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## Changes in the Sexual Dimorphism of Triatominae in the Transition from Natural to Artificial Habitats

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*A shift from large to small average sizes was observed in Triatoma infestans and Rhodnius domesticus in field and domestic (or laboratory) conditions of life. It was more pronounced in the female specimens, leading to a subsequent reduction of sexual size dimorphism. This feature is discussed in terms of genetic and populational changes occurring from natural to artificial habitats, in particular those related to population densities. Sexual size dimorphism is then recommended as a new character to be used in the study of species of Triatominae adapting to domestic ecotopes.*

Key words: Triatominae - sexual size dimorphism - silvatic - domestic - laboratory

Triatominae (Hemiptera, Reduviidae) are blood sucking insects widely distributed in Latin America, where several are important vectors of *Trypanosoma cruzi*, causative agent of Chagas disease (American trypanosomiasis). Most species occur in silvatic ecotopes, associated with small nest-building mammals or birds, but several have made the transition to domestic habitats where they feed on humans and domestic animals, transmitting *T. cruzi*.

Many species can be reared in the laboratory by feeding on laboratory animals such as mice, rabbits or chickens, and various authors have reported a general reduction in average size of specimens reared over several generations (Szumlewicz 1976, Zeledón 1981).

We have also noted a reduction in the sexual dimorphism of head measurements associated with captive breeding of Triatominae, and report here a study of this phenomenon in *Rhodnius domesticus* (Neiva & Pinto).

In spite of its name, *R. domesticus* seems to be entirely silvatic, confined to Atlantic forest ecotopes along the Brazilian coast. Although frequently infected with *T. cruzi*, and also *T. rangeli*, it has little epidemiological significance (Leal et

al. 1961, Galvão et al. 1962, Schlemper Jr et al. 1985, Steindel et al. 1991). Colonies of *R. domesticus* originating from the Island of Santa Catarina have been reared at the Federal University of Santa Catarina since 1984, and the species was recently collected again from its natural bromeliad habitats on the Island. Our morphometric comparisons of the laboratory and silvatic samples were prompted by the idea that any metric changes might parallel similar differences that we have seen in silvatic and domestic populations of *Triatoma infestans* (Klug) in Bolivia (Dujardin & Casini 1996, Dujardin et al. 1997a,b).

We add to the *R. domesticus* study a similar study in *T. infestans*, comparing sexual size dimorphism between its silvatic and domestic ecotopes of Bolivia, and we report related observations from published data on *R. prolixus* (Harry 1992).

### MATERIALS AND METHODS

We examined 22 adult *R. domesticus*, 14 direct from the field collections and 8 from the laboratory colony of the same geographic origin. For comparison we also reexamined published data on the related *R. prolixus* Stal (Harry 1992, 1994), and *T. infestans* (Dujardin et al. 1997b). For each of the *R. domesticus* specimens we made 12 head measurements as indicated in Fig. 1. Means and standard deviations for each character (Table I) were compared by sex and biotope using the non-parametric test of Kruskal and Wallis (1952) and each comparison was tested for significance using the sequential Bonferroni test as described by Sokal and Rohlf (1995).

We made a similar comparison for silvatic and domestic *T. infestans* from Bolivia using the five

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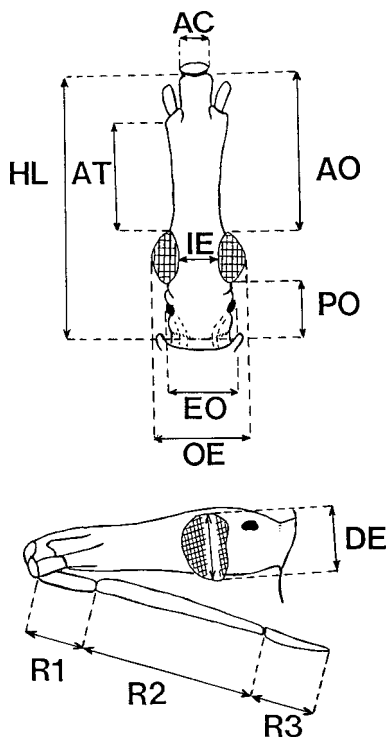


