

SHORT COMMUNICATION

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An investigation of the TH01 locus in a population from northern Thailand

Received: 9 December 1996 / Received in revised form: 7 April 1997

Abstract The STR locus HUMTH01 was studied in 110 unrelated Thais from the area of Chiang Mai in North Thailand. By using PCR and vertical PAGE, six alleles were identified and the frequencies ranged from 0.005 to 0.400. The allele frequency distribution in this population showed significant differences from a Japanese population and other ethnic populations but was similar to Asians in the USA and Australia. The genotype distribution meets Hardy-Weinberg expectations. The average power of exclusion (in no-parent and one-parent cases) and the discriminating power (DP) were calculated to be 0.3020, 0.4761 and 0.8722 respectively.

Key words HUMTH01 · STR · PCR · Northern Thais · Population data

Introduction

Since the analysis of the polymorphism at the HUMTH01 locus appeared beneficial in human identification and paternity testing (Wiegand et al. 1993; Hochmeister et al. 1994; Kubat et al. 1995; Martin et al. 1995; Pfitzinger et al. 1995; Sjerps et al. 1995), allele and genotype data on this locus were explored in a northern Thai population.

Materials and methods

DNA was extracted from blood of 110 unrelated individuals living around Chiang Mai in North Thailand. The first 80 blood samples were extracted according to the method of Miller et al. (1988), the remaining other 30 samples according to Walsh et al. (1991). The

primers used in PCR were those described by Gill et al. (1992), the cycling conditions as suggested by Puers et al. (1993). PAGE of PCR products was done according to Sajantila et al. (1994) and the gels were silver-stained (Budowle et al. 1991). Alleles were designated by comparison with an allelic ladder (consisting of alleles 6, 7, 8, 9, and 9.3/10) made by the "crush and soak" method (Sambrook et al. 1989). Alleles 9.3 and 10 were pooled and reassigned as allele 10. The discriminating power and power of exclusion were calculated according to Fisher (1951) and Garber and Morris (1983) respectively. The Hardy-Weinberg equilibrium (HWE) analysis was based on a comparison of observed and expected genotypes according to Dickinson-Gibbons (1976) and determination of the unbiased estimate of the expected heterozygosity (Nei and Roychoudhury 1974). The frequency profile comparison between different ethnic populations was carried out by using RxC contingency tables.

Results and discussion

In this population sample from Thailand, six alleles were found with frequencies ranging from 0.005 to 0.40 (Table 1). Allele number 9 was the most frequent and 15 genotypes were observed in northern Thais. Genotype 7/9 was found to be the most common ($f = 0.327$). The population data for locus HUMTH01 showed no significant deviation from HWE ($0.10 > p > 0.05$, d.f. = 2) and were similar to Asians in USA (Puers et al. 1993) ($0.9 > p > 0.75$) and

Table 1 Allele frequencies at the TH01 locus in northern Thais, Japanese from central Japan, US-Caucasians, and US-Blacks

Allele	Northern Thais	Japanese ^a	US-Caucasians ^b	US-Blacks ^b
5	–	–	0.005	–
6	0.118	0.242	0.226	0.135
7	0.305	0.286	0.159	0.370
8	0.045	0.049	0.110	0.211
9	0.400	0.382	0.143	0.146
10 (9.3 + 10)	0.127	0.041	0.357	0.137
11	0.005	–	–	0.007

^a Obtained from Nagai et al. (1996)

^b Obtained from Puers et al. (1993)

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Australia (van Oorschot et al. 1994) ($0.5 > p > 0.25$), but different from a central Japanese population ($p < 0.001$) (Nagai et al. 1996) and other major ethnic populations (US-Caucasians, $p < 0.001$; US-Blacks, $p < 0.001$). The discriminating power and the average power of exclusion were 0.8722, 0.3020 (no-parent case) and 0.4761 (one-parent case) respectively. This study shows that the TH01 system may have slightly less power for forensic purposes when used in the Thai population than in Caucasian populations (Puers et al. 1993; Hammond et al. 1994; Nellemann et al. 1994; Hochmeister et al. 1994; Pestoni et al. 1995; Sjerps et al. 1995). We suggest that local population data should be collected because there may be significant differences in allele distribution not only between major ethnic groups but even within Asian populations.

Acknowledgements This study was supported by donations to the Forensic Genetics Unit, Department of Forensic Medicine, Faculty of Medicine, Chiang Mai University. We thank Professor Dr. Torpong Sanguanserm Sri who allowed us to use laboratory equipment donated by the Volkswagen-Foundation

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