Blaps sp.
Tribolium sp.

(Ed.)

DUNDEE, ANGUS
 UNIVERSITY OF DUNDEE, DEPARTMENT OF NATURAL HISTORY

Only the stock unique to this laboratory is listed.

Wild stock

1. Tribolium castaneum - Kenya. Collected in December, 1967, from stored maize in the Nairobi district.

(Ed.)

DUNDEE
 DUNDEE UNIVERSITY
 DEPARTMENT OF BIOLOGICAL SCIENCES

	<u>Origin</u>
<u>Tribolium castaneum</u>	Slough
<u>T. confusum</u>	
<u>T. madens</u>	
<u>T. destructor</u>	
<u>T. brevicornis</u>	
<u>T. castaneum</u> - M..S.G	Sekelleff
<u>T. castaneum</u> - M.S.G.	Slough
<u>T. confusum</u> - Black mutant	Slough
<u>T. confusum</u> - Pearl Eye	Slough
<u>Ephestia cautella</u>	Slough
<u>Gnathocerus maxillosus</u>	Slough
<u>G. germutus</u> - Slough, New Zealand	Egypt
<u>Latheticus oryzae</u> - Normal and Dark	Slough
<u>Sitophilus zeamais</u>	Slough

F. L. Waterhouse

ENDINBURGH
UNIVERSITY OF ENDINBURGH, INSTITUTE OF ANIMAL GENETICS

Tribelium castaneum

- A. Wild type strain
 - 1. Chicago wild type
- B. Mutant strains
 - 1. Microphthalmic (Me)
 - 2. microcephalic, pearl (mc, p)
 - 3. Bar eye, sooty (Be s/ts)
 - 4. squint (sq)

Stocks obtained from Berkeley, California.

(Ed.)

ENDINBURGH, SCOTLAND
DEPARTMENT OF AGRICULTURE AND FISHERIES FOR SCOTLAND
AGRICULTURAL SCIENTIFIC SERVICES, EAST CRAIGS

Tribelium castaneum Herbst.

Wild type strain of unknown origin, collected from imported feedstuffs.

Tribelium confusum J. duV.

Wild type strain of unknown origin, collected from imported feedstuffs.

(Ed.)

GLASGOW, SCOTLAND
UNIVERSITY OF GLASGOW
DEPARTMENT OF ZOOLOGY

Wild type Strains:

Wild strain of *T. castaneum* from the Infestation Department
Ministry of Agriculture & Fisheries, Glasgow

T. castaneum
ppas partially pointed abdominal sternites

LONDON
 QUEEN ELIZABETH COLLEGE, DEPARTMENT OF BIOLOGY

Bruchus pectinicornis
Latheticus oryzae
Stenophilus granarius
Tenebrio molitor
Tribelium anaphe
Tribelium castaneum
Tribelium madens
Trogoderma

All insects are derived from the Pest Infestation Laboratory, Slough, Bucks.

(ed.)

NEWCASTLE UPON TYNE.
 THE UNIVERSITY OF NEWCASTLE UPON TYNE, SCHOOL OF AGRICULTURE

Tribelium castaneum

A. Wild type

1. pearl (p)
2. black (b), tawny (b^t)
3. antennapedia (ap)
4. paddle (pd)
5. red (r)

Tribelium confusum

A. Wild type

1. ebony (e²)
2. pearl (p)

All stocks derived from cultures at the Insectary of the Pest Infestation
 Central Laboratory, Slough, Bucks.

(ed.)

SLOUGH, BUCKS
 MINISTRY OF AGRICULTURE, FISHERIES AND FOOD
 THE INSECTARY OF THE PEST INFESTATION CONTROL LABORATORY

ORDER Family (-subfamily) Genus (sub genus), species.	COMMON NAME	COUNTRY OF ORIGIN OF STOCK	CULTURE MEDIUM	REARING TEMPERATURE °C
Cucujidae				
<u>Cryptolestes capensis</u> (Waltl)			10	25
<u>Cryptolestes ferrugineus</u> (Steph.)	Rust-red grain beetle		10	30
<u>Cryptolestes pusilloides</u> (Steel & Howe)		(Canada)	10	25
<u>Cryptolestes pusillus</u> (Schon.)	Flat grain beetle		10	25
<u>Cryptolestes turcicus</u> (Grouv.)			10	25
<u>Cryptolestes ugandae</u> (Steel & Howe)		(E. Africa)	10	25
Curculionidae				
<u>Sitophilus granarius</u> (L.)	Grain weevil	(Russia)	1	25
<u>Sitophilus oryzae</u> (L.)	Rice weevil	Britain	1	25
<u>Sitophilus zeamais</u> Motsch.	Maize weevil		1	25
Dermestidae				
<u>Anthrenocerus australis</u> (Hope)	Australian carpet beetle	(Britain)	20	25
<u>Anthrenus</u> (Anthrenus) <u>flavipes</u> LeC. (=vorax Waterh.)	Furniture carpet beetle		20	30
<u>Anthrenus</u> (Nathrenus) <u>verbasci</u> (L.)	Varied carpet beetle	Britain	20	20
<u>Anthrenus</u> (Anthrenodes) <u>sarnicus</u> Mroczkowski			35	20
<u>Anthrenus</u> (Florilinus) <u>olgae</u> Kalik			20	20
<u>Attagenus</u> sp? (<u>alfieri</u> of Hinton 1945)		Kenya	17	25
<u>Attagenus fasciatus</u> (Thunberg) (=gloriosae (Fabricius))		Botswana	18	25
<u>Attagenus unicolor</u> Brahm (=megatoma (F.) & piceus (Ol.) nec. Thb.)	Black carpet beetle		20	30
<u>Attagenus pellio</u> (L.)	Fur beetle	Britain	20	20
<u>Dermestes ater</u> Deg.	Black larder beetle	Britain	21a	25
<u>Dermestes frischii</u> Kug.	Hide beetle	(Nigeria)	21a	25
<u>Dermestes haemorrhoidalis</u> Kuster		Britain	21a	25
<u>Dermestes lardarius</u> L.	Bacon beetle	Britain	21a	25
<u>Dermestes maculatus</u> Deg.	Leather beetle		21a	25

ORDER Family (-subfamily) Genus (sub genus), species.	COMMON NAME	COUNTRY OF ORIGIN OF STOCK	CULTURE MEDIUM
Gelechiidae			
<u>Sitotroga cerealella</u> (Oliv.)	Angoumois grain moth		1
Tineidae			
<u>Tinea columberiella</u> Wocke	Case bearing clothes moth		20
<u>Tinea flavescens</u> Haworth		Britain	20
<u>Tineola bisselliella</u> (Humm.)	Common clothes moth		20
<u>Tinea Pellionella</u> (L.)			20
COLEOPTERA			
Anobiidae			
<u>Lasioderma serricorne</u> (F.)	Cigarette beetle		6
<u>Stegobium paniceum</u> (L.)	Biscuit beetle		6
Antribidae			
<u>Araecerus fasciculatus</u> (Deg.)			39
Bostrichidae			
<u>Rhyzopertha dominica</u> (F.)	Lesser grain borer		1
Bruchidae			
<u>Acanthoscelides obtectus</u> (Say)	Dried Bean beetle	W. Africa	27
<u>Callosobruchus analis</u> (F.)			29
<u>Callosobruchus chinensis</u> (L.)	Cowpea weevil		29
<u>Callosobruchus maculatus</u> (F.)		Sierra Leone	29
<u>Callosobruchus rhodesianus</u> (Pic)			29
<u>Caryedon serratus</u> (Oliv.)	Groundnut seed beetle		26a
(=gonagra (F.))			
<u>Zabrotes subfasciatus</u> (Boh.)			28
Cerylonidae			
<u>Murmidius ovalis</u> (Beck)		Ceylon	13
Cleridae			
<u>Necrobia rufipes</u> (Deg.)	Copra beetle		22
<u>Necrobia ruficollis</u> (F.)			40

Stock Lists

I. Wild type strains

ORDER Family (-subfamily) Genus (sub genus), species.	COMMON NAME	COUNTRY OF ORIGIN OF STOCK	CULTURE MEDIUM	REARING TEMPERATURE °C.
DICTYOPTERA				
Blattidae				
<u>Blatta orientalis</u> L.	Oriental cockroach		18a	27
<u>Blattella germanica</u> (L.)	German cockroach		18a	27
<u>Periplaneta americana</u> (L.)	American cockroach		18a	27
<u>Periplaneta australasiae</u> (F.)	Australian cockroach		18a	27
DIPTERA				
Muscidae				
<u>Musca domestica</u> L.	Housefly	Britain	25	27
HYMENOPTERA				
Formicidae				
<u>Monomorium pharaonis</u> (L.)	Pharaoh's ant	Britain	33	27
Braconidae				
<u>Bracon hebetor</u> Say		America	31	25
LEPIDOPTERA				
Pyralidae - Pyralinae				
<u>Pyralis farinalis</u> (L.)	Meal Moth		5	25
Pyralidae - Phycitinae				
<u>Ephestia (Anagasta)</u>	Mediterranean	Britain	8a	25
<u>Kuehniella</u> (Zell.)	flour moth	Britain	8a	25
<u>Ephestia (Cadra)</u>	Tropical warehouse	(S. Africa)	8a	25
<u>cautella</u> (Walk.)	moth			
<u>Ephestia (Ephestia)</u>	Warehouse moth	Britain	8a	25
<u>elutella</u> (Hubn.)				
<u>Ephestia (Cadra)</u>		Cyprus	34a	30
<u>calidella</u> (Guen.)				
<u>Ephestia (Cadra)</u>		Cyprus	34a	30
<u>figulilella</u> Gregs.				
<u>Plodia interpunctella</u> (Hubn.)	Indian meal moth	Britain	8a	25
Pyralidae - Galleriidae				
<u>Achroia grisella</u> (F.)	Lesser wax moth		16a	25
<u>Galleria mellonella</u> (L.)	Honeycomb moth		16a	25
<u>Paralipsa gularis</u> (Zell.)			26	25

ORDER Family (-subfamily) Genus (sub genus), species.	COMMON NAME	COUNTRY OF ORIGIN OF STOCK	CULTURE MEDIUM	REARING TEMPERATURE °C
<u>Dermestes peruvianus</u> Castelnau		Britain	21a	25
<u>Trogoderma anthrenoides</u> (Sharp)		U.S.A.	2	30
<u>Trogoderma glabrum</u> (Herbst)		U.S.A.	2	30
<u>Trogoderma granarium</u> Everts	Khapra beetle	(Britain)	2	30
<u>Trogoderma grassmanii</u> Beal		U.S.A.	18	30
<u>Trogoderma inclusum</u> LeC.	Larger cabinet beetle		10	25
<u>Trogoderma irroratum</u> Reitt.		Egypt	2	30
<u>Trogoderma variabile</u> Ballion (=parabile Beal)		U.S.A.	2	30
<u>Trogoderma simplex</u> Jayne		U.S.A.	18	30
<u>Trogoderma sternale plagifer</u> Casey		New Mexico	32	30
Mycetophagidae				
<u>Typhaea stercorea</u> (L.)	Hairy grain beetle	Nigeria	4	25
Nitidulidae				
<u>Carpophilus dimidiatus</u> (F.)	Corn-sap beetle		14	25
<u>Carpophilus hemipterus</u> (L.)	Dried fruit beetle		15	25
<u>Carpophilus marginellus</u>			23a	25
Trogossitidae				
<u>Lophocateres pusillus</u> (Klug.)	Siamese grain beetle		11	30
<u>Tenebroides mauritanicus</u> (L.)	The Cadelle	Pakistan	12	30
Ptinidae				
<u>Gibbium psylloides</u> (Czemp)	Hump spider beetle	Britain	17a	20
<u>Mezium affine</u> Boield.		Britain	17a	20
<u>Mezium americanum</u> Lap.	American spider beetle		17a	20
<u>Niptus hololeucus</u> (Fald.)	Golden spider beetle	Britain	17a	20
<u>Pseudeurostus hilleri</u> (Reitt.)		Britain	17a	20
<u>Ptinus clavipes</u> Panz.	Brown spider beetle	Britain	17a	20
<u>Ptinus pusillus</u> Sturm.			17a	20
<u>Ptinus sexpunctatus</u> Panz.			17a	20
<u>Ptinus tectus</u> Boield	Australian spider beetle		19a	25
<u>Stethomezium squamosum</u> Hint.	African spider beetle	Britain	17a	20
<u>Ptinus unicolor</u> (P. & M.)		Kenya	17a	20
<u>Trigonogenius globulus</u> Sol.	Globular spider beetle	Ireland	17a	20
<u>Trigonogenius particularis</u> Pic		Kenya	18a	25

ORDER Family (-subfamily) Genus (sub genus), species.	COMMON NAME	COUNTRY OF ORIGIN OF STOCK	CULTURE MEDIUM	REARING TEMPERATURE °C
Silvanidae				
<u>Ahasveras advena</u> (Waltl)	Foreign grain beetle	(West Africa)	10	25
<u>Cathartus quadricollis</u> (Guer.)	Square-necked grain beetle	W. Africa	10	25
<u>Oryzaeophilus mercator</u> (Fauv.)	Merchant grain beetle		10	25
<u>Oryzaeophilus surinamensis</u>	Saw-toothed grain beetle		10	25
Tenebrionidae				
<u>Alphitobius diaperinus</u> (Panz.)	Lesser mealworm		7	25
<u>Alphitobius laeviagatus</u> (F.)	Black fungus beetle		7	25
<u>Alphitobius viator</u> Muls. & God.		Sierra Leone	7	25
<u>Alohitophagus bifasciatus</u> (Say)	Two-banded fungus beetle	Britain	5	25
<u>Gnathocerus cornutus</u> (F.)	Broad-horned flour beetle		17	25
<u>Gnathocerus maxillosus</u> (F.)	Slender horned flour beetle		6	25
<u>Latheticus oryzae</u> Waterh.	Long headed flour beetle		6	30
<u>Palorus laesicollis</u> (Fairm.)		Kenya	24	25
<u>Palorus ratzeburgii</u> (Wissm.)	Small-eyed flour beetle		6	25
<u>Palorus subdepressus</u> (Woll.)	Depressed flour beetle	Turkey	7	25
<u>Tenebrio molitor</u> L.	Yellow mealworm		10a	25
<u>Tenebrio obscurus</u> F.	Dark mealworm		10a	25
<u>Tribolium anaphe</u> Hint.		Nigeria	17	25
<u>Tribolium brevicornis</u> LeC.		U.S.A.	23	25
<u>Tribolium castaneum</u> (Herbst)	Rust-red flour beetle	Britain	23	25
<u>Tribolium confusum</u> Duv.	Confused flour beetle		23	25
<u>Tribolium destructor</u> Uytt.	Dark flour beetle	(Holland)	17	25
<u>Tribolium madens</u> (Charp.)	Black Flour beetle	(Yugoslavia)	17	25
Languriidae				
<u>Pharaxonotha kirschi</u> (Reitt.)			6a	25

The letter "a" after a number indicates that drinking water is added to the culture either in the form of damp blotting paper or as a corked tube of water containing a wick of blotting paper.

CULTURE MEDIA

No.	Food	Weight Rat (Ounces)
1.	Wheat	
2.	Wheat + wheatfeed	
3.	Wheat + wheatfeed + glycerol	7:3
4.	Wheat + wheatfeed + glycerol on a damp pad of cotton wool	7:3:1
5.	Wheat + wheatfeed on a damp pad	7:3:1
6.	Wheatfeed + yeast	7:3:1
7.	Wheatfeed + yeast on a damp pad	10:1
8.	Wheatfeed + yeast + glycerol	10:1
9.	Wheatfeed + yeast + glycerol on a damp pad	10:1:3
10.	Wheatfeed + rolled oats + yeast	10:1:3
11.	Wheatfeed + rolled oats + yeast + groundnuts	5:5:1
12.	Wheatfeed + rolled oats + yeast + groundnuts + cork	5:5:1:1
13.	Wheatfeed + rolled oats on a damp pad	5:5:1:1
14.	Rolled oats + yeast	2:1
15.	Rolled oats + yeast + dates	10:1
16.	Wheatfeed + rolled oats + yeast + glycerine + honey + brood comb	6:1:6
17.	Wheatfeed + fishmeal + yeast	5:5:1:2
18.	Wheatfeed + rolled oats + fishmeal + yeast	8:4:1
19.	Fishmeal + yeast	5:5:2:1
20.	Fishmeal + yeast + flannel	16:1
21.	Fishmeal + yeast + bacon ends	16:1
22.	Fishmeal + yeast + bacon ends + cheese	16:1
23.	Wholemeal flour + yeast	16:1
24.	Wheatfeed + rolled oats + flour + yeast	12:1
25.	Wheatfeed + grassmeal + yeast + shortex	3:3:3:1
26.	Groundnuts	20:10:1
27.	Haricot beans	
28.	Butter beans	
29.	Cowpeas + dried green peas	
30.	Liver + sugar and water	1:1
31.	Moth culture (Sub-family Phycitinae)	
32.	Fishmeal + Yeast + dried cockroaches	
33.	Liver, swiss roll and honey	
34.	Wheatfeed + glucose + yeast	
35.	Wheatfeed + fishmeal + yeast + cholesterol	5:2:1
36.	Crushed dog biscuit + yeast	8:8:1:1
37.	Bread and butter	20:1
38.	Sweet biscuits	
39.	Maize	
40.	Wood sawdust + bones	

1. Resistant strains

A. Tribolium castaneum (Tenebrionidae)

- | | |
|---------------------------------|--------------------|
| 1. DDT resistant | South Africa, 1960 |
| 2. Non-specific resistant | Australia, 1968 |
| 3. Lindane specific resistant | Zambia, 1970 |
| 4. Malathion specific resistant | Nigeria, 1963 |
| 5. Insecticide susceptible | Unknown, 1970 |

II. Mutants

<u>MUTANT STOCKS</u>	<u>MUTATION</u>	<u>COUNTRY OF ORIGIN OF STOCK</u>	<u>MEDIA</u>	<u>TEMP.</u>
Bostrichidae				
<u>Rhyzopertha dominica</u> (F.)	Black		1	30
Bruchidae				
<u>Callosobruchus maculatus</u> (F.)	Giant		29	30
Cucujidae				
<u>Cryptolestes pusillus</u> (Schön)	Black	Trinidad	10	30
Dermestidae				
<u>Dermestes maculatus</u> Deg.	Black/ Brown	Australia	21	25
Nitidulidae				
<u>Carpophilus dimidiatus</u> (F.)	Pearl-eyed		10	25
Silvanidae				
<u>Ahasveras advena</u> (Waltl)	Black	Britain	10	25
<u>Oryzaephilus surinamensis</u> (L.)	Small	Burma	10	25
Tenebrionidae				
<u>Latheticus oryzae</u> Waterh.	Dark		6	30

<u>MUTANT STOCKS</u>	<u>MUTATION</u>	<u>COUNTRY OF ORIGIN OF STOCK</u>	<u>MEDIA</u>	<u>TEMP.</u>
<u>Tribolium confusum</u> Duv.	Black			
<u>T. confusum</u>	Pearl-eyed	Malta	23	25
<u>T. confusum</u>	Pearl-eyed	Britain	23	25
<u>T. confusum</u>	Pearl-eyed		23	25
<u>T. confusum</u>	Black and Pearl-eyed		23	25

(Ed.)

