

TRIBOLIUM INFORMATION BULLETIN

VOLUME 39

1999

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Tribolium Information Bulletin, Volume 39

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NOTE

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ACKNOWLEDGMENTS

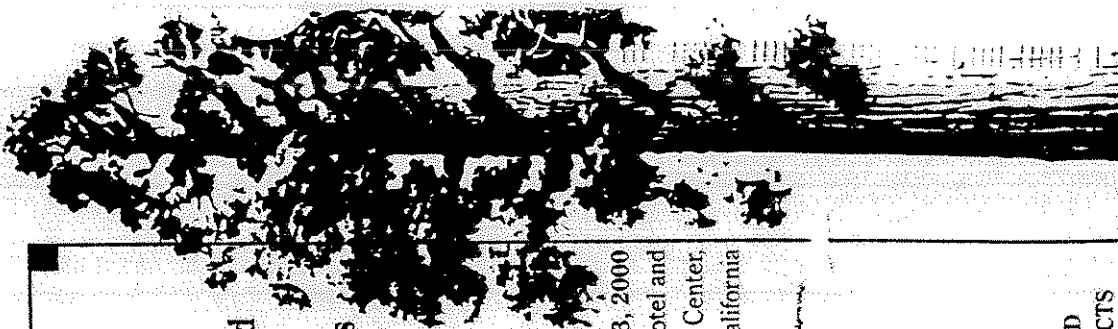
THE EDITOR IS INDEBTED TO BARBARA SOKOLOFF AND ELAINE SOKOLOFF FOR ASSISTANCE IN THE PREPARATION AND DISTRIBUTION OF TIB 39. I ALSO THANK FREDERIC ARMAND AND ILIA ADAME FOR TYPING PART OF THE REVIEW OF TRIBOLIUM BREVICORNIS.

ANNOUNCEMENT I

FOR SALE

A SMALL NUMBER OF SETS OF SOKOLOFF'S THREE VOLUME THE BIOLOGY OF TRIBOLIUM WITH SPECIAL EMPHASIS ON GENETIC ASPECTS IS AVAILABLE FROM THE EDITOR ON A FIRST COME-FIRST SERVED BASIS AND ONLY AS A FULL THREE VOLUME SET.

PRICE: \$180/SET (INCLUDING POSTAGE, HANDLING & INSURANCE)



International Conference on Controlled Atmosphere and Fumigation in Stored Products

Oct. 29 - Nov. 3, 2000
Radisson Hotel and Convention Center,
Fresno, California

Second Announcement

REGISTRATION AND CALL FOR ABSTRACTS

J.G. Leesch
CAF ORGANIZING COMMITTEE
USDA-ARS-HCRL
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DR. ALEXANDER SOKOLOFF
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92407-2318 60

OFFICIAL LANGUAGE

The official language of the conference will be English.

FRESNO

The city of Fresno is located in the central San Joaquin Valley of California midway between San Francisco and Los Angeles. It is the heart of agricultural production for the state and represents a significant amount of the production in the United States. Major agricultural products include fresh fruit such as citrus, peaches, plums, pears, apples, apricots, grapes and durable products such as cereals, cotton, walnuts, almonds, pistachios, raisins, prunes and other dried fruit.

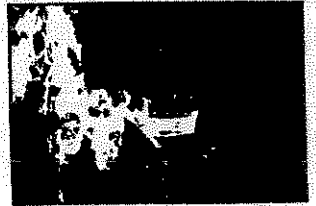
October is the end of the busy harvesting season in the Valley with only some walnuts and persimmons remaining to be processed. The weather is changing from the heat of summer to the mild climate of fall. Leaves are beginning to change to golden yellows and brilliant reds. Near Fresno are such popular sights as Yosemite National Park, Sequoia National Forest (home of the Giant Redwoods) and Kings Canyon National Park. A post conference tour to the Yosemite National Park can be arranged, if enough people are interested.

Fresno is easily accessible from either San Francisco, Los Angeles, Phoenix and Salt Lake City by air with several flights each day to connecting flights in those cities.



PERMANENT COMMITTEE

J Banks, Australia
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 R. Noyes U.S.A.
 Ch. Reichnuth, Germany
 B. E. Ripp, Australia
 M. Sidik, Indonesia

**ABOUT THE CONFERENCE**

The International Conference on Controlled Atmosphere and Fumigation in Stored Products (formerly the International Conference on Controlled Atmosphere and Fumigation in Grain Storage) is the leading international meeting reporting on advances in research and development on gaseous treatments applied to protect commodities in storage. These commodities include cereal grains, legumes, root crops, dried fruits and vegetables, and other agricultural products including bulbs and flowers, excluded are CA treatments for quality preservation of fresh agricultural products. The permanent Committee of the International CAF Conference (in a meeting held during the Conference in Cyprus, 1996) selected the Horticultural Crops Research Laboratory (HCRL), USDA, ARS, in Fresno, to host the next Conference, because the HCRL has been instrumental in implementing advanced dried fruit and nut processing and storage technologies under hot and dry climate conditions of California.

This conference will be held under the cloud of current worldwide pressure to reduce or phase out use of fumigants in general, and methyl bromide in particular. These constraints pose new challenges to technologically advanced countries and in particular to countries heavily relying on the use of conventional fumigants.

CONFERENCE OBJECTIVES

- *To report on advances in research and development and the current status of controlled atmosphere and fumigation for insect pests, microflora and quality control in stored products.*
- *To evaluate and discuss the potential alternative control methods in view of the threat to reduce or phase out use of chemical fumigants in general and methyl bromide in particular.*
- *To evaluate the technical, environmental and economical advantages of, and examine the potential for, emerging controlled atmosphere technologies.*
- *To provide a forum for interchange of ideas between government authorities, research agencies and industry.*
- *To enable the industry to display the latest commodity treatment, monitoring, handling and storage equipment, facilities and services.*
- *To discuss, disseminate information, and allow transfer of technologies applying controlled atmosphere and fumigation.*

CONFERENCE FORMAT

Speakers may present papers covering different aspects of research and application of controlled atmosphere and fumigation techniques on stored products. The tentative titles of the sessions are:

1. Biological responses of arthropods to treatment with CA and/or fumigation.
2. Biological responses of microflora to treatment with CA and/or fumigation.
3. Influence of CA and/or fumigation on quality preservation of stored products.

4. Physical and chemical processes in controlled atmosphere CA and/or fumigation.

5. Application methodology of CA and/or fumigation, including use of carbon dioxide under increased pressure.

6. Sealing techniques and methods of determining gas-tightness.

7. Integrated commodity management methods with CA and/or fumigation.

8. Potential threats to conventional CA and/or fumigation (regulatory insect resistance).

9. Quarantine and regulatory issues pertaining to use of CA and/or fumigation.

10. Sampling and trapping to monitor insect populations in relation to CA and/or fumigation.

A Poster Session will constitute an integral part of the program and will allow presentation of a wide variety of specific research projects related to the above sessions.

PROCEEDINGS

Oral and poster presentations will be considered for publication in the Proceedings. The Proceedings will be fully edited. Full manuscripts of each presentation/poster will be submitted by authors at the conference.

LOCAL ORGANIZING COMMITTEE

Jim G. Leesch, *Chairman*
 Libby Fouse, John Hendon, Gary Obernuth,
 Jane Tebbets, Pat Vail, Larry Zettler

PARTICIPANTS

The Conference will be of benefit to a wide range of interests associated with the preservation of grain, oilseeds, dried fruit and similar commodities in storage including:

Dried Fruit and Nut Industries, Grain Handlers, Food Processing Companies, Equipment, Materials and Chemical Suppliers, Engineers and Storage Designers, Pest Control Officers, Food Technologists, Entomologists, Mycologists, Food Company Directors and Managers, Researchers, Government/Regulatory Agencies.

FOR ADDITIONAL INFORMATION

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ACCOMMODATION INFORMATION (Requires Action):

The hotel of the CAF Conference is the Radisson Hotel & Conference Center at 2233 Ventura Street, Fresno, CA 93721.

The room rate is \$87.00 per night for all rooms (up to 4 persons occupancy) plus hotel tax which is currently 12%.

Reservations are to be made by each participant directly with the hotel. For those not guaranteeing a room with a credit card, the first nights room charge must be paid in advance to the hotel payable by either check (North America only) or money order (all in US\$). There is a 72 hour cancellation policy. The rate for the Conference will apply 3 nights before and after the conference for participants who wish to stay in Fresno either before or after the conference. The hotel will not guarantee the room rate after October 18, 2000. You can make reservations by calling (800) 333-3333 or (559) 268-1000 and mentioning the CAF Conference and its dates. You can also register by internet at www.radisson.com or by mailing or faxing this form, along with your credit card number, check or money order to:

Radisson Hotel & Convention Center • 2233 Ventura Street • Fresno, CA 93721 • USA • Fax: (559) 268-72

Name: _____ Phone: _____

Address: _____

City: _____ State: _____ Country: _____ Zip: _____

Check-in date: _____ Check-out date: _____

Number of rooms requested: _____ Number of people: _____

Room type (single or double) requested: _____ () Smoking () Non-Smoking

Amount enclosed US\$ (Minimum \$100.00): _____

Credit Card #: _____ Expiration date: _____

Name of Card (Visa, MasterCard, American Express etc.): _____

Authorized Signature: _____ Date: _____

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CAF ORGANIZING COMMITTEE
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ANNOUNCEMENT II

INTERNATIONAL CONFERENCE ON CONTROLLED ATMOSPHERE AND FUMIGATION IN STORED PRODUCTS

APPLICATION FOR PRESENTATION

POSTER PRESENTATIONS: Posters should be prepared to fit in a space measuring 122 cm by 122 cm (4ft by 4ft). Format for the posters is the same as that for oral presentations (i.e. Title, Abstract, Introduction, Materials and Methods, Results, Discussion and Conclusions). They will be published in the Proceedings.

ORAL PRESENTATIONS: Oral presentations will be limited to 15 or 20 minutes depending on the number of presentations to be made. Presentations can be made with either overhead transparencies or 35mm slides. An overhead projector and a slide carousel projector with slide trays will be provided.

DEADLINE TO RECEIVE ABSTRACTS FOR PRESENTATIONS AND POSTER: JANUARY 31, 2000. Abstracts can be transmitted by e-mail to: Dr. J. Leesch JLeesch@gnis.net

INSTRUCTIONS FOR ABSTRACT PREPARATION: Abstracts of presentations and posters will be presented in the Program that will be distributed to participants upon registration at the beginning of the Conference. Therefore a standard size and uniformity will be required of each abstract. Abstracts should be typed (preferably using a word processor such as Word or WordPerfect for Windows). The size of the abstract should be no longer than 1 page and the total height and width of the text on the page should be within a measurement of 15 cm wide by 22 cm high. The top line should be the title of the presentation (poster in block (all capital) letters. The second line should list the author(s) with the last name in block letters. The third and fourth line should be the full address of the institution or affiliation of the author(s). The text should follow using font size of 10 or 12 point and a plain style such as Times Roman or Courier. The abstract text is limited to 250 words and should not contain tables and figures.

INDUSTRIAL DISPLAYS: Space for an exhibition of scientific, technical and commercial equipment will be provided by the Conference.

If you are interested in exhibiting during the conference, please contact the Organizing Committee to receive information as to available space, fees and time for exhibition.

OFFICIAL LETTER OF INVITATION: To facilitate efforts to obtain financial support for attending the Conference, an official letter of invitation will be sent to participants upon written request of the Secretariat. The letter does not obligate the Organizing Committee to pay for either travel or other expenses incurred by the participants. However, participants who cannot get financial aid to attend the Conference may write the Organizing Committee to obtain information as to the availability of funds to assist in travel and other expenses.

CAF REGISTRATION FORM

Last Name _____ First Name _____ Middle Initials _____

Name for Badge _____ Title: Prof () Dr () Mr () Ms ()

Company/Organization _____

Address _____

City _____ State _____ Country _____ Postal Code _____

Phone: _____ Fax: _____ E-mail: _____

Accompanied by _____ (Last Name) _____ (First Name)

REGISTRATION FEES (in U.S. Dollars)

Registration until April 30, 2000	(US\$ 350)	=	_____
Registration after April 30, 2000	(US\$ 425)	=	_____
Accompanying Person**	(US\$ 60)	=	_____
Additional Conference Tour Tickets	(US\$ 55) (2 tours available during conference)	=	_____
Additional Banquet Tickets	(US\$ 35)	=	_____
Post Conference Tour to Yosemite	(US\$ 60) X No. Attending _____	=	_____
TOTAL =			_____

IF PAYING BY CREDIT CARD, PLEASE COMPLETE THIS PART.

I agree to pay the sum of total fees above for US\$ _____

This is for registration and optional tour deposits (if applicable) for the CAF Conference.

My credit card is: () Visa () MasterCard () American Express

Card Number _____

Cardholder's Name: _____ Expiration Date: _____

Signature _____ Date: _____

(Please be sure that all information supplied is legible. We cannot be responsible for errors due to inability to read information provided.)



* Registration Fees cover: Admission to the Conference, 1 Program with abstracts, welcoming reception at the Radisson Hotel, 1 ticket to the banquet at the Radisson Hotel, beverages and snacks during breaks, 1 ticket to each of the 2 Conference tours, Proceedings of the Conference (book format).
 **Accompanying Person Fees cover: Admission to the Conference, ticket to the welcoming reception at the Radisson Hotel, and Spouses Program.

OVER for accommodation information that requires action.



XXI International Congress of Entomology Iguassu Falls - Brazil August 20-26, 2000

Dear entomologist

As you are probably aware, Brazil will host the next International Congress of Entomology, from August 20 to 26, 2000, at one of the most beautiful world sites, Iguassu Falls. The Organizing Committee, composed of entomologists from several countries, is doing its best to make the XXI ICE one of the most important entomological events ever held.

Besides the formal congress program, scientists will have the chance to participate of satellite entomological events. Up to the end of June, we are receiving suggestions for symposia, and if you want to contribute, send your suggestion to the president of the Scientific Committee Programme, Dr F. Moscardi, at moscardi@cnpso.embrapa.br. We will also provide social and touristic activities for both participants and accompanying persons. For more information, please visit our homepage at <http://www.embrapa.br/ice>.

During the last two years we have organized a database of more than 12.000 addresses of entomologists, from all over the world, including yourself. If you want to continue receiving information on the XXI ICE, please confirm your interest through our homepage, looking for the icon "Subscribe to the congress mailing list now" on the bottom of the front page. Or directly, on the address <http://centeio.sede.embrapa.br/ice/ICE.nsf/MailingList?OpenForm>.

In case you do not have access to the Internet, please fill the form below and send it to us, to continue receiving XXI ICE information. If we do not receive your confirmation, we will assume that you are not interested in participating of the XXI ICE and your name will be dropped from our mailing list.

Thank you for your help, and we hope to seeing you during the Congress

Family Name	_____
First Name	_____
Institution	_____
Address	_____
City, ZIP	_____
State	_____
Country	_____

Our address
XXI International Congress of Entomology
Caixa Postal. 231 - 86001-970 Londrina PR - BRAZIL

**P R E L I M I N A R Y A N N O U N C E M E N T
G E N E T I C S C O N F E R E N C E • M A Y 7 - 1 1 , 2 0 0 0**

PRESERVING THE GENETIC HERITAGE OF LIFE ON EARTH.

The Zoological Society of San Diego and its research division, the Center for Reproduction of Endangered Species, will be hosting an international conference to consider the values, benefits, means, and options for conserving the genetic heritage of life on Earth. In January, you will be sent a registration brochure with more details of this upcoming conference. Conference dates are May 7-11, 2000. For more information visit us at: www.sandiegozoo.org/cres/genetic_conference.html



ZOOLOGICAL SOCIETY OF SAN DIEGO®

G E N E T I C R E S O U R C E S F O R T H E N E W C E N T U R Y

3rd Announcement

GSC 2000

**The 2000 Annual Meeting of the
Genetics Society of Canada**

will be in

Vancouver

with the Genetics Society of America

on the Campus of the University of British Columbia

June 14 - 17, 2000

The 2000 Annual Meeting of the Genetics Society of Canada will be held Vancouver B.C.. For GSC members, there will also be the usual student poster and travel bursar competitions, the award lectures and a banquet. Accommodation for the meeting will be available at downtown hotels or on the campus at the University of British Columbia.

For further information please see the inside of this Bulletin issue, visit the new meeting WWW site at,

<http://www.zoology.ubc.ca/genetics/2000>

and keep an eye on your mailbox. Further inquiries about the meeting can be directed to:

Tony Griffiths
agriff@unixg.ubc.ca

Genetics 2000: Visions of the future

A joint meeting of the
Genetics Society of Canada and the
Genetics Society of America.

On the campus of the
University of BC, Vancouver, Canada,
June 14 - 17, 2000

As society enters the new millenium, genetics is on the brink of an era that will see large changes in the ways that genetic research is carried out, and the ways in which society perceives this research. The focus of this extraordinary meeting is best summed up by the word "vision". In plenary session, senior geneticists will present their vision of the themes that will occupy geneticists into the new millenium. These discussions will be relevant to all geneticists, including students.

There will also be regular poster sessions for presenting current research. Some presenters may be invited to participate in the plenary sessions.

Plenary sessions will include the following areas:

Changing concepts in genetics.
Genetics and the human organism.
Genetics in human affairs.
The future of model genetic organisms.
Industry versus academy in the future of genetic research.
The future of genetics education.

So far, the following individuals have accepted invitations to speak:

Patricia Baird (University of British Columbia)
David Botstein (Stanford University)
James Crow (University of Wisconsin)
Julian Davies (University of British Columbia)
John Doebly (University of Wisconsin)
Clare Fraser (The Institute for Genome Research)
Bill Gelbart (Harvard University)
Anthony Griffiths (University of British Columbia)
Dan Hartl (Harvard University)
Mary Claire King (University of Washington)

Chuck Langley (University of California, Davis)
Maurice Moloney (University of Calgary)
Arno Motulski (University of Washington)
Stephen O'Brien (National Cancer Institute)
Charles Scriver (McGill University)

Acomodation and registration costs are very affordable. GSC members pay a reduced rate. (Now would be a good time to join!) The abstract and late registration deadline is March 31, 2000.

Registration, acomodation and abstracts are processed through the Genetics 2000 website:

<http://www.zoology.ubc.ca/genetics/2000>

For more information, please contact the head of the Local Organizing Committee:

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The participation in discussion of the first topic of discussion entitled "Interactions in Tribolium: Competition or predator-prey?" was very disappointing. Instead of a forum there was one response, enough only for a dialogue. Because the Editor considers the subject worth discussing, the subject suggested for the open forum remains open for discussion, and it will be for the next two years.

As usual, the Editor reminds subscribers that the very existence of the TIB is dependent not only on subscription, but also on contributions to the Newsletter. Please be as generous of your time as possible by responding when calls for contribution arrive in your hands. This includes not only research notes but also revision of personnel in your lab, stock lists, and lists of current bibliography.

TRIBOLIUM NEWS EXCHANGE MEMBER LIST

A small group of Tribolium investigators have initiated a Tribolium News Exchange. Its purpose is to provide an informal forum to exchange ideas, techniques and suggestions about Tribolium.

If you wish to join the Tribolium News Exchange Group, contact Margaret Bloch Qazi at her email address:
tribolium@emerald.tufts.edu

Announcement VII

At long last we have received permit for importation of 3 species of Tribolium: *T. anaphe*, *T. audax* and *T. destructor*. If you are interested in obtaining samples of these beetles in the near future contact me by fax or e-mail.

TOBI THOMAS (BIOLOGICAL SERVICE ORGANIZATION)

A biological service organization specializing in the importation of insects for biological control. We have a long history of successful work in this field and are now expanding our services to include the importation of Tribolium species.

For more information on our services and the Tribolium species mentioned above, please contact me by fax or e-mail. We are currently accepting orders for these species and will be shipping them in the near future.

June 1999

Tribolium News Exchange - ELECTRONIC MAIL ADDRESSES.

The following includes an updated list of people subscribed to the Tribolium News Exchange. The purpose of this group is to provide an informal forum to exchange ideas, techniques and suggestions about Tribolium. To send electronic mail to the group, address it to: tribolium@emerald.tufts.edu

If you have suggestions of other people not included on this list who you think would be interested in participating, please send their addresses to me.

Happy beetling,

Margaret Bloch Qazi

People on the Tribolium News Exchange E-mail List:

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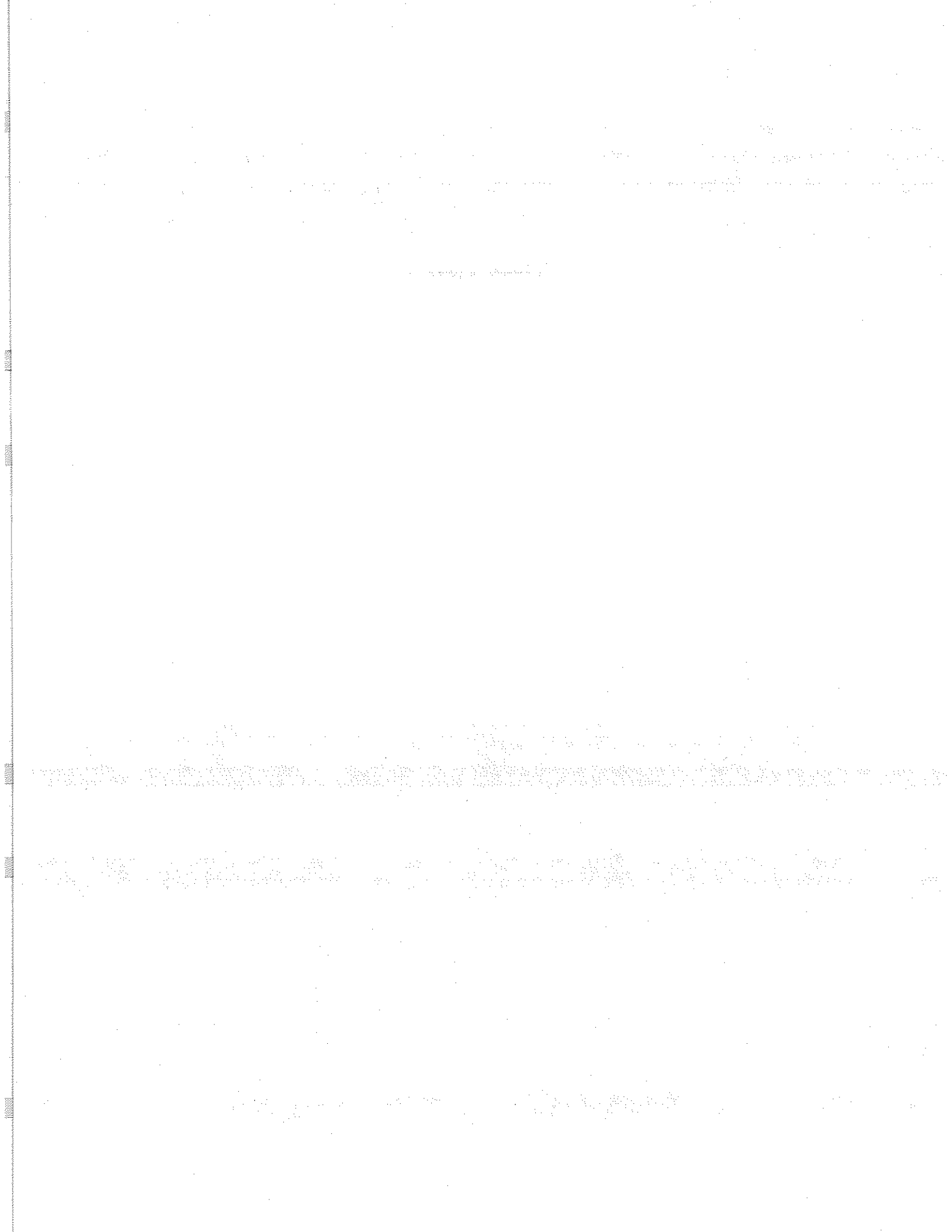
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STOCK LISTS



RIVER FOREST, ILLINOIS
ROSARY COLLEGE
DEPARTMENT OF NATURAL SCIENCES

I. Wild type strains

A. Tribolium castaneum

1. "Chicago" (originally from Thomas Park)
2. "Brazil" (originally from Rio de Janeiro; also known as cI)
3. "Arkansas" (originally from Michael Wade)

B. Tribolium confusum

1. "Chicago" (originally from Thomas Park)
2. "Circle" (collected in Chicago)
3. bI (derived from "Chicago")
4. bII (derived from "Chicago")
5. bIII (derived from "Chicago")
6. bIV (derived from "Chicago")

II. Mutant strains

A. Tribolium castaneum

1. "Chicago" black (derived from "Chicago")

B. Tribolium confusum

1. "Chicago" black (derived from "Chicago")

David M. Craig

GAINESVILLE, FLORIDA

ARS, USDA

P.O. BOX 14565

INSECT ATTRACTANTS, BEHAVIOR AND BASIC BIOLOGY LABORATORY.

<u>Attagenus megatoma</u>	black carpet beetle
<u>Cadra cautella</u>	almond moth
<u>Cylas formicarius elegantulus</u>	sweet potato weevil
<u>Lasioderma serricorne</u>	cigarette beetle
<u>Oryzaephilus surinamensis</u>	sawtoothed grain beetle
<u>Parameylois 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>	

(Ed.).

KINGSTON, RHODE ISLAND 02881
UNIVERSITY OF RHODE ISLAND
DEPARTMENT OF ZOOLOGY

Tribolium castaneum

Purdue Foundation	via Purdue
Black Foundation	via Purdue
Corn oil unsaturated fatty acid sensitive (cos)	

Tribolium confusum

Chicago	Park 1955
black	via San Bernardino
pearl	via San Bernardino

<u>Tribolium madens</u>	via San Bernardino
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<u>Tribolium brevicornis</u>	via San Bernardino
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(Ed.).

LAFAYETTE, INDIANA 47907
 PURDUE UNIVERSITY
 ANIMAL SCIENCES DEPARTMENT

Tribolium castaneum

I. Wild type strains

A. Foundation "+" - originated in 1954 at Purdue University from a broad genetic base and maintained with no artificial selection and minimal breeding.

B. Foundation s - Same genetic base as Foundation "+", but genetically marked with the sooty mutant (s).

C. Foundation b - Originated in 1959 at Purdue University with a broad genetic base unrelated to Foundation "+", no artificial selection, minimal inbreeding, and genetically marked with the black mutant (b).

D. Foundation p - Originated in 1959 at Purdue University with a broad genetic base unrelated to Foundation "+" and b, no selection, minimal inbreeding, and genetically marked with the pearl mutant (p).

W.M. MUIR

LEXINGTON, KENTUCKY
 UNIVERSITY OF KENTUCKY
 AGRICULTURAL EXPERIMENT STATION

I. Base populations

- | | |
|--------------------------------|--------|
| 1. Purdue + foundation | Purdue |
| 2. Purdue s foundation (sooty) | Purdue |
| 3. Purdue b foundation (black) | Purdue |
| 4. Purdue p foundation (pearl) | Purdue |

II. Synthetic strains -- with a history of long-term selection for increased pupa weight but maintained in population cages without selection pressure but discrete generations.

- | | |
|----------|-----------------|
| 1. MRS-1 | Minnesota, 1970 |
| 2. MRS-2 | Minnesota, 1970 |
| 3. P | Purdue, 1976 |

4. C

Davis, 1976

III. Synthetic strain IS from a cross of CSI-10 X EI inbred lines, maintained in population cages with extremely large
 1. IS - From a cross of CSI-10 X ei inbred lines, maintained in population cages with extremely large population size and random mating for 28 generations.

(Ed.).

MANHATTAN, KANSAS
 KANSAS STATE UNIVERSITY
 DEPARTMENT OF ENTOMOLOGY

LEPIDOPTERA

Phycitidae: Cadra cautella and Plodia interpunctella

Gelechiidae: wild and red eyed strains.

Pyralidae: Corcyra cephalonica

COLEOPTERA

Anobiidae: Lasioderma serricorne and Stegobium paniceumRostrichidae: Rhyzopertha dominicaBruchidae: Callosobruchus maculatusCucujidae: Cryptolestes ferrugineus, C. pusillus,Curculionidae: Sitophilus granarius, S. oryzae, and two strains of S. zeamais.Dermeestidae: Trogoderma inclusum, Attagenus megatomaOstomatidae: Tenebroides mauritanicusPtinidae: Gibbium psyllodesSilvanidae: Ahasverus advena, Oryzaephilus surinamensis, D. mercator

Tenebrionidae:

Palorus ratzeburgi, Kansas 1965

Tenebrio molitor, Kansas

Tenebrio obscurus Manhattan, Kansas, 1971

Tribolium castaneum, Kansas

Tribolium confusum, Kansas

Valerie Wright

MANHATTAN, KANSAS 66502

U.S. GRAIN MARKETING RESEARCH LABORATORY

Tribolium castaneum

I. Insecticide-resitant strains

1. 6A-1, malathion-specific, collected in Georgia, 1980
2. NC-1, malathion-specific, collected in North Carolina. From W.C. CAMPBELL.
3. Kano, malathion-specific, collected in northern Nigeria, 1961. From W.R. Wilkin.
4. CTC 12, nonspecific, oxidase type, collected in Kingaroy, Australia, 1968. From W.R. Wilkin.
5. TC 95, nonspecific. From B.R. Champ.
6. DDT C, DDT-resistant, collected in South Africa, 1959. From D.G. Blackman.
7. Rmal-2 allelic to Rmal-1
8. Kdiel--Resistant to lindane, dieldrin and other cyclodienes, linkage group not determined.

II. Mutant strains

(see next pages)

Tribolium castaneum:

Mutant / Strain	Mutant Origin	Full Name or description	Linkage		Stock Origin
			Group	Stocks	
35-17	Manhattan	dominant lethal	2	35-17/Ag4,Stm	Manhattan
3P1	Purdue	crossover suppressor	3	3P1/au14	Purdue
3P2	Purdue	crossover suppressor	3	3P2/au14	Purdue
A(Ag1),Stm	Manhattan	abdominal (from Ag), cis Stm	2	A(Ag1),Stm /ptID60	Manhattan
A(Ag2)	Manhattan	abdominal (from Ag)	2	A(Ag2)/ Ag4,Stm	Manhattan
A(mc)	Manhattan	abdominal (from mc)	2	A(mc),p/Strm,Cx5	Manhattan
A10	Manhattan	Abdominal 10	2	A10 / Ey	Manhattan
A10,mxpA10	Manhattan	Abdominal 10, mxp fr. A10	2	A10,mxpA10/Ag5,Stm	Manhattan
A12	Manhattan	Abdominal 12	2	A12/Ey	Manhattan
A14,Ey	Manhattan	Abdominal 14, Stm cis	2	A14,Ey / Ag4,Stm	Manhattan
A15, Stm	Manhattan	Abdominal 15, Stm cis	2	A15,Stm/Ey	Manhattan
A20 Rdiel	Unknown	Dieldrin resistant	2	A20 Rdiel	Unknown
A4	Manhattan	Abdominal 4	2	A4/Strm,Ag5	Manhattan
A8	Manhattan	Abdominal 8	2	A8/Strm,Cx5	Manhattan
A83	Manhattan	Abdominal 83	2	A83/Strm	Manhattan
ab	Bogota	antenna bifurcada	9	ab,pas30,p	Manhattan
ab	Bogota	antenna bifurcada	9	ab/ab	Bogota, Colombia
ab (IN20)	Parkside	inbred line, 20 generations, from ab	?	ab(IN20)/ab(IN20)	U.Wisc., Parkside
Abidjan	Ivory Coast	wild-type strain	-	Abidjan	Ivory Coast, 19??
AD100,Stm,Cx5	Manhattan	Notched gena,Stm,Cx5 (cis)	2	AD100,Stm,Cx5/Es1	Manhattan
Ag	Manhattan	Antennagalea	2	Ag/Es1	Manhattan
Ag	Manhattan	Antennagalea	2	Ag/Stb	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/35-17	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/A ⁹²	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/SK14	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/Es1	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/sp	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/vve	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/X(ab-1s)	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/X-31	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/X-47	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (cis)	2	Ag4,Stm/X-83	Manhattan
Ag4, Stm	Manhattan	Antennagalea 4, Stm (revertant)	2	Ag4,Stm ^R /Es1	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	Ag5,Stm/A4	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	A10,mxpA10/Ag5,Stm	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	Ag5,Stm/A14,Ey	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	Ag5,Stm/Es1	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	Ag5,Stm/Es2	Manhattan
Ag5, Stm	Manhattan	Antennagalea 5, Stm (cis)	2	Ag5,Stm/GoP14	Manhattan
AgPin	Manhattan	Antennagalea (Pinhead)	2	AgPin/Strm,Cx5	Manhattan
Ahd	Purdue	Arrowhead	8	Ahd/Ahd	Purdue
ap	Englert	antennapedia	8	ap, b	Manhattan
ap	Englert	antennapedia	8	ap, sq2	Manhattan
ap	Englert	antennapedia	8	ap,sq/ap,sq,Bald	Manhattan
ap	Englert	antennapedia	8	MMS (s,rb,ap,au,mas)	Manhattan
Api	Manhattan	Antennapalpus	2	Api,apt,mas,pas	Manhattan
Api	Manhattan	Antennapalpus	2	Api/Api	Manhattan
apt	Sokoloff & Hoy	alate prothorax	2	apt, pas	San Bernardino
apt	Sokoloff & Hoy	alate prothorax	2	b, apt, sa, c	Manhattan
apt	Sokoloff & Hoy	alate prothorax	2	Quint(mxp,apt,mas,pas,ub)	Manhattan
au	Hoy	aureate	3	b(t),p,lod,au,msg	Manhattan
au	Hoy	aureate	3	au,lod,isoline (JS)	Purdue
au	Hoy	aureate	3	au, lod, p	San Bernardino
au	Hoy	aureate	3	mas, p,au	Manhattan
au	Hoy	aureate	3	MMS (s,rb,ap,au,mas)	Manhattan
au ¹⁴	Purdue	aureate 14, lethal	3	3P1/au ¹⁴	Purdue
au ¹⁴	Purdue	aureate 14, lethal	3	3P2/au ¹⁴	Purdue
au ¹⁴	Purdue	aureate 14, lethal	3	3.2 Bamp/au ¹⁴	Purdue
au ²	Manhattan	aureate	3	au ²	Manhattan
b	Sokoloff	black body color	3	b	San Bernardino
b	Sokoloff	black body color	3	b, ap	Manhattan
b	Sokoloff	black body color	3	b, apt, sa, c	Manhattan
b(i-2)	Purdue	black body color	3	b(i-2)	Purdue
b(M)	Purdue	black body color	3	b(M)	Purdue
b(New)	Manhattan	black, dominant	3	b(New)/b(ST)	Manhattan
b(ST)	Manhattan	black, dominant	3	b(ST)/Chr	Manhattan
b(ST)	Manhattan	black, dominant	3	b(ST)/b(New)	Manhattan
b(t)	Dyte & Blackman	tawny body color	3	b(t)	San Bernardino
b(t)	Dyte & Blackman	tawny body color	3	b(t),p,lod,au,msg	Manhattan
ba	Manhattan	broken antennae	2	ba, mxp, apt, pas30	Manhattan
Bald	Manhattan	Bald (reduced setiferous pits)	8	Bald	Manhattan
Bald	Manhattan	Bald (reduced setiferous pits)	8	Bald,ap,sq/ap,sq	Manhattan
Bamp27	Manhattan	Blunt anterior metasternal, projection 27	3	Bamp27/+,au/au	Manhattan
Bamp27	Manhattan	Blunt anterior metasternal, projection 27	3	M1/M1,Bamp27/+	Manhattan
Bamp27,au	Manhattan	Blunt anterior metasternal, projection 27, au (cis)	3	Bamp27/+,au/au	Manhattan
Bamp29	Manhattan	Blunt anterior metasternal, projection 29	3	Bamp29/+	Manhattan
Bamp31	Manhattan	Blunt anterior metasternal, projection 31	3	Bamp31/+	Manhattan
Bamp58	Manhattan	Blunt anterior metasternal, projection 58	3	Bamp58/+	Manhattan

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1999

Manhattan, Kansas

Bamp ¹	Purdue	Blunt anterior metasternal: projection J-1	3	Bamp ¹ /+	Purdue
BampSp	Manhattan	Blunt anterior metasternal: projection Sp	3	BampSp/+	Manhattan
Bang-1	Bangladesh	wild-type strain	-	Bang-1	Bangladesh, 1989
Bang-2	Bangladesh	wild-type strain	-	Bang-2	Bangladesh, 1979
Banos	Ecuador	wild-type strain	-	Banos	Ecuador, 19??
Be	Lasley & Sokoloff	Bar eye	4	Be/+	San Bernadino
Be	Lasley & Sokoloff	Bar eye	4	Be/+ s/s	San Bernadino
Berlin	Germany	wild-type strain	-	Berlin	Germany, 19??
bge	Manhattan	bug-eyed	?	bge	Manhattan
Bha-4 (slight sq)	India	wild-type strain	-	Bha-H	India, 1988
Bha-B (squinty)	India	wild-type strain	-	Bha-B	India, 1988
Blakely	Georgia	wild-type strain	-	Blakely	Georgia, 1993
BMT Lab	Beaumont	wild-type strain	-	BMT Lab	Beaumont, 1974?
Bordeaux	France	wild-type strain	-	Bordeaux	France, 19??
box	Manhattan	box (abdominal)	2	box / Es	Manhattan
BRM	Texas	wild-type strain	-	BRM	Texas, 1988
BRZ-4	Brazil	wild-type strain	-	BRZ-4	Brazil, 1987
BRZ-5	Brazil	wild-type strain	-	BRZ-5	Brazil, 1988
BRZ-6	England	wild-type strain	-	BRZ-6	England, 1943
BT-15	Bangladesh	wild-type strain	-	BT-15	Bangladesh, 1981
c	Eddleman	chestnut eye	7	b, apt, sa, c	Manhattan
c	Eddleman	chestnut eye	7	Nppc	San Bernadino
c	Eddleman	chestnut eye	7	sa,c	San Bernadino
Causey-S	S. Carolina	wild-type strain	-	Causey-S	S. Carolina, 1991
chl	Manhattan	confusum-like	?	chl	Manhattan
Cg	Manhattan	Cleft gular (sutures)	?	Cg/+	Manhattan
Chr	Ackermann	Charcoal body color	3	Chr/b(ST)	Manhattan
ChrE	Manhattan	Charcoal (Elytra indented)	3	ChrE/+	Manhattan
co	Manhattan	cola body color	9	co,p	Manhattan
co	Manhattan	cola body color	9	co,Pyr-R	Manhattan
co	Manhattan	cola body color	9	Se,co,p/+,co,p	Manhattan
COL-1	Colombia	wild-type strain	-	COL-1	Colombia, 1987
COL-2	Colombia	wild-type strain	-	COL-2	Colombia, 1989
CR-1	Costa Rica	wild-type strain	-	CR-1	Costa Rica, 19??
Crab	Manhattan	Crab (warped legs)	7	Crab/PL4	Manhattan
CRR1-1	India	wild-type strain	-	CRR1-1	India, 1989
CRR1-2	India	wild-type strain	-	CRR1-2	India, 1983
CTC-4	Australia	wild-type strain	-	CTC-4	Australia, 1965
CTC-485	Australia	wild-type strain	-	CTC-485	Australia, 1988
Cx20	Manhattan	Cephalothorax 20	2	Cx20/Es1	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/A8	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/A(mc),p	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/AgPin	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/Es1	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/Lu	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/Mcs1R5	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/Ski4R3	Manhattan
Cx5,Stm	Manhattan	Cephalothorax 5, Stm (cis)	2	Cx5,Stm/Ski6R1	Manhattan
Cx5,Stm,AD100	Manhattan	Cephalothorax 5, Stm, notched gena	2	Cx5,Stm,AD100/Es1	Manhattan
Cx6	Manhattan	Cephalothorax 6	2	Cx6/Es1	Manhattan
Dch1	Sokoloff	Dachshund 1	2,9	Dch1/Es1	Manhattan
Dch1	Sokoloff	Dachshund 1	2,9	Dch1/Lu	Manhattan
Dch1	Sokoloff	Dachshund 1	2,9	Dch1/Ski6	Manhattan
Dch3	Manhattan	Dachshund 3	2,9	Dch3/ Ey	Manhattan
Dch4	Manhattan	Dachshund 4	2	Dch4 / Es	Manhattan
Det43	Manhattan	Divergent elytral tips	4,5	Det43/+	Manhattan
Df(Dch1)	Manhattan	Deficiency (from Dch1)	2	Df(Dch1)/Ey	Manhattan
Df1-3/Ey	Manhattan	Deficiency	2	Df1-3/Dp/Es1	Manhattan
Df1-3/Ey	Manhattan	Deficiency	2	Df1-3/Ey	Manhattan
Df1-5/Ey	Manhattan	Deficiency	2	Df1-5/Ey	Manhattan
dms	Manhattan	distorted metasternal suture	?	dms	Manhattan
Dp	Manhattan	Duplication (from Dch1)	2	Dp/Es1/Df(Dch)	Manhattan
Dp	Manhattan	Duplication (from Dch1)	2	Dp/Es1/Df1-3	Manhattan
Dp	Manhattan	Duplication (from Dch1)	2	Dp/Es1/pas30	Manhattan
Dp	Manhattan	Duplication (from Dch1)	2	Dp/Ey/Ey	Manhattan
DpLu	Manhattan	Duplication (from Lu)	2	DpLu/Ey	Manhattan
DpSpa	Manhattan	Duplication (from Spa)	2	DpSpa/Es1/pas30	Manhattan
Ds	Oregon State U.	Displaced sternellum	4	Ds/+	Manhattan
ds(euD)	Manhattan	displaced sternellum (from euD)	?	ds(euD)	Manhattan
Ds(New)	Manhattan	Displaced sternellum	?	Ds(New)/+	Manhattan
ds-X	Manhattan	displaced sternellum, x-linked	4?X	ds-X	Manhattan
Dwi-1	India	wild-type strain	-	Dwi-1	India, 1989
Dwi-1 #189	India	wild-type strain	-	Dwi-1 #189	India
Dwi-3 (dark body color)	India	wild-type strain	-	Dwi-3 (dark body color)	India, 1989
Dwi-3 #191	India	wild-type strain	-	Dwi-3	India
Dwi-3 isoline (ST)	India	inbred isoline	-	Dwi-3 isoline (ST)	India/Manhattan
Em.A16s	Manhattan	Enlarged mentum, abdominal (cis)	2	Em.A16s/Stb	Manhattan
Er	Manhattan	Eye reduced	2	Er/mxpD1,Ski6	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	AD100,Stm,Cx5/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ag+RplD1/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ag/Es1	Manhattan

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Manhattan, Kansas

Es	Manhattan	Extra sclerite (abdominal)	2,4	Ag4, Stm/Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ag5, Stm/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR102-1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR2	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR2a-2	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR3a-1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR4a(Dp)/Ey	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR5a	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR6a-1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR6a-2	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/AR8a	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/ARA3	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/ARA4	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	box / Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Es1/Cx5, Stm	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Cx6/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Cx20/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Dch1/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Dch3/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Det43/Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Dp/Es1/Df(Dch)	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Dp/Es1/Df1-3	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Dp/Es1/pas30	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	DpSpa/Es1/pas30	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	DpLu (Es1, Sk16)/Ey	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ey/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	g/Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	GoPL6/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	GoPL10/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	GoPL11/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	GoPL14/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Hw/mxpX9, Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ip69/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	LuR1a/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Mc-2, Ubx1/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	mxp8/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	mxp19/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	mxp170/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	mxpX9, Es1/Ey	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ns, Stm/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	ptD16, Stm/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	ptD57, Stm/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Spa/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Stb/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Stbd/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Stm+RSplID/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Stm-Es1/+NDJ	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	StmR1/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	StmR2/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	StmR5/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	StmR6/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	StmR, Ag4/Es1	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	tr/Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ubx1/Es	Manhattan
Es	Manhattan	Extra sclerite (abdominal)	2,4	Ubx2, Stm/Es	Manhattan
Es(Sk16)	Manhattan	Extra sclerite (from Sk16)	2	Es(Sk16)/+	Manhattan
Es(Sk16)	Manhattan	Extra sclerite (from Sk16)	2	Es(Sk16)GoPL4	Manhattan
Es1+R1	Manhattan	Extra sclerite revertant 1	2	Es1+R1/Ey	Manhattan
Es1+R9	Manhattan	Extra sclerite revertant 9	2	Es1+R9/Ey	Manhattan
Es2	Manhattan	Extra sclerite 2	2	Es2/Ag5	Manhattan
Estill	S. Carolina	wild-type strain	-	Estill	S. Carolina, 19??
eu	Lasley & Sokoloff	extra urogomphi	2	eu	San Bernadino
eu	Lasley & Sokoloff	extra urogomphi	2	eu, apt, mas	Manhattan
eu	Lasley & Sokoloff	extra urogomphi	2	eu, mas	Manhattan
euD	Manhattan	Extra urogomphi (male sterile)	2	euD/+	Manhattan
Ey	Manhattan	eyeless	2,5	A10 / Ey	Manhattan
Ey	Manhattan	eyeless	2,5	A12/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	A15, Stm/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Dch3 / Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Df(Dch1)/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Df1-3/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Df1-5/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	DpLu/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Ey/Es1	Manhattan
Ey	Manhattan	eyeless	2,5	LuR1a/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	LuRpII/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Mcs1R1/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	Mcs1R2/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	mxpD1, Sk16/Ey	Manhattan
Ey	Manhattan	eyeless	2,5	mxpX9, Es1/Ey	Manhattan

