

Evolutionary genetics of the insular lacertid lizard *Podarcis tiliguerta*: genetic structure and population heterogeneity in a geographically fragmented species

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Podarcis tiliguerta is an insular Mediterranean lacertid lizard endemic to Corsica, Sardinia and many neighbouring small islands. The genetic structure and population heterogeneity of the species were studied by means of allozyme electrophoresis at 20 presumptive gene loci. The observed heterozygosity (H_o) decreases southwards towards more arid climatic regimes. No severe reduction in genetic variability was found in samples from the tiny satellite islands, except for a population inhabiting a very small island off the south-western coast of Sardinia (Meli Island), in which about 80 per cent of the genetic variability was lost. Population heterogeneity analysis carried out by the estimation of Wright's F -statistics demonstrated substantial genetic differentiation among populations. The value of F_{ST} (0.460) exceeds values known for other lizard species. F -statistics and genetic distance data show that genetic variation is distributed into three geographically coherent population groups. The first group includes populations from the northern part of the range (Corsica), the second includes populations from the small islands off the south-eastern coast of Corsica (Cerbicale and Lavezzi), and the third comprises populations from the southern part of the range (Sardinia and Meli Island). The pattern of genetic variability and the apparent clinal variation of alleles at a few loci (*Idh-1*, *Gapd*, *Gpi*) indicate that the subdivided genetic structure of *P. tiliguerta* is moulded by the interplay of stochastic processes and agents selectively affecting allele frequency changes.

Keywords: allozyme electrophoresis, population heterogeneity, genetic differentiation, gene flow, *Podarcis tiliguerta*, Lacertidae.

Introduction

Island species generally do not behave as single, infinitely large panmictic populations. This is because the species range is divided into ecologically diverse subranges, each with its own selective factors. In these species the pattern of genetic differentiation among subdivisions is influenced by the size of the subdivisions, by differences in the modes of selection among subdivisions, and by the degree of migration among them. Because these forces contribute to the deviation from Hardy–Weinberg proportions, at any moment population genetic structure can be

described by measuring this deviation within populations and the amount of differentiation among populations (Wright, 1965, 1978). Although in recent years allozyme polymorphisms have been instrumental in studying the genetic structure of island populations on micro- and macrogeographical scales and in estimating the levels of gene flow between insular populations of various groups of organisms (see for example Koehn & Eanes, 1978; Ayala, 1982; Berry, 1983; Selander & Whittman, 1983; Larson *et al.*, 1984), to date quantitative surveys focusing on the amounts of genetic variation in insular Mediterranean lacertid lizards (Lacertidae) are scarce (Gorman *et al.*, 1975; Capula, 1990, 1994a), despite the fact that these reptiles are widespread and rather common.

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In this paper we use allozyme electrophoresis to estimate the distribution of genetic variation within and among populations, the genetic diversity and the level of gene flow in the insular Mediterranean lacertid lizard *Podarcis tiliguerta*. This species has been previously studied to evaluate its genetic divergence from some close relatives (Lanza *et al.*, 1977; Mayer, 1981; Guillaume & Lanza, 1982), but population genetic data are still lacking. *Podarcis tiliguerta* is a member of a lacertid genus that includes several species endemic to Mediterranean islands. It is a polytypic species found on Corsica, Sardinia and many neighbouring small islands (Arnold & Burton, 1978). *Podarcis tiliguerta* is at an intermediate stage in the taxon cycle (Wilson, 1961) in that it has a complete pattern of distribution (no uninhabited islands) and shows strong interisland geographical variation in morphology and coloration (Lanza & Poggesi, 1986).

Materials and methods

Sampling

Samples of *P. tiliguerta* used in this study were obtained from 14 localities; three from Corsica, four from the Cerbicale Islands (SE Corsica), one from Lavezzi Island (S Corsica), five from Sardinia, and one from Meli Island (SW Sardinia). For interspecific comparison five samples of the closely related species *P. muralis* and *P. sicula* were also employed. The geographical origin and the number of specimens analysed per population are indicated in Fig. 1a,b.

To avoid killing animals or injurious biopsy, approximately 1 cm of the tail of each lizard was taken off following the suggestion by Mayer & Tiedemann (1985). The piece of tail was then kept in Eppendorf reaction tubes (2 mL) and stored below -70°C until electrophoretic analysis.