SLOUGH, BUCKS, U.K.
TROPICAL STORED PRODUCTS CENTRE, MINISTRY OF OVERSEAS DEVELOPMENT

I. Wild type strains

COLEOPTERA

Anobiidae

Lasioderma serricorne

Cyprus, 1964

Silvanidae

Oryzaephilus surinamensis
Oryzaephilus surinamensis
Oryzaephilus surinamensis (bicornis)
Oryzaephilus surinamensis (Small)

Crete, 1964
 Cyprus, 1964
 Crete, 1964
 Far East, 1967

LEPIDOPTERA

Phycitidae

Cadra cautella
Cadra cautella
Cadra figulilella
Plodia interpunctella
Plodia interpunctella

Cyprus, 1964
 Rhodesia, 1965
 Cyprus, 1967
 South Africa, 1965
 N. Nigeria, 1965

(Ed.)

Stock Lists

INDIA

GORAKHPUR, U.P.
UNIVERSITY OF GORAKHPUR, DEPARTMENT OF ZOOLOGY

Wild type strain

1. Tribolium castaneum from local godowns.

(Ed.)

HISSAR, HARAYANA
PUNJAB AGRICULTURAL UNIVERSITY, DEPARTMENT OF GENETICS

I. Wild type strains (Tribolium castaneum)

1. IZT I
2. MAD I
3. PAU I
4. PAU II
5. Chicago wild
6. Brazil
7. Inbred lines in 8th. generation of full sibbing.

via Sokoloff, Berkeley
via Sokoloff, Berkeley

II.. Mutant strains (Tribolium castaneum)

S-8	<u>Py</u>	via Sokoloff, Berkeley
S-12	<u>P</u>	via Sokoloff, Berkeley
S-20	<u>Me</u>	via Sokoloff, Berkeley
S-24	<u>Squint</u>	via Sokoloff, Berkeley
S-26	<u>sa</u>	via Sokoloff, Berkeley
S-28	<u>mc</u>	via Sokoloff, Berkeley
S-35	<u>py r</u>	via Sokoloff, Berkeley
S-53	<u>jet</u>	via Sokoloff, Berkeley
S-71	<u>sa</u>	via Sokoloff, Berkeley
S-74	<u>ju</u>	via Sokoloff, Berkeley
S-81	<u>Bes</u>	via Sokoloff, Berkeley
S-90	<u>Fy r Mr</u>	via Sokoloff, Berkeley
S-100	<u>b Mo</u>	via Sokoloff, Berkeley
S-154	<u>Be Fta</u>	via Sokoloff, Berkeley
S-248	<u>Fta c ca</u>	via Sokoloff, Berkeley
S-253	<u>lod p</u>	via Sokoloff, Berkeley
S-304	<u>Msg</u>	via Sokoloff, Berkeley
S-313	<u>ser py r</u>	via Sokoloff, Berkeley
S-325	<u>Fta</u>	via Sokoloff, Berkeley
S-333	<u>Spa</u>	via Sokoloff, Berkeley
S-341	<u>r</u>	via Sokoloff, Berkeley
S-346	<u>Fas-3</u>	via Sokoloff, Berkeley
S-483	<u>pd</u>	via Sokoloff, Berkeley

(Ed.)

BAHAUDU SHAH TAFAR MARJ, NEW DELHI-1
MAULANA AZAD MEDICAL COLLEGE, DEPARTMENT OF BIOCHEMISTRY

T. castaneum

Wild strain of local origin

(Ed.)

JABALPUR, MADHYA PRADESH
J.N. AGRICULTURAL UNIVERSITY
COLLEGE OF VETERINARY SCIENCE & A.H.
DEPARTMENT OF ANIMAL BREEDING & GENETICS

1. Random Stocks: R-1, R-2, R-3, R-4, R-5, R-6, R-7, R-8, R-9, R-10.
PAU-1 (HSR-Wild).
2. Inbred Lines: I-1, I-2, I-3, I-4, I-5, I-6, I-7, I-8, I-9, I-10.

These stocks have been inbred for 19 generations.

3. Mutant stocks: S-1 Chi-wild
S-8 py
S-10 p
S-12 Chi b/b, Chi +/b, Chi +/+
S-53 jet
S-100 b Mo
S-248 Fta c Ca
S-304 msg
S-313 ser py r
S-333 Spa
S-341 r

ISRAEL

TEL AVIV, ISRAEL
TEL AVIV UNIVERSITY, DEPARTMENT OF ZOOLOGY

1. Stock list

Tribolium castaneum:

Wild type
++ (Purdue)

Mutants
black (bb)
sooty (ss)
extra urogomphi (eu)

Tribolium confusum

Mutants
black (mcGill) (bb)

Two other strains listed in TIB-15 were lost by infection.

All strains except eu were obtained from Dr. R. R. Sokal's laboratory, Stony Brook, N.Y. USA in 1970. The CS bb and CS ss stocks were recently contaminated by wild type beetles and therefore contain some wild-type genetic background.

Dr. David Wool

ITALY

PAVIA

UNIVERSITY PAVIA, CENTRO DE GENETICA

1. Tribolium confusum Duval, wild strain obtained from Professor A. Keck, Biological Institut, Regensburg.
2. id. id., strain of recent colonization from specimens collected in Pavia, small, difficult colony.

JAPAN

KYOTO

KYOTO UNIVERSITY, FACULTY OF AGRICULTURE
ENTOMOLOGICAL LABORATORY

1. Wild type

Bruchidae

Callosobruchus chinensis

Callosobruchus maculatus

Zabrotes bifasciatus

Kyoto

11 other districts in Japan
Louisiana, U.S.A.
California, U.S.A.
(Fresno, Lab., U.S.D.A.)

Burma

Israel

Thailand

Hong Kong

Tenebrionidae

Tribolium castaneum

Tribolium confusum

Kyoto

2. Mutant strain

Black mutant from a strain of Callosobruchus chinensis

MISIMA, SIZUOKA-KEN
NATIONAL INSTITUTE OF GENETICS

No stock list available.

(Ed.)

MIYAZAKI
MIYAZAKI UNIVERSITY, DEPARTMENT OF BIOLOGY

Alphitobius diaperinus--wild type strains
Callosobruchus shinensis--Kyoto strains
Martianus dermestoides--wild type strains
Palorus ratzeburgi--wild type strains
Sitophilus oryzae --wild type strains
Sitophilus zeamais--wild type strains
Tenebrio obscurus--wild type strains
Tribolium castaneum--wild type strains
Tribolium confusum--wild type strains

(Ed.)

OKAYAMA, JAPAN
OKAYAMA UNIVERSITY
FACULTY OF AGRICULTURE
LABORATORY OF APPLIED ENTOMOLOGY

Wild type

<u>Alphitobius diaperinus</u>	Miyazaki
<u>Callosobruchus chinensis</u>	Hiroshima Kyoto
<u>Gnathocerus cornutus</u>	Miyazaki
<u>Lasioderma serricorne</u>	Miyazaki
<u>Latheticus oryzae</u>	Miyazaki
<u>Oryzaeophilus mercator</u>	Miyazaki
<u>Oryzaeophilus surinamensis</u>	Miyazaki
<u>Palorus ratzeburgi</u>	Miyazaki
<u>Palorus subdepressus</u>	Miyazaki
<u>Rhizopertha dominica</u>	Miyazaki
<u>Sitophilus oryzae</u>	Miyazaki
<u>Sitophilus zeamais</u>	Miyazaki
<u>Stegobium paniceum</u>	Miyazaki
<u>Tenebrio obscurus</u>	Miyazaki
<u>Tribolium castaneum</u>	Miyazaki
<u>Tribolium confusum</u>	Miyazaki

T. Yoshida

MEXICO

CHAMPINGO
CAMPO EXPERIMENTAL "EL HORNO"

Tribolium castaneum
Tribolium confusum

Both cultures have long been maintained in our rearing chambers.
Their source is unknown.

(Ed.)

THE NETHERLANDS

AMSTERDAM, THE NETHERLANDS
ROYAL TROPICAL INSTITUTE
DEPT. AGRIC. RES.

<u>Cryptolestes ferrugineus</u>	P.I.L.
<u>Cryptolestes pusilloides</u>	P.I.L.
<u>Cryptolestes capensis</u>	P.I.L.
<u>Latheticus oryzae</u>	unknown
<u>Gnathocerus maxillosus</u>	Malawi, 1971
<u>Gnathocerus cornutus</u>	Malawi, 1971
<u>Carpophilus dimidiatus</u>	Malawi, 1971
<u>Caryedon serratus</u>	Senegal, 1970
<u>Callosobruchus maculatus</u>	unknown
<u>Tribolium castaneum</u> - Lindane resistant	Malawi, 1970
<u>Tribolium castaneum</u> - Lindane resistant + non-specific malathion resistance	
<u>Tribolium castaneum</u> (susceptible)	Malawi, 1971
<u>Tribolium confusum</u> (susceptible)	Malawi, 1971
<u>Sitophilus oryzae</u> (susceptible)	Malawi, 1971
<u>Sitophilus zeamais</u> I Lindane resistant	Malawi, 1971
<u>Sitophilus zeamais</u> II Lindane resistant	Malawi, 1971
<u>Oryzaephilus surinamensis</u> (small strain)	Thailand, 1972
<u>Oryzaephilus surinamensis</u>	unknown
<u>Oryzaephilus mercator</u> (small strain)	Thailand, 1970
<u>Oryzaephilus mercator</u>	Germany
<u>Trogoderma granarium</u>	Sudan, 1970
<u>Necrobia rufipes</u>	Ivory Coast, 1970
<u>Sitotroga cerealella</u>	unknown
<u>Plodia interpunctella</u>	China
<u>Corcyra cephalonica</u>	Ivory Coast
<u>Ephestia cautella</u>	Nigeria

NEW ZEALAND

NELSON
DEPARTMENT OF SCIENTIFIC AND INDUSTRIAL RESEARCH
ENTOMOLOGY DIVISION

- Stegobium paniceum--from infested rat food pellets at Otago University, Dunedin
- Oryzaephilus surinamensis--from infested rat food pellets at Otago University, Dunedin
- Gnathocerus cornutus--from infested rat food pellets at Otago University, Dunedin
- Sitophilus oryzae--from spaghetti in galleys of overseas ships at Port Nelson
- Sitophilus zeamais--from rice in galleys of overseas ships at Port Nelson

(Ed.)

PRIVATE BAG, HAMILTON
RUAKURA AGRICULTURAL RESEARCH CENTRE, DEPARTMENT OF AGRICULTURE

Tribolium castaneum

1. Wild type strains derived from imported strain from Edinburgh.
2. Mutant strain carrying the chromosome II mutant pearl (p) and obtained from Tribolium Stock Center, Berkeley, California.

A. R. Quartermain

PRIVATE BAG, PALMERSTON NORTH
DEPARTMENT OF SCIENTIFIC AND INDUSTRIAL RESEARCH
GRASSLAND DIVISION

Tribolium castaneum

1. Heavy and light population resulting from 18 generations of selection for increased and decreased pupal weight.
2. Strong, moderate and weak populations resulting from 20 generations of within-family selection for increased pupal weight at three selection intensities.

(Ed.)

PORTUGAL

LISBON

LABORATORIO DA DEFESA FITOSSANITARIA DOS PRODUTOS ARMAZENADOS
MINISTERIO DA ECONOMIA

The laboratory maintains the following cultures in the breeding room at 25° - 27° C and 65 - 70% R. H. The origin of the culture, the year of commencement and the culture media are given for each insect species.

<u>Acanthoscelides obtectus</u> (Say)--white bean	Coimbra, 1968
<u>Anagasta kuhniella</u> (Zell.)--bran and glycerine	Carcavelos, 1968
<u>Cadra cautella</u> (Walk.)--decorticated almonds	Algarve, Portugal, 1969
<u>Callosobruchus maculatus</u> (F.)--California black eye	Alcobaca, 1968
<u>Gnathocerus cornutus</u> (F.)--whole-meal flour and yeast	Portugal, 1969
<u>Lasioderma serricorne</u> (F.)--bran and dried yeast	Lisboa, 1964
<u>Oryzaephilus surinamensis</u> (L.)--broken wheat	Portugal, 1960
<u>Plodia interpunctella</u> (Hübner)--bran and glycerine	Carcavelos, 1968
<u>Rayzopertha dominica</u> (F.)--wheat	S. Tomé, W. Africa, 1969
<u>Sitophilus granarius</u> (L.)--wheat	Portugal, 1969
<u>Sitophilus oryzae</u>	Portugal, 1969
<u>Sitotroga cerealella</u> (Oliv.)--barley	Portugal, 1968
<u>Tenebroides mauritanicus</u> (L.)--broken maize, bran and dried yeast	Portugal, 1967
<u>Tribolium castaneum</u> (Herbst)--flour	Bissau (Guiné), 1957
<u>Zabrotes subfasciatus</u> --white bean	Lisboa, 1968

(Ed.)

SPAIN

MADRID

INSTITUTO NACIONAL DE INVESTIGACIONES AGRARIAS
LABORATORIO DE GENETICA DE POBLACIONES

Tribolium castaneum

A. Wild type strains

1. Consejo	CSIC, Madrid, Spain, 1964
2. Purdue	Purdue, USA, 1964
3. Edinburgh 1	Edinburgh, Scotland, 1970
4. Edinburgh 2	Edinburgh, Scotland, 1970
5. Campanario	Campanario, Spain, 1973

B. Mutant type strains

6. Black Purdue

Purdue, USA, 1964

C. Experimental lines

Originated from the "Consejo" strain and selected for egg laying performance through 42 generations.

		<u>Selected for</u>		<u>Temperature</u>
7.	AN - I	high performance	at	33° C
8.	AN - II	high performance	at	33° C
9.	AF - I	high performance	at	28° C
10.	AF - II	high performance	at	28° C
11.	AT - I	high performance	at	38° C
12.	AT - II	high performance	at	38° C
13.	BN - I	low performance	at	33° C
14.	BN - II	low performance	at	33° C
15.	BF - I	low performance	at	28° C
16.	BF - II	low performance	at	28° C
17.	BT - I	low performance	at	38° C
18.	BT - II	low performance	at	38° C
19.	RN - I*	high cross performance	at	33° C
20.	SN - I*	high cross performance	at	33° C
21.	RN - II	high cross performance	at	33° C
22.	SN - II	high cross performance	at	33° C
23.	RF - I	high cross performance	at	28° C
24.	SF - I	high cross performance	at	28° C
25.	RF - II	high cross performance	at	28° C
26.	SF - II	high cross performance	at	28° C
27.	RT - I	high cross performance	at	38° C
28.	ST - I	high cross performance	at	38° C
29.	RT - II	high cross performance	at	38° C
	ST - II	high cross performance	at	38° C

* R & S corresponding lines were selected through a reciprocal recurrent selection.

31 - 38 Inbred lines with 26 generations of full sibbing.

		High performance at different levels of select.						
39.	ATD-I	"	"	"	"	"	"	"
40.	ATD-II	"	"	"	"	"	"	"
41.	BTD-I	"	"	"	"	"	"	"
42.	BTD-II	"	"	"	"	"	"	"
43.	CTD-I	"	"	"	"	"	"	"
44.	CTD-II	"	"	"	"	"	"	"
45.	DTD-I	"	"	"	"	"	"	"
46.	DTD-II	"	"	"	"	"	"	"
47.	ETD-I	"	"	"	"	"	"	"
48.	ETD-II	"	"	"	"	"	"	"
49.	FTD-I	"	"	"	"	"	"	"
50.	FTD-II	"	"	"	"	"	"	"

D. Mutants

	<u>Source and date</u>
51. antennapedia <u>ap</u> , VIII	Purdue, 1964
52. Bar eye <u>Be</u> , IV	Purdue, 1968
53. Black <u>b</u> , III	Sokoloff, 1964
54. chestnut <u>c</u> , VII	Purdue, 1964
55. cerdeban <u>cd</u> , III	Purdue, 1964
56. Diferencial <u>Df</u> , IV	Purdue, 1964
57. fussed antennal segm. -2 <u>fas-2</u> , IV	Sokoloff, 1968
58. ivory <u>i</u> , ?	Purdue, 1964
59. jet <u>j</u> , V	Purdue, 1964
60. juvenile urogenophy <u>ju</u> , IV	Purdue, 1964
61. light ocular diaph. <u>lod</u> , III	Purdue, 1964
62. maroon <u>m</u> , V	Purdue, 1964
63. microcephalic <u>mc</u> , V	Purdue, 1964
64. paddle <u>pd</u> , I	Purdue, 1964
65. pearl <u>p</u> , II	Sokoloff, 1968
66. pegleg <u>pg</u> , II	Purdue, 1968
67. pink <u>pPk</u> , II	Purdue, 1968
68. pygmy <u>py</u> , I	Purdue, 1968
69. red <u>r</u> , I	Purdue, 1968
70. ring <u>rg</u> , I	Purdue, 1964
71. rose, <u>rs</u> , I	Purdue, 1964
72. ruby <u>rb</u> , ?	Purdue, 1964
73. Short antenna <u>Sa</u> , VII	Purdue 1. 1964
74. short elytra <u>sh</u> , VIII	Purdue, 1964
75. sooty <u>s</u> , IV	Purdue, 1964
76. spotted <u>sp</u> , I	Purdue, 1964
77. squint <u>sq</u> , VIII	Purdue, 1964
78. white <u>w</u> , ?	Purdue, 1964
79. wine <u>rw</u> , I	Purdue, 1968
80. eye mutant ?	Madrid, 1967
81. elytra mutant ?	Madrid, 1967
82. melanetic stink gland-like ?	Madrid, 1968

Tribolium confusum

A. Wild type strains

83. Campanario	Campanario, Spain, 1973
84. Santa Maria	Santa Maria, Spain, 1973
85. Mandayona	Mandayona, Spain, 1973

B. Mutant type strains

86. black <u>b</u> , III	Sokoloff, 1968
87. creased abdominal sternites <u>cas</u> , II	Sokoloff, 1968
88. ebony-2 <u>e-2</u> , II	Sokoloff, 1968

Stock Lists

YUGOSLAVIA

ZAGREB, KACICEVA 9
INSTITUTE FOR PLANT PROTECTION
AGRICULTURAL FACULTY

I. Wild type strain

LEPIDOPTERA

Gelechiidae

Sitotroga cerealella (Oliv.)

Phycitidae

Anagasta kubniella Zell.

COLEOPTERA

Bostrichidae

Rhizopertha dominica (F.)

Bruchidae

Acanthoscelides obtectus (Say)

Cucujidae

Cryptolestes spp. (Species not yet identified, but ferrugineus
and pusillus are present)

Curculionidae

Sitophilus zeamais Motsch.

Sitophilus oryzae (L.)

Sitophilus granarius (L.)

Dermestidae

Attagenus megatoma (F.)

Attagenus piceus (Oliv.)

Trogoderma granarium Everts

Ostomatidae

Tenebrioides mauritanicus (L.)

Ptinidae

Mezium spp. (species not yet identified)

Silvanidae

Oryzaephilus surinamensis (L.)

Oryzaephilus surinamensis (L.) v. bicornis

Oryzaephilus mercator (Fauv.)

Stock Lists

Tenebrionidae

Gnathocerus cornutus (F.) |
Palorus spp. (species not yet identified but ratzeburgi and
subdepressus are present)
Tenebrio molitor L.
Tribolium castaneum (Herbst).
Tribolium confusum Duv.

All insects are originated from storehouses and mills from Croatia, Yugoslavia. They are reared in a lab under constant circumstances during 3-4 years. Only species Trogoderma granarium is of unknown origin, collected from imported foodstuffs. This species is not found in Yugoslavia yet.