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Ey	Manhattan	eyeless	2.5	ptfD60/Ey	Manhattan
Ey	Manhattan	eyeless	2.5	SkI4R2/Ey	Manhattan
Ey A14	Manhattan	Eyeless, Abdominal 14 (cis)	2	Ag5,Stm/Ey,A14	Manhattan
Ey-Lethal-Free	Manhattan	lethal free from Eyeless	NA	Ey-Lethal-Free	Manhattan
fa	Manhattan	fused antennae	5?	fa	Manhattan
fe	Manhattan	folded elytra	?	fe	Manhattan
Fl	Purdue	fused funnicle	6	Fl/+	Purdue
FFM-C	Georgia	wild-type strain	-	FFM-C	Georgia, 1993
fs(sa)	Manhattan	short antennae, female sterile	?	fs(sa)	Manhattan
FSS2	England	wild-type strain	-	FSS2	England, 1943
Fta	Sokoloff & St. Hilaire	Fused tarsi and antennae	?	Fta/+	San Bernadino
g	Manhattan	glossy	2	g/Es	Manhattan
g	Manhattan	glossy	2	g,pas30	Manhattan
Ga-1	Georgia	Georgia 1, wild type	NA	Ga-1	Georgia
Ga-2	Georgia	Georgia 2, Ga-1 inbred 20 generations	NA	Ga-2	U.Wisc.-Parkside
Ga-2 iso-M1	Georgia	Georgia 2, isolate to M1/M1	NA	Ga-2 iso-M1	U.Wisc.-Parkside
Ga-9s	Georgia	Georgia back-X to s 9X, sel. For Rmai gene	4	Ga-9s	Manhattan
Ger-1	Germany	wild-type strain	-	Ger-1	Germany, <1989
Gi	Sokoloff & Brownlee	Giant (body size)	NA	Gi/Gi	San Bernadino
Go	Manhattan	Goliath (body size)	7	Go/H	Manhattan
Go	Manhattan	Goliath (body size)	7	Go/+_c/c	Manhattan
Go	Manhattan	Goliath (body size)	7	Go/+_b,sa,c/b,sa,c	Manhattan
GoPL10	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL10/Es1	Manhattan
GoPL11	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL11/Es1	Manhattan
GoPL14	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL14/Es1	Manhattan
GoPL4	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL4/Ag5,Stm	Manhattan
GoPL4	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL4/Crab	Manhattan
GoPL4	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL4/Es(Skl6)	Manhattan
GoPL6	Manhattan	Goliath-derived crossover suppressor	7.2	GoPL6/Es1	Manhattan
GW-13	Australia	wild-type strain	-	GW-13	Australia, 1977
GW-3	Australia	wild-type strain	-	GW-3	Australia, 1988
GW-4	Australia	wild-type strain	-	GW-4	Australia, 1965
h	Dawson	hazel eye	4	h, s	San Bernadino
H-1 (ST)	Parkside	H-factor	?	H-1	U.Wisc.-Parkside
H-2 (ST)	Parkside	H-factor	?	H-2	U.Wisc.-Parkside
Heng-5	Thailand	wild-type strain	-	Heng-5	Thailand, 1989
HO-TCS	Singapore	wild-type strain	-	HO-TCS	Singapore, 1989
HO-TJC, #121	Singapore	wild-type	-	HO-TJC	Singapore
Hw	Manhattan	Hairy wing	2	Hw/Es,mxPX9	Manhattan
I	Bartlett	ivory (eye color)	?	i,lod	San Bernadino
Ibad-2cf	Nigeria	wild-type strain	-	Ibad-2cf	Nigeria, 1987
Is	Manhattan	Incomplete sternellum	?	Is/+	Manhattan
ISR-1	Israel	wild-type strain	-	ISR-1	Israel, 1988
ISR-2	Israel	wild-type strain	-	ISR-2	Israel, 1988
J	Park	jet, body color	5	j,mc	San Bernadino
J	Park	jet, body color	5	mc,rb,j	Manhattan
J	Park	jet, body color	5	rb,j	San Bernadino
J2 (Z-4)	Manhattan	jet, body color	5	j2	Manhattan
Japan #1	Japan	wild-type strain	-	Japan #1	Japan, <1978
Japan #2	Japan	wild-type strain	-	Japan #2	Japan, 1988
Japan #4	Japan	wild-type strain	-	Japan #4	Japan, 1988
ju	Sokoloff	juvenile urogomphi	4?	ju,eu,b	Manhattan
Kent (small eyes)	England	wild-type strain	-	Kent (small eyes)	England, 1977
Lab-S Rusty	Manhattan	Lab strain, rusty, wild-type	NA	Lab-S Rusty	Manhattan
LF-3 (JS)	Purdue	Lethal free strain	3	LF-3 (JS)	Purdue
Little Rock	Arkansas	wild-type strain	-	Little Rock	Arkansas, 1988
lod	Sokoloff	light optical diaphragm	3	au,lod isolate (JS)	Purdue
lod	Sokoloff	light optical diaphragm	3	au,lod,p	San Bernadino
lod	Sokoloff	light optical diaphragm	3	b(t),p,lod,au,msg	Manhattan
lod	Sokoloff	light optical diaphragm	3	i,lod	San Bernadino
lod	Sokoloff	light optical diaphragm	3	lod,p	San Bernadino
lod	Sokoloff	light optical diaphragm	3	M1,au,lod,p	Manhattan
lod	Sokoloff	light optical diaphragm	3	mc(eg),lod,p	Manhattan
lod	Sokoloff	light optical diaphragm	3	Rd(HD),lod,p	Manhattan
lod	Sokoloff	light optical diaphragm	3	Rd,mc,lod,p	Manhattan
ip69	Manhattan	labiopedia 69	2	ip69/Es1	Manhattan
Lu	Manhattan	Lucifer (dorsal head horns)	2	Lu / Stm,Cx5	Manhattan
Lu	Manhattan	Lucifer (dorsal head horns)	2	Lu,SkI6/Stb	Manhattan
Lu	Manhattan	Lucifer (dorsal head horns)	2	Lu, au/Dch1	Manhattan
LuR1a	Manhattan	Lucifer revertant	2	LuR1a/Es1	Manhattan
LuR1a	Manhattan	Lucifer revertant	2	LuR1a/Ey	Manhattan
m.l. 9.14	Manhattan	(Male linked)	2	9.14 (male linked)	Manhattan
M1	Manhattan	Medea 1	3	M1Big III, p	Manhattan
M1	Manhattan	Medea 1	3	M1 - iso 3B1 (G)	Manhattan
M1	Manhattan	Medea 1	3	M1 isolate (JS)	Manhattan
M1	Manhattan	Medea 1	3	M1,au,M3	Manhattan
M1	Manhattan	Medea 1	3	M1,au,p,lod	Manhattan
M1	Manhattan	Medea 1	3	M1,b	Manhattan
M1	Manhattan	Medea 1	3	M1/M1, Bamp27/+	Manhattan
M3	Manhattan	Medea 3	3	M3,au	Manhattan

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M3	Manhattan	Medea 3	3	M1,au,M3	Manhattan
mas	Hoy & Sokoloff	missing abdominal sternite	2	mas	San Bernadino
mas	Hoy & Sokoloff	missing abdominal sternite	2	mas, p,au	Manhattan
mas	Hoy & Sokoloff	missing abdominal sternite	2	ptl, mas, pas	Manhattan
mas	Hoy & Sokoloff	missing abdominal sternite	2	Quint(mxp, apt, mas, pas, ub)	Manhattan
mas	Hoy & Sokoloff	missing abdominal sternite	2	MMS (s,rb,mas,ap,au)	Manhattan
mas2	Manhattan	missing abdominal sternite 2	2?	mas2	Manhattan
mc	Sokoloff & Lasley	microcephalic	5	j,mc	San Bernadino
mc	Sokoloff & Lasley	microcephalic	5	mc,rb,j	Manhattan
mc(eg)	Sokoloff & Lasley	microcephalic (eye growth variant)	5	mc(eg),p,lod	Manhattan
mc(eg)	Sokoloff & Lasley	microcephalic (eye growth variant)	5	Rd,mc(eg),lod,p	Manhattan
Mc-2,Ubx1	Manhattan	Microcephalic-2,Ultrathorax(cis)	2	Mc-2,Ubx1/Es1	Manhattan
Mcs1	Manhattan	Miscadestral sclerite	2	Mcs1/Strn	Manhattan
Mcs1R1	Manhattan	Miscadestral sclerite, revertant 1	2	Mcs1R1/Ey	Manhattan
Mcs1R2	Manhattan	Miscadestral sclerite, revertant 2	2	Mcs1R2/Ey	Manhattan
Mcs1R4	Manhattan	Miscadestral sclerite, revertant 4	2	Mcs1R4/mxpNG	Manhattan
Mcs1R5	Manhattan	Miscadestral sclerite, revertant 5	2	Mcs1R5/Strn,Cx5	Manhattan
Mek-1	China	wild-type strain	-	Mek-1	China, 1987
Mo	Sokoloff	Microphthalmic	6	Mo/+	San Bernadino
Montreal	Montreal	wild-type strain	-	Montreal	Montreal, 1973
mrg	Sokoloff & Hoy	melanotic stink gland	7	b(t),p,lod,au,msg	Manhattan
mxp	Sokoloff	maxillopedia	2	ba, mxp, apt, pas30	Manhattan
mxp	Sokoloff	maxillopedia	2	mxp, apt, pas30	Manhattan
mxp	Sokoloff	maxillopedia	2	Ubx1, mxp, apt/A10, mxpA10	Manhattan
mxp	Sokoloff	maxillopedia	2	Quint(mxp, apt, mas, pas, ub)	Manhattan
mxp170	Manhattan	maxillopedia 170, lethal	2	mxp170/Es1	Manhattan
mxp19	Manhattan	maxillopedia 19, lethal	2	mxp19/Es1	Manhattan
mxp8	Manhattan	maxillopedia 8, lethal	2	mxp8/Es1	Manhattan
mxpD1,Sk16/Ey	Manhattan	Maxillopedia, dom. 1, Sk16 (cis)	2	mxpD1,Sk16/Ey	Manhattan
mxpNG	Manhattan	maxillopedia, Notched Gena, lethal	2	mxpNG/Es1	Manhattan
mxpNG	Manhattan	maxillopedia, Notched gena	2	Mcs1R4/mxpNG	Manhattan
mxpX9, Es	Manhattan	lethal maxillopedia, Es (cis)	2,4	mxpX9,Es1/Ey	Manhattan
NDG-2 (#59)	Manitoba	Wild-type	NA	NDG-2 (#59)	Manitoba
NDJ-11	Hawaii	wild-type strain	-	NDG-11	Hawaii, 1976
NDJ-13	Vancouver	wild-type strain	-	NDG-13	Vancouver, 1976
NDJ-3	Manitoba	wild-type strain	-	NDG-3	Manitoba, 1987
NDJ-6 (some white eye)	Minnesota	wild-type strain	-	NDJ-6 (some white eye)	Minnesota, 1982
NIG-1 (red eye)	Nigeria	wild-type strain	-	NIG-1 (red eye)	Nigeria, 1988
Npp	Hoy	Non-punctate prothorax	?	Nppc (Sokl 428)	San Bernadino
p	Park	pearl eye	9	p	San Bernadino
p	Park	pearl eye	9	ab,pas30,p	San Bernadino
p	Park	pearl eye	9	au, lod, p	San Bernadino
p	Park	pearl eye	9	b(t),au,lod, p,msg	Manhattan
p	Park	pearl eye	9	co,p	Manhattan
p	Park	pearl eye	9	lod, p	San Bernadino
p	Park	pearl eye	9	M1Big III, p	Manhattan
p	Park	pearl eye	9	mas, p,au	Manhattan
p	Park	pearl eye	9	mc(eg),lod, p	Manhattan
p	Park	pearl eye	9	pas30, p	Manhattan
p	Park	pearl eye	9	Rd,mc(eg),p	Manhattan
p	Park	pearl eye	9	Rd,mc(eg),lod, p	Manhattan
p	Park	pearl eye	9	Rd(HD),lod, p	Manhattan
p	Park	pearl eye	9	Se,co,p/+ co,p	Manhattan
p	Park	pearl eye	9	Se,p/+ p	Manhattan
PAK-1	Pakistan	wild-type strain	-	PAK-1	Pakistan, 1979
PAK-2 (dark body color)	Pakistan	wild-type strain	-	PAK-2 (dark body color)	Pakistan, 1979
PAK-3	Pakistan	wild-type strain	-	PAK-3	Pakistan, 1988
pas	Sokoloff	pointed abdominal sternite	2	apt, pas	San Bernadino
pas	Sokoloff	pointed abdominal sternite	2	ptl, mas, pas	Manhattan
pas	Sokoloff	pointed abdominal sternite	2	Quint(mxp, apt, mas, pas, ub)	Manhattan
pas30	Manhattan	pointed abdominal sternite 30	2	ab, pas30, p	Manhattan
pas30	Manhattan	pointed abdominal sternite 30	2	ba, mxp, apt, pas30	Manhattan
pas30	Manhattan	pointed abdominal sternite 30	2	mxp, apt, pas30	Manhattan
pas30	Manhattan	pointed abdominal sternite 30	2	ub, pas30	Manhattan
pd	Park & Frank	paddle antenna	X	py, pd, pit	San Bernadino
pep	Manhattan	peppered cuticle	X	pep	Manhattan
pnk (NDG-2)	Manhattan	pink eye, from NDG-2	?	pnk (NDG-2)	Manhattan
pnk (Tiw-1 iso-43)	Manhattan	pink eye, from Tiw-1 iso-43	?	pnk (Tiw-1 iso-43)	Manhattan
PRC-Nanj	China	wild-type strain	-	PRC-Nanj	China, 1989
PRC-Ning	China	wild-type strain	-	PRC-Ning	China, 1989
Pruz +	Poland	wild-type strain	-	Pruz +	Poland, 1988
Pruz-1	Poland	wild-type strain	-	Pruz-1	Poland, 1963
Ps	Manhattan	Pinched sternellum	2	Ps/Rd(CS)	Manhattan
PS-129	India	wild-type strain	-	PS-129	India, 1984
pte	Sokoloff	platinum eye	X	py, pd, pit	San Bernadino
ptl	Lasley & Sokoloff	prothoraxless	2	ptl	San Bernadino
ptl	Lasley & Sokoloff	prothoraxless	2	ptl, mas, pas	Manhattan
ptl(Rd)	Manhattan	prothoraxless from Rd stock	2	ptl(Rd)	Manhattan
ptD16,Strn	Manhattan	Dom. prothoraxless 16, Strn (cis)	2	ptD16,Strn/Es1	Manhattan
ptD2	Manhattan	Dom. prothoraxless 2	2	ptD2/Stb	Manhattan
ptD26Y,Strn	Manhattan	Dom. prothoraxless 26, Y-linked	2,Y	ptD26Y,Strn/+	Manhattan

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ptD57, Strm	Manhattan	Dom. prothoraxless 57, Strm (cis)	2	ptD57, Strm/Es1	Manhattan
ptD60	Manhattan	dominant prothoraxless 60	2	A(Ag1), Strm /ptD60	Manhattan
ptD60	Manhattan	dominant prothoraxless 60	2	ptD60/Ey	Manhattan
py	Lasley	pygmy	X	py, pd, plt	San Bernadino
py2	Manhattan	pygmy 2	X	py2	Manhattan
Pyr-R	Peter Collins	Pyrethroid resistant	9	co, Pyr-R	Peter Collins
QTC 279 (Pyr-R)	Peter Collins	Pyrethroid resistant	9?	QTC 279 (Pyr-R)	Peter Collins
Raj-1	India	wild-type strain	-	Raj-1	India, 1<1979
Ram-B	India	wild-type strain	-	Ram-B	India, 19??
Ramsey (MT '88)	Minnesota	wild-type strain	-	Ramsey (MT '88)	Minnesota, 1988
Rap	Manhattan	Recurved anterior pronotum	2	Rap	Manhattan
rb	Deweese	ruby eye	5	mc, rb, j	Manhattan
rb	Deweese	ruby eye	5	MMS (s,rb,ap,au,mas)	Manhattan
rb	Deweese	ruby eye	5	rb, j	San Bernadino
Rd	Dawson	Reindeer, homozygous viable	2	Rd	San Bernadino
Rd	Dawson	Reindeer, homozygous viable	2	Rd, mas, p	Manhattan
Rd	Dawson	Reindeer, homozygous viable	2	Rd, mc, p	Manhattan
Rd	Dawson	Reindeer, homozygous viable	2	Rd, pas30	Manhattan
Rd(CS)	Manhattan	Reindeer, crossover suppressor	2	Ps/Rd(CS)	Manhattan
Rd(HD)	Manhattan	Reindeer (honey-dipper style)	2	Rd(HD)	Manhattan
Rd(HD)	Manhattan	Reindeer (honey-dipper style)	2	Rd(HD), lod, p	Manhattan
Rdiei BC9 Lab-S	Unknown	Dieldrin resistant from Lab-S	NA	Rdiei BC9 Lab-S	Unknown
REJ-1	Philippines	wild-type strain	-	REJ-1	Philippines, 19??
RINI-3	India	wild-type strain	-	RINI-3	India, 1989
RINI-4	India	wild-type strain	-	RINI-4	India, 19??
s	Bartlett, Bell & Shideler	sooty	4	s	San Bernadino
s	Bartlett, Bell & Shideler	sooty	4	h, s	San Bernadino
s	Bartlett, Bell & Shideler	sooty	4	Be, s	San Bernadino
s	Bartlett, Bell & Shideler	sooty	4	Ga-9s	Georgia, 1993
s	Bartlett, Bell & Shideler	sooty	4	MMS (s,rb,ap,au,mas)	Manhattan
sa	Sokoloff	short antenna	?	b, apt, sa, c	Manhattan
sa	Sokoloff	short antenna	?	Go, b, sa, c	Manhattan
sa	Sokoloff	short antenna	?	sa, c	San Bernadino
Sa-8	Manhattan	Short antenna-8	?	Sa-8	Manhattan
sa-X	Manhattan	short antenna, X-linked	X	sa-X	Manhattan
Se	Manhattan	Short elytra	9	Se	Manhattan
Se	Manhattan	Short elytra	9	Se, co, p	Manhattan
Se	Manhattan	Short elytra	9	Se, p	Manhattan
se 46	Purdue	short elytra 46	?	se 46	Purdue
Se12	Purdue	Short elytra 12	?	Se12	Purdue
Se-2	Manhattan	Short elytra 2	8	Se-2	Manhattan
Shellman	Georgia	wild-type strain	-	Shellman	Georgia, 1993
Sk12s	Manhattan	Socketless spontaneous 2	2	Sk12s/Stb	Manhattan
Sk14	Manhattan	Socketless 4	2	Sk14/Ag4, Strm	Manhattan
Sk14R2	Manhattan	Socketless 4, revertant 2	2	Sk14R2/Ey	Manhattan
Sk14R3	Manhattan	Socketless 4, revertant 3	2	Sk14R3/Strm, Cx5	Manhattan
Sk16	Manhattan	Socketless 6	2	Sk16/Strm, Cx5	Manhattan
Sk16R1	Manhattan	Socketless 6, revertant 1	2	Sk16R1/Strm, Cx5	Manhattan
small	Purdue	small body size	?	small	Purdue
Sok 16	California	wild-type strain	-	Sok 16, (TC16, Veracruz)	California, 19??
Sok 19	California	wild-type strain	-	Sok 19, (TC19, Berkeley)	California, 19??
Sok 22	California	wild-type strain	-	Sok 22, (TC22)	California, 19??
Sok 25	California	wild-type strain	-	Sok 25, (TC25, ex NY)	California, 19??
Sok 4	California	wild-type strain	-	Sok 4, (TC4, Davis)	California, 19??
Sok 8	California	wild-type strain	-	Sok 8, (TC8, McGill)	California, 19??
Solet	Israel	wild-type strain	-	Solet	Israel, 1979
sp	Manhattan	shoulder pads	2	sp/Strm, Ag4	Manhattan
Spa	Sokoloff & Hoy	Spatulate antennae	2,4	Spa/Es1	Manhattan
sq (Tiw-1)	India	squint (from Tiw-1)	?	sq (Tiw-1)	India
sq(euD)	Manhattan	squint (from euD)	?	sq(euD)	Manhattan
sq	Bywaters	squint eye	8	Bald, ap, sq/ap, sq	Manhattan
sq	Bywaters	squint eye	8	sq	San Bernadino
sq2	Manhattan	squint eye 2	8	ap, sq2	Manhattan
sq-B	Burma	squint (from Burma)	?	sq-B	Burma
Stb	Manhattan	Stubby antennae	2,X	Ag/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	Em, A16s/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	Ey, pasN/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	Es/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	Lu, Sk16/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	ptD2/Stb	Manhattan
Stb	Manhattan	Stubby antennae	2,X	Sk12s/Stb	Manhattan
Stbd	Manhattan	Stubboid (short antennae)	2	Lu/Stbd	Manhattan
Stbd	Manhattan	Stubboid (short antennae)	2	Stbd/Es	Manhattan
Strm	Manhattan	Stumpy	2	Strm/Strm	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	X-83/Strm, Ag4	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	X-47/Strm, Ag4	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	vve/Strm, Ag4	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	sp/Strm, Ag4	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	g/Strm, Ag4	Manhattan
Strm, Ag4	Manhattan	Strm, Antennagalea 4	2	X-31/Strm, Ag4	Manhattan

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Stm, Ag5	Manhattan	Stm, Antennagalea 5	2	A4/Stm, Ag5	Manhattan
Stm, Ag5	Manhattan	Stm, Antennagalea 5	2	A10, mxpA10/Stm, Ag5	Manhattan
Stm, Ag5	Manhattan	Stm, Antennagalea 5	2	A14, Ey/Stm, Ag5	Manhattan
Stm, Ag5	Manhattan	Stm, Antennagalea 5	2	Es2/Stm, Ag5	Manhattan
Stm, Ag5	Manhattan	Stm, Antennagalea 5	2	GoPL4/Stm, Ag5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	A8/Stm, Cx5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	AgPin/Stm, Cx5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	Lu / Stm, Cx5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	AD100, Stm, Cx5/Es1	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	Sk4R3/Stm, Cx5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	Sk6R1/Stm, Cx5	Manhattan
Stm, Cx5	Manhattan	Stm, Cephalothorax 5, cis	2	Stm, Cx5/Es1	Manhattan
Stm, Ns	Manhattan	Stm, Narrow sternellum (cis)	2	Stm, Ns/Es1	Manhattan
Stm+RSptID	Manhattan	Stm spontaneous revertant, pti (dominant)	2	Stm+RSptID/Es1	Manhattan
Stm-Es1/+NDJ	Manhattan	Non-disjunction	?	Stm-Es1/+NDJ	Manhattan
StmR1	Manhattan	Stm revertant 1	2	StmR1/Es1	Manhattan
StmR2	Manhattan	Stm revertant 2	2	StmR2/Es1	Manhattan
StmR5	Manhattan	Stm revertant 5	2	StmR5/Es1	Manhattan
StmR6	Manhattan	Stm revertant 6	2	StmR6/Es1	Manhattan
Stm-Ski6/+NDJ	Manhattan	Non-disjunction	?	Stm-Ski6/+NDJ	Manhattan
Sylvania	Sylvania, GA	wild-type strain	-	Sylvania	Sylvania, GA
T(Y,3)	Manhattan	Translocation Y-3	Y,3	T(Y,3)	Manhattan
T(Y,4)	Manhattan	Translocation Y-4	Y,4	T(Y,4)	Manhattan
tar	Manhattan	anterior melanotic stink glands	2	tar	Manhattan
tib	Manhattan	tibialess (from ab)	9?	tib	Manhattan
Tiw-1	India	?	NA	Tiw-1	India
Tiw-1 (iso 43)	India	Tiw-1 isolate	NA	Tiw-1 (iso 43)	India
Tiw-1(iso 43) pink	India	pink eye from Tiw-1	NA	Tiw-1(iso 43) pink	India
Tiw-5	India	wild-type strain	-	Tiw-5	India, 1989
Tiw-6	India	wild-type strain	-	Tiw-6	India, 1989
tr	Manhattan	trembler, recessive lethal	2,4	Es/tr	Manhattan
tr	Manhattan	trembler, homozygous viable	2,4	tr	Manhattan
ub	Manhattan	unbuckled	2	ub	Manhattan
ub	Manhattan	unbuckled	2	ub.pas30	Manhattan
ub	Manhattan	unbuckled	2	Quint(ub, mxp, apt, mas, pas)	Manhattan
ue	Manhattan	unsclerotized elytra	?	ue	Manhattan
Ug-1	Uganda	wild-type strain	-	Ug-1	Uganda, 1989
UG-3	Tanzania	wild-type strain	-	UG-3	Tanzania, 1986
Utx(New)	Manhattan	Ultrathorax (New)	2	Utx(New)/+	Manhattan
Utx1	Manhattan	Ultrathorax	2	Utx1/Es	Manhattan
Utx1	Manhattan	Ultrathorax	2	Utx1/Utx1	Manhattan
Utx2, Stm	Manhattan	Ultrathorax 2, Stm (cis)	2	Utx2, Stm/Es1	Manhattan
Vienna (GA '93)	Georgia	wild-type strain	-	Vienna (GA '93)	Georgia, 1993
vwe	Manhattan	vestigial wings and elytra	2	vwe/Stm, Ag4	Manhattan
w	Eddleman & Bell	white eye	4	w	San Bernadino
Waunakee (WI '92)	Wisconsin	wild-type strain	-	Waunakee (WI '92)	Wisconsin, 1992
WI-1	Wisconsin	wild-type strain	-	WI-1	Wisconsin?
X(ab-1s)	Manhattan	Lethal revertant from ab	9	X(ab-1s)/Ag4, Stm	Manhattan
X-31	Manhattan	lethal 31	2	X-31/Ag4, Stm	Manhattan
X-47	Manhattan	lethal 47	2	X-47/Stm, Ag4	Manhattan
X-83	Manhattan	Lethal 83	2	X-83/Stm, Ag4	Manhattan
Z-1	Alabama	wild-type strain	-	Z-1	Alabama, 1988
Z-2 (occ. dk. red eye)	Oklahoma	wild-type strain	-	Z-2 (occ. dk. red eye)	Oklahoma, 1988
Z-3 (#30)	Kankakee	wild-type strain	-	Z-3 (#30)	Kankakee, IL
Z-4 (occ. dark body)	Iowa	wild-type strain	-	Z-4 (occ. dark body)	Iowa, 1988
Z-5	Minnesota	wild-type strain	-	Z-5	Minnesota, 1988
Z-7	S. Carolina	wild-type strain	-	Z-7	S. Carolina, 1988

Tribolium confusum:

BA50 - cf	Kansas	Tribolium confusum	-	BA50 - cf	Kansas, ~1986-87
HP70 - cf	Kansas	Tribolium confusum	-	HP70 - cf	Kansas, ~1986-87
MN61 - cf	Kansas	Tribolium confusum	-	MN61 - cf	Kansas, ~1986-87
PAK-3-cf	Pakistan	Tribolium confusum	-	PAK-3-cf	Pakistan, 1988
P-Ning -cf	China	Tribolium confusum	-	P-Ning -cf	China, 1989
T. confusum (apt, mas, st)	San Bernadino	T. cf. (alate prothorax, missing abd. stem, st)	?	T. confusum (apt, mas, st)	San Bernadino
T. confusum (b, au, lod, p)	San Bernadino	T. cf. (black, aureate, light optical diaph., pearl)	?	T. confusum (b, au, lod, p)	San Bernadino
T. confusum (lod, p)	San Bernadino	light optical diaphragm, pearl	?	T. confusum (lod, p)	San Bernadino
T. confusum (PRC)	P. R. China	Tribolium confusum	-	T. confusum (PRC)	P. R. China
ThaiB-cf (tan eye)	Thailand	Tribolium confusum	-	ThaiB-cf (tan eye)	Thailand, 19??
UG-2 cf	Uganda	Tribolium confusum	-	UG-2 cf	Uganda, 1989

Other Species:

Gnathocerus cornutus	?	wild-type strain	-	Gnathocerus cornutus	?
Longheaded flour beetle	?	wild-type strain	-	Longheaded flour beetle	?
T. brevicornis	Manhattan	Tribolium brevicornis	-	T. brevicornis	Manhattan
T. freemani	Japan	Tribolium freemani	-	T. freemani	Japan
T. madag.	Manhattan	Tribolium madag.	-	T. madag.	Manhattan

SAVANNAH, GEORGIA
 STORED-PRODUCT INSECTS RESEARCH AND DEVELOPMENT LABORATORY

I. Wild type strains

A. Lepidoptera

- | | |
|------------------------------------------|------------------------------------------|
| | N.C. |
| 1. <u>Cadra cautella</u> (Walker) | Tifton, Ga. |
| 2. <u>Plodia interpunctella</u> (Hubner) | Modesto, Ca. |
| 3. <u>Sitotroga cerealella</u> (Olivier) | Manhattan, Ka.
Can., and Durham, N.H. |

b. Coleoptera

- | | |
|-----------------------------------------------|----------------------------------|
| 1. <u>Attagenus megatoma</u> (Fab.) | CSMA strains |
| 2. <u>Callosobruchus maculatus</u> (Fab.) | Fresno, ca. |
| 3. <u>Cryptolestes ferrugineus</u> (Stephens) | S. Carolina |
| 4. <u>Lasioderma serricornis</u> (Fab.) | Unknown |
| 5. <u>Oryzaephilus mercator</u> (Fauvel) | Unknown |
| 6. <u>Oryzaephilus surinamensis</u> (L.) | Manhattan, Kan. |
| 7. <u>Rhyzopertha dominica</u> Fab.) | Unknown |
| 8. <u>Sitophilus granarius</u> (L.) | Manhattan, Kan. |
| 9. <u>S. oryzae</u> (L.) | Ark., Calif., Kan., La. |
| 10. <u>S. zeamais</u> Motschulsky | Estill, S.C. |
| 11. <u>Stegobium paniceum</u> (L.) | Madison, Wis. |
| 12. <u>Tribolium castaneum</u> (Herbst) | Unknown |
| 13. <u>Tribolium confusum</u> duVal | Manhattan, Kan. |
| 14. <u>Trogoderma glabrum</u> (Herbst) | Madison, wis.,
Riverside, Ca. |

II. Mutant strains. None

Richard T. Arbogast, Laboratory Director.

South Orange, New Jersey
 Seton Hall University
 Department of Biology

T. castaneum

Wild Type Strains

Seton Hall-1

McGill, via California State

Synthetic Strains

Pearl Foundation, via Purdue University

Black Foundation, via Purdue University

Mutant Strains

Paddle (pd) via Cal State U., S.B.

Ho ho
 Red (R) Via Cal State U., S.B.

White (w) Via Cal State U., S.B.

short antenna (ca) Via Oregon State

Short antenna (Sa) Via Purdue University

Tribolium confusum Via Carolina Biological Supply

Elint Krause

Storrs, CT 06269
University of Biology

Tribolium castaneum

Ga-1

sooty

Charcoal

RR strain from Costantino

Oryzaephilus surinamensis

J.S. Bancroft

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

I. Wild type strains

A. Coleoptera strains

Dermeestidae

<u>Attagenus megatoma</u> (F.)	Madison, Wis., 1975, Savannah, Ga. 1974
<u>Trogoderma variabile</u> Ballion	field collected, Mn. 1972

Cucujidae

<u>Oryzaephilus surinamensis</u> (L)	
<u>Oryzaephilus mercator</u> (Fauvel)	
<u>Cryptolestes pusillus</u> (Schoenherr)	Manhattan Ka. 1967
<u>Cryptolestes ferrugineus</u> (Stephens)	Unknown

Silvanidae

<u>Ahasverus advena</u> Waltl.	Minnesota
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Tenebrionidae

<u>Cyaneus angustus</u> (LeConte)	Winnipeg; Minnesota
<u>Tribolium castaneum</u> (Herbst)	Corvallis, Ore
<u>Tribolium confusum</u> duVal	Unknown
<u>Tenebrio molitor</u>	Carolina Biological, 1984

Anobiidae

<u>Lasioderma serricorne</u> (Fab.)	Savannah, Ga.
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Bostrichidae

<u>Rhizopertha dominica</u> (F.)	Manhattan, Ka.
<u>Prostephanus truncatus</u> (Horn)	Unknown

Curculionidae

<u>Sitophilus granarius</u> (L.)	Unknown
<u>S. oryzae</u> (L.)	"

B. Lepidoptera

Pyralidae

<u>Anagasta kuehniella</u> (Zeller)	Savannah, Ga.
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Gelechiidae

<u>Sitotroga cerealella</u> (Oliver)	Savannah, Ga.
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(Ed.)

St. Paul, Minnesota 55108
University of Minnesota
Department of Entomology
Stored-Grain Pest Management Program

Eight species of stored-product beetles and two species of moths are maintained in the laboratory. These species include: Angoumois grain moth, flat grain beetle, Indian meal moth, larger grain borer, lesser grain borer, merchant grain beetle, red flour beetle, red flour beetle, rusty grain beetle, rice weevil, and sawtoothed grain beetle.

The Angoumois grain moth was obtained in June 1993 from Community Research Service, Kentucky State University, Kentucky. All other species were obtained in January 1992 from the Department of Entomology, Kansas State University, Manhattan, Kansas. Except for the merchant grain beetle, all species originated from farm-stored grain. The origin of merchant grain beetles is unknown.

Areas of research:

Developing and validating sampling schemes for insects associated with farm-stored grain.

Evaluating nonchemical alternatives for suppressing stored-grain traits.

Modeling population trends of insects from life-history traits.

Rhadriraju Subramanyam, Ph. D.

San Bernardino, CA 92407
California State University
Biology Department

Tribolium brevicornis

Wild Type Strains

1. Original Waterman Population #2
2. Original Waterman Population #1
3. Waterman
4. Waterman
5. Waterman
6. Riverside
7. Riverside
8. Waterman
9. Waterman
10. Waterman
11. Waterman Canyon +/-
12. Waterman Canyon II
- 12'. Riverside
13. Riverside
- 13' Riverside

Mutant Strains.

15. Red eyed mutation and fas-1
16. Strong Rg (reduced gena) fas-1
17. msg-like melanotic stink glands?
18. Short elytra (sshe)
19. she (te-like)
20. short elytra (se)
21. light ocular diaphragm? (lod)
22. incomplete mesosternum (ims)
23. creased abdominal sternites (cas)

Tribolium castaneumWild type strains

1. Chicago
4. Davis
8. McGill
12. Sacramento
16. Yucaipa
19. Synthetic, marked with sooty
22. Wildtype
23. New York

Mutant strains

- | | |
|----------------------------|---------------------------------------------------|
| 31. pg, (p) | pegleg, pearl |
| 33. ros apt | roseate, alate prothorax |
| 38. r | red |
| 51. dve, pd | divergent elytra, paddle |
| 53. pd, py | paddle, pygmy |
| 55. py, r | pygmy, red |
| 56. msg, py, r | pygmy, red, melanotic stink glands |
| 59. r sp | red, Spotted |
| 68. Malta p | Malta pearl |
| 70. pg | peg leg |
| 74. mas, p (pg) | missing abdominal segments, pearl, pegleg |
| 82. b Chicago | black, Chicago |
| 83. b McGill | black, McGill, UPF background |
| 91. lod, p | light ocular diaphragm, pearl |
| 93. Gi | Giant |
| 94. Gi, ptl | Giant, prothoraxless |
| 96. mt | mottled |
| 99. b | tawny |
| 100. b | dusky |
| 101. ap, rb, au, mc, s | antennapedia, ruby, aureate, microcephalic, sooty |
| 105. fas-2 | fused antennal segments-2 |
| 120. spiral | spiral arrangement of all three parts of the body |
| 124. Be, s | Bar eye, sooty |
| 139. mc | microcephalic |
| 143. fas-3a | fused antennal segments-3a |
| 150. rb | ruby |
| 161. Sa, c, mxp | Short antenna, chestnut, maxillopedia |
| 196. mas | missing abdominal segments |
| 220. Rd, p, knp | Reindeer, pearl, knobby prothorax |
| 256. weird | weird eggs |
| 272. supergiant | |
| 276. Davis low body weight | |
| 295. p, pd | pearl, paddle |
| 338. pd, py, p | paddle, pygmy, Pearl |
| 381. b, ptl | black, prothorax-less |
| 392. j-2 | jet-2 from Beeman |
| 421. Rd, ptl, p | Reindeer, prothorax-less, pearl |
| 436. au, mc | aureate, microcephalic |
| 444. i, locd, Mo | ivory, light ocular diaphragm, Microphthalmic |

Stock Lists

- 63. ble (rby)
- 65. ble. e (cas. sti)
- 71. bif
- 73. bit. es (elb)
- 79. cas. sti
- 85. Chi +/- ex N.Y.
- 87. Chi +/-
- 89. cru
- 90. cru. Hg
- 93. dim'd eye
- 95. dpe
- 97. di (strong)
- 101. di. r2
- 103. dt (see umb)
- 105. dt. es
- 109. dt. p
- 111. e
- 114. e Winnipeg
- 115. e. fas-3
- 117. e McGill
- 119. e L & H
- 121. e2 (fas. p. sti)
- 123. e (fas-1)
- 124. e-2. p
- 125. ele
- 128. e2. p fas-2
- 129. es
- 130. e2-lod
- 131. es. fas
- 133. es (car)
- 142. fas-1
- 145. fas-2
- 147. fas-2. di. msg
- 149. fas-2. lod. p
- 150. fas-2
- 151. fas-2. msg
- 152. fas-4
- 153. fas-3 Yugo
- 154. fro
- 157. lod. p.
- 159. lod. rs
- 161. mag
- 163. +/- McDonald
- 165. +/- McGill
- 167. msg (sp)
- 170. msg inbred
- 171. msg inbred 113 generations
- 172. Hg (msg. e-2. p)
- 176. Hg. es (apt. msg)
- 177. Hg. es
- 180. fas-2. lod. p. msg
- 181. N.Y. +/- (msg.sti)
- 182. Npp-like (weak) r/r
- 183. ov-like
- 185. ov-like. Sp
- 187. p (sti. cas)

Stock Lists

448 ap., Chr	antennapedia, Charcoal
464. i, lod	ivory, light ocular diaphragm
471 R-mal	malathion resistant
478 Spa, p	Spatulate, pearl
484. mxp (ap)	maxillopedia, antennapedia
485. fas-like	fused antennal segments-like
487 by syn	black synthetic
488. (au), lod, p	aureate., light ocular diaphragm, pearl
491. fas	fused antennal segments
494 Ag/Es'	Argentum/ Es'
S 6	
495. Dch /ey	Dachshund, eyeless
496. Stbd/Es'	
497. pt/ey	
490. Stm/ey	

T. confusum

1. apt, msg
2. apt, msg, r
4. au, p
6. au, msg, rus (p)
7. b. (sh., cas. sp)
8. bI
10. b. fas-2
11. b. fas-3, r
14. b. lod, au, p (sp)
15. b. p
17. b. rus
19. bI
23. sh., spl
25. b. sp
29. b. twa
33. b-2
35. b (cas. sti. r)
36. b (fas-3)
37. b, r (cas)
39. b Chicago/ b McGill
41. b Donner
43. b Georgia
47. b McGill
49. b McGill ex N.Y.
51. b McGill (syn)
53. b McGill fas
57. b SSM, sp
58. b. spl
59. b. syn
60. b. circle +/-

- 188. p (dre)
- 189. p Slough
- 192. py-like
- 193. ptl (msg)
- 194. +/+ Redlands
- 195. r (msg, sp)
- 196. riboflavinless. p
- 199. ru (sti, cas)
- 200. r Zagreb (msg, cas, Fas)
- 203. rby (cas, msg)
- 205. rus (cas, sti, msg)
- 206. rus-like (inbred)
- 210. black found in 206
- 211. Sacramento +/-
- 213. San Bernardino +/-
- 215. sh (Berkeley)
- 219. sp (spl)
- 221. spl-1 Sok (ble, sti)
- 223. sti (msg, cas)
- 225. stl
- 227. soy adapted
- 229. synthetic +/-
- 230. pearl robp:avonless (msg)
- 231. thu (msg)
- 233. twa
- 234. umb
- 236. XI (sh, sp)
- 238. +/- Yugo
- 239. msg.
- 250. +/- Japan
- 251. autosomal lethal nr b.

Tribolium freemani

1. +/+	Wild type	Slough
1a. +/+	Wild type	Slough
1b. +/+	Wild type	Slough
2. +/+	Wild type	Japan
2a. +/+	Wild type	Japan
2b. +/+	Wild type	Japan
3. b/b	black	San Bernardino
3a. b/b	black	San Bernardino
4. cas	creased abdominal segments	San Bernardino
4a. cas	" " "	" "
4b. cas dve	cr. abd. seg., divergent elytra	" "
5. cor	corrugated elytra	" "
6. ju	juvenile urogomphi	" "
7 fas-1	fused antennal segs.-1.	" "
8. mdl	median line on abd. sternites	" "
8a "	" " " "	" "
9. ov-like	overshot-like	" "
9a "	"	" "
10. sc	scar	" "
10a. sc	"	" "
10b. sc	"	" "
11. vt	vaulted elytra	" "
12. ims	incomplete mesosternum	" "

A. Sokoloff

WASHINGTON, D.C. 20204
 DEPARTMENT OF HEALTH, EDUCATION AND WELFARE
 DIVISION OF MICROBIOLOGY

Coleoptera

Anobiidae

Stegobium paniceum (L.)

Anthribidae

Araecerus fasciculatus (Deg.) (poor condition; may be dead).

Bostrichidae

Rhyzopertha dominica (F.)

Bruchidae

Acanthoscelides obtectus (Say)

Cleridae

Necrobia rufipes (Deg.)

Cucujidae

Ahasverus advena (Waltl)

Cryptolestes ferrugineus (Steph.). Poor condition, may be dead.

C. pusillus (Schon.)

C. turcicus (Grouv.)

Oryzaephilus surinamensis (Linnaeus)

Curculionidae

Sitophilus granarius (L.)

S. zeaeis Motschulsky

Deræstidae

Anthrenus flavipes LeC. Weak culture

Anthrenus verbasci (Linnaeus)

Deræstes maculatus De Geer

Trogoderma variabile Ballion

Ostomidae

Sibium psylloides (Czemp.)

Silvanidae

Ahasverus advena (Waltl.)

Oryzaephilus surinamensis

Tenebrionidae

Alphitobius diaperinus (Panz.)

Gnathocerus maxillosus (F.)

Palorus ratzeburgi (Wissm.)

Tribolium brevicornis (LeConte)

T. castaneum (Herbst)

T. confusum Duv.

T. destructor Uytt. --weak culture, may be diseased.

T. madens (Charpentier)

M. Nakashima

AUSTRALIA

Burnley, Victoria
 Victoria Plant Research Institute
 Department of Agriculture

COLEOPTERA

Tribolium castaneum

Wild type strains
 Malathion specific resistant strain
 Malathion non-specific strain

Tribolium confusum

Wild type strains
 Malathion specific strain

Oryzaephilus surinamensis

Wild type strain
 Malathion resistant strain

Oryzaephilus mercatorAlphitobius diaperinusCryptolestes ferrugineusGnathocerus cornutusGnathocerus maxillosusLatheticus oryzaeRhyzopertha dominicaSitophilus granariusSitophilus oryzaeSitophilus zeamaisTenebroides mauritanicus

LEPIDOPTERA

BURLINGTON, NORTH CAROLINA
CAROLINA BIOLOGICAL SUPPLY COMPANY

Tribolium castaneum

1. black
2. jet
3. pearl
4. Wild
5. High body weight
6. Low body weight

Tribolium confusum

1. Wild

(Ed.).

BURLINGTON, VERMONT 05401
UNIVERSITY OF VERMONT
DEPARTMENT OF ZOOLOGY
STEVENS/GOODNIGHT LAB

T. confusum

bI
bII
bIII
bIV
b-Chicago b/b
b-Chicago
b-Circle
b-yugo-Illinois b/b
b-yugo-Illinois +/-
bSM
b-yugo-Kentucky
b-McGill
b-Thailand
b-Nigeria
b-Pakistan

T. castaneum

cI
cSM-+/+
cCM-b/b
cIV-a
c-Brazil
c-Costa Rica
c-Thailand
c-Spain
c-Israel

Oryzaephilus
surinamensis

L. Stevens

CARBONDALE, ILLINOIS 62901
SOUTHERN ILLINOIS UNIVERSITY AT CARBONDALE
DEPARTMENT OF ZOOLOGY

Tribolium castaneum

I. Wild type strains

1. Purdue + Foundation

II. Mutant strains

1. antennapedia (ap)

D.C. Englert

Chicago, Illinois 60637-1573
The University of Chicago
Department of Ecology and Evolution

Stock lists

I. Wild type strains

A. Tribolium castaneum

1. c+, "Chicago" (from Thomas Park)
2. c-ARK, Arkansas
3. c-YUGO, Yugoslavia, now Croatia
4. c-Texas
5. c-BS, collected in Naperville, IL, on birdseed
6. c-Infantes, Spain
7. c-Jerez, Spain
8. c-Campanaro, Spain
9. c-Osaka, Japan
10. c-Nigeria

B. Tribolium confusum (≠ infected with Wolbachia pipientis)

11. b+, "Chicago" from Thomas park)
12. b-I, inbred strain derived from (1).
13. b-II, inbred strain
14. b-III, " "
15. b-IV " "
16. b-YUGO, Yugoslavia, now Croatia
17. b-YUGO, " "
18. b-Illinois
19. b-Mississippi
10. b-Nigeria

Michael J. Wade Norman T. Johnson

Ephestia cautellaEphestia figulellaGalleria mellonellaPlodia interpunctella

P. Williams

Indooroopilly, Queensland 4068, Australia
 Queensland Department of Primary Industries
 Plant Protection Unit

Coleoptera

Oryzaephilus surinamensis

Wild type strains

VOS 48	insecticide susceptible	Victoria
QOS 42	fenitrothion susceptible	Queensland
QOS 115	chlorpyrifos-methyl-R a	Queensland

Rhyzopertha dominica

Wild type strains

QRD 369	phosphine-resistance	Queensland
QRD 14	insecticide susceptible	Queensland
QRD 2	multiresistant	Queensland
QRD 63	multiresistant	Queensland
QRD 318	pyrethroid-resistant	Queensland

Sitophilus oryzae

Wild type strains

LS 2	insecticide susceptible	Queensland
QSO 56	multi-resistant	Queensland
CSO 231	multi-resistant	W. Australia
QSO 388	phosphine-resistant	Queensland

Tribolium castaneum

Wild type strains

QTC 4	insecticide susceptible	Queensland
QTC 279	pyrethroid insecticide resistant	Queensland
QTC 285	multi-resistant, composite strain	Queensland
QTC 12	non-specific malathion resistant	Queensland
QTC 34	malathion specific-resistant	Queensland
QTC 320	phosphine-resistant	Queensland

Lepidoptera

Queensland

Ephestia cautella Wild

Patrick J. Collins, Senior Entomologist

**ECOLOGY OF FIELD AND STORED PRODUCT PESTS SECTION
AGRICULTURE AND AGRI-FOOD CANADA
WINNIPEG RESEARCH CENTRE
195 DAFOE ROAD
WINNIPEG, MANITOBA, R3T 2M9**

STOCKLIST

SPECIES		ORIGIN	
COLEOPTERA			
1.	<i>Acanthoscelides obtectus</i>	Phillips, Wis	1993
2.	<i>Ahasverus advena</i>	Argyle, MB	1991
3.	<i>Callosobruchus maculatus</i>	Phillips, Wis	1993
4.	<i>Cryptolestes ferrugineus</i>	Manitoba, MB	1991
5.	<i>Cryptolestes pusillus</i>	Lac du Bonnet, MB	1988
6.	<i>Cryptolestes turcicus</i>		1971
7.	<i>Cynaenus angustus</i>	Minnesota, MN	1982
8.	<i>Lasioderma serricorne</i>	Winnipeg, MB	1984
9.	<i>Liposcelis bostrychophilus</i>	Winnipeg, MB	1994
10.	<i>Oryzaephilus mercator</i>	Winnipeg, MB	1994
11.	<i>Oryzaephilus surinamensis</i>	Landmark, MB	1991
12.	<i>Prostephanus truncatus</i>	Mexico City, Mexico	1977
13.	<i>Rhyzopertha dominica</i>	Manitoba	1993
14.	<i>Sitophilus granarius</i>		
15.	<i>Sitophilus oryzae</i>	Coal Lake, AB	1992
16.	<i>Sitophilus zeamais</i>		
17.	<i>Stegobium paniceum</i>	Winnipeg, MB	1993
18.	<i>Tenebrio molitor</i>	Winnipeg, MB	1980
19.	<i>Tribolium audax</i>		
20.	<i>Tribolium castaneum</i>	Manitoba	1991

SPECIES		ORIGIN
The following <i>Tribolium castaneum</i> mutant strains were received in November, 1985 from Dr. Sokoloff's laboratory at California State University.		
21.	Culture S38	red eye
22.	Culture S351	red eye, pygmy, fused antennal segments
23.	Culture S156	microphthalmic
24.	Culture S136	jet (dark body)
25.	Culture S113	sooty (dark body)
26.	Culture S63	pearl eye
27.	Culture S165	chestnut eye
28.	Culture S148	maroon eye
29.	Culture S38	paddle (antennae fused, flattened)
30.	<i>T. castaneum</i>	abbreviated appendages (aa), missing abdominal <i>sternites</i> (mas)
31.	<i>T. castaneum</i>	Rio Desago Malathion resistance
The following mutant strains of <i>Tribolium castaneum</i> have had no linkage analysis:		
32.	malathion-specific resistance	
33.	black body and pearl eyes	
The following mutant strains of <i>Tribolium confusum</i> have had no linkage analysis.		
34.	red eyes	
35.	black body	
36.	<i>Tribolium confusum</i>	Winnipeg, MB 1994
37.	<i>Tribolium madens</i>	
38.	<i>Trogoderma variabile</i>	
39.	<i>Typhaea stercorea</i>	Manitoba 1991
LEPIDOPTERA		
1.	<i>Plodia interpunctella</i>	Winnipeg, MB 1990
2.	<i>Sitotroga cerealella</i>	Kansas 1982

Dr. Noel D.G. White
Section Head

Gembloux agricultural University - Unit of general and applied Zoology
2, Passage des déportés - B-5030 Gembloux, Belgium
zoologie@fsagx.ac.be

Dr Eric Haubruge & Ludovic Arnaud

Insect	Strain	Origin	Year	From
Bostrychidae				
<i>Prostephanus truncatus</i>	Togo	Togo	1993	-
	Dalaba	Guinea-Konakry	1996	-
<i>Rhizopertha dominica</i>				
Insecticide susceptible	Canada	Canada	1991	P. Fields
Insecticide resistant	Methyl bromide, phosphine	Kenya		P. Golob
Bruchidae				
<i>Callosobruchus maculatus</i>	Senegal	Senegal	1989	D. Seck
	Campinas (black strain)	Brazil	1975	O. Legros
Curculionidae				
<i>Sitophilus granarius</i>	Belgium	Belgium	1991	-
<i>Sitophilus zeamais</i>	Senegal	Senegal	1995	G. Pierrard
	Dimbokro	Ivory Coast	1998	-
Gryllidae				
<i>Gryllus bimaculatus</i>		Spain		E.H. Morrow
Tenebrionidae				
<i>Tribolium anaphe</i>		Nigeria	1956	Slough, UK
<i>Tribolium audax</i>		Canada	1969	Slough, UK
<i>Tribolium brevicornis</i>				A. Sokoloff
<i>Tribolium confusum</i>	Dalaba	Guinea-Conakry	1996	-
	Hoielaert	Belgium	1999	-
<i>Tribolium castaneum</i>				
Insecticide susceptible	Abidjan	Ivory Coast	1989	F. Fleurat-Leussard
	Lab-S	USA		R. Beeman
	Japan	Japan		H. Nakakita
	Mozambique	Mozambique		N. White
	Ex-maff	UK	1991	P. Golob
Insecticide resistant	A20 Rdiel (dieltrin, lindane)	USA		R. Beeman
	Argyle, malathion-specific	Canada	1992	N. White
	CTC-12 (malathion, cross resistant)	Australia	1968	D. Wool
	Dalaba, malathion-specific	Guinea-Conakry	1996	-
	Dimbokro (malathion-specific, lindane)	Ivory-Coast	1997	-
	Ga-1, malathion-specific	Geogia, USA	1980	R. Beeman
	Kano, malathion-specific	Nigeria	1961	D. Wool
	Landmark, malathion-specific	Canada	1991	N. White
	Pakistan, malathion-specific & lindane	Pakistan		P. Golob
	Paulo d'Amico (malathion-specific)	Canada	1976	N. White
	PRm, malathion-specific	Philippines	1989	P. Golob
	Rio desago, malathion-specific	Canada	1976	N. White
	Steinback (malathion-specific)	Canada	1989	N. White
	Sun Chong (malathion, cross resistant)	Canada	1976	N. White
	Thailand (malathion-specific & phosphine)	Thailand	1989	P. Golob
	Waseco, malathion-specific	Canada	1982	N. White
Mutant strain	Black Jack	-	1993	-
<i>Tribolium destructor</i>		Ethiopia	1968	Slough, UK
<i>Tribolium freemani</i>		Japan	1980	H. Nakakita
<i>Tribolium madens</i>				A. Sokoloff

SANTA FE DE BOGOTA, D.C.,
UNIVERSIDAD NACIONAL DE COLOMBIA
FACULTAD DE CIENCIAS
DEPARTAMENTO DE BIOLOGIA
APDO. AEREO #14470

Tribolium castaneum

I. Wild type strains

NAME	ORIGIN	DATE
1. ABBC	Synthetic, Bogota	1982
2. Apulo	Apulo (Cund.) Col.	1982
3. Bogota	Inst. Publ. Health, Bogota, Col.	1978 1981
4. Bucaramanga	Bucaramanga, (Sant.)	1981
5. Cartagena	Cartagena, Bol., Col	1980
6. Fusa	Fusagasuga, Cund. Col	1986
7. Honda	Honda, tol. Col.	1986

II. Domestic mutants

Mutant strains discovered in Colombia

NAME	SYMBOL	LINKAGE GROUP	ORIGIN	DATE OF ENTRY
	N			
8. Antennapedia	ap	VIII	Bog.	1981
9. Argentum eyes	ae	I	Bog	1993
10. Bifurcated antenna	ab	II	Bog.	1980
	N			
11. Black	b	III	Bog.	1983
12. colossal pupae	cp	?	Bog.	1993
	b			
13. Charcoal	Chr	III	Bog.	1979
14. Disjuncted elytra	ed	?	Bog.	1990
15. Fused antennameres	af	?	Bog.	1980
16. Glass legs	pv	?	Bog.	1980
17. Globose antenna	Ag	VII	Bog.	1989
18. Light eyes-1	oc	?	Bog.	1990
19. Light eyes-2	?	?	Bog.	1990
20. Light eyes-3	?	?	Bog.	1991
21. Light eyes-4	?	?	Bog.	1993
22. Metathoracic scar	sc	III	Bog.	1983
	V			
23. Miniature appendaged	ma	I	Bog.	1981
24. Narrow eyes	oje	?	Bog.	1980
25. Red eyes	or	?	Bog.	1986
26. White eye	obl	IV	Bog.	1982

III. Imported mutants from Tribolium Stock Center, 1985

	D			
27. Antennapedia	ap	VIII		
28. Black	b	III	33. Pearl eye	p II
29. Charcoal	Chr	III	34. Platinum eye	pte I
30. Miniature appendaged	ma	I	35. Pygmy	py I
31. Microcephalic	mc	V	36. Short antenna	Sa VII
32. Microphthalmic	Mo	VI	37. Sooty	s IV

DENMARK

LYNGBY

STATENS SKADEDYRLABORATORIUM

(DANISH PEST INFESTATION LABORATORY)

Anthrenus museorumA. voraxAttagenus smirnoviA. unicolor (piceus)A. woodroffeiDermaestes hemorrhoidalisLasioderma serricorneOryzaephilus surinamensisProstephanus truncatusPtinus tectusSitophilus granariusS. oryzaeStegobium (Sitodrepa) paniceumTenebrio molitorThylodrias contractusTribolium confusumT. destructorTrogoderma angustumT. granarium

K. Arevad and H. Mourier

FRANCE

VILLEURBANE (LYON) RHONE
INSTITUT NATIONAL DES SCIENCES APPLIQUEES
LABORATOIRE DE BIOLOGIE

A. Wild type strains

1. Sitophilus granarius L.
2. S. oryzae L.
 - a. FB strain (La Reunion)
 - b. SFr strain (lyon) (56,500+3,000 ovarian symbiotes)
 - c. W strain (Villeurbane) (22,700+1500 ovarian symbiotes)
3. S. zeamais Mots--from PIL, Slough

B. Selected lines of Sitophilus oryzae

1. SS/Sfr strain: aposymbiotic strain (0 ovarian symbiotes)
obtained from Sfr
2. LL strain (slow development) (42,000+3000 ovarian symbiotes)
3. RR strain (fast development) (80,000+5000 ovarian symbiotes)

P. Nardon

(No updated list available, Ed.).