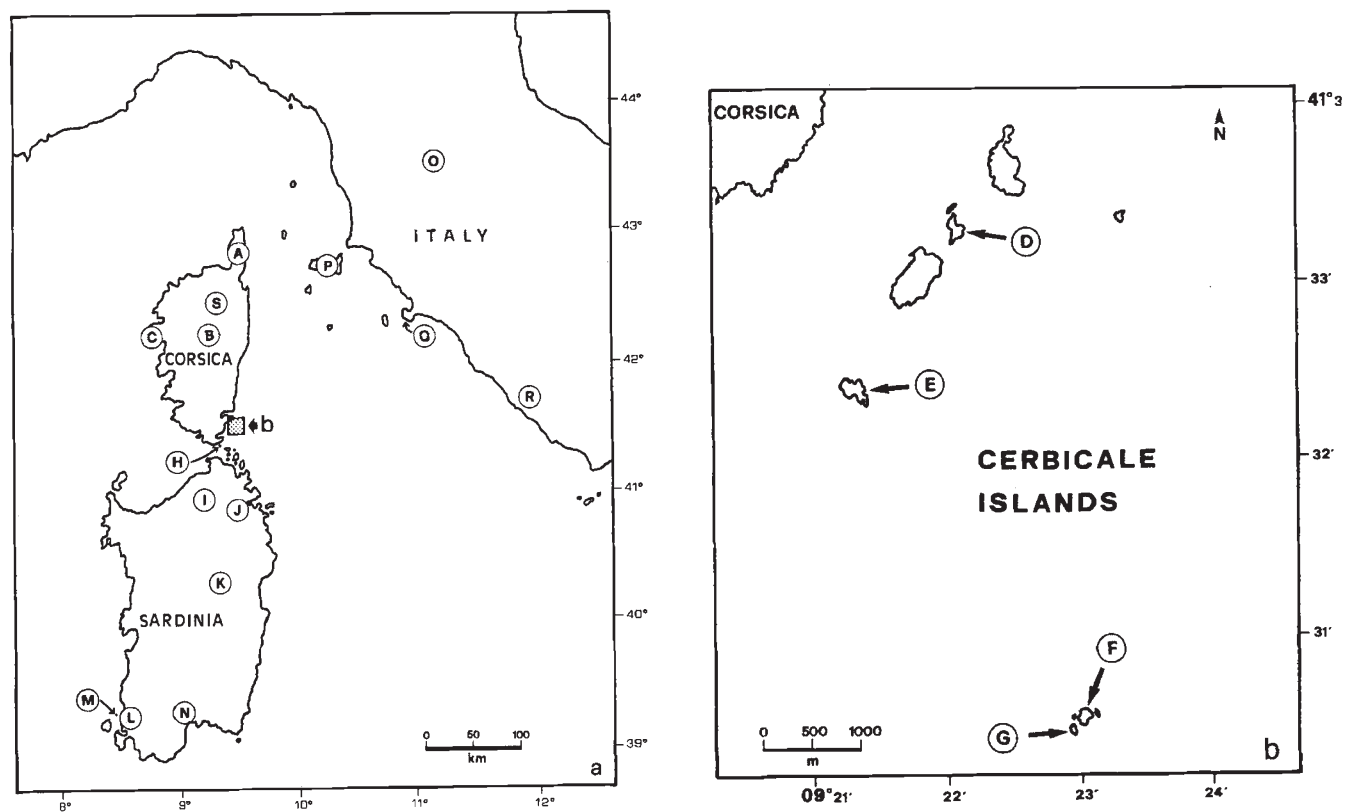


Fig. 1 Maps of Sardinia, Corsica, Italy (a) and Cerbicale Islands (b) showing sampling localities (sample size (N) is given in parentheses). *Podarcis tiliguerta*: A, Sisco ($N = 8$); B, Vizzavona ($N = 14$); C, Cargèse ($N = 12$); D, Maestro Maria ($N = 8$); E, Pietricaggiosa ($N = 15$); F, Toro Grande ($N = 12$); G, Toro Piccolo ($N = 7$); H, Lavezzi ($N = 8$); I, Tempio Pausania ($N = 19$); J, Olbia ($N = 14$); K, Nuoro ($N = 7$); L, Capo Giordano ($N = 5$); M, Meli ($N = 7$); N, Cagliari ($N = 4$); *Podarcis muralis*: O, Firenze ($N = 15$); P, Elba ($N = 9$); Q, Monte Argentario ($N = 17$); *Podarcis sicula*: R, Roma ($N = 9$); S, Caporalino ($N = 10$).

Electrophoresis

Standard horizontal starch gel electrophoresis was performed on tail muscle tissue; parts of this tissue were crushed in 0.1 mL of distilled water. Gene products for the following 20 presumptive enzyme loci were analysed: α Gpd, Ldh-1, Ldh-2, Mdh-1, Mdh-2, Me-1, Me-2, Idh-1, Idh-2, 6Pgd, Gapd, Sod-1, Got-1, Got-2, Ck, Ak, Mpi, Gpi, Pgm-1, Pgm-2. The buffer systems used and electrophoretic procedures are given in Table 1. The staining techniques used were those described by Capula (1990).

Isozymes were numbered in order of decreasing mobility from the most anodal; allozymes were named numerically according to their mobility relative to the commonest one found in a reference population of *P. sicula* from Rome (Italy), indicated as 100 (>100 = faster mobility; <100 = slower mobility).

Analysis

Genotypic and allelic frequencies were determined by direct counts from allozyme phenotypes, and the resulting data were analysed by various statistical methods to describe the genetic structure of the *P. tiliguerta* populations. Genotypic proportions expected on the basis of Hardy–Weinberg equilibrium were calculated by Levene's (1949) formula for small samples. The statistical significance of departures from Hardy–Weinberg equilibrium was estimated using a test for calculating exact significance probabilities, analogous to Fisher's exact test (Haldane, 1954; Elston & Forthofer, 1977). To determine whether the heterogeneity in the genotypic distribution reflects differences in allele frequencies, the variation in genic proportions among populations was subjected to a contingency χ^2 analysis (Workman & Niswander, 1970). The genetic variability of populations was estimated using the following parameters: mean number of alleles per locus (A); proportion of polymorphic loci, at the 99 per cent level (P); observed mean heterozygosity per locus (H_o); expected mean heterozygosity per locus (H_e) (unbiased estimate; Nei, 1978).

The distribution of genetic variation within and among populations was assessed using Wright's F -statistics (Wright, 1965, 1978). Statistical significance of Wright's standardized variance in allele frequencies (F_{ST}) was tested by the χ^2 test:

$\chi^2 = 2N_t F_{ST}(k-1)$, with $(k-1)(s-1)$ degrees of freedom, where N_t is the total sample size, k is the number of alleles, and s is the number of subdivi-

sions (Workman & Niswander, 1970). A hierarchical analysis of population differentiation was also performed using the formulation of Wright (1978).