II. Mutants

Tribolium confusum

Chromosome III

Yugoslavian black (=bZ)--Yugoslavia 1969 (report of A.
 Sokoloff, TIB 13)

Zlatko Korunic

CZECHOSLOVAKIA

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Sitophilus granarius L.
S. eryzae L.
Rhizopertha dominica F.
Trogoderma granarium Ev.
Tribolium confusum Duv.
T. destructor Uytt.
Tenebrio molitor L.
Oryzaephilus surinamensis L.
O. mecatator Fauv.
Carpophilus hemipterus L.
Stegobium paniceum L.
Acanthoscelides obtectus Say
Anagasta kuhniella Zell.
Cadra cautella Wik.
Plodia interpunctella Hub.

New Mutants

Tribolium castaneum

squint-like (sql) Report of A. A. Dewees. This is an autosomal recessive of good penetrance, expressivity, and viability found at an estimated gene frequency of approximately .14 in a sack of contaminated white flour. Tests of allelism indicate that the gene is not an allele of squint (sq) but it has the identical phenotype. Ommatidia are absent in the adult but the ocular diaphragm is visible through the exoskeleton and has a considerably reduced or absent ocular foramen giving rise to the squint-eye appearance. Linkage studies are presently under way.

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Feeding and oviposition preference in *T. audax*, *T. madens*, *T. castaneum*,
T. confusum and *T. brevicornis*

As part of a laboratory experiment in the Population Genetics and Ecology course we introduced 50 unsexed beetles of each of the above species into three plastic choice chambers such as those designed and illustrated by Inouye (196_). The choice chambers contained equal quantities of two media: corn plus yeast (C Y); soy plus yeast (S Y) and standard medium, i. e. wheat plus yeast (W Y). The media were separated by a plastic divider. Sizable holes in this divider allowed free passage of the beetles from one side of the chamber to the other. The beetles were counted and placed on the surface of the two flours in the middle of the chamber. The chambers were placed in an incubator maintained at 32° C., and about 60 per cent R. H. for a period of 48 hours. After this period the contents of each side of the chamber were aspirated into a vial, the material was then sifted with a coarse screen to remove the adults and a fine screen to isolate the eggs. The adults were then sexed and counted and the number of eggs laid in each medium w_p s recorded.

The results are summarized in Table 1. A preliminary analysis of these limited data leads to the following conclusions:

For adult medium preference,

1. Given the choice of WY and SY all the species except *T. brevicornis* are found more often in WY than in SY. *T. brevicornis* adults were found about equally distributed in the two media.

2. Given the choice of CY and WY, *T. confusum*, *T. madens* and *T. audax* (all in the *confusum* species group) seem to prefer the corn plus yeast over the wheat plus yeast medium. *T. castaneum*, on the other hand, is found more often in the wheat plus yeast medium. *T. brevicornis* is found about equally frequently in these two media.

3. Given the choice of CY and SY, *T. audax*, *T. madens*, *T. castaneum* and *T. confusum* adults seldom are found in soy plus yeast. *T. brevicornis*, on the other hand, is found equally frequently in the two media.

By itself, the occurrence of the adults in a medium at a given instant (in this case when the two media are separated) does not necessarily mean a preference for that particular medium. A stronger case can be advanced if the number of eggs found in the medium corresponds to the prevalence of adults.

According to the data in Table 1:

4. Given the choice of WY and SY *T. castaneum*, *T. audax* and *T. madens* prefer to lay their eggs in wheat than in soy. *T. confusum* accepts both media as an oviposition site, while *T. brevicornis* seems to prefer

soy to wheat medium as an egg-laying site.

5. Given the choice of WY and CY, T. castaneum prefers wheat, T. audax and T. confusum clearly prefer corn to wheat, and T. madens and T. brevicornis use the two media about equally frequently as an oviposition site.

6. Given the choice of CY and SY it is clear that T. audax, T. madens, T. castaneum and T. confusum females prefer CY to deposit their eggs. The data for T. brevicornis, on the other hand, reinforces the conclusion that the preferred medium by females of this species appears to be soy.

A previous study (Sokoloff, 1972 and Sokoloff and Cavataio, 1972) showed that T. brevicornis is unusual in that it can survive for as long as nine months without food or water. This study shows that T. brevicornis, in contrast to the other species, can exploit soy flour in spite of the presence of toxic substances which affect the developmental rate of Tribolium castaneum and T. confusum.

(For review see Sokoloff, Franklin, Overton and Ho, 1966).

Table 1. Food choice and oviposition of 5 species of Tribolium.

Media	<u>T. brevicornis</u>			<u>T. madens</u>			<u>T. audax</u>		
	♂	♀	eggs	♂	♀	eggs	♂	♀	eggs
WY	11	10	41	14	25	51	22	13	222
SY	16	10	67	5	6	12	3	5	27
WY	10	11	48	6	14	49	10	7	39
CY	12	13	36	15	12	58	14	16	188
CY	13	11	57	28	20	87	25	24	278
SY	14	13	106	2	0	0	0	0	0

Media	<u>T. castaneum</u>			<u>T. confusum</u>		
	♂	♀	eggs	♂	♀	eggs
WY	17	24	225	12	12	38
SY	0	1	9	1	7	49
WY	13	16	175	7	8	61
CY	6	12	15	12	20	117
CY	14	26	31	13	36	186
SY	0	1	3	1	0	3

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Life-span productivity in *Tribolium* for two single cross hybrids and parental inbreds.

The flour beetle *Tribolium* has become widely recognized as an excellent biological model for testing new concepts in ecology and genetics. Most selection studies to date with *Tribolium* have examined polygenic traits determined largely by additive gene effects. (e.g. developmental rate, larval weight at a fixed age, and pupal weight). In general, the results are in agreement with selection theory based on an additive gene model (Bell, 1969).

A major problem area today in population genetics concerns non-additive gene effects resulting from dominance and/or epistasis. The nature and utilization of this portion of the total genetic variation in a population is not well understood from either a theoretical or an experimental basis.

For *Tribolium* to serve effectively in model experiments relating to non-additive gene effects, appropriate polygenic traits are essential. Drawing from earlier experience with *Drosophila* (Bell, et al. 1955; Brown and Bell 1961) we have investigated certain components of fitness in *Tribolium castaneum* as possible models of nonadditive gene effects.

Our first study in this area examined life-span egg production, viability, and fertility of two unrelated single cross hybrids (C♀ x B♂ and Va♀ x K♂) and their respective parental inbred lines B, C, K and Va. Newly emerged males and females of the various genotypes were assigned in single pairs to individual creamers containing 2g of egg collection medium (95% whole wheat flour + 5% brewers' yeast mixture screened through a 0.21mm mesh). Daily egg collections were made for the first three weeks followed by periodic observations thereafter. Eggs were counted at 15X magnification and incubated for subsequent hatchability observations. All sterile matings were excluded from the results as summarized in Figure 1.

Even though the numbers observed for each genotype were small certain results were clearly manifested. Non-additive gene effects in terms of heterosis ($F_1 - MP$) were observed for three components of life-span productivity: (1). The hybrids reached sexual maturity or initiated egg lay some 3 days earlier than their inbred parents; (2). Hybrid egg lay was more than twice that of inbreds and the peak rate of lay was sustained longer, and (3). The hybrids lived longer than their parental inbreds. Among the former, 81% were still productive at the end of the 90 day test period while only 26% of the inbreds survived.

Another trait in *Tribolium* which has potential as a non-additive gene model is biomass, an aggregate trait which combines body weight

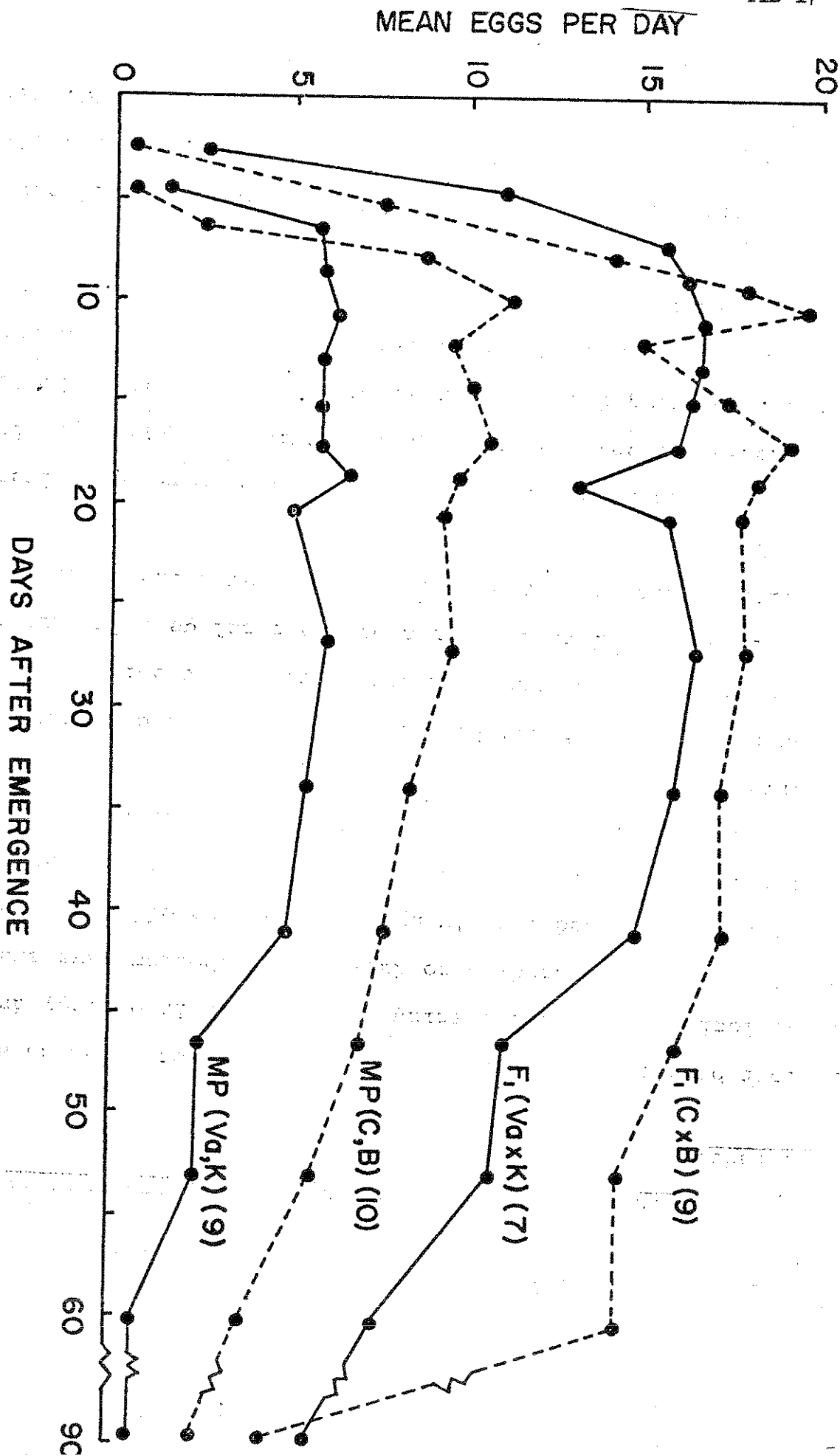


Figure 1. Daily rate of egg lay for fecundated females contrasting single cross (F₁) hybrids and mid-parental (MP) inbreds (Plotted as 2-day averages for the first 20 days and on a weekly basis thereafter. Number observed shown for each genotype).

with the fitness trait "number of offspring". Krause and Bell (1972) investigated biomass in two unrelated heterogeneous base populations (Purdue black and pearl) and found that 75-80% of the phenotypic variance for the fitness component was determined by rate of egg lay. Genetic effects on 24-hr. egg lay were largely non-additive.

More recently, Orozco and Bell (1974) found considerable additive genetic variance ($h^2=0.36 \pm .03$) for 4-day virgin egg lay in the Consejo Base Population of Tribolium when performance was measured in an optimum environment (33°C). However estimates for additive gene effects were considerable smaller ($h^2=.25 \pm .03$) for egg lay under severe stress (28°C), and non-additive effects assumed significant proportions (15%).

The above studies suggest that egg lay in Tribolium can serve as a useful biological model of non-additive gene effects. In addition, the introduction of optimum and stress testing environments provides an interesting genotype x environment model.

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Effect of Presence of Males on Egg Production in Tribolium castaneum.

A number of authors Ruano and Orozco (1956), Espejo and Orozco (1966), Kummer (1960) have shown that presence of males is essential for egg production in insects. Tagarro and Rico (1966) studied the effect of removing the male on egg laying rate of fecundated females of Tribolium castaneum. They mated a number of females to an individual male immediately after emergence and separated them on the 7th day. The egg production of individual female was recorded for 4 days. There was no significant drop in the rate of laying during this period.

Ruano and Orozco (1966) reported egg production of fecundated females of Tribolium castaneum as 67 ± 11 eggs and virgin females 18 ± 7 eggs for a period of 4 days between 7th and 11th day after emergence. Their findings indicated that presence of male affected the number of eggs laid by each female.

Espejo and Orozco (1966) reported that in Tribolium castaneum when fecundated females were deprived of their males the egg laying rate was maintained at a high level for a long period of time, the reduction in number of eggs was very slow. While when virgin females were mated they achieved immediately a very high egg laying rate, as high as in females mated from the beginning.

Kummer (1960) studied the effect of presence of male or male secretions on rate of egg laying in Drosophila melanogaster. The main conclusions of his experiments were that the process of copulation alone was not important, in fact both the secretion of paragonia and that of the testes were essential if full egg production by females had to be achieved. From these experiments there was an indication that presence of males might be necessary for complete egg production in females. In view of this an experiment was planned to study the effect of presence of males on egg production in females of Tribolium castaneum.

Material and Methods

The foundation population (PAU-I) of Tribolium castaneum was used in these experiments. This population was maintained in the laboratory for over a period of two years in mass cultures.

Culture media used consisted of 95% wheat flour and 5% yeast with vitamin mixture (A₁B₂ & D₃). The whole wheat flour was sieved through a filter of (0.7 mm pore size) sterilized in an oven for one hour at 150°C, yeast and vitamin mixture was then added to the sterilized flour. This mixture was mixed in a shaker at high speed, bottled and kept sealed in refrigerator for use.

The larvae were collected from the foundation population and kept in fresh media. The cultures were maintained at 32°C + 1°C and 70% relative humidity. Fifty newly emerged virgin females (0-4 hours old) were collected. Each female was randomly allotted to a replicate. Such two replicates were generated. Males were assigned to each female at random in replicate (II). The females of replicate (IV) were not provided with any males. The animals were raised on the egg media. The pair matings in replicate (II) and virgin females in replicate (IV) were raised on in P.V. 5g glass vials in 2g media.

Twenty four hour egg collection was recorded every day at 10 A.M. The eggs were transferred to new vial daily. The egg production was recorded for 31 days. On 31st day males were separated from replicate (II) and provided to virgin females in replicate (IV).

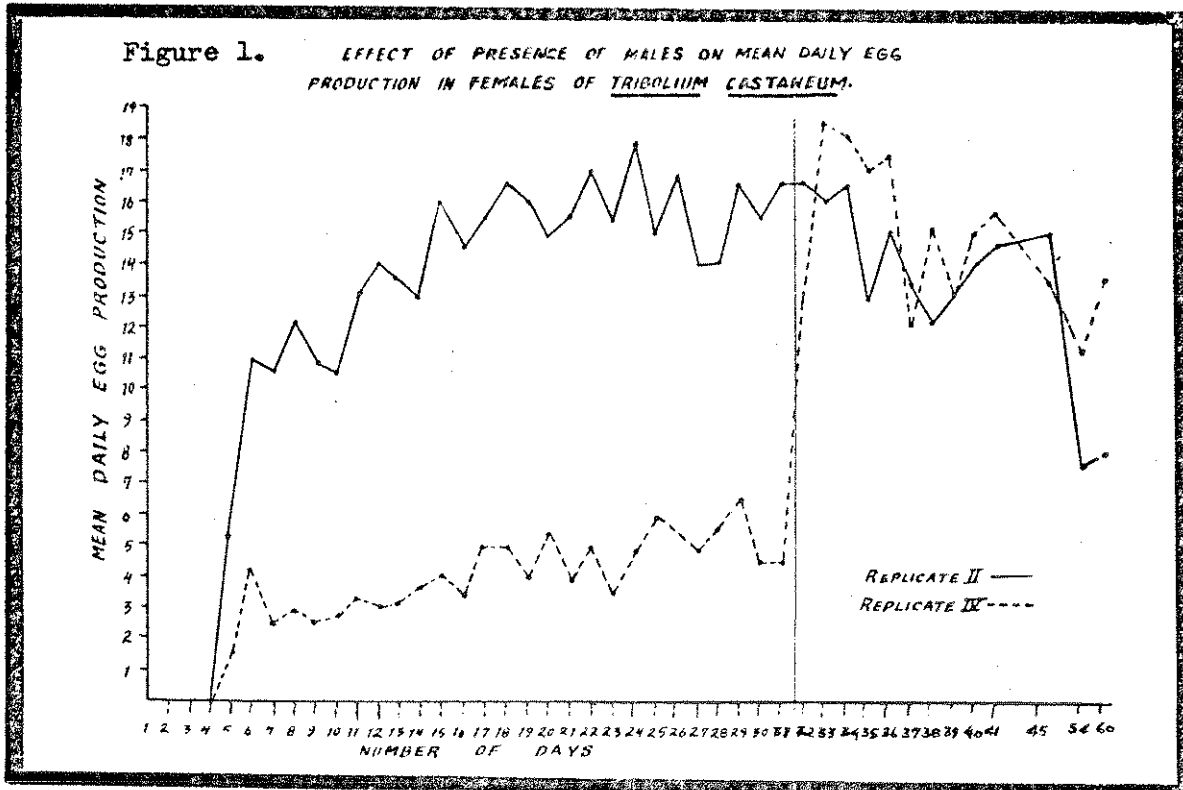
The egg collection was recorded daily for a further period of 10 more days. After 10 days egg production was recorded for a period of 24 hrs. every third day upto 60 days.

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Results

It was observed that those females which were provided with males as pair mates, produced nine eggs per day in the first week as against 2.8 eggs by virgin females (Table 1 and Fig. 1). Both the groups showed increase in egg production upto the 5th week. Females with males (pair mated group) showed three-fold increase in egg production compared to the virgin group. After a period of 31 days (after 5th week) the situation was reversed, the egg production in virgin group which was provided now with males shot up by three-folds (16.60) in the 6th week and then levelled off around the average egg production of pair mated group. The egg production in the pair mated group whose males were removed fell steadily and levelled off, around the level of egg production of virgin group (after three to four weeks) while the virgin group increased their production within a period of twenty four hours.



Discussion

It was observed that there was a significant difference in egg production between two treatments (pair mated/virgin group). The difference was three-fold under similar set of controlled environmental conditions. The only conclusion which can be drawn was that males are necessary for complete egg laying. This was in conformity with the finding of Ruano and Orozco (1966) who reported a three-fold difference in virgin and pair mated group, even though their findings were based on the average of 4 day production between 7th and 11th day of egg laying. Tagarro and Rico (1966) could however not find any significant difference between the two sets in contradiction of Ruano and Orozco (1966) from the same laboratory. They observed that if matings were done after emergence and kept for 7 days and recorded for 4 days after separation, they found no significant drop in egg production. It may be observed here

Table 1. Effect of presence of males on egg production of females in Tribolium castaneum.

No. of replicates	No. of eggs produced each week									
	1	2	3	4	5	6	7	8	9	10
1. (i) II (Males provided)	9.00	11.60	13.31	15.79	16.20					
(ii) II (Males removed)						12.27	13.70	15.00	7.7	7.9
2. (i) IV (Virgin females)	2.80	2.93	4.56	4.81	4.10					
(ii) IV (Males provided)						16.60	14.63	13.40	11.3	13.7

that it takes 96 hours for the first egg to be formed (Bhat, 1970). Consequently mating immediately after emergence is of no consequence. The peak egg production in Tribolium castaneum was reported to be from 7th to 11th day (Bhat, 1971), it would not be unusual to expect no fall in egg production time. It took roughly three to four weeks for egg production to fall to the level of virgin group.