ZOOLOGISCHES INSTITUT I
(ZOOLOGIE) DER ALBERT LUDWIGS UNIVERSITÄT
D 78 FREIBURG IM BREISGAU
KATHARINENSTRASSE 20

Wild type strains

1. Oryzaephilus surinamensis Freiburg
2. Tribolium castaneum San Bernardino
3. T. confusum San Bernardino

Mutant strains (All from San Bernardino)

- A. Tribolium castaneum
 4. alate prothorax (apt)
 5. Bar eye (Be)
 6. black (Brazil background)
 7. black (Chicago background)
 8. Dachs (Dch)
 9. Fused tarsi and antennae (Fta)
 10. Microphthalmic (Mo)
 11. nude (nd)
 12. pygmy (py)

13. short antenna (sa)
14. Short antenna (Sa-2)
15. sooty (s)
16. Spatulate antenna (Spa)
- weird eggs (wd)

- B. Tribolium confusum
 18. black-3 (b-3)
 19. ebony (e)
 20. ebony-2 (e-2)
 21. McGill black (Mc6b)

K. Sander

MUNICH,
BAYER. LANDESANSTALT FÜR BODENKULTUR
UND PFLANZENBAU, ABT. PFLANZENSCHUTZ

Coleoptera

Bruchidae--Acanthoscelides obtectus (Say)

Cucujidae--Cryptolestes turcicus Grouv. Munich, 1966

Ptinidae

Gibbium psyllodes (Czemp)

Regensburg, 1960

Ptinus tectus (Boi.)

Munich, 1972

Silvanidae

Oryzaephilus mercator (Fauv.) Munich, 1966
O. surinamensis (L) ? 1971
 Munich (cont'd)

Tenebrionidae

Gnathocerus cornutus (F.) MUNICH, 1966
Tribolium castaneum ? 1971
T. confusum Duv. Munich, 1960
T. destructor Nyttenb. " 1957

Lepidoptera

Phycitidae--Ephestia kuehniella (Zell.) " 1966

E. Naton.

GERMANY

D-80333 München
 Institut für Zoologie
 Luisenstrasse 14

WILD TYPE

Tribolium castaneum

MUTANTS provided by A. Sokoloff

Tribolium castaneum

Bar eye, sooty (Be, s)
 Black, microcephalic pearl (b,mc,p)
 Microcephalic (mc)
 Microcephalic aureate (mc, au)
 Microphthalmic (Mo)
 Squint (sq)

Tribolium confusum

Diminished eye (dim)

INDIA

NEW DELHI
 INDIAN AGRIC. RESEARCH INSTITUTE
 DIVISION OF ENTOMOLOGY
 INSECT GENETICS LAB.

STOCK LIST

STRAIN	RESIST LEVEL	REARING MEDIA
1. Malathion-resist.	>x200	common wheat flour charged with tech malathion.
2. lindane-resist.	>x100	c. w. f. charged with tech. lindane.
3. DDT-RESISTANT	>x100	c.w.f. charged with tech ddt.
4. pirimiphosmethyl resistant	>x100	C.W.F. CHARGED WITH tech. pirimiphosmethyl
5. phosphine-resistant	> 6.3	c.w.f.
6. delta-methrin resist	>2819.3	cwf charged with tech deltamethrin
7. fenitrothion-resist.	>25.96	c.w.f.
8. susceptible	-	c.w.f.
9. black mutant	-	"

Tribolium confusum

10. susceptible	-	c.w.f.
11. nigrat- melanic mutant	-	"

J.D. Saxena.

Tel Aviv University, Israel

Tribolium castaneum -wild type :

- | | |
|------------|-----------------|
| 1) Ishaaya | Israel |
| 2) CTC-12 | Slough, England |
| 3) Kano-C | Slough, England |

-mutants

- | | | |
|-------------|------------------|-------------------------------|
| 1) csbb | black | Tribolium stock center |
| 2) cs pearl | pearl | Tribolium stock center |
| 3) cs mc | microcephalic* | recovered from cs pearl stock |
| 4) cs eu | extra urogomphi* | recovered from csbb stock |
| 5) cs pygmy | pygmy | Tribolium stock center |

* it is uncertain if the mutations are still maintained in the stocks.

Tribolium confusum - wild type

- | | |
|---------------|------------------------|
| 1) CF Chicago | Tribolium stock center |
|---------------|------------------------|

-mutants

- | | | |
|----------|-------------|------------------------|
| 1) CF bb | black | Tribolium stock center |
| 2) CF xl | extra large | Tribolium stock center |

Tribolium brevicornis

- | | |
|-----------------|------------------------|
| 1) Riverside ++ | Tribolium stock center |
|-----------------|------------------------|

Tribolium freemani

- | | |
|-------------------|-------------------|
| 1) Tsukuba strain | Gembloux, Belgium |
|-------------------|-------------------|

Prof. David Wool,

JAPAN

NATIONAL FOOD RESEARCH INSTITUTE
 MINISTRY OF AGRICULTURE, FORESTRY AND FISHERIES
 2-1-2 KANNONDAI, YATABE-MACHI
 TSUKUBA-GUN, IBARAKI-KEN 305

Psocoptera

Liposcelidae

Liposcelis bostrychophilus Badonel Wild

Liposcelis entomophilus (Enderlein) Wild

Trogidae

Lepinotus reticulatus Enderlein Wild

Coleoptera

Anobiidae

Lasioderma serricorne (Fabricius) Wild

Stegobium paniceum (L.) Wild

Ptinidae

Gibbium equinoctiale Boieldieu Wild

Bostrichidae

Rhyzopertha dominica (Fabricius) Wild

Dinoderus minutus (Fabricius) Wild

Cucujidae

Cryptolestes turcicus Wild

Cryptolestes pusilloides (Steel & Howe) Wild

Silvanidae

Oryzaephilus surinamensis (L.) Wild

Tenebrionidae

Alphitobius diaperinus (Panzer) wild

Gnathocerus cornutus (Fabricius) Wild (Okayama str.)

Palorus ratzeburgi (Wissmann) Wild

Tribolium castaneum (Herbst) Wild

T. confusum Jacquelin du Val Wild

T. freemani Hinton Wild

Tenebrio molitor L.

Bruchidae

Callosobruchus chinensis (L.) Wild

Anthribidae

Araecerus fasciculatus Degeer Wild

Rhynchophoridae

Sitophilus zeamais Motschulsky Wild

Sitophilus oryzae (L.) Wild

Lepidoptera

Pyralidae

Ephestia cautella (Walker) Wild

E. kuhniella (Zeller) Wild

Plodia interpunctella Wild

Corcyra cephalonica Wild

Gelechiidae

Sitotroga cerealella (Olivier) Wild

OKAYAMA
LABORATORY OF APPLIED ENTOMOLOGY
COLLEGE OF AGRICULTURE
OKAYAMA UNIVERSITY

1. Wild type strains

COLEOPTERA

1. <u>Alphitobius diaperinus</u>	Miyazaki
2. <u>Callosobruchus chinensis</u>	Okayama
3. <u>C. maculatus</u>	
4. <u>Gnathocerus cornutus</u>	Miyazaki
5. <u>Lasioderma serricorne</u>	Okayama
6. <u>Latheticus oryzae</u>	Miyazaki
7. <u>Oryzaephilus surinamensis</u>	Miyazaki
8. <u>Palorus ratzeburgii</u>	Miyazaki
9. <u>P. subdenressus</u>	Miyazaki
10. <u>Rhyzopertha dominica</u>	Miyazaki
11. <u>Sitophilus oryzae</u>	Okayama
12. <u>S. zeamais</u>	Okayama
13. <u>Tenebrio molitor</u>	Okayama
14. <u>Tenebroides mauritanicus</u>	Okayama
15. <u>Tribolium castaneum</u>	Miyazaki
16. <u>T. confusum</u>	Miyazaki
17. <u>T. freemani</u>	

HYMENOPTERA

1. <u>Anisopteromalus calandrae</u>	Okayama
2. <u>Choetospila elegans</u>	Okayama
3. <u>Lariophagus distinguendus</u>	Okayama

Toshiharu Yoshida

INSTITUTE OF BIOLOGICAL SCIENCES
UNIVERSITY OF TSUKUBA
SAKURA-MURA, IBARAKI
300-31 JAPAN

Bruchidae

Callosobruchus chinensis

13 wild type strains from different localities in Japan
and abroad

Black colored mutant derived from Shusenji strain.

- cC Mainland China
- fC Fukushima, Japan
- hC Hirosaki, Japan
- h1C Hirosaki, Japan

jC Kyoto, Japan, 1936
 mC Morioka, Japan
 nC Niigata, Japan, 1964
 pC Punjab, India
 sCb1 Shusenji black mutant
 tC Tokyo (Nishigahara, Nat. Inst. Agr.), Japan
 taC Tsukuba, Japan
 taC2 Tsukuba, Japan
 tsC Tsukuba, Japan
 yC Taisha, Japan

C. maculatus

12 wild type strains from different localities in the world.

aQ U.S.A. (probably Louisiana).
 bQ Burma
 cQ Fresno Lab., USDA, Calif., U.S.A.
 eQ Thailand
 fQ Thailand
 nQ Ohio, U.S.A.
 rQ
 tQ Tel Aviv, Israel (Dept. Plant Prot., Stored Prod. Res. Res. Lab.)
 kQ Kyoto, Japan
 mQ Kansas State Univ., Manhattan, KS, U.S.A.
 sQ Savannah Lab, USDA, Georgia, U.S.A.

C. analis From United Kingdom
C. phaseoli From United Kingdom
Zabrotes subfaciatus From Africa
Acanthoscelides obtectus From California, U.S.A.
 Hymenoptera
 Braconidae
Heterospilus prosopidis from Hawaii, U.S.A.
 Pteromalidae
Anisopteromalus calandrae, Japan
Chaetospila elegans from United Kingdom
Dinarmus basalis from India

K. Fujii

PAKISTAN

LAHORE
University of the Punjab (New Campus)
Department of Zoology

Tribolium castaneum

- a) Pak Wild type strain
- b) CTC 12 Malathion resistant
- c) FSS II Multi organophosphorus susceptible

PEOPLE'S REPUBLIC OF CHINA

Beijing
Beijing Agricultural University
Dept of Animal Science

Tribolium castaneum

Wild type strains

1. Base population for quantitative genetics, Guelph, 1987.
2. Inbreeding line--Beijing, 1987

Mutant strains: pygmy

1. Base population maintained with no artificial selection and minimum of inbreeding--Guelph, 1987
2. Inbreeding line--Beijing, 1987.

Lao Zhang

POLAND.

International Centre of Ecology,
Polish Academy of Sciences
05-092 Lomianki, Dziekanów Leśny near Warsaw
Poland

Stock list:

T. confusum Duval, strain: HV
T. castaneum Herbst, strain: cl

MPus
P. B. J.

Stock Lists

SPAIN

MADRID

INSTITUTO NACIONAL DE INVESTIGACIONES AGRARIAS
DEPARTAMENTO DE GENETICA CUANTITATIVA Y MEJORA ANIMAL

Tribolium castaneum

A. wild type strains

1. Consejo	C.S.I.C. 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
6. Coronada	La Coronada, Spain	1976
7. Andujar	Andujar, Spain	1975
8. Jerez	Jerez, Spain	1975
9. Osuna	Osuna, Spain	1975
10. Carpio	Carpio, Spain	1975
11. Jafo	Jafo, Israel	1975
12. Beer-Sheba	Beer-Sheba, Israel	1975

B. Mutant type strains

13. Black Purdue	Purdue, USA,	1964
------------------	--------------	------

C. Experimental lines

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

	selected for	Temperature (oC)
14. AN-I	high performance at	33
15. AN-II	" "	33
16. AF-I	" "	28
17. AF-II	" "	28
18. AT-I	" "	38
19. AT-II	" "	38
20. BN-I	low performance at	33
21. BF-I	" "	28
22. BF-II	" "	28
23. BT-I	" "	38
24. BT-II	" "	38

Stock Lists

25. RN-I*	high cross performance at	33
26. SN-I*	" " " "	33
27. RN-II	" " " "	33
28. SN-II	" " " "	33
29. RF-I	" " " "	28
30. SF-I	" " " "	28
31. RF-II	" " " "	28
32. SF-II	" " " "	28
33. RT-I	" " " "	38
34. ST-I	" " " "	38
35. RT-II	high cross performance at	38
36. ST-II	" " " "	
37. CTD-I	high performance at diff. levels of selection	
38. CTD-II	" " " "	
39. DTD-I	" " " "	
40. DTD-II	" " " "	
41. ETD-I	" " " "	
42. ETD-II	" " " "	
43. FTD-I	" " " "	
44. FTD-II	" " " "	

D. mutants

45. antennapedia ap, VIII	Purdue, 1964
46. diferencial Df, IV	Purdue, 1964
47. fused antennal segments-2 fas-2 IV	Sokoloff, 1968
48. ivory i ?	Purdue, 1964
49. paddle, pd i	Purdue, 1964
50. pearl p II	Sokoloff, 1968
51. pegleg pg II	Purdue, 1968
52. pygmy py I	Purdue, 1968
53. rose rs I	Purdue, 1964
54. ruby rb ?	Purdue, 1964
55. short elytra sh VIII	
56. squint sq VIII	Purdue, 1964
57. white w ?	Purdue, 1964
58. wine r I	Purdue, 1968
59. eye mutant ?	Madrid, 1967
60. maroon m V	Purdue, 1977
61. melanotic stink glands--like	Madrid, 1968
62. sooty s IV	Sokoloff, 1977
63. chestnut c VII	Sokoloff, 1977
64. microcephalic mc V	Sokoloff, 1977
65. Microphthalmic Mo VI	Sokoloff, 1977
Pk	
66. pink p II	Sokoloff, 1977
67. Bar eye Be IV	Sokoloff, 1977
68. prothoraxless ptl IX	Sokoloff, 1977
69. light ocular diaphragm lod III	Purdue, 1968
70. black B III	Sokoloff, 1977

Tribolium confusum**A. Wild type strains**

71. Coronada La Coronada, Spain

B. Mutants

72. creased abdominal sternites cas II Sokoloff, 1968

73. ebony-2 e-2 II Sokoloff, 1968

Ma. C. Fuentes

REPUBLIQUE DU SENEGAL Bambey,**MINISTERE DE L'AGRICULTURE**

Institut Sénégalais

De Recherches Agricoles

Centre National de la Recherche Agronomique

1. Stock list*Tribolium Castaneum* (Wild type strains)
genetic origin Bambey (Senegal): Dr. Dogo SECK
Chef ISRA/CNBA

UNITED KINGDOM

University of Newcastle upon Tyne

United Kingdom,

Faculty of Agriculture and Biological Sciences,

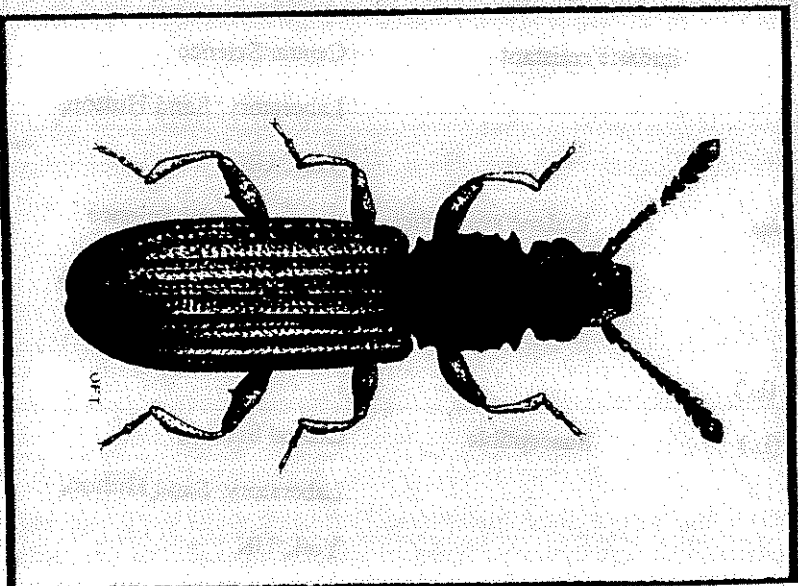
Department of Agricultural and Environmental Science, University of Newcastle upon Tyne, NE1 7RU, UK.

<u>Species/Strains</u>	<u>Status</u>	<u>Derived from</u>
I Wild type strains		
A. <i>Tribolium castaneum</i>		
1. Ph-1	malathion specific resistant	Dr. Freeman, NRI, UK
2. FSS-II	malathion susceptible	Central Science Laboratory, Sand Hutton, York, UK.
B. <i>Tribolium confusum</i>		
1. <i>Tribolium confusum</i>	malathion susceptible	Central Science Laboratory, Sand Hutton, York UK.
C. <i>Sitophilus granarius</i>		
1. 1022 A	lindane resistant	Central Science Laboratory, Sand Hutton, York, UK.
2. <i>Sitophilus granarius</i>	lindane susceptible	Central Science Laboratory, Sand Hutton, York, UK.
D. <i>Sitophilus oryzae</i> (L.)		
1. <i>Sitophilus oryzae</i> (L.)	Susceptible	Central Science Laboratory, Sand Hutton, York, UK.



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Stock Lists

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Other services

- **Identification** - CSL runs courses on identification and on storage and public health pests. We run an identification service and a wide range of insect identification cards is available. Apply to the Librarian at CSL Harpenden (address below) for identification cards.
- **Advice** - on the care and maintenance of cultures can be provided.
- **Mites** - CSL can supply cultures of mites, for which a separate leaflet is available.

Contacts

Mrs C Trowe or Mrs S Henderson
Central Science Laboratory,
MAFF, London Road, Slough
Berks, SL3 7HJ UK

Telephone : 0753 534626
International code : +44 -753 534626
Fax : 0753 824058
International Fax : +44 -753 824058

The Librarian,
Central Science Laboratory,
MAFF, Hatchung Green
Harpenden, Herts, AL5 2BD UK

Telephone : 0582 715241
International code : +44 -528 715241
Fax : 0582 762178
International Fax : +44 -582 762178

Species currently available

Coleoptera

- | | | |
|----|------------------------------------------------|-----------------------------------------------|
| 47 | <i>Ahisserus nitens</i> | <i>Dermestes maculatus</i> pearl-eye mutant |
| | <i>Alphitobius diaperinus</i> | <i>Dermestes pennsylvanicus</i> |
| | <i>Anthrenus austriacus</i> | <i>Gibbium nequinoctiale</i> |
| | <i>Anthrenus flavipes</i> | <i>Gnathocerus cornutus</i> |
| | <i>Anthrenus picturatus hintoni</i> | <i>Gnathocerus maxillosus</i> |
| | <i>Anthrenus sarnicus</i> | <i>Lasioderma serricorne</i> |
| | <i>Anthrenus verbasci</i> | <i>Lasioderma serricorne</i> black mutant |
| | <i>Attagenus brunneus</i> | <i>Latheticus oryzae</i> |
| | <i>Attagenus cyphonoides</i> | <i>Mezium affine</i> |
| | <i>Attagenus fasciatus cinnamomeus</i> | <i>Mezium americanum</i> |
| | <i>Attagenus misidiosus</i> | <i>Niphus hololeucus</i> |
| | <i>Attagenus pello</i> | <i>Oryzaephilus acuminatus</i> |
| | <i>Attagenus rufiventris</i> | <i>Oryzaephilus mercator</i> |
| | <i>Attagenus smaragdus</i> | <i>Oryzaephilus surinamensis</i> |
| | <i>Attagenus unicolor canadensis</i> | <i>Oryzaephilus surinamensis</i> small mutant |
| | <i>Attagenus unicolor japonicus</i> | <i>Palorus cerylonoides</i> |
| | <i>Attagenus unicolor simulans</i> | <i>Palorus feticola</i> |
| | <i>Attagenus unicolor unicolor</i> | <i>Palorus genalis</i> |
| | <i>Attagenus woodroffei</i> | <i>Palorus ratzeburgii</i> |
| | <i>Attagenus fasciatus fasciatus</i> | <i>Palorus subdepressus</i> |
| | <i>Calosobruchus maculatus</i> | <i>Prostephanus truncatus</i> |
| | <i>Carpophilus dimidiatus</i> | <i>Pseudeurostus hilleri</i> |
| | <i>Carpophilus dimidiatus</i> pearl-eye mutant | <i>Pinus claryps</i> |
| | <i>Carpophilus hemipterus</i> | <i>Pinus exulans</i> |
| | <i>Coelopalorus foenicollis</i> | <i>Pinus pusillus</i> |
| | <i>Cryptolestes capensis</i> | <i>Pinus sexpunctatus</i> |
| | <i>Cryptolestes ferrugineus</i> | <i>Pinus tectus</i> |
| | <i>Cryptolestes pusillus</i> | <i>Rhyzopertha dominica</i> |
| | <i>Cryptolestes pusillus</i> fuscus | <i>Sitophagus hololeptoides</i> |
| | <i>Cryptolestes turcicus</i> | <i>Sitophilus granarius</i> |
| | <i>Cryptolestes turcicus</i> red-eye mutant | <i>Sitophilus oryzae</i> |
| | <i>Cryptolestes ugandae</i> | <i>Sitophilus zeamais</i> |
| | <i>Dermestes ater</i> | <i>Sphaerius gibboides</i> |
| | <i>Dermestes frischeri</i> | <i>Siegbium panicum</i> |
| | <i>Dermestes haemorrhoidalis</i> | <i>Sitona squamosum</i> |
| | <i>Dermestes lardarius</i> | <i>Tenebrio molitor</i> |
| | <i>Dermestes maculatus</i> | <i>Tenebrio obscurus</i> |
| | | <i>Typhlus unicolor</i> |
| | | <i>Tribolium confusum</i> |

Stock Lists

- | | |
|--|-----------------------------------------|
| | <i>Tribolium anaphae</i> |
| | <i>Tribolium aurum</i> |
| | <i>Tribolium brevicorne</i> |
| | <i>Tribolium castaneum</i> |
| | <i>Tribolium castaneum</i> black mutant |
| | <i>Tribolium destructor</i> |
| | <i>Tribolium freemani</i> |
| | <i>Tribolium mindens</i> |
| | <i>Trigonogenus globulus</i> |
| | <i>Trigonogenus particularis</i> |
| | <i>Trogoderma angustum</i> |
| | <i>Trogoderma anthrenoides</i> |
| | <i>Trogoderma glutinum</i> |
| | <i>Trogoderma granarium</i> |
| | <i>Trogoderma grassmanni</i> |
| | <i>Trogoderma inclusum</i> |
| | <i>Trogoderma irritatum</i> |
| | <i>Trogoderma ornatum</i> |
| | <i>Trogoderma sternale plagifer</i> |
| | <i>Trogoderma variabile</i> |
| | <i>Trogoderma varium</i> |
| | <i>Typhaea stercoraria</i> |
-
- | | |
|--|------------------------------|
| | Dictyoptera |
| | <i>Blattella orientalis</i> |
| | <i>Blattella germanica</i> |
| | <i>Diploptera punctata</i> |
| | <i>Periplaneta americana</i> |
-
- | | |
|--|------------------------------|
| | Lepidoptera |
| | <i>Ephestia cautella</i> |
| | <i>Ephestia kuehniella</i> |
| | <i>Galleria mellonella</i> |
| | <i>Plodia interpunctella</i> |
| | <i>Sitotroga cerealella</i> |
| | <i>Tinea pellionella</i> |
| | <i>Tineola bisselliella</i> |
-
- | | |
|--|---------------------------|
| | Thysanura |
| | <i>Leptisma saccharum</i> |
-
- | | |
|--|-----------------------------|
| | Hymenoptera |
| | <i>Monomorium pharaonis</i> |
-
- | | |
|--|----------------------------------|
| | Psocoptera |
| | <i>Liposcelis bostrychophila</i> |
| | <i>Liposcelis subfuscus</i> |
| | <i>Liposcelis paucus</i> |
| | <i>Leptothrips pumilus</i> |
| | <i>Trogium pulsatarium</i> |
-
- | | |
|--|--------------------------------|
| | Hemiptera |
| | <i>Aphis fabae</i> |
| | <i>Aphis gossypii</i> |
| | <i>Brevicoryne brassicae</i> |
| | <i>Macrostiphum euphorbiae</i> |
| | <i>Mizus persicae</i> |
| | <i>Nasonovia ribisnigri</i> |
| | <i>Phorodon humuli</i> |
| | <i>Rhopalosiphum padi</i> |
| | <i>Sitobion avenae</i> |

Availability

Please give two weeks notice. Although most species can be supplied within two weeks, those that breed slowly may take longer to supply. We will advise you if there is likely to be a delay.

CENTRAL SCIENCE LABORATORY

LONDON ROAD
SLOUGH
BERKS U K
SL3 7HJ

TEL: 44 1753 534626
FAX 44 1753 82405

Insects mentioned below are bred in controlled environmental conditions and, as far as possible, free from disease. All new stocks pass through a quarantine procedure before acceptance into the main insectary. This list was last updated January 1995. The country of origin and year of receipt at this laboratory are shown against the strains where this information is known. For some of the older strains such information is not known. Please note that some strains do not have a name, especially if only one strain of a species is held. Where more than five strains of a species are held, full details are not given. (However, full details of all mutant strains held are given). Please write to me or Carol TROWE for further details on any aspect of this list and with any requests for specimens. The latter will be met where sufficient are available, but a charge will have to be made.

CHRISTINE B MUGGLETON (Mrs)

INSECT DATABASE FOR TRIBOLIUM INFORMATION BULLETIN

Genus, species, sub-species.	Strain	Place of origin	Year received
COLEOPTERA			
<i>Ahasverus advena</i>	6 strains from 2 countries, many differing in their susceptibility to pesticides		
<i>Alphitobius diaperinus</i>	6 strains all from Britain, many differing in their susceptibility to pesticides		
<i>Anthrenocerus australis</i>		Britain	1933
<i>Anthrenus flavipes</i>			
<i>Anthrenus flavipes seminiveus</i>			
<i>Anthrenus picturatus hintoni</i>		Russia	1977
<i>Anthrenus sarnicus</i>	Wiltshire	Britain	1966
<i>Anthrenus verbasci</i>		Britain	1951
<i>Attagenus brunneus</i>	Canada		
<i>Attagenus brunneus</i>	Spain		
<i>Attagenus cyphonoides</i>		Spain	
<i>Attagenus fasciatus fasciatus</i>		Tashkent	1976
<i>Attagenus fasciatus cinnamomeus</i>		New S. Wales	1972
<i>Attagenus insidiosus</i>		Botswana	1965
<i>Attagenus insidiosus</i>		Kenya	
<i>Attagenus pello</i>		Britain	1950
<i>Attagenus rufiventris</i>		Botswana	1970
<i>Attagenus smimovi</i>		Kenya	1962
<i>Attagenus unicolor canadensis</i>		N. America	1980
<i>Attagenus unicolor japonicus</i>		Japan	1956
<i>Attagenus unicolor simulans</i>		U.S.S.R.	1976
<i>Attagenus unicolor unicolor</i>			pre 1958
<i>Attagenus woodroffeii</i>	Sweden	Sweden	1978
<i>Attagenus woodroffeii</i>	Finland	Finland	1965
<i>Callosobruchus maculatus</i>			
<i>Carpophilus dimidiatus</i>		USA	pre 1958
<i>Carpophilus hemipterus</i>			1962
<i>Coelopalorus foveicollis</i>		Trinidad	1972
<i>Cryptolestes capensis</i>			1961
<i>Cryptolestes ferrugineus</i>	24 strains all from Britain, many differing in their susceptibility to pesticides		
<i>Cryptolestes pusilloides</i>		Canada	1944
<i>Cryptolestes pusillus</i>			
<i>Cryptolestes pusillus fuscus</i>		Trinidad	1960
<i>Cryptolestes turcicus</i>			pre 1958
<i>Cryptolestes ugandae</i>		E. Africa	1954
<i>Dermestes ater</i>		Britain	1953
<i>Dermestes frischii</i>		Nigeria	pre 1958

<i>Dermestes haemorrhoidalis</i>		Britain	1962
<i>Dermestes lardarius</i>		Britain	pre 1958
<i>Dermestes maculatus</i>	Chittagong	Chittagong	1975
<i>Dermestes peruvianus</i>		Britain	1961
<i>Gibbium aequinoctiale</i>		Britain	1937
<i>Gnatocerus cornutus</i>			pre 1958
<i>Gnatocerus cornutus</i>			
<i>Gnatocerus maxillosus</i>			pre 1958
<i>Lasioderma serricorne</i>			pre 1958
<i>Latheticus oryzae</i>			pre 1958
<i>Mezium affine</i>		Britain	pre 1958
<i>Mezium americanum</i>			1960
<i>Niptus hololeucus</i>		Britain	pre 1958
<i>Oryzaephilus acuminatus</i>		Sri Lanka	
<i>Oryzaephilus mercator</i>			pre 1958
<i>Oryzaephilus mercator</i>	9127 Pickering	Britain	1994
<i>Oryzaephilus surinamensis</i>	54 strains from 4 countries, many differing in their susceptibility to pesticides		
<i>Palorus cerylonoides</i>		Indonesia	
<i>Palorus ficicola</i>	1168	Nigeria	
<i>Palorus ficicola</i>	1176	Nigeria	
<i>Palorus genalis</i>		Guyana	
<i>Palorus ratzeburgii</i>		Britain	1960
<i>Palorus subdepressus</i>		Turkey	1956
<i>Prostephanus truncatus</i>		Tanzania	1981
<i>Pseudeurostus hilleri</i>		Britain	1940
<i>Ptinus clavipes</i>		Britain	1954
<i>Ptinus exulans</i>		Britain	1971
<i>Ptinus pusillus</i>			pre 1958
<i>Ptinus sexpunctatus</i>			pre 1958
<i>Ptinus tectus</i>	Wild	Britain	1975
<i>Ptinus tectus</i>	PICL		1960
<i>Ptinus tectus</i>	Birkenhead	Britain	1975
<i>Rhyzopertha dominica</i>	7 strains from 3 countries, many differing in their susceptibility to pesticides		
<i>Sitophagus hololeptoides</i>		Trinidad	1972
<i>Sitophilus granarius</i>	11 strains from 3 countries, many differing in their susceptibility to pesticides		
<i>Sitophilus oryzae</i>	5 strains from 4 countries, many differing in their susceptibility to pesticides		
<i>Sitophilus zeamais</i>			pre 1958
<i>Sitophilus zeamais</i>	U.S.A.	U.S.A.	1982
<i>Sphaericus gibboides</i>		Britain	1976
<i>Stegobium paniceum</i>			1959
<i>Stethomezium squamosum</i>		Britain	1976
<i>Tenebrio molitor</i>			pre 1958



CENTRAL SCIENCE LABORATORY

Insect Cultures Order Form

Name:

Address:

.....

.....

Tel No: Fax No:

Species	Quantity	Live/ Dead	Adult/ Larvae	Price
Subtotal				
(Note: VAT is not payable for orders paid from outside UK) VAT				
Post & Packing				2.50
TOTAL				

Latest date required: (see note about availability):

Signature: Date:

Please send or fax this order to: Mrs C Trowe
 Central Science Laboratory
 London Road
 SLOUGH, Berkshire, SL3 7HJ UK
 Fax: 0753 824058
 International code +44 -753 824058

(Cheques payable to Central Science Laboratory)

<i>Tenebrio obscurus</i>			pre 1958
<i>Tipnus unicolor</i>		Kenya	pre 1958
<i>Tribolium anaphe</i>		Nigeria	1956
<i>Tribolium audax</i>		Canada	1969
<i>Tribolium brevicornis</i>		U.S.A.	
<i>Tribolium castaneum</i>	9 strains from 3 countries, many differing in their susceptibility to pesticides		
<i>Tribolium confusum</i>	W-44		
<i>Tribolium confusum</i>	Lab. susc.		
<i>Tribolium confusum</i>			1962
<i>Tribolium destructor</i>	African	Ethiopia	1968
<i>Tribolium freemani</i>		Japan	1980
<i>Tribolium madens</i>		Yugoslavia	1959
<i>Trigonogenius globulus</i>		Ireland	1961
<i>Trigonogenius particularis</i>		Kenya	1962
<i>Trogoderma angustum</i>		Germany	1975
<i>Trogoderma anthrenoides</i>		U.S.A.	1957
<i>Trogoderma glabrum</i>		U.S.A.	1959
<i>Trogoderma granarium</i>		Britain	
<i>Trogoderma granarium</i>		Britain	pre 1958
<i>Trogoderma grassmani</i>		U.S.A.	1976
<i>Trogoderma inclusum</i>			pre 1958
<i>Trogoderma irroratum</i>		Egypt	1959
<i>Trogoderma ornatum</i>		U.S.A.	1974
<i>Trogoderma sternale plagifer</i>		New Mexico	1966
<i>Trogoderma variabile</i>		U.S.A.	1965
<i>Trogoderma varium</i>		Korea	1970
<i>Typhaea stercorea</i>	Datchet	Britain	1980
THYSANURA			
<i>Lepisma saccharina</i>		Britain	1978
LEPIDOPTERA			
<i>Epehestia cautella</i>		Cyprus	1969
<i>Epehestia cautella</i>	Brown/Yellow	Florida	
<i>Epehestia cautella</i>	Bedstock		
<i>Epehestia elutella</i>	Lab.		
<i>Epehestia elutella</i>	Millwall	Britain	1969
<i>Epehestia kuehniella</i>	Welsh Buffer Depot	Britain	
<i>Epehestia kuehniella</i>	Rhydymwyn	Britain	1988
<i>Epehestia kuehniella</i>		Britain	1949
<i>Galleria mellonella</i>	B	U.S.A.	1992
<i>Galleria mellonella</i>		U.S.A.	1987
<i>Plodia interpunctella</i>	88	Turkey	1977
<i>Plodia interpunctella</i>		Britain	1968
<i>Plodia interpunctella</i>	121	Chicago	1977
<i>Plodia interpunctella</i>	102	Tanzania	1977
<i>Sitotroga cerealella</i>	623	U.S.A.	1972
<i>Sitotroga cerealella</i>	A68	Nepal	1981
<i>Tinea pellionella</i>		Britain	1989
<i>Tineola bisselliella</i>	U.S.A. Lab. strain		
<i>Tineola bisselliella</i>	U.K. Wild strain		
<i>Tineola bisselliella</i>		Britain	1989

MUTANTS

Carpophilus dimidiatus
Cryptolestes turcicus
Dermestes maculatus
Dermestes maculatus
Lasioderma serricorne
Oryzaephilus mercator
Oryzaephilus surinamensis

pearl-eye
 Red-eye mutant
 Black-brown
 Pearl-eye
 Black mutant
 0779 pearl-eye
 small
 484 -sp eye, lod
 484-sp eye
 484 black dd
 black

Australia
 Australia
 U.S.A.
 Pacific Islands
 East Pakistan
 speckled eye, light
 ocular diaphragm
 speckled eye
 1964
 1964
 1975
 1978
 1964
 1994
 1983

Tribolium castaneum

SLOUGH, BUCKS, U.K.
TROPICAL DEVELOPMENT AND RESEARCH INSTITUTE (FORMERLY TPI)
STORAGE DEPARTMENT
OVERSEAS DEVELOPMENT ADMINISTRATION
PEST BIOLOGY AND INSPECTION SECTION

TROPICAL DEVELOPMENT AND RESEARCH INSTITUTE (TDRI)

The Tropical Development and Research Institute (TDRI) was formed 1 April, 1983, following the amalgamation of the Tropical Products Institute and the Centre for Overseas Pest Research. The Director of the Institute is Dr. Malcolm Thain who was formerly Director of the Tropical Products Institute.

The Institute, part of the Overseas Development Administration and funded from the aid programme, will provide technical assistance to developing countries. The budget will total over eight million pounds in the financial year 1983/84.

TDRI will continue to work on post-harvest technology and pest and vector management for the benefit of developing countries, by controlling the pests harmful to agriculture, stored products and public health, and by improved processing, storage and marketing of agricultural fisheries products.

The main emphasis of its work in scientific research and development, marketing, information, advice and training will centre on the improvement of food supplies in accordance with the major objectives of the British overseas aid programme. Work will also continue on certain non-food crops of particular importance to developing countries. These activities will be carried out, as at present, in the UK and overseas in countries throughout the developing world.

Since post harvest technology and pest and vector management are broad and varied subjects, TDRI will concentrate its activities in those areas where it has a comparative advantage in terms of experience, knowledge and cost-effectiveness. Close cooperation will continue with government organizations, universities and industry in developing countries, the UK and other industrialized countries, and with multilateral and bilateral aid agencies.

Requests from developing country governments qualifying for British aid will be channelled through the Overseas Development Administration, which may commission TDRI to carry out the work if it lies within the scope of its terms of reference, and if resources are available. In addition, TDRI may, subject to the claims on its resources commissioned by ODA, accept contracts for relevant work on behalf of developing countries from multilateral aid agencies and other organizations.

TDRI is based in London, although relocation to a new site outside the central London area is under consideration. It currently employs over 450 staff.

Requests for information, advice, investigations or training should be sent to:

The Director
Tropical Development and Research Institute
56-62 Gray's Inn Road
London WC1X 8LU
England (Telephone 01-242 5412)

All stocks are maintained at 27 degrees centigrade and 70% R.H. The stocks listed below are those currently maintained for ongoing research projects. Other storage pest species are kept in culture from time to time for training or short research projects.

1. Wild type strains

A. Coleoptera

Rostrichidae

1. Prostephanus truncatus -- Mexico, Tanzania

Bruchidae

1. Acanthoscelides obtectus -- Swaziland; Turkey
2. Callosobruchus analis -- MAFF Lab., Slough; Indonesia
3. Callosobruchus chinensis -- Nepal; Kenya
4. Callosobruchus maculatus -- Brazil, 2 strains; Nigeria, 2 strains; Oman; Senegal; Sierra Leone; Turkey; Upper Volta; Yemen.
5. Caryedon serratus -- Unknown
6. Zabrotes subfasciatus -- Uganda (collected from cowpeas and bred on cowpeas); Colombia.

Curculionidae

1. Sitophilus oryzae -- Peru (pulse-feeding strain breeding on split peas)
2. S. zeamais -- Mexico

B. Lepidoptera

Galleriinae: Corcyra cephalonica -- Malawi

Gellechiidae: Sitotroga cerealella -- Sudan

Phycitinae: Ephestia cautella -- Brazil

CHEMICAL CONTROL SECTION

(stocks of some major beetles pests are maintained, under selection pressure with insecticide where necessary, in order to enable the FAO recommended methods for the detection and measurement of resistance to be carried out. Incoming strains from abroad are screened and the methods are demonstrated in training programs.)

Wild type strains

Coleoptera

Bostrichidae

Prostephanus truncatus--Strains tested for phosphine resistance: Botswana; Indonesia; Mali (8 strains) Nepal; Nigeria; Pakistan (2 strains) Singapore; Sri Lanka (4 strains); Tunisia; Zimbabwe.

Bruchidae

Acanthoscelides obtectus -- Ethiopia

Callosobruchus chinensis -- India

Curculionidae

Sitophilus oryzae -- Insecticide-susceptible strain (reference strain) -- via MAFF Lab, Slough

S. oryzae -- Malathion and lindane resistant strain (A.76) -- via MAFF Lab., Slough.

Tenebrionidae

Tribolium castaneum -- Multiple insecticide-resistant strain (CTC 12) -- australia

T. castaneum -- Malathion-specific resistant strains (Kano C) -- Nigeria

T. castaneum -- Insecticide-susceptible strain (reference strain) -- MAFF Lab, Slough

Dr. F. F. Pevett
Deputy Head of Department

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I. Wild type strains

A. Coleoptera

Anobiidae

- | | |
|---------------------------------|------------|
| 1. <u>Lasioderma serricorne</u> | a. Unknown |
| 2. <u>Stegobium paniceum</u> | a. ex-MAFF |

Bostrichidae

- | | |
|----------------------------------|-------------------------|
| 1. <u>Dinoderus distinctus</u> | a. Tanzania |
| 2. <u>D. minutus</u> | a. Indonesia |
| 3. <u>D. porcellus</u> | a. Togo |
| 4. <u>Prostephanus truncatus</u> | a. Costa Rica |
| | b. Mexico (3 strains) |
| | c. Nigeria |
| | d. Tanzania (4 strains) |
| | e. Togo |
| | f. Kenya |
| 5. <u>Rhyzopertha dominica</u> | a. Ex-MAFF |
| | b. Angola† |
| | c. Kenya (3 strains)††† |
| | d. Mali † |
| | e. Morocco † |
| | f. Nepal† |
| | g. Sri Lanka |

Bruchidae

- | | |
|---------------------------------------|-------------------------|
| 1. <u>Acanthoscelides obtectus</u> -- | a. Colombia (2 strains) |
| | b. Uganda |
| | c. Zimbabwe |
| 2. <u>Callosobruchus analis</u> -- | a. MAFF Lab. |
| 3. <u>Callosobruchus chinensis</u> -- | a. Indonesia |
| 4. <u>Callosobruchus maculatus</u> -- | a. Uganda |
| 5. <u>Carvedon serratus</u> | a. India |
| 6. <u>Zabrotes subfasciatus</u> -- | b. Uganda |

Curculionidae

1. Sitophilus oryzae

i. Normal strains

- a. Ex-MAFF
- b. India
- c. Morocco
- d. Zimbabwe

ii. Pulse-feeding

2. S. zeamais --

- a. Ex-MAFF
- b. India

Dermeestidae

1. Dermeestes ater

- a. Ex-MAFF

2. D. maculatus

- a. Jamaica

3. Trogoderma granarium

- a. India
- b. Sudan

Histeridae

1. Teretriosoma nigrescens

- a. Mexico

Lophocateridae

1. Lophocateres pusillus

- a. Philippines

Silvanidae

1. Ahasverus advena

- a. Ex-MAFF

2. Oryzaephilus sp.

- a. Kenya (4 strains)

3. Oryzaephilus surinamensis

- a. Ex-MAFF

Tenebrionida

1. T. castaneum

- a. Ex-MAFF
- b. Botswana†
- c. Indonesia (2 strains)
- d. Kenya ‡
- e. Mali ‡
- f. Mozambique
- g. Pakistan†
- h. Philippines +
- i. Sri Lanka
- j. Thailand (3 strains)†††+
- k. Zimbabwe (2 strains)†

2. Latheticus oryzae

- a. Ex-MAFF

3. Gnatocerus cornutus

- a. Ex-MAFF

4. Palorus subdepressus

- a. Ex-MAFF

Key

† Number of strains which have to date been found to be Phosphine resistant.

+ Malathion resistance noted.