Fig. 1- OE: outer distance between eyes; IE: inner distance between eyes; EO: external distance between ocelli; AO: antocular distance; PO: postocular distance (excluding neck); HL: head length; DE: diameter of the eye; AT: antenniferous tubercle; R1, R2 and R3: lengths of first, second and third rostral segments, respectively, and AC: width of the anteclypeus.

head measurements retained by Dujardin et al. (1997b) as giving the best accuracy and precision of measurement: OE, PO, HL, AT and R2 (Fig. 1).

We could also carry out multivariate analysis using a covariance-matrix based principal component analysis on log-transformed data from *R. domesticus* and *T. infestans*. To avoid matrix singularities, two measures of *R. domesticus* (IE and HL, Fig. 1) were discarded because of their partial overlap with other measures. The remaining variables were all significantly correlated with the first principal component (PC1) which contributed 64% to the overall variation. PC1 could therefore be accepted as a general indicator of size (Bookstein 1989, Dos Reis et al. 1990) so that the resulting factor maps (Fig. 2) clearly illustrate global size differences in the populations analyzed.

**RESULTS**

Females from the field collected sample of *R. domesticus* were significantly different from males at almost all measurements, whereas no such difference between males and females was found in the laboratory specimens (Table II, left). This was

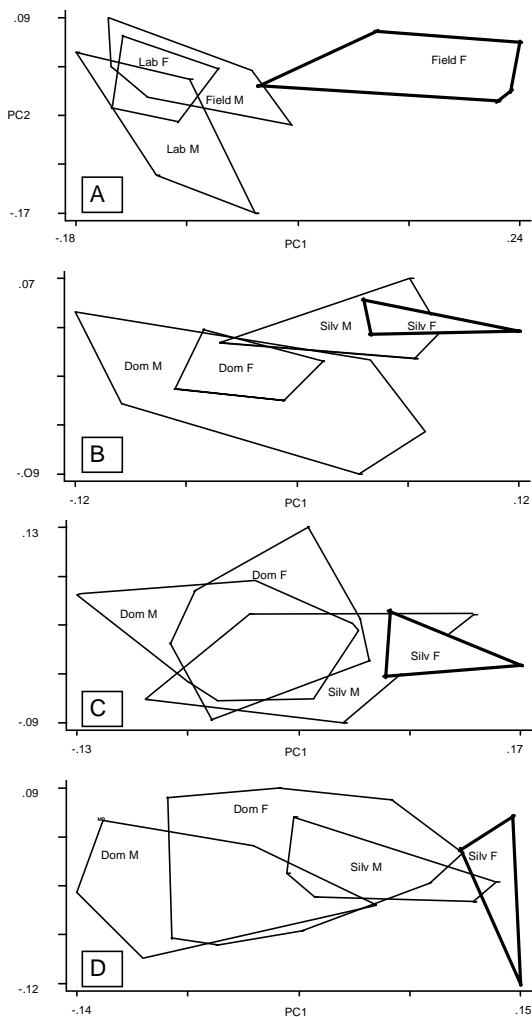


Fig. 2 : factor maps corresponding to *Rhodnius domesticus* (A, top) and three samples of *Triatoma infestans* (B, C, D). Female (F) and male (M) specimens of *R. domesticus* (A, top), either reared in laboratory (Lab F, Lab M) or recently collected in the field (Field F, Field M), are projected onto the first (PC1, 64%) and second (PC2, 14%) principal components. The three remaining factor maps (B, C, D) are female (F) and male (M) specimens of *T. infestans*, collected in house (Dom F, Dom M) or in silvatic ecotopes (Silv F, Silv M), projected onto PC1 and PC2; B, sample collected at Laguna Angostura in 1983; C, sample collected at Jamach'Uma in 1993 and D, sample collected at Jamach'Uma in 1996 (see Dujardin et al. 1997b). Convex hulls enclose specimens for each category. These four factor maps illustrate three main features in both *R. domesticus* and *T. infestans*: (i) no or poor sexual differentiation in laboratory or domestic habitat (compare Lab M and Lab F in *R. domesticus*, Dom M and Dom F in *T. infestans*); (ii) significant sexual dimorphism in field, silvatic habitat (compare Field M and Field F in *R. domesticus*, Silv M and Silv F in *T. infestans*), and (iii) an important reduction of female head dimensions from field to either laboratory (compare Field F and Lab F in *R. domesticus*) or domestic habitats (compare Silv F and Dom F in *T. infestans*). Notice that in the 1996 sample (D, bottom), there was also a size difference between silvatic and domestic male *T. infestans*.