The genetic relationships among the populations studied were evaluated using Nei's (1972) standard genetic identity (I) and standard genetic distance (D). All genetic variability, F -statistics, and genetic distance measures were calculated by the computer program BIOSYS-1 (Swofford & Selander, 1981).

To test the correlation between genetic and geographical distances, a Mantel test was used as advocated by Smouse *et al.* (1986). Tests were performed using the software NTSYS 1.50 (Rohlf, 1989).

Genotype data at 20 electrophoretic loci were analysed for rates of gene flow (Nm) in *P. tiliguerta* populations. Quantitative estimates of Nm were derived using the indirect method of Wright (1978). In this method, assuming neutrality of the alleles studied and an equilibrium between genetic drift and migration in an infinite island model, Nm is related to F_{ST} by the formula $F_{ST} = 1/(1+4Nm)$, where N is the local population size and m is the average rate of immigration.

An estimation of phenetic relationships among populations was obtained by generating a phenogram of all samples by means of the unweighted pair-group method with arithmetic averaging (UPGMA) based on the matrix of Nei's genetic distances (Sneath & Sokal, 1973).

Results

Pattern of variation

Five of the 20 presumptive gene loci analysed (25 per cent) were found to be monomorphic and fixed for the same allele in all the samples studied (α Gpd, Mdh-2, Sod-1, Ck, Ak). The allele frequencies at the other 15 variable loci are given in the Appendix. Six loci (30 per cent) were locally (≥ 5 populations) and strongly polymorphic (Ldh-1, Me-1, 6Pgd, Gapd, Got-1, Gpi). All other loci (45 per cent) were weakly polymorphic. As far as *P. tiliguerta* is concerned, the number of polymorphic loci was nine (26 alleles) in the populations from Corsica, six (21 alleles) in the populations from the Cerbicale and Lavezzi islands, and nine (29 alleles) in the samples from Sardinia and Meli Island. Four unique alleles (Slatkin, 1987) were detected in the Corsican populations, two in the populations from the Corsican satellite islands, and nine in the Sardinian populations (Fig. 2).

The results of the contingency χ^2 analysis are given in Table 2. The analysis reveals that ten out of

Table 1 Enzymatic proteins examined and electrophoretic conditions employed

Enzyme	Migration + = anodal - = cathodal	Buffer system*	V/cm	Time (h)	References
Glycerol-3-phosphate dehydrogenase α GPD (EC 1.1.1.8)	+	3	8	6	Ayala <i>et al.</i> (1972)
Lactate dehydrogenase LDH (EC 1.1.1.27)	+	4	7	6	Brewer & Sing (1970)
Malate dehydrogenase MDH (EC 1.1.1.37)	+	3	8	5	Shaw & Prasad (1970)
Malic enzyme ME (EC 1.1.1.40)	+	1	8	6	Ayala <i>et al.</i> (1972)
Isocitrate dehydrogenase IDH (EC 1.1.1.42)	+	3	8	5	Shaw & Prasad (1970)
6-Phosphogluconate dehydrogenase 6PGD (EC 1.1.1.44)	+	3	8	6	Shaw & Prasad (1970)
Glyceraldehyde-3-phosphate dehydrogenase GAPD (EC 1.2.1.12)	+	2	7	6	Ayala <i>et al.</i> (1972)
Superoxide dismutase SOD (EC 1.15.1.1)	+	1	8	5	Selander <i>et al.</i> (1971)
Glutamate-oxaloacetate transaminase GOT (EC 2.6.1.1)	+	3	9	6	Selander <i>et al.</i> (1971)
Creatine kinase CK (EC 2.7.3.2)	-	1	8	5	Ayala <i>et al.</i> (1972)
Adenylate kinase AK (EC 2.7.4.3)	+	1	8	5	Ayala <i>et al.</i> (1972)
Mannose-6-phosphate isomerase MPI (EC 5.3.1.8)	+	2	8	4	Harris & Hopkinson (1976)
Glucose-6-phosphate isomerase GPI (EC 5.3.1.9)	-	3	8	6	Selander <i>et al.</i> (1971)
Phosphoglucomutase PGM (EC 5.4.2.2)	+	4	8	6	Brewer & Sing (1970)

Enzymes are arranged by Enzyme Commission Number (EC).

*Buffer systems used: (1) Continuous Tris/citrate (Selander *et al.*, 1971); (2) Tris/versene borate (Brewer & Sing, 1970); (3) Phosphate/citrate (Harris, 1966); (4) Tris/maleate (modified from Brewer & Sing, 1970).

thirteen polymorphic loci exhibit statistically significant heterogeneity in the allele frequencies (see Table 2). This result indicates that there are highly significant differences among the gene pools of the *P. tiliguerta* populations, suggesting considerable local genetic differentiation. At three loci (*Idh-1*, *Gapd*, *Gpi*) displaying large differences in allele frequencies among populations, six alleles show significant departures from a random distribution (Sokal & Oden's (1978) spatial autocorrelation analysis), indicating apparent clinal variation in allele frequencies. In the case of the *Idh-1* locus, the *Idh-1*¹⁰⁸ allele is fixed in the populations from the Corsican satellite islands and Sardinia, decreasing in frequency towards the northern part of the range; *Idh-1*¹⁰² is the predominant allele in the populations

from Corsica. At the *Gapd* locus, where two electrophoretic alleles occur at high frequencies, i.e. *Gapd*⁹⁴ and *Gapd*¹⁰⁰, the *Gapd*¹⁰⁰ allele predominates in the Corsican populations, whereas the *Gapd*⁹⁴ allele is present at higher frequencies in most of the other samples. At the *Gpi* locus, two alleles occur at high frequencies: *Gpi*⁹⁵ and *Gpi*¹⁰³. The *Gpi*⁹⁵ allele is nearly fixed in the populations from Sardinia, including Meli Island, and its frequency decreases towards the north, whereas *Gpi*¹⁰³ is the predominant allele in the samples from Corsica and Corsican satellite islands (see Appendix). The alleles at the remaining loci exhibit random geographical patterns.