‡ Pirimiphos methyl resistance noted.

B. Lepidoptera

Pyralidae

1. Corcyra cephalonica a. Ex-MAFF
2. Ephestia cautella a. Ex-MAFF
3. Ephestia elutella b. Ethiopia
a. Ex-MAFF

Gellechiidae:

1. Sitotroga cerealella a. Sudan

CHEMICAL CONTROL SECTION

(stocks of some major beetles pests are maintained, under selection pressure with insecticide where necessary, in order to enable the FAO recommended methods for the detection and measurement of resistance to be carried out. Incoming strains from abroad are screened and the methods are demonstrated in training programs.)

Wild type strains

Coleoptera

Bostrichidae

Prostephanus truncatus--Strains tested for phosphine resistance: Botswana; Indonesia; Mali (8 strains) Nepal; Nigeria; Pakistan (2 strains) Singapore; Sri Lanka (4 strains); Tunisia; Zimbabwe.

Bruchidae

- Acanthoscelides obtectus -- Ethiopia
Callosobruchus chinensis -- India

Curculionidae

- Sitophilus oryzae -- Insecticide-susceptible strain (reference strain) -- via MAFF Lab, Slough
S. oryzae -- Malathion and lindane resistant strain (A.76) -- via MAFF Lab., Slough.

Tenebrionidae

- Tribolium castaneum -- Multiple insecticide-resistant strain (CTC 12) -- australia
T. castaneum -- Malathion-specific resistant strains (Kano C) -- Nigeria
T. castaneum -- Insecticide-susceptible strain (reference strain) -- MAFF Lab, Slough

Dr. Chris P. Haines

Stock Lists

YUGOSLAVIA

INSTITUTE FOR BIOLOGICAL RESEARCH
"SINIŠA STANKOVIĆ"
UNIVERSITY OF BELGRADE
DEPARTMENT OF INSECT PHYSIOLOGY & BIOCHEMISTRY

1. *Morimus funereus*, L. (Cerambycidae - Coleoptera), wild type, Fruška Gora & Derdap, Serbia (geographic origin)
2. *Cerambyx cerdo*, L. (Cerambycidae - Coleoptera), wild type, Fruška Gora, Serbia
3. *Tenebrio molitor*, L. (Tenebrionidae - Coleoptera), wild type, Fruška Gora, Serbia
4. *Lymantria dispar*, L. (Lymantriidae - Lepidoptera), wild type, Despotovac, Serbia

Dr. Zlatko Prolic, Ph. D.

R.W. BEEMAN'S TRIBOLIUM HOME PAGE

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A. Sokoloff
Biology Department
California State University
San Bernardino, California 92407.

***R.W. Beeman's Tribolium Home Page**

For those who are not aware that Tribolium has a Home Page prepared by R.W. Beeman and accessible at the address <http://bru.usgmr1.ksu.edu/beeman/tribolium> it is included in this issue of the Tribolium Information Bulletin. It was last edited August '3, 1998, and it contains information on techniques on how to handle beetles in the laboratory as well as an up-to-date description of mutants and their linkage relationships, something which was also available from their extensive stock lists published in the TIB. On behalf of its readers, the Editor thanks Dr. Beeman and his group for their efforts in making the Home Page as current as possible.

Welcome to the *Tribolium* Home Page

This site contains data and articles about the genetics of the red flour beetle, *Tribolium castaneum*, and related species. Work being done in Dr. Beeman's laboratory involves both standard and molecular approaches.

- **Standard Genetics**

- **Beetle Handling**
- **Linkage Maps**
- **Mutants**
- **Medea: Maternal-Effect Selfish Genes**
- **Insecticide Resistance in *Tribolium***

- **Download these Web Pages for local use on your PC.**



[To Dr. Beeman's Page](#)
[To Biological Research Unit](#)

Send comments or questions to
beeman@usgmr1.ksu.edu
haas@usgmr1.ksu.edu

Last Edited: August 13, 1998



BEETLE WRANGLING TIPS

(An Introduction to the Care and Handling of *Tribolium castaneum*)

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Sub culturing

Paper Transfer

Scoop or Spoon Transfer

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Pan Sorting (after sieving)

Use of Topping

Sub culturing Schedule

Diseased Stocks

Trouble Shooting

Trouble Prevention

Disease & Mites

Developmental Rates of *Tribolium castaneum*

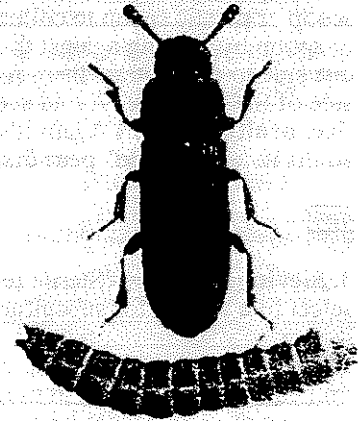
Sexing *Tribolium*

Pupae - Materials

Pupae - Methods

Adults - Materials

Adults - Methods



A. Subculturing

1. Paper transfer- The use of paper strips to transfer adult beetles from an older stock jar to a new one is the quickest and easiest method. Paper strips approximately 5" X 1" are used for sub culturing pint and quart jars. In a bottle with many beetles on the surface of the flour, put the paper strip into the mass of beetles, and wait for them to cover the bottom 1/4 - 1/3 of the strip. (If the jar has fewer adults on top, tilt the jar slightly to one side. Adults will gather in the low side, where you can collect them on the paper strip.) Then quickly but carefully withdraw the strip from the first jar and insert it into the jar of new flour. Shake the paper strip and tap it against the sides of the jar to remove the beetles. Repeat the process until the desired number of beetles is transferred. Discard that paper strip and use a fresh one for the next jar you subculture.

Smaller 5" X 3/4" paper strips are used for sub culturing square bottles or vials. The smaller strips may be cut even narrower for easier insertion into the smaller containers. Alternately, a 3/4" strip can be "bowed" along the narrow edge with the fingers to provide easier insertion into the vial and a more effectively shaped paper strip surface for collecting a smaller population of beetles which are being tilted to one side of the curved surface of a vial (or corner of a bottle) to concentrate them.

(Note: Use one clean strip of paper for each culture jar sub cultured. It's best not to lay the strip down on any surface while sub culturing a jar because of the possibility of a stray egg, larva or adult clinging to it, and being introduced into your jar as a contaminant.)

Use paper transfer whenever possible...it helps prevent transfer of disease via equipment if disease is a problem, and minimizes the possibility of contamination from a stray egg or small larva left in the sieve

or pan. It also selects for the healthiest, most vigorous beetles (with the exception of stocks of beetles with short /defective legs that have difficulty climbing a paper strip.

It's a good idea to spot check each stock sub cultured at the time of each subculture. Just place an extra 10 beetles in a petri dish, cool on ice, and inspect the beetles for proper phenotype. Discard the beetles used for spot check. (If your stock is very small, and every beetle counts, save them, but be very conscientious of good "ot;t,;sterile technique." (i.e., "bang" each petri dish lid and bottom on the tabletop before each use to dislodge any stray eggs or larvae.)

TOP

2. Scoop or spoon transfer- For Dch-3, and other mutants with very short / defective legs, use a small scoop or plastic spoon to collect adults from one jar or bottle and transfer to another. "Sterilize" the scoop or spoon by rapping against the table top several times on both sides. Tilt the bottle so adults move to one side to concentrate them for scooping. Scoop carefully to prevent mashing any beetles against the side of the container. Avoid scooping flour as much as possible. (You just want to collect ~~not~~ **live, healthy** adults.) Again, it's a good idea to spot check each stock as you subculture it. Place about 10 adults in a "sterilized" petri dish as mentioned above.

TOP

3. Sieving transfer ("Sterile technique")- When paper or scoop transfers are not possible, sieve and select live beetles for subculture by using the following protocol:

Bang sieves, receiving pans, and aluminum sorting pans firmly and thoroughly on wastebasket lid immediately before and immediately after use. Bang the plastic transfer funnel lip sharply on the tabletop or wastebasket lid several times.

Inspect banged equipment visually for presence of clinging larvae or adults. If larvae are stuck in the sieve, try to dislodge by additional banging. If this fails, gently poke at them with a brush to encourage them to go on through or withdraw, whichever is the shorter route to getting out. Be careful not to damage them while they are caught in the sieve. If they bleed onto the sieve, their blood and body fluids will corrode the screen.

"Squeegee" sterilize brushes between thumbnail and index finger before using each time.

Always sieve into a receiving pan, never onto the table top! Sieve any flour which contains larvae as quickly as possible, with continuous agitation dump siftings immediately into sorting pan to minimize the opportunity for larvae to try to crawl through screen and get stuck. For those caught in screen, dislodge first by banging sieve against receiving pan (first up-side-down, then right-side-up). Dislodge any remaining larvae by poking or "tickling" through screen gently with brush. Don't use ~~any~~ lateral brushing action to dislodge stuck larvae --- rough treatment can squash larvae, and hemolymph from injured larvae can corrode screen of sieve!!

After sieving diseased stocks, wipe down the sieve and receiving pan with alcohol and dry completely (place on a heat source such as a scope light source, or top of hot incubator, to evaporate excess moisture and solvent)

TOP

4. Pan sorting (after sieving)

a. Adults - Count or sort the beetles collected in the aluminum pan by brushing adults into a petri dish with a small to medium sized brush. If your sample has a very large number of adults in it, flying beetles can be a problem. (Beetles seem to get more excited and want to fly away when crowded, or when conditions are hot and humid.) You can minimize the problem by first putting all the collected beetles in one or more petri dishes and place lids on the dishes. Then return smaller portions of beetles to the

aluminum pan for sorting a bit at a time.

b. Pupae - If collecting pupae from a jar with a spoon, you can exclude many adults by tilting the jar to one side. Adults will move to the low side, and you can scoop from the center (Be sure to "sterilize" the spoon first by wiping off and rapping it against the table top several times on both sides!). Sieve, then brush adults and larvae into one petri dish, and brush pupae into another dish.

Note: Sorting adults, pupae, or larvae with a brush is easier if accumulations of exuvia (castoff skins) are first removed. One method to remove them is by gently blowing them out of the pan, using a side-to-side and near-to-far sweeping motion with your breath, blowing them into a waste basket. It usually takes 3 to four "sweepings" to get most of the exuvia out. (Be careful to blow gently enough that only exuvia, and some dead adults are blown out --not the live adults, pupae and larvae. Dead beetles and exuvia are lighter than live ones and careful blowing helps to separate them.)

Another way of separating pupae from adults and larvae is to sift the whole jar, place the adults, larvae and pupae (the siftings), into a petri dish or other clean container, then work with small amounts of the siftings. For each lot, blow off the skins, then shake down the adults and pupae, leaving the larvae. Pour the adults and pupae onto a petri dish lid in a covered sieve receiving pan, and let the adults run off, leaving mostly pupae. Exuvia can also be removed by vacuuming the siftings from a quart jar before placing in the aluminum sorting pan.

TOP

5. Use of topping - Topping (coarsely ground wheat) is used to give beetles traction on the flour so they can right themselves if they fall onto their backs (while many beetles in a container can help each other get up, a lone beetle can get stranded on its back and starve to death!). Use topping if:

a. Population density is low due to disease or mutation.

b. Adults have impaired ability to right themselves due to a mutation affecting leg size or shape. For instance, it is wise to use topping with stocks of *Dch-3* since they can't get around as well as beetles with normal sized and shaped front legs, and since they have lower fertility than other strains.

TOP

6. Subculturing schedule - If using a 30°C incubator temperature, subculture heavily used stocks weekly. Other stocks may be subcultured every other week or monthly.

7. Diseased stocks - Diseased stocks should be subcultured every two days to dilute the disease organisms. **Transfer only live beetles!** Dead or moribund beetles should be discarded.

TOP

B. TROUBLE SHOOTING

If a stock is not producing progeny, check the following:

1. Are there any adults still alive? If there are live adults, check to see if they are all males (some disease seems to plug up and kill the females first).
2. Is there evidence of disease.... dead, dried, and sometimes darkened, larvae and pupae? ("Licorice stick" is a good description of dead larvae's appearance. Dead pupae appear discolored and mummified, and are often chewed on by the adults.)
3. Are there mites in with the adults, or clinging to the adults? To differentiate between grain mites, psocids, and parasitic mites, you can look at this [web page](#) to see what grain mites and psocids look like.

- Parasitic mites tend to hang all over the adults, sometimes to the point of giving them a frosted look, and also hide under the wings and elytra. They seem to prefer female beetles, possibly as a way of being near eggs which they may feed on.
- A permanent or long term cure is possible. Follow this link to view the section on parasitic mites in the "Disease & Mites" part of this guide.

TOP

C. TROUBLE PREVENTION

1. Keep all containers of beetles or culture flour closed or covered when not being used or worked with.
2. Bang pans and sieves up-side-down vigorously against wastebasket lid before and after each use to remove any remaining eggs or small larvae.
3. Wipe off and rap spoons and scoops against table top before each use.
4. "Squeegee" sterilize brushes before each use.
5. If beetle adults, larvae or pupae are found on the table top as a result of sieving, discard unless you saw it fall and are 100% certain of its origin! (It helps to begin with a spotless working surface and floor. This helps increase the probability of an accurate recovery of a dropped or spilled beetle. It does not insure against accidentally picking up a "fly-in" in place of the intended beetle!).
6. Don't house beetles in airtight containers, and don't push corks tightly into mouths of vials. Insects need fresh air!

TOP

D. Disease & Mites

Eggs may also be collected on Gold Medal flour (or other equally fine flour), and a new stock begun from the debris-free eggs. Allow the adults to lay eggs on the fine flour for 24 hour periods of time. Each day, collect the eggs by double sieving. This method involves using two sizes of sieves, a #25 and a #50, stacked one on top of the other. The #25 is placed on top, with the #50 between the #25 and the receiving pan. Adults remain on the #25 sieve and can be placed temporarily in a covered, sterilized petri dish. The eggs will be retained on the #50 sieve, and can then be transferred to a clean petri dish. (Alternately, if the two sieves being used are warped and difficult to separate after sifting, egg collection can be done in two separate siftings: separate the adults from the flour using only the #25 sieve first, then sieve out the eggs using only the #50 sieve.) All extraneous material (frass, debris) can then be removed from the collected eggs using a small brush. Put cleaned eggs in jar or bottle of fresh flour for development. This works for ridding a stock of mites as well as disease.

Parasitic mites can easily retard or destroy an otherwise healthy stock. The mites hang all over the adults, sometimes to the point of giving them a frosted look. They seem to prefer females. A permanent or long term cure can be achieved, with a lot of work.

1. Initially, a subset of adults needs to be cleaned. This means putting them on ice and removing the mites with a vacuum probe or aspirator. Mites are persistent, and can also hide (safely) under the elytra.
2. When the beetles are recovered, put them in fine flour with topping for egg collection.
3. Collect the eggs 1-3 days later (depending on the number of adults ovipositing).
4. Now come the hardest part. Put the eggs on some dark paper or other good-contrast surface, under the microscope. With an insect pin and a small vial of ethanol, remove EVERYTHING that is not a plump, healthy egg. Dip the head of the pin in the ethanol, and then blot up the trash. Swish and repeat.
5. Be careful of mites that are feeding on the eggs, as they swell up almost egg-size, and the egg wraps around them as it is depleted. Also look for loose mites roving the surface. Roll the eggs over.
6. Put the now "sterile" eggs in new flour. Keep infested cultures and healthy cultures separate. Disinfect your equipment.

[TOP](#)

E. Developmental Rates of *Tribolium castaneum*

Rearing Temperature	30°C	34°C
Egg	3 days	2 days
Larva	20 days	15 days
Pupa	4 days	3 days
<u>Reproductive Maturation</u>	<u>5 days</u>	<u>4 days</u>
Total time egg to egg	32 days	24 days

(The reproductive lifetime is 3-4 months for females and 4-6 months for males. Isolated males have been known to live for up to a year)

Note: At 22°C, development is much slower.

[TOP](#)

F. Sexing *Tribolium*

Separating the sexes is necessary in order to run a number of genetics tests. Both adults and pupae can be sexed. If the intended cross must be a virgin cross, it is necessary to sex the beetles as pupae to insure no previous mating has taken place. Following are some materials and methods which have worked well in our laboratory, plus suggested alternate materials you might use.

[TOP](#)

1. **Pupae** (Sexing beetles as pupae rather than as adults is easiest since pupae move very little compared to the adults, and do not need to be immobilized by cooling them on ice.)

a. **Materials** (Microscope, light source, working surface, manipulating tools)

Microscope: A stereoscope is needed to sex the pupae. You will want to be able to magnify the pupae by at least 20 - 30X. A zoom lens stereoscope is very handy.

Light source: A good light source will reduce eyestrain if you are going to be looking at many pupae. The best is a fiberoptics light system. It is both a cooler light source than conventional lights, and those on gooseneck pipes can be aimed at exactly the area you need to focus on. We use a fiber optics light with two light pipes. If you use a standard light, be careful not to overheat your pupae by having the light source too close to them.

Working surface: A small plate of a non-static generating material (approx. 3" x 4") is very handy for separating the sexes. We use a 3" x 4" piece of styrofoam backed posterboard. This has a thickness of about 1/4" which makes it easy to pick up, is light weight, and has a smooth surface to work on. We have chosen a deep blue color, since that color provides a good contrast to the color of the pupae. Any dark color will do. Light colors should be avoided because they cause a glare from the lights.

Manipulating tools: A small natural bristle brush can be used to move the pupae on the "plate".

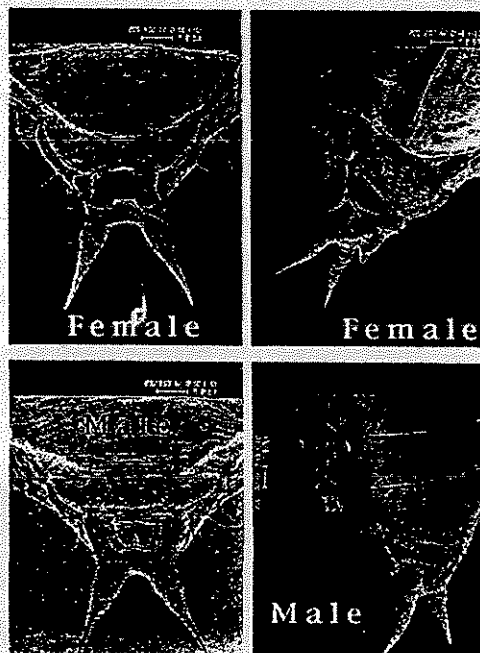
Alternately, a commercial or a homemade vacuum probe can be used to manipulate the pupae. We use a version available through the Jensen Tool catalog, which is hooked up to the vacuum system in our building. (The same probe could also be connected to small electrical vacuum pump). A much simpler version can be made from a plastic drinking straw, a 2-foot piece of flexible rubber tubing (approx. 1/8" internal diameter), and a plastic pipette tip. In this case, the vacuum is supplied by the user's mouth.

Other: Plastic petri dishes or other small containers can be used to temporarily hold the pupae both before and after sexing. These same containers, or small bottles or vials which contain about 1" of flour can be used to hold the pupae until they eclose to adulthood. Any container used for this purpose should have a lid which would keep the wandering adults from escaping. (The lid also needs to have small air holes placed in the top if it is a very tight fitting lid. Petri dish lids do not need air holes.) A plastic funnel is handy for pouring pupae or adults from a sorting pan into a bottle or jar.

TOP

b. Methods

1. Tilt dish and tap some onto the sexing plate.
2. Using a small brush or vacuum probe, line up the pupae in a horizontal line about half way down the plate (have them all facing the same direction, i.e. all heads up or all heads down).
3. As you look at each pupa under the microscope to determine the sex, brush one sex into a new line above the original line, and the other sex into a new line below the original line. When this process is done, you should have two new lines in place of the old one; one with males and one with females. Use the diagrams at right to identify males and females. (Tip: Ignore the two pointed structures at the very end of the pupa - these are the urogomphi, not the genital papillae. The female papillae, which are much larger than those of the male, are two finger-like structures just anterior to the pointed urogomphi. The male papillae are enough smaller that they look like just fingertips rather than fingers.)
4. Double check your work. To do this, it is important to brush one of the sexes off of the plate into a petri dish or other container while you double check the other sex. (Pupae can't walk, but they can roll or squirm, and if you leave both sexes on the plate while double checking them, one could squirm into the wrong line or group, undoing your careful sexing!) Look at each pupae again to verify its sex, placing missexed pupae in the correct group. You might want to check your pupae a third time just to be sure while you're getting the hang of it.
5. Label your containers and place each sex in a separate container with flour to allow them to eclose to adulthood. (You can use them for crosses once the adults have darkened to brown.)



TOP

2. Adults

a. Materials (ice bucket, ice block)

Ice bucket: Any container which can hold crushed ice can be used to precool adult beetles. The

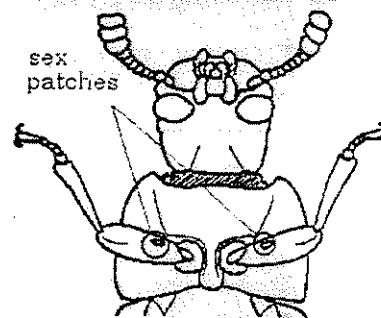
small styrofoam boxes or buckets which are used for picnics or fishing are perfect for this.

Ice block: This is used to keep the beetles immobile while you are looking at them under the microscope. We use small, flat plastic tissue culture bottles which we fill with water about 1/4" at a time until they are mostly full. (Don't fill any container completely full because it will crack or burst when frozen!) Any low-profile container which can hold crushed ice could be used; for instance, a large petri dish full of crushed ice. The pre-cooled beetles are then placed in a smaller, low-profile container (such as a smaller petri dish lid), and this smaller container of beetles is placed on the larger low-profile container filled with ice. You should be able to place this assemblage under your stereoscope. The beetles should remain immobile long enough for you to be able to sex them. (Tip: The ice eventually melts, allowing the beetles to "wake up", so you will want to limit the number of beetles you sex at onetime to a number compatible with the "staying power" of your cooling equipment.)

TOP

b. Methods

1. Collect adults from stock (using methods mentioned above in Wrangling Tips) and place in a covered petri dish or other container for temporary holding. Put the container of beetles on the crushed ice in the ice bucket to precool them before putting them on the ice block.
2. Tap a small number of adults from the petri dish into a smaller flat container on the ice block.
- 3.
4. Line up the adults as with the pupae above, and separate them into two new lines according to sex. Use the diagram at right to distinguish between males and females. The males have a small patch of short bristles on the inside of the first pair of legs, about 1/3 the distance out from the bases. If the patches have picked up flour, they will appear like two domes of flour or flour paste, and will be fairly easy to see. If they have not yet picked up flour, they will appear as slightly darker, textured spots on the legs. (Changing the angle of light or changing the position of the beetle can help make the patches more visible if you are having trouble seeing them.)
5. Recheck the sexes while still on the ice block.
6. Brush each sex into a separate petri dish or other covered holding container until all the beetles are sexed.
7. (Reminder: Work with only a small number of beetles at a time. This will allow you to do the sorting and rechecking before the ice block starts to melt and the beetles wake up and try to walk off!)
8. Use the beetles for your crosses. If they are sexed as adults, they are usually used right away rather than being held in flour like the pupae mentioned above. If you do place them in containers for use later, remember that the females are probably already fertilized and will be producing offspring in their jar.



Male RFB

TOP

Last Edited: August 13, 1998



Linkage Groups

- [LG 1-4 \(large image\)](#)
- [LG 5-9 \(large image\)](#)

- [LG 1 = X](#)
- [LG 2](#)
- [LG 3](#)
- [LG 4](#)
- [LG 5](#)
- [LG 6](#)
- [LG 7](#)
- [LG 8](#)
- [LG 9](#)

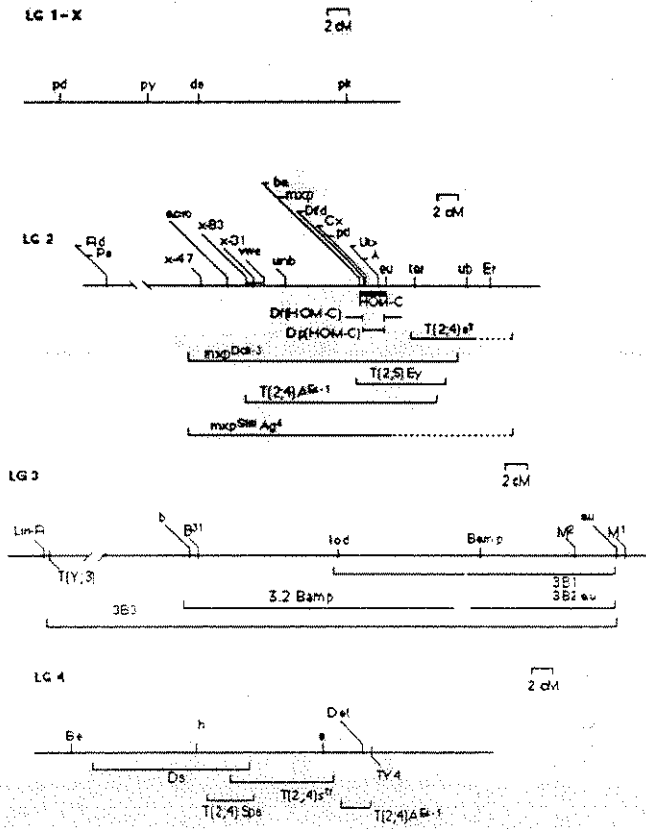
Last Edited: August 13, 1998





Tribolium castaneum chromosomes 1-4

Select a Linkage Group:



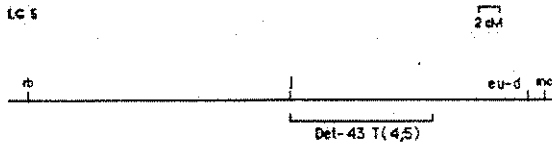
Last Edited: August 13, 1998



Tribolium castaneum chromosomes 5-9

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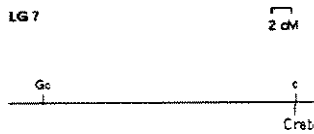
LC 5



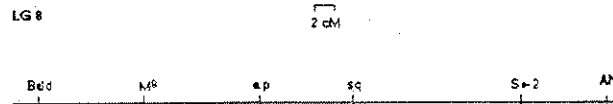
LC 6



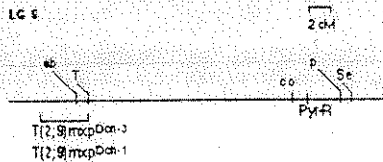
LG 7



LG 8



LG 9

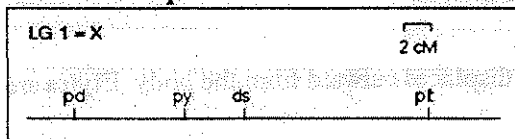


Last Edited: August 13, 1998



Tribolium castaneum Linkage Group 1

Clickable Map



Mutant Name/note

ds	displaced sternellum
pd	paddle
plt	platinum eye
py	pygmy

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ds-X (displaced sternellum, X-linked)

- **Structure affected:** Sternellum on ventral prothorax and elytra
- **Linkage Group:** 1=X
- **Origin:** Spontaneous, from Stm,Cx5/Ey; s/s stock
- **Description:** The sternellum is shortened and slightly displaced outward from the body. Elytra are short, exposing the dorsal tip of the abdomen.

Last Edited: August 13, 1998



pd (paddle)

- **Structure affected:** Antennae.
- **Linkage Group:** $I = X$
- **Origin:**
- **Description:** Antennal segments are fused, giving antenna a "canoe paddle" appearance.

Last Edited: August 13, 1998



plt (platinum eye)

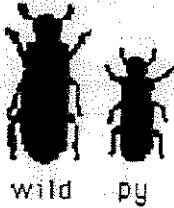
- **Structure affected:** eye
- **Linkage Group:** 1=X
- **Origin:** spont.
- **Description:** white-eye

Last Edited: August 13, 1998



py (pygmy)

- **Structure affected:** global
- **Linkage Group:** 1=X
- **Origin:** spontaneous
- **Description:** Body mass reduced by one-half. All proportions normal.

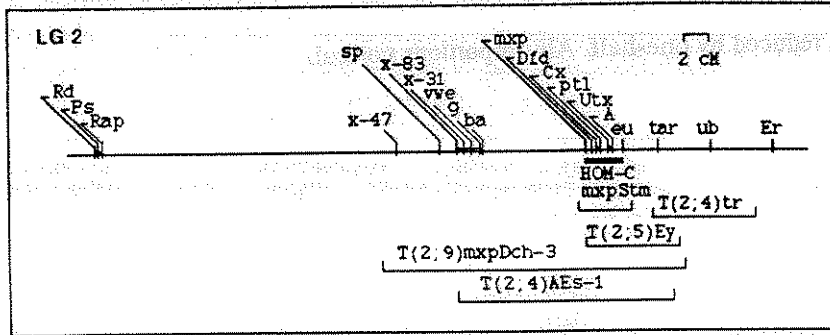


Last Edited: August 13, 1998



Tribolium castaneum Linkage Group 2

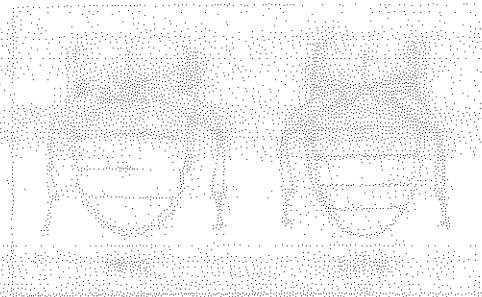
Clickable Map



Mutant	Name/note
A	Abdominal
ba	broken antenna
Cx	Cephalothorax
Dfd	Deformed
Er	Eyes reduced
eu	extra urgomphi
glossy	glossy cuticle
mxp	maxillopedia
mxpStm	Stumpy
Ps	Pinched sternellum
ptl	prothoraxless
Rap	Recurved anterior pronotum
Rd	Reindeer
sp	shoulder pads
T(2;4)AEs-1	Extra sclerite
T(2;4)tr	tremblor
T(2;5)Ey	Eyeless
T(2;9)mxdCh-3	Dachsund
tar	tar
ub	unbuckled

<u>Utx</u>	Ultrathorax
<u>vwe</u>	vestigial wings and elytra
<u>x-31</u>	lethal
<u>x-47</u>	lethal
<u>x-83</u>	lethal

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A (Abdominal)

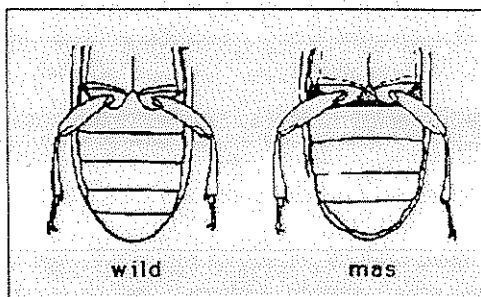
LG 2

ES-1 (Extra sclerite)

- **Structure affected:** Ventral abdominal segment 2.
- **Linkage Group:** T(2;4)
- **Origin:**
- **Description:**
 - Homeotic transformation of ventral part of abdominal segment 2 (normally forming the socket of the coxae of the third pair of legs) towards segment 3.
 - Excellent crossover suppressor and balancer (second only to Ey in usefulness).
 - No crossover suppr. between HOM-C and Rd.
 - Ag/Es1 spontaneously generates viable Df(HOM-C) gametes at a frequency of 1/1000 when outcrossed (see Ag).
 - Generated euD when outcrossed (BB, p. 32).
 - Translocation demonstrated cytologically, both cis and trans with Spa.

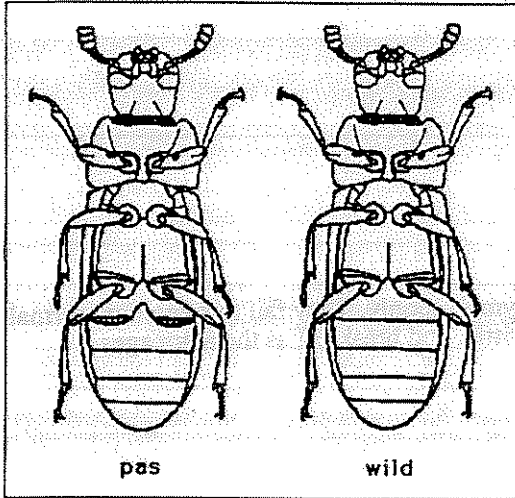
mas (missing abdominal sternite)

- **Structure affected:** 3rd abdominal sclerite
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** mas is an abdominal 3 to abdominal 2 transformation.



pas (pointed abdominal sternite)

- **Structure affected:** 4th abdominal sclerite
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** pas is an abdominal 4 to abdominal 3 transformation.

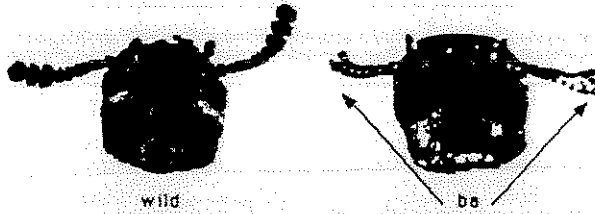


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ba (broken antenna)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** Ethylmethanesulfonate-induced.
- **Description:** Antennae appear defective in hemolymph supply. They fail to sclerotize normally after adult eclosion, then become melanotic and brittle, and break off as the adult ages.



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Cx (Cephalothorax)

LG 2

Cx

- **Structure affected:** Labium and prothorax.
- **Linkage Group:** 2
- **Origin:** Gamma radiation-induced.
- **Description:** Complex homeotic transformations of labium and prothorax.



Ag

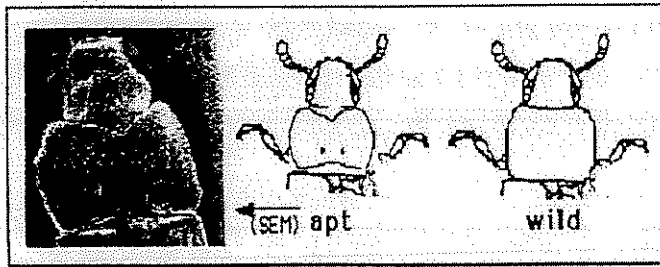
- **Structure affected:** Antenna.
- **Linkage Group:** 2
- **Origin:** G, GA-1.
- **Description:** A galea-like projection is found on the basal segment (scape) of the antenna. (Ag/Es1 spontaneously generates viable Df(HOM-C) gametes at a frequency of 1/1000 when outcrossed. These uncover Cx, ptl, Utx and A, but complement mxp and eu).

apt

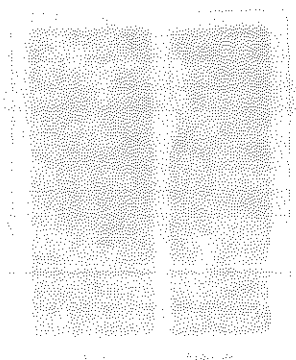
- **Structure affected:** dorsal pronotum.
- **Linkage Group:** 2
- **Origin:** spontaneous; A. Sokoloff, Berkeley, CA.
- **Description:** The dorsal pronotum is taking on characteristics of the dorsum of the next most posterior segment, the mesothorax (T2). In its strongest expression, the anterior margin of the pronotum has a large midline indentation, and the posterior midline is beginning to look like the T2 scutellum (the little roundly triangular structure just posterior to the pronotum).

Tribolium Cx (Cephalothorax)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg2/2cx.html>



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Dfd (Deformed)

- **Structure affected:** No gene-specific mutants known. Identified as molecular homolog of Drosophila Deformed gene.
- **Linkage Group:** 2
- **Origin:**
- **Description:**

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Er (Eyes reduced)

- **Structure affected:** Eyes and surrounding head capsule
- **Linkage Group:** 2
- **Origin:** Gamma irradiation of Ag mutant
- **Description:**
 - First recovered as an Mc on an Ag chromosome. The two genes later segregated away from each other.
 - The male first recovered was F1 of Ag/Es.
 - Head is reduced posterior to the genal shelf. Most of the dorsal component of the eye is missing. Ventral expression of eye is more complete, though somewhat reduced due to a reduction of that portion of the head capsule itself. **In contrast to Ey, Er most often has good bilateral expression of the ventral eye.**

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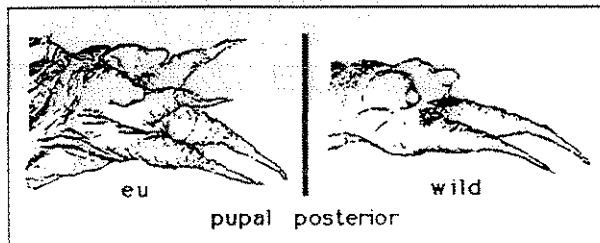
eu (extra urigomphi)

LG 2

Description/Notes

eu (extra urigomphi)

- **Structure affected:** urogomphi (paired "horns" at posterior tip of abdomen of larvae and pupae).
- **Linkage Group:** 2
- **Origin:** Spontaneous.
- **Description:** Supernumerary pair of urogomphi develop via homeotic transformation of abdominal segment 9 toward 10.



euD (dominant allele)

- **Structure affected:** Posterior abdominal segments (A10 & A11)
- **Linkage Group:** T(2; 5)
- **Origin:** Gamma irradiation of Rd.mas,p males, Beeman Lab, USGMRL, Manhattan, KS
- **Description:**
 - Translocation: T(2;5), confirmed cytologically
 - Male sterile
 - extra urogomphi (unilaterally or bilaterally), found in larvae and pupae
 - Genital papillae of male and female pupae are abnormal
 - Male aedeagus non-rotated, rendering males functionally sterile
 - Female ovipositors with split lateral sclerite, causing dorsal-ventral flattening of ovipositor
 - euD/eu beetles have reiterated genital papillae in the females, and lack an aedeagus in males
 - Appears to be hyper-mutator stock

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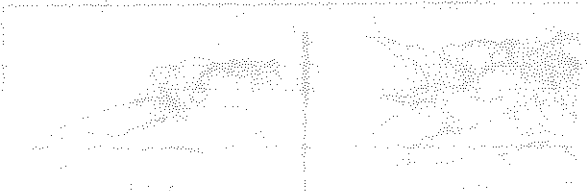
Tribolium; g (glossy)

<http://bru.usgmlr.ksu.edu/beeman/tribolium/lg2/2g.html>

g (glossy)

- **Structure affected:** cuticle, global
- **Linkage Group:** 2
- **Origin:** EMS mutagenesis of sooty. Dch3 /Ey chromosome extraction (Dch3/unb)
- **Description:**
 - Color has been found to be "light pumpkin", compared to wild-type.
 - The exterior surface, with the exception of elytra, has a higher reflectivity than normal, due to a reduction of the "surface microsculpture" between the setiferous pits.
 - The elytra are sometimes divergent (usually not).
 - T1 epimera tend to be slightly incomplete, not quite extending as far under the sternellum as with wild-type.
 - Ventral sclerites also have an imperfect T1-T2 juncture at the T2 coxae.

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(tribolium) (glossy) (larva)

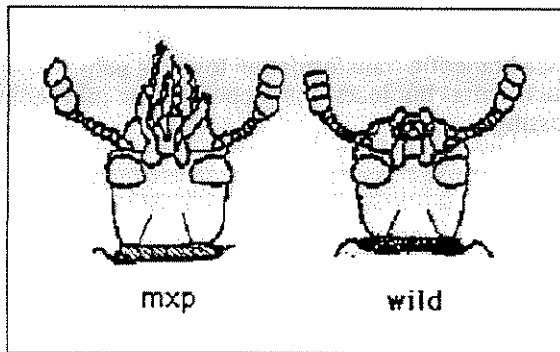


mxp (maxillopedia)

LG 2

mxp (maxillopedia)

- **Structure affected:** maxillary & labial palps
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** mxp/mxp causes transformation of the labial and maxillary palps into legs.



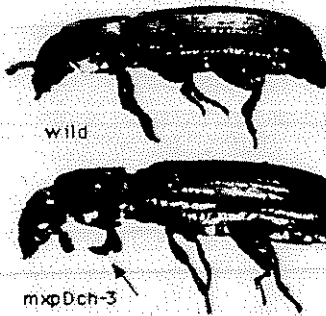
Dch-1 (Dachshund)

- **Structure affected:** Antennae and legs.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** Dominant, gain-of-function (GOF). Antennae & legs shorter than normal (partially transformed towards palp) in heterozygotes. Dch-1/Dch-3 heteroallelic adults have complete and dramatic transformation of legs into palps.

Dch-3 (Dachshund)

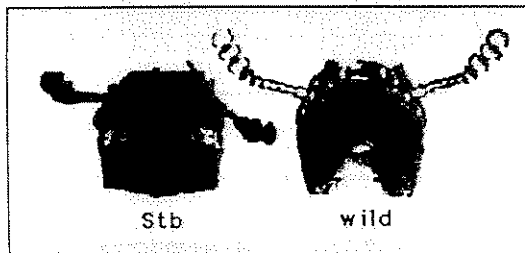
- **Structure affected:** Prothorax.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** GOF. Prothoracic legs are dramatically reduced. Remainder of prothorax is reduced to a lesser extent, presumably via a homeotic transformation toward labial or maxillary segment. Effect restricted to prothorax. Antennae are normal.

Tribolium: mxp (maxillopedia)

<http://bru.usgml.ksu.edu/beeman/tribolium/lg2/2mxp.html>

Stb (Stubby)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** EMS, GA-1
- **Description:** GOF. Funicle of antennae reduced via homeotic transformation towards palp. Sometimes behaves as an X-linked trait (upon outcrossing) but loses this property upon inbreeding, only to regenerate the property again upon outcrossing.



Stm (Stumpy)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** Ethylmethane sulfonate.
- **Description:** GOF. Club and funicle of antennae reduced.
 - Near-lethal with lethal mxp alleles
 - Complements viable mxp.
 - Homozygous stock is fertile.
 - Other stocks balanced with Stm are only slightly leaky.
 - Lab-S RFLP matches Stm RFLP in a clone from the A gene, so Stm probably is a Lab-S chromosome.
 - No translocation found cytologically by Giovanni Mocelin.

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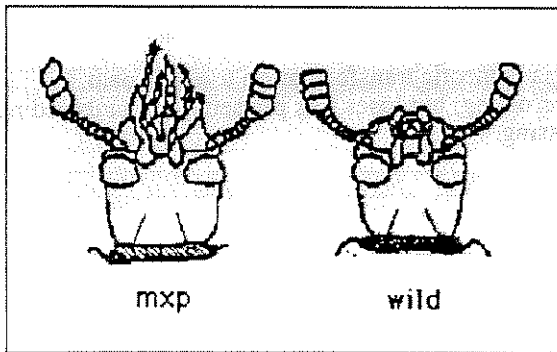


mxp (maxillopedia)

LG2

mxp (maxillopedia)

- **Structure affected:** maxillary & labial palps
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** mxp/mxp causes transformation of the labial and maxillary palps into legs.



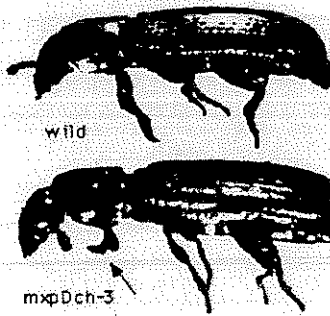
Dch-1 (Dachshund)

- **Structure affected:** Antennae and legs.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** Dominant, gain-of-function (GOF). Antennae & legs shorter than normal (partially transformed towards palp) in heterozygotes. Dch-1/Dch-3 heteroallelic adults have complete and dramatic transformation of legs into palps.

Dch-3 (Dachshund)

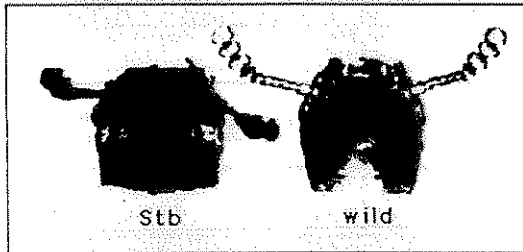
- **Structure affected:** Prothorax.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** GOF. Prothoracic legs are dramatically reduced. Remainder of prothorax is reduced to a lesser extent, presumably via a homeotic transformation toward labial or maxillary segment. Effect restricted to prothorax. Antennae are normal.

Tribolium; mxp (maxillopedia)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg2/2mxp.html>

Stb (Stubby)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** EMS, GA-1
- **Description:** GOF. Funicle of antennae reduced via homeotic transformation towards palp. Sometimes behaves as an X-linked trait (upon outcrossing) but loses this property upon inbreeding, only to regenerate the property again upon outcrossing.



Stm (Stumpy)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** Ethylmethane sulfonate.
- **Description:** GOF. Club and funicle of antennae reduced.
 - Near-lethal with lethal mxp alleles
 - Complements viable mxp.
 - Homozygous stock is fertile.
 - Other stocks balanced with Stm are only slightly leaky.
 - Lab-S RFLP matches Stm RFLP in a clone from the A gene, so Stm probably is a Lab-S chromosome.
 - No translocation found cytologically by Giovanni Mocelin.

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Ps (Pinched sternellum)

- **Structure affected:** T1 sternellum and dorsum, T3 antecoxal sutures, basal segments of legs, antennae and maxillary palps.
- **Linkage Group:** 2
- **Origin:** Reversion of Skill / Stm balanced mutants
- **Description:**
 - The original beetle was Ps and Stm.
 - Setae are commonly found on coxae, as are setae and spikes on the antennal scape and maxillary palps.
 - Prothorax has dorsal dents and bulges, and both a dorsal and a ventral anterior midline dip.
 - T3 antecoxal sutures are disrupted about 2/3 the way out from midline.

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Tribolium: ptl (prothoraxless)

<http://bru.usgmrl.ksu.edu/beeman/tribolium/lg2/2ptl.html>

ptl (prothoraxless)

- **Structure affected:** prothorax
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** pronotum is reduced in size, and prothoracic legs are stunted and malformed.

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Rap (Recurved anterior pronotum)

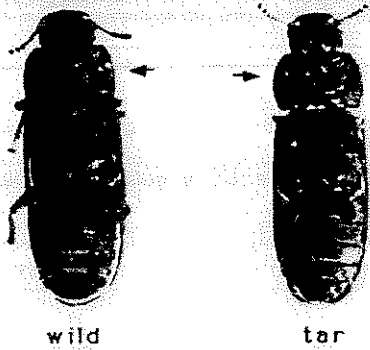
- **Structure affected:** Dorsal pronotum (T1)
 - **Linkage Group:** 2
 - **Origin:** Spontaneous mutant found in the Ga-1 (wild-type) stock
 - **Description:**
 - Dorsal pronotum has anterior midline dip, bilateral shallow dents, and enlarged antero-lateral margins.
 - This dominant is homozygous viable. The phenotype very strong in homozygotes, but easily recognizable in heterozygotes.
-

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tar

- **Structure affected:** anterior quinone glands
- **Linkage Group:** 2
- **Origin:** EMS
- **Description:** The anterior stink glands are darkly pigmented, usually a red-brown to purple-brown rather than the normal clear yellow, and seem incapable of secretion. Posterior stink glands are not affected.

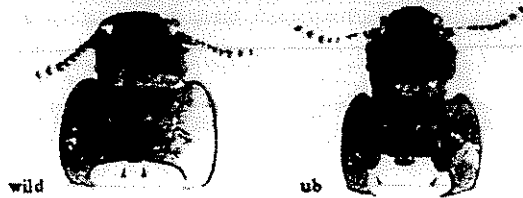


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ub (unbuckled)

- **Structure affected:** Prothorax, appendages.
- **Linkage Group:** 2
- **Origin:** Beeman Lab, USGMRL, Manhattan, KS. Spontaneous mutant recovered from F2 of ab X pas30,p cross.
- **Description:**
 - T1 epimera deflected ventro-posteriorly.
 - T2 metepimera are displaced from the plane of surrounding sclerites ("flaps").
 - Antenna, legs, female genital stylii, and mouthparts are more slender than those of wild-type.
 - Genital papillae of female pupae have a small, round sclerotization at the ventral midline anterior margin. Expression ranges from a barely visible dot to an enlarged structure taking up the full middle third of the papillar base. These larger ones usually have a darker sclerotized center. (Larvae have not been seen with these dots).