also shown in the comparison by biotope, whereby males showed no significant metric differences but females showed a drastic size reduction going from field to laboratory populations (Table II, right). We reexamined the data presented by Harry (1992) on samples of *Rhodnius* collected from various localities in Venezuela. These data showed a lower level of sexual dimorphism in the six domestic samples of *R. prolixus*, and a pronounced dimorphism in the silvatic strain collected directly from palm tree

habitats (determined as *R. robustus* by Harry 1994) (detailed results not shown).

Using the same Bonferroni tests, significant sexual dimorphism was detected only in the silvatic population of *T. infestans*, not in the domestic specimens, and the reduction in size of the head measurements between silvatic and domestic specimens was again more pronounced for females (Table III).

These changes in sexual size dimorphism were verified also after multivariate analyses: in all cases,

TABLE I  
Metric properties of *Rhodnius domesticus*

	Females				Males			
	Field		Laboratory		Field		Laboratory	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
OE	32	0.9	29	1.0	31	1.2	29	0.5
IE	14	0.4	13	1.0	13	0.4	13	0.5
EO	22	0.8	21	0.5	21	0.0	21	0.0
AO	41	1.0	38	1.7	37	2.0	39	0.9
PO	33	1.6	29	0.8	30	0.6	30	0.6
HL	74	2.4	68	2.5	68	1.8	69	2.2
DE	21	1.2	19	0.0	19	1.0	20	1.7
AT	28	1.1	27	0.8	26	0.9	27	1.3
R1	18	0.5	16	0.5	17	0.5	17	1.3
R2	56	1.5	52	1.3	52	1.8	52	0.6
R3	17	0.5	16	1.0	16	0.5	15	1.0
AC	20	1.0	18	0.5	19	0.8	18	0.5

Mean and standard deviation (Stdev) in graduation units for 12 head measurements (OE to AC, see Fig. 1) for field and laboratory samples of female and male *R. domesticus*. All measurements were performed by the same investigator (TC) at 25x (OE, IE, EO, AO, PO, HL, DE, AT, R1, R2, R3) and 40x (PO, AC) magnification using a monocular micrometer. One micrometer graduation represented 20 μ for the measurement PO and AC (50x magnification), and 40 μ for the other head measurements (25x magnification).

TABLE II  
Univariate analysis in *Rhodnius domesticus*

	Between males and females				Between field and laboratory			
	Laboratory		Field		Females		Males	
	P	Bonf	P	Bonf	P	Bonf	P	Bonf
OE	0.3496	NS	0.0316	NS	0.0054	S	0.0189	NS
IE	0.1692	NS	0.0019	S	0.0189	S	0.1155	NS
EO	0.3173	NS	0.0093	S	0.0201	S	1.0000	NS
AO	0.8817	NS	0.0039	S	0.0149	S	0.2352	NS
PO	0.3428	NS	0.0054	S	0.0079	S	0.2225	NS
HL	0.3807	NS	0.0032	S	0.0120	S	0.3850	NS
DE	0.2155	NS	0.0179	NS	0.0280	S	0.2188	NS
AT	0.8770	NS	0.0038	S	0.0536	S	0.1175	NS
R1	0.7568	NS	0.0014	S	0.0044	S	1.0000	NS
R2	0.3428	NS	0.0049	S	0.0077	S	0.8236	NS
R3	0.1776	NS	0.0026	S	0.0363	S	0.1461	NS
AC	0.6171	NS	0.0612	NS	0.0280	S	0.6446	NS

