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Allele frequencies of 6 STR loci (TH01, TPOX, CSF1PO, D13S317, D16S539 and D7S820) in São Tomé e Príncipe (West Africa)

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Resumo As frequências alélicas de seis marcadores autossómicos do tipo *short tandem repeat* (STR) (TH01, TPOX, CSF1PO, D13S317, D7S820 e D16S539) foram estimadas numa amostra populacional (N=53-85) de São Tomé e Príncipe (África Ocidental). Não foram encontrados desvios significativos ao equilíbrio de Hardy-Weinberg em qualquer dos *loci*. A distribuição de frequências e valores de heterozigotia são semelhantes aos encontrados noutras populações Africanas sub-Saarianas.

Palavras-chave São Tomé e Príncipe; short tandem repeats (STRs); genética populacional.

Abstract Allele frequencies of six autosomal short tandem repeat (STR) markers (TH01, TPOX, CSF1PO, D13S317, D7S820 and D16S539) were estimated in a population sample (N=53-85) of unrelated individuals from São Tomé e Príncipe (West Africa). No deviations from Hardy-Weinberg equilibrium were observed for all loci. Allele frequency distribution and heterozygosity values were similar with those from other sub-Saharan African populations.

Key words: São Tomé e Príncipe; short tandem repeats (STRs); population genetics.

Introduction

Microsatellites or short tandem repeats (STRs) are abundant and widespread in the human genome and can be easily analyzed by automated typing procedures. They are highly discriminating and useful genetic markers in several human fields including linkage analysis (Cashman *et al.*, 1995) and identity testing (Butler, 2004). Microsatellites are also valuable tools in anthropological research, scoring genetic variability in human populations (Rosenberg *et al.*, 2002).

São Tomé e Príncipe, a former Portuguese colony, is a small archipelago located in the Gulf of Guinea (West Africa) inhabited by populations descendent from African slaves brought from the continental mainland. Presently, over 180000 inhabitants live in the two major islands of the archipelago, distributed by three main population groups: Angolares, Forros and Tongas (Trovoadá *et al.*, 2007). Several autosomal microsatellite data have been already reported for the Saotomean population (Gusmão *et al.*, 2001; Pereira *et al.*, 1999; 2000).

In this study, the distribution pattern of six autosomal loci (TH01, TPOX, CSF1PO, D13S317, D7S820 and D16S539) currently used in commercial kits for identification purposes was analyzed in a population sample from São Tomé e Príncipe providing additionally information on population genetic diversity. The genetic relationship between the sampled population and other African populations from the mainland continent was also examined.

Material and methods

Unrelated individuals (N=53-85) from the São Tomé e Príncipe archipelago were typed for TH01 (N=85), TPOX (N=80), CSF1PO (N=70), D13S317 (N=57), D7S820 (N=53) and D16S539 (N=57) polymorphic STRs. After informed consent, genomic DNA was extracted from blood samples using the chelex method.

Two PCR multiplex were performed, one for TH01, TPOX, CSF1PO *loci* and the second for D13S317, D7S820 and D16S539 *loci*, using primers labeled with Cy5. Primer sequences were available in <http://www.cstl.nist.gov/biotech/strbase>. PCR amplification was carried out using the Multiplex Qiagen Kit (Qiagen) under the following conditions: initial activation step at 95°C for 15 min, followed by 30 cycles of 95°C – 45 s; 60°C – 45 s and 72°C – 1 min. The PCR products were analyzed in ReproGel High Resolution (GE Healthcare) using the automatic sequencer ALFexpress II (Amersham Pharmacia Biotech). Allele sizes were determined automatically with the software ALFwin Fragment Analyzer 1.00 by comparison with ladders and using control samples provided in the GenePrint STR Multiplex System (for TH01, TPOX, CSF1PO *loci*) and GenePrint SilverSTR III System (for D13S317, D7S820 and D16S539 *loci*) (Promega Corporation).

Allele frequencies, heterozygosity values and exact P values for accordance with Hardy-Weinberg equilibrium (Guo and Thompson, 1992) and for statistical significance on genetic differentiation between populations (Raymond and Rousset, 1995) were calculated using the software package Arlequin, vs. 3.01 (Excoffier *et al.* 2005; <http://cmpg.unibe.ch/software/arlequin3/>). Allele frequencies from Cabinda (Beleza *et al.*, 2004), Mozambique (Alves *et al.*, 2004), Equatorial Guinea (Alves *et al.*, 2005) and Guinea-Bissau (Gonçalves *et al.*, 2002) were used for genetic differentiation tests with São Tomé e Príncipe population sample, applying the Bonferroni correction for multiple tests (Hochberg, 1988). A neighbor-joining tree was constructed using the PHYLIP3.5c software package (Felsenstein, 1993) based on Reynolds Genetic distance (Reynolds *et al.*, 1983) with a bootstrap of 10000 resamples, using population data from different Portuguese regions, Central Portugal (Manco *et al.* 2007-2008), North Portugal (Pinheiro *et al.*, 2005), Madeira (Fernandes *et al.*, 2002), Azores (Velosa *et al.*, 2002), and from other African populations, Cabinda (Beleza *et al.*, 2004), Mozambique (Alves *et al.*, 2004), Equatorial Guinea (Alves *et al.*, 2005) and Guinea-Bissau (Gonçalves *et al.*, 2002).

Results and discussion

Allele frequencies for the 6 STR loci in the population sample of São Tomé e Príncipe are displayed in Table 1. No deviations from Hardy-Weinberg expectations were observed for all *loci*. Population allele distribution and heterozygosity values were similar to other geographic neighboring populations confirming the typical sub-Saharan African genetic profile of the Saotomean population.

When comparing allele frequency data from São Tomé e Príncipe to other 4 African continental mainland populations from Cabinda (Beleza *et al.*, 2004), Mozambique (Alves *et al.*, 2004), Equatorial Guinea (Alves *et al.*, 2005) and Republic of Guinea-Bissau (Gonçalves *et al.*, 2002) several significant differences were found (exact $P < 0.05$) but applying the Bonferroni correction (SISA) the significance level lowered to 0.0127 and significant values were only found with Mozambique at *loci* TH01 (exact $P = 0.007 \pm 0.001$) and TPOX (exact $P = 0.000 \pm 0.000$), and with Guinea-Bissau at *loci* TH01 (exact $P = 0.000$), which seems to reflect the colonization history of São Tomé e Príncipe that received most part of slaves from the West African coast neighboring the archipelago.

Table 1. Allele frequencies and other statistical parameters for the 6 STRs in a population sample from São Tomé e Príncipe.

Allele	TH01 (N=85)	TPOX (N=80)	CSF1PO (N=70)	D7S820 (N=57)	D13S317 (N=53)	D16S539 (N=57)
5						0.009
6	0.053	0.056				
7	0.359	0.050	0.071			
8	0.359	0.300	0.050	0.254	0.009	0.018
9	0.141	0.300	0.029	0.123		0.263
9.3	0.070					
10	0.018	0.075	0.336	0.342	0.028	0.088
11		0.181	0.164	0.193	0.377	0.307
12		0.038	0.279	0.079	0.387	0.211
13			0.071		0.142	0.079
14				0.009	0.047	0.026
15					0.009	
<i>p</i>	0.626	0.180	0.912	0.636	0.855	0.149
Ho	0.741	0.763	0.843	0.737	0.698	0.825
He	0.719	0.781	0.775	0.766	0.698	0.784

N: number of samples; *p*: Hardy-Weinberg equilibrium, exact test; Ho: observed heterozygosity; He: expected heterozygosity;

A neighbor-joining tree was constructed based upon the 6 STR allele frequencies in our sampled population and from populations from Central Portugal (Manco *et al.*, 2007/2008), North Portugal (Pinheiro *et al.*, 2005), Madeira (Fernandes *et al.*, 2002), Azores (Velosa *et al.*, 2002), Cabinda (Beleza *et al.*, 2004), Mozambique (Alves *et al.*, 2004), Equatorial Guinea (Alves *et al.*, 2005) and Guinea-Bissau (Gonçalves *et al.*, 2002) (Figure 1).

Figure 1. Neighbor-joining tree based on allele frequencies of the 6 STR loci (TH01, TPOX, CSF1PO, D13S317, D7S820 and D16S539) in the studied population sample from São Tomé e Príncipe (STP), and in different Portuguese regions (C Port – Central Portugal, N Port – North Portugal, Madeira, Azores) and other African populations (Cabinda, Mozambiq – Mozambique, Guin Eq – Equatorial Guinea and Guin Bis – Guinea-Bissau) with previous reported data.

The topology of the tree unequivocally differentiates African and European populations. Among the African cluster, São Tomé e Príncipe is closer to Cabinda and Equatorial Guinea consistent with the low significant differences at STR loci. The population from Central Portugal shows close genetic proximity with the neighbor mainland North Portuguese population than to Madeira and Azores archipelagos in accordance to the regional distances.

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