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Utx (Ultrathorax)

- **Structure affected:** Elytra, mesothorax (T2)
- **Linkage Group:** 2
- **Origin:** EMS mutagenesis using Lab-S or GA-1 wild-type strains.
- **Description:**
 - Dominant with warped elytra (gain-of-function, based on dosage analysis) and protruding T2 epimera ("flaps").
 - Recessive A1 to T3 transformation (loss-of-func?) seen in homozygous embryos.
 - Utx1 acts as a lethal only in the Utx1/Es1 bal. stock. Using Stm, Ey or mxpNG-1 as balancer, apparent Utx1 homozygotes are generated. These have "membranous antecoxae" (recessive) phenotype in addition to warped elytra and flaps. We cannot reconcile this phenomenon with the lethal embryo (A1 to T3) phenotype seen in the Utx1/Es1 stock. Utx1 derived from "homozygous" stock regains lethality when placed opposite Es1!
 - Recombination with apt, but not A.

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vwe (vestigial wings and elytra)

- **Structure affected:** wings & elytra
- **Linkage Group:** 2
- **Origin:** EMS
- **Description:** Wings & elytra of pupae are extremely reduced & vestigial. Pupae unable to eclose. Occasional adult escapers with well-differentiated (but miniature) wings & elytra.

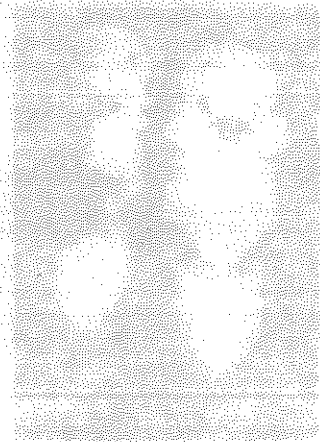


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This is a lethal mutation lacking a visible phenotype

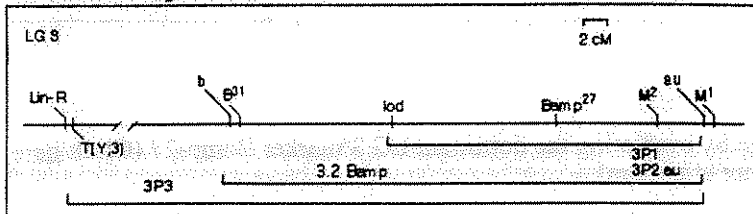
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Tribolium castaneum Linkage Group 3

Clickable Map



Mutant	Name
3.2 Bamp	(3P2 based crossover suppressor w/ 40+ cM range)
3P1	(Bamp-27 based crossover suppressor w/ 30 cM range)
3P2	(Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au)
3P3	(Bamp-27 based crossover suppressor w/ 45+ cM range)
au	aureate
b	black body - b itself is incompletely recessive, but other alleles are completely recessive.
Bamp-27	Blunt abdominal and metathoracic points
Bamp-31	Blunt abdominal and metathoracic points
Lin-R	Lindane resistance
lod	light ocular diaphragm
M¹	Medea
M²	Medea
T(Y;3)	Translocation

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3B. . . family of Bamp-27 based LG 3 crossover suppressors

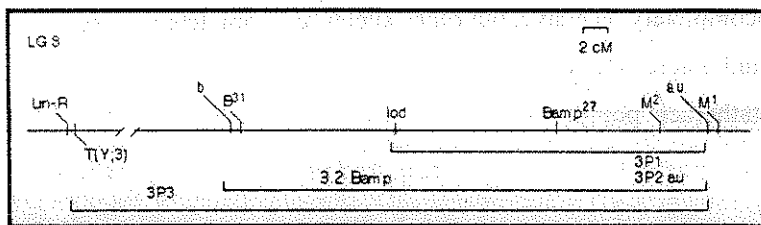
LG 3

Description/notes

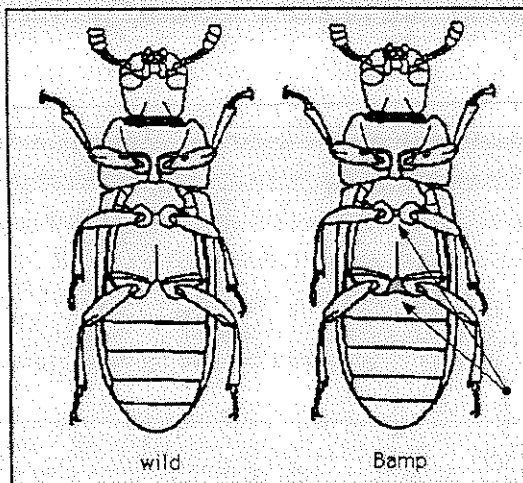
These crossover suppressors were made by irradiating the dominant LG 3 marker Bamp-27 (Click [here](#) to go to the Bamp-27 page, or see picture at bottom).

- **3P1** has the dominant Bamp phenotype, and covers approximately 30 cM (See map below).
- **3P2** is essentially 3P1, with a new or recombined recessive au mutation attached (Click [here](#) to go to the au page, or see picture at bottom).
- **3P3** is one of the largest crossover suppressors available in *Tribolium*, covering approximately 15% of the genome. It allows approximately 1% single or double recombinants along most of its length. It has only the dominant Bamp phenotype (See map below)
- **3.2 Bamp** is a radiation-extended 3P2 crossover suppressor. Therefore it has the dominant Bamp and the recessive au. It covers about 40 cM. In the presence of 3.2, M1 and b recombine 2%, apparently at the b end of the 3.2 Bamp CS.

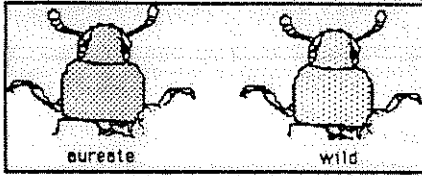
LG 3 linkage map



Bamp-27 phenotype (dominant)



au phenotype (recessive)



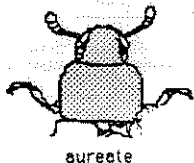
Last Edited: August 13, 1998





au (aureate)

- **Structure affected:** entire surface of cuticle
- **Linkage Group:** 3
- **Origin:** spontaneous
- **Description:** setae (hairs) on general body surface are 2-3 times as dense as normal, giving a frosted or hairy appearance.



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b (black)

- **Structure affected:** cuticle
- **Linkage Group:** 3
- **Origin:** spontaneous
- **Description:** black body color.

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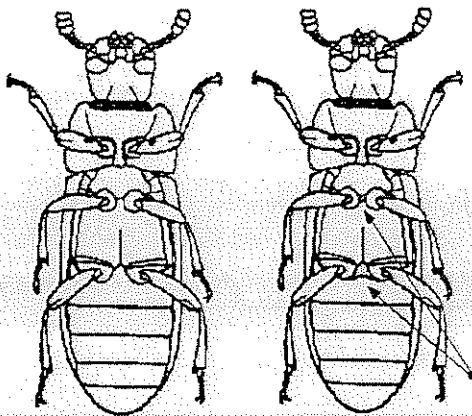


Bamp-27 (Blunt abdominal and metathoracic projections)

LG 3

Bamp-27 (Blunt abdominal and metathoracic projections)

- **Structure affected:**
- **Linkage Group: 3**
- **Origin:** Gamma irradiation of Ga-1
- **Description:**
 - The anterior midline projection of the T3 (third thoracic) sternum, which usually forms a firm junction with the posterior midline of the T2 sternum at the T2 legs, lacks its usual point, leaving a small gap at the T2-T3 juncture.
 - The anterior midline projection of the A3 (third abdominal) sternum, which usually forms a firm junction with the posterior midline of the T3 sternum at the T3 legs, has a reduced point, leaving a noticeable gap at the T3-A3 juncture.
 - This was the irradiated chromosome for the 33B ... family of LG 3 crossover suppressors.
 - Very fertile.
 - No translocations found by cytology.



wild

Bamp

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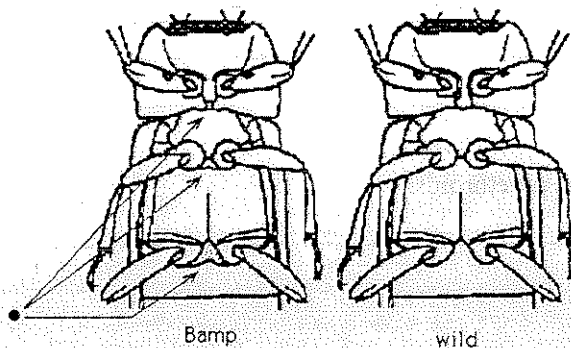
Bamp-31 (Blunt abdominal and metathoracic projections)

LG 3

Bamp-31 (Blunt abdominal and metathoracic projections)

Blunt abdominal and metathoracic points

- **Structure affected:** T1 Sternellum and A3 point.
- **Linkage Group:** 3
- **Origin:** ChrE (lethal)
- **Description:**
 - The most obvious and easily identified feature of Bamp31 is its shortened, narrowed T1 sternellum. The blunted abdominal and metathoracic projections are more subtle than that found in other Bamp alleles. This allele also commonly has disrupted gular sutures and divergent, "rumped" elytra.



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Insecticide Resistance

Lin-R (Lindane resistance)

- **Linkage Group:** 3
 - **Origin:** spontaneous
 - **Description:** resistant to lindane and cyclodiene insecticides because of a mutation in the GABA(A) receptor.
-

Rmal (Malathion resistance)

- **Linkage Group:** 6
 - **Origin:** spontaneous
 - **Description:** resistant to malathion and phenthoate insecticides because of a modification in a carboxylesterase enzyme.
-

Pyr-R (Pyrethroid resistance)

- **Linkage Group:** 9
 - **Origin:** spontaneous
 - **Description:** resistant to alpha-cyano synthetic pyrethroids.
-

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lod (light ocular diaphragm)

- **Structure affected:**
- **Linkage Group:** 3
- **Origin:** Spontaneous
- **Description:** Ocular diaphragm (ring around outer perimeter of eye) is unpigmented rather than the normal black pigmentation. Can only be recognized in mutant eye color background, such as pearl, ruby, etc.

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Medea

MEDEA is an acronym for Maternal-Effect Dominant Embryonic Arrest



Medea-killed larvae

Medea factors all share several characteristics:

- They breed true through the female line.
- They segregate in the male.

When a heterozygous Medea female (M/+) is crossed to a wild type male (+/+), the M gene and its homolog segregate normally. However, all progeny that do not inherit the Medea allele die at or shortly after egg hatch. The lethality is maternal, but the "rescue" is zygotic. The rescuing M allele can be inherited from either parent.

There have been four well-studied Medea factors. Of these, two (M-1 and M-4) are currently maintained at the Tribolium Stock Center. Almost all M strains in the field carry M-4. Of these, about a third also carry M-1. M-4 is the only Medea factor present in North American and European strains, being found in about half of them. Australian and Indian strains are almost devoid of Medea factors. South American, Asian, and African strains often have 2 or more M factors.

M¹ (Medea)

- **Linkage Group:** 3
- **Description:** The first and most-studied Medea factor.

M² (Medea)

- **Linkage Group:** 3
 - **Description:** This Medea factor faded away and is no longer detectable.
-

M³ (Medea)

- **Linkage Group:** 8
 - **Description:** This Medea factor faded away and is no longer detectable.
-

M⁴ (Medea)

- **Linkage Group:** Unknown
 - **Description:** This Medea has an interesting distribution within the United States.
-

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T(Y;3)

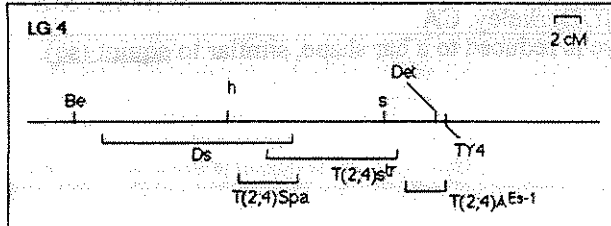
- **Structure affected:** None
 - **Linkage Group:** T(Y;3)
 - **Origin:** gamma-induced on a Ga-1 chromosome
 - **Description:**
 - Translocation between LG3 and the Y chromosome
 - Mutant always carried by males.
 - Translocation demonstrated cytologically.
-

Last Edited: August 13, 1998



Tribolium castaneum Linkage Group 4

Clickable Map



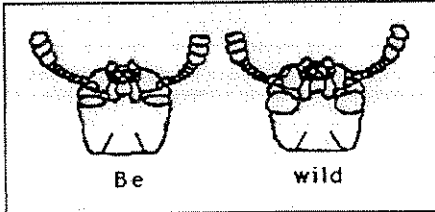
Mutant	Name/note
<u>Be</u>	Bar eye
<u>h</u>	hazel eye
<u>s</u>	sooty
<u>Det-43</u>	Divergent elytral tips
<u>T(Y;4)</u>	Translocation
<u>Ds</u>	Displaced sternellum
<u>T(2;4)Spa</u>	Spatulate
<u>T(2;4)tr</u>	tremblor
<u>T(2;4)AEs-1</u>	Eyeless

Last Edited: August 13, 1998



Be (Bar eye)

- Structure affected: eye
- Linkage Group: 4
- Origin: Spontaneous, dominant. A. Sokoloff, Berkeley, CA.
- Description: The number of facets in the eye is reduced: a bar shape, similar to squint (sq).



Last Edited: August 13, 1998



h (hazel)

- **Structure affected:** eye color
- **Linkage Group:** 4
- **Origin:** spontaneous
- **Description:** hazel or tan eye color, allelic with "white"

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Tribolium: s (sooty)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg4/4s.html>

s (sooty)

- **Structure affected:** cuticle
- **Linkage Group:** 4
- **Origin:** spontaneous
- **Description:** cuticle is dark blackish brown, instead of the normal rust-red.

Last Edited: August 13, 1998



Det-43 (Divergent elytral tips)

- **Structure affected:** Elytra
- **Linkage Group:** T(4; 5)
- **Origin:** Gamma-induced on a GA-1 chromosome, Beeman Lab, US Grain Marketing Research Lab., Manhattan, KS.
- **Description:**
 - Elytra are divergent at the tips and have a characteristic "knob and bend" at about 2/3 their length at the lateral margins. This characteristic is fully penetrant.
 - It is a T(4;5) translocation.

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Tribolium T(Y;4)

[http://bru.usgmr1.ksu.edu/beeman/tribolium/lg4/4t\(y;4\).html](http://bru.usgmr1.ksu.edu/beeman/tribolium/lg4/4t(y;4).html)

T(Y;4)

- **Structure affected:** None
 - **Linkage Group:** T(Y;4)
 - **Origin:** gamma-induced on a Ga-1 chromosome
 - **Description:**
 - Translocation between LG4 and the Y chromosome
 - Translocation demonstrated cytologically.
-

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Ds (displaced sternellum)

- **Structure affected:** T1 (prothoracic) sternellum, elytra, global
- **Linkage Group:** 4
- **Origin:** Spontaneous, from Peter Dawson's lab, Oregon State Univ.
- **Description:**
 - Sternellum is shortened and displaced outwardly from the body wall.
 - The elytra are slightly shortened.
 - Length of appendages and of the overall body are slightly reduced.
 - Reduces the recombination frequency between h and s from around 17% to 7.7%.
 - No evidence of chromosome translocation detected in tests for pseudolinkage or upon cytological exam.
 - Most likely associated with an inversion with one breakpoint between h and s and the other breakpoint on the opposite side of h.

Last Edited: August 13, 1998



Spa (Spatulate antenna)

- **Structure affected:** Antenna
 - **Linkage Group:** T(2; 4)
 - **Origin:** Alexander Sokoloff lab, Berkeley, CA
 - **Description:** Antennal club and funicle fusion, giving antenna a shortened look.
-

Last Edited: August 13, 1998



tr (tremblor)

- **Structure affected:** None (behavior affected)
- **Linkage Group:** T(2; 4)
- **Origin:** EMS mutagenesis of sooty
- **Description:**
 - **Translocation:** t(2;4)
 - Originally, adults had a tremorous gait. This trait has lessened with time.
 - The stock remains balanced with Es, but while balanced with Ey, viable homozygotes were recovered which are sooty colored.

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A (Abdominal)

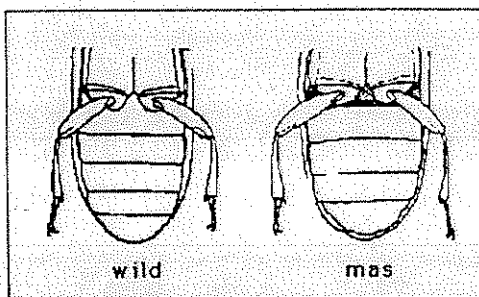
LG2

ES-1 (Extra sclerite)

- **Structure affected:** Ventral abdominal segment 2.
- **Linkage Group:** T(2;4)
- **Origin:**
- **Description:**
 - Homeotic transformation of ventral part of abdominal segment 2 (normally forming the socket of the coxae of the third pair of legs) towards segment 3.
 - Excellent crossover suppressor and balancer (second only to Ey in usefulness).
 - No crossover suppr. between HOM-C and Rd.
 - Ag/Es1 spontaneously generates viable Df(HOM-C) gametes at a frequency of 1/1000 when outcrossed (see Ag).
 - Generated euD when outcrossed (BB, p. 32).
 - Translocation demonstrated cytologically, both cis and trans with Spa.

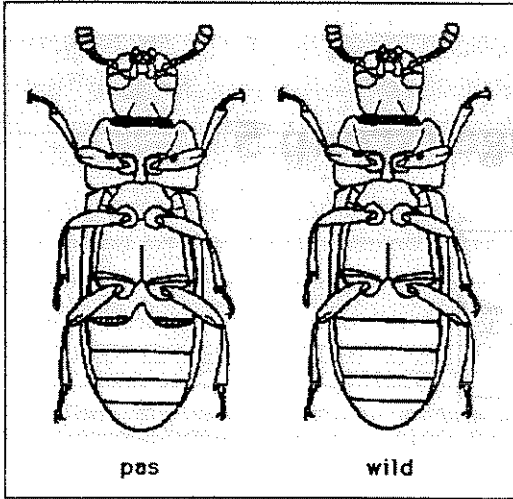
mas (missing abdominal sternite)

- **Structure affected:** 3rd abdominal sclerite
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** mas is an abdominal 3 to abdominal 2 transformation.



pas (pointed abdominal sternite)

- **Structure affected:** 4th abdominal sclerite
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** pas is an abdominal 4 to abdominal 3 transformation.

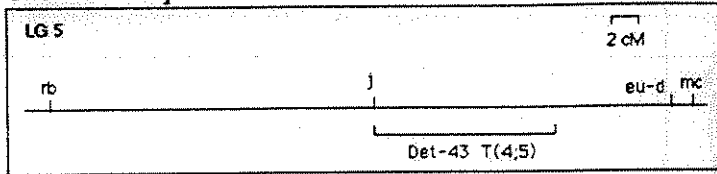


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Tribolium castaneum Linkage Group 5

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Mutant	Name/note
<u>Det-43</u>	Divergent elytral tips
<u>eu-D</u>	extra urgomphi
<u>j</u>	jet
<u>mc</u>	microcephalic
<u>T(2;5) Ey</u>	Eyeless
<u>rb</u>	ruby

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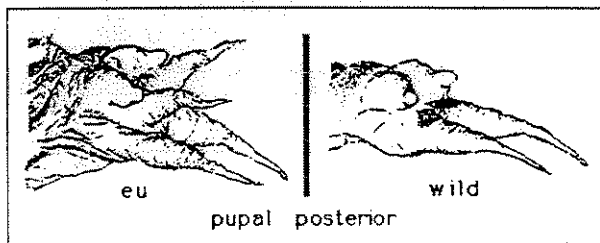
eu (extra urigomphi)

LG.2

Description/Notes

eu (extra urigomphi)

- **Structure affected:** urogomphi (paired "horns" at posterior tip of abdomen of larvae and pupae).
- **Linkage Group:** 2
- **Origin:** Spontaneous.
- **Description:** Supernumerary pair of urogomphi develop via homeotic transformation of abdominal segment 9 toward 10.



euD (dominant allele)

- **Structure affected:** Posterior abdominal segments (A10 & A11)
- **Linkage Group:** T(2, 5)
- **Origin:** Gamma irradiation of Rd,mas,p males, Beeman Lab, USGMRL, Manhattan, KS
- **Description:**
 - Translocation: T(2;5), confirmed cytologically
 - Male sterile
 - extra urogomphi (unilaterally or bilaterally), found in larvae and pupae
 - Genital papillae of male and female pupae are abnormal
 - Male aedeagus non-rotated, rendering males functionally sterile
 - Female ovipositors with split lateral sclerite, causing dorsal-ventral flattening of ovipositor
 - euD/eu beetles have reiterated genital papillae in the females, and lack an aedeagus in males
 - Appears to be hyper-mutator stock

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Tribolium: j (jet body)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg5/5j.html>



j (jet body)

- **Structure affected:** Cuticle, global
- **Linkage Group:** 5
- **Origin:** Alexander Sokoloff lab, Berkeley, CA
- **Description:** Jet-black body color

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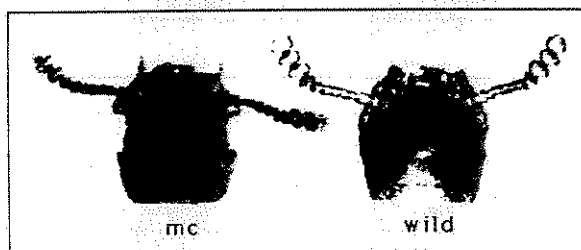


mc (microcephalic)

LG 5

mc (microcephalic)

- **Structure affected:** eyes and head
- **Linkage Group:** 5
- **Origin:** spontaneous (from Sokoloff)
- **Description:**
 - Width of head is reduced posterior to genal shelf.
 - Eyes are variably reduced, ranging from slight reduction in eye size and number of facets, to complete reduction with no facets. Not bilaterally uniform.
 - One strain occasionally has an "eye-growth" which appears on a sclerotized encroachment into the anterior edge of the eye. The growth ranges from very small and fine, and often appears segmented in its largest and strongest expression.



Ey (Eyeless)

- **Structure affected:** eyes and head
- **Linkage Group:** T(2, 5)
- **Origin:** Gamma irradiation of GA-1 (wild-type).
- **Description:**
 - Dominant allele of mc on LG 5, with a similar range of expression (see mc). Ey is a T(2;5).
 - The lethality seems to be associated with LG 2, since Dp(2)/Ey/Ey is viable (where Dp is derived from mxpDch/Es1).
 - Ey is a good HOM-C balancer, (good crossover suppression, fertile, fully penetrant, heterozygotes extremely viable, homozygous lethal). Only one recombinant (out of ca. 2000) has been observed, placing Ey closer to the mxp-apt region than to the A region.

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Tribolium: rb (ruby eyes)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg5/5rb.html>



rb (ruby eyes)

- **Structure affected:** Eyes
- **Linkage Group:** 5
- **Origin:** Alexander Sokoloff lab, Berkeley, CA
- **Description:** Reddish eye color.

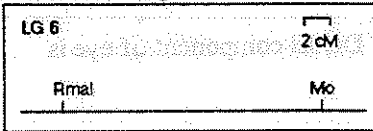
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Tribolium castaneum Linkage Group 6

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Mutant Name/note

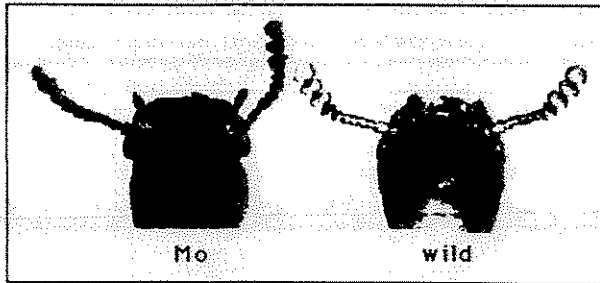
- Mo Micro-ophthalmic
- Rmal Resistance to malathion

Last Edited: August 13, 1998



Mo (Micro-ophthalmic)

- **Structure affected:** Head and eye
- **Linkage Group:** 6
- **Origin:** Alexander Sokoloff, Univ. California at Berkeley
- **Description:** Width of head capsule reduced behind genal shelf. Dorsal component of eye is reduced or missing. Good bilateral expression.



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Insecticide Resistance

Lin-R (Lindane resistance)

- **Linkage Group:** 3
 - **Origin:** spontaneous
 - **Description:** resistant to lindane and cyclodiene insecticides because of a mutation in the GABA(A) receptor.
-

Rmal (Malathion resistance)

- **Linkage Group:** 6
 - **Origin:** spontaneous
 - **Description:** resistant to malathion and phenthoate insecticides because of a modification in a carboxylesterase enzyme.
-

Pyr-R (Pyrethroid resistance)

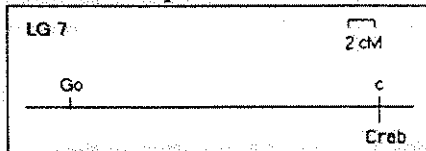
- **Linkage Group:** 9
 - **Origin:** spontaneous
 - **Description:** resistant to alpha-cyano synthetic pyrethroids.
-

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Tribolium castaneum Linkage Group 7

Clickable Map



Mutant Name/note

c chestnut eye

Crab Crab legs

Go Goliath

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c (chestnut eye)

- **Structure affected:** Eye
- **Linkage Group:** 7
- **Origin:** Alexander Sokoloff, University of California at Berkeley
- **Description:** Red-brown colored eye

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Crab

- **Structure affected:** legs
- **Linkage Group:** 7
- **Origin:** EMS mutagenesis, 1986 (Beeman lab, Manhattan, KS)
- **Description:**
 - Tibia shortened, thickened, and bowed, giving them a crab-like look.
 - Male "sex patches" on T1 femur occasionally found on T1 tibia.
 - Linked to c, (chestnut eye color) (0%)

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Go (Goliath)

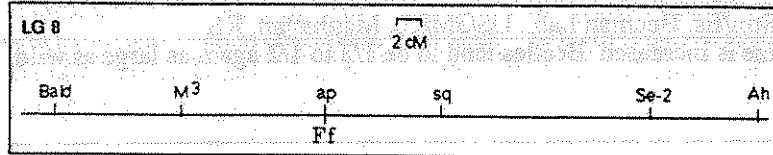
- **Structure affected:** Body size, global
- **Linkage Group:** 7
- **Origin:** Gamma-induced in Stm/Es. Beeman Lab., USGMRL, Manhattan, KS
- **Description:** Overall body size is increased. Beetles tend to be 1/3 to 1/2 again as large as wild type siblings.

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Tribolium castaneum Linkage Group 8

Clickable Map



Mutant Name/note

Ah	Arrowhead
ap	antennapedia
Ff	Fused funicle
Bald	Bald
M³	Medea
Se-2	Split elytra
sq	squint

Last Edited: August 13, 1998



Ah (Arrowhead)

- **Structure affected:** Eye and head.
- **Linkage Group:** 8
- **Origin:** Giovani Mocelin at Jeff Stuart's Lab, Purdue University
- **Description:**
 - This mutant looks like a moderate version of Ey, with extreme expression of all eye facets missing found only rarely.
 - Ah first showed up when males from isogenic line M1/M1 (from Big IV/LII) were irradiated and crossed to MMS females.
 - It maps on LG8 in this order: Bald-ap-sq-Ah, about 25 cM from sq (it may be very near Se-2).
 - The original Ah was a female.
 - They appear to be homozygous viable.

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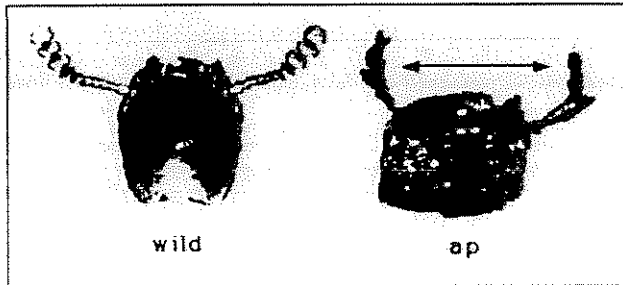


ap (antennapedia)

LG 8

ap (antennapedia)

- **Structure affected:** Antennae.
- **Linkage Group:** 8
- **Origin:** Spontaneous.
- **Description:** Homeotic transformation of antenna to leg.



F (Fused funicle)

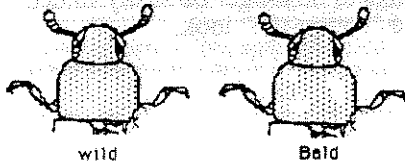
- **Structure affected:** Antennae.
- **Linkage Group:** 8
- **Origin:** Ff originated from an experiment in which Ds (Lg4) was irradiated with gamma rays.
- **Description:**
 - May be a dominant ap allele.
 - Dominance confirmed in a outcross to Ga-1. Penetrance is ~100%.
 - Characteristic phenotype - normal club, funnicle with fusion to usually 4-5 segments. The distal 1-2 segments are enlarged to a size intermediate between that of a funnicle and club segment, giving the club an enlarged 4-segment look usually.

Last Edited: August 13, 1998



Bald

- **Structure affected:** Entire cuticle.
- **Linkage Group:** 8
- **Origin:** EMS, Lab-S or Ga-13.
- **Description:** Patches of setae (cuticular hairs) missing over entire surface of adult body. Gives overall glossy appearance to cuticle.



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Se-2 (Short elytra 2)

- **Structure affected:** Elytra
- **Linkage Group:** 8
- **Origin:** From ab stock, Scott Thomson, (while working in Beeman lab)
- **Description:**
 - Elytral tips are divergent, exposing wings and membranous dorsal abdomen.
 - Enough wing surface is usually exposed to cause the wings to no longer be neatly folded underneath. Wings are often rumpled, giving the beetle a "cotton-tail" look.
 - Elytra are rarely "short".

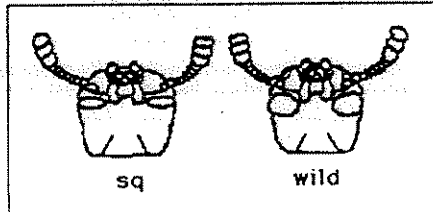
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Tribolium sq (squint)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg8/8sq.html>

sq (squint)

- **Structure affected:** eye
- **Linkage Group:** 8
- **Origin:** Spontaneous recessive, A. Sokoloff, Berkeley, CA.
- **Description:** The number of facets in the eye is reduced, giving the eye a "squinty" look.

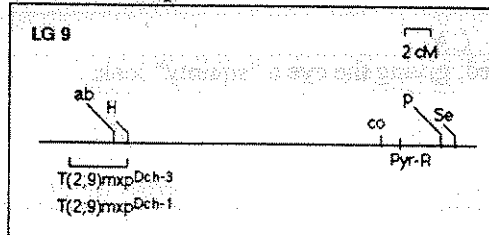


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Tribolium castaneum Linkage Group 9

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Mutant	Name/note
<u>ab</u>	antenna bifurcata
<u>co</u>	cola body
<u>p</u>	pearl eye
<u>Pyr-R</u>	Pyrethroid resistance
<u>Se</u>	Split elytra
<u>T(2;9) mxpDch-1</u>	Dachsund
<u>T(2;9) mxpDch-3</u>	Dachsund
<u>H</u>	H factor

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ab (antenna bifurcata)

- **Structure affected:** Antennae and trochanter of legs
- **Linkage Group:** 2
- **Origin:** Colombia
- **Description:**
 - The antennae are branched, usually at the pedicel (second segment out from head). Size of the branch varies from a small projection, up to 1/3 the length of the antenna. The projection sometimes appears to be segmented.
 - Spikes on the trochanter usually appear at basal edge and vary in size. Some appear segmented. Less commonly found than branches on antennae.
 - Male sterile.

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Tribolium; co (cola body)

<http://bru.usgmr1.ksu.edu/beeman/tribolium/lg9/9co.html>

co (cola body)

- **Structure affected:** cuticle
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** dark brown body color

Last Edited: August 13, 1998



p (pearl eye)

- **Structure affected:** Eyes
- **Linkage Group:** 2
- **Origin:** Alexander Sokoloff, University of California at Berkeley
- **Description:** White colored eye

Last Edited: August 13, 1998



Insecticide Resistance

Lin-R (Lindane resistance)

- **Linkage Group:** 3
 - **Origin:** spontaneous
 - **Description:** resistant to lindane and cyclodiene insecticides because of a mutation in the GABA(A) receptor.
-

Rmal (Malathion resistance)

- **Linkage Group:** 6
 - **Origin:** spontaneous
 - **Description:** resistant to malathion and phenthoate insecticides because of a modification in a carboxylesterase enzyme.
-

Pyr-R (Pyrethroid resistance)

- **Linkage Group:** 9
 - **Origin:** spontaneous
 - **Description:** resistant to alpha-cyano synthetic pyrethroids.
-

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Se (Short elytra)

- **Structure affected:** Elytra
- **Linkage Group:** 9
- **Origin:** Spontaneous, from Ey/Stm X Rd.mas,p (Beeman lab, Manhattan, KS).
- **Description:**
 - Expression is variable, ranging from very short elytra noticeably rounded posteriorly, to almost normal length with a subtle posterior rounding.
 - Closely linked to p (pearl eye)

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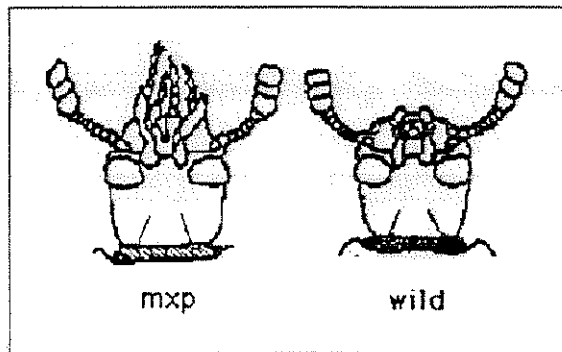


mxp (maxillopedia)

LG 2

mxp (maxillopedia)

- **Structure affected:** maxillary & labial palps
- **Linkage Group:** 2
- **Origin:** spontaneous
- **Description:** mxp/mxp causes transformation of the labial and maxillary palps into legs.

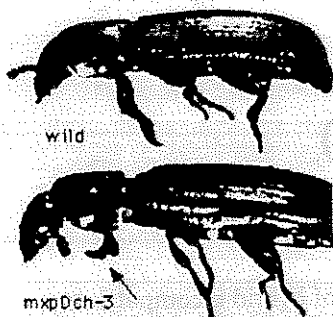


Dch-1 (Dachshund)

- **Structure affected:** Antennae and legs.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** Dominant, gain-of-function (GOF). Antennae & legs shorter than normal (partially transformed towards palp) in heterozygotes. Dch-1/Dch-3 heteroallelic adults have complete and dramatic transformation of legs into palps.

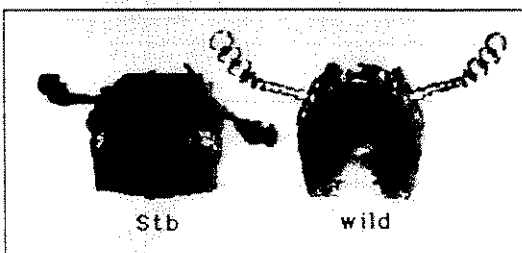
Dch-3 (Dachshund)

- **Structure affected:** Prothorax.
- **Linkage Group:** T(2;9)
- **Origin:** Radiation-induced.
- **Description:** GOF. Prothoracic legs are dramatically reduced. Remainder of prothorax is reduced to a lesser extent, presumably via a homeotic transformation toward labial or maxillary segment. Effect restricted to prothorax. Antennae are normal.



Stb (Stubby)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** EMS, GA-1
- **Description:** GOF. Funicle of antennae reduced via homeotic transformation towards palp. Sometimes behaves as an X-linked trait (upon outcrossing) but loses this property upon inbreeding, only to regenerate the property again upon outcrossing.



Stm (Stumpy)

- **Structure affected:** Antennae.
- **Linkage Group:** 2
- **Origin:** Ethylmethane sulfonate.
- **Description:** GOF. Club and funicle of antennae reduced.
 - Near-lethal with lethal mxp alleles
 - Complements viable mxp
 - Homozygous stock is fertile
 - Other stocks balanced with Stm are only slightly leaky
 - Lab-S RFLP matches Stm RFLP in a clone from the A gene, so Stm probably is a Lab-S chromosome.
 - No translocation found cytologically by Giovanni Mocelin.

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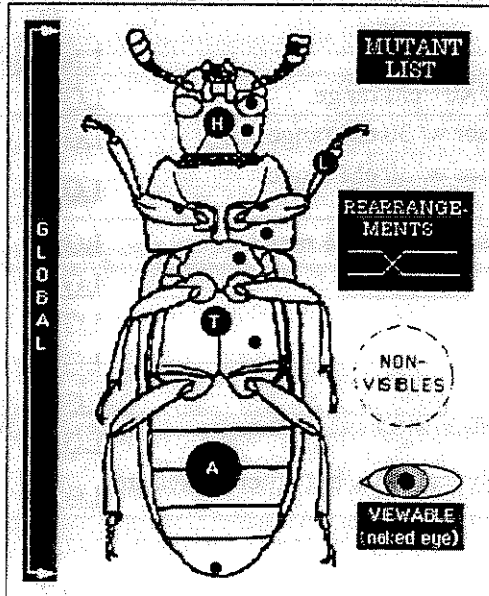
H (Hybrid incompatibility factor)

- **Structure affected:**
- **Linkage Group:** 2
- **Origin:** Tiw-1 strain (India)
- **Description:** There are three kinds of strains involved in this type of hybrid incompatibility, namely H strains, neutral strains and non-permissive (NP) strains. When an H male is crossed to a neutral female, F1 hybrids are viable. However, when an H male is crossed to an NP female at 25 C, the progeny all die as larvae. Incompatibility is less severe at 32 C. Crosses between neutral and NP strains are fully compatible.

Last Edited: August 13, 1998

Mutants by Region Affected

Clickable Image



- [Whole Mutant List](#)
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 - [Shape](#)
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- [Thorax](#)
 - [T1](#)
 - [T2 \(not elytra\)](#)
 - [T3](#)
 - [Elytra](#)
 - [Legs](#)
- [Abdomen-general](#)
- [Abdomen-genital region](#)
- [Elytra](#)
- [Gross \(naked eye/hand lens\)](#)
- [Legs](#)
- ["Invisible" \(lethals; resistance; Medea\)](#)
- [Rearrangements](#)
- [Global](#)

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Mutant List in Alphabetical Order

Mutant	Name/note	Linkage Group
<u>3.2 Bamp</u>	3P2 based crossover suppressor w/ 40+ cM range.	LG 3
<u>3P1</u>	Bamp-27 based crossover suppressor w/ 30 cM range.	LG 3
<u>3P2</u>	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	LG 3
<u>3P3</u>	Bamp-27 based crossover suppressor w/ 45+ cM range.	LG 3
<u>A</u>	Abdominal	LG 2
<u>ab</u>	antenna bifurcata	LG 9
<u>Ah</u>	Arrowhead	LG 8
<u>ap</u>	antennapedia	LG 8
<u>au</u>	aureate	LG 3
<u>b</u>	black body - b itself is incompletely recessive, but other alleles are completely recessive	LG 3
<u>ba</u>	broken antenna	LG 2
<u>Bald</u>	Bald	LG 8
<u>Bamp-27</u>	Blunt abdominal and metathoracic points	LG 3
<u>Bamp-31</u>	Blunt abdominal and metathoracic points	LG 3
<u>Be</u>	Bar eye	LG 4
<u>c</u>	chestnut	LG 7
<u>co</u>	cola body	LG 9
<u>Crab</u>	Crab legs	LG 7
<u>Cx</u>	Cephalothorax	LG 2
<u>Det-43</u>	Divergent elytral tips	T(4; 5)
<u>ds</u>	displaced sternellum	LG 1 = X
<u>Ds</u>	Displaced sternellum	LG 4
<u>Er</u>	Eyes reduced	LG 2
<u>eu</u>	extra urigomphi	LG 2
<u>eu-D</u>	extra urigomphi	T(2; 5)
<u>glossy</u>	glossy cuticle	LG 2
<u>Go</u>	Goliath	LG 7
<u>h</u>	hazel	LG 4
<u>H</u>	Hybrid incompatibility factor	LG 9
<u>j</u>	jet body	LG 5
<u>Lin-R</u>	Lindane resistance	LG 3
<u>lod</u>	light ocular diaphragm	LG 3
<u>M¹</u>	Medea factor	LG 3

Tribolium Mutant List

<http://bru.usgmr1.ksu.edu/beeman/tribolium/regions/mut-list.html>

<u>M²</u>	Medea factor	LG 3
<u>M⁴</u>	Medea factor	unknown
<u>M⁹</u>	Medea factor	LG 8
<u>mc</u>	microcephalic	LG 5
<u>Mo</u>	Micro-optthalmic	LG 6
<u>mxp</u>	maxillopedia	LG 2
<u>mxpStb</u>	Stubby	LG 2
<u>mxpStm</u>	Stumpy	LG 2
<u>p</u>	pearl	LG 9
<u>pd</u>	paddle	LG 1 = X
<u>plt</u>	platinum	LG 1 = X
<u>Ps</u>	Pinched sternellum	LG 2
<u>ptl</u>	prothoraxless	LG 2
<u>py</u>	pygmy	LG 1 = X
<u>Pyr-R</u>	Pyrethroid resistance	LG 9
<u>rb</u>	ruby	LG 5
<u>Rd</u>	Reindeer	LG 2
<u>Rmal</u>	Resistance to malathion	LG 6
<u>s</u>	sooty body	LG 4
<u>Se</u>	Split elytra	LG 9
<u>Se-2</u>	Split elytra	LG 8
<u>sp</u>	shoulder pads	LG 2
<u>Spa</u>	Spatulate	T(2; 4)
<u>sq</u>	squint	LG 8
<u>T(2;4) AEs-1</u>	Extra sclerite	T(2; 4)
<u>T(2;4) tr</u>	tremblor	T(2; 4)
<u>T(2;5) Ey</u>	Eyeless	LG 5
<u>T(2;9) mxpDch-1</u>	Dachsund	T(2; 9)
<u>T(2;9) mxpDch-3</u>	Dachsund	T(2; 9)
<u>T(Y;3)</u>	translocation	LG 3
<u>T(Y;4)</u>	translocation	LG 4
<u>tar</u>	tar	LG 2
<u>ub</u>	unbuckled	LG 2
<u>Utx</u>	Ultrathorax	LG 2
<u>vwe</u>	vestigial wings and elytra	LG 2

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Head

Mutant	Name/note	Linkage Group
<u>ab</u>	antenna bifurcata	<u>LG 9</u>
<u>Ah</u>	Arrowhead	<u>LG 8</u>
<u>ap</u>	antennapedia	<u>LG 8</u>
<u>ba</u>	broken antenna	<u>LG 2</u>
<u>Be</u>	Bar eye	<u>LG 4</u>
<u>c</u>	chestnut	<u>LG 7</u>
<u>Er</u>	Eyes reduced	<u>LG 2</u>
<u>Ff</u>	Fused funicle	<u>LG 8</u>
<u>h</u>	hazel	<u>LG 4</u>
<u>lod</u>	light ocular diaphragm	<u>LG 3</u>
<u>mc</u>	microcephalic	<u>LG 5</u>
<u>Mo</u>	Micro-ophthalmic	<u>LG 6</u>
<u>mxp</u>	maxillopedia	<u>LG 2</u>
<u>mxpStb</u>	Stubby	<u>LG 2</u>
<u>mxpStm</u>	Stumpy	<u>LG 2</u>
<u>p</u>	pearl	<u>LG 9</u>
<u>pd</u>	paddle	<u>LG 1 = X</u>
<u>plt</u>	platinum	<u>LG 1 = X</u>
<u>Ps</u>	Pinched sternellum	<u>LG 2</u>
<u>rb</u>	ruby	<u>LG 5</u>
<u>Rd</u>	Reindeer	<u>LG 2</u>
<u>sp</u>	shoulder pads	<u>LG 2</u>
<u>Spa</u>	Spatulate	<u>T(2; 4)</u>
<u>sq</u>	squint	<u>LG 8</u>
<u>T(2;5)Ey</u>	Eyeless	<u>LG 5</u>
<u>T(2;9) mxpDch-1</u>	Dachsund	<u>T(2; 9)</u>

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Head-antennae

Mutant	Name/note	Linkage Group
ab	antenna bifurcata	<u>LG 9</u>
ap	antennapedia	<u>LG 8</u>
ba	broken antenna	<u>LG 2</u>
Ff	Fused funicle	<u>LG 8</u>
pd	paddle	<u>LG 1 = X</u>
mxpStb	Stubby	<u>LG 2</u>
mxpStm	Stumpy	<u>LG 2</u>
Rd	Reindeer	<u>LG 2</u>
Spa	Spatulate	T(2; 4)
T(2;9) mxpDch-1	Dachsund	T(2; 9)

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Head-eyes

Mutant	Name/note	Linkage Group
Ah	Arrowhead	<u>LG 8</u>
Be	Bar eye	<u>LG 4</u>
c	chestnut	<u>LG 7</u>
Er	Eyes reduced	<u>LG 2</u>
h	hazel	<u>LG 4</u>
lod	light ocular diaphragm	<u>LG 3</u>
mc	microcephalic	<u>LG 5</u>
Mo	Micro-ophthalmic	<u>LG 6</u>
p	pearl	<u>LG 9</u>
plt	platinum	<u>LG 1 = X</u>
rb	ruby	<u>LG 5</u>
sq	squint	<u>LG 8</u>
T(2;5)Ey	Eyeless	<u>T(2; 5)</u>

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Head-shape

Mutant	Name/note	Linkage Group
<u>Ah</u>	Arrowhead	<u>LG 8</u>
<u>Bamp-31</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>Cx</u>	Cephalothorax	<u>LG 2</u>
<u>Er</u>	Eyes reduced	<u>LG 2</u>
<u>mc</u>	microcephalic	<u>LG 5</u>
<u>Mo</u>	Micro-ophthalmic	<u>LG 6</u>
<u>T(2;5)Ey</u>	Eyeless	<u>LG 5</u>

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Head-mouthparts

Mutant Name/note	Linkage Group
<u>mxp</u> maxillopedia	<u>LG 2</u>
<u>Ps</u> Pinched sternellum	<u>LG 2</u>
<u>sp</u> shoulder pads	<u>LG 2</u>

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Thorax

Mutant	Name/note	Linkage Group
<u>3.2 Bamp</u>	3P2 based crossover suppressor w/ 40+ cM range.	<u>LG 3</u>
<u>3P1</u>	Bamp-27 based crossover suppressor w/ 30 cM range.	<u>LG 3</u>
<u>3P2</u>	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	<u>LG 3</u>
<u>3P3</u>	Bamp-27 based crossover suppressor w/ 45+ cM range.	<u>LG 3</u>
<u>acro</u>	acromegaly	<u>LG 2</u>
<u>ap</u>	antennapedia	<u>LG 8</u>
<u>apt</u>	alate prothorax	<u>LG 2</u>
<u>Bamp-27</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>Bamp-31</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>Cx</u>	Cephalothorax	<u>LG 2</u>
<u>Det-43</u>	Divergent elytral tips	<u>T(4; 5)</u>
<u>Ds</u>	Displaced sternellum	<u>LG 4</u>
<u>ds</u>	displaced sternellum	<u>LG 1 = X</u>
<u>mxp</u>	maxillopedia	<u>LG 2</u>
<u>Ps</u>	Pinched sternellum	<u>LG 2</u>
<u>ptl</u>	prothoraxless	<u>LG 2</u>
<u>Rap</u>	Recurved anterior pronotum	<u>LG 2</u>
<u>Se</u>	Split elytra	<u>LG 9</u>
<u>sp</u>	shoulder pads	<u>LG 2</u>
<u>tar</u>	tar	<u>LG 2</u>
<u>ub</u>	unbuckled	<u>LG 2</u>
<u>Utx</u>	Ultrathorax	<u>LG 2</u>
<u>vwe</u>	vestigial wings and elytra	<u>LG 2</u>

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Thorax-T1

Mutant	Name/note	Linkage Group
apt	alate prothorax	<u>LG 2</u>
Bamp-31	Blunt abdominal and metathoracic points	<u>LG 3</u>
Cx	Cephalothorax	<u>LG 2</u>
ds	displaced sternellum	<u>LG 1 = X</u>
Ds	Displaced sternellum	<u>LG 4</u>
mxp	maxillopedia	<u>LG 2</u>
Ps	Pinched sternellum	<u>LG 2</u>
ptl	prothoraxless	<u>LG 2</u>
Rap	Recurved anterior pronotum	<u>LG 2</u>
sp	shoulder pads	<u>LG 2</u>
tar	tar	<u>LG 2</u>
ub	unbuckled	<u>LG 2</u>