It was observed that the physical presence of male was not important, but copulation followed by injection of semen was important because after the males were removed from the pair mated group in the 6th week, there was no immediate decrease in egg production of this group. It took almost three to four weeks for a significant drop in egg production to be registered. It was argued that if only the physical presence of males was important for enhancing egg production then the egg production would drop immediately after the separation, instead decrease in egg production was very slow, hence physical presence of male as a factor was not important and the injection of semen was necessary for enhancing egg production. These results are in conformity with those of Kummer (1960) who reported that both the secretion of paragonia and that of testes were essential if full egg production by females had to be achieved in Drosophila melanogaster.

It was further observed that virgin group when provided with males showed an immediate rise in egg production within 24 hours and then levelled off at the level of pair mated group, indicating that something must have transferred from the males to the virgin females and enhanced the egg production. Similar results have been reported by Espejo and Orozco (1966).

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The egg production curve in flour beetle *Tribolium castaneum*.

The egg production curve was studied on the basis of average daily production per surviving female. The curve is generally J-shaped and clearly divisible into two portions. One characterized by an initial increase upto the 8th day followed by a stable and persistent plateau which is essentially linear. Regression functions were fitted to both the portions of the curve from 4th to 7th and 8th to 31st day. There was a linear increase from 4th to 7th day by 3.82 eggs per day. From 8th to 31 day of life the value of regression coefficient was 0.118. The egg production from 5th to 10th day was correlated with total life production in the following order: 5th to 7th, 6th to 8th, 7th to 9th, 8th to 10th, 5th to 10th, 6th to 10th and 7th to 10th. In addition rank correlations were also estimated. The value of product moment and rank correlation coefficients was highest between 8th to 10th day period and the life time production. These were also the peak days of egg production. Each ovariole produced 1.02 eggs per day.

The total number of eggs produced over the life time period was correlated with ovariole number. The product moment correlation coefficient was estimated as 0.59 which was significantly different from zero. It was concluded on the basis of this study that egg production on days 8, 9 and 10th could fairly represent the egg production on life time basis.

Introduction

Flour beetles of the genus *Tribolium* constitute important primary and secondary pests in all kinds of cereal products. For this reason, they have attracted the attention of investigators with a broad spectrum of interests. Since Chapman (1924) used and stressed the usefulness of *Tribolium* in population studies, *Tribolium castaneum* has received considerable attention from geneticists, ecologists, morphologists and students of evolution because of the interesting material available in beetles and their suitability to various phases of research.

Egg production is highly variable and slight changes in environment greatly influence its expression. Studies on the egg production curve have been reported by Robertson (1957), Narain (1962) in *Drosophila melanogaster* and by Bhat (1961) in *Drosophila ananassae*. The shape of the curve has generally been described as that of an inverted J. Regression equations were fitted by these authors to see if the larger part of the curve was generally linear. The egg production per day was of the order of 32.1 with peak egg production on 4th, 5th and 6th day of emergence in both species.

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Material and Methods

The foundation population of Tribolium castaneum used in these experiments was obtained from the Department of Genetics, P. A. U., Hissar. The line used in this study was P. A. U. -1 foundation population, maintained in the laboratory for over a period of two years in mass cultures.

The animals were raised in whole wheat medium which consisted of 95% wheat flour and 5% yeast and vitamin mixture (A₁, B₂ and D₃). The whole wheat flour was sieved through a sifter (0.7 mm pore size) sterilized in an oven for one hour at 250°C. Yeast and vitamin mixture were in a shaker at high speed, bottled and kept sealed in a refrigerator until ready for use.

The larvae were collected from the foundation population of Tribolium castaneum and kept in fresh medium. The cultures were maintained in the temperature cabinet at 32 ± 1°C and 70% relative humidity.

The pupae were collected and sexed. These were watched for the emergence of the virgin adults. The freshly emerged adults were whitish in colour and could be easily identified; Hundreds of such females (0-4 hours old) were collected. Each female was randomly allotted to a replicate. Three such replicates were generated. Males were assigned to each female at random in three replicates (I, II and III).

The animals raised on the media wire sieved through a filter with a pore size of 0.125 mm.

The pair matings (one male and one female) were raised on in P.V. 5 g glass vials in 2 g media.

24 hours egg collection was recorded every day at 10:00 a.m. by shifting the contents of each vial through a sieve with the pore size of 0.125 mm. Since the grain size of media was 0.125 mm, the media would pass through with ease and the eggs would remain in the sieve, the egg size being 0.30 mm. They could not pass through. The eggs were transferred to a thick black paper, counted and recorded. The animals were then transferred to new vials daily. The egg production was recorded daily for 31 days.

Results

Daily egg production in three replicates (I, II and III) each of 20 females were studied for a period of 31 days. The results are detailed in table 1.

Table 1. Mean daily egg production for first twelve days of life and subsequently based on weekly totals in Tribolium castaneum.

Number of replicates		Number of Days of Life											
		1	2	3	4	5	6	7	8	9	10	11	12
I	(\bar{X})					2.9	11.0	11.7	10.6	11.4	11.1	8.8	12.4
II	(\bar{X})					5.3	11.1	10.6	12.1	10.9	10.5	12.9	14.0
III	(\bar{X})				2.2	12.0	8.2	16.2	15.0	14.0	13.3	16.2	14.0
	(\bar{X})				0.7	6.8	10.2	12.3	12.6	11.9	11.5	13.0	13.6

		Number of Weeks				
		1	2	3	4	5
I	(\bar{X})	8.60	10.91	13.47	13.19	14.60
II	(\bar{X})	9.00	11.60	13.31	15.79	16.20
III	(\bar{X})	9.70	14.50	14.01	11.10	14.10
	(\bar{X})	7.5	12.30	14.57	14.30	15.23

The beetles on an average laid 381.14 ± 27.9 eggs for a period of 31 days. The egg production curve, drawn on the basis of average daily production per surviving female, is presented in figure 1.

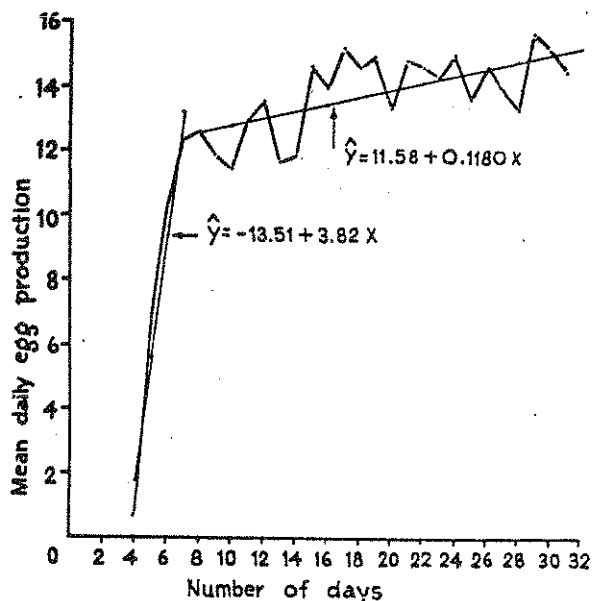


Fig.1. Mean daily egg production in Tribolium castaneum.

From this figure and table 1 it was observed that in one replication eggs were laid during the fourth day and in other two, on the fifth day.

The egg production increased progressively during the first few days, reaching the peak on the 8th day. The average egg production per female per day was found to be 12.6, 11.9, 11.5, 13.0 on 8th, 9th, 10th and 11th day of egg laying. Thereafter, the production fluctuated between 14.30 to 15.23 eggs per day. The properties of egg production curve were studied. The curve is generally J shaped and clearly divisible into two portions. One characterized by an initial increase upto the 8th day, followed by a fluctuant curve which was essentially linear. Regression lines were fitted to both portions of the curve, from 4th to 7th day and 8th to 31st day. The regression equations are detailed below:

<u>Days</u>	<u>Regression coefficient</u>	<u>Regression equation</u>
4 - 7	3.820	$Y = -13.51 + 3.820 x$
8 - 31	0.118	$Y = 11.58 + 0.118 x$

Y = Number of eggs per female per day.
X = Number of days.

The curve shows a linear increase from 4th to 7th day by 3.82 eggs per day. From 8th to 31st day of life, the value of regression coefficient was 0.118 suggesting only slight change in egg production after the 8th day of life.

The egg production from 5th to 10th day was correlated with total life time production in the following order: 5th to 7th, 6th to 8th, 7th to 9th, 8th to 10th, 5th to 10th, 6th to 10th, and 7th to 10th day. The results are detailed in table 2.

Table 2. Product moment and rank correlation coefficients between various periods and life time production in Tribolium castaneum.

<u>S.No.</u>	<u>Period (days)</u>	<u>Correlation coefficient</u>	<u>Rank correlation</u>
1.	5, 6, 7	0.538**	0.585
2.	6, 7, 8	0.677**	0.690
3.	7, 8, 9	0.714**	0.575
4.	8, 9, 10	0.777**	0.837
5.	5 to 10	0.694**	0.742
6.	6 to 10	0.731**	0.799
7.	7 to 10	0.743**	0.781

** p 0.01

The 8th, 9th and 10th day of period had the highest (0.777) correlation coefficient with the life time production. It appears that during this period eggs laid will fairly estimate the life time egg production of these females.

At the end of the experiment the females were sacrificed and ovariole number was recorded. A total of 8124 eggs were laid by twenty two females for 31 days. The total number of ovarioles for these 22 females was 255. The total number of eggs for thirty one days produced by each ovariole was calculated to be 31.84 eggs. Total number of eggs produced by each ovariole per day was 1.02 eggs which work out to about one egg produced by one ovariole in a period of twenty four hours.

The total number of eggs produced over the life time period was correlated with ovariole number. The product moment correlation coefficient was estimated as 0.59 which was significantly different from zero.

Discussion

The egg production curve showed an initial progressive increase during the first few days reaching the peak on the 8th day. Thereafter, the production fluctuated between 14.3 to 15.2 eggs per day. Like most production curves, it is slightly J shaped. The initial rise in egg production was quickly followed by a fluctuant curve which was essentially linear. The regression coefficient of egg production on days from 4th to 7th day was 3.82 and from the 8th to 31st day was 0.118. The egg production increased at the rate of 3.82 eggs per day from 4th to 7th day and thereafter at the rate of 0.118 eggs per day.

Sonleitner (1965) reported fecundity and fertility in Tribolium castaneum at 3 days intervals upto 18th day of age. The number of eggs reported per day in the first week was 17.7, in the second week 21.3 and the maximum number of eggs produced per day was 23.2 eggs.

In these experiments the number of eggs produced per day in the first week and second week was 11.9 and 13.0 and the maximum number of eggs per day was 15.23. It seem that the stock used by Sonleitner (1965) was of higher egg producing potential. The results are in agreement with those of Ruano and Orzoco (1966). They reported 16.8 eggs in females which were provided with males. Fuentes and Ruano (1967) studied the effect of different temperatures on egg production in Tribolium castaneum. Maximum number of eggs were laid by females between 32° to 36°C.

The egg production from 5th to 10th day period of life was correlated with total life production (based on 31 days) in the various orders (Table 2). Combined 8th, 9th and 10th day egg production had the highest product moment and rank correlations with life time production. These were also the peak days of egg production.

Similar results were reported in Drosophila melanogaster by Robertson (1957) and Narain (1962) and in Drosophila ananasse by Bhat (1961). These authors reported rank order correlations between first 10 days and life time production. It was suggested that egg production was maximum on 4th, 5th, and 6th day and this could predict the total life time production. Since in Tribolium castaneum the product moment correlation and rank correlation was highest between the period 8th, 9th and 10th day and the total life time production, it could easily be used as an index to predict the life time production. The value of regression coefficient after the 8th day of life was small, which suggests that there is very little increase in egg production after the initial stability is reached.

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Bioenergetics in Tribolium Populations.

The bioenergetics of Tribolium populations were studied at three temperatures. Single-species populations containing either 52 T. confusum or 52 T. castaneum adults and mixed species populations containing 26 T. confusum and T. castaneum adults were reared at 25°, 30°, and 35° C. Caloric values of food consumed; caloric values of eggs, larvae, pupae, and adults; and caloric expenditures for respiration were determined at 10, 20, 30, 40, and 50 days.

Single-species population data indicated that T. castaneum consumed more food and had consistently higher total caloric values in living components (sum of egg, larva, pupa, and adult caloric values) at all temperatures. Respiration caloric values indicated that T. castaneum was metabolically more active. These data from single species populations indicate that T. castaneum is inately "superior" at all temperatures (higher metabolic rate, more rapid increase in population biomass). Furthermore, the differences between T. castaneum and T. confusum increased with increasing temperature.

Mixed-species population data indicated that interspecific interactions modify the outcome observed for single species. Effects of interspecific interactions were determined by subtracting biomass caloric values of single-species populations from biomass caloric values of single-species populations. T. castaneum was the predicted survivor in populations reared at 35° and 30° C. In populations reared at 35° C, both species had lower caloric values when reared together. T. castaneum had higher caloric values, whereas T. confusum had lower caloric values when mixed species values were compared to single species values in populations reared at 30°C. Experiments conducted at 25° C were not of sufficient duration to predict the surviving species. There was the suggestion however, that T. confusum gained in caloric biomass accumulation at the expense of T. castaneum in mixed-species populations.

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*A preliminary study of the decreased sexual competitiveness of X-irradiated males of *T. castaneum*.

Loss of sexual competitiveness frequently has been observed in male insects irradiated for sterile-male release tests for control of harmful insects. Thus if "sterilized" males are tested in a 4 : 1 : 1 ratio with normal males and females, 80% reduction in the number of viable progeny should be expected; actually, the reduction often is less than 50%. Most insect species for which sterile-male release measures have been proposed are relatively short-lived and/or lacking in somatic cell turnover in the adult stage. It is not known whether or not the decline in competitiveness represents a manifestation of the same injury which leads to (or hastens) death after higher doses, nor whether some recovery eventually occurs. Indeed, there have been few attempts to elucidate the mechanisms involved, although there is some evidence (Nair and Bhakthan, 1969) that there is damage to the flight muscles of irradiated flies. Apart from practical implications, the phenomenon is of great interest because it appears to represent the expression of radiation injury to nonproliferative tissues, which are regarded as highly radioresistant. If decreased competitiveness occurs in sublethally-irradiated males of groups such as Coleoptera, Hemiptera, or Orthoptera, it may be possible to dissociate this manifestation of injury from the lesions involved in acute lethality, which in these groups appears to stem from damage to proliferative cells, viz., the cells which renew the lining of the midgut (Ducoff, 1972). It may also be possible to detect and score recovery of competitiveness with the passage of time.

We have, accordingly, initiated a series of experiments to determine competitiveness of male *Tribolium* at various times after exposure to ionizing radiation under various conditions and patterns of exposure. Various numbers of irradiated males are placed with 1 nonirradiated male and 1 nonirradiated female for 3 successive 2-day periods at 30°; the adults are removed and the vials of flour-yeast medium with deposited progeny are incubated for several weeks, until the larvae are large enough for convenient counting. The productivity, P , of a given type of mating is the mean number of viable progeny/female/day. Using a slight modification of the method of Fried (1971), the total productivity, P_T , of a test mating is:

$$P_T = \frac{P_N + R C P_R}{1 + R C}$$

in which P_N and P_R are the respective mean productivities, in single-pair matings, of nonirradiated and of irradiated males; 1 and R represent the respective number of nonirradiated and of irradiated males in a test mating; and C is the competitiveness of the irradiated males. Competitiveness, then, is readily calculated, as

$$C = \frac{P_N - P_T}{R(P_T - P_R)}$$

Using T. castaneum exposed to 8100 R of 280 kVp X-rays and test ratios of 3 and of 6 irradiated males/1 nonirradiated male and 1 nonirradiated female, we find that males tested during the first week after exposure were about 50% competitive, but males tested for the first time in the sixth week were only 20% competitive. All acute deaths following midlethal irradiation occur within the first 4 weeks, so a significant amount of injury remains, presumably in nonproliferative tissues. (Work supported by USPHS grant CA 13779).

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Sire, Dam and Age Effects on Offspring Number in *Tribolium castaneum*.

Progeny were obtained from eighty-five single pair matings of *Tribolium* selected at random from a large population maintained with a minimum of inbreeding, by sampling eggs from the parents for forty-eight hours following the time when they reached 35 days of age. Pupae from all matings were separated by sex and classed by family size in five groups (15-22, 23-30, 31-38, 39-46 and 47-54 offspring per family). Male and female progeny from each class were selected at random and mated as pupae in a five by five mating plan, from the low class to the high class for sires and from the low class to the high class for dams with one mating per cell. Ten replicates were obtained by sampling eggs from different sets of parents in the mating plan at 28, 33, 34, 35, 40, 41, 42, 47, 48 and 49 days of age to establish replicates one to ten respectively.

The effects of sire, dam, age and their first order interactions were tested for significance using X^2_r values from Friedman's 2 Way Analysis of Variance by Ranks (Bradley 1968).

The simple effect of sire family size on offspring number is not significant (Table 1, $\alpha > 0.250$); however, the sire by dam family size interaction is significant (Table 1, $\alpha < 0.050$). The interaction of sire and dam family size on offspring number is due to the lowest two dam family sizes (15-22, 23-30) in each sire family size class. As sire family size levels increase from S_1 to S_5 , the relationship of the mean number of offspring in the lower two dam family size classes changes. The value of the mean offspring family size for the second lowest dam family class (23-30) minus the lowest dam family class (15-22) changes from negative to higher positive values (ie. -15.4, +5.0, +9.5, +9.7, +13.9) across sire classes.

Dam family size has a highly significant effect on offspring number (Table 1, $\alpha < 0.010$). Mean offspring family size rises across the lower three dam family size classes (Table 1, $D_1, D_2, D_3, \alpha < 0.025$); with the middle class (31-38, D_3) as the pivotal point, the mean offspring family size plateaus (Table 1, $D_3, D_4, D_5, \alpha > 0.950$).

The effect of age of parents at time of mating on offspring number was highly significant (Table 1, $\alpha < 0.0005$). The mean size of offspring family increases rapidly from maturity to a peak at approximately thirty-five days of age, then drops off slowly to the end of reproductive life.

Some non-mutant specimens of *T. castaneum* of different age groups were dissected; in newly emerged soft bodied adults I found persistent pupal ventral muscles (as well as tergal muscles) overlying loosely all the ventrites and underneath them the normal adult ventral muscles in the mobile ventrites 3-5 as well as some small and sometimes asymmetrical bundles in the anterior connate ventrites 1 & 2, like those of the last 3 ventrites (Fig. 4). In older adults, neither the persistent pupal nor the peculiar adult muscles of the connate ventrites were found. Similar features were seen also in young adults of *Tenebrio obscurus*; these phenomena will be discussed further in a study of abdominal structure in adult Coleoptera, now in progress.

Table 1.