As to the diagnostic loci (at the 0.99 per cent level) between *P. tiliguerta* and the two other *Podarcis* species, four (20 per cent) displayed fixation of

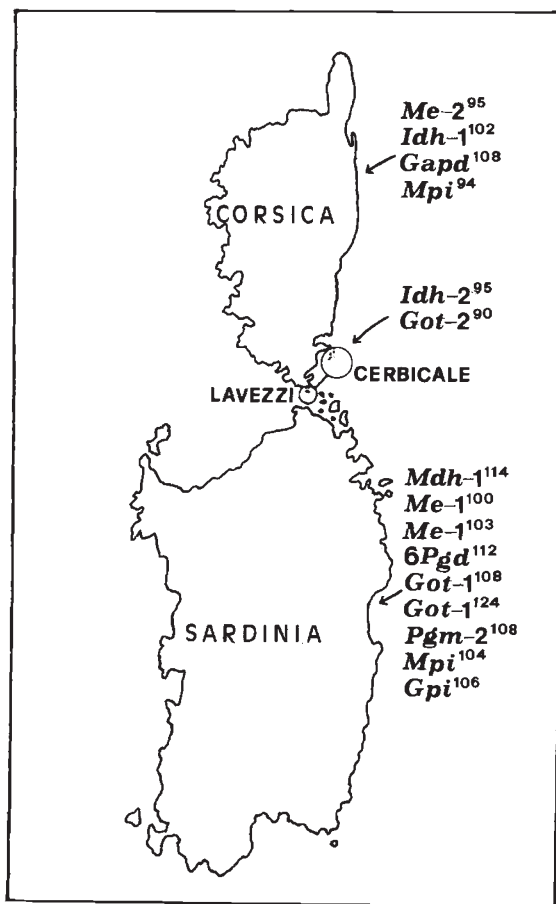


Fig. 2 Alleles specific to the *Podarcis tiliguerta* populations from Corsica, Corsican satellite islands (Cerbicale and Lavezzi), and Sardinia.

alternative alleles between *P. tiliguerta* and *P. sicula* (*Idh-1*, *Got-1*, *Mpi*, *Gpi*). *Podarcis tiliguerta* and *P. muralis* showed alternative alleles at two loci (*Mpi*, *Gpi*) and highly differentiated allele frequencies at two further loci (*Idh-1*, *Got-1*). It must be noticed that two (*Idh-1*, *Gpi*) out of the four loci displaying fixation of alternative alleles among the three species also exhibit clinal variation in allele frequencies in *P. tiliguerta* populations.

Significant deviations from Hardy-Weinberg equilibrium in the direction of heterozygote deficiencies were found for the following populations and loci (in parentheses): Vizzavona (*6Pgd*, $P < 0.05$); Pietri-caggiosa (*Got-2*, $P < 0.003$); Olbia (*Me-1*, $P = 0.001$; *Gapd*, $P < 0.01$).

Genetic variability

The measures of genetic variability used here are given in Table 3. The overall mean number of alleles

Table 2 Chi-square values resulting from contingency χ^2 analysis at the polymorphic loci among populations of *Podarcis tiliguerta*

Locus	No. of alleles	χ^2	d.f.	P
<i>Ldh-1</i>	2	82.742	13	**
<i>Mdh-1</i>	2	5.812	13	NS
<i>Me-1</i>	4	135.780	39	**
<i>Me-2</i>	2	59.222	13	**
<i>Idh-1</i>	2	154.213	13	**
<i>Idh-2</i>	2	46.746	13	**
<i>6Pgd</i>	3	87.274	26	**
<i>Gapd</i>	3	126.192	26	**
<i>Got-1</i>	4	61.416	39	*
<i>Got-2</i>	2	21.321	13	NS
<i>Mpi</i>	3	64.021	26	**
<i>Gpi</i>	3	171.681	26	**
<i>Pgm-2</i>	2	18.782	13	NS
Total		1035.202	273	**

d.f. = degrees of freedom; * $P < 0.01$; ** $P < 0.001$; NS = nonsignificant.

per locus in *P. tiliguerta* was 1.2, ranging from 1.0 (Meli Island) to 1.5 (Tempio Pausania and Nuoro, Sardinia). The proportion of polymorphic loci (P) showed a similar trend, ranging from 0.05 (Meli Island) to 0.35 (Tempio Pausania and Nuoro; Sardinia) and averaging 0.22. The overall mean observed heterozygosity (H_o) was 0.066 and ranged from 0.007 (Meli Island) to 0.106 (Vizzavona, Corsica). The mean values of percent polymorphism (P) and heterozygosity (H_o) detected in the populations from the small Corsican islands ($P = 0.20$; $H_o = 0.072$) were similar to those found in the samples from Sardinia ($P = 0.24$; $H_o = 0.058$). On the other hand, the samples from Corsica showed relatively higher values of average polymorphism and heterozygosity ($P = 0.27$; $H_o = 0.090$). The sample from Meli Island represents an exception, as it is characterized by a severe reduction in genetic variability ($P = 0.05$; $H_o = 0.007$).