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Thorax-T2

Mutant	Name/note	Linkage Group
<u>3.2 Bamp</u>	3P2 based crossover suppressor w/ 40+ cM range.	LG 3
<u>3P1</u>	Bamp-27 based crossover suppressor w/ 30 cM range.	LG 3
<u>3P2</u>	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	LG 3
<u>3P3</u>	Bamp-27 based crossover suppressor w/ 45+ cM range.	LG 3
<u>acro</u>	acromegaly	LG 2
<u>ap</u>	antennapedia	LG 8
<u>Bamp-27</u>	Blunt abdominal and metathoracic points	LG 3
<u>Det-43</u>	Divergent elytral tips	T(4; 5)
<u>ds</u>	displaced sternellum	LG 1 = X
<u>Ds</u>	Displaced sternellum	LG 4
<u>Se</u>	Split elytra	LG 9
<u>Se-2</u>	Split elytra	LG 8
<u>Utx</u>	Ultrathorax	LG 2
<u>vwe</u>	vestigial wings and elytra	LG 2

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Thorax-T3

Mutant	Name/note	Linkage Group
3.2 Bamp	3P2 based crossover suppressor w/ 40+ cM range.	<u>LG 3</u>
3P1	Bamp-27 based crossover suppressor w/ 30 cM range.	<u>LG 3</u>
3P2	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	<u>LG 3</u>
3P3	Bamp-27 based crossover suppressor w/ 45+ cM range.	<u>LG 3</u>
Bamp-27	Blunt abdominal and metathoracic points	<u>LG 3</u>
Bamp-31	Blunt abdominal and metathoracic points	<u>LG 3</u>
Ps	Pinched sternellum	<u>LG 2</u>
vwe	vestigial wings and elytra	<u>LG 2</u>

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Elytra

Mutant Name/note	Linkage Group
<u>Det-43</u> Divergent elytral tips	T(4; 5)
<u>Ds</u> Displaced sternellum	LG 4
<u>Se</u> Split elytra	LG 9
<u>Se-2</u> Split elytra	LG 8
<u>sp</u> shoulder pads	LG 2
<u>Utx</u> Ultrathorax	LG 2
<u>vwe</u> vestigial wings and elytra	LG 2

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Legs

Mutant	Name/note	Linkage Group
<u>ab</u>	antenna bifurcata	LG 9
<u>Crab</u>	Crab legs	LG 7
<u>Rd</u>	Reindeer	LG 2
<u>T(2;9) mxpDch-1</u>	Dachsund	T(2; 9)
<u>T(2;9) mxpDch-3</u>	Dachsund	T(2; 9)

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Abdomen-general

Mutant	Name/note	Linkage Group
<u>3.2 Bamp</u>	3P2 based crossover suppressor w/ 40+ cM range.	<u>LG 3</u>
<u>3P1</u>	Bamp-27 based crossover suppressor w/ 30 cM range.	<u>LG 3</u>
<u>3P2</u>	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	<u>LG 3</u>
<u>3P3</u>	Bamp-27 based crossover suppressor w/ 45+ cM range.	<u>LG 3</u>
<u>A</u>	Abdominal	<u>LG 2</u>
<u>Bamp-27</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>sp</u>	shoulder pads	<u>LG 2</u>
<u>T(2;4) AEs-1</u>	Extra sclerite	<u>T(2, 4)</u>

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Abdomen-genital

Mutant Name/note	Linkage Group
Δ Abdominal	LG 2
<u>eu</u> extra ur/gomphi	LG 2
<u>eu-D</u> extra ur/gomphi	T(2; 2)

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Gross (naked eye; hand lens)

Mutant	Name/note	Linkage Group
<u>A</u>	Abdominal	<u>LG 2</u>
<u>b</u>	black body - b itself is incompletely recessive, but other alleles are completely recessive.	<u>LG 3</u>
<u>Be</u>	Bar eye	<u>LG 4</u>
<u>co</u>	cola body	<u>LG 9</u>
<u>Crab</u>	Crab legs	<u>LG 7</u>
<u>Det-43</u>	Divergent elytral tips	<u>T(4; 5)</u>
<u>Go</u>	Goliath	<u>LG 7</u>
<u>h</u>	hazel	<u>LG 4</u>
<u>Mo</u>	Micro-ophthalmic	<u>LG 6</u>
<u>p</u>	pearl	<u>LG 9</u>
<u>plt</u>	platinum	<u>LG 1 = X</u>
<u>Ps</u>	Pinched sternellum	<u>LG 2</u>
<u>py</u>	pygmy	<u>LG 1 = X</u>
<u>rb</u>	ruby	<u>LG 5</u>
<u>Rd</u>	Reindeer	<u>LG 2</u>
<u>s</u>	sooty body	<u>LG 4</u>
<u>j</u>	jet body	<u>LG 5</u>
<u>Se-2</u>	Split elytra	<u>LG 8</u>
<u>sp</u>	shoulder pads	<u>LG 2</u>
<u>T(2;9) mxpDch-3</u>	Dachsund	<u>T(2; 9)</u>
<u>ywe</u>	vestigial wings and elytra	<u>LG 2</u>

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"Invisible" mutants

Rearrangements (no marker)

Mutant Linkage Group

TCY:3 LG 3

TCY:4 LG 4

Hybrid Incompatibility

Mutant Name/note	Linkage Group
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<u>H</u> Hybrid incompatibility factor	<u>LG 9</u>
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Insecticide Resistance

Mutant Name/note	Linkage Group
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<u>Lin-R</u> Lindane resistance	<u>LG 3</u>
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<u>Pyr-R</u> Pyrethroid resistance	<u>LG 9</u>
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<u>Rmal</u> Resistance to malathion	<u>LG 6</u>
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MEDEA (maternal effect)

Mutant Linkage Group

M¹ LG 3

M² LG 3

M³ LG 8

M⁴

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Rearrangements

Mutant	Name/note	Linkage Group
<u>3.2 Bamp</u>	3P2 based crossover suppressor w/ 40+ cM range.	<u>LG 3</u>
<u>3P1</u>	Bamp-27 based crossover suppressor w/ 30 cM range.	<u>LG 3</u>
<u>3P2</u>	Bamp-27 based crossover suppressor w/ 30 cM range, and recessive au.	<u>LG 3</u>
<u>3P3</u>	Bamp-27 based crossover suppressor w/ 45+ cM range.	<u>LG 3</u>
<u>A</u>	Abdominal	<u>LG 2</u>
<u>Bamp-27</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>Bamp-31</u>	Blunt abdominal and metathoracic points	<u>LG 3</u>
<u>Cx</u>	Cephalothorax	<u>LG 2</u>
<u>Def-43</u>	Divergent elytral tips	<u>T(4; 5)</u>
<u>eu-D</u>	extra urigomphi	<u>LG 5</u>
<u>mxpStm</u>	Stumpy	<u>LG 2</u>
<u>Spa</u>	Spatulate	<u>T(2; 4)</u>
<u>T(2;4) AEs-1</u>	Extra sclerite	<u>T(2; 4)</u>
<u>T(2;4) tr</u>	tremblor	<u>T(2; 4)</u>
<u>T(2;5)Ey</u>	Eyeless	<u>T(2; 5)</u>
<u>T(2;9) mxpDch-1</u>	Dachsund	<u>T(2; 9)</u>
<u>T(2;9) mxpDch-3</u>	Dachsund	<u>T(2; 9)</u>
<u>T(Y;3)</u>	Translocation	<u>LG 3</u>
<u>T(Y;4)</u>	Translocation	<u>LG 4</u>

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Global

Mutant Name/note		Linkage Group
au	aureate	<u>LG 3</u>
b	black body	<u>LG 3</u>
Bald	Bald	<u>LG 8</u>
co	cola body	<u>LG 9</u>
ds	displaced sternellum	<u>LG 1 = X</u>
glossy	glossy cuticle	<u>LG 2</u>
Go	Goliath	<u>LG 7</u>
j	jet body	<u>LG 5</u>
py	pygmy	<u>LG 1 = X</u>
s	sooty body	<u>LG 4</u>
sp	shoulder pads	<u>LG 2</u>

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Medea

MEDEA is an acronym for Maternal-Effect Dominant Embryonic Arrest



Medea-killed larvae

Medea factors all share several characteristics:

- They breed true through the female line.
- They segregate in the male.

When a heterozygous Medea female (M/+) is crossed to a wild type male (+/+), the M gene and its homolog segregate normally. However, all progeny that do not inherit the Medea allele die at or shortly after egg hatch. The lethality is maternal, but the "rescue" is zygotic. The rescuing M allele can be inherited from either parent.

There have been four well-studied Medea factors. Of these, two (M-1 and M-4) are currently maintained at the Tribolium Stock Center. Almost all M strains in the field carry M-4. Of these, about a third also carry M-1. M-4 is the only Medea factor present in North American and European strains, being found in about half of them. Australian and Indian strains are almost devoid of Medea factors. South American, Asian, and African strains often have 2 or more M factors.

M¹ (Medea)

- **Linkage Group:** 3
- **Description:** The first and most-studied Medea factor.

M² (Medea)

- **Linkage Group:** 3
 - **Description:** This Medea factor faded away and is no longer detectable.
-

M³ (Medea)

- **Linkage Group:** 8
 - **Description:** This Medea factor faded away and is no longer detectable.
-

M⁴ (Medea)

- **Linkage Group:** Unknown
 - **Description:** This Medea has an interesting distribution within the United States.
-

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TRIBOLIUM INFORMATION BULLETIN 39

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1954

RE: [Illegible Title]

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Tenebrio and other Tenebrionidae, 1993-1999. Key to Subjects.

1. Anatomy, Histology and Morphology
2. Behavior and Behavioral Ecology and Evolutionary Ecology
3. Cytology and Fine Structure
4. Tissue Culture, Embryology and Development
5. Ecology and Population Biology
6. General
7. Genetics and Animal Breeding
8. Insecticides, Insecticide Resistance, Attractant and Repellents
9. Irradiation and Use of Isotopes
10. Nutrition
11. Parasitology and Symbiosis
12. Pests
13. Biochemistry, Physiology and Molecular Biology
14. Space and Aerial Ecology
15. Speciation and Evolutionary Biology
16. Statistical Methods and Mathematical Models
17. Taxonomy
18. Technique
19. Teratology

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RESEARCH, TEACHING AND TECHNICAL NOTES

RAPD-Based Genetic Linkage Maps of *Tribolium castaneum*

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ABSTRACT

A genetic map of the red flour beetle (*Tribolium castaneum*) integrating molecular with morphological markers was constructed using a backcross population of 147 siblings. The map defines 10 linkage groups (LGs), presumably corresponding to the 10 chromosomes, and consists of 122 randomly amplified polymorphic DNA (RAPD) markers, six molecular markers representing identified genes, and five morphological markers. The total map length is 570 cM, giving an average marker resolution of 4.3 cM. The average physical distance per genetic distance was estimated at 350 kb/cM. A cluster of loci showing distorted segregation was detected on LG9. The process of converting RAPD markers to sequence-tagged site markers was initiated: 18 RAPD markers were cloned and sequenced, and single-strand conformational polymorphisms were identified for 4 of the 18. The map positions of all 4 coincided with those of the parent RAPD markers.

Notes - Research, Teaching and Technical

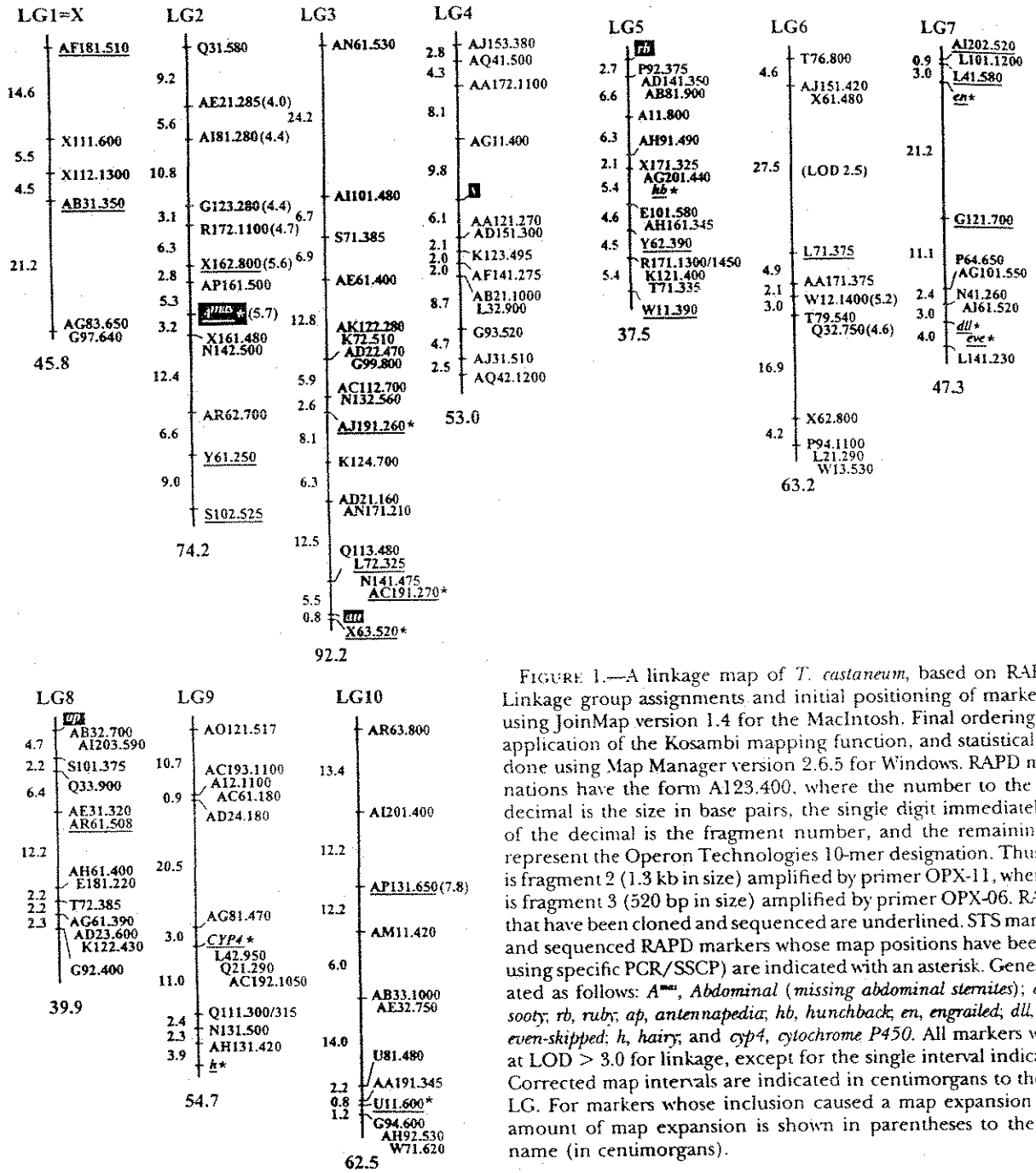


FIGURE 1.—A linkage map of *T. castaneum*, based on RAPD marker Linkage group assignments and initial positioning of markers was done using JoinMap version 1.4 for the MacIntosh. Final ordering of marker application of the Kosambi mapping function, and statistical analysis was done using Map Manager version 2.6.5 for Windows. RAPD marker designations have the form A123.400, where the number to the right of the decimal is the size in base pairs, the single digit immediately to the left of the decimal is the fragment number, and the remaining characters represent the Operon Technologies 10-mer designation. Thus, X112.130 is fragment 2 (1.3 kb in size) amplified by primer OPX-11, whereas X63.52 is fragment 3 (520 bp in size) amplified by primer OPX-06. RAPD markers that have been cloned and sequenced are underlined. STS markers (cloned and sequenced PCR/SSCP) are indicated with an asterisk. Genes are abbreviated as follows: A^{mut}, Abdominal (missing abdominal sternites); au, *auvate*; sooty; rb, *ruby*; ap, *antennapedia*; hb, *hunchback*; en, *engrailed*; dll, *distalless*; e, *even-skipped*; h, *hairy*; and *cyp4*, *cytochrome P450*. All markers were mapped at LOD > 3.0 for linkage, except for the single interval indicated on LG Corrected map intervals are indicated in centimorgans to the left of each LG. For markers whose inclusion caused a map expansion of >4.0, the amount of map expansion is shown in parentheses to the right of the name (in centimorgans).

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Homeotic evidence for an appendicular labrum in *Tribolium castaneum*

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Introduction

The labrum has long been a subject of controversy and lively debate (Rempel, 1975). It has most often been considered a cuticular structure of the acron, though some have suggested that it is a segmental appendage.

We report the first arthropod mutation associated with a homeotic transformation of the labrum. In the gamma irradiation induced *Tribolium castaneum* mutant, *Antennagalea-5* (Ag^5), both antennal and labral structures are transformed to resemble gnathal appendages. Our results suggest that the labrum is a fused structure composed of two pairs of appendage endites and is serially homologous to the gnathal appendages.

Materials and Methods

Beetles were reared at 30° C on whole-wheat flour containing 5% (w/w) brewer's yeast. The *maxillopedia-Stumpy / Abdominal-Miscaudal sclerotization* (mxp^{Sim} / A^{McsI}) balanced stock (Beeman et al. 1989) was used for the reversion mutagenesis. mxp^{Sim} / A^{McsI} males ca. 1 wk old were dosed with 4 kR of gamma irradiation (15 minute exposure), then divided into 5 groups of 285 each (1425 total) and held for 2 days at 30°C. 195 virgin *sooty* (*s*) females ca. 2 weeks old were then added to each of the five groups of males. Males were discarded after two days and females were allowed to oviposit for several weeks. F1 adults were screened for reversion (loss) of either of the dominant mutations mxp^{Sim} (homeotic transformation of antennae to maxillary palps) or A^{McsI} (homeotic transformation of ventral abdominal segment 8 to segment 7) and would be recognized by their wild-type phenotype. In the absence of genetic reversion, all progeny would carry either the mxp^{Sim} or the A^{McsI} mutation. In addition to revertants, new dominant mutations were also detected. These mutants were outcrossed to *Gal* to remove incidental mutations, with progeny crossed to the balancer chromosome *Abdominal-Extra sclerite* (A^{Esl}) (Beeman et al., 1989) to establish balanced mutants stocks.

Specimens were prepared for laser confocal microscopy as follows: Frozen or 70% ethanol-preserved adult beetle heads were manually cleaned of debris or cleared in NaOH and rinsed, then blotted dry on tissue paper and mounted directly on double-stick tape on a 75 x 25-mm glass microscope slide. Mouthparts and antennae were dissected from cleared or uncleared specimens and mounted in Euparal (ASCO Laboratories) on a 75 x 25-mm glass microscope slide.

A Zeiss LSM400 laser confocal microscopy system was used to image most specimens. Optical sections ranged from 40 to 570 um and were recorded in the 4 line average mode with picture size of 512 X 512 pixels. Some specimens were imaged using a Spot Camera II (Diagnostics) digital camera on an Olympus SZH-10 stereoscope or an Olympus BX60 compound microscope. Images were archived and digitally processed using Corel Photo Paint, then arranged and labeled using Corel Draw.

Results and Conclusions

This reversion mutagenesis generated the new dominant mutant, *Antennagalea-5* (Ag^5) on the mxp^{Sim} chromosome. In Ag^5 , the distal antennal scape bears setae borne on a dorsal enlargement. The Ag^5 scape also has a membranous base rather than the normal sclerotized "knob".

Dominant modifications of the anterior region of the adult head capsule are also found. In Ag^5 heterozygotes, the anterior rim of the head is always indented at the epicranial arms due to the shortening of the clypeus, exposing portions of the dorsal labrum, mandibles and antennae normally covered by the anterior rim.

The labrum also undergoes grossly detectable morphological transformations in ~2.7 % of the beetles. Changes range from a midline longitudinal membranous strip and/or a slight elongation of the labrum to a striking transformation of both the labrum and clypeus into enlarged complex structures which can include a strongly sclerotized tooth-like tip (Figs. 2 & 3). The most strongly expressed tooth-like tips resemble the distal mandible. Palp-like structures have not been found in any transformed labrum examined thus far, suggesting that the wild-type labrum probably consists of the two endites only.

The apparent appendicular nature of the labrum raises a variety of interesting questions. The labrum has often been assumed to be a part of the acron. However, by definition, the acron is a structure that has no appendages. The labrum therefore must not be located on the acron, and likely represents a segment. Butt (1960) proposed that the labrum is the fused appendages of the intercalary segment. Matsuda (1960) pointed out that the existence of such an labro-intercalary segment would mean that "this segment would carry two pairs of appendages"; a complete pair from the labrum and another complete pair from the intercalary segment. By definition, each insect segment has only one pair of appendages (Manton, 1977; Kukalova-Peck, 1992). However, if the labrum is composed of endites only (galea and lacinia homologs) as our study proposes, and if the transitory appendages of the intercalary segment represent only the repressed palps (telopodites), then the sum of labral and intercalary appendages equals just one complete appendage pair. This configuration eliminates the basis of Matsuda's protest.

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Figures

(Left and right arrows show labrum and clypeus)

Fig. 1
Wild-type head, lateral view

Fig. 2
Ag5 with transformed labrum

Fig. 3
Ag5, close-up of transformed labrum



Dose-mortality responses of the flour beetles to Triflumuron and Cyromazine

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1. Introduction

The red flour beetle, *Tribolium castaneum* Herbst and the confused flour beetle, *Tribolium confusum* DuVal are major pests of the stored food commodities throughout the world (Sokoloff, 1972). At present control of the stored-product insect pests by pathogens and parasitoids, botanicals and insect growth regulators are gaining importance over the conventional insecticides to avoid the presence of the chemical residue in the human food (Wilkin and Fishwick, 1981, Mondal, 1993) and incidence of insect resistance to insecticides (Dyte, 1974; Dyte and Blackman, 1967, 1970; Champ and Dyte, 1976; Metcalf, 1980; Georghiou and Mellon, 1983; Champ, 1986).

The insect growth regulators (IGRs) being non-persistent, biodegradable and more selective to the insect species than the conventional pesticides (Menn and Henrick, 1981), have attracted attention of the entomologists to use these compounds against the stored products insect pests. According to the mode of action, IGRs are categorized into three classes, viz., juvenile hormone analogues, chitin synthesis inhibitors and ecdysone agonists (Willis, 1974; Staal, 1975; Wing and Aller, 1990). As a whole the IGRs retard growth and development of the insects. Among these compounds the chitin synthesis inhibitors disturb or even abort moulting in the developing insects, and these insects either become crippled or die. The early larvae of several stored-product insects when exposed for a longer period to low doses of these compounds produced significant larval mortality, which is a cumulative toxic effect on different larval instars (Mian and Mulla, 1982; Ishaaya and Yablonski, 1987). At very low doses these compounds affected larval development and population growth in *T. castaneum* (Mondal and Port, 1995; Mondal *et al.*, 1998). However, toxicity of these compounds decreases with larval age when exposed for a shorter period (Carter, 1975; Mian and Mulla, 1982; Mayuravalli and Reddy, 1986).

In the present study toxicity of two chitin synthesis inhibitors viz. Triflumuron and Cyromazine against *T. castaneum* and *T. confusum* was determined.

2. Materials and Methods

2.1. Insects used and culture:

Species	Strain	Origin of Culture	Food media used	Rearing temperature ⁰ C
<i>T. castaneum</i>	Normal	Institute of Biological Sciences, Rajshahi University, Bangladesh.	Whole wheat flour and Brewers yeast (19:1)	30
<i>T. confusum</i>	FSS II CTC 12	Slough Laboratory, UK Slough Laboratory, UK Institute of Biological Sciences, Rajshahi University, Bangladesh.		

2.2. IGRs used:

Chemical name	Common name	Code name	Company	Formulation
Triflumuron	Baycidal/Alsystin	BAY SIR 8415	Bayer AG	25% WP
Cyromazine	Larvadex/Trigard/ Vetetazine	CGA 72 662	Ciba Geigy	98% WP

2.3. Preparation of doses: Different doses of the test compounds were made by adding appropriate weight of each compound with the standard food medium and mixing them thoroughly with an electrical blender.

2.4. Screening of doses: A preliminary screening of the doses of each compound was performed on different larval instars, neonates and adults, exposed for different period (section 2.5) to determine 0% and 100% mortality. Then a series of five doses were chosen for each larval instar and adult stage of the beetles.

2.5. Exposure period: Different stages of the beetles were exposed to the treatment for following period:

- (i) Larvae from first to sixth instars (Mondal, 1984), of both species were exposed to different doses of both compounds for 48 hours.
- (ii) Newly hatched larvae (neonates) of both species were exposed to the treatment up to pupation.
- (iii) Newly emerged adults were exposed to the treated food separately. The adults were exposed for 10 and 20 days to triflumuron and cyromazine treated food respectively.

2.6. Methodology: Individuals of each larval instar, neonates, sexed and unsexed adults were kept in separate flat bottom glass vial (50x25 mm) containing approximately 0.5g of either treated or untreated food. The top of the vial was plugged with cotton.

2.7. Replication:

- (i) Fifty individuals of each larval instar, neonates and adults of FSS II and CTC 12 strains of *T. castaneum* were exposed to either untreated or triflumuron treated food. The experiments were replicated five times.

- (ii) Forty individuals of each larval instars and neonates of normal laboratory strains of *T. castaneum* were exposed to either untreated or cyromazine treated food. Twenty adults were used in similar way. All the experiments were replicated five times.
- (iii) Thirty individuals of each larval instars, neonates and adults were exposed either to triflumuron treated or untreated food. The experiments were replicated five times.
- (vi) Forty individuals of each larval instars and neonates of normal laboratory strains of *T. confusum* were exposed to either untreated or cyromazine treated food. Twenty adults were used in similar way. All the experiments were replicated five times.

2.8. Analysis of data: The percentage mortality data were subjected to statistical analysis (Busvine, 1971) and the dose-mortality response was expressed as a median lethal dose (LD_{50}). The percentage mortality obtained was corrected using Abbott's formula (Abbott, 1925) wherever necessary.

3. Results

3.1. Larvae: Older larvae of both *T. castaneum* and *T. confusum* showed tolerance to triflumuron treated food at 48 hours exposer (Table 1). The first instar larvae of FSS II and CTC 12 strains of *T. castaneum* and the second instar larvae of *T. confusum* were found to be more susceptible to triflumuron. CTC 12 strain of *T. castaneum* was more tolerant to triflumuron than FSS II strain. Both strains of *T. castaneum* were found to be more tolerant to the compound compared to *T. confusum*.

Tolerance to cyromazine increased with the larval age of both species of the beetle (Table 2). The second instar larvae were most susceptible to cyromazine. *T. confusum* larvae showed more tolerance to the compound than *T. castaneum* larvae.

3.2. Neonates: In long exposer the LD_{50} for both species of the beetles was very low in triflumuron treated food, and comparatively higher doses of cyromazine were required to obtain the LD_{50} level (Table 3). *T. confusum* larvae were found to be tolerant to both compounds than *T. castaneum* larvae at long exposer.

3.3. Adults: The males of both species were found to be more susceptible to both compounds than the females. Adult CTC 12 beetles were more tolerant to triflumuron than adult FSS II beetles. *T. confusum* adults were found to be more resistant than *T. castaneum* adults in both treatments (Table 4).

4. Discussion

Chitin synthesis inhibitors (CSIs) act against the larval stage of insects, which usually fail to survive due to incomplete moulting or disrupted cuticle formation (Fox, 1990). Both triflumuron and cyromazine are stomach toxicants, and need the larvae to feed on the treated food. If the larvae feed on these compounds at the beginning of instar, the lethal effect appears during any of the next moults (Hammann and Sirrenberg, 1980; Zobelein *et al.*, 1980). Hence at short exposer period higher doses of these compounds are needed for the toxic effect compared to the dose required at longer exposer. Similar results were reported by Ishaaya *et al.* (1984) and Ishaaya and Yablonski (1987) when *T. confusum* larvae were treated with other CSI compounds. Young larvae are normally susceptible to CSI treatments (Mian and Mulla, 1982). Susceptibility of the second instar larvae of the laboratory strains of both species as obtained in the present study, might be due to the voracious nature of these larvae than the younger ones. The older larvae generally show tolerance to CSI compounds (Neal, 1974; Rathburn and Boike, 1975; Busvine *et al.*, 1976;

Hammann and Sirrenberg, 1980, Retnakaran and Wright, 1987; Natarajan *et al.*, 1988; DeMark and Bennett, 1989), possibly due to their better ability to metabolize these compounds (Neumann and Guyer, 1987; Clarke and Jewess, 1990).

By virtue of mode of action, CSI compounds would not be lethal to the adults (Grosscurt and Jongmsa, 1987). In the present study at higher doses of the compounds and long exposer adult mortality was observed in both species. In adults CSI compounds very often disrupt the gut and associated organs (Clarke *et al.*, 1977; Becker, 1978,1980; Mitsui *et al.*, 1984; Soltani, 1984, Pelsue, 1985 and Parween, 1997), resulting digestive disorders (Ishaaya and Ascher, 1977), starvation and ultimate death (Soltani, 1984; Neumann and Guyer, 1988). Moreover, triflumuron treated food disrupted the reproductive systems in both sexes of adult *T. castaneum* (Parween, 1996). All these effects might have resulted in the lethal effects in adults, those fed on triflumuron or cyromazine treated food in the present experiment.

5. Conclusion

Both Triflumuron and Cyromazine can control the flour beetles effectively at low doses if the young larvae are allowed to feed on treated food for whole larval life. A short treatment to either the larvae or to the adults would not be effectively toxic and will need very high dose. The confused flour beetles were found to be slightly tolerant to the CSI treatment. However, the multi-resistant strain (CTC 12) of *T. castaneum* was found to be slightly tolerant to triflumuron than the susceptible strain (FSS II) (Table 5). The present result indicates that both Triflumuron and Cyromazine may be used in the management of *Tribolium* which is important from both the Integrated Pest Management (IPM) and Global Environment Protection (GEP) point of views.

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Table 1. Dose-mortality responses of larvae of FSS II and CTC 12 strains of *T. castaneum* and normal *T. confusum* exposed to triflumuron for 48 hours.

Species & strain	Larval Instar	LD ₅₀ (ppm)	95% confidence limit (ppm)		Slope	χ^2 value (DF)
			upper	lower		
<i>T. castaneum</i>						
FSS II	1st	68.15	88.03	52.76	3.04	48.78(3)
	2nd	6263.19	7566.68	5184.26	4.83	67.79(3)
	3rd	7792.67	8697.50	6981.96	6.03	30.59(3)
	4th	16078.03	20337.82	12710.47	3.86	76.08(3)
	5th	20054.55	22082.83	18213.00	6.08	28.78(3)
	6th	28069.00	29572.19	266625.37	11.15	15.24(2)
CTC 12	1st	1596.34	1817.79	1401.86	6.31	55.49(3)
	2nd	6830.88	8434.25	5532.32	3.32	46.19(3)
	3rd	17672.16	19644.85	15897.57	8.66	70.28(3)
	4th	18126.26	19444.58	16897.33	10.75	44.99(3)
	5th	22255.09	23882.76	20738.31	11.63	25.07(2)
	6th	45966.42	50883.58	41524.41	8.03	45.24(3)
<i>T. confusum</i>						
FSS II	1st	10.98	12032.29	10026.88	2.32	2.007(3)
	2nd	357.04	388.29	323.29	2.27	6.707(3)
	3rd	5116.37	5547.43	4718.81	2.42	3.31(3)
	4th	8770.19	9983.91	7704.02	3.18	13.115(3)
	5th	11863.74	12901.00	10909.88	2.28	5.37(3)
	6th	15769.14	17926.11	13871.71	2.66	8.95(3)

Table 2. Dose-mortality responses of larvae of normal strains of *T. castaneum* and *T. confusum* exposed to cyromazine for 48 hours.

Species	Larval Instar	LD ₅₀ (ppm)	95% confidence limit		Slope	χ^2 value (DF)
			upper	lower		
<i>T. castaneum</i>						
FSS II	1st	27202.60	32108.70	23046.12	2.87	17.81(3)
	2nd	483.68	545.04	429.23	1.61	0.54(3)
	3rd	3848.40	4313.20	3433.69	1.72	1.04(3)
	4th	8129.59	8129.59	9945.19	1.86	10.65(3)
	5th	18038.37	18038.37	16214.90	1.83	3.78(3)
	6th	39528.56	39528.56	44901.33	3.02	11.41(3)
<i>T. confusum</i>						
FSS II	1st	27553.02	32679.44	23230.79	2.89	19.38(3)
	2nd	500.79	563.58	445.01	1.63	0.41(3)
	3rd	4171.61	4673.95	3723.25	1.73	0.40(3)
	4th	8684.01	10505.24	7178.53	1.89	9.66(3)
	5th	18376.46	20563.97	16421.64	1.73	5.50(3)
	6th	39799.99	42510.89	37261.99	2.99	6.08(3)

Table 3. Dose-mortality responses of neonates of *T. castaneum* and *T. confusum* exposed to triflumuron and cyromazine up to pupation

Species & strain	Compound	LD ₅₀ (ppm)	95% confidence limit (ppm)		Slope	χ ² value (DF)	
			upper	lower			
<i>T. castaneum</i> FSS II	Triflumuron	1.81x10 ⁻⁵	5.24x10 ⁻⁵	6.26x10 ⁻⁵	1.12	73.68(2)	
		CTC 12	1.54x10 ⁻⁴	3.36x10 ⁻⁴	7.11x10 ⁻⁵	0.79	28.94(3)
		<i>T. confusum</i>	0.153	0.137	0.17	3.76	11.62(3)
<i>T. castaneum</i>	Cyromazine	152.86	177.63	131.55	1.97	4.48(3)	
<i>T. confusum</i>		168.98	196.37	145.41	1.92	2.83(3)	

Table 4. Dose-mortality responses of adults of *T. castaneum* and *T. confusum* exposed to triflumuron and cyromazine for 10 and 20 days respectively.

Species & strain	Sex	LD ₅₀ (ppm)	95% confidence limit (ppm)		Slope	χ ² value (DF)
			upper	lower		
Triflumuron						
<i>T. castaneum</i> FSS II	Male	12643.77	13620.66	11736.92	7.93	24.11(3)
	Female	12833.97	13782.86	11950.42	6.47	21.37(3)
	Unsexed	12080.44	13579.90	10746.55	5.95	39.39(3)
CTC 12	Male	15470.18	17215.39	13901.87	6.89	33.59(3)
	Female	17888.70	18944.68	16891.57	5.78	18.78(3)
	Unsexed	15386.49	17396.23	13608.93	5.43	31.69(3)
<i>T. confusum</i>	Male	21956.70	23824.74	20235.12	3.51	7.34(3)
	Female	23232.45	25260.13	21367.52	3.35	3.50(3)
	Unsexed	20566.12	23701.46	17845.54	3.56	9.17(3)
Cyromazine						
<i>T. castaneum</i>	Male	56197.17	65471.70	47870.85	2.79	8.09(3)
	Female	57580.57	68388.15	48480.89	2.81	9.30(3)
	Unsexed	55947.23	66680.83	46941.42	2.76	9.53(3)
<i>T. confusum</i>	Male	58426.82	64112.92	53245.03	2.97	6.25(3)
	Female	59449.07	65493.08	53962.85	2.83	6.04(3)
	Unsexed	56645.43	66806.22	48030.03	2.77	8.44(3)

Table 5: Resistance of the larvae and adults of the resistant CTC-12 strain of *T. castaneum* to triflumuron

Life stage	Exposure period	LD50 values		E.R.F.*
		FSS II	CTC-12	
Larva				
1st instar	48 hrs	68.15	1596.34	23.42
2nd "	"	6263.197	6830.88	1.09
3rd "	"	7792.67	17672.16	2.27
4th "	"	16078.03	18126.26	1.13
5th "	"	20054.55	22255.09	1.11
6th "	"	28069.00	45966.42	1.64
Adult				
Male	10 days	12643.77	15470.18	1.22
Female	"	12833.97	17888.70	1.39
Unsexed	"	12080.44	15386.49	1.27
Neonate	16 days	2.28-06	2.85-05	12.50

* Estimated Resistant Factor = $\frac{\text{LD}_{50} \text{ for resistant strain}}{\text{LD}_{50} \text{ for susceptible strain}}$

STABILIZATION OF RICE BRAN BY PHYSICO-CHEMICAL TREATMENTS DURING INSECT INFESTATION

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Summary

Rice bran contain 16-20% oil. The present study shows that the chemical and heat treatments significantly ($P < 0.001$) reduced the insect infestation of rice bran by *Tribolium castaneum* Herbst and *Tribolium confusum* Duval compared to control. The process based on the principle that lipase activity will be low at low p^H of hydrochloric acid at 40 litre / ton of bran for lowering the p^H of rice bran from 6.0 to 4.0. This simple method which takes less than 4 minutes for a batch of 15kg will be useful for stabilization of rice bran in rice mills where steam or electricity is unavailable.

INTRODUCTION

The bran is the most important by-product of the rice milling industry and is valued as a cattle feed, it also yields an edible oil. It comprises the germ, the pericarp and aleurone layer and is often found mixed with varying quantities of husk. The bran contains up to 25% oil used for edible and other purposes. The oil is extracted in large quantities in Japan and to a lesser extent in the USA. In India rice bran oil industry has made rapid strides during the recent years. Rice bran is as good an edible oil as refined ground nut and cotton seed oil and has better keeping qualities due to the presence of antioxidants (α and γ tocopherols). It is suitable for use as a salad and cooking oil and in the preparation of shortenings and vanaspati (hydrogenated oil). Rice bran oil is reported to prevent cancer in the liver of rats. The bran should be extracted soon after it is milled, other wise it deteriorates quickly; an extremely active lipase begins to act as soon as the bran is removed from rice and caused free fatty acid in the oil to increase rapidly at the rate of 1% per hour during the first few hours of the storage of the bran. Heating of the bran immediately after it is milled at about 100°C for a period of 2 hours is found to inactivate the lypolytic enzyme system and increase the storage life of the bran considerably (Anon, 1966). Several problems of raw bran management have restricted production of edible grade rice bran oil. One of the problem is high lipase activity in the bran which quickly hydrolyzes the oil into fatty acids. Another problem for storing the rice bran is high insect infestation in the stores. These two problems are need to be solved by adapting the methods of stabilization of rice bran which is described in the present experiment. Rice bran has considerable potential as a contributor to world oil supply.

MATERIALS AND METHODS

Freshly milled raw rice bran, containing an average of 15% oil was obtained from a local rice mill. Commercial hydrochloric acid (28-30%) was used for acid treatment.

Acid treatment: Rice bran was spread in a layer of about 5 cm thick and the required amount of hydrochloric acid was sprayed on and mixed well by hand using protective gloves.

Heat treatment: 5kg bran of IRRI-8 was heated in an oven at temperature 105°C for 3 hours to prevent the lipase activity during storage.

Analytical Method: Oil content of rice bran with free fatty acid (FFA) content of extracted oil were determined at every 10 days interval by using AOCS (American Oil Chemical Society) method (Anon, 1973). The pH of bran was determined at every 10 days intervals with an aqueous suspension of the bran (10g/100ml) using an Ellico pH meter.

Insect infestation study: 100 gm of rice bran from acid and heat treatments were kept individually in a gunny bag with 40 adults of *Tribolium castaneum* Herbst and *Tribolium confusum* Duval. 100gm of fresh rice bran with same number of the two species of *Tribolium* was kept individual in a gunny bag and this was considered as control. Six replications were taken for each treatment. After every 30 days the total number of larvae of each species from each treatment was removed by sieving the bran. This was continued up to 3 months at room temperature. All the gunny bags were placed in a wooden tray and covered by steel frame against rodent damage.

RESULTS AND DISCUSSION

Rice bran lipase is known to have a pH optimum of 7.5-8.0 (Sastry, 1973) with activity declined with either an increase or decrease in pH. The pH of rice bran had to be lowered to 4.0 to achieve a commercially acceptable low level of enzyme activity. The pH of commercial bran varies from 6.0 to 6.9 and about 4 ml of hydrochloric acid (28-30% strength) was required for 100g of rice bran (or 40 litre/ton of bran) to lower the pH to 4.0. In all the experiments on acid stabilization of rice bran, a minimum of 40ml of hydrochloric acid per kg of bran was used. The efficacy of hand mixing of hydrochloric acid with rice bran in controlling its lipase activity as measured by increasing in the free fatty acid (FFA) content of the bran oil (Table-2). FFA content of oil from untreated raw bran increased rapidly to 52.7 after 90 days of storage but the increase was much slower from 2.8 to 9.3 after 90 days storage in the hydrochloric acid treated bran. The observed increase in FFA from 2.8 to 9.3% probably resulted from non uniformity in distribution of hydrochloric acid in the bran mass with some pockets of bran remaining at higher pH. This was evident from variations of up to one pH unit among different portions of the treated bran. Increasing the quantity of hydrochloric acid from 40mg/kg to 50ml/kg of bran did not further improve (FFA increase from 3.0 to 7.4 in 90 days) the efficacy of the process. Acid stabilized bran was not easily infested. In the present experiment 40 adults insects of *T. castaneum* and *T. confusum* were individually introduced into acid stabilized, heat stabilized and untreated bran (100gm each). Rapid proliferational of *T. castaneum* and *T. confusum* were observed within a month in heat stabilized (803 and 476 larvae respectively) and untreated (833 and 525 larvae respectively). The bran turned dark and moldy after about two months. The acid treated bran on the other hand had only 144 and 104 larvae of *T. castaneum* and *T. confusum* respectively and showed no viable mold growth after three months of storage. It is concluded from this experiment that the occurrence of the larvae of *T. castaneum* and *T. confusum* was highly significant in both the treatments on individual month basis except in case of occurrence of *T. confusum* in first month (Table-1). Acid stabilization appears to facilitate extraction of the crude oil.

Feed quality of bran does not appear to be affected by the acid treatment. Prabhakar and Venkatosh (1986) assessed the feed quality of acid stabilized bran for poultry, using 264 layers over a period of one year at 20, 30 and 40% levels of bran in the composite diet. They found no statistically significant difference among acid stabilized, heat stabilized and untreated bran with regard to feed consumption, egg weight, albumin and yolk indices, shell thickness and Haugh units. However, at all three levels of bran in the diet, the layers fed acid stabilized bran always recorded less feed consumption per egg (137.69g/egg) compared to heat stabilized (168.0g/egg) or untreated (150.0g/egg) bran. Also the number of hen house eggs was always higher for the acid stabilized bran fed group (6410 in 13 weeks, mean wt 58.8±0.95) or compared to heat stabilized 6167, mean wt 58.8±0.95) or untreated (6109, mean wt 57.6±6.95) bran. This simple chemical method for stabilization of rice bran might provide an answer to the problems of handling raw bran in developing countries (like Bangladesh) with numerous small rice mills which lack adequate steam and electricity and where stabilization of rice bran using heat is expensive.

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Table 1. Effect of heat and acid treatments to the rice bran on the occurrence of the larvae of *Tribolium castaneum* Herbst and *Tribolium confusum* Duval.

No. of observations (month)	Number of Larvae							
	<i>T. castaneum</i>				<i>T. confusum</i>			
	Acid treatment	Heat treatment	Normal	F-values	Acid treatment	Heat treatment	Normal	F-values
Mean ± S. E.	Mean ± S. E.	Mean ± S. E.		Mean ± S. E.	Mean ± S. E.	Mean ± S. E.		
1 st	2.66 ± 0.494	38.0 ± 5.2599	30.30 ± 3.480	25.89 ^{***}	3.33 ± 0.6146	9.833 ± 2.340	10.50 ± 3.939	2.19 ^{9N.S.}
2 nd	34.66 ± 3.480	189.50 ± 20.408	223.0 ± 13.214	50.26 ^{***}	23.00 ± 2.394	124.66 ± 14.79	103.83 ± 12.53	28.8 ^{3***}
3 rd	34.833 ± 4.04	173.833 ± 13.99	163.33 ± 19.08	31.17 ^{***}	25.66 ± 2.836	103.66 ± 5.469	148.0 ± 6.93	133.9 ^{7***}

N.S. = Not significant

*** = Highly significant (P.0.001)

Table 2. Efficacy of hand mixing of hydrochloric acid for stabilization of rice bran.

Period (days)	Normal		Treated with HCL 40ml/kg bran	
	pH	FFA Oleic acid/100g oil	pH	FFA Oleic acid/ 100g oil
0	6.99	2.3	3.9	2.8
10	6.80	15.5	4.0	3.0
20	6.70	17.5	4.1	3.5
30	6.6	27.4	4.3	4.2
40	6.4	35.0	4.3	4.5
50	6.3	47.5	4.2	4.7
60	6.2	49.0	4.3	5.3
70	6.1	50.5	4.2	6.3
80	6.1	57.5	4.1	8.3
90	6.0	52.7	4.1	9.3

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*Tribolium brevicornis—a review.

Tribolium brevicornis is a species of Tribolium first reported by John LeConte (1865, 1866) from the vicinity of Fort Tejon, California. Originally it was named Aphanotus brevicornis, but Hinton (1948) in his revision of the genus included it in the genus Tribolium.

Davidson (1893) and Nininger (1916) recorded the beetles in the nests of Xylocopa, and Hicks (1929) from nests of another bee, Anthidium mormonum fragariellum. Linsley, (1943, 1944) found that nests of Xylocopa orpifex discovered in a decaying log were infested with Tribolium brevicornis. No numerical data were provided.

Okumura and Strong (1965) collected T. brevicornis in Alpine and Mono counties in California, and in a subsequent study Strong (1970) trapped this beetle near barns in nine (unspecified) localities in California. Again, there was no mention of the numbers of these beetles. Although T. brevicornis appears to be primarily Californian, it has been found in Parma, Idaho. In this area, the alfalfa bee, Megachile pacifica is raised commercially in "bee boards" (pieces of wood drilled with holes set out to provide nesting sites for the bee) to increase the yield of alfalfa seed. Tribolium brevicornis has become a major pest, destroying many nests of the alfalfa bee. T. brevicornis may have been imported into Idaho in lumber originating from California (R.W. Smith, personal communication).

Cultures of Tribolium have been maintained in the laboratory at the University of Idaho, Moscow Idaho, and at the University of California, Riverside, California for many years. These domesticated strains will be referred to as the Riverside and the Idaho strains, and they are available at the Tribolium Stock Center, California State University, San Bernardino, California.

A third strain of T. brevicornis was discovered by accident by the author during a field collecting trip of the author and his Entomology class to Waterman Canyon, just North of the city of San Bernardino. While the students dispersed in all directions to collect insects in the creek or on the banks of Waterman Creek, the author looked for insects under the bark of trees. Under the bark of an alder tree Alnus rhombifolia downed during a previous winter storm, I found some beetles that looked familiar. Instead of dropping them into an alcohol vial I put them in a dry vial to examine more closely in the laboratory. They looked similar in color and general body shape as T. brevicornis from Riverside, and the compound eye being bisected into two lobes by the gena in the two samples seem to indicate that the beetles at Waterman Canyon were indeed T. brevicornis. Matings between Waterman Canyon males with Riverside females showed that the beetles were compatible to produce F1, and the crosses between F1 beetles produced F2 which were perfectly fertile was further evidence that the two strains belonged to the same species and that the new strain could be maintained in the laboratory on standard flour beetle feed (whole wheat flour plus brewer's yeast in the proportion of 19:1 by weight).

When it became evident that these beetles thrived in such a medium, sizable populations of the Waterman strain were obtained. These populations, referred as Waterman Canyon II and Waterman Canyon III gave us the opportunity to study the manner in which feral populations become domesticated. In other words, can feral strains, feeding on immature stages of Xylocopa and perhaps on live and even dead adults of the carpenter bee in nature adapt readily to a laboratory feed on which they have never fed?

Two portions of the dead alder tree about a meter in length were taken to the laboratory and sectioned with a band saw into strips 3 X 3 cm in width and height, exposing the galleries of the carpenter bee later identified as Xylocopa tabaniformes orpifex. Within these galleries were found a large number of Tribolium brevicornis adults and a few larvae and pupae. Some of these larvae were extracted from the dead bodies of carpenter bees on which they were feeding. In the laboratory the feral beetles were placed in beetle medium and their fecundity and fertility determined. Some samples of eggs collected in egg farms in which the flour has been dyed with neutral red were exposed separately to males and females to determine the rate of cannibalism. Other samples of eggs were placed in different nutrients, and their developmental rates to the adult stage and the viability from egg to the adult determined.