Values and Significance Levels for the Effects of Sire, Dam and Age on Offspring Number

Comparison	d.f.	χ^2_r
S ₁ , S ₂ , S ₃ , S ₄ , S ₅	4	5.34
D ₁ , D ₂ , D ₃ , D ₄ , D ₅	4	14.14***
D ₁ , D ₂ , D ₃	2	8.15**
D ₃ , D ₄ , D ₅	2	0.60
A ₁ , A ₂ , A ₃ , A ₄ , A ₅	9	183.18***
A ₆ , A ₇ , A ₈ , A ₉ , A ₁₀		
A ₂ , A ₃ , A ₄ , A ₅ , A ₆	8	17.6 **
A ₇ , A ₈ , A ₉ , A ₁₀		
Dam x Age Interaction	4	4.45
Sire x Age Interaction	4	3.68
Sire x Dam Interaction	4	10.16*

* $P(\alpha) \leq 0.05$; ** $P(\alpha) \leq 0.025$; *** $P(\alpha) \leq 0.01$

NOTE: For $N > 10$ and/or $k > 3$, χ^2_r is distributed as χ^2 . $k = \text{d.f.} + 1$,

N = number of replicates, $P(\alpha)$ = Probability of type 1 error.

S₁ = 15-22 Sire Family Size Class, S₂ = 23-30 Sire Family Size Class,

S₃ = 31-38 Sire Family Size Class, etc.

D₁ = 15-22 Dam Family Size Class, D₂ = 23-30 Dam Family Size Class,

D₃ = 31-38 Dam Family Size Class, etc.

A₁ = Parental age of 28 days at time of sample mating,

A₂ = Parental age of 33 days at time of sample mating; A₃ = Parental age of 34 days at time of sample mating, etc.

Table 2. Average Offspring Family Size for each Dam Family Size within each Sire Family Size with the Overall Means of each Class.

	DAM FAMILY SIZE					Mean Offspring Family Size for Each Sire Class
	15-22 (D ₁)	23-30 (D ₂)	31-38 (D ₃)	39-46 (D ₄)	47-54 (D ₅)	
15-22 (S ₁)	29.9	14.5	27.7	22.0	27.9	24.36
SIRE 23-30 (S ₂)	21.2	26.2	34.3	33.1	29.5	28.86
FAMILY 31-38 (S ₃)	24.0	33.5	26.7	27.5	31.2	27.92
SIZE 39-46 (S ₄)	17.5	27.2	30.3	24.7	30.1	26.04
47-54 (S ₅)	16.3	30.2	35.1	31.7	30.4	29.14
Mean Offspring Family Size for Each Dam Class	21.78	26.32	30.82	28.80	29.42	

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Table 3. Average Offspring Family Size for each Age of Parents at Time of Mating

Mean Offspring Family Size for Each Parental Age	Age of Parents at Time of Mating									
	(A ₁) 28	(A ₂) 33	(A ₃) 34	(A ₄) 35	(A ₅) 40	(A ₆) 41	(A ₇) 42	(A ₈) 47	(A ₉) 48	(A ₁₀) 49
13.96		34.64	35.20	35.36	29.96	28.56	25.80	22.24	19.80	27.88

Speaking of the ventral longitudinal muscles of the ventrites 1 & 2, it is possible that these muscles are normal and functional in teneral adults, providing the insect with some flexing capacity while the ventrites are still soft enough. As soon as they have become more or less sclerotised, which is marked with the changes in the color of the ventrites from amber to reddish-brown, all the pupal and ventral longitudinal adult muscles of the ventrites 1 & 2 may be absorbed since they are no more effective. Such muscles might be termed transient ventral longitudinal muscles.

None of the dissected specimens of ppas mutants were newly emerged adults and the ventral longitudinal muscles in any of the first 2 ventrites of ppas mutants are not similar in form to those of the corresponding segments of the newly emerged non-mutants, which rather resemble those of the ventrite 3.

I am indebted to Dr. R. A. Crowson of Zoology Department, Glasgow University, for his invaluable encouragement and assistance. My thanks are also due to Dr. A. Sokoloff and G. Hosie who supplied the material of ppas mutants and non-mutants of T. castaneum respectively.

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*Comparison of Methods of Counting Tribolium Eggs.

Introduction

Genetic experiments with Tribolium, particularly selection experiments, often involve counting large numbers of eggs. Therefore, it would be advantageous to mechanize counting. To increase the efficiency, while maintaining a high degree of accuracy, hand counting, as the standard, was compared with machine counting, using one of two automated "bacteria colony" counters, in two experiments.

Materials and Methods

Varying numbers of eggs, ranging from about 5 to about 125, were sifted from flour medium and placed at random into 24 vials. The eggs in each vial were counted eight times, twice (determinations 1 and 2) by each of two operators (operators 1 (O_1) and 2 (O_2)) using each of two methods (method 1 (M_1), hand; and method 2 (M_2), machine).

Experiment 1:

For each hand count, the contents (eggs and residual flour particles) of each vial were poured into a linear egg counting stage (Muir and Grossman, 1973); the eggs were counted under a microscope, and the contents returned to the same vial. (With some experience, operators can usually distinguish eggs from flour particles.) The machine counter was a BioTran Automated Colony Counter,¹ Model CIII, including a CV-10 viewing monitor. For this method, the operator placed the contents of each vial into a plastic petri dish and placed it on the Counter. He then made two determinations before the contents were returned to the vial. After each determination, the petri dish was shaken slightly, and again placed on the Counter for another reading. By using the viewing monitor, the operator took care that the eggs were not clustered at the perimeter of the dish and were not touching each other. The specification of the machine is that it will count all particles as small as 0.2 mm in diameter.

Experiment 2:

The second experiment was similar to the first, with about the same range in numbers of eggs per vial. Counting by hand involved two steps; first, counting the eggs in each vial, using the linear egg counting stage under a microscope, then counting the number of flour particles for a count of total (eggs + flour) particles. The flour particles that were counted were those judged by the operator to be of sufficient size to be counted also by the machine. The machine counter was the Petri-Scan Bacteria Colony Counter.² Its specification is that all particles over 0.25 mm in diameter will be counted.

Statistical Analysis:

Because eggs or flour particles might have been lost while transferring them to and from a vial, a linear factor, called sequence, was included in the analysis. The sequence of the eight counts per vial differed in the two experiments, so they were analyzed separately.

Square-roots of egg and total particle numbers were analyzed because the square-root transformation tended to make the error variances homogeneous according to Bartlett's test (Ostle, 1963).

Results and Discussion

Experiment 1:

The BioTran counted significantly ($P < .05$) more eggs (or total particles) than the hand counting method (Tables 1 and 2). However, this difference was not consistent over vials, as indicated by the significant ($P < .01$) vial by method (V x M) interaction.

¹BioTran, Model CIII is manufactured by New Brunswick Scientific Co., Inc., 1130 Somerset Street, New Brunswick, N.J. 08903.

²Petri-Scan is manufactured by American Instrument Company, Box 477, Barrington, Ill. 60010.

To study whether the counting methods have a constant difference over the range of numbers of eggs counted, a linear regression analysis of machine counts on hand counts was performed on the transformed data. The correlation between methods of counting was .99, and the linear regression coefficient of the transformed data for machine count on hand count was $.953 \pm .033$, not significantly ($P > .05$) different from 1.00 (Figure 1). This indicates that for every additional 100 eggs counted by hand, this machine counted about 91 eggs. The machine count was higher than the hand count up to about 80-85 eggs, after which the two methods counted about the same number of eggs.

Variation between determinations by operator 1 using the machine (.0335) was not significantly different from that between determinations by operator 2 using the machine (.0473) (Table 2), and there was no significant interaction between methods and operators. Thus the machine may be used by either operator with the same accuracy and also with the same precision.

Experiment 2:

The Petri-Scan counted significantly ($P < .01$) fewer total particles than the hand counting method (Tables 1 and 2). This difference was not consistent over vials as evidenced by the significant ($P < .01$) interaction of vials (quantity of eggs or particles) with counting method.

Linear regression analyses of machine counts on hand counts were performed on the mean $\sqrt{\text{eggs}}$ (Figure 2) and mean $\sqrt{\text{total particles}}$ (Figure 3). In both analyses the correlation between methods of counting was about .96. The linear regression coefficients were $.783 \pm .043$, and $.733 \pm .045$, respectively. This indicates that for every additional 100 eggs or total particles counted by hand, the Petri-Scan counted, respectively, about 61 or about 54 total particles. Both regression coefficients were significantly ($P < .01$) different from 1.00.

There appeared to be a significant difference ($P < .01$) between operators in the number of total particles counted; operator 1 counted an average of about 35 particles per vial, while operator 2 counted an average of only about 33.5 particles per vial. This difference may reflect the variation in judgement of the operators as to what sized particle was to be counted by the hand method.

Table 1. Least-squares means of square-root of egg and total particle counts per vial for each method by experiments.

Vial No.	Experiment 1				Experiment 2				
	√Eggs / Total Particles		Difference (1 - 2)		√Eggs / Total Particles		Difference (1 - 2)		
	Method 1 Hand	Method 2 BioTran	Method 1 Hand	Method 2 Petri-Scan	Method 1 Hand	Method 2 Petri-Scan	Method 1 Hand	Method 2 Petri-Scan	
1	7.0521	6.8361	.2160	7.6807	6.9616	.7192	8.3034	6.9616	1.3418
2	4.1819	4.2709	-.0890	4.3582	4.0277	.3304	4.7395	4.0277	.7118
3	6.5741	6.5137	.0554	6.8368	5.9213	.9156	7.4135	5.9213	1.4922
4	5.1228	5.3136	-.1908	9.7045	7.2787	2.4258	10.1681	7.2787	2.8894
5	8.9719	8.9405	.0314	7.2786	5.5407	1.7380	7.6801	5.5407	2.1394
6	7.9213	7.9989	-.0776	3.7393	3.1831	.5562	3.8707	3.1831	.6876
7	4.1828	4.3008	-.1180	12.6277	10.2648	2.3630	13.7588	10.2648	3.4940
8	8.8728	9.0662	-.1934	6.8163	5.6507	1.1656	7.7633	5.6507	2.1126
9	6.5763	6.7071	-.1308	6.0142	5.2903	.7240	6.8373	5.2903	1.5470
10	5.9154	5.9368	-.0214	6.7820	5.0979	1.6840	7.4485	5.0979	2.3506
11	4.0615	3.9673	.0942	5.3614	4.0423	1.3192	6.0821	4.0423	2.0398
12	2.4494	2.4986	-.0492	6.1657	4.8622	1.3036	7.3404	4.8622	2.4782
13	9.0666	8.8972	.1694	5.9776	4.7349	1.2426	6.4761	4.7349	1.7412
14	6.1643	6.2641	-.0998	2.5968	1.9920	.6048	3.2394	1.9920	1.2474
15	6.4222	6.5760	-.1538	5.3538	4.9442	.4096	5.9694	4.9442	1.0252
16	5.8722	6.1400	-.2678	3.1213	2.4062	.7156	3.7378	2.4062	1.3316
17	5.3847	5.2667	.1180	6.8344	5.6052	1.2292	7.6440	5.6052	2.0388
18	4.3298	4.5530	-.2232	5.9146	4.7168	1.1978	6.5744	4.7168	1.8576
19	9.9740	9.6414	.3326	4.5774	4.3896	.1878	5.1328	4.3896	.7432
20	4.4700	5.8914	-1.4214	6.0205	5.2191	.8014	6.4417	5.2191	1.2226
21	10.6846	10.7684	-.0838	4.9238	3.6582	1.2656	5.4420	3.6582	1.5833
22	7.5816	3.4842	-.9026	6.2191	6.0791	.1400	6.6171	6.0791	.5380
23	5.4772	5.8716	-.3944	5.4313	4.4929	.9384	5.5899	4.4929	1.0970
24	2.1769	2.6457	-.4688	5.2883	4.7757	.5122	5.6521	4.7757	.8764
X	6.2286	6.3898	-.1612	6.0677	5.0473	1.0204	6.6551	5.0473	1.6078
ΣX	.45704	.44131	.07474	.41977	.33926	.12343	.44523	.33926	.15120

Table 2. Analysis of variance of square-root of egg and total particle counts for each experiment.

<u>Experiment 1</u>				<u>Experiment 2</u>			
Mean Squares				Mean Squares			
<u>Source</u>	<u>d.f.</u>	$\sqrt{\text{Eggs}}$	$\sqrt{\text{Total Particles}}$	<u>Source</u>	<u>d.f.</u>	$\sqrt{\text{Eggs}}$	$\sqrt{\text{Total Particles}}$
Sequence (S)	1	.2755**		Sequence (S)	1	7.9060**	10.6200**
Method (M)	1	1.2462*		Method (M)	1	49.9725**	124.0922**
Operator (O)	1	.0352		Operator (O)	1	.2066	.8021**
M x O	1	.0340		M x O	1	.0004	.2125
O x Det (D)/M ₁	1	.0077		Det (D) x M	1	3.6746**	3.1012**
D/(O ₁ x M ₂)	1	.0335		D x O	1	2.1438**	2.2239**
D/(O ₂ x M ₂)	1	.0473		D x M x O	1	.0497	.0380
Vials (V)	23	38.4810**		Vials (V)	23	27.2340**	28.9819**
V x S	23	.0284		V x S	23	.3295**	.3421**
V x M	23	.2681**		V x M	23	.7312**	1.0974**
V x O	23	.0232		V x O	23	.0566	.0624
V x M x O	23	.0246		V x M x O	23	.0430	.0619
V x O x D/M ₁	23	.0081		V x D x M	23	.3276*	.3274*
V x D/(O ₁ x M ₂)	23	.0242		V x D x O	23	.0496	.0610
V x D/(O ₂ x M ₂)	23	.0229		V x D x M x O	23	.0879	.1032
Total	191			Total	191		

** Significant at 1% level of probability.

* Significant at 5% level of probability.

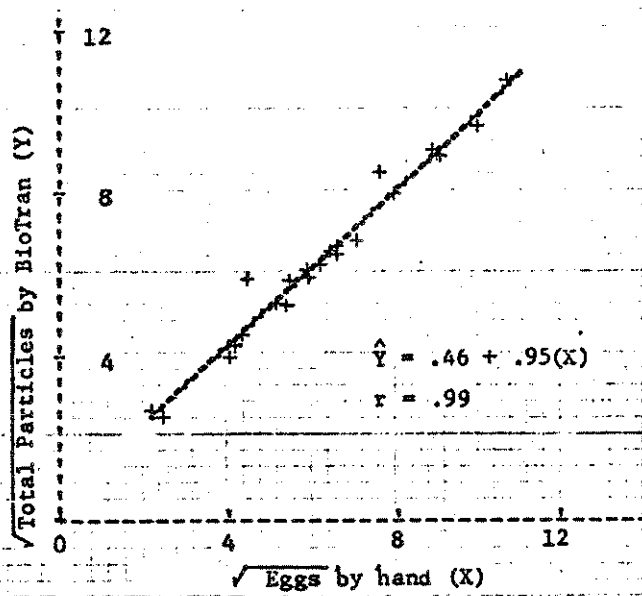


Figure 1. Regression of $\sqrt{\text{total particles}}$ counted by the BioTran on $\sqrt{\text{eggs}}$ counted by hand.

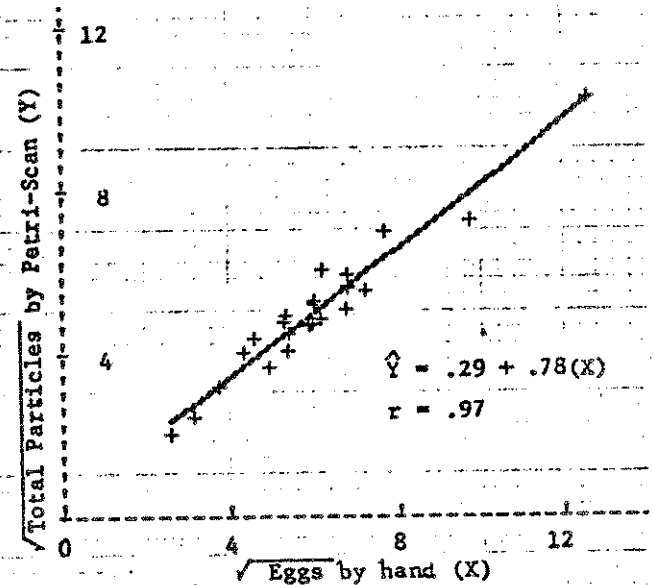


Figure 2. Regression of $\sqrt{\text{total particles}}$ counted by the Petri-Scan on $\sqrt{\text{eggs}}$ counted by hand.

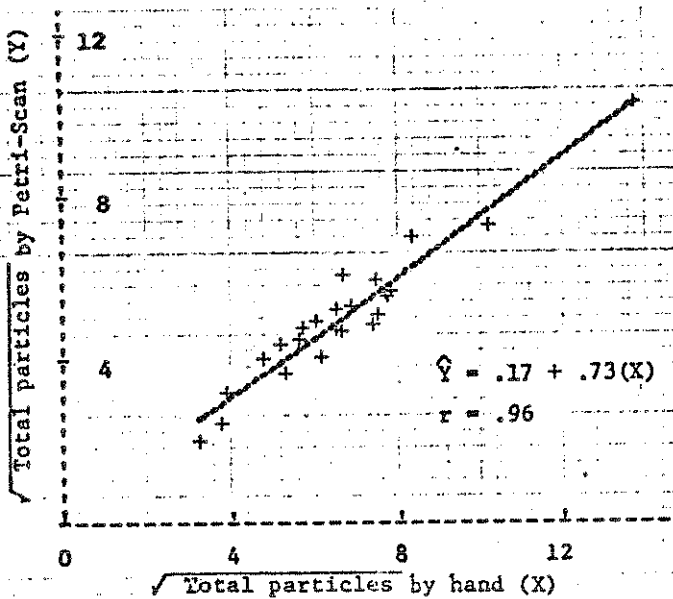


Figure 3. Regression of $\sqrt{\text{total particles}}$ counted by the Petri-Scan on $\sqrt{\text{total particles}}$ counted by hand.

Conclusions

Counts using the BioTran machine agree with hand counts better than do counts using the Petri-Scan machine. Over a range of eggs, numbering from 5 to 125, the BioTran counted 91% of the eggs counted by the hand method. Operators probably will not differ in their counts, no matter which method they use, if they are experienced in distinguishing eggs from flour particles.

Although there may be some bias in machine counting, using the machine increases efficiency by reducing operator fatigue, allowing larger quantities of eggs to be counted, and allowing a change in operators without significantly affecting the results.

Acknowledgments

The BioTran Automated Colony Counter was provided through the courtesy of Mr. Ted Shields, Sales Engineer, New Brunswick Scientific Co., Inc. The Petri-Scan was provided through the courtesy of Mr. William T. Robinson, Sales Engineer, American Instrument Company. We thank Dr. H. W. Norton for his assistance in the analysis of the data, and Clyde Anderson for his technical assistance.

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*The Influence of the ppas Mutation on the Abdominal Musculature of Tribolium castaneum.

Sokoloff (The Genetics of Tribolium and Related species, Academic Press, New York; 1966) has pointed out that ppas mutants of Tribolium castaneum have homeotic features in that the subsequent ventrites tend to become similar to the 1st ventrite, having a middle projection corresponding to the intercoxal process of the 1st ventrite and also having an anterior articulating membrane. These features suggested that some kind of abnormalities might be expected in the ventral musculature of the ppas mutants.

Through the kindness of Dr. A. Sokoloff I received some adult ppas mutants and through Mr. G. Hosie of the Infestation Department, Ministry of Agriculture Fisheries, Glasgow, some non-mutant specimens of Tribolium castaneum; I dissected nearly 20 random specimens (of both sexes and different ages) of ppas, and some non-mutants for comparison.