The levels of polymorphism and heterozygosity found in the samples of *P. muralis* ($P = 0.35$; $H_o = 0.059$) were similar to the ones observed in *P. tiliguerta*, whereas little genetic variability was found in the samples of *P. sicula* ($P = 0.05$; $H_o = 0.015$). This latter result is in agreement with those reported by Capula (1990, 1994a,b,c), who also found very low levels of percentage polymorphism and heterozygosity in several *P. sicula* populations from peninsular Italy and Sicily.

Table 3 Genetic variability parameters in *Podarcis tiliguerta* populations

Genetic variability parameters	Corsica			Cerbicale Islands						Sardinia				
	A	B	C	D	E	F	G	H	I	J	K	L	M	N
Mssl	7.8	9.9	9.8	7.7	12.9	12.0	6.8	6.3	16.5	12.6	6.8	5.0	7.0	4.0
<i>A</i>	1.2	1.4	1.3	1.3	1.3	1.1	1.1	1.2	1.5	1.4	1.5	1.1	1.0	1.1
<i>P</i>	20	30	30	25	25	15	15	20	35	25	35	15	5	10
H_o	0.063	0.106	0.101	0.081	0.072	0.042	0.073	0.093	0.082	0.041	0.100	0.030	0.007	0.038
(SE)	0.031	0.044	0.038	0.040	0.033	0.027	0.046	0.044	0.037	0.020	0.042	0.016	0.007	0.027
H_e	0.068	0.118	0.117	0.082	0.080	0.060	0.060	0.098	0.091	0.079	0.110	0.043	0.018	0.041
(SE)	0.034	0.047	0.042	0.036	0.035	0.034	0.037	0.045	0.041	0.037	0.043	0.026	0.018	0.031

Mssl, mean sample size per locus; *A*, mean number of alleles per locus; *P*, mean proportion of polymorphic loci; H_o , observed mean heterozygosity; H_e , expected mean heterozygosity (unbiased estimate; Nei, 1978) (SE, standard error). For geographical origin of populations see Fig. 1.

Population heterogeneity

Table 4 provides estimates of F_{IS} , F_{IT} and F_{ST} among *P. tiliguerta* populations. Eight of the 13 single locus F_{ST} values were statistically significant, suggesting high genetic differentiation among populations. F_{ST} values were high for the loci *Ldh-1* and *Me-1*, and noticeably high for the loci *Idh-1* and *Gapd*, which exhibit alleles partially diagnostic between the populations from Corsica and the Corsican satellite islands, and *Gpi*, which displays significant differences in allele frequencies between the populations from Sardinia and those from Corsica and the Corsican islands. The mean F_{ST} value is 0.460, indicating that 46 per cent of genetic variation in *P. tiliguerta* is attributable to differentiation among populations. As expected, the estimates of F_{IT} tend to mirror those of F_{ST} , although there are discrepancies at certain loci. The mean F_{IT} value was 0.488, indicating a deficiency of heterozygotes within the species. F_{IS} values are low (see Table 4), presumably because Hardy–Weinberg proportions are maintained within populations by random mating. Wright's (1978) hierarchical analysis combined across loci in all populations shows that 22 per cent of genetic differentiation (0.217) is attributable to divergence among the three population groups (Corsica, Corsican satellite islands, Sardinia). Genetic differentiation among populations within each group is slightly higher (0.257).

Genetic distance

Within *P. tiliguerta*, high values of genetic distance were found, *D* ranging from 0.003 (between Maestro Maria and Toro Grande samples) to 0.199 (between Sisco and Cagliari samples), and averaging 0.071

Table 4 Summary of *F*-statistics at 13 loci for *Podarcis tiliguerta* populations

Locus	F_{IS}	F_{IT}	F_{ST}
<i>Ldh-1</i>	0.117	0.430	0.355*
<i>Mdh-1</i>	-0.027	-0.002	0.024 NS
<i>Me-1</i>	0.087	0.388	0.330*
<i>Me-2</i>	0.556	0.661	0.236*
<i>Idh-1</i>	-0.080	0.731	0.751*
<i>Idh-2</i>	-0.333	-0.018	0.236*
<i>6Pgd</i>	0.141	0.378	0.276*
<i>Gapd</i>	-0.023	0.502	0.514*
<i>Got-1</i>	0.077	0.191	0.123 NS
<i>Got-2</i>	0.355	0.403	0.075 NS
<i>Mpi</i>	0.077	0.203	0.136 NS
<i>Gpi</i>	-0.104	0.663	0.694*
<i>Pgm-2</i>	-0.077	-0.005	0.067 NS
Mean	0.052	0.488	0.460

Asterisk denotes statistical significance for F_{ST} as determined by the χ^2 -test (see text).