Flying ability was measured by the technique devised by Lerner and Inouye and described in Sokoloff (1977).

All of the above attributes were compared with those obtained from the Riverside and Idaho populations strains which are considered as being highly domesticated, since they have been kept in their respective laboratories for many generations. Samples of feral and domesticated strains were measured to obtain total body length and body weight. The beetles were then introduced in flour, allowed to lay eggs,

and the eggs were placed in corresponding vials. Upon hatching, the larvae were placed in one of three diets: a carbohydrate diet consisting of whole wheat flour; 2) a protein diet consisting of ground, dehydrated insects (Tenebrio, Tribolium and Apis larvae) and 3) a carbohydrate and protein diet made of combining diets 1 and 2. The beetles were reared in an incubator maintained at 32°C and about 70% R.H. The adult body weight and total body length measurements were continued into the F1 and F2 of the feral population to assess the variability of these organisms when reared under optimal conditions. Other methods used will be described as needed.

Results

1. Size

All T. brevicornis collected at Waterman Canyon as adults were sexed and weighed, and subsequently total body length was determined by etherizing the beetles and measuring them with an ocular micrometer. These measurements were later compared with those from samples of the Riverside and Idaho strains, and with the F1 and F2 descendants of the Waterman strain reared under controlled optimal laboratory conditions. The results are shown in Tables 1 and 2.

The data indicate that the mean body size of feral beetles of both sexes collected at the Waterman Canyon compare well within the mean size of the domesticated Riverside strain. However, it should be noted that the feral strain, as shown by the coefficient of variation (and as expected) is much more variable in size. This is true for the samples designated as Waterman Canyon II P1 and Waterman Canyon III P1. The F1 and the F2 descendants of the Waterman Canyon, reared under optimal conditions, have considerable increase in body size, and a great reduction in variability.

2. Size in relation to diet.

It is noteworthy that T. brevicornis males are significantly larger than females in respect to weight and body length. This is unusual since T. castaneum and T. confusum females are generally larger than males, as shown by Howe (1968) for Tribolium and Sokoloff (1955) for species of the pseudoobscura subgroup of Drosophila. It also seems unusual for insects in general, since females in species of insects will be larger than males. (Mulder and Sokoloff, 1982).

Tables 3 and 4 summarize body weight and body length when the beetles are reared on different diets. The greatest values are obtained when the beetles are reared on whole wheat (without yeast) or on ground dry insects. An improvement in body weight is observed when the beetles are reared on a mixture of ground dry insects and whole wheat, but the body weight on this mixture is not comparable to that obtained on whole wheat flour plus yeast.

3. Hatchability in different diets

The results are shown in Table 5. Larvae reared on whole wheat or on a mixture of dried insects and whole wheat give comparable values—about 60 per cent of the larvae become adults in these two media. Larvae reared on dried insects alone have a low survival value (about 11 per cent of the larvae become adults) but these adults are of size comparable to that of beetles reared on whole wheat alone. (see Tables 3 and 4)

Roberts (1975) determined the developmental period of Tribolium brevicornis in quart Mason jars filled with one of three media (CY, SY and WY) placing 1000 eggs in each jar. The containers were then placed in an incubator maintained at 32°C and 60% R.H. They were left untouched for 21 days. From then on the contents of each jar were sifted and the adults counted and removed. This procedure was repeated daily until all the larvae had either died or emerged as adults. The length of time involved was different for each medium.

The mean and standard deviation for development in CY was 39 +/- 7.86 days, in WY was 47.2 +/- 7.34 days and in SY was 52 +/- 7.34 days.

4. Fertility

Fertility of the feral strains seems to be of the same order of magnitude as that of domesticated strains such as the Riverside strain as shown in Table 6.

5. Egg cannibalism

Preliminary comparisons show that the Waterman canyon strain is much more cannibalistic than the domesticated Idaho and Riverside strains. Exposure of 50 eggs of *T. brevicornis* to 9 adult beetles of each sex resulted in the loss of 24 per cent of the eggs when the adults were from the feral strain. In comparable experiments only 4 per cent of the eggs were cannibalized by the Riverside strain and 0 per cent by the Idaho strain.

6. Dispersing ability

This was tested by the method of Prus (1963). The Waterman Canyon strain showed a greater tendency to disperse: 47 %. The F1, reared on standard medium, had a lowered tendency to disperse (about 23%). The Idaho and Riverside strains did not show any tendency to disperse out of the test vials.

Another criterion for dispersal, flying ability, was tested using the simple method of Lerner and Ho: Beetles to be tested are placed in a glass petri dish and introduced in a plastic bread or shoe box. At the opposite end of the box is placed another glass petri dish containing food. Flour beetles cannot climb out of the open glass dishes. If any beetles are found outside of the petri dish in the shoe box or in the petri dish containing flour, the beetles must have either hopped out of their dish or flown out to reach the dish containing food. None of the beetles in any of the strains tested showed any tendency to fly when the beetles are tested at room temperature, although when beetles are found to fall on their backs they are capable of, and will, extend their membranous wings.

7. Dispersion and Nutrition.

Roberts (1975) investigated dispersion of *T. brevicornis* as a function of two parameters: 1) population density, and 2) nutritional source provided. He used the technique of Prus (1963) to assess the effect of crowding and the effect of medium on dispersal. Into three sets of vials 4 grams of a medium WY, CY, or SY and beetles at a density of 2, 8, 12, 25, 50, 75, 100 and 200. were introduced. Each vial was connected to an empty vial by a piece of vinyl tubing, with a slender piece of thread running through it. One end of the thread lay on the surface of the medium. The other end hung half way down in the empty vial. At the end of 24, 48, and 72 hours the vials were examined, and the number of beetles recorded as a measure of dispersion. The results are shown in the accompanying Tables 9-11. His experiments show definite differences in the dispersal rates of the beetles for the various media examined. Dispersion rates in WY are the lowest followed by CY and SY. This dispersion undoubtedly reflects to some degree the preferences of *T. brevicornis*. There is dispersion in every trial for SY, with rates of .0138 beetles / hr. to .7500/hr. These rates remain relatively low until a population density of 50 beetles is reached, and from that point on dispersion takes place at a rapid pace. In WY there is relatively low dispersal until a density of 75 beetles is reached, and from that point on dispersion takes place at a rapid pace. WY had a dispersal rate 50-75% lower than either of the other two media. The dispersal from CY was generally intermediate between the other two extremes. Only when the initial population reached 200 did the rate of dispersal exceed that of SY.

These results may be attributable to the fact that these beetles were raised in a WY environment and subsequently the beetles, when placed in any other food source, were unsatisfied and attempted to locate the food source with which they were accustomed. These results are in conflict with those anticipated as a result of the findings in the first area of investigation. It is however noteworthy that while the dispersal rates are high for SY, they are definitely lower for this organism than they are for other species of *Tribolium*. Furthermore, the dispersal figures for SY are not much higher than those for CY, which is an acknowledged food source for this species.

8. Resistance to starvation and lack of water

To test these attributes, pupae were isolated in an empty petri dish and allowed to become adults. 100 of these adults, who were never fed or given a drink, were placed in a petri dish, the dish was covered and placed in a plastic box. The beetles were kept at room temperature (maintained at 25°C and about 60% R.H) and checked periodically for mortality. This experiment was performed twice. The first indicated that the beetles can live over 200 days without being given food or water (Sokoloff, 1971). The second experiment confirmed the conclusions that this species is highly resistant to starvation. At the end of 63 days there were 89 live and 3 dead beetles; at 69 days there were 88 live and 2 dead; at 76 days 87 live and 2 dead; at 90 days, 87 live and the 2 dead beetles which had been eaten except for their heads. After 104 days there were 86 live and 3 dead. After 119 days there were still 86 live but only 2 dead beetles, one of which was intact and the other almost completely devoured except for the head. Fifty percent of the beetles died between days 210 and 217. The longest survivor died at the end of nine months. With an LD50 of about 210 days, T. brevicornis is the species most resistant to starvation. T. castaneum has an LD50 of less than 44 days, T. confusum less than 38 days, and T. madens less than 50 days (Sokoloff and Cavataio, 1971).

9. Production of quinones.

Among the species of Tribolium analyzed by R.K. Ladisch for quinones and hydroquinones (see Table 15.6 in Sokoloff, 1975, for complete data) T. brevicornis is the species which secretes these organic substances most actively. On the basis of mean micrograms per beetle the 20 specimens for each species of Tribolium tested give the following values: (the first values are Quinones and the second Hydroquinones, respectively: T. anaphe: 73.1 + 6.73 and 6.9 + 0.44; T. brevicornis 225.7 + 28.72 and 15.76 + 1.04; T. castaneum 39.5 + 3.43 and 2.38 + 0.25; T. confusum 33.08 + 2.62 and 2.72 + 0.2; T. destructor 135.5 + 12.29 and 4.65 + 0.34; and T. madens 197.7 + 15.18 and 7.9 + 0.86. Thus, in terms of body weight units, T. brevicornis is the species of Tribolium which produced the highest amounts of quinones and hydroquinones.

10. Effect of etherization

Etherization of five minutes will anaesthetize the beetles sufficiently to obtain sex ratio of the sample, or to obtain the phenotype of wild type and mutants in genetics crosses. Etherization has little effect on fecundity, normal levels of egg-laying activity being reestablished in three days. Longer exposures to ether cause a delay in egg-laying activity—the longer the exposure to ether, the greater the period of recovery. LD50 for surviving exposure to ether is about 25 minutes.

Roberts (1975) studied the effect of exposure to ether on mortality and fecundity of T. brevicornis. He isolated 425 pupae and allowed them to become adults. When the beetles were 7-14 days old he subdivided the females into eight groups, each containing 50 of them. 50 males were added to each group. The eight groups were then etherized with anesthesia grade ether (J.T. Baker Chemical Company): for 2, 5, 7, 10, 15, 20, 22, and 25 min., respectively.

After sufficient time for revival had been given (24 hours) the number of dead in each group was counted and the percentage of mortality determined. Every 24 hours the cultures were examined, the eggs were counted, and the mean number of eggs per female determined. This process was carried out for 13 days, until all the females had returned to their normal egg laying schedules.

Roberts found that T. brevicornis females, under optimal conditions (32°C and 70% R. H.) lay 3.38 eggs /24 hour period, which is much less than the egg-laying rate determined by Gray, 1948 and Park and Frank, 1948 who determined an egg laying rate of 15 eggs/day for T. castaneum and T. confusum.

Exposure to ether for 5 minutes had little effect on fecundity of the beetles, and within three days after exposure the females had recovered their normal egg-laying rate. Time in excess of five minutes produced a marked decrease in the number of eggs laid per day. In general the longer the time of exposure,

to ether, the more dramatic the reduction in fecundity, and the greater the time required for the females to recover to their normal egg-laying rate.

The LD50 of etherization in T. brevicornis is 25 minutes, about the same time observed by Sokoloff (1960a, b) for T. castaneum.

11. Effect of diet on survival and development

Roberts (1975) studied the effects of three media : corn plus yeast (CY), wheat plus yeast (WY) and soy plus yeast (SY) on survival and development of T. brevicornis to the adult stage. He concluded that there are definite and discrete differences in the developmental rates of Tribolium brevicornis in the three media examined (CY, WY, and SY): The most rapid development occurred in beetles raised in CY. The earliest emergence occurred at 28 days and the latest at 55 days of development. The mean developmental period in WY was 47.2 +/- 9.49 days, with extremes being 36 and 65 days. Development was by far the longest in SY, with a mean of 52 +/- 7.34 days required for the beetles to reach adulthood, the extremes being the same as in CY. It is interesting to note that the developmental period in CY is still considerably longer (18 days) than that for either T. castaneum or T. confusum.

The percentage of eggs surviving to adulthood varied also with the medium. Those in WY had the greatest chance of reaching adulthood, with 62.5% of the eggs attaining this stage of development, while those in CY had considerably less chance ((37% chance of becoming adults), and those in SY had even a lower probability (31.7%). There are of course many causes of larval mortality, and since the conditions in all three environments were alike we would ordinarily expect to see the same level of emergence. Since this was not the case, we can assume that some of the discrepancies may be due to differences in the media themselves. It appears then that WY is the best medium for the development of this species, followed by CY and finally SY. This experiment shows that Tribolium brevicornis can develop in a medium consisting of SY. However, it would seem that this medium does not provide the necessary nutritional factors for rapid development: in contrast to the more rapid development of this organism in the other two media examined.

12. Feeding and oviposition site preferences in Tribolium

Mickel (1946) and Standish (1947), working respectively with Tribolium confusum and T. castaneum reported that these two species of flour beetles could not tolerate or successfully produce offspring in soybean products. This work was verified by Fraenkel et al (1950) and Lin and Richards (1952). Albers et al. (1974) showed the same dislike in two other species of Tribolium: T. madens and T. audax. Albers et al (1974) showed that T. brevicornis, when given the choice of selecting between corn plus yeast (CY), wheat plus yeast (WY) and soy plus yeast.

(S plus Y) has a distinct preference for SY over the other offerings. These preferences of T. audax, T. brevicornis, T. castaneum, T. confusum and T. madens were determined by introducing beetles of these species into choice plastic chambers in which the media to be chosen were introduced in equal parts into adjacent compartments of equal size separated by a divider which would separate the flour but allowed passage of the beetles from one to the other compartment as they wished. The chambers were placed in an incubator maintained at 32°C and about 60% relative humidity 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 sieve 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 was recorded. The results are summarized in Table 17. Analysis of the data led to the following conclusions:

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

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 medium 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.

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

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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:

Given the choice of WY and SY, T. castaneum, T. audax and T. madens prefer to lay their eggs in wheat rather 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.

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.

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.

It appears that the major reason for the marked aversion to this medium is the presence of a crystalline trypsin inhibitor, which composes approximately 2% of the total weight of a given soy sample. Bowman (1944) and Ham and Standstedt (1945) were the first to successfully demonstrate the ability of soluble fractions of raw soybean to inhibit trypsin. This substance was later isolated by Kunitz (1946). Additional work by Liener (1953) and Almquist and Merritt (1953) working with vertebrates provided a great wealth of evidence for the presence of anti-trypsins in soybean. Like, Fraenkel and Liener (1954) determined that of all the inhibitors present in soy, it the antitrypsins that were the most effective in halting growth and development in all species of Tribolium. Building upon work first commenced by Borchers and Akerson (1952) they discovered that the addition of 20% yeast to soy produced beneficial effects upon the growth of Tribolium. According to most insect nutritionists, this is due to an antagonism between the principle in raw soybean which is toxic to the larvae and the nutrients present in yeast. Most entomologists believe that pests of stored products possess specific sensory mechanisms which enable them to select yeast particles at the expense of the ration. Soybean inhibitors which are toxic to insects have made soy immune and basically impregnable to attack from flour beetles.

Using the preliminary findings of Sokoloff et al, 1966, Sokoloff 1975, and Albers et al (1974) Roberts carried out experiments concentrating on T. brevicornis. In his experiments, Roberts isolated 2000 T. brevicornis pupae according to sex in a jar containing a great excess of WY medium. The pupae were allowed to develop into adults and to mature for 10 days. 50 beetles (25 of each sex) were then introduced into each of four plastic containers such as those designed by Inouye (1965). Each of these choice chambers contained equal quantities of two media arranged in the following combinations: CY-WY, CY-SY, WY-SY, SY-S. The media were separated by a plastic divider, which had sizeable perforations to allow the free passage of the beetles from one side to the other. The fifty beetles were placed in the center of the container, with approximately 25 beetles on each side of the divider. The chambers were then closed with lids containing small openings for ventilation and placed in an incubator at 32°C and a relative humidity of 60% for a period of 48 hours. After the time interval had expired the contents of each side of the chamber were aspirated into a vial. The medium was then sifted with a coarse sieve to remove the adults and the number of eggs laid in each medium recorded. Altogether there were ten replicate choice chambers set up for each combination of media. The data are shown in Tables 12-15. The results were as follows:

13. Preference of Medium in Adults

The data showing the preference of medium by adults are given in Tables 12-15. The results can be summarized as follows:

1. Given the choice of CY and WY a majority of the beetles showed a preference for CY. With a mean 28.3 +/- 1.86 in that medium as compared with 21.8 +/- 1.93 beetles in the WY. The females had a distinct preference for CY over the other offering, while the males were comparatively indifferent
2. Given the choice of CY and SY the beetles were found fairly equally distributed, with about 25.2 +/- 2.29 beetles in CY as compared to 24.8 +/- 2.29 in the SY. In this case neither sex showed a preference for either medium.
3. Given the choice of WY and SY, T. brevicornis mostly chooses SY. A mean of 27.4 +/- 1.74 beetles were found in the SY in contrast to 22.6 +/- 1.76 in the WY. The males appeared to be apathetic in choosing a medium, while the females showed a marked preference for SY.
4. Given the choice of SY and S, the beetles were found to demonstrate no preference for either medium. There was a mean of 25.2 +/- 1.52 beetles in S as compared to 24.8 +/- 1.89 in SY. Neither sex showed a specific preference for either medium.

14. Oviposition Site Preference

1. Given the choice between WY and CY, T. brevicornis showed greater preference for CY than for WY. With a mean of 105.6 +/- 3.76 eggs deposited in CY to that of 82.9 +/- 4.38 in WY.
2. Given the choice between CY and SY T. brevicornis is found to lay its eggs about evenly in the two media, with a mean of 93.5 +/- 4.75 eggs being laid in Cy to 96.4 +/- 4.94 in SY.
3. Given the choice of WY and SY, T. brevicornis has an overwhelming preference for SY with a mean of 103.1 +/- 4.23 eggs being laid in SY as compared with 68.3 +/- 3.62 eggs laid in WY.
4. Given the choice between SY and S, T. brevicornis reveals no partiality. Its eggs are deposited equally throughout the two media with a mean of 90.7 +/- 3.19 eggs laid in SY as compared with 89.9 +/- 4.2 eggs in S.

From the data, which largely corroborate those obtained by Albers et al (1974), Roberts concludes that T. brevicornis has the following preference for the media studied :

1. Nutritional preference: CY>SY>S>WY.
2. Ovipositional preference: CY>SY>S>WY

The results clearly show that T. brevicornis can tolerate , survive and flourish on soybean products, as well as or better than in other media as compared to other species.

Sokoloff et al. (1965) referred to Orr and Watt's (1957) table summarizing the amino acid, mineral and vitamin content of the four flour food sources used in this experiment. The table shows that soy is richer or as rich as yeast and the other two grains, corn and wheat in amino acids and certain minerals as alluded to by Block and Bolling (1951) Block and Weiss (1956) and Watt and Merrill (1963), However soy contains various toxic substances:

Soyin 2.5%
 Trypsin inhibitor 2%
 A Ph 4.2 extract of soybean
 A Kaolin treated acid extract of soybean
 Ethyl alcohol soluble inhibitor
 An alkaline extract of soybean and Soyin (from Lipke, 1954)

These substances affect the developmental rate and greatly impede the digestive processes of T. castaneum, T. confusum, T. audax and T. madens and thus make it very poor nutritional and ovipositional site for these species.

From the results obtained by Roberts, and others mentioned above, it appears that T. brevicornis possesses some unique qualifications that allows it to exploit this medium more effectively than the other species of Tribolium investigated.

14. Exposure to radiation

The egg is the stage most sensitive to radiation (1,000 rads).

15. T. brevicornis as host to Hymenolepis diminuta

T. brevicornis is a potential host of Hymenolepis diminuta. Mankau (1977) has shown that size is a factor in the infection Tribolium spp. and Eleodes sp. by Hymenolepis diminuta.

16. Cytology of T. brevicornis

S. G. Smith (1952) conducted cytological studies in 191 species of Coleoptera distributed in 127 genera, 66 families and 51 superfamilies. He concluded that the primitive number in these beetles the primitive number of chromosomes is nine pairs of autosomes, an X of the size of the autosomes, and a minute y, both being V-shaped and associated during maturation divisions at two terminal contact points in the form of a parachute. The formula $9AA + X + y$ is characteristic of the Tenebrionoidea as a whole, but a number of species have fewer chromosomes. They are considered to be derived species.

Within the genus T. castaneum has 9 pairs of autosomes, an X about the size of the autosomes, and a small y, these two appearing in the form of a parachute, conforming with the primitive condition; T. madens has also this primitive formula, but in addition has a small number of supernumeraries which differ from such species as Diabrotica in that they are able to form bivalents in the spermatocyte (Smith 1956, 1960); T. confusum and T. destructor are derived species, the first having 8 pairs of autosomes, a large X, and a large Y (neo-X and neo-Y); and the size of the neo-X and neo-Y being again reduced in T. destructor. (For further details see Sokoloff, 1972). Halstead (1969) compared the cytology of T. audax a species closely related to T. madens and found no detectable differences (both having nine pairs of autosomes plus three pairs of supernumeraries. Plus Xyp (parachute shaped)

Moore and Sokoloff (1982) have found that Tribolium brevicornis and Tribolium anaphe have nine pairs of autosomes. Both species had a somatic chromosome number of 18. There was no evidence of a heteromorphic sex chromosome pair in these two species. See Fig. 00.

Discussion

The data gathered in the present study suggest that the domesticated and feral strains of T. brevicornis differ in three of the attributes tested: body size, cannibalism and dispersion.

The feral population has a greater difference in body size. This difference is primarily in the coefficient of variability, the Waterman Canyon strain exhibiting much greater variability than the domesticated Riverside and Idaho strains. The means of the three strains are comparable, which indicates that, overall, the feral strain is able to obtain sufficient food during the development to produce beetles comparable in size to those reared in the laboratory under optimal conditions. It is noteworthy that among the beetles derived from nature, a certain proportion are very small beetles-about one ninth of one tenth of the weight of the largest beetles. This small size apparently is due to environmental and not to genetic causes because both sexes are represented among the small beetles, and furthermore, when small females are isolated they produce F1 comparable in size to the progeny produced by larger females.

The feral population showed a much greater tendency toward egg-cannibalism (20% of the eggs were destroyed by the Waterman Canyon strain) than the domesticated strains (less than 8% for the Riverside strain and 0% for the Idaho strain). This is particularly surprising for the Idaho strain, since beetles in Moscow, Idaho, have become predaceous of larvae of Megachile pacifica, the alfalfa leafcutter bee (which commercial growers raise in an attempt to improve the yield of alfalfa).

The feral population showed a much greater tendency to disperse than the Riverside or Idaho strain. When the feral population was compared with their F1, it was clear that the latter did not disperse as readily as the parental population.

In respect to the other attributes (length of development, fecundity, fertility, mortality of the juvenile stages, etc.) the three strains give similar results.

Successful stored-products pests, according to Howe (1962) need the ability to eat dry foods and use and conserve the water produced in the breakdown of carbohydrates. On the basis of evidence derived from the laboratory and from scant observations in the field, it is highly probable that the various species of Tribolium are omnivorous. Initially, they probably lived under the bark of trees, and Howe (l.c.) and this reporter (Sokoloff, 1974) believe that the beetles were primarily carnivorous, eating as larvae or adults, eggs and pupae of their own or other species of insects. They were secondarily herbivorous, perhaps consuming fungi and bacteria as a source of vitamins and minerals and perhaps carbohydrates as a source of metabolic water.

Our attempts at providing a clearcut answer to the question of the role of T. brevicornis in nature, in the decaying log biocoenosis, were not completely successful. T. brevicornis has also started to move as predators into another man-made environment, the bee-boards where the alfalfa leafcutter bee Megachile pacifica raises its young (Polk, 1979). This illustrates the great plasticity of these insects: they can survive as carnivores (on the larvae or pupae in bee nests), or as herbivores (on wheat flour or other stored products) equally well.

Summary

Data are presented to show that, a normal inquiline of Xylocopa tabaniformes orpifex differs from domesticated strains in body size (feral populations more variable than domesticated strains) and two behavioral traits: cannibalism and dispersal (feral strains are more cannibalistic and have greater tendency to disperse).

The primary ecological niche of T. brevicornis is not completely established, owing to the necessity to disturb the habitat by sawing into the log. However, the differences in behavior and the gut contents of these insects suggest that in nature these beetles play either the role of predators or scavengers of Xylocopa.

Interestingly, T. brevicornis was found in Xylocopa nests but did not occur in a beehive of Apis mellifera found only three meters away in a downed sycamore, even though the latter would provide a richer source of food than the nests of Xylocopa. Apparently the hygienic behavior of the honey bee is very effective, and discourages the establishment of T. brevicornis in bee nests while Xylocopa bees have not developed this hygienic behavior. In any case, adults and larvae of T. brevicornis have been found in the galleries of Xylocopa. Larvae of T. brevicornis have been collected from pupae of Xylocopa on which the larvae of brevicornis were feeding. Furthermore, recently collected adult beetles from dead bodies of Xylocopa contained fragments of Xylocopa exoskeletons in their digestive tracts. The evidence gathered so far indicates that T. brevicornis is probably a secondary or tertiary consumer, engaging in scavenging, predatory and possibly cannibalistic activities within the decaying log biocoenosis. (Sokoloff et al. 1983).

T. brevicornis and T. audax have only 9 pairs of chromosomes. In both species there is no evidence of a distinguishing heteromorphic X-Y pair.

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Table 1

Body length of various strains of *T. brevicornis* reared in standard medium at 32°C and 70% R.H.

Strain	Males					Females				
	N	\bar{m}	σ	SE	CV(%)	N	\bar{m}	σ	SE	CV(%)
Riverside	61	6.257	.4414	.0565	7.05	60	5.952	.3073	.0397	5.16
Waterman Canyon II P ₁	49	5.963	1.002	.143	16.80	50	6.146	.7299	.1032	11.88
Waterman Canyon III P ₁	84	5.542	.906	.0989	16.35	88	5.602	.7900	.08422	14.10
Waterman Canyon II F ₁	90	6.709	.3798	.0400	5.66	90	6.672	.3784	.03989	5.67
Waterman Canyon II F ₂	61	6.8	.3710	.0475	5.46	60	6.59	.3900	.5035	5.92

Table 2

Body weight of various strains of *T. brevicornis* reared in standard medium at 32°C and 70% R.H.

Strain	Males					Females				
	N	\bar{m}	σ	SE	CV(%)	N	\bar{m}	σ	SE	CV(%)
Riverside	61	9.879	1.783	.2282	18.05	60	8.422	1.434	.1851	17.03
Waterman Canyon II P ₁	49	9.780	3.930	.5615	40.18	50	9.912	2.977	.4210	30.03
Waterman Canyon III P ₁	84	8.140	4.0762	.4448	50.08	88	6.715	3.020	.3219	44.97
Waterman Canyon II F ₁	90	12.783	1.768	.1863	13.83	90	12.784	2.024	.2134	15.83
Waterman Canyon II F ₂	61	13.948	2.171	.2780	15.56	60	12.975	1.796	.2319	14.04

Table 3

Body weight of T. brevicornis (Waterman Canyon II strain) reared under different diets at 32°C and 70% R.H.

	Males					Females				
	N	\bar{m}	σ	SE	CV(%)	N	\bar{m}	σ	SE	CV(%)
A. Whole Wheat	32	7.37	1.67	.2952	22.66	26	6.92	2.01	.3942	28.11
B. Ground Dry Insects	25	7.30	2.31	.462	31.64	20	7.15	2.59	.5791	36.22
C. Mixture of A + B	35	9.65	2.29	.4092	23.95	25	9.09	2.53	.506	27.83
D. Whole Wheat + Yeast	22	12.973	1.613	.3439	12.43	20	12.545	1.330	.2974	10.60

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Table 4

Body length of various strains of T. brevicornis reared in standard medium at 32°C and 70% R.H.

	Males					Females				
	N	\bar{m}	σ	SE	CV(%)	N	\bar{m}	σ	SE	CV(%)
A. Whole Wheat	29	5.88	.500	.9285	8.50	29	5.73	.402	.07466	7.02
B. Ground Dry Insects	22	5.688	.529	.1128	9.30	23	5.655	.500	.1043	8.84
C. Mixture of A + B	30	6.109	.441	.08052	7.22	30	6.032	.502	.06714	8.32

Notes - Research, Teaching and Technical

Table 5

Hatchability of T. brevicornis larvae reared in different diests at 32°C and 70% R.H.

	Initial Larvae (N)	Dead (N)	Survival (Larva to adult) (%)
A. Whole Wheat	100	42	58
B. Dried Insects	400	355	11.25
C. A & B	100	40	60

Table 6

Fertility of various T. brevicornis strains incubated at 32°C and 70% R.H.

Strain	Sample Size	Survival to larval stage	Fertility (%)
Riverside	89	58	65
Waterman Canyon I	110	84	76
Waterman Canyon II	156	96	61.5

Table No. 1. Dispersion from Wheat plus Yeast

# beetles per test	Dispersion after 24 hours	Dispersion after 48 hours	Dispersion after 72 hours
2	0	0	0
8	0	0	0
12	0	0	0
25	0	0	1
50	7	7	8
75	9	11	17
100	12	16	18
200	8	21	31

Table No. 2. Dispersion from Soy plus Yeast

# beetles per test	Dispersion after 24 hours	Dispersion after 48 hours	Dispersion after 72 hours
2	1	1	1
8	0	0	1
12	2	2	2
25	1	1	2
50	13	20	24
75	31	40	43
100	36	39	48
200	26	50	54

Table No. 3. Dispersion from CY

# beetles per test	Dispersion after 24 hours	Dispersion after 48 hours	Dispersion after 72 hours
2	1	1	1
8	2	2	2
12	0	0	0
25	1	3	3
50	3	3	12
75	9	13	18
100	23	28	32
200	32	55	59

Table No. 10 Dispersal in terms of Percent Analysis

# beetles per test	% beetles leaving SY	% beetles leaving WY	% beetles leaving CY
2	50%	-	50%
8	12.5%	-	25%
12	16.6%	-	-
25	8%	4%	12%
50	48%	16%	24%
75	57.3%	22.6%	24%
100	48%	18%	32%
200	27%	15.5%	29.5%

Table No. 11 Dispersal rate of *T. brevicornis* in different media as a function of density at the end of 72 hours.

# beetles in vial	SY	WY	CY
2	.0138/hr	-	-
8	.0138/ hr	-	.0138/ hr
12	.0277/ hr	-	.2111/ hr
25	.0277/ hr	.0138/ hr	.0416/ hr
50	.3333/ hr	.1111/ hr	.1666/ hr
75	.5970/ hr	.2361/ hr	.2500/ hr
100	.6666/ hr	.2500/ hr	.4400/ hr
200	.7500/ hr	.4300/ hr	.8190/ hr

Table No. 12, Choice Chamber WY-CY

	Wheat plus Yeast				Corn plus Yeast			
	males	females	Total	eggs	males	females	Total	Eggs
1	11	13	24	99	14	12	26	127
2	13	15	28	78	12	10	22	89
3	9	12	21	95	16	13	29	97
4	12	5	17	37	13	20	33	103
5	12	9	21	66	13	16	29	111
6	13	13	26	110	12	12	24	101
7	14	6	20	64	11	19	30	99
8	13	13	26	114	12	12	24	93
9	12	9	21	113	13	17	30	104
10	9	5	14	53	16	20	36	132
Total	118	100	218	829	132	151	283	1056

X and standard deviation:

	WY	CY
Males	11.8 +, -1.19	13.2 +, - 1.23
Females	9.9 +, - 1.88	15.1 +, - 1.91
Total	21.8 +, - 1.93	28.5 +, - 1.96
Eggs	82.9 +, - 4.38	105.6 +, - 3.76

Table No. 13, Choice Chamber CY-SY

	Corn plus Yeast				Soy plus Yeast			
	males	females	total	eggs	males	females	total	Eggs
1	12	11	23	103	13	14	27	109
2	15	6	21	71	10	19	29	134
3	7	5	12	49	18	20	38	79
4	14	19	33	128	11	6	17	82
5	21	8	29	103	4	17	21	114
6	11	15	26	103	14	10	24	123
7	18	12	30	95	7	13	20	85
8	11	10	21	102	14	15	29	112
9	8	21	29	122	17	4	21	54
10	13	15	28	59	12	10	22	72
total	130	122	252	935	120	128	248	964

X and standard deviation:

	CY	SY
Males	13 +, - 1.91	12 +, - 1.88
Females	12.2 +, - 2.17	12.8 +, - 2.18
Total	25.2 +, - 2.29	24.8 +, - 2.29
Eggs	93.5 +, - 4.75	96.4 +, - 4.94

Table No. 14. Choice Chamber WY-SY

	Wheat plus Yeast				Soy plus Yeast			
	males	females	total	eggs	males	females	total	eggs
1	13	9	22	68	12	16	28	131
2	12	8	20	59	13	17	30	142
3	11	8	19	66	14	17	31	98
4	11	18	29	85	14	7	21	72
5	13	11	24	64	12	14	26	105
6	11	7	18	51	14	18	32	112
7	15	6	21	52	10	19	29	106
8	15	11	26	88	10	14	24	98
9	14	11	25	91	11	14	25	95
10	12	10	22	59	13	15	28	72
Total	127	99	226	683	123	151	274	1031

X and standard deviation:

	WY	SY
Males	12.2 +/- 1.20	12.3 +/- 1.20
Females	9.9 +/- 1.59	15.5 +/- 1.59
Total	22.6 +/- 1.74	27.4 +/- 1.74
Eggs	68.3 +/- 3.62	103.1 +/- 4.23

Table No. 15. Choice Chamber SY-S

	Soy plus Yeast				Soy			
	males	females	total	eggs	males	females	total	eggs
1	13	13	26	96	12	12	24	89
2	12	13	25	97	13	12	25	90
3	18	14	32	89	7	11	18	84
4	12	11	23	69	13	14	27	48
5	12	14	26	98	13	11	24	67
6	11	8	19	89	14	17	31	137
7	11	13	24	90	14	12	26	106
8	14	11	25	105	11	14	25	109
9	15	16	31	102	10	9	19	84
10	10	7	17	92	15	18	33	85
Total	128	120	248	907	122	130	252	899

X and standard deviation:

	SY	S
Males	12.8 +/- 1.39	12.2 +/- 1.43
Females	12.0 +/- 1.59	13.0 +/- 1.66
Total	24.8 +/- 1.89	25.2 +/- 1.52
Eggs	90.7 +/- 3.19	89.9 +/- 4.28

Table 16 Amino Acid, Mineral and Vitamin Content of Seven Foods in mg/100g Edible Portion

Ingredient	Food						
	Corn	Brown rice	White rice	Soybean flour (full-fat)	Whole wheat flour	White wheat (patent)	Brewer's yeast (dried)
Amino Acid*							
Tryptophan	47	81	82	541	164	129	710
Threonine	311	294	298	1547	383	302	2353
Isoleucine	361	352	356	2112	577	483	2398
Leucine	1011	646	655	3030	862	809	3226
Lysine	225	296	300	2483	365	239	3300
Methionine	145	135	137	528	203	138	836
Cystine	101	102	103	698	292	210	548
Phenylalanine	246	237	240	1226	495	348	1384
Tyrosine	354	377	382	1943	657	577	1902
Valine	398	524	531	2062	616	453	2723
Arginine	275	432	438	2842	636	466	2250
Histidine	161	126	128	937	271	210	1251
Alanine	776	-	-	1616	465	317	3456
Aspartic Acid	968	354	359	4766	725	455	5232
Glutamic Acid	1377	1027	1041	7211	4156	3653	6334
Glycine	265	513	520	1641	812	381	2427
Proline	651	363	368	2641	1389	1254	1983
Serine	441	381	386	2565	614	542	2657
Minerals							
Ash	800	1200	500	4600	1700	430	7100
Calcium	6	32	24	199	41	16	210
Phosphorus	(164)	221	94	558	372	87	(1753)
Sodium	(1)	9	5	1	3	2	(51)
Potassium	-	214	92	1660	370	95	1894
Vitamins							
Thiamine	0.20	0.34	0.07	0.85	0.55	0.06	15.61
Riboflavin	0.06	0.05	0.03	0.31	0.12	0.05	4.28
Niacin	1.4	4.7	1.6	2.1	4.3	0.9	37.9

* Adapted from ORR and Watt (1957). The above foods have the following protein content and nitrogen conversion factors: corn - 7.8% protein, n * 6.25; rice, brown - 7.5% protein, n*5.95; rice, polished - 7.6% protein, n*5.95; soybean flour (full-fat) - 35.9% protein, n*5.71; wheat flour-whole - 13.3% protein, n*5.83; patent flour - 10.5% protein, n*5.70; and brewer's yeast (dried) - 7.4% protein, n*6.25 (here the total n is equivalent to 36.9 protein).

Notes - Research, Teaching and Technical

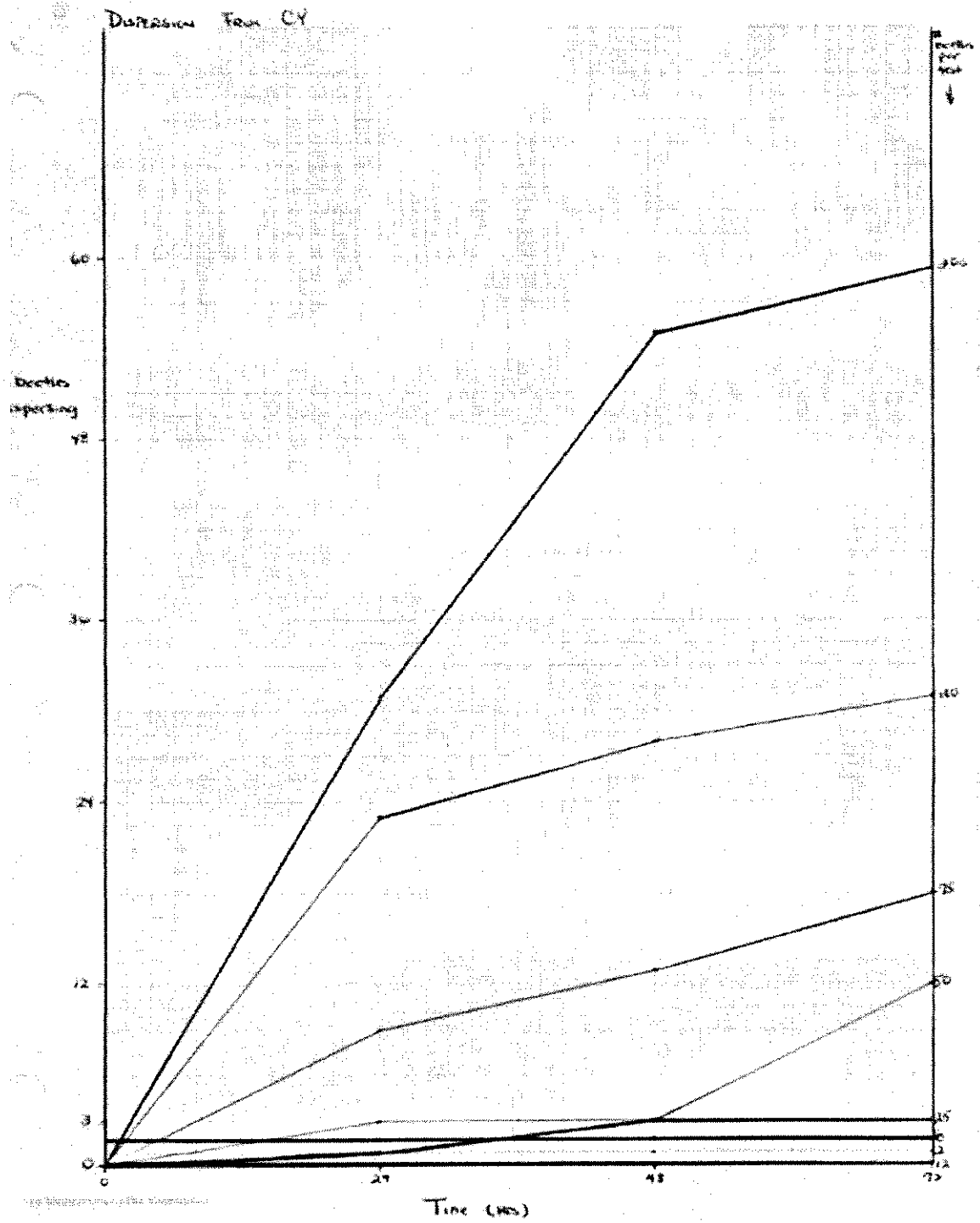
Table 16a Amino Acid, Mineral and Vitamin Content of Seven Foods in mg/100g Edible Portion

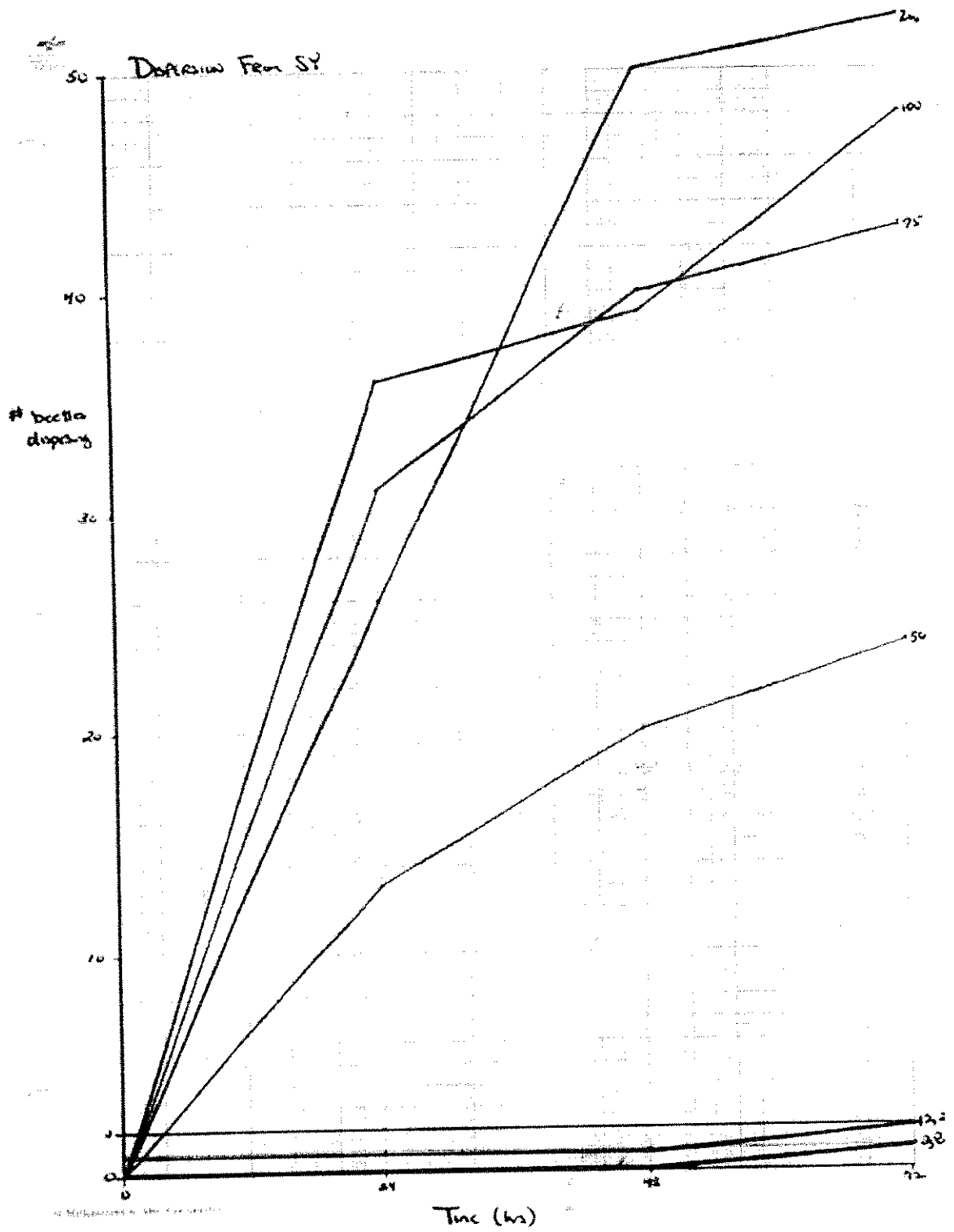
Ingredient	Food						
	Corn	Brown rice	White rice	Soybean flour (full-fat)	Whole wheat flour	White wheat (patent)	Brewer's yeast (dried)
Amino Acid*							
Tryptophan	47	81	82	541	164	129	710
Threonine	311	294	298	1547	383	302	2353
Isoleucine	361	352	356	2112	577	483	2398
Leucine	1011	646	655	3030	862	809	3226
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Methionine	145	135	137	528	203	138	836
Cystine	101	102	103	698	292	210	548
Phenylalanine	246	237	240	1226	495	348	1384
Tyrosine	354	377	382	1943	657	577	1902
Valine	398	524	531	2062	616	453	2723
Arginine	275	432	438	2842	636	466	2250
Histidine	161	126	128	937	271	210	1251
Alanine	776	-	-	1616	465	317	3456
Aspartic Acid	968	354	359	4766	725	455	5232
Glutamic Acid	1377	1027	1041	7211	4156	3653	6334
Glycine	265	513	520	1641	812	381	2427
Proline	651	363	368	2641	1389	1254	1983
Serine	441	381	386	2565	614	542	2657
Minerals							
Ash	800	1200	500	4600	1700	430	7100
Calcium	6	32	24	199	41	16	210
Phosphorus	(164)	221	94	558	372	87	(1753)
Sodium	(1)	9	5	1	3	2	(51)
Potassium	-	214	92	1660	370	95	1894
Vitamins							
Thiamine	0.20	0.34	0.07	0.85	0.55	0.06	15.61
Riboflavin	0.06	0.05	0.03	0.31	0.12	0.05	4.28
Niacin	1.4	4.7	1.6	2.1	4.3	0.9	37.9

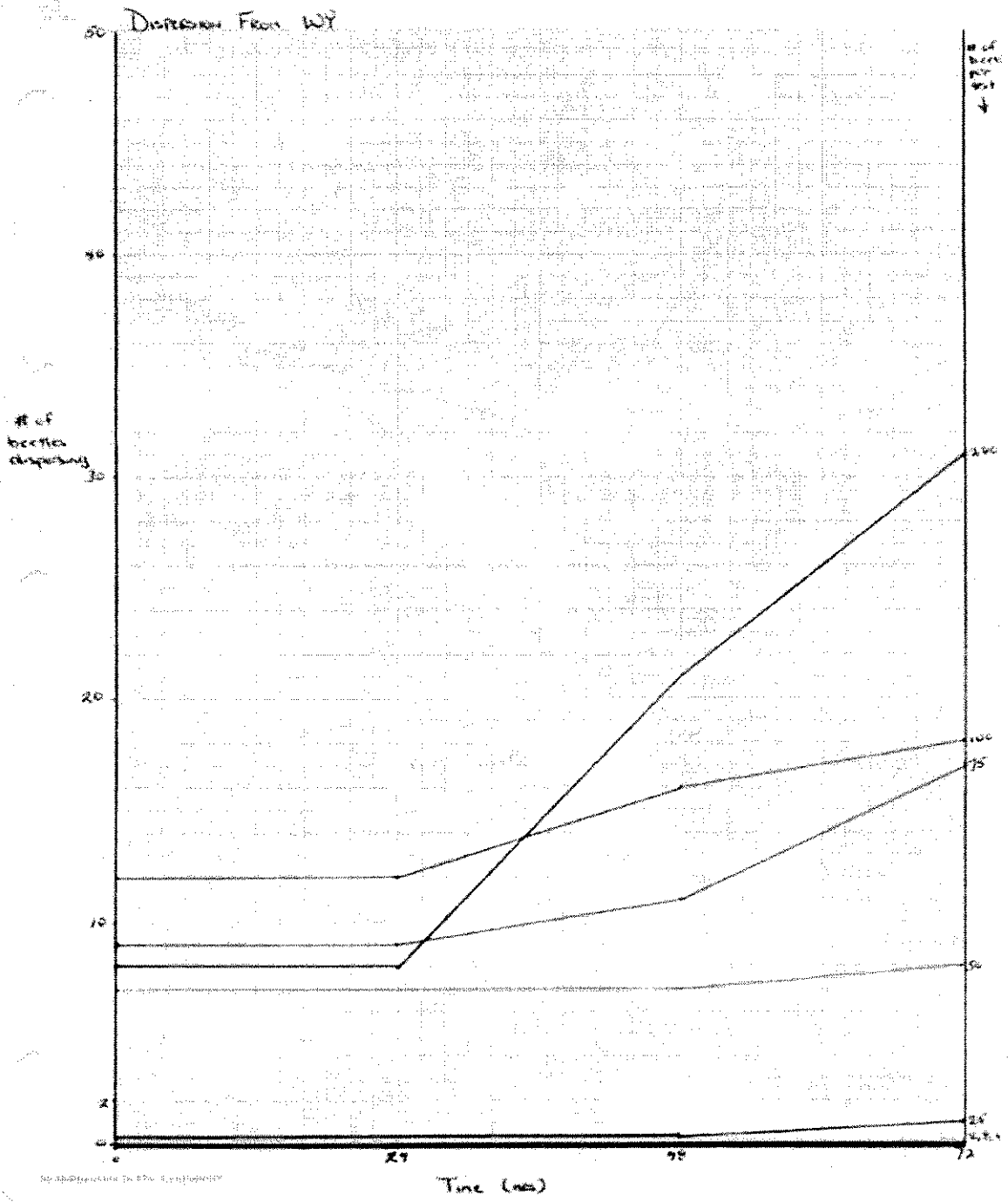
* Adapted from ORR and Watt (1957). The above foods have the following protein content and nitrogen conversion factors: corn - 7.8% protein, n * 6.25; rice, brown - 7.5% protein, n*5.95; rice, polished - 7.6% protein, n*5.95; soybean flour (full-fat) - 35.9% protein, n*5.71; wheat flour-whole - 13.3% protein, n*5.83; patent flour - 10.5% protein, n*5.70; and brewer's yeast (dried) - 7.4% protein, n*6.25 (here the total n is equivalent to 36.9 protein).