As seen in Fig. 1, the non-mutant ones have the first 3 ventrites connate and first 2 in fully mature specimens lack the ventral longitudinal muscles which are present in the ventrites 3, 4, 5; these muscles in a segment actually move the segment one behind. The ventral longitudinal muscles of nearly all the dissected specimens of the ppas mutants differed more or less from those of the normal type and the homeotic modification of the ventrite 2 was well expressed. In one of them (Fig. 3) the ventrites 2-4 showed the modification in decreasing degree posteriorly. As far as the musculature is concerned, this specimen in comparison with the other ppas mutants has the extremest muscular abnormality, with median ventral longitudinal muscles in the ventrites 1 & 2, similar to those normally in front of the ventrite 1, while the ventrite 3 has lost them, as is the normal condition of the ventrites 1 & 2; this specimen also lost the ventral lateral oblique muscles altogether. Many intermediate types were found between this extremest mutant and non-mutant type. In these intermediate types, the abnormalities involved are modifications, often asymmetrical, in the orientation and position of the ventral muscles. The modifications in their orientation are changes in the direction and location of the muscles so that the normally longitudinal muscles have become oblique or vice versa and the normal positions of the muscles have also been displaced. The structural modifications are the coalescence and lengthening of the muscle bundles e.g. (Fig. 2). From these observations it appears that in ppas mutant of this species, the ventral musculature shows homeotic tendencies towards the abdominal base. This tendency is more evident in the extremest ppas mutant (Fig. 3) than the intermediate mutants.

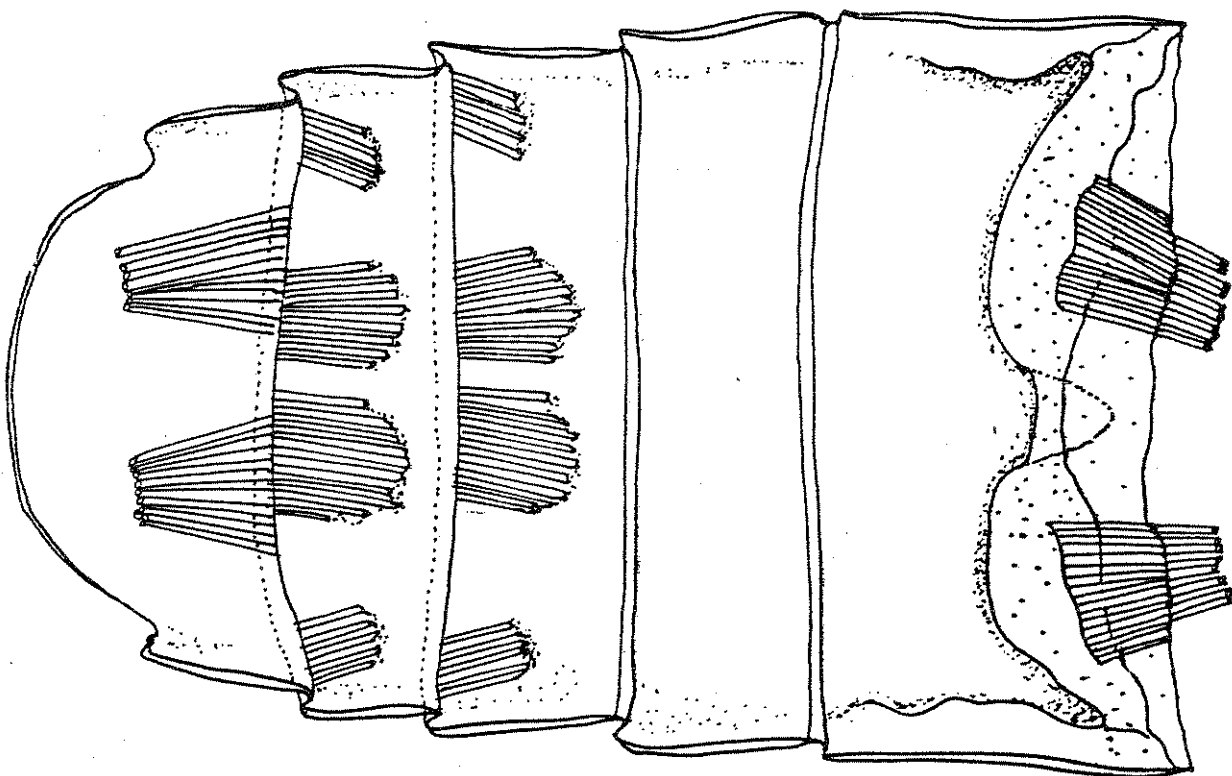


Fig. 1. The ventral abdominal musculature of a fully mature non-mutant Tribolium castaneum.

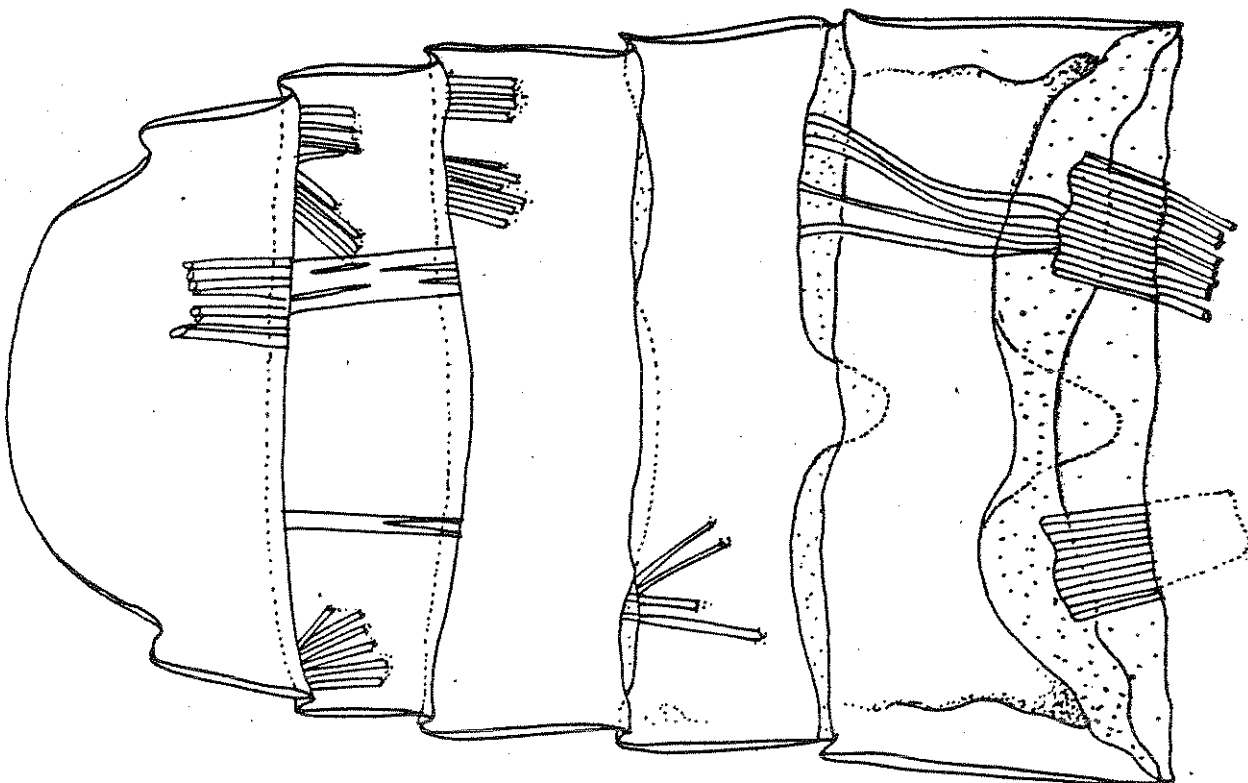


Fig. 2. The ventral abdominal musculature of a fully mature intermediate ppas mutant of T. castaneum.

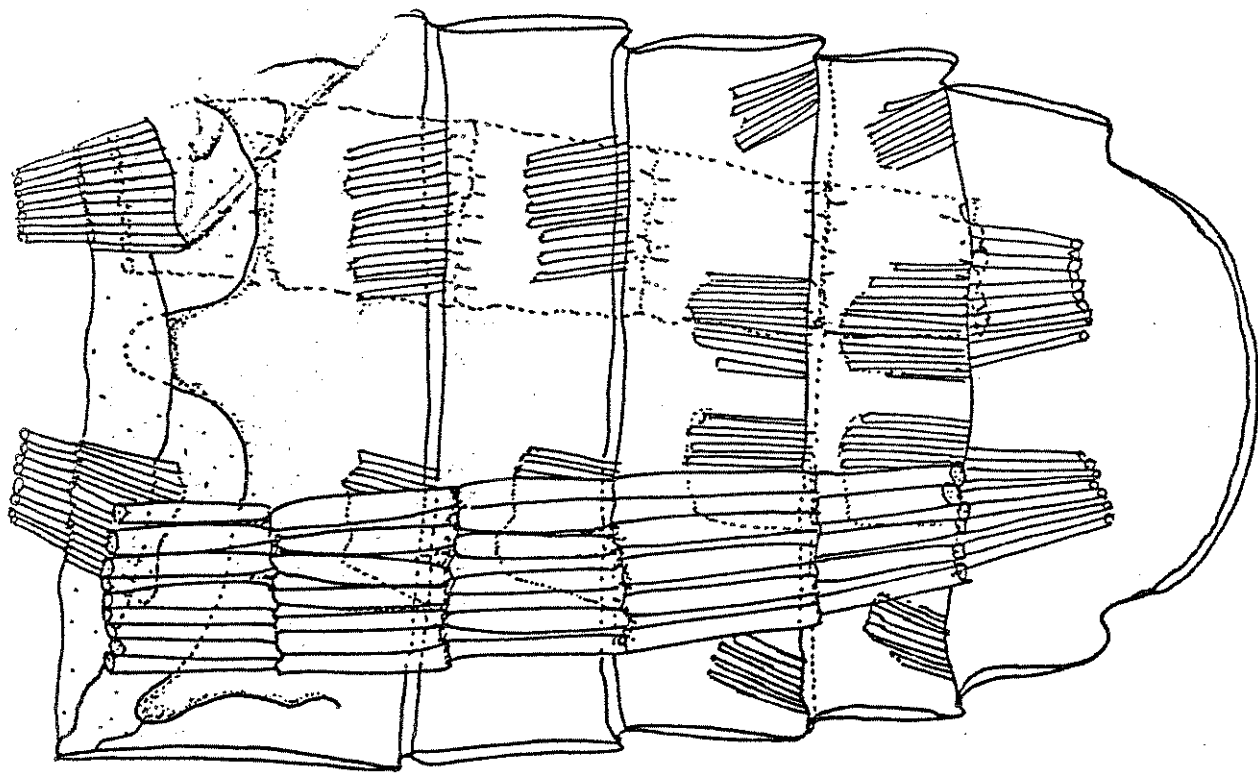


Fig. 4. The ventral abdominal musculature of a newly emerged non-mutant adult T. castaneum; The thick loosely overlying persistent pupal muscles are omitted on the right hand side, notice the transient ventral muscles in the ventrites 1 & 2.

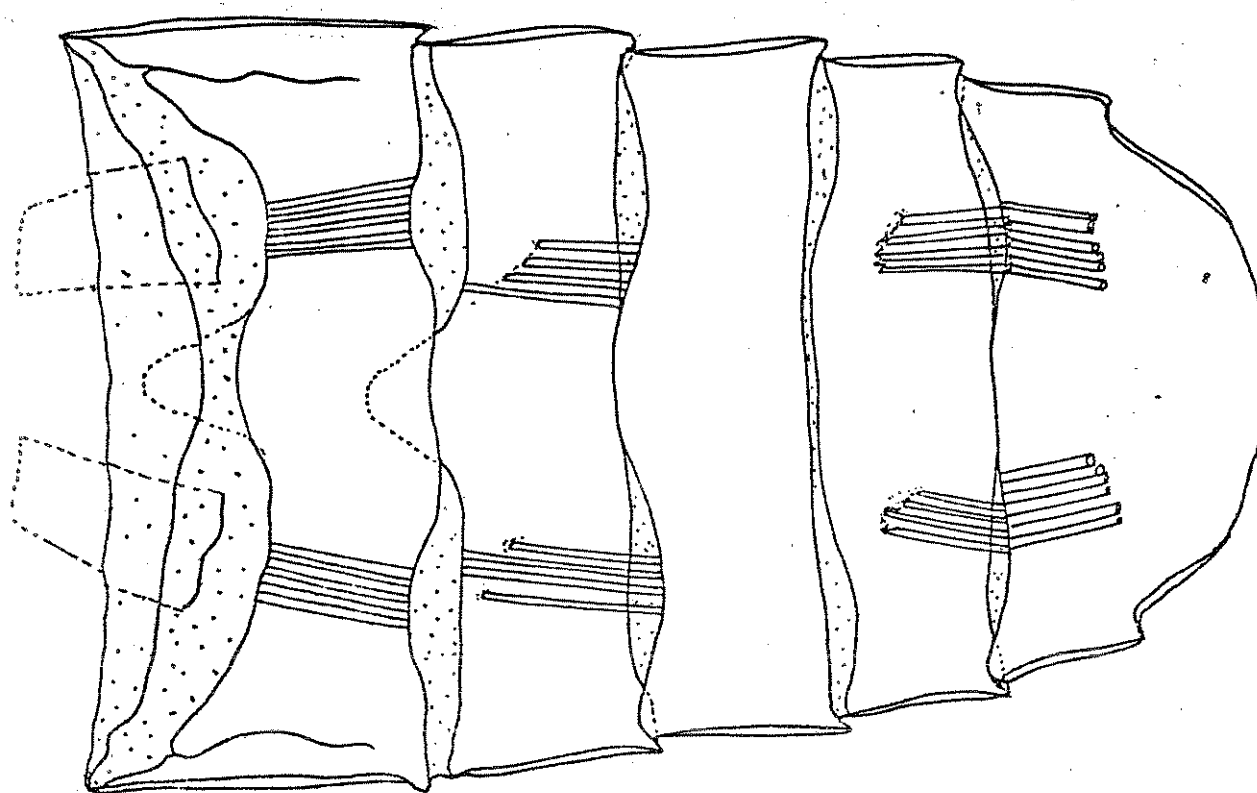


Fig. 3. The ventral abdominal musculature of an extreme ppas mutant of T. castaneum.

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*Dietary and Density Dependant Factors in the Induction of Population Autocide of Flour Beetles, *Tribolium confusum*.

The response of the adult *Tribolium confusum* Jacquelyn duVal to stress imposed by malathion and crowding was investigated. Stress imposed by malathion concentrations of 1.0 ppm or greater with population densities of 75 or more adult beetles per 3 grams of flour was adequate to initiate the production of gas with resulting death of all beetles. A color change in the flour (white to pink) was a secondary effect resulting from the presence of quinone gas.

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*Selection for changes in developmental period in *Tribolium castaneum* (Chicago Black strain).

In conjunction with other ongoing research it became necessary to investigate the amount of genetic variation in egg to adult developmental period maintained in the Chicago Black strain of *Tribolium castaneum*. This paper will report on the results of five generations of artificial selection for both fast and slow developmental period.

Materials and Methods

The Chicago black (bb) beetles were obtained from stocks maintained in our laboratory. Throughout the experiment they were reared on sifted whole wheat flour mixed with 5% brewers yeast at 25°C and 70% R. H. Four egg farms were set up and 400 one-day old eggs from each farm were used to set up each of the four lines that made up generation 0. The cultures were always raised in 40 grams of flour in half-pint Mason jars. After 22 days the cultures were sifted and all pupae removed to separate vials corresponding to their respective cultures. If any, larvae remained, the cultures were sifted again every four days until all had pupated. The vials containing pupae were checked daily and all emerging adults were removed to holding vials. At the end of generation 0 two of the lines were each subdivided into fast and slow lines by taking the 10% fastest and slowest from each line and allowing them to mate with their cohorts for a day in 40 grams of fresh flour. The other two lines were designated controls. All the emerged adults from the particular control line were mixed and 10% taken at random to form the parents of the next generation. Thus generation 1 consisted of two fast lines, two slow lines, and two control lines.

Table 1. - Egg to adult developmental time -- mean and variance and sample sizes.

Generation	<u>Fast Lines</u>		<u>Slow Lines</u>		<u>Control Lines</u>	
	1	2	1	2	1	2
0. F	25.741	25.687	25.741	25.687	25.740	25.646
6 ²	1.446	0.839	1.446	0.839	1.370	0.956
n	352	367	352	367	319	356
1. F	25.803	25.446	26.304	26.720	25.978	26.152
6 ²	2.385	0.568	1.474	3.888	0.451	1.193
n	213	195	253	300	178	223
2. F	26.073	25.301	26.784	27.405	26.007	26.100
6 ²	2.70	0.527	2.204	4.676	0.453	0.715
n	96	153	139	159	140	110
3. F	25.387	25.457	26.939	29.031	26.329	26.584
6 ²	0.588	0.606	2.548	4.367	0.939	1.787
n	150	164	164	97	213	154
4. F	24.984	25.380	28.321	31.312	25.386	26.201
6 ²	0.842	2.649	2.466	10.450	0.754	2.424
n	185	108	131	138	44	169
5. F	25.005	25.056	28.738		25.463	26.258
6 ²	0.816	2.179	3.235		1.300	2.894
n	186	144	103		123	155

It should be noted that all new generations were started at least 5 days after the last adult emerged to insure that the youngest adults were mature. This regime was continued for the rest of the experiment except that due to a population decline the parents of subsequent generations consisted of the 20 fastest, slowest, or randomly selected beetles. This kept population size at about 150. It was hoped that maintaining the population so low eliminated any changes in developmental period due to density effects.

Results and discussion

A summary of the data is given in table 1. Note that only four generations were run in slow line number 2. As can be seen from generation 0 the bb strain has a developmental period averaging a little less than 26 days. The distribution is faintly bimodal; a major mode occurring on day 26 and a very much smaller mode 2-3 days later. This is very similar to the distribution for this strain found by Kence (1973). Previous selection experiments on developmental period in T. castaneum have shown a relatively rapid response to selection for slower development and a less dramatic response for faster development, Dawson (1965, 1966). Wool (1969), on the other hand, found that he could select for faster developmental time in bb rather easily but found no response to selection for slower developmental period.

In the work reported on in this paper the response to selection was highly asymmetric. The response to selection for slow development was apparent after one generation and by the end of the experiment the mean had shifted an average of nearly four days. The response to selection for fast development was neither as rapid nor as clear cut. There was a trend toward faster development but this was not found statistically significant. A single classification analysis of variance was performed by grouping all mean developmental times into either control, fast, or slow. The results were significant ($F_s = 22.743$, $P < .001$), showing that selection had differentiated the groups. Further analysis revealed no significant difference in Fast vs. Control ($F_s = 2.388$, $P < .10$) and a large difference in Slow vs. Control ($F_s = 28.946$, $P < .001$).

The individual population variances (see Table 1) were similarly grouped and Druskal-Wallis tests performed separately on each selected group vs. the control group. As was the case with the means there was no difference between the Fast and the Control (adj H = .0309, $P < .5$) and a highly significant difference between the Slow and the Control (adj H = 12.444, $P < .005$). Thus it seems that selection for slower developmental period acts not only by shifting the mean of the population but by increasing its variance.

The response to slow selection can be explained by Kence's (1973) work in which he hypothesizes a polymorphism in number of larval instars which results in the bimodality in developmental period. He assumes that in bb the morph for higher number of instars is maintained at low levels. The results reported here tend to support these assumptions.

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*Multifactor Ecological Effects on the Fitness of Tribolium Populations.

This research was done to gain information on the effects of nutrients, radiation, and competition studied singly or in combinations on the productivity of Tribolium castaneum and Tribolium confusum populations. The response of Tribolium populations to stress in a laboratory situation may supply explanations for their responses in nature.