* $P < 0.0001$; NS = nonsignificant.

across all populations. This is because (i) the samples from the Corsican satellite islands were genetically differentiated from those inhabiting the mother island (average $D = 0.083$), and (ii) high intraspecific genetic distances were found comparing Sardinian (including Meli Island) populations with those from Corsica (average $D = 0.113$) and from the small Corsican islands (average $D = 0.077$). On the other hand, low values of standard genetic distance were found (i) among the five satellite Corsican islands (average $D = 0.023$), (ii) within Corsica (average $D = 0.046$), and (iii) within Sardinia (average $D = 0.039$). However, these latter

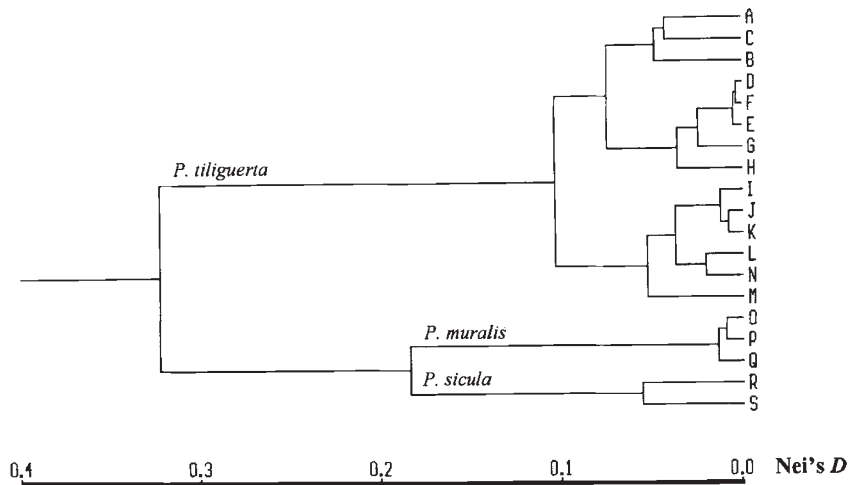


Fig. 3 Phenogram generated by UPGMA cluster analysis based on Nei's (1972) standard genetic distances among populations of *Podarcis tiliguerta*, *P. muralis*, and *P. sicula* (cophenetic correlation = 0.945). For geographical origin of populations see Fig. 1.

values were larger than those usually detected within the genus *Podarcis* among local populations of the same species (see for example Mayer, 1981; Capula, 1990, 1994a,b), indicating at least a certain amount of genetic subdivision for the populations inhabiting the major islands.

The comparison between genetic distances and geographical distances (Mantel test) reveals a positive correlation, significantly different from zero, for the total sample of *P. tiliguerta* (matrix correlation (normalized Mantel statistic): $r = 0.629$, $P < 0.004$).

As to the interspecific genetic distances, they ranged from $D = 0.181$ to $D = 0.318$, falling into the range obtained from comparisons between well recognized biological species of the genus *Podarcis* (Mayer & Tiedemann, 1982; Capula, 1990, 1994a,c). *Podarcis muralis*, considered to be phylogenetically related to *P. tiliguerta*, appears to be remarkably differentiated from the latter ($D = 0.300$), and is more closely related to *P. sicula* ($D = 0.181$). On the other hand, the comparison between *P. tiliguerta* and *P. sicula* gave an average Nei's genetic distance ($D = 0.318$) very close to that found between the former species and *P. muralis*.

Gene flow

The rate of gene flow (Nm) was calculated at various levels, from within each of the three major groups indicated by genetic distance data and cluster analysis to the total sample of *P. tiliguerta*. Estimates of Nm based on Wright's method are low; Nm among Sardinian samples is 0.500, whereas when comparing Sardinian and Corsican samples Nm reduces to 0.284. Rates of gene flow are slightly

higher among the samples from Corsica ($Nm = 0.845$), but Nm reduces to 0.435 when comparing the populations from Corsica with those from the Corsican satellite islands. The Nm value for the total sample of *P. tiliguerta* is 0.293.

Cluster analysis

The genetic relationships among the populations studied are depicted in Fig. 3. The UPGMA clustering procedure revealed three main clusters, corresponding to the three species analysed, in the phenogram constructed on the basis of the matrix of Nei's standard genetic distances. Within the cluster of *P. tiliguerta*, the existence of three subclusters should be noted. The first subcluster includes the populations from Corsica (A–C), the second contains the four samples from the Cerbicale Islands (D–G) and the sample from the Island of Lavezzi (H), and the third includes the six populations from Sardinia (I–L, N) and Meli Island (M). Cophenetic correlation between the matrix of genetic distances and the derived phenogram was rather high (0.945).

Discussion

The results of our analyses on allozyme variation in *P. tiliguerta* indicate that genetic polymorphism is relatively high in this species. The level of heterozygosity ($H_o = 0.066$) is larger than (i) the average estimated by Capula (1990) for nine species of *Podarcis* ($H_o = 0.053$) and (ii) the average calculated by Nevo (1978) for 17 species of reptiles ($H_o = 0.047$). The highest values of heterozygosity are found in the samples from Corsica, whereas the lowest ones are