Table 17. Food choice and oviposition of 5 species of Tribolium

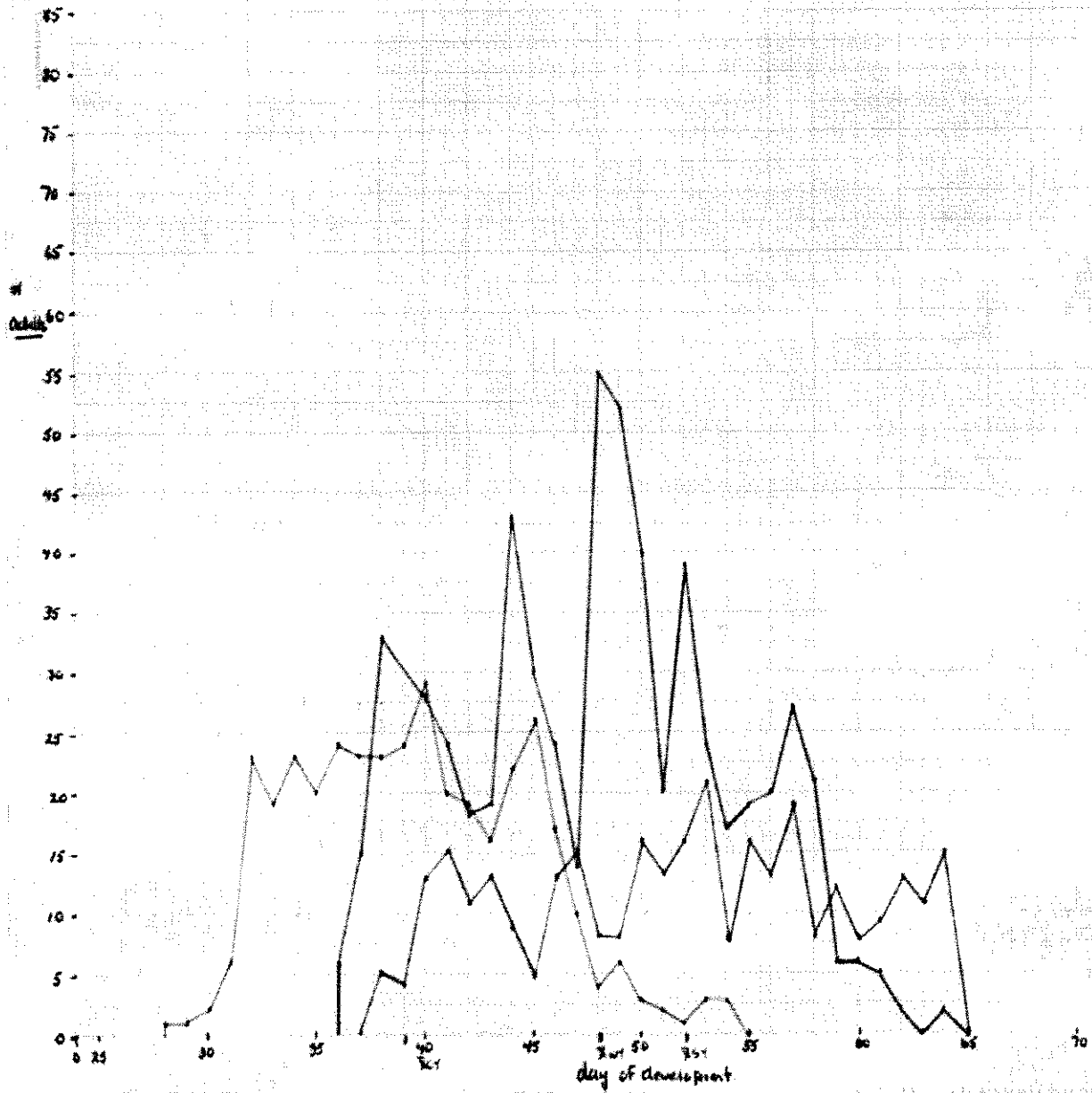
	<u>T. audax</u>			<u>T. brevicornis</u>			<u>T. castaneum</u>			<u>T. confusum</u>			<u>T. madens</u>		
	M	F	Eggs	M	F	Eggs	M	F	Eggs	M	F	Eggs	M	F	Eggs
<u>Media</u>															
WY	22	13	222	11	10	41	17	24	225	12	12	38	14	25	51
SY	3	5	27	16	10	67	0	1	9	1	7	49	5	6	12
WY	10	7	39	10	11	48	13	16	175	7	8	61	6	14	49
CY	14	16	188	12	13	36	6	12	15	12	20	117	15	12	58
CY	25	24	278	13	11	57	14	26	31	13	36	186	28	20	87
SY	0	0	0	14	13	106	0	1	3	1	0	3	2	0	0

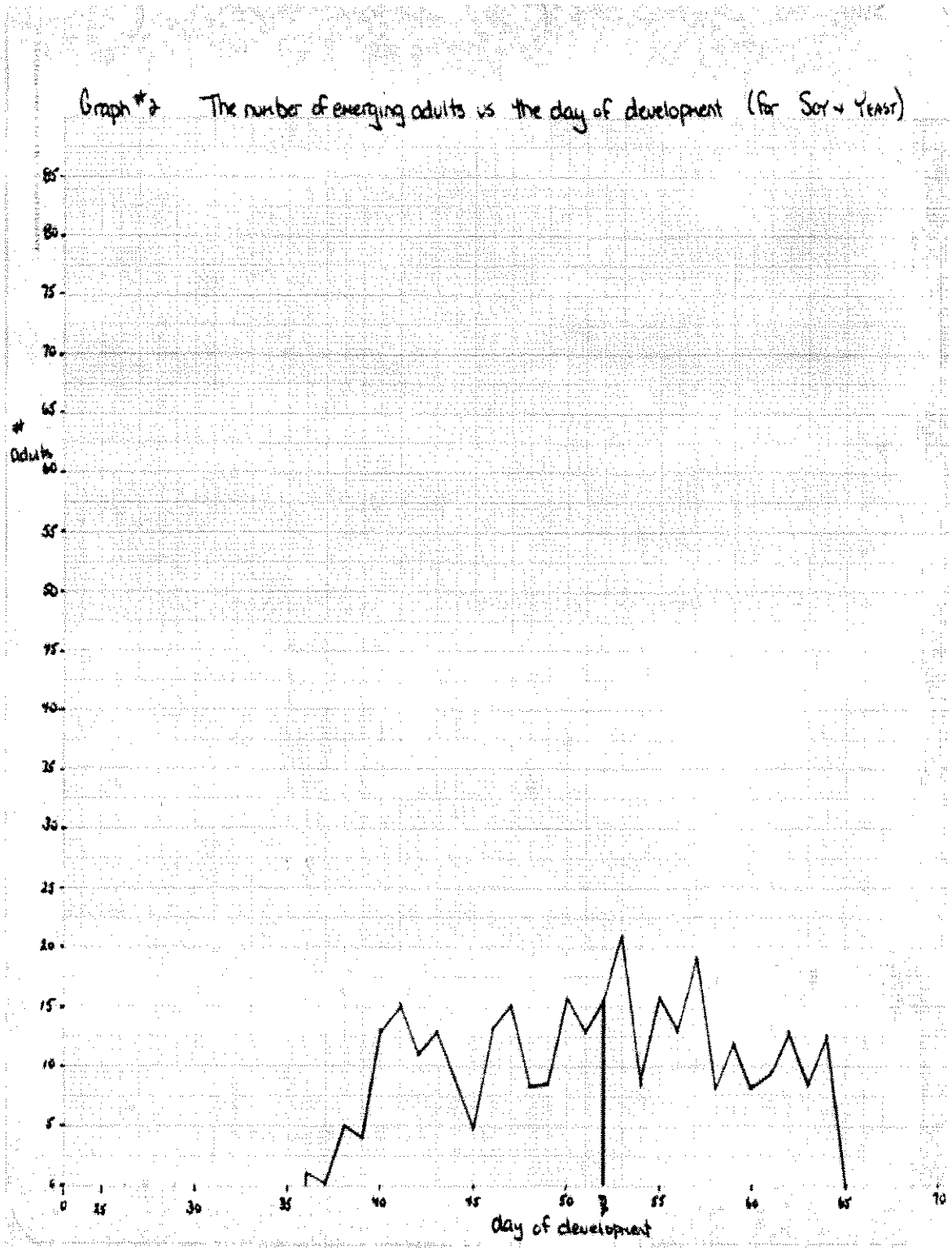




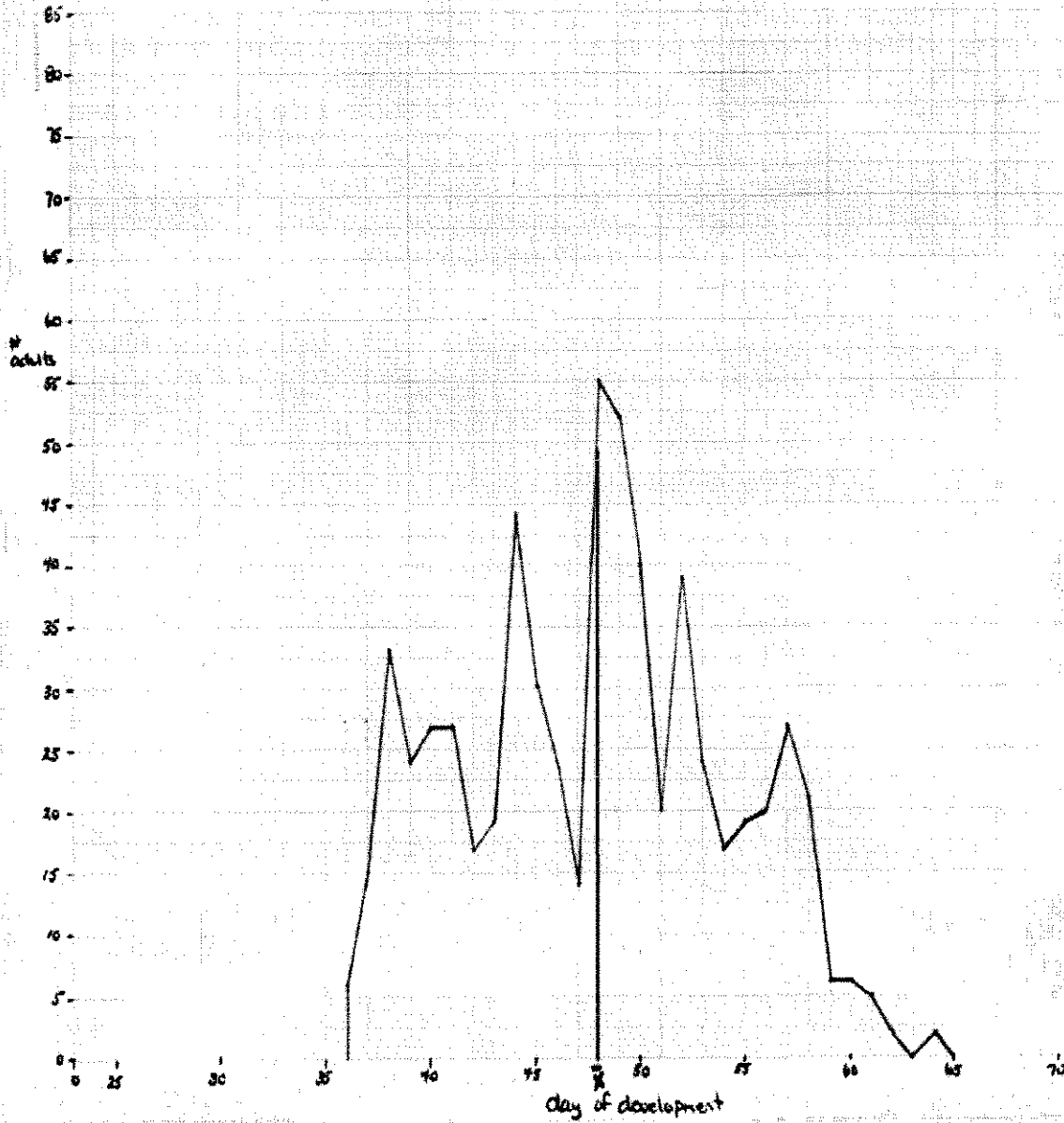


Graph #1 The number of emerging adults vs. the day of development. (for each media studied)

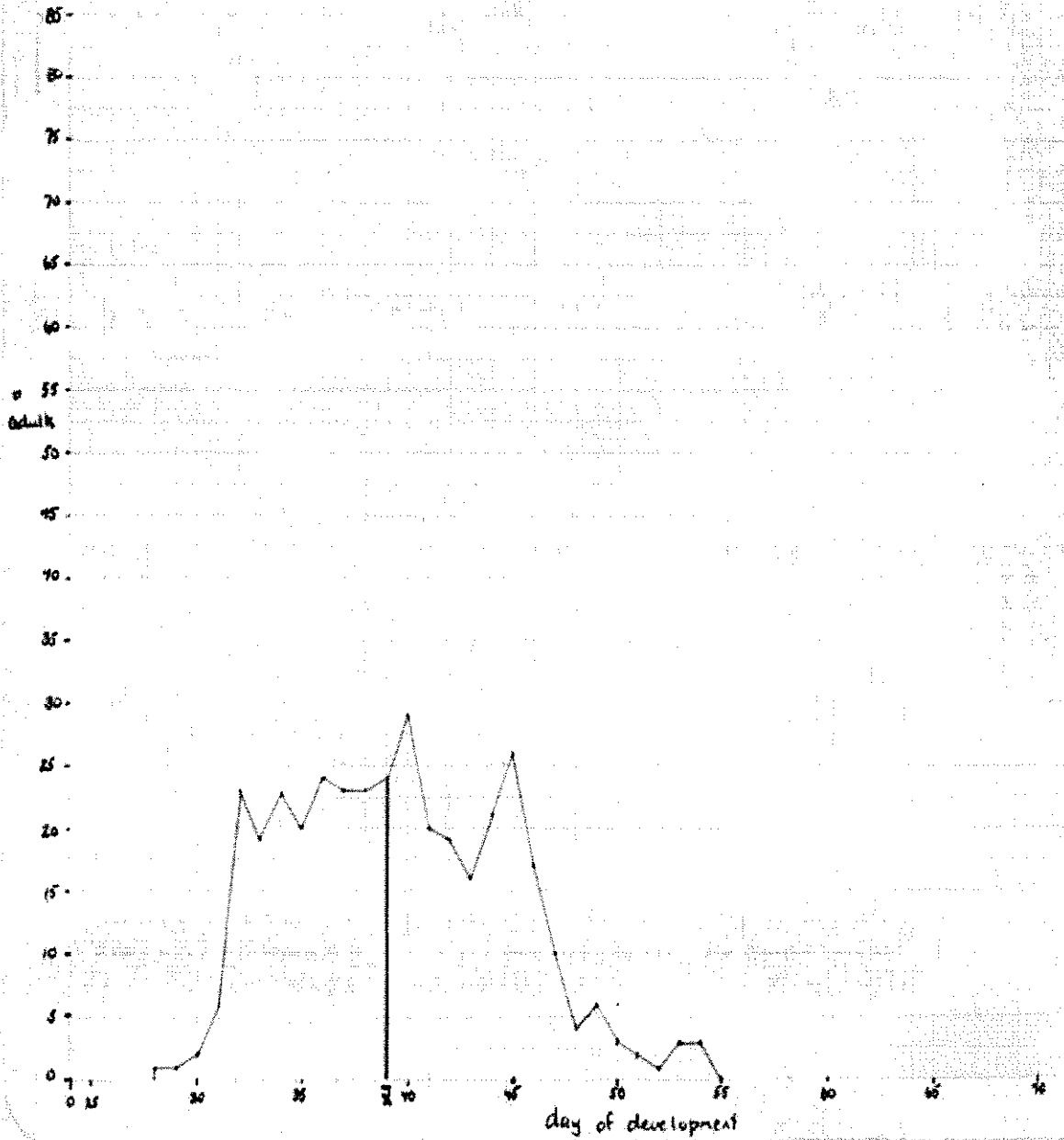


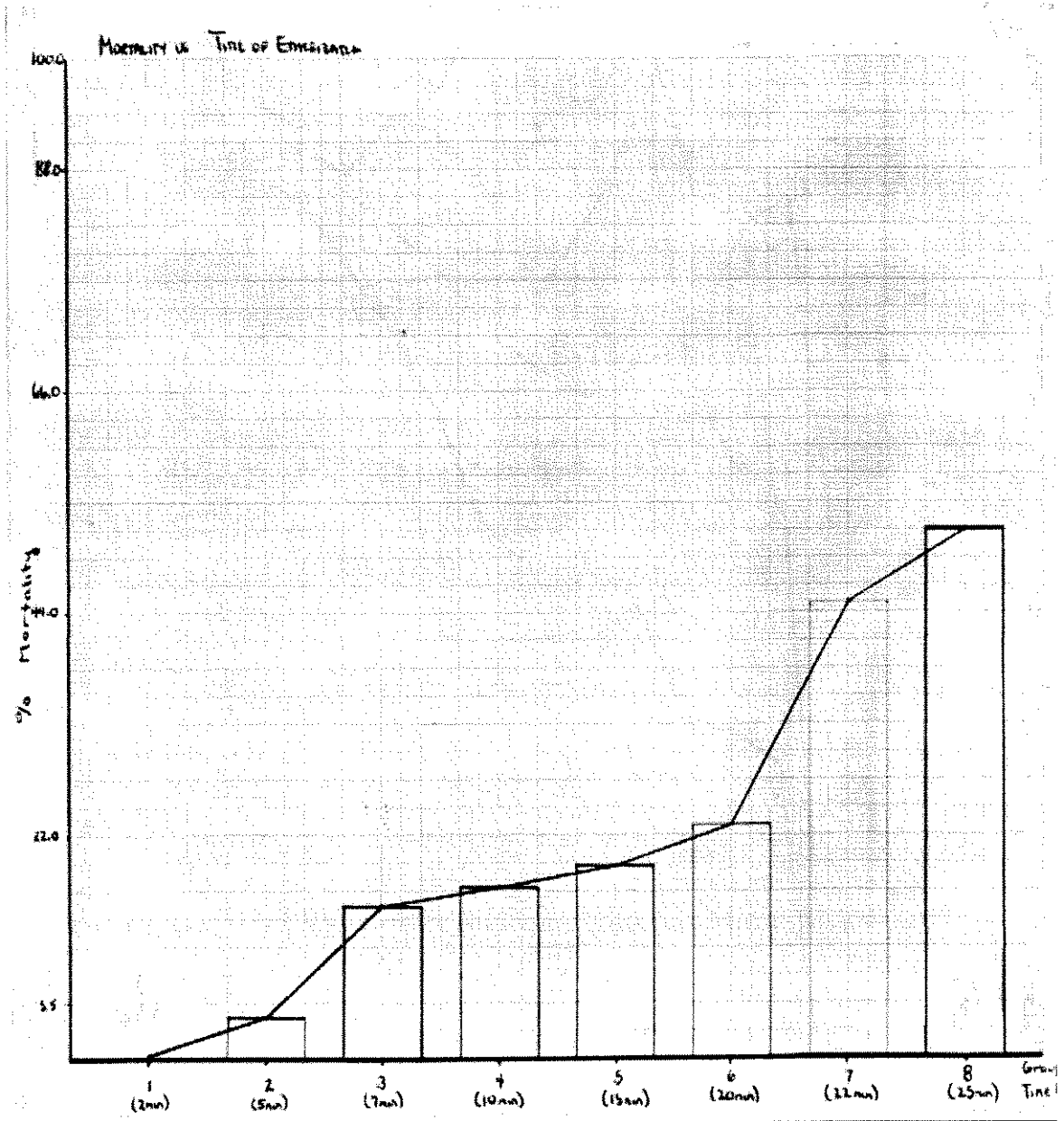


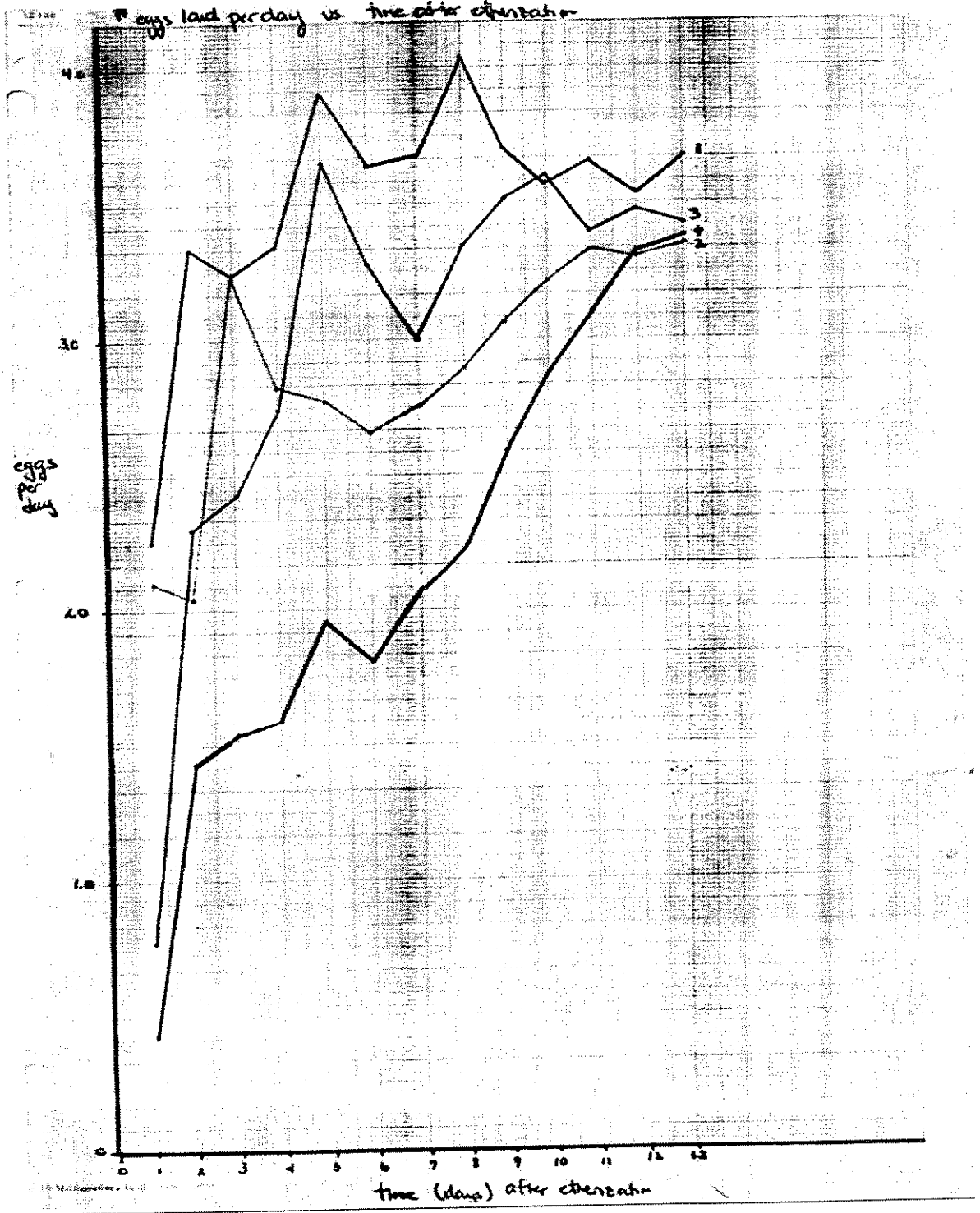
Graph #3 The number of emerging adults vs. the day of development (for wheat + yeast)



Graph #4 The number of emerging adults vs. the day of development (for Corn + Yeast)







I. Determination of Mean Egg Laying Schedule 400 females examined over 72 hour period.

Day	Loss	\bar{x} eggs per female	
1	1353	3.38 ± .66	$\bar{x} = 3.38 \pm .64$
2	1383	3.46 ± 1.02	
3	1326	3.31 ± .916	

II. Results

Group	Time After (hr)	# Dead	% Mortality	Days after Ectoparasitization												
				1	2	3	4	5	6	7	8	9	10	11	12	13
1	2	0	0	2.22	3.34	3.24	3.36	3.40	3.64	3.68	3.68	3.70	3.58	3.61	3.54	3.68
2	5	4	4	2.10	2.04	3.25	2.83	2.77	2.66	2.75	2.89	3.06	3.21	3.33	3.29	3.35
3	7	15	15	2.67	2.30	2.42	2.74	3.65	3.18	3.00	3.35	3.51	3.61	3.39	3.48	3.42
4	10	17	17	1.42	1.42	1.54	1.89	1.95	1.81	2.07	2.24	2.59	2.88	3.09	3.31	3.38
5	15	19	19	2.00	2.25	1.08	1.30	1.43	1.40	1.55	1.85	2.22	2.31	2.70	2.45	3.28
6	20	23	23	1.42	2.03	1.79	1.82	2.89	3.11	3.39	3.29	3.18	3.66	3.37	3.15	3.16
7	22	45	45	1.78	1.42	1.57	1.96	2.96	3.03	3.18	2.89	3.46	3.79	3.54	3.46	3.36
8	25	32	32	1.66	1.58	1.85	1.25	1.00	1.63	2.05	2.29	2.79	3.08	2.88	3.25	3.46



Figure 1. Side view of first meiotic metaphase in a cell from the testes of T. brevicornis showing bi-partite chromosomes.

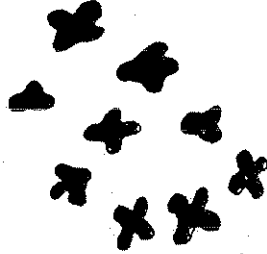


Figure 2. Side view of first meiotic metaphase in a cell from the testes of T. brevicornis showing quadripartite chromosomes.



Figure 3. Polar view of spermatogonial mitosis in the testes of T. anaphe.

Sokoloff, A., Sirotnik, B. and Beeman, C. California State University, San Bernardino, 92404.

*Length and weight in Tribolium brevicornis Le Conte.

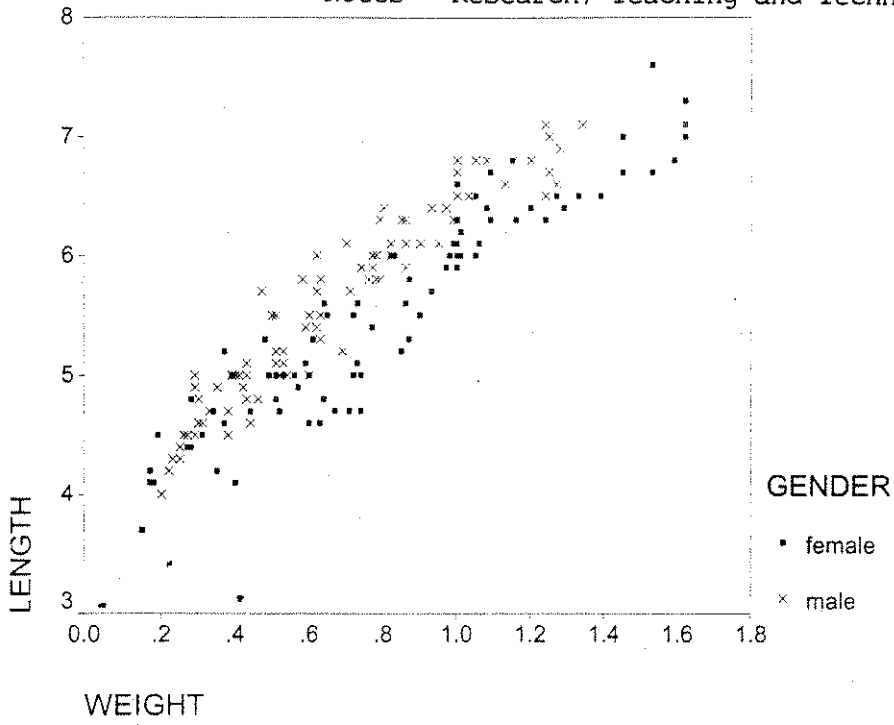
Tribolium brevicornis is one of 33 species known in Tribolium whose biological characteristics have been fairly well determined (See review by Sokoloff in this issue of the Tribolium Information Bulletin). In studying its biological attributes it has been found that males seemed larger than females, which is unusual, since in Insecta females are generally larger than males. (see Sokoloff 1955 and Sokoloff 1972 for information on this topic for Tribolium species and Drosophila species in the pseudoobscura subgroup. It will be seen that males are generally smaller than females).

Measurements of length of a small sample of beetles seemed to verify this visual impression. Since size can be determined by length of the whole animal and its weight, we decided to have data on both of these traits, and thus to be able to decide which data are less arduous to obtain. To this end a sample of 88 males and about 83 females was obtained from a culture, separated by sex, anesthetized, and each weight and length obtained from every individual in the sample. Using Pearson's Correlation the correlation for the two sexes separately and for all cases is significant is significant at the .01 level (2-tailed) as shown in Tables 1, 2, and 3 and depicted in Fig. 1.

By visual inspection (not statistical testing) males of a particular weight tend to be larger than females of the same weight. The difference is slight.

From a practical standpoint, it is probably much easier and more accurate to obtain individuals' weights with a good microbalance than to determine length of each beetle with an ocular micrometer.

Notes - Research, Teaching and Technical



Correlations

GENDER = male

Correlations^a

		LENGTH	WEIGHT
LENGTH	Pearson Correlation	1.000	.953**
	Sig. (2-tailed)	.	.000
	N	88	88
WEIGHT	Pearson Correlation	.953**	1.000
	Sig. (2-tailed)	.000	.
	N	88	88

** . Correlation is significant at the 0.01 level (2-tailed).

a. GENDER = male

GENDER = female

Correlations^a

		LENGTH	WEIGHT
LENGTH	Pearson Correlation	1.000	.940**
	Sig. (2-tailed)	.	.000
	N	83	83
WEIGHT	Pearson Correlation	.940**	1.000
	Sig. (2-tailed)	.000	.
	N	83	83

** . Correlation is significant at the 0.01 level (2-tailed).

a. GENDER = female

Correlations

Correlations -- ALL CASES

		LENGTH	WEIGHT
LENGTH	Pearson Correlation	1.000	.917**
	Sig. (2-tailed)	.	.000
	N	171	171
WEIGHT	Pearson Correlation	.917**	1.000
	Sig. (2-tailed)	.000	.
	N	171	171

** . Correlation is significant at the 0.01 level (2-tailed).

Notes - Research, Teaching and Technical

Grain Marketing and Production Research Center
 Biological Research Unit
 1515 College Avenue
 Manhattan, KS 66502
 Telephone: 785-776-2704/FAX 785-537-5584
James E. Throne, Research Leader

RESEARCH HIGHLIGHTS AND TECHNOLOGY TRANSFER FOR 1998-1999*We have collaborated with...*

INDUSTRIAL FUMIGANTS AND WHITMIRE, INC., to evaluate a new encapsulated formulation of cyfluthrin as a wheat protectant. (Franklin H. Arthur)

THERMO TRILOGY CORP., Columbia, MD, to evaluate neem products for storage pest control. (James E. Baker)

RALSTON PURINA, INTERNATIONAL PAPER CO., AND SEALED AIR CORP. to develop insect resistant packaging. (Michael A. Mullen)

In molecular research...

GENOME MAP FOR STORED-PRODUCT INSECT PEST. Constructed a complete genome molecular map of the red flour beetle using randomly amplified polymorphic (RAPD) DNA markers, visible mutant markers, and specific gene markers. The map is now being used to facilitate map-based cloning of important insect control genes. (Richard W. Beeman)

NEW MECHANISM OF INSECT RESISTANCE TO BIOPESTICIDE DETERMINED. A proteinase-mediated mechanism of insect resistance to the entomocidal toxins of *Bacillus thuringiensis* has been identified and characterized. Resistant insects lack a proteolytic enzyme that is critical for activation of the precursor forms of the toxins. This knowledge will be helpful in developing strategies for managing resistance to biopesticides in the field. (Brenda Oppert and Karl J. Kramer)

Development of ...

THE USE OF NEAR-INFRARED REFLECTANCE SPECTROSCOPY for detection of insect larvae in grain kernels, for disinfestation of insect-infested grain, and for identification of insects. (Floyd E. Dowell/James E. Baker/James E. Throne)

THE USE OF SPATIAL MODEL OF RUSTY GRAIN BEETLE DENSITY and bin temperature to simulate effects of time of aeration, bin size, and latitude on insect populations in stored wheat. Starting automatic aeration controllers at harvest suppressed insects below economic levels until the spring. (Paul W. Flinn and David W. Hagstrum)

BIOCHEMICAL TECHNIQUES TO EVALUATE PROTEINASES IN MIXTURES. Insect proteinases are studied to identify those that may be targeted by biopesticides. Techniques were developed to promote rapid and efficient identification of digestive proteinases in moth and beetle pests of stored products. Researchers from areas other than agriculture have used these assays to study proteinases prior to purification. (Brenda Oppert)

A NEW PHEROMONE TRAP FOR THE INDIANMEAL MOTH. This trap can be used in public areas, but is hidden from view. (Michael A. Mullen and Alan K. Dowdy)

Control of insect pests using...

HEAT AND DIATOMACEOUS EARTH. Laboratory tests indicate that the use of diatomaceous earth in combination with heat has the potential of reducing the temperature or time necessary to effect adequate insect management. (Alan K. Dowdy)

AREA-WIDE IPM FOR SUPPRESSION OF INSECT PESTS IN STORED WHEAT. A 5-year Area-Wide IPM project was recently funded by ARS to determine whether more uniform application of insect pest management across the marketing system could reduce insect problems in stored wheat. The program should

Notes - Research, Teaching and Technical

reduce the frequency of pesticide application, the cost of pest management, and the risk of insect problems.
(David W. Hagstrum/Paul W. Flinn/James E. Throne/Frank H. Arthur/Alan K. Dowdy/Michael A. Mullen)

OPEN FORUM

Note: This paper, published in 1996 in the *Tribolium Information Bulletin* 36:83-85 has been shortened and somewhat modified to fit a two-page requirement for the TIB On-Line

Sokoloff, A. Biology Department, California State University, San Bernardino, CA 92407.

*Interactions in Tribolium: Competition or predator-prey?

Population biologists have developed classification systems to define rigorously social interactions between lower organisms. Some interactions between associated populations are of benefit (+), other interactions are harmful (-), and others are neither beneficial nor harmful (0). In commensalism of two species one benefits while the other is not harmed (+/0). In competition both species are harmed in some way (-/-). In predator-prey or parasite-host interactions one species benefits and the other is harmed (+/-) (one species serves as food for the other). We are concerned here only in the last two interactions.

As the reviews of King and Dawson (1972) and Sokoloff (1975) have summarized, the late Thomas Park and his students and collaborators studied interactions between Tribolium castaneum (CS) and T. confusum (CF). He concluded that the interaction between these stored-product pests was competition: one of the species or the other was eliminated depending on the environmental conditions used. In the mid-sixties Park *et al* (1965) and Sokoloff and Lerner (1967) independently came to the conclusion that the interaction observed when these two species are placed in the same vial is a predator-prey interaction and not (as originally assumed by Park and his collaborators and others) a competition interaction. Sokoloff and Lerner thought that under certain conditions (such as rearing CS and CF in whole wheat flour enriched with brewer's yeast at 29° C and 70% R.H. the interaction is one of mutual predation because food is present in abundance and regularly renewed, and under these conditions CS is the winner. Under the same conditions, but utilizing other media such as corn, CS was eliminated by CF. Again the amount of food is probably in excess, since once CS is eliminated CF experiences a threefold increase in population size. But here the possibility that competition has occurred cannot be ruled out, because certain nutritional requirements are in limited supply in corn. Evidence that a shortage of these requirements causes CS to become a more active cannibal was obtained by Inouye and Lerner (1965).

At the time when these experiments were carried out neither Park *et al* (1965) nor Sokoloff and Lerner (1967) had any experimental basis to show that temperature may be a useful guide to resolve what kind of interaction (competition or predator-prey) is prevailing in the experiment. Bowker (1978) showed in her measurements of energetics of populations of single and mixed species of CF and CS that when beetles are reared at 25° or 30° C predator-prey interactions predominate, while at 35°C competition interactions predominate. Unfortunately, her paper did not attract the attention of Triboliumists: Her paper is not cited by any of the papers on competition or other interaction studies in the last 20 years.

In my opinion, recent students of interactions in Tribolium species, judging from the contents and their titles and the temperatures at which the experiments have been carried out, have misidentified the type of interaction they are observing, perhaps because of an inadequate search of the available literature. To a certain extent reliance on the literature published and available in data bases leads to errors in interpretation such as those I have described here. I have made available the facilities of the *Tribolium Information Bulletin* as an open forum to discuss the topic. So far only one Triboliumist has shown interest, but the open forum will remain open for a couple of years.

Literature Cited

- Bowker, L.S. 1979. The energetics in populations of Tribolium confusum and Tribolium castaneum. *Environm. Entomol.* 15:1264-1267.
- Inouye, N. and Lerner, I.M. 1965. Competition between Tribolium species (Coleoptera: Tenebrionidae) on several diets. *J. stored Prod. Res.* 1:186-191.
- King, C. E. and Dawson, P.S. 1972. Population biology and the Tribolium model. *Evol. Biol.* 5:133-227.
- Park, T., Mertz, D. B., Grodzinski, W. and Prus, T. 1965. Cannibalistic predation in populations of flour beetles. *Physiol. Zool.* 38:289-321.
- Sokoloff, A. 1975. The Biology of Tribolium with Special Emphasis on Genetic Aspects. Oxford Univ. Press, Vol. 2.
- Sokoloff, A. and Lerner, I.M. 1967. Laboratory ecology and mutual predation of Tribolium species. *AmerNat.* 101:261-276.

The following is Dr. Charles Goodnight's opinion about the topic "Interactions in Tribolium: Competition or predator-prey" prepared for the Open Forum.

The interaction between Tribolium confusum and T. castaneum clearly has elements of both competition and predation. It is hard not to consider the eating of one species by another to be anything other than predation. However, when distinguishing between these two processes it is perhaps more important to consider whether the dynamics of the system are better modeled as competition or as predation. As an evolutionary biologist I will not embarrass myself by expressing an opinion on the ecological dynamics. From an evolutionary perspective the evolution of the interaction appears to be best considered to be one of competition (either T. confusum or T. castaneum surviving) at the individual, group and community level. I am doing this selection both on communities (both species transferred between generations) and on systems where only one of the species can evolve, with the other drawn from a stock population each generation. Those data are still very preliminary. As before I am finding no evidence of evolution by individual selection, however there is evidence that group or community selection is changing the outcome of competition.

Several years ago we published a study of the effect of coexistence on the interaction between T. castaneum and T. confusum (Goodnight and Craig, 1996). In this study we set up 10 two species communities and 10 pairs of single species populations. These lines were maintained for 18 generations using discrete generation husbandry (see Goodnight and Craig 1996 for details). The advantage of discrete generation husbandry is that the two communities coexist with neither species going extinct. Nevertheless they interact sufficiently that there is ample opportunity for co-evolution to occur. At the end of 18 generations we set up continuous "Park style" competition (Park, 1948). The two species communities were allowed to compete, and the single species populations were combined into two species communities for the first time. We set 15 replicates for each lineage. Under this continuous form of husbandry it is inevitable that one of the species will go extinct, although the last community had both species surviving for 1000 days.

The outcome of this experiment was that there was no evidence that the two species communities co-evolved, that is there was no significant amount in the treatment component in the outcome of competition. However, there was a huge variation among lineages within treatments in the outcome of the competitive interaction. From this we conclude that there was no consistent evolution of the outcome of competition by individual selection, but the outcome would evolve if group or community selection were imposed. Note that our design could not distinguish between a lack of evolution by individual selection and a balanced "red queen" situation in which the competitive ability of the two species was evolving, but the overall outcome was not changing.

I am currently testing the prediction that competitive outcome can evolve by group and community selection, but not individual selection in an experiment in which I am selecting for the outcome of competition (either T. confusum or T. castaneum surviving) at the individual, group and community level. I am doing this selection both on communities (both species transferred between generations) and on systems where only one of the species can evolve, with the other drawn from a stock population each generation. These data are still preliminary. As before, I am finding no evidence of

I am currently testing the prediction that competitive outcome can evolve by group and community selection, but not individual selection in an experiment in which I am selecting for the outcome of competition

Although certainly not definitive, these experiments suggest to me that from an evolutionary perspective the two species T. castaneum/T. confusum communities are behaving as competitive systems rather than predator-prey systems. The reason for this is that the predator-prey systems are directional, with one species (the predator) benefiting at the expense of the other species (the prey). It seems likely that I could have so a directional change due to individual selection in a directional system, but perhaps not in a competitive system. It is particularly telling that there is no change in the outcome of competition due to individual even when only one of the species is allowed to evolve (Goodnight unpublished data). In a predator-prey system I would suspect that either becoming a more efficiency predator or more resistant predation would be a reasonable outcome. On the other hand, in a competitive system it is often argued that intraspecific competition should be as intense a selective force as interspecific competition. Thus, the failure of competitive outcome to evolve due to individual selection is perhaps less surprising.

Literature Cited

Goodnight, C. J. and D.M. Craig. 1996. The effect of coexistence on competitive outcome in Tribolium castaneum and T. confusum. *Evolution* 50:1241-1250.

Park, T. 1948. Experimental studies of interspecies competition. I. Competition between populations of flour beetles, Tribolium confusum Duval and Tribolium castaneum Herbst, *Ecological Monographs* 18:265-305.

GEOGRAPHICAL DIRECTORY

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NOTE: An asterisk denotes the individual who, as far as known, has worked or is working on Coleoptera. The plus sign (+) before the geographical locality indicates there was no current contribution. Since the information was obtained from previous issues of TIB, there is no guarantee that the information is accurate.

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pests, insecticide resistance. (7, 12)

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Bengston, M., Ph.d. Grain protectants, fumigants, stored
products pest management.
Collins, P. J., Ph.d. Insecticide and fumigant resistance,
fumigation, stored products pest management.
Daglish, G. J. Ph.D. Grain protectants, fumigants, stored
products pest management.

Melbourne, Victoria
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Braby, M.F. (8, 12, 18)
Williams, P., B.Sc., A.R.C.S., Ph.D., D.I.C., M.I. Biol.
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 genetics. (7)
 Wattiaux, J.M., Ph.D. Charge de Recherches au F.N.R.S. Age
 effects. (13)

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 13)

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Dr. David Abramson: Mycotoxicology, Analytical Chemistry.
 Mr. Colin J. Demianyk: Biologist; Insects Ecology, Industry
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 Dr. Paul G. Field: Entomology, Physiology, Insect Behavior.
 Dr. John T. Mills: Mycology, Ecology.
 Dr. Noel D.G. White Section Head: Entomology, Insecticides,
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Nunez, F. Genetics of Tribolium. Natural insecticides.

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- | | |
|-----------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Tautz, Diethard (Professor) | Evolution and development; Segmentation in <i>Drosophila</i> , <i>Tribolium</i> and zebrafisch. |
| Schroeder, Reinhard (Postdoc) | Head- and terminal gap genes; antibodies, antisense, CALI technique. |
| Wolff, Christian (Postdoc) | Function of <i>Tribolium</i> upstream elements in <i>Drosophila</i> . |
| Klingler, Martin (Res. Associate) | Embryonic lethal mutations in <i>Tribolium</i> affecting segmentation, appendage formation; transposon-mediated transformation; evolution of enhancer elements. |
| Berghammer, Andreas (Ph.D. stud.) | Transformation of <i>Tribolium</i> |
| Bucher, Gregor (Ph.D. student) | <i>Tribolium</i> gap-gene mutants |
| Maderspacher, Florian (Ph.D. st.) | <i>Tribolium</i> pair-rule mutants |

Martin Klingler

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GREAT BRITAIN

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List of members working on *Tribolium* and other coleopterous species.

1. Name: **Abdul Rahman Dhuyo**

Degree held: M.Sc. (Agri. Entomology)

Field of work: Biological control of *Prostephnus* and *Tribolium* species.

2. Name: **Abdur Rauf**

Degree held: M.Sc. (Agri. Entomology)

Field of work: Enzymatic mechanism of resistance to insecticides in *Tribolium* and *Sitophilus* species.

3. Name: **Fazalullah.M Bughio**

Degree held: M.Sc. (Agri. Entomology)

Field of work: Rice grain resistance to *Tribolium* and *Sitophilus* species.

4. Name: **Ovais Safader Pathan**

Degree held: M.Sc. (Agri. Entomology)

Field of work: Wheat grain resistance to *Tribolium* species.

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Compton, J.A.F. B.Sc., MSc. Farm and village level storage management

Dales, M.J. B. Sc. PhD. Control of stored-product insects (Chemical dusts, botanicals, IG's), aFumigation with carbon dioxide.

Donaldson, I. M.Phil. Farming systems, training, project management

Farnell, G. B. Sc. M.Phil. Farming systems, training, project management

Freeman, N. Biology and behavior of stored product pests.

Giles, F.H., B.Sc., D.I.C., Dip.Agric.Sci., M.I.Biol. Stored products technology. Biology and control of insect pests. (12)

Golob, F., B.Tech., Ph.D. Control of stored product pests, especially with insecticides. (8, 12, 18)

Gudrups, I. B.Sc. PhD F.R.E.S. Biology and Behavior of stored products pests.

Haines, C.F., B.Sc., Ph.D. Stored products entomology and acarology. Control by pheromones and natural predators. (2, 6, 12, 17)

Hodges, R.J., B.Sc., Ph.D. Stored products entomology. Control and inspection procedures using pheromones. (12)

Orchard, J. Ph. D. Biochemist, resistance of grains to insect attack.

Freyett, P.F., B.Sc., Ph.D., D.I.C., A.R.C.S., M.I. Biol. Deputy Head of Centre. Biology and control of storage insects. (6, 12, 17)

Taylor, R.W.D. B. Sc. Control of stored-product pests, especially with fumigants.

Wright, M. B.Sc., M.Sc., PhD. Storage technologist. Project management.

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Binns, T.J.	Insecticide efficacy and resistance
Cogan, P.M.	Insect monitoring
Cox, Dr. F.D.	Stored product entomology
Hills, Dr. K.A.	Fumigation, controlled atmospheres, gene-
Muggleton, Dr. J.	Insecticide resistance, population genetics
Prickett, Dr. A.J.	Insecticides, resistance statistics (6, 7,
Walter, Miss C.	Insecticide biochemistry

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- Dr. Carlos Juan - Molecular phylogeny and phylogeography of Tenebrionidae and Chrysomelidae. Satellite DNA and heterochromatin of Tenebrionidae.
- Dr. Claudia Garin (Argentinian postdoc) - Molecular phylogeny of Chrysolina and Oreina (Chrysomelidae).
- Mr. Juan Pons - Satellite DNA, heterochromatin and evolution of Pimelia (Tenebrionidae), particularly those of the Canary Islands.
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Graduate Students: Teresa Short (molecular analysis of HOMC, *mxp*)
Marcé Lorenzen (transformation of Tribolium)
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