The results indicated a progressive reduction in productivity for the P₁ adults of T. castaneum and T. confusum as the gamma (⁶⁰Co) exposures increased. The sterilizing dose for each species was between 3 and 6 kR at 36 ± 2 C and 56 ± 10% relative humidity. With stress from radiation or competition T. castaneum consistently was reproductivity fitter than T. confusum; whereas, the opposite was true without the stress.

A progressive reduction in productivity for both species was noted as the percent of cottonseed flour increased. The life cycle from egg to adult was apparently prolonged on cottonseed flour. The flour beetles were not consuming the cottonseed flour because of palatability and/or consistency. Dissection of the dead adults revealed empty and shriveled digestive systems.

As the gamma radiation dose increased and the proportion of cottonseed flour increased, the numbers of F₁ progeny of T. castaneum and T. confusum decreased. The radiation effects obscured the flour effects in modifying the fitness of Tribolium, although the flour proportions did contribute to the overall productivity decreases.

The 9 kR exposure was the LD₁₀₀ for parental adults of T. castaneum on 100% cottonseed flour and T. confusum on all flour proportions. An increase in F₁ mortality was observed with increased population density.

Fitness parameters, such as productivity and lethality, for single- and mixed-species populations of T. castaneum and T. confusum varied inversely to the gamma radiation exposure and/or the proportion of cottonseed flour.

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*Larval length and width measurements as an index of growth in Tribolium castaneum.

Many indices have been devised to quantify the growth process of Tribolium. Measurements include the duration of metamorphosis (Park, et. al, 1961), adult, pupal and larval weight (Bell, 1969) and estimation of biomass (King and Dawson, 1972). In our laboratory we have quantified growth by length and width measurements, similar to Brindley (1930) with T. confusum. There are several reasons for choosing these measurements:

(1) Ocular micrometer readings are easy to obtain and record and are only slightly more time consuming than sexing pupae, (2) It is an inexpensive system and maintenance is minimal as opposed to weight measurements which require the maintenance of sensitive balances, (3) It offers the opportunity of quantitatively assessing on an individual basis, and (4) most importantly, the system provides an accurate means of describing growth during the very early stages of Tribolium development. The following study is presented as an example of the kind of information that can be obtained via these measurements and in fact suggests that only one measurement is needed (length or width) to monitor Tribolium development.

The genetic material used during this study were two populations of T. castaneum, the unsaturated fatty acid sensitive population (cos/cos) described by Costantino and Rowe (1972) and the Purdue Foundation population (+/+). Eggs from both the sensitive and normal populations were collected over a 24 hour period and distributed in groups of 25 to creamers containing diet 0 (percentage composition: wheat flour 95, and brewers yeast 5), diet 3 (percentage composition: wheat flour 92, brewers yeast 5, and corn oil 3), and diet 5 (percentage composition: wheat flour 90, brewers yeast 5, and corn oil 5) and maintained at $33 \pm 1^\circ\text{C}$ in low ($35 \pm 5\%$) and high ($77 \pm 3\%$) relative humidity. Seventy-two hours after egg collection, larvae from all environmental combinations were counted and eight randomly chosen individuals were measured (Scully, 1972). In order to assure accurate measurement, larvae were placed in the freezer compartment of a refrigerator for approximately two minutes prior to measurement. This inhibited movement of the larvae for several minutes and did not affect viability. In all cases, larvae were measured with an ocular micrometer to the nearest .005 of a millimeter; the total magnification was x .8. (Actual number of millimeters = recorded number of millimeters divided by .8). The results of each day represent an independent sample of individuals.

Tables 1a and b contain the mean and standard error of the mean for measurements of larval length and width as recorded for low relative humidity. The length parameter exhibits a greater range of values during early development than does width. This difference in phenotypic variance suggests that larval length may be a more informative measure of the developmental growth of a population than is width.

The analysis of variance for the first nine days of these data is presented in Tables 2a and b. The analysis performed on length measurements agrees with that performed on width. All components are statistically meaningful. Because growth occurred over the observation period the differences due to Days were significant. The concentration of corn oil significantly affected larval growth as did the genotype of the population. The Days by Oil interaction is an indication that the response to the concentration of corn oil changes with time while the Days by Genotype interaction reflects how differently both populations grow with respect to time. The Oil by Genotype interaction points out the different phenotypic responses of both genotypes.

Table 1 (A) Mean length of larvae and standard error of the mean of populations of T. castaneum grown at low relative humidity on different concentrations of corn oil media. Each mean represents an independent sample of 8 individuals. Given in millimeters (total magnification x .8; observed number of millimeters = actual number of millimeters / .8). (B) Mean width of larvae and standard error of the mean of the same populations cultured under the same conditions.

(A)	Days After Hatching	Sensitive Diet			Normal Diet		
		0	5%	3%	0	5%	3%
1	0.775±.016	0.787±.016	0.756±.011	0.781±.019	0.800±0	0.781±.009	
2	0.800±.021	0.775±.027	0.731±.016	0.835±.024	0.819±.033	0.762±.016	
3	0.940±.034	0.945±.043	0.831±.027	1.045±.028	1.025±.031	0.955±.050	
4	1.110±.012	1.010±.033	0.935±.027	1.035±.024	1.100±.027	1.055±.034	
5	1.294±.072	1.006±.013	1.000±.008	1.417±.046	1.400±.041	1.256±.065	
6	1.500±.026	1.255±.047	0.985±.036	1.490±.037	1.375±.082	1.370±.042	
7	1.695±.075	1.210±.018	1.085±.051	1.860±.095	1.935±.069	1.910±.073	
8	1.905±.084	1.400±.104	1.110±.050	2.145±.058	2.150±.128	1.875±.067	
9	2.760±.118	1.430±.093	1.314±.114	3.060±.189	3.065±.066	3.025±.155	
10	3.280±.170	1.806±.148	1.336±.066	3.600±.192	3.435±.135	2.970±.110	

(B)	Days After Hatching	Sensitive Diet			Normal Diet		
		0	5%	3%	0	5%	3%
1	.131±.013	.137±.012	.136±.011	.150±.009	.131±.013	.131±.009	
2	.169±.013	.150±.013	.144±.006	.169±.009	.162±.008	.156±.011	
3	.180±.008	.160±.007	.162±.012	.185±.008	.175±.008	.185±.011	
4	.200±.000	.165±.100	.170±.008	.190±.007	.205±.012	.190±.012	
5	.200±.012	.167±.012	.172±.009	.217±.008	.244±.013	.189±.011	
6	.230±.008	.200±.007	.170±.008	.225±.017	.240±.019	.240±.010	
7	.285±.013	.185±.011	.190±.012	.270±.011	.250±.013	.280±.008	
8	.300±.010	.210±.007	.175±.015	.325±.015	.280±.022	.315±.017	
9	.450±.027	.240±.010	.221±.018	.510±.022	.495±.019	.475±.032	
10	.495±.017	.275±.013	.221±.010	.540±.025	.500±.019	.455±.020	

Table 2 A) Analysis of variance of larval length of the sensitive and normal genotypes cultured in low relative humidity.

Source of Variation	Degrees of Freedom	Mean Squares	F-ratio
Days (D)	8	22.622	378.0*
Oil (O)	2	6.234	104.1*
Genotype (G)	1	10.860	346.9*
D X O	16	0.878	14.7*
D X G	8	2.322	38.8*
O X G	2	3.129	52.3*
D X O X G	16	0.517	8.6*
Residual	378	0.060	
Total	431		

*Significant at the 0.05 level of probability.

B) Analysis of variance of larval width of the sensitive and normal genotypes cultured in low relative humidity.

Source of Variation	Degrees of Freedom	Mean Squares	F-ratio
Days (D)	8	0.445	263.5*
Oil (O)	2	0.132	78.4*
Genotype (G)	1	0.450	266.7*
D X O	16	0.017	10.2*
D X G	8	0.052	30.8*
O X G	2	0.086	50.6*
D X O X G	16	0.010	6.0*
Residual	378	0.002	
Total	431		

*Significant at the 0.05 level of probability.

Table 3. Correlation coefficients of mean larval length and width.

<u>Genotype</u>	<u>Diet</u>	<u>Low Relative Humidity</u>			<u>High Relative Humidity</u>		
		0	3	5	0	3	5
(cos/cos)		.992	.981	.962	.989	.670	.989
(+ / +)		.989	.973	.991	.994	.991	.980

The correlation coefficients of mean larval length with larval width (Table 3) in all cases but one are greater than .95 indicating a strong linear relationship between these measurements. Predictions of larval width based on measured larval length should be good and the volume of a particular organism easily estimated. The exception in this study is very likely attributable to sampling error.

In summary, it appears as if larval length and width measurements provide easily acquirable yet accurate information on the growth of Tribolium larvae. Analysis based on length yields similar information as analysis based on width. The linear correlation coefficients of mean length and mean width are on the order of .95 justifying the use of only one of the parameters as a measure of growth. It is suggested that larval length be measured as this parameter is the most sensitive to changes in time and to environmental differences.

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A strain of small *Oryzaephilus mercator* Fauvel from Thailand.

Introduction

The occurrence of a strain of small *Oryzaephilus surinamensis* (L.) in the Far East and its biology has been described by Aitken (1966). Results from crosses between the small and a normal sized strain showed that the inheritance of size was under polyfactorial control and that size was inherited through both parents, the female parent having a greater influence. In 1970 some small *Oryzaephilus mercator* Fauvel were collected from a shipment of polished rice imported from Thailand to Holland.

This strain was cultured on rolled oats at 25°C and 70% R.H. since its collection.

The inheritance of size of this strain has been studied.

Experiments and results

End 1972 pronota were measured of 50 males and 50 females of the strain of small beetles (called S) and of a laboratory strain of normal size (called L) according to the method described by Aitken (1966).

Strain L was cultured for a large number of generations on rolled oats at 25°C and 70% R.H.

Crosses were made between strains S and L and a number of back crosses were carried out. Of the progeny of the various crosses the pronota of 50 females were measured. The results are shown in Table I. They are similar to those found by Aitken (1966). Differences in size between both strains are obvious, also when they are bred on a nutrient-rich food rolled oats. The size of the F_1 of interstrainal crosses lays between that of the parental strains.

In both sexes of the F_1 there is a bias towards the large condition which is more pronounced when the female parent is of the large strain. The progeny of the F_1 inter se cross is smaller than of the F_1 hybrids but a larger size is reached when the female grandparent was large.

The influence of the female parent is also reflected in the progeny of the various back crosses.

Discussion

Some preliminary experiments showed that the developmental periods of the S and L strain with rolled oats as food at 25°C and 30°C and 70% R. H. were the same. The fertility and fecundity of the hybrids also does not seem to be impaired. The question arises therefore how a small strain originates and how it maintains its identity.

It is supposed that the small strain in the long run is better adapted to a poor diet than strains of normal size and so a small strain can exist.

On a more nutritious food like rolled oats however, it is likely that a small strain will be gradually swamped by a strain of normal size. To prove this assumption experiments have to be carried out on the productivity and competition of both strains on deficient diets.

Table I

Length of pronota in strains S and L, inter strain crosses and back crosses bred on rolled oats at 25°C and 70% R.H.

Strain or cross	mean length mm.		standard deviation	
	♂♂	♀♀	♂♂	♀♀
L	0.73	0.71	0.041	0.032
L x S	0.69	0.68	0.030	0.022
L x S inter se	0.66	0.66	0.043	0.027
L x (L x S)	0.71	0.69	0.034	0.029
S x (L x S)	0.64	0.64	0.026	0.030
(L x S) x L	0.72	0.71	0.031	0.028
(L x S) x S	0.65	0.63	0.036	0.030
S	0.59	0.59	0.019	0.028
S x L	0.67	0.67	0.024	0.026
S x L inter se	0.64	0.64	0.033	0.031
L x (S x L)	0.68	0.67	0.030	0.027
S x (S x L)	0.63	0.62	0.033	0.035
(S x L) x L	0.70	0.69	0.029	0.028
(S x L) x S	0.64	0.64	0.039	0.027

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*Selection against homzygotes in a variable environment. A preliminary experiment with *Tribolium*.

Introduction

Some data from Nature and a few experimental results reported so far indicate that even changes of very short duration relative to generation time may produce important genetic changes in populations. For example, Band and Ives (1961, and later publications summarized by Band 1972), reported a positive correlation between the range of diurnal temperature fluctuation and the frequency of lethal and semi-lethal chromosomes in a natural population of *Drosophila melanogaster*. Beardmore (1961) concluded from his study of bristle number in *Drosophila* populations that a diurnal change in temperature was highly effective in maintaining genetic variance. Beardmore and Levine (1963) reported that larval viability of a population after 19 generations in a diurnally-fluctuating environment was higher than three other populations maintained in 3 different but constant conditions over the same period. Long (1970) compared populations after many generations in one constant and three fluctuating environments with a period of 1, 32 and 96 days respectively. Using a multiple index of fitness, he was able to show that the population from the diurnally-changing conditions had the highest fitness.

Increased fitness is often related to an increase in the proportion of heterozygotes, which are also more homeostatic and may survive in a wider range of environmental conditions. On the other hand, many natural populations are inbred - and inbreeding tends to increase the proportion of homozygotes in the population.

It, therefore, seemed of interest to expose populations, for a length of time, to variable environment under very close inbreeding, and compare their fitness with populations of the same initial genetic constitution under the same mating system, but maintained in a constant environment.

Materials and Methods

A wild type strain of *Tribolium castaneum*, originally obtained from Dr. R. R. Sokal's laboratory in Stony Brook, L. I., N. Y., was used in this experiment. Offspring of two randomly-chosen pairs of adults (selected from the stock in the pupal stage) were pair-mated in vials containing 1 g. of medium (flour and brewer's yeast in the ratio 100:5). Each pair founded one replicate line for the experiment, by ovipositing for 7 days.

Ten of these lines were maintained in a standard (constant) environment (30°C, 70% RH) denoted C, and 30 lines in a variable environment, denoted V. These were exposed to a series of treatments.

The experiment lasted 6 generations. Every generation, beginning when first detected, offspring pupae were recovered from the vials twice weekly, separated by sex, and held separate until adults emerged.

To minimize the loss of experimental lines from occasional sterility of parent adults, 3 replicate pairs chosen at random, were set up for 7 days oviposition each line if enough offspring were available. From these, 1 fertile pair was chosen to continue the line.

Six treatments were decided upon arbitrarily ("no treatment" being one of the list, Table 1) and were given to the V lines in generations 1-5 twice every week, beginning with the termination of oviposition and ending when pupation began. No treatments were given at generation 6.

Calculations: Two parameters, associated with fitness and likely to be affected by inbreeding, were calculated. (1) Productivity (defined as the number of pupae obtained from a pair of adults ovipositing for 7 days in 1 g. of medium). This parameter is the combined result of parent fecundity and offspring survival to pupation. (2) Median developmental period of offspring from egg to pupa. Developmental period data were analyzed without transformation. Square-root transformation of pupal counts were used in the analysis of productivity.

Results

The mean productivity of the experimental groups is illustrated in fig. 1 (left). It is instructive to compare the productivity of C lines to that of V lines at generation 6 in which all lines were maintained in the constant environment. In this generation the mean productivity of V lines was significantly higher than the C lines ($P < 0.05$ by t test).

The median developmental periods of V and C are illustrated in fig. 1 (right). The developmental period of the V group was generally longer than that of the C group. However, in generation 6, when all lines were tested in the constant environment, the average median for V was shorter than for C, although the difference between them was just barely significant.

At generation 3, hybrid lines were made by crossing between C lines. (C x C). At generation 4, the following single-pair crosses were made: C x C, V x V, and C x V. In addition, the offspring of the cross in the preceding generation were sibmated (A) and cross-mated (B). For each cross, several replicates (8 to 16) were made. All

crosses were maintained in the constant environment. Examination of the data reveals a significant difference in productivity between group B (cross-mated) and their best parents C_I ($P < 0.01$), but no such difference in group A. Productivity of B was almost as high as in the beginning of the experiment. (fig. 2a).

There were no significant differences in developmental period between hybrids in the different crosses, but it was shorter than that of the inbred lines ($P < 0.01$). (fig. 2b).

Discussion

The results indicate that the sib-mated V lines, with a history of 5 generations of exposure to unpredictable short-term treatments, were more fit (as indicated by their higher productivity and shorter developmental time) than the C lines, which were of the same origin and practiced the same mating system but maintained in the constant environment, in which C and V were compared at generation 6.

Sib-mating, with subsequent sampling of but one pair of offspring to continue a line, could be expected to lead to rapid homozygotization of the genome in each line, exposing deleterious alleles to selection pressures. Moreover, even homozygosity per se should be more deleterious in a suboptimal variable environment than in a favourable constant one, because homozygous individuals may lack the physiological flexibility to adapt to variable conditions (Lerner, 1954). Therefore, populations living in variable environments should include a higher proportion of heterozygous individuals than in constant environments. We conclude that homozygous individuals were less likely to be included as parents in the V lines than in the C lines, and therefore were selected against in the variable environment. This process has reduced the rate of homozygotization in the V lines, resulting in their better fitness. This conclusion is supported by the fact that outcrossing at generation 5 (fig. 2) has restored productivity to its originally high level, and that hybridization reduced developmental time of all crosses compared to the inbred lines.

Table 1. Experimental treatments for the V group. To obtain the complete sequence of experimental history, the generations should follow each other in sequence. No treatments were given during the oviposition and pupation periods.

Treatment symbols:

- 0 no treatment
- 1 cold treatment: 2°C for 1 hour
- 2 heat treatment: 38°C for 24 hours
- 3 starvation treatment: 24 hours without food, at 30°C
- 4 competition treatment: 7 days competition with adult males of T. confusum (black mutant)
- 5 competition treatment - same as 4 but for 10 days.

WEEKS

	1	2	3	4			
Gen. 1	Oviposition No treatment	1	0	2	0	Pupation No treatment	
Gen. 2		3	0	3	3		2
Gen. 3		0	0		4		0
Gen. 4		0	2	1	1		3
Gen. 5		0		5			0
Gen. 6		0	0	0	0		0

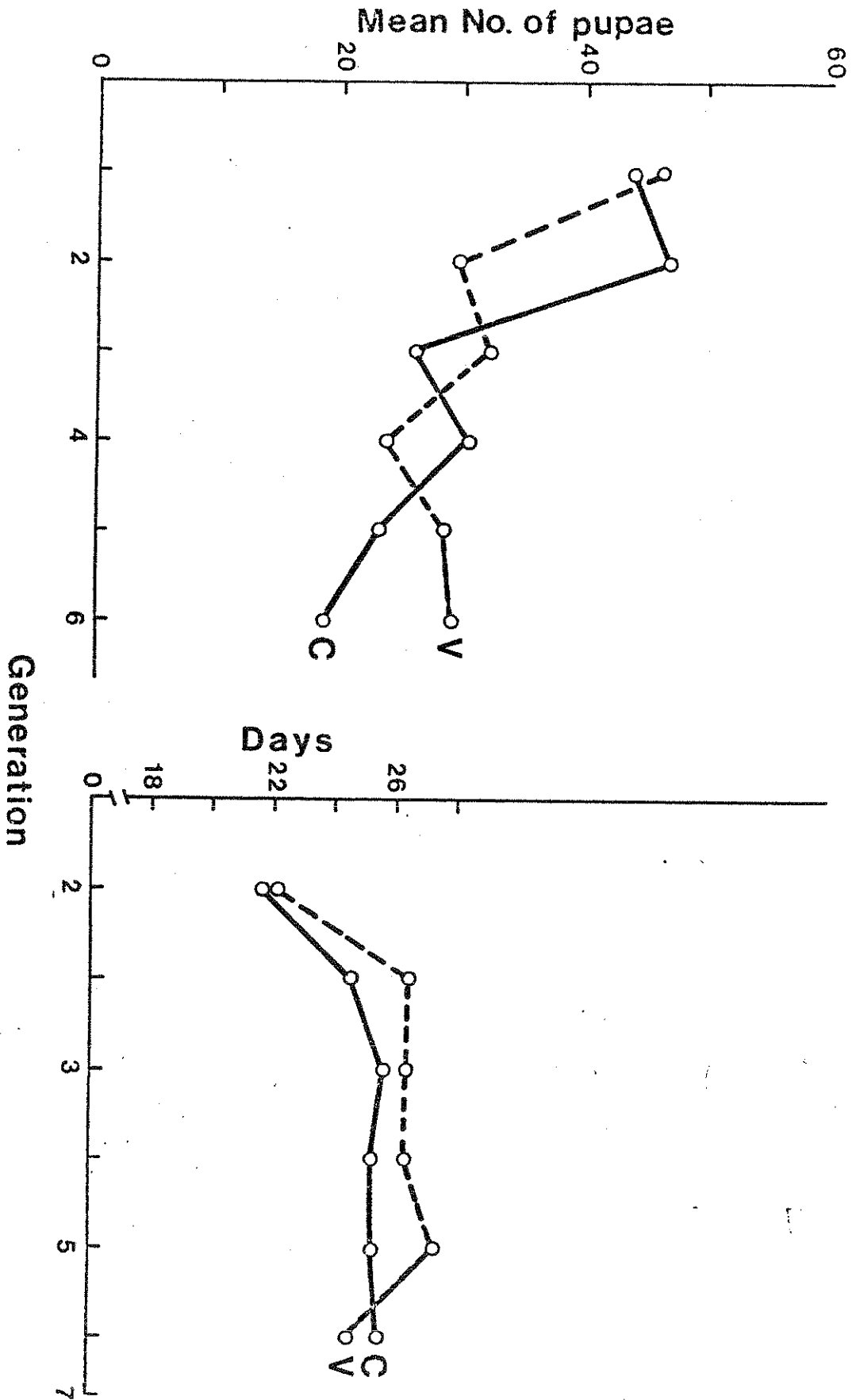


Fig. 1. (left) (right)

Mean productivity of the 2 experimental groups
Average median egg-to-adult developmental period
(in days) of the V and C groups.

(Variable environment: broken lines
Constant environment: solid line)

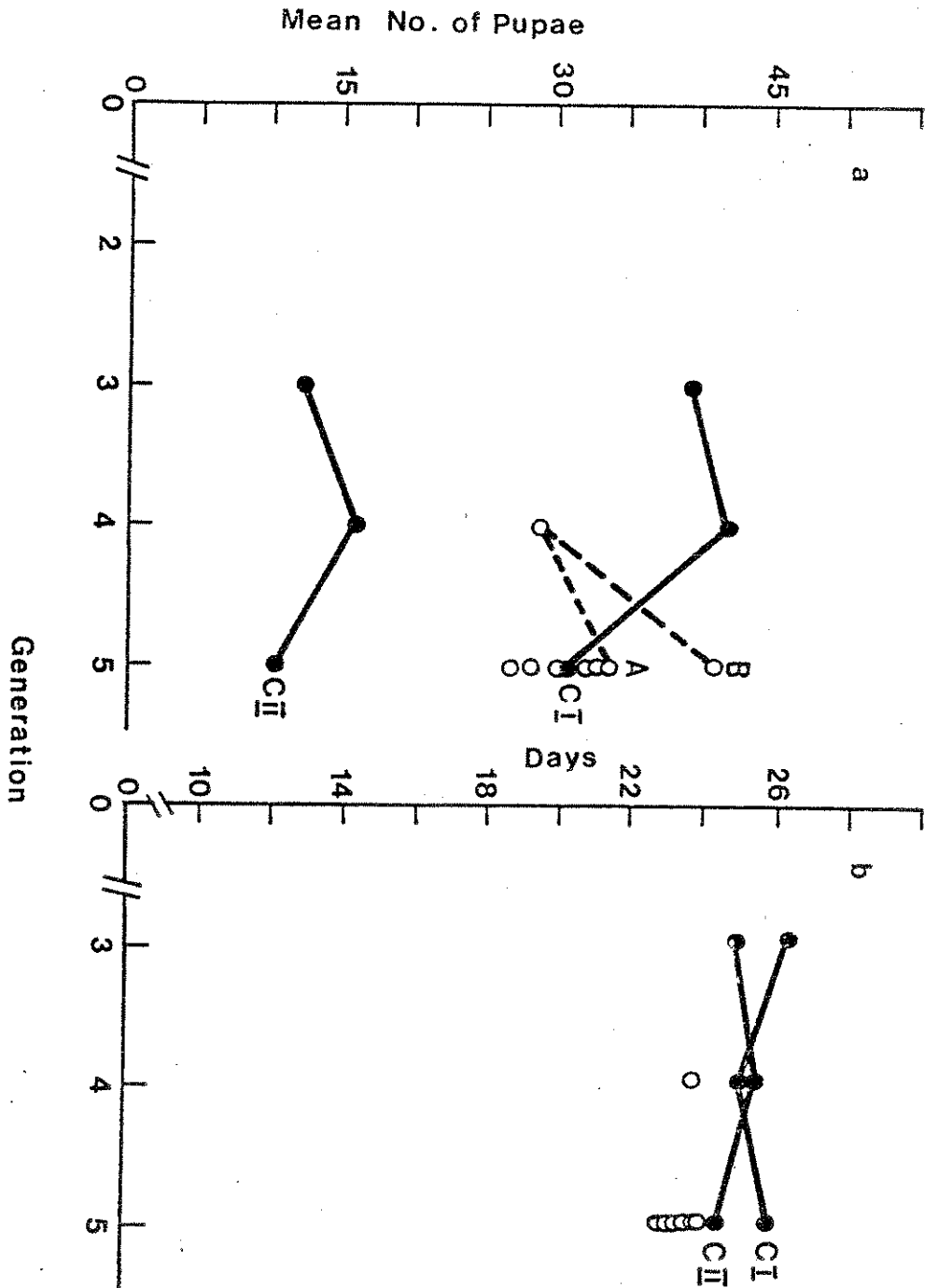


Fig. 2. Productivity (a) and developmental period (b) of the hybrid lines compared with their inbred parents CI and CII, at generations 3 - 5. Solid circles - inbred lines. Open circles - hybrids. A and B are the sib-mated (A) and cross-mated (B) offspring of the hybrids obtained at generation 4. (See text).

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The influence of the male on female egg-laying of "Tribolium castaneum".

A series of experiments were carried out in order to study the influence of the male of Tribolium castaneum on the egg laying of the female to whom it has been mated. Some preliminar work on this subject has been reported by Orozco et al. (1971).

Insects from the Consejo population (Orozco and Bell, 1974) were used, and they were kept at 33°C and 70% relative humidity. All matings were individually made.

The following effects were considered:

L_V = Egg laying of virgin females.

L_F = Egg laying of fecundated females.

L_M = Egg laying of virgin females in the presence of a male which was separated from the female by means of a wire netting placed in the corresponding vial.

L_D = Egg laying of females mated to males previously irradiated in a caesium bomb at the following dosages: $D_1 = 16$ Krds, $D_2 = 20$ Krds, $D_3 = 30$ Krds. All dosages completely sterilized the males.

L_C = Egg laying of females mated to Tribolium confusum males. These matings were all unviable.

Three types of experiments each with several replications were carried out. In each experiment virgin and fecundated females were compared with females submitted to other treatments as shown in Table 1.

Table 1. Effects tested in each experiment.

Experiment	Effects tested						Number of replications
	L _V	L _F	L _{D₁}	L _{D₂}	L _{D₃}	L _M	
I	x	x	x		x		3
II	x	x				x	2
III	x	x		x		x	4

Table 2 presents the egg laying average figures for the four treatments of experiment I (both by replications and for the pooled data), together with the mean squares of the corresponding analyses of variance.

Table 2. Four-days average egg laying with its standard error and mean squares from analyses of variance of Experiment I.

		Rep. 1	Rep. 2	Rep. 3	Pool
average values per treatment	L _F	76.38±1.39	48.83±1.98	73.23±2.15	66.15±1.29
	L _{D₁}	53.97±1.57	30.82±1.38	42.37±1.79	42.39±1.07
	L _{D₃}	56.03±1.58	31.76±1.76	41.87±1.90	43.22±1.18
	L _V	37.67±1.45	21.26±1.61	22.20±1.82	27.04±0.93
mean squares of ANOV	Repli- cates	-	-	-	49,481.21**
	Treat- ments	25,179.10**	21,055.51**	56,644.50**	76,220.59**
	R x T	-	-	-	2,737.36**
	Resi- dual	223.71	258.16	323.14	266.81

** (P < 0.01)

No differences were found between doses of irradiation and the lay of the females mated to those irradiated males was intermediate between those of virgin and fecundated females with differences in both cases highly significant (P < 0.01). The significant effect found for the replication x treatment component of variance in the pooled analysis is of a quantitative type rather than a qualitative one since the same pattern was found in the four replicates for the performance of the four treatments.

Table 3.- Four-days average egg laying with its standard error and mean squares from the analyses of variance of Experiment II.

		Rep. 1	Rep. 2	Pool
average values for treatment	L _F	57.23±2.04	36.75±1.48	46.99±1.45
	L _M	34.25±1.57	15.94±0.89	25.58±1.14
	L _V	25.37±1.69	15.16±0.95	20.33±1.06
mean squares of ANOVA	Replicates	-	-	38,034.44**
	Treatments	27,033.24**	14,671.42**	39,754.49**
	R x T	-	-	1,950.17**
	Residual	315.65	136.95	227.83

** (P < 0.01)

Table 3 is as Table 2 but for the data from experiment II. In this case the egg laying of virgin females separated from the male by a wire screen placed in the vial was in both replicates far below the lay of fecundated females. However, although the difference L_M - L_F was not significant in replicate 2, it was highly significant (P < 0.01) but small (8.88 eggs) in replicate 1. In the pooled analysis the difference L_M - L_F = 5.25 eggs was not significant even at the 0.10 level, but in this case the highly significant effect for the interaction replication x treatment reflects the contradictory results obtained in both replications.

The average laying figures for the four replicates of experiment III and their pool are presented in Table 4, with the mean squares for the corresponding analyses of variance. All the effects in these analyses were highly significant (P < 0.01). The reason for the significance of the interaction are the small differences in ranking order between L_V, L_C and L_M when considering the four replications.

The treatment L_F was always superior to L_{D2} and the performance of L_{D2} always larger than those for the other three treatments, with highly significant differences (P < 0.01) in each replication and also in the pooled analysis. Comparing L_C and L_M with L_V does not produce important differences in some replicates but in other cases it gives significant differences even though they are smaller than those observed with respect to L_F and L_D. Table 5 presents all the set of differences for the pooled analysis and from them it can be deduced that the performances of L_V, L_C and L_M are not significantly different. Only in the first replication of this experiment III, L_M and L_C were larger than L_V; but given the results in the other three replications and in the pooled analysis, it seems reasonable to state that L_F L_D L_C = L_M = L_V.

Table 4.- Two-days average egg laying with its standard error and mean squares from the analyses of variance of Experiment III.

	Replications				Pool
	1	2	3	4	
LF	33.02±0.78	33.56±0.72	31.88±0.94	31.70±0.97	32.54±0.43
LD ₃	30.01±0.86	19.97±0.83	19.22±0.57	22.02±0.90	22.83±0.45
LM	19.84±0.83	12.79±0.80	11.11±0.46	9.73±0.59	13.53±0.42
LC	16.21±0.70	9.73±0.51	10.26±0.51	12.11±0.76	12.07±0.32
LV	10.33±0.71	11.03±0.73	14.03±0.67	11.01±0.92	11.62±0.38
Replicates	-	-	-	-	2,403.00**
Treatments	8,971.75**	9,558.62**	7,800.71**	8,035.50**	32,012.54**
R x T	-	-	-	-	784.68**
Residual	60.30	51.44	46.90	63.63	55.35

** (P < 0.01)

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Cannibalism among different developmental forms of the black and pearl foundation Tribolium populations.

Wool (1969) had recognized that cannibalism is an important factor in the regulation of competing Tribolium populations. Kence (1971) and Miller and Kence (1973) studying cannibalism between different genotypes at the black locus observed preferential cannibalism of one genotype over the others. However, Sellin and Krause (1973) using mixed cultures of Short antennae (Sa-2) and wild type could not detect any preferential cannibalism although random observations previously supported Kence's findings.

Park et. al. (1965) have shown that cannibalism can occur at many stages of the beetle's development: larva-egg, larva-pupa, adult-larva, adult-pupa and adult-egg. To investigate the cannibalism preferences of different developmental forms, a preliminary study involving black and pearl foundation populations was made. Krause and Bell (1972) had found these populations differed in a number of basic traits, in particular, the pearl larva were 50% larger than the black at 13 days. Competitive situations between larva-pupa, adult-pupa and adult-larva were investigated under various amounts of media.

Material and Methods

Different stages of development were collected from mass matings of 10 male and 10 females. Five bottles each of the black (black body color) and pearl (pearl eye color) foundation populations of Tribolium castaneum were maintained on standard media (95% sifted whole wheat flour and 5% dried Brewer's yeast) in an incubator set at 91±1°F and 70±2% R.H. These matings were transferred into fresh media every 24 hours to maintain similar age limits of the developmental forms used: larva (13 days), pupa (18 days) and adults (25 days).

Table 5.- Differences between treatments and their significance for the pool analysis of experiment III. (Newman Keuls sequential test).

	L _V	L _C	L _M	L _{D₃}
L _F	20.92**	20.47**	19.01**	9.71*
L _{D₃}	11.21**	10.76**	9.30**	
L _M	1.91	1.46		
L _C	0.45			

** (P > 0.01)

From the analysed experiments it may be concluded that:

a) Given the well known fact that the virgin females lay less eggs than the fecundated ones (about one third) it has been shown clearly that females mated to irradiated sterile males perform intermediate.

b) However, the lay of females mated to males of another species (Tribolium confusum) was not enhanced and their lay was equal to the virgin ones.

c) Finally, when females were near a male of the same species but physical copulation was prevented, it seemed that the egg laying was not increased, at least to a great extent. This result is not very conclusive even with the obtainment of two positive cases out of the six tested, although this may reflect the complexity of the treatment.

If males of Tribolium confusum copulate with females of Tribolium castaneum as indicated by Sokoloff (1972) it is necessary to think that the stimulation to lay from the copulation itself with males of the same species is different and positive.

In this way the increase in egg laying could be due to two facts: first the copulation and corresponding physiological factors and afterwards the fertilization of the eggs.

The possible stimulus for increasing lay due to the presence of the male near the female through some influence of pheromones is therefore in some doubtful state after the results obtained in experiments I and III. The slight increase in two replicates is not enough to conclude that this stimulus is a positive one.

Six combinations of competitive cannibalism were observed between the black and pearl populations: larva (predator) - pupa (prey); adult (predator) - pupa (prey); adult (predator) - larva (prey). Each combination maintained in 3/4 oz. creamer consisted of 25 prey and 10 predators which were kept together for four days. Four creamers were used for each combination, resulting in a total of 100 potential prey. The number of prey surviving were used as a measure of cannibalism. To differentiate cannibalism from livability, a control group of 100 beetles were observed simultaneously. Three replicates of the experiment were used, each under different environmental conditions: no media, little media (just enough media to cover the bottom of the creamer) and excess media (10 q. per creamer).

Chi-square tests were used to compare the various parameters studied.

Results and Discussion

The results of the various predator-prey combinations between different developmental forms, between genotypes within a single species and between various environmental regimes are shown in Table 1a. With exception, the excess media regime results in no appreciable cannibalism as compared with the more stressful conditions of little or no media. Since beetles are a tunneling organism, the excess media regime provides both sufficient food and decreases the chance encounter between predator and prey.

Kence (1973) observed genotypic differences in cannibalistic behavior between larva and pupa. In our study (Table 1a) the survival of the pupa were significantly lowered from that of the controls (Table 1b), particularly in the no and little media environments. The controls here serve as a measure of livability of the prey under the various environments. With exception, similar lowering of pupal survival was observed (Table 1a) when adults were used as predators. However, with the larva-pupa situations, the pearl were the better cannibalizers while the black adults are better cannibalizers than the pearl in reference to pupa survival.

When adults were used as predators and larva as prey (assuming the larva are the prey, although conceivably it could be reversed), decreases in larval survival was observed on both no and little media regimes. However, in reference to the no media regimes, the controls (Table 1b) had poor viability, probably due to the necessity of food for complete development. Again the black adults had better appetites.

Ideally, it would be desirable to study preferential cannibalism of one genotype versus the other. An error in the experimental design became evident in analyzing the data when no information from the control combination of black predator on black prey and pearl on pearl were obtained. This limits the interpretations in

reference to preferential cannibalism. From this preliminary study, it seems that the pearl larva, which are larger in size, are better cannibalizers of the black pupa. However, size differences are probably not the only causes of observed difference in response between the genotypes since black adults are better cannibalizers of pearl larva and pupa than the reverse combination.

Table la. Number of prey surviving (of 100 organisms) exposed to different predator-prey combinations under various environmental regimes

Predator	Prey	Environment regimes		
		No media	Little media	Excess media
Black larva	Pearl pupa	52	90	94
Pearl larva	Black pupa	13	84	93
Black adults	Pearl pupa	8	13	65
Pearl adults	Black pupa	45	26	99
Black adults	Pearl larva	1	81	90
Pearl adults	Black larva	13	98	100

Table lb. Number of survivors (of 100 organisms) corresponding respectively to the prey (in Table la.) exposed to various environmental regimes.

	Environment regimes		
	No media	Little media	Excess media
Pearl pupa	98	92	93
Black pupa	98	95	98
Pearl pupa	97	100	98
Black pupa	100	99	93
Pearl larva	8	100	99
Black larva	40	99	100

Summary

Preferential cannibalism was observed, particularly, under the stressful conditions of little or no media. The pearl larva were better cannibalizers of the black pupa than the reverse combination. Part of this differential response may be accounted for by the pearl larva being larger in size. In addition, the black adults were observed to be better cannibalizers of both the pearl pupa and larva.

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NOTES - TECHNICAL

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A container for studying oviposition rate in the absence of egg cannibalism.

Sonleitner (1961) described a plastic replica of a shell vial (similar in construction to the fractionable shell vial of Ghent, 1966) which he used to study oviposition rate of beetles on the surface of the flour medium. A single pair of beetles confined in such a device exhibits an oviposition rate only slightly less than that of a single pair in 4 grams of medium in a shell vial. Because all eggs are oviposited below the subsurface mesh, they are safe from cannibalism. Thus the device enables one to study the oviposition rate of single pairs under fairly normal conditions but in the absence of egg cannibalism. Controlled numbers of eggs may be fed to the pair by introducing them onto the surface of the medium above the mesh. Such introduced eggs should be marked (Sonleitner, 1961) so that they may be distinguished from those produced by the pair.

Such a tower is made from several short sections of lucite tube (1 inch in diameter 1/16th inch wall thickness, 1/2 inch long.) The bottom section is cemented to a small square of plexiglass which serves as a base. The top section has a piece of OOX bolting cloth cemented across its lower end. When the bottom section is overfilled with medium and the top section placed over it and fastened in place with Scotch tape, the bolting cloth should be buried just beneath the surface of the medium. The beetles are then placed in the top section on the surface of the medium. Eggs are recovered by taking the tower apart and sifting the medium in the bottom section.

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