Non-parametric comparisons (Kruskal & Wallis 1952) at 12 head measurements (OE to AC, see Fig. 1) of *R. domesticus*, exploring sexual dimorphism in the laboratory and in the field collected samples (see columns "Laboratory" and "Field"), and biotope differences for each sex (see columns "Females" and "Males"). P, level of significance; Bonf, sequential Bonferroni test (after Sokal & Rohlf 1995, p. 241).

TABLE III  
Univariate analysis in silvatic and domestic *Triatoma infestans*  
(data from Dujardin et al. 1997b)

	Between males and females				Between silvatic and domestic			
	Domestic (40, 33)		Silvatic (20, 10)		Females (33, 10)		Males (40, 20)	
	P	Bonf	P	Bonf	P	Bonf	P	Bonf
OE	0.3299	NS	0.0907	NS	0.1951	NS	0.6801	NS
PO	0.0380	NS	0.2231	NS	0.0001	S	0.0123	S
HL	0.0258	NS	0.0056	S	0.0011	S	0.0201	NS
AT	0.0281	NS	0.0543	NS	0.0039	S	0.0008	S
R2	0.1193	NS	0.5669	NS	0.0105	S	0.1084	NS

Non-parametric comparisons (Kruskal & Wallis 1952) at 5 head measurements (see Dujardin et al. 1997b) of *T. infestans*, exploring sexual dimorphism in domestic and silvatic samples (see columns "Domestic" and "Silvatic"), and ecotope differences for each sex (see columns "Females" and "Males"). P, level of significance; Bonf, sequential Bonferroni test (after Sokal & Rohlf 1995, p. 241).

sexual dimorphism was reduced in domestic or laboratory populations compared to silvatic populations, and the changes seemed mainly due to a reduction in size of the females (Fig. 2).

#### DISCUSSION

The existence of sexual dimorphism in Triatominae is well known, with females, on average, being larger than males (Lent & Wygodzinsky 1979). No previous study has examined the fate of this dimorphism in different habitats, although a general reduction in body size has been noted for bugs reared over successive generations in the laboratory (Szumlewicz 1976, Zeledón 1981). This has been suggested to be due to a longer generation time in natural habitats, as a result of several incomplete bloodmeals delaying the successive moults (Zeledón et al. 1970, Zeledón 1981), but more simply could be due to improved survivorship of smaller individuals in the laboratory (Schofield 1996). Other observations also show that domestic specimens, on average, tend to be smaller than their silvatic conspecifics (Harry 1994, Dujardin et al. 1997a,b), which could again be due to improved survivorship in domestic habitats. The present study provides a quantitative analysis of these changes, and reveals that in the material examined the reduction in average size could be mainly due to size reductions in female bugs, leading to a corresponding reduction in sexual dimorphism.

We believe that, more than a direct relation with silvatic and domestic habitats, our observation could reveal the effect of population density on bugs size. Indeed, in a population near its carrying capacity - which is generally the case in domestic or laboratory populations - each individual would

get less blood because of competition and would then be smaller, especially among the females with greater food requirements. Due to higher survivorship in domestic or laboratory colonies, smaller individuals would survive and the average size decrease, especially in females.

According to that hypothesis, and as long as the population density generally increases from silvatic to domestic habitats, we can expect modifications in sexual dimorphism in the transition of bugs from silvatic to domestic habitats, and suggest that this feature could be used in the study of this transition in other species.

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