observed in some samples from Sardinia. The populations from the Corsican satellite islands are characterized by intermediate values of H_o . This indicates that in *P. tiliguerta* H_o basically decreases southwards in the direction of both decreasing rainfall and increasing air temperatures. Although the lowest level of genetic variability was observed in a very small island (Meli), generally there was no significant correlation between island size and heterozygosity (Spearman's rank-correlation). Our data allow us to compare the genetic variability of 'parent' island populations with that of small island populations. In all, six small islands associated with two parent islands (Corsica, Sardinia) were sampled. None of the populations inhabiting the five small islands off the south-eastern coast of Corsica showed a markedly reduced genetic variability compared with that of the parent island, i.e. the so-called 'small island effect' (Gorman *et al.*, 1975). The levels of polymorphism and heterozygosity detected either on the larger (Lavezzi, 0.729 km²; Pietricaggiosa, 0.046 km²) or on the tiny satellite islands (Maestro Maria, 0.028 km²; Toro Grande, 0.016 km²; Toro Piccolo, 0.005 km²), although lower than those found on Corsica ($P = 0.20$ vs. $P = 0.27$; $H_o = 0.072$ vs. $H_o = 0.090$), were more than we might expect from the size of these small islands (see for example Gorman *et al.*, 1975; Gorman & Kim, 1976; Guillaume & Lanza, 1982; Capula, 1994a,c). Moreover, these populations were genetically differentiated from those inhabiting Corsica (average Nei's $D = 0.083$). As the Cerbicale Islands are geographically very close to Corsica (≈ 3 km from Punta Cerbicale, SE Corsica), the possibility of several arrivals of lizards from the parent island via passive transport cannot be excluded. Such arrivals could explain the preservation of the relatively high genetic variability observed in the samples from these small islands. A different situation is found in the sample from Meli Island (SW Sardinia), which is characterized by a severe reduction in genetic variability compared with that observed in the populations from the parent island. This could be caused by genetic drift phenomena, as supported by the fact that this population (i) inhabits a tiny fringing island (< 0.01 km²), (ii) has no unique alleles and is characterized by a predominance of fixed alleles at each locus, (iii) is genetically very similar (Nei's $D = 0.029$) to the geographically closest population (≈ 1 km) from Capo Giordano (SW Sardinia) (Fig. 1).

The genetic heterogeneity analysis demonstrates substantial genetic differentiation among local populations of *P. tiliguerta*, with the level of genetic subdi-

vision exceeding values known for other lizard species (McKinney *et al.*, 1972; Sites & Greenbaum, 1983; Sarre *et al.*, 1990). The estimated standardized variance in gene frequency (F_{ST}) for the total sample is highly significant, with a value (0.460) much higher than that calculated by Capula (1994a) for the insular lacertid lizard *P. wagleriana* ($F_{ST} = 0.153$) and very high for vertebrates in general (see for example Gorman *et al.*, 1975; Wright, 1978; Schonewald-Cox *et al.*, 1983; Raghianti & Wake, 1986). This indicates that 46 per cent of the gene diversity was between populations, and 54 per cent was within populations.

F -statistics and genetic distance data show that, at the scale of the study, genetic variation in *P. tiliguerta* is distributed into three population groups. These groups are geographically coherent, as indicated by UPGMA cluster analysis (Fig. 3). The first group includes the populations from the northern part of the range (Corsica), the second includes the populations from the small islands off the south-eastern coast of Corsica (Cerbicale and Lavezzi), and the third comprises the populations from the southern part of the range (Sardinia and Meli Island). Genetic distances found among the three population groups are much higher than those usually detected within the genus *Podarcis* among local populations of the same species (see Mayer, 1981; Capula, 1990, 1994a,b,c), the average Nei's D values ranging from 0.077 to 0.113. On the other hand, these values fall below those normally encountered comparing populations of well recognized biological species (Thorpe, 1983), as showed by the comparison with *P. muralis* and *P. sicula* samples (Fig. 3).

Several conditions may be responsible for population structuring and differentiation in the geographically fragmented *P. tiliguerta*, such as the occurrence of genetic drift phenomena, the influence of different selective regimes, and the mode and level of gene flow. Although the distribution of genetic diversity among *P. tiliguerta* populations appears to be random at most loci within the area studied, and genetic drift might account for the possible loss of alleles in the populations inhabiting the smallest fringing island (Meli Island), this in no way implies that alleles at all loci are behaving neutrally and that differentiation can occur only by genetic drift. The hypothesis that different selective regimes actively favour different alleles, at least at a few loci, cannot be excluded. The latter possibility is supported by the observation that genetic differentiation among populations results mainly from large differences in allele frequencies at a few loci (*Idh-1*, *Gapd*, *Gpi*). Marked differentiation at these loci may be the

result of range fragmentation produced by late Pleistocene geological events, when Sardinia, Corsica, and the other small satellite islands (linked to each other until the last phase of the Würmian glacial period (Caloi *et al.*, 1988)) were physically separated by sea channels resulting from marine ingression (Lanza, 1972, 1983). However, because the alleles at the *Idh-1*, *Gapd* and *Gpi* loci show significant departure from random distribution and appear to vary clinally across the populations studied, it is possible that most of the polymorphisms at these loci are being selected.

Reduced gene flow is another important factor promoting genetic differentiation between geographically isolated populations (Slatkin, 1987). Therefore we expected to find a relatively high correlation between geographical and genetic distances and low or very low levels of gene flow among the three population groups in this study. The results of our analyses are congruent with these expectations, as we found a positive and significant correlation between genetic and geographical distances. This correlation is an expression of isolation by distance at the scale of the study. Moreover, the rates of gene flow among the three groups of populations are noticeably low (Nm ranging from 0.284 to 0.435), indicating highly significant independence of local demes. It must be stressed that when the estimated average number of migrants per generation (Nm) is <0.5 , local demes are supposed to be largely unconnected under any model of gene flow (Nagylaki, 1983). This situation corresponds well to the case of the island model of population differentiation (Wright, 1940, 1943), in which strong isolation and large genetic differences among populations are involved.

In summary, the genetic structure of *P. tiliguerta* appears to be more subdivided than that of other lizards studied to date. Although gene diversity analysis cannot provide much information about the relative significance of selectionist or neutralist causes of this subdivision, the pattern of genetic variability and the apparent clinal variation of alleles at some loci suggest that the genetic structure of populations is moulded by the interplay of stochastic processes (geographical isolation, genetic drift) and agents selectively affecting allele frequency changes (local bioclimatic regimes).

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Appendix Allele frequencies at 15 variable loci in *Podarcis tiliguerta* (A–N), *P. muralis* (O–Q), and *P. sicula* (R, S) populations. For geographical origin of populations see Fig. 1

Locus	Allele	Population																			
		A	B	C	D	E	F	G	H	I	J	K	L	M	N	O	P	Q	R	S	
<i>Ldh-1</i>	80	1.000	0.964	1.000	0.312	0.500	0.500	1.000	0.286	0.605	0.536	0.643	1.000	1.000	0.875	0.000	0.000	0.000	0.000	0.000	
	100	0.000	0.036	0.000	0.688	0.500	0.500	0.000	0.714	0.395	0.464	0.357	0.000	0.000	0.000	0.125	1.000	1.000	1.000	1.000	
<i>Ldh-2</i>	90	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.000	0.000	0.000	
	100	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.944	1.000	1.000	1.000	
<i>Mdh-1</i>	100	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.974	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	
	114	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.026	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
<i>Me-1</i>	94	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.200	
	100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.053	0.214	0.000	0.000	0.000	0.000	0.042	0.000	0.000	1.000	0.800	
	103	0.000	0.100	0.000	0.000	0.000	0.000	0.000	0.000	0.026	0.000	0.143	0.000	0.000	0.000	0.000	0.188	0.000	0.000	0.000	
	106	0.312	0.800	0.214	0.188	0.233	0.167	0.429	0.429	0.605	0.715	0.500	0.700	0.000	1.000	0.417	0.562	0.958	0.000	0.000	0.000
	110	0.688	0.100	0.786	0.812	0.767	0.833	0.571	0.571	0.316	0.071	0.357	0.300	1.000	0.000	0.500	0.250	0.000	0.000	0.000	0.000
	112	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.041	0.000	0.042	0.000	0.000	0.000
<i>Me-2</i>	95	0.000	0.000	0.250	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
	100	1.000	1.000	0.750	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.944	1.000	1.000	1.000	
	105	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.000	0.000	0.000	
<i>Idh-1</i>	100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	1.000	1.000	
	102	1.000	0.700	0.333	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.958	0.944	1.000	0.000	0.000	
	108	0.000	0.300	0.667	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.000	0.000	0.000	0.000	0.000	
	110	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.000	0.000	0.000	
	120	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.042	0.000	0.000	0.000	0.000	
<i>Idh-2</i>	95	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.250	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
	100	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.750	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	
<i>6Pgd</i>	93	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.200	0.000	
	100	0.688	0.769	1.000	1.000	1.000	1.000	1.000	1.000	0.921	0.962	0.929	1.000	1.000	0.500	1.000	1.000	1.000	0.800	1.000	
	106	0.312	0.231	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.038	0.071	0.000	0.000	0.500	0.000	0.000	0.000	0.000	0.000	
	112	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.079	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	

Population

Locus	Allele	A	B	C	D	E	F	G	H	I	J	K	L	M	N	O	P	Q	R	S	
<i>Gapd</i>	94	0.000	0.500	0.000	0.875	0.800	1.000	0.625	0.250	0.633	0.250	0.214	1.000	1.000	1.000	0.000	0.000	0.000	0.000	0.000	
	100	1.000	0.400	0.833	0.125	0.200	0.000	0.375	0.750	0.367	0.750	0.786	0.000	0.000	0.000	1.000	1.000	1.000	1.000	1.000	
	108	0.000	0.100	0.167	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
<i>Got-1</i>	90	0.188	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.026	0.036	0.072	0.000	0.214	0.000	1.000	0.944	1.000	0.000	0.000	
	97	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.000	0.000	0.000	
	100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	1.000	
	104	0.812	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.974	0.928	0.857	1.000	0.786	1.000	0.000	0.000	0.000	0.000	0.000	0.000
	108	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.036	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
124	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.071	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
<i>Got-2</i>	85	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.029	0.000	
	90	0.000	0.000	0.000	0.063	0.100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
	100	1.000	1.000	1.000	0.937	0.900	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.971	1.000	
<i>Mpi</i>	94	0.000	0.000	0.208	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
	98	1.000	1.000	0.792	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.929	0.900	1.000	1.000	0.000	0.000	0.000	0.000	0.000	
	100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	1.000	1.000	1.000	1.000	1.000	
	104	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.071	0.100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
<i>Gpi</i>	92	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.250	0.000	0.000	0.000	
	95	0.063	0.250	0.250	0.250	0.091	0.250	0.071	0.000	0.974	1.000	1.000	0.900	1.000	1.000	0.000	0.000	0.000	0.000	0.000	
	100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	1.000	0.750	1.000	1.000	1.000	
	103	0.937	0.750	0.750	0.750	0.909	0.750	0.929	1.000	0.000	0.000	0.000	0.100	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
	106	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.026	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	
<i>Pgm-1</i>	93	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.059	0.000	0.000	
	100	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.944	0.941	1.000	1.000	
<i>Pgm-2</i>	95	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.029	0.000	
	100	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	0.929	1.000	1.000	1.000	1.000	0.778	0.942	1.000	1.000	
	108	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.071	0.000	0.000	0.000	0.000	0.056	0.000	0.000	0.000	
	115	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.166	0.029	0.000	0.000	