

Announcement of Population Data  
Genetic polymorphisms of 15 STR loci in two Tibetan  
populations from Tibet Changdu and Naqu, China

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Received 23 January 2006; received in revised form 10 March 2006; accepted 10 March 2006

Available online 18 April 2006

### Abstract

The allelic distribution of 15 short tandem repeat (STR) loci included in the AmpFI STR<sup>®</sup> Identifier<sup>™</sup> kit was examined in 100 Changdu Tibetan and 118 Naqu Tibetan unrelated individuals living in the Tibet Province, PR China. The distribution of these observed genotypes was not significantly different from the expected distribution according to Hardy–Weinberg equilibrium.

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**Keywords:** Forensic science; Short tandem repeats (STR); Genetic polymorphism; Population data; Tibetan; Chinese

**General information on the studied population:** Changdu area and Naqu area are Tibetan inhabitancy under the jurisdiction of Tibet, abutting on Sichuan Province and Qinghai Province of China, respectively. People residing in these areas speak in unique dialect and have distinct lifestyle. The geographical distribution of Tibetan in Tibet Province is shown in Fig. 1 [1]. We detected 15 highly polymorphic short tandem repeats (STR) of the two populations and compared the difference between them and other populations living in China.

**Population:** Blood samples were collected from randomly selected individuals in the regions.

**DNA extraction:** Genomic DNA was extracted using the Chelex-100 protocol as described by Walsh et al. [2] and quantified spectrophotometrically.

**PCR:** Target DNA was amplified following the manufacturer's instruction (AmpFI STR<sup>®</sup> Identifier<sup>™</sup> PCR Amplification, Applied Biosystems) with some modifications: total reaction volume was reduced from 25  $\mu$ l recommended to 8  $\mu$ l, in which was included 0.5–1.0 ng DNA, 3  $\mu$ l reaction mix, 1.8  $\mu$ l prime set and 0.25  $\mu$ l AmpliTaq Gold DNA polymerase (5 U/ $\mu$ l).

**Typing:** ABI Prism<sup>®</sup> 3100 Genetic Analyzer using the recommended protocol (AmpFI STR<sup>®</sup> Identifier<sup>™</sup> PCR Amplification, Applied Biosystems) electrophoresis results were analyzed by data collection software (Version 1.1), Genescan analysis (Version 3.7) and Genotyper (Version 3.6) software (Applied Biosystems).

**Results:** See Tables 1 and 2.

**Quality control:** Laboratory internal control standards and kit controls.

**Data analysis:** Allele frequencies (since autosomal co-dominant) were computed using the gene counting method. The potential usefulness of the studied markers for forensic studies in the Changdu Tibetan and Naqu Tibetan was assessed by calculating discrimination power (DP) [3], probability of paternity exclusion (PE) [4] and the heterozygosity value ( $H$ ) [5]. The STR DP, PE and  $H$  were calculated with the allelic frequency as shown in Tables 1 and 2. R  $\times$  C contingency test was employed for pairwise interpopulation comparisons [6,7]. Excel program and SPSS 11.5 for windows software were used for analysis of data.

**Access to the data:** The complete data are available to any interested researchers via mail from author ([klongli@yahoo.com](mailto:klongli@yahoo.com)).

**Other remarks:** The population samples were highly polymorphic. There was less genetic diversity among Changdu Tibetan and Naqu Tibetan populations than in the Chinese